

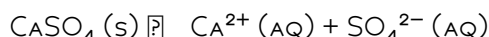
THE PRODUCT OF THE MOLARITIES OF THE DISSOLVED IONS, RAISED TO A POWER EQUAL TO THE ION'S COEFFICIENT

THE PRODUCT OF THE MOLARITIES OF THE DISSOLVED IONS, RAISED TO A POWER EQUAL TO THE ION'S COEFFICIENT IS A FUNDAMENTAL CONCEPT IN CHEMICAL EQUILIBRIUM, PARTICULARLY IN THE CONTEXT OF SOLUBILITY EQUILIBRIA AND THE CALCULATION OF SOLUBILITY PRODUCTS. IT SERVES AS A QUANTITATIVE MEASURE THAT ALLOWS CHEMISTS TO PREDICT WHETHER A SALT WILL PRECIPITATE FROM A SOLUTION OR REMAIN DISSOLVED UNDER SPECIFIC CONDITIONS. UNDERSTANDING THIS PRINCIPLE REQUIRES A COMPREHENSIVE EXPLORATION OF IONIC INTERACTIONS, EQUILIBRIUM PRINCIPLES, AND THE MATHEMATICAL FORMULATION OF SOLUBILITY PRODUCTS (K_{sp}). THIS ARTICLE AIMS TO DELVE DEEPLY INTO THIS CONCEPT, EXPLORING ITS THEORETICAL FOUNDATIONS, APPLICATIONS, AND SIGNIFICANCE IN CHEMICAL ANALYSIS.

INTRODUCTION TO IONIC EQUILIBRIA AND SOLUBILITY PRODUCTS

UNDERSTANDING DISSOLUTION AND PRECIPITATION

WHEN A SALT DISSOLVES IN WATER, IT DISSOCIATES INTO ITS CONSTITUENT IONS. FOR EXAMPLE, CONSIDER THE DISSOLUTION OF CALCIUM SULFATE:



THE PROCESS IS DYNAMIC, WITH IONS CONTINUALLY GOING INTO SOLUTION AND RECOMBINING TO FORM THE SOLID SALT. THE EXTENT OF DISSOLUTION DEPENDS ON THE INHERENT PROPERTIES OF THE SALT, TEMPERATURE, AND SOLUTION CONDITIONS. WHEN THE SOLUTION BECOMES SATURATED, THE RATES OF DISSOLUTION AND PRECIPITATION ARE EQUAL, ESTABLISHING AN EQUILIBRIUM.

THE CONCEPT OF THE SOLUBILITY PRODUCT CONSTANT (K_{sp})

THE SOLUBILITY PRODUCT CONSTANT, DENOTED AS K_{sp} , IS A KEY PARAMETER THAT CHARACTERIZES THE EQUILIBRIUM BETWEEN A SPARINGLY SOLUBLE SALT AND ITS IONS IN SOLUTION. IT IS DEFINED AS THE PRODUCT OF THE MOLAR CONCENTRATIONS OF THE IONS, EACH RAISED TO THE POWER OF ITS COEFFICIENT IN THE BALANCED DISSOLUTION EQUATION:

FOR CALCIUM SULFATE:

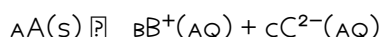
$$K_{sp} = [\text{Ca}^{2+}]^1 \times [\text{SO}_4^{2-}]^1$$

THIS EXPRESSION IS VALID AT EQUILIBRIUM, SERVING AS A THRESHOLD: IF THE IONIC PRODUCT EXCEEDS K_{sp} , PRECIPITATION OCCURS; IF IT IS LESS, THE SALT CONTINUES TO DISSOLVE; IF EQUAL, THE SYSTEM IS AT EQUILIBRIUM.

MATHEMATICAL FORMULATION OF THE IONIC PRODUCT

GENERAL EXPRESSION FOR IONIC PRODUCT

FOR ANY SALT WITH A DISSOLUTION REACTION:



THE IONIC PRODUCT (ALSO CALLED THE ION ACTIVITY PRODUCT, IAP, WHEN ACTIVITY COEFFICIENTS ARE CONSIDERED) IS GIVEN BY:

$$IAP = [B^+]^B \times [C^{2-}]^C$$

THIS PRODUCT IS DIRECTLY COMPARABLE TO K_{sp} AT EQUILIBRIUM; THEIR RATIO INDICATES THE SYSTEM'S TENDENCY TO PRECIPITATE OR DISSOLVE.

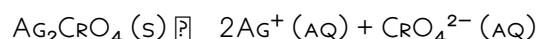
RAISING MOLARITIES TO THE POWER OF COEFFICIENTS

THE CORE PRINCIPLE IN THE TITLE REFERS TO THE MATHEMATICAL OPERATION OF RAISING EACH IONIC CONCENTRATION TO THE POWER CORRESPONDING TO ITS COEFFICIENT IN THE BALANCED DISSOLUTION REACTION. THIS IS CRUCIAL BECAUSE:

- IT ACCOUNTS FOR THE STOICHIOMETRY OF THE DISSOLUTION.
- IT ENSURES THE IONIC PRODUCT ACCURATELY REFLECTS THE EQUILIBRIUM STATE.
- IT ALLOWS FOR THE PREDICTION OF PRECIPITATION OR DISSOLUTION BASED ON CONCENTRATION MEASUREMENTS.

FOR EXAMPLE:

- FOR Ag_2CrO_4 :



THE IONIC PRODUCT IS:

$$IAP = [Ag^+]^2 \times [CrO_4^{2-}]^1$$

AT EQUILIBRIUM, THIS EQUALS THE SOLUBILITY PRODUCT CONSTANT, K_{sp} .

SIGNIFICANCE OF THE ION COEFFICIENTS IN THE CALCULATION

WHY RAISE CONCENTRATIONS TO THE POWER OF THEIR COEFFICIENTS?

THE REASON FOR RAISING ION CONCENTRATIONS TO THE POWER OF THEIR COEFFICIENTS STEMS FROM THE LAW OF MASS ACTION, WHICH STATES THAT AT EQUILIBRIUM, THE RATIO OF THE PRODUCT OF THE ACTIVITIES OF PRODUCTS TO REACTANTS REMAINS CONSTANT. WHEN TRANSLATING THIS INTO MOLAR CONCENTRATIONS (ASSUMING IDEAL BEHAVIOR), THE EXPONENTS REFLECT THE MOLECULAR RATIOS IN THE DISSOCIATION REACTION.

THIS MATHEMATICAL OPERATION:

- ENSURES THE CALCULATION ACCURATELY REFLECTS THE STOICHIOMETRY.
- MAINTAINS CONSISTENCY WITH THERMODYNAMIC PRINCIPLES.
- PROVIDES A STANDARDIZED METHOD TO COMPARE IONIC PRODUCTS WITH K_{sp} .

IMPLICATIONS IN PRECIPITATION AND DISSOLUTION

- IF THE IONIC PRODUCT EXCEEDS K_{sp} , THE SOLUTION IS SUPERSATURATED, AND A PRECIPITATE WILL FORM.
- IF IT EQUALS K_{sp} , THE SOLUTION IS SATURATED, AND THE SYSTEM IS AT EQUILIBRIUM.
- IF IT IS LESS THAN K_{sp} , THE SOLUTION IS UNSATURATED, AND DISSOLUTION CAN OCCUR.

APPLICATIONS OF THE CONCEPT IN CHEMICAL ANALYSIS

PREDICTING PRECIPITATION REACTIONS

BY CALCULATING THE IONIC PRODUCT RAISED TO THE APPROPRIATE POWERS, CHEMISTS CAN:

- DETERMINE WHETHER A PRECIPITATE WILL FORM UNDER SPECIFIC CONDITIONS.
- CONTROL THE PURITY OF SOLUTIONS BY ADJUSTING CONCENTRATIONS.
- DESIGN SEPARATION PROCESSES IN ANALYTICAL CHEMISTRY.

CALCULATING SOLUBILITY AND K_{sp}

IN EXPERIMENTAL SETTINGS, THE IONIC CONCENTRATIONS ARE MEASURED, AND THE IONIC PRODUCT IS COMPUTED:

1. MEASURE MOLAR CONCENTRATIONS OF IONS IN SOLUTION.
2. RAISE EACH CONCENTRATION TO THE POWER OF ITS COEFFICIENT.
3. MULTIPLY THESE VALUES TO OBTAIN THE IONIC PRODUCT.
4. COMPARE WITH KNOWN K_{sp} TO ASSESS SATURATION STATUS.

THIS PROCESS ENABLES THE DETERMINATION OF UNKNOWN SOLUBILITY PRODUCTS OR THE VERIFICATION OF EQUILIBRIUM CONDITIONS.

COMPLEX ION FORMATION AND ITS EFFECT ON IONIC PRODUCTS

IN SOME CASES, IONS MAY FORM COMPLEXES, ALTERING THE FREE ION CONCENTRATIONS AND THUS THE IONIC PRODUCT. CHEMISTS MUST ACCOUNT FOR COMPLEXATION EQUILIBRIA WHEN APPLYING THE PRINCIPLE, OFTEN MODIFYING THE BASIC CALCULATION TO INCLUDE ACTIVITY COEFFICIENTS OR COMPLEX STABILITY CONSTANTS.

LIMITATIONS AND CONSIDERATIONS

ROLE OF ACTIVITY COEFFICIENTS

- IN REAL SOLUTIONS, IONS DO NOT BEHAVE IDEALLY.
- ACTIVITY COEFFICIENTS ACCOUNT FOR INTERACTIONS BETWEEN IONS.
- THE IONIC PRODUCT SHOULD IDEALLY BE CALCULATED USING ACTIVITIES RATHER THAN MOLAR CONCENTRATIONS FOR ACCURACY.

TEMPERATURE DEPENDENCE

- K_{sp} VARIES WITH TEMPERATURE.
- THE IONIC PRODUCT MUST BE COMPARED AT THE SAME TEMPERATURE TO MAKE VALID PREDICTIONS.

Non-Ideal Solutions and Ionic Strength

- High ionic strength solutions deviate significantly from ideal behavior.
- Corrections are necessary for precise calculations.

Conclusion

The concept of raising the molarities of dissolved ions to a power equal to their coefficients in the dissolution reaction is a cornerstone of chemical equilibrium analysis. It ensures that the calculation of the ionic product accurately reflects the stoichiometry of the dissociation process, providing a reliable tool for predicting solubility, understanding precipitation phenomena, and designing chemical processes. Mastery of this principle allows chemists to quantitatively analyze complex systems, optimize conditions for purification or separation, and deepen their understanding of ionic interactions in solution. Recognizing the importance of this mathematical operation emphasizes the elegance and consistency of chemical principles rooted in the law of mass action and thermodynamics.

Frequently Asked Questions

What does the term 'product of the molarities of the dissolved ions raised to a power equal to the ion's coefficient' refer to in chemistry?

It refers to the expression used in calculating the ion activity product (IAP) for a sparingly soluble compound, where each ion's molarity is raised to the power of its coefficient in the balanced dissolution equation.

How is the ion activity product (IAP) different from the solubility product constant (K_{sp})?

While both involve the product of ion concentrations raised to their coefficients, K_{sp} is a constant at a given temperature representing equilibrium, whereas the IAP can vary with ion concentrations and indicates whether a solution is saturated, undersaturated, or supersaturated.

Why is it important to raise the molarity of each ion to the power of its coefficient in solubility calculations?

Because the equilibrium expression for sparingly soluble salts includes each ion's concentration raised to its stoichiometric coefficient, ensuring the calculation accurately reflects the ionic interactions and the conditions of saturation.

Can you give an example of calculating the product of molarities raised to their coefficients for a compound?

Yes. For calcium carbonate, $\text{CaCO}_3(\text{s}) \rightleftharpoons \text{Ca}^{2+}(\text{aq}) + \text{CO}_3^{2-}(\text{aq})$, the product is $[\text{Ca}^{2+}]^1 \times [\text{CO}_3^{2-}]^1 = [\text{Ca}^{2+}][\text{CO}_3^{2-}]$. Since coefficients are 1, the product is simply the product of their molarities.

How does the ion product relate to the solubility product constant (K_{sp})?

The ion product (IAP) is calculated similarly to K_{sp}; if IAP exceeds K_{sp}, the solution is supersaturated and precipitation may occur; if it's less, the solution is unsaturated.

WHAT ROLE DOES TEMPERATURE PLAY IN THE CALCULATION OF THIS PRODUCT?

TEMPERATURE AFFECTS THE SOLUBILITY AND, CONSEQUENTLY, THE MOLARITIES OF IONS, AS WELL AS THE VALUE OF K_{sp} . HOWEVER, THE CALCULATION OF THE PRODUCT ITSELF DEPENDS ON CURRENT ION CONCENTRATIONS, WHICH VARY WITH TEMPERATURE.

IS THE 'PRODUCT OF MOLARITIES RAISED TO THE POWER OF THE ION'S COEFFICIENT' APPLICABLE ONLY TO SOLUBLE SALTS?

NO, IT'S PRIMARILY USED FOR SPARINGLY SOLUBLE SALTS AND IN SOLUBILITY EQUILIBRIA, BUT SIMILAR PRINCIPLES APPLY IN OTHER EQUILIBRIUM CALCULATIONS INVOLVING IONIC COMPOUNDS.

HOW CAN UNDERSTANDING THIS PRODUCT HELP IN PREDICTING PRECIPITATION REACTIONS?

BY CALCULATING THE ION PRODUCT AND COMPARING IT TO K_{sp} , CHEMISTS CAN PREDICT WHETHER A SALT WILL PRECIPITATE OUT OF SOLUTION UNDER GIVEN CONDITIONS.

WHAT ARE COMMON MISCONCEPTIONS ABOUT THE PRODUCT OF MOLARITIES RAISED TO ION COEFFICIENTS?

A COMMON MISCONCEPTION IS CONFUSING THE ION PRODUCT WITH THE SOLUBILITY PRODUCT CONSTANT; THE ION PRODUCT VARIES WITH CONCENTRATION, WHILE K_{sp} IS A FIXED VALUE AT A SPECIFIC TEMPERATURE REPRESENTING EQUILIBRIUM.

ADDITIONAL RESOURCES

THE PRODUCT OF THE MOLARITIES OF THE DISSOLVED IONS, RAISED TO A POWER EQUAL TO THE ION'S COEFFICIENT IS A FUNDAMENTAL CONCEPT IN CHEMISTRY, PARTICULARLY IN THE STUDY OF CHEMICAL EQUILIBRIA AND SOLUBILITY. THIS EXPRESSION IS CENTRAL TO UNDERSTANDING HOW IONIC COMPOUNDS BEHAVE IN AQUEOUS SOLUTIONS AND FORMS THE BASIS FOR CALCULATING SOLUBILITY PRODUCTS, WHICH IN TURN INFLUENCE A WIDE RANGE OF PRACTICAL APPLICATIONS—FROM PHARMACEUTICAL FORMULATIONS TO ENVIRONMENTAL CHEMISTRY. MASTERY OF THIS CONCEPT PROVIDES CHEMISTS AND STUDENTS ALIKE WITH A POWERFUL TOOL TO PREDICT WHETHER A COMPOUND WILL PRECIPITATE FROM SOLUTION OR REMAIN DISSOLVED UNDER SPECIFIC CONDITIONS.

UNDERSTANDING ION PRODUCTS AND THEIR SIGNIFICANCE

WHAT IS THE ION PRODUCT?

AT ITS CORE, THE PRODUCT OF THE MOLARITIES OF THE DISSOLVED IONS, RAISED TO A POWER EQUAL TO THE ION'S COEFFICIENT, IS AN EXPRESSION USED TO DESCRIBE THE IONIC EQUILIBRIUM OF SPARINGLY SOLUBLE SALTS. WHEN A SALT DISSOLVES IN WATER, IT DISSOCIATES INTO ITS CONSTITUENT IONS:



THE ION PRODUCT (OFTEN DENOTED AS Q) IS CALCULATED BY MULTIPLYING THE MOLAR CONCENTRATIONS OF THE IONS, EACH RAISED TO THE POWER CORRESPONDING TO THEIR COEFFICIENTS IN THE DISSOCIATION EQUATION:

$$Q = [\text{A}^{x+}]^1 \times [\text{B}^{-}]^x$$

WHY IS IT IMPORTANT?

- PREDICTING PRECIPITATION: COMPARING Q WITH THE SOLUBILITY PRODUCT CONSTANT (K_{sp}) TELLS US WHETHER A SALT WILL PRECIPITATE OR DISSOLVE FURTHER.

- UNDERSTANDING EQUILIBRIUM DYNAMICS: IT HELPS IN ANALYZING THE DIRECTION OF CHANGE IN A SATURATED SOLUTION.
- DESIGNING CHEMICAL PROCESSES: IT GUIDES IN CONTROLLING CONDITIONS TO FAVOR OR PREVENT PRECIPITATION IN INDUSTRIAL SETTINGS.

THE ROLE OF THE SOLUBILITY PRODUCT CONSTANT (K_{sp})

DEFINITION AND CALCULATION

THE SOLUBILITY PRODUCT CONSTANT (K_{sp}) IS THE EQUILIBRIUM CONSTANT FOR THE DISSOLUTION OF A SPARINGLY SOLUBLE SALT:



$$[K_{sp} = [\text{A}^{x+}]^x \times [\text{B}^{-}]^x]$$

NOTE THAT THE MOLAR CONCENTRATIONS ARE THOSE AT EQUILIBRIUM, SPECIFICALLY IN A SATURATED SOLUTION.

COMPARING Q AND K_{sp}

- WHEN $(Q < K_{sp})$: THE SOLUTION IS UNSATURATED; MORE SALT CAN DISSOLVE.
- WHEN $(Q = K_{sp})$: THE SOLUTION IS SATURATED; EQUILIBRIUM IS ESTABLISHED.
- WHEN $(Q > K_{sp})$: THE SOLUTION IS SUPERSATURATED; PRECIPITATION IS LIKELY TO OCCUR.

THIS COMPARISON IS CRUCIAL FOR CONTROLLING AND MANIPULATING SOLUTION CONDITIONS.

CALCULATING THE PRODUCT OF IONIC MOLARITIES

STEP-BY-STEP PROCESS

1. IDENTIFY THE DISSOCIATION EQUATION:

WRITE THE BALANCED DISSOCIATION FOR THE SALT.

2. DETERMINE ION CONCENTRATIONS:

OBTAIN MOLARITIES OF EACH ION IN THE SOLUTION AT EQUILIBRIUM.

3. APPLY THE EXPONENTIATION:

RAISE EACH ION CONCENTRATION TO THE POWER OF ITS COEFFICIENT IN THE DISSOCIATION EQUATION.

4. CALCULATE THE PRODUCT:

MULTIPLY ALL THESE VALUES TOGETHER TO GET THE ION PRODUCT (Q).

EXAMPLE: SILVER CHLORIDE (AgCl)

DISSOCIATION:



- COEFFICIENTS ARE BOTH 1, SO:

$$[Q = [\text{Ag}^{+}] \times [\text{Cl}^{-}]]$$

IF IN A SATURATED SOLUTION:

$$\begin{aligned} [\text{Ag}^+] &= 1.3 \times 10^{-5} \text{ M} \\ [\text{Cl}^-] &= 1.3 \times 10^{-5} \text{ M} \end{aligned}$$

THEN:

$$Q = (1.3 \times 10^{-5}) \times (1.3 \times 10^{-5}) = 1.69 \times 10^{-10}$$

COMPARE WITH K_{sp} TO DETERMINE THE SOLUTION'S STATUS.

RAISING THE MOLARITIES TO THE POWER OF THE ION'S COEFFICIENT

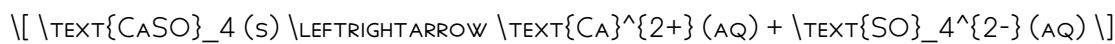
SIGNIFICANCE OF THE COEFFICIENT

IN DISSOCIATION EQUATIONS, COEFFICIENTS INDICATE THE RATIO OF IONS PRODUCED PER FORMULA UNIT DISSOLVED. WHEN CALCULATING THE IONIC PRODUCT:

- EACH ION'S MOLARITY IS RAISED TO THE POWER OF ITS COEFFICIENT.
- THIS ACCOUNTS FOR THE STOICHIOMETRY OF THE DISSOCIATION.

PRACTICAL IMPLICATIONS

- FOR SALTS WITH MULTIPLE IONS AND HIGHER COEFFICIENTS, THE CALCULATION BECOMES MORE COMPLEX BUT FOLLOWS THE SAME PRINCIPLES.
- FOR EXAMPLE, CALCIUM SULFATE (CaSO_4) DISSOCIATES AS:



- BOTH IONS HAVE COEFFICIENTS OF 1, SO THE PRODUCT IS SIMPLY $[\text{Ca}^{2+}] \times [\text{SO}_4^{2-}]$.
- FOR MORE COMPLEX SALTS LIKE ALUMINUM HYDROXIDE:



- THE IONIC PRODUCT IS:

$$Q = [\text{Al}^{3+}] \times [\text{OH}^-]^3$$

APPLICATIONS AND PRACTICAL CONSIDERATIONS

PRECIPITATION AND PURIFICATION

- BY MANIPULATING ION CONCENTRATIONS, CHEMISTS CAN INDUCE OR PREVENT PRECIPITATION.
- FOR EXAMPLE, ADDING COMMON IONS CAN SHIFT THE IONIC PRODUCT ABOVE OR BELOW K_{sp} , CAUSING SELECTIVE PRECIPITATION.

ENVIRONMENTAL CHEMISTRY

- UNDERSTANDING THE SOLUBILITY OF METAL SALTS INFLUENCES WATER TREATMENT PROCESSES.
- THE FORMATION OF INSOLUBLE SALTS CAN REMOVE POLLUTANTS.

PHARMACEUTICAL CHEMISTRY

- SOLUBILITY INFLUENCES DRUG FORMULATION AND BIOAVAILABILITY.

- CALCULATIONS INVOLVING THE IONIC PRODUCT HELP OPTIMIZE CONDITIONS FOR DRUG STABILITY.

INDUSTRIAL PROCESSES

- SCALE FORMATION IN PIPES AND REACTORS DEPENDS ON IONIC PRODUCT CALCULATIONS.
- PROPER CONTROL OF ION CONCENTRATIONS PREVENTS EQUIPMENT FOULING.

FACTORS AFFECTING THE IONIC PRODUCT

TEMPERATURE

- USUALLY, K_{sp} VARIES WITH TEMPERATURE, INFLUENCING THE IONIC PRODUCT.
- HIGHER TEMPERATURES CAN INCREASE OR DECREASE SOLUBILITY.

COMMON IONS

- THE PRESENCE OF COMMON IONS SHIFTS EQUILIBRIUM (LE CHATELIER'S PRINCIPLE), AFFECTING THE IONIC PRODUCT AND SOLUBILITY.

pH OF THE SOLUTION

- ACIDIC OR BASIC CONDITIONS ALTER ION CONCENTRATIONS, ESPECIALLY FOR SALTS INVOLVING WEAK ACIDS OR BASES.

SUMMARY OF KEY POINTS

- THE PRODUCT OF THE MOLARITIES OF THE DISSOLVED IONS, RAISED TO A POWER EQUAL TO THE ION'S COEFFICIENT, IS A VITAL CALCULATION FOR UNDERSTANDING SOLUBILITY AND EQUILIBRIUM.
- THIS PRODUCT, COMPARED TO THE K_{sp} , INDICATES WHETHER A SALT WILL PRECIPITATE OR DISSOLVE.
- ACCURATE CALCULATION REQUIRES KNOWLEDGE OF THE DISSOCIATION EQUATION, ION CONCENTRATIONS, AND STOICHIOMETRY.
- THE CONCEPT IS WIDELY APPLICABLE ACROSS CHEMISTRY FIELDS, FROM INDUSTRIAL MANUFACTURING TO ENVIRONMENTAL MANAGEMENT.

FINAL THOUGHTS

MASTERING THE CALCULATION AND INTERPRETATION OF THE PRODUCT OF THE MOLARITIES OF THE DISSOLVED IONS, RAISED TO A POWER EQUAL TO THE ION'S COEFFICIENT, IS ESSENTIAL FOR ANYONE WORKING WITH IONIC SOLUTIONS. WHETHER PREDICTING PRECIPITATION, DESIGNING CHEMICAL PROCESSES, OR UNDERSTANDING ENVIRONMENTAL PHENOMENA, THIS FUNDAMENTAL CONCEPT PROVIDES A QUANTITATIVE FRAMEWORK THAT BRIDGES THEORETICAL CHEMISTRY WITH REAL-WORLD APPLICATIONS. BY DEVELOPING A SOLID GRASP OF HOW TO PERFORM AND ANALYZE THESE CALCULATIONS, CHEMISTS CAN MAKE INFORMED DECISIONS THAT INFLUENCE SAFETY, EFFICIENCY, AND INNOVATION ACROSS COUNTLESS SCIENTIFIC AND INDUSTRIAL DOMAINS.

The Product Of The Molarities Of The Dissolved Ions Raised To A Power Equal To The Ion S Coefficient

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