ACIDS AND BASES

The two theories that we will discuss are the Theory and the Theory and the **Theory.** Theory.

The Proton in Water: Theory

Swedish chemist Arrhenius defined

• an acid as a substance that ionizes in water to give _________ ions, and

· a base as a substance that ionizes in water to give *index* ions.

Hydrochloric acid, HCl, is a $\frac{1}{100}$ acid, and is very soluble in water. It

ionizes into its component ions in the following manner:

 $HCl (q) \rightarrow H^+(aq) + Cl^-(aq)$

The hydrogen ion interacts strongly with a lone pair of electrons on the oxygen of a water molecule. The resulting ion, H_3O^* is called the

<u>ion. Iomann iom.</u>

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The $\frac{1}{\sqrt{1-\frac{1}{\sqrt{1$

ACIDIC solutions are formed when an acid transfers a $\frac{1}{\sqrt{1-\frac{1}{n}}}$ to water. The reaction of HCI with water can be written in either of the following ways: $HCl (aq) + H₂O (l) \rightarrow H₃O+ (aq) + Cl⁻(aq)$ HCl (aq) $\rightarrow H^*$ (aq) + $Cl^-(aq)$

The Bronsted-Lowry Concept of Acids and Bases

• _______________ are proton donors, and

• Change of the proton acceptors.

In the example above, the HCI acts as a Brønsted Acid by ______________________ a proton to water. Water in turn acts as a Brønsted Base by ________________ a proton from the HCI.

Water can act as an acid or a base. When it reacts with the HCI it acts as a base.

In the example below it is acting as an $\frac{1}{\sqrt{1-\frac{1}{n}}}$, a proton $\frac{1}{\sqrt{1-\frac{1}{n}}}$.

 $NH_3(aq) + H_2O(l) \rightarrow NH_4^+(aq) + OH^-(aq)$

Here, H2O acts as a Brønsted acid by ____________ a proton to NH3 which acts as a Bronsted base.

Using the Arrhenius definition, we say that the resulting solution is because it contains OH- ions, thus we say that the NH3 molecule is basic (a

<u>____________________</u> acceptor).

All Arrhenius acids are also Brønsted acids.

All Arrhenius bases are also Bronsted bases.

ACID & BASE STRENGTH

Acids and Bases as Equilibrium Reactions Acids and bases are an important sub-section of _____________ reactions. The of strong acids and bases through reaction with water is assumed to go to completion (figure 1).

However, the majority of acids are _______ and only a fraction of the weak acid present in aqueous solution actually reacts with water. Most of the acid remains in its state.

Salts

In dilute aqueous solutions, salts are completely dissociated into ions. Eg. MgCl_{2 (aq)} \rightarrow _______ + ___ Aqueous solutions of some salts like NaCl are neutral ($pH = __\$). Some aqueous solutions of salts are $\frac{1}{\sqrt{2}}$ (such as AlCl₃, pH < 7). Other solutions of salts are basic. An example is $\frac{1}{12}$ the pH > 7.

How can you predict whether an aqueous solution of a salt is acidic, basic or neutral?

PRAC Hydrolysis of

You need to know if the ions of the salt will react with ________. In other words, do any of the ions undergo

in Foundations shows which anions and cations will cause acidic and basic solutions.

Complete the pH of Salt Solutions PRACTICAL

Generalizations you can make from the results table: 1. Anions from strong acids do not and therefore produce neutral salts. Salts 2A.doc

2. Cations from strong bases do not hydrolyse and therefore produce salts.

3. Basic anions are those that will react with ________ to produce hydroxide ions.

Basic anions are derived from weak acids. Salts of these anions are called

4. Acidic anions are those that react with water to produce ______________ ions. Acidic cations can be derived from weak bases and aquated metal cations.

Conjugate Acid-Base Pairs

Let's look at the reaction of $NH₃$ and $H₂O$ again: (1) $NH_3 + H_2O \rightarrow NH_4^+ + OH^-$ The reverse of this reaction is:

(2)

In this case, NH4 + acts as an acid which a proton to OH- . 0H $acts as a$ $\qquad \qquad$

An acid and a base that are related by the gain and loss of a proton are called a ______________________acid-base pair. For example, NH4* is the ___________ ______ of NH₃ , and NH₃ is the ______________________________ of NH₄+. Every acid has associated with it a **constantly asset that the base.**

Likewise, every base has associated with it a $\frac{1}{1-\frac{1}{2}}$ acid.

For any reaction: $HX + H₂O \rightarrow H₃O⁺ + X⁻$

If HX is a strong acid it will give up its proton *______________*, this makes X⁻ a ___________ base because it has less affinity for the proton. The forward reaction is favored, mainly in solution.

If HX is a $\frac{1}{\sqrt{1-\frac$ X will have a high affinity for a _________, and X is a _________ base than water. The reverse reaction is favored, mainly _______________ of HX in solution. **In summary,**

• the stronger an acid the **intercontage in the stronger** its conjugate base.

• the stronger a base the **intercontage in the stronger of the stronger** acid.

Amphoteric metals, aluminium, chromium and zinc Amphoteric metals react with both _______________ and ___________ solutions. They react with metal _______________ solutions, such as sodium hydroxide, to form $\frac{1}{\sqrt{1-\frac{1$ *2A1(s) +* 20H-(aq) + 6H20(1) ---> $A|(OH)_{3}(s) + OH(aq)$ --->

Yll Chemistry Answers:

Page 1 Page 2

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Arrhenius Red Bronsted-Lowry Conducts Arrhenius Sour Hydronium Metals
Hydroxide Hydroc Hydroxide Hydrogen
Strong Garbonate Hydronium/Hydrogen Oxides
Hydrogen Hydrox Hydrogen Hydroxides
H⁺ Plue H+ Blue Proton Conducts
Acids Conducts Acids Bitter
Bases amphe Donating Acids Receiving/accepting Salt
Acid Met Acid Water
Donor bonne Donor
Donating chemical chemical Donating ionisation
Basic in the second series of the series Basic weak
Proton malay

Carbonates Amphoteric molecular/original pH equilibrium pH concentration pH solvent (aka. H₂O) pH weaker

Page 3 H^* OH \degree or O^{2-} Mq^{2+} C -7 Acidic Ba(OH)2 Water Ionization Hydrolyse/React **Neutral** Water Bases Hydronium/H+

Page 4

 NH_4 ⁺ + OH⁻ \rightarrow NH₃ + H₂O donates base conjugate conjugate acid conjugate base conjugate conjugate easily/readily weak protons/hydronium ions weak proton stronger consists/molecules weaker strong strong weak

Yll Chemistry **Conjugate Acid-Base Pairs**

For the following equations identify (circle with a key) the acid - base conjugate pairs.

\n- 1.
$$
CH_3COOH_{(aq)} + H_2O_{(l)} \rightleftharpoons CH_3COO^{-}(aq) + H_3O^{+}(aq)
$$
\n- 2. $HSO_4^{-}(aq) + CO_3^{2-}(aq) \rightleftharpoons SO_4^{2-}(aq) + HCO_3^{-}(aq)$
\n- 3. $H_3PO_{4(aq)} + 3NaOH_{(aq)} \rightarrow Na_3PO_{4(aq)} + 3H_2O_{(l)}$
\n- 4. $NH_{3(aq)} + HCl_{(aq)} \rightarrow NH_4Cl_{(aq)}$
\n- 5. $Na_2CO_{3(aq)} + H_2O_{(l)} \rightleftharpoons 2Na^{+}(aq) + HCO_3^{-}(aq) + OH^{-}(aq)$
\n- 6. $NH_{3(aq)} + H_2O_{(l)} \rightleftharpoons NH_4^{+}(aq) + OH^{-}(aq)$
\n- 7. $OH^{-}(aq) + HNO_{3(aq)} \rightarrow NO_3^{-}(aq) + H_2O_{(l)}$
\n- 8. $OH^{-}(aq) + HCO_3^{-}(aq) \rightleftharpoons CO_3^{2-}(aq) + H_2O_{(l)}$
\n- 9. $H^{+}(aq) + OH^{-}(aq) \rightleftharpoons H_2O_{(l)}$
\n- 10. $H_2C_2O_{4(aq)} + NH_{3(aq)} \rightleftharpoons NH_4^{+}(aq) + HC_2O_4^{-}(aq)$
\n

For the following complete the equation and then identify the conjugate acid base pairs.

- 11. $H_2S_{(aq)} + OH^-(aq) \rightarrow$ 12. $H_2PO_4^-(aq) + CO_3^2^-(aq) \rightleftharpoons$ 13. $HClO_{(aq)} + NH_{3(aq)} \rightleftharpoons$
- 14. $HF_{(aq)} + O^{2-}(aq) \rightleftharpoons$
- 15. $HNO_{3(aq)} + HCO_{3(aq)} \rightarrow$

16.
$$
HCl_{(aq)} + NaOH_{(aq)} \rightarrow
$$

17.
$$
H_3O^{\dagger}(aq) + HSO_4(qq) \rightleftharpoons
$$

18.
$$
HCO_{3}^{-}(aq) + H_{2}O(l) \rightleftharpoons
$$

- 19. $HF_{(aq)} + NH_{3(aq)} \rightarrow$
- 20. $NH_4^+(aq) + H_2O(1) \rightarrow$

V11 Chemistry Conjugate Acid-Base Pairs- Answers

$ACID \rightarrow BASE$ $BASE \rightarrow ACID$

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 $CH_3COOH_{(aq)} + H_2O_{(l)} \rightleftharpoons CH_3COO_{(aq)} + H_3O_{(aq)}$ HSO_4 ⁻(aq) + CO_3 ²⁻(aq) \Rightarrow SO_4 ²⁻(aq) + HCO_3 ⁻(aq) $H_3PO_{4(aq)}$ + 30H⁻(aq) \rightarrow H_2PO_{4} ⁻(aq) + H₂O(I) $NH_{3(aq)} + HCl_(aq) \rightarrow NH₄⁺ + Cl_(aq)$ CO_3^2 ⁻(aq) + $H_2O(1)$ \rightleftharpoons HCO₃⁻(aq) + OH⁻(aq) $NH_{3(aq)} + H_2O_{(1)} \rightleftharpoons NH_4^+(aq) + OH^-(aq)$ $OH^-_{(aq)}$ + $HNO_{3(aq)} \rightarrow NO_3^-_{(aq)}$ + $H_2O_{(l)}$ $OH^-_{(aq)} + \underline{HCO_3^-_{(aq)}} \rightleftharpoons \underline{CO_3^{2-}_{(aq)}} + H_2O_{(1)}$ $H_3O^+(aq)}$ + OH $(aq) \rightleftharpoons 2H_2O(1)$ $H_2C_2O_{4(aq)}$ + NH_{3(aq)} \rightleftharpoons NH₄⁺(aq) + <u>HC₂O₄</u>-(aq)

The pairs are highlighted as similar species below: $H_2S_{(aq)}$ + OH⁻(aq) $\rightarrow H_2O_{(1)}$ + HS⁻(aq) $H_2PO_4^{-}$ _(eq) + CO_3^{2-} _(eq) \Rightarrow HPO_4^{2-} + HCO_3^{-} $HClO_{(aq)}$ + NH_{3(aq)} \rightleftharpoons ClO⁻_(aq) + NH₄⁺_(aq) $HF_{(aq)} + O^{2-}(aq) \rightleftharpoons F_{(aq)}^-(aq) + OH_{(aq)}$ $HNO_{3(aq)} + HCO_{3(aq)} \rightarrow NO₃⁻ + H₂CO_{3(aq)}$ $HCl_{(aq)}$ + NaOH $_{(aq)} \rightarrow$ NaCl $_{(aq)}$ + H₂O₍₁₎ $H_3O^+(_{(aq)}$ + HSO₄⁻(aq) $\rightleftharpoons H_2O_{(l)}$ + H₂SO_{4(aq)} HCO^{3-} (aq) + H₂O(1) \rightleftharpoons H₂CO_{3(aq)} + OH⁻(aq) $HF_{(aq)} + NH_{3(aq)} \rightarrow F_{(aq)} + NH_{4}^{+}(aq)$ $NH_4^+(qq)$ + $H_2O(1)$ $\rightarrow NH_3(qq)$ + $H_3O^+(qq)$

Amphoteric Substances

Base + Acid With acid dissolve \rightarrow salt + water (as for any metal hydroxide)

 $A[(OH)_3(s) + H^*(aq) \rightarrow$

 $Cr(OH)_3(s) + H^*(aq) \rightarrow$

 $Zn(OH)₂(s) + H[*](aq) \rightarrow$

Amphoteric metals + Acids With acids dissolve \rightarrow salt + hydrogen gas (as for any reactive metal)

 $A|(s) + H^*(aq) \rightarrow$

 $Cr(s) + H^*(aq) \rightarrow$

 $Zn(s) + H^*(aq) \rightarrow$

Metal Oxide + Acids With acids dissolve \rightarrow salt + water (as for any metal oxide)

 $A|_2O_3(s) + H^*(aq) \rightarrow$

 $Cr_2O_3(s) + H^*(aq) \rightarrow$

 $ZnO(s) + H^{+}(aq) \rightarrow$

ANSWERS

 $A|(OH)_3(s) + 3H'(aq) \rightarrow Al^{3*}(aq) + 3H_2O(l)$ $Cr(OH)_3(s) + 3H^*(aq) \rightarrow Cr^{3*}(aq) + 3H_2O(l)$ $Zn(OH)₂(s) + 2H^t(aq) \rightarrow Zn²⁺(aq) + 2H₂O(l)$ $2A|(s) + 6H'(aq) \rightarrow 2A|^{3}(aq) + 3H_2(q)$ $2Cr(s) + 6H'(aq) \rightarrow 2Cr^{3}(aq) + 3H_2(q)$ $Zn(s) + 2H^{(aq)} \rightarrow Zn^{2}(aq) + H_2(q)$ $Al_2O_3(s) + 6H^*(aq) \rightarrow 2Al^{3*}(aq) + 3H_2O(l)$ $Cr_2O_3(s) + 6H^*(aq) \rightarrow 2Cr^{3*}(aq) + 3H_2O(l)$ $ZnO(s) + 2H^{*}(aq) \rightarrow Zn^{2*}(aq) + H_{2}O(l)$

Yll Chemistry REVISION: *MC* **SECTION**

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- 1. Which one of the following is **not** an acid-base reaction?
	- *A* $NH_{3(q)}$ + $HCl_{(q)}$ \rightarrow $NH_4Cl_{(s)}$
	- B NH₄Cl_(s) + Ca(OH)_{2(s)} + heat \rightarrow NH_{3(g)} + CaCl_{2(ag)} + H₂O_(l)
	- C (NH₄)₂CO_{3(s)} + heat \rightarrow 2NH_{3(g)} + 2H₂O_(g) + CO_{2(g)}
	- $D Ba(OH)_{2(s)} + 2NH_4SCN_{(s)} \rightarrow Ba^{2+}(aq) + 2SCN-(aq)+ 2H_2O(l) + 2NH_3(aq)$
- 2. Which one of the following is a **conjugate acid-base pair?**

A HNO3 and H2O B H₃O⁺ and OH-C NH3 and OH- D HNO₃ and NO₃-

- 3. Which of the following correctly describes the *difference between a strong acid and a weak acid?*
	- (a) All solutions of weak acids are poor conductors of electricity but solutions
	- of strong acids are always good conductors of electricity.
	- (b) Concentrated solutions can only be prepared from strong acids.
	- (c) The degree of ionisation for strong acids is greater than for weak acids.
	- (d) Weak acids are less soluble than strong acids.
- 4. Which of the conjugate acid-base pairs is *not* correctly listed?

Acid Conjugate base

- (a) NH₃ NH₂⁺
- (b) 504^2 ⁻ HSO_4 ⁻
- (c) $HCO₃$ ⁻ $CO₃$ ²-
- (d) $HPO₄²PO₄³$
- 5. Water taken from a swimming pool had a pH of 9.5. Which of the following substances could be added to the pool water to bring the pH closer to 7?
	- (a) $Na_2CO_{3(s)}$ (b) $\text{NaCl}_{(s)}$ (c) $Na_3PO_{4(aq)}$ (d) HCl(aq)

6. In which of the following equations *is* the *underlined* substance acting as an *acid?*

(a) $C_6H_5COO^-$ + $H_2SO_3 \rightarrow HSO_3^-$ + C_6H_5COOH (b) $2Na + 2H_2O \rightarrow 2Na^+ + 2OH^- + H_2$ (c) HCO₃ + H₂O \rightarrow H₂CO₃ + OH (d) NH₄⁺ + OH⁻ \rightarrow NH₃ + H₂O

7. The hydrogen carbonate ion, $HCO₃$ (_{ag}), may act as an acid or a base in aqueous solution. In which one of the equations below is it acting as an acid?

A HCO_3 ⁻ (aq) + $H_2O(1)$ \rightarrow $H_2CO_3(1)$ + OH ⁻ (aq) $B HCO₃(aq) + H₃O⁺(aq) \rightarrow H₂CO_{3(aq)} + H₂O₍₁₎$ *C* $HCO_3(qq) + H_2O(q) \rightarrow CO_3^{2}(qq) + H_3O(qq)$ D HCO₃ (_{aq}) + OH⁻(_{aq}) → H₂CO_{3(aq}) + O²⁻(_{aq})

8. Acid ionisation or dissociation constants (K_a) provide information as to the extent to which an acid ionises / dissociates in aqueous solution. The larger the value the stronger the acid, the lower the weaker:

The table below shows acid dissociation constants, (K_a) , for four acids, measured at 25°C.

Which is the WEAKEST acid shown in the table above? A, HF $B, HSO₄$ $C, CH₃COOH$ D, HCIO

ANSWERS

Yll Chemistry **SHORT ANSWER**

NET IONIC EQUATIONS *AGAIN!*

Write equations for any reactions that occur in the following situations. If no reaction occurs write *"no reaction".* In each case describe *in full* what you would observe, including any colours, odours, precipitates (give the colour), gases evolved (give the colour or describe as colourless).

Your equations should refer only to the actual species involved. These species may be **ions** [for example; Ag+(.0], **molecules** [for example; NH3(9), NH3(aq), CH3COOH(aq)1 or **solids** [or precipitates for example; B0504(s), *Cu(s),* $Na₂CO_{3(s)}$].

1.

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(a) Concentrated hydrochloric acid is poured onto solid Magnesium Carbonate.

EQUATION:

OBSERVATION:

(b) Nitric Acid poured onto a black solid copper oxide.

EQUATION:

OBSERVATION:

(C) *A* piece of Calcium metal is dropped into a Sulfuric Acid solution EQUATION:

OBSERVATION:

(d) Solid zinc is added to 2.0 mol L^{-1} hydrochloric acid solution. EQUATION:

OBSERVATION:

2.

a. Ammonia solution and acetic acid (ethanoic acid) are weak electrolytes and are not good conductors of electricity. But when you add them to each other the resulting solution has a high electrical conductivity. **Explain** this result.

[3 points!]

b. Give an explanation for each of the following observation. Solid lithium bromide, LiBr(s) is a *non conductor of electricity* and so too is liquid water H20(1) yet the combination of LiBr(s) and H20(1) produces a *very good conducting mixture.*

[3 points!]

1. Determine how many mL of 3.00 mol L^{-1} sodium hydroxide are required to neutralise 30.0 mL of 2.00 mol L-1hydrochloric acid solution.

2. 2.00 g of sodium hydroxide was dissolved in water and the solution made up to 2.50 x 10²mL. Determine:

i. the concentration of the solution

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ii. the concentration of the solution in $q L^{-1}$

(iii. the volume of 0.100 mol L^{-1} sulfuric acid required to neutralise 25.0 mL of the solution and

iv. the mass of pure sulfuric acid required for complete neutralisation.

3. A solution of crystalline oxalic acid $(H_2C_2O_4$. $2H_2O)$, a diprotic acid, is prepared by dissolving 12.6 g of the acid in 4.00 L of water. 20.0 mL of this solution is required to completely neutralise 36.0 mL of a potassium hydroxide solution. Calculate the concentration in mol L^{-1} and strength in $q L^{-1}$ of the potassium hydroxide solution.

4. 0.700 g of an ammonium salt was dissolved in distilled water and then 25.0 mL of a 1.10 mol L $^{-1}$ sodium hydroxide solution was added. The solution was boiled. All ammonia gas was expelled. After cooling it was found that 30.0 mL of 0.450 mol L^{-1} hydrochloric acid was required to neutralise the *excess* base. Determine the percentage of ammonia in the salt.

5. Determine the percentage purity of a sample of zinc if 0.325 *g* required 38.0 mL of 0.250 mol L 'hydrochloric acid for a complete reaction.

6. 0.100 *g* of calcium carbonate was dissolved in 25.0 mL of hydrochloric acid solution, and it was found that 5.50 mL of sodium hydroxide solution were required to neutralise the excess acid. A second titration showed that 27.5 mL of sodium hydroxide

neutralised 25.0 mL of the hydrochloric acid solution. Determine the concentration of the sodium hydroxide and hydrochloric acid solutions.

7. (EXTENSION) In a method of volumetric determination of sulfate ion, the SO_4^2 -ion is precipitated as PbS0₄ by the addition of Pb²⁺. The PbSO₄ is then analysed for Pb and the number of moles of SO₄²-ion is equal to the number of moles of Pb^{2+} in the precipitate.

The amount of lead is determined by titration using a compound symbolised as H4Y. [This compound is actually called 1,2-diaminoethane-N,N,N',N'-tetraethanoic acid.]

A known amount of H4Y *is* added to the precipitate, bringing it to solution.

 $PbSO_4(s) + H_4Y \rightarrow PbY^2$ - (aq) + SO_4^2 - (aq) + 4 H^+ (aq) Then the amount of excess H4Y is determined by titration with standard lead nitrate.

 Pb^{2+} + H₄Y \rightarrow PbY²⁻_(aq) + 4 H⁺

From this, the amount of Pb^{2+} in the $PbSO_4$, and hence the amount of $504²$ ion in the sample, can be calculated.

The following results were obtained for a sample of ground water analysed as above.

Volume of ground water sample: : **10.00 L**

Volume of 0.1000 mol L⁻¹ H₄Y added to the precipitate: 25.00 mL

Volume of 0.1000 mol L -1Pb(NO3)2 required to titrate excess H4Y: **8.26 mL**

- a. Calculate the total number of moles of H4Y added, to the precipitate. [1 mark]
- b. Calculate the number of moles of H_4 Y in excess. [2 marks]
- c. Calculate the number of moles of H_4 Y that combined with the Pb504.

[2 marks]

d. Calculate the concentration of the SO_4^2 - ion in the ground water.

[3 marks]

Suggest a source of sulfate in the ground water resulting from agricultural activity.

Yll Chemistry Answers: 1. 20.0 mL 2. 0.200 mol L $^{-1}$, 8.00 gL $^{-1}$, 25.0 mL, 0.245g 3. 0.028 mol L⁻¹, 1.56 gL⁻¹ 4. 34.0% 5. 95.6% 6. 0.100 mol L -1,0.0900mo1 7. ask me © (a) (b) (c) (d)

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 $\label{eq:2.1} \frac{1}{\sqrt{2}}\int_{\mathbb{R}^3}\frac{1}{\sqrt{2}}\left(\frac{1}{\sqrt{2}}\right)^2\frac{1}{\sqrt{2}}\left(\frac{1}{\sqrt{2}}\right)^2\frac{1}{\sqrt{2}}\left(\frac{1}{\sqrt{2}}\right)^2\frac{1}{\sqrt{2}}\left(\frac{1}{\sqrt{2}}\right)^2.$

 $\label{eq:2.1} \frac{1}{\sqrt{2}}\int_{0}^{\infty}\frac{1}{\sqrt{2\pi}}\left(\frac{1}{\sqrt{2\pi}}\right)^{2}d\mu\left(\frac{1}{\sqrt{2\pi}}\right)\frac{d\mu}{d\mu}d\mu\left(\frac{1}{\sqrt{2\pi}}\right).$ \sim $\label{eq:2.1} \mathcal{L}(\mathcal{L}^{\text{max}}_{\mathcal{L}}(\mathcal{L}^{\text{max}}_{\mathcal{L}}),\mathcal{L}^{\text{max}}_{\mathcal{L}^{\text{max}}_{\mathcal{L}}})$

 $\mathcal{L}^{\text{max}}_{\text{max}}$ and $\mathcal{L}^{\text{max}}_{\text{max}}$