

SUPPOSE THAT TWO HYDROXIDES, MOH AND M(OH)₂, BOTH HAVE A K_{SP} OF 2.15 × 10⁻¹² AND THAT INITIALLY BOTH CATIONS

SUPPOSE THAT TWO HYDROXIDES, MOH AND M(OH)₂, BOTH HAVE A K_{SP} OF 2.15 × 10⁻¹² AND THAT INITIALLY BOTH CATIONS PROVIDES AN INTRIGUING SCENARIO FOR UNDERSTANDING SOLUBILITY EQUILIBRIA, PARTICULARLY HOW THE SOLUBILITY PRODUCTS (K_{SP}) INFLUENCE THE DISSOLUTION BEHAVIOR OF DIFFERENT HYDROXIDES. IN THIS ARTICLE, WE EXPLORE THE IMPLICATIONS OF HAVING TWO HYDROXIDES WITH IDENTICAL K_{SP} VALUES, ANALYZE THEIR SOLUBILITY, AND UNDERSTAND THE FACTORS THAT AFFECT THEIR EQUILIBRIUM IN AQUEOUS SOLUTIONS. THIS DISCUSSION IS FUNDAMENTAL FOR STUDENTS AND CHEMISTS INTERESTED IN SOLUBILITY EQUILIBRIA, IONIC INTERACTIONS, AND THE PRINCIPLES GOVERNING PRECIPITATION REACTIONS.

UNDERSTANDING K_{SP} AND ITS SIGNIFICANCE

WHAT IS K_{SP}?

THE SOLUBILITY PRODUCT CONSTANT, K_{SP}, IS AN EQUILIBRIUM CONSTANT THAT DESCRIBES THE EXTENT TO WHICH A SPARINGLY SOLUBLE COMPOUND DISSOLVES IN WATER. IT IS DEFINED FOR THE EQUILIBRIUM:



THE K_{SP} EXPRESSION IS:

$$K_{\text{SP}} = [\text{M}^{+}]^n [\text{OH}^{-}]^n$$

WHERE THE BRACKETS DENOTE MOLAR CONCENTRATIONS AT EQUILIBRIUM.

A SMALLER K_{SP} INDICATES A LESS SOLUBLE COMPOUND, WHEREAS A LARGER K_{SP} SIGNIFIES HIGHER SOLUBILITY.

IMPLICATION OF IDENTICAL K_{SP} VALUES

HAVING TWO DIFFERENT HYDROXIDES, MOH AND M(OH)₂, BOTH WITH THE SAME K_{SP} VALUE OF 2.15 × 10⁻¹², IS UNUSUAL. THIS EQUALITY SUGGESTS THAT, AT EQUILIBRIUM, BOTH COMPOUNDS HAVE SIMILAR SOLUBILITY LIMITS IN WATER, EVEN THOUGH THEIR CHEMICAL FORMULAS DIFFER. HOWEVER, THE DIFFERING STOICHIOMETRY OF THEIR DISSOLUTION REACTIONS INFLUENCES THEIR INDIVIDUAL SOLUBILITY AND THE CONCENTRATIONS OF IONS IN SOLUTION.

ANALYZING THE DISSOLUTION OF MOH AND M(OH)₂

DISSOLUTION REACTIONS

THE DISSOLUTION REACTIONS FOR EACH HYDROXIDE ARE:

1. MOH:



THE K_{SP} EXPRESSION:

$$K_{sp}^{(1)} = [\text{M}^+][\text{OH}^-]$$

2. $\text{M}(\text{OH})_2$:



THE K_{SP} EXPRESSION:

$$K_{sp}^{(2)} = [\text{M}^{2+}][\text{OH}^-]^2$$

SINCE BOTH HAVE THE SAME K_{SP} VALUE:

$$K_{sp}^{(1)} = K_{sp}^{(2)} = 2.15 \times 10^{-12}$$

WE CAN ANALYZE THEIR SOLUBILITIES BASED ON THEIR ION CONCENTRATIONS.

CALCULATING SOLUBILITY (S)

LET'S DEFINE THE MOLAR SOLUBILITY OF EACH COMPOUND:

- For MOH, S_1 = MOLAR CONCENTRATION OF M^+ (AND OH^-) IN SOLUTION AT EQUILIBRIUM.
- For $\text{M}(\text{OH})_2$, S_2 = MOLAR CONCENTRATION OF M^{2+} IONS; THE HYDROXIDE ION CONCENTRATION WILL BE $2 \times S_2$.

FOR MOH:

$$[\text{M}^+] = [\text{OH}^-] = S_1$$

USING THE K_{SP}:

$$K_{sp}^{(1)} = S_1 \times S_1 = S_1^2$$

$$S_1 = \sqrt{K_{sp}^{(1)}} = \sqrt{2.15 \times 10^{-12}} \approx 1.46 \times 10^{-6} \text{ M}$$

FOR $\text{M}(\text{OH})_2$:

$$[\text{M}^{2+}] = S_2$$

$$[\text{OH}^-] = 2 S_2$$

APPLYING THE K_{SP}:

$$K_{sp}^{(2)} = S_2 \times (2S_2)^2 = S_2 \times 4 S_2^2 = 4 S_2^3$$

SOLVE FOR S_2 :

$$S_2^3 = \frac{K_{sp}^{(2)}}{4} = \frac{2.15 \times 10^{-12}}{4} = 5.375 \times 10^{-13}$$

$$S_2 = \sqrt[3]{5.375 \times 10^{-13}} \approx 8.1 \times 10^{-5} \text{ M}$$

SUMMARY OF SOLUBILITIES:

- MOH: $S_1 \approx 1.46 \times 10^{-6} \text{ M}$
- $\text{M}(\text{OH})_2$: $S_2 \approx 8.1 \times 10^{-5} \text{ M}$

DESPITE BOTH HAVING THE SAME K_{SP}, $\text{M}(\text{OH})_2$ IS SIGNIFICANTLY MORE SOLUBLE BECAUSE ITS STOICHIOMETRY ALLOWS FOR A HIGHER TOTAL MOLAR CONCENTRATION OF IONS IN SOLUTION.

IMPACT OF INITIAL CATION CONCENTRATIONS

INITIAL CONDITIONS AND COMMON ION EFFECT

SUPPOSE INITIALLY BOTH CATIONS, M^+ AND M^{2+} , ARE PRESENT IN SOLUTION AT CERTAIN CONCENTRATIONS. THE PRESENCE OF THESE IONS INFLUENCES THE SOLUBILITY OF THEIR RESPECTIVE HYDROXIDES VIA THE COMMON ION EFFECT:

- COMMON ION EFFECT: THE ADDITION OF M^+ OR M^{2+} SHIFTS THE DISSOLVING EQUILIBRIUM TOWARD THE SOLID FORM, DECREASING SOLUBILITY.
- WHEN BOTH CATIONS ARE INITIALLY PRESENT, THE SOLUBILITY OF EACH HYDROXIDE REDUCES FURTHER, DEPENDING ON THE CONCENTRATION OF THE IONS ALREADY IN SOLUTION.

PREDICTING PRECIPITATION AND DISSOLUTION BEHAVIOR

IF THE INITIAL CONCENTRATIONS ARE LOW, BOTH HYDROXIDES CAN DISSOLVE UNTIL THEIR RESPECTIVE SOLUBILITY LIMITS. HOWEVER:

- IF THE INITIAL M^+ CONCENTRATION IS HIGH, MOH 'S SOLUBILITY WILL BE SUPPRESSED.
- IF THE INITIAL M^{2+} CONCENTRATION IS HIGH, $M(OH)_2$ 'S SOLUBILITY WILL BE SUPPRESSED.

IN SOME CASES, ADDING EXCESS M^{2+} IONS COULD CAUSE $M(OH)_2$ TO PRECIPITATE OUT OF SOLUTION, EVEN IF INITIALLY DISSOLVED, AFFECTING THE EQUILIBRIUM.

COMPARATIVE ANALYSIS OF HYDROXIDE SOLUBILITY

FACTORS INFLUENCING SOLUBILITY

SEVERAL FACTORS INFLUENCE THE SOLUBILITY OF HYDROXIDES, INCLUDING:

- IONIC STRENGTH: INCREASED IONIC STRENGTH GENERALLY DECREASES SOLUBILITY.
- pH OF THE SOLUTION: HIGHER pH (MORE OH^-) CAN CAUSE PRECIPITATION OF HYDROXIDES.
- COMPLEX FORMATION: THE FORMATION OF COMPLEXES WITH OTHER IONS CAN ALTER SOLUBILITY.
- TEMPERATURE: GENERALLY, INCREASING TEMPERATURE INCREASES SOLUBILITY, BUT SPECIFIC BEHAVIOR DEPENDS ON ENTHALPY CHANGES.

REAL-WORLD APPLICATIONS

UNDERSTANDING THE SOLUBILITY OF HYDROXIDES WITH IDENTICAL K_{SP} VALUES IS CRITICAL IN MANY PRACTICAL FIELDS:

- WATER TREATMENT: PRECIPITATING HEAVY METALS AS HYDROXIDES.
- PHARMACEUTICALS: ENSURING PROPER SOLUBILITY OF HYDROXIDE FORMS OF DRUGS.
- CHEMICAL MANUFACTURING: CONTROLLING PRECIPITATION IN INDUSTRIAL PROCESSES.

CONCLUSION

IN SUMMARY, EVEN WHEN TWO HYDROXIDES LIKE MOH AND $M(OH)_2$ SHARE THE SAME K_{SP} VALUE, THEIR SOLUBILITY BEHAVIORS CAN DIFFER SIGNIFICANTLY DUE TO THEIR STOICHIOMETRY AND THE RESULTING ION CONCENTRATIONS AT EQUILIBRIUM. THE MOLAR SOLUBILITY CALCULATIONS REVEAL THAT $M(OH)_2$ IS MORE SOLUBLE THAN MOH , OWING TO THE QUADRATIC DEPENDENCE OF ITS SOLUBILITY PRODUCT ON HYDROXIDE ION CONCENTRATION. WHEN INITIAL CATIONS ARE PRESENT, THE SOLUBILITY IS FURTHER INFLUENCED BY THE COMMON ION EFFECT, AFFECTING PRECIPITATION AND DISSOLUTION DYNAMICS. A

THOROUGH UNDERSTANDING OF THESE PRINCIPLES IS ESSENTIAL FOR PREDICTING AND CONTROLLING SOLUBILITY IN VARIOUS CHEMICAL AND INDUSTRIAL PROCESSES.

BY GRASPING THESE CONCEPTS, CHEMISTS CAN BETTER MANIPULATE CONDITIONS TO FAVOR DISSOLUTION OR PRECIPITATION, OPTIMIZE REACTIONS, AND DESIGN EFFECTIVE SEPARATION PROCESSES. THE STUDY OF SOLUBILITY EQUILIBRIA REMAINS A CORNERSTONE OF INORGANIC CHEMISTRY, WITH FAR-REACHING APPLICATIONS ACROSS SCIENCE AND INDUSTRY.

FREQUENTLY ASKED QUESTIONS

WHAT DOES THE SIMILAR K_{sp} VALUE OF 2.15×10^{-12} FOR MOH AND $M(OH)_2$ INDICATE ABOUT THEIR SOLUBILITY?

IT INDICATES THAT BOTH HYDROXIDES HAVE VERY LOW SOLUBILITY IN WATER, WITH THEIR EQUILIBRIUM CONCENTRATIONS OF IONS BEING EXTREMELY SMALL.

HOW DOES THE DIFFERENCE IN STOICHIOMETRY BETWEEN MOH AND $M(OH)_2$ AFFECT THEIR ION CONCENTRATIONS AT EQUILIBRIUM?

SINCE MOH DISSOCIATES INTO ONE M^+ AND ONE OH^- ION, WHILE $M(OH)_2$ DISSOCIATES INTO ONE M^{2+} AND TWO OH^- IONS, THEIR ION CONCENTRATIONS AT EQUILIBRIUM WILL DIFFER ACCORDINGLY, REFLECTING THEIR DISSOCIATION RATIOS.

GIVEN EQUAL INITIAL CONCENTRATIONS OF M^+ AND OH^- , WHICH HYDROXIDE IS MORE LIKELY TO PRECIPITATE FIRST IN A SOLUTION?

BOTH HYDROXIDES ARE EXTREMELY INSOLUBLE; HOWEVER, $M(OH)_2$, WITH THE SAME K_{sp} , MAY TEND TO PRECIPITATE UNDER CONDITIONS FAVORING THE FORMATION OF M^{2+} IONS, ESPECIALLY IF THE M^{2+} CONCENTRATION INCREASES.

HOW CAN THE COMMON ION EFFECT INFLUENCE THE SOLUBILITY OF THESE HYDROXIDES?

THE PRESENCE OF ADDITIONAL M^+ OR OH^- IONS IN SOLUTION WILL SUPPRESS THE DISSOLUTION OF BOTH HYDROXIDES DUE TO LE CHATELIER'S PRINCIPLE, FURTHER DECREASING THEIR SOLUBILITY.

IF THE INITIAL CONCENTRATION OF M^+ IONS IS INCREASED, HOW DOES THIS AFFECT THE SOLUBILITY OF $M(OH)_2$?

INCREASING M^+ CONCENTRATION CAN SHIFT THE EQUILIBRIUM TO FAVOR THE SOLID FORM, DECREASING THE SOLUBILITY OF $M(OH)_2$ DUE TO THE COMMON ION EFFECT.

WHAT IS THE SIGNIFICANCE OF THE K_{sp} VALUES BEING SO LOW FOR BOTH HYDROXIDES, AND HOW DOES IT IMPACT THEIR PRECIPITATION BEHAVIOR?

THE VERY LOW K_{sp} VALUES MEAN BOTH HYDROXIDES ARE HIGHLY INSOLUBLE, SO THEY TEND TO PRECIPITATE READILY WHEN THEIR CONSTITUENT IONS REACH CERTAIN THRESHOLD CONCENTRATIONS IN SOLUTION.

HOW WOULD THE PRESENCE OF A COMMON ION, SUCH AS ADDITIONAL OH^- , INFLUENCE THE SOLUBILITY OF MOH AND $M(OH)_2$?

ADDING MORE OH^- IONS WOULD DECREASE THE SOLUBILITY OF BOTH HYDROXIDES BY SHIFTING THEIR DISSOLUTION EQUILIBRIA TO FAVOR THE SOLID FORM, REDUCING THEIR ION CONCENTRATIONS IN SOLUTION.

CAN THE SOLUBILITY PRODUCT CONSTANTS HELP DETERMINE THE SOLUBILITY OF THESE HYDROXIDES IN MOLAR TERMS?

YES, KNOWING THE K_{sp} ALLOWS CALCULATION OF MOLAR SOLUBILITY BY SETTING UP EQUILIBRIUM EXPRESSIONS BASED ON THEIR DISSOCIATION RATIOS, PROVIDING QUANTITATIVE INSIGHT INTO THEIR SOLUBILITY IN WATER.

ADDITIONAL RESOURCES

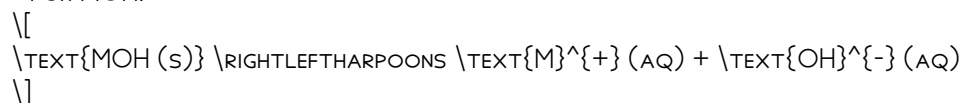
UNDERSTANDING THE SOLUBILITY OF HYDROXIDES: AN IN-DEPTH ANALYSIS OF MOH AND $M(OH)_2$

WHEN EXPLORING THE FASCINATING WORLD OF INORGANIC CHEMISTRY, PARTICULARLY THE SOLUBILITY OF METAL HYDROXIDES, THE CONCEPTS SURROUNDING THEIR SOLUBILITY PRODUCT CONSTANTS (K_{sp}) ARE FUNDAMENTAL. IN THIS DETAILED REVIEW, WE FOCUS ON A HYPOTHETICAL SCENARIO INVOLVING TWO HYDROXIDES, MOH AND $M(OH)_2$, BOTH SHARING AN IDENTICAL K_{sp} VALUE OF 2.15×10^{-12} . THIS SCENARIO PROVIDES A RICH FOUNDATION FOR UNDERSTANDING SOLUBILITY EQUILIBRIA, THE BEHAVIOR OF IONS IN AQUEOUS SOLUTIONS, AND THE IMPLICATIONS OF INITIAL CATION CONCENTRATIONS.

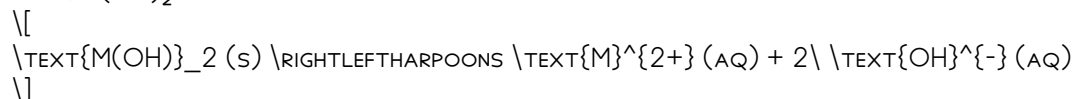
INTRODUCTION TO HYDROXIDES AND THEIR DISSOLUTION EQUILIBRIA

HYDROXIDES ARE COMPOUNDS FORMED BY THE COMBINATION OF METAL CATIONS WITH HYDROXIDE IONS (OH^-). THEIR SOLUBILITY IN WATER IS GOVERNED BY DYNAMIC EQUILIBRIUM PROCESSES, WHICH CAN BE REPRESENTED BY THEIR DISSOCIATION REACTIONS:

- For MOH :



- For $M(OH)_2$:



THE SOLUBILITY PRODUCT CONSTANT, K_{sp} , QUANTIFIES THE SOLUBILITY OF THESE COMPOUNDS AT EQUILIBRIUM. IT IS EXPRESSED AS:

- For MOH :

$$K_{sp} = [\text{M}^+][\text{OH}^-]$$

- For $M(OH)_2$:

$$K_{sp} = [\text{M}^{2+}][\text{OH}^-]^2$$

THE IDENTICAL K_{sp} VALUES SUGGEST THAT AT EQUILIBRIUM, THE PRODUCT OF ION CONCENTRATIONS REACHES THE SAME NUMERICAL VALUE FOR BOTH COMPOUNDS, DESPITE DIFFERENCES IN THEIR STOICHIOMETRY AND POTENTIAL SOLUBILITY BEHAVIORS.

IMPLICATIONS OF IDENTICAL K_{SP} VALUES FOR DIFFERENT HYDROXIDES

WHEN TWO HYDROXIDES HAVE THE SAME K_{SP}, BUT DIFFERENT CHEMICAL FORMULAS, SEVERAL INTERESTING IMPLICATIONS ARISE:

1. COMPARISON OF SOLUBILITIES

- DEFINITION: SOLUBILITY (s) REFERS TO THE MAXIMUM AMOUNT OF COMPOUND THAT CAN DISSOLVE IN WATER TO REACH EQUILIBRIUM.

- For MOH:

$$s_{\text{MOH}} = [\text{M}^{+}] = [\text{OH}^{-}]$$

BECAUSE ONE MOLE OF MOH DISSOLVES TO PRODUCE ONE MOLE OF M⁺ AND ONE MOLE OF OH⁻.

- For M(OH)₂:

$$s_{\text{M(OH)}_2} = [\text{M}^{2+}] = s_{\text{M(OH)}_2}$$

BUT THE HYDROXIDE ION CONCENTRATION WILL BE TWICE THE MOLAR SOLUBILITY BECAUSE EACH FORMULA UNIT PRODUCES TWO HYDROXIDE IONS:

$$[\text{OH}^{-}] = 2 \times s_{\text{M(OH)}_2}$$

2. SOLVING FOR SOLUBILITY

GIVEN THE K_{SP} VALUES:

- For MOH:

$$K_{\text{SP}} = s_{\text{MOH}} \times s_{\text{MOH}} = s_{\text{MOH}}^2$$

$$s_{\text{MOH}} = \sqrt{K_{\text{SP}}} = \sqrt{2.15 \times 10^{-12}} \approx 1.46 \times 10^{-6}, \text{ mol/L}$$

- For M(OH)₂:

$$K_{\text{SP}} = [\text{M}^{2+}][\text{OH}^{-}]^2$$

$$= s_{\text{M(OH)}_2} \times (2 s_{\text{M(OH)}_2})^2 = s_{\text{M(OH)}_2} \times 4 s_{\text{M(OH)}_2}^2 = 4 s_{\text{M(OH)}_2}^3$$

$$s_{\text{M(OH)}_2}^3 = \frac{K_{\text{SP}}}{4}$$

$$s_{\text{M(OH)}_2} = \left(\frac{K_{\text{SP}}}{4} \right)^{1/3} \approx \left(\frac{2.15 \times 10^{-12}}{4} \right)^{1/3}$$

$$\approx (5.375 \times 10^{-13})^{1/3} \approx 8.1 \times 10^{-5}, \text{ mol/L}$$

- HYDROXIDE ION CONCENTRATION:

$$[\text{OH}^{-}] = 2 \times s_{\text{M(OH)}_2} \approx 2 \times 8.1 \times 10^{-5} = 1.62 \times 10^{-4}, \text{ mol/L}$$

THIS COMPARISON REVEALS THAT $M(OH)_2$ IS SIGNIFICANTLY MORE SOLUBLE THAN MOH UNDER IDENTICAL K_{SP} CONDITIONS, OWING TO THE STOICHIOMETRY AND THE NATURE OF THE DISSOCIATION.

IMPACT OF INITIAL CATION CONCENTRATIONS ON SOLUBILITY EQUILIBRIA

THE INITIAL CONCENTRATIONS OF M^+ AND M^{2+} IONS IN SOLUTION CAN SIGNIFICANTLY INFLUENCE THEIR RESPECTIVE HYDROXIDE SOLUBILITIES THROUGH COMMON ION EFFECTS AND SHIFTS IN EQUILIBRIUM.

1. COMMON ION EFFECT

- THE PRESENCE OF M^+ OR M^{2+} IONS IN SOLUTION CAN SUPPRESS FURTHER DISSOLUTION OF MOH OR $M(OH)_2$, RESPECTIVELY. THIS OCCURS BECAUSE:
 - FOR MOH :
 - ELEVATED M^+ CONCENTRATIONS SHIFT THE EQUILIBRIUM LEFT, REDUCING SOLUBILITY.
 - FOR $M(OH)_2$:
 - ELEVATED M^{2+} CONCENTRATIONS ALSO SHIFT THE EQUILIBRIUM LEFT, DECREASING SOLUBILITY.

2. QUANTITATIVE ANALYSIS OF COMMON ION EFFECT

SUPPOSE INITIAL CONCENTRATIONS ARE:

- FOR M^+ : $([M^+]_{INITIAL})$
- FOR M^{2+} : $([M^{2+}]_{INITIAL})$

THE SOLUBILITY PRODUCT EXPRESSIONS BECOME:

- FOR MOH :
$$K_{SP} = ([M^+]_{INITIAL} + s) \times ([OH^-]_{INITIAL} + s)$$
BUT IF INITIAL M^+ IS SIGNIFICANT, THE EFFECTIVE SOLUBILITY (s_{MOH}) DECREASES.

- FOR $M(OH)_2$:
$$K_{SP} = ([M^{2+}]_{INITIAL} + s) \times (2s + [OH^-]_{INITIAL})^2$$
SIMILAR REASONING APPLIES.

3. PRACTICAL IMPLICATIONS

IN AQUEOUS ENVIRONMENTS WHERE M^+ OR M^{2+} ARE ALREADY PRESENT:

- THE SOLUBILITY OF MOH AND $M(OH)_2$ WILL BE REDUCED COMPARED TO PURE WATER.
- THIS EFFECT INFLUENCES PRECIPITATION AND DISSOLUTION PROCESSES IN NATURAL WATERS AND INDUSTRIAL APPLICATIONS.

THERMODYNAMIC AND KINETIC CONSIDERATIONS

1. THERMODYNAMICS

- THE K_{sp} VALUE REFLECTS THE THERMODYNAMIC EQUILIBRIUM; IT IS TEMPERATURE-DEPENDENT.
- THE PROVIDED K_{sp} (2.15×10^{-12}) IS INDICATIVE OF VERY LOW SOLUBILITY, CHARACTERISTIC OF INSOLUBLE OR SPARINGLY SOLUBLE HYDROXIDES.
- BY COMPARING K_{sp} VALUES, CHEMISTS CAN PREDICT WHICH HYDROXIDE WILL PRECIPITATE FIRST UNDER CERTAIN CONDITIONS.

2. KINETICS

- SOLUBILITY IS ALSO INFLUENCED BY REACTION KINETICS:
- NUCLEATION AND CRYSTAL GROWTH RATES DETERMINE HOW QUICKLY A HYDROXIDE PRECIPITATES.
- FOR HYDROXIDES WITH VERY LOW SOLUBILITY, SLOW DISSOLUTION CAN INFLUENCE THEIR BEHAVIOR IN SOLUTIONS, ESPECIALLY IN DYNAMIC SYSTEMS.

APPLICATIONS AND PRACTICAL RELEVANCE

UNDERSTANDING THE SOLUBILITY BEHAVIORS OF HYDROXIDES HAS BROAD APPLICATIONS:

- ENVIRONMENTAL CHEMISTRY:
 - PREDICTING HEAVY METAL HYDROXIDE PRECIPITATION IN WATER TREATMENT.
 - MANAGING WATER pH AND METAL ION CONCENTRATIONS TO PREVENT HARMFUL SOLUBILITY.
- INDUSTRIAL PROCESSES:
 - DESIGNING PROCESSES FOR METAL RECOVERY OR REMOVAL VIA HYDROXIDE PRECIPITATION.
 - CONTROLLING pH LEVELS TO OPTIMIZE SOLUBILITY AND PRECIPITATION.
- GEOCHEMISTRY:
 - EXPLAINING MINERAL FORMATION AND STABILITY IN NATURAL SETTINGS.
 - MODELING THE SOLUBILITY OF METAL HYDROXIDES IN GEOLOGICAL FORMATIONS.

SUMMARY AND KEY TAKEAWAYS

- BOTH MOH AND $\text{M}(\text{OH})_2$ SHARE AN IDENTICAL K_{sp} VALUE OF 2.15×10^{-12} , BUT THEIR SOLUBILITY PROFILES DIFFER DUE TO STOICHIOMETRY.
- SOLUB

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