APPENDIX-A

A Table of Greek letters

Upper case	Lower case	In English
A	α	alpha
В	β	beta
Γ	γ	gamma
Δ	δ	delta
Е	ε	epsilon
Z	ζ	zeta
Н	η	eta
Θ	θ	theta
I	ι	iota
K	κ	kappa
Λ	λ	lambda
M	μ	mu
N	υ	nu
Ξ	ξ	csi
0	0	omicron
П	π	pi
P	ρ	rho
Σ	σ	sigma
T	τ	tau
Υ	υ	upsilon
Φ	ф	phi
Ψ	Ψ	psi
X	χ	chi
Ω	ω	omega

APPENDIX-B

Useful Physical Constant in Chemistry

Constant	Symbol	Value
Acceleration due to gravity	g	9.8 m s ⁻²
Atomic mass unit	amu, m _u or u	$1.66 \times 10^{-27} \text{ kg}$
Avogadro's Number	N, N _A	$6.022 \times 10^{23} \text{mol}^{-1}$
Bohr radius	r _o	0.529×10^{-10} m
Boltzmann constant	k	$1.38 \times 10^{-23} \mathrm{J \ K^{-1}}$
Electron charge to mass ratio	$\frac{-e}{m_e}$	$-1.7588 \times 10^{11} \text{ C kg}^{-1}$
Electron classical radius	r _e	$2.818 \times 10^{-15} \mathrm{m}$
Electron mass energy(J)	m _e c ²	$8.187 \times 10^{-14} \mathrm{J}$
Electron mass energy (MeV)	$m_e c^2$	0.511 MeV
Electron rest mass	m _e	$9.109 \times 10^{-31} \mathrm{kg}$
Faraday constant	F	$9.649 \times 10^{4} \text{Cmol}^{-1}$
Fine-structure constant	α	7.297×10^{-3}
Gas constant	R	8.314 J mol ⁻¹ K ⁻¹
Gravitational constant	G	$6.67 \times 10^{-11} \mathrm{Nm^2 Kg^{-2}}$
Neutron mass energy	m _n c ²	1.505 × 10 ⁻¹⁰ J or 939.565 MeV
Neutron rest mass	m _n	$1.675 \times 10^{-27} \mathrm{kg}$

Neutron-electron mass ratio	$\frac{\mathrm{m_n}}{\mathrm{m_e}}$	1838.68
Neutron-proton mass ratio	$\frac{m_n}{m_p}$	1.0014
Permeability of a vacuum	μ_0	$4\pi \times 10^{-7} \text{NA}^{-2}$
Permittivity of a vacuum	ϵ_0	$8.854 \times 10^{-12} \mathrm{F m^{-1}}$
Planck constant	h	$6.626 \times 10^{-34} \mathrm{J \ s}$
Proton mass energy	$m_p c^2$	1.503 × 10 ⁻¹⁰ J or 938.2 <mark>72</mark> MeV
Proton rest mass	m_p	$1.6726 \times 10^{-27} \mathrm{kg}$
Proton-electron mass ratio	$\frac{m_p}{m_e}$	1836.15
Rydberg constant	R_{e}	$1.0974 \times 10^7 \mathrm{m}^{-1}$
Speed of light in vacuum	c	$2.9979 \times 10^8 \mathrm{m/s}$

APPENDIX-C

Most Popular Chemists and their Contributions AMEDEO AVOGADRO 1776 – 1856

The first scientist to realize that elements could exist in the form of molecules rather than as individual atoms; originator of Avogadro's law.

JACOB BERZELIUS 1779 – 1848

A founder of modern chemistry: the first person to measure accurate atomic weights for the chemical elements; discovered three elements: cerium, thorium and selenium; devised the modern symbols for elements; described how chemical bonds form by electrostatic attraction.

NIELS BOHR 1885 – 1962

Founded quantum mechanics when he remodeled the atom that electrons occupied 'allowed' orbits around the nucleus while all other orbits were forbidden; architect of the Copenhagen interpretation of quantum mechanics.

ROBERT BOYLE 1627 – 1691

Transformed chemistry from a field bogged down in alchemy and mysticism into one based on measurement. He defined elements, compounds, and mixtures; and he discovered the first gas law – Boyle's Law.

LAWRENCE BRAGG 1890 - 1971

Discovered how to locate the positions of atoms in solids using X-ray diffraction, enabling scientists to build 3D models of the atomic arrangements in solids. The discovery was arguably the most significant experimental breakthrough of twentieth century science.

HENNIG BRAND 1630 – 1710

Discovered phosphorus, becoming the first named person in history to discover a chemical element.

GEORG BRANDT 1694 – 1768

The first named person in history to discover a new metal – cobalt; was one of the first scientists to condemn alchemy, publicly demonstrating tricks used by alchemists to make people think they could make gold.

ROBERT BUNSEN 1811 – 1899

Discovered cesium and rubidium; discovered the antidote to arsenic poisoning; invented the zinc-carbon battery and flash photography; discovered how geysers operate.

ERWIN CHARGAFF 1905 – 2002

Chargaff's rules paved the way to the discovery of DNA's structure.

MARIE CURIE 1867 - 1934

Co-discovered the chemical elements radium and polonium; made numerous pioneering contributions to the study of radioactive elements; carried out the first research into the treatment of tumors with radiation.

JOHN DALTON 1766 – 1844

Dalton's Atomic Theory is the basis of chemistry; discovered Gay-Lussac's Law relating gases' temperature, volume, and pressure; discovered the law of partial gas pressures.

DEMOCRITUS (C. 460 — C. 370 BC)

Devised an atomic theory featuring tiny particles always in motion interacting through collisions; advocated a universe containing an infinity of diverse inhabited worlds governed by natural, mechanistic laws rather than gods; deduced that the light of stars explains the Milky Way's appearance; discovered that a cone's volume is one-third that of the cylinder with the same base and height.

EMPEDOCLES (C. 490 - C. 430 BC)

An ancient theory of natural selection; mass conservation; and the four elements which are now often misattributed to Aristotle.

MICHAEL FARADAY 1791 - 1867

Discovered electromagnetic induction; devised Faraday's laws of electrolysis; discovered the first experimental link between light and magnetism; carried out the first room-temperature liquefaction of a gas; discovered benzene.

ROSALIND FRANKLIN 1920 - 1958

Provided much of the experimental data used to establish the structure of DNA; discovered that DNA can exist in two forms; established that coal acts as a molecular sieve.

WILLARD GIBBS 1839 - 1903

Gibbs invented vector analysis and founded the sciences of modern statistical mechanics and chemical thermodynamics.

GEORGE DE HEVESY 1885 – 1966

Discovered element 72, hafnium. Pioneered isotopes as tracers to study chemical and biological processes; discovered how plants and animals utilize particular chemical elements after they are taken in as nutrients.

FRED HOYLE 1915 - 2001

Established that most of the naturally occurring elements in the periodic table were made inside stars and distributed through space by supernova explosions.

IRENE JOLIOT-CURIE 1897 – 1956

Co-discovered how to convert stable chemical elements into 'designer' radioactive elements; these have saved millions of lives and are used in tens of millions of medical procedures every year.

MARTIN KLAPROTH 1743 – 1817

Discovered the chemical elements uranium, zirconium, and cerium – naming the first two of these elements; verified the discoveries of titanium, tellurium and strontium, again naming the first two.

STEPHANIE KWOLEK 1923 – 2014

Invented kevlar, the incredibly strong plastic used in applications ranging from body armor to tennis racquet strings.

ANTOINE LAVOISIER 1743 – 1794

A founder of modern chemistry; discovered oxygen's role in combustion and respiration; discovered that water is a compound of hydrogen and oxygen; proved that diamond and charcoal are different forms of the same element, which he named carbon.

ERNEST LAWRENCE 1901 – 1958

Invented the cyclotron, used by scientific teams in his laboratories to discover large numbers of new chemical elements and isotopes. Founded big science.

IANE MARCET 1769 – 1858.

Author of *Conversations on Chemistry*, a unique textbook for its time written for people with little formal education, such as girls and the poor. The book inspired Michael Faraday to overcome his poor origins to become a great scientist.

DMITRI MENDELEEV 1834 – 1907

Discovered the periodic table in a dream. Utilized the organizing principles of the periodic table to correctly predict the existence and properties of six new chemical elements.

HENRY MOSELEY 1887 – 1915

Proved that every element's identity is uniquely determined by its number of protons, establishing this is the true organizing principle of the periodic table; correctly predicted the existence of four new chemical elements; invented the atomic battery.

GIULIO NATTA 1903 - 1979

Discovered how to produce polymer chains with orderly spatial arrangements – i.e. stereoregular polymers.

ALFRED NOBEL 1833 – 1896

Invented dynamite, the blasting cap, gelignite, and ballistite; grew enormously wealthy manufacturing explosives; used his wealth to bequeath annual prizes in science, literature, and peace.

HANS CHRISTIAN OERSTED 1777 – 1851

Discovered electromagnetism when he found that electric current caused a nearby magnetic needle to move; discovered piperine and achieved the first isolation of the element aluminum.

LOUIS PASTEUR 1822 – 1895

The father of modern microbiology; transformed chemistry and biology with his discovery of mirror-image molecules; discovered anaerobic bacteria; established the germ theory of disease; invented food preservation by pasteurization.

LINUS PAULING 1901 - 1994

Maverick giant of chemistry; formulated valence bond theory and electronegativity; founded the fields of quantum chemistry, molecular biology, and molecular genetics. Discovered the alpha-helix structure of proteins; proved that sickle-cell anemia is a molecular disease.

MARGUERITE PEREY 1909 – 1975

Discovered francium, the last of the naturally occurring chemical elements to be discovered – all elements since have been produced artificially.

WILLIAM PERKIN 1838 – 1907

At age 18 started the synthetic dye revolution when his discovery of mauveine brought the once formidably expensive color purple to everyone. Perkins' revolution took the world by storm, transforming textiles, foods and medicine.

C. V. RAMAN 1888 - 1970

Discovered that light can donate a small amount of energy to a molecule, changing the light's color and causing the molecule to vibrate. The color change acts as a 'fingerprint' for the molecule that can be used to identify molecules and detect diseases such as cancer.

WILLIAM RAMSAY 1852 – 1916

Predicted the existence of the noble gases and discovered or was first to isolate every member of the group; created the world's first neon light.

ERNEST RUTHERFORD 1871 – 1937

The father of nuclear chemistry and nuclear physics; discovered and named the atomic nucleus, the proton, the alpha particle, and the beta particle; discovered the concept of nuclear half-lives; achieved the first laboratory transformation of one element into another.

GLENN SEABORG 1912 TO 1999

Took part in the discovery of ten of the periodic table's chemical elements. His work on the electronic structure of elements led to the periodic table being rewritten.

Appendix 3

HERMANN STAUDINGER 1881 – 1965

Founded macromolecular chemistry when he established that molecules made of hundreds of thousands of atoms exist; demonstrated that synthetic polymers can make fibers similar to natural fibers; discovered polyoxymethylene; discovered pyrethroid natural insecticides.

J.J. THOMSON 1856 - 1940

Discovered the electron; invented one of the most powerful tools in analytical chemistry – the mass spectrometer; obtained the first evidence for isotopes of stable elements.

HAROLD UREY 1893 - 1981

Discovered deuterium; showed how isotope ratios in rocks reveal past Earth climates; founded modern planetary science; the Miller-Urey experiment demonstrated that electrically sparking simple gases produces amino acids – the building blocks of life.

ALESSANDRO VOLTA 1745 – 1827

Pioneer of electrical science; invented the electric battery; wrote the first electromotive series; isolated methane for the first time; discovered a methane-air mixture could be exploded using an electric spark – the basis of the internal combustion engine.

SERGEI WINOGRADSKY 1856 - 1953

Founded microbial ecology; discovered chemosynthetic life forms which obtain energy from chemical reactions rather than from sunlight; discovered nitrogen-fixing bacteria in soil that make nitrates available to green plants.

APPENDIX-D

	18	Heium	10	Ne S	Neon 20.180	18	Argon 39.948	36	Krypton	54	Xe	131.3	98	Radon 222.018	118	Oganesson (294)	71	Lutetium	103	Lr	Lawerncium (262)		
		17	6	H.	Fluorine 18.998	17	Chlorine 35.453	35	Bromine	53	Iodine	126.904	82	At Astatine 209.987	117	Ts Tennessine (294)	20	Yb	102	Š	Nobelium 259.101	7	ıme
		16	8	0	Oxygen 15.999	16	Sulphur 32.056	34	Selenium	52	Tellurium	127.6	84	Polonium 208.982	116	LV In Livermorium (293)	69	Tm	101	Md	fendelevium 258.1	objects A	
		15				_	Phosphorus	_				_		Bismutt 208.980	115	Moscovium	89	Er				Lantha-	nide
		14	9	<u>ن</u>	Carbon 12.011	14	Silicon 28.086	32	Germanium	50	Sn	118.711	82	Pb Lead 207.2	114	FI Flerovium (289)		Ho				oble	Gas
(13	5	B	Boron 10.811	13	Aluminium 26,982	31	Gallium		Indium	2	M	Ti Thallium 204.383	113	Nihonium (286)		Dy Indiana	-	_	-		
nent						12	IIB 2B	30	Zinc	48	Cd	112.414	80	Hg Mercury 200.592	112	m Copernicium N					_	Halogon	9
Elen						=	199	29	Copper	47	Age	107.9	79	Au Gold 197.0	111	Roentgenium		Terbium		Bk	_	Non-	metal
f the						10	م	28	Nickel	46	Pd	106.42	28	Pt Platinum 195.085	110	Ds Darmstadtium (281)	64	Gadolinium	96	CE	Curium 247.070		metal
ple o						6	VIIIV 8	27	Cobalt	45	Rhodium Thodium	102.906	72	Ir Iridium 192.217	109	Mt Meitherium [778]	63	Europium	95	Am	Americium 243.061		
lic Ta						œ	, (,	56	Fe	_		\rightarrow		Osmium 190.23		Hassium N	 62	Samarium	94	Pu	Plutonium 244.064	Basic	Metal
Periodic Table of the Elements						1	VÍIB 7B	25	Mn	_				Rhenium		Bohrium	19	Pm	93	aN	Veptunium 237.064	Fransition	Metal
							VÏB 6B							Tungsten 183.84		Seaborgium		Neodymium					
						ור	VB 5B	23	Vanadium	+-		-		Tantalum 180.948	105	_ =	26	Praseodymium Ne	91		я	Alkaline	Earth
							IÝB 4B	⊢				_		Hafnium 178.49		Rutherfordium (26.1)						Alkali	Metal
							JIIB 3B	\vdash		_		\rightarrow			89-103	Rt		Cerium Cerium	-	Th			
		ZA ZA	4	Be	9.012			-		_				Barium 137.328		Radium		Lanthanum	68	Ac	Actiniu 227.028		
П	11A 1A	H Hydrogen 1,008	8	i.	6.941 B	11		_		_		-			_	Francium I	anthanide	Series	:	Actinide	Series		

APPENDIX-E

Important Organic Chemical Reactions

Sandmeyer Reaction

The Sandmeyer reaction is a chemical reaction which is used to synthesize aryl halides from aryl diazonium salts. This reaction is a method for substitution of an aromatic amino group by preparing diazonium salt that is followed by its displacement and copper salts often catalyze it.

$$ArN_{2}^{+}X^{-} \xrightarrow{CuCl/HCl} ArCl+N_{2}$$

$$CuBr/HBr \longrightarrow ArBr+N_{2}$$

$$CuCN/KCN \longrightarrow ArCN+N_{2}$$

Gattermann Reaction

Bromine and Chlorine can be present in the benzene ring by preparing the benzene diazonium salt solution with similar halogen acid present with copper powder. This is the Gattermann Reaction.

$$ArN_2^+X^- \xrightarrow{Cu/HCl} ArCl+N_2 + CuX$$

$$Cu/HBr \xrightarrow{Cu/HBr} ArBr+N_2 + CuX$$

Balz-Schiemann Reaction

When arene-diazonium chloride is prepared with fluoroboric acid, arene diazonium fluoroborate is precipitated and decomposes to yield aryl fluoride which on heating.

$$Ar \overset{+}{N_2} Cl^- + \overset{+}{HBF_4} \underset{\text{prior oboric}}{\longrightarrow} Ar - \overset{+}{N_2} BrF_4^- \overset{\text{Heat}}{\longrightarrow} Ar - F + BF_3 + N_2$$

Finkelstein Reaction

In the Finkelstein Reaction Alkyl iodides are prepared easily by the reaction of alkyl chlorides with Nal in dry acetone.

$$R - X + NaI \rightarrow R - I + NaX$$
 $(X = CI, Br)$

Swarts Reaction

When alkyl chloride is heated in the presence of a metallic fluoride like AgF, Hg₂F₂, SbF₃ or CoF₂, we get alkyl fluoride The reaction is specifically used to prepare alkyl fluorides.

$$H_3C - X + AgF \rightarrow H_3C - F + AgX$$
 $(X = Cl, Br)$

Wurtz Reaction

When Alkyl halides get reacted with sodium with dry ether, we get hydrocarbons that include the double number of carbon atoms present in the halide. This is known as the Wurtz Reaction.

$$\begin{array}{ll} CH_{3}Br+2Na+BrCH_{3} \stackrel{dry\; ether}{\longrightarrow} CH_{3}-CH_{3}+2NaBr\\ \\ Bromomethane & Ethane \\ CH_{3}Br+2Na+BrC_{2}H_{5} \stackrel{dry\; ether}{\longrightarrow} C_{2}H_{5}-C_{2}H_{5}\\ \\ Bromoethane & n-Bu\; tan\; e \end{array}$$

Wurtz-Fittig Reaction

When a mixture of alkyl halide and aryl halide gets treated with sodium in dry ether, we get an alkyl arene.

$$X + Na + RX \xrightarrow{Ether} R + NaX$$

Fittig Reaction

Aryl halides prepared with sodium in dry ether to give analogous compounds where two aryl groups joined.

$$2 + Na + Na + 2NaX$$

Friedel-Crafts Alkylation Reaction

Benzene is prepared with an alkyl halide in the presence of anhydrous aluminum chloride to give alkylbenzene.

$$+ C_2H_5CI \xrightarrow{Anhyd.AlCl_3} + HCI$$
Ethylbenzene

Friedel-Crafts Acylation Reaction

We get acyl benzene when an acyl halide is reacted with benzene in the presence of Lewis acids.

$$+ CH3COCI \xrightarrow{Anhyd.AlCl3} + HCl$$

$$Acetophenone$$
(13.77)

$$\begin{array}{c}
COCH_{3} \\
+ (CH_{3}CO)_{2}O \\
Acetic anhydride
\end{array}$$

$$\begin{array}{c}
Anhyd.AlCl_{3} \\
\Delta
\end{array}$$

$$+ CH_{3}COOH$$

Reimer-Tiemann Reaction

When preparing phenol with chloroform in the presence of sodium hydroxide, –CHO group is present at the ortho position of the benzene ring which results into salicylaldehyde.

Appendix 5

Kolbe's Reaction

Phenol reacts with sodium hydroxide to give sodium phenoxide which then reacts with carbon dioxide in acidic medium to give 2-hydroxybenzoic acid.

Rosenmund Reduction

When Acyl chloride is hydrogenated to an aldehyde over a catalyst, known as Rosenmund catalyst which is either palladium or barium sulfate. The catalyst is poisoned with either sulphur or quinoline in order to prevent further reduction of aldehyde to alcohol.

Stephen Reaction

Nitriles with stannous chloride in the presence of hydrochloric acid reduced to the corresponding imine and give the corresponding aldehyde after hydrolysis.

$$RCN + SnCl_2 + HCl \longrightarrow RCH = NH \xrightarrow{H_3O^+} RCHO$$

Etard Reaction

Chromyl chloride oxidizes methyl group to get chromium complex which on hydrolysis provides corresponding benzaldehyde.

$$H_3O^+$$
 CHC

Benzaldehyde

Gatterman - Koch Reaction

Benzene is prepared with carbon monoxide and hydrogen chloride in the presence of anhydrous aluminium chloride to give benzaldehyde.

Clemmensen Reduction

In Clemmensen reduction, Carbonyl group of aldehydes and ketones on treatment with zinc amalgam and concentrated hydrochloric acid reduced to CH2 group.

$$C = O \xrightarrow{\text{Zn-Hg}} CH_2 + H_2O$$

Wolff Kishner Reduction

Carbonyl group of aldehydes and ketones on treatment with hydrazine produces hydrazone which on heating with potassium hydroxide in a high boiling solvent (ethylene glycol) and reduce to $-CH_2$ – group.

$$C = O \xrightarrow{NH_2NH_2} C = NNH_2 \xrightarrow{KOH/ethylene glycol} heat$$
 $C = O \xrightarrow{CH_2+N_2} C = NNH_2 \xrightarrow{CH_2+N_2} C = NNH_2$

Tollens' test

Heating an aldehyde with fresh prepared ammoniacal silver nitrate solution produces a bright silver mirror due to the formation of silver metal.

RCHO+2[Ag(NH₃)]₂+3OH⁻
$$\rightarrow$$
 RCOO +2Ag + 2H₂O+4NH₃
Silver mirror

Fehling's test

Fehling's solution A (aqueous copper sulfate) and Fehling solution B (alkaline sodium potassium tartrate) are mixed in equal amounts before the test. A reddish brown precipitate is obtained when an aldehyde is heated with Fehling's reagent.

R-CHO +
$$2Cu^{2+}$$
 +5OH⁻ \longrightarrow RCOO⁻ + Cu_2O +3 H_2O
Reddish-brown ppt

Aldol reaction

Aldehydes and ketones having one α -hydrogen undergo a reaction in the presence of dilute alkali as the catalyst to produce α -hydroxy aldehydes or β -hydroxy ketones.

(i) Aldol condensation

Aldol and Ketol lose water to provide α , β -unsaturated carbonyl compounds which are aldol condensation products.

$$\begin{array}{c} 2\,\text{CH}_3-\text{CHO} & \stackrel{\text{dil. NaOH}}{\longleftrightarrow} \text{CH}_3-\text{CH}-\text{CH}_2-\text{CHO} \\ & \text{OH} \\ & 3\,\text{-Hydroxybutanal} \\ & \text{(Aldol)} \\ \\ & \frac{\Delta}{-\text{H}_2\text{O}} & \text{CH}_3-\text{CH}=\text{CH}-\text{CHO} \\ & \text{But}-2\text{-enal} \\ & \text{(Aldol condensation product)} \\ \end{array}$$

$$\begin{array}{c} 2\,\mathrm{CH_3} - \mathrm{CO} - \mathrm{CH_3} & \xrightarrow{\mathrm{Ba(OH)_2}} & \mathrm{CH_3} - \overset{\mathrm{CH_3}}{\mathrm{C}} - \mathrm{CH_2CO} - \mathrm{CH_2} \\ \text{Propanone} & \mathrm{OH} \\ \text{(Ketol)} \end{array}$$

4-Hydroxy-4-methylpentan - 2 - one

$$\xrightarrow{\Delta} CH_3 CH_3 CH_3 - C = CH - CO - CH_3$$
4 -Methylpent -3 -en - 2 -one
(Aldol condensation product)

(ii) Cross aldol condensation

Aldol condensation is carried out between two different aldehydes and ketones. It gives a mixture of four prod-

ucts if both of them includes
$$\alpha$$
-hydrogen atoms.
CH₃ -CH=CH-CHO + CH₃CH₂ -CH=CHCHO | CH₃ | 2-Methylbut-2-enal

CH₃CHO
$$+$$
 CH_3 CH₂CHO
$$CH_3$$
CH₂CHO
$$CH_3$$
CH₃-CH=CH-CHO
$$Dut-2-enal$$
But-2-enal

Cannizzaro Reaction

Aldehydes without α-hydrogen atom undergo self-oxidation and reduction reaction when prepared with concentrated alkali. The reaction is known as disproportionation reaction.

Kolbe electrolysis

In Kolbe electrolysis, An aqueous solution of sodium or potassium salt of a carboxylic acid gives alkane containing an even number of carbon atoms on electrolysis.

2CH₃COONa+2H₂O Electrolysis CH₃- CH₃+2CO₂+H₂+2NaOH Sodium acetate

Hell-Volhard-Zelinsky (HVZ) Reaction

Carboxylic acids having a α-hydrogen are halogenated at the α -position give α -halo carboxylic acids on treatment with chlorine or bromine in the presence of small amount of red phosphorus.

$$\begin{array}{c} \text{R-CH}_2\text{-COOH} & \xrightarrow{\text{(i)}\ X_2\text{Red phosphorus}} & \text{R-CH-COOH} \\ & \text{(ii)}\ \text{H}_2\text{O} & | \\ & X \\ & X = \text{Cl, Br} \\ & \alpha - \text{Halocarboxylic acid} \end{array}$$

Gabriel Phthalimide Synthesis

Phthalimide prepared with ethanolic potassium hydroxide produces potassium salt of phthalimide when heated with alkyl halide followed by alkaline hydrolysis forms the corresponding primary amine.

$$\begin{array}{c|c} O & O & O & O \\ \hline \parallel & & & & \\ \hline C & N & \text{KOH} & \hline C & NK^- & R-X & \hline C & N-R \\ \hline \parallel & & & & \\ O & O & O \\ \hline Phthalimide & N-Alkylphthalmide \\ \end{array}$$

$$\begin{array}{c|c}
O \\
\parallel \\
C \\
N-R \\
\hline
N-R \\
\hline
N-R \\
NaOH(aq)
\\
\hline
O \\
NaOH(aq)
\\
C \\
C \\
C \\
O \\
Na^+ \\
(1^\circ amine)
\\
O
\end{array}$$

Hoffmann Bromamide Degradation Reaction

An amide upon heating with bromine in presence of sodium hydroxide produces primary amine. Migration of an alkyl or aryl group takes place from carbonyl carbon of the amide to the nitrogen atom. The amine so produced includes one carbon less than that present in the amide.

O
$$\parallel$$
 R-C-NH₂ +Br₂ +4NaOH \longrightarrow R-NH₂+Na₂CO₃+2NaBr+2H₂O

Carbylamine Reaction

Aliphatic and aromatic primary amines when heated with chloroform and ethanolic potassium hydroxide produces isocyanides or carbyl amines which are foul smelling substances.

Hinsberg's Test

Benzenesulfonyl chloride (C₆H₅SO₂Cl) reacts with primary and secondary amines to produce sulphonamides.

The reaction of benzene-sulfonyl chloride with primary amine yields N-ethyl benzene-sulfonyl amide. The hydrogen attached to the nitrogen in sulphonamide is strongly acidic due to the presence of strong electron withdrawing sulfonyl group. Hence, it is soluble in alkali.

$$\begin{array}{c|c}
O & O \\
\parallel & O \\
-S-Cl+H-N-C_2H_5
\end{array}$$

$$\begin{array}{c|c}
O & \parallel \\
-S-N-C_2H_5+HC \\
\parallel & \mid \\
O & H
\end{array}$$

N-Ethyl benzene sulphonamide(Soluble in alƙali)

(ii) In the reaction with a secondary amine, N, N-diethylbenzenesulfonamide is formed. Since N,N-diethyl benzene sulphonamide does not contain any hydrogen atom attached to a nitrogen atom, it is not acidic and hence insoluble in alkali.

$$\begin{array}{c|c} O \\ O \\ S \\ -S \\ -Cl \\ +H \\ -N \\ -C_2 \\ H_5 \end{array} \rightarrow H_3 \\ C \\ -D \\ -S \\ -N \\ -C_2 \\ H_5 \\ +HCl \\ O \\ C_2 \\ H_5 \\ N.N-Diethylbenzenesulphonamide$$

(iii) Tertiary amines do not react with benzenesulphonyl chloride.

Coupling Reactions

Benzene diazonium chloride gets reacted with phenol in which the phenol molecule at its para position is mixed with the diazonium salt to give p-hydroxyazobenzene.

APPENDIX-F

Important ReagentsList of Organic Reagents

Aqueous NaOH	Reflux Nucleophilic substitution, converts haloalkanes to alco		
Mg in dry ether	Reflux	Used to make Grignard reagents with haloalkanes.	
PCl ₅	Room temperature	Chlorinating agent, reacts with OH group in alcohols and carboxylic acids.	
HNO ₃ and H ₂ SO ₄	55°C	Adds NO ₂ group into benzene ring.	
Cl ₂ and AlCl ₃	Warm gently	Adds Cl group into benzene ring.	
CH ₃ CH ₂ Cl and AlCl ₃	Warm gently	Adds CH ₃ CH ₂ group into benzene ring.	
HCl and NaNO ₂	Below 5°C	Forms diazonium salts with phenylamine.	

HCl and NaNO ₂	Below 5°C	Forms diazonium salts with phenylamine.			
Name of Reagent	Conditions	Example of its Use			
K ₂ Cr ₂ O ₇ with conc. H ₂ SO ₄	Warm gently	Oxidising agent, used commonly for oxidising secondary alcohols to ketones.			
Excess conc. H ₂ SO ₄	Heat to 170°C	Dehydrating agent, used to dehydrate alcohols to alkenes.			
Cl ₂ (g)	Ultra violet light	Free radical reaction, used to convert alkanes to haloalkanes.			
Br ₂ in CCl ₄	Room temperature, in the dark	Electrophilic addition, converts alkenes to dihaloalkanes.			
H ₂ (g)	Nickel catalyst, 300°C and 30 atmospheric pressure	Hydrogenating agent, used to convert benzene to cyclohexane.			
H ₂ (g)	Nickel catalyst, 150°C	Reducing agent, used to convert alkenes to alkanes			
Tin in hydrochloric acid	Reflux	Reducing agent for converting nitrobenzene to phenylamine.			
Acidified KMnO ₄	Room temperature	Oxidising agent, converts alkenes to diols.			
NaOH in ethanol	Reflux	Elimination reaction, converts haloalkanes to alkenes.			
NaOH (Intramolecular Cannizaro) reaction	NA	$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$			
(i) Aluminium isobutoxide (ii) Acetone (Oppenaur Oxidation)		$\begin{array}{c c} CH_3 & CH_3 & Al \\ H_3C-C-OH & CH_3 & H_3C \\ H & H_3C & C=O \\ H_3C & C=O \end{array}$			
RO ⁻ (Claisen Schmidt Reaction)		H_3C — C — H + H			
(i) Acetic anhydride (ii) Sodium acetate (Perkin's reaction)	,	$\begin{array}{c c} O & H_3C-CH_2 & O \\ H & H_3C-CH_2 & CH=CH-C-OH \\ \hline O & Cinnamic acid \\ H_3C-C-ONa & CH=CH-C-OH \\ \hline \end{array}$			
(i) Aluminium isobutoxide (ii) Propan-2 ol (MPV Reduction)		Reduces ketone to alcohol C = O Aluminium iso butoxide propan-2-ol H			

	T	
Cannizzaro Reaction		This is reaction of compounds which don't have alpha hydrogen. O H
Cross Cannizaro reaction		$\begin{array}{c} O \\ \\ \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \end{array} \begin{array}{c} O \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ \\ $
Anhydrous HI contains no water		$ \begin{array}{c} CH_3 \\ CH_3 - C - O - CH_3 \xrightarrow{HI} CH_3 - C - I + CH_3OH \\ CH_3 \\ CH_3 \end{array} $
(i) CHCl ₃ - Alc KOH (ii) H ₃ O ⁺ (Riemann Tiemann reaction)		OH CHO Salicyldehyde
(i) CO ₂ (ii) H ₃ O ⁺ (Kolbe's reaction)	High Pressure	O-Na OH O
AlCl ₃ (Fries rearrangement)	Heat	OH O
(i) K ₂ S ₂ O ₈ (ii) H ₃ O ⁺ (Elb's persulphate oxidation)		OH OH OH OH
(i) Fused NaOH (ii) H ₃ O ⁺ (Dow's process)	High Pressure	CI OH
KMnO ₄ , H ⁺ (or) K ₂ Cr ₂ O ₇ , H ⁺ (or) H ₂ CrO ₄		Oxidises alcohol to acid
PCC (Pyridinium chloro chromate)	Solvent CS ₂	Restricted oxidation of alcohol. Forms aldehyde.
MnO ₂ special oxidising agent for alcohol		OH C=O OH OH MnO ₂ selectively oxidizes primary or secondary allylic or benzylic alcohols to carbonyl compounds.
HI		
For 1 degree carbon		$H_3C \longrightarrow O$ $CH_3 + HI \longrightarrow H_3C \longrightarrow H_3C-I$ OH

For 3 degree carbon		$CH_3-O-C(CH_3)_3 + HI = (CH_3)_3C-I + CH_3OH$
Conc HI contains very less water		$\begin{array}{c} CH_3 \\ H_3C \stackrel{ }{\longrightarrow} CH_3 \stackrel{H1}{\longrightarrow} H_3C \stackrel{H_3C}{\longrightarrow} I + H_3C-OH \\ CH_3 & CH_3 \end{array}$
(i) Alc KOH (ii) NaNH ₂		CI CI H H H H H H H
X ₂ /CCI ₄		Adds both X on compound having double or triple bond
Cold Dil KMnO ₄		$ \begin{array}{cccccccccccccccccccccccccccccccccccc$
Hot KMnO₄/OH⁻		$R \xrightarrow{\qquad} R_i \xrightarrow{\qquad} R \xrightarrow{\qquad} R_i \xrightarrow{\qquad} O$
CF ₃ SO ₃ ⁻	4	Super Leaving Group
LiAlH ₄ –ether		Reduces ester to alcohol
NaBH ₄ – ROH (protic solvent)		Reduces aldehydes and ketones to alcohol but cannot act on ester
Cr ₂ O ₃ -Cu ₂ O	Heat, High pressure	Causes cleavage and reduction of ester $RCOOR_1 = RCH_2OH + R_1-OH \longrightarrow R \qquad OH + R_1-OH$
OsO ₄ + H ₂ O-NaHSO ₃ or Cold Dilute KMnO ₄ (Hydroxylation)		Gives syn vicinal diol from alkene HO OH C=C C-C
Per-formic acid (Anti Hydroxylation)		Gives anti vicinal diol HO $C = C$ OH
HBr in presence of H ₂ O ₂ (Only for HBr)		Anti Markovnikov's $R \xrightarrow{CH_2} R \xrightarrow{Br}$
Hot Alkaline KMnO ₄		Replaces "=" with either C = O or COOH $R \xrightarrow{Q} R_{1} R_{1} \longrightarrow R \xrightarrow{Q} R_{1}$ $QH R_{1}$
CH ₂ -I ₂ , ZnCu		Adds methyl group in cyclic manner R R R R R

	1	
Alcoholic KOH	Heat	Causes dehydrohalogenation of alkyl halides to form alkenes. HX from compound and adds double bond
Zn dust		Convert vicinal dihalides to alkenes.
H ₂ Pd/BaSO ₄ or S-Quinoline (Lindlar's catalyst) or BH ₃ - THF		Reduces alkyne to <i>cis</i> -alkene. It also reduces acid halides to aldehydes.
$Ph_3P = CH_2$		Converts aldehyde or ketones to alkene. The reaction is known as Wittig Reaction. $C = O \xrightarrow{Ph_3P = CH_2} C = CH_2$
NOX		H_3C $N=O$ $N=O$ O O O O O O O O O
OMDM (Oxymercuration demercuration) (i) Hg (OAc) ₂ + THF-H ₂ O (ii) NaBH ₄ -OH		H_3C CH_3 OH
NBS		Substitutes allyic carbon with aldehyde CH ₃ NBS NBS
(i) NH ₂ -NH ₂ (ii) RO	heat	Reduces carbonyl group to ketone or aldehyde
H ₃ PO ₂	Heat	Removes diazo group N≡N-Cl H₃PO₂ H₃PO₂
LiAH ₄ or NaBH ₄ or Ph ₃ SnH		Reduction
R ₂ CuLi (Lithium dialkylcuprate)	7	Removes halogen from RX and adds R. The reaction is known as Corey-House synthesis. The reaction is used to produce an alkane. The R- from R_2 CuLi displaces -X from R-X to produce an alkane.
Mg-ether		Adds Mg between R & X
Red P + HI		Reduces alcohols, aldehydes, ketones and carboxylic acids to alkanes.

Name	Condition	Example
$Br_2 + CS_2$		Causes mono-bromination of phenol to produce p-bromophenol.
$Br_2 + H_2O$	Compound should be more activated than benzene	Produces 2,4,6-Tribromophenol from phenol. OH OH Br./H,O Br

Appendix 11

Sn + HCl or Fe + HCl	Reduces nitro group O N NH ₂ Sn+HCl
CuCl + HCl	Removes Diazo group The N-Cl Cucl HCl
NH ₄ SH or Na ₂ S	Special regents which reduce only –NO ₂ group to –NH ₂
$CrO_2Cl_2 + H_3O^+$	Converts toluene to benzaldehyde

APPENDIX-G

Important Minerals, Compositions & their Chemical Formula

Mineral	Composition		Remarks
Apatite	Calcium phosphate	$Ca_{10}(PO_4)_6X_2$ (X is F, Cl, or (OH).]	Main mineral in phosphate rock
Asbestos	Hydrated magnesium silicate	Mg ₆ (Si ₄ O ₁₂)(OH) ₃	In form of long fibres
Baryte	Barium sulphate	BaSO ₄	Filler for pigments
Betonies	A clay mineral	(Al.Mg) ₈ (Si ₄ O ₁₀) ₃ (OH) ₁₀ •12H ₂ O	Agglomeration additive
Borax	Sodium borate	Na ₂ B ₄ O ₇ •10H ₂ O	
Clay	Hydrated aluminium silicates		Used in paper making
Cryolite	Sodium aluminium fluoride	Na ₃ AlF ₆	Low melting point
Diamond - industrial	Crystalline carbon	С	The hardest mineral
Diatomite	Hydrated silica	SiO ₂ (H ₂ O) _n	Marine fossils, large surface area
Feldspar	A mineral group	K, Al silicates	
Fluorspar	Calcium fluoride	CaF ₂	Main source of fluorine
Garnet	A group of silicates that crystal- lize in the cubic system	Mg ₃ Fe ₂ Si ₃ O ₁₂	Abrasives, gemstones
Graphite	Carbon (crystalline)	С	
Gypsum	Calcium sulphate	CaSO ₄ •2H ₂ O	
Kaolinite	A clay mineral	Al ₄ (Si ₄ O ₁₀)(OH) ₈	
Limestone	Calcium carbonate	CaCO ₃	
Magnetite	Magnesium carbonate	MgCO ₃	
Marble	Calcium carbonate	CaCO ₃ crystalline	
Mica		K, Al silicates	
Nepheline syenite	Sodium aluminium silicate		

Potash	Potassium chloride and carbonate	KCl, K ₂ CO ₃	Fertilizer
Pumice	Silicate		Porous, light, volcanic rock, large surface area
Quartz	Silica	SiO ₂	
Salt	Sodium chloride NaCl		
Sand and gravel	Silica	SiO ₂	
Sulfur	Sulfur	S	(-)
Talc	Hydrated magnesium silicate	Mg ₃ (Si ₄ O ₁₀)(OH) ₂	Also known as soapstone
Trona	Sodium carbonate	Na ₂ CO ₃ •NaHCO ₃ •2H ₂ O	
Vermiculite	Hydrated silicates	(Mg,Fe ²⁺ ,Fe ³⁺) ₃ [(Al,Si) ₄ O ₁₀] (OH) ₂ ·4H ₂ O	Expands and swells on heating
Zeolite	Hydrated alkali alumino silicates	$Na_2(AlO_2)_x(SiO_2)_y \bullet nH_2O$	Ion exchanger

APPENDIX-H

Important Metals & their Ores

S.No	Metal	Ores
1	Aluminium(Al)	Bauxite, Corundum, Feldspar, Cryolite, Alunite, Kaolin
2	Antimony(Sb)	Stibnite
3	Barium(Ba)	Barytes
4	Bismuth(Bi)	Bismuthate
5	Cadmium(Cd)	Greenockite
6	Calcium(Ca)	Dolomite, Calcite, Gypsum, Fluorspar, Asbestos
7	Cobalt(Co)	Smelite
8	Copper(Cu)	Cuprite, Copper glance ,Copper pyrites
9	Gold(Au)	Calaverite, Sylvenites
10	Iron(Fe)	Hematite, Limonite, Magnetite, Siderite, Iron pyrite, Copper pyrites
11	Lead(Pb)	Galena
12	Magnesium(Mg)	Magnesite, Dolomite, Epsom salt, Kieserite, Carnalite
13	Manganese(Mn)	Pyrolusite, Magnate
14	Mercury(Hg)	Cinnabar
15	Nickel(Ni)	Millerite
16	Potassium(K)	Nitrate(saltpetre), Carnallite
17	Silver(Ag)	Ruby silver, Horn silver
18	Sodium(Na)	Chile saltpetre , Trona, Borax, Common salt
19	Strontium(Sr)	Strontianite, Silestone
20	Tin(Sn)	Cassiterite
21	Uranium(U)	Carnallite, Pitch blende
22	Zinc(Zn)	Zinc blende, Zincite, Calamine