



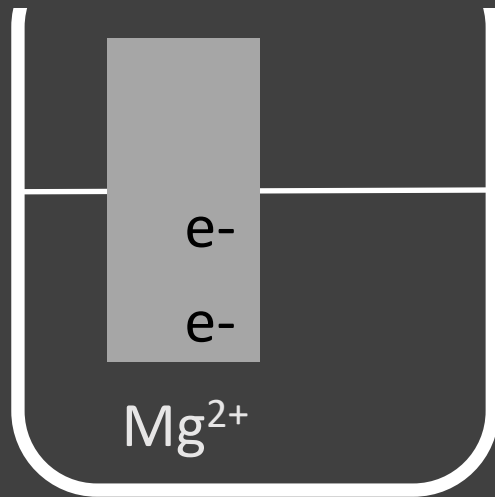
A2 Physical Chemistry

Measuring Electrode Potentials

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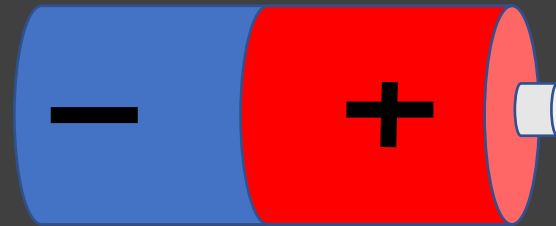
Measuring Electrode Potentials

We charge that collects on the electrode is known as the electrode potential and the Mg^{2+}/Mg equilibrium is known as a half cell.



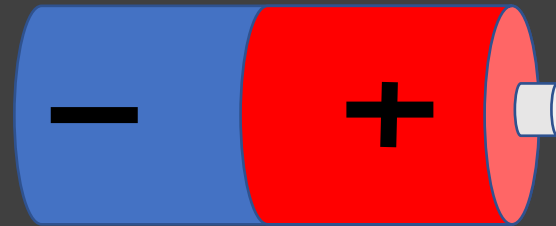
However in order to measure the electrode potential for Mg you need to form a complete circuit.

Measuring Electrode Potentials



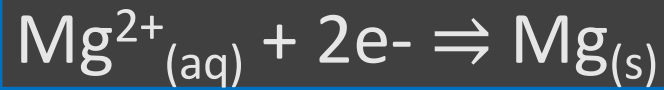
We can compare our situation with a battery. A battery has a negative and positive end.

Measuring Electrode Potentials

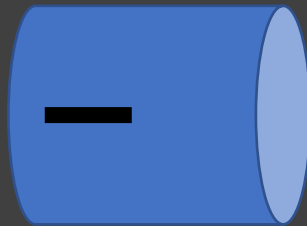


We can compare our situation with a battery. A battery has a negative and positive end. The voltage is the difference in electrical potential between the two ends.

Measuring Electrode Potentials



Electrode Potential A

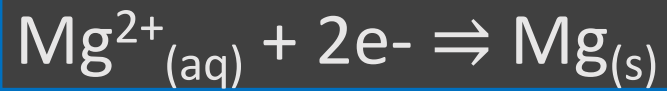


We can compare our situation with a battery. A battery has a negative and positive end. The voltage is the difference in electrical potential between the two ends.

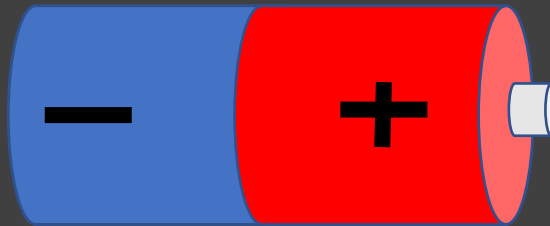
Currently with our Mg^{2+}/Mg system we only have one end.

Therefore we have to have another half cell in order to measure the electrode potential of the Mg^{2+}/Mg half cell.

Measuring Electrode Potentials



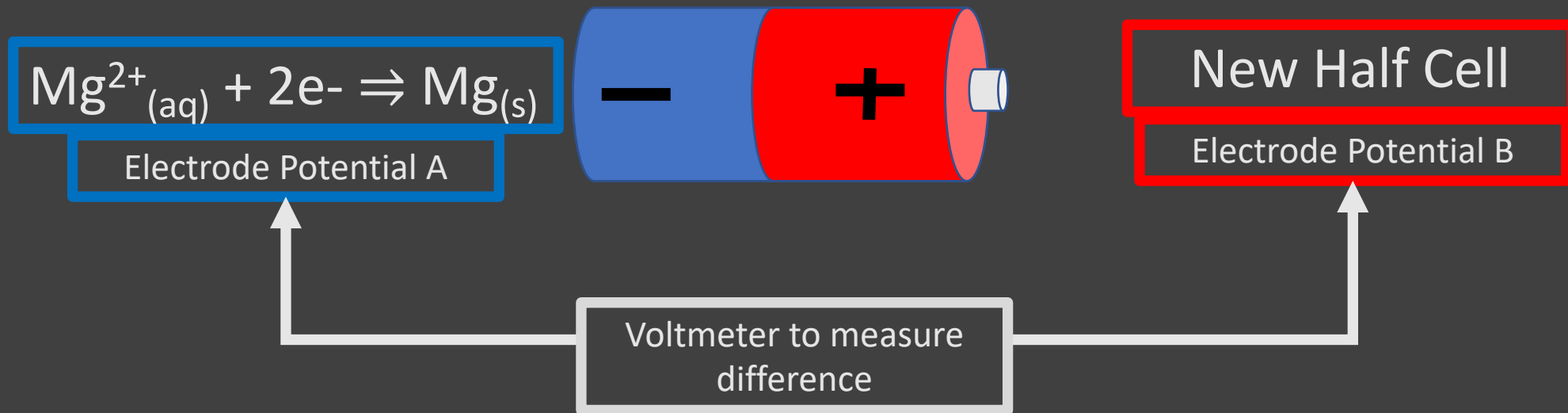
Electrode Potential A



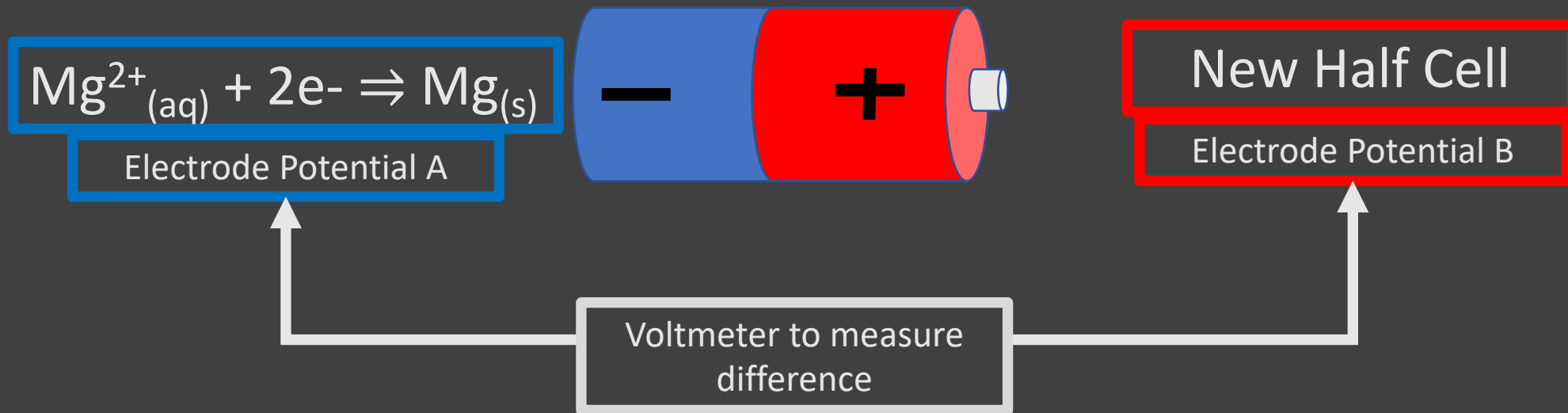
New Half Cell

Electrode Potential B

Measuring Electrode Potentials

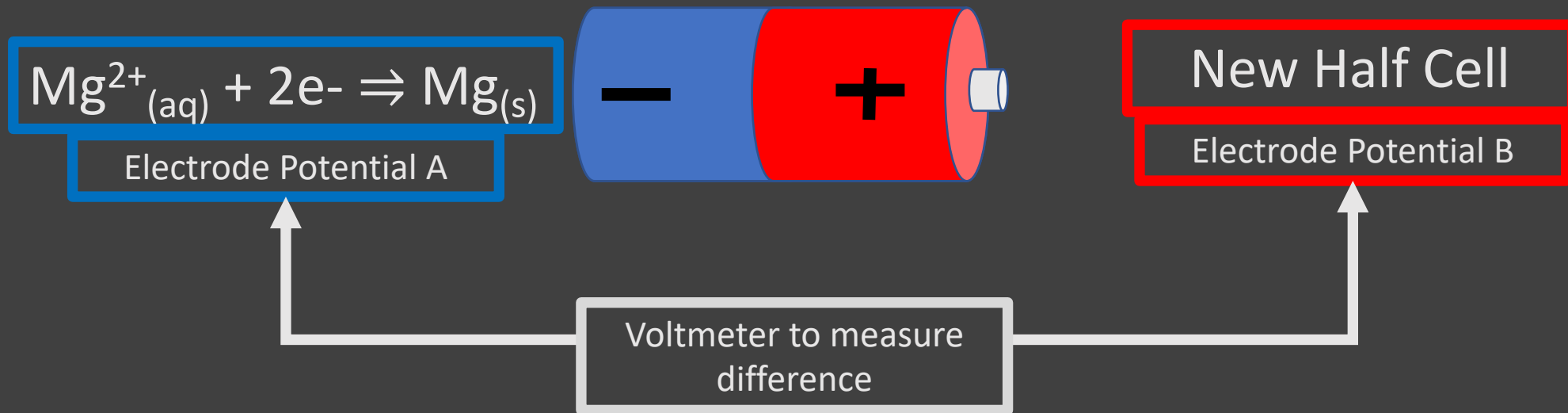


Measuring Electrode Potentials



$$\text{Electrode Potential A} = \text{Voltage} - \text{Electrode Potential B}$$

Measuring Electrode Potentials

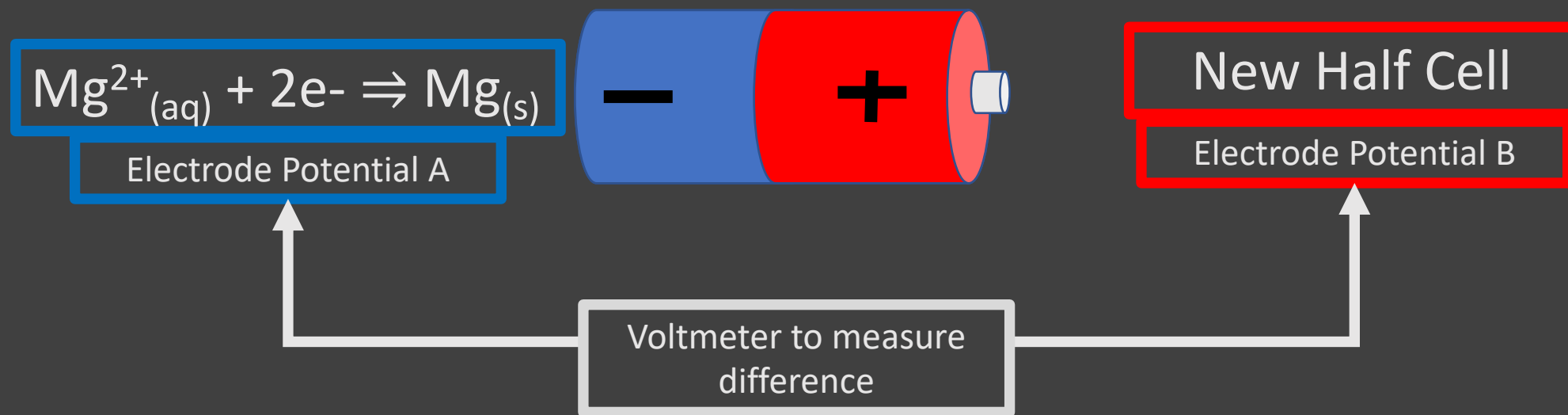


$$\text{Electrode Potential A} = \text{Voltage} - \text{Electrode Potential B}$$

However how we will know the electrode potential of the Mg^{2+}/Mg electrode as the other half cell?

If we don't know Electrode Potential B, we cannot find Electrode Potential A.

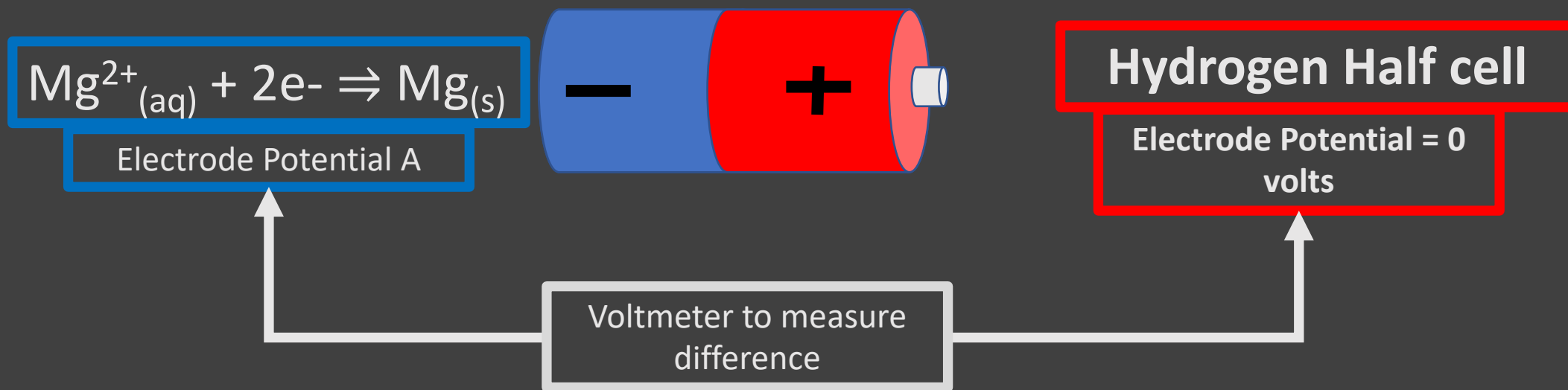
Measuring Electrode Potentials



$$\text{Electrode Potential A} = \text{Voltage} - \text{Electrode Potential B}$$

We deal with this by setting the defining the electrode potential of the $2\text{H}^+/\text{H}_2$ half cell as 0 volts. Electrode potentials of all other half cells are then quoted compared to the hydrogen half cell.

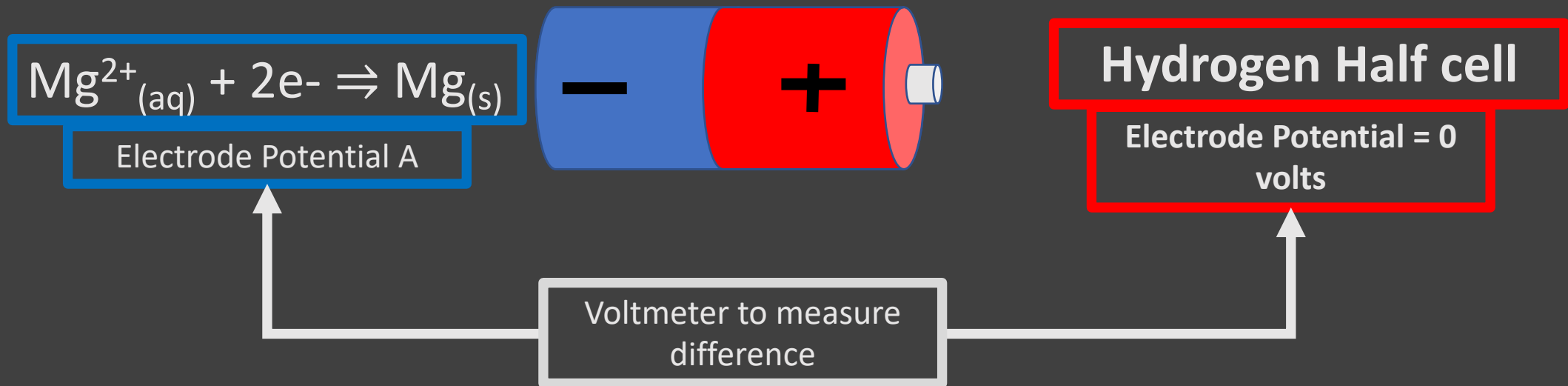
Measuring Electrode Potentials



$$\text{Electrode Potential A} = \text{Voltage} - \text{Electrode Potential B}$$

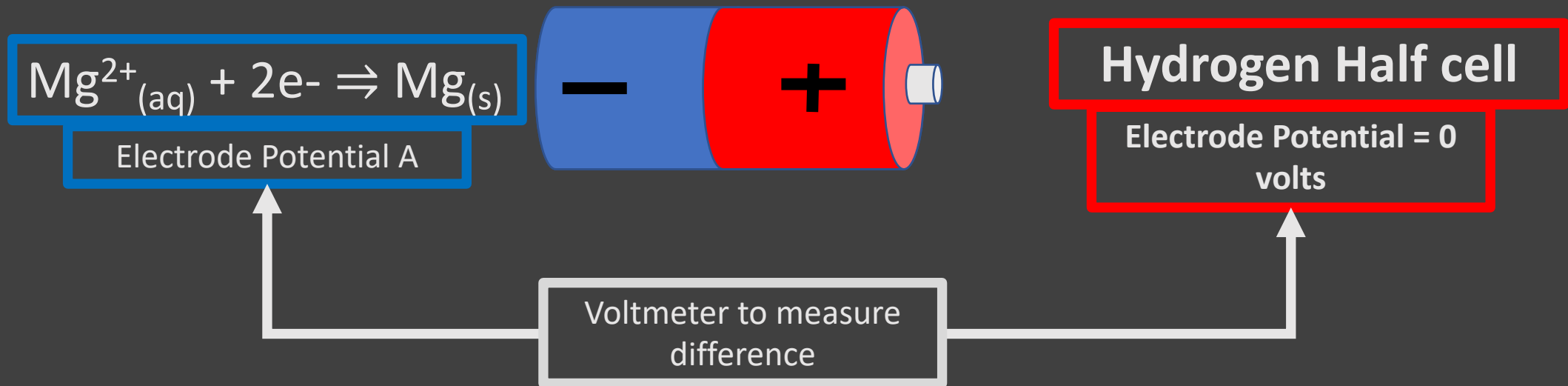
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Measuring Electrode Potentials



$$\text{Electrode Potential A} = \text{Voltage} - 0$$

Measuring Electrode Potentials



Electrode Potential A = Voltage

Hydrogen Half Cell



Hydrogen Half Cell



No metal to form a circuit?

How do we deal with gases?

What concentration of $\text{H}^+_{(\text{aq})}$ to use?

No metal to form a circuit?

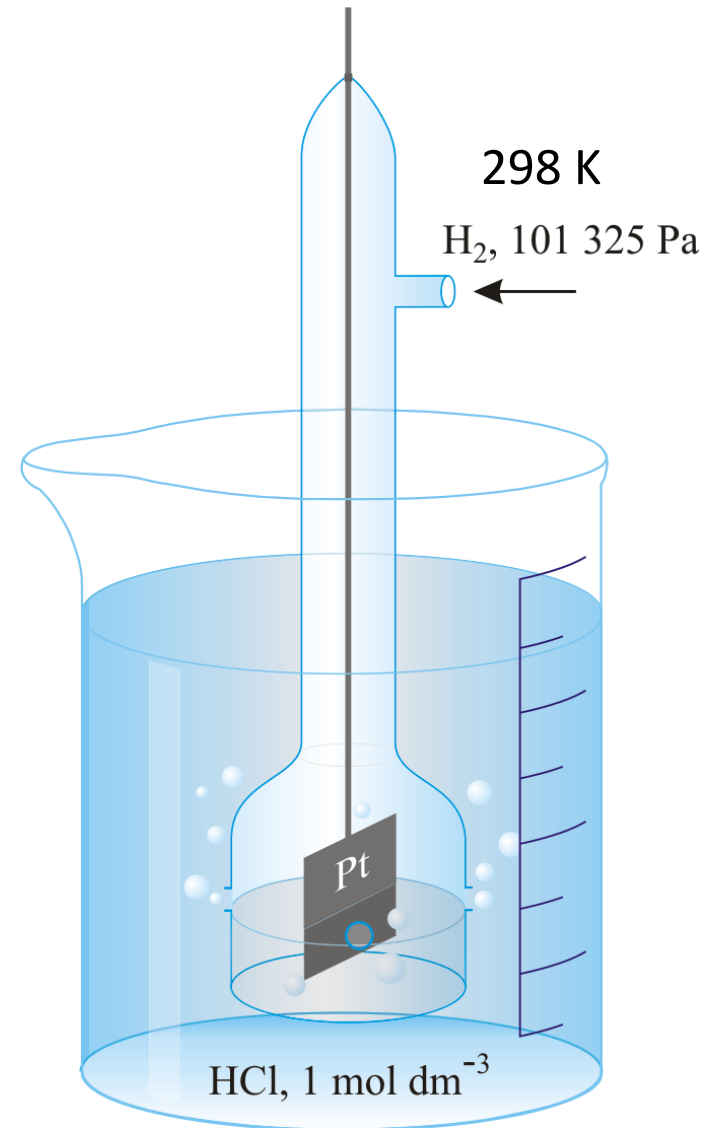
How do we deal with gases?

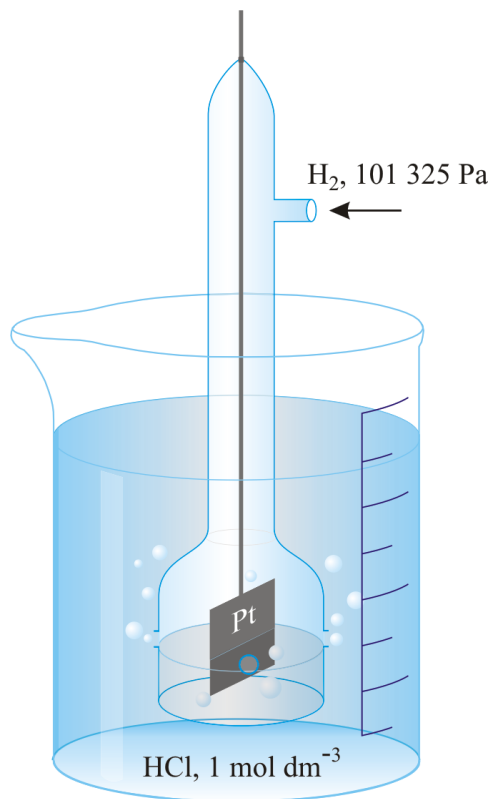
What concentration of $\text{H}^+_{(\text{aq})}$ to use?

Hydrogen Half Cell

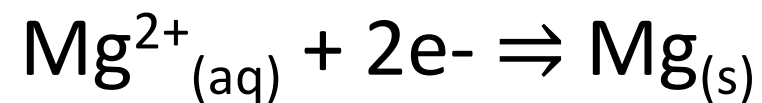
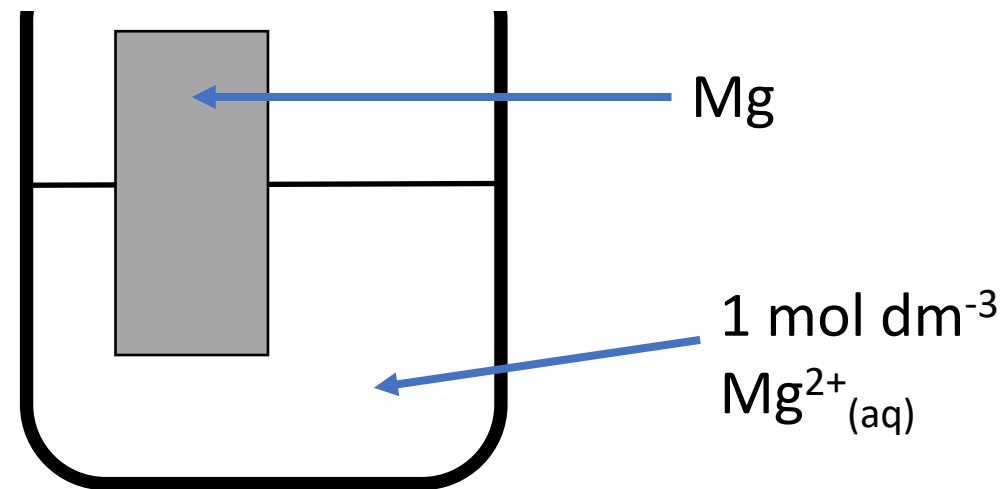


Standard Electrode Potential = 0 volts

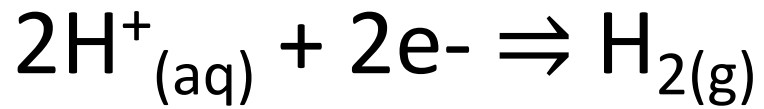
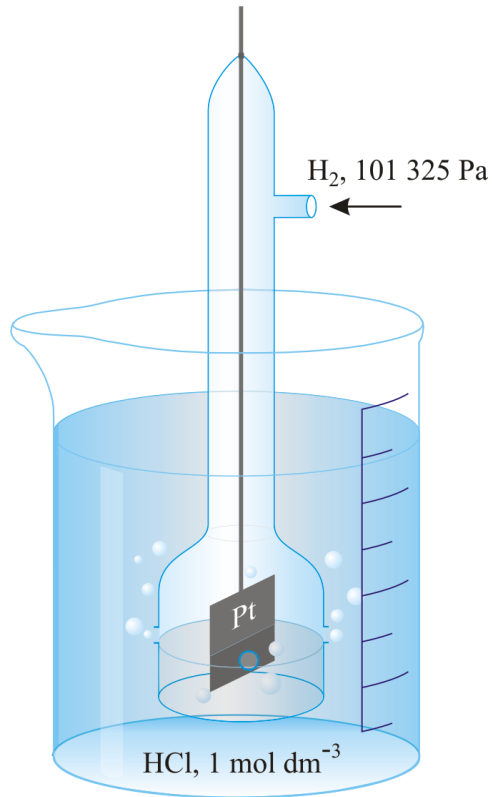




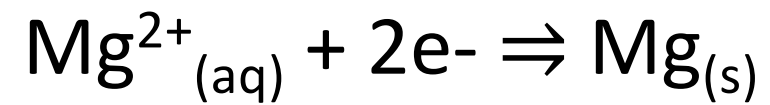
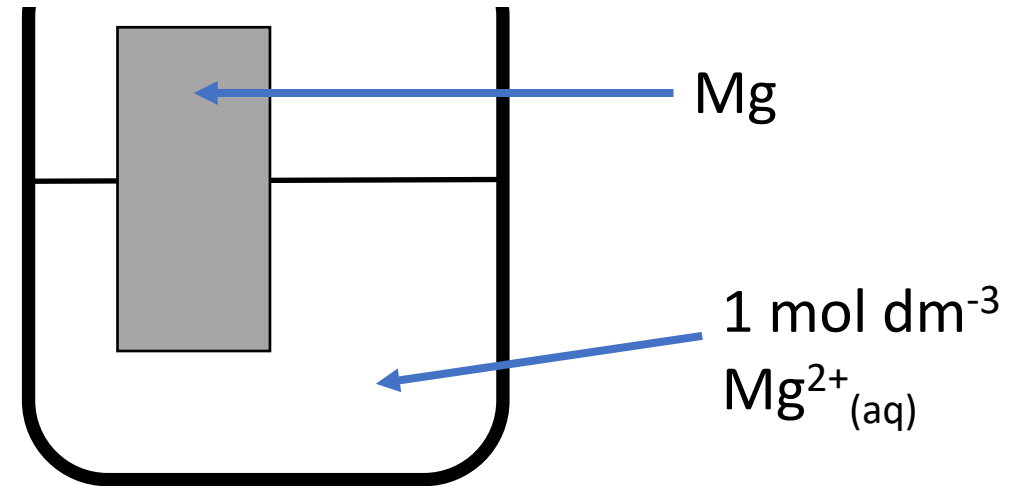
Standard Electrode Potential = 0 volts

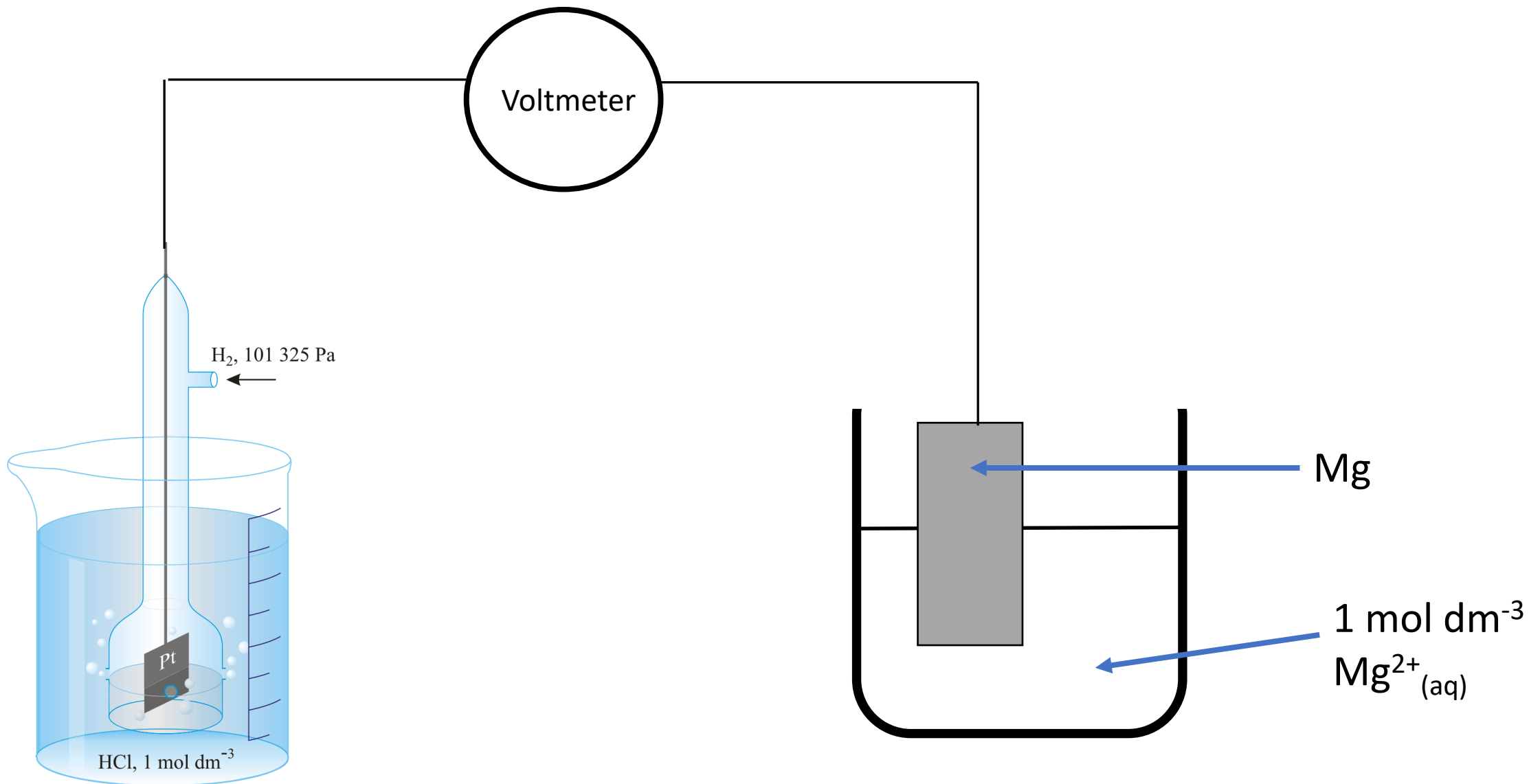


Voltmeter

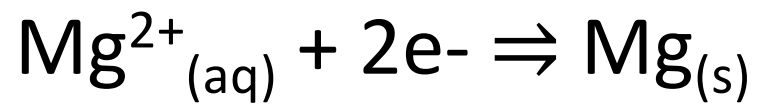


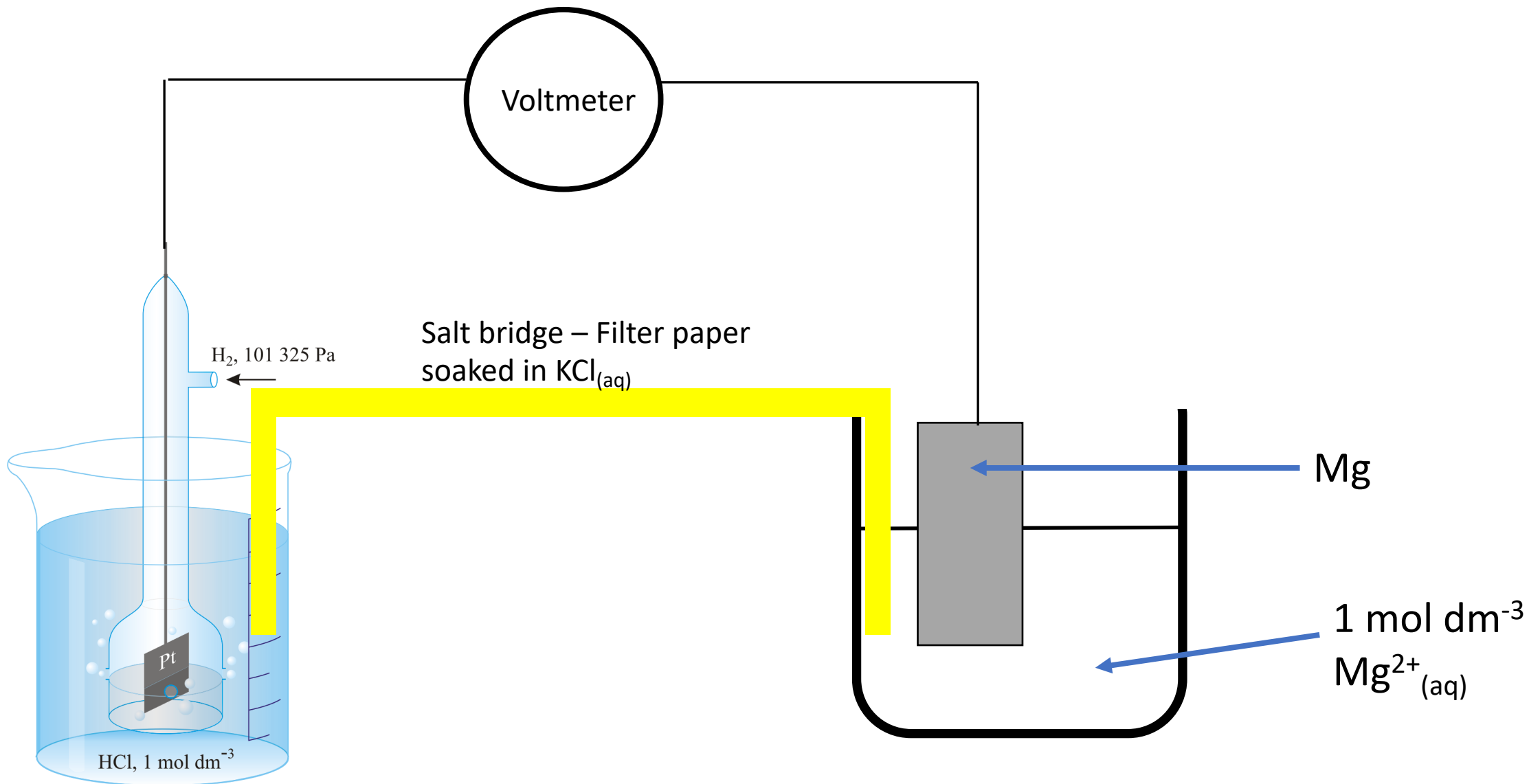
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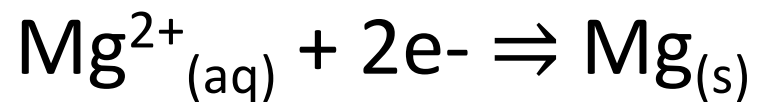


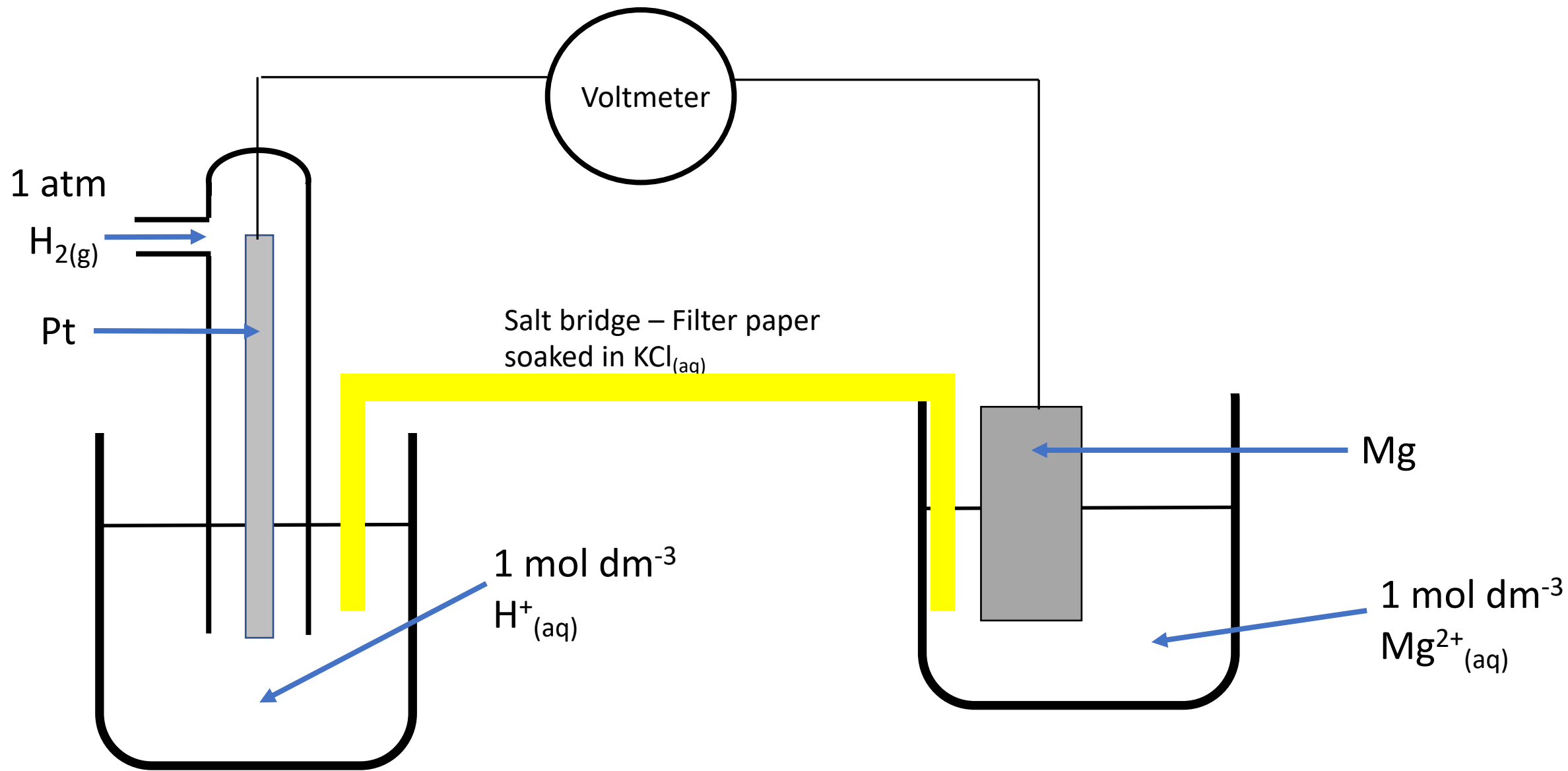
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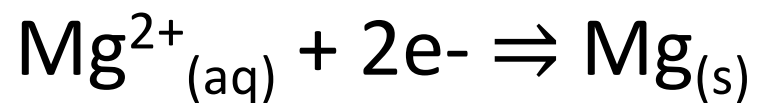


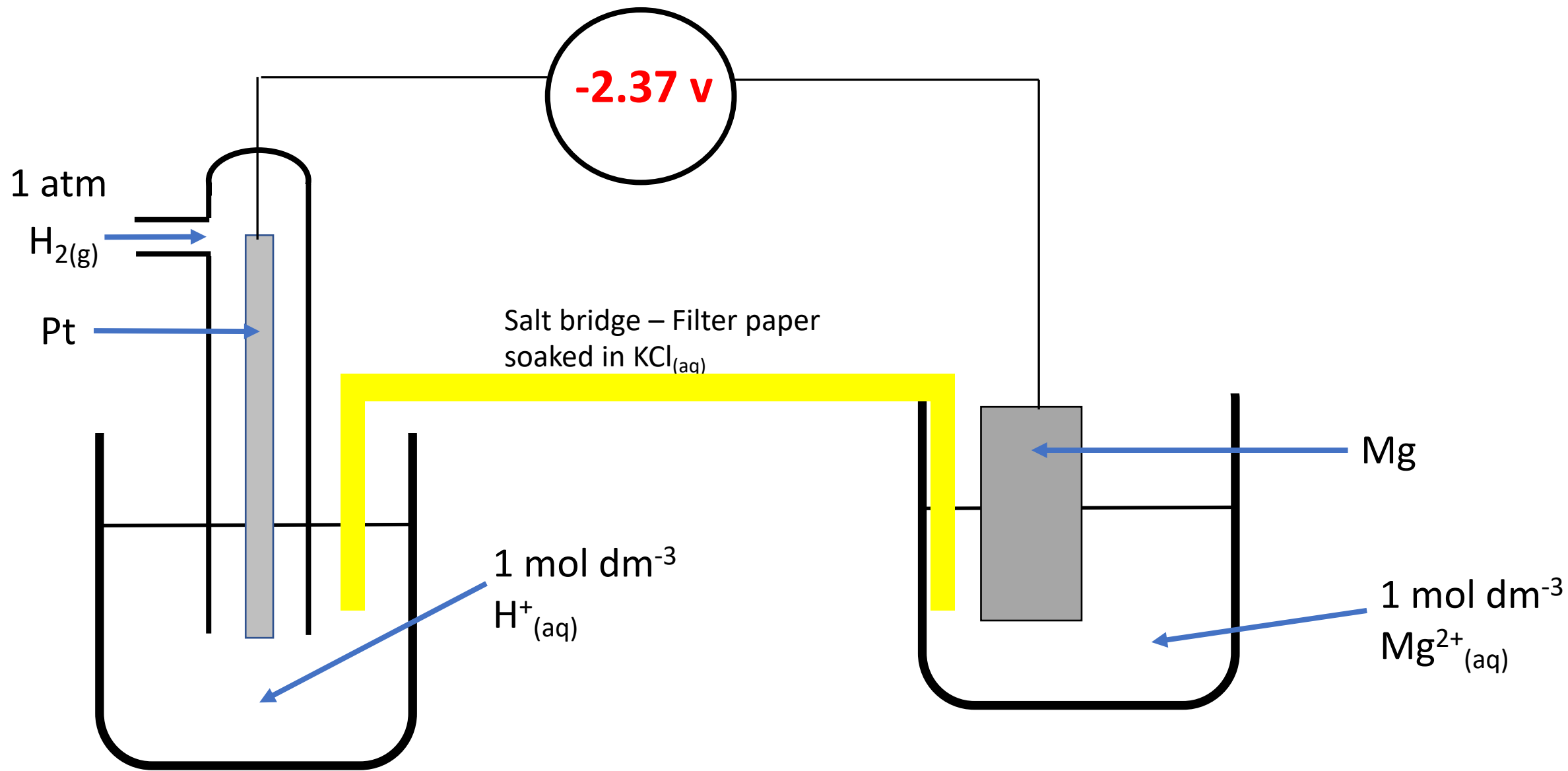
Standard Electrode Potential = 0 volts



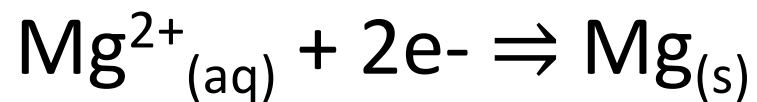


Standard Electrode Potential = 0 volts

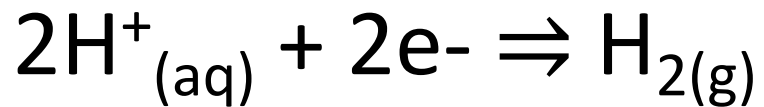
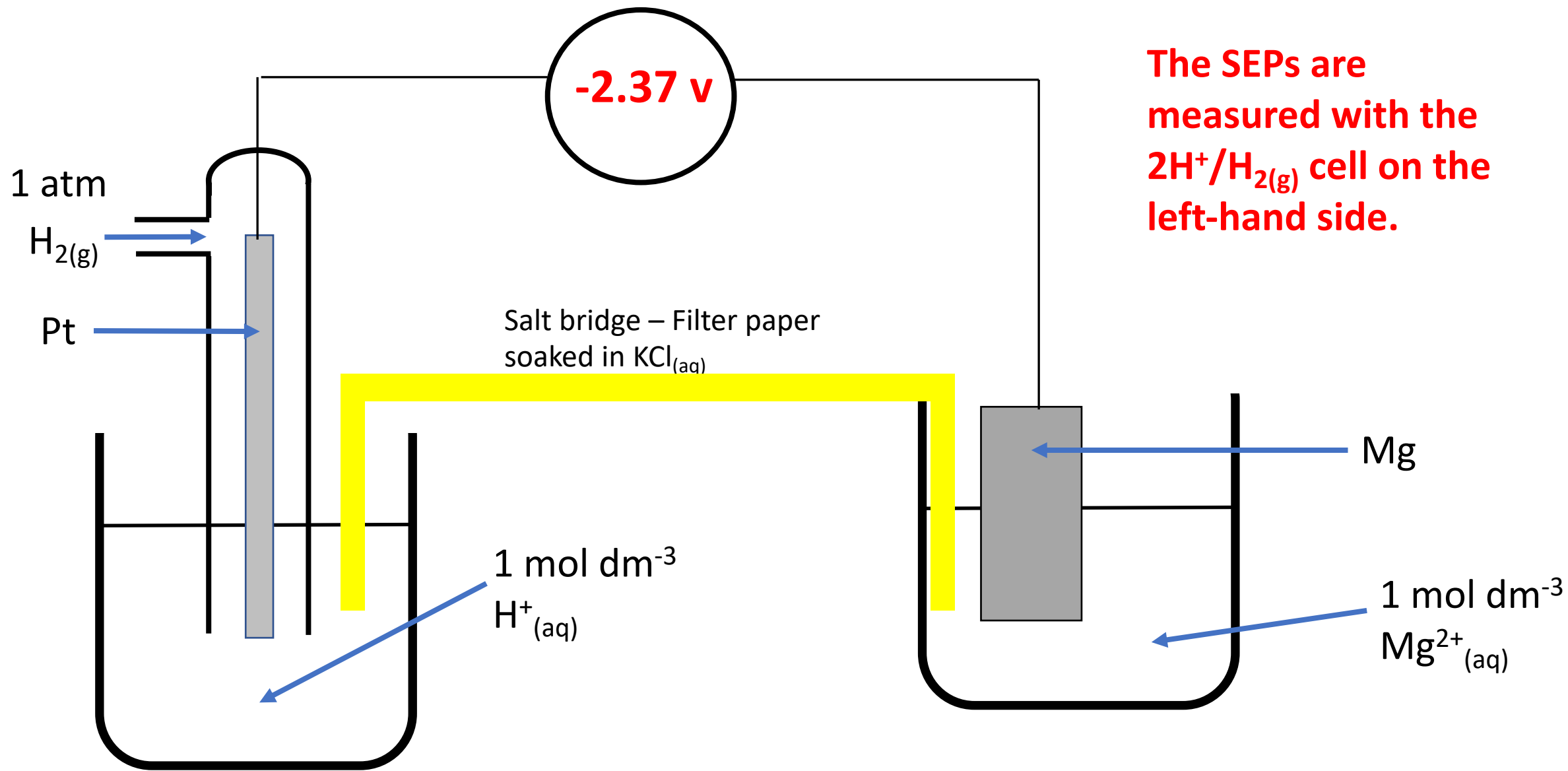




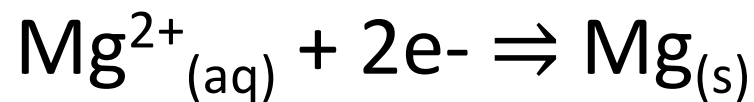
Standard Electrode Potential = 0 volts



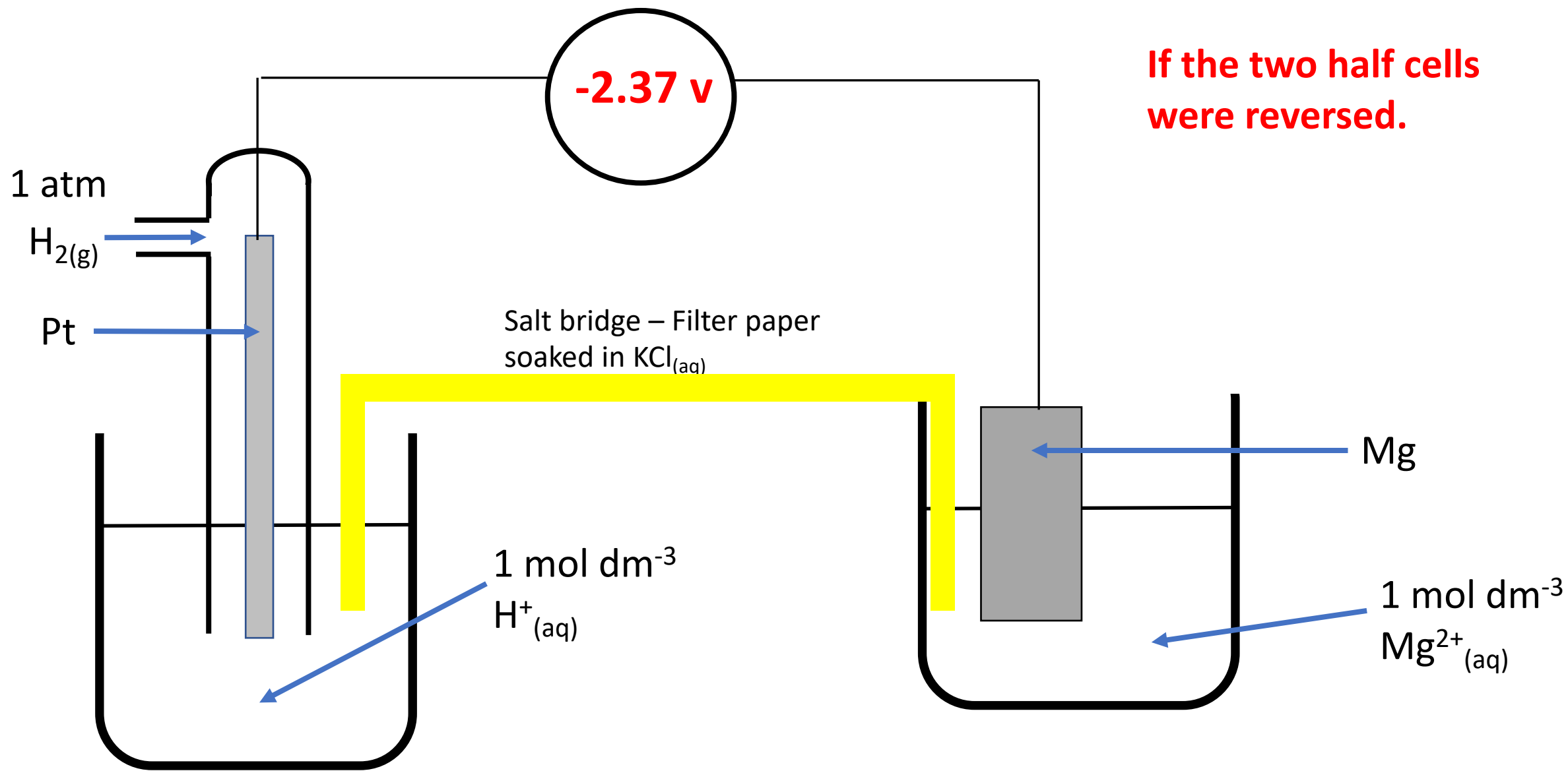
Standard Electrode Potential = ?



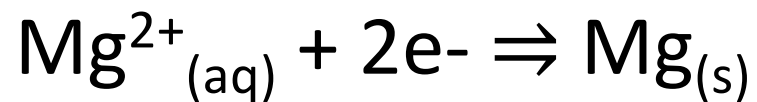
Standard Electrode Potential = 0 volts



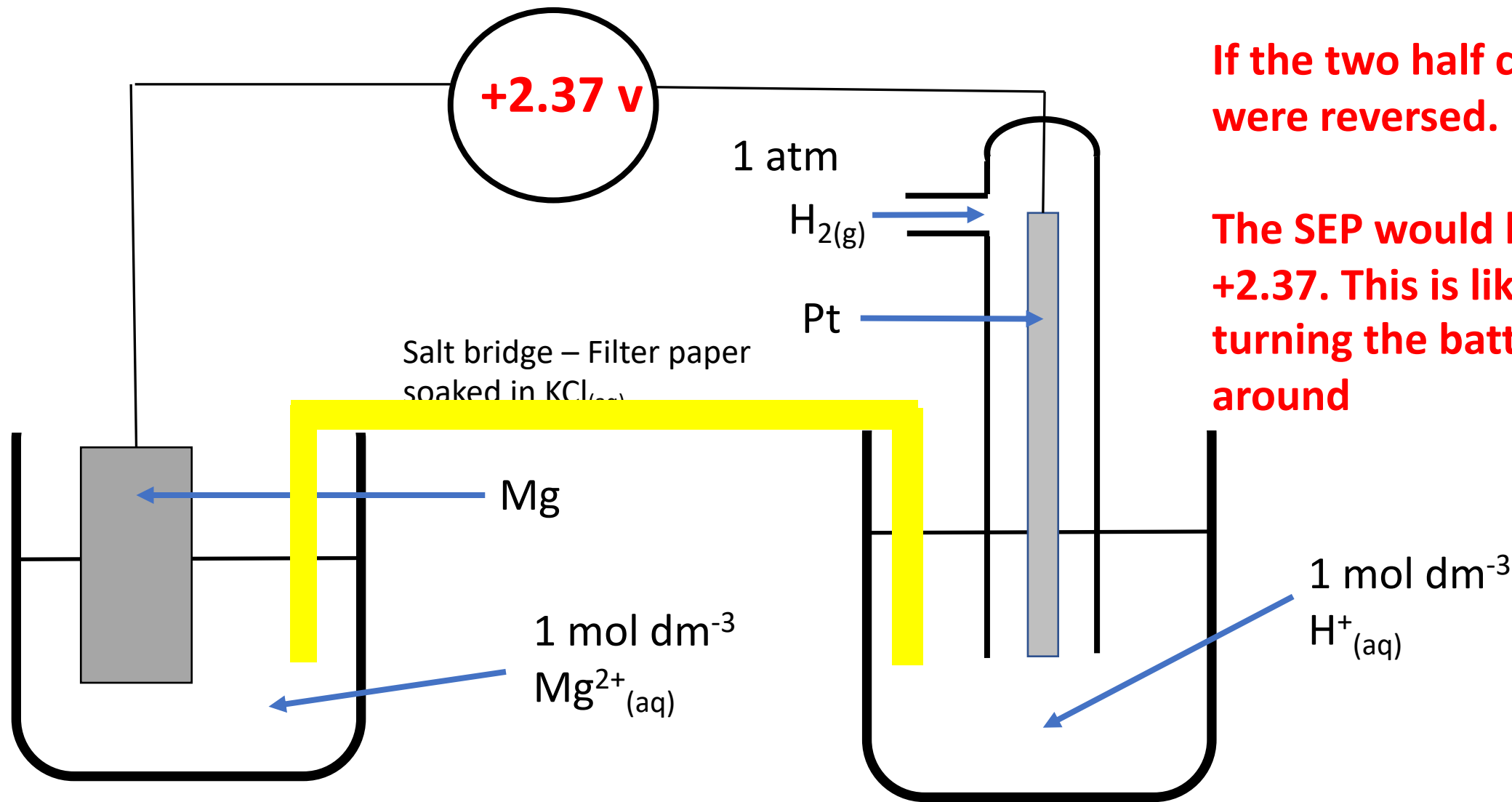
Standard Electrode Potential = ?



Standard Electrode Potential = 0 volts

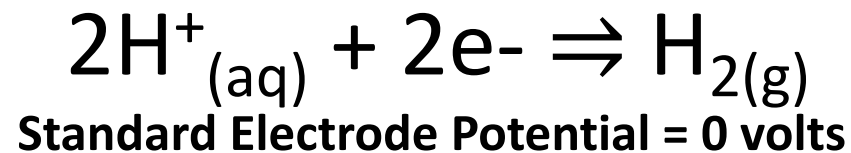
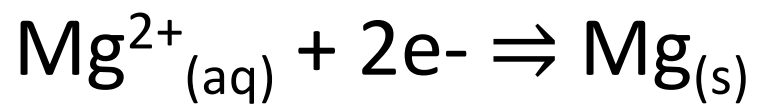


Standard Electrode Potential = ?



If the two half cells were reversed.

The SEP would be +2.37. This is like turning the battery around



Standard Electrode Potentials

Always written as a
reduction half equation –
i.e. electrons on the left
hand side

Half equation	SEP/V
$\text{K}^+(\text{aq}) + \text{e}^- \rightarrow \text{K}(\text{s})$	-2.93
$\text{Ca}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Ca}(\text{s})$	-2.87
$\text{Na}^+(\text{aq}) + \text{e}^- \rightarrow \text{Na}(\text{s})$	-2.71
$\text{Mg}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Mg}(\text{s})$	-2.37
$\text{Al}^{3+}(\text{aq}) + 3\text{e}^- \rightarrow \text{Al}(\text{s})$	-1.66
$\text{Zn}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Zn}(\text{s})$	-0.76
$\text{Fe}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Fe}(\text{s})$	-0.44
$\text{Pb}^{2+}(\text{aq}) + 2\text{e}^- \rightarrow \text{Pb}(\text{s})$	-0.13
$2\text{H}^+(\text{aq}) + 2\text{e}^- \rightarrow \text{H}_2(\text{g})$	0.00
$\text{Cu}^+(\text{aq}) + \text{e}^- \rightarrow \text{Cu}(\text{s})$	0.15
$\text{Ag}^+(\text{aq}) + \text{e}^- \rightarrow \text{Ag}(\text{s})$	0.80
$\text{Au}^{3+}(\text{aq}) + 3\text{e}^- \rightarrow \text{Au}(\text{s})$	1.50

Compared to



Less easily reduced – less likely to gain electrons

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Compared to
 $2\text{H}^+(\text{aq})/\text{H}_2(\text{g})$

Less easily reduced – less likely to gain electrons

More easily reduced – more likely to gain electrons

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