

Chemistry class 11<sup>th</sup> Important Questions**1. Atomic Radius (Size)****General trend**

- Increases down a group
- Decreases across a period (left → right)

**Usual order (same period)**

Group 1 &gt; Group 2 &gt; Group 13 &gt; Group 14 &gt; Group 15 &gt;

Group 16 &gt; Group 17

**Key exceptions**

- N > O > F**  
Due to strong electron-electron repulsion in compact p-orbitals
- Ga < Al**  
Because of **d-block contraction**
- Pb ≈ Sn** (sometimes Pb smaller)  
Due to **lanthanide contraction**

**2. Ionic Radius****General trend**

- Cations < neutral atom**
- Anions > neutral atom**
- Increases down a group

**Isoelectronic series order**

Greater nuclear charge → smaller ion

 $\text{N}^{3-} > \text{O}^{2-} > \text{F}^- > \text{Ne} > \text{Na}^+ > \text{Mg}^{2+} > \text{Al}^{3+}$ **3. Ionization Energy (IE)****General trend**

- Increases across a period
- Decreases down a group

**Usual order (period 2)** $\text{Li} < \text{Be} < \text{B} < \text{C} < \text{N} < \text{O} < \text{F} < \text{Ne}$ **Important exceptions**

- Be > B**  
(filled 2s orbital is more stable)
- N > O**  
(half-filled p<sup>3</sup> is more stable than p<sup>4</sup>)
- Al < Mg**
- S < P**

**4. Electron Affinity (EA)****General trend**

- Becomes **more negative** across a period
- Less negative down a group

**Expected order** $\text{Cl} > \text{F} > \text{Br} > \text{I}$ **Exceptions**

- Cl > F**  
(small size of F causes e<sup>-</sup>-e<sup>-</sup> repulsion)
- N, Be, Mg, noble gases ≈ zero or positive EA**
  - N: half-filled p<sup>3</sup>
  - Be/Mg: filled s<sup>2</sup>
  - Noble gases: complete octet

**5. Electronegativity (EN)****General trend**

- Increases across a period
- Decreases down a group

**Highest to lowest (common)** $\text{F} > \text{O} > \text{N} > \text{Cl} > \text{Br} > \text{I} > \text{S} > \text{C} > \text{P}$ **Exceptions / notes**

- N > Cl** (sometimes surprises students)
- Noble gases usually **excluded**
- F is the most electronegative element**

**6. Metallic Character****General trend**

- Increases down a group
- Decreases across a period

**Order (period 3)** $\text{Na} > \text{Mg} > \text{Al} > \text{Si} > \text{P} > \text{S} > \text{Cl}$ **Exception**

- Al > Mg** in some chemical reactions (due to high charge density)

**7. Oxidizing & Reducing Power****Oxidizing power (non-metals)** $\text{F}_2 > \text{Cl}_2 > \text{Br}_2 > \text{I}_2$ **Reducing power (metals)** $\text{Cs} > \text{Rb} > \text{K} > \text{Na} > \text{Li}$ **Exception**

- Li is strongest reducing agent in aqueous solution**  
(high hydration enthalpy)

**8. Shielding Effect****General trend**

- Increases down a group
- Almost constant across a period

**Exception**

- d and f electrons shield poorly**  
→ causes:
  - Lanthanide contraction
  - High density & high IE in transition metals

**9. Inert Pair Effect****Observed in**

- Heavier p-block elements (Group 13–16)**

**Effect**

Lower oxidation states become more stable down the group

**Example**Group 13: +3 → +1 (Tl<sup>+</sup> most stable)Group 14: +4 → +2 (Pb<sup>2+</sup> stable)**10. Diagonal Relationship**

Elements diagonally adjacent show similar properties:

 $\text{Li} \leftrightarrow \text{Mg}$  $\text{Be} \leftrightarrow \text{Al}$  $\text{B} \leftrightarrow \text{Si}$ Reason: similar **charge density and size****11. Lanthanide Contraction****What it is**

<p><b>1. Acidic strength of hydrides (H–X)</b>  <b>Across a period (left → right)</b>        Acidity <b>increases</b> with electronegativity:  <math>\text{CH}_4 &lt; \text{NH}_3 &lt; \text{H}_2\text{O} &lt; \text{HF}</math>  <b>Down a group (top → bottom)</b>        Acidity <b>increases</b> with size (weaker H–X bond):  <math>\text{HF} &lt; \text{HCl} &lt; \text{HBr} &lt; \text{HI}</math></p> <p><b>2. Acidic strength of oxoacids (same central atom)</b>        More oxygen atoms → stronger acid  <math>\text{HClO} &lt; \text{HClO}_2 &lt; \text{HClO}_3 &lt; \text{HClO}_4</math>        Reason: greater <b>–I effect</b> and better resonance stabilization.</p> <p><b>3. Oxoacids with different central atoms (same oxidation state)</b>        Higher electronegativity → stronger acid  <math>\text{HClO}_4 &gt; \text{HBrO}_4 &gt; \text{HIO}_4</math></p> <p><b>4. Acidic strength of binary acids of non-metals</b>        General order:  <math>\text{HF} &lt; \text{HCl} &lt; \text{HBr} &lt; \text{HI}</math>  <math>\text{H}_2\text{O} &lt; \text{H}_2\text{S} &lt; \text{H}_2\text{Se} &lt; \text{H}_2\text{Te}</math>  <math>\text{NH}_3 &lt; \text{PH}_3 &lt; \text{AsH}_3 &lt; \text{SbH}_3</math>        Key idea: <b>bond strength decreases down the group</b></p>	<p>Gradual decrease in atomic &amp; ionic radii from <b>La → Lu</b></p> <p><b>Consequences</b></p> <ul style="list-style-type: none"> <li>• <math>\text{Zr} \approx \text{Hf}</math> in size</li> <li>• High density &amp; melting point of 5d elements</li> <li>• <math>\text{Ga} &lt; \text{Al}</math> (size anomaly)</li> </ul> <p><b>5. Acidic strength of metal hydroxides</b>  <b>Alkali metals</b>  <math>\text{LiOH} &lt; \text{NaOH} &lt; \text{KOH} &lt; \text{RbOH} &lt; \text{CsOH}</math>  <b>Alkaline earth metals</b>  <math>\text{Mg}(\text{OH})_2 &lt; \text{Ca}(\text{OH})_2 &lt; \text{Sr}(\text{OH})_2 &lt; \text{Ba}(\text{OH})_2</math>        (As metallic character increases, basicity increases → acidity decreases)</p> <p><b>6. Acidic strength of oxides</b>  <b>Period 3 oxides</b>  <math>\text{Na}_2\text{O} &lt; \text{MgO} &lt; \text{Al}_2\text{O}_3 &lt; \text{SiO}_2 &lt; \text{P}_2\text{O}_5 &lt; \text{SO}_3 &lt; \text{Cl}_2\text{O}_7</math> <ul style="list-style-type: none"> <li>• Metal oxides → basic</li> <li>• Non-metal oxides → acidic</li> <li>• <b><math>\text{Al}_2\text{O}_3</math> is amphoteric</b></li> </ul></p> <p><b>7. Acidic strength of organic acids</b>  <b>Carboxylic acids</b>        Electron-withdrawing groups increase acidity:  <math>\text{HCOOH} &gt; \text{CH}_3\text{COOH} &gt; \text{C}_2\text{H}_5\text{COOH}</math>  <math>\text{CF}_3\text{COOH} &gt; \text{CCl}_3\text{COOH} &gt; \text{CH}_3\text{COOH}</math>  <b>Phenols (substituent effect)</b>  <math>\text{p-NO}_2\text{-phenol} &gt; \text{phenol} &gt; \text{p-CH}_3\text{-phenol}</math></p> <p><b>8. Acidic nature of hydrogen halides in water</b>        Strong acids:  <math>\text{HCl} \approx \text{HBr} \approx \text{HI}</math>        Weak acid:  <math>\text{HF}</math>        Exception reason: <b>very strong H–F bond</b></p> <p><b>9. Acidic strength of polyprotic acids (same acid)</b>        Successive ionization:  <math>\text{H}_3\text{PO}_4 &gt; \text{H}_2\text{PO}_4^- &gt; \text{HPO}_4^{2-} &gt; \text{PO}_4^{3-}</math></p>
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