

INORGANIC CHEMISTRY CHEAT SHEET

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1. Periodic Trends & Effective Nuclear Charge

Slater's Rules for Effective Nuclear Charge (Z^*):

$$Z^* = Z - S$$

where S is the shielding constant:

- Electrons in same group (ns, np): count 0.35 each (0.30 for $1s$).
- Electrons in $(n-1)$ shell: count 0.85 each.
- Electrons in $(n-2)$ or lower shells: count 1.00 each.
- For d or f electrons: same group 0.35, lower groups 1.00.

General Periodic Trends:

- Atomic Radius:** Increases down, decreases across $\rightarrow$.
- Ionization Energy (IE):** Increases across $\rightarrow$, decreases down. *Exceptions:* $IE_1(\text{Be}) > IE_1(\text{B})$ ($2s^2$ vs $2p^1$), $IE_1(\text{N}) > IE_1(\text{O})$ (half-filled $2p^3$).
- Electron Affinity (EA):** Most negative for Halogens. $\text{Cl} > \text{F}$ due to inter-electronic repulsion in compact $2p$ of F.
- Electronegativity (χ):** $\text{F}(4.0) > \text{O}(3.5) > \text{N/Cl}(3.0) > \text{Br}(2.8) > \text{C/S/I}(2.5) > \text{H/P}(2.1)$.

2. VSEPR Theory & Geometries

Steric Number (SN):

$$\text{SN} = \frac{1}{2} [V + M - C + A]$$

V : valence e^- , M : monovalent atoms, C : cationic charge, A : anionic charge.

SN	bp	lp	Geometry	Example
2	2	0	Linear (180°)	$\text{BeCl}_2, \text{CO}_2$
3	3	0	Trig. Planar (120°)	BF_3
3	2	1	Bent ($< 120^\circ$)	$\text{SO}_2, \text{NO}_2^-$
4	4	0	Tetrahedral (109.5°)	$\text{CH}_4, \text{NH}_4^+$
4	3	1	Trig. Pyramidal	$\text{NH}_3, \text{PCl}_3$
4	2	2	Bent (104.5°)	$\text{H}_2\text{O}, \text{OF}_2$
5	5	0	Trig. Bipyramidal	PCl_5
5	4	1	Seesaw	SF_4
5	3	2	T-shaped	ClF_3
5	2	3	Linear	$\text{XeF}_2, \text{I}_3^-$
6	6	0	Octahedral (90°)	SF_6
6	5	1	Sq. Pyramidal	BrF_5
6	4	2	Sq. Planar	XeF_4

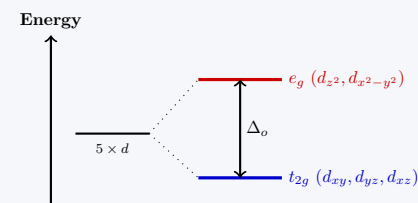
Bent's Rule: More electronegative substituents prefer hybrid orbitals with **less** s -character (axial positions in TBP). Lone pairs prefer orbitals with **more** s -character (equatorial positions in TBP).

3. Coordination Chemistry & CFT

Crystal Field Splitting (Δ):

- Octahedral (O_h):** d -orbitals split into t_{2g} (d_{xy}, d_{yz}, d_{xz}) at $-0.4\Delta_o$ and e_g ($d_{z^2}, d_{x^2-y^2}$) at $+0.6\Delta_o$.
- Tetrahedral (T_d):** Inverted splitting: e lower ($-0.6\Delta_t$), t_2 higher ($+0.4\Delta_t$).

$$\Delta_t = \frac{4}{9}\Delta_o$$



Spectrochemical Series (Weak to Strong Field):



Magnetic Moment (μ_s):

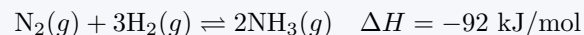
$$\mu_s = \sqrt{n(n+2)} \text{ B.M.} \quad (n = \text{number of unpaired } e^-)$$

Jahn-Teller Distortion: Occurs when degenerate orbitals are unsymmetrically occupied (strongest in high-spin d^4 , low-spin d^7 , d^9 in O_h).

4. Industrial Processes & Main Group

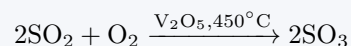
Key Chemical Processes:

- **Haber-Bosch Process (Ammonia):**

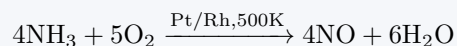


Cat: Fe with $\text{K}_2\text{O}/\text{Al}_2\text{O}_3$ promoter
(450°C, 200 atm).

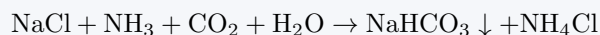
- **Contact Process (Sulfuric Acid):**



- **Ostwald Process (Nitric Acid):**



- **Solvay Process (Soda Ash):**



5. Metallurgy & Ellingham Diagrams

Principles of Metallurgy:

- **Calcination:** Heating in absence of air (carbonates $\rightarrow$ oxides).
- **Roasting:** Heating in excess air (sulfides $\rightarrow$ oxides).
- **Zone Refining:** Based on principle that impurities are more soluble in molten metal than solid (Si, Ge).
- **Mond Process (Ni):** $\text{Ni} + 4\text{CO} \xrightarrow{330\text{K}} \text{Ni}(\text{CO})_4 \xrightarrow{450\text{K}} \text{Ni} + 4\text{CO}$.
- **Van Arkel Process (Zr, Ti):** $\text{Zr} + 2\text{I}_2 \rightarrow \text{ZrI}_4 \xrightarrow{1800\text{K}} \text{Zr} + 2\text{I}_2$.

Ellingham Diagram Guidelines:

- Plots ΔG° vs T for oxide formation.
- Slope = $-\Delta S^\circ$. Positive slope for most metals ($M + \text{O}_2 \rightarrow \text{MO}$).
- Any metal line **below** another can reduce the oxide of the upper metal at that temperature.

6. Qualitative Analysis & Flame Colors

Flame Test Colors:

- Li^+ : Crimson Red Na^+ : Persistent Yellow
- K^+ : Lilac / Violet Ca^{2+} : Brick Red
- Sr^{2+} : Crimson Ba^{2+} : Apple Green
- Cu^{2+} : Blue-Green Pb^{2+} : Pale Blue

Qualitative Analysis Cation Groups:

- **Grp I** ($\text{Pb}^{2+}, \text{Ag}^+, \text{Hg}_2^{2+}$): Precipitate as Chlorides with dil. HCl.
- **Grp II** ($\text{Cu}^{2+}, \text{Cd}^{2+}, \text{Bi}^{3+}, \text{Pb}^{2+}, \text{Hg}_2^{2+}$): Precipitate as Sulfides in acidic H_2S .
- **Grp III** ($\text{Fe}^{3+}, \text{Al}^{3+}, \text{Cr}^{3+}$): Precipitate as Hydroxides with $\text{NH}_4\text{OH} + \text{NH}_4\text{Cl}$.
- **Grp IV** ($\text{Zn}^{2+}, \text{Mn}^{2+}, \text{Ni}^{2+}, \text{Co}^{2+}$): Precipitate as Sulfides in basic $\text{H}_2\text{S}/\text{NH}_4\text{OH}$.
- **Grp V** ($\text{Ba}^{2+}, \text{Sr}^{2+}, \text{Ca}^{2+}$): Precipitate as Carbonates with $(\text{NH}_4)_2\text{CO}_3$.

7. Organometallics & 18-Electron Rule

18-Electron Rule:

Total Valence Electrons (TVE) = Metal Valence e^- + Ligand Electrons

Stable transition metal organometallic complexes usually have 18 electrons.

Common Ligand Electron Counts (Neutral Method):

- $1e^-$ donors: H, CH_3 , Cl, Br, I
- $2e^-$ donors: CO, PR_3 , NH_3 , H_2O , $\text{CH}_2 = \text{CH}_2$
- $3e^-$ donors: η^3 -allyl
- $4e^-$ donors: 1,3-butadiene
- $5e^-$ donors: η^5 -Cp (Cyclopentadienyl)
- $6e^-$ donors: η^6 -benzene