

ORDER THESE CHEMICAL SPECIES BY INCREASING PH OF AN 0.1M AQUEOUS SOLUTION OF EACH. THAT IS, IMAGINE MAKING

ORDER THESE CHEMICAL SPECIES BY INCREASING PH OF AN 0.1M AQUEOUS SOLUTION OF EACH. THAT IS, IMAGINE MAKING A SERIES OF SOLUTIONS WITH DIFFERENT CHEMICAL SPECIES AT A CONCENTRATION OF 0.1M EACH. UNDERSTANDING HOW VARIOUS SUBSTANCES INFLUENCE THE PH OF THEIR AQUEOUS SOLUTIONS IS FUNDAMENTAL IN CHEMISTRY, ESPECIALLY WHEN PREDICTING THE ACIDITY OR BASICITY OF UNKNOWN SOLUTIONS OR DESIGNING BUFFERS. THIS ARTICLE PROVIDES A COMPREHENSIVE GUIDE TO RANKING CHEMICAL SPECIES BASED ON THEIR PH LEVELS WHEN DISSOLVED IN WATER AT EQUAL MOLAR CONCENTRATIONS, ALONG WITH EXPLANATIONS OF THE UNDERLYING PRINCIPLES GOVERNING THESE BEHAVIORS.

INTRODUCTION TO PH AND CHEMICAL SPECIES IN AQUEOUS SOLUTIONS

THE PH SCALE MEASURES THE ACIDITY OR ALKALINITY OF A SOLUTION, RANGING FROM 0 (MOST ACIDIC) TO 14 (MOST BASIC), WITH 7 BEING NEUTRAL. WHEN A CHEMICAL SPECIES DISSOLVES IN WATER, IT CAN EITHER DONATE PROTONS (H^+ IONS), ACCEPT PROTONS, OR REMAIN NEUTRAL, THEREBY AFFECTING THE PH OF THE SOLUTION.

KEY FACTORS INFLUENCING THE PH OF A SOLUTION INCLUDE:

- WHETHER THE COMPOUND IS AN ACID, BASE, OR NEUTRAL
- THE STRENGTH OF ACIDS AND BASES (STRONG VS. WEAK)
- THE DEGREE OF IONIZATION/DISSOCIATION IN WATER
- THE MOLECULAR PROPERTIES OF THE SPECIES INVOLVED

IN AN AQUEOUS SOLUTION, STRONG ACIDS AND STRONG BASES DISSOCIATE COMPLETELY, LEADING TO SIGNIFICANTLY DIFFERENT PH VALUES COMPARED TO WEAK ACIDS AND WEAK BASES, WHICH ONLY PARTIALLY DISSOCIATE.

UNDERSTANDING ACIDIC AND BASIC CHEMICAL SPECIES

TO ACCURATELY ORDER CHEMICAL SPECIES BY INCREASING PH, WE NEED TO UNDERSTAND THE BEHAVIOR OF COMMON ACIDS AND BASES IN WATER.

STRONG ACIDS

- FULLY DISSOCIATE IN WATER
- EXAMPLES: HYDROCHLORIC ACID (HCl), HYDROBROMIC ACID (HBr), HYDROIODIC ACID (HI), NITRIC ACID (HNO_3), SULFURIC ACID (H_2SO_4)
- RESULT IN HIGH CONCENTRATION OF H^+ IONS; THUS, VERY LOW PH (<3)

WEAK ACIDS

- PARTIALLY DISSOCIATE
- EXAMPLES: ACETIC ACID (CH_3COOH), FORMIC ACID ($HCOOH$), OXALIC ACID

- RESULT IN MODERATE H^+ CONCENTRATIONS; pH TYPICALLY BETWEEN 3 AND 6

STRONG BASES

- FULLY DISSOCIATE IN WATER
- EXAMPLES: SODIUM HYDROXIDE ($NaOH$), POTASSIUM HYDROXIDE (KOH), BARIUM HYDROXIDE ($Ba(OH)_2$)
- LEAD TO HIGH OH^- CONCENTRATIONS; pH TYPICALLY ABOVE 12

WEAK BASES

- PARTIALLY ACCEPT PROTONS
- EXAMPLES: AMMONIA (NH_3), METHYLAMINE, ANILINE
- RESULT IN MODERATE OH^- CONCENTRATIONS; pH TYPICALLY BETWEEN 8 AND 11

NEUTRAL COMPOUNDS

- NEITHER DONATE NOR ACCEPT PROTONS
- EXAMPLES: PURE WATER, SALTS DERIVED FROM STRONG ACIDS AND STRONG BASES (E.G., $NaCl$)
- pH OF AROUND 7

ORDER OF CHEMICAL SPECIES BY INCREASING pH

WHEN 0.1M SOLUTIONS OF VARIOUS CHEMICAL SPECIES ARE PREPARED, THEIR pH VALUES REFLECT THEIR ACID-BASE STRENGTHS. THE ORDER FROM MOST ACIDIC (LOWEST pH) TO MOST BASIC (HIGHEST pH) GENERALLY FOLLOWS THE STRENGTH OF THEIR ACID/BASE BEHAVIOR.

TYPICAL ORDER:

1. STRONG ACIDS (LOWEST pH)
2. WEAK ACIDS
3. NEUTRAL SOLUTIONS
4. WEAK BASES
5. STRONG BASES (HIGHEST pH)

STEP-BY-STEP EXAMPLE: RANKING COMMON CHEMICAL SPECIES

LET'S ANALYZE SPECIFIC COMMON SUBSTANCES AT 0.1M CONCENTRATION TO ILLUSTRATE THEIR RELATIVE pH VALUES.

STRONG ACIDS

- EXAMPLE: 0.1M HCl
- DISSOCIATES COMPLETELY

- $\text{pH} \approx 1$

WEAK ACIDS

- EXAMPLE: 0.1M ACETIC ACID (CH_3COOH)
- PARTIALLY DISSOCIATES
- APPROXIMATE $\text{pH} \approx 2.9\text{--}3.0$

NEUTRAL COMPOUNDS

- EXAMPLE: 0.1M NaCl
- DOES NOT AFFECT pH SIGNIFICANTLY
- $\text{pH} \approx 7$

WEAK BASES

- EXAMPLE: 0.1M AMMONIA (NH_3)
- PARTIALLY ACCEPTS PROTONS
- $\text{pH} \approx 11.2$

STRONG BASES

- EXAMPLE: 0.1M NaOH
- FULLY DISSOCIATES
- $\text{pH} \approx 13$

THUS, THE ORDER FROM LOWEST TO HIGHEST pH IS:

$\text{HCl} < \text{ACETIC ACID} < \text{NaCl} \approx 7 < \text{AMMONIA} < \text{NaOH}$

FACTORS AFFECTING pH BEYOND CONCENTRATION

WHILE THE ABOVE EXAMPLES ASSUME IDEAL BEHAVIOR, SEVERAL FACTORS CAN INFLUENCE THE pH OF SOLUTIONS:

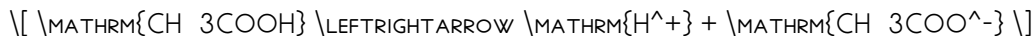
- IONIZATION CONSTANT (K_a OR K_b): WEAK ACIDS/BASES HAVE SPECIFIC DISSOCIATION CONSTANTS THAT DETERMINE THEIR DEGREE OF IONIZATION.
- TEMPERATURE: INCREASING TEMPERATURE CAN SHIFT DISSOCIATION EQUILIBRIA, AFFECTING pH .
- PRESENCE OF OTHER IONS: IONIC STRENGTH CAN INFLUENCE ACTIVITY COEFFICIENTS, SLIGHTLY AFFECTING pH .
- POLYPROTIC ACIDS: ACIDS LIKE SULFURIC ACID RELEASE MORE THAN ONE PROTON, AFFECTING pH CALCULATIONS.

CALCULATING pH FOR WEAK ACIDS AND BASES

CALCULATING THE pH OF WEAK ACIDS AND BASES INVOLVES EQUILIBRIUM EXPRESSIONS:

WEAK ACID EXAMPLE: ACETIC ACID (CH_3COOH)

THE DISSOCIATION:



THE EXPRESSION FOR K_a :

$$K_a = \frac{[\text{H}^+][\text{CH}_3\text{COO}^-]}{[\text{CH}_3\text{COOH}]}$$

ASSUMING INITIAL CONCENTRATION $C = 0.1 \text{ M}$ AND x AS THE CONCENTRATION OF H^+ :

$$K_a = \frac{x^2}{C - x}$$

SINCE K_a FOR ACETIC ACID IS ABOUT (1.8×10^{-5}) , SOLVING FOR x GIVES THE H^+ CONCENTRATION, AND THUS pH:

$$\text{pH} = -\log [\text{H}^+]$$

SIMILARLY, FOR BASES, THE K_b EXPRESSION APPLIES.

IMPLICATIONS FOR LABORATORY AND INDUSTRIAL APPLICATIONS

UNDERSTANDING HOW TO ORDER CHEMICAL SPECIES BY pH IS CRUCIAL IN VARIOUS APPLICATIONS:

- BUFFER PREPARATION: SELECTING WEAK ACIDS/BASES TO BUFFER SOLUTIONS AT DESIRED pH
- ENVIRONMENTAL CHEMISTRY: ASSESSING SOIL OR WATER ACIDITY
- PHARMACEUTICALS: DESIGNING DRUG FORMULATIONS WITH SPECIFIC pH REQUIREMENTS
- CHEMICAL MANUFACTURING: CONTROLLING pH DURING REACTIONS AND PROCESSING

SUMMARY: KEY TAKEAWAYS

- THE pH OF A SOLUTION DEPENDS HEAVILY ON WHETHER THE COMPOUND IS AN ACID, BASE, OR NEUTRAL, AND ON ITS STRENGTH.
- STRONG ACIDS AND BASES HAVE pH VALUES NEAR 1 AND 13, RESPECTIVELY, DUE TO COMPLETE DISSOCIATION.
- WEAK ACIDS AND BASES HAVE INTERMEDIATE pH VALUES, INFLUENCED BY THEIR DISSOCIATION CONSTANTS.
- NEUTRAL SALTS LIKE NaCl DO NOT SIGNIFICANTLY AFFECT pH, MAINTAINING NEAR 7.
- THE ORDER OF CHEMICAL SPECIES BY INCREASING pH AT 0.1M CONCENTRATION GENERALLY FOLLOWS: STRONG ACIDS < WEAK ACIDS < NEUTRAL SALTS < WEAK BASES < STRONG BASES.

CONCLUSION

IN CONCLUSION, WHEN ASKED TO ORDER CHEMICAL SPECIES BY INCREASING pH OF A 0.1M AQUEOUS SOLUTION, IT'S ESSENTIAL TO UNDERSTAND THEIR ACID-BASE NATURE AND DISSOCIATION BEHAVIOR. RECOGNIZING THE DIFFERENCE BETWEEN STRONG AND WEAK ACIDS/BASES AND THEIR RESPECTIVE DISSOCIATION CONSTANTS ALLOWS FOR ACCURATE PREDICTIONS OF THEIR pH VALUES. THIS UNDERSTANDING IS VITAL FOR CHEMISTS, ENVIRONMENTAL SCIENTISTS, AND INDUSTRY PROFESSIONALS WHO MANIPULATE SOLUTIONS TO ACHIEVE SPECIFIC pH CONDITIONS FOR VARIOUS APPLICATIONS.

BY MASTERING THESE PRINCIPLES, YOU CAN CONFIDENTLY DETERMINE THE RELATIVE pH LEVELS OF DIVERSE CHEMICAL SPECIES AND EFFECTIVELY DESIGN SOLUTIONS TAILORED TO YOUR NEEDS.

FREQUENTLY ASKED QUESTIONS

HOW DOES THE NATURE OF THE CHEMICAL SPECIES AFFECT THE pH OF A 0.1M AQUEOUS SOLUTION?

THE NATURE DETERMINES WHETHER THE SPECIES IS ACIDIC, BASIC, OR NEUTRAL. ACIDIC SPECIES RELEASE H^+ IONS, LOWERING pH; BASIC SPECIES RELEASE OH^- IONS, INCREASING pH; NEUTRAL SPECIES HAVE LITTLE EFFECT ON pH.

WHICH CHEMICAL SPECIES AMONG ACIDS, BASES, OR SALTS WILL HAVE THE HIGHEST pH IN A 0.1M AQUEOUS SOLUTION?

STRONG BASES, SUCH AS NaOH, WILL HAVE THE HIGHEST pH, TYPICALLY CLOSE TO 13-14, COMPARED TO ACIDS OR NEUTRAL SALTS AT THE SAME CONCENTRATION.

WHAT IS THE EXPECTED ORDER OF pH FOR THE FOLLOWING SPECIES: HCl, NaOH, NH_4Cl , AND CH_3COOH IN 0.1M SOLUTIONS?

ORDER FROM LOWEST TO HIGHEST pH: HCl (ACIDIC), CH_3COOH (WEAK ACID), NH_4Cl (WEAKLY ACIDIC SALT), NaOH (STRONG BASE).

WHY DOES A WEAK ACID LIKE ACETIC ACID HAVE A HIGHER pH THAN A STRONG ACID AT THE SAME CONCENTRATION?

BECAUSE WEAK ACIDS DO NOT IONIZE COMPLETELY, RESULTING IN FEWER H^+ IONS IN SOLUTION, THUS A HIGHER pH COMPARED TO STRONG ACIDS WHICH FULLY IONIZE.

HOW CAN YOU PREDICT THE ORDER OF INCREASING pH FOR VARIOUS CHEMICAL SPECIES IN AQUEOUS SOLUTIONS?

BY CONSIDERING THEIR ACID-BASE PROPERTIES, STRENGTH OF ACIDS/BASES, AND THEIR DISSOCIATION CONSTANTS (K_A OR K_B), YOU CAN RANK THEM ACCORDINGLY.

WOULD A SALT DERIVED FROM A STRONG ACID AND STRONG BASE HAVE A NEUTRAL pH IN SOLUTION? WHY?

YES, SUCH SALTS TYPICALLY PRODUCE A NEUTRAL SOLUTION WITH pH CLOSE TO 7 BECAUSE THEIR IONS DO NOT HYDROLYZE SIGNIFICANTLY IN WATER.

IF MAKING SOLUTIONS OF DIFFERENT CHEMICAL SPECIES AT 0.1M, HOW WOULD YOU EXPERIMENTALLY DETERMINE THEIR pH ORDER?

USE A CALIBRATED pH METER TO MEASURE THE pH OF EACH SOLUTION AND THEN COMPARE THE VALUES TO ESTABLISH THE INCREASING ORDER.

ADDITIONAL RESOURCES

ORDER THESE CHEMICAL SPECIES BY INCREASING pH OF AN 0.1M AQUEOUS SOLUTION OF EACH: AN IN-DEPTH INVESTIGATION

INTRODUCTION

UNDERSTANDING HOW DIFFERENT CHEMICAL SPECIES INFLUENCE THE pH OF AQUEOUS SOLUTIONS IS FUNDAMENTAL TO THE FIELDS OF CHEMISTRY, ENVIRONMENTAL SCIENCE, AND ENGINEERING. THE pH OF A SOLUTION—A MEASURE OF ITS ACIDITY OR ALKALINITY—IS DETERMINED BY THE NATURE AND STRENGTH OF THE SOLUTES DISSOLVED IN WATER. WHEN PREPARING 0.1 M SOLUTIONS OF VARIOUS CHEMICAL SPECIES, THEIR INTRINSIC ACID-BASE PROPERTIES DICTATE WHETHER THE SOLUTION WILL BE ACIDIC, NEUTRAL, OR BASIC, THEREBY AFFECTING THE pH.

THIS ARTICLE AIMS TO SYSTEMATICALLY ANALYZE AND ORDER A SET OF CHEMICAL SPECIES BASED ON THEIR EXPECTED pH WHEN DISSOLVED AT A CONCENTRATION OF 0.1 M IN WATER. SUCH AN ORDERING IS NOT ONLY ACADEMICALLY INTERESTING BUT ALSO PRACTICALLY RELEVANT FOR APPLICATIONS RANGING FROM INDUSTRIAL PROCESSES TO ENVIRONMENTAL ASSESSMENTS.

FUNDAMENTAL CONCEPTS UNDERPINNING pH DETERMINATION

BEFORE DELVING INTO SPECIFIC SPECIES, IT IS ESSENTIAL TO REVISIT CORE PRINCIPLES THAT GOVERN pH:

- STRONG ACIDS AND BASES DISSOCIATE COMPLETELY IN WATER, SIGNIFICANTLY INFLUENCING pH.
- WEAK ACIDS AND BASES DISSOCIATE PARTIALLY, WITH THE EXTENT GOVERNED BY THEIR DISSOCIATION CONSTANTS (K_A AND K_B).
- SALTS DERIVED FROM STRONG ACIDS AND STRONG BASES TYPICALLY PRODUCE A NEUTRAL SOLUTION ($\text{pH} \sim 7$).
- SALTS FROM WEAK ACIDS OR WEAK BASES CAN HYDROLYZE IN WATER, LEADING TO SOLUTIONS THAT ARE EITHER SLIGHTLY ACIDIC OR BASIC.

COMMON CHEMICAL SPECIES FOR ANALYSIS

FOR THIS EXPLORATION, CONSIDER THE FOLLOWING CHEMICAL SPECIES, REPRESENTATIVE OF VARIOUS ACID-BASE BEHAVIORS:

- STRONG ACID: HYDROCHLORIC ACID (HCl)
- WEAK ACIDS: ACETIC ACID (CH_3COOH), CARBONIC ACID (H_2CO_3)
- STRONG BASE: SODIUM HYDROXIDE (NaOH)
- WEAK BASES: AMMONIA (NH_3), ANILINE ($\text{C}_6\text{H}_5\text{NH}_2$)
- SALTS FROM STRONG ACIDS AND BASES: SODIUM CHLORIDE (NaCl)
- SALTS FROM WEAK ACIDS OR BASES: AMMONIUM CHLORIDE (NH_4Cl), SODIUM ACETATE (CH_3COONa)
- INORGANIC OXYANIONS/HYDROGEN COMPOUNDS: HYDROGEN SULFATE (H_2SO_4), HYDROGEN CARBONATE (HCO_3^-)

METHODOLOGY FOR DETERMINING pH ORDER

TO ORDER THESE SPECIES BY INCREASING pH, WE ANALYZE THEIR ACID-BASE PROPERTIES:

1. IDENTIFY WHETHER THE SPECIES IS A STRONG ACID/BASE OR WEAK ACID/BASE.
2. CALCULATE OR ESTIMATE THE pH OF 0.1 M SOLUTIONS BASED ON DISSOCIATION CONSTANTS.
3. CONSIDER HYDROLYSIS EFFECTS FOR SALTS.
4. COMPARE THE RESULTING pH VALUES TO ESTABLISH THE ORDER.

DEEP DIVE: ACIDIC, NEUTRAL, AND BASIC SOLUTIONS

1. STRONG ACID: HYDROCHLORIC ACID (HCL)

HCL IS A STRONG MONOPROTIC ACID THAT DISSOCIATES COMPLETELY:



IN A 0.1 M SOLUTION, THE CONCENTRATION OF H^+ IONS IS APPROXIMATELY 0.1 M, LEADING TO:

$$\text{pH} = -\log[\text{H}^+] = -\log(0.1) = 1$$

EXPECTED pH: ~ 1

2. WEAK ACID: ACETIC ACID (CH_3COOH)

ACETIC ACID IS A WEAK ACID WITH A DISSOCIATION CONSTANT:

$$K_A \approx 1.8 \times 10^{-5}$$

THE DISSOCIATION EQUILIBRIUM:



FOR A 0.1 M SOLUTION, THE APPROXIMATION:

$$[\text{H}^+] \approx \sqrt{K_A \times C} = \sqrt{(1.8 \times 10^{-5} \times 0.1)} \approx \sqrt{1.8 \times 10^{-6}} \approx 1.34 \times 10^{-3} \text{ M}$$

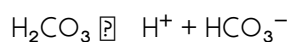
THUS,

$$\text{pH} \approx -\log(1.34 \times 10^{-3}) \approx 2.87$$

EXPECTED pH: ~ 2.9

3. WEAK ACID: CARBONIC ACID (H_2CO_3)

CARBONIC ACID IS A WEAK DIPROTIC ACID WITH FIRST DISSOCIATION:



$$K_{A1} \approx 4.3 \times 10^{-7}$$

SIMILAR CALCULATION:

$$[\text{H}^+] \approx \sqrt{K_{A1} \times C} \approx \sqrt{(4.3 \times 10^{-7} \times 0.1)} \approx \sqrt{4.3 \times 10^{-8}} \approx 6.55 \times 10^{-4} \text{ M}$$

$$\text{pH} \approx -\log(6.55 \times 10^{-4}) \approx 3.18$$

EXPECTED PH: ~3.2

4. NEUTRAL SALT: SODIUM CHLORIDE (NaCl)

NaCl ORIGINATES FROM A STRONG ACID (HCl) AND A STRONG BASE (NaOH). IT DOES NOT HYDROLYZE:



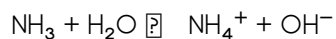
SOLUTION IS NEUTRAL:

$$\text{pH} \approx 7$$

5. WEAK BASE: AMMONIA (NH₃)

AMMONIA IS A WEAK BASE WITH $K_B \approx 1.8 \times 10^{-5}$.

THE HYDROLYSIS:



CALCULATING OH⁻ CONCENTRATION:

$$K_B = [\text{NH}_4^+][\text{OH}^-] / [\text{NH}_3]$$

ASSUMING X IS THE CONCENTRATION OF OH⁻:

$$K_B \approx x^2 / 0.1$$

$$x^2 \approx K_B \times 0.1 = 1.8 \times 10^{-5} \times 0.1 = 1.8 \times 10^{-6}$$

$$x \approx \sqrt{1.8 \times 10^{-6}} \approx 1.34 \times 10^{-3} \text{ M}$$

THEN,

$$\text{pOH} = -\log(1.34 \times 10^{-3}) \approx 2.87$$

$$\text{pH} = 14 - \text{pOH} \approx 11.13$$

EXPECTED PH: ~11.1

6. WEAK BASE: ANILINE (C₆H₅NH₂)

ANILINE IS A WEAK BASE WITH A K_B AROUND 4.3×10^{-10} , MUCH WEAKER THAN AMMONIA.

HYDROLYSIS:



CALCULATIONS SUGGEST A VERY LOW CONCENTRATION OF OH⁻:

$$x \approx \sqrt{K_B \times C} = \sqrt{4.3 \times 10^{-10} \times 0.1} \approx 2.07 \times 10^{-6} \text{ M}$$

$$\text{pOH} \approx 5.68$$

pH $\approx$ 8.32

EXPECTED pH: $\sim$ 8.3

7. SALT OF WEAK ACID AND STRONG BASE: SODIUM ACETATE (CH_3COONa)

SODIUM ACETATE IS A SALT DERIVED FROM ACETIC ACID AND SODIUM HYDROXIDE. IT HYDROLYZES:



THE BASE HYDROLYSIS CONSTANT (K_B OF ACETATE):

$$K_A \text{ OF ACETIC ACID } \approx 1.8 \times 10^{-5}$$

$$K_B \text{ (FOR ACETATE)} = K_W / K_A \approx 1.0 \times 10^{-14} / 1.8 \times 10^{-5} \approx 5.56 \times 10^{-10}$$

HYDROLYSIS:

$$[\text{OH}^-] \approx \sqrt{K_B \times C} = \sqrt{5.56 \times 10^{-10} \times 0.1} \approx 7.46 \times 10^{-6} \text{ M}$$

$$\text{pOH} \approx 5.13$$

$$\text{pH} \approx 8.87$$

EXPECTED pH: $\sim$ 8.9

8. SALT OF WEAK ACID AND WEAK BASE: AMMONIUM CHLORIDE (NH_4Cl)

NH_4Cl IS DERIVED FROM A WEAK BASE (NH_3) AND A STRONG ACID (HCl). ITS HYDROLYSIS:



THE K_A OF NH_4^+ :

$$K_A \approx K_W / K_B \text{ OF } \text{NH}_3 \approx 1.0 \times 10^{-14} / 1.8 \times 10^{-5} \approx 5.56 \times 10^{-10}$$

CALCULATING H_3O^+ CONCENTRATION:

$$[\text{H}_3\text{O}^+] \approx \sqrt{K_A \times C} \approx \sqrt{5.56 \times 10^{-10} \times 0.1} \approx 7.46 \times 10^{-6} \text{ M}$$

$$\text{pH} \approx -\log(7.46 \times 10^{-6}) \approx 5.13$$

EXPECTED pH: $\sim$ 5.1

SUMMARIZING THE pH VALUES

SPECIES	APPROXIMATE pH
HYDROCHLORIC ACID (HCL)	$\sim$ 1.0

Order These Chemical Species By Increasing Ph Of An 0 1m Aqueous Solution Of Each That Is Imagine Making

Order These Chemical Species By Increasing Ph Of An 0 1m Aqueous Solution Of Each That Is Imagine Making

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