1. The following is a simple representation of an organic synthesis process known as the Diels-Alder reaction. Draw electron pushing arrows on the reactants to make the product.

or

2. Rank the following cations in order from longest (\#1) to shortest wavelength (\#3). (Hint: energy is inversely proportional to wavelength)

3. Which of the following gas molecules will have the lowest root-mean-square speed at $150^{\circ} \mathrm{C}: \mathrm{SF}_{6}, \mathrm{CCl}_{4}$, or $\mathrm{H}_{2} \mathrm{O}$

Using the equation " $\mathrm{v}=\operatorname{sqrt}(3 \mathrm{RTM})$ " we know the substance with the largest molar mass will have the lowest root-mean-square speed. $\mathrm{SF}_{6}$ has a molar mass of $146.06 \mathrm{~g} / \mathrm{mol}^{2}, \mathrm{CCl}_{4}$ has a molar mass of $153.82 \mathrm{~g} / \mathrm{mol}$, and $\mathrm{H}_{2} \mathrm{O}$ has a molar mass of $18.02 \mathrm{~g} / \mathrm{mol}$. Thus, $\mathbf{C C l}_{4}$ has the lowest root-mean-square speed at $150^{\circ} \mathrm{C}$.

1. A sample of nitrogen gas effuses at a rate four times that of an unknown gas. What is the molecular weight of the unknown gas in $\mathrm{g} / \mathrm{mol}$ ? (Hint: $\frac{r_{1}}{r_{2}}=\sqrt{\frac{M_{2}}{M_{1}}}$ )
Graham's law of effusion: r1/r2 = sqrt(M2/M1)
Rate of effusion of $N_{2}=4=r 1$, Rate of effusion of unknown gas $=1=r 2$
Molecular mass of $\mathrm{N}_{2}=28.0134 \mathrm{~g} / \mathrm{mol}=\mathrm{M} 1$, Molecular mass of unknown gas $=\mathrm{x}=\mathrm{M} 2$
Molecular mass of unknown gas $=\mathrm{x}=(\mathrm{r} 1 / \mathrm{r} 2)^{\wedge} 2 * \mathrm{M} 1=\mathbf{4 4 8 . 2 1 4 4} \mathbf{g} / \mathbf{m o l}$
2. A 14.5 L closed vessel contains $1.49 \mathrm{~g} \mathrm{H}_{2} \mathrm{O}, 4.32 \mathrm{~g} \mathrm{CO}_{2}$, and $3.21 \mathrm{~g} \mathrm{SO}_{3}$. At $40^{\circ} \mathrm{C}$, what is the pressure in the vessel (in atm)? Assume these gases behave ideally.

$$
\begin{aligned}
& P V=n R T \\
& P=\frac{n R T}{V} \\
& 1.49 \mathrm{~g} \mathrm{H} 2 \mathrm{O} \times \frac{1 \mathrm{~mol} \mathrm{H}_{2} \mathrm{O}}{18.02 \mathrm{~g}}=0.083 \mathrm{~mol} \mathrm{H} \mathrm{H} \\
& 4.32 \mathrm{~g} \mathrm{CO}_{2} \times \frac{1 \mathrm{~mol} \mathrm{CO}}{44.01 \mathrm{~g}}=0.098 \mathrm{~mol} \mathrm{CO} 2 \\
& 3.21 \mathrm{~g} \mathrm{SO}_{3} \times \frac{1 \mathrm{~mol} \mathrm{SO}}{80.06 \mathrm{~g}}=0.040 \mathrm{~mol} \mathrm{SO} 3 \\
& \begin{aligned}
\text { Total mol of gas } & =0.083+0.098+0.040 \\
& =0.221 \mathrm{~mol}
\end{aligned} \\
& P=\frac{n R T}{V}=\frac{(0.221)(0.08206)(313.15)}{14.5} \\
& =0.392 \mathrm{~atm}
\end{aligned}
$$

3. Considering the ideal gas as the system, calculate the heat transferred to/from the surroundings $\left(\mathrm{q}_{\text {surroundings }}\right)$ in the process below. This expansion is performed against a constant external pressure of 1 atm .

4. In the conversion of acetyl CoA (shown below) to carbon dioxide, is carbon oxidized or reduced?


Oxidized
2. Suppose you are conducting an experiment to study two silver salts, AgBr and AgCl . They have solubility product constants of $7.7 * 10^{-13}$ and $1.6 * 10^{-10}$, respectively. You begin with a solution of $0.10 \mathrm{M} \mathrm{Br}^{-}$and $0.10 \mathrm{M} \mathrm{Cl}^{-}$and gradually add in $\mathrm{Ag}^{+}$. What percent of $\mathrm{Br}^{-}$remains in solution when AgCl first precipitates?

AgBr will start to precipitate when $\left[\mathrm{Ag}^{+}\right]\left[\mathrm{Br}^{-}\right]=7.7 * 10^{-13}$
$\left[\mathrm{Ag}^{+}\right]=7.7 * 10^{-13} / 0.10=7.7 * 10^{-12}$
AgCl will start to precipitate when $\left[\mathrm{Ag}^{+}\right]\left[\mathrm{Cl}^{-}\right]=1.6 * 10^{-10}$
$\left[\mathrm{Ag}^{+}\right]=1.6 * 10^{-10} / 0.10=1.6 * 10^{-9}$
When AgCl begins to precipitate, $\left[\mathrm{Ag}^{+}\right]=1.6 * 10^{-9},\left[\mathrm{Ag}^{+}\right]\left[\mathrm{Br}^{-}\right]=7.7 * 10^{-12},[\mathrm{Br}-]=4.8$ * $10^{-4}$ (remaining in solution, not precipitated)
$\%$ remaining in solution when AgCl first precipitates $=4.8 * 10^{-4} / 0.10 * 100 \%=\mathbf{0 . 4 8} \%$
3. Billy conducts an electron diffraction experiment. There are 2 slit openings lined up top to bottom with the appropriate width for diffraction. Billy will fire trillions of electrons upon the slits. How many detectors should Billy use if they want the diffraction pattern to appear as if there is only a single slit?
1

1. The photoelectric effect describes the emission of electrons from a material, oftentimes a metal, when electromagnetic radiation hits its surface. The equation $K E_{\max }=E_{p}-\phi$ describes the maximum kinetic energy of the ejected electrons in terms of $E_{p}$, the energy of the incoming photon, and $\phi$, the work function. $\phi$ for gold is $8.17 \cdot 10^{-19} \mathrm{~J}$ and light with a frequency of $1.00 \cdot 10^{15} \mathrm{~s}^{-1}$ is shined on a gold plate. If there are ejected electrons, what is the speed of the ejected electrons? If there are no ejected electrons, write "no electrons are ejected."

## No electrons ejected

2. Pu-244 undergoes beta-decay at a half-life of 16 hours. If a chemist starts with a sample of 512 g of Pu-244 and allows it to undergo beta-decay, how much of the original sample remains after 4 days?

## 8 grams

3. Calculate the pH of a $1.00 \times 10^{-2} \mathrm{M} \mathrm{H}_{2} \mathrm{SO}_{4}$ solution. $\left(K_{a 1}\right.$ is very large, consider the first dissociation step as totally complete; $K_{a 2}=1.2 \times 10^{-2}$ )
First dissociation: $\left[H^{+}\right]=1.00 \times 10^{-2} \mathrm{M}$
Second dissociation:
Ice Table:

|  | $\mathrm{HSO}_{4}^{-}$ | $\mathrm{SO}_{4}^{2-}$ | $H^{+}$ |
| :---: | :---: | :---: | :---: |
| I | $1.00 \times 10^{-2} \mathrm{M}$ | 0 | $1.00 \times 10^{-2} \mathrm{M}$ |
| C | -x | +x | +x |
| E | $\left(1.00 \times 10^{-2}-x\right)$ | x | $\left(1.00 \times 10^{-2}+\mathrm{x}\right)$ |
| $\begin{aligned} & K_{a 2}=1.2 \times 10^{-2}=\frac{\left(1.00 \times 10^{-2}+x\right) \cdot x}{\left(1.00 \times 10^{-2}-x\right)} \\ & x=-0.026524 \text { or } 0.004524 \\ & {\left[H^{+}\right]=1.00 \times 10^{-2}+0.004524=0.014524 \mathrm{M}} \\ & p H=-\log 0.014524 \simeq \mathbf{1 . 8 3 8} \end{aligned}$ |  |  |  |

1. How many more chlorine atoms are present in a 3-chloropentane molecule compared to a 2-chloropentane molecule?

0
2. An 80 mL aqueous solution contains 0.7 g of hydrofluoric acid (initial amount) dissolved in water. Calculate the pH of the solution $\left(\mathrm{K}_{\mathrm{a}}\right.$ of $\mathrm{HF}=6.6 \times 10^{-4}$ at $25^{\circ} \mathrm{C}$; $\mathrm{MW} \mathrm{HF}=$ $20.01 \mathrm{~g} / \mathrm{mol}$ ).

$$
0.7 \mathrm{~g} \mathrm{HF}\left(\frac{1 \mathrm{~mol}}{20.01 \mathrm{~g}}\right)=0.03498 \mathrm{~mol} \mathrm{HF} \quad \frac{0.03498 \mathrm{~mol}}{0.08 \mathrm{~L}}=0.437 \mathrm{M} \mathrm{HF}
$$

|  | $\mathrm{HF}(\mathrm{aq})+\mathrm{H}_{2} \mathrm{O}(l) \rightleftharpoons \mathrm{H}_{3} \mathrm{O}^{+}(\mathrm{aq})+\mathrm{F}^{-}$ |  |  |
| :---: | :---: | :---: | :---: |
| I | 0.437 M | - | 0 |
| E | $0.437-\mathrm{x}$ | - | x |

$\mathrm{K}_{\mathrm{a}}=6.6 \times 10^{-4}=\frac{(x)(x)}{0.437-x}$
use method of successive approximation $\rightarrow \approx 6.6 \times 10^{-4}=\frac{(x)(x)}{0.437}$
$\mathrm{x}=0.0167=\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]$
$\mathrm{pH}=-\log (0.0167)=\mathbf{1 . 7 7 - 1 . 7 8}$
3. The percent dissociation of a solution with an initial amount of 0.3 M of nitrous acid is $1.3 \%$. A 0.3 M solution of hydrocyanic acid has the same percent dissociation. Calculate the pOH of the solution with hydrocyanic acid.

$$
\begin{aligned}
& \frac{H 3 O}{0.3} \cdot 100=1.3 \% \text { for } \mathrm{HNO}_{2} \\
& {\left[\mathrm{H}_{3} \mathrm{O}^{+}\right]=0.0039 \mathrm{M} \text { for } \mathrm{HNO}_{2}=\left[\mathrm{H}_{3} \mathrm{O}^{+}\right] \text {for } \mathrm{HCN}} \\
& \mathrm{pH}=-\log (0.0039)=2.41 \\
& \mathrm{pOH}=14-2.41=\mathbf{1 1 . 5 9}
\end{aligned}
$$

1. $2 \mathrm{SO}_{2(\mathrm{~g})}+\mathrm{O}_{2(\mathrm{~g})} \rightarrow 2 \mathrm{SO}_{3(\mathrm{~g})}$ is one of the essential reactions involved in the industrial production of $\mathrm{H}_{2} \mathrm{SO}_{4}$. Suppose at the beginning of the reaction, there are 100 L of mixed gases: $7 \%$ is $\mathrm{SO}_{2}, 11 \%$ is $\mathrm{O}_{2}, 82 \%$ is $\mathrm{N}_{2}$. At the end of the reaction, the total volume of the gas becomes 97.2 L . What percentage of $\mathrm{SO}_{2}$ was converted? (Assume that the total pressure and temperature remain constant throughout the reaction, and the gasses behave ideally) (3 significant figures)
Since the total pressure and temperature remain constant
$\frac{n_{i}}{V_{i}}=\frac{n_{f}}{V_{f}} \Rightarrow n_{f}=0.972 n_{i}$
$\Delta n=(1-0.972) n_{i}=0.028 n_{i}$
$\Delta n\left(\mathrm{SO}_{2}\right)=2 \times \Delta n=0.056 n_{i}$
$\% n\left(\mathrm{SO}_{2}\right)_{\text {converted }}=\frac{0.056 n_{i}}{0.11 n_{i}}=50.9 \%$
2. You are performing a titration to determine the concentration of the weak monoprotic acid HA $\left(\mathrm{K}_{\mathrm{a}}=2.0 * 10^{-3}\right)$ using a strong monoprotic base B with a concentration of 1 M $\left(\mathrm{K}_{\mathrm{b}}=8.6 * 10^{3}\right)$. You know that the initial concentration of HA is 1.6 M and that the equivalence point is reached when you add exactly 50 mL of B . Determine the pH value of the solution at the equivalence point. (Temperature $=25^{\circ} \mathrm{C}$ )
$V(H A)_{i}=\frac{1 \times 0.05}{1.6}=0.03125 \mathrm{~L}$
$V(\text { Solution })_{e q}=0.03125 L+0.05 L=0.08125 L$
$n\left(A^{-}\right)_{\text {eq }}=0.05 L \times 1 M=0.05 \mathrm{~mol}$
$\left[A^{-}\right]_{e q}=\frac{0.05 \mathrm{~mol}}{0.08125 \mathrm{~L}}=0.615384615 \mathrm{M}$
$K_{b}(H A)=2.0 \times 10^{-3}=\frac{x^{2}}{0.615384615 M-x} \rightarrow x=0.035082321$
$p H=14-p O H=14-(-\log (0.035082321))=12.54508832 \simeq 12.55$
3. $\mathrm{A} \mathrm{Na}_{2} \mathrm{SO}_{3}$ solid is exposed to the air for an extended period of time. Some of the $\mathrm{Na}_{2} \mathrm{SO}_{3}$ is oxidized by the oxygen in the air. To determine the purity of the $\mathrm{Na}_{2} \mathrm{SO}_{3}$, a 3.5 g sample is taken from the solid and reacted with excess $\mathrm{H}_{2} \mathrm{SO}_{4} .560 \mathrm{~mL}$ of gas is produced under standard conditions. What is the purity in percent of the $\mathrm{Na}_{2} \mathrm{SO}_{3}$ solid?

$$
\begin{aligned}
& n\left(\mathrm{SO}_{2}\right)=0.56 \mathrm{~L} \div 22.4 \mathrm{~L} / \mathrm{mol}=0.025 \mathrm{~mol} \\
& m\left(\mathrm{Na}_{2} \mathrm{SO}_{3}\right)=0.025 \mathrm{~mol} \times 126.043 \mathrm{~g} / \mathrm{mol}=3.151075 \mathrm{~g} \\
& \% \mathrm{Na}_{2} \mathrm{SO}_{3}=\frac{3.151075 \mathrm{~g}}{3.5 \mathrm{~g}} \times 100 \% \simeq 90.0 \%
\end{aligned}
$$

1. Calculate the pH once 5 mL of 1 M HCl is added to 500 mL of a buffer solution of 0.4 M $\mathrm{CH}_{3} \mathrm{COOH}$ (a weak acid) and $0.1 \mathrm{M} \mathrm{CH}_{3} \mathrm{COO}^{-}$. (Given: $\left.\mathrm{K}_{\mathrm{b}}\left(\mathrm{CH}_{3} \mathrm{COO}^{-}\right)=5.56 \times 10^{-10}\right)$

Find moles of $\mathrm{CH}_{3} \mathrm{COOH}$ and $\mathrm{CH}_{3} \mathrm{COO}-$
$\mathrm{CH}_{3} \mathrm{COOH}: 0.5^{*} 0.4=0.2$ moles
$\mathrm{CH}_{3} \mathrm{COO}-: ~ 0.1 * 0.5=0.05$ moles

Find moles of HCl
$0.005^{*} 1=0.005$ moles

Find pKa
$10^{\wedge}-14 / 5.56^{*} 10^{\wedge}-10=1.8^{*} 10^{\wedge}-5$
$-\log \left(1.8^{*} 10^{\wedge}-5\right)=5.1$

Henderson Hasselbach
$5.1-\log (0.2+0.005 / 0.05-0.005)=4.438$
2. What happens to the boiling point of water when you dissolve rock climbing chalk (magnesium carbonate) in it?
The boiling point increases
3. A certain amount of $\mathrm{SO}_{2}(\mathrm{a} \mathrm{L})$ and $\mathrm{H}_{2} \mathrm{~S}(\mathrm{bL})$ are mixed under room temperature and normal atmospheric pressure. They react to form yellow solids (S) and water. If the volume of the gases after reaction is only $1 / 4$ of the volume before the reaction, what is the ratio of $a: b$ ? (There are two possible ratios, list both of them).
Answer:

a. $\mathrm{SO}_{2}$ is the limiting reactant

$$
\frac{b-2 a}{a+b}=1 / 4 \rightarrow a: b=1: 3
$$

b. $H_{2} S$ is the limiting reactant

$$
\frac{a-1 / 2 \times b}{a+b}=1 / 4 \rightarrow a: b=1: 1
$$

1. A mixture contains $\mathrm{Na}_{2} \mathrm{~S}, \mathrm{Na}_{2} \mathrm{SO}_{3}, \mathrm{Na}_{2} \mathrm{SO}_{4}$. Knowing that there is $38 \%$ of sulfur in this mixture (by mass). What is the percentage of oxygen (by mass)?

$$
\begin{gathered}
\% \mathrm{Na}=38 \% \times \frac{2 \times 22.990 \mathrm{~g} / \mathrm{mol}}{32.066 \mathrm{~g} / \mathrm{mol}}=54.49 \% \\
\% \mathrm{O}=100 \%-38 \%-54.49 \%=7.51 \%
\end{gathered}
$$

2. $\mathrm{A} / \mathrm{an}$ $\qquad$ is a mixture of two compounds with well-defined boiling points that cannot be separated through fractional distillation.

Azeotrope
3. Write the chemical formula of Cobalt (III) oxalate $\left(\mathrm{Co}_{2}\left(\mathrm{C}_{2} \mathrm{O}_{4}\right)_{3}\right)$

1. Given the following reaction: $1 / 2 \mathrm{~N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{NO}_{2}(\mathrm{~g}) \quad \Delta \mathrm{H}_{\mathrm{f}}=+33.2 \mathrm{~kJ}$

If you have 2.1 L (at STP) of $\mathrm{N}_{2}$, which is the limiting reagent, how much heat is needed for the reaction to use up all the $\mathrm{N}_{2} .(1 \mathrm{~mol}=22.4 \mathrm{~L}$ at STP$)$
Answer key:
$1 / 2 \mathrm{~N}_{2}(\mathrm{~g})+\mathrm{O}_{2}(\mathrm{~g}) \rightarrow \mathrm{NO}_{2}(\mathrm{~g}) \quad \Delta \mathrm{H}_{\mathrm{f}}=+33.2 \mathrm{~kJ}$
If you have 2.1 L (at STP) of $\mathrm{N}_{2}$, which is the limiting reagent, how much heat is needed for the reaction to use up all the $\mathrm{N}_{2} ?(1 \mathrm{~mol}=22.4 \mathrm{~L}$ at STP$)$
2.1 $\mathrm{L} \mathrm{N}_{2} \times 1 \mathrm{~mol} \mathrm{~N}_{2} / 22.4 \mathrm{~L} \mathrm{~N}_{2}=0.093725 \mathrm{~mol} \mathrm{~N}_{2}$
$0.093725 \mathrm{~mol} \mathrm{~N}_{2} \quad \mathrm{x} \quad 1 \mathrm{~mol}$ of rxn / $1 / 2 \mathrm{~mol} \mathrm{~N}_{2} \quad \mathrm{x} \quad 33.2 \mathrm{~kJ} / 1 \mathrm{~mol} \mathrm{rxn}=\mathbf{6} .225 \mathrm{~kJ}$
2. An unknown gas AB decomposes into A and B in the following reaction:

$$
\mathrm{AB}_{(\mathrm{g})} \rightleftharpoons \mathrm{A}_{(\mathrm{g})}+\mathrm{B}_{(\mathrm{g})}
$$

10 g of AB is sealed into an evacuated 1.000 L container at $100^{\circ} \mathrm{C}$. After equilibrium is reached, the total pressure in the container is 3.50 atm . Given the molar mass of AB is $134.96 \mathrm{~g} / \mathrm{mol}$, calculate the equilibrium constant K for the decomposition reaction above.

```
    10gAB}\times\frac{1\textrm{mol}}{134.96\textrm{g}}=7.41\times1\mp@subsup{0}{}{-2}\textrm{mol
    \rho = \frac { n R T } { V } = \frac { ( 7 . 4 1 \times 1 0 ^ { - 2 } ) ( 0 . 0 8 2 0 6 ) ( 3 7 3 \mathrm { K } ) } { 1 . 0 0 0 \mathrm { L } } = 2 . 2 7 \mathrm { ath }
    AB }\vec{A+B}\quad3.50=(2.27-X)+x+
I 2.27 0 0 0 x cel.23
C-x +x +k
E 2.27-x x x
E 1.04ata l.23atm 1.23atm
    K=}\frac{(.23)(1.23)}{c.04}=1.45
```


## Answer: 1.455

3. Consider the following decomposition reaction of an unknown gas AB :

$$
2 \mathrm{AB}_{(\mathrm{g})} \rightleftharpoons \mathrm{A}_{2(\mathrm{~g})}+\mathrm{B}_{2(\mathrm{~g})}
$$

50 g of AB is sealed into a 1.5 L evacuated container and heated to $500^{\circ} \mathrm{C}$. The equilibrium constant of this reaction at this temperature is $2.59 \times 10^{-5}$. Find the equilibrium partial pressure of AB to 3 significant figures given the total pressure at equilibrium is 6.45 atm .


```
C-2x+x+x n=6.45
E n-2x +x +x
    2.54\times1\mp@subsup{0}{}{-5}=\frac{\mp@subsup{x}{}{2}}{(6.45-2x\mp@subsup{)}{}{2}}
        2.54\times10-5}=\frac{\mp@subsup{x}{}{2}}{6.4\mp@subsup{5}{}{2}
        x=3.28\times10-2
    n-2x=6.38
```


## HARD PACKET 10

1. The following is the equations and $\mathrm{K}_{\mathrm{a}}$ values of the dissociation of an unknown weak diprotic acid, $\mathrm{H}_{2} \mathrm{~A}$ :

$$
\begin{array}{ll}
\mathrm{H}_{2} \mathrm{~A}+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{HA}^{-}+\mathrm{H}_{3} \mathrm{O}^{+} & \mathrm{K}_{\mathrm{a} 1}=4.3 \times 10^{-7} \\
\mathrm{HA}-+\mathrm{H}_{2} \mathrm{O} \rightleftharpoons \mathrm{~A}^{2-}+\mathrm{H}_{3} \mathrm{O}^{+} & \mathrm{K}_{\mathrm{a} 2}=5.6 \times 10^{-11}
\end{array}
$$

A 0.12 M solution of $\mathrm{H}_{2} \mathrm{~A}$ is titrated with 0.1 M solution of NaOH . Given that 150 mL of the NaOH solution is required to reach the first equivalence point of $\mathrm{H}_{2} \mathrm{~A}$, what is the pH of the solution at the second equivalence point?

```
Mokn of wroll for frit eq panut =0.150L\times0.1 =0.015mol
    V of H2A: 0.015m0d= (0.12M)(V)
        V=0.125L
    At seoond eq. pount, all H2A}->\mp@subsup{H}{2}{22}\mathrm{ so 0.015 pul Al }\mp@subsup{A}{}{2-
        total V ef soln = 0.125L+0.150L+0.150L = 0.425L
[\mp@subsup{A}{}{2-}]=\frac{0.005\textrm{m}}{0.425L}=3.53\times1\mp@subsup{0}{}{-2}\textrm{M}
H2O+\mp@subsup{\textrm{H}}{}{2-}\rightleftharpoons\mp@subsup{\textrm{OH}}{}{-}+\mp@subsup{\textrm{HA}}{}{-}
                Fk}=\frac{1\mp@subsup{0}{}{-14}}{5.6\times1\mp@subsup{0}{}{-4}}=1.79\times1\mp@subsup{0}{}{-4
    3.53\times10-2 
        1.79\times10 -4}=\frac{\mp@subsup{x}{}{2}}{3.53\times1\mp@subsup{0}{}{-2}-x
        1.79\times10-4}\approx\frac{\mp@subsup{x}{}{2}}{3.53\times1\mp@subsup{0}{}{-2}
        4~2.51\times10-3
\mathrm{ suresue }->x=2.426\times1\mp@subsup{0}{}{-3}=[2H]
pH=14+\operatorname{log}(2.426\times1\mp@subsup{0}{}{-3})=11.38
```

2. According to solvent leveling, what is the strongest acid that can exist in water?
$\mathrm{H}_{3} \mathrm{O}^{+}$
3. According to solvent leveling, what is the strongest base that can exist in water? $\mathrm{OH}^{-}$
