

Nasimov A.M., Tashpulatov X.Sh.

Umumiy kimyo

2-Qism

- *Anorganik kimyo
- *Organik kimyo
- *Ma'lumotnomalar

KIRISH

“Kadrlar tayyorlash milliy dasturi” tizimida ta’limda uzlusizlikni ta’minlash bo’g’ini yoshlarni qiziqishi va iqtidorini e’tiborga olib, kasbga yo’naltirish hisoblanadi. Shundan kelib chiqib tabiiy va tibbiyot yo’nalishida Akademik litseylar o’quvchilari uchun ko’plab qo’llanma va darsliklar chop etilgan. Ularda asosan oliy oq’uv yurtlariga kirish uchun nazariy ma’lumotlar va amaliy mashg’ulotlar berilgan.

Qo’lingizdagi qo’llanma esa Akademik litseylarning tabiiy yo’nalishi 3 yillik o’quv rejasi asosida tuzilgan bo’lib, unda umumiyl, anorganik va organik kimyodan nazariy ma’lumotlar, masalalarning yechimlari keltirilgan. Uni tayyorlashda mavzuni yoritish uchun iloji boricha qisqa va to’liqroq ma’lumot berishga harakat qilindi.

Hozirda kirish imtihonlarida test topshiriqlarini yechish usullari bo’yicha juda ko’p qo’llanmalar mavjudligiga qaramasdan, ularning ko’pchiligidagi masalalar yechish turli xil proportsiyalar tuzishga asoslanadi. Bu esa o’quvchida bir tomonlama, ya’ni faqat masalani yechimini topishdan boshqa maqsad qoldirmasligiga olib kelishi mumkin. Shuning uchun biz qo’llanmada iloji boricha test topshiriqlarini yechishda kimyoviy formulalardan foydalanishga, kattaliklarni to’g’ri keltirishga harakat qildik. Qo’llanmada barcha kattaliklar Xalqaro birliklar sistemasi (SI) va belgilashlariga mos tushadi.

Ushbu qo’llanmani Kimyodan mavzulashtirilgan testlar to’plami bilan birgalikda foydalanish maqsadga muvofiq. Chunki unda mustaqil yechish uchun masalalar keltirilmagan.

Qo’llanmadan Akademik litsey o’quvchilari, kirish imtihonlariga tayyorlanuvchi abituriyentlar va repetitorlar foydalanishlari mumkin.

Qo”llanmani ko’rib chiqib o’z fikrlarini bildirganligi uchun mualliflar SamQXI qoshidagi 1 son Akademik litseyi kimyo fani o’qituvchisi G’ofurov A. ga, anorganik kimyo kafedrasi dotsenti, kimyo fanlari nomzodi, dotsent Lutfullayev E.L. larga chuqur minnatdorlik bildiradi.

Mualliflar ushbu qo’llanma bo’yicha barcha fikr va mulohazlarni quyidagi manzilda mammuniyat bilan qabul qiladi. Samarqand shahri, 140104, Universitet xiyoboni ko’chasi 15, Samarqand davlat universiteti, Kimyo bo’limi, Anorganik kimyo kafedrasi, 408 xona.

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2-Qism

**ANORGANIK
KIMYO
REAKSIYALARDA**

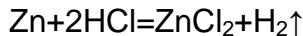
ANORGANIK KIMYO REAKSIYALARDA

Mavzu: Vodorod – H₂

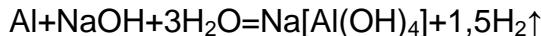
Olinishi:

Laboratoriya da:

1. Suyultirilgan kislotalarga metallar ta'siridan:



2. Amfoter metallarga ishqor ta'sir ettirib:



Sanoatda:

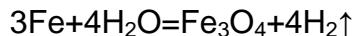
1. Metanni konversiyalab:



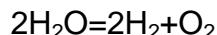
2. Suv bug'ini cho'g'langan koksga ta'sir ettirib:



3. Cho'g'langan temirga suv bug'i ta'siridan:

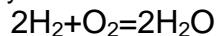


4. Suvni elektroliz qilib:



Kimyoviy xossalari:

1. Kislorodda yorqin havorang alanga berib yonadi:



2. Galogenlar, N₂ va S bilan oson ta'sirlashadi:



3. Ko'pchilik metal oksidlarini qaytaradi(vodorodotermiya):



4. Ishqoriy va ishqoriy yer metallari bilan gidridlar hosil qiladi:

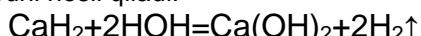


5. Molekulyar vodorod VIIIB guruh metallariga yaxshi yutiladi. Masalan, 1 hajm Pd 700 hajm vodorodni yutadi.

Mavzu: Gidridlar – EH_x

1. Tuzsimon gidridlar – ishqoriy va ishqoriy-yer metallarining gidridlari: LiH, NaH, KH, CsH, RbH, CaH₂, SrH₂ va BaH₂.

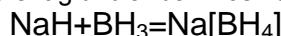
Ular suv bilan ta'sirlashib vodorodni hosil qiladi:



2. Kovalent gidridlar – IVA, VA, VIA va VIIA guruh elementlarining gidridlaridir. CH₄, SiH₄, NH₃, PH₃, H₂O, H₂S, HF, HCl, HBr, HJ, ASH₃, BeH₂ va BH₃.

3. Metal gidridlar – IB-VIIIB guruh elemntlarining gidridlaridir. Masalan, Pd-H₂.

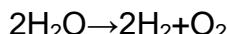
4. Kompleks gidridlar – tuzsimon va kovalent gidridlaridan hosil bo'ladi.



Mavzu: Suv – H₂O

Kimyoviy xossalari:

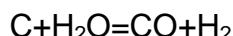
1. Suv 1000°C dan yuqorida parchalanadi:



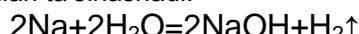
2. Galogenlar bilan ta'sirlashadi:



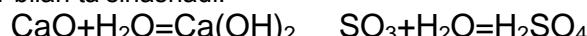
3. Cho'g'langan ko'mir bilan ta'sirlashadi:



4. Ishqoriy va ishqoriy-yer metallari bilan ta'sirlashadi:

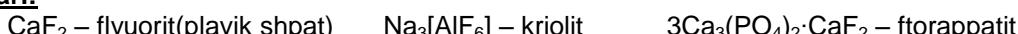


5. Asosli va kislotali oksidlar bilan ta'sirlashadi:



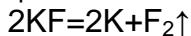
Mavzu: Ftor – F₂

Minerallari:



Olinishi:

1. KF ning HF dagi suyuqlanmasini elektroliz qilib olinadi:



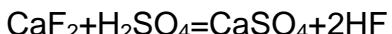
Kimyoviy xossalari:

1. Ftor juda aktiv metalmas. U faqatgina bevosita kislород va azot bilan ta'irlashmaydi.

Mavzu: Vodorod ftorid – HF

Olinishi:

Flyuoritga kons. H_2SO_4 ta'siridan olinadi:



Kimyoviy xossalari:

1. Plavik kislota shishani yemiradi:



2. AgF suvda yaxshi eriydi.

Mavzu: Xlor – Cl₂

Minerallari:

KCl – silvin

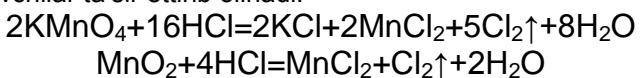
KCl·MgCl₂·6H₂O – karnallit

KCl·MgSO₄·3H₂O – kainit

Olinishi:

Laboratoriya da:

1. Xlorid kislotaga oksidlovchilar ta'sir ettirib olinadi:



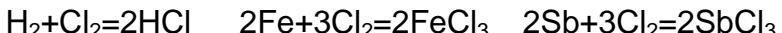
Sanoatda:

1. Osh tuzi eritmasini elektroliz qilib olinadi:

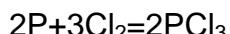


Kimyoviy xossalari:

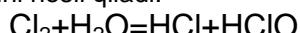
1. Xlor juda kuchli oksidlovchi. U vodorod va metallar bilan kuchli ta'sirlashadi:



2. U metalmaslar bilan ham ta'sirlashadi:



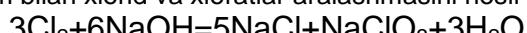
3. Xlor suvda erib kislotalar aralashmasini hosil qiladi:



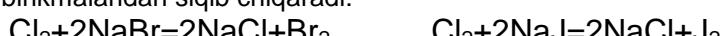
4. Xlor sovuq ishqor eritmalari bilan xlorid va gipoxloritlar aralashmasini hosil qiladi:



5. Xlor qaynoq ishqor eritmalari bilan xlorid va xloratlar aralashmasini hosil qiladi:



6. Xlor brom va yodni birikmalaridan siqib chiqaradi:



Mavzu: Vodorod xlorid – HCl

Olinishi:

1. Sintez usuli bilan olinadi:



2. Osh tuziga kons. H_2SO_4 ta'sir ettirib olinadi:



Kimyoviy xossalari:

1. Xlorid kislota kuchli kislota hisoblanadi:



2. 3 hajm kons.HCl va 1 hajm kons. HNO_3 aralashmasiga "zar suvi" yoki "podsho arog'i" deyiladi. Unda Au va Pt eriydi:



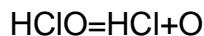
3. Kuchlanishlar qatorida H_2 dan o'ngda turgan metallar HCl da erimaydi.

Mavzu: Xorning kislородли birikmlari

Kislородли kislotalari: HClO , HClO_2 , HClO_3 , HClO_4

Kimyoviy xossalari:

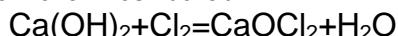
1. Gipoxlorit kislota juda kuchsiz kislota. Parchalanganda atomar kislород ajralib chiqqanligi uchun juda kuchli oksidlovchi:



2. Gipoxloritlarni olish uchun xlor sovuq ishqor eritmasida eritiladi:



3. Ohak eritmasiga xlor yuborilsa xlorli ohak hosil bo'ladi:



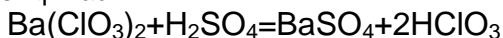
4. Xlorli ohak havoda va kislotalar ta'siridan parchalanadi:



5. ClO₂ ishqorlarda erib disproporsilanadi:



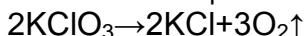
6. Xlorat kislota quyidagicha hosil qilinadi:



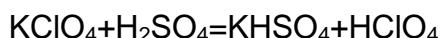
7. Bertolle tuzi katalizatorsiz 400°C da parchalanganda xlorid va perxloratlar hosil bo'ladi:



8. Bertolle tuzi MnO₂ ishtirokida xlorid va kislordni hosil qiladi:



9. Perxlorat kislota – HClO₄ eng kuchli kislordli kislota hisoblanib, u perxloratlarga kons. H₂SO₄ ta'siridan olinadi:

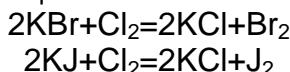


10. Perxloratlar oksidlovchilik xossasini namoyon qilmaydi.

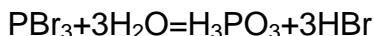
Mavzu: Brom va yod – Br₂/J₂

Olinishi:

1. Bromidlarni/yodidlarni xlor bilan siqb chiqarib olinadi:

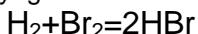


2. HBr sintez yoki PBr₃ gidrolizidan olinadi:



Kimyoviy xossalari:

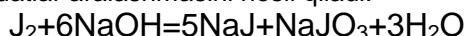
1. Brom vodorod bilan qizdirilganda reaksiyaga kirishadi:



2. Brom kuchli oksidlovchi hisoblanadi:



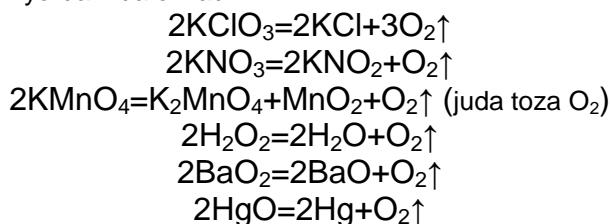
3. Yod ishqorda erib yodid va yodatlar aralashmasini hosil qiladi:



Mavzu: Kislород – O₂

Olinishi:

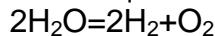
1. Parchalash reaksiyalari yordamida olinadi:



2. Ishqoriy metallar peroksidlarini CO₂ bilan ta'siridan:



3. Sanoatda havoni fraksiyalab, yoki suvni elektroliz qilib:



Kimyoviy xossalari:

1. Moddalar O₂ da yonganda oksidlar hosil bo'ladi:



2. Barcha organik moddalar O₂ da yonadi:

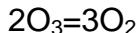


3. Fosfor O₂ da yonganda 2 xil oksid hosil bo'ladi:



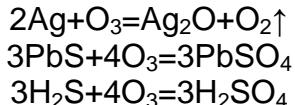
Olinishi:

1. Ozonatorda olinadi:



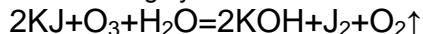
Kimyoviy xossalari:

1. Ozon juda kuchli oksidlovchi:



Sifat reaksiya:

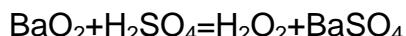
1. KJ eritmasi orqali o'tkazilganda binafsha rangli yod cho'kmasi hosil bo'ladi:



Mavzu: Vodorod peroksid – H₂O₂

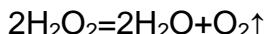
Olinishi:

1. BaO₂ ga H₂SO₄ ta'siridan:

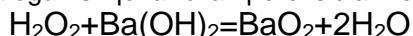


Kimyoviy xossalari:

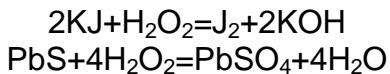
1. MnO₂ katalizatori ishtirokida oson parchalanadi:



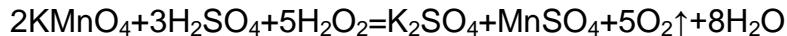
2. H₂O₂ kuchsiz kislota xossasiga ega. Ishqorlar bilan peroksidlar hosil qiladi:



3. H₂O₂ kuchli oksidlovchi hisoblanadi:



4. H₂O₂ kuchli oksidlovchilar bilan oksidlanadi:



Mavzu: Oltingugurt – S

Minerallari:

FeS – temir sulfid; FeS₂ – pirit; ZnS – rux aldamasi; CaSO₄·2H₂O – gips; CaSO₄ – angidrit; BaSO₄ – barit; Na₂SO₄·10H₂O – Glauber tuzi; MgSO₄·7H₂O – taxir tuz.

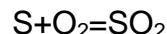
Olinishi:

1. Sanoatda quyidagicha olinadi:

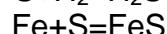
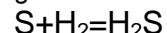


Kimyoviy xossalari:

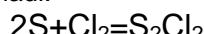
1. Oltingugurt kislorodda yonib SO₂ ni hosil qiladi:



2. Oltingugurt metallar va vodorod bilan qizdirilgan ta'sirlashadi:



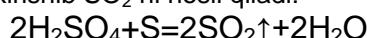
3. S qizdirilganda xlor bilan reaksiyaga kirishadi:



4. S qaynoq ishqorda disproporsialanadi:



5. S kons. H₂SO₄ bilan reaksiyaga kirishib SO₂ ni hosil qiladi:



Mavzu: Vodorod sulfid – H₂S

Olinishi:

1. FeS ni HCl bilan ta'siridan:

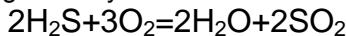


2. Sintez usul bilan:

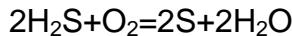


Kimyoviy xossalari:

1. U to'la yonganda ko'kish alanga berib yonadi:



2. U chala yonganda S hosil bo'ladi:



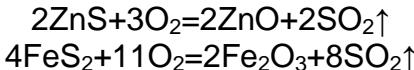
Mavzu: Oltingugurt(IV)oksid - SO_2

Olinishi:

1. S yoki H_2S ning yonishidan:



2. Sulfid va piritni yoqib olinadi:

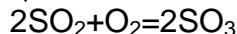


3. Misning konsentrangan sulfat kislota bilan ta'siridan:



Kimyoviy xossalari:

1. U kislotalar ishtirokida yonib SO_3 ni hosil qiladi:



2. U suvda erib kuchsiz kislota – sulfit kislotani hosil qiladi:



Mavzu: Oltingugurt (VI) oksid – SO_3

Olinishi:

1. SO_2 ning oksidlanishidan hosil bo'ladi:



Kimyoviy xossalari:

1. U suv bilan sulfat kislotani hosil qiladi. Bu reaksiya kuchli ekzotermik:



2. Sulfat kislotada erib oleumni hosil qiladi. Uning asosiy tarkibi pirosulfat kislota hisoblanadi:



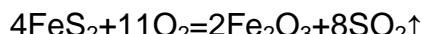
Mavzu: Sulfat kislota – H_2SO_4

Olinishi:

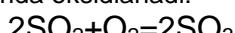
1. Nitroza

2. Kontakt. Bu 2 usul bir-biri bilan faqat SO_2 ning oksidlanish jarayonida farq qiladi:

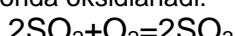
1-bosqich. Sulfid rudalari yoqiladi:



2-bosqich. Nitroza usulida SO_2 NO katalizatorida oksidlanadi:



Kontakt usulida SO_2 V_2O_5 katalizatorida oksidlanadi:



3-bosqich. SO_3 kons. sulfat kislotaga yuttiriladi:

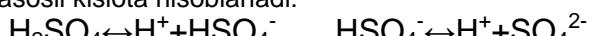


Oleum suvga yuttiriladi:

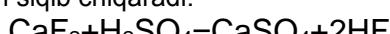


Kimyoviy xossalari:

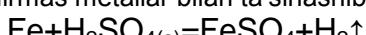
1. Sulfat kislota kuchli ikki asosli kislota hisoblanadi:



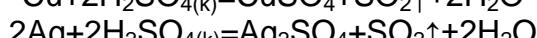
2. U ko'pchilik kislotalarni tuzlaridan siqib chiqaradi:

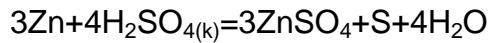


3. Suyultirilgan H_2SO_4 ko'pchilik nodirmas metallar bilan ta'sirlashib vodorod ajralib chiqadi:

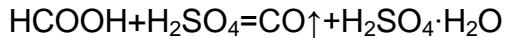
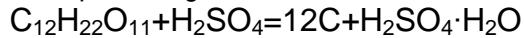


4. Konsentrangan sulfat kislotadan hech qanday metal vodorodni siqib chiqara olmaydi. Chunki u kuchli oksidlovchi.





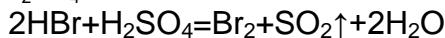
5. Konsentrangan sulfat kislota ko'pchilik organik moddalardan suvni tortib oladi:



6. U hatto ko'mirni ham oksidlashi mumkin:



7. Turli qaytaruvchilar kons. H_2SO_4 ta'sirida oksidlanadi:



Mavzu: Tiosulfat kislota – $\text{H}_2\text{S}_2\text{O}_3$

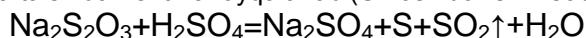
Olinishi:

1. Tiosulfatlar sulfitlarga oltingugurt qo'shib qizdirib olinadi:



Kimyoviy xossalari:

1. Tiosulfatlar sulfat kislota ta'siridan eritma loyqalanadi (S hosil bo'lism hisobiga):



Mavzu: Selenat kislota – H_2SeO_4

Kimyoviy xossalari:

1. Selenat kislota juda kucli oksidlovchi. Unda hatto oltin eriydi:

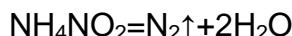


Mavzu: Azot – N_2

Olinishi:

Laboratoriya da.

1. NH_4NO_2 ni parchalab olinadi:



2. $(\text{NH}_4)_2\text{Cr}_2\text{O}_7$ ni parchalab olinadi:

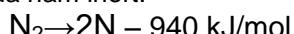


Sanoatda.

Havoni fraksion kondensatlab va distillab olinadi. Bunda birinchi bo'lib azot haydaladi.

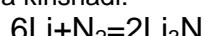
Kimyoviy xossalari:

Azot molekulasi barqarorligi uchun u juda ham inert.

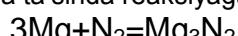


U passivligi jihatidan inert gazlardan keyin ikkinchi o'rinda turadi:

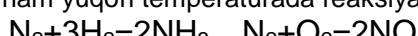
1. U faqat Li bilan xona sharoitida reaksiyaga kirishadi:



2. U qolgan aktiv metallar bilan temperatura ta'sirida reaksiyaga koirshib nitridlar hosil qiladi:



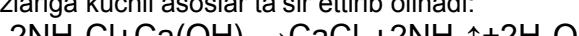
3. U vodorod va metalmaslar bilan ham yuqori temperaturada reaksiyaga kirishadi:



Mavzu: Ammiak – NH_3

Olinishi:

1. Laboratoriya da ammoniy tuzlariga kuchli asoslar ta'sir ettirib olinadi:



2. Nitridlarning parchalanishidan:



3. Sanoatda sintez usuli bilan olinadi:



Kimyoviy xossalari:

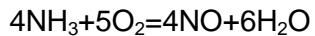
1. Suvdag'i eritmasi asos xossasiga ega:



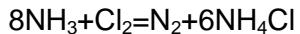
2. Ammiak katalizatorsiz yonganda N_2 hosil bo'ladi:



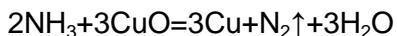
3. U katalizator ishtirokida yonganda NO ni hosil qiladi:



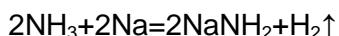
4. Ammiakning suvli eritmasi xlor ishtirokida oksidlanadi:



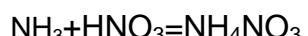
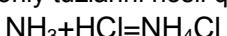
5. CuO ta'sirida oksidlanadi:



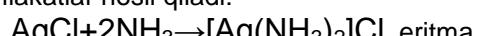
6. Aktiv metallar bilan amidlar hosil bo'ladi:



7. Kislotalar ta'sirida ammoniy tuzlarini hosil qiladi:



8. NH₃ d-metallar tuzlari bilan ammiakatlar hosil qiladi:



Mavzu: Gidrazin – N₂H₄

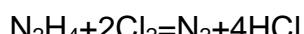
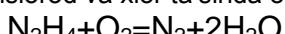
Olinishi:

1. NH₃ ni NaOCl bilan oksidlab olinadi:



Kimyoviy xossalari:

1. U kislorod va xlor ta'sirida oksidlanadi:



Mavzu: Gidroksilamin – NH₂OH

Olinishi:

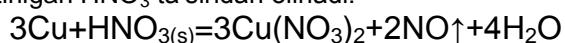
1. HNO₃ ning katod qaytarilishidan hosil bo'ladi.



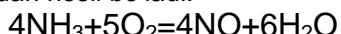
Mavzu: Azot (II) oksidi – NO

Olinishi:

1. Labaratoriya misga suyultirilgan HNO₃ ta'siridan olinadi:



2. Sanoatda NH₃ ni katalitik oksidlanishidan hosil bo'ladi:



Kimyoviy xossalari:

1. U havoda qo'ng'ir rangli gaz hosil qiladi:



Mavzu: Nitrit kislota – HNO₂

Kimyoviy xossalari:

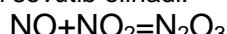
1. Nitrit kislota kuchli oksidlovchilar ta'siridan oksidlanib nitratlarga o'tadi:



Mavzu: Azot(III)oksid – N₂O₃

Olinishi:

1. NO va NO₂ ning ekvimolyar aralashmasini sovutib olinadi:



Mavzu: Azot(IV)oksid – NO₂

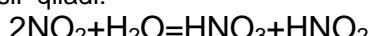
Olinishi:

1. Cu ni kons.HNO₃ da eritib olinadi:

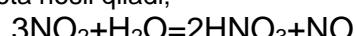


Kimyoviy xossalari:

1. Suvda erib kislotalar aralashmasini hosil qiladi:



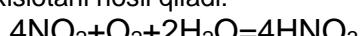
2. Agar u issiq suvda erisa faqat bitta kislota hosil qiladi;



3. Ishqorlarda erib tuzlar aralashmasini hosil qiladi.



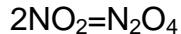
4. U kislorod ishtirokida suvda erib nitrat kislotani hosil qiladi:



Mavzu: Azot(IV)oksidi dimeri - N₂O₄

Olinishi:

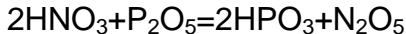
1. NO₂ ning sovushidan hosil bo'ladi. Bunda qo'ng'ir rang yo'qoladi:



Mavzu: Azot(V)oksid – N₂O₅

Olinishi:

1. Nitrat kislotani degidratlab olinadi:



Kimyoviy xossalari:

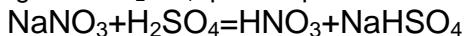
1. U suvda erib nitrat kislotani hosil qiladi:



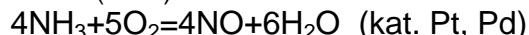
Mavzu: Nitrat kislota – HNO₃

Olinishi:

1. Laboratoriyada quruq nitratlarga kons. H₂SO₄ qo'shib qizdirib olinadi:



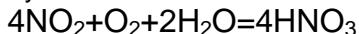
2. Sanoatda Ostvald usulida olinadi (1915):



U keyin katalitik oksidlanadi:

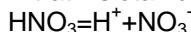


Hosil bo'lgan NO₂ va O₂ suvgaga yuttiriladi:

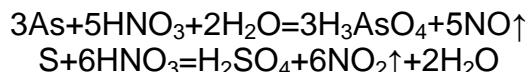


Kimyoviy xossalari:

Nitrat kislota kuchli oksidlovchi kislota



1. Nitrat kislota deyarli barcha metallar (Au, Pt, Os, Ir, Nb, Ta va W dan tashqari) va metalmaslar bilan ta'sirlashadi;



2. Standart elektrod potensial qatorida vodoroddan keyin turgan metallar kons. HNO₃ da erib NO₂ ni hosil qiladi:



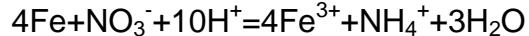
Agar suyultirilgan nitrat kislota ishlatisa NO hosil bo'ladi:



3. Zn – Pb qatorida metallar NO ni hosil qiladi:

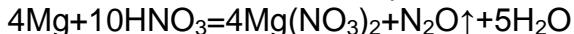


Juda suyultirilganda k-ta bilan NH₄⁺ hosil bo'ladi;



Kons HNO₃ ta'sirida Fe, Al, Cr passivlashadi:

4. Li – Mg qatoridagi metallar kons. HNO₃ bilan N₂O hosil qiladi:



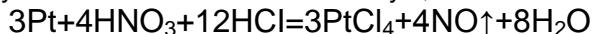
Juda suyultirilgan HNO₃ bilan NH₄NO₃ ni hosil qiladi



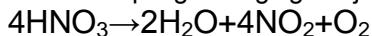
Faqat Co erkin azorgacha qaytariladi:



5. 1 mol kons. HNO₃ va 3 mol kons. HCl aralashmasi juda kuchli oksidlovchi hisoblanadi. U "zar suvi" yoki "podsho arog'i" deyladi. U hatto Au va Pt ni oksidlaydi;



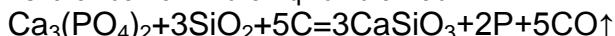
6. Nitrat kislota yorug'lik ta'sirida parchalanib qo'ng'ir rangli gaz ajratadi:



Mavzu: Fosfor – P

Olinishi:

Kalsiy fosfatni qum ishtirokida ko'mir bilan qizdirib olinadi:



Kimyoviy xossalari:

1. Uishqoriy va ishqoriy yer metallari bilan ta'sirlashib fosfidlar hosil qiladi:



2. Fosfor metalmaslar bilan ta'sirlashib qaytaruvchilik xossasini namoyon qiladi:



3. U kons. HNO_3 bilan ta'sirlashadi:

**Olinishi:**

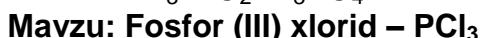
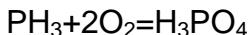
1. Kalsiy fosfidni gidroliz qilib olinadi:

**Kimyoviy xossalari:**

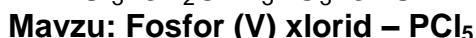
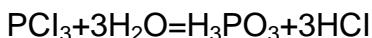
1. U NH₃ ga qaraganda kuchsiz asos xossasini namoyon qiladi:



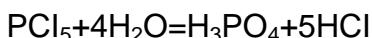
2. U havoda fosfat kislotagacha oksidlanadi:



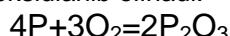
U gidrolizga uchraydi:



U gidrolizga uchraydi:

**Olinishi:**

1. Fosfor kislorod yetishmagan muhitda oksidlanib olinadi:

**Kimyoviy xossalari:**

1. U havoda qizdirilganda oksidlanadi:



2. U suvda fosfit kislotani hosil qiladi:

**Kimyoviy xossalari:**

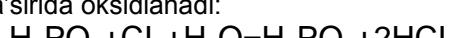
1. U ishqorlar bilan 2 xil tuzlarni hosil qiladi:



2. Fosfit kislota qizdirilganda disproprisilanadi:



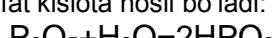
3. Fosfit kislota oksidlovchilar ta'sirida oksidlanadi:

**Olinishi:**

1. Fosforni mo'l kislorodda yoqib olinadi;

**Kimyoviy xossalari:**

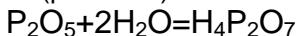
1. P₂O₅ soviq suvda eriganda metafosfat kislota hosil bo'ladi:



2. P₂O₅ issiq suvda eriganda ortofosfat kislota hosil bo'ladi:



3. Bu 2 mol suv bilan ta'sirlashsa difosfat (pirofosfat) kislota hosilbo'ladi;



Mavzu: Ortofosfat kislota - H_3PO_4

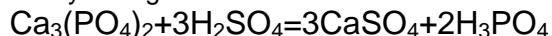
Olinishi:

1. Laboratoryada fosfat kislota fosforni 32% li nitrat kislotada eritib olinadi:

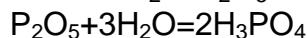
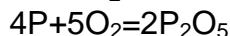
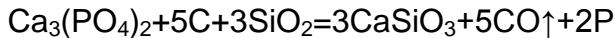


2. Sanoatda fosfat kislota ekstraksiyon va termik usulda olinadi:

a) Ekstraksiyon usulda maydalangan fosforit sulfat kislotada eritiladi:

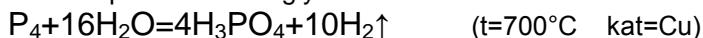


b) Termik usulda fosforitdan fosfor olinadi. Olingan fosfor kislorodda oksidlantirilib P_2O_5 olinadi. P_2O_5 suvda eritilib fosfat kislota olinadi:



Termik usulda olingan fosfat kislota ekstraksiya usulida olingan kislotaga qaraganda tozaligi va yuqori konsentratsiyali bo'lishi bilan ajralib turadi.

3. Kelajakda fosfat kislota chiqindisiz texnologiya asosida olinishi mumkin:

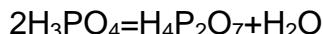


Kimyoviy xossalari:

1. U uch xil tuzlarni hosil qiladi:



2. U $200^{\circ}C$ da qizdirilganda pirofasfat kislota hosil bo'ladi:



Mavzu: Fosforli o'g'itlar

Fosforli o'g'itlarga quyidagilar kiradi:

1. Fosforit yoki appatit – $Ca_3(PO_4)_2$ (16-35% P_2O_5)

2. Oddiy superfosfat :



3. Qo'sh superfosfat:



4. Pretsipitat:



Mavzu: Azot saqlovchi o'g'itlar

1. Ammofos- $NH_4H_2PO_4$

2. Diammofos- $(NH_4)_2HPO_4$

3. Karbamid- $CO(NH_2)_2$

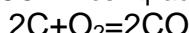
Karbamid yuqori bosimda ammiakni karbonat angidrid bilan reaksiyasidan hosil bo'ladi:



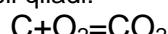
Mavzu: Uglerod

Kimyoviy xossalari:

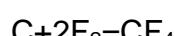
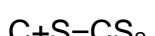
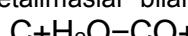
1. Uglerod kislorod yetishmagan muhitda CO ni hosil qiladi.



2. Uglerod kislorodda to'la yonib CO_2 ni hosil qiladi:



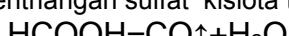
3. U ko'pchilik metallmaslar bilan ta'sirlashadi:



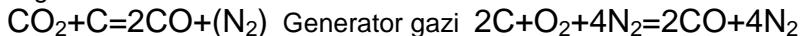
Mavzu: Uglerod(II)oksid – CO

Olinishi:

Laboratoriyada chumoli kislotaga konsentrangan sulfat kislota ta'siridan olinadi:



1. Sanoatda quyidagi usullar bilan olinadi:

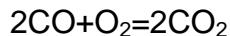


2. Shuningdek suv gazi ko'inishida olinadi:

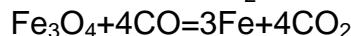
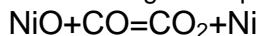


Kimyoviy xossalari:

1. U havoda ko'kish alanga berib yonadi:



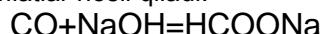
2. U juda ko'p metallarning oksidlarini erkin metallargacha qaytaradi(karbotermiya):



3. CO ba'zi d- metallar bilan birikib karbonillar $\{\text{Me}(\text{CO})_x\}$ hosil qiladi:



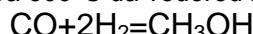
4. CO ishqorlar bilan ta'sirlashib formiatlar hosil qiladi.



5. CO xlор va oltingugurt bilan ta'sirlashadi;



6. U ZnO kat. ishtirokida yuqori bosim va 300°C da vodorod bilan birikib metanolni hosil qiladi:



Mavzu: Uglerod(IV)oksid – CO_2

Olinishi:

Laboratoriya da CaCO_3 ga HCl ta'siridan Kipp apparatida olinadi:



1. Sanoatda ohaktosh kuydirilganda hosil bo'ladi:



2. Barcha organik moddalar va ko'mir yonganda ham CO_2 hosil bo'ladi:



Kimyoviy xossalari:

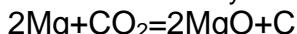
1. Uglerod bilan qaytarilganda CO hosil bo'ladi:



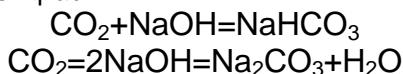
2. Suvda erib kuchsiz kislota karbonat kislotani hosil qiladi:



3. CO_2 yonishga yordam bermasa ham aktiv metallar unda yonadi:

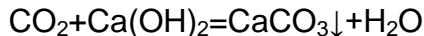


4. Ishqorlarda erib karbonatlarni hosil qiladi:



Sifat reaksiya:

CO_2 ajralib chiqayotganligi $\text{Ca}(\text{OH})_2$ yoki $\text{Ba}(\text{OH})_2$ orqali shu gazni o'tkazib aniqlanadi. Bunda eritma loyqalanadi:

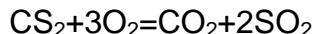


Mavzu: Uglerod disulfid – CS_2

Ko'mirni oltingugurt bilan qizdirib olinadi:



U tez alanganuvchan modda:

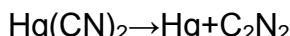


Mavzu: Galogenidlari – CF_4



Mavzu: Uglerodning azotli birikmalari

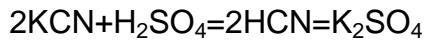
1. Disian – C_2N_2



2. Vodorod sianid – HCN

Olinishi:

1. Sianidlarga kislota ta'siridan olinadi:

**Boshqa birikmlari:**

Sianat kislota – H-O-C≡N

Izosianat kislota – H-N=C=O

Tiosianat kislota – H-S-C≡N (Rodanidlar)

Izotiosianat kislota – H-N=C=S

Mavzu: Karbidlar

- Atsetilenidlar – ishqoriy/ishqoriy-yer metallarining karbidlari.

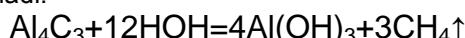
Li_2C_2 , Na_2C_2 , MgC_2 , CaC_2 , SrC_2 , BaC_2

Ular gidrolizidan atsetilen hosil bo'ladi:



- Metanidlar – Al_4C_3 va Be_2C

Ular gidrolizidan metan hosil bo'ladi:



- Olmossimon karbidlar – B_3C va SiC . Charxtoshlarda ishlataladi:

- Fe_3C – sementit.

Mavzu: Karbonat kislota – H_2CO_3 **Olinishi:**

- CO_2 ni suvg'a yuttirib olinadi;

**Kimyoviy xossalari:**

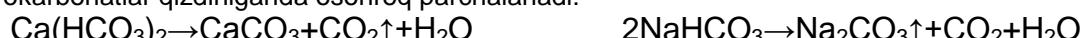
- Karbonat kislota kuchsiz ikki asosli kislota

Mavzu: Karbonatlar**Kimyoviy xossalari:**

- Li_2CO_3 dan boshqa ishqoriy metallarning karbonatlari parchalanmasdan suyuqlanadi. Qolganlari qizdirilganda parchalanadi:



- Gidrokarbonatlar qizdirilganda osonroq parchalanadi:

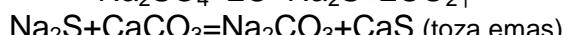
**Mavzu: Soda ishlab chiqarish**

Na_2CO_3 – soda, kalsinirlangan soda; NaHCO_3 – ichimlik soda

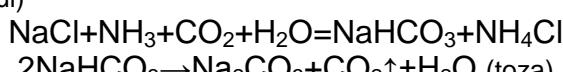
$\text{Na}_2\text{CO}_3 \cdot 10\text{H}_2\text{O}$ – kristall soda

Olinishi:

- Leblan usuli:



- Solvey usuli (Ammiakli usul)



- Elektrolitik usul:

**Mavzu: Kremniy - Si****Olinishi:**

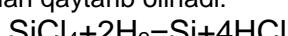
- SiF_4 ni K bilan qaytarib olinadi:



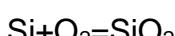
- SiO_2 ni Mg yoki Al bilan qaytarib olinadi:



- Juda toza kremniy SiCl_4 ni vodorod bilan qaytarib olinadi:

**Kimyoviy xossalari:**

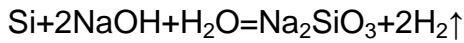
- Yuqori temperaturada kislorodda yonadi:



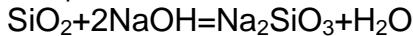
- Faqat ftor bilan odatdag'i sharoitda ta'sirlashadi:



- Issiq ishqorlarda eriydi:



4. SiO_2 ishqorlarda erib silikatlarni hosil qiladi:

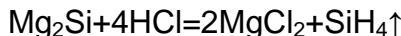


5. Uni faqat ftorid kislota eritadi:



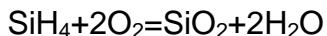
Olinishi:

1. Magniy silitsidga xlorid kislota ta'siridan olinadi:

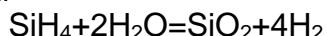


Kimyoviy xossalari:

1. Silan havoda o'z-o'zidan alangalanadi:

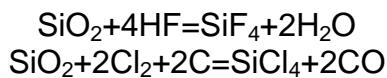


2. U suv bilan quyidagicha ta'sirlashadi:



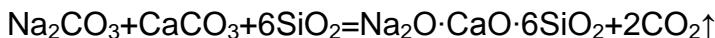
Mavzu: Kremniyning galogenli birikmalari – $\text{Si}\Gamma_4$

Olinishi:



Mavzu: Shisha

Olinishi:



$\text{K}_2\text{O} \cdot \text{CaO} \cdot 6\text{SiO}_2$ – Eruvchan shisha

Mavzu: Metallarning olinish usullari

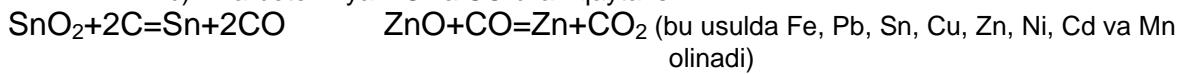
Metallarning olinishi 3 ga bo'linadi:

1. Pirometallurgiya – yuqori temperaurada biror qaytaruvchi bilan qaytarish. Qaytaruvchiga ko'ra yana farqlanadi:

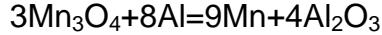
a) Vodorodotermiya – H_2 bilan qaytarish:



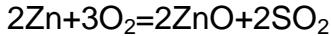
b) Karbotermiya – C va CO bilan qaytarish:



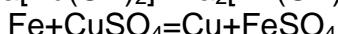
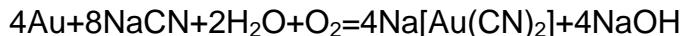
c) Alyumotermiya – Al bilan qaytarish:



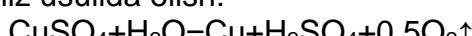
Pirometallurgiyada agar metal sulfid holida bo'lisa, avval yoqilib oksid holiga keltiriladi:



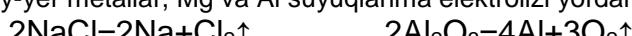
2. Gidrometallurgiya – avval metal eruvchan holga keltirilib eritmaga o'tkaziladi, so'ngra biror aktivroq metal bilan qaytariladi:



3. Elektrometallurgiya – elektroliz usulida olish:



Ishqoriy metallar, ishqoriy-yer metallar, Mg va Al suyuqlanma elektrolizi yordamida olinadi:



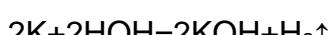
Mavzu: Ishqoriy metallar

Kimyoviy xossalari:

1. Kislorodda yonib faqat Li oksid hosil qilsa, qolganlari peroksid va nadperoksidlar hosil qiladi:



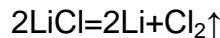
2. Suv bilan shiddatli ta'sirlashadi:



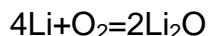
Mavzu: Litiy – Li

Olinishi:

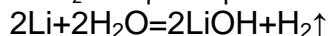
LiCl va KCl suyuqlanmasini elektroliz qilib olinadi:

**Kimyoviy xossalari:**

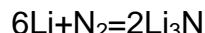
1. Kislородда yониб oксид hosil qiladi:



2. Suv bilan odatdagи sharoitda ta'sirlashib H_2 ni siqib chiqaradi:



3. Li azot bilan xona temperaturasida ta'sirlashasi:



4. Li barcha galogenlar, S va H_2 bilan ta'sirlashadi:



5. Li magniya diagonal o'xshash element. Uning karbonati, fosfat va fosfiti kam eriydi. Li_2CO_3 qizdirilganda parchalanadi:

**Mavzu: Natriy – Na****Minerallari:**

NaCl – tosh tuz

$\text{NaCl} \cdot \text{KCl}$ – silvinit

$\text{Na}_2\text{SO}_4 \cdot 10\text{H}_2\text{O}$ – Glauber tuzi

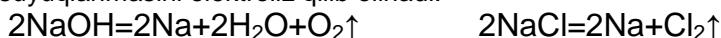
$\text{Na}_3[\text{AlF}_6]$ – Kriolit

NaNO_3 – Chili selitrasи

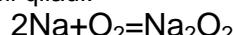
$\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O}$ – bura

Olinishi:

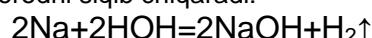
1. NaOH yoki NaCl suyuqlanmasini elektroliz qilib olinadi:

**Kimyoviy xossalari:**

1. Kislород bilan ta'sirlashib peroksid hosil qiladi:



2. Suv bilan shiddatli ta'sirlashib vodorodni siqib chiqaradi:



3. Galogenlar, S va H_2 bilan ta'sirlashadi:

**Mavzu: Kaliy – K****Minerallari:**

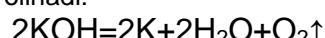
KCl – silvin

$\text{KCl} \cdot \text{MgCl}_2 \cdot 6\text{H}_2\text{O}$ – karnallit

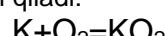
$\text{KCl} \cdot \text{MgSO}_4 \cdot 3\text{H}_2\text{O}$ – kainit

Olinishi:

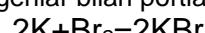
1. KOH suyuqlanmasini elektroliz qilib olinadi:

**Kimyoviy xossalari:**

1. Kislородда yониб asosan superoksidni hosil qiladi:



2. Suv bian yonish orqali ta'sirlashadi. Galogenlar bilan portlaydi:

**Mavzu: Berilliy – Be****Olinishi:**

1. BeF_2 ni vakuumda Ca bilan qizdirib olinadi:

**Kimyoviy xossalari:**

1. Kislородda qizdirilganda yonadi:



2. Suyultirigan kislotalarda eriydi:



3. Ishqorlar bilan berillatlar hosil qiladi:

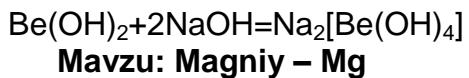


4. Berilliy karbid metanid hisoblanadi:



5. $\text{Be}(\text{OH})_2$ amfoterlik xossasini namoyon qiladi;





Minerallari:

MgCO_3 – magnezit

$\text{CaCO}_3 \cdot \text{MgCO}_3$ – dolomit

$\text{KCl} \cdot \text{MgCl}_2 \cdot 6\text{H}_2\text{O}$ – karnallit

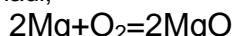
Olinishi:

1. MgCl_2 suyuqlanmasini elektroliz qilib olinadi:



Kimyoviy xossalari:

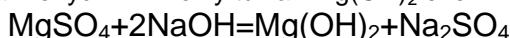
1. Havoda Mg sirti oksid parda bilan qoplanadi;



2. Sovuq suv bilan sekin, issiq suv bilan tez ta'sirlashib, H_2 ni siqib chiqaradi:



3. Mg(OH)_2 oq kukun, suvda kam eriydi. Ammoniy tuzlari Mg(OH)_2 cho'kmsi tushishiga xalaqit beradi:



Mavzu: Kalsiy – Ca

Minerallari:

CaCO_3 – bor, ohaktosh, marmar, kalsit

CaF_2 – flyuorit, plavik shpat

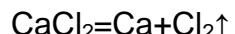
$\text{CaCO}_3 \cdot \text{MgCO}_3$ – dolomit

$\text{Ca}_3(\text{PO}_4)_2$ – fosforit

$\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$ – gips

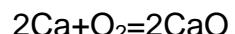
Olinishi:

1. CaCl_2 suyuqlanmasini elektroliz qilib olinadi:



Kimyoviy xossalari:

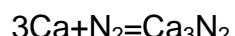
1. Kislorodda yonib oksid hosil qiladi:



2. Suv bilan shiddatli ta'sirlashib ohakli suvni hosil qiladi:



3. Azot ta'sirida nitrid hosil qiladi:



Mavzu: Kalsiy birikmalari

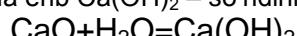
Olinishi:

1. CaO ohaktoshni parchalab olinadi:

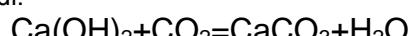


Kimyoviy xossalari:

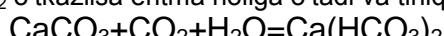
1. CaO – so'ndirilmagan ohak. U suvda erib Ca(OH)_2 – so'ndirilgan ohakni hsoil qiladi:



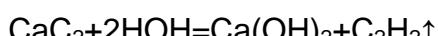
2. Ca(OH)_2 CO_2 ta'sirida loyqalanadi:



3. CaCO_3 suspenziyasi orqali CO_2 o'tkazilsa eritma holiga o'tadi va tiniqlashadi:



4. CaC_2 atsetilenid hisoblanadi:



Mavzu: Suvning qattiqligi va uni yo'qotish usullari

Vaqtinchalik qattiqlikni yo'qotish:

1. Qaynatib yo'qotiladi:

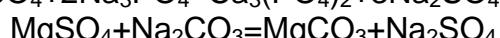
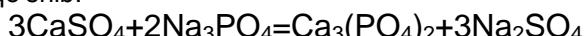


2. Ohak qo'shib yo'qotiladi:



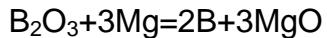
Doimiy qattiqlikni yo'qotish:

1. Soda yoki natriy fosfat qo'shib:

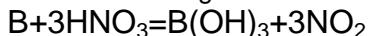


2. Kationit yoki anionitdan foydalilanadi.

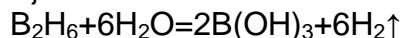
Mavzu: Bor – B

Olinishi:**Kimyoviy xossalari:**

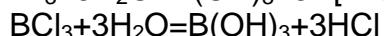
1. Bor suyultilgan kislotalarda erimaydi. U konsentrlangan nitrat kislotada eriydi:



2. Diboran B_2H_6 gidrolizdan vodorod ajraladi:



3. BF_3 gidrolizidan tetraftorborat kislotasi, qolganlaridan galogenovodorodlar hosil bo'ladi:



Mavzu: Alyuminiy – Al

Minerallari:

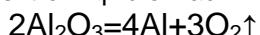
$\text{Na}_3[\text{AlF}_6]$ – kriolit

Al_2O_3 – korund

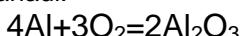
$\text{Al}_2\text{O}_3 \cdot 2\text{H}_2\text{O}$ – bokosit

Olinishi:

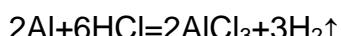
1. Al_2O_3 ning kriolitdagi suyuqlanmasini elektroliz qilib olinadi:

**Kimyoviy xossalari:**

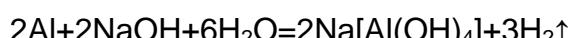
1. Al havoda yupqa oksid pardasi bilan qoplanadi:



2. Suyultirilgan kislotalarda eriydi:

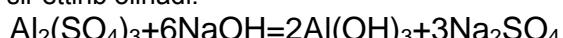


3. Ishqorlarda oson eriydi:



4. Kons. HNO_3 da erimaydi;

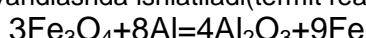
5. $\text{Al}(\text{OH})_3$ tuzlariga ishqor ta'sir ettirib olinadi:



6. Al(OH)₃ amfoter;



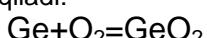
7. Granula holidagi Al metallarni payvandlashda ishlataladi(termit reaksiysi):



Mavzu: Germaniy – Ge

Kimyoviy xossalari:

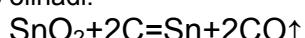
1. Ge kislordoda yonganda (IV) oksid hosil qiladi:



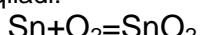
Mavzu: Qalay – Sn

Olinishi:

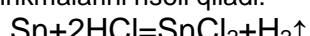
1. SnO_2 ni 1000°C ko'mir bilan qaytarib olinadi:

**Kimyoviy xossalari:**

1. Sn kislordoda yonganda (IV) oksid hosil qiladi:



2. Suyultirilgan kislotalarda erib Sn(II) birikmalarini hsoil qiladi:



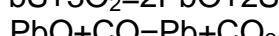
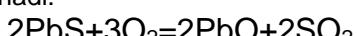
3. Kons. sulfat kislotada erib Sn(IV) birikmalarini hsoil qiladi:



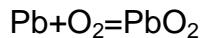
Mavzu: Qo'rg'oshin – Pb

Olinishi:

1. Asosan pirometallurgiya usulida olinadi:

**Kimyoviy xossalari:**

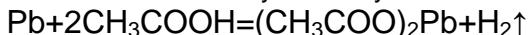
1. Havoda oksidlanguanda PbO_2 ni hosil qiladi:



2. Nitrat kislotada oson eriydi:



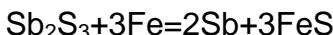
3. Suyultirilgan kislotalarda, masalan sırka kislotada yaxshi eriydi:



Mavzu: Surma va vismut – Sb va Bi

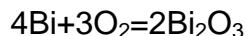
Olinishi:

1. Pirometallurguya usuli bilan olinadi:



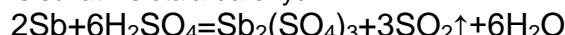
Kimyoviy xossalari:

1. Sb va Bi kislroddha yonadi:

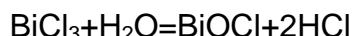


2. Sb va Bi elektrod potensiallar qatorida H_2 dan o'ngda joylashganligi uchun suyultirilgan kislotalarda erimaydi.

3. Sb va Bi kons. nitrat va kons. sulfat kislotalarda eriydi:



4. Sb va Bi ning galogenidlari gidrolizga uchraydi:



Mavzu: Temir – Fe

Minerallari:

Fe_3O_4 – magnetit, magnitli temirtosh

Fe_2O_3 – gematit, qizil temirtosh

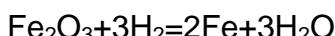
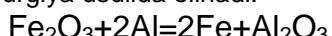
FeCO_3 – siderit

FeS_2 – pirit

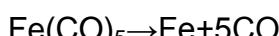
FeS – temir kolchedani

Olinishi:

1. Asosan pirometallurgiya usulida olinadi:



2. Juda toza temir karbonillardan olinadi:



Kimyoviy xossalari:

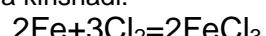
1. Temir havoda oksidlanadi:



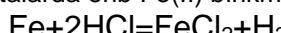
2. Temir yuoqri temperaturada suv bug'lari bilan ta'sirlashadi:



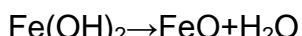
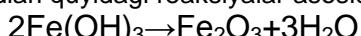
3. Xlor va S bilan reaksiyaga kirishadi:



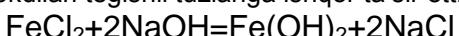
4. Suyultirilgan kislotalarda erib Fe(II) birikmalarini hosil qiladi:



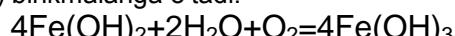
5. Temirning oksidlari quyidagi reaksiyalar asosida olinadi:



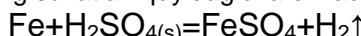
6. Gidrokdilari tegishli tuzlariiga ishqor ta'sir ettirib olinadi:



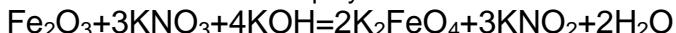
7. $\text{Fe}(\text{OH})_2$ oson oksidlanib Fe(III) birikmalariga o'tadi:



8. Temirning sulfatlari quyidagicha olinadi:



9. Temirning (+VI) birikmlari temir birikmalarini ishqoriy muhitda oksidlanishidan hosil bo'ladi:



10. Sariq qon tuzi va qizil qon tuzi quyidagi reaksiyalar asosida olinadi:



11. Sariq qon tuzi va qizil qon tuzi Fe^{3+} va Fe^{2+} ionlarini aniqlashda ishlatalidi:



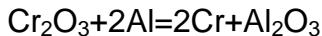
Mavzu: Xrom – Cr

Minerallari:

$\text{Cr}_2\text{O}_3 \cdot \text{FeO}$ – xromit, xromli temirtosh PbCrO_4 – krokoit

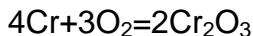
Olinishi:

1. Alyumotermiya usulida olinadi:

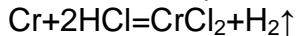


Kimyoviy xossalari:

1. Oksidlanganda Cr_2O_3 hosil bo'ladi:

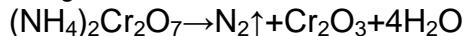


2. Suyultirilgan kislotarlarda erib Cr(II) birikmalarini hosil qiladi:

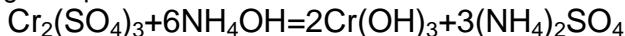


3. Sovuq kons. HNO_3 da erimaydi. Chunki passivlashadi.

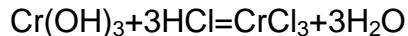
4. Cr_2O_3 ammoniy bixromat parchalanganda hosil bo'ladi:



5. Cr(OH)_3 xrom(III) tuzlariga ishqor ta'sir ettirib olinadi:



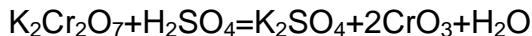
6. Cr(OH)_3 amfoter hisoblanadi:



7. Xrom(II) birikmali beqaror. CrO Cr(OH)_2 parchalanganda hosil bo'ladi:



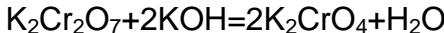
8. Xrom(VI) oksid kaliy bixromat eritmasiga kons. H_2SO_4 ta'siridan yorqin qizil ignasimon kristall holida cho'kadi:



9. H_2CrO_4 – xromat kislota. O'rtacha kuchdag'i ikki asosli kislota.

10. $\text{H}_2\text{Cr}_2\text{O}_7$ – bixromat kislota. Kuchli ikki asosli kislota.

11. Xromatlar ishqoriy va neytral muhitda barqaror. Kislotali muhitda bixromatlarga o'tadi:



12. Xromatlar kuchli oksidlovchilar hisoblanadi. Kislotali muhitda Cr(III) birikmalariga qaytariladi:



Mavzu: Molibden – Mo

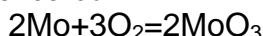
Olinishi:

1. Alyumotermiya yoki vodorodotermiya usulida olinadi:



Kimyoviy xossalari:

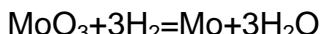
1. Yuqori temperaturada kislorod bilan ta'sirlashadi:



Mavzu: Volfram – W

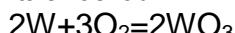
Olinishi:

1. Vodorodotermiya usulida olinadi:



Kimyoviy xossalari:

1. Juda yuqori temperaturada kislorod bilan ta'sirlashadi:



2. Deyarli hech qaysi kislotada erimaydi.

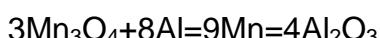
Mavzu: Marganets – Mn

Minerallari:

MnO_2 – pirolyuzit Mn_3O_4 – gausmanit MnS – marganets aldamasi

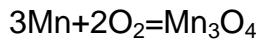
Olinishi:

1. Alyumotermiya usulida olinadi:



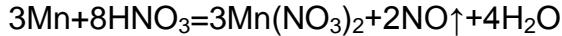
Kimyoviy xossalari:

1. Havoda yonganda Mn_3O_4 hosil bo'ladi:



2. Suyultirilgan kislotalarda erib Mn(II) birikmalarini hosil qiladi:
$$\text{Mn} + 2\text{HCl} = \text{MnCl}_2 + \text{H}_2 \uparrow$$

3. Suyultirilgan nitrat kislotada eriydi:

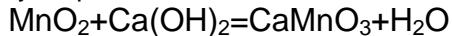


4. Mn(II) birikmlari asos xossaga ega. Mn(OH)₂ suvda yaxshi erimasa ham kuchli asos hisoblanadi:
$$\text{MnCl}_2 + 2\text{NaOH} = \text{Mn}(\text{OH})_2 + 2\text{NaCl}$$

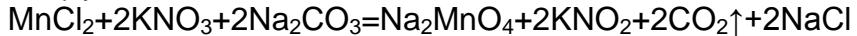
5. MnO₂ oksidlovchilik xossasini namoyon qiladi:



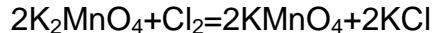
6. MnO₂ amfoterlik xossasini namoyon qiladi:



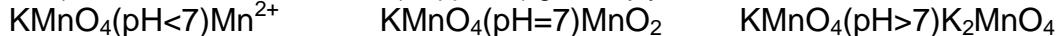
7. Mn(VI) birikmalari quyi birikmlarini oksidlanishidan hosil bo'ladi:



8. Mn(VII) birikmalari uning quyi birikmalarining oksidlanishidan hosil bo'ladi:



9. KMnO₄ kuchli oksidlovchi hisoblanadi. Kislotali muhitda Mn²⁺ (*rangsiz*), neytral muhitda MnO₂ (*qo'ng'ir cho'kma*) va kislotali muhitda MnO₄²⁻ (*to'q yashil*) gacha qaytariladi:



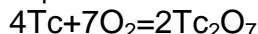
10. KMnO₄ qizdirilganda parchalanadi:



Mavzu: Texnetsiy va Reniy – Tc va Re

Kimyoviy xossalari:

1. Kislorodda yonganda +7 birikmlarini hsoil qiladi:



2. Reniy elektrod potensiallar qatorida H₂ dan o'ngda turadi. Shuning uchun suyultirilgan kislotalarda erimaydi:



3. HReO₄ kuchli kislota.

Mavzu: Rux – Zn

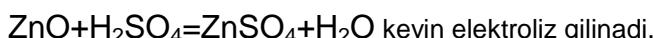
Olinishi:

1. Ikki xil usulda: quruq va eritmada olinadi:

Quruq:

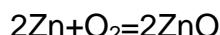


Eritmada:



Kimyoviy xossalari:

1. Kislorodda oksidlanadi:



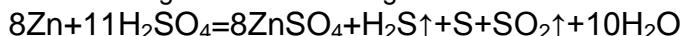
2. Suyultirilgan kislotalarda tez eriydi:



3. Rux amfoter bo'lganligi uchun ishqorlarda eriydi:



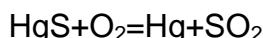
4. Konsentrangan sulfat kislotada eriganda cho'kma va gazlar aralshmasi hosil bo'ladi:



Mavzu: Simob – Hg

Olinishi:

1. Kinovarni yoqib olinadi:



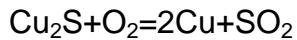
Mavzu: Mis – Cu

Minerallari:

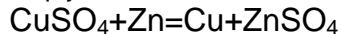
CuFeS₂ – xalkopirit Cu₂S – xalkozin, mis yaltirog'i CuCO₃·Cu(OH)₂ – malaxit

Olinishi:

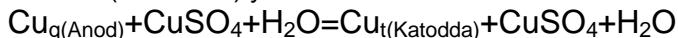
1. Sulfid rudalarini yoqib olinadi:



2. Mis tuzlarini eritmada Fe yoki Zn bilan qaytarib olinadi:

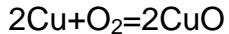


3. Toza mis eritma elekrtolizidan (rafinirlash) yo'li bilan olinadi:



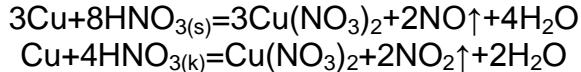
Kimyoviy xossalari:

1. Havoda qizdirilganda qorayib CuO ni hosil qiladi:

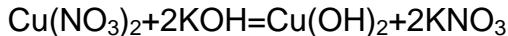


2. Standart elektrod potensiallar qatorida H₂ dan o'ngda turganligi uchun suyultirilgan kislotalarda erimaydi.

3. Har qanday konsentratsiyali nitrat kislotada eriydi:



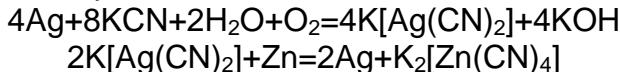
4. Cu(OH)₂ ko'k paxtasimon chokma holida cho'kadi:



Mavzu: Kumush – Ag

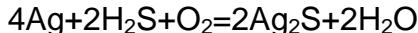
Olinishi:

1. Sulfid rudalaridan siyanidli usul yordamida olinadi:



Kimyoviy xossalari:

1. Ag nodir metal hisoblanadi. H₂S bilan kislород ishtirokida ta'sirlashadi:



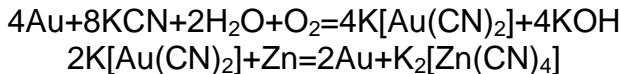
2. Har qanday konsentratsiyali HNO₃ da eriydi:



Mavzu: Oltin – Au

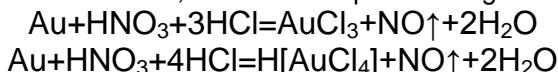
Olinishi:

1. Siyanidli usul yordamida olinadi:



Kimyoviy xossalari:

1. Oltin eng nodir metal hisoblanadi. U havo, kislota va ishqorlar ta'siriga chidamli. Zar suvida eriydi:



Mavzu: Titan – Ti

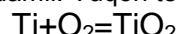
Olinishi:

1. Titan quyidagicha olinadi:

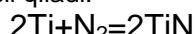


Kimyoviy xossalari:

1. Titan suv va mineral kislotalar ta'siriga chidamli. Yuqori temperaturada kislород bilan ta'sirlashadi:



2. Yuqori temperaturada azot bilan nitrid hosil qiladi:



3. Granula holida HNO₃ da eriydi:



4. Granula holida issiq sulfat kislotada eriydi:



5. Kons.H₂SO₄ bilan qizdirilganda reaksiyaga kirishadi:



3-Qism

**ORGANIK
KIMYO**

*** Nazariy ma'lumot**

***Ma'lumotlar**

ORGANIK KIMYO

I BOB. ORGANIK BIRIKMALAR TUZILISHI NAZARIYASI VA ORGANIK REAKSIYALARING TURLARI.

Mavzu: Organik kimyo faniga kirish.

“Organik kimyo” fani termini 1808 yil shved olimi Bersellius tomonidan fanga kiritilgan.

XVIII asr o’rtalarida juda kam organik moddalar o’rganilgan. Masalan, sırka chumoli, benzoy va qahrabo kislotalar, Sheele tabiiy moddalardan vino, sut, limon, olma kislotalarni va glitserinni ajratib olgan.

Dastlabki organik sintezlar quyidagilar:

1824 yil Vyoler ditsiandan oksalat kislotani sintez qildi, 1828 yil ammoniy sianatdan mochevinani sintez qildi.

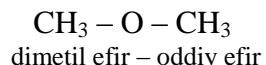
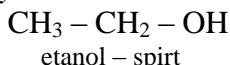
1845 yil Kolbe sırka kislotani, Bertlo chumoli kislotani, etil spirit, C_2H_2 , benzol, metan va yog’ni sintez qildi.

Ta’rif: “Organik kimyo – uglerod brikmalarining kimyosidir”.

1861 yil rus olimi A.M. Butlerov “organik birikmalar tuzilishi nazariyasini” e’lon qiladi. Uning asosiy hollari quyidagilar:

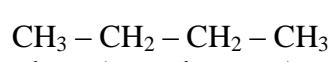
1. Organik birikmalarda uglerod atomlari o’zaro ma’lum tartibga ko’ra birikadi.
2. Moddaning kimyoviy xossasi uning miqdoriy va sifat tarkibiga bog’liq.
3. Agar bitta molekula bir xil miqdordagi va bir xil molekulyar massaga ega bo’lgani bilan, tuzilishi har xil bo’lsa, u holda izomeriya hodisasi kuzatiladi.
4. Moddaning kimyoviy xossasi modda molekulasi tarkibidagi atomlar guruhi o’zaro ta’siriga bog’liq.

Ta’rif: Tarkibi va molekulyar massasi bir xil, tuzulishi va xossalari har xil bo’lgan moddalar izomerlar deyiladi.

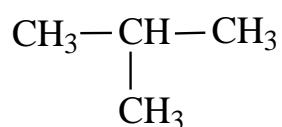


Izomeryaning bir qancha turlari bor:

- 1) *Uglerod skeleti bo’yicha izomeriya:*

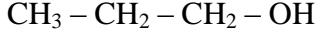


n-butan (tarmoqlanmagan)

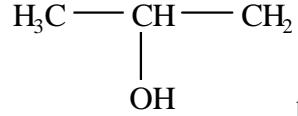


izobutan (tarmoqlangan)

- 2) *Holat izomeriya, funksional guruh holatiga ko’ra izomeriya:*

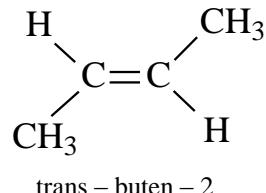
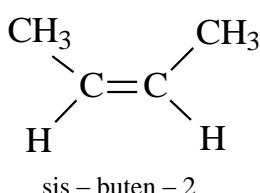


propanol -1



propanol -2

- 3) *Geometrik izomeriya (sis-, trans - izomeriya), qo’shbog’ tutgan organik birikmalarda kuzatiladi:*

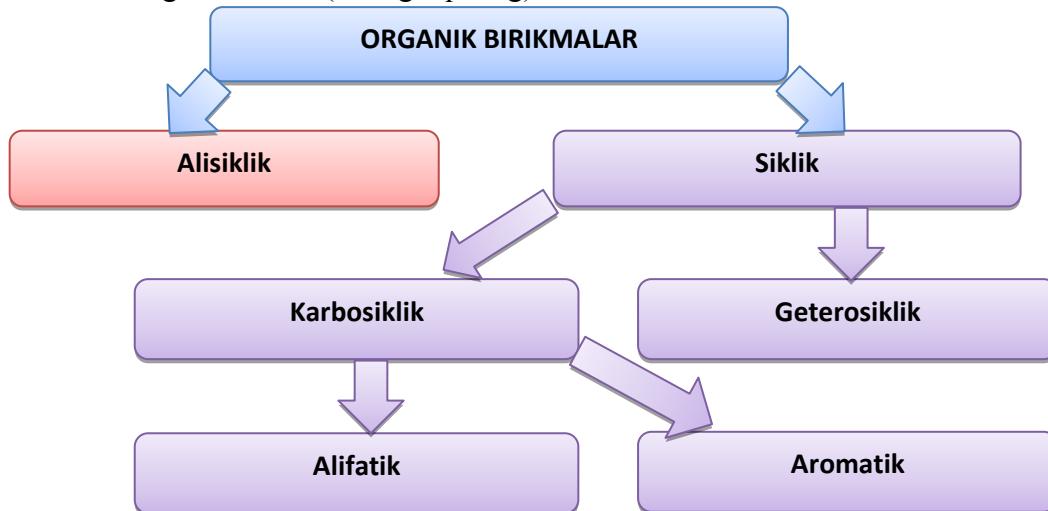


- 4) *Optik izomeriya* – ular qutblangan yorug'lik nuruni o'ng yoki chap tomoniga buradi. Agar o'ngga bursa – D, chapga bursa – L izomer deyiladi.

Mavzu: Organik birirkmalarning klassifikatsiyasi.

Organik birikmalar turlicha klassifikatsiyalanadi.

1. Alisiklik – siklik bo'lмаган birikmalar.
2. Siklik birikmalarga bo'linadi (ilovaga qarang).



Mavzu: Organik reaksiyalarning turlari.

Reaksiyalarni klassifikatsiyalashda kimyoviy o'zgarish va ayni molekuladagi bog' uzilishi hisobga olinadi.

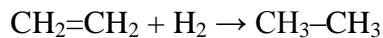
Kimyoviy o'zgarishlarga ko'ra reaksiyalar quyidagicha bo'linadi:

- 1) Birikish
- 2) Ajralish
- 3) O'rın olish.

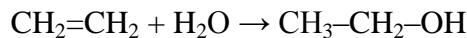
Birikish reaksiyalarida asosan to'yinmagan ya'ni karrali bog'ga ega molekulalar ishtirok etadi va qo'shbog' yoki uchbog' uziladi. Bunday reaksiyalar **A** (*Addition*) harfi bilan belgilanadi.

Bunday reaksiyalarning bir qancha turlari bor:

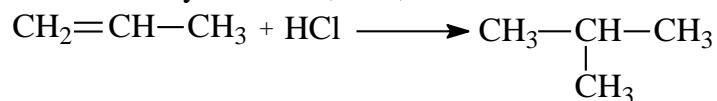
- a) *Gidrogenlanish* – ya'ni Ni, Pt yoki Pd katalizatori ishtirokida vodorod birikish reaksiyasi:



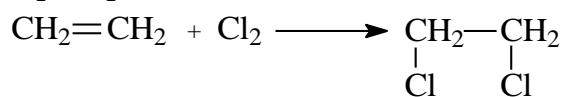
- b) *Gidratlanish* – suv molekulalarining birikishi. Masalan, etilen gidratlanganda etanol hosil bo'ladi:



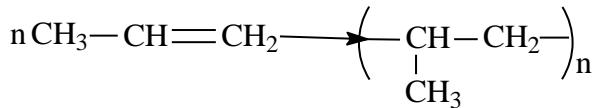
- c) *Galogenvodorod* birikish reaksiyalari – HF, HCl, HBr va HJ birikishi:



- d) *Galogenlanish* – F₂, Cl₂, Br₂ va J₂ birikishi:



- e) *Polimerlanish* reaksiyasi – ya'ni kichik molekulyar massali to'yinmagan molekulalardan yuqori molekulyar birikmalar (YuMB) hosil bo'lish reaksiyalari:



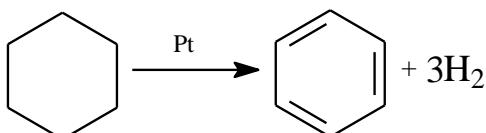
propilen polipropilen

f) *Siklobirikish reaksiyasi* – masalan, Dills-Alder reaksiyasi.

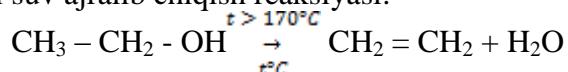
Ajralish reaksiyalarida atom yoki atomlar guruhi ajralib chiqadi va karrali bog'lar hosil bo'ladi. Bunday reaksiyalar **E** (*Elimination*) harfi bilan belgilanadi.

Ajralash reaksiyalarining bir qancha turlari bor:

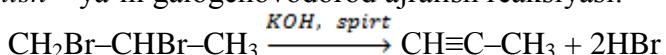
a) *Degidrogenlanish* – ya'ni vodorod ajralish reaksiyasi:



b) *Degidratlanish* – ya'ni suv ajralib chiqish reaksiyasi:



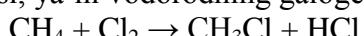
c) *Degidrogalogenlanish* – ya'ni galogenovodorod ajralish reaksiyasi:



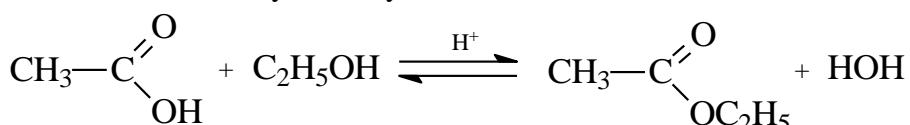
O'rin olish reaksiyalarida molekuladagi atom yoki atomlar guruhi boshqasiga almashinadi. Bunday reaksiyalar **S** (*Substitution*) harfi bilan belgilanadi.

O'rin olish reaksiyalarini organik kimyoda muhim o'rin tutadi.

a) *Galogen almashinish* reaksiyasi, ya'ni vodorodning galogenga almashinishi:

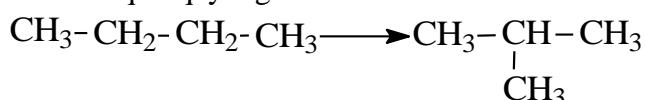


b) *Kondensatsiya* reaksiyasi – quyi molekulyar birikma ajralishi bilan amalga oshadigan reaksiyalar. Masalan, eterifikatsiya reaksiyasi:



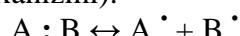
c) *Polikondensatsiya* reaksiyasi – polimerlanishdan farq qilib, quyi molekulyar birikmlar ta'siridan polimer hosil bo'lishi bilan birga past molekulyar massali modda qo'shimcha sifatida hosil bo'ladi. Masalan, fenolformaldegid smolasi, kapron va h.k.lar.

Yuqoridagi 3 ta reaksiyadan tashqari qayta guruhlansih va izomerlanish reaksiya ham uchraydi.



Bog'ning uzilishi va qayta hosil bo'lishiga ko'ra reaksiyalar 2 ga bo'linadi:

1) *Gomogen uzilish* (erkin radikal mexanizmi):



2) *Geterogen uzilish* (ion mexanizm)



II BOB. UGLEVODORODLAR.

Mavzu: Uglevodorodlar.

Ta’rif: Tarkibida faqat uglerod va vodorod atomlari tutgan organik moddalarga uglevodorodlar deyiladi.

Uglevodorodlar 2 ga bo’linadi

1. To’yingan uglevodorodlar – a) alkanlar b) sikloalkanlar
2. To’yinmagan uglevodorodlar – a) alkenlar b) alkinlar c) alkadiyenlar d) polienlar e) arenlar.

Mavzu: Alkanlar

Ta’rif: Tarkibida faqat σ – bog’lar tutgan uglevodorodlarga alkanlar deyiladi.

Ularning umumiy formulasi C_nH_{2n+2} .

Gibrildlanishi – sp³.

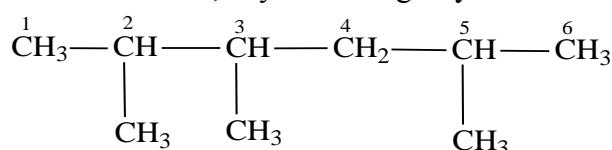
Ta’rif: Uglerod soni ortishi bilan bir yoki bir necha – CH_2 guruhiga farq qiladigan va xossalari o’xshash moddalarga gomologlar deyiladi.

Alkanlarning gomologik qatori

Formulasi	Nomi	Agregat holati	Formulasi	Nomi	Agregat holati
C_1H_4	metan		C_8H_{18}	oktan	
C_2H_6	etan	Gaz	C_9H_{20}	nonan	Suyuq
C_3H_8	propan		$C_{10}H_{22}$	dekan	
C_4H_{10}	butan		$C_{15}H_{32}$	pentadekan	
C_5H_{12}	pentan		$C_{20}H_{42}$	eykozan	Qattiq
C_6H_{14}	geksan	Suyuq	$C_{100}H_{202}$	gektan	
C_7H_{16}	geptan				

Izomeriyasi va nomenklaturasi: alkanlarni IUPAC bo'yicha nomlashda quyidagiga e'tibor beriladi:

- 1) Eng uzun uglerod zanjiri tanlanadi.
- 2) Tarmoqqa yaqin tomondan raqamlanadi va radikal nomi ko'rsatiladi.
- 3) Dastlab kichik radikal, keyin kattarog'i aytiladi.

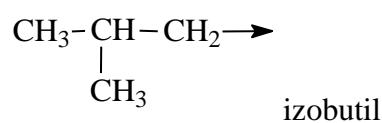
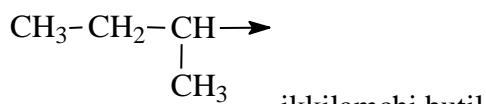


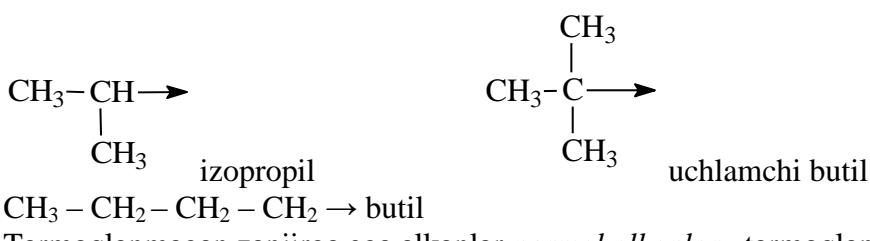
2,3,5-trimetilgeksan

Radikallar bu alkandan vodorod yo'qolganda qolgan qoldiq hisoblanadi va C_nH_{2n+1} formulaga muvofiq keladi.

$CH_3 \rightarrow$ metil
 $CH_3-CH_2 \rightarrow$ etil

$CH_3-C H_2-CH_2 \rightarrow$ propil





Tarmoqlanmagan zanjirga ega alkanlar *normal alkanlar*, tarmoqlanganlari *izo – alkanlar* deyiladi.

Alkanlarda izomeriya – Izomeriya butandan boshlanadi.

$\text{CH}_3 - \text{CH}_2 - \text{CH}_2 - \text{CH}_3$

n-butan

$\begin{array}{c} \text{CH}_3-\text{CH}-\text{CH}_3 \\ | \\ \text{CH}_3 \end{array}$

2 – metil propan (izobutan)

tuzulishi izomeriasi (zanjir) hisoblanadi.

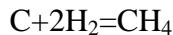
Uglerod soni ortishi bilan izomerlar soni ortadi. Masalan pentanda 3 ta, heptanda 9 ta, oktanda – 18, dekanda – 75.

Tabiatda uchrashi:

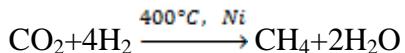
Alkanlarning asosiy manbai – tabiiy gaz va neft hisoblanadi.

Olinishi:

- 1) Uglerodni gidrogenlab (H_2 qo'shib):

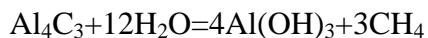


- 2) CO_2 va CO ni katalitik gidrogenlab olinadi:

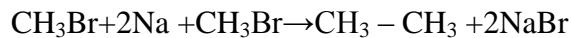
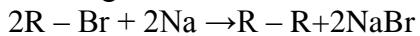


Laboratoriya

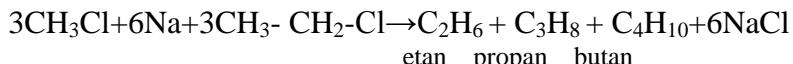
- 1) Al_4C_3 ni gidrolizlab:



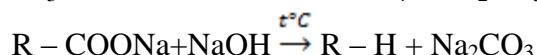
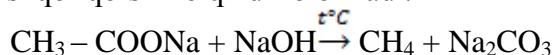
- 2) Vyurs reaksiysi, ya'ni galogenalkenlarga Na ta'siridan olinadi.



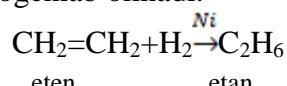
brommetan etan



- 3) Karbon kislota tuzlarini ishqor qo'shilib qizdirib olinadi.



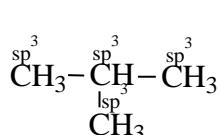
- 4) To'yinmagan uglevodorodlarni gidrogenlab olinadi.



Fizikaviy xossalari:

Alkanlarda uglerod soni ortishi bilan suyuqlanish va qaynash temperaturasi ortadi. Tarmoqlanmagan alkanlar tarmoqlangan alkanlarga qaraganda yuqori suyuqlanish va qaynash temperurasiga ega. Alkanlar suvda erimaydi.

Molekula tuzulishi:



Alkanlarda uglerod atomlari faqat sp^3 gibridlangan holda bo'ladi.

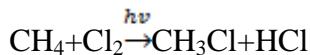
1) $sp^3 - 4 \times 4 = 16$ ta 2) $s - 10$ ta
ga teng.

valent burchagi $109^\circ 28'$

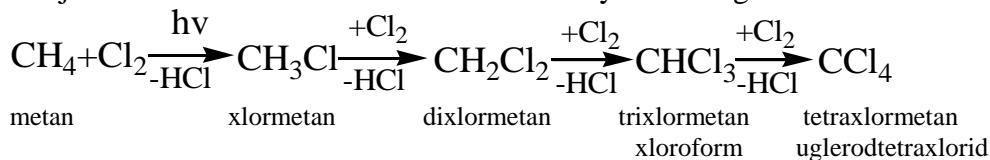
Kimyoviy xossalari:

Alkanlar “parafin” lar ham deyiladi va ancha inert hisoblanadi. Alkanlarda barcha uglerod atomlari vodorod atomlari vodorod atomi bilan to’yinganligi uchun birikish reaksiyalariga kirishmaydi.

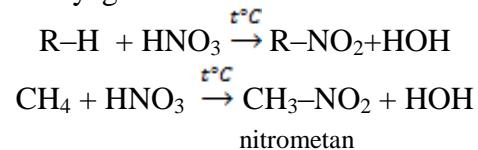
- 1) Alkanlar galogenlar bilan oson ta’sirlashadi:



- 2) Reaksiya zanjir mexanizmi asosida boradi va buni Semyonov ochgan.

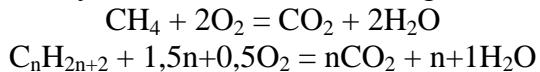


- 3) Alkanlar nitrat kislota bilan reaksiyaga kirishadi:

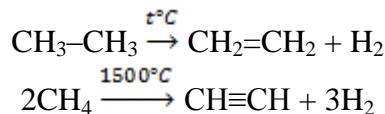


bu reaksiyaga Konovalov reaksiyasi deyiladi.

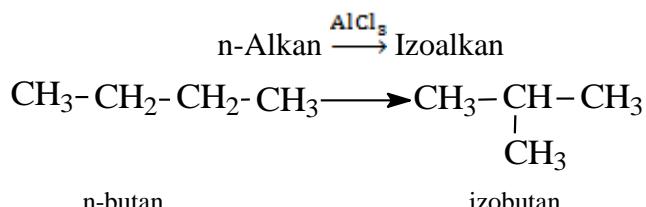
- 4) Alkanlar kislorod ta’sirida oson yonadi. Bunda karbonat angidrid va suv hosil bo’ladi:



- 5) Alkanlar yuqori temperaturada degidrogenlanadi (H_2 molekulasi ajralib chiqib karrali bog’lar hosil bo’ladi):



- 6) Alkanlar $AlCl_3$ yoki $AlBr_3$ ishtirokida qizdirilganda izomerланади. Bunda zanjir tarmoqlanadi:



Ishlatilishi:

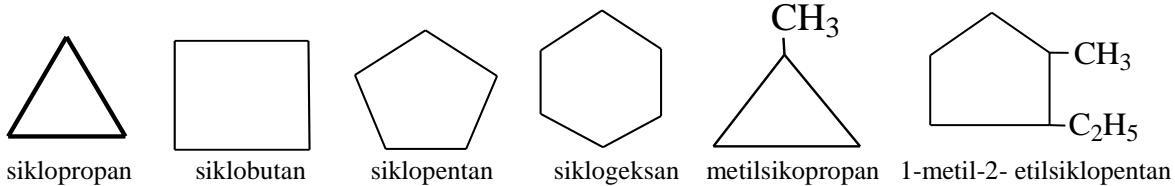
Alkanlar eng asosiysi yoqilg’ilardan biri hisoblanadi. Masalan, CH_4 tabiiy gazning 90-95% gacha qismini tashkil qiladi. Geksan va xloroform qutbsiz erituvchi sifatida, CCl_4 o’t o’chirishda ishlataladi.

Mavzu: Sikloalkanlar.

Ta’rif: Uglerod atomlari o’zaro faqat σ -bo’lgan bilan bog’langan, yopiq xalqali uglevodorodlarga sikloalkanlar deyiladi.

Ularning umumiy formulasi: C_nH_{2n} . Gibridlanishi – sp^3

Nomlanishi: Sikloalkanlarni nomlashda zanjirdagi uglerod soniga “siklo” – so’zi qo’shiladi.

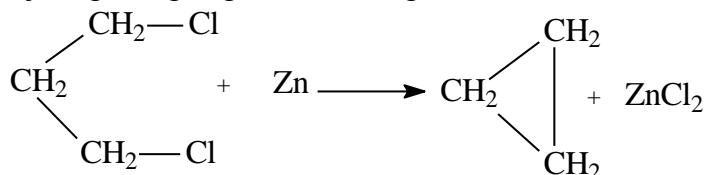


Tabiatda uchrashi:

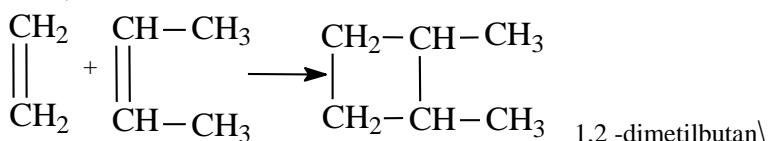
Sikloalkanlar va ularning hosilalari neft va o'simliklar tarkibida uchraydi. Rus olimi V.V. Markovnikov neftdan C_5H_{10} va C_6H_{12} larni ajratib oldi va bu sinfni kashf qiladi.

Olinishi:

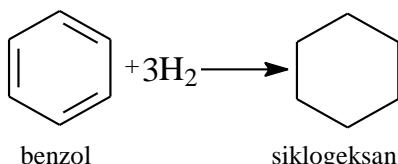
- Ajratilgan digalogenli hosilalarga Zn ta'siridan olinadi.



- Alkenlarni dimerlab olinadi:



- Benzolni gidrogenlab olinadi:



Fizikaviy xossalari:

C_3H_6 , C_4H_8 va C_5H_{10} rangsiz gazlar, C_6H_{12} suyuqlik hisoblanadi.

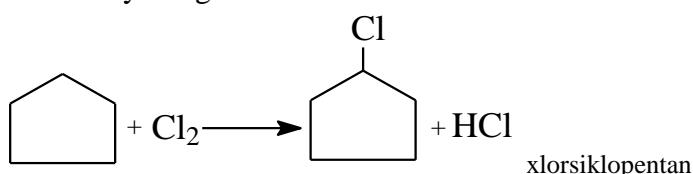
Kimyoiy xossalari:

Sikloalkanlarning dastlabki vakillari bo'lgan siklopropan va siklobutan birikish reaksiyalariga oson kirishadi. Bunda zanjir ochiladi. (Bayer nazariyasi)

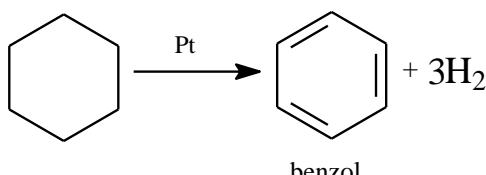
- Sikloalkanlar gidrogenlanganda alkanlar hosil bo'ladi.



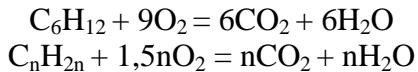
- Siklopantan va siklogeksan ko'proq to'yiganlik xossasini nomoyon qiladi. Shuning uchun o'r'in olish reaksiyalariga oson kirishadi:



- Siklogeksan 300°C da Pt katalizatorida benzolni hosil qiladi:



- 4) Sikloalkanlar yonganda suv va karbonat angidrid hosil bo'ladi:



Ishlatilishi:

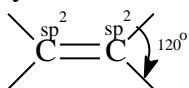
Siklopropan meditsinada og'riqni qoldirish uchun, siklobutan va siklopantan organik sintezda, siklogeksan qutbsiz erituvchi sifatida ishlatiladi. $\text{C}_6\text{H}_6\text{Cl}_6$ – geksaxlorsiklogeksan qishloq xo'jaligida insektitsid sifatida ishlatiladi. Siklopropan narkoz sifatida ishlatiladi.

Mavzu: Alkenlar.

Ta'rif: Tarkibida bitta qo'shbog' tutgan to'yinmagan uglevodorodlarga alkenlar deyiladi.

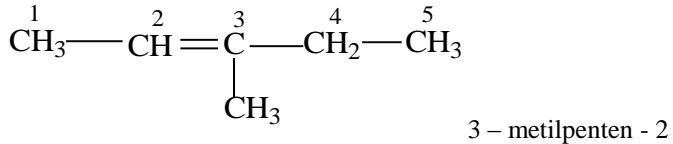
Ularning umumiy formulasi: C_nH_{2n}

Gibrildanishi



Nomlanishi:

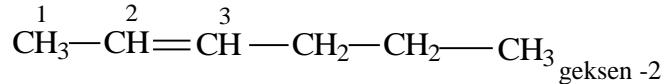
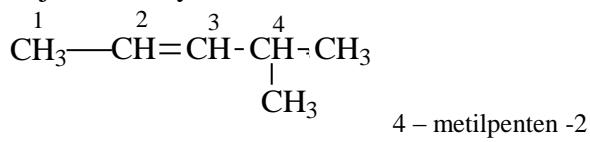
Alkenlarni nomlash uchun tarkibida qo'shbog' tutgan eng uzun uglevodorod zanjiri tanlanib, qo'shbog' yaqin tomondan raqamlanadi va uglevodorod nomiga –en qo'shimchasi qo'shiladi.



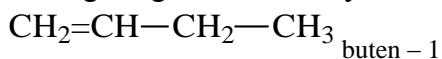
Izomeriyasi:

Alkenlar uchun zanjir izomeriyasi, qo'shbog'ning holat izomeriyasi, shuningdek geometrik izomeriya o'rinni.

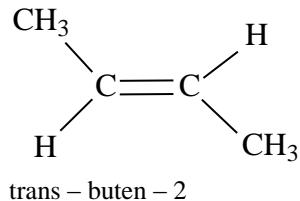
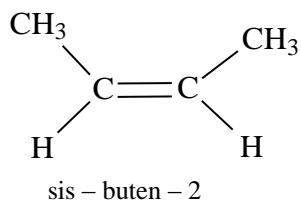
1) Zanjir izomeriya



2) Qo'shbog'ning holat izomeriyasi

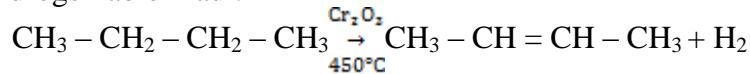


3) Geometrik izomeriyada qo'shbog'da turli o'rinxbosarlar joylashganda qo'shbog' 1 – o'rinda joylashganda yuzaga keladi.

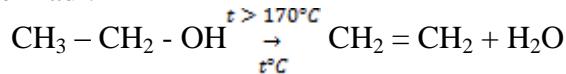


Olinishi:

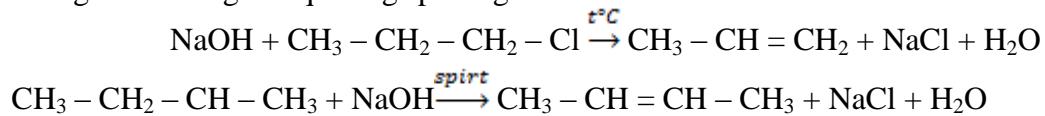
1) Alkanlarni degidrogenlab olinadi:



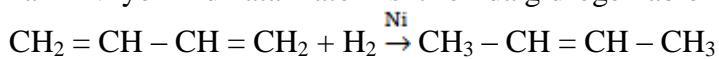
2) Spirtlarni degidratlab olinadi:



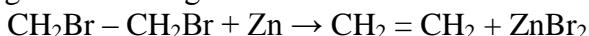
3) Galogenalkanlarga ishqorning spirtdagi eritmasi ta'sir ettirib olinadi:



4) Diyenlar va alkinlarni Ni yoki Pd katalizatori ishtirokida gidrogenlab olinadi:



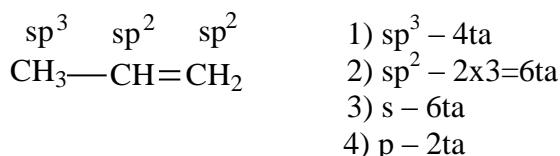
5) Vitsinal digalogenli uglevodorodlarga Zn ta'siridan olinadi:



Fizikaviy xossalari:

$\text{C}_2\text{H}_4 - \text{C}_4\text{H}_8$ – gaz, undan yuqori vakillari suyuqliklardir. Etilen va propilen kuchsiz hidga ega.

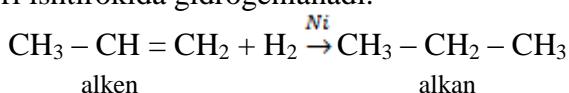
Molekulada tuzulishi:



Kimyoviy xossalari:

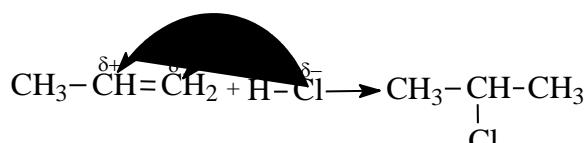
Alkenlarda bitta qo'shbog' bo'lganligi uchun ular to'yinmaganlik xossasini namoyon qiladi va birikish reaksiyalariga oson kirishadi.

1) Alkenlar Ni katalizatori ishtirokida gidrogenlanadi.

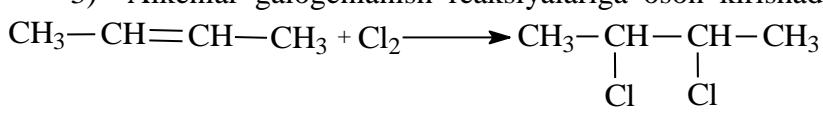


2) Alkenlar vodorod galogenlar bilan ta'sirlashadi. Bunda Markovnikov qoidasiga binoan reaksiya sodir bo'ladi.

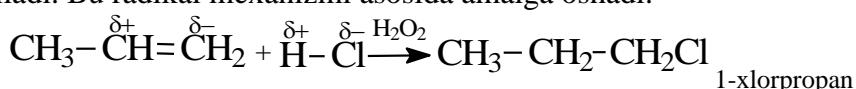
Ta'rif: *Vodorod ko'p gidrogenlangan uglerodga, galogen kam gidrogenlangan uglerodga birikadi.*



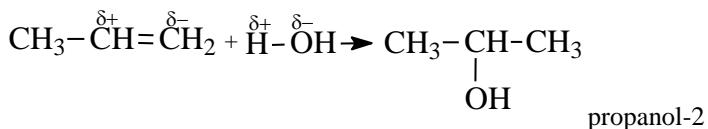
3) Alkenlar galogenlanish reaksiyalariga oson kirishadi. Xlor bromli suvni rangsizlantiradi:



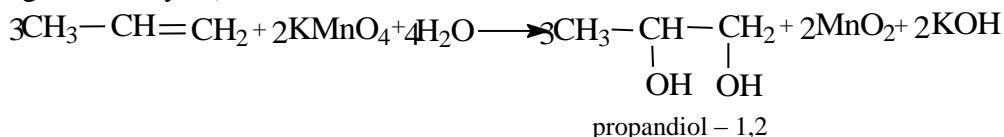
4) Alkenlarga vodorod galogenidlar H_2O_2 ishtirokida biriksa, bunda Markovnikov qoidasiga teskari birikish kuzatiladi. Bu radikal mexanizmi asosida amalga oshadi.



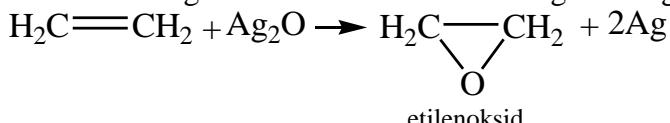
5) Alkenlar H_2SO_4 ishtirokida suv bilan birikib spirtlarni hosil qiladi.



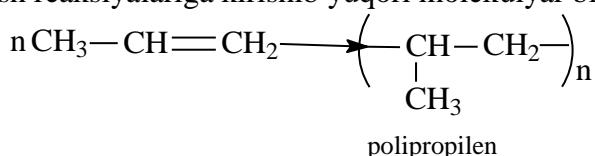
6) Alkenlar KMnO_4 eritmasi bilan oksidlanganda ikki atomli spirtlar hosil bo'ladi. Bunda eritma rangsizlanadi(Vagner reaksiyasi):



7) Alkenlar Ag tuzlari ishtirokida oksidlanganda organik oksidlar hosil bo'ladi:



8) Alkenlar polimerlanish reaksiyalariga kirishib yuqori molekulyar birikmalarini hosil qiladi:



Ishlatilishi:

Etilen etanol olishda, polietilen, polivinylchlorid, etilenglikol olishda; propilen polipropilen olishda, fenol va atseton olishda, glitserin olishda ishlatiladi. Buten kauchik ishlab chiqarishda ishlatiladi.

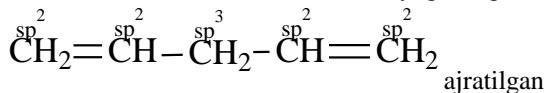
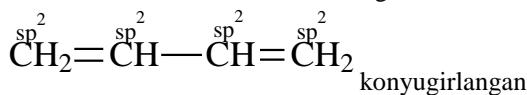
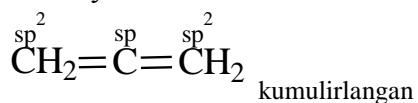
Mavzu: Alkadiyenlar. Diyen uglevodorodlari.

Ta'rif: Tarkibida 2 ta qo'shbog' bo'lgan ochiq zanjirli uglevodorodlar sinfiga alkadiyenlar deyiladi.

Ularning umumiy formulasi: $\text{C}_n\text{H}_{2n-2}$; Gibrildanishi – sp^2

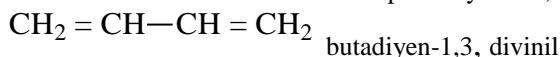
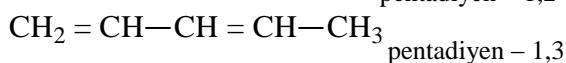
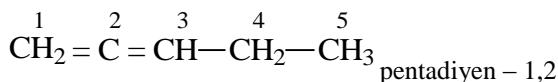
Shuningdek ular alkinlarga izomer hisoblanadi.

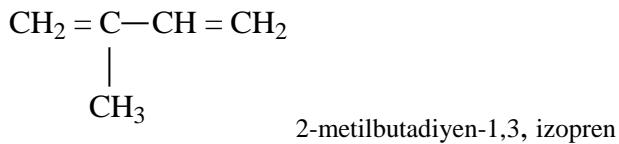
Alkadiyenlar molekulasida ikkita qo'shbog' ajratilmagan yoki ajratilgan bo'lishi.



Nomlanishi va izomeriyasi:

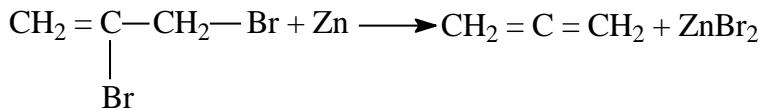
Alkadiyenlar nomlash uchun 2 ta qo'shbog' tutgan eng uzun uglevodorod zanjiri tanlanib qo'shbog' raqami ko'rsatiladi va asosiy uglevodorod nomiga -diyen qo'shimchasi qo'shiladi.



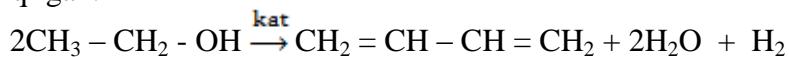


Olinishi:

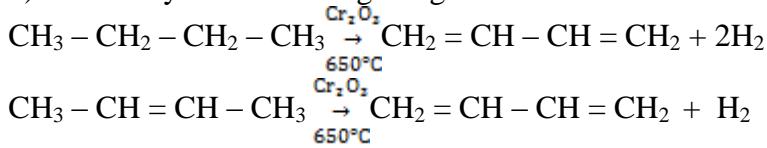
- 1) Alkadiyenlarni digalogenalkenlarga Zn ta'siridan olish mumkin:



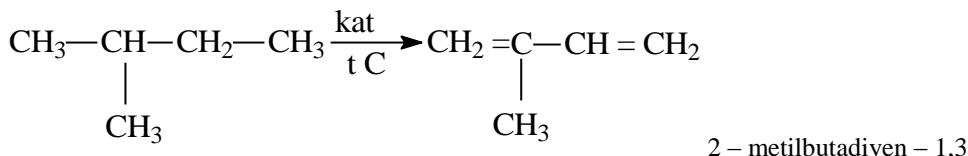
- 2) Butadiyen – 1,3 etil spirtni $\text{Al}_2\text{O}_3 + \text{ZnO}$ katalizatorida qizdirib olinadi. Buni 1927 yil S.V. Lebedev taklif qilgan:



- 3) Buten yoki butenni digedrogenlab olinadi:



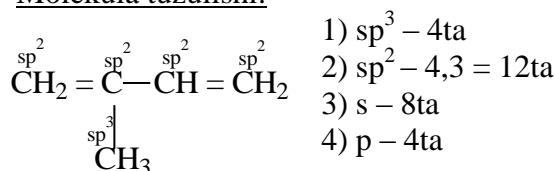
- 4) Izopren izopentenni yoki 2 – metil butenni degidrogenlab olinadi.



Fizikaviy xossalari:

Alkadiyenlarning dastlabki vakillari gaz yoki suyuqliklardir.

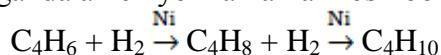
Molekula tuzulishi:



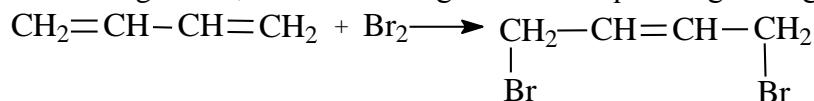
Kimyoviy xossalari:

Alkadiyenlarda 2 ta qo'shbog' bo'lganligi uchun ular to'yinmaganlik xossasini namoyon qiladi va birikish reaksiyalariga oson kirishadi.

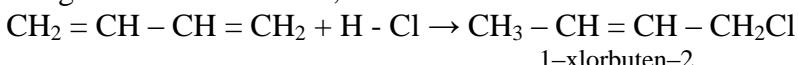
- 1) Alkadiyenlar gidrogenlanganda alken yoki alkanlar hosil bo'ladi:



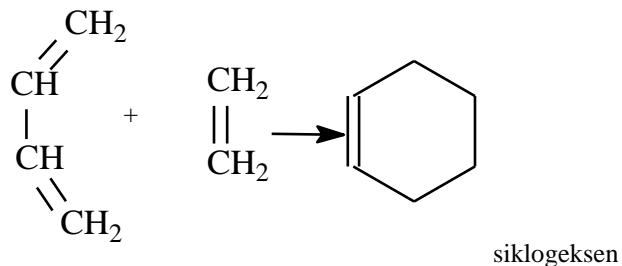
- 2) Alkadiyenlar bromlanganda 1,4 birikish amalga oshadi va qo'shbog' o'rtaga ko'chadi.



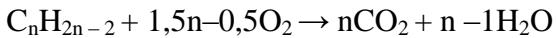
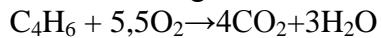
- 3) Galogenvodorodlar ham 1,4 birikadi:



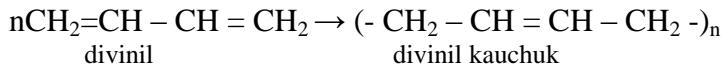
- 4) Alkenlar bilan birikib sikloalkanlar hosil qiladi. Bunga Dills–Alder reaksiyasi deyiladi:



- 5) Alkadiyenlar yonganda suv va karbonat angidrid hosil bo'ladi:



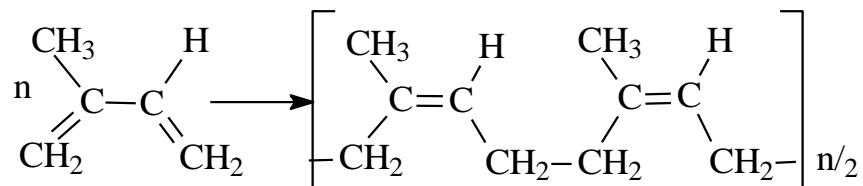
- 6) Alkadiyenlar polimerlanish reaksiyasiga kirishib yuqori molekulyar birikmalarini hosil qiladi:



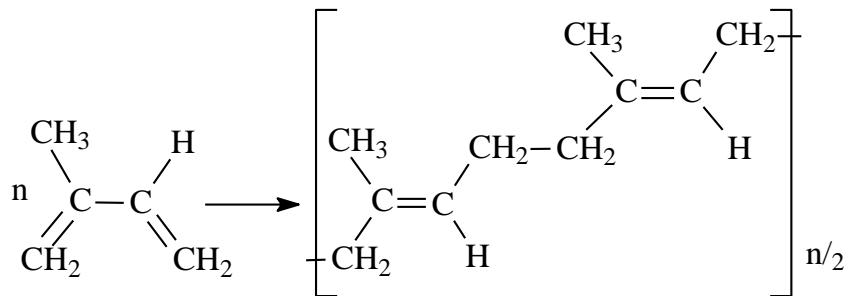
Ishlatilishi:

Ularning polimerlari kauchuk sifatida ishlatiladi. Kauchukning bir necha turlari bor.

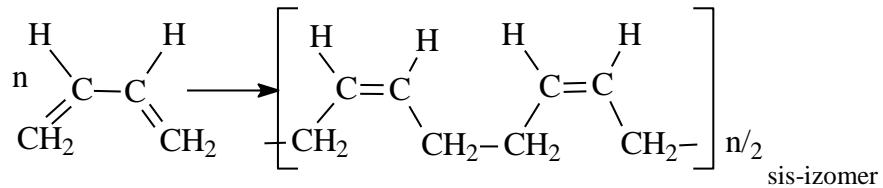
- 1) Tabiiy kauchuk, ya'ni sis – izopren kauchuk:

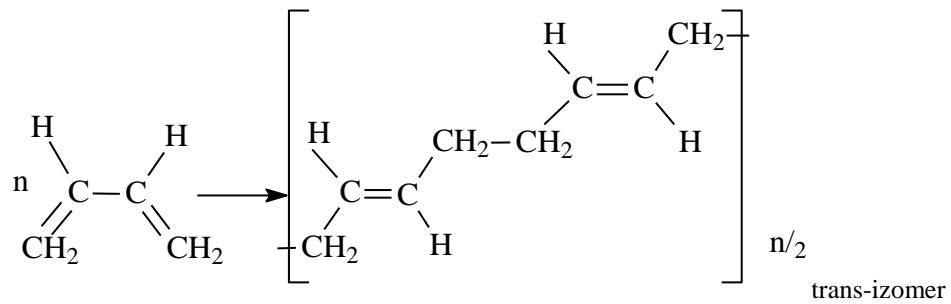


- 2) Trans-izopren kauchuk yoki guttapercha:

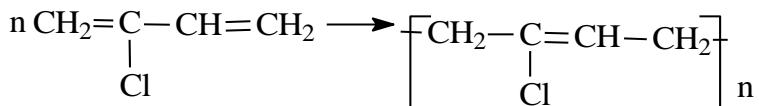


- 3) Divinil kauchuk – butadiyen-1,3 kauchuk:

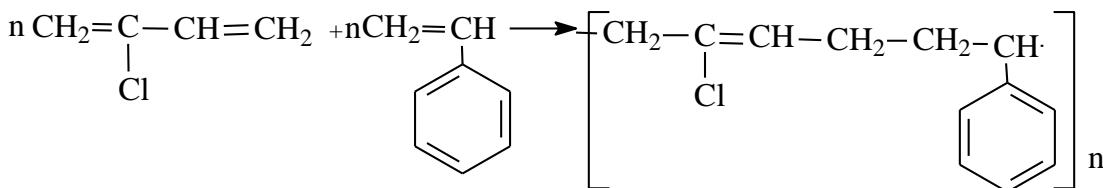




4) Xloropren kauchuk:



5) Butadiyen stirol kauchuk:



Tabiiy kauchuk asosan sis-izoprendan tuzulgan. U Braziliyadagi Geveya daraxtidan olinadi. Tabiiy kauchukning molekulyar massasi 200000 dan 400000 gacha bo'ladi.

Kauchukning xossalarini yaxshilash uchun unga ishlov beriladi, ya'ni vulkanlanadi. Bunda oltingugurt qo'shib qizdiriladi. Agar 50% dan kam S qo'shilsa rezina, 50% dan ko'p S qo'shilsa ebonit hosil bo'ladi. Shuningdek kauchukni "sovuv vulkanlash" uchun S_2Cl_2 ishlatiladi. Tabiiy kauchuk rezina olish uchun, sintetik kauchuk xo'jalik mollari (shisha, penoplast, rezina asboblar) olishda ishlatiladi.

Mavzu: Alkinlar. Atsetilen qatori uglevodorodlari.

Ta'rif: *Tarkibida bitta uchbog' tutgan ochiq zanjirli to'yinmagan uglevodorodlarga alkinlar deyiladi.*

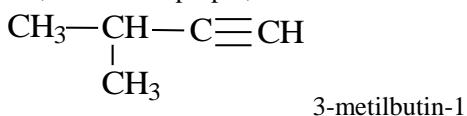
Ularning umumiy formulasi: $\text{C}_n\text{H}_{2n-2}$

Gibrildlanishi – sp $\begin{array}{c} \text{sp} \\ | \\ - \text{C} \equiv \text{C} - \end{array}$

Nomlanishi va izomeriyasi:

Alkinlarni nomlash uchun tarkibida uchbog' tutgan eng uzun uglevodorod zanjiri tanlanib, uchbog' yaqin tomondan raqamlanadi va – in qo'shimchasi qo'shiladi.

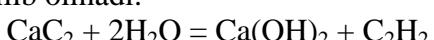
$\text{CH} \equiv \text{CH}$	$\text{CH}_3 - \text{C} \equiv \text{CH}$	$\text{CH} \equiv \text{C} - \text{C}_2\text{H}_5$	$\text{CH}_3 - \text{C} \equiv \text{C} - \text{CH}_3$
etin, atsetilen	propin, metilatsetilen	butin, etilatsetilen	butin-2, dimetilatsetilen



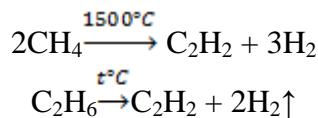
Olinishi:

Alkinlar asosan parchalanish reaksiyalari yordamida olinadi.

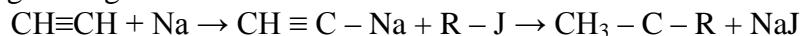
1) Atsetilen CaC_2 ni gidroliz qilib olinadi:



2) Metan yoki etanni piroliz qilib olinadi:



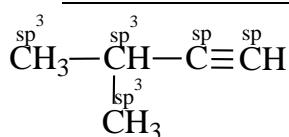
3) Atsetilenning gomologlari atsetilenni alkillab olinadi:



Fizikaviy xossalari:

C_2H_2 efir hidiga ega rangsiz gaz, C_4H_6 dan boshlab suyuqlik, $\text{C}_{12}\text{H}_{22}$ dan boshlab qattiq moddalardir.

Molekula tuzulishi:

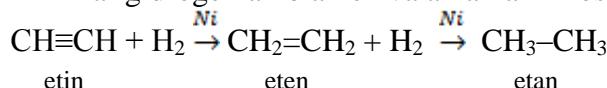


- 1) $\text{sp}^3 - 3 \times 4 = 12$ ta
- 2) $\text{sp} - 2 \times 2 = 4$ ta
- 3) $\text{s} - 8$ ta
- 4) $\text{p}-4$ ta

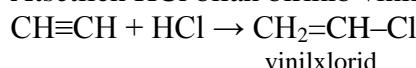
Kimyoviy xossalari:

Alkinlarda 2 ta π bog' bo'lganligi uchun ular birikish reaksiyalariga oson kirishadi.

1) Alkinlar gidrogenlanib alken va alkanlarni hosil qiladi:



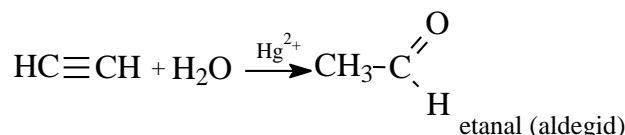
2) Atsetilen HCl bilan birikib vinilxloridni hosil qiladi:



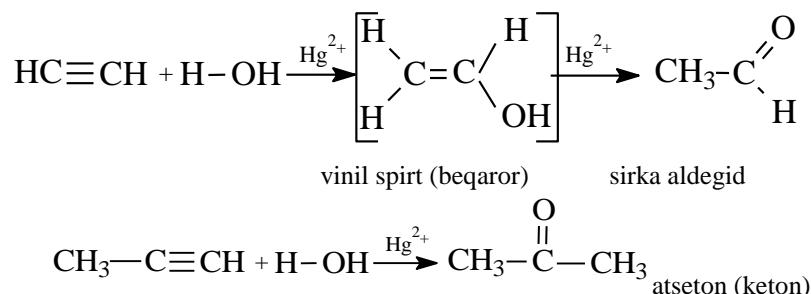
3) $\text{CH}_3-\text{C}\equiv\text{CH} + \text{Br}_2 \rightarrow \text{CH}_3-\text{CBr}=\text{CHBr} \rightarrow \text{CH}_3-\text{CBr}_2-\text{CHBr}_2$

1,2-dibrompropen 1,1,2,2-tetrabrompropan

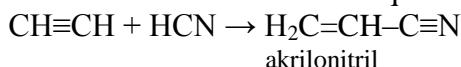
4) Alkinlar HgSO_4 katalizatori ishtirokoda suv bilan birikadi. Bunda atsetilendenan *sirka aldegid*, qolgan vakillaridan *ketonlar* hosil bo'ladi. Bu reaksiyani M.G.Kucherov 1881 yil ochgan (Kucherov reaksiyasi).



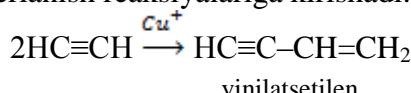
Reaksiya mexanizmi:

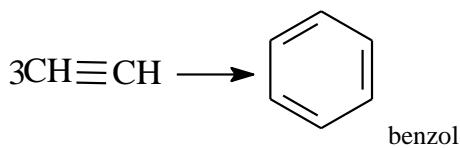


5) Alkinlar vodorod sianid bilan birikib alkilonitrillarni hosil qiladi:

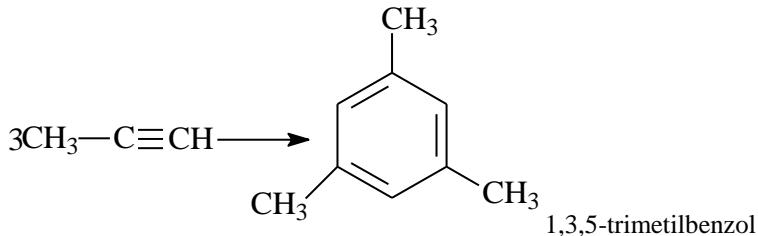


6) Alkinlar dimerlanish va polimerlanish reaksiyalariga kirishadi:

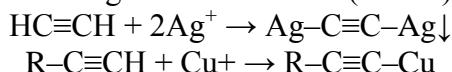




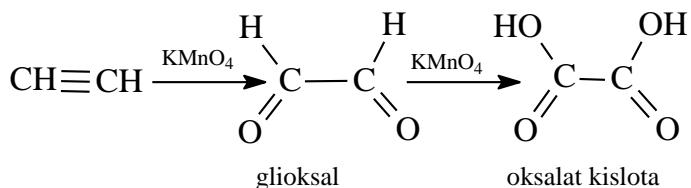
Bu reaksiyaga *Zelinskiy* reaksiyasi deyiladi.



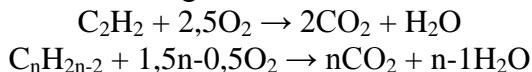
- 7) Alkinlar og'ir metallarning (I) valentli birikmalari bilan atsetilenidlarni hosil qiladi. Bunday reaksiya amalga oshishi uchun uchbog' birinchi o'rinda (chetda) joylashishi kerak:



- 8) Atsetilenning oxirgi oksidlanish mahsuloti oksalat kislota hisoblanadi:



- 9) Alkinlar yonganda suv va karbonat angidrid hosil bo'ladi:



Ishlatilishi:

Atsetilen PVX, poliakrilonitril, xloropren kauchuk, butadiyen kauchuk, sirka aldegid va sirka kislota olishda ishlatiladi. Atsetilen yonganda kuchli issiqlik ajralib chiqqanligi uchun u metallarni payvandlashda va qirqishda foydalaniлади.

Mavzu: Neftni qayta ishlash. Kreking jarayoni.

Neft – turli uglevodorodlar aralashmasidan iborat yoqilg'i sifatida ishlatiladigan foydali qazilma hisoblanadi.

Uning tarkibidagi uglevodorodlar soniga qarab 250-300°C da qaynaydi.

Neftning quyidagi turlari bor:

- 1) Metan neftlari – ko'proq alkanlar tutadi;
- 2) Naften neftlari – ko'proq sikloalkanlar tutadi;
- 3) Aralash neftlar – tarkibida alkan, naften va aromatik uglevodorodlar tutadi.

Shuningdek neft tarkibida suv va yo'ldosh gazlar tutadi.

Neftning o'rtcha zichligi 0,9g/ml atrofida bo'ladi.

Neft fraksion qayta ishlash yo'li bilan qayta ishlanadi. Ya'ni ular qaynash temperaturasiga ko'ra ajratiladi. Bu jarayon *rektifikatsion kolonnada* olib boriladi. Bu kolonnada bug'langan neft tarelkalarda kondensatlanadi.

Neftning asosiy fraksiyalari quyidagilar.

- 1) Benzin;
- 2) Kerosin;

- 3) Gazoyl;
- 4) Mazut.

Benzin fraksiyasi 150°C gacha olib boriladi. Bunda asosan uglerod soni 5-9 gacha bo’lgan yengil uglevodorodlar ajraladi. Bunda 40-70°C da gazolin yoki *petroleyn efiri*, keyin qaynash temperaturasi 70-120°C bo’lgan *benzin* ajraladi. Benzin fraksiyasida ichki yonuv dvigatellarida ishlataladigan *aviatsiya* va *avtomobil benzini* olinadi.

Keyingi fraksiyada 120-140°C da qaynaydigan *ligroin* hosil bo’ladi. Ligroin dizel yoqilg’isi sifatida ishlataladi.

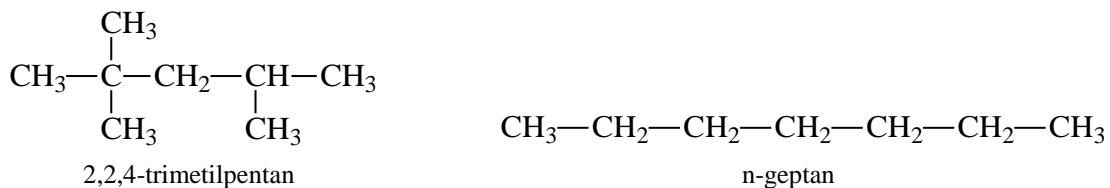
Kerosin fraksiyasida uglerod atomlari soni 9-11 gacha bo’lgan uglevodorodlar bo’ladi. Kerosin traktorlar uchun yoqilg’i va uy-ro’zg’or ihslarida ishlataladi.

Gazoyl fraksiyasida uglerod atomlari soni 11-16 gacha bo’lgan uglevodorodlar bo’ladi.

Mazut fraksiyasida atomlari soni 16 tadan ko’p bo’lgan uglevodorodlar bo’ladi. Mazut(qoramoy) qayta haydaladi. Bundan *solyar moylari* va *surkov moylari* hosil bo’ladi.

Qoramoy haydalangan qolgan qoldiq *gudron* deyiladi va u *asfalt* sifatida ishlataladi.

Benzinning eng muhim xusuiyati – *oktan soni* hisoblanadi. ya’ni izooktan va geptanga nisbatan olinadi.



Geptan qanchalik ko’p bo’lsa, benzin shuncha ko’p *detonatsiyalanadi*. Uning oldini olish uchun antideetonator tetraeilqo’rg’oshin (TEQ) $\text{Pb}(\text{C}_2\text{H}_5)_4$ qo’shiladi. Motor tubiga cho’kkan qo’rg’oshinni tozalash uchun 1,2-dibrometan qo’shiladi va qo’rg’oshin PbBr_4 holida atmosferaga chiqariladi. Bu esa atmosfera havosni ifloslanishiga sabab bo’ladi.

Kreking.

Neftdan olinayotgan benzin barcha iste’molchilar talabini qondirishga yetmaydi. Shuning uchun yuqori uglerod soniga ega uglevodorodlar *piroliz* qilinadi. Bu jarayon *kreking* (*crack - parchalash*) ham deyiladi.

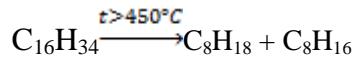
Krekingni quyidagicha ifodalash mumkin:



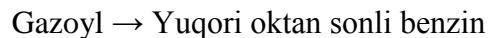
Kreking 2 xil bo’ladi:

- 1) Termik kreking;
- 2) Katalistik kreking.

Termik krekingda temperatura 450°C dan yuqori bo’ladi. Bunda yuqori uglerosli alkanlardan quyi alkan va alken hosil bo’ladi:



Katalistik kreking 450°C dan past temperaturada olib boriladi. Bunda katalizator sifatida AlCl_3 va alyumosilikatlar ishlataladi.



Neftdan ajratib olingan benzinning oktan sonini oshirish uchun *rifroming* jarayonidan foydalaniladi. Riformingda tarmoqlanmagan uglevodorodlar tarmoqlangan uglevodorodlarga aylanadi. Bunda katalizator sifatida Al_2O_3 qo’llaniladi.

Mavzu: Aromatik uglevodorodlar – Arenlar.

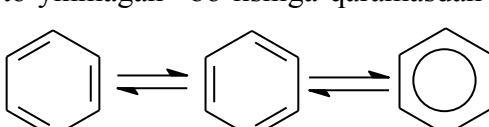
Ta’rif: Tarkibida aromatik xalqa, ya’ni benzol xalqasi tutgan uglevodorodlarga aromatik uglevodorodlar ya’ni arenlar deyiladi.

Umumiyl formulasi: C_nH_{2n-6} . $n \geq 6$.

1825 yil M.Faradey toshko’mirdan benzolni ajratib oladi va unda C va H atomlari 1:1 nisbatda ekanligini aniqlaydi.

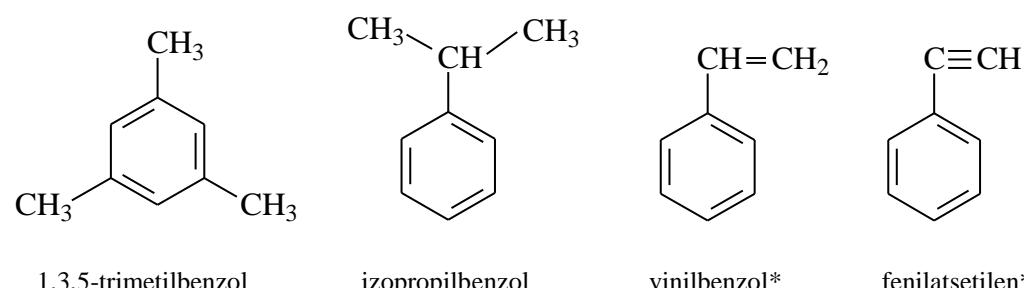
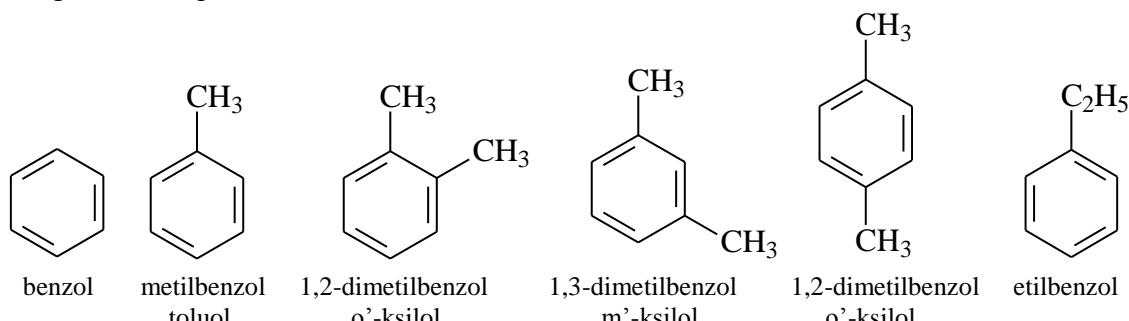
Libix bu moddaga benzol nomini beradi.

Benzol o’ziga xos hidga ega edi va u juda qiziqarli xossalarni namoyon qiladi. U juda “to’ymagan” bo’lishiga qaramasdan birikish reaksiyalariga juda qiyin kirishadi va o’rin olish olish reaksiyalariga oson kirishadi. Shuning uchun benzol va hosilalari *aromatik uglevodorodlar* deb nomlanadi.

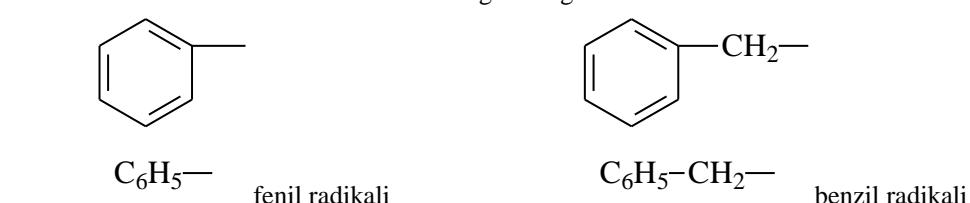
 Benzolning tuzilishini 1865 yil nemis olimi A.Kekule taklif qilgan.

Nomlanishi va izomeriyasi:

Benzol va uning hosilalari benzol yadrosiga nisbatan nomlanadi. Bunda avval kichik radikal, so’ngra kattaroq radikal nomlanadi.



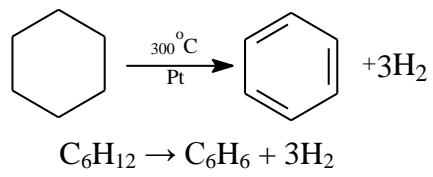
*vinilbenzol va fenilsetilen benzol gomologgi emas.



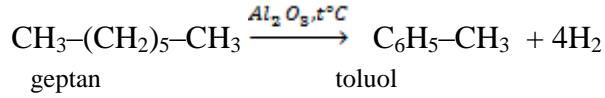
Olinishi:

Dastlab arenlar toshko’mirni quruq haydab olingan. Benzol va fenol neft tarkibida ham uchraydi.

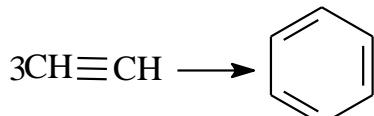
- Arenlar sikloalkanlarni degidrogenlab olinadi:



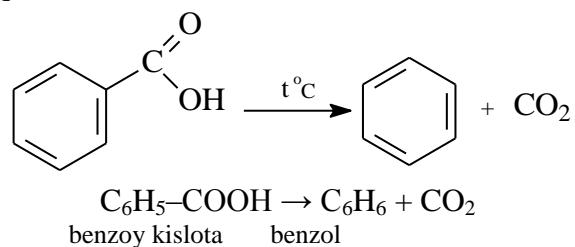
2) Alkanlarni degidrogenlab olinadi:



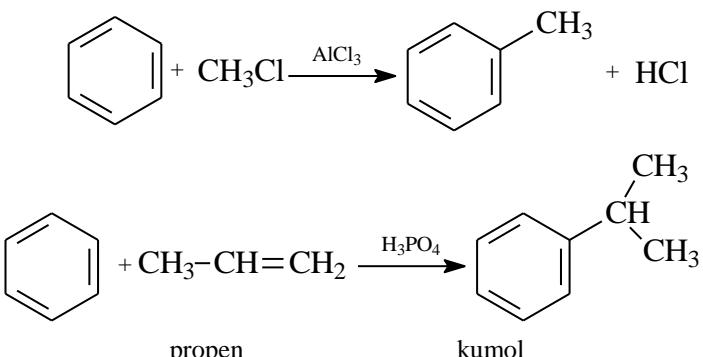
3) Atsetilenni trimerlab olinadi (Zelinskiy reaksiyasi):



4) Aromatik kislotalarni qizdirib olinadi (dekarboksillash):



5) Benzol gomologlari Fridel-Krafts bo'yicha alkillab olinadi:

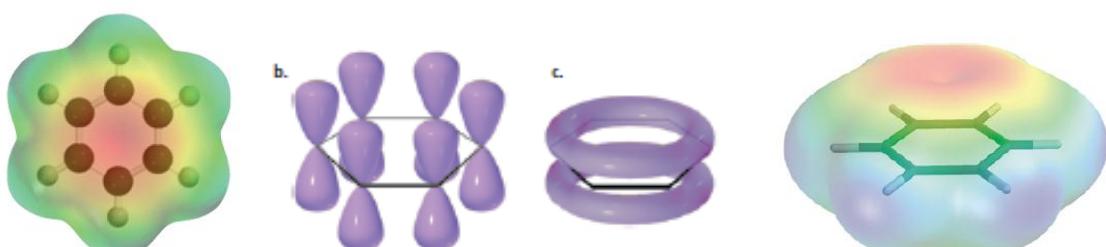


Fizikaviy xossalari:

Benzol va gomologlari rangsiz suyuqliklar hisoblanadi. Yuqori vakillari kristall moddalardir.

Molekula tuzilishi:

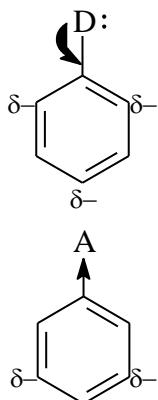
Benzol yadrosidagi barcha uglerod atomlari sp^2 gibridlangan holda bo'ladi.



1) $sp^2 - 6 \times 3 = 18$ ta

2) $p - 6$ ta

3) $s - 6$ ta



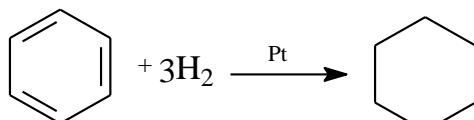
Benzol yadrosiga 2 xil o'rribosarlar ta'sir qiladi.

- 1) I tur o'rribosarlar – *elektrodonor* guruuhlar. σ' - va p' - holat elektron bulutini zichligini kuchaytiragi. Ularga $-\text{CH}_3$ (Alk), $-\text{CH}=\text{CH}_2$, $-\text{OH}$, $-\text{NH}_2$, $-\text{F}$, $-\text{Cl}$, $-\text{Br}$ va $-\text{J}$ kiradi.
- 2) II tur o'rribosarlar – *elektroakseptor* guruuhlar. Ular m' - holatga yo'naltiradi. Ularga $-\text{NO}_2$, $-\text{CHO}$, $-\text{COOH}$, $-\text{CN}$ va $-\text{SO}_3\text{H}$ kiradi.

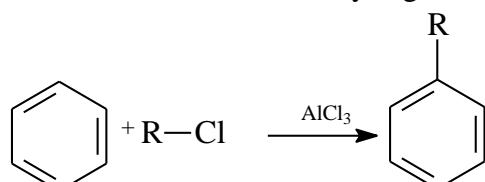
Kimyoviy xossalari:

Benzol va uning gomologlari uchun *o'rin olish* reaksiyasi xarakterli. Chunki aromatik xalqa barqarorlikni ta'minlaydi.

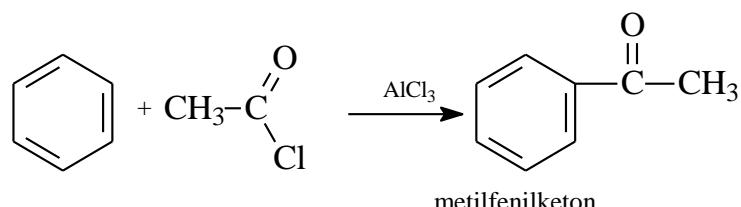
- 1) Benzol gidrogenlanganda siklogeksan hosil bo'ladi:



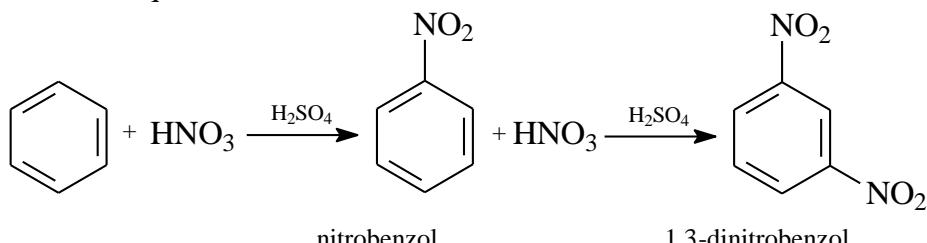
- 2) Benzol galogenoalkanlar bilan alkillanish reaksiyasiga kirishadi:



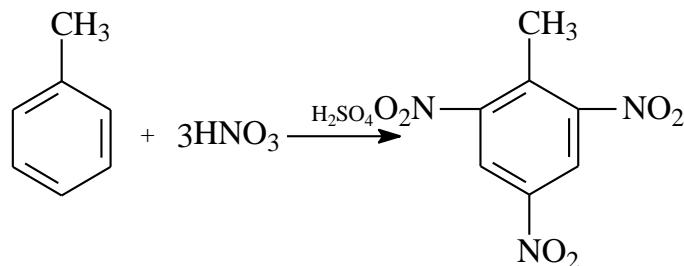
- 3) Benzol Fridel-Krafts katalizatori ishtirokida xlorangidridlar bilan atsillanib arilketonlarni hosil qiladi:



- 4) Benzol va uning gomologlari nitrolovchi aralashma (HNO_3 + H_2SO_4) ta'sirida nitrobirikmlarni hosil qiladi:

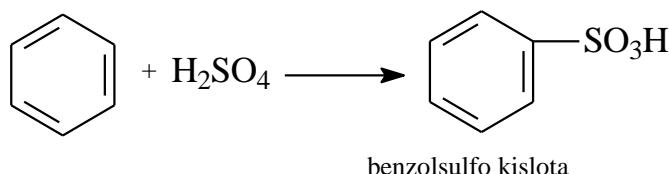


- 5) Toluol nitrolanganda trinitrotoluol (TNT) hosil bo'ladi. TNT kuchli portlovchi modda:



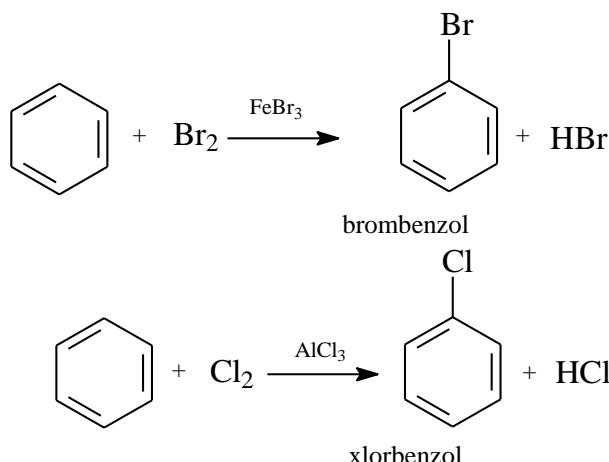
2,4,6-trinitrotoluol – TNT

- 6) Benzol kons. H_2SO_4 bilan qizdirilganda reaksiyaga kirishadi:

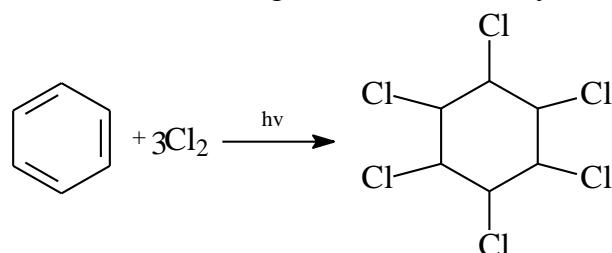


- 7) Benzol va uning gomologlari galogenlanish reaksiyasiga oson kirishadi.

a) Reaksiya FeBr_3 yoki AlCl_3 ishtirokida borganda o'r'in olish reaksiyasi sodir bo'ladi:

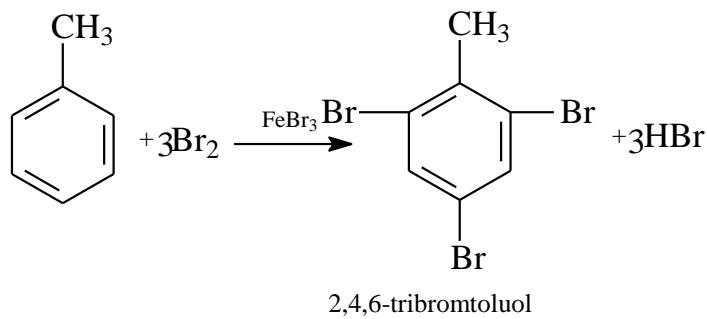


b) Benzol ultrabinafsha nurlar ta'sirida xlorlanganda birikish reaksiyasi sodir bo'ladi:

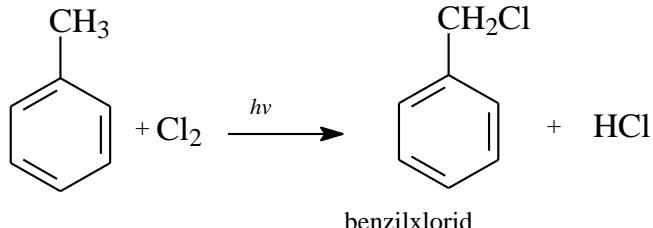


geksaxlorsiklogeksan, geksaxloran

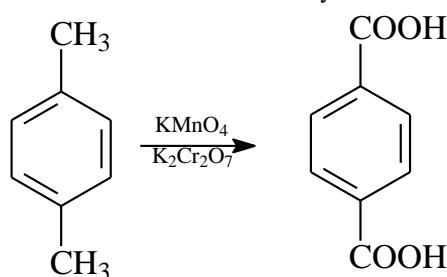
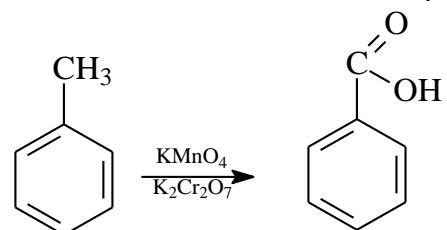
c) Toluol katalizator ishtirokida galogenlanganda trigalogenli hosila hosil bo'ladi:



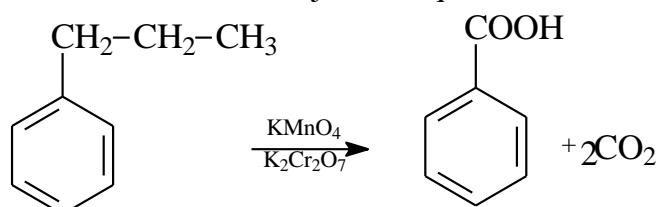
d) Toluol quyosh nurlari ishtirokida xlorlanganda xlor yon zanjirga birikadi:



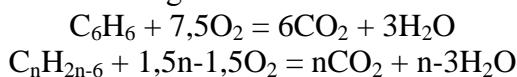
- 8) Benzol hosilalari oksidlanganda xalqadagi o'rindbosarlar soniga ko'ra aromatik mono- va dikarbon kislotalar hosil bo'ladi. Oksidlovchi sifatida KMnO_4 yoki $\text{K}_2\text{Cr}_2\text{O}_7$ qo'llaniladi:



Shunisi muhimki, bitta o'rindbosarli hosilada uglerod sonidan qat'iy nazar benzoy kislota hosil bo'ladi. Qolgan uglerodlar CO_2 ko'rinishida ajralib chiqadi:



- 9) Arenlar yonganda suv va karbonat angidrid hosil bo'ladi:



Ishlatilishi:

Benzol erituvchi sifatida ishlatiladi. Unda yog', kauchuk va laklar yaxshi eriydi. Kumol atseton va fenol olishda oraliq mahsulot hisoblanadi. Stirol polistirol kauchuk olishda ishlatiladi. Geksaxloran insektitsid sifatida ishlatiladi.

III BOB. KISLOROD SAQLOVCHI ORGANIK BIRIKMALAR.

Mavzu: Spirtlar.

Ta’rif: Tarkibida uglevodorod radikali bilan bog’langan gidroksil guruh $-OH$ tutgan organik birikmalar sinfiga spirtlar deyiladi.

Spirtlar radikal tabiatiga ko’ra to’yingan va to’yinmagan spirtlarga bo’linadi. Gidroksil guruh soniga ko’ra esa bir va ko’p atomli spirtlarga bo’linadi.

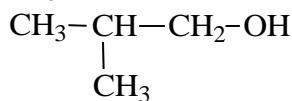
To’yingan bir atomli spirtlarning umumiy formulasi $C_nH_{2n+1}OH$.

Nomlanishi va izomeriyasi:

Spirtlarni nomlash uchun gidroksil guruh yaqin tomondan raqamlanib, tegishli alkan nomiga *-ol* qo’shimchasi qo’shiladi.

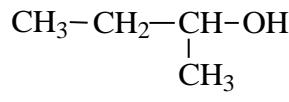
CH_3OH methanol, metil spirt

$CH_3-CH_2-CH_2-CH_2-OH$ butanol-1, butil spirt



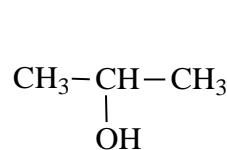
C_2H_5OH etanol, etil spirt

2-metilpropanol-1, izopropil spirt

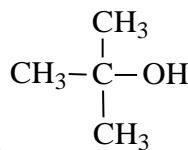


$CH_3-CH_2-CH_2-OH$ propanol-1, propil spirt

butanol-2, ikkilamchi butil spirt



propanol-2, izopropil spirt

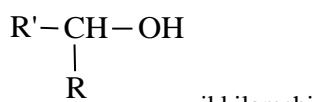


2-metilpropanol-2, uchlamchi butil spirt

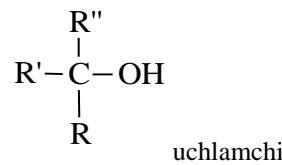
$C_5H_{11}-OH$ pentanol, amil spirt

Gidroksil guruh bilan bog’langan uglerod xarakteriga ko’ra birlamchi, ikkilamchi va uchlamchi spirtlarga bo’linadi:

$R-CH_2-OH$ birlamchi



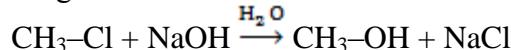
ikkilamchi



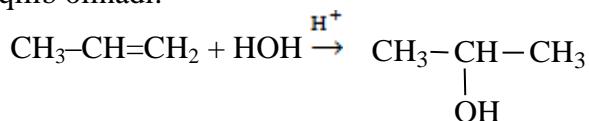
uchlamchi

Olinishi:

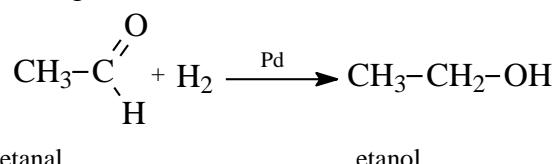
1) Galogenalkanlarga ishqorning suvli eritmasi ta’sir ettirib:

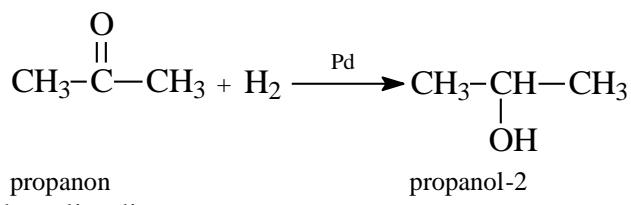


2) Alkenlarni gidroliz qilib olinadi:

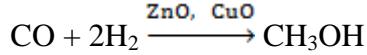


3) Aldegid va ketonlarni gidrogenlab olinadi. Aldegidlar gidrogenlanganda birlamchi, ketonlar gidrogenlanganda ikkilamchi spirtlar hosil bo’ladi:





- 4) Metanol sintez gazidan olinadi:



- 5) Etanol glyukozaning spirtli bijg'ishidan hosil qilinadi:

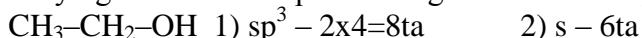


Fizikaviy xossalari:

Spirtlarning quyi vakillari rangsiz suyuqliklar, yuqori vakillari kristall moddalardir. Ular o'zaro va suv molekulalari bilan molekulalararo vodorod bog'lanish hosil qiladi.

Molekula tuzilishi:

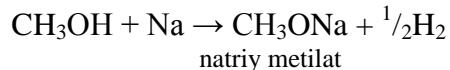
To'yingan bir atomli spirtlarda uglerod atomlari sp^3 gibridlangan holda bo'ladi:



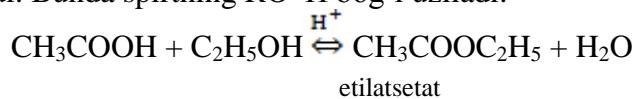
Kimyoviy xossalari:

Alkanollar uchun gidrosil guruh $-\text{O}-\text{H}$ va radikal bo'yicha boradigan reaksiyalar farq qiladi.

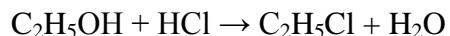
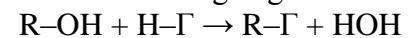
- 1) Alkanollar juda kuchsiz kislota xossasini namoyon qiladi. ular ishqoriy metallar bilan alkogolyatlar hosil qiladi:



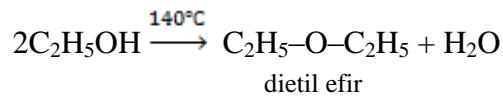
- 2) Spirtlar karbon kislotalar bilan ta'sirlashib murakkab efirlarni hosil qiladi. reaksiya kislotali muhitda amalga oshadi. Bunda spirtning $\text{RO}-\text{H}$ bog'i uziladi:



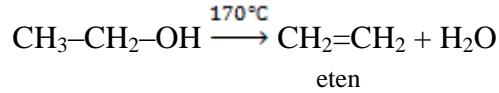
- 3) Spirtlarga galogenovodrodlar ta'siridan alkilgalogenidlar hosil bo'ladi:



- 4) Spirtlar sulfat kislota ishtirokida 140°C da qizdirilganda molekulalararo degidratlanib oddiy efirlar hosil bo'ladi:

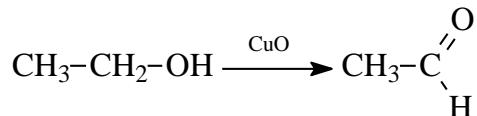


- 5) Spirtlar sulfat kislota ishtirokida 170°C da qizdirilganda ichki molekulyar degidratlanib alkenlar hosil bo'ladi:

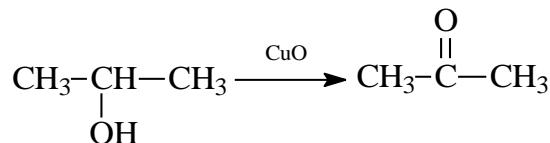


- 6) Spirtlar CuO yoki ZnO ishtirokida oksidlanganda karbonil birikmalar hosil bo'ladi.

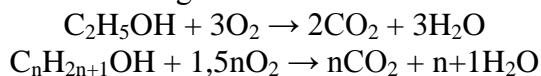
- a) Birlamchi spirtlar oksidlanganda aldegidlar hosil bo'ladi:



- b) Ikkilamchi spirtlardan esa ketonlar hosil bo'ladi:



7) Spirtlar yonganda suv va karbonat angidrid hosil bo'ladi:



Ishlatilishi:

Methanol asosan erituvchi sifatida va formaldegid olish uchun ishlatiladi. U juda kuchli zahar (5ml ko'r qiladi, 30ml o'limga olib keladi).

Etanol oziq-ovqat sanoatida, erituvchi sifatida, Lebedev usulida butadiyen kauchuk olishda, sirka aldegid va etilatsetat olishda ishlatiladi.

Propanol va butanol lak va kraskalarni eritishda ishlatiladi.

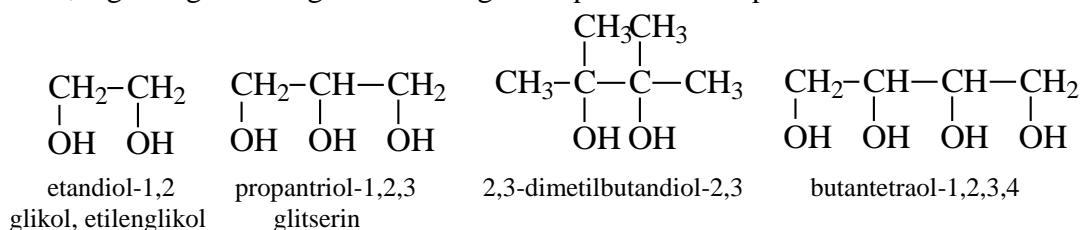
Mavzu: Ko'p atomli spirtlar.

Ta'rif: Tarkibida ikki yoki undan ortiq gidroksil guruh $-OH$ tutgan organik moddalarga ko'p atomli spirtlar deyiladi.

Umumiyl formulasi: R(OH)_n $n= 2, 3, 4\dots$

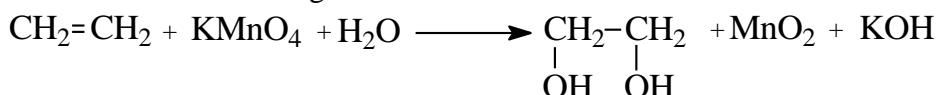
Nomlanishi va izomeriyasi:

Ko'p atomli spirlarni nomlash uchun gidroksil guruhlar tutgan eng uzun uglevodorod zanjiri tanlanib, tegishli gidroksil guruhlar soniga $-ol$ qo'shimchasi qo'shiladi.

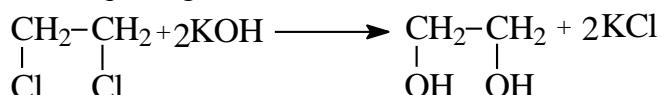


Olinishi:

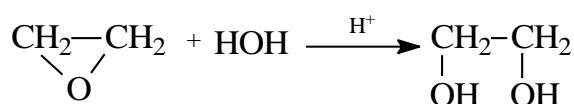
1) Etilenglikol etilenni KMnO_4 ning suvli eritmasi bilan oksidlab olinadi:



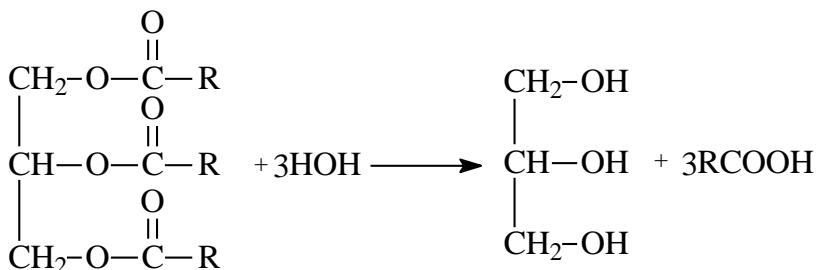
2) Glikol 1,2-dixloretanni ishqorning suvli eritmasi bilan ta'siridan olinadi:



3) Etilenoksidni kislotali muhitda suv bilan ta'siridan olinadi:



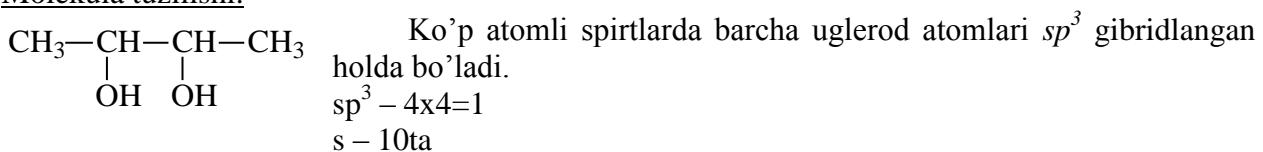
4) Glitserin yog'larni gidroliz qilib olinadi:



Fizikaviy xossalari:

Etilenglikol rangsiz qovushqoq suyuqlik. Suv bilan har qanday nisbatda aralashadi. Zaharli. Glitserin qovushqoq shirin ta'mli rangsiz suyuqlik. Suv bilan har qnday nisbatda joylashadi.

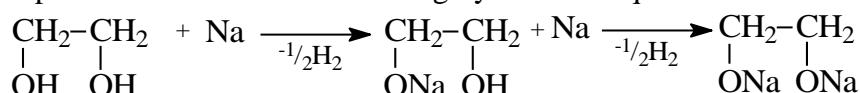
Molekula tuzilishi:



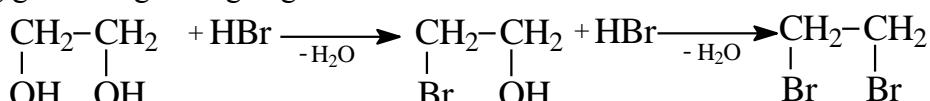
Kimyoviy xossalari:

Ko'p atomli spirtlar bir atomli spirtlar kabi xossalarga ega.

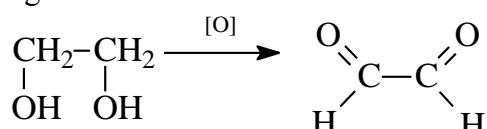
- 1) Ko'p atomli spirtlar aktiv metallar bilan alkogolyatlar hosil qiladi:



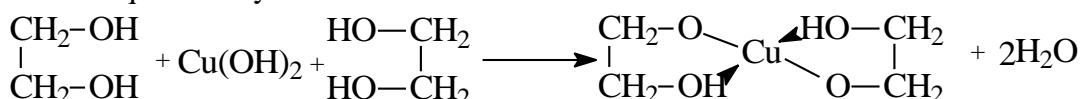
- 2) Ularning hidroksil guruhi galogenovodorodlar ta'sirida almashinadi:



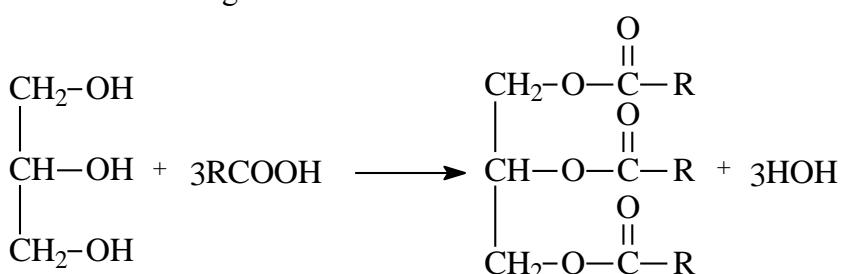
- 3) Etilenglikol oksidlanganda glioksal hosil bo'ladi:



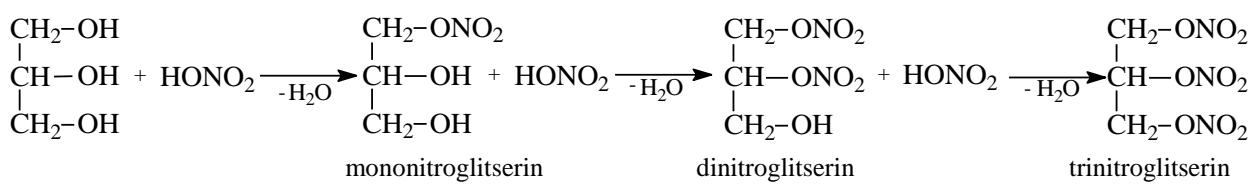
- 4) Glikollar yangi tayyorlangan $\text{Cu}(\text{OH})_2$ bilan kompleks tuz hosil qiladi. bu reaksiyadan glikollarni aniqlashda foydalaniladi.



- 5) Glitserin anorganik va organik kislotalar bilan murakkab efirlarni hosil qiladi. bunda mono, di- va tri- almashgan efirlar hosil bo'ladi:

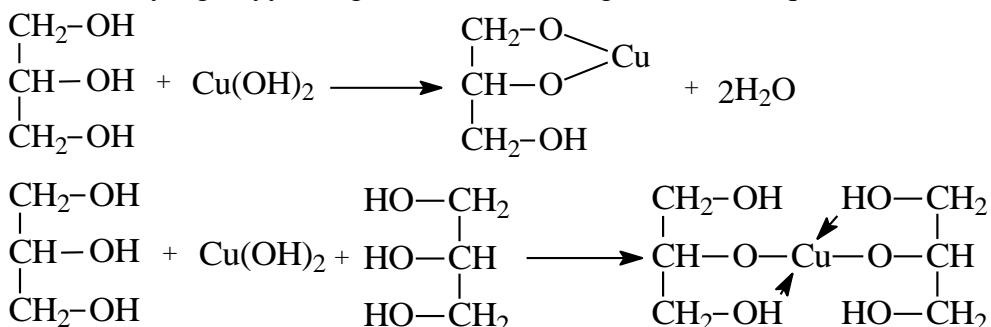


- 6) Glitserin nitrolanganda nitroefirlar hosil bo'ladi:



Trinitroglycerin kuchli portlovchi modda. Undan dinamit tayyorlanadi (A.Nobel).

7) Glitserin yangi tayyorlangan $\text{Cu}(\text{OH})_2$ bilan glitserat hosil qiladi:



Ishlatilishi:

Etilenglikol antifriz, tormoz suyuqligi, poliefir tola (lavsan) olishda ishlatiladi. Glitserinni 1779 yili Sheele olgan. U dinamit tayyorlashda, poliefirlar olishda, kosmetika sanoati va meditsinada ishlatiladi.

Mavzu: Oddiy efirlar.

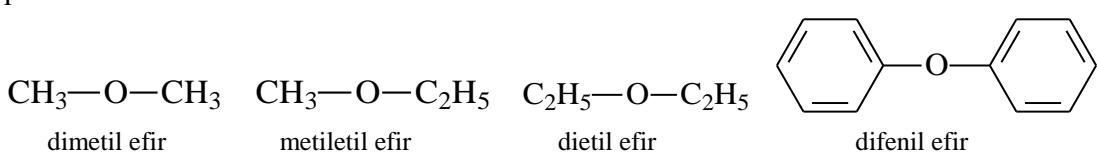
Ta’rif: Ikkita uglevodorod radikali kislorod ko’prigi (-O-) orqali bog’langan organik moddalar sinfiga oddiy efirlar deyiladi.

Umumiyl formulasi: R-O-R'

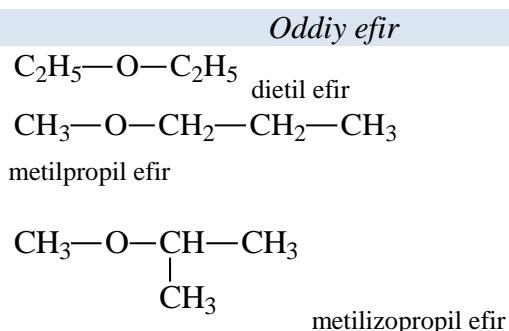
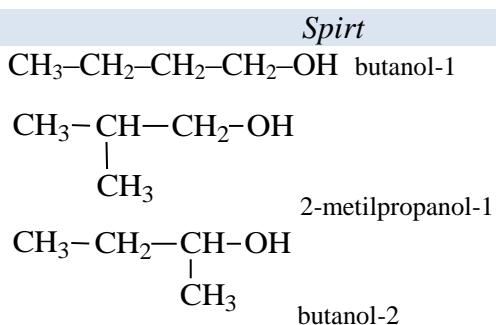
R – radikal to’yingan, to’yinmagan yoki aromatik bo’lishi mumkin.

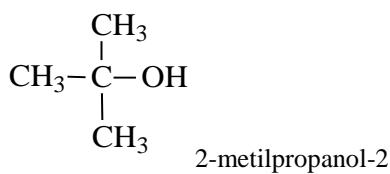
Nomlanishi va izomeriyasi:

Oddiy efirlarni nomlash uchun avval kichik radikal, so’ngra kattaroq radikal nomi aytilib, *efir* so’zi qo’shiladi.



To’yingan efirlar to’yingan bir atomli spirtlarga izomer hisoblanadi. Masalan, $\text{C}_4\text{H}_{10}\text{O}$ formulaga quyidagi spirt va oddiy efirlar mos keladi:



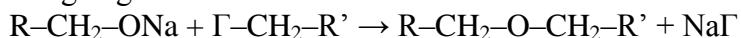


Olinishi:

- 1) Spirtlarni pastroq temperaturada ichki molekulayar degidratlab olinadi:



- 2) Alkogolyatlarga alkilgalogenidlari ta'sir ettirib olinadi:



Fizikaviy xossalari:

Oddiy efirlar oz'iga xos hidli, rangsiz suyuqliklar hisoblanadi. Ularning qaynash temperaturasi tegishli spirtlarnikiga qaraganda past. Chunki ularda vodorod bog'lanish yo'q.

Kimyoviy xossalari:

Oddiy efirlar asosan donor-akseptor bog'lanish hisobiga moddalar hosil qiladi.

Ishlatilishi:

Dietil efir metalorganik birikmalarni eritishda erituvchi sifatida va meditsinada umumiy narkoz sifatida ishlatiladi.

Mavzu: Fenollar.

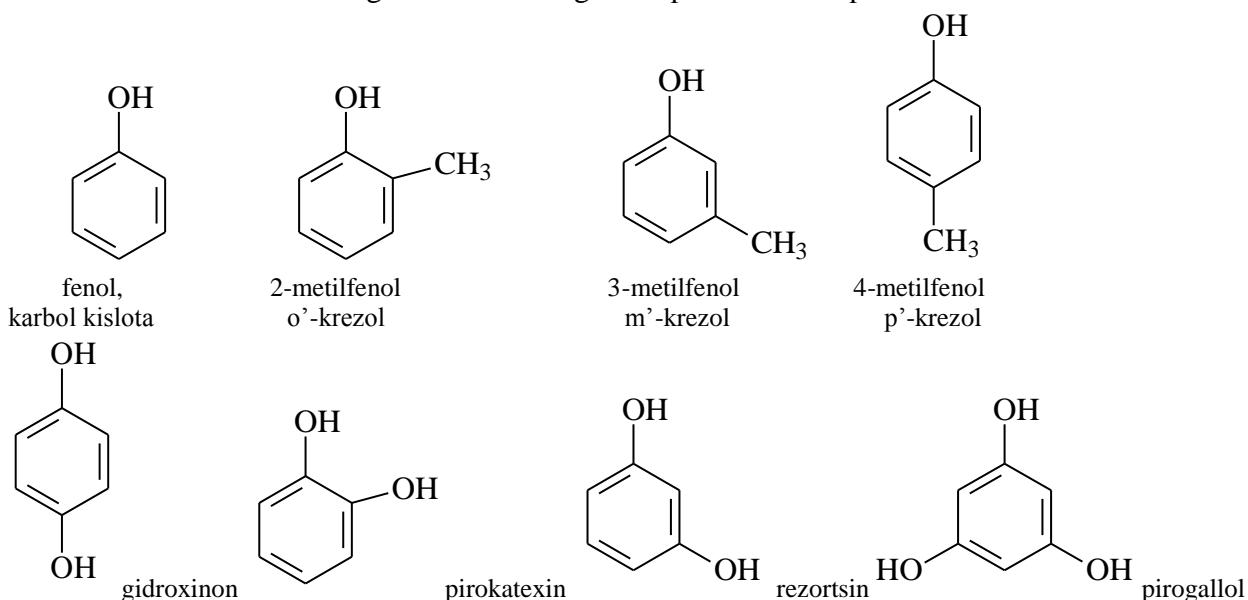
Ta'rif: Benzol yadrosi bilan bevosita bog'langan gidroksil guruh $-\text{O}-\text{H}$ ga ega organik moddalar sinfiga fenollar deyiladi.

Ularning umumiy formulasi: $\text{Ar}-\text{OH}$.

Benzol yadrosi uglerodlari sp^2 gibriddlangan.

Nomlanishi va izomeriyasi:

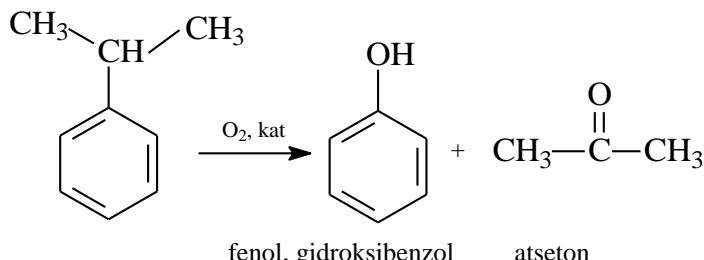
Fenollarni nomlash uchun tegishli aren nomiga $-ol$ qo'shimchasi qo'shiladi.



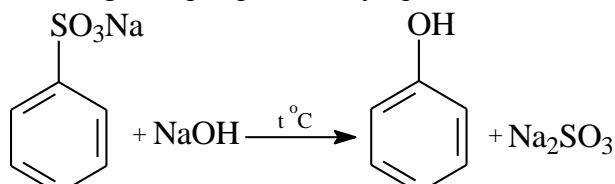
Olinishi:

Fenol va uning gomologlari toshko'mir smolasidan, shuningdek sintez usulida olinadi.

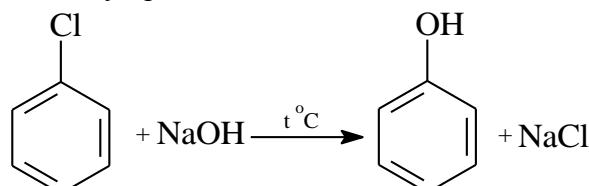
- 1) Fenol kumolni katalitik oksidlab olinadi:



- 2) Aromatik sulfokislota tuzlariga ishqor qo'shib suyuqlantirib olinadi:



- 3) Fenol xlorbenzolga 350°C da yuqori bosimda NaOH eirtmasi ta'sir ettirib olinadi:

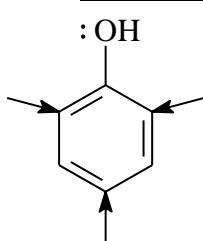


Fizikaviy xossalari:

Ko'pchilik fenollar rangsiz moddalar hisoblanadi. Ular o'ziga xos hidga ega.

Molekula tuzilishi:

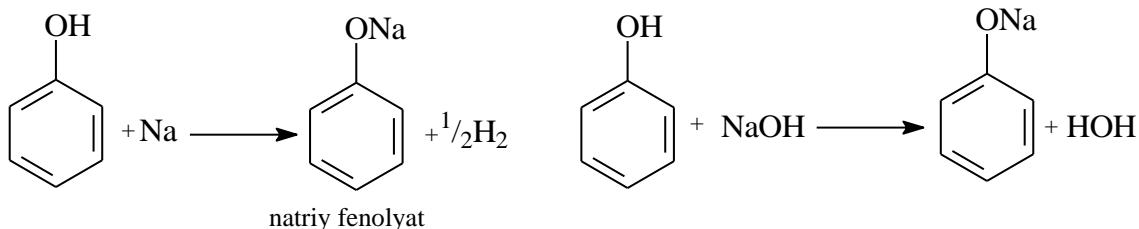
Fenol molekulasida benzol yadrosi uglerodlari sp^2 gibridlangan holda bo'ladi. OH guruh I tur o'rinxbosar (elektrodonor) hisoblanadi. Shuning uchun o'- va p'-holatlari aktiv bo'ladi.



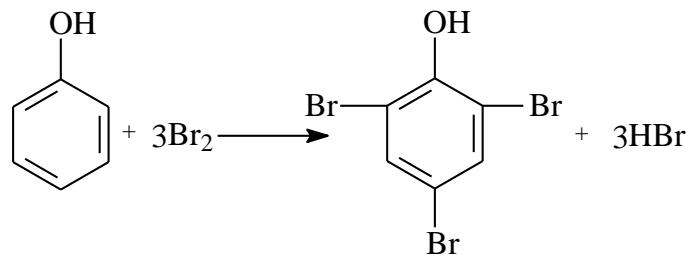
Kimyoiy xossalari:

Fenollar uchun benzol yadrosiga va gidroksil guruh bo'yicha amalga oshadigan reaksiyalar farqlanadi.

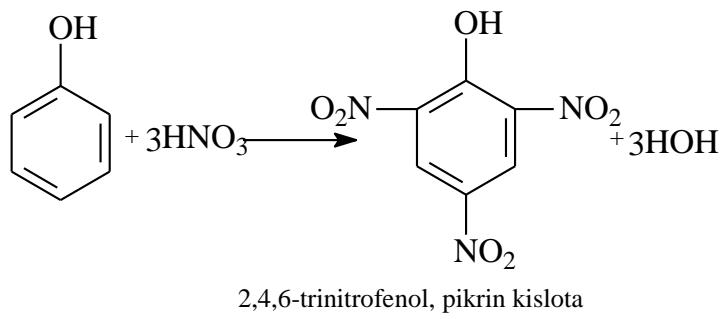
- 1) Fenollar spirlarga qaraganda kuchliroq kislotalik xossasini namoyon qiladi. Shuning uchun ular ishqoriy metallar va ishqorlar bilan ta'sirlashib fenolyatlarni hosil qiladi:



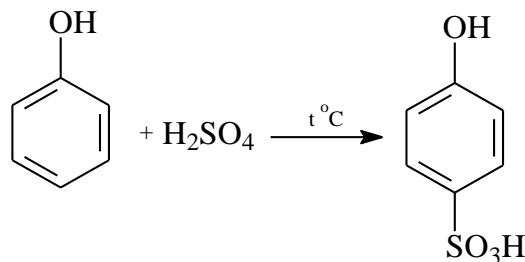
- 2) Fenollar osonlik bilan galogenlanadi:



- 3) Fenol kons. nitrat kislota bilan ta'sirlashib trinitrofenol – kuchli portlovchi moddani hosil qiladi:

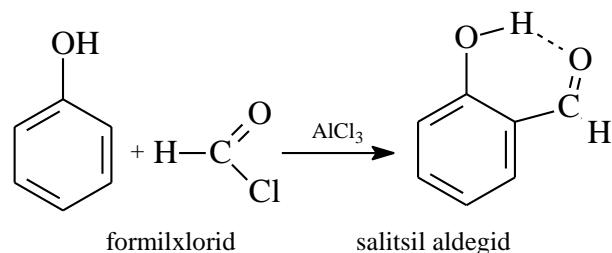


- 4) Fenol kons. sulfat kislota bilan qizdirilganda reaksiyaga kirishadi:

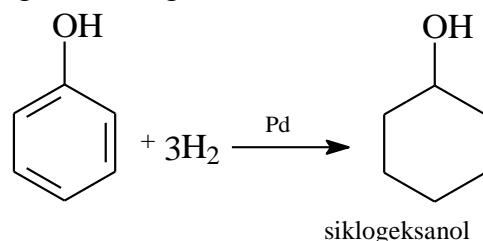


p'-fenolsulfokislota

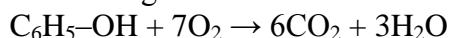
- 5) Fenol Fridel-Krafts katalizatori ishtirokida osonlik bilan atsillanadi:



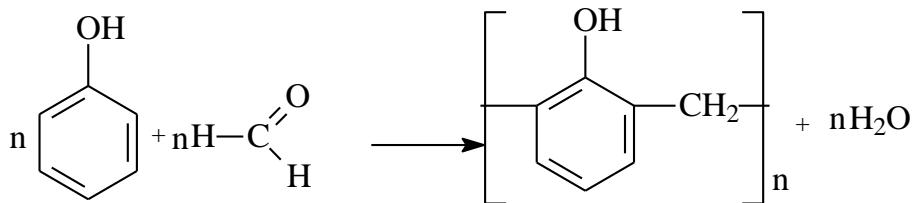
- 6) Fenol katalitik gidrogenlanganda siklogeksanol hosil bo'ladi:



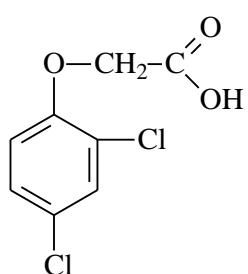
- 7) Fenollar yonganda suv va karbonat angidrid hosil bo'ladi:



- 8) Fenol metanal bilan sopolimerlanib fenolformaldegid smolasini hosil qiladi:



Ishlatilishi:



Fenolning ko'p qismi fenolformaldegid smolasasi olsih uchun ishlatiladi.

Fenol shuningdek antisептик сифатида ham ishlatiladi. U terini ya'llig'laydi.

Krezollar bo'yоqlar сифатида ishlatiladi. Salitsil kislotasi farmasevtikada qo'llaniladi. 2,4-dixlorfenoksisirka kislotasi va uning natriyli tuzi gerbitsid сифатида ishlatiladi.

Sifat reaksiya:

Eritmada fenol borligi FeCl_3 eritmasi bilan aniqlanadi. Bunda binafsha rangli kompleks hosil bo'ladi.

Mavzu: Karbonil birikmalar.

Ta'rif: Tarkibida karbonil guruh $\text{C}=\text{O}$ tutgan birikmlarga karbonil birikmalar deyiladi.

Karbonil birikmalar 2 ga bo'linadi:

- 1) Aldegidlar
- 2) Ketonlar.

Mavzu: Aldegidlar.

Ta'rif: Tarkibida aldegid $\text{C}=\text{O}$ guruh tutgan organik moddalarga aldegidlar deyiladi.

Umumiy formulasi: $\text{R}-\text{C}(\text{H})=\text{O}$

Nomlanishi va izomeriyasi:

Aldegidlarni nomlash uchun karbonil guruh tutgan eng uzun uglevodorod zanjiri tanlanib, aldegid guruh uglerodi 1 raqamli deb olinadi. Tegishli uglevodorod nomiga *-al* qo'shimchasi qo'shiladi.

$\text{H}-\text{C}(\text{H})=\text{O}$
metanal, formaldegid
chumoli aldegid

$\text{CH}_3-\text{C}(\text{H})=\text{O}$
etanal
sirka aldegid

$\text{CH}_3-\text{CH}_2-\text{C}(\text{H})=\text{O}$
propanal
propion aldegid

$\text{CH}_3-\text{CH}_2-\text{CH}_2-\text{C}(\text{H})=\text{O}$
butanal
moy aldegid

$\text{CH}_3-\text{CH}(\text{CH}_3)-\text{C}(\text{H})=\text{O}$
2-metylpropanal
izomoy aldegid

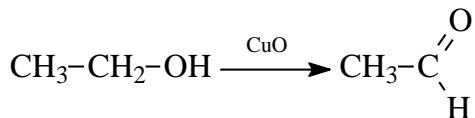
$\text{CH}_2=\text{CH}-\text{C}(\text{H})=\text{O}$
propenal
akrolein

$\text{CH}_3-\text{CH}=\text{CH}-\text{C}(\text{H})=\text{O}$
buten-2-al
kroton aldegid

$\text{C}_6\text{H}_5-\text{C}(\text{H})=\text{O}$
benzaldegid

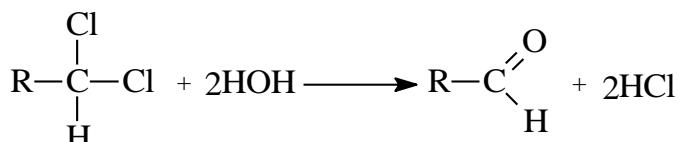
Olinishi:

- 1) Birlamchi spirtlarni oksidlab olinadi:

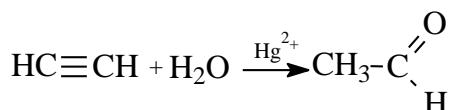


- 2) Formaldegid metanni 450°C da AlPO_4 yoki NO_x katalizatorlari ishtirokida oksidlab olinadi:
- $$\text{CH}_4 \xrightarrow{[\text{O}]} \text{HCOH} + \text{H}_2\text{O}$$

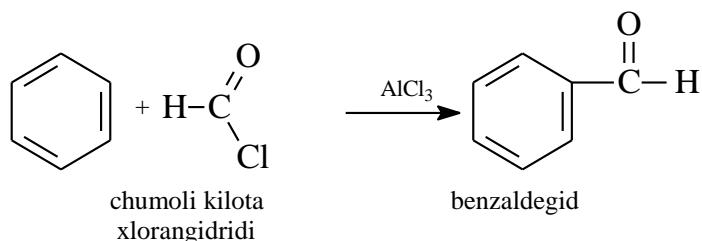
- 3) Digalogenalkanlarni gidroliz qilib olinadi:



- 4) Atsetilenni Kucherov reaksiyasi asosida hidratlab olinadi:



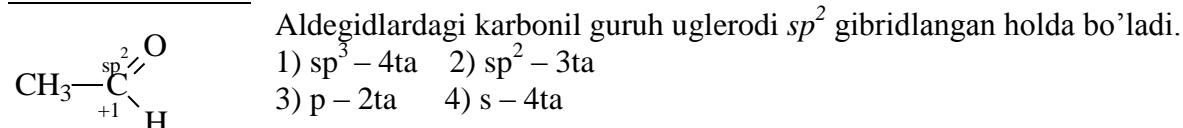
- 5) Benzaldegid benzolga AlCl_3 katalizatori ishtirokida formilxlorid ta'siridan olinadi:



Fizikaviy xossalari:

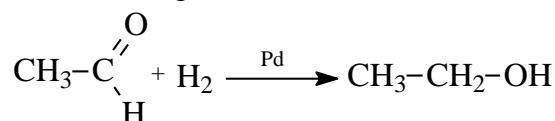
To'yingan aldegidlar rangsiz o'ziga xos hidli suyuqliklardir. Formaldegid o'tkir hisli gaz.

Molekula tuzilishi:

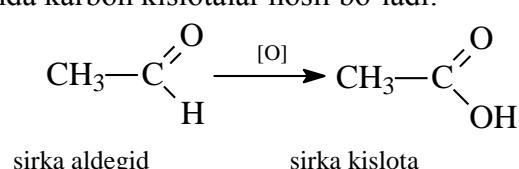


Kimyoviy xossalari:

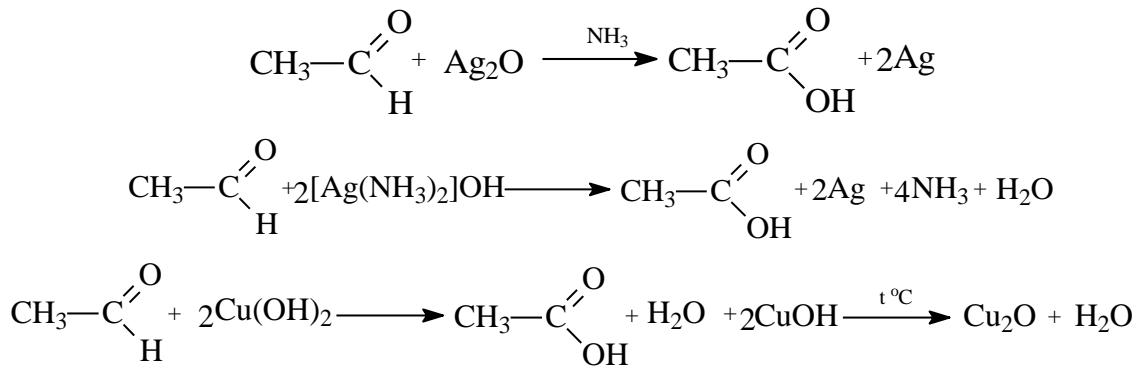
- 1) Aldegidlar qaytarilganda birlamchi spirtlar hosil bo'ladi:



- 2) Aldegidlar oksidlanganda karbon kislotalar hosil bo'ladi:



- 3) Aldegidlar uchun sifat reaksiya bu – ularning Ag₂O ning ammiakdagi eritmasi yoki yangi tayyorlangan Cu(OH)₂ bilan oksidlanish reaksiyasidir. Birinchi reaksiyaga “kumush ko’zgu” reaksiyasi deyiladi:



- 4) Formaldegid polimerlanish reaksiyasiga kirishganda poliformaldegid hosil bo’ladi:
 $n\text{CH}_2\text{O} \rightarrow (-\text{CH}_2-\text{O}-)_n$

Ishlatilsishi:

Formaldegid fenolformaldegid smolasi olishda, uning 35-40% li suvli eritmasi – formalin biopreparatlarni saqlashda ishlatiladi.

Atsetalsegid sirkal kislota va sirkal angidridi olishda ishlatiladi.

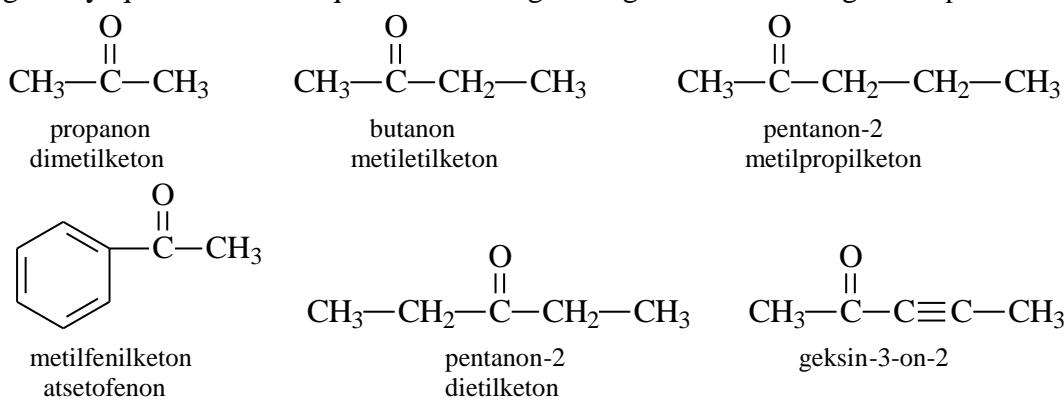
Akrolein atseton va glitserin olishda ishlatiladi.

Mavzu: Ketonlar.

Ta’rif: Tarkibi R-C(=O)-R' bo’lgan organik moddalarga ketonlar deyiladi.

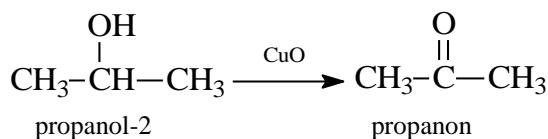
Nomlanishi va izomeriyasi:

Ketonlarni nomlash uchun karbonil guruh tutgan eng uzun uglevodorod zanjiri tanlanib, karbonil guruh yaqin tomonidan raqamlanadi va tegishli uglevodorod nomiga –on qo’shimchasi qo’shiladi.

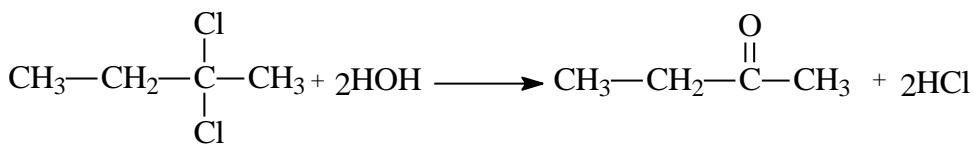


Olinishi:

- 1) Ikkilamchi spirlarni oksidlاب olinadi:



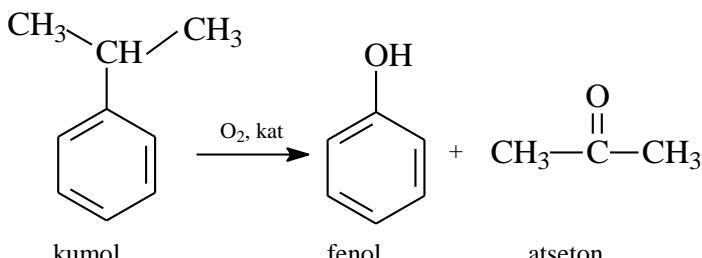
- 2) Digalogenalkanlarni gidroliz qilib olinadi:



2,2-dixlorbutan

butanon

3) Atseton kumolni oksdilab olinadi:

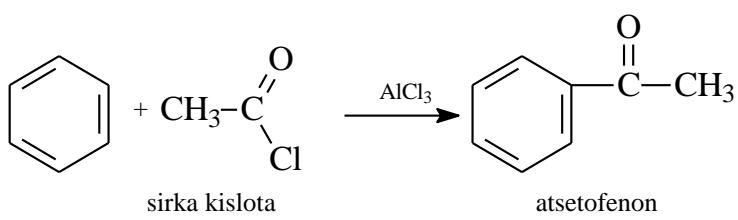


kumol

fenol

atseton

4) Aromatik ketonlar benzolni atsillab olinadi:



sirka kislota
xlorangidridi

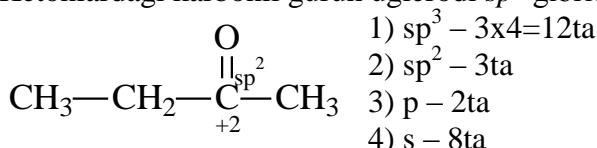
atsetofenon

Fizikaviy xossalari:

Atseton yoqimli hidli rangsiz suyuqlik.

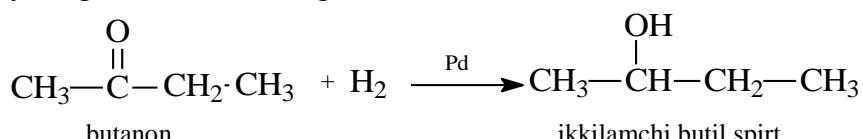
Molekula tuzilishi:

Ketonlardagi karbonil guruh uglerodi sp^2 gibridlangan holda bo'ladi.



Kimyoviy xossalari:

1) Ketonlar qaytarilganda ikkilamchi spirtlar hosil bo'ladi:



2) Ketonlar oksidlanishga chidamli. Ular "kumush ko'zgu" reaksiyasiga kirishmaydi.

Ishlatilishi:

Atseton nitrosellyuloza va boshqa moddalarni eritishda erituvchi sifatida ishlatiladi. Shuningdek u kinoplenka va portlovchi moddalar olishda ishlatiladi.

Mavzu: Karbon kislotalar.

Ta'rif: Tarkibida uglevodorod radikali bilan bog'langan karboksil guruh $\text{---C}(\text{O})\text{OH}$ tutgan organik moddalar sinfiga karbon kislotalar deyiladi.

Ularning umumiyl formulasi: $\text{R}-\overset{\substack{\text{O} \\ || \\ \text{sp}^2}}{\text{C}}-\text{OH}$

Karbon kislotalar quyidagicha bo'linadi:

- 1) Monokarbon kislotalar (to'yangan, to'yinmagan, aromatik);
- 2) Dikarbon kislotalar (to'yangan, to'yinmagan, aromatik);
- 3) Karbon kislotalarning hosilalalri (galogen, gidroksi, aminokarbon kisloatalar).

Nomlanishi va izomeriyasi:

Karbon kislotalarni nomlash uchun karboksil guruh uglerodi 1 deb olinib, tegishli uglevodorod nomiga *kislota* so'zi qo'shib aytiladi.

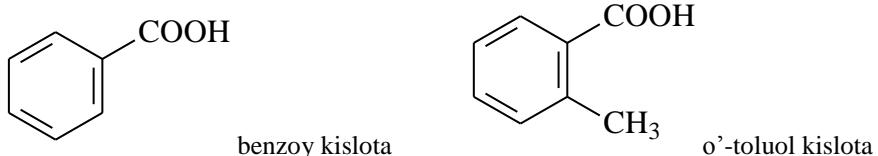
To'yangan monokarbon

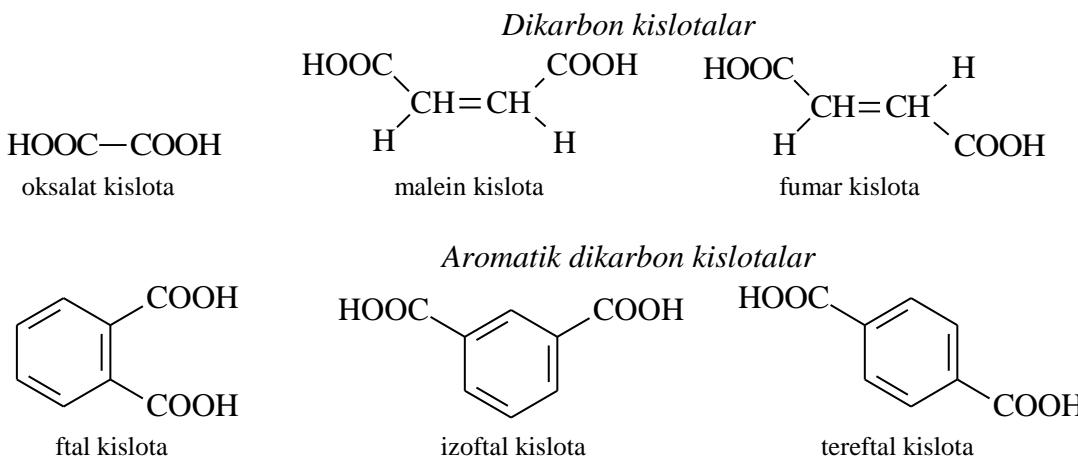
$\text{H}-\text{C}(=\text{O})-\text{OH}$	$\text{CH}_3-\text{C}(=\text{O})-\text{OH}$	$\text{CH}_3-\text{CH}_2-\text{C}(=\text{O})-\text{OH}$	$\text{CH}_3-\text{CH}_2-\text{CH}_2-\text{C}(=\text{O})-\text{OH}$
metan kislota chumoli kislota	etan kislota sirka kislota	propan kislota propion kislota	butan kislota moy kislota
$\text{CH}_3-\overset{\text{CH}_3}{\underset{ }{\text{CH}}}-\text{C}(=\text{O})-\text{OH}$	$\text{CH}_3-(\text{CH}_2)_3-\text{C}(=\text{O})-\text{OH}$		$\text{CH}_3-\overset{\text{CH}_3}{\underset{ }{\text{CH}}}-\text{CH}_2-\text{C}(=\text{O})-\text{OH}$
2-metilpropan kislota izomoy kislota	pentan kislota valerian kislota		3-metilbutan kislota izovalerian kislota
$\text{CH}_3-(\text{CH}_2)_4-\text{C}(=\text{O})-\text{OH}$	$\text{CH}_3-(\text{CH}_2)_{14}-\text{C}(=\text{O})-\text{OH}$	$\text{CH}_3-(\text{CH}_2)_{16}-\text{C}(=\text{O})-\text{OH}$	
geksan kislota kapron kislota	geksadekan kislota palmitin kislota		oktadekan kislota stearin kislota

To'yinmagan monokarbon

$\text{CH}_2=\overset{\text{CH}_3}{\underset{ }{\text{C}}}-\text{COOH}$	$\text{CH}_3-\text{CH}=\text{CH}-\text{COOH}$
propen kislota akril kislota	2-metilpropen kislota metakril kislota
$\text{CH}_3-(\text{CH}_2)_7-\text{CH}=\text{CH}-(\text{CH}_2)_7-\text{COOH}$	buten-2 kislota kroton kislota
yoki $\text{C}_{17}\text{H}_{33}\text{COOH}$ olein kislota	yoki $\text{C}_{17}\text{H}_{33}\text{COOH}$ linol kislota
$\text{C}_5\text{H}_{11}-\text{CH}=\text{CH}-\text{CH}_2-\text{CH}=\text{CH}-(\text{CH}_2)_7-\text{COOH}$	
$\text{C}_2\text{H}_5-\text{CH}=\text{CH}-\text{CH}_2-\text{CH}=\text{CH}-\text{CH}_2-\text{CH}=\text{CH}-(\text{CH}_2)_7-\text{COOH}$	
yoki $\text{C}_{17}\text{H}_{29}\text{COOH}$ linolen kislota	$\text{CH}\equiv\text{C}-\text{COOH}$ propin kislota

Aromatik monokarbon kislotalar

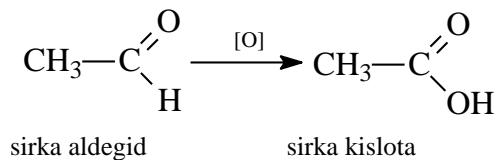




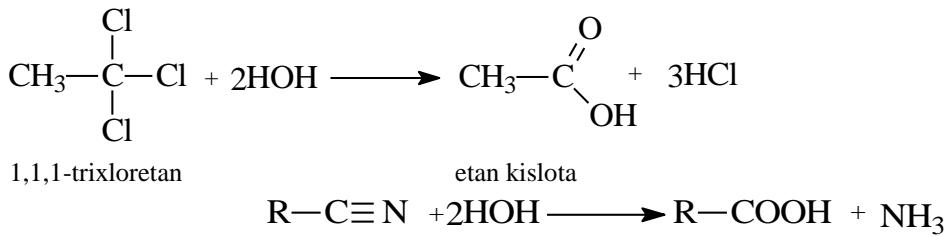
Olinishi:

Ko'pchilik organik moddalar, masalan, spirtlar va aldegidlarning oxirgi oksidlanish mahsuloti karbon kislotalari hisoblanadi.

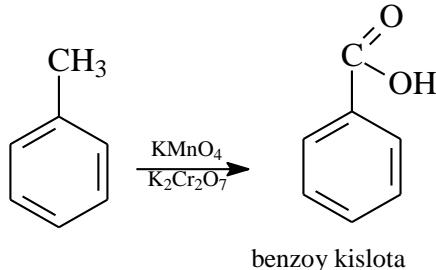
1) Aldegidlarni oksidlab olinadi:



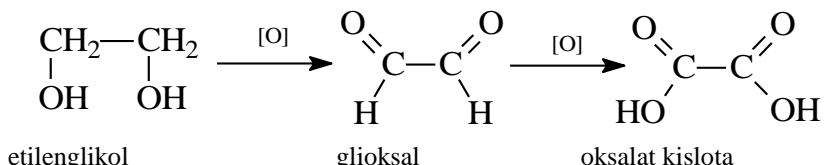
2) Alkil triklorli hosilalarni yoki nitrillarni gidroliz ailib olinadi:



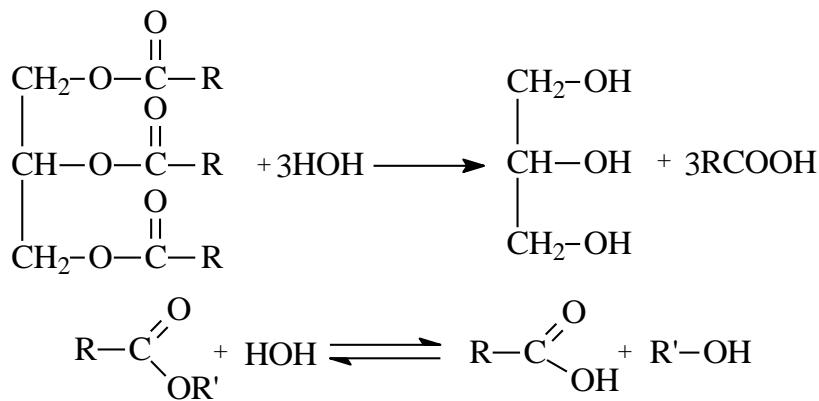
3) Benzol hosilalari oksidlanganda benzoy kislota hosil bo'ladi:



4) Etilenglikol oksidlanganda oksalat kislota hosil bo'ladi:



5) Yog'lar va murakkab efirlar gidrolizlanganda ham karbon kislotalari hosil bo'ladi:



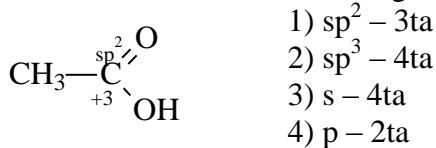
Fizilaviy xossalari:

Karbon kislotalarning quyi vakillari suyuq rangsiz moddalardir. Yuqoti vakillari rangsiz kristall moddalardir. Molekulada uglerod soni ortishi bilan ularning suvda eruvchanligi kamayadi.

Karbon kislotalarda molekulalararo vodorod bog'lanish mavjud bo'lganligi uchun ularning qaynash temperaturasi tegishli murakkab efirlarnikiga qaraganda yuqori.

Molekula tuzilishi:

Karbon kislotalarda karboksil guruh uglerodi sp^2 gibridlangan holda bo'ladi:

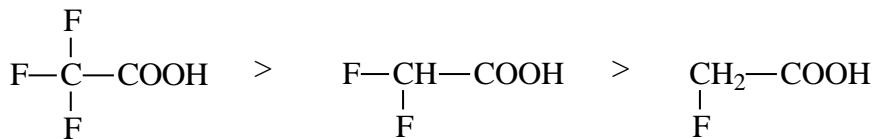
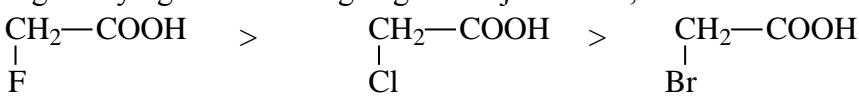


Kimyoviy xossalari:

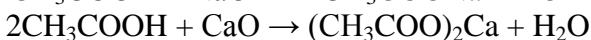
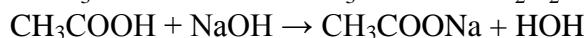
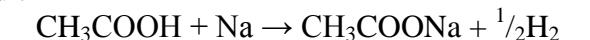
Karbon kislotalar kislotalik xossasini namoyon qiladi. To'yigan karbon kislotalarda radikalda uglerod soni ortishi bilan kislotalik kuchi kamayadi:



Agar to'yigan radikalda galogen mavjud bo'lsa, kislota kuchi ortadi:



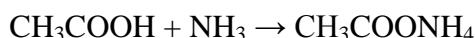
- 1) Karbon kislotalar metallar, metal oksidlari va ishqorlar bilan ta'sirlashib karbon kislota tuzlarini hosil qiladi:



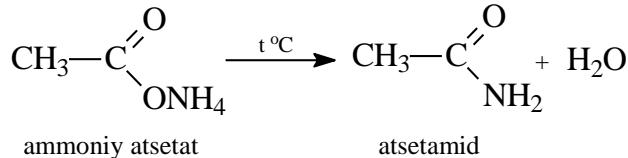
- 2) Hosil bo'lgan karbon kislota tuzlari ishqor bilan qizdirilganda dekarboksillanish reaksiysi amalga oshadi:



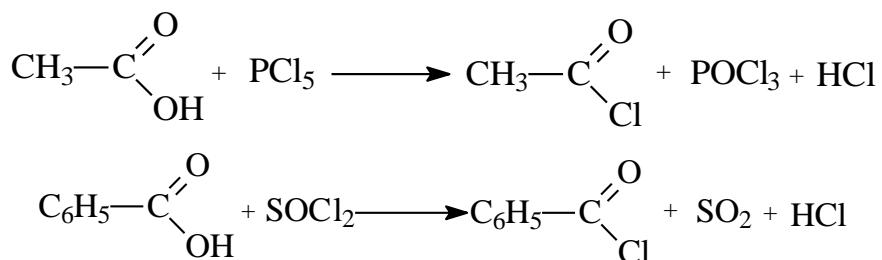
- 3) Karbon kislotalar shuningdek NH_3 va NH_4OH bilan ta'sirlashib ammoniy tuzlarini hosil qiladi:



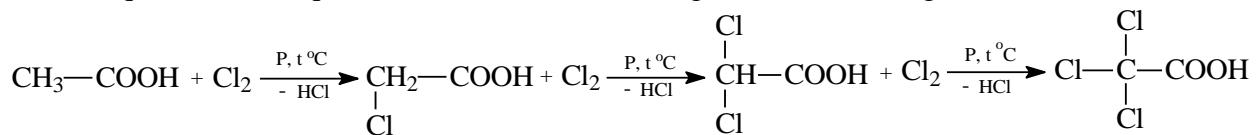
Ammoniy tuzlari 200°C da qizdirilganda degidratlanib amidlarni hosil qiladi:



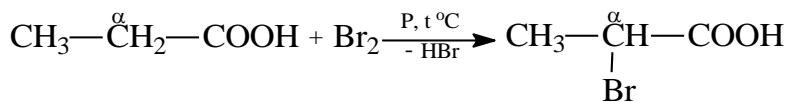
- 4) Karbon kisloatalarga PCl_5 yoki tionilxlorid SOCl_2 ta'sir ettirilganda xlorangidridlar hosil bo'ladi:



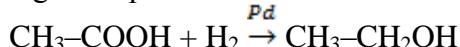
- 5) Issiq sirkə kislota qizil fosfor ishtirokida xlorlanganda xlor radikalga birikadi:



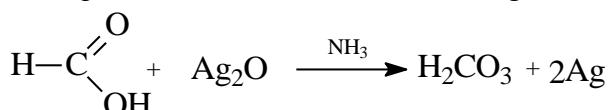
sirkə kislota xlorsirkə kislota dixlorsirkə kislota trixlorsirkə kislota
Qolgan gomologlari ham galogenlanganda dastlab faqat α -holat galogenlanadi:



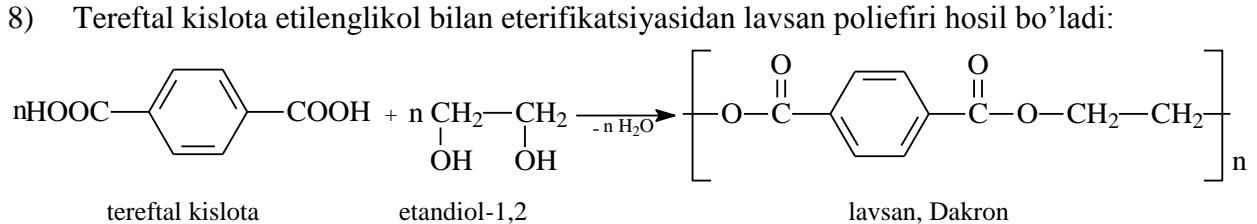
- 6) Karbon kislotalar qaytarilganda spirtlar hosil bo'ladi:



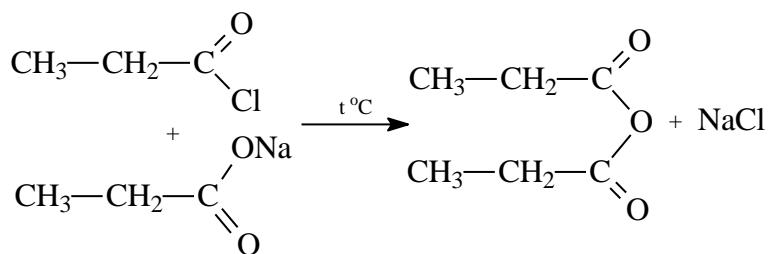
- 7) Karbon kislotalardan faqat chumoli kislota "kumush ko'zgu" reaksiyasiga kirishadi:



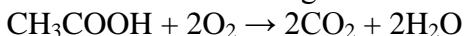
- 8) Tereftal kislota etilenglikol bilan eterifikatsiyasidan lavsan poliefiri hosil bo'ladi:



- 9) Karbon kislota xlorangidridlari tegishli tuzlar bilan qizdirilganda angidridlar hosil bo'ladi:

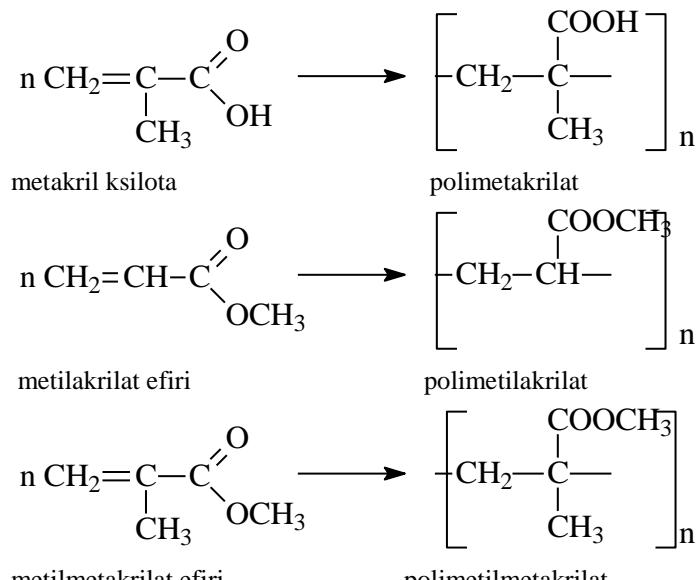


10) Karbon kislotalar yonganda suv va karbonat angidrid hosil bo'ladi:



Ishlatilishi:

Chumoli kislota junga ishlov berishda, sirka kilsota sirka va sirka angidrid (undan atsetat shoyisi va aspirin olinadi) olishda ishlatiladi. Palmitin va stearin kislotalar sovun ichlab chiqarishda ishlatiladi. Akril va metakril kislotaning metil efirlari organik shisha (pleksiglas) ishlab chiqarishda ishlatiladi:



Olein va linol kislotalar moylar tarkibiga kiradi. Benzoy kislota konserva va dori-darmon sanoatida ishlatiladi.

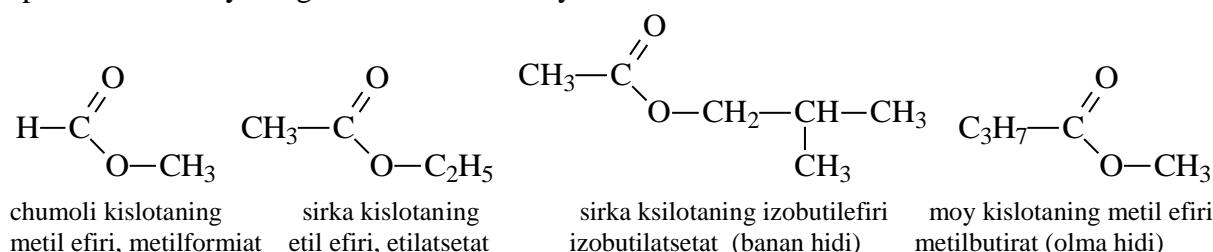
Mavzu: Murakkab efirlar.

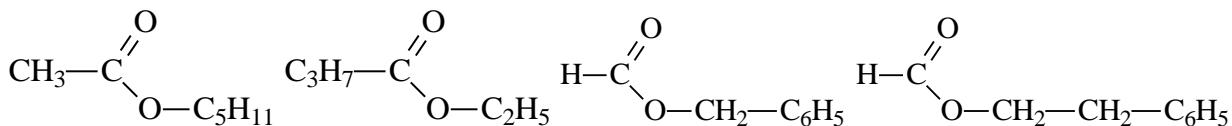
Ta'rif: Karbon kislotalar va spirlardan hosil bo'lgan karbon kislota hosilalariga murakkab efirlar deyiladi.

Ularning umumiyl formulasi: $\text{R}-\overset{\text{O}}{\underset{\text{O}}{\text{C}}}-\text{O}-\text{R}'$

Nomlanishi va izomeriyasi:

Murakkab efirlarni nomlash uchun tegishli kislota nomiga spirt nomi qo'shib *efiri* deyiladi, yoki spirt nomidan keyin tegishli kislota nomi aytildi.





sirka kislotaning izoamil
efiri, izoamilatsetat
(nok hidi)

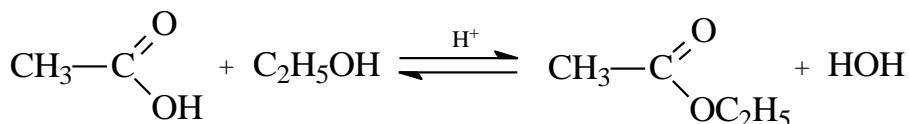
moy kislotaning
etil efiri
etilbutirat (ananas hidi)

chumoli kislotaning
benzil efiri, benzilformiat
(jasmin hidi)

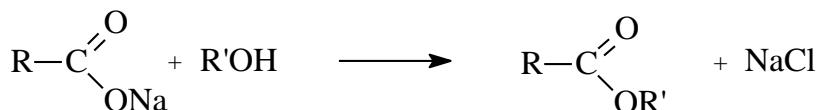
chumoli kislotaning feniletil efiri
feniletilformiat
(xrizantema hidi)

Olinishi:

Murakkab efirlar asosan eterifikatsiya reaksiyasi yordamida olinadi:



Shuningdek karbon kislota tuzlariga galogenoalkanlar ta'siridan olinadi:

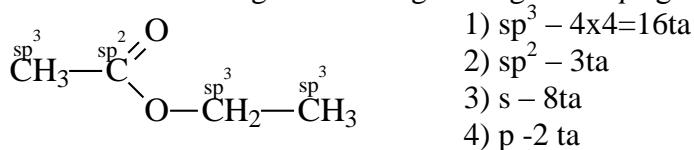


Fizikaviy xossalari:

Murakkab efirlar rangsiz, yoqimli hidli suyuqliklar hisoblanadi. Ularda vodorod bog'anish mavjud emas.

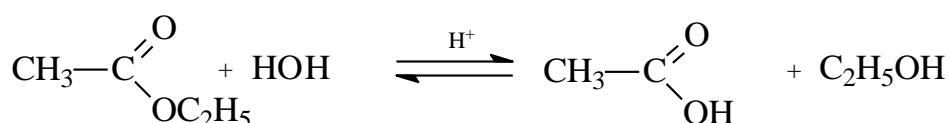
Molekula tuzilishi:

Murakkab efirlardagi karboksil guruh uglerodi sp^2 gibridlangan holda bo'ladi.

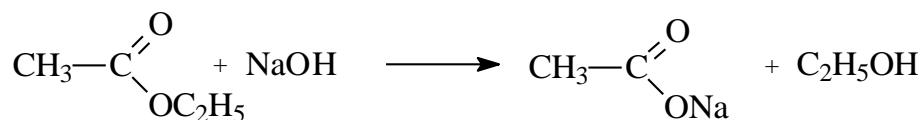


Kimyoviy xossalari:

- 1) Murakkab efirlar kislotali muhitda qaytar gidrolizga uchraydi. Bunda tegishli kislota va spirit hosil bo'ladi:

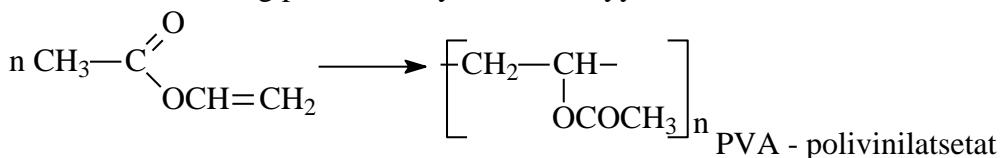


- 2) Murakkab efirlar ishqorlar bilan qaytmas gidrolziga uchraydi. Bunda karbon kislota tuzi va spirit hosil bo'ladi:



Ishlatilishi:

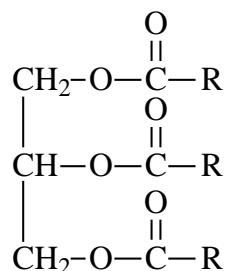
Murakkab efirlar yoqimli hidga ega bo'lganligi uchun mevalarning sun'iy hidlarini hosil qilishda ishlatiladi. Vinilatsetatning polimeri kley va laklar tayyorlashda ishlatiladi.



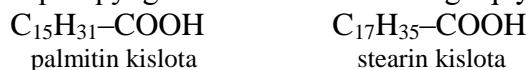
Polimetilmetakrilat organik shisha sifatida ishlatiladi.

Mavzu: Yog'lar.

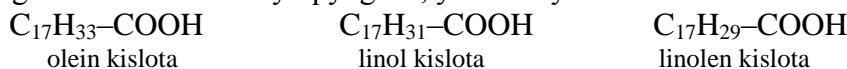
Ta'rif: Glitserin va yuqori yog' kislotalarining murakkab efirlariga yog'lar deyiladi.
Ularning umumiy formulasi:



To'yingan yog' kislotalaridan qattiq yog'lar hosil bo'ladi. Ularga quyidagilar kiradi:



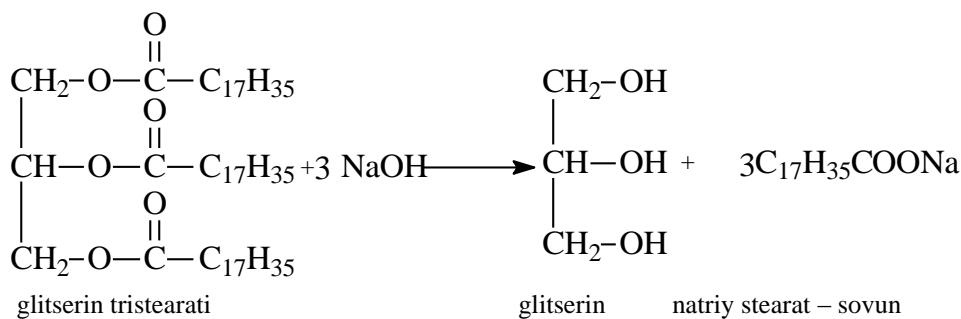
To'yinmagan yog' kisloatalaridan suyuq yog'lar, ya'ni moylar hosil bo'ladi:



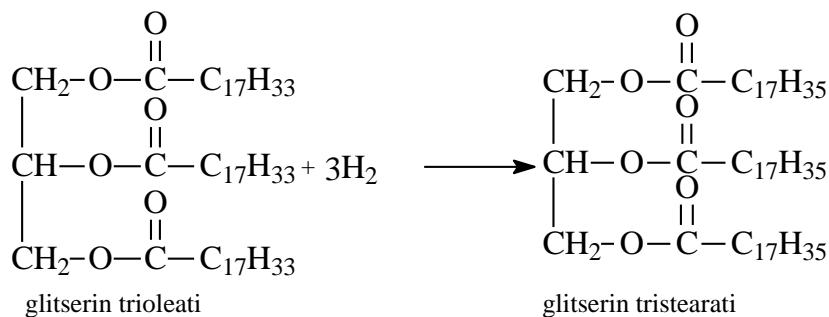
Hayvon yog'lari to'yingan, o'simlik yog'lari to'yinmagan kislotalaridan hosil bo'ladi.

Yog'larga ishqor ta'sir ettirilganda glitserin va karbon kislotaning ishqorli tuzi hosil bo'ladi.

Bunga "yog'larning Sovunlanishi" reaksiyasi deyiladi.



Suyuq moylar gidrogenlanganda qattiq yog'lar hosil bo'ladi. Bu reaksiyaga *margarinlash* reaksiyasi deyiladi:



Yog' kislotalarining kaliyli tuzlari suyuq bo'ladi.

Yog' kislotalarining Ca li va Mg li tuzlari suvda erimaydi. Shuning uchun "qattiq suv"da sovun ishlatib bo'lmaydi. Bunda sintetik yuvish vositalari – sulfokislotalarning natriyli tuzlari ishlatiladi.

Mavzu: Uglevodlar.

Ta’rif: Tarkibida gidroksil va karbonil guruhga ega organik moddalarga uglevodlar deyiladi.

Umumiyl formulasi: $C_n(H_2O)_m$.

Uglevodlar monosaxaridlar, disaxaridlar va polisaxaridlarga bo’linadi.

Mavzu: Monosaxaridlar.

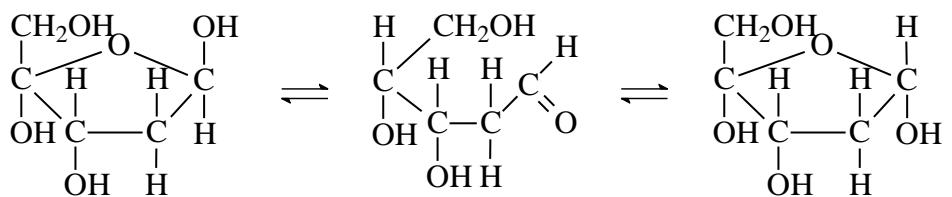
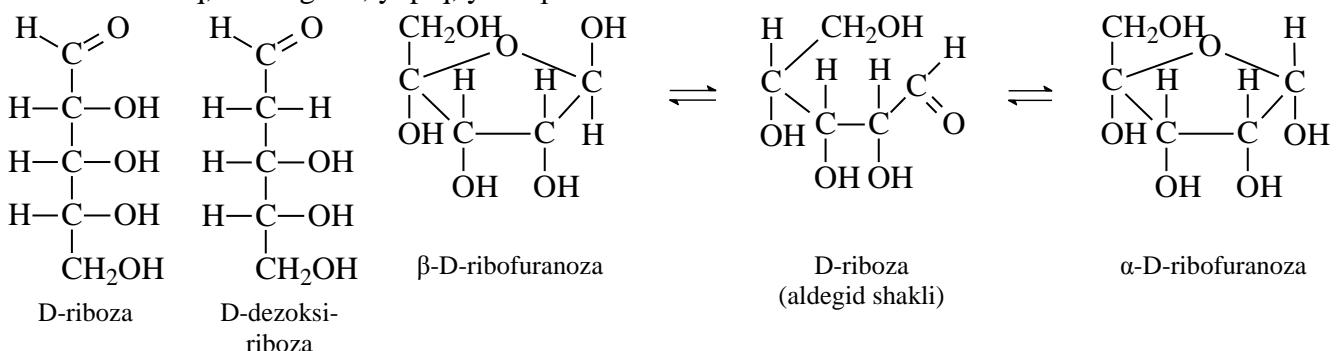
Monosaxaridlarda bitta uglevod molekulasi mavjud bo’ladi. Ular tarkibidagi uglerod soniga ko’ra tetroza, pentoza va geksozalarga bo’linadi.

Shuningdek, tarkibidagi karbonil guruh tabiatiga qarab aldoza va ketoza bo’linadi.

	Aldoza	Ketoza
Pentoza	$\begin{array}{ccccccc} & CH_2 & - & CH & - & CH & - \\ & & & & & & \\ OH & & OH & & OH & & OH \\ & & & & & & \end{array}$ $\begin{array}{ccccc} & & C=O & & \\ & & \diagdown & & \\ & & OH & & \end{array}$	$\begin{array}{ccccccc} & CH_2 & - & CH & - & CH & - \\ & & & & & & \\ OH & & OH & & OH & & OH \\ & & & & & & \end{array}$ $\begin{array}{ccccc} & & C=O & & \\ & & & & \\ & & O & & OH \end{array}$
Geksoza	$\begin{array}{ccccccc} & CH_2 & - & CH & - & CH & - \\ & & & & & & \\ OH & & OH & & OH & & OH \\ & & & & & & \end{array}$ $\begin{array}{ccccc} & & C=O & & \\ & & \diagdown & & \\ & & OH & & \end{array}$	$\begin{array}{ccccccc} & CH_2 & - & CH & - & CH & - \\ & & & & & & \\ OH & & OH & & OH & & OH \\ & & & & & & \end{array}$ $\begin{array}{ccccc} & & C=O & & \\ & & & & \\ & & O & & OH \end{array}$

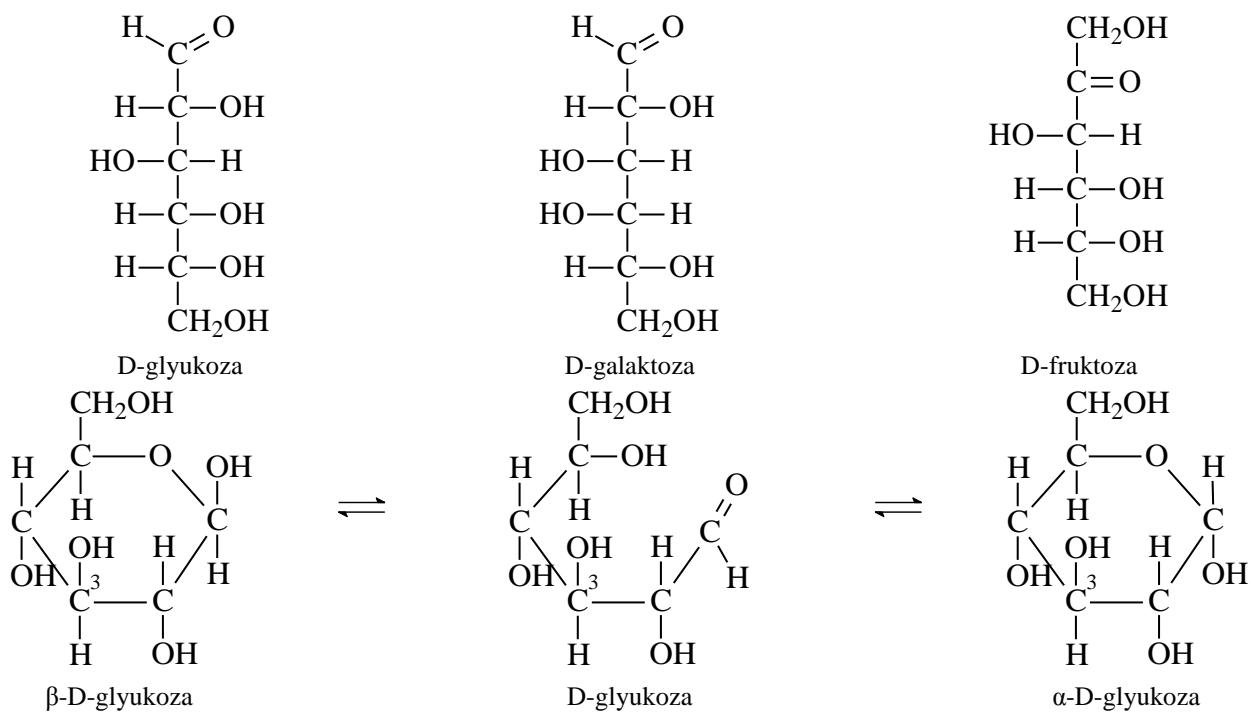
Pentozalar $C_5H_{10}O_5$ formulaga muvofiq keladi. Ularning asosiy vakillari riboza va dezoksiriboza hisoblanadi.

Uar ochiq, shiningdek, yopiq, ya’ni poluatsetal shaklda ham bo’ladi:



RNK da ribozaning siklik β -shakli, DNK da dezoksiribozaning siklik β -shaklida bo’ladi.

Geksozalar $C_6H_{12}O_6$ formulaga muvofiq keladi. Ularning asosiy vakillari glyukoza, fruktoza va galaktoza hisoblanadi.

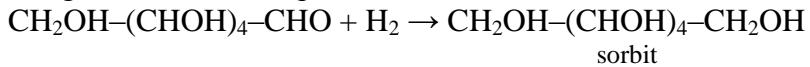


Fizikaviy xossalari:

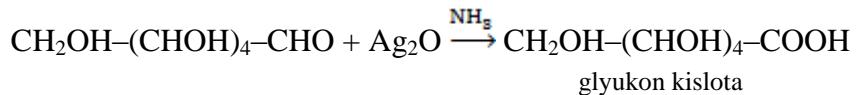
Glyukoza va fruktoza rangsiz shirin ta'mli moddalardir. Ular suvda yaxshi eriydi.

Kimyoiy xossalari:

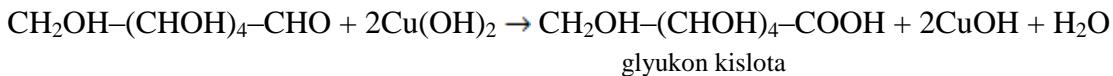
- 1) Glyukoza qaytarilganda olti atomli spirit hosil bo'ladi:



- 2) Glyukoza aldegidlar kabi "kumush ko'zgu" reaksiyasiga kirishadi. Bunda glyukon kislota hosil bo'ladi:



- 3) Glyukoza shuningdek yangi tayyorlangan $\text{Cu}(\text{OH})_2$ bilan ham aldegidlar kabi ta'sirlashadi:



- 4) Glyukoza uchun 4 xil bijg'ish xarakterli:

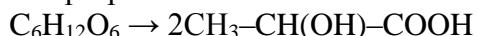
a) Spiritli bijg'ish – etanol hosil bo'ladi:



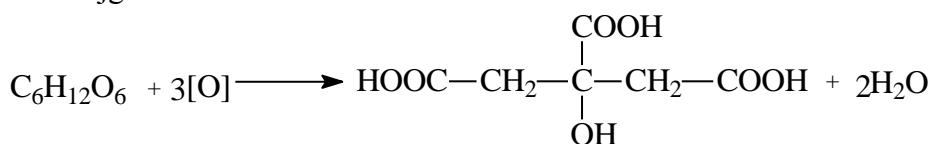
b) Moy kislotali bijg'ish – butan kislota hosil bo'ladi:



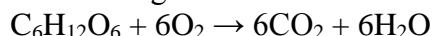
c) Sut kislotali bijg'ish – 2-gidroksipropan kislota hosil bo'ladi:



d) Limon kislotali bijg'ish:

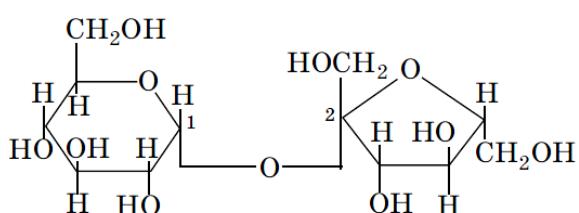


- 5) Glyukoza yonganda suv va karbonat angidrid hosil bo'ladi:

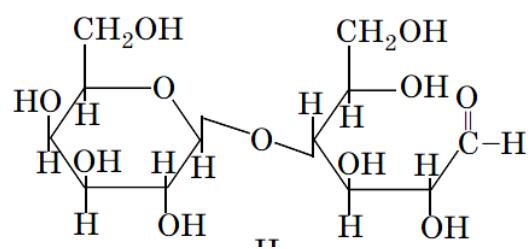
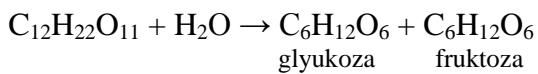


Mavzu: Disaxaridlar-C₁₂H₂₂O₁₁.

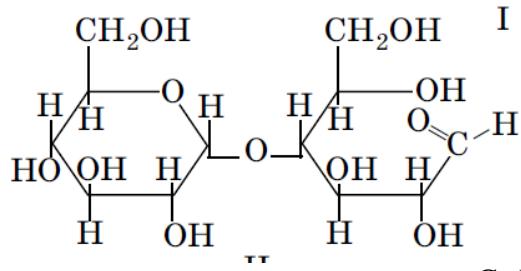
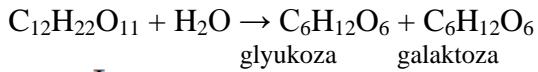
Disaxaridlar ikkita monosaxaridlardan hosil bo'lgan. Ularga saxaroza, maltoza va lakoza kiradi.



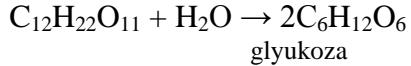
Saxaroza α -D-glyukoza va β -D-fruktozadan tuzilgan. Shuning uchun saxaroza gidrolizidan glyukoza va fruktoza hosil bo'ladi.



Lakoza β -D-galaktoza va β -D-glyukozadan tuzilgan. Shining uchun lakoza gidrolizidan galatoza va glyukoza hosil bo'ladi.



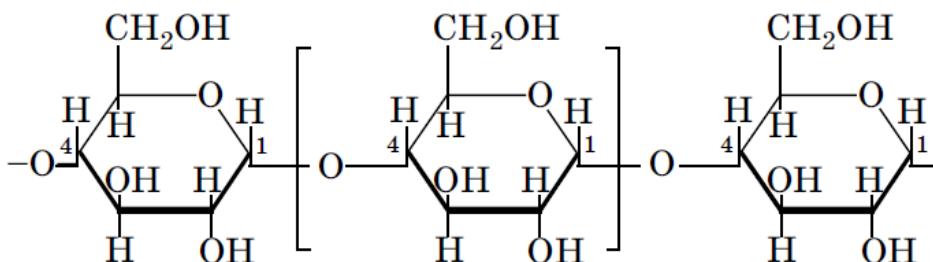
Maltoza ikki molekula α -D-glyukozadan tuzilgan. Shing uchin maltoza gidrolizidan faqat glyukoza hosil bo'ladi.



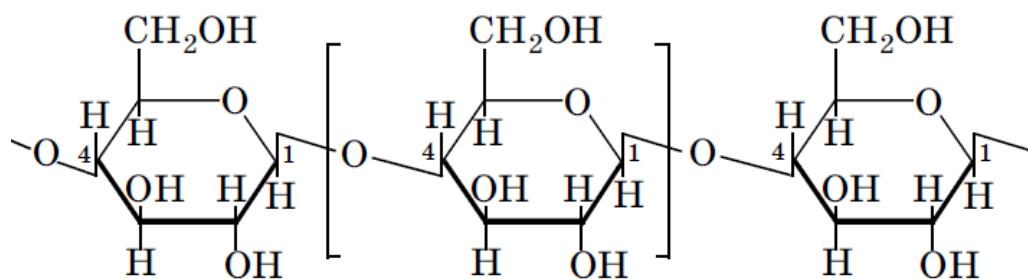
Mavzu: Polisaxaridlar – (C₆H₁₀O₅)_n.

Polisaxaridlarga kraxmal, sellyuloza va glikogen kiradi.

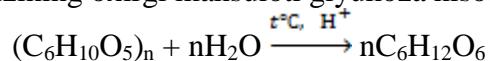
Kraxmal α -D-glyukozalar zanjiridan tuzilgan:



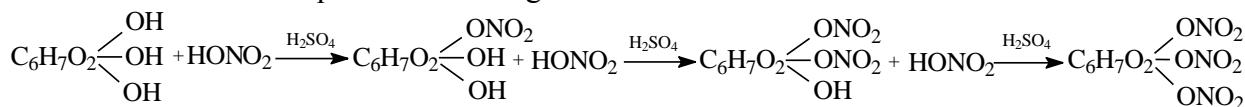
Sellyuloza β -D-glyukozalar zanjiridan tuzilgan:



Kraxmal va sellyuloza gidrolizining oxirgi mahsulotlari glyukoza hisoblanadi:

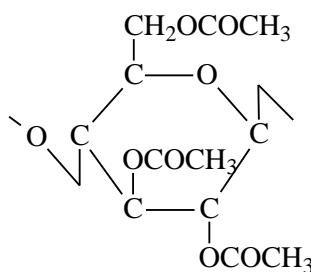


Sellyuloza molekulasida 3 ta hidroksil guruh bo'ganligi uchun u nitrat kislota bilan mono-, di- va trinitrohosilalarni hosil qiladi. Hosil bo'lган mahsulotlar murakkab efir hisoblanadi:



Trinitroselyuloza – piroksilin tutunsiz porox sifatida ishlatiladi.

Shuningdek sellyuloza sirka kislota bilan atsetat efirlar hosil qiladi:
bu efirlar sun'iy atsetat tolalar olishda ishlatiladi:



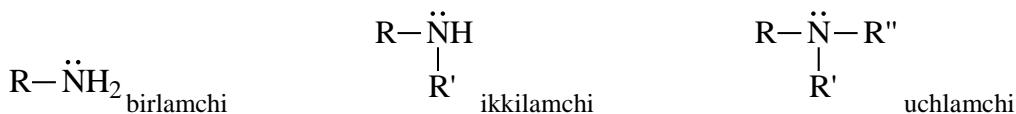
IV BOB. AZOT SAQLOVCHI ORGANIK BIRIKMALAR.

Mavzu: Aminlar.

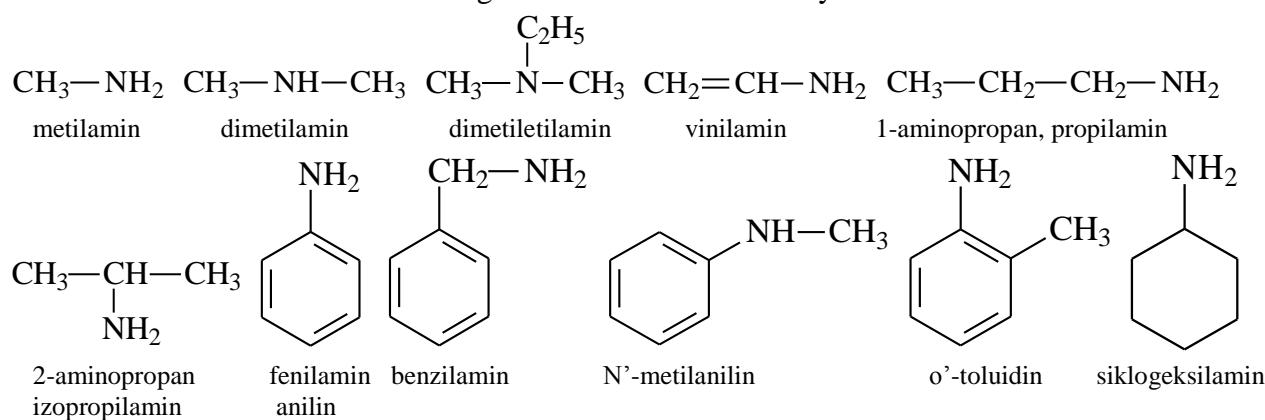
Ta’rif: Ammiak molekulasidagi bitta, ikkita yoki uchta vodorod atomining uglevodorod radikaliga almashinishidan hosil bo’lgan organik moddalar sinfiga aminlar deyiladi.

Nomlanishi va izomeriyasi:

Aminlar ammiakdagi vodorod almashinish darajasiga ko’ra birlamchi, ikkilamchi va uchlamchi aminlarga bo’linadi:



Aminlarni nomlash uchun azotga tutash radikallar nomi aytildi:

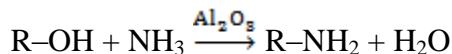


Olinishi:

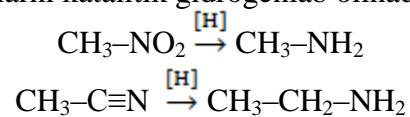
- 1) Ammiak va aminlarni galgenalkanlar bilan alkillab olinadi:



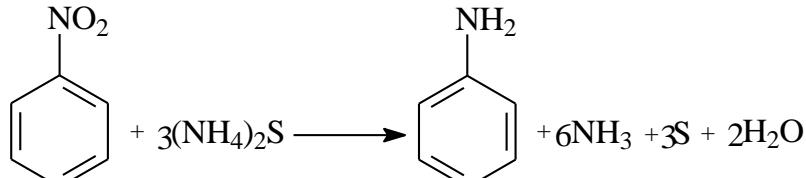
- 2) Spirlarning Al_2O_3 katalizatori ishtirokida ammiak bilan ta’siridan olinadi:



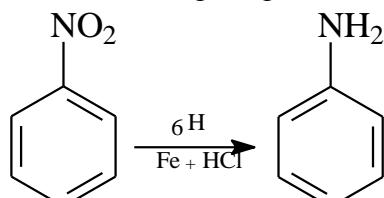
- 3) Nitrobirkimlarni yoki nitrillarni katalitik gidrogenlab olinadi:



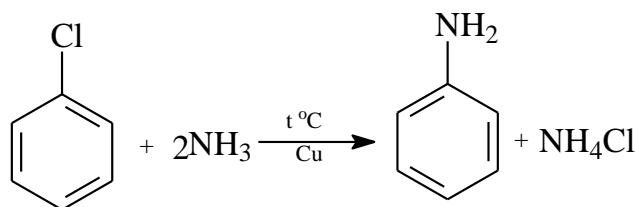
- 4) Anilinni birinchi bo’lib 1842 yil Zinin nitrobenzolni ammoniy sulfid bilan qaytarib olgan:



- 5) Hozirgi vaqtida anilin nitrobenzolni katalitik gidrogenlab olinadi:



- 6) Galogenalkanlarga ammiak ta’sir ettirib olinadi:



Fizikaviy xossalari:

Alkinaminlar rangsiz gaz yoki suyuqliklardir. Anilin o'ziga xos hidli moysimon suyuqlik.

Aminlarda molekulalararo vodorod bog'lanish mavjud.

Yuqori aminlar aminokislolar parchalanishidan hosil bo'lganligi uchun ular aynigan baliq hidiga ega.

Molekula tuzilishi:

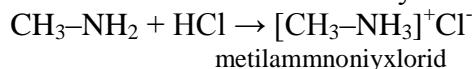
Aminlarda gibriddanish radikal tabiatiga bog'liq.

Kimyoviy xossalari:

Aminlar asoslik xossasini namoyon qiladi. Alifatik aminlar ammiakdan kuchliroq asos hisoblansa, aromatik aminlarning asosligi ammiaknikidan kuchsiz:

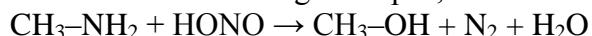


- 1) Aminlar mineral kislolar bilan ta'sirlashib alkilammoniy tuzlarini hosil qiladi:

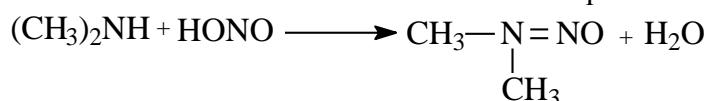


- 2) Aminlar nitrit kislota bilan ta'sirlashadi:

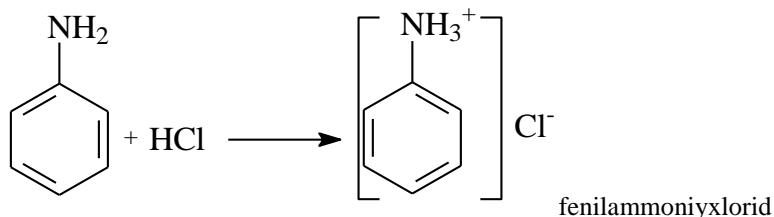
a) Birlamchi aminlar nitrit kislota bilan ta'sirlashganda spirt, azot va suv hosil bo'ladi:



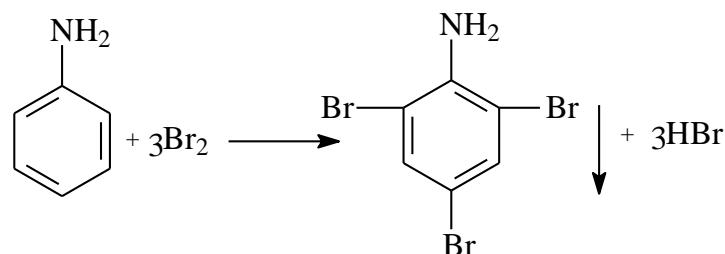
b) Ikkilamchi aminlar nitrit kislota ta'sirida nitrozaaminlarni hosil qiladi:



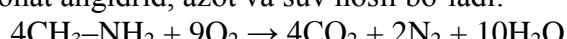
- 3) Anilin kislolar bilan xuddi alifatik aminlardek ta'sirlashadi:



- 4) Aminoguruh I tur o'rribosar hisoblanganligi uchun reaksiyon markazni o' va p' holatga yo'naltiradi:



- 5) Aminlar yonganda karbonat angidrid, azot va suv hosil bo'ladi:

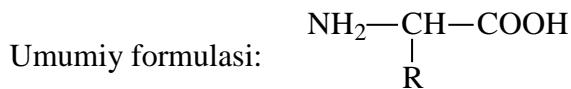


Ishlatilishi:

Anilin bo'yoqlar va portlovchi moddalar olishda ishlatiladi. Metilamin va dimetilamin dori ishlab chiqarishda ishlatiladi.

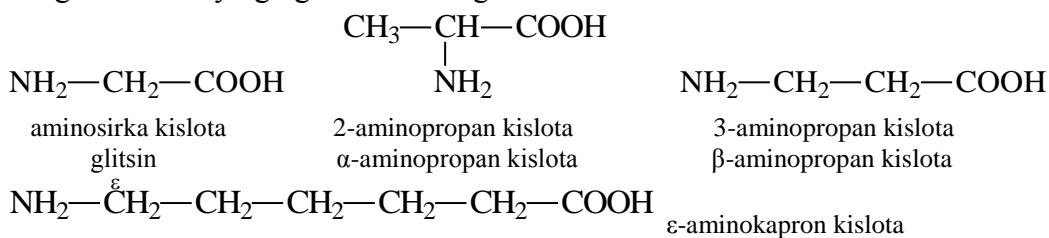
Mavzu: Aminokislotalar.

Ta'rif: Tarkibida ham amino $-NH_2$ ham karboksil $-COOH$ guruh tutgan organik moddalarga aminokislotalar deyiladi.



Nomlanishi va izomeriyasi:

Aminokislotalarda karboksil guruh 1 deb olinib, aminoguruh joyi qo'shib aytildi. Shuningdek karboksil guruhdan keying uglerod α holatga nisbatan nomlanadi:



Eng ko'p uchraydigani bu α -aminokislotalar hisoblanadi:

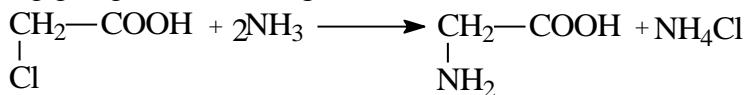
<i>Nº</i>	<i>Nomi</i>	<i>Formulasi</i>	<i>Belgisi</i>	<i>Harfli belgisi</i>
1	Glitsin	$\text{NH}_2-\text{CH}_2-\text{COOH}$	Gly	G
2	Alanin	$\begin{array}{c} \text{CH}_3-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Ala	A
3	Valin	$\begin{array}{c} \text{CH}_3-\text{CH}-\text{CH}-\text{COOH} \\ \qquad \\ \text{CH}_3 \qquad \text{NH}_2 \end{array}$	Val	V
4	Izoleysin	$\begin{array}{c} \text{CH}_3-\text{CH}_2-\text{CH}-\text{CH}-\text{COOH} \\ \qquad \\ \text{CH}_3 \qquad \text{NH}_2 \end{array}$	Ile	I
5	Leysin	$\begin{array}{c} \text{CH}_3-\text{CH}-\text{CH}_2-\text{CH}-\text{COOH} \\ \qquad \\ \text{CH}_3 \qquad \text{NH}_2 \end{array}$	Leu	L
6	Fenilalanin	$\begin{array}{c} \text{C}_6\text{H}_5-\text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Phe	F
7	Serin	$\text{HO}-\text{CH}_2-\text{CH}-\text{COOH}$ $\qquad \qquad $ $\qquad \qquad \text{NH}_2$	Ser	S
8	Treonin	$\begin{array}{c} \text{CH}_3-\text{CH}-\text{CH}-\text{COOH} \\ \qquad \\ \text{OH} \qquad \text{NH}_2 \end{array}$	Thr	T

9	Lizin	$\begin{array}{c} \text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Lys	K
10	Arginin	$\begin{array}{c} \text{HN}=\text{C}-\text{NH}-(\text{CH}_2)_3-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Arg	R
11	Asparagin kislota	$\begin{array}{c} \text{HOOC}-\text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Asp	D
12	Glutamin kislota	$\begin{array}{c} \text{HOOC}-\text{CH}_2-\text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Glu	E
13	Sistein	$\begin{array}{c} \text{HS}-\text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Cys	C
14	Tirozin	$\begin{array}{c} \text{HO}-\text{C}_6\text{H}_4-\text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Tyr	Y
15	Prolin	$\begin{array}{c} \text{CH}-\text{COOH} \\ \\ \text{NH} \end{array}$	Pro	P
16	Triptofan	$\begin{array}{c} \text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Trp	W
17	Gistidin	$\begin{array}{c} \text{N} \\ \\ \text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH} \end{array}$	His	H
18	Metionin	$\begin{array}{c} \text{CH}_3-\text{S}-\text{CH}_2-\text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Met	M
19	Glutamin	$\begin{array}{c} \text{H}_2\text{N}-\text{C}(=\text{O})-\text{CH}_2-\text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Gln	Q
20	Asparagin	$\begin{array}{c} \text{H}_2\text{N}-\text{C}(=\text{O})-\text{CH}_2-\text{CH}-\text{COOH} \\ \\ \text{NH}_2 \end{array}$	Asn	N

Olinishi:

Aminokislolar oqsillarning monomeri hisoblanadi. Shuning uchun ular oqsil gidrolizidan hosil bo'ladi.

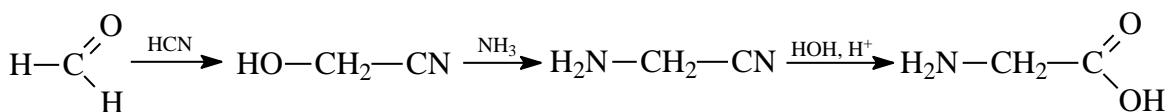
1) Karbon kislotaning gelogenli hosilalariga ammiak ta'siridan olinadi:



2-xlorsirka kislota

2-aminoxirka kislota

2) Glitsinni formaldegiddan olish mumkin:

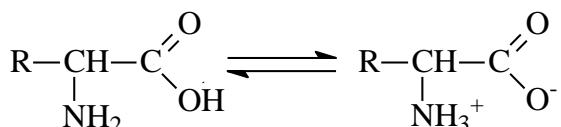


Fizikaviy xossalari:

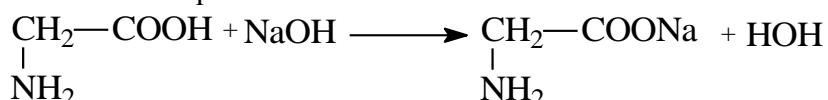
Aminokislotalar rangsiz kristall moddalar bo'lib, suvda yaxshi eriydi. Organik erituvchilarda kam eriydi. Ko'pchiligi shirin ta'mli.

Kimyoviy xossalari:

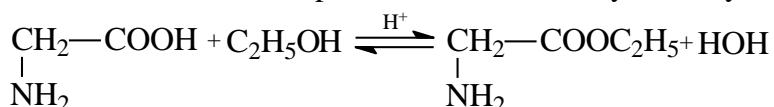
Aminokislotalarda ham amino (asos), ham karboksil (kislota) guruh bo'lganligi uchun ular ichki tuz – betainlarni hosil qiladi:



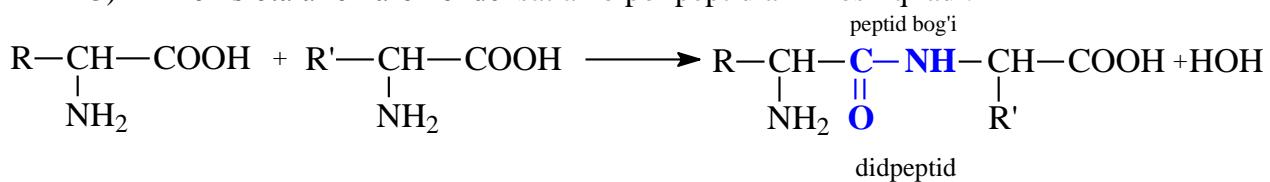
1) Aminokislotalar kuchsiz kislotalik xossasini namoyon qiladi. Ular ishqorlar bilan ta'sirlashib tuzlarni hosil qiladi:



2) Aminokislotalar kislotalar ishtirokida spirtlar bilan eterifikatsiya reaksiyasiga kirishadi:



3) Aminokislotalar o'zaro kondensatlanib polipeptidlarni hosil qiladi:



Ishlatilishi:

Aminokislotalar muhim biologik moddalar hisoblanadi. Ular oqsillarning asosiy tarkibidir. ε-aminokapron kislota kaprolaktam olishda ishlatiladi. Kaprolaktam poliamid materiallar (naylon) olishda ishlatiladi.

Mavzu: Oqsillar.

Ta'rif: Bir-birlari bilan peptid bog'lari orqali bog'langan α-aminokislotalardan tashkil topgan tabiiy biopolimerlarga oqsillar deyiladi.

Oqsillar tarkibiga ko'ra 2 ga bo'linadi:

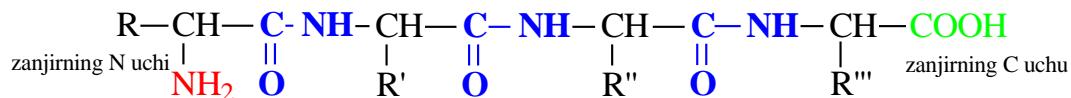
- 1) Proteinlar.
- 2) Proteidlar.

Proteinlar faqat aminokislotalar qoldig'idan tuzilgan. Ular albuminlar (ular suvda eriydi, sut, tuxum va qon oqsili kiradi), globulinlar (suvda erimaydi, qon globilinlari va mushak oqsili – miozin kiradi), skleroproteinlar (suvda erimaydi, keratin, teri oqsili va fibroin kiradi)ga bo'linadi.

Proteidlar – murakkab oqsillar, aminokislotalar va oqsil bo'lмаган qoldiqlardan tuzilgan. Ularga fosfoproteidlar (fosfat kislota qoldig'i tutgan oqsil, masalan, sut oqsili kazein), glikoproteidlar (uglevod

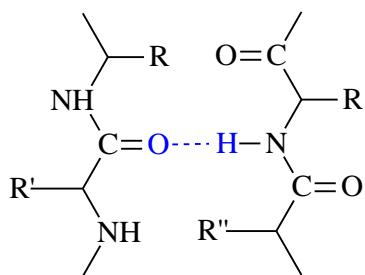
qoldig'i tutgan oqsillar), xromoproteidlar (rangli moddalar, masalan, hemoglobin), nukleoproteidlar (nuklein kislota qoldig'i tutgan oqsillar) kiradi.

Oqsillar faqat α -aminokislotalar qoldig'idan tuzilgan:



Oqsillar 4 xil strukturada mavjud bo'ladi:

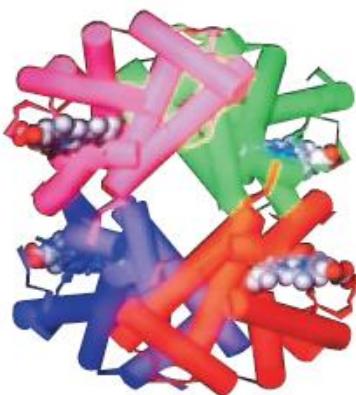
- 1) Birlamchi;
- 2) Ikkilamchi;
- 3) Uchlamchi;
- 4) To'rtlamchi.
- 1) Oqsillarning birlamchi strukturasi aminokislotalarning chiziqli zanjiridir:



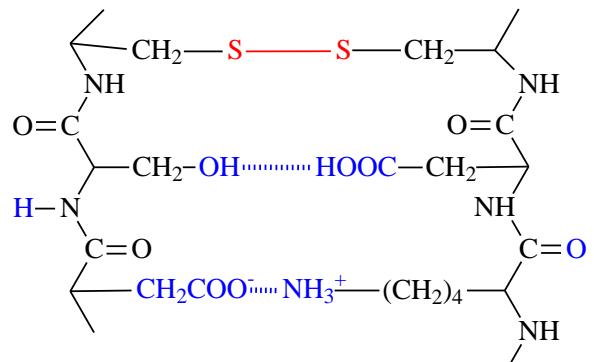
Phe-Val-Asn-Gln-His-Leu-Cys-Gly-Ser-His-Leu-Val-Glu-Ala-Leu-Tyr-Leu-Val-Cys

- 2) Oqsillarning ikkilamchi strukturasida aminokislotalar molekulasidagi karbonil va amino guruhlar orasida vujuga kelgan vodorod bog'lari tufayli spiral fazoviy shaklga keladi:
- 3) Oqsilning uchlamchi strukturasi polipeptid zanjiridagi funksional guruhlari o'zaro ta'sridan vujudga keladi.

Bunda karboksil va aminoguruhdan tuz ko'prigi, karboksil va hidroksil guruhdan murakkab efir ko'prigi, oltingugurt atomlari disulfid ko'prigi, shuningdek hidrofob ta'sirlar natijasida hosil bo'ladi.



- 4) Oqsilning to'rtlamchi strukturasida bir nechta oqsil molekulalari yig'ilib, murakkab tuzsilishni hosil qiladi:



Oqsillarning barqarorligi turlicha hisoblanadi. Ular kislotali yoki ishqoriy muhitda gidrolizlanib, α -aminokislotalarni hosil qiladi.

Oqsillar tarkibiga turli funksional guruhlarning kirganligi rangli reaksiyalar yordamida aniqlanadi.

- 1) *Biuret* reaksiyasi – oqsilga ishqoriy muhitda CuSO_4 eritmasi qo'shilganda binafsha rang hosil bo'ladi. Bu peptid bog'ga sifat reaksiya hisoblanadi.
- 2) *Ksantoprotein* reaksiyasi – aromatik va geteroatomli xalqali oqsillarga kons. HNO_3 qo'shilganda, sariq rangga kiradi (xalqaning nitrolanishi). Ishqor qo'shilganda sariq rang zarg'aldoq rangga kiradi (fenilalanin va tirozin).
- 3) *Millon* reaksiyasi – oqsilga $\text{Hg}(\text{NO}_3)_2$ va nitrat kislota ta'sirida qizil-qo'ng'ir cho'kma hosil bo'ladi.
- 4) Agar oqsil tarkibida oltingugurt tutsa, qo'rg'oshin tuzlari ta'sirida eritmada qora PbS cho'kmasi hosil bo'ladi.

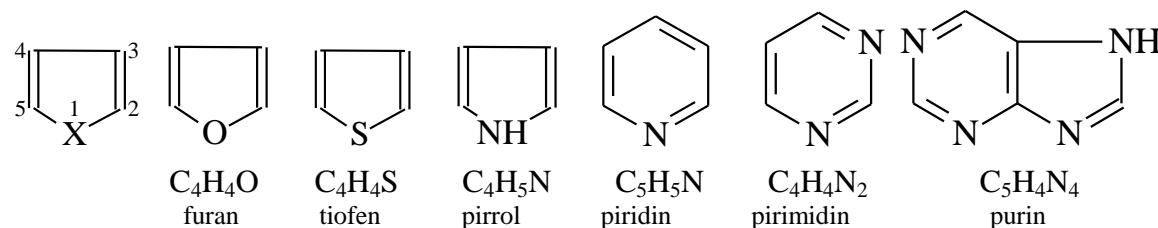
Oqsillar hayotning asosi hisoblanadi. Masalan, gemoglobin oqsili ($C_{738}H_{1166}O_{208}S_2Fe_4$), insulin gormoni $C_{254}H_{377}N_{65}O_{75}S_6$ oqsillar shular jumlasiga kiradi.

Mavzu: Geterosiklik birikmalar.

Ta’rif: Xalqa tarkibida uglerod atomidan boshqa element atomlari (N , S , O) tutgan moddalarga geterosiklik birikmalar deyiladi.

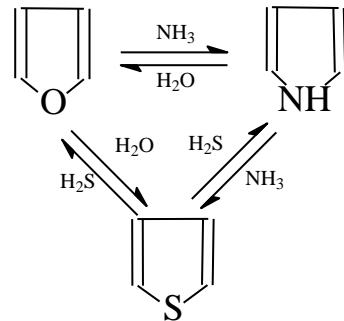
Nomlanishi va izomeriyasi:

Geterosiklda geteroatom 1 deb raqamlanib, sikl nomi aytildi:



Olinishi:

Al_2O_3 katalizatori sihtirokida ularning bir-biriga aylanishini quyidagicha tasvirlash mumkin:

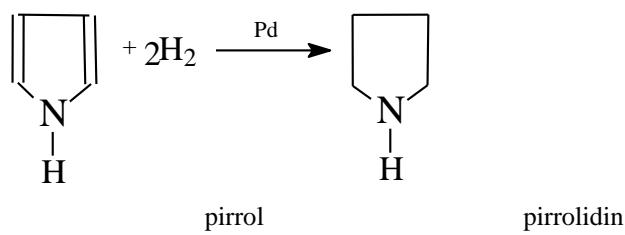


Fizikaviy xossalari:

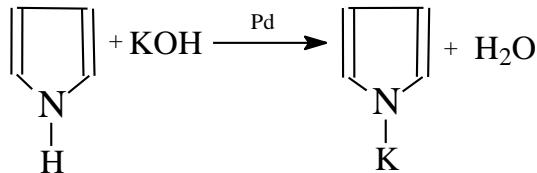
Pirrol, tiofen va furan yoqimli hidli rangsiz suyuqliklardir.

Kimyoiy xossalari:

1) Ularning barchasi gidrogenlanganda qo’shbog’ uziladi:



2) Pirrolning vodorodi nisbatan erkin. Shuning uchun u kuchsiz kislota xossasini namoyon qiladi:

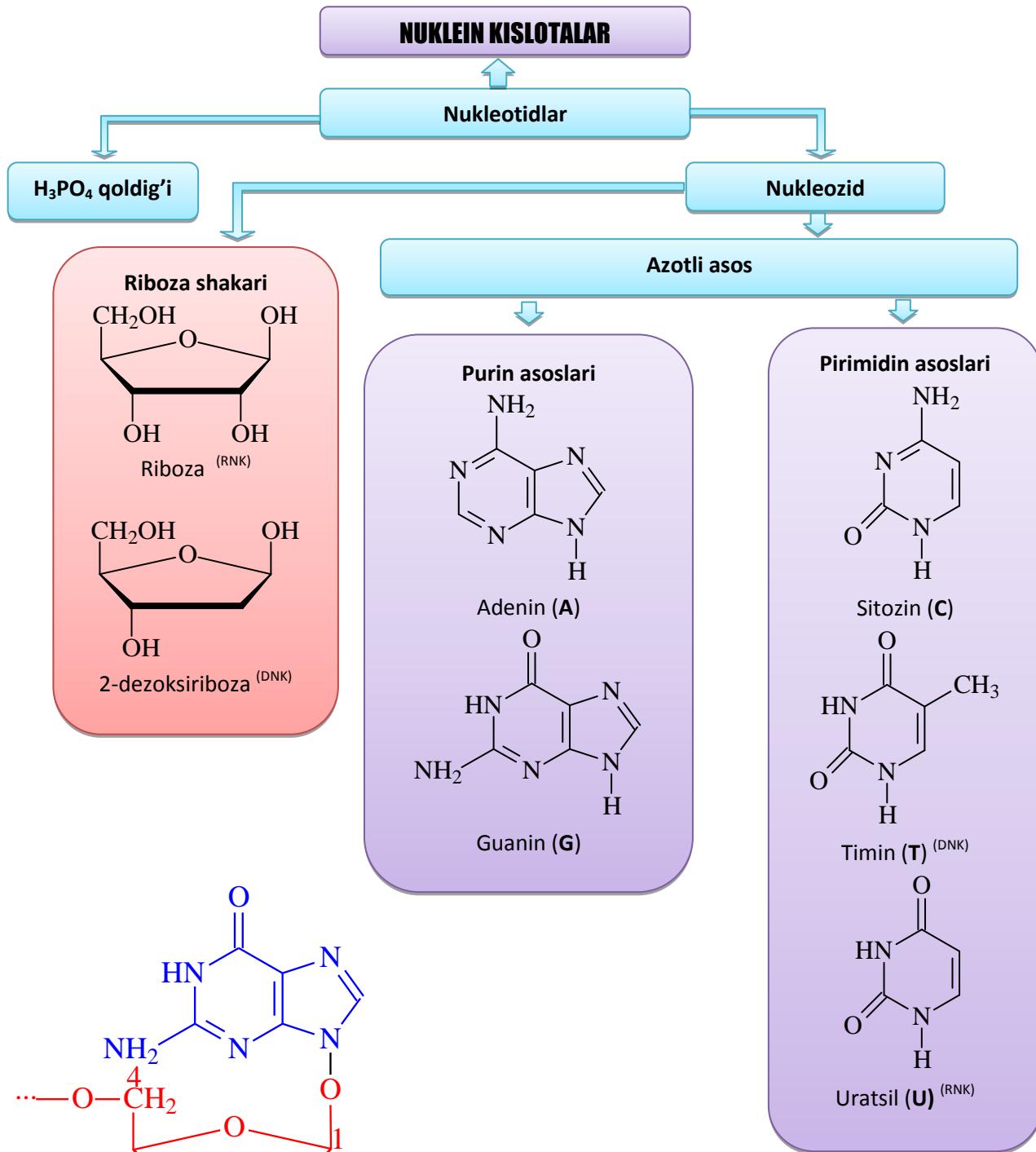


Ishlatilishi:

Pirrol va furan organik sintezda ishlatiladi. Piridin organik erituvchi sifatida ishlatiladi.

Mavzu: Nuklein kislotalar.

Ta’rif: Nukleotid monomerlarining makromolekulasiga nuklein kislotalar deyiladi.



Nukleozidlар – bu purin va primidin asoslalarining riboza yoki dezoksiriboza bilan birikmasidir.

Ribonuklein kislota molekulasini quyidagicha tasvirlash mumkin:

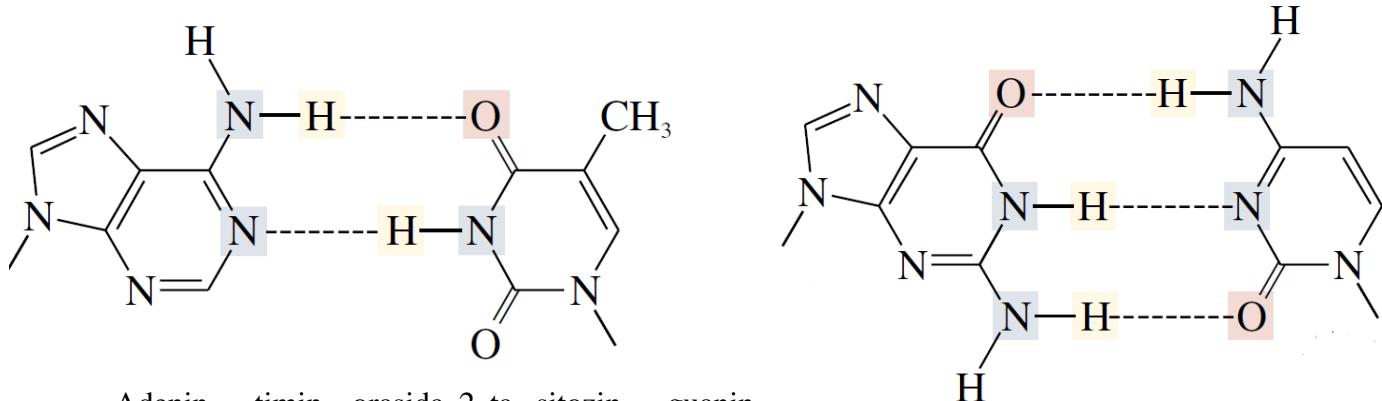
Zanjirda azotli asos bilan pentozaning 1'-uglerodi, fosfat kislota qoldig'i bilan 3'-uglerodi va keyingi nukleozid bilan 5'-ulerod orqali bog'langan.

Hujayra tarkibida ribonuklein kislota (RNK) va dezoksiribonuklein kislota (DNK) farqlanadi.

RNK tarkibida *riboza*, fosfat kislota qoldig'i, adenin, guanin, sitozin va *uratsil* bo'ladi. D NK tarkibida esa *dezoksiriboz*a, fosfat kislota qoldig'i, adenin, guanin, sitozin va *timin* bo'ladi.

DNK zanjiri qo'shaloq spiral tuzilishga ega. Buni 1953 yilda (1962 yilgi Nobel mukofoti sovrindorlari D.Uotson va F.Krik ochishgan. Bu ikki spiral bir-birlari bilan purin va pirimidin asoslari orasida vujudga keladigan *vodorod bog'lanish* orqali bog'langan bo'ladi. Molekulalararo vodorod bog'lanish *komplementar* (mos) juftlarni hosil qildi.

Ya'ni adenin – guanin (A – T) jufti va sitozin – guanin (C – G) juftlari.



Adenin – timin orasida 2 ta, sitozin – guanin orasida 3 ta vodorod bog'i mavjud.

Inson D NK si tarkibida purin va pirimidin asoslarining ulushi quyidagicha:

Purin	Pirimidin	Asos nisbati
Adenin (A) 30,3%	Timin (T) 30,3%	A/T=1,00
Guanin (G) 19,5%	Sitozin (C) 19,9%	G/C=0,98
Jami purinlar: 49,8%	Jami pirimidinlar: 50,1%	

RNK da timin asosiga uratsil asosi mos keladi.

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4. Francis A. Carey. Organic Chemistry, USA, McGraw-Hill, 2000.
5. CRC Handbook of Chemistry and Physics, USA, CRC, 2003.
6. Полинг Л. «Общая химия», М.: Мир, 1974.
7. “Kimyo-mavzulashtirilgan testlar to’plami”, Buxoro, 2009.

4-Qism

ILOVALAR

***Ma'lumotlar**

***Jadvallar**

***Kattaliklar**

ILOVALAR

VALENTLIK

O'zgarmas valentli elementlar:

I	Li, Na, K, Rb, Cs, H, F, Ag
II	Be, Mg, Ca, Sr, Ba, Zn, Cd, O
III	B, Al

Guruh valentliklari:

I valentli

-OH	Gidroksil
-NO ₂	Nitrit
-NO ₃	Nitrat
-F (Cl, Br, J)	Galogenid
-FO (Cl, Br, J)	Gipo+Γ+it
-FO ₂ (Cl, Br, J)	Γ+it
-FO ₃ (Cl, Br, J)	Γ+at
-FO ₄ (Cl, Br, J)	Per+Γ+at
-CN	Sianid
-MnO ₄	Permanganat
-NH ₄	Ammoniy

II valentli

=S	Sulfid
=SO ₃	Sulfit
=SO ₄	Sulfat
=CO ₃	Karbonat
=SiO ₃	Silikat
=CrO ₄	Xromat
=Cr ₂ O ₇	Bixromat, dixromat
=MnO ₄	Manganat
=B ₄ O ₇	Tetraborat
=HPO ₃	Fosfit
=S ₂ O ₃	Tiosulfat

O'zgaruvchan valentli elementlar:

II-IV	C, Si, Sn, Pb
III-V	P, As
II-IV-VI	S, Se, Te
I-III-V-VII	Cl, Br, J
I-II-III-IV	N
II-III-VI	Cr, Mo
II-III-IV-VI-VII	Mn
II-III	Fe, Co, Ni
I-II	Cu, Hg
I-III	Au, Tl

III valentli

≡N	Nitrid
≡P	Fosfid
≡PO ₄	Fosfat
≡AsO ₃	Arsenit
≡AsO ₄	Arsenat
≡BO ₃	Borat

IV valentli

P ₂ O ₇	Pirofosfat
SiO ₄	Ortosilikat

ANORGANIK BIRIKMALARNING ASOSIY SINFLARI

Valentligi	Oksidlar	Asoslar	Kislotalar	Tuzlar
				Me_nAc_m
I	E_2O (Na_2O)	$MeOH$ ($NaOH$)		
II	EO (CaO)	$Me(OH)_2$ ($Ca(OH)_2$)		
III	E_2O_3 (Al_2O_3)	$Me(OH)_3$ ($Al(OH)_3$)		
IV	EO_2 (NO_2)			
V	E_2O_5 (N_2O_5)			
VI	EO_3 (SO_3)			
VII	E_2O_7 (Cl_2O_7)			

VALENTLIK ASOSIDA FORMULARLAR TUZISH

Me* I valentli, Ac** I valentli – hech qanday indeks qo'yilmaydi.

MeAc – $NaCl$, $KMnO_4$, $AgNO_3$;

Me I valentli, Ac II valentli – metaldan so'ng 2 indeksi qo'yiladi.

Me₂Ac – Na_2SO_4 , Li_2CO_3 , K_2MnO_4 ;

Me I valentli, Ac III valentli – metaldan so'ng 3 indeksi qo'yiladi.

Me₃Ac – Na_3PO_4 , Li_3BO_3 , Cs_3AsO_3 ;

Me II valentli, Ac I valentli – kislota qoldig'idan so'ng 2 indeksi qo'yiladi.

MeAc₂ – $CaCl_2$, $Mg(ClO_4)_2$, $Fe(CN)_2$;

Me II valentli, Ac II valentli – hech qanday indeks qo'yilmaydi.

MeAc – $CaSO_4$, $MgCO_3$, CdS ;

Me II valentli, Ac III valentli – valentliklar almashtirilib indeksiga qo'yiladi.

Me₃Ac₂ – $Ca_3(PO_4)_2$, Mg_3P_2 , $Sr_3(AsO_3)_2$;

Me III valentli, Ac I valentli – kislota qoldig'idan so'ng 3 indeksi qo'yiladi.

MeAc₃ – $AlCl_3$; $Fe(CN)_3$, $Al(JO_3)_3$;

Me III valentli, Ac II valentli – valentliklar almashtirilib indeksiga qo'yiladi.

Me₂Ac₃ – $Al_2(SO_4)_3$, Cr_2S_3 , $Al_2(CO_3)_3$;

Me III valentli, Ac III valentli – hech qanday indeks qo'yilmaydi.

MeAc – $AlPO_4$, $FeAsO_3$, AlN ;

KIMYODA QO'LLANILADIGAN O'ZGARMAS KATTALIKLAR

$N_A = 6,02 \cdot 10^{23}$	Avogadro soni
$m_e = 9,11 \cdot 10^{-31} \text{ kg}$	Elektronning absolyut massasi
$q_e = 1,6 \cdot 10^{-19} \text{ Kl}$	Elektronning kulonlardagi zaryadi
$1 \text{ u.b.} = 1,66 \cdot 10^{-24} \text{ g}$	u.b. = uglerod birligi
$F = 96500 \text{ Kl}$	Faradey soni
$R = 8,314$	Universal gaz doimiysi
$T_0 = 273 \text{ K}$	Normal sharoitdagi absolyut temperatura
$P_0 = 101,325 \text{ KPa}$	Normal sharoitdagi bosim
$V_M = 22,4 \text{ L}$	Gazning molyar hajmi

KIMYOVIY TENGLAMALAR BOYICHA PROPORTSIYALAR TUZISH*



*proportsiyada ayni birlik tagidan o'sha birlik yozilishiga e'tibor beriladi.

ELEMENTLARNING ELEKTRON KONFIGURATSIYASINI TUZISH

$1s^2 2s^2 2p^6 3s^2 3p^6 4s^2 3d^{10} 4p^6 5s^2 4d^{10} 5p^6 6s^2 4f^{14} 5d^{10} 6p^6 7s^2 5f^{14} 6d^{10} 7p^6$

KIMYODA QO'LLANILADIGAN BELGILASHLAR

Belgisi	Ma'nosi	Birligi*	Belgisi	Ma'nosi	Birligi
m	Massa	[g, kg, t, mg]	D_X	Biror gazning X gazga nisbatan zichligi	
V	Hajm	[l, ml, m ³]	t	Selsiy shkalasida temperatura	[°C]
ρ	Zichlik	[g/ml]	T	Absolyut temperatura	[K]
n	Modda miqdori	[mol]	P	Bosim	[kPa, atm, sim.ustuni]
A_r	Nisbiy atom massa	[m.a.b]	ω	Massa ulush	[%]
M_r	Nisbiy molekulyar massa	[m.a.b]	φ	Hajmiy ulush	[%]
M	Molyar massa	[g/mol]	v	Reaksiya tezligi	[mol/l·sek]
τ	Reaksiya vaqtি	[sek, min, soat]	γ	Reaksiya tezligining temperatura koeffitsienti	
K_M	Muvozanat konstantasi		C_M	Molyar konsentratsiya	[mol/l, M]
C_N	Normal konsentratsiya	[N, mol-ekv/l]	C_m	Molyal konsentratsiya	[m]
T	Eritma titri	[g/ml]	K_D	Dissosilanish konstantasi	
α	Dissosilanish darajasi	[%]	pH	Vodorod ko'rsatkich	
k	Moddaning elektrokimyoiy ekvivalenti	[mg/Kl]	q	Tok miqdori	[Kl]
t	Elektroliz vaqtি	[sekund]	I	Tok kuchi	[A]
E	Kimyoviy ekvivalent		η	Reaksiya unumi	[%]
Q	Issiqlik miqdori	[J]	ΔH	Entalpiya o'zhgarishi	[KJ/mol]

* birliklarning asosiy qo'llaniladigani qalin shriftda berilgan

KIMYOVİY REAKSİYA TENGLAMALARINI TUZISH

Kimyoviy reaksiyalar tenglamalarini tuzishda quyidagi hollar inobatga olinadi:

- Birikish reaksiyalarida oddiy moddalar o'zaro valentliklariga muvofiq murakkab moddani hosil qiladi:



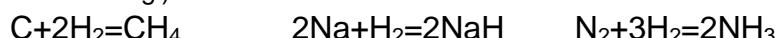
- Oddiy moddalar kislroroda yonganda metallardan faqat 1 ta oksid(Na va K da peroksid va nadperoksid) hosil bo'ladi:



- Metalmaslar yonganda kislrod kam bo'lsa quiy oksid(chala yonish), kislrod mo'l bo'lsa yuqori oksid(to'la yonish) hosil bo'ladi:



- Vodorod bilan ko'pchilik metalmaslar va faqat ishqoriy/ishqoriy-yer metallari kirishib gidridlar hosil qiladi. (EH_x -x element valentligi):



- Suvdan odatdagি sharoitda faqat ishqoriy va ishqoriy-yer metallari vodorodni siqib chiqarib tegishli asoslarni hosil qiladi. Qolgan metallar odatdagи sharoitda suv bilan ta'sirlashmaydi(erimaydi):



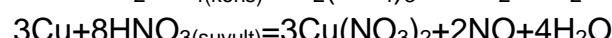
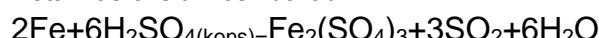
- Qolgan metal va metalmaslar faqat qizdirilgandagina suv bilan ta'sirlashib tegishli oksid va vodorodni hosil qiladi:



- Aktiv metallar (elektrod potensiallar qatorida H_2 dan chapda joylashgan) kislotalar bilan ta'sirlashganda vodorodni siqib chiqaradi va tegishli tuzlar hosil bo'ladi:



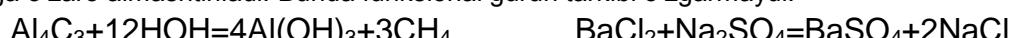
- Nitrat kislotadan va kons. H_2SO_4 dan hech qaysi metal vodorodni siqib chiqara olmaydi. Bunday reaksiyalarda tuz, suv va metalmas oksidi hosil bo'ladi:



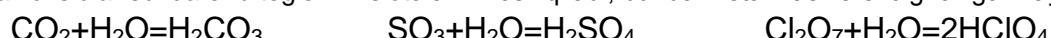
- O'rın olish reaksiyalarida tegishli metal yoki metalmasning o'rni boshqasiga almashinadi:



- Almashinish reaksiyalarini tenglamalarini tuzishda musbat qism musbat qismga, manfiy qism ham manfiya o'zaro almashtiriladi. Bunda funksional guruh tarkibi o'zgarmaydi:



- Kislotali oksidlar suvda erib tegishli kislotalarni hosil qiladi, bunda metalmas valentligi o'zgarmaydi:



- Asosli oksidlardan faqat ishqoriy va ishqoriy yer metallari odatdagи sharoitda suvda erib tegishli ishqorlarni hosil qiladi:



- Murakkab modda yonganda (oksidlanganda) uning tarkibiy qismlari yonib tegishli oksidlarni hosil qiladi:



Metallar elektrod potensiallar qatori

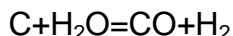
Li, K, Ba, Ca, Na, Mg, Al, Mn, Zn, Cr, Fe, Cd, Co, Ni, Sn, Pb (H₂) Cu, Hg, Ag, Pt, Au

Aktivlik kamayadi

KIMYOVIY MASALALAR YECHISHDA ENG KO'P UCHRAYDIGAN REAKSIYALAR

TENGLAMALARI

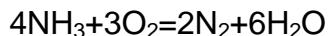
1. Suv gazining hosil bo'lishi:



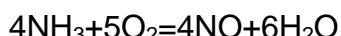
2. Kaliy permanganatning parchalish reaksiysi:



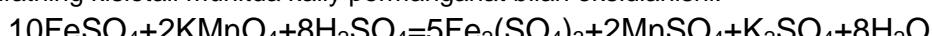
3. Ammiakning katalizatorsiz oksidlanishi:



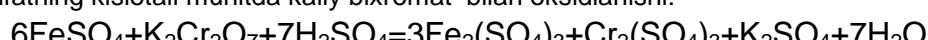
4. Ammiakning katalizator (Fe/Pt) ishtirokida oksidlanishi:



5. Temir(II)sulfatning kislotali muhitda kaliy permanganat bilan oksidlanishi:



6. Temir(II)sulfatning kislotali muhitda kaliy bixromat bilan oksidlanishi:



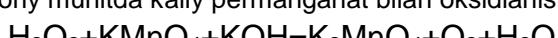
7. Xlorning sovuq ishqor eritmasida erishi:



8. Xlorning qaynoq ishqor eritmasida erishi:



9. Vodorod peroksidning ishqoriy muhitda kaliy permanganat bilan oksidlanishi:



10. Ozonning kaliy yodid eritmasi bilan ta'siri:



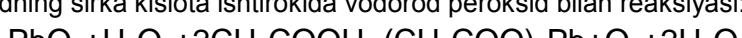
11. Oltinning "zar suvi" ("shox arog'i")da erishi:



12. Oltinning "zar suvi" ("shox arog'i")da erishi (kompleks kislota hosil bo'lishi):



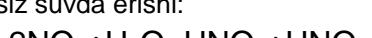
13. Qo'rg'oshin(IV)oksidning sirka kislota ishtirokida vodorod peroksid bilan reaksiysi:



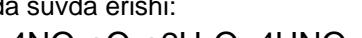
14. Natriy okslataning sulfat kislota ishtirokida kaliy permanganat bilan oksidlanishi:



15. Azot(IV)oksidning kislород ishtirokisiz suvda erishi:



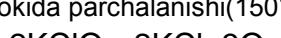
16. Azot(IV)oksidning kislород ishtirokida suvda erishi:



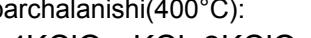
17. Metanolning sulfat kislota ishtirokida kaliy permanganat bilan oksidlanishi:



18. Kaliy xloratning MnO₂ katalizatori ishtirokida parchalanishi (150°C):



19. Kaliy xloratning katalizatorsiz termik parchalanishi (400°C):



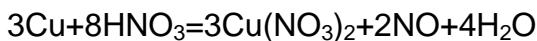
20. Ammoniy bixromatning parchalanishi:



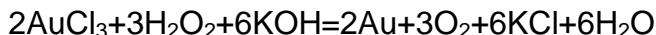
21. Misning konsentrangan nitrat kislotada erishi:



22. Misning suyultirilgan nitrat kislotada erishi:



23. Oltin(III)xloridning ishqoriy muhitda vodorod peroksid bilan ta'siri:



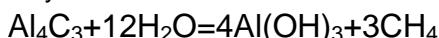
24. Oltin(III)xloridning ishqoriy muhitda metanal bilan ta'siri:



25. Glyukozaning spirtli bijg'ishi:



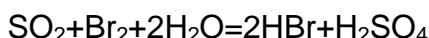
26. Alyuminiy karbidning gidrolizi reaksiyasi:



27. Kalsiy karbidning gidrolizi reaksiyasi:



28. Oltingugurt(VI)oksidning bromli suv bilan oksidlanishi:



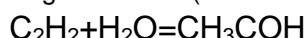
29. Kaliy permanganatning xlorid kislota bilan ta'siri:



30. Marganes(IV)oksidning xlorid kislotada erishi:



31. Atsetilenning Hg^{2+} katalizatori ishtirokida gidratlanishi (Kucherov reaksiyasi):



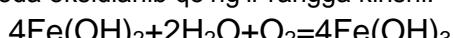
32. Metanning 1500°C da pirolizi reaksiyasi:



33. Piritni kuydirish reaksiyasi:



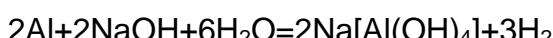
34. Temir(II)gidroksidning nam havoda oksidlanib qo'ng'ir rangga kirishi:



35. Kalsiy fosfatdan fosfor olish reaksiyasi:



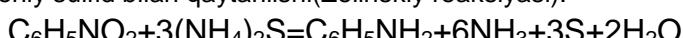
36. Alyuminiyning ishqor eritmasida erishi va tetragidroksoalyuminat tuzi hosil bo'lishi (H_2 ajralishi bilan boradi):



37. Metilaminning yonish reaksiyasi:



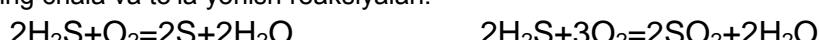
38. Nitrobenzolning ammoniy sulfid bilan qaytarilishi (Zelinskiy reaksiyasi):



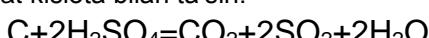
39. Sulfit kislota/oltingugurt(IV)oksid va vodorod sulfidning o'zaro ta'siridan cko'kma-S hosil bo'lishi:



40. Vodorod sulfidning chala va to'la yonish reaksiyalari:



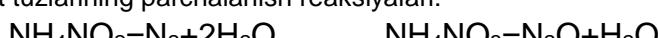
41. Uglerodning konsentrangan sulfat kislota bilan ta'siri:



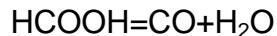
42. Mis(II)oksidning ammiak ishtirokida qaytarilishi:



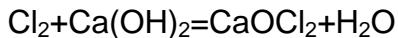
43. Ammoniy nitrit va nitrat tuzlarining parchalanish reaksiyalari:



44. Chumoli kislotaning konsentrangan sulfat kislota ta'siridan parchalanishi:



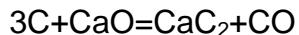
45. Xlorli ohakning hosil bo'lish reaksiyasi:



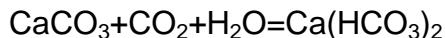
46. Kaliy bixromatning xlorid kislotada erishi:



47. Kalsiy karbidning olinishi:



48. Kalsiy karbonat suspenziyasi orqali CO_2 o'tkazilganda eritma tiniqlashishi:



49. Metal nitratlarning parchalanishi qonuniyati:

49.1. Me kuchlanishlar qatorida Mg dan chapda:



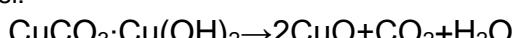
49.2. Me kuchlanishlar qatorida Mg bilan Cu orasida:



49.3. Me kuchlanishlar qatorida Cu dan o'ngda:



50. Malaxitning parchlanish reaksiyasi:



51. Fosforning suyultirilgan nitrat kislotada erishi:



52. Fosforning konsentrangan nitrat kislotada erishi:



IZOTOP ARALASHMALARIDAN MOLEKULA KOMBINATSIYALARINI HISOBBLASH JADVALI*

Izotop Indeks	1	2	3	4
1	1	2	3	4
2	1	3	6	10
3	1	4	10	20
4	1	5	15	35

***Izoh.** Masalan, bizdan uchta kislorod izotoplariidan ^{16}O , ^{17}O va ^{18}O hosil bo'lishi mumkin bo'lган ozon molekulasi kombinatsiyalarini so'ralsin. Izotop qatoridan 3 va ozon indeksidagi 3 kesishgan joyda 10 kombinatsiya soni turibdi.

Murakkab moddada element kombinatsiyalari o'zaro ko'paytiriladi. Masalan, 3 ta vodorod va 2 ta kislorod izotopidan hosil bo'lган suv molekulasi kombinatsiyalari soni $6 \times 2 = 12$ ga teng.

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KIMYOVIY MASALALAR YECHISH FORMULALARI

1	Atom/molekulaning absolyut massasini toppish	$m_a = A_r \cdot 1,66 \cdot 10^{-24} \text{ g}$
2	Modda miqdorini hisoblash(massa bo'yicha)	$n = \frac{m}{M}$
3	Modda miqdorini hisoblash(hajm bo'yicha)	$n = \frac{V}{22,4}$
4	Modda miqdorini hisoblash(molekula soni bo'yicha)	$n = \frac{N}{N_A}$
5	Modda zichligini topish	$\rho = \frac{m}{V}$
6	Birlashgan gaz qonuni	$\frac{P_0 V_0}{T_0} = \frac{P_1 V_1}{T_1}$
7	Ideal gazning holat trnglamasi	$PV = nRT$
8	Mendeleyev-Klapeyron tenglamasi	$PV = \frac{m}{M} RT$ $PM = \rho RT$
9	Gazlar aralashmasining hajmiy ulushini topish	$\bar{M} = \frac{M_1 x + M_2 (100 - x)}{100}$
10	Oddiy moddaning ekvivalentini topish	$E = \frac{A_r}{valentlik}$
11	Murakkab moddaning ekvivalentini topish	$E_{Oksid} = \frac{M_{Oksid}}{Um val}$ $E_{Asos} = \frac{M_{Asos}}{Val_{Metal}}$ $E_{Kislota} = \frac{M_{K-ta}}{Kislota negizi}$ $E_{Tuz} = \frac{M_{Tuz}}{Um val}$
12	Kimyoviy reaksiya tezligi	$v = \pm \frac{\Delta c}{\Delta t} = \pm \frac{\Delta n}{V \Delta t}$
13	Kimyoviy reaksiya tezligiga konsentratsiya ta'siri	$v = k \cdot C_A \cdot C_B$
14	Kimyoviy reaksiya tezligiga bosim ta'siri	$v = k \cdot P_A \cdot P_B$
15	Vant-Goff qoidasi	$\frac{v_{t_2}}{v_{t_1}} = \gamma^{\frac{t_2 - t_1}{10}}$ $\frac{v_{t_2}}{v_{t_1}} = \frac{t_1}{t_2}$
16	Kimyoviy muvozanat konstantasi	$K_M = \frac{C_C^c \cdot C_D^d}{C_A^a \cdot C_B^b}$
17	Eritma foiz konsentratsiyasini hisoblash	$\omega = \frac{m_{modda}}{m_{eritma}} \cdot 100\%$
18	Molyar konsentratsiyani hisoblash	$C_M = \frac{n}{V}$
19	Normal konsentratsiyani hisoblash	$C_N = \frac{m}{E V}$

20	Foiz konsentratsiyadan molyarlikka o'tish	$C_M = \frac{10 \omega \rho}{M}$
21	Foiz konsentratsiyadan normalikka o'tish	$C_N = \frac{10 \omega \rho}{E}$
22	Ervchanlik va massa ulush bog'liqligi	$\omega = \frac{S}{S + 100} 100\%$
23	Eritma titrini hisoblash	$T = \frac{N E}{1000}$
24	Dissosilanish darajasini hisoblash	$\alpha = \frac{n}{N} 100\%$
25	Ostvaldning suyultirish qonuni	$K_D = \frac{\alpha^2 c}{1 - \alpha}$ $\alpha = \sqrt{\frac{K_D}{c}}$
26	Vodorod ko'rsatkichni hisoblash	$pH = -\lg[H^+]$
27	Faradeyning 1-qonuni	$m = \frac{E I t}{F}$

ANORGANIK KIMYODA SIFAT REAKSIYALAR

Kationlar				
No	Kation	Reagent	Reaksiya tenglamasi	Kuzatilayotgan hodisa
1	Ag^+	Cl^-	$\text{Ag}^+ + \text{Cl}^- = \text{AgCl} \downarrow$	Oq cho'kma
2	Ba^{2+}	SO_4^{2-}	$\text{Ba}^{2+} + \text{SO}_4^{2-} = \text{BaSO}_4 \downarrow$	Oq cho'kma
3	Fe^{2+}	$[\text{Fe}(\text{CN})_6]^{3-}$	$\text{Fe}^{2+} + [\text{Fe}(\text{CN})_6]^{3-} = \text{Fe}_3[\text{Fe}(\text{CN})_6]_2$	Toq ko'k cho'kma (Turnbul ko'ki)
4	Fe^{3+}	$[\text{Fe}(\text{CN})_6]^{4-}$	$\text{Fe}^{3+} + [\text{Fe}(\text{CN})_6]^{4-} = \text{Fe}_4[\text{Fe}(\text{CN})_6]_3$	Toq ko'k cho'kma (Berlin zangorisi)
5	H^+ (k-ta)	Metiloranj	Indikator rangi sariqdan qizil rangga o'tadi	
		Lakmus	Eritma rangi qizilga o'zgaradi	
6	NH_4^+	Ishqorlar	$\text{NH}_4^+ + \text{OH}^- = \text{NH}_3 \uparrow + \text{H}_2\text{O}$	Ammiak hidi
Anionlar				
1	Cl^-	Ag^+ (AgNO_3)	$\text{Ag}^+ + \text{Cl}^- = \text{AgCl} \downarrow$	Oq cho'kma
2	CO_3^{2-}	Kislotalar	$\text{CO}_3^{2-} + 2\text{H}^+ = \text{CO}_2 \uparrow + \text{H}_2\text{O}$	Gaz ajraladi
3	OH^- (ishqor)	Fenolftalein	Indikator rangi malina rangga o'tadi	
		Lakmus	Eritma rangi ko'kka o'zgaradi	
4	S^{2-}	Pb^{2+}	$\text{Pb}^{2+} + \text{S}^{2-} = \text{PbS} \downarrow$	Qora cho'kma
5	SO_4^{2-}	Ba^{2+} (BaCl_2)	$\text{Ba}^{2+} + \text{SO}_4^{2-} = \text{BaSO}_4 \downarrow$	Oq cho'kma
Anorganik moddalar				
1	CO_2	$\text{Ca}(\text{OH})_2$ eritmasi (ohakli suv)	$\text{CO}_2 + \text{Ca}(\text{OH})_2 = \text{CaCO}_3 \downarrow + \text{H}_2\text{O}$ yana davom ettirilsa $\text{CaCO}_3 + \text{CO}_2 + \text{H}_2\text{O} = \text{Ca}(\text{HCO}_3)_2$	Avval oq cho'kma, keyin yana eriydi
2	NH_3	HCl	$\text{NH}_3 + \text{HCl} = \text{NH}_4\text{Cl}$	Oq tutun

ATOM ORBITALLARINING GIBRIDLANISHI

Nº	Gibridlanish turi	Xarakteristik belgisi	Molekula fazoviy tuzilishi	Molekula fazoviy shakli	Misollar
Standart gibridlanish holati					
1	sp	Faqat 2 ta σ bog' VB 180°	Chiziqli		Alkinlar, $BeCl_2$, CO_2 , (CO , N_2), HCN
2	sp^2	Faqat 3 ta σ bog' VB 120°	Uchburchakli		Alkenlar va diyenlar qo'shbog' uglerodi, BF_3 , $AlCl_3$, SO_3 , NO_3^- , CO_3^{2-}
3	sp^3	Faqat 4 ta σ bog' VB $109^\circ 28'$	Tetraedr		CH_4 va alkanlar, NH_4^+ , SO_4^{2-} , PO_4^{3-} , ClO_4^- , MnO_4^{2-} , MnO_4^- , SiO_2
4	sp^3d	Faqat 5 ta σ bog' VB $3 \times 120^\circ$ va $2 \times 90^\circ$	Trigonal bipiramida		PCl_5 , SbF_5 , XeO_3F_2
5	sp^3d^2	Faqat 6 ta σ bog' VB $6 \times 90^\circ$	Oktaedrik		SF_6 , XeO_2F_4
Nostandart gibridlanish holati					
6	sp^3	3 ta σ bog' va 1 ta taqsimlanmagan elektron jufti VB $107,3^\circ$	Uchburchakli piramida		NH_3 , H_3O^+ , ClO_3^- , XeO_3
7	sp^3	2 ta σ bog' va 2 ta taqsimlanmagan elektron jufti VB $104,5^\circ$	Burchakli		H_2O , ClO_2^- , XeO_2
8	sp^3	1 ta σ bog' va 3 ta taqsimlanmagan elektron jufti VB 100°	Chiziqli		Galogenovodorodlar $H\Gamma$ ($\Gamma=F$, Cl , Br , J)
9	sp^2	2 ta σ bog' va 1 ta taqsimlanmagan elektron jufti VB 120°	Burchakli		SO_2

ORGANIK BIRIKMALAR KLASSEFIKATSİYASI

Nº	Sinf nomi		Umumiyl formulasi	Xarakteristik belgisi	Sifat reaksiya	Gibridlaniishi
1	C_nH_m	Alkanlar	C_nH_{2n+2}	Faqat σ bog'li to'yigan ochiq zanjirli		sp^3
2		Alkenlar	C_nH_{2n}	1 ta qo'shbog'li (=) ochiq zanjirli	Br_2 li suv	sp^2
3		Alkadiyenlar	C_nH_{2n-2}	2 ta qo'shbog'li (=) ochiq zanjirli	Br_2 li suv	sp^2
4		Alkinlar	C_nH_{2n-2}	1 ta uchbog'li (≡) ochiq zanjirli	Ag_2O/NH_3	sp
5		Arenlar	C_nH_{2n-6}	Aromatik xalqali yopiq zanjirli		sp^2
6		Sikloalkanlar	C_nH_{2n}	Faqat σ bog'li to'yigan yopiq zanjirli		sp^3
7	Spirtlar		R-OH	OH guruh alkil radikaliga bog'langan		sp^3
	a) Bir aromli Oddiy efirlar		R-OH	1 ta OH guruh alkil radikaliga bog'langan		sp^3
	b) Ko'p atomli		$R(OH)_n$	1 dan ortiq OH guruh alkil radikaliga bog'langan	$Cu(OH)_2$	sp^3
8	Oddiy efirlar Bir atomli spirtlar		R-O-R	Alkil radikallar -O- ko'prigi orqali bog'langan		sp^3
9	Fenollar		Ar-OH	OH guruh aromatik xalqa(benzol)ga bog'langan	$FeCl_3$	sp^2
10	Galogen hosilalar		R-Γ	Galogenlar(F, Cl, Br, I) radikalga bog'langan		sp^3
11	Karbonil birikmalar					
	a) Aldegidlar Ketonlar		$\begin{array}{c} O \\ \\ R-C-H \end{array}$	Tarkibida aldegid guruhi mavjud	Ag_2O/NH_3	sp^2
	b) Ketonlar Aldegidlar		$\begin{array}{c} O \\ \\ R-C-R' \end{array}$	Alkil yoki aromatik radikallar o'zaro -CO- ko'prigi orqali bog'langan		sp^2
12	Karboksil birikmalar					
	a) Karbon kislotalari Murakkab efirlar		$\begin{array}{c} O \\ \\ R-C-OH \end{array}$	Radikallar carboksil guruhiga bog'langan kislota xossasiga ega moddalar		sp^2
	b) Murakkab efirlar Karbon kislotalari		$\begin{array}{c} O \\ \\ R-C-OR' \end{array}$	Karbon kislota vodorodi radikalga eterifikatsiya reaksiyasi orqali almashigan		sp^2
13	Aminokislotalar		$\begin{array}{c} O \\ \\ R-CH(NH_2)-C-OH \end{array}$	Aminoguruh (NH₂) karbon kislota radikaliga bog'langan kislota hosilari		sp^2
14	Nitrobirimkalar		R-NO ₂	Nitroguruh (NO₂) radikalga bog'langan		sp^2
15	Aminlar		R-NH ₂	Ammiak vodorodi bir yoki bir nechta radikalga almashigan		sp^3
16	Uglevodlar		$C_n(H_2O)_m$	Tarkibida gidroksil (OH) va karbonil (C=O) guruhi mavjud		
17	Geterosiklik birikmalar			Xalqada ugleroddan tashqari geteroatom (N, O, S) tutgan yopiq zanjirli		sp^2

Izoh: Sinfning yuqorisida o'zaro izomer moddalar sinfi ko'rsatilgan

BA'ZI KIMYOVİY MODDALARNING TARİXIY NOMİ

1. **Akril kislota** = propen kislota –
 $\text{CH}_2=\text{CH}-\text{COOH}$
2. **Aktinoidlar** – Th-Lr
3. **Akrolein** = propenal –
 $\text{CH}_2=\text{CH}-\text{COH}$
4. **Alebastr** – $\text{CaSO}_4 \cdot 2\text{H}_2\text{O}$
5. **Alyumokaliyli achchiqtosh** –
 $\text{K}_2\text{SO}_4 \cdot \text{Al}_2(\text{SO}_4)_3 \cdot 24\text{H}_2\text{O}$
6. **Amil spirit** – $\text{C}_5\text{H}_{11}\text{OH}$
7. **Ammiakli selitra** – NH_4NO_3
8. **Angidrit** – CaSO_4
9. **Anilin** – $\text{C}_6\text{H}_5\text{NH}_2$
10. **Antixlor** – $\text{Na}_2\text{S}_2\text{O}_3 \cdot 5\text{H}_2\text{O}$
11. **Asbest** – $3\text{MgO} \cdot 2\text{SiO}_2 \cdot 2\text{H}_2\text{O}$
12. **Atsetilen** – C_2H_2
13. **Atseton** – $\text{CH}_3\text{-CO-CH}_3$
14. **Barit** – BaSO_4
15. **Berlin zangorisi** – $\text{Fe}_4[\text{Fe}(\text{CN})_6]_3$
16. **Berolle tuzi** – KClO_3
17. **Boksit** – $\text{Al}_2\text{O}_3 \cdot n\text{H}_2\text{O}$
18. **Bura** – $\text{Na}_2\text{B}_4\text{O}_7 \cdot 10\text{H}_2\text{O}$
19. **Chili selitrası** – NaNO_3
20. **Diammofos** – $(\text{NH}_4)_2\text{HPO}_4$
21. **Dolomit** – $\text{CaCO}_3 \cdot \text{MgCO}_3$
22. **Ervchan shisha** –
 $\text{K}_2\text{O} \cdot \text{CaO} \cdot 6\text{SiO}_2$
23. **Etilenglikol** = etandiol-1,2 –
 $\text{CH}_2(\text{OH})-\text{CH}_2(\text{OH})$
24. **Fenolformaldegid smolası** -
Fenol+Metanal – $\text{C}_6\text{H}_5\text{-OH}+\text{HCOH}$
25. **Flyuorit** – CaF_2
26. **Fosfin** – PH_3
27. **Fosforit** – $\text{Ca}_3(\text{PO}_4)_2$
28. **Fosgen** – COCl_2
29. **Fumar kislota** – trans
 HOOC-CH=CH-COOH
30. **Generator gazi** - $\text{CO}+\text{N}_2$
31. **Glauber tuzi** – $\text{Na}_2\text{SO}_4 \cdot 10\text{H}_2\text{O}$
32. **Glinozyom** – Al_2O_3
33. **Glitserin** = propantriol-1,2,3 –
 $\text{CH}_2(\text{OH})-\text{CH}(\text{OH})-\text{CH}_2(\text{OH})$
34. **Glitsin** = aminosırka kislota –
 $\text{NH}_2\text{-CH}_2\text{-COOH}$
35. **Glyukon kislota** – $\text{C}_6\text{H}_{12}\text{O}_7$
36. **Hind selitrası** – KNO_3
37. **İchimlik soda** – NaHCO_3
38. **Izopren** – $\text{CH}_2=\text{C}(\text{CH}_3)\text{-CH=CH}_2$
39. **Izosianat kislota** – HCNO
40. **Javel suvi** – $\text{KCl}+\text{KClO}$
41. **Kalomel** – Hg_2Cl_2
42. **Kalsit, bo'r, ohaktosh, marmar** –
 CaCO_3
43. **Kaolin** – $\text{Al}_2\text{O}_3 \cdot 2\text{SiO}_2 \cdot 2\text{H}_2\text{O}$
44. **Kapron kislota** = geksan kislota –
 $\text{C}_5\text{H}_{11}\text{-COOH}$
45. **Karbamid/mochevina** – $\text{CO}(\text{NH}_2)_2$
46. **Karbol kislota=fenol**
47. **Karborund** – SiC
48. **Kinovar** – HgS
49. **Korund** – Al_2O_3
50. **Kraxmal/sellyuloza** – $\text{C}_6\text{H}_{10}\text{O}_5$
51. **Krezollar** = metilfenollar –
 $\text{C}_6\text{H}_4(\text{CH}_3)\text{OH}$
52. **Kriolit** – $\text{Na}_3[\text{AlF}_6]$
53. **Kristall soda** – $\text{Na}_2\text{CO}_3 \cdot 10\text{H}_2\text{O}$
54. **Kroton aldegid** = buten-2-al –
 $\text{CH}_3\text{-CH=CH-COOH}$
55. **Ksilol-o'**, m' va p'= dimetilbenzollar –
 $\text{C}_6\text{H}_4(\text{CH}_3)_2$
56. **Kumol** = izopropilbenzol –
 $\text{C}_6\text{H}_5\text{-CH}(\text{CH}_3)_2$
57. **Kuporos moyi** – $\text{kons. H}_2\text{SO}_4$
58. **Kvarts/qum** – SiO_2
59. **Labborak suvi** – $\text{NaCl}+\text{NaClO}$
60. **Lantanoidlar** – Ce-Lu
61. **Lavsan** = Tereftal kislota
+Etielnglikol – $\text{HOOC-C}_6\text{H}_4\text{-COOH}+\text{CH}_2(\text{OH})-\text{CH}_2(\text{OH})$
62. **Linol kislota** – $\text{C}_{17}\text{H}_{31}\text{COOH}$

63. **Linolen kislota** - $C_{17}H_{29}COOH$
64. **Magnezit** – $MgCO_3$
65. **Magnitli temirtosh** – Fe_3O_4
66. **Malaxit** – $CuCO_3 \cdot Cu(OH)_2$
67. **Malein kislota** - sis
 $HOOC-CH=CH-COOH$
68. **Metakril kislota** –
 $CH_2=C(CH_3)-COOH$
69. **Mis kuporosi** – $CuSO_4 \cdot 5H_2O$
70. **Mor tuzi** – $(NH_4)_2Fe(SO_4)_2 \cdot 6H_2O$
71. **Nodir gazlar** – He, Ne, Ar, Kr, Xe, Rn
72. **Nodir metallar** – Ag, Au, Pt. Ru. Rh, Ir, Os
73. **Norvegiya selitrası** – $Ca(NO_3)_2$
74. **Novshadil spirit** – NH_4OH
75. **O'tish metallari** – d- va f- elementlar
76. **O'yuvchi kaliy** – KOH
77. **O'yuvchi natriy/kaustik soda** - NaOH
78. **Og'ir metallar** ($\rho > 5 \text{ g/sm}^3$) – Mn, Fe, Co, Ni, Cu, Zn, Cd, Hg, Sn, Pb
79. **Oksalat kislota** – $H_2C_2O_4$
80. **Olein kislota** – $C_{17}H_{33}COOH$
81. **Oleum** – $H_2SO_4 \cdot nSO_3$
82. **Oson suyuqlanuvchan metallar**
 $(t_s < 1000^\circ C)$ – Li-Cs, Mg,Ca, Al
83. **Palmitin kislota** – $C_{15}H_{31}COOH$
84. **Pikrin kislota** = 2,4,6-trinitrofenol -
 $C_6H_2(OH)(NO_2)_3$
85. **Piridin** - C_5H_5N
86. **Pirit** – FeS_2
87. **Pirogallol** =trigidroksibenzol –
 $C_6H_3(OH)_3$
88. **Pirolyuzit** – MnO_2
89. **Plavik kislota** – HF
90. **Qaldiroq gaz** – $2H_2+O_2$
91. **Qiyin suyuqlanuvchan metallar**
 $(t_s > 1000^\circ C)$ – Ti, Ir, Hf, W, Nb, Ta, Cr, Re
92. **Qizil qon tuzi** – $K_3[Fe(CN)_6]$
93. **Qo'rg'oshin shakari** –
 $Pb(CH_3COO)_2 \cdot 3H_2O$
94. **Qora metallar** – Fe va uning qotishmalari
95. **Quruq muz** – CO_2
96. **Rangli metallar** – Ag, Au, Cu, Mn, Co, Ni
97. **Rux aldamasi** – ZnS
98. **Sariq qon tuzi** – $K_4[Fe(CN)_6]$
99. **Saxaroza/maltoza/laktoza** –
 $C_{12}H_{22}O_{11}$
100. **Shisha** – $Na_2O \cdot CaO \cdot 6SiO_2$

MUNDARIJA

II KITOB

ANORGANIK KIMYO REAKSIYALARDA

Vodorod(3). Gidridlar(3). Suv(3). Ftor(3). Vodorod ftorid(4). Xlor(4). Vodorod xlorid(4). Xloring kislородли биримлари(4). Brom va yod(5). Kislород(5). Ozon(6). Vodorod peroksid(6). Oltingugurt(6). Vodorod sulfid(6). Oltingugurt(IV)oksid(7). Oltingugurt(VI)oksid(7). Sulfat kislota(7). Tiosulfat kislota(8). Selenat kislota(8). Azot(8). Ammiak(8). Gidrazin(9). Gidroksilamin(9). Azot(II)oksid(9). Nitrit kislota(9). Azot(III)oksid(9). Azot(IV)oksid(9). Azot(IV)oksidi dimeri(10). Nitrat kislota(10). Fosfor(10). Fosfin(11). Fosfor(III)xlorid(11). Fosfor(V)xlorid(11). Fosfit kislota(11). Fosfor(V)oksid(11). Ortofosfat kislota(12). Fosforli o'g'itlar(12). Azot saqlovchi o'g'itlar(12). Uglerod(12). Uglerod(II)oksid(12). Uglerod(IV)oksid(13). Uglerod disulfid(13). Galogenidlari(13). Uglerodning azotli birimlari(13). Karbidlar(14). Karbonat kisota(14). Karbonatlar(14). Soda ishlab chiqarish(14). Kremniy(14). Silan(15). Kremniyning galogenidlari(15). Shisha(15). Metallarning olinish usullari(15). Ishqoriy metallar(15). Litiy(15). Natriy(16). Kaliy(16). Berilliyl(16). Magniy(17). Kalsiy(17). Kalsiy birimlari(17). Suvning qattiqligi va uni yo'qotish usullari(17). Bor(17). Alyuminiy(18). Germaniy(18). Qalay(18). Qo'rg'oshin(18). Surma va vismut(19). Temir(19). Xrom(20). Molibden(20). Volfram(20). Marganets(20). Texnetsiy va reniy(21). Rux(21). Simob(21). Mis(21). Kumush(22). Oltin(22). Titan(22).

ORGANIK KIMYO.

I BOB. ORGANIK BIRIKMALAR TUZILISHI NAZARIYASI VA ORGANIK REAKSIYALARING TURLARI.

Organik kimyo faniga kirish(24). Organik birimlarning klasifikatsiyasi(25). Organik reakiyalarning turlari(25).

II BOB. UGLEVODORODLAR.

Uglevodorodlar(27). Alkanlar(27). Sikloalkanlar(29). Alkenlar(31). Alkadiyenlar. Diyen uglevodorodlari(33). Alkinlar. Atsetilen qatori uglevodorodlari(36). Neftni qayta ishlash. Kreking jarayoni(38). Aromatik ugelvodorodlar – Arenlar(39).

III BOB. KISLOROD SAQLOVCHI ORGANIK BIRIKMALAR.

Spirtlar(44). Ko'p atomli spirtlar(47). Oddiy efirlar(49). Fenollar(50). Karbonil birimlalar(53). Aldegidlar(53). Ketonlar(55). Karbon kislotalar(56). Murakkab efirlar(). Yog'lar(21). Uglevodlar(22). Monosaxaridlar(23). Disaxaridlar-C₁₂H₂₂O₁₁(24). Polisaxaridlar-(C₆H₁₀O₅)_n(25).

IV BOB. AZOT SAQLOVCHI ORGANIK BIRIKMALAR.

Aminlar(26). Aminokislotalar(27). Oqsillar(28). Geterosiklik birimlalar(29). Nuklein kislotalar(30).

FOYDALANILGAN ADABIYOTLAR RO'YXATI(76).

ILOVALAR.

Valentlik(77). Anorganik birimlarning asosiy sinflari(79). Valentlik asosida formulalar tuzish(79). Kimyoda qo'llaniladigan o'zgarmas kattaliklar(80). Kimyoviy tenglamalar bo'yicha proportsiyalar tuzish(80). Elementlarning elektron konfiguratsiyasini tuzish(80). Kimyoda qo'llaniladigan belgilashlar(81). Kimyoviy reaksiya tenglamalarini tuzish(82). Kimyoviy masalalar yechishda eng ko'p uchraydigan reaksiyalar tenglamalari(83). Izotop aralashmalaridan molekula kombinatsiyalarini hisoblash jadvali*(85). Kimyoviy masalalar yechish formulalari(86). Anorganik

kimyoda sifat resksiyalar(87). Atom orbitallarining gibridlaniши(88). Organik birikmalar klassifikatsiyasi(89). Ba'zi kimyoviy moddalarning tarixiy nomi(90).