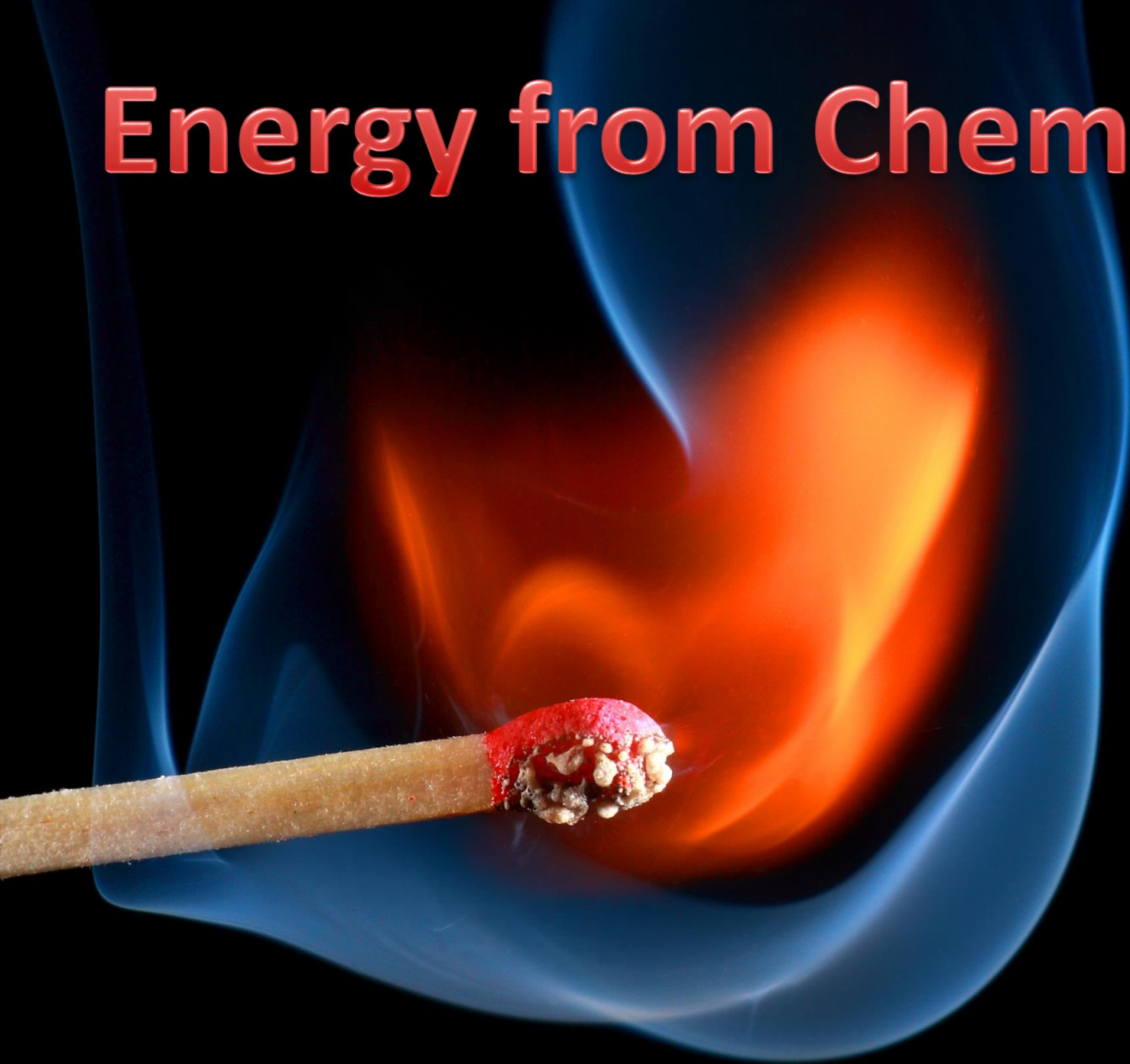
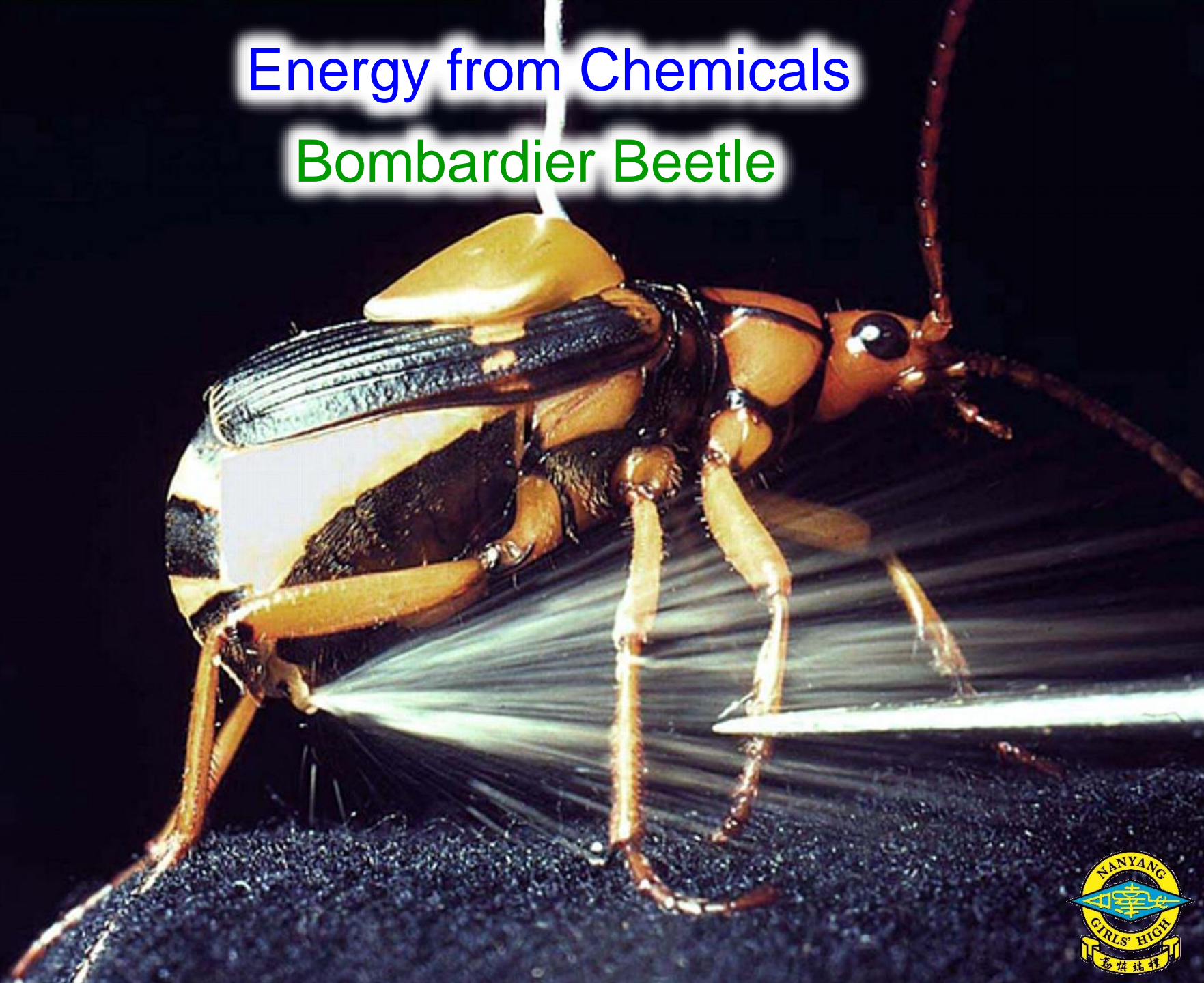


# Energy from Chemicals



# Energy from Chemicals

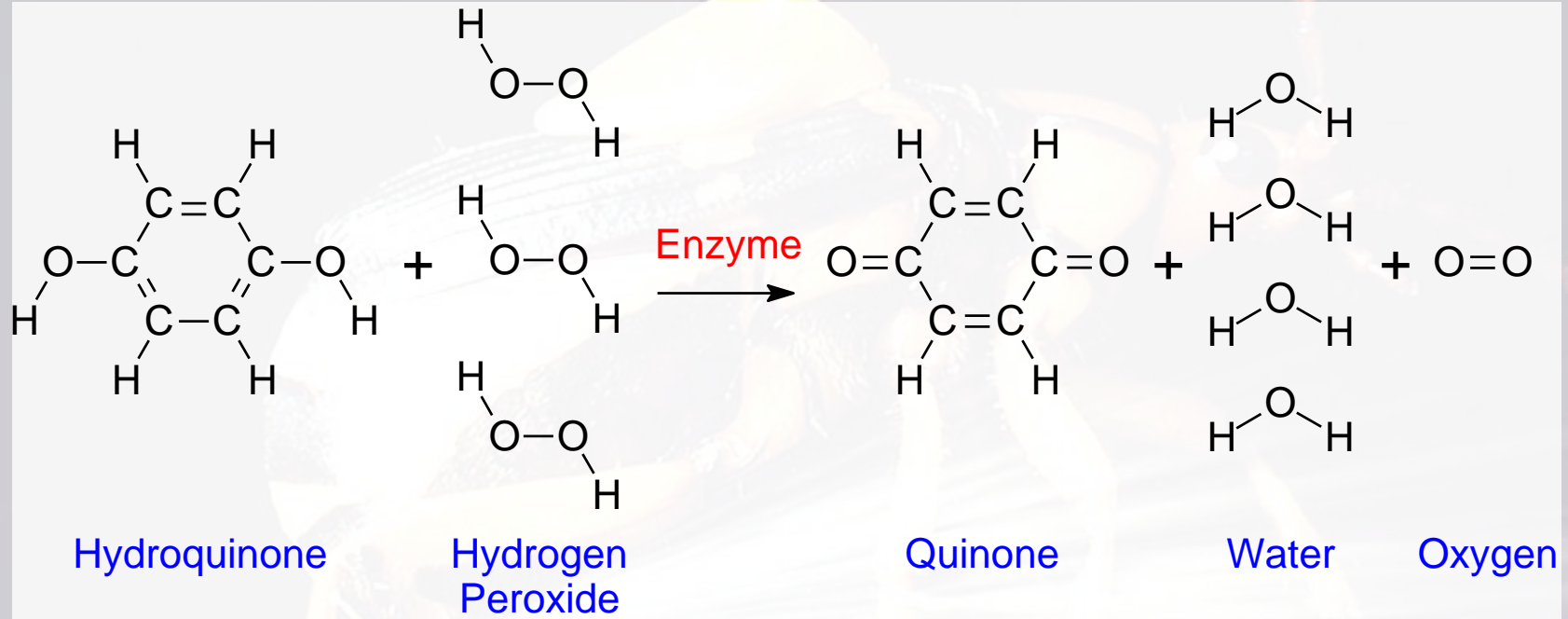
## Bombardier Beetle





# Energy from Chemicals

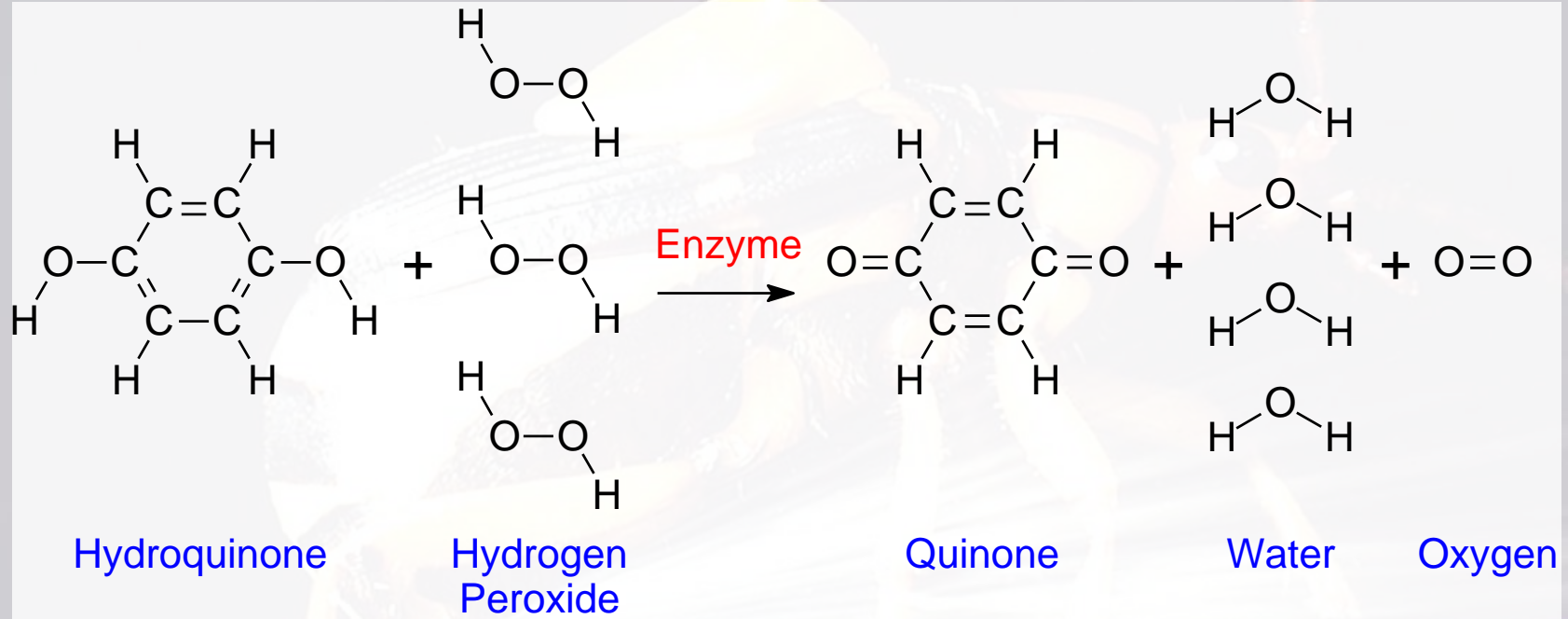
## Bombardier Beetle



- This is an *exothermic* reaction that *releases* approximately *400 kJ of energy* for every mole of hydroquinone that reacts.

# Energy from Chemicals

## Bombardier Beetle



- Result? Whatever is attacking the bombardier beetle is sprayed with a *hot* and *corrosive liquid*.



# Energy from Chemicals



Chemical reactions are always accompanied by a *change* in energy.

- What are some generalisations about *change*?

# Energy from Chemicals

- Some examples of generalisations that can be made:
  - **Change:**
    - Change is inevitable (unavoidable).
    - Change can have positive consequences or negative consequences.
      - Change can be reversible or irreversible.
      - Change can be steady, cyclic, random or chaotic.



# Energy from Chemicals

- **Enduring Understandings:**

- Chemical reactions (change) can be spontaneous or non-spontaneous.
- Spontaneous reactions (changes) release energy which can be used to do work. Non-spontaneous reactions (changes) require energy in order to take place.





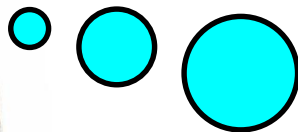
# Energy from Chemicals

- **Essential Questions:**

- How are matter and energy related?
- Why are some chemical reactions exothermic and others endothermic?
- Why do some chemical reactions occur spontaneously, while others do not?
- Where does the heat and light that is given off by an exothermic reaction originate from, and where does it go to?



# Energy from Chemicals



What must I know  
and understand  
about *energy*  
*changes* in  
Chemistry?

# Energy from Chemicals

## Learning Outcomes

### Candidates should be able to:

- a) Describe the meaning of enthalpy change in terms of exothermic ( $\Delta H$  negative) and endothermic ( $\Delta H$  positive) reactions.
- b) Represent energy changes by energy profile diagrams, including reaction enthalpy changes and activation energies.
- c) Describe bond breaking as an endothermic process and bond making as an exothermic process.
- d) Explain overall enthalpy changes in terms of the energy changes associated with the breaking and making of covalent bonds.
- e) Describe hydrogen, derived from water or hydrocarbons, as a potential fuel, reacting with oxygen to generate electricity directly in a fuel cell (details of the construction and operation of a fuel cell are not required).

- Singapore Examinations and Assessment
- Board University of Cambridge International Examinations
- Ministry of Education Singapore





# Energy from Chemicals



What are the different types of energy changes that can take place in Chemistry?

- **Exothermic:** Releases energy into the surroundings.
- **Endothermic:** Absorbs energy from the surroundings.

# Energy from Chemicals – Exothermic

Exo is Greek for “outside.”

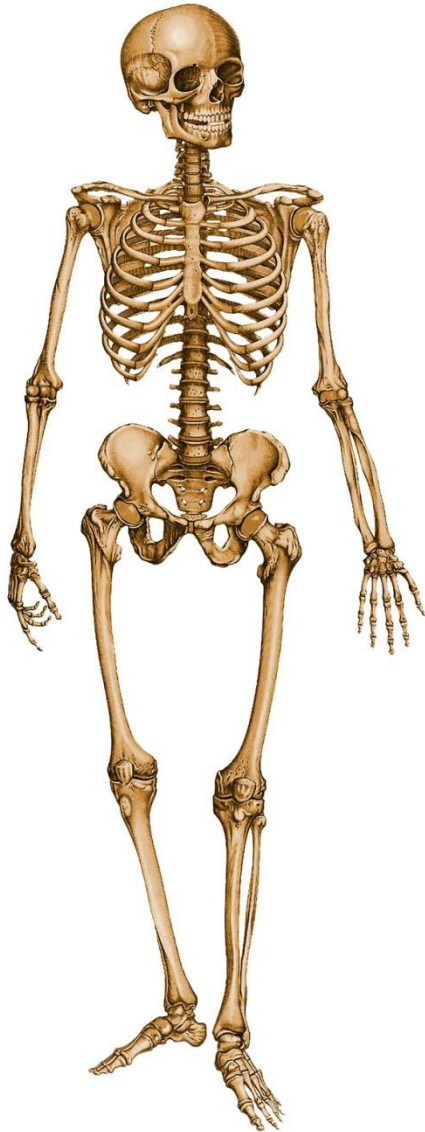
Thermasi is Greek for “to heat.”

The literal meaning of exothermic is therefore “to heat outside.”

Exo is a common term used throughout science. For example, insects are described as having an exoskeleton, which literally means “outside skeleton.” This accurately describes the animal’s hard outer casing.



# Energy from Chemicals – Endothermic



Endo is Greek for “**inside.**”  
Thermasi is Greek for “**to heat.**”  
The literal meaning of endothermic  
is therefore “**to heat inside.**”

Endo is a common term used  
throughout science. For example,  
mammals are described as having  
an endoskeleton, which literally  
means “**inside skeleton.**”



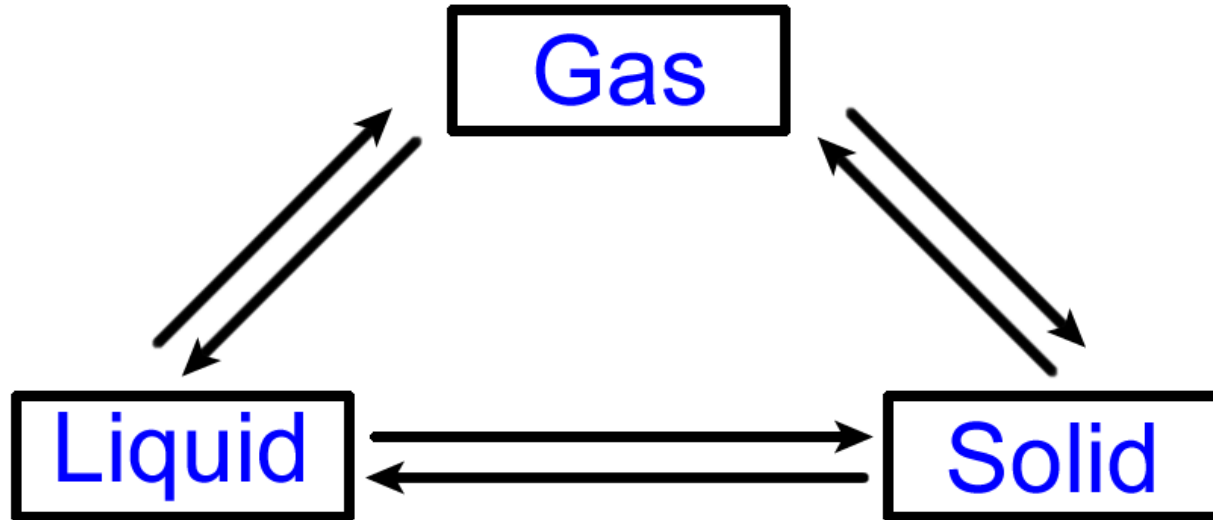


# Energy from Chemicals

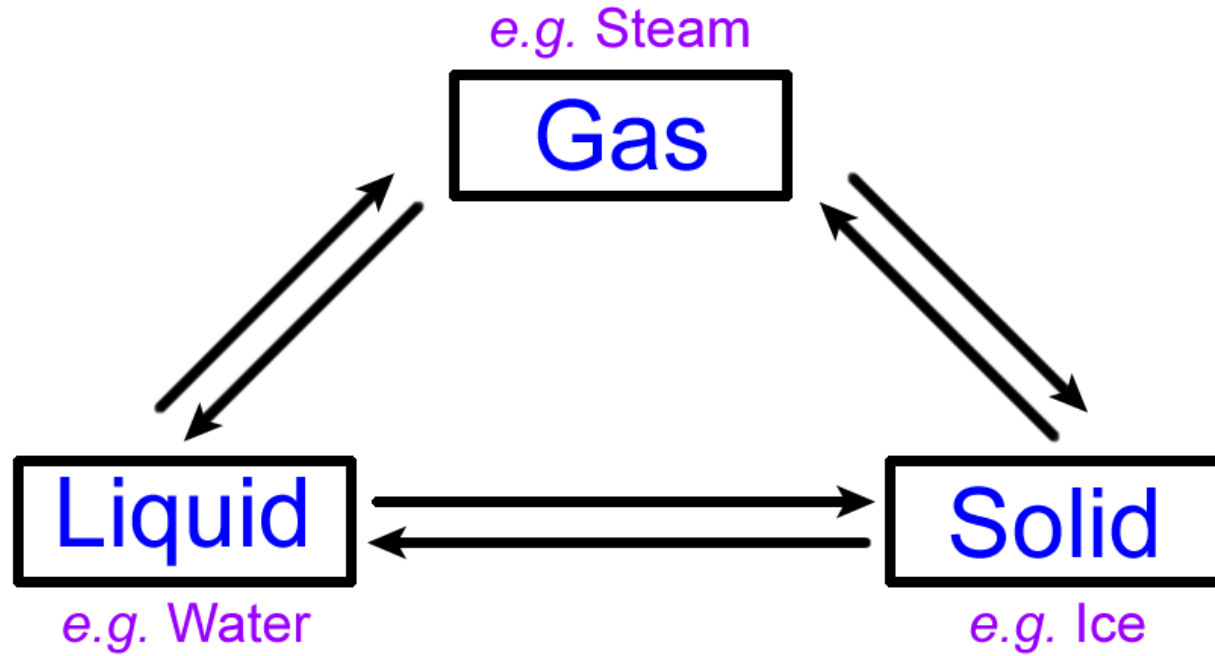


I remember from *kinetic particle theory* that changes in state also involve energy changes.

# Energy from Chemicals



# Energy from Chemicals

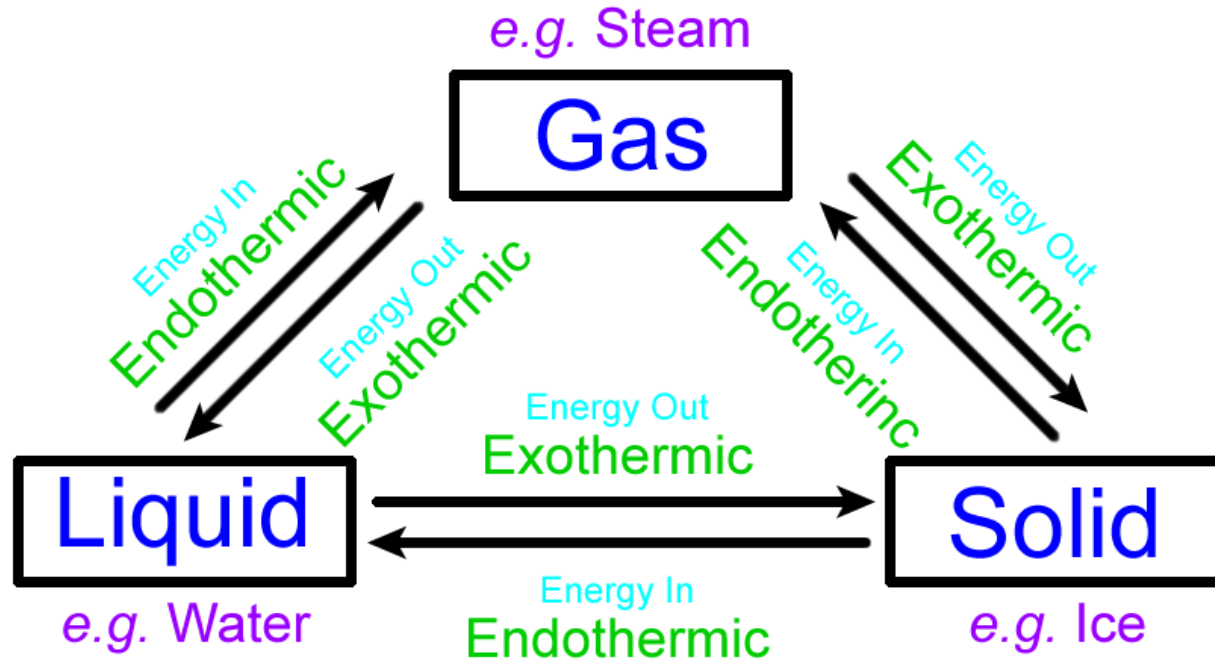


Remember:





# Energy from Chemicals



Remember:



# Energy from Chemicals

Phase changes involve a change in energy content of the system. Alcohol feels cold on the skin because the vaporisation of a liquid is an **exothermic / endothermic** process. As the liquid alcohol evaporates to form alcohol vapour, its energy content **increases / decreases** i.e. energy flows **into / out of** the system. Energy is **taken from / given to** the surroundings (the immediate surroundings is the skin), whose temperature therefore **increases / decreases**.

Steam at 100 °C is said to cause a worse burn than boiling water at the same temperature because the condensation of a vapour to a liquid is an **exothermic / endothermic** process and energy flows **into / out of** the system **from / into** the surroundings. Once again the immediate surroundings include the skin whose temperature is therefore **decreased / increased**.



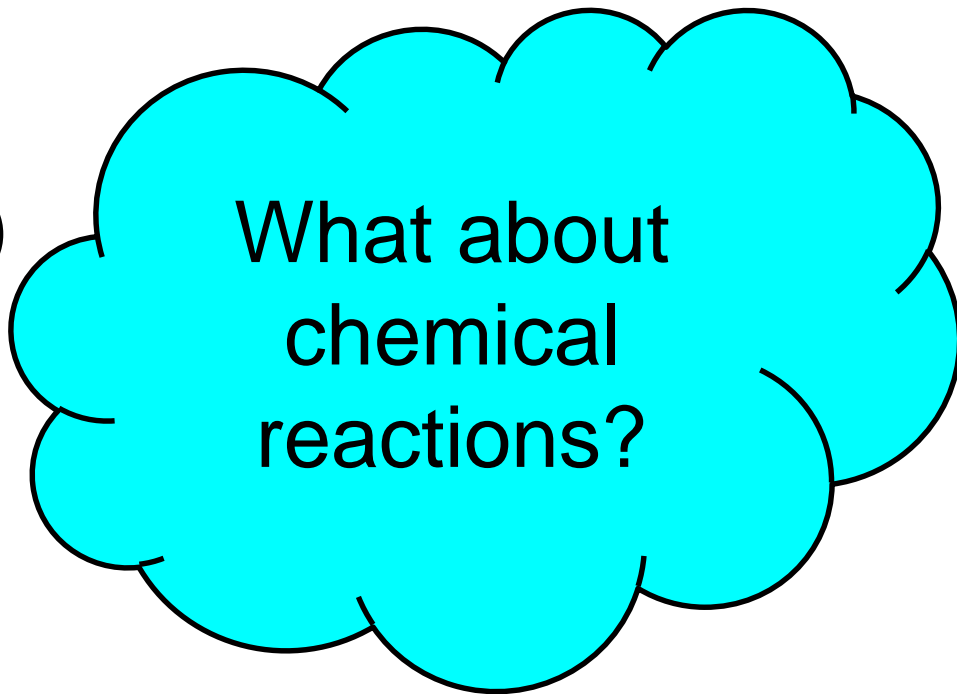
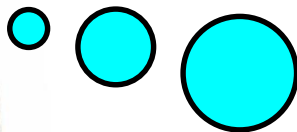
# Energy from Chemicals

Phase changes involve a change in energy content of the system. Alcohol feels cold on the skin because the vaporisation of a liquid is an ~~exothermic~~ / **endothermic** process. As the liquid alcohol evaporates to form alcohol vapour, its energy content **increases** / ~~decreases~~ i.e. energy flows **into** / ~~out of~~ the system. Energy is **taken from** / ~~given to~~ the surroundings (the immediate surroundings is the skin), whose temperature therefore ~~increases~~ / **decreases**.

Steam at 100 °C is said to cause a worse burn than boiling water at the same temperature because the condensation of a vapour to a liquid is an **exothermic** / ~~endothermic~~ process and energy flows ~~into~~ / **out** of the system ~~from~~ / **into** the surroundings. Once again the immediate surroundings include the skin whose temperature is therefore ~~decreased~~ / **increased**.



# Energy from Chemicals



What about  
chemical  
reactions?



# Exothermic Reactions



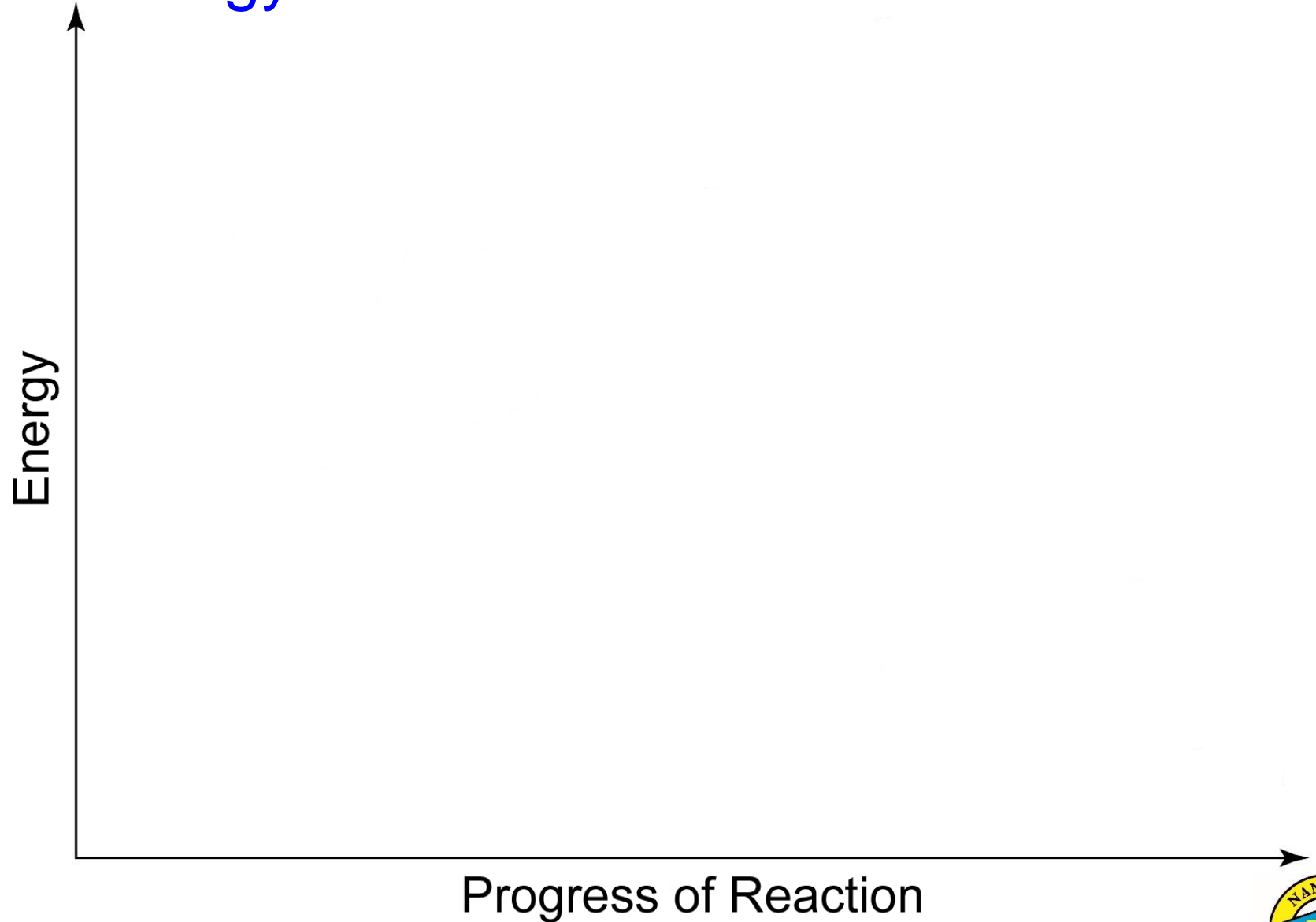
# Energy from Chemicals

## Important Things to Note...

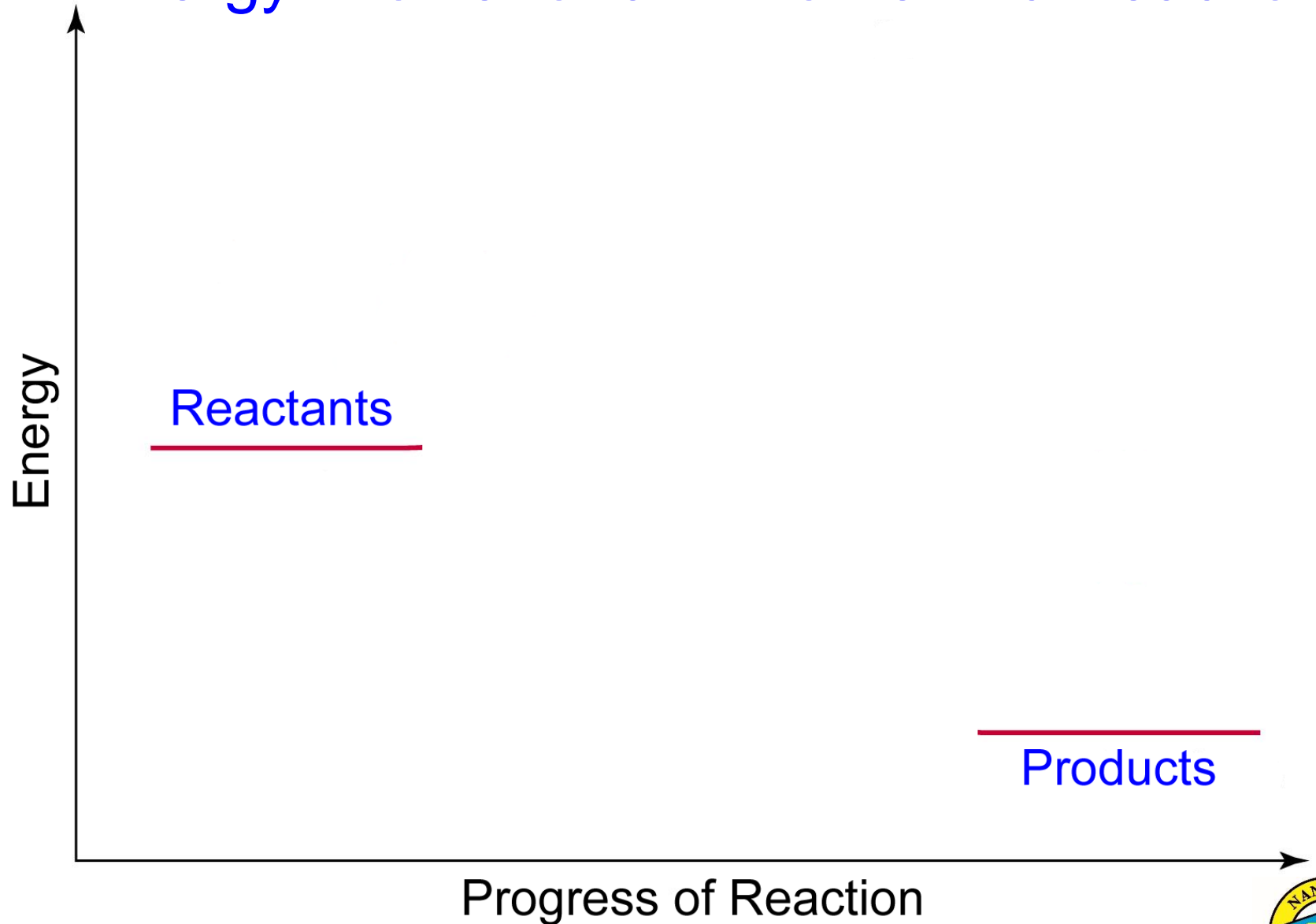
- $\Delta H$  means **change in enthalpy**.
- An **exothermic** change **releases** energy into the surroundings. Numerical values of  $\Delta H$  are **negative**.
- During an **exothermic** reaction, the temperature of the surroundings **increases**.



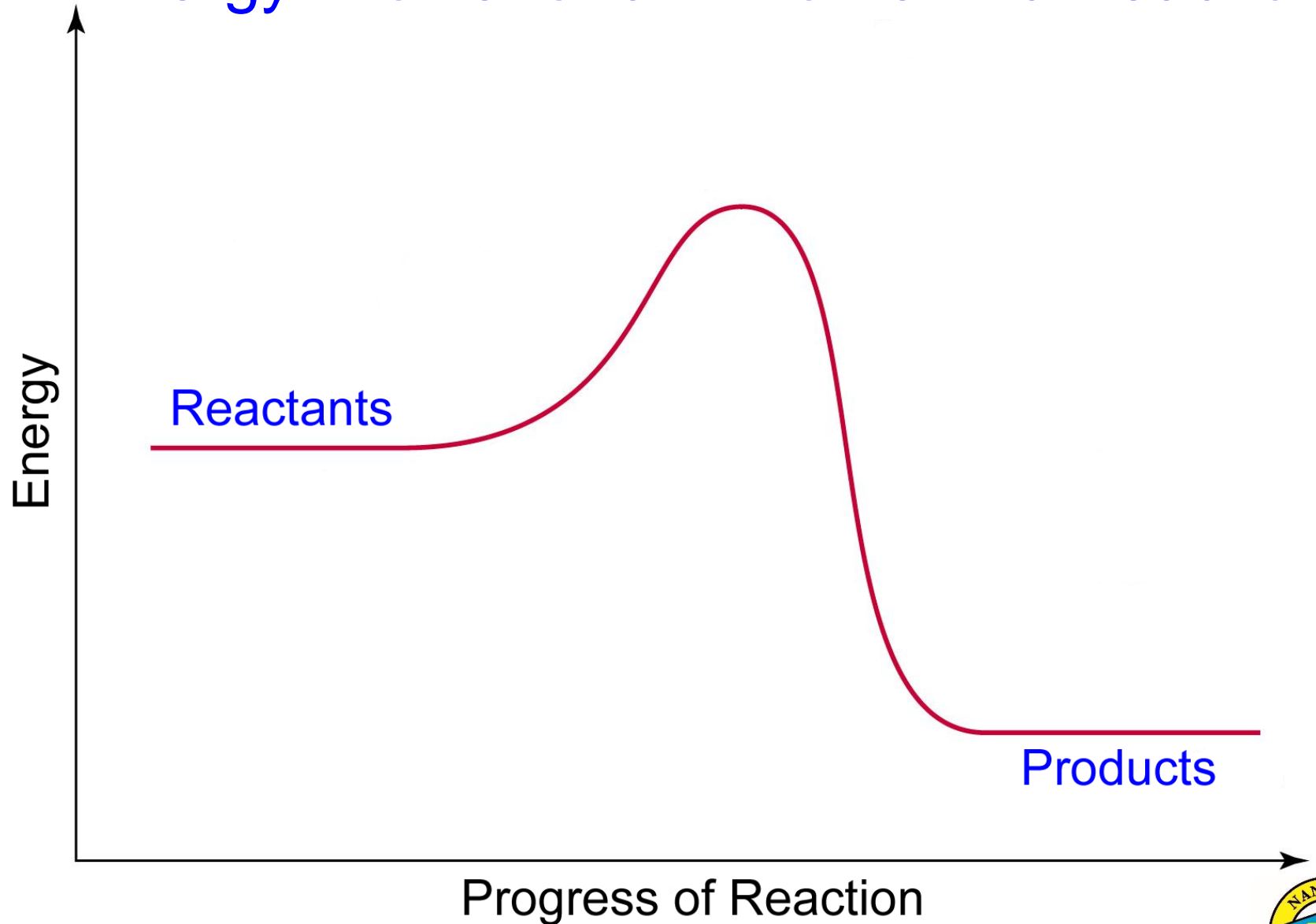
# Energy Profile for an Exothermic Reaction



# Energy Profile for an Exothermic Reaction

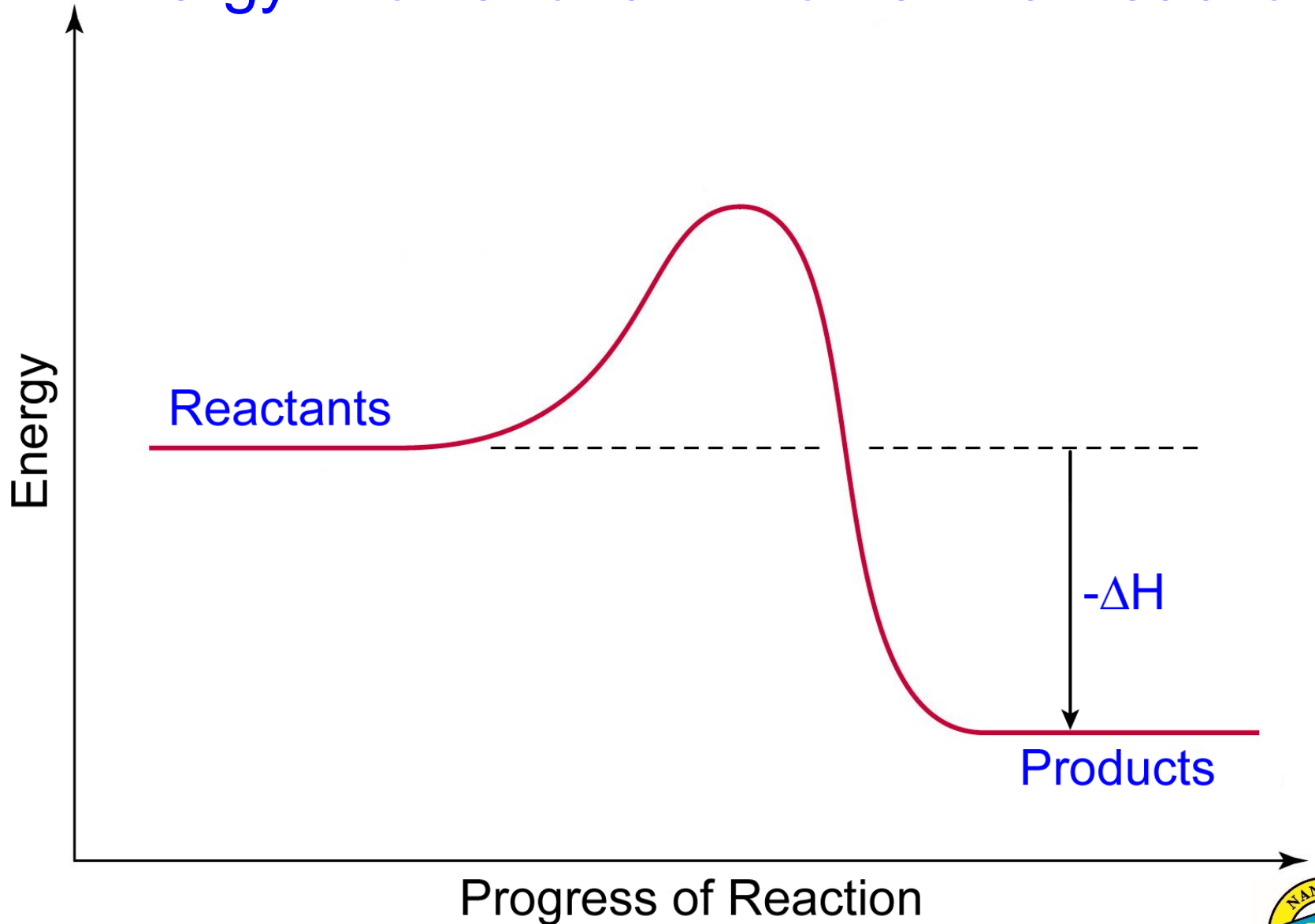


# Energy Profile for an Exothermic Reaction

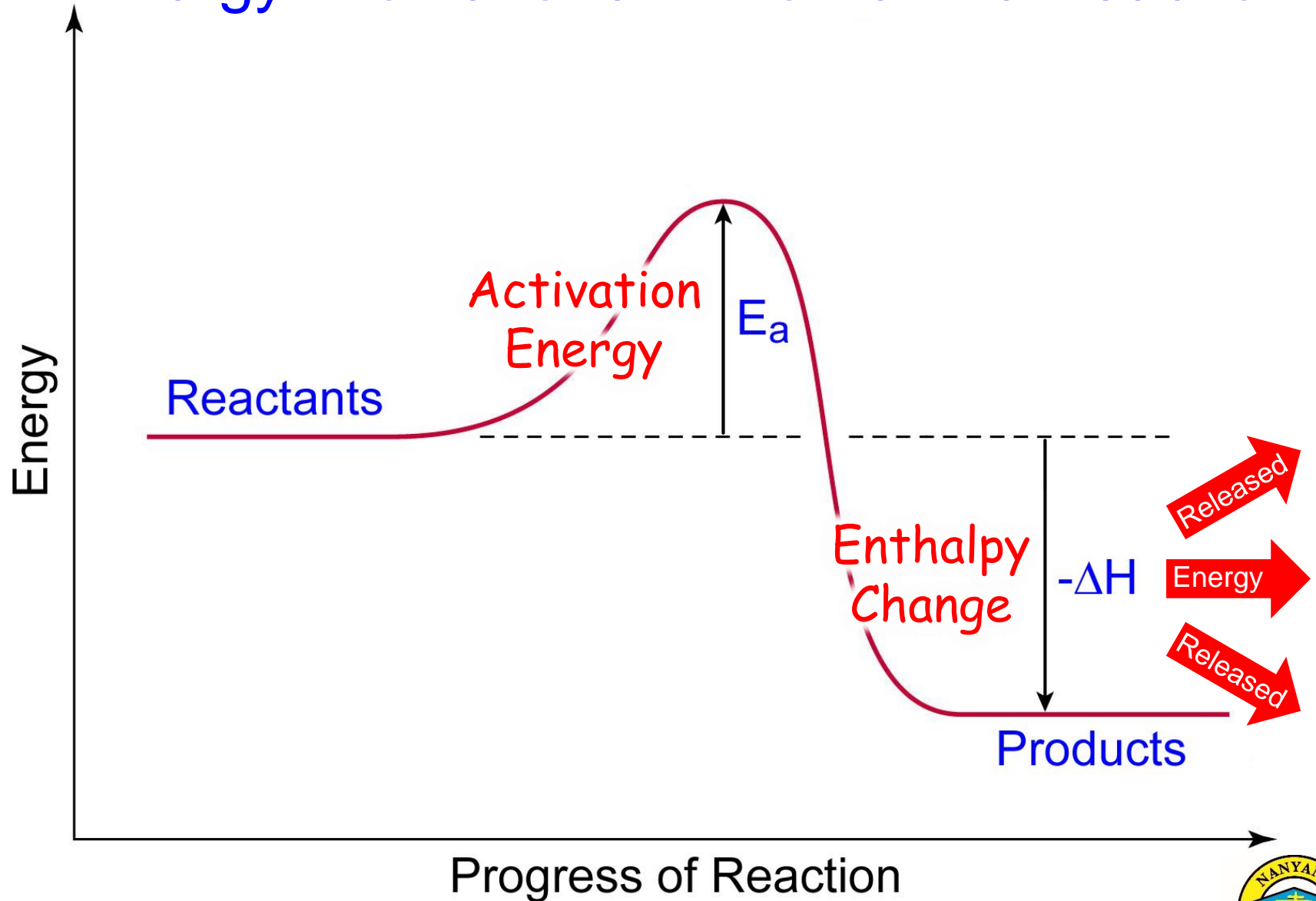




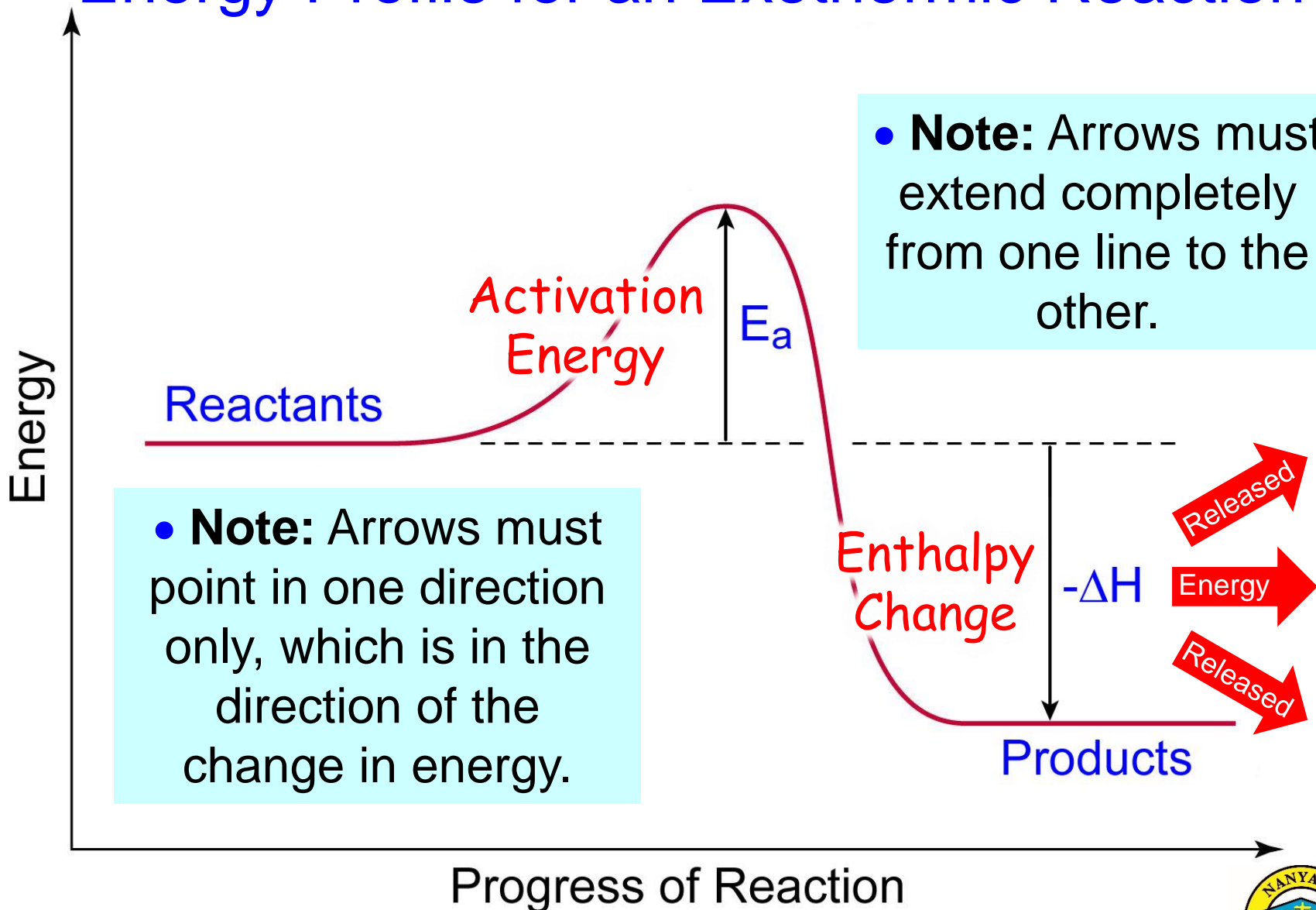
# Energy Profile for an Exothermic Reaction



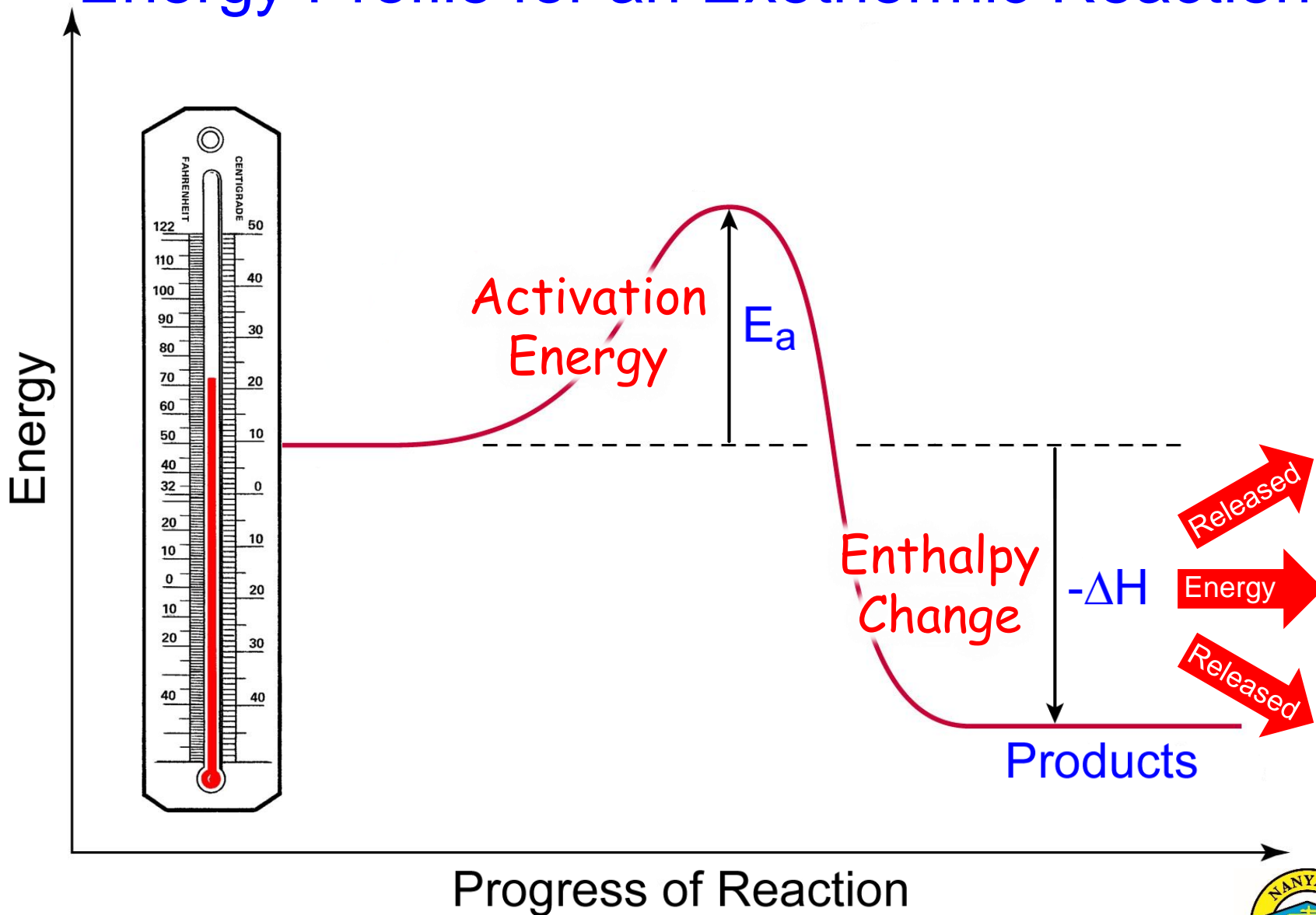
# Energy Profile for an Exothermic Reaction



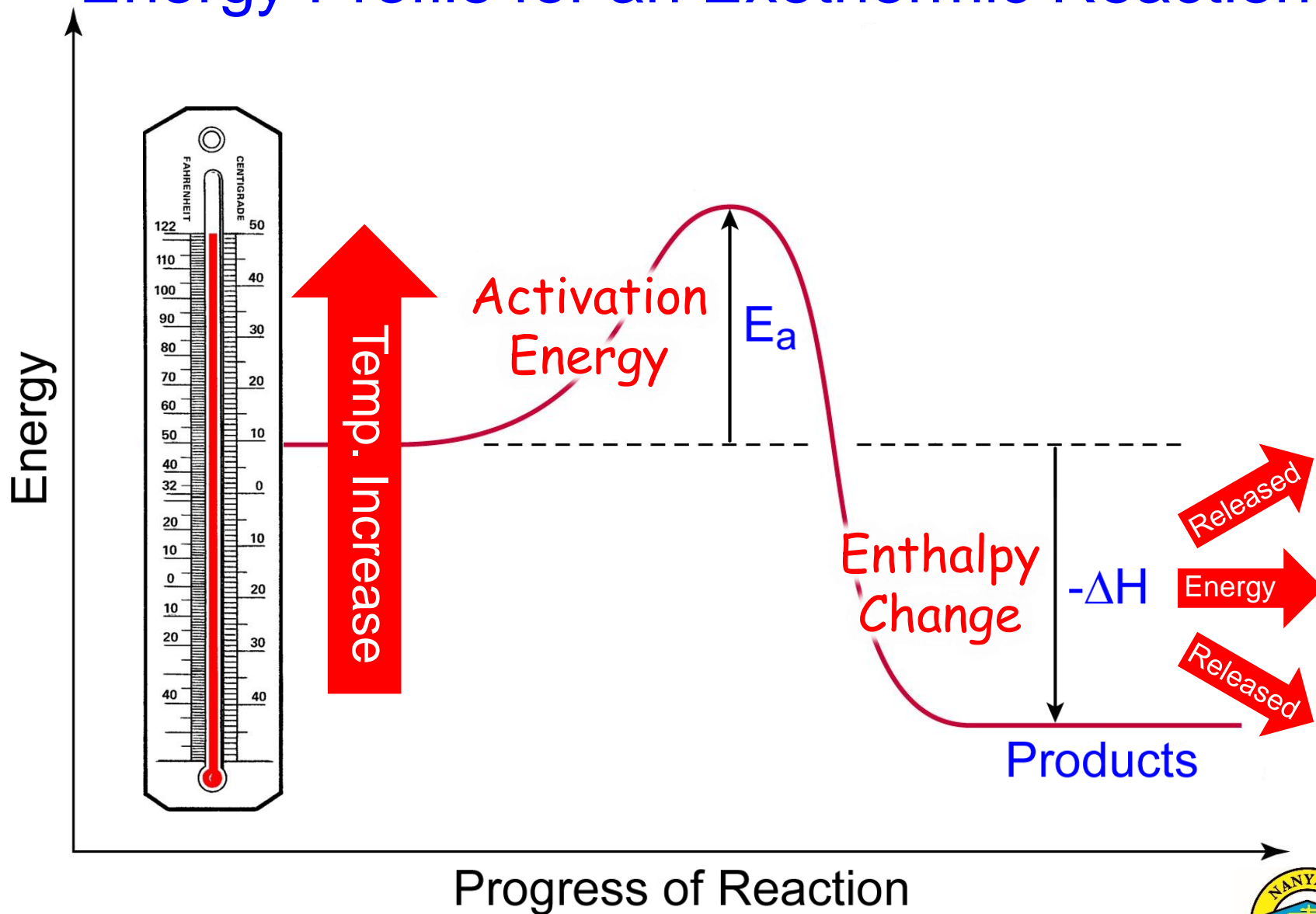
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# Energy Profile for an Exothermic Reaction

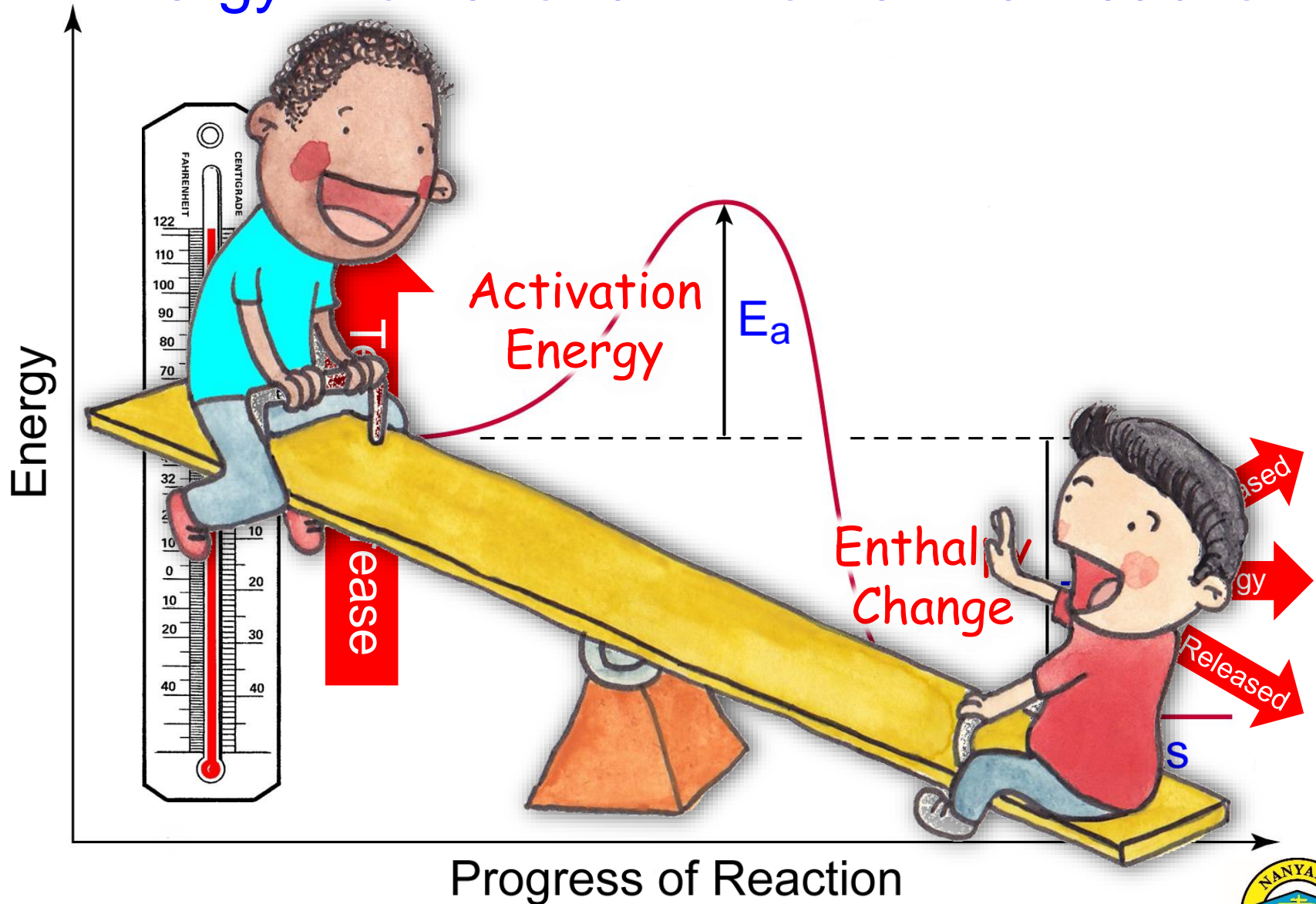


# Energy Profile for an Exothermic Reaction





# Energy Profile for an Exothermic Reaction





# Energy from Chemicals

- It is very important to remember that the thermometer is part of the *surroundings* and *not* part of the *reaction*!
- As a consequence, the thermometer measures the *temperature* (or *energy*) of the *surroundings* and not the *temperature* (or *energy*) of the *reaction*!

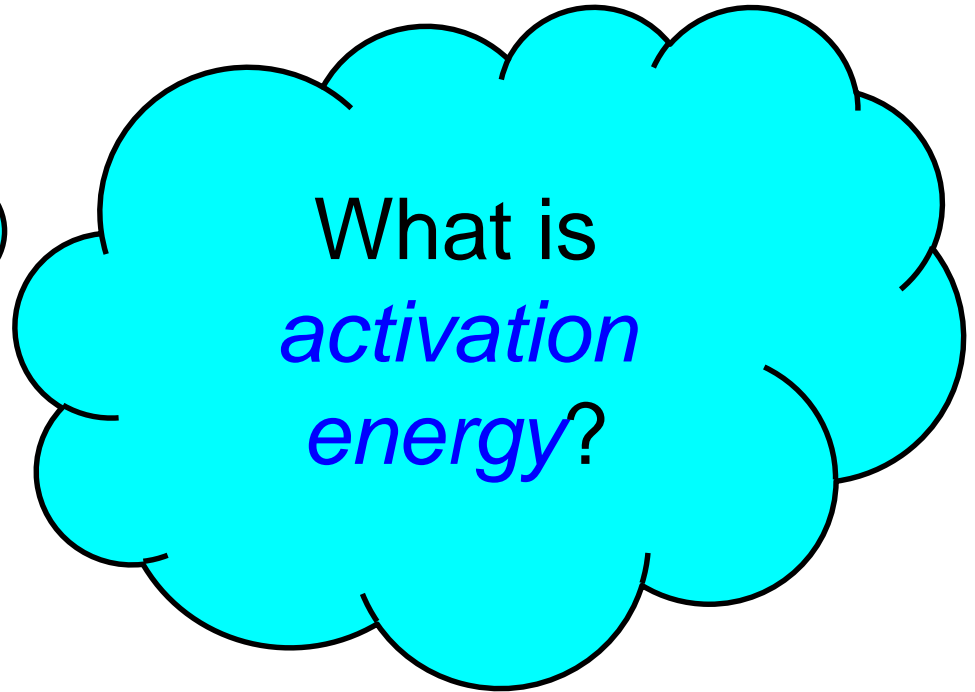
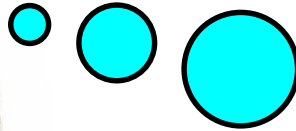


# Energy from Chemicals

- The particles in a reaction do not all react at once.
- People running a marathon run at different speeds and cross the finishing line at different times.
- It is the same for the particles in a reaction. The particles move at different speeds, with different amounts of kinetic energy, and will collide and react at different times.



# Energy from Chemicals



What is  
*activation*  
*energy?*

# Activation Energy



- Activation energy is the minimum amount of energy that must be supplied to a chemical in order for a reaction to take place.



# Activation Energy



- For example, the phosphorus and sulfur in a match head will not react with the oxygen in the air until the match is struck against the sandpaper on the side of the match box.

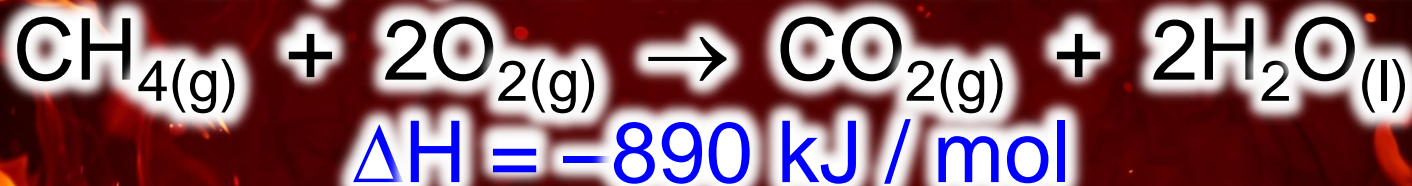
# Activation Energy



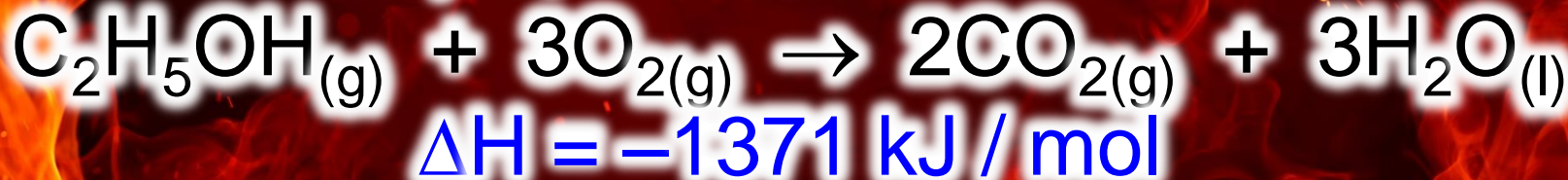
- Friction between the match head and sandpaper generates heat energy, which is used to break chemical bonds in the molecules of phosphorus, sulfur and oxygen, thus allowing them to react.

# Exothermic Reactions

- All combustion reactions (where a fuel reacts with oxygen) are exothermic.
- For example, the combustion of methane:



- For example, the combustion of ethanol:





# Endothermic Reaction



# Energy from Chemicals

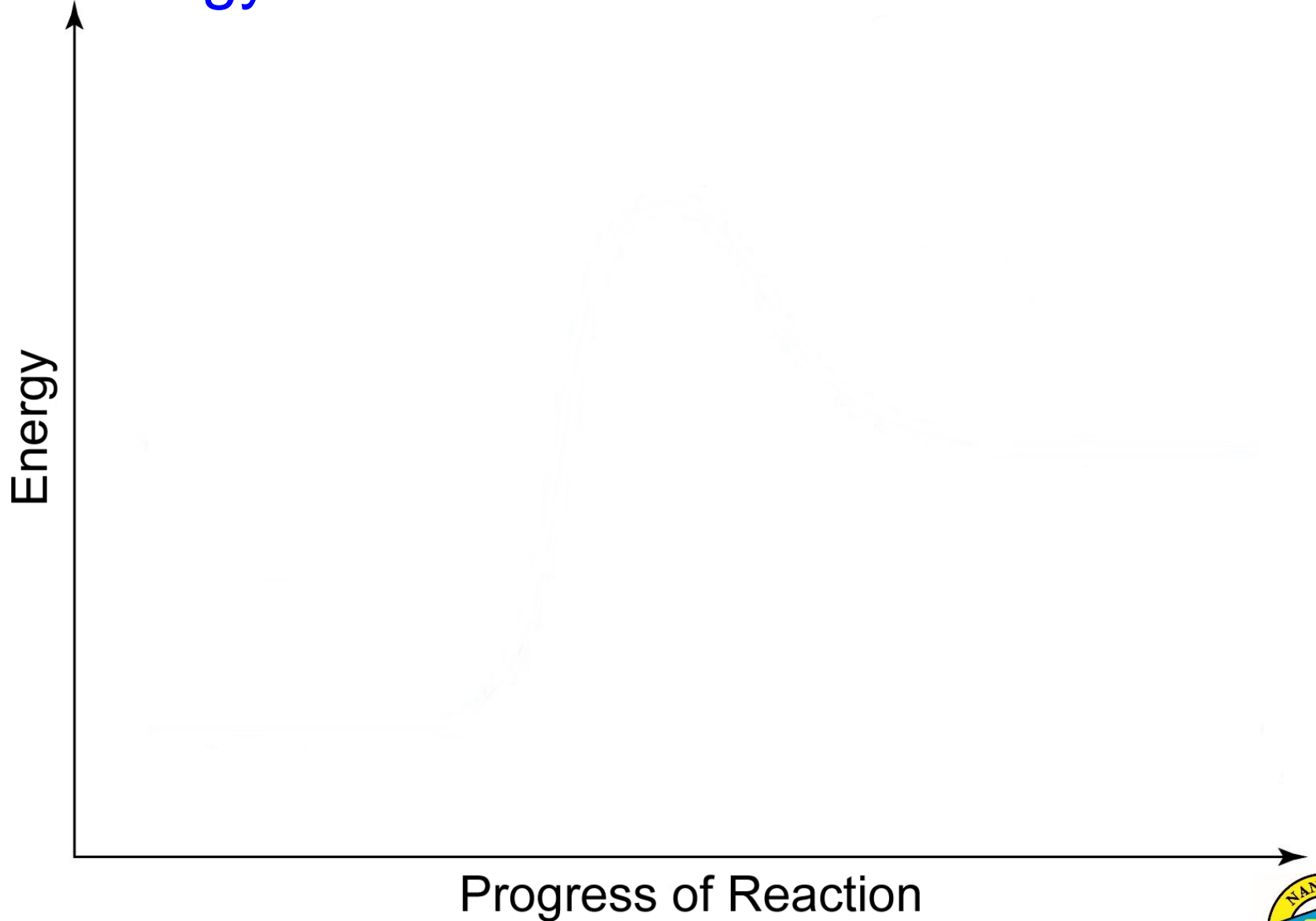
## Important Things to Note...

- $\Delta H$  means **change in enthalpy**.
- An **endothermic** change **absorbs** energy from the surroundings. Numerical values of  $\Delta H$  are **positive**.
- During an **endothermic** reaction, the temperature of the surroundings **decreases**.

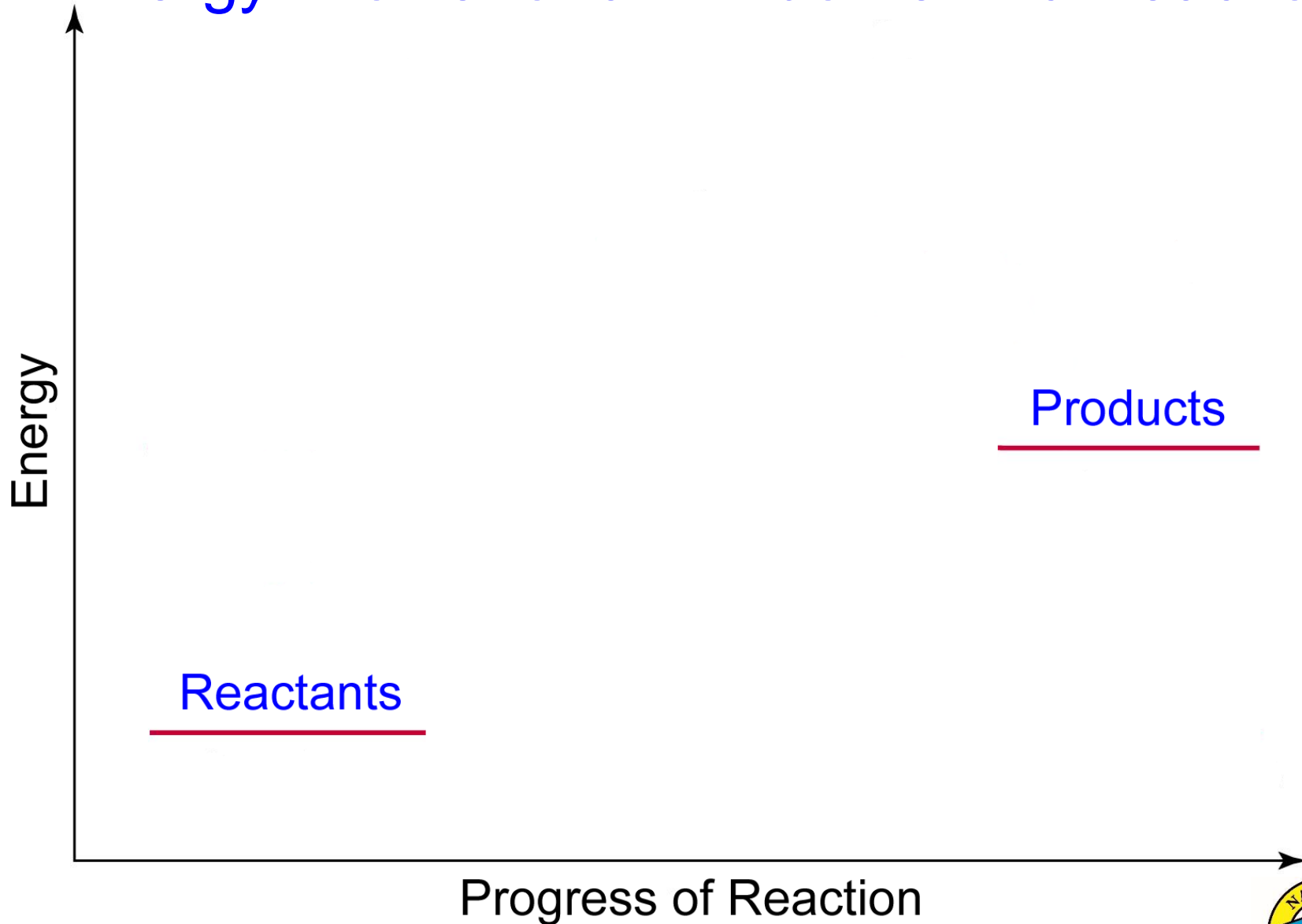




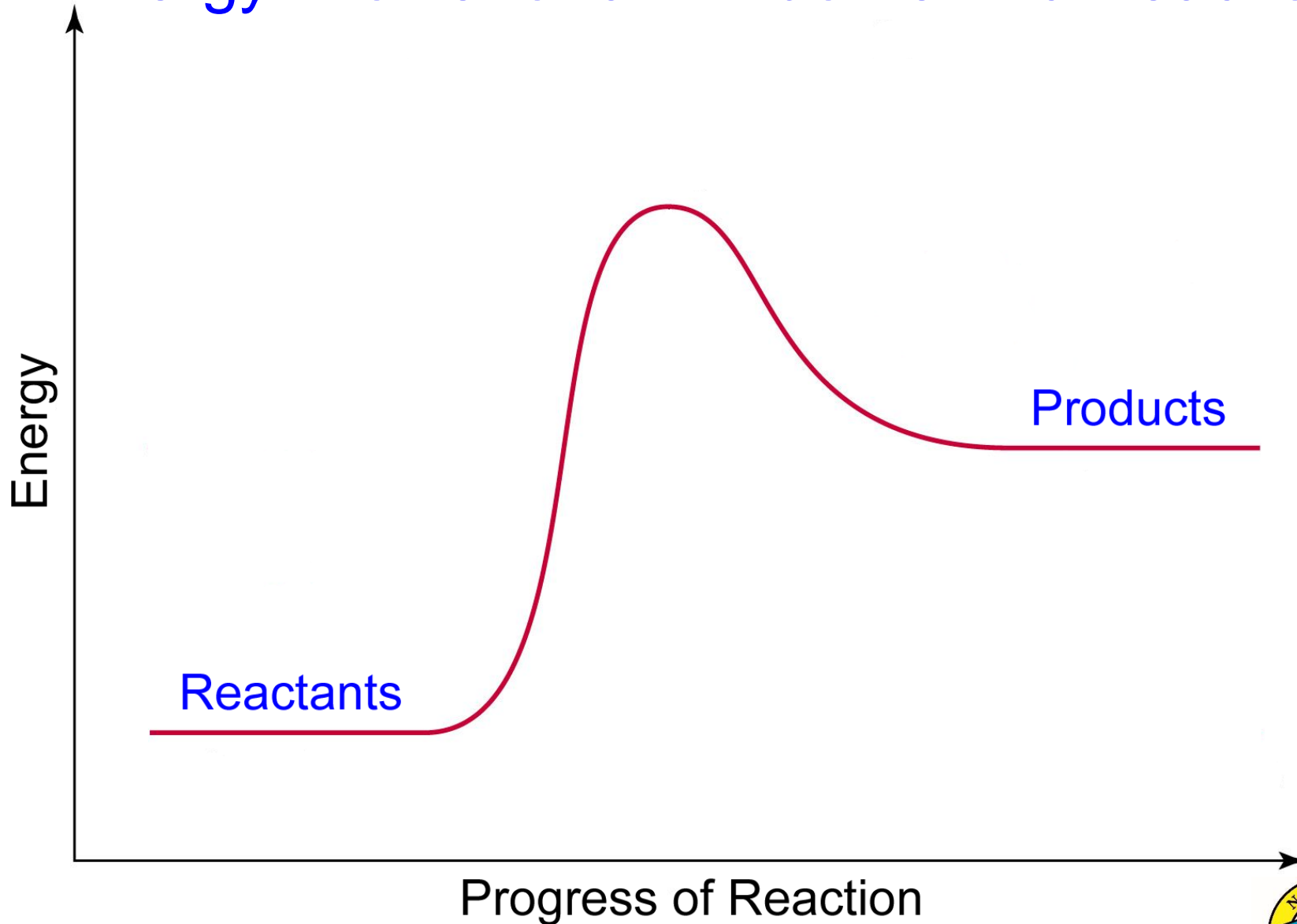
# Energy Profile for an Endothermic Reaction



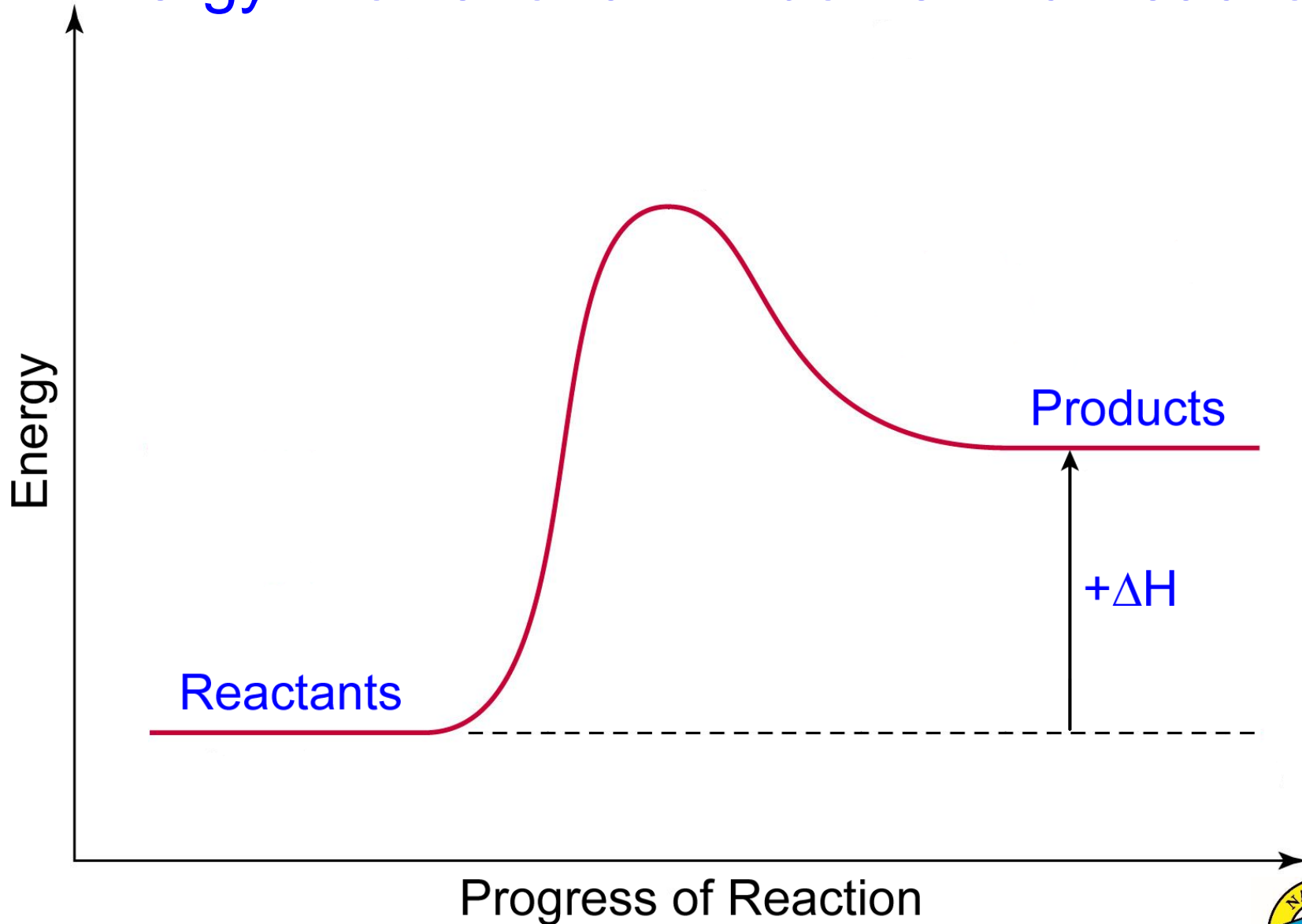
# Energy Profile for an Endothermic Reaction



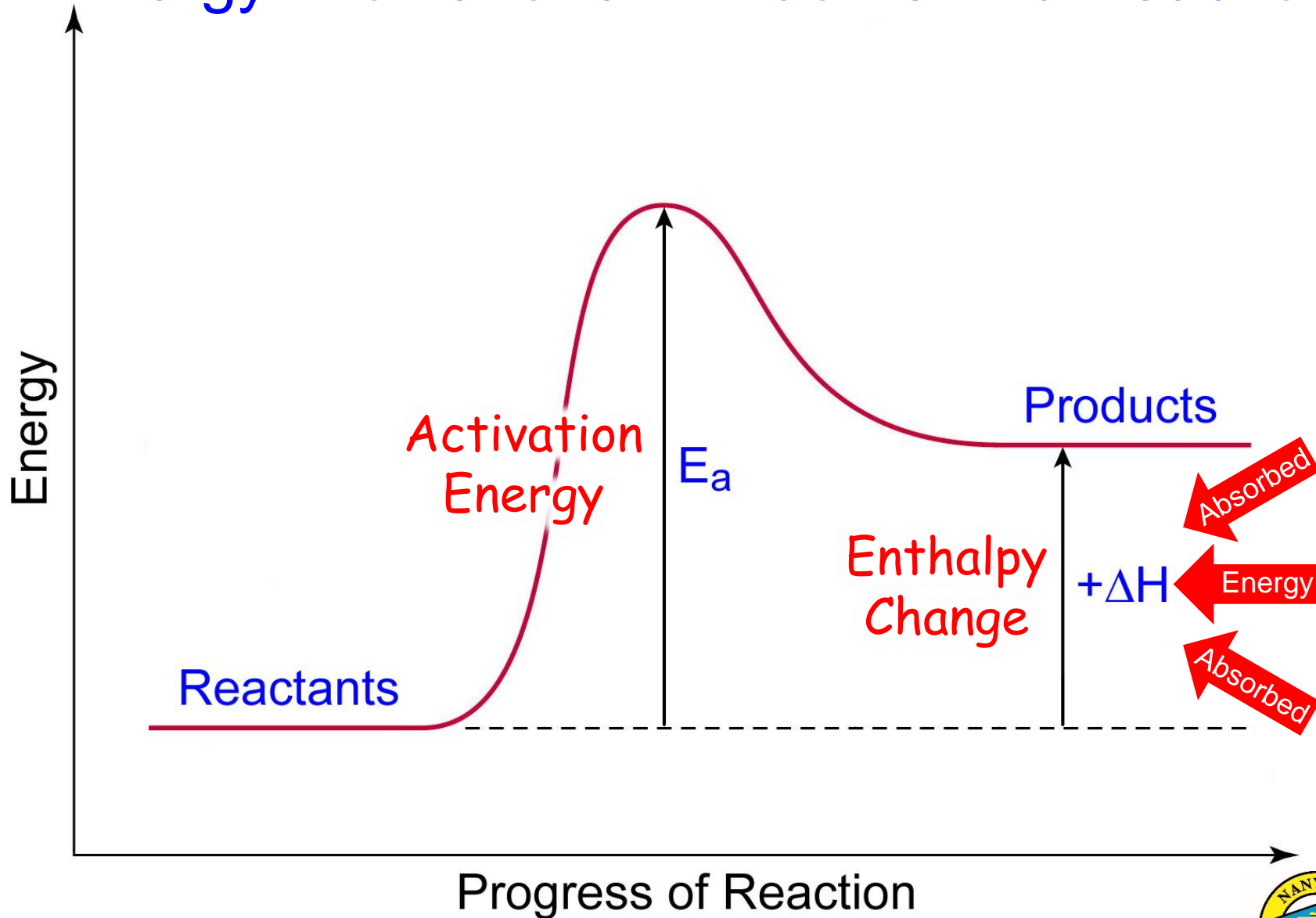
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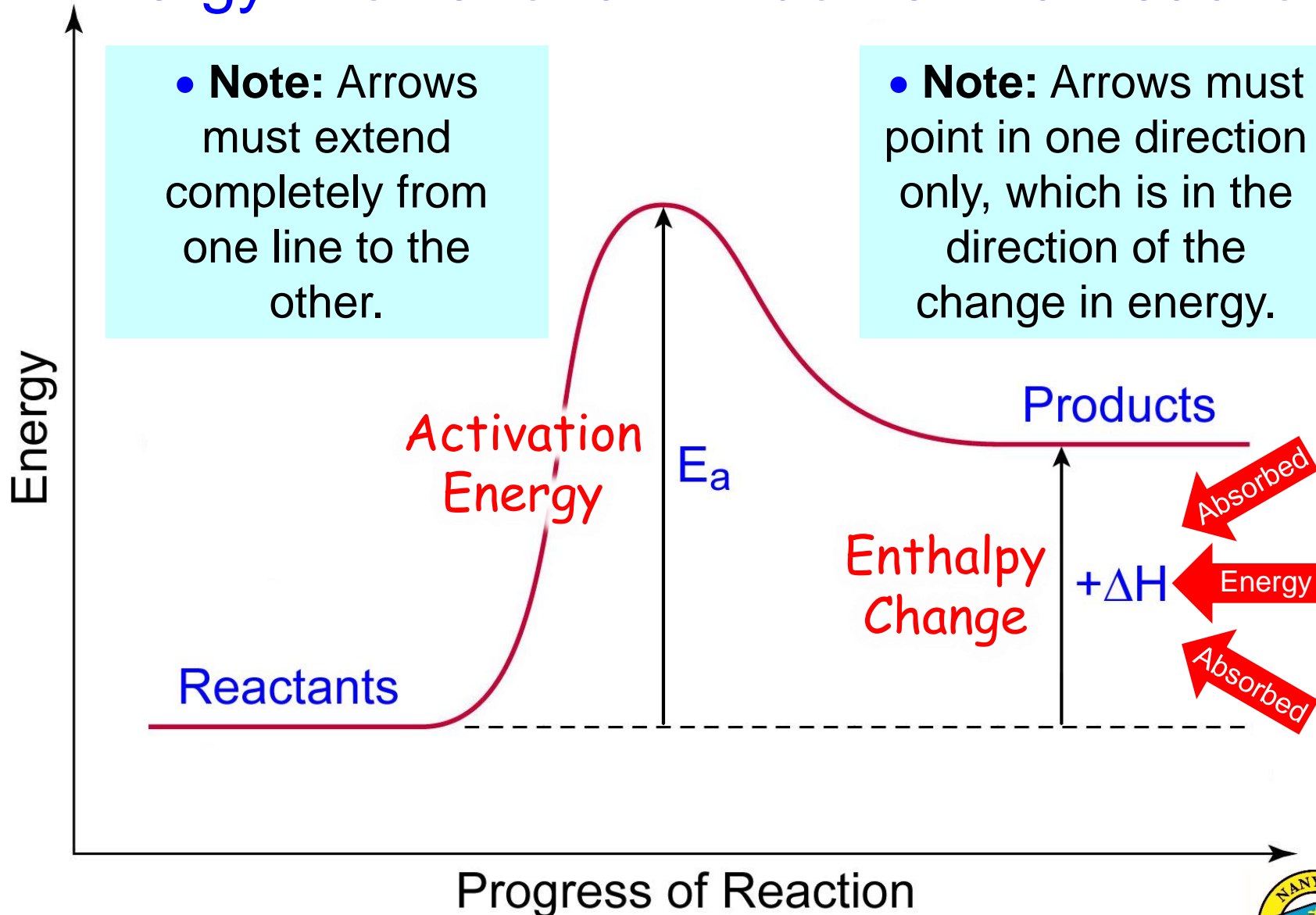
# Energy Profile for an Endothermic Reaction



# Energy Profile for an Endothermic Reaction

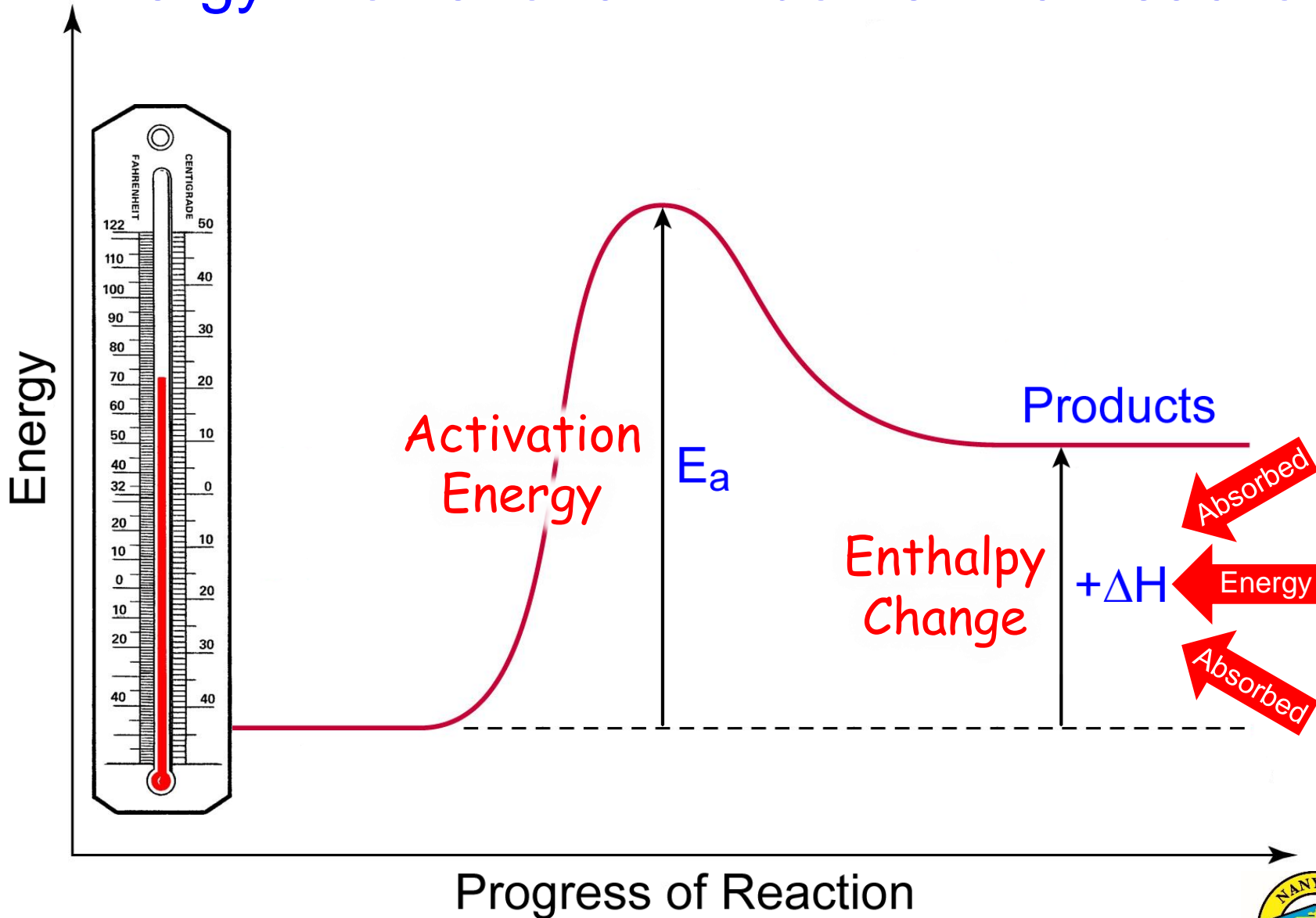


# Energy Profile for an Endothermic Reaction

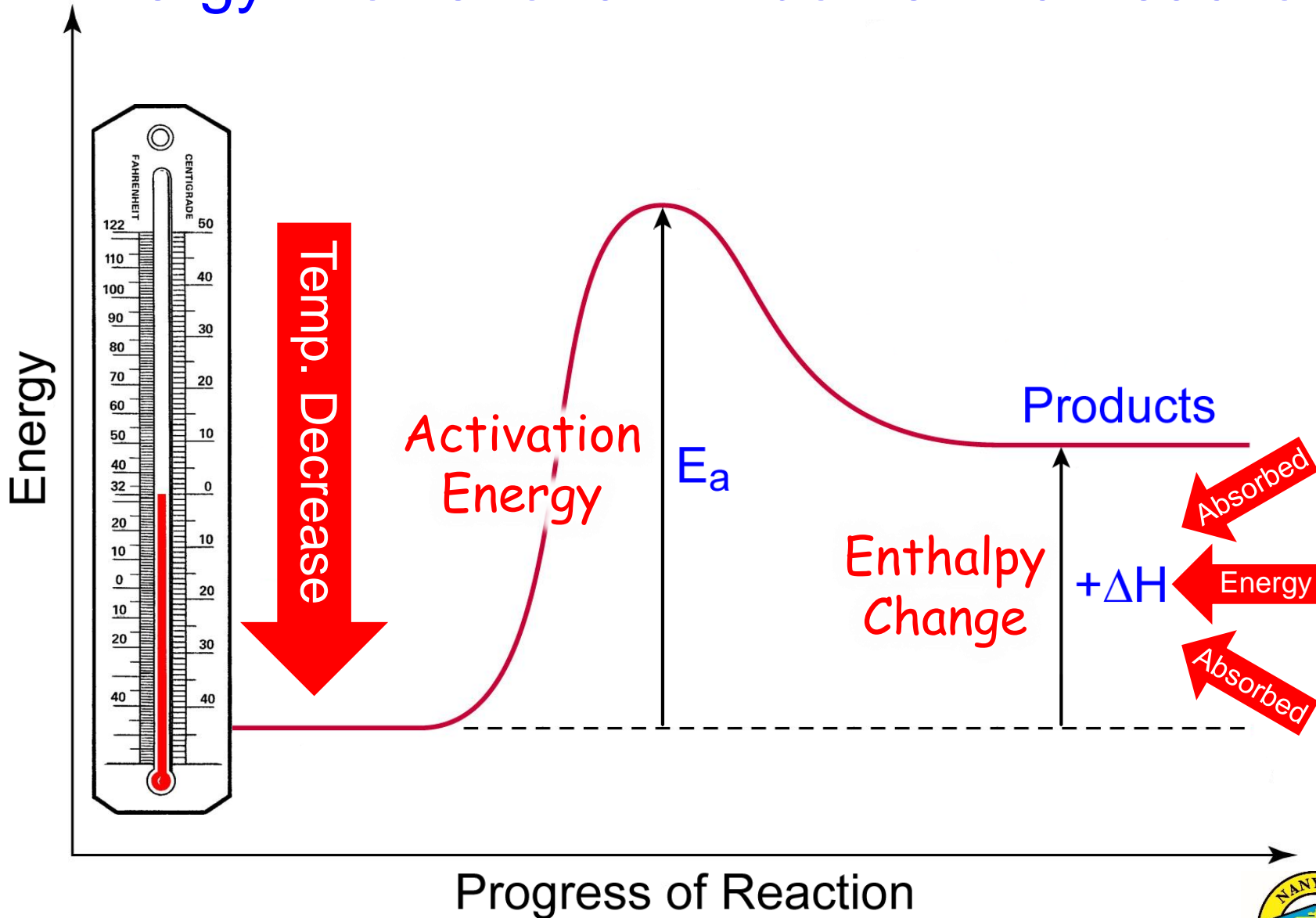




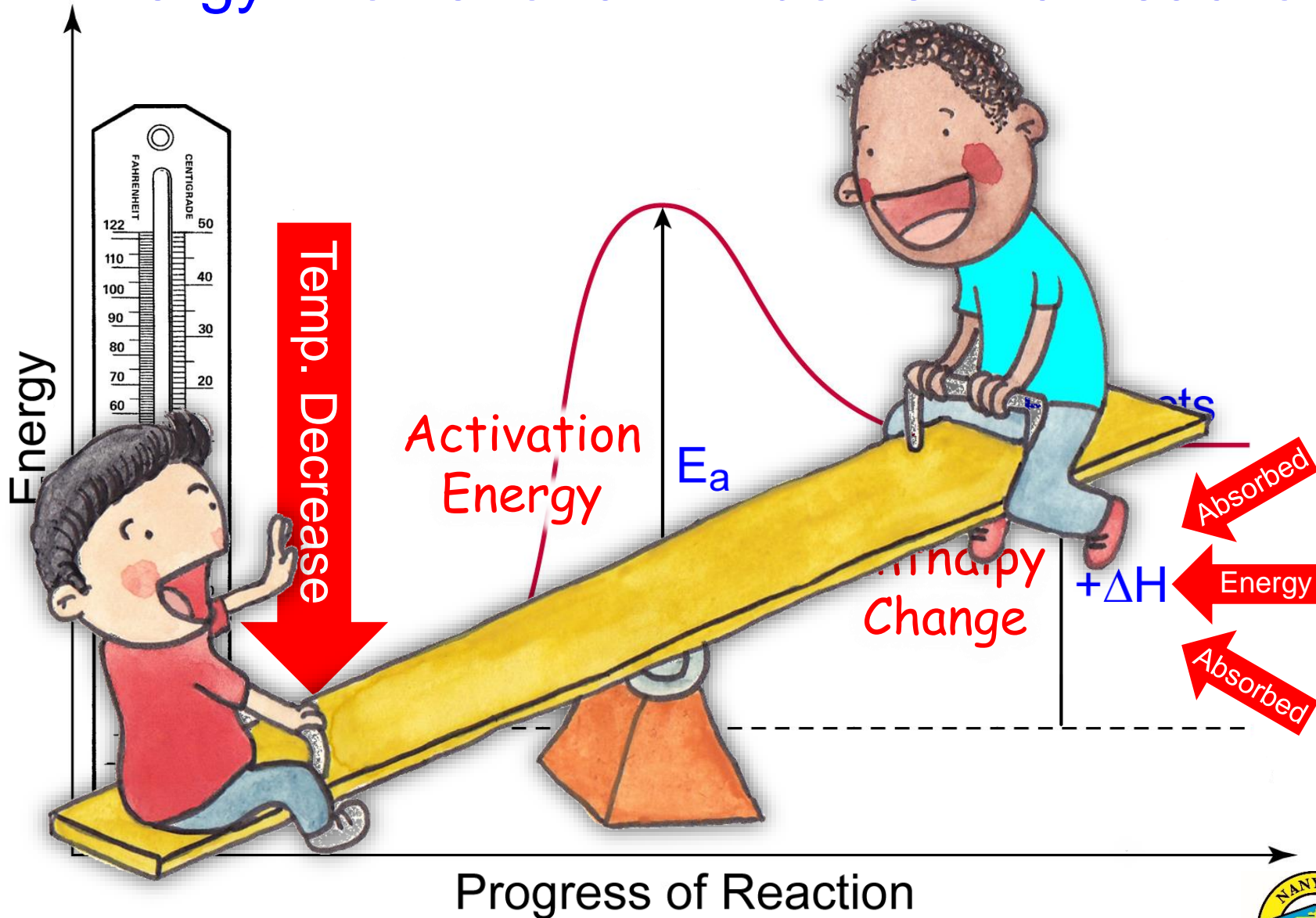
# Energy Profile for an Endothermic Reaction



# Energy Profile for an Endothermic Reaction



# Energy Profile for an Endothermic Reaction



# Energy from Chemicals

Chemical reactions are accompanied by energy changes. When the energy content of a chemical system decreases during a reaction, the energy content of the products is **greater / less** than that of the reactants. The reaction is said to be **exothermic / endothermic** and a **rise / fall** in temperature is observed. The value of the enthalpy change of the reaction ( $\Delta H$ ) is **positive / negative**.

When the temperature falls during a reaction, the reaction is said to be **exothermic / endothermic**. The energy content of the products is **greater / less** than that of the reactants *i.e.* the energy content of the system has **increased / decreased** during the reaction.  $\Delta H$  for the reaction is **positive / negative**.



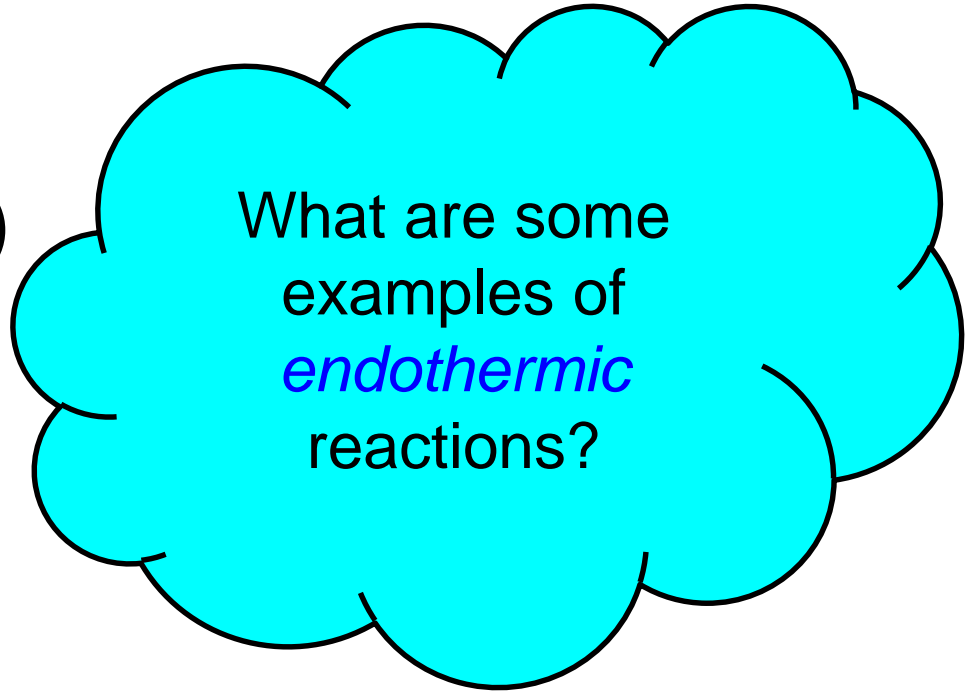
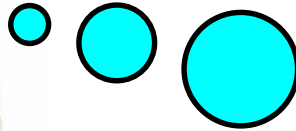
# Energy from Chemicals

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# Energy from Chemicals



What are some  
examples of  
*endothermic*  
reactions?



# Endothermic Reactions

- Photosynthesis:

carbon dioxide + water  $\rightarrow$  glucose + oxygen



- Thermal Decomposition:

calcium carbonate  $\rightarrow$  calcium oxide + carbon dioxide



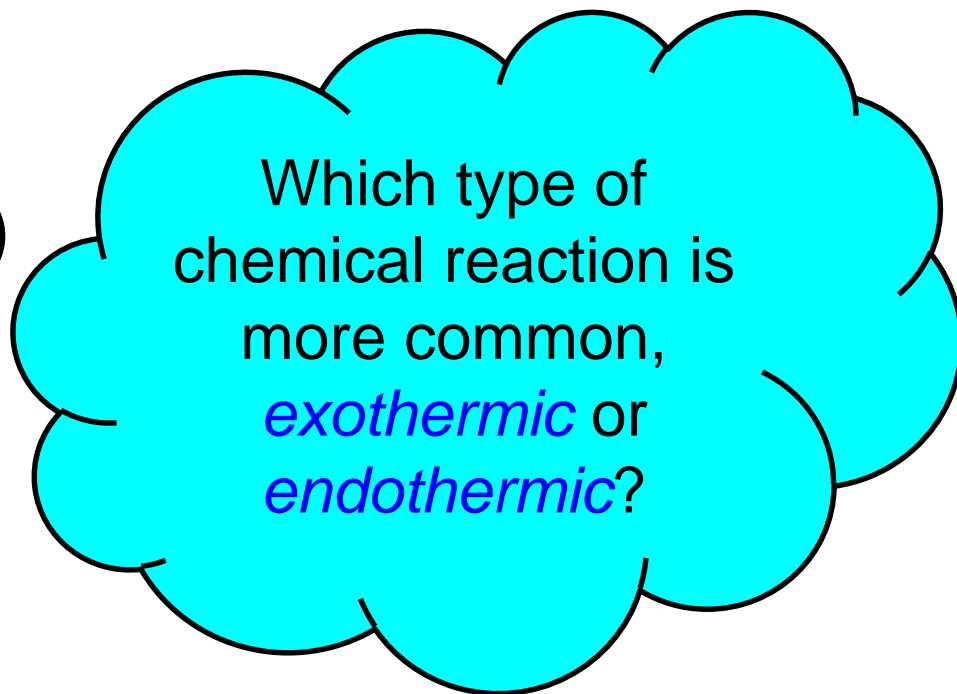
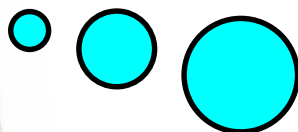
- Cracking Alkanes:

decane  $\rightarrow$  octane + ethene





# Energy from Chemicals



Which type of  
chemical reaction is  
more common,  
*exothermic* or  
*endothermic*?

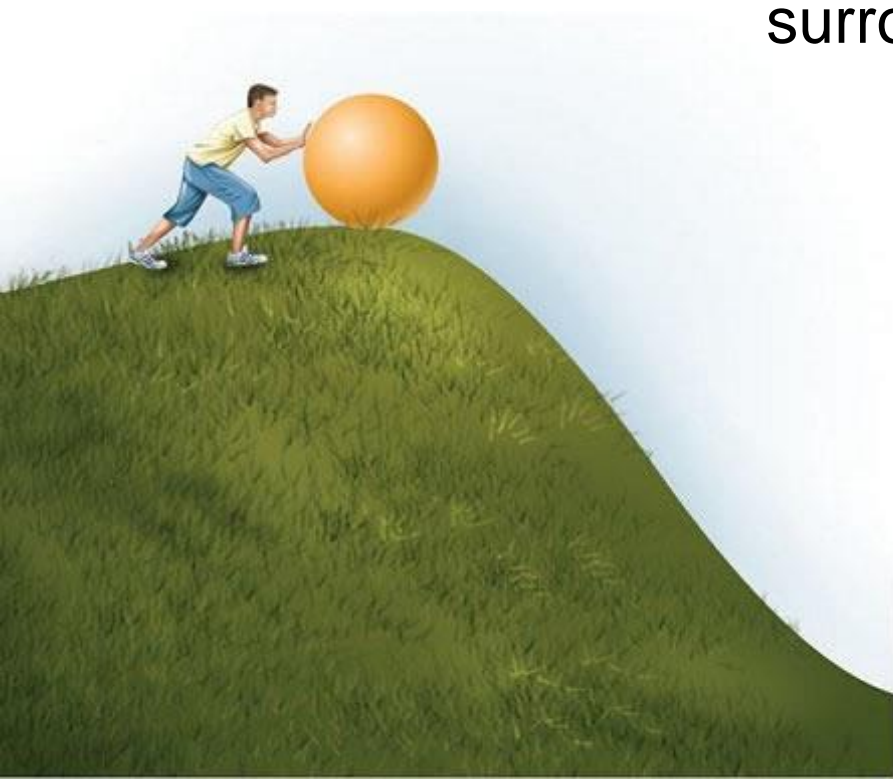
# Exothermic and Endothermic Reactions

- Chemical reactions take place in order for chemicals to **reduce their energy** and become **more stable**.
  - It is natural for chemicals to go from a state of high energy to a state of low energy, just as it is natural for a ball to roll down a hill.



# Exothermic and Endothermic Reactions

- In an **exothermic** reaction, high energy reactants form low energy products, with energy given off to the surroundings.
- In an **endothermic** reaction, low energy reactants form high energy products, with energy absorbed from the surroundings.



# Exothermic and Endothermic Reactions

- In terms of energy changes, **exothermic** reactions are more **favorable** than **endothermic** reactions.
- **Exothermic** reactions are therefore more **common** than **endothermic** reactions.



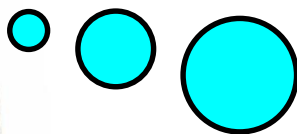


# Exothermic and Endothermic Reactions

- **Question:** How do these diagrams illustrate the concept of **activation energy**?
- **Answer:** The person must use a **specific amount of energy** to roll the ball from the left-hand-side to the top of the hill. Only then is the ball free to roll down the opposite side.



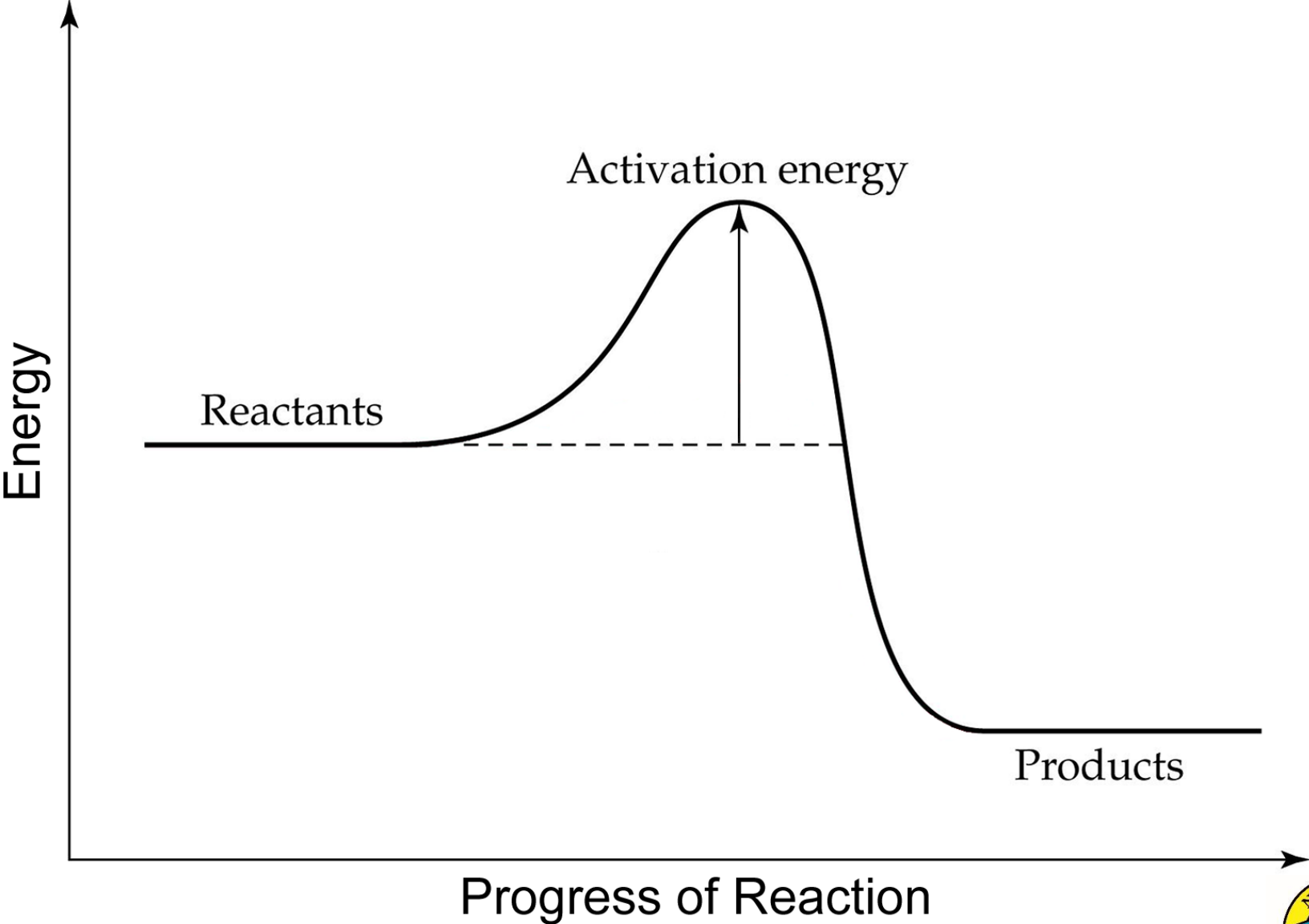
# Energy from Chemicals



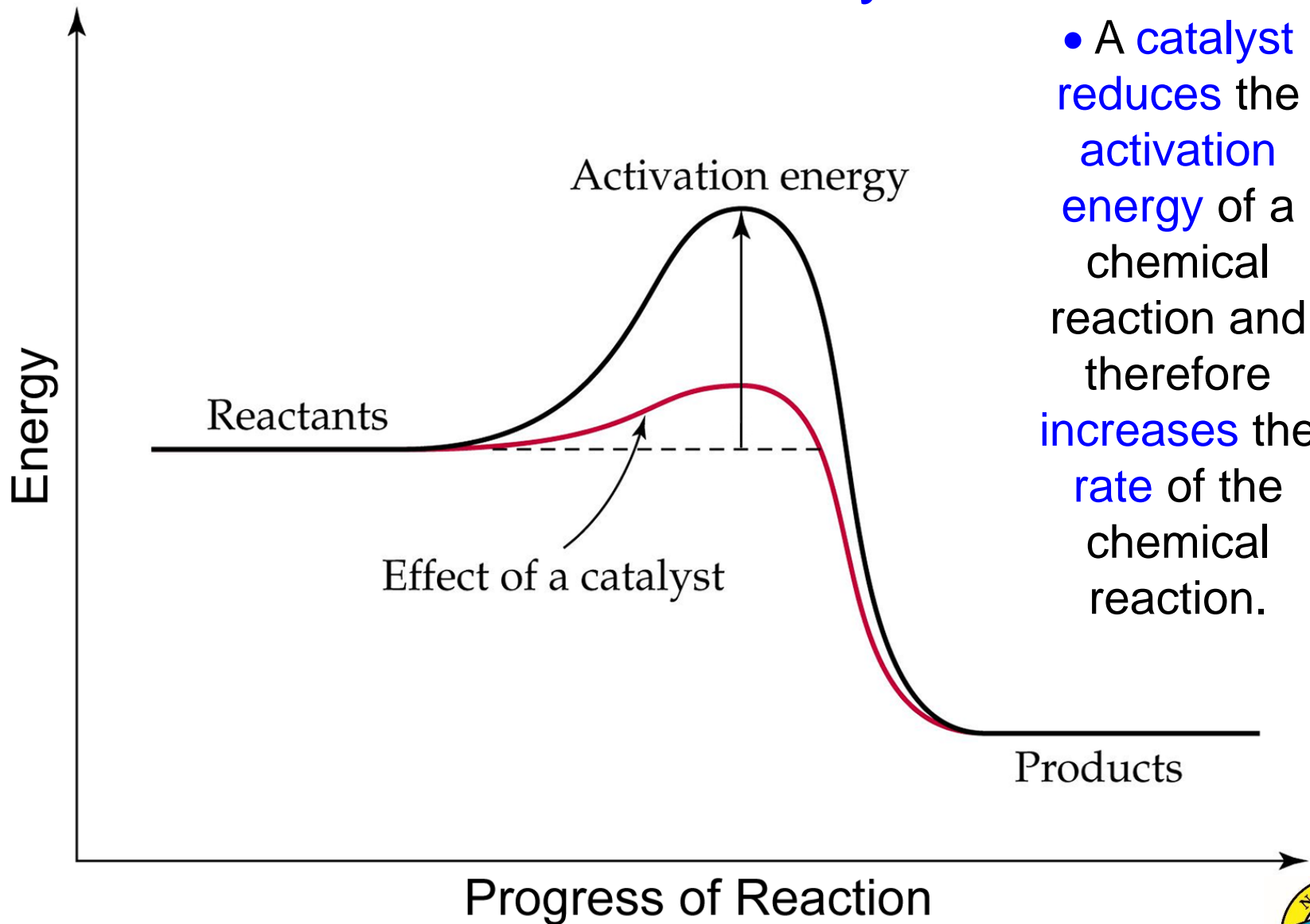
What effect does a *catalyst* have on the *activation energy* of a chemical reaction?



# Effect of a Catalyst

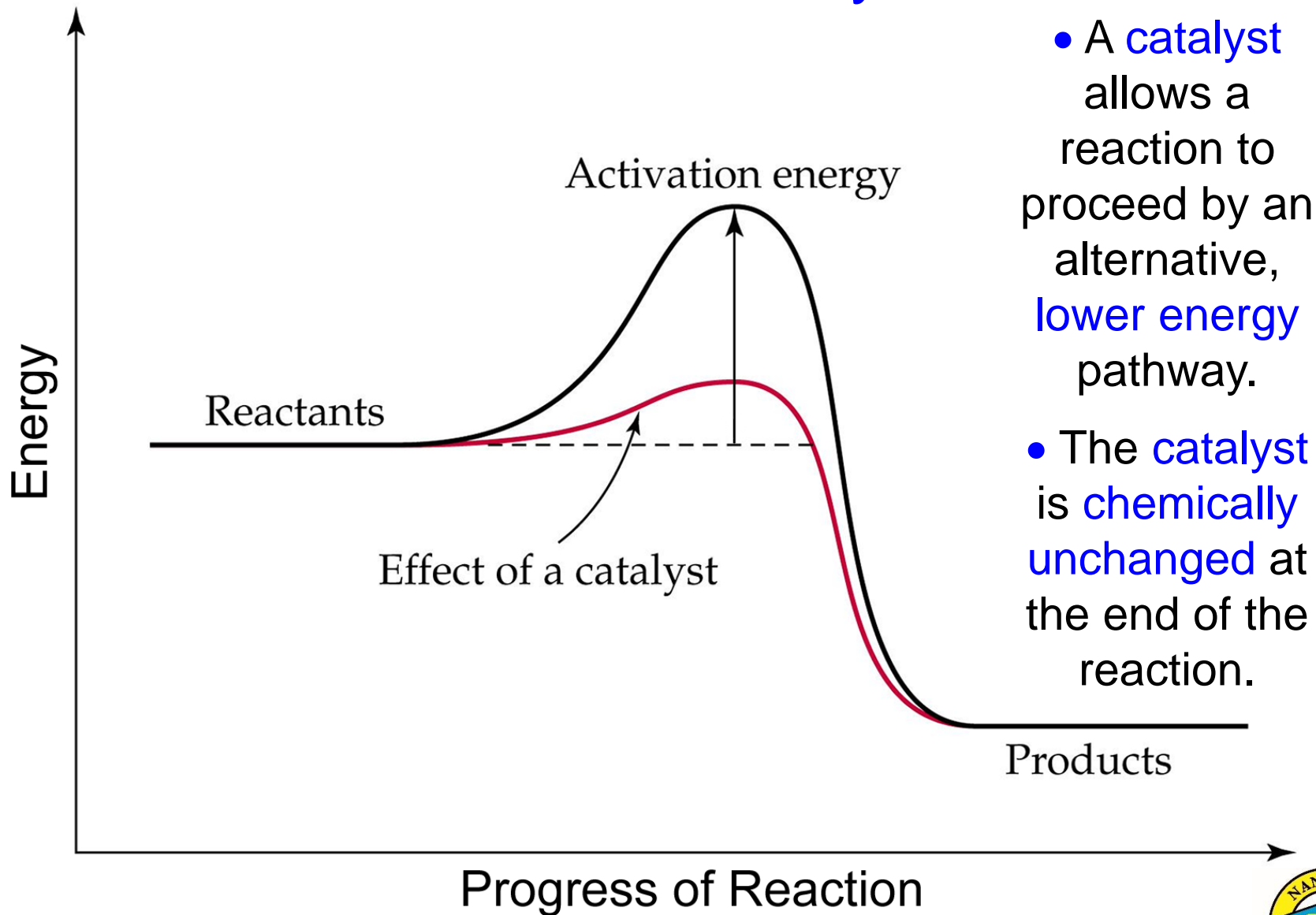


# Effect of a Catalyst



- A catalyst reduces the activation energy of a chemical reaction and therefore increases the rate of the chemical reaction.

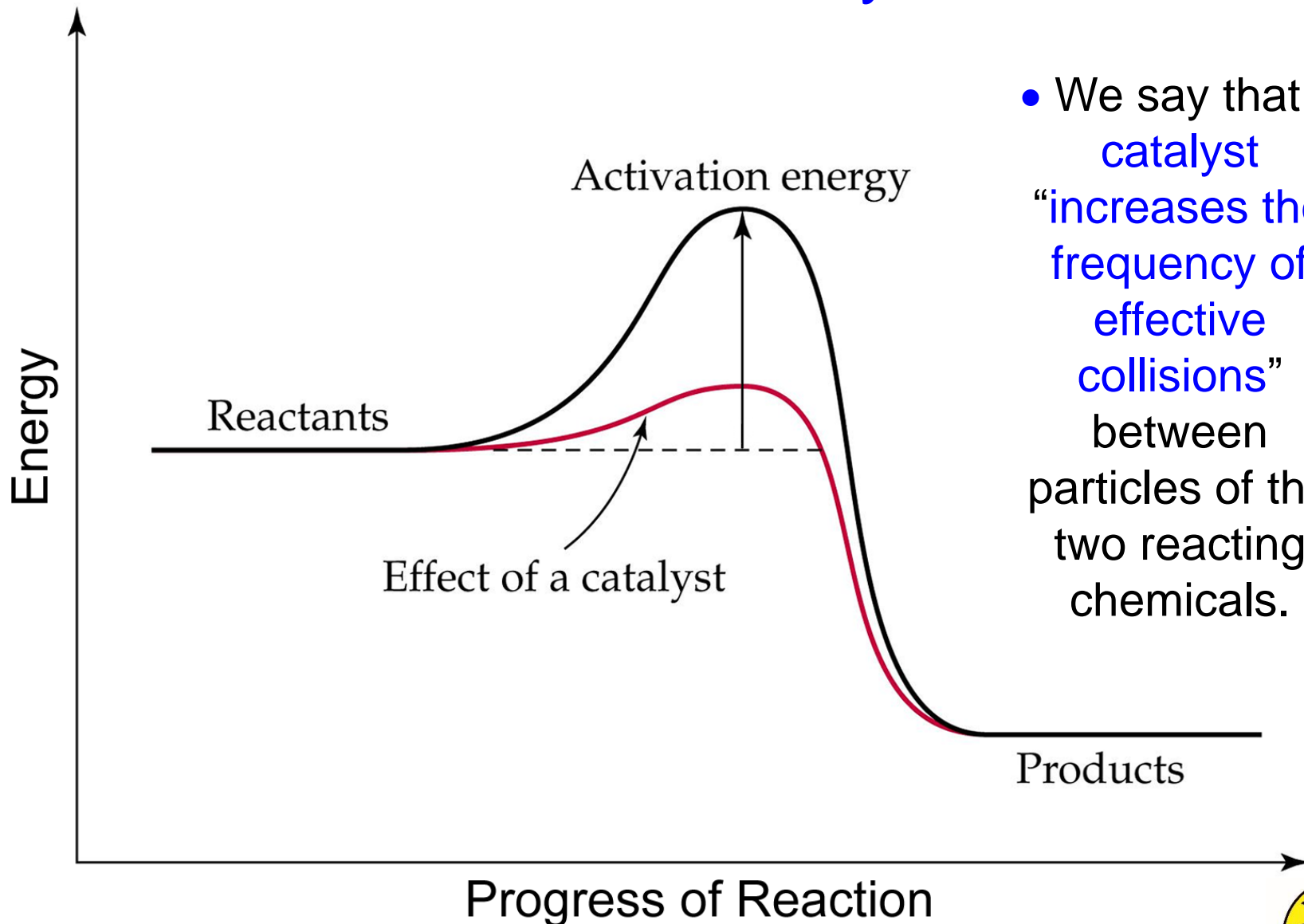
# Effect of a Catalyst



- A catalyst allows a reaction to proceed by an alternative, lower energy pathway.
- The catalyst is chemically unchanged at the end of the reaction.

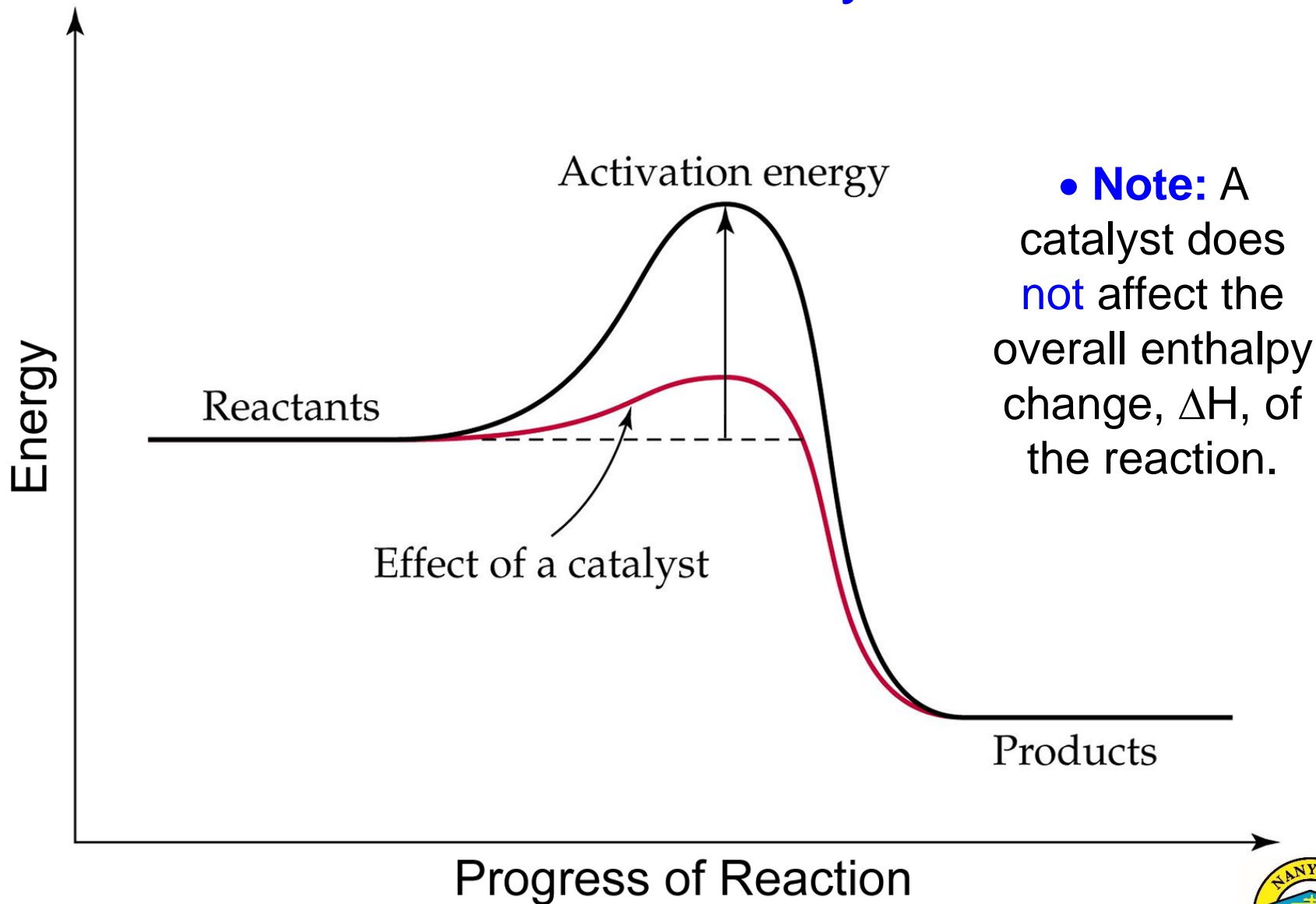


# Effect of a Catalyst

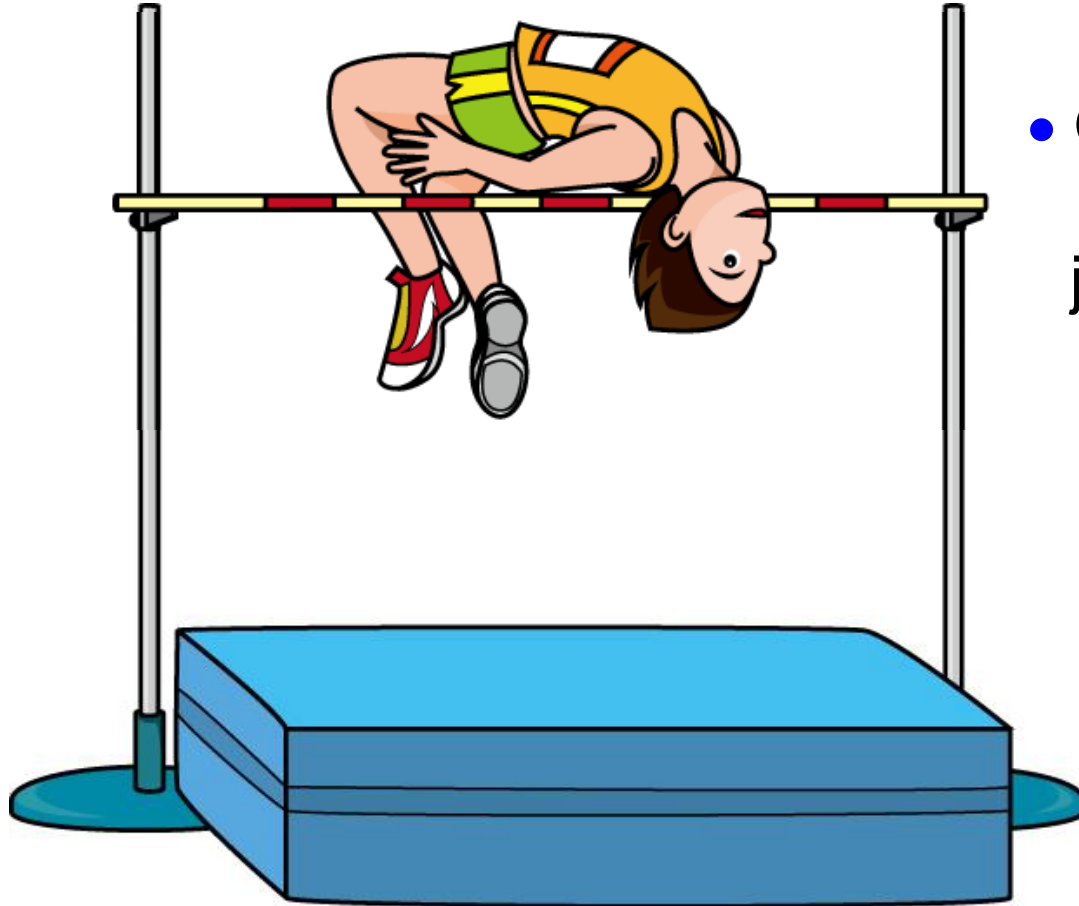


- We say that a catalyst “increases the frequency of effective collisions” between particles of the two reacting chemicals.

# Effect of a Catalyst

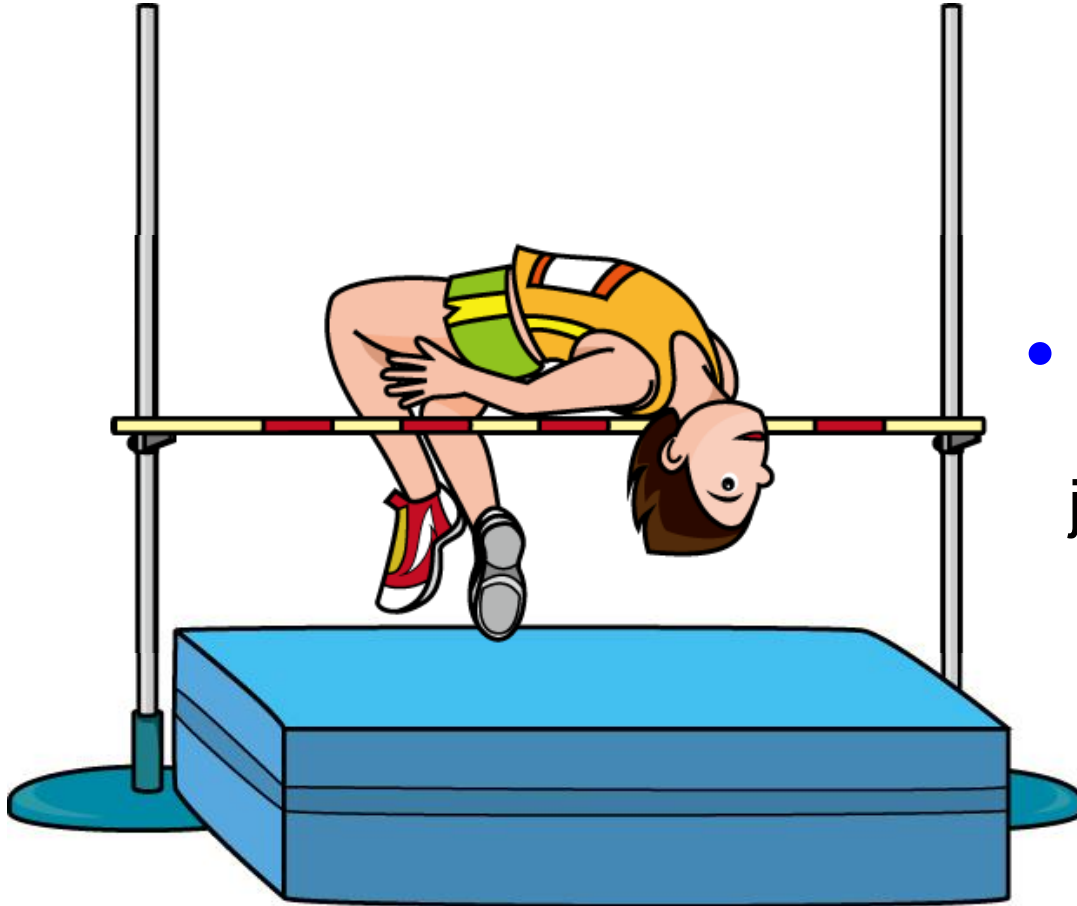


# Effect of a Catalyst



- Only 3 out of 30 students can jump this high.

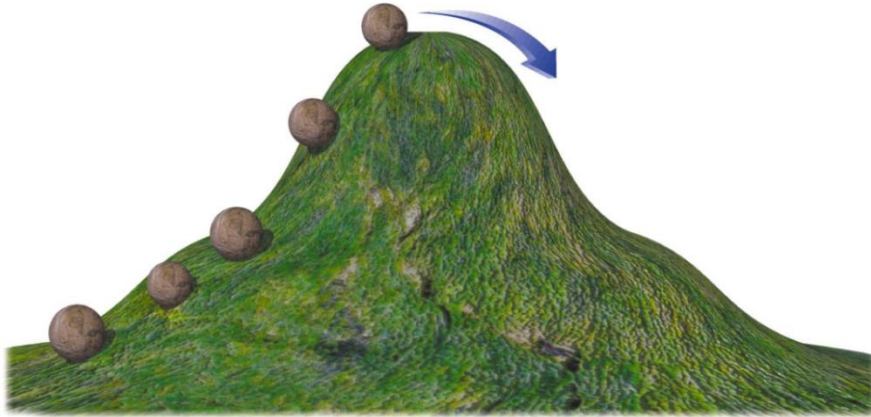
# Effect of a Catalyst



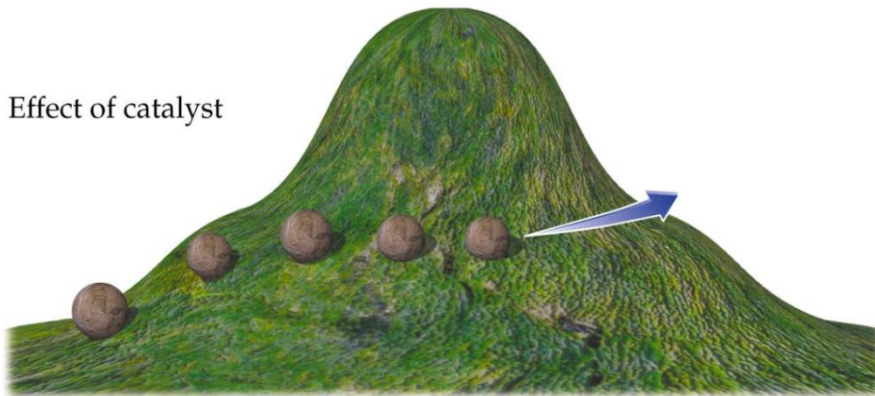
- But 24 out of 30 students can jump this high.



# Effect of a Catalyst

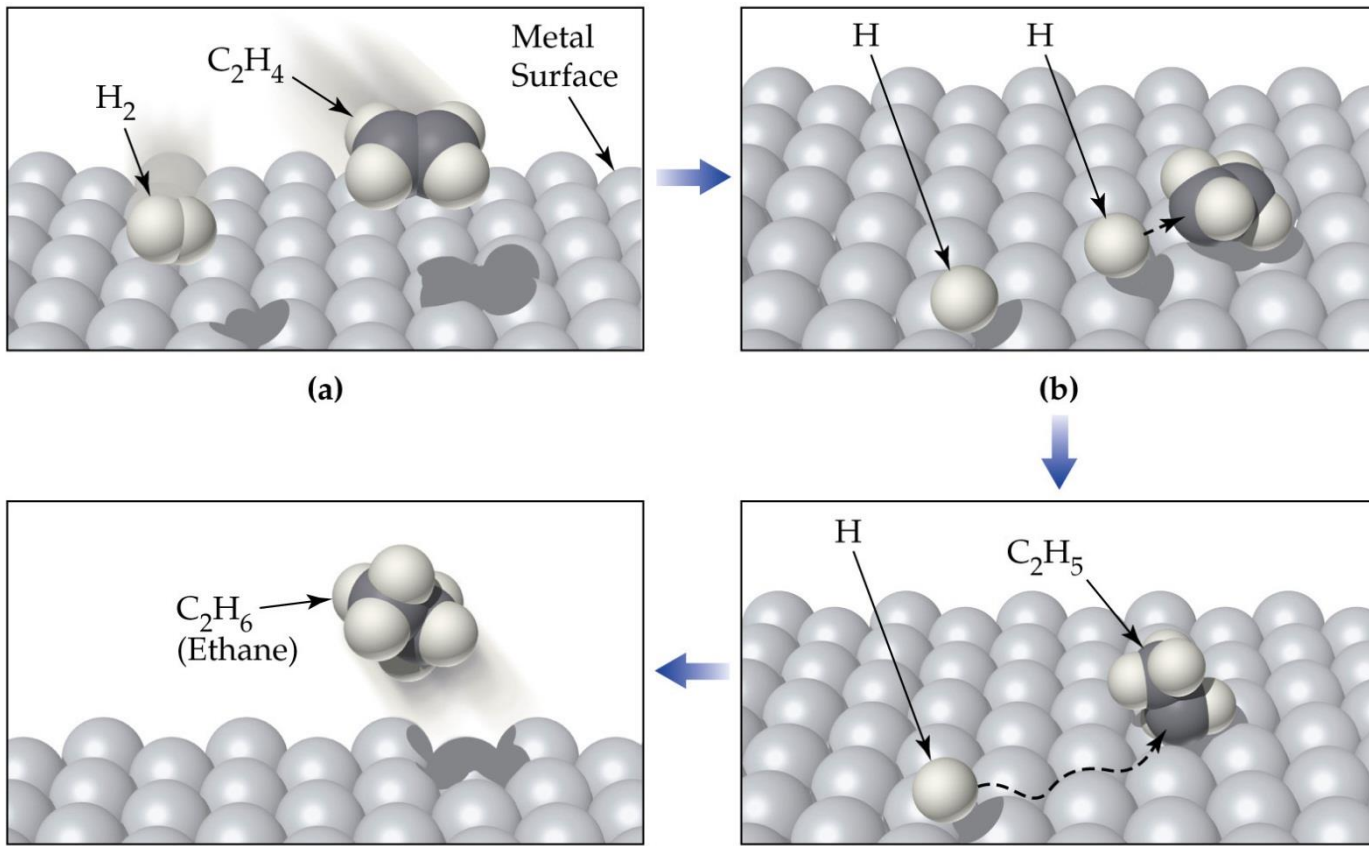


- Analogy for activation energy. **Less energy** is required to push the boulders **around** the side of the hill compared to the energy that is required to push the boulders **over** the hill.



- Using the **same amount of energy**, more boulders can be pushed around the side of the hill compared to the number that can be pushed over the hill.

# Effect of a Catalyst

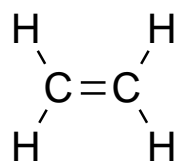


(a)

(b)

(d)

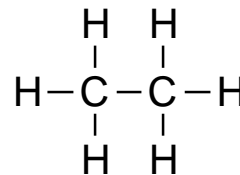
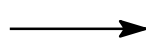
(c)



Ethene



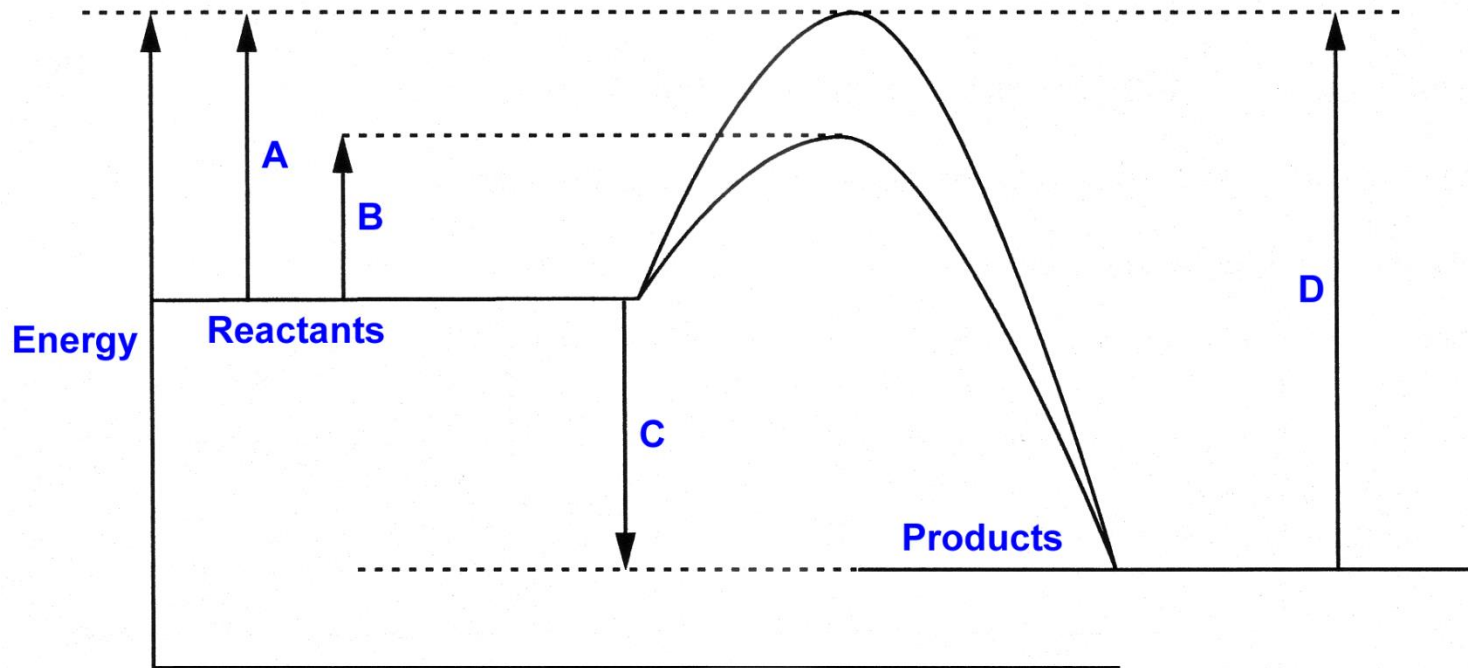
Hydrogen



Ethane

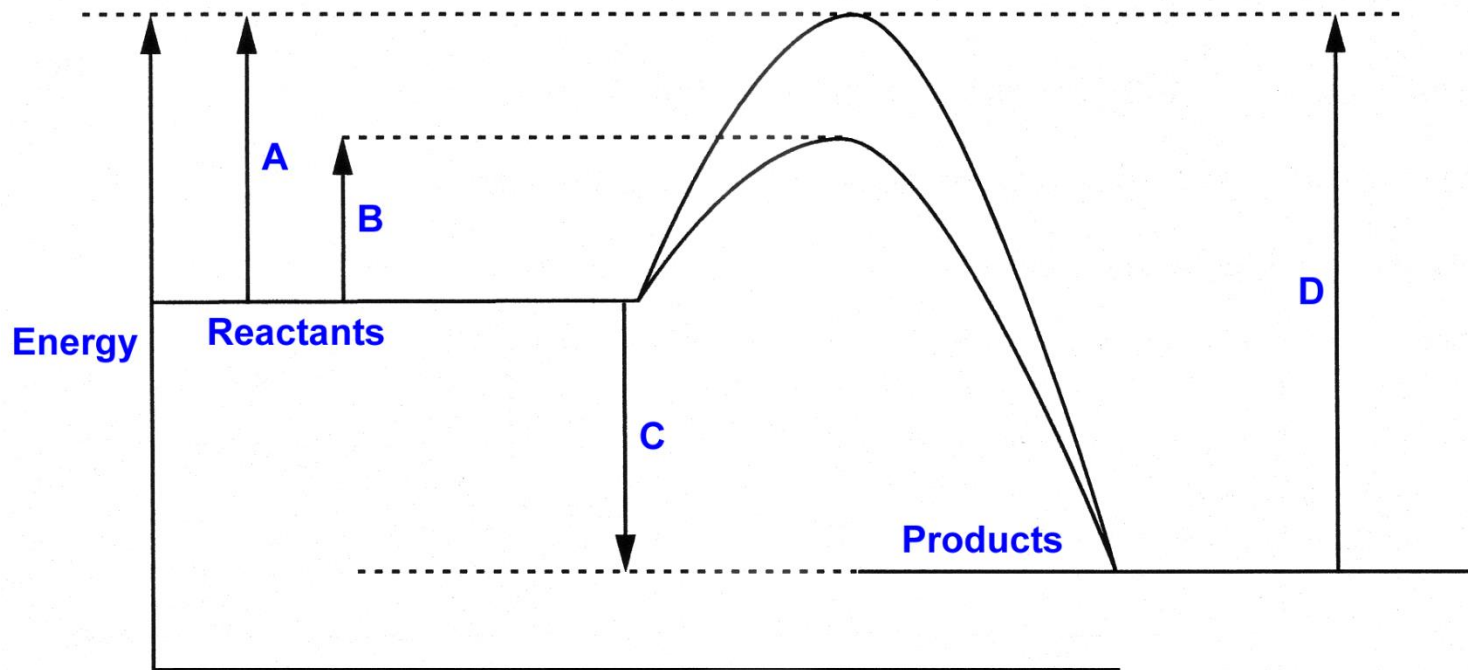
# Assessment for Learning

- Study the energy change diagram shown below.
  - Identify energy changes **A**, **B**, **C** and **D**.



# Assessment for Learning

- Study the energy change diagram shown below.
  - Identify energy changes **A**, **B**, **C** and **D**.



**A** Activation energy of the forward reaction.

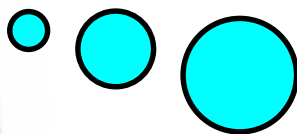
**B** Activation energy of the forward reaction with a catalyst.

**C** Enthalpy change ( $\Delta H$ ) of the forward reaction.

**D** Activation energy of the backward reaction.



# Energy from Chemicals



How can I *calculate*  
the overall energy  
change of a  
reaction?



# Energy from Chemicals

- Imagine that you were given \$50 (+50).
- You spend \$30 (-30).
- You will be in *credit* by \$20  
 $(+50) + (-30) = +20$ .
- Overall, you have “*gained money from the surroundings*”.





# Energy from Chemicals

- Imagine that you were given \$50 (+50).
- You spend \$80 (−80).
- You will be in *debt* by \$30  
 $(+50) + (-80) = -30$ .
- Overall, you have “*lost money to the surroundings*”.



# Energy Change Calculations

Reactions can be considered to take place in two stages:

- 1) Bonds between atoms in the reactants are *broken*.
- 2) New bonds between atoms are *formed* to produce the products.



# Energy Change Calculations

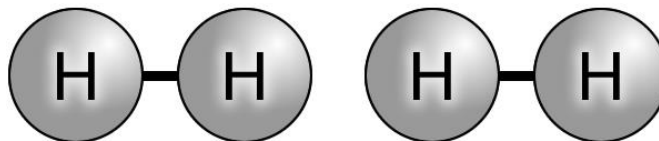
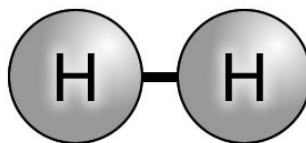
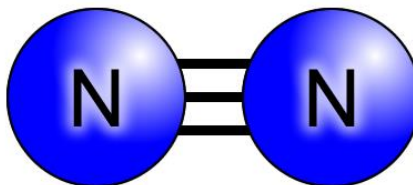
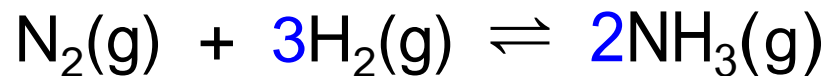
## Important Things to Note...

- Bond **breaking** *absorbs energy* and is therefore **endothermic**.  
Numerical values of  $\Delta H$  for bond **breaking** are **positive**.
- Bond **formation** *releases energy* and is therefore **exothermic**.  
Numerical values of  $\Delta H$  for bond **formation** are **negative**.



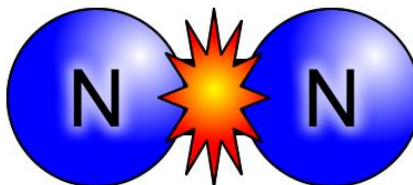
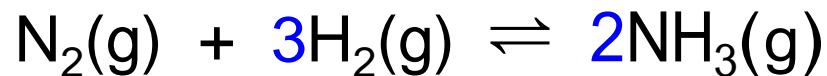
# Energy Change Calculations

nitrogen + hydrogen  $\rightleftharpoons$  ammonia

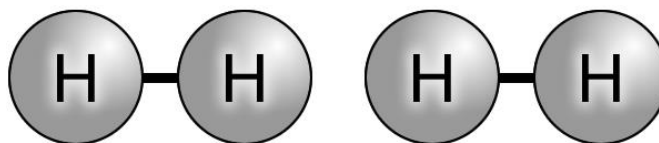
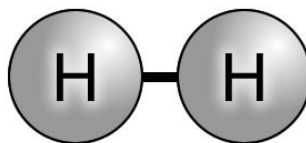


# Energy Change Calculations

nitrogen + hydrogen  $\rightleftharpoons$  ammonia

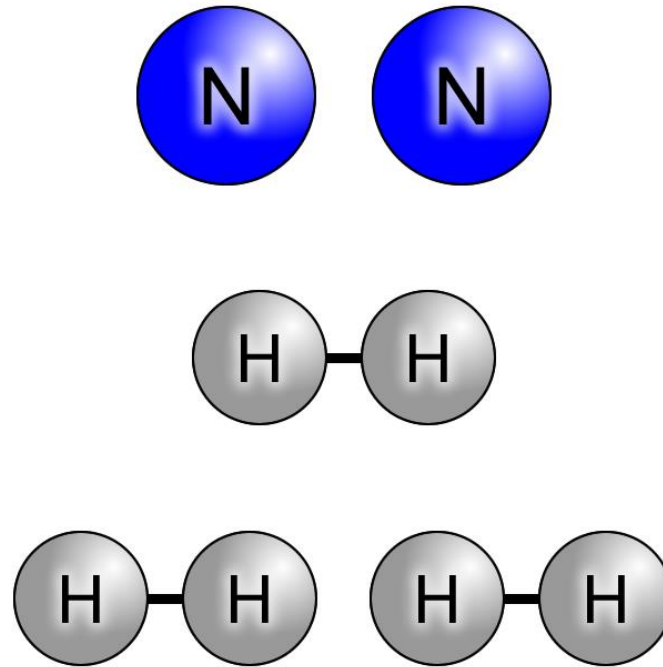
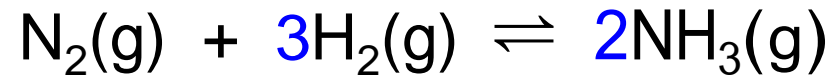


1) Bond breaking  
*absorbs* energy.  
 *$\Delta H$  is positive.*



# Energy Change Calculations

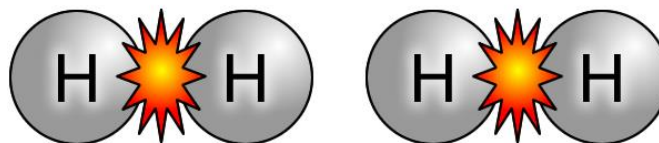
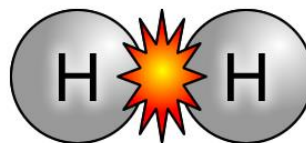
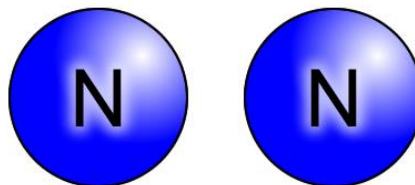
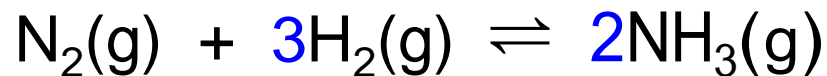
nitrogen + hydrogen  $\rightleftharpoons$  ammonia





# Energy Change Calculations

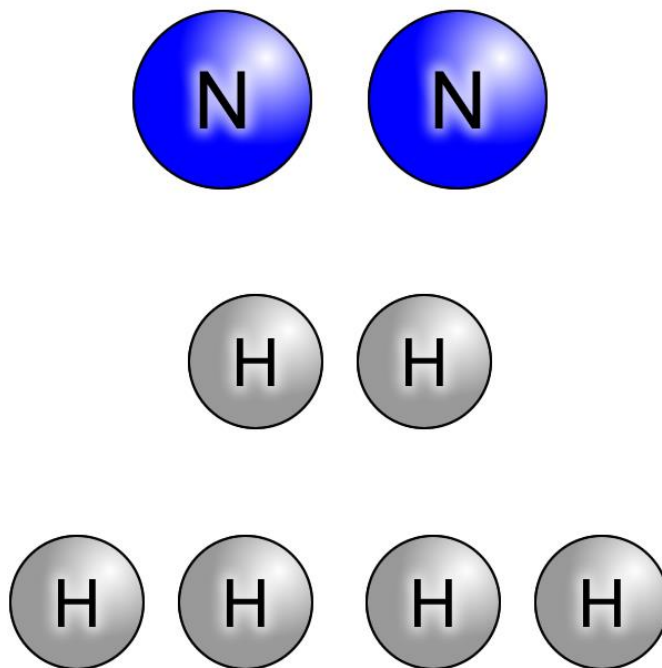
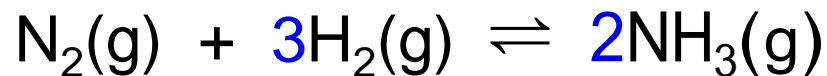
nitrogen + hydrogen  $\rightleftharpoons$  ammonia



1) Bond breaking  
*absorbs* energy.  
 *$\Delta H$  is positive.*

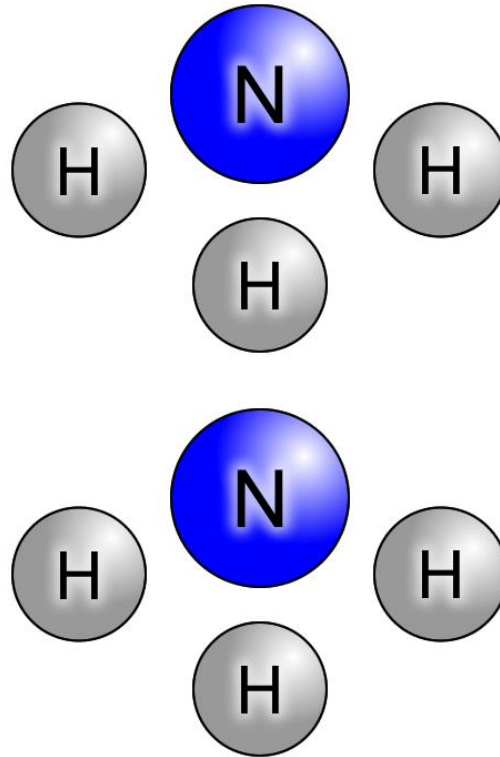
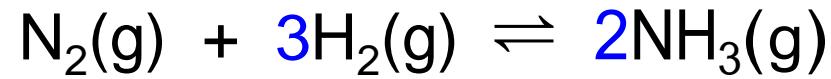
# Energy Change Calculations

nitrogen + hydrogen  $\rightleftharpoons$  ammonia



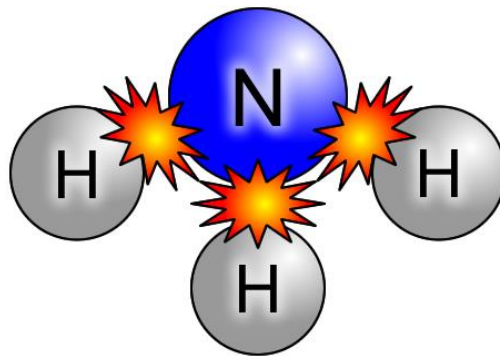
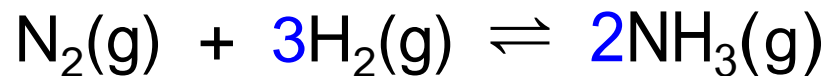
# Energy Change Calculations

nitrogen + hydrogen  $\rightleftharpoons$  ammonia

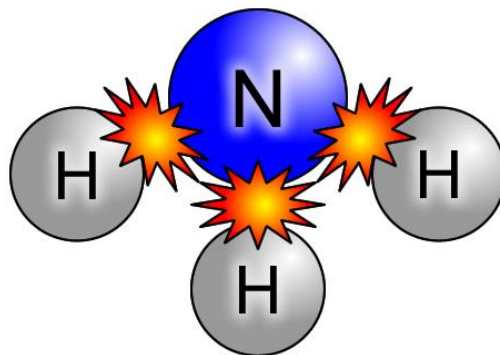


# Energy Change Calculations

nitrogen + hydrogen  $\rightleftharpoons$  ammonia

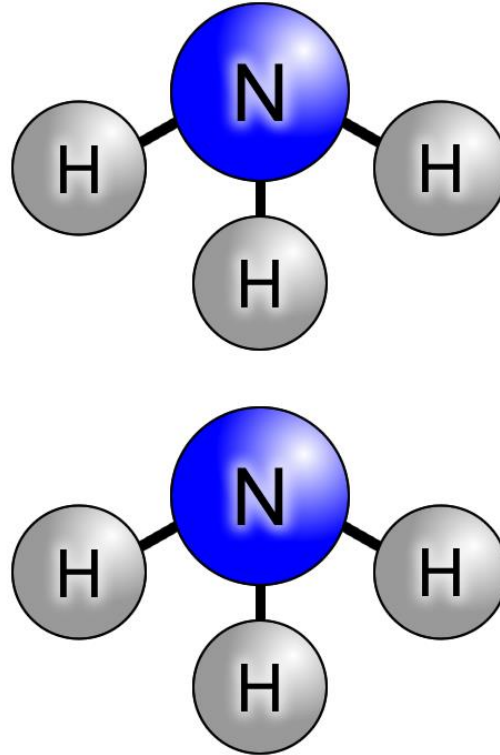
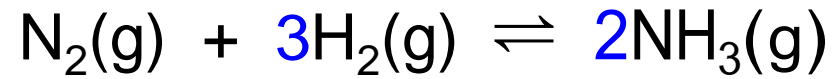


2) Bond formation  
*releases* energy.  
 *$\Delta H$  is negative.*



# Energy Change Calculations

nitrogen + hydrogen  $\rightleftharpoons$  ammonia



# Energy Change Calculations

## Important Things to Note...

Overall enthalpy change of a chemical reaction ( $\Delta H$ )	=	Total amount of energy absorbed to break existing chemical bonds ( $+\Delta H$ )	+	Total amount of energy released when new chemical bonds are formed ( $-\Delta H$ )
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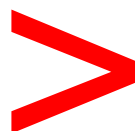




# Energy Change Calculations

If...

Total amount of energy absorbed to break existing chemical bonds  
( $+\Delta H$ )



Total amount of energy released when new chemical bonds are formed  
( $-\Delta H$ )

Overall, energy is absorbed and the reaction will be **endothermic**, i.e.  $\Delta H$  will be positive.



# Energy Change Calculations

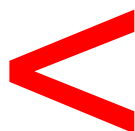
- *Energy absorbed* to *break* bonds in the reactants is *greater* than the *energy released* when new bonds are *formed* in the products.
- Overall, the reaction *absorbs energy* from the surroundings. The reaction is therefore *endothermic*.



# Energy Change Calculations

If...

Total amount of energy absorbed to break existing chemical bonds  
( $+\Delta H$ )



Total amount of energy released when new chemical bonds are formed  
( $-\Delta H$ )

Overall, **energy is released** and the reaction will be **exothermic**, *i.e.*  $\Delta H$  will be negative.



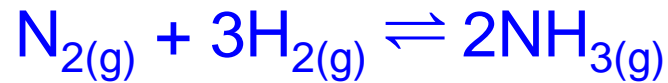
# Energy Change Calculations

- *Energy absorbed* to *break* bonds in the reactants is *less* than the *energy released* when new bonds are *formed* in the products.
- Overall, the reaction *releases energy* into the surroundings. The reaction is therefore *exothermic*.

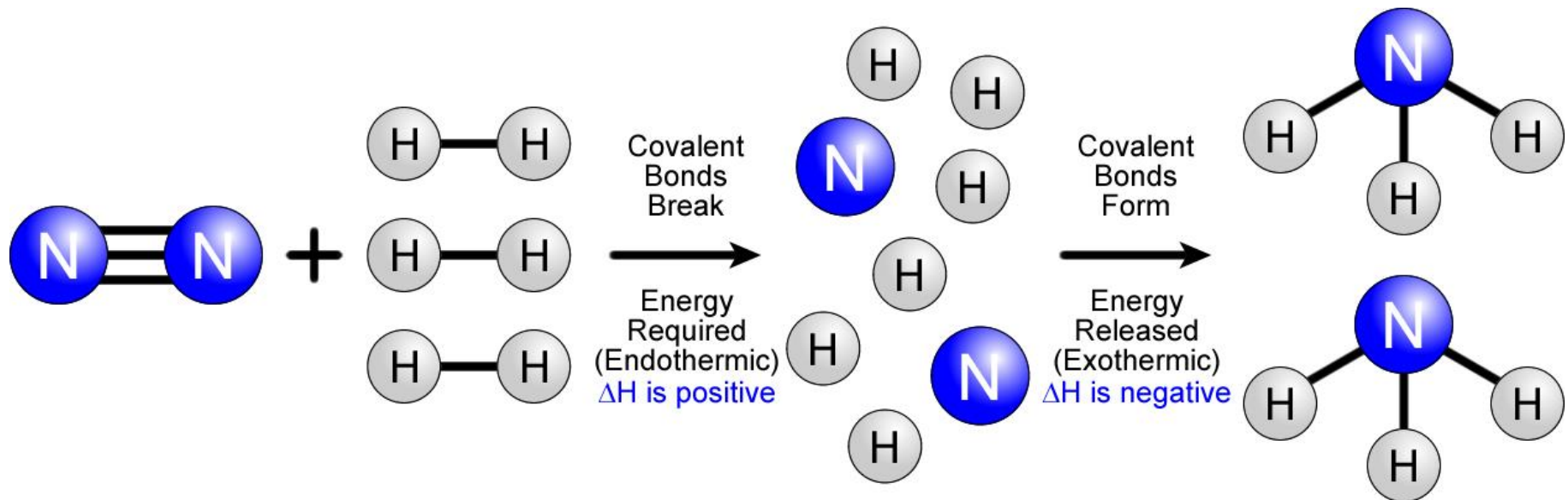


## Energy Change Calculations – Example #1:

The Reaction Between Nitrogen and Hydrogen to form Ammonia

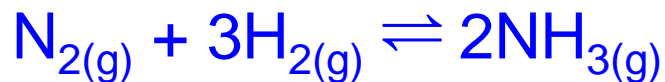


Overview of the Reaction:



## Energy Change Calculations – Example #1:

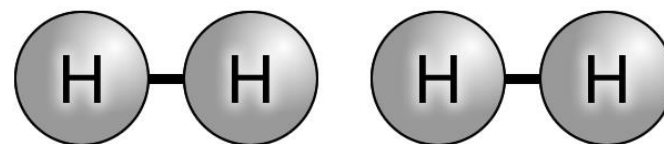
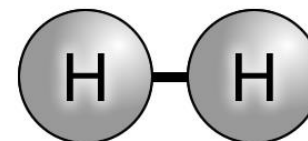
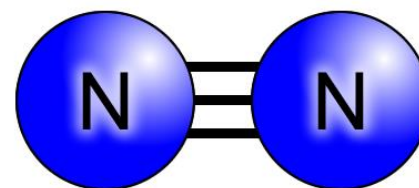
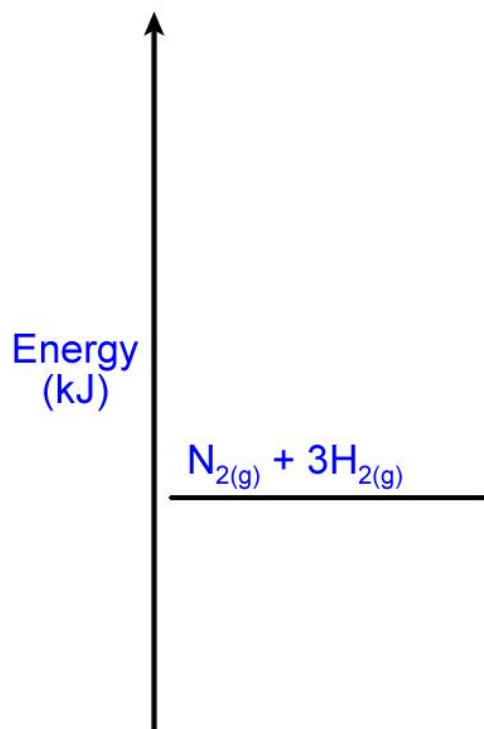
The Reaction Between Nitrogen and Hydrogen to form Ammonia



$\text{N}\equiv\text{N} = 944 \text{ kJ/mol}$

$\text{H}-\text{H} = 436 \text{ kJ/mol}$

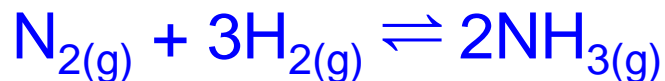
$\text{N}-\text{H} = 388 \text{ kJ/mol}$





## Energy Change Calculations – Example #1:

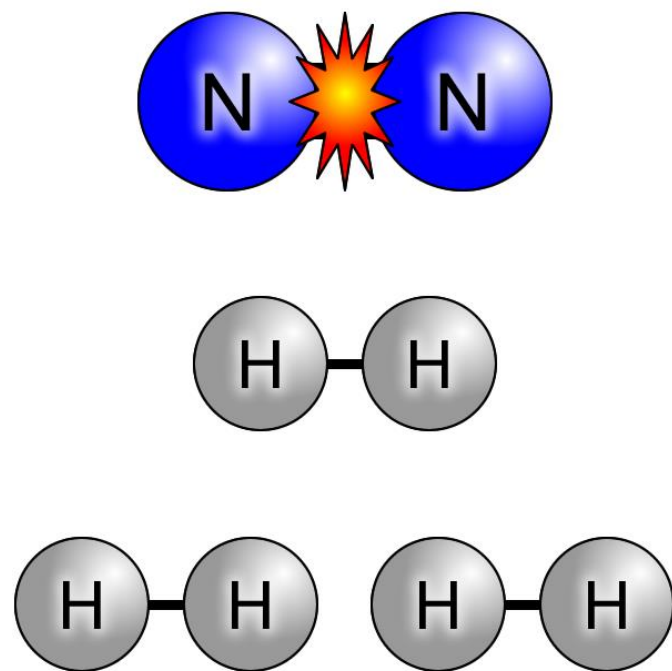
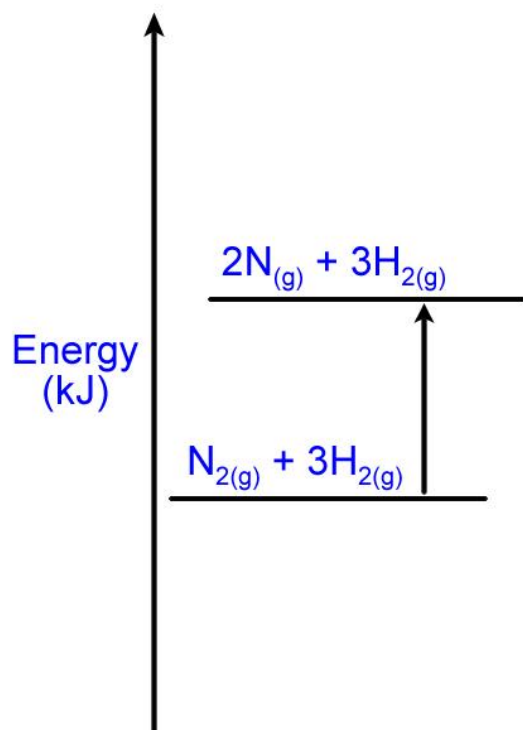
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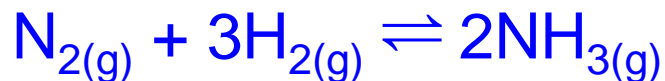
$\text{N}-\text{H} = 388 \text{ kJ/mol}$



1  $\times$   $\text{N}\equiv\text{N}$  bond is broken forming 2  $\times$  N atoms.

## Energy Change Calculations – Example #1:

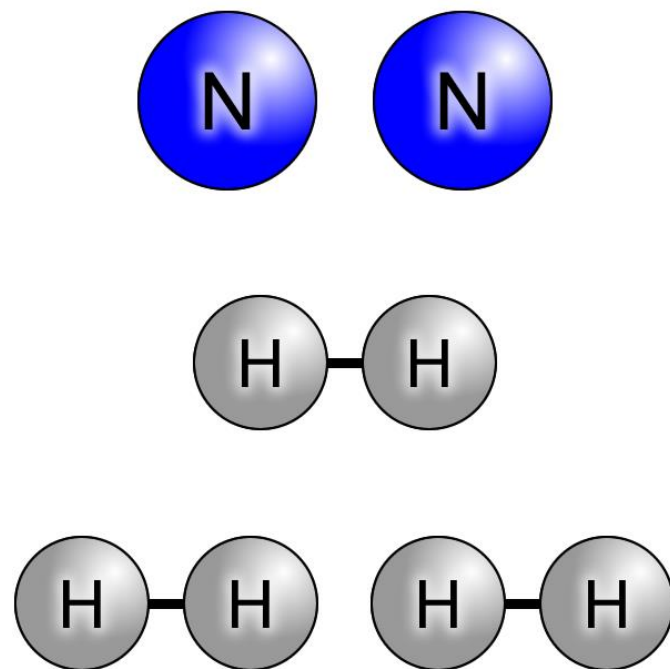
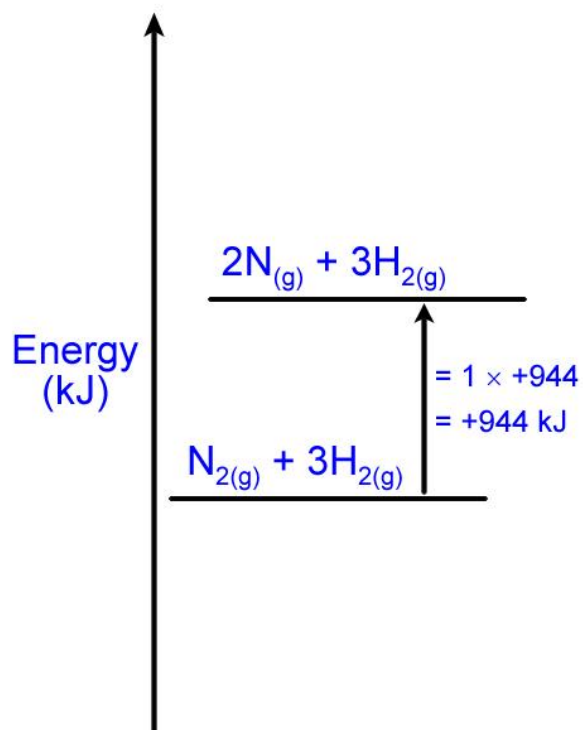
The Reaction Between Nitrogen and Hydrogen to form Ammonia



$\text{N}\equiv\text{N} = 944 \text{ kJ/mol}$

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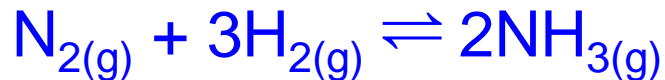
$\text{N}-\text{H} = 388 \text{ kJ/mol}$



Bond breaking is *endothermic*:  $\Delta H$  for this change is *positive*.

## Energy Change Calculations – Example #1:

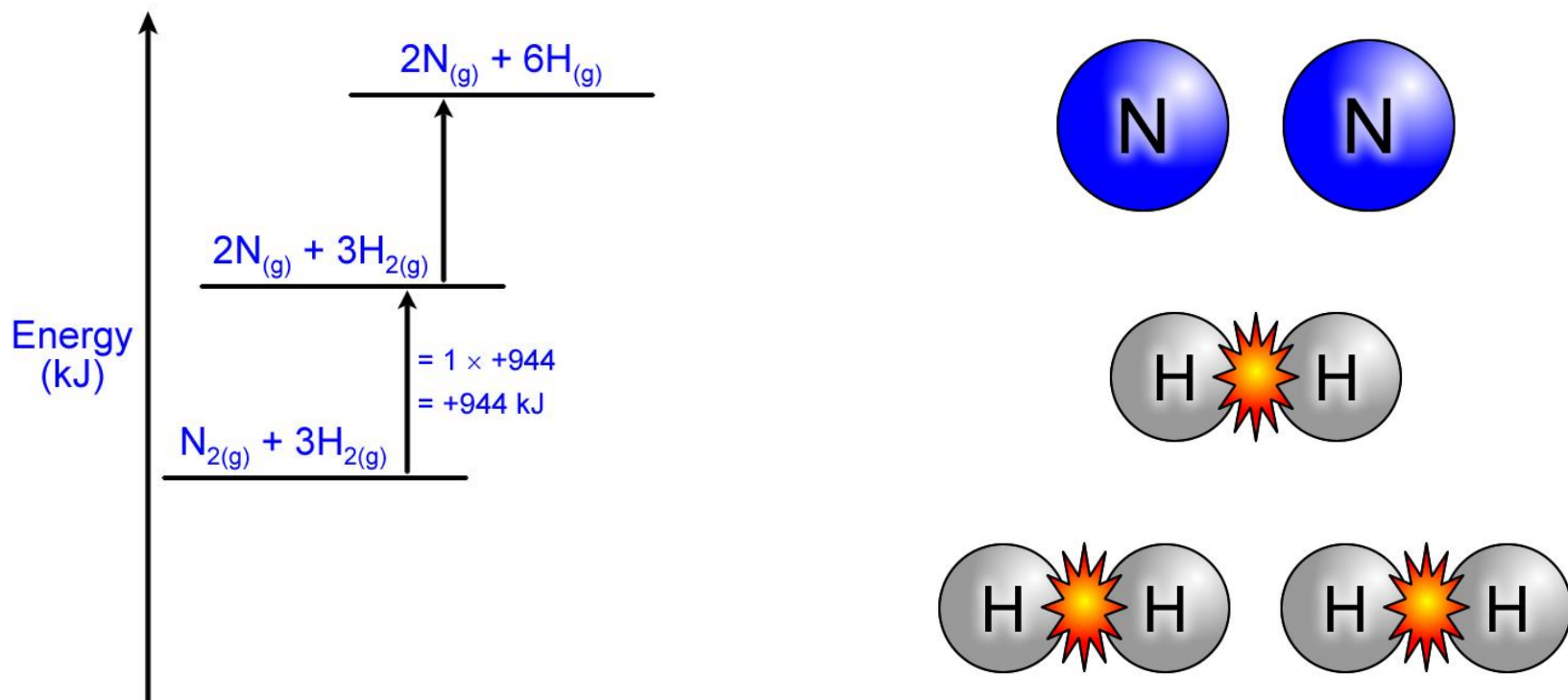
The Reaction Between Nitrogen and Hydrogen to form Ammonia



$\text{N}\equiv\text{N} = 944 \text{ kJ/mol}$

$\text{H}-\text{H} = 436 \text{ kJ/mol}$

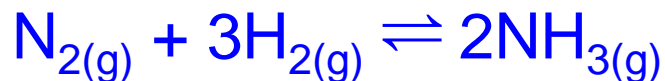
$\text{N}-\text{H} = 388 \text{ kJ/mol}$



$3 \times \text{H}-\text{H}$  bonds are broken forming  $6 \times \text{H}$  atoms.

## Energy Change Calculations – Example #1:

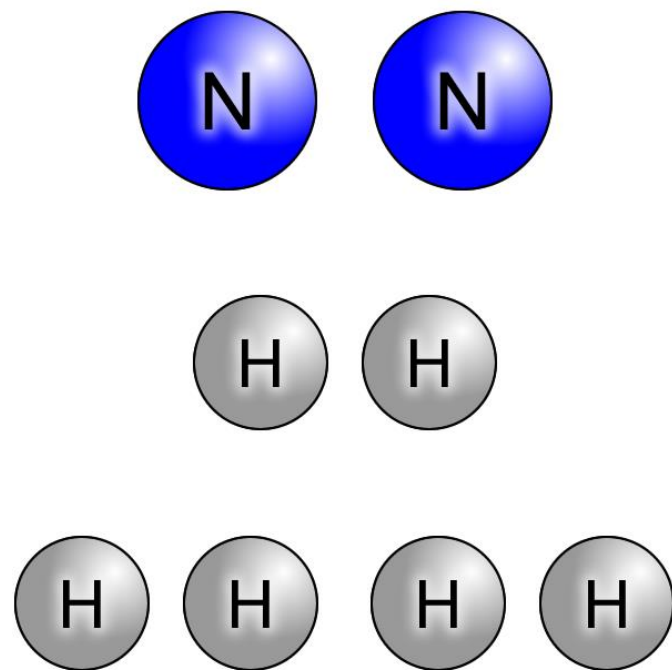
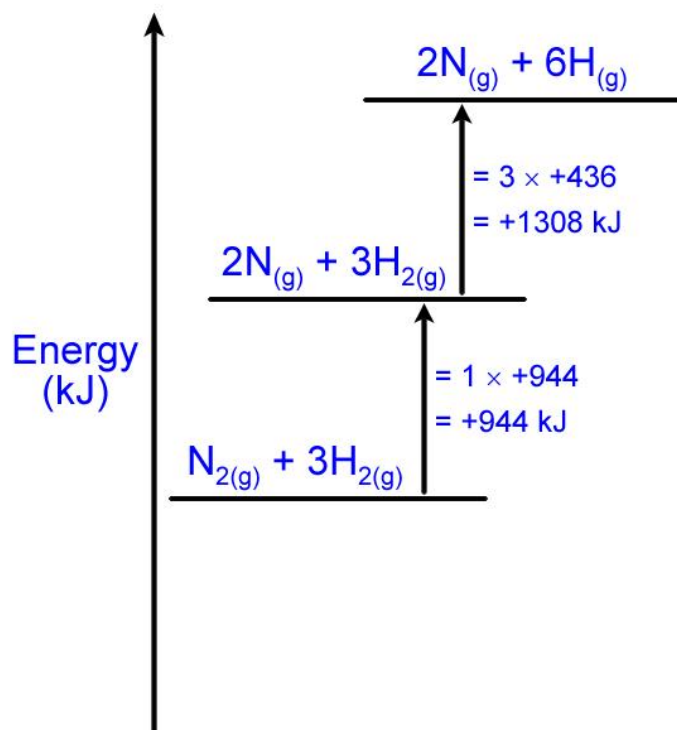
The Reaction Between Nitrogen and Hydrogen to form Ammonia



$\text{N}\equiv\text{N} = 944 \text{ kJ/mol}$

$\text{H}-\text{H} = 436 \text{ kJ/mol}$

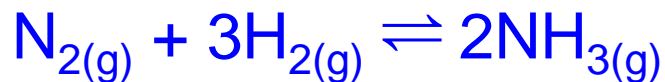
$\text{N}-\text{H} = 388 \text{ kJ/mol}$



Bond breaking is *endothermic*:  $\Delta H$  for this change is *positive*.

# Energy Change Calculations – Example #1:

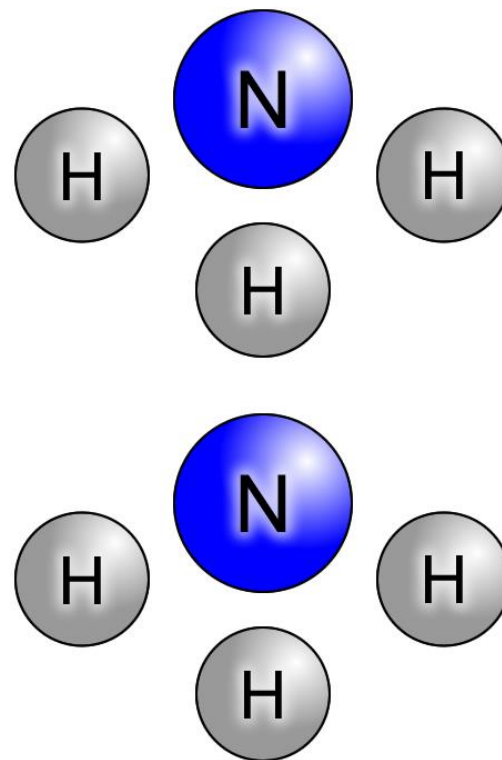
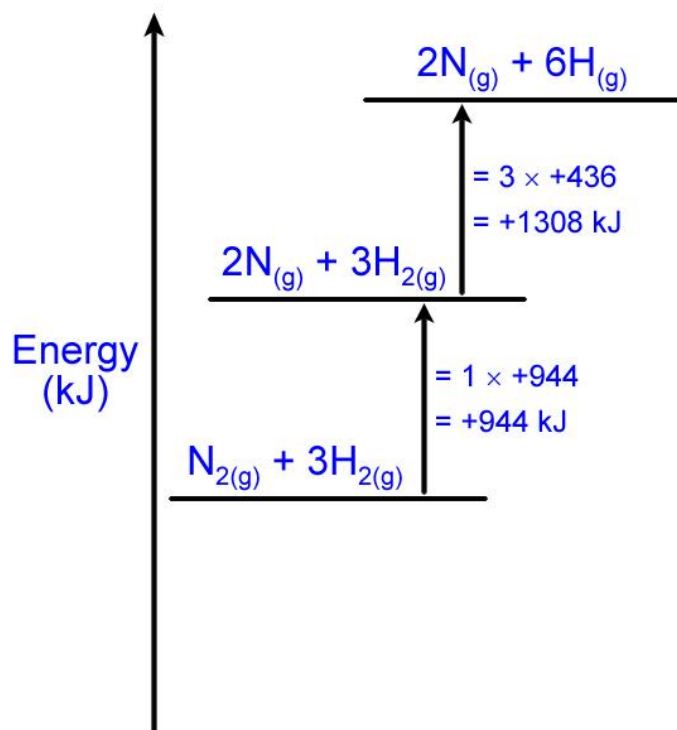
The Reaction Between Nitrogen and Hydrogen to form Ammonia



$\text{N}\equiv\text{N} = 944 \text{ kJ/mol}$

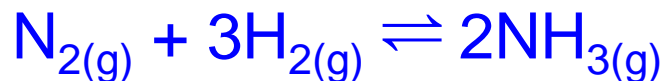
$\text{H}-\text{H} = 436 \text{ kJ/mol}$

$\text{N}-\text{H} = 388 \text{ kJ/mol}$



## Energy Change Calculations – Example #1:

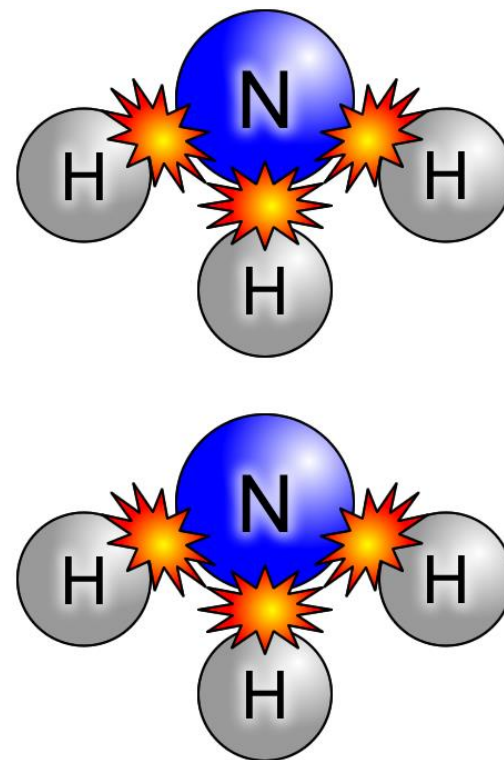
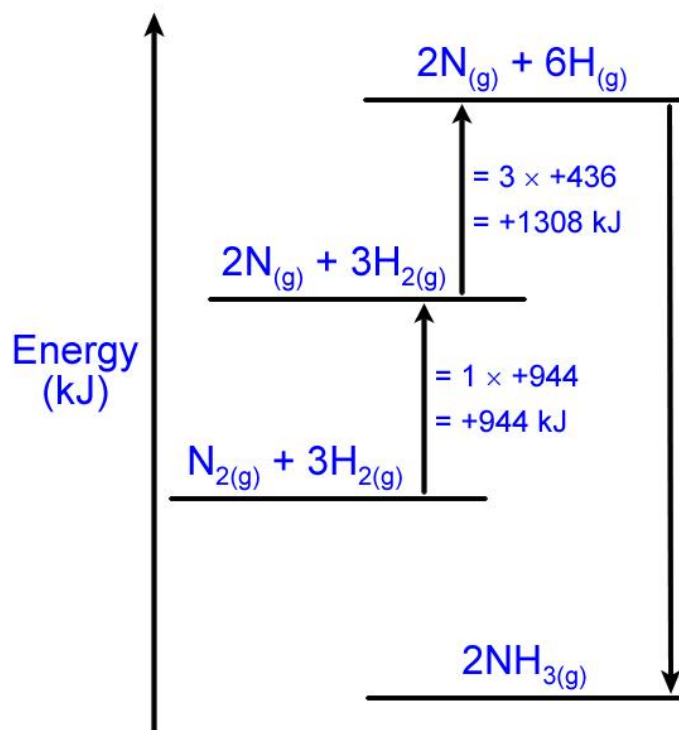
The Reaction Between Nitrogen and Hydrogen to form Ammonia



$\text{N}\equiv\text{N} = 944 \text{ kJ/mol}$

$\text{H}-\text{H} = 436 \text{ kJ/mol}$

$\text{N}-\text{H} = 388 \text{ kJ/mol}$

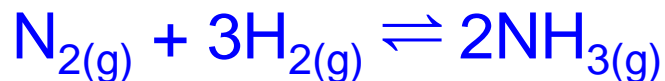


$6 \times \text{N}-\text{H}$  bonds are formed creating  $2 \times \text{NH}_3$  molecules.



## Energy Change Calculations – Example #1:

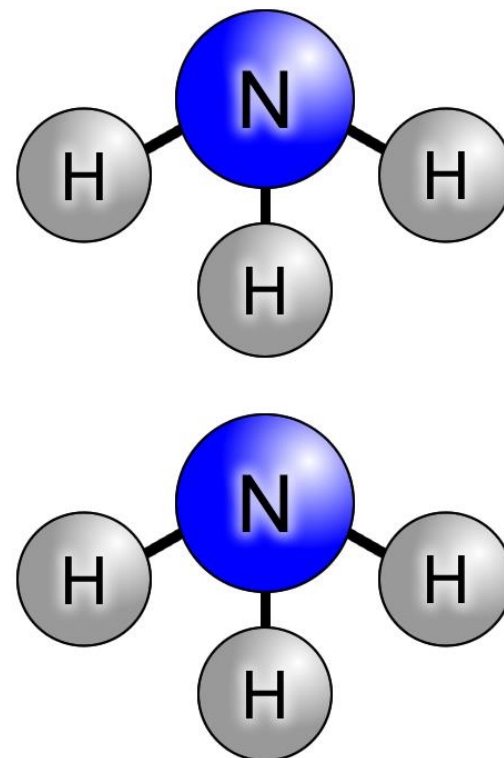
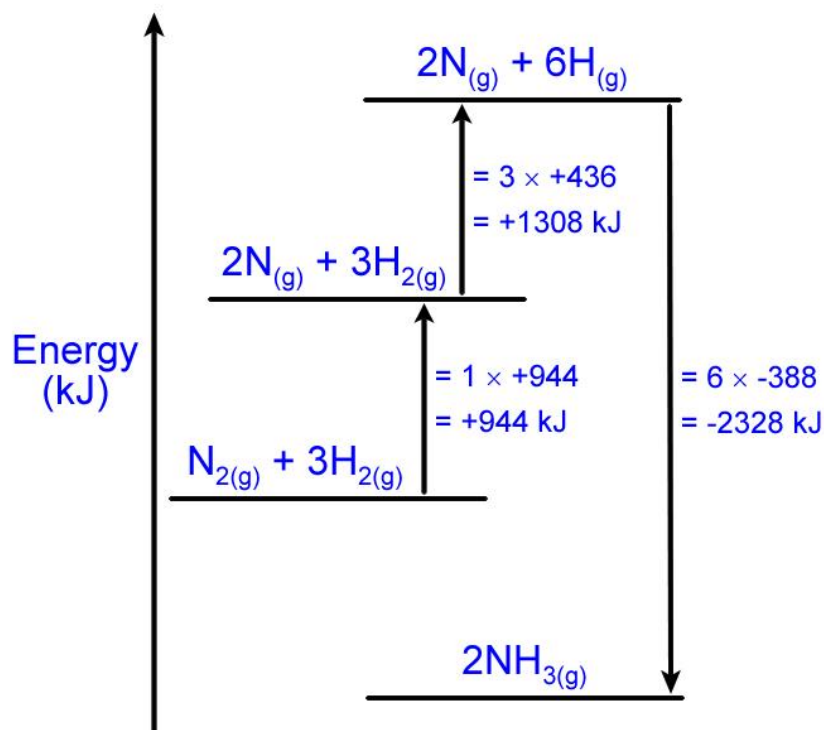
The Reaction Between Nitrogen and Hydrogen to form Ammonia



$\text{N}\equiv\text{N} = 944 \text{ kJ/mol}$

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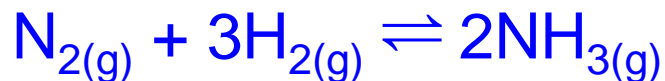
$\text{N}-\text{H} = 388 \text{ kJ/mol}$



Bond formation is *exothermic*:  $\Delta\text{H}$  for this change is *negative*.

## Energy Change Calculations – Example #1:

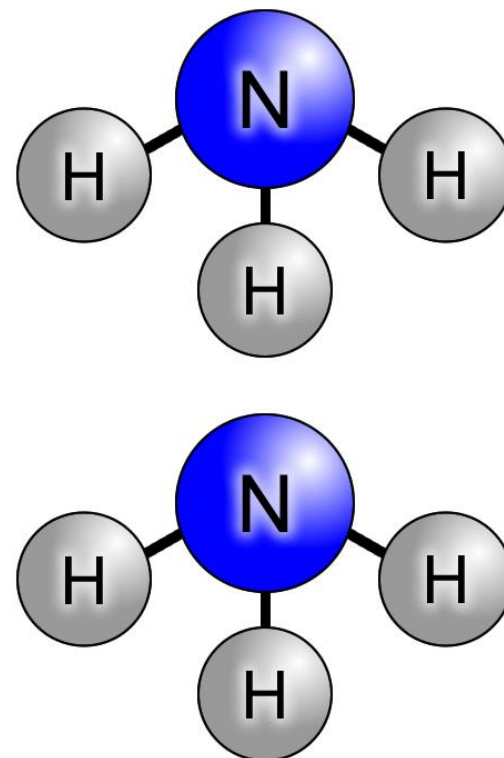
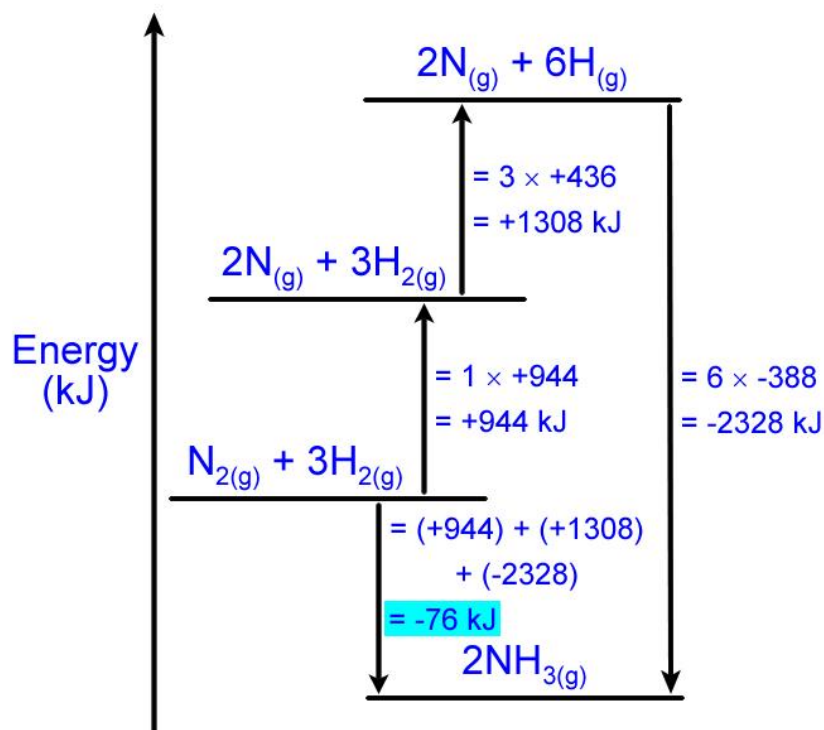
The Reaction Between Nitrogen and Hydrogen to form Ammonia



$\text{N}\equiv\text{N} = 944 \text{ kJ/mol}$

$\text{H}-\text{H} = 436 \text{ kJ/mol}$

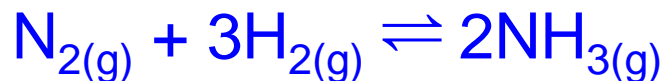
$\text{N}-\text{H} = 388 \text{ kJ/mol}$



$$(+944) + (+1308) + (-2328) = -76.0 \text{ kJ}$$

## Energy Change Calculations – Example #1:

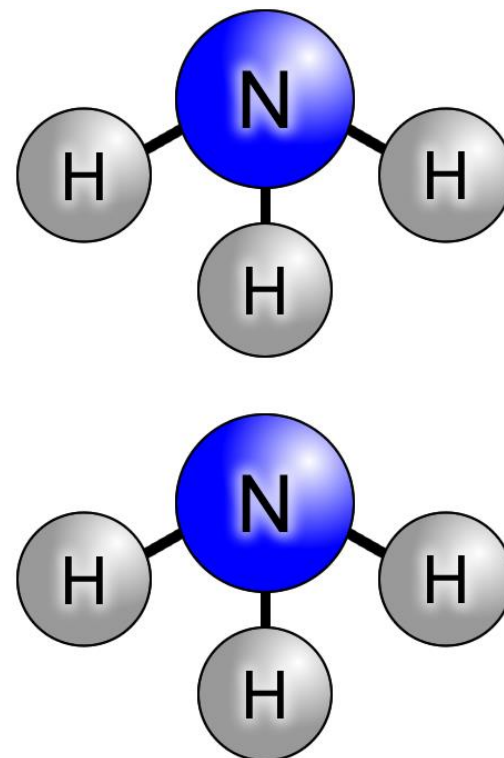
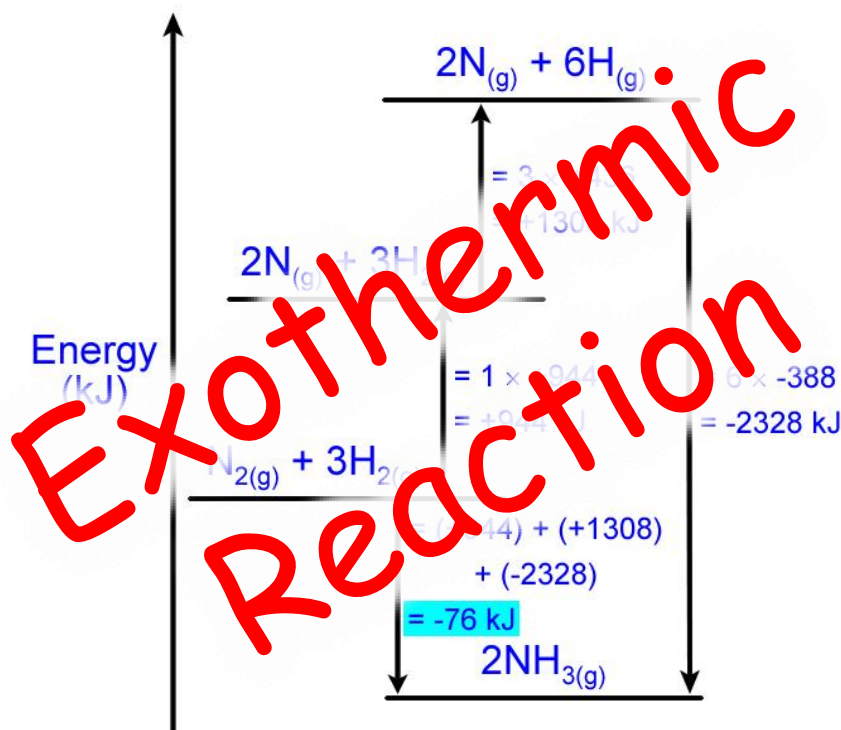
The Reaction Between Nitrogen and Hydrogen to form Ammonia



$\text{N}\equiv\text{N} = 944 \text{ kJ/mol}$

$\text{H}-\text{H} = 436 \text{ kJ/mol}$

$\text{N}-\text{H} = 388 \text{ kJ/mol}$



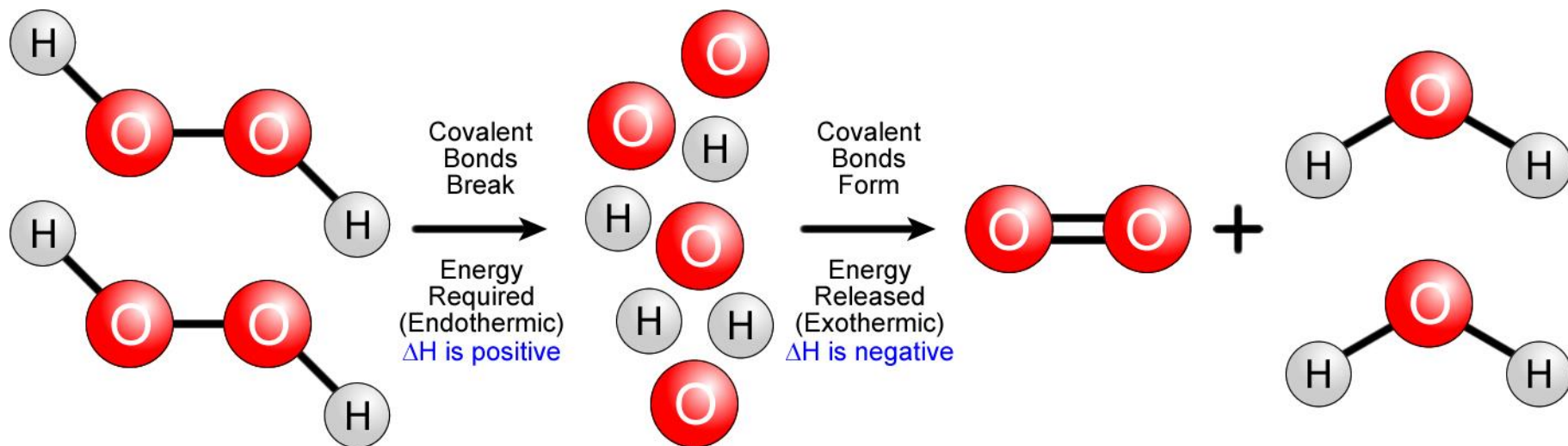
$$(+944) + (+1308) + (-2328) = \text{-76.0 kJ}$$

## Energy Change Calculations – Example #2:

The Decomposition of Hydrogen Peroxide to form Oxygen and Water



Overview of the Reaction:



## Energy Change Calculations – Example #2:

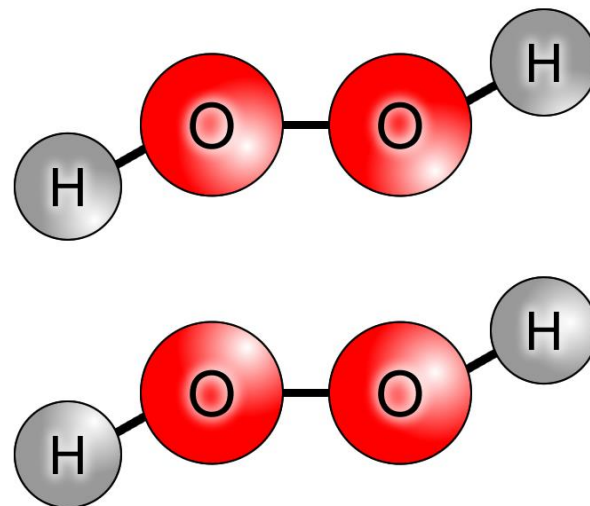
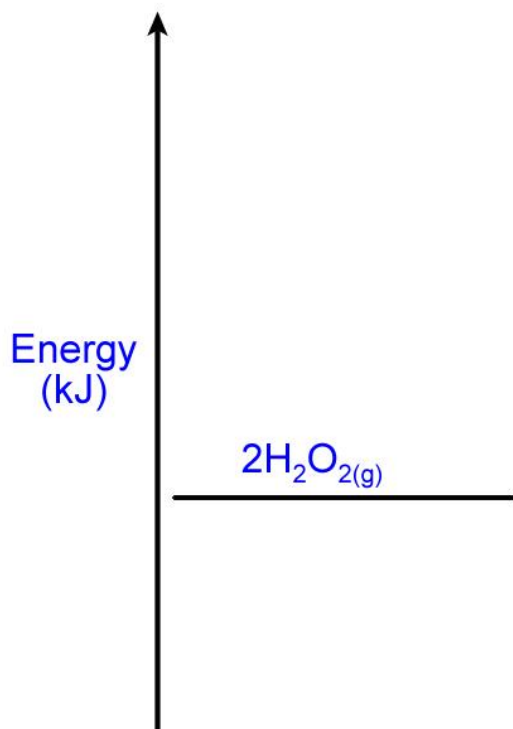
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

O–H = 463 kJ/mol

O=O = 496 kJ/mol



## Energy Change Calculations – Example #2:

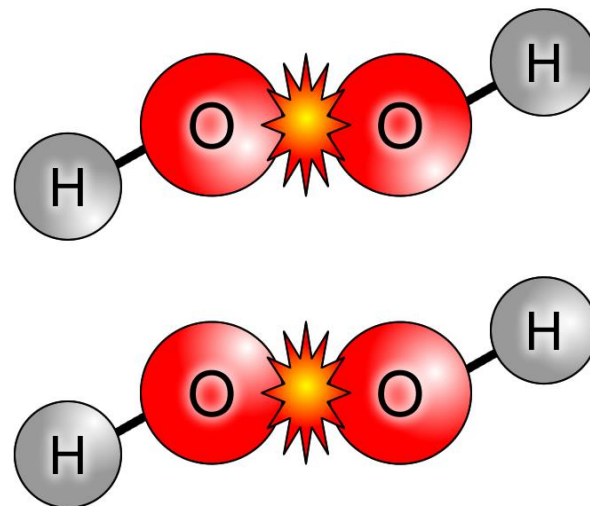
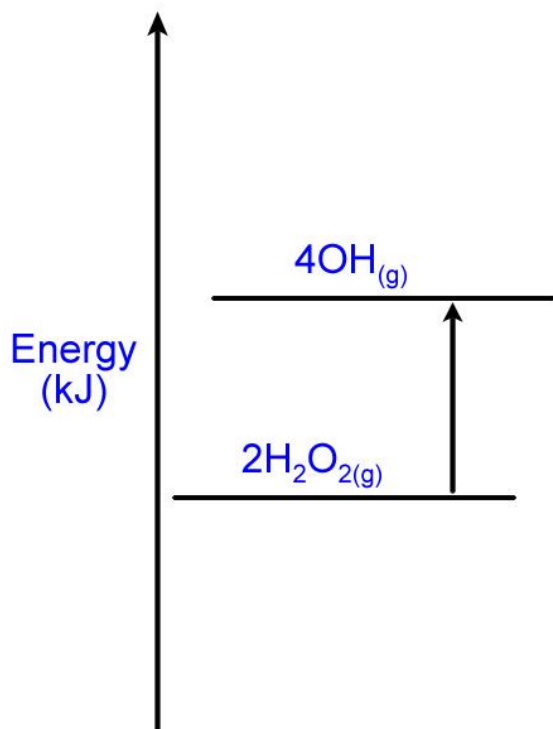
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

O–H = 463 kJ/mol

O=O = 496 kJ/mol



$2 \times \text{O–O}$  bond are broken forming  $4 \times \text{OH}$  molecules.

## Energy Change Calculations – Example #2:

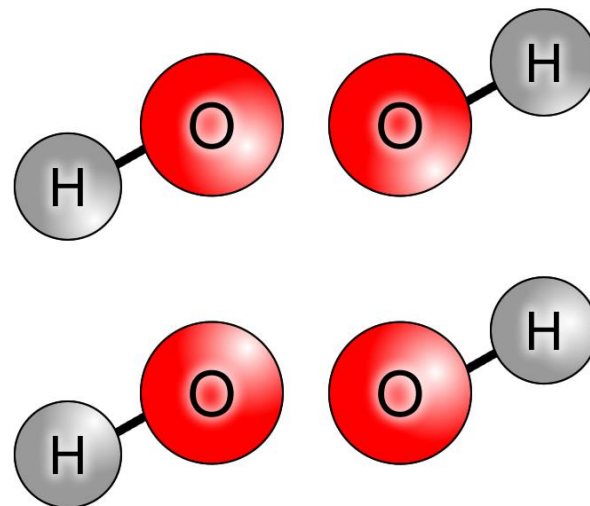
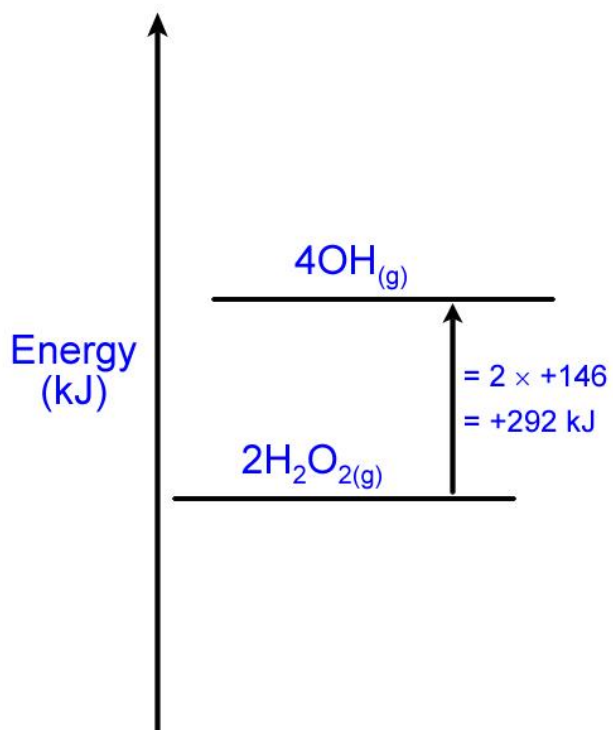
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

O–H = 463 kJ/mol

O=O = 496 kJ/mol



Bond breaking is *endothermic*:  $\Delta H$  for this change is *positive*.



## Energy Change Calculations – Example #2:

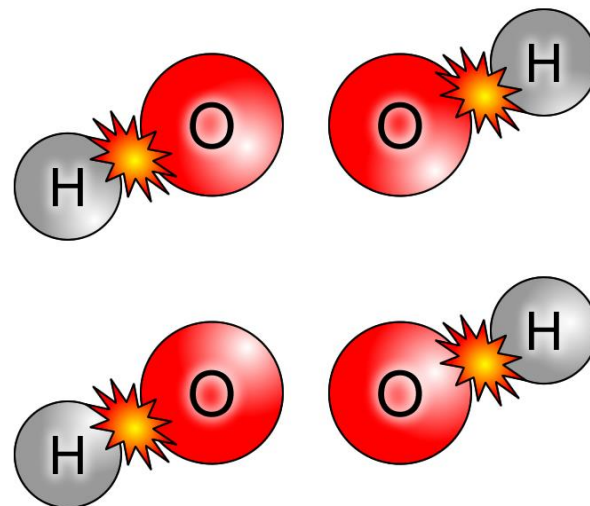
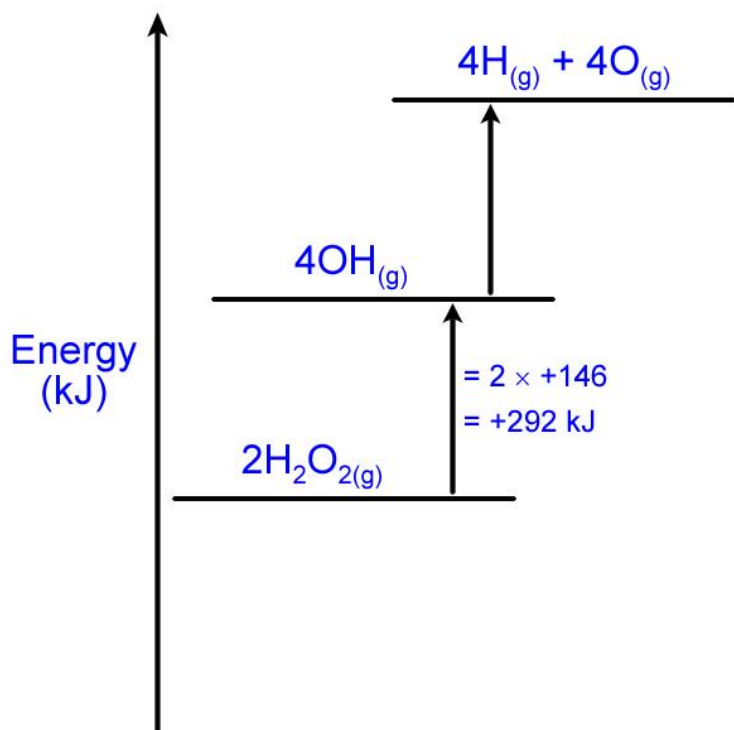
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

O–H = 463 kJ/mol

O=O = 496 kJ/mol



$4 \times \text{O–H}$  bonds are broken forming  $4 \times \text{O}$  and  $4 \times \text{H}$  atoms.

## Energy Change Calculations – Example #2:

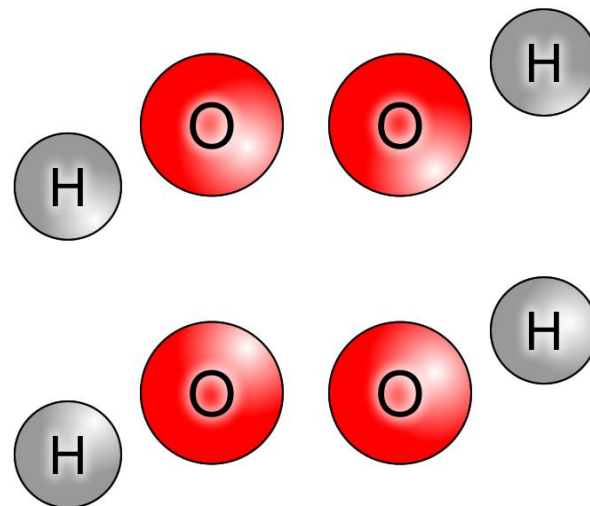
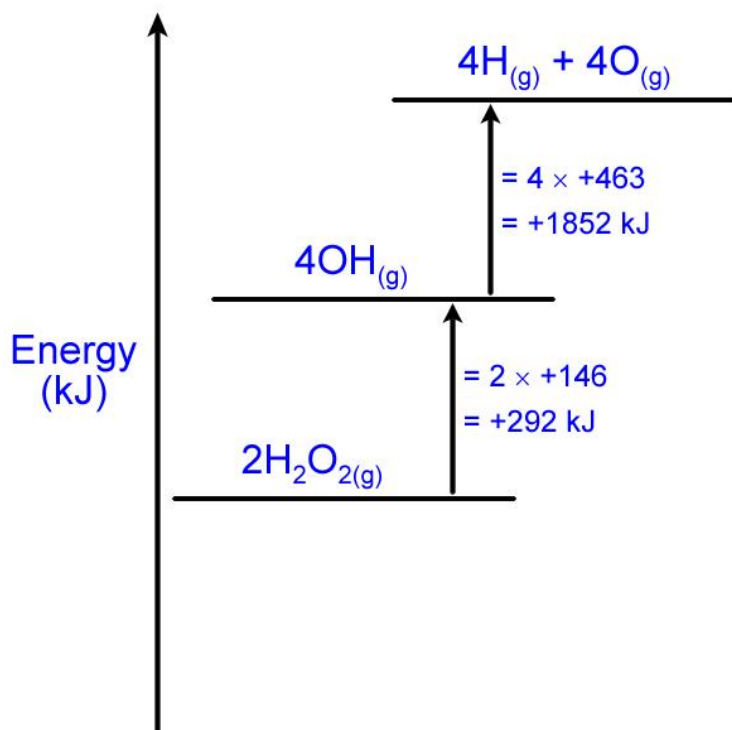
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Bond breaking is *endothermic*:  $\Delta\text{H}$  for this change is *positive*.

## Energy Change Calculations – Example #2:

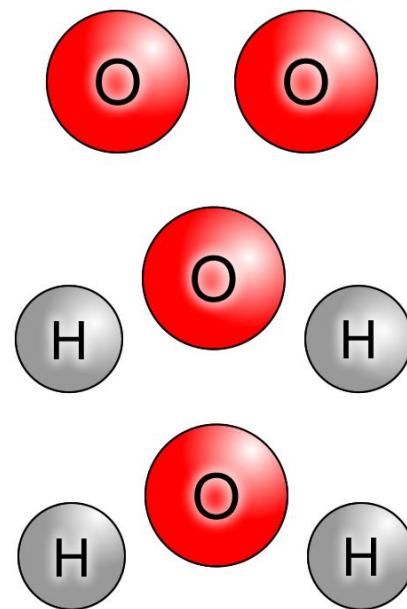
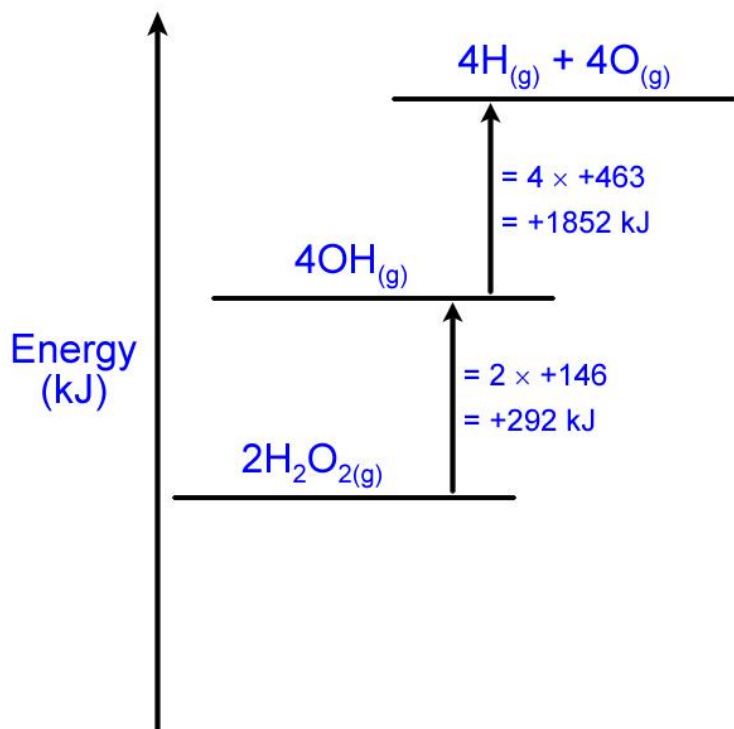
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

O–H = 463 kJ/mol

O=O = 496 kJ/mol



## Energy Change Calculations – Example #2:

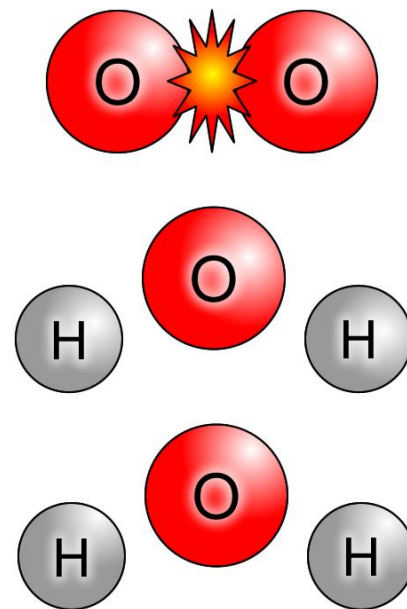
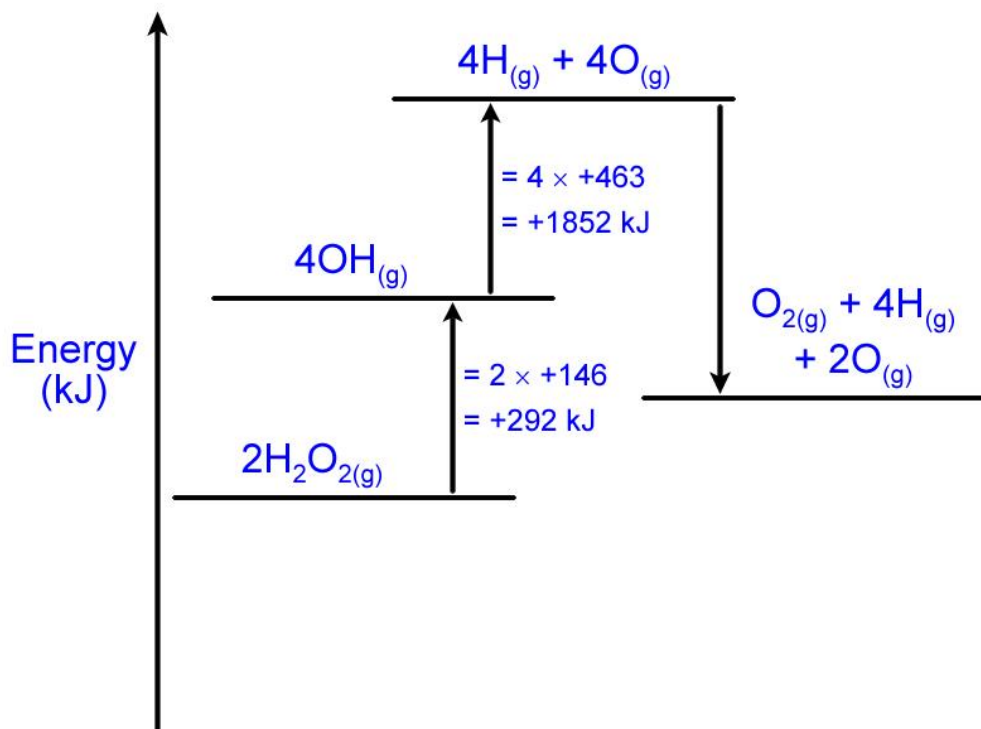
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

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O=O = 496 kJ/mol



1 × O=O bond is formed creating 1 × O<sub>2</sub> molecule.

## Energy Change Calculations – Example #2:

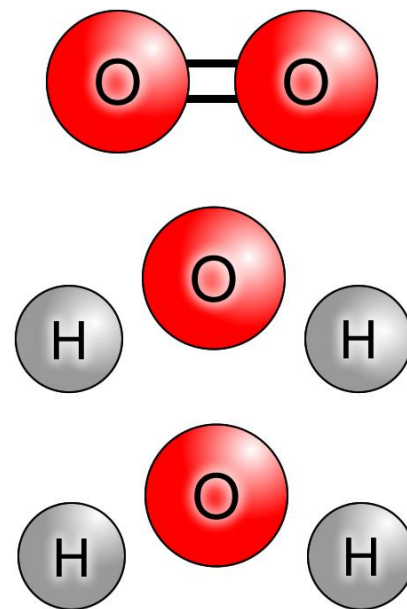
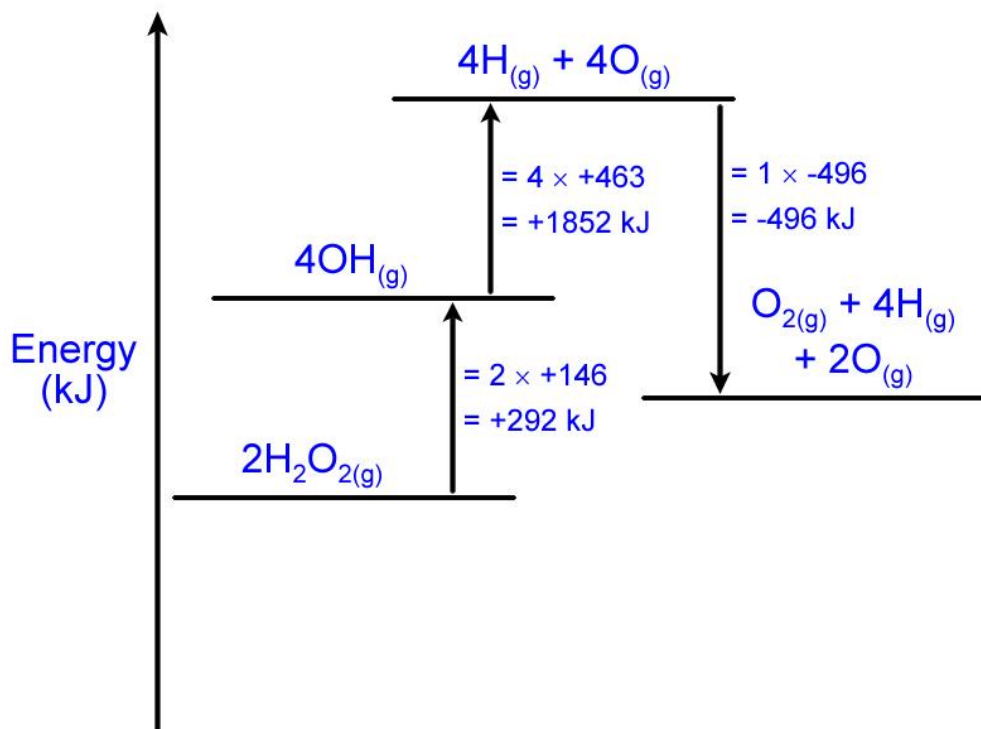
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

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Bond formation is *exothermic*:  $\Delta\text{H}$  for this change is *negative*.

## Energy Change Calculations – Example #2:

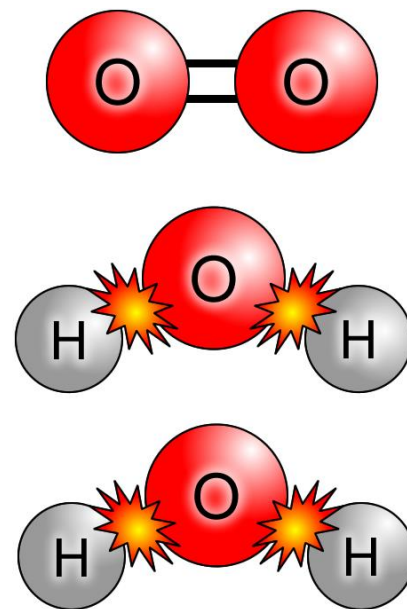
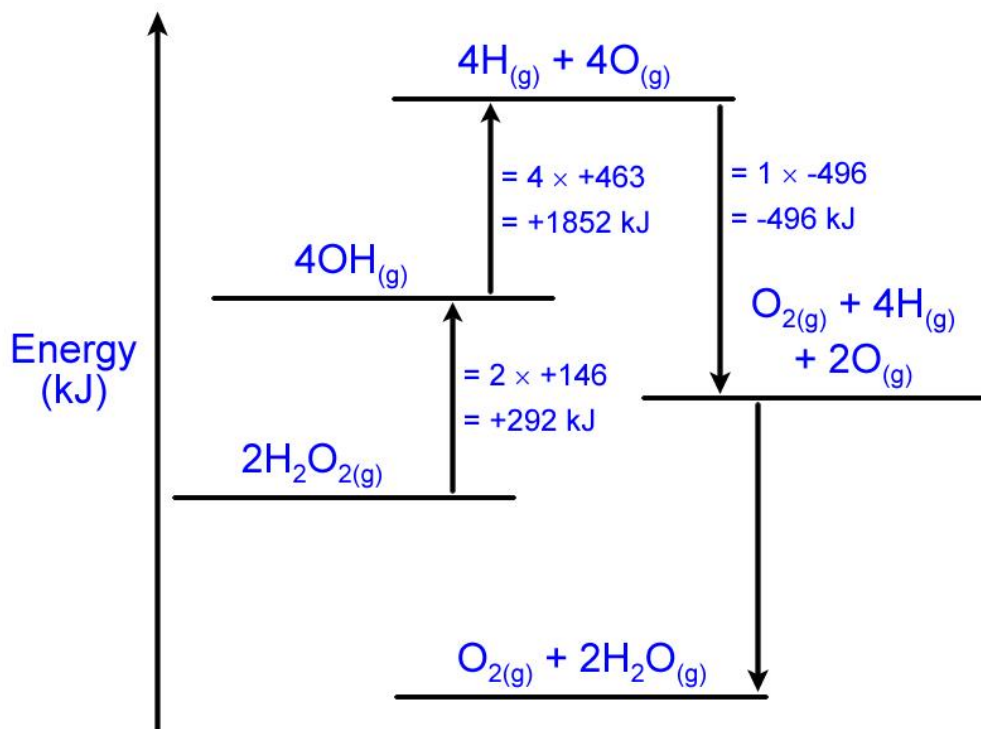
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

O–H = 463 kJ/mol

O=O = 496 kJ/mol



$4 \times \text{O–H}$  bonds are formed creating  $2 \times \text{H}_2\text{O}$  molecules.

## Energy Change Calculations – Example #2:

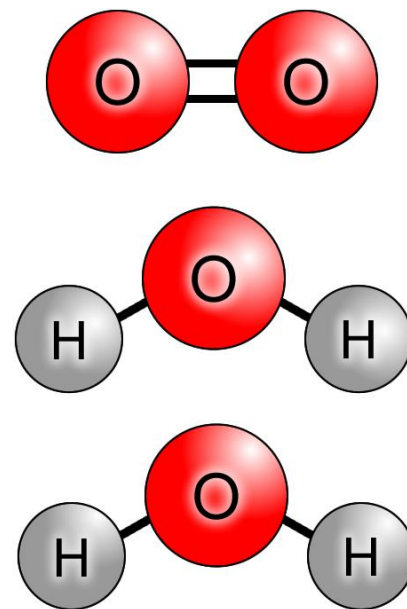
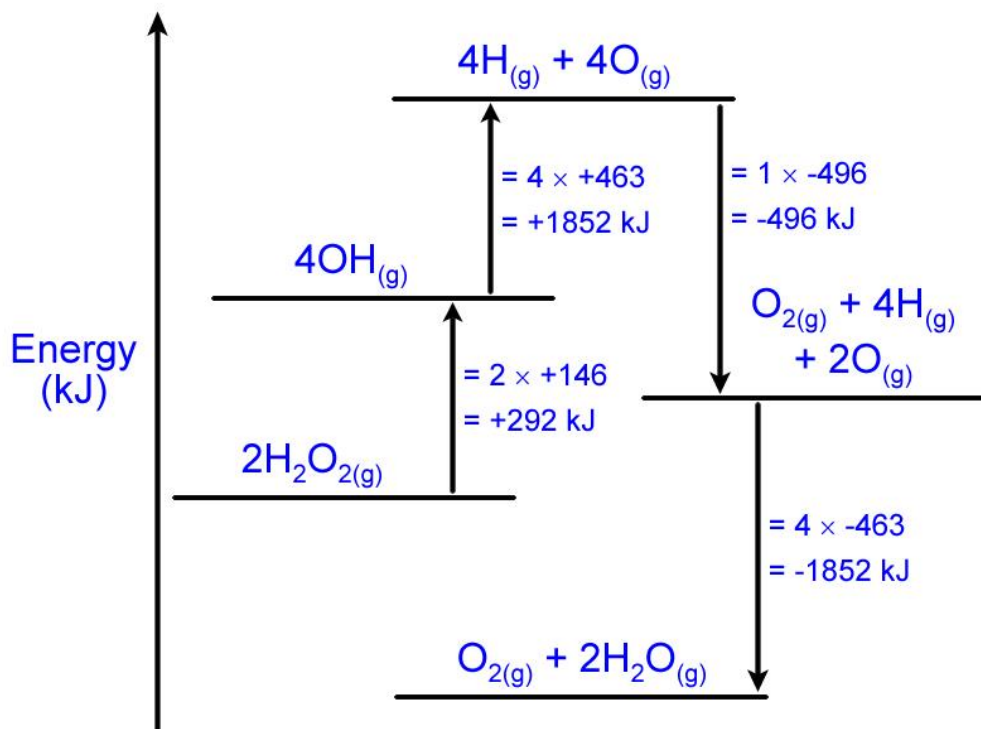
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



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## Energy Change Calculations – Example #2:

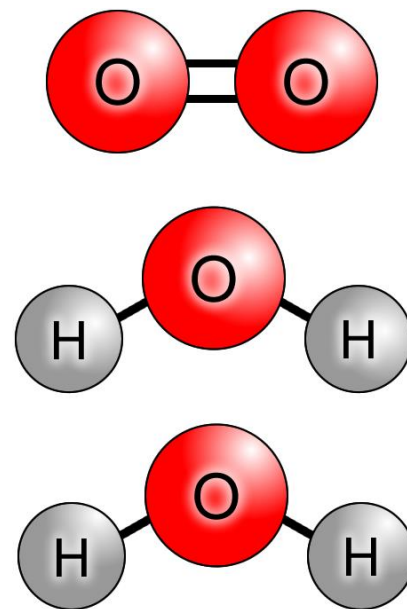
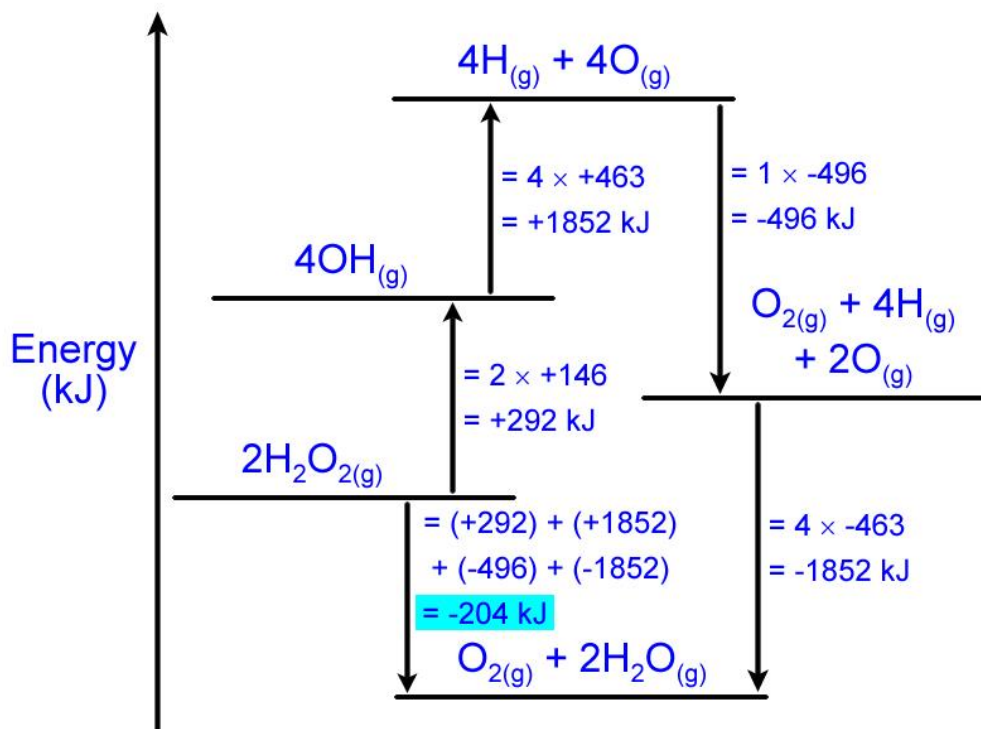
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

O–H = 463 kJ/mol

O=O = 496 kJ/mol



$$(+292) + (+1852) + (-496) + (-1852) = -204 \text{ kJ}$$

## Energy Change Calculations – Example #2:

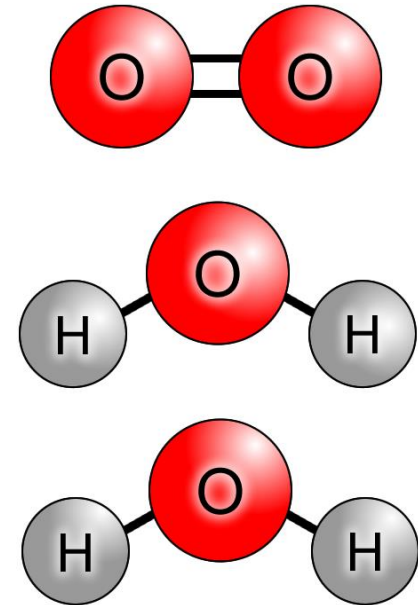
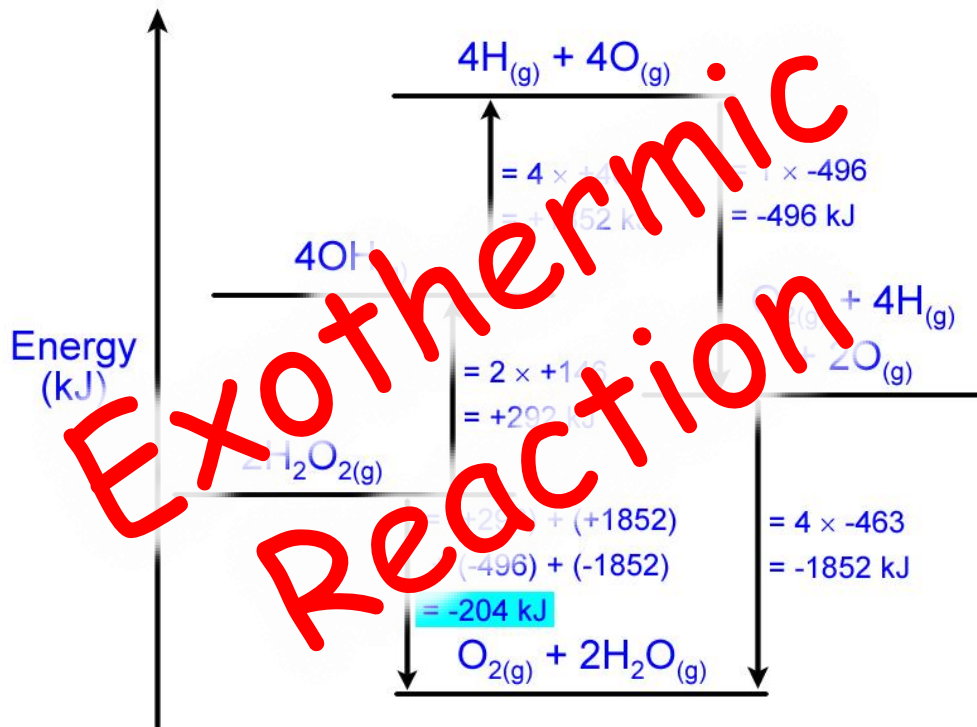
The Decomposition of Hydrogen Peroxide to form Oxygen and Water



O–O = 146 kJ/mol

O–H = 463 kJ/mol

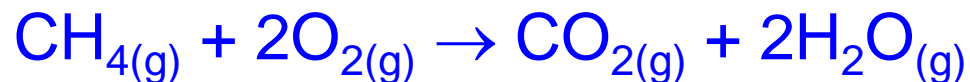
O=O = 496 kJ/mol



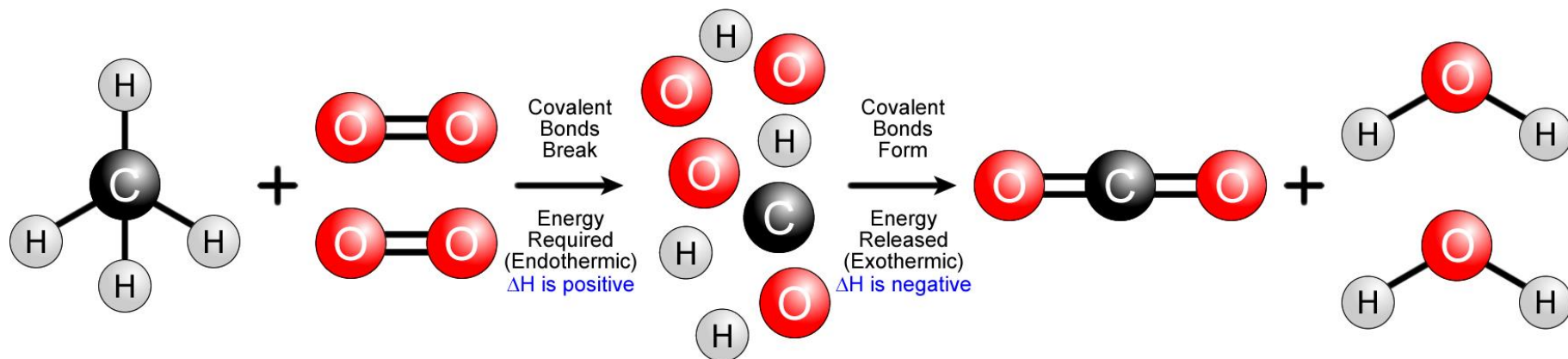
$$(+292) + (+1852) + (-496) + (-1852) = \text{-204 kJ}$$

## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water

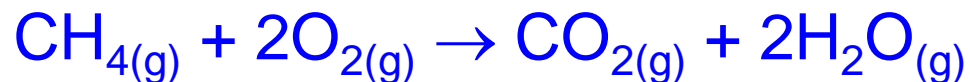


Overview of the Reaction:

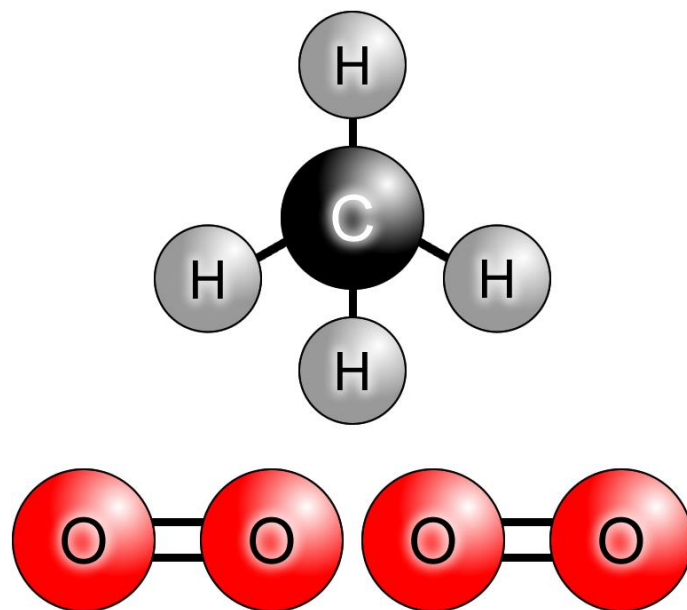
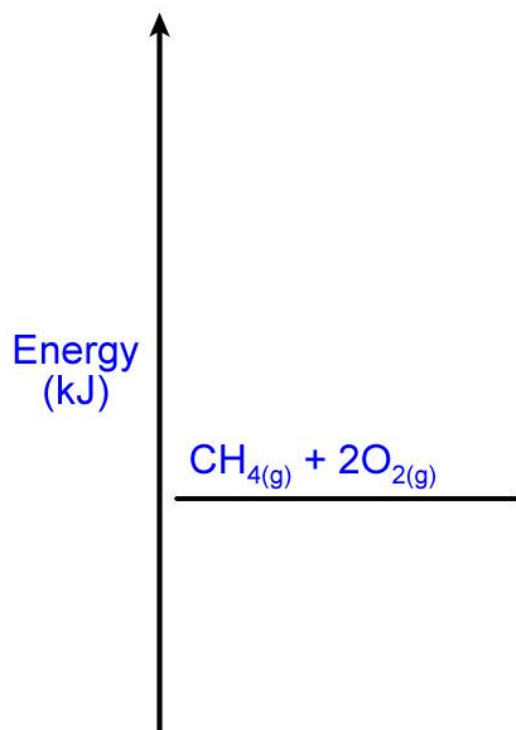


## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water

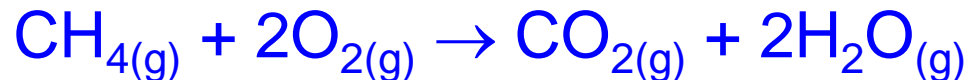


C–H = 412 kJ/mol    O=O = 496 kJ/mol    C=O = 743 kJ/mol    O–H = 463 kJ/mol

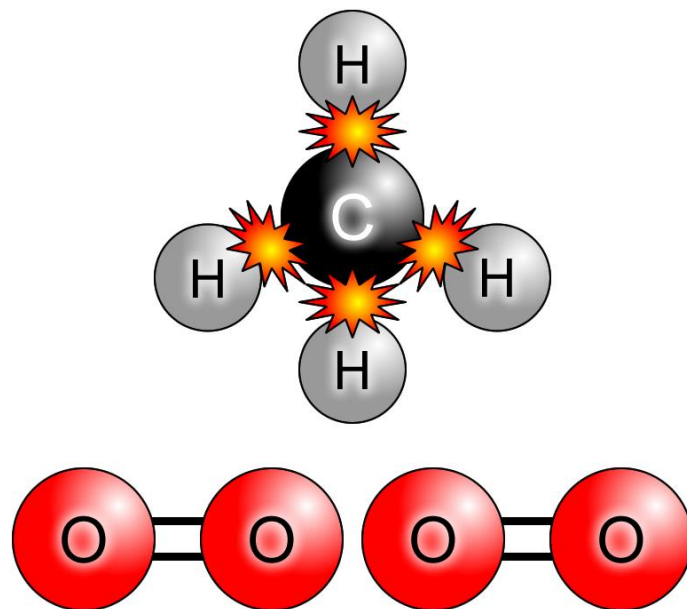
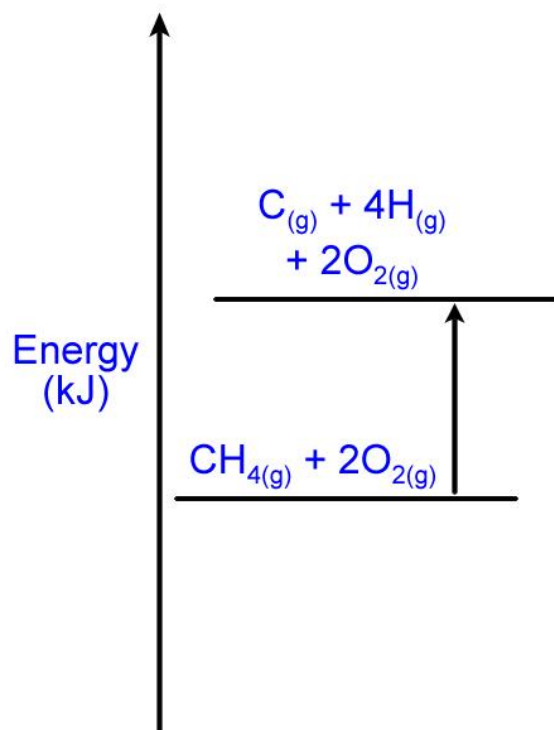


## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



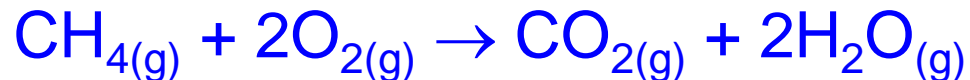
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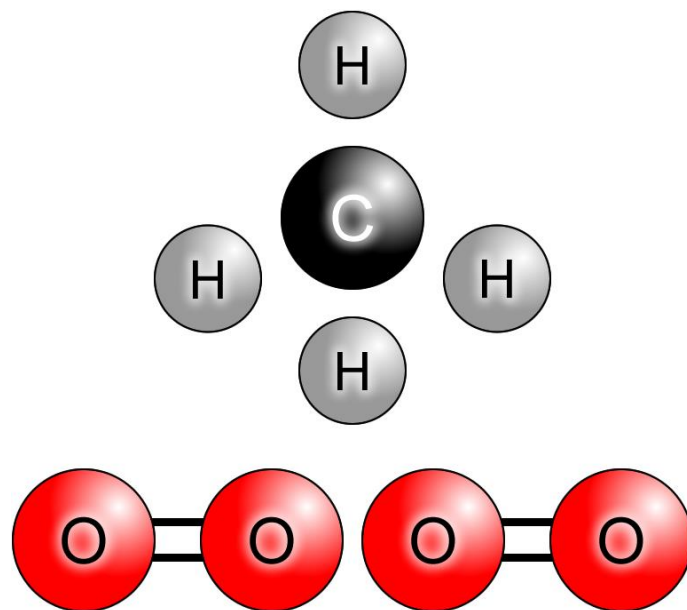
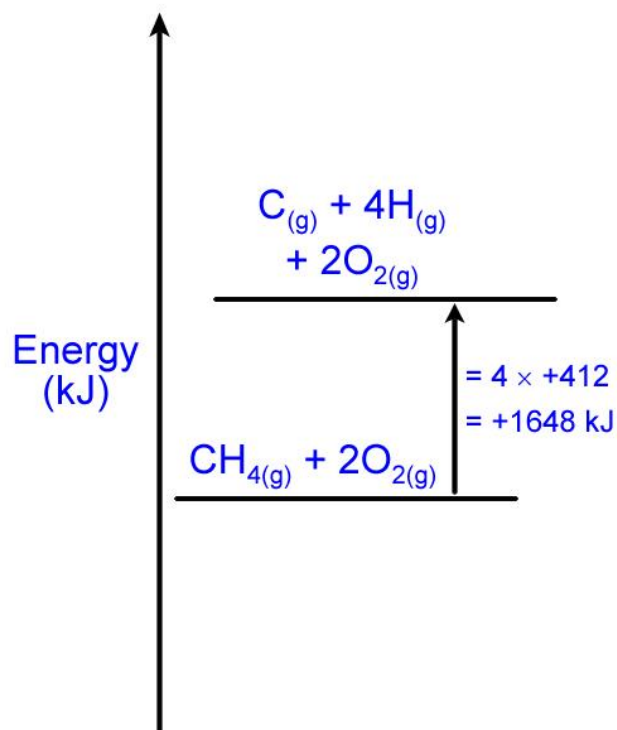
$4 \times \text{C-H}$  bonds are broken forming  $1 \times \text{C}$  atom and  $4 \times \text{H}$  atoms.

## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



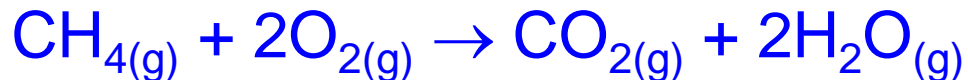
C–H = 412 kJ/mol    O=O = 496 kJ/mol    C=O = 743 kJ/mol    O–H = 463 kJ/mol



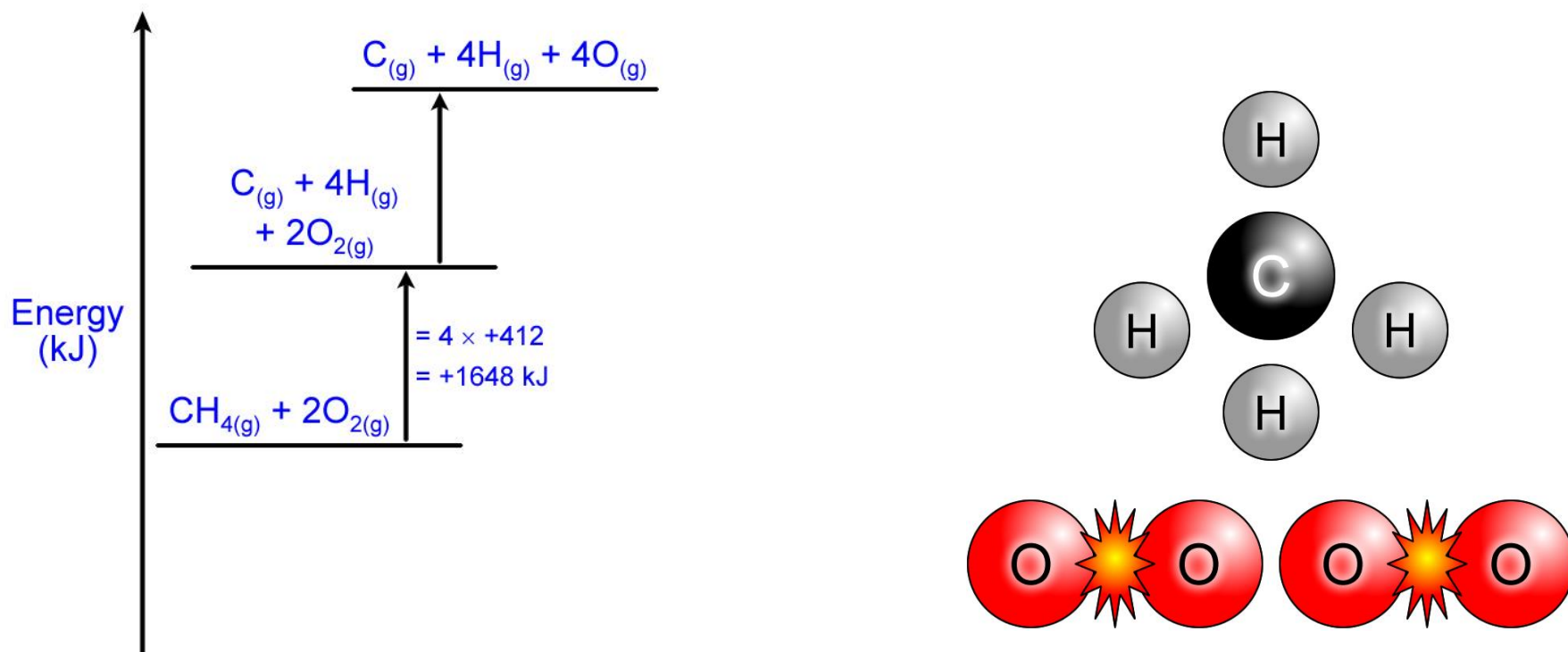
Bond breaking is *endothermic*:  $\Delta H$  for this change is *positive*.

## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



C–H = 412 kJ/mol    O=O = 496 kJ/mol    C=O = 743 kJ/mol    O–H = 463 kJ/mol

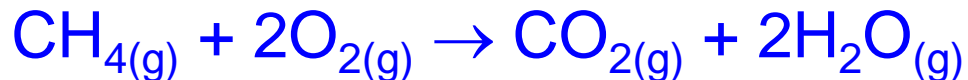


$2 \times \text{O}=\text{O}$  bonds are broken forming  $4 \times \text{O}$  atoms

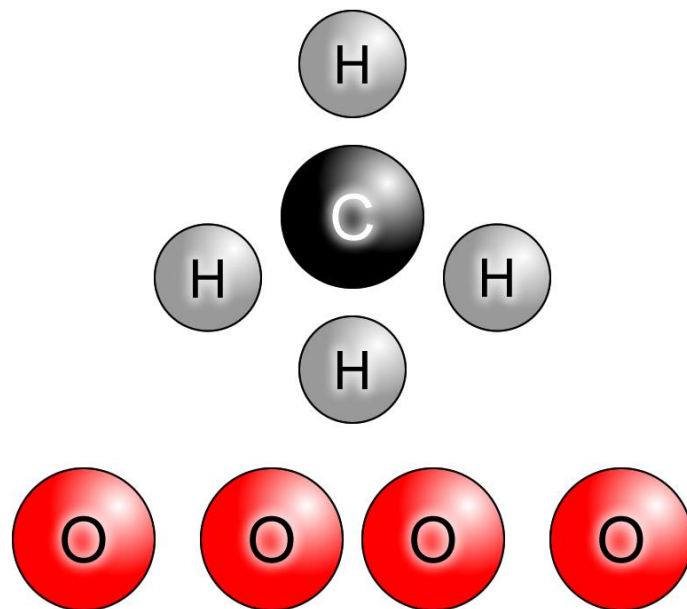
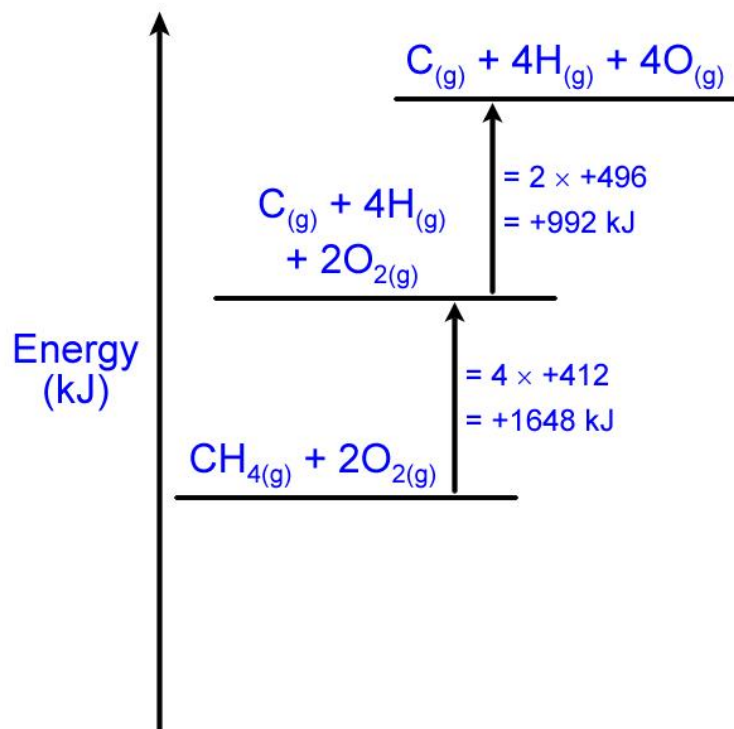


## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



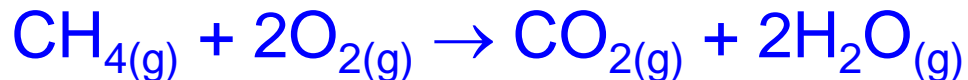
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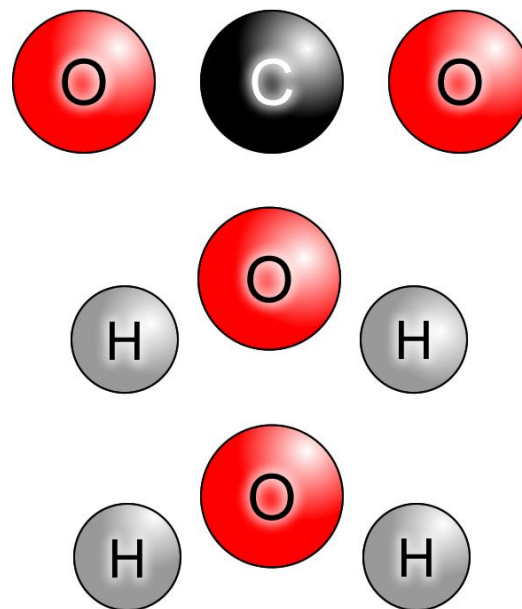
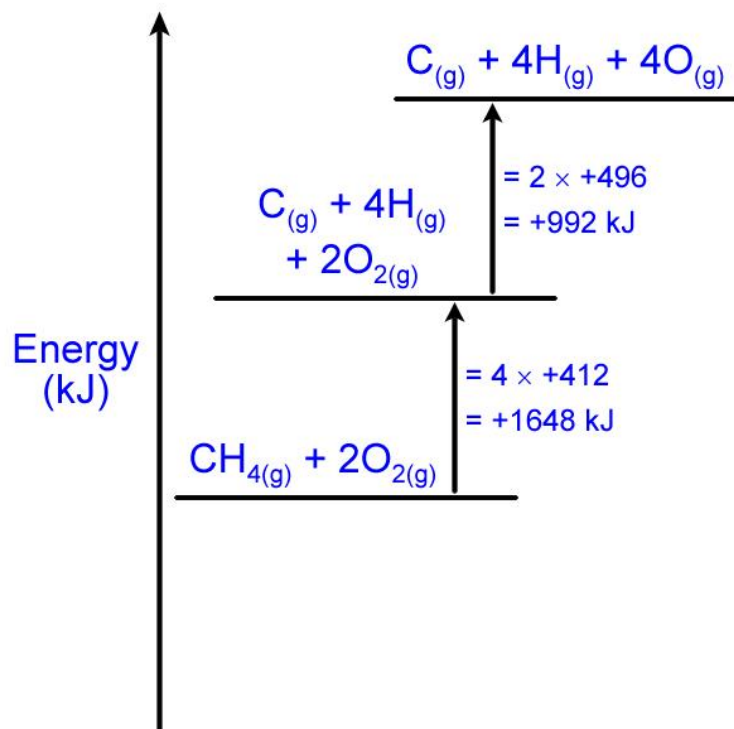
Bond breaking is *endothermic*:  $\Delta H$  for this change is *positive*.

## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water

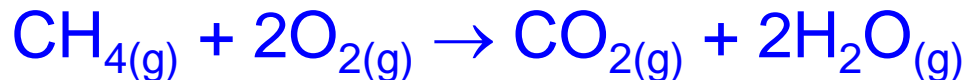


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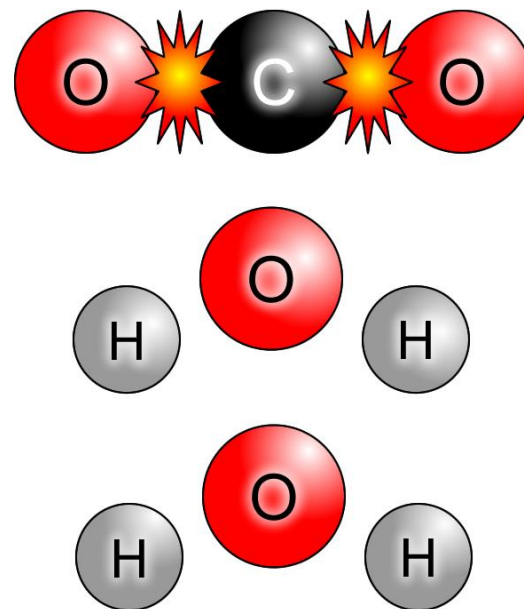
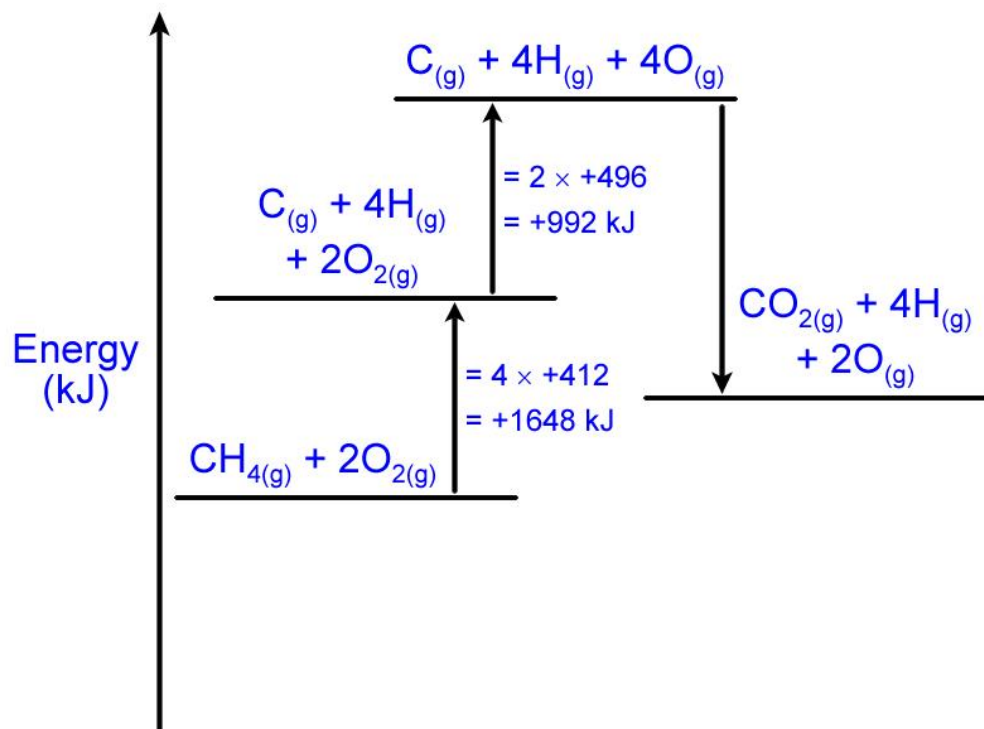


## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



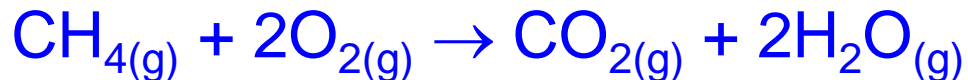
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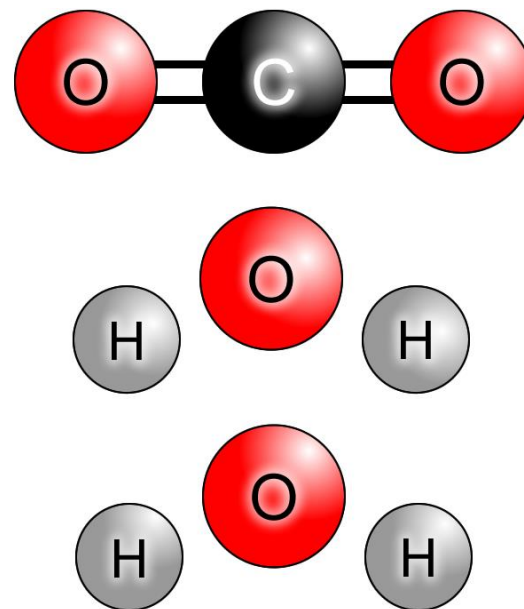
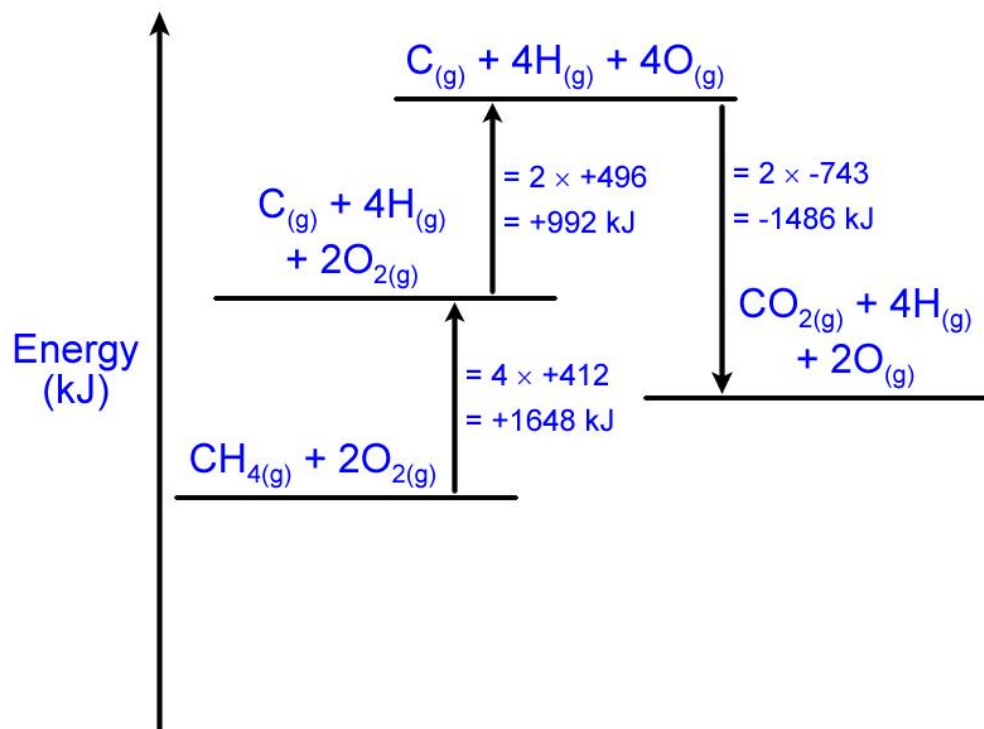
2 × C=O bonds are formed creating 1 × CO<sub>2</sub> molecule.

## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



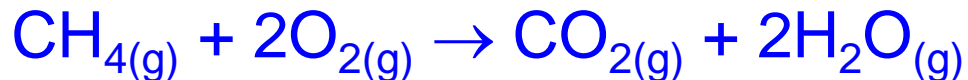
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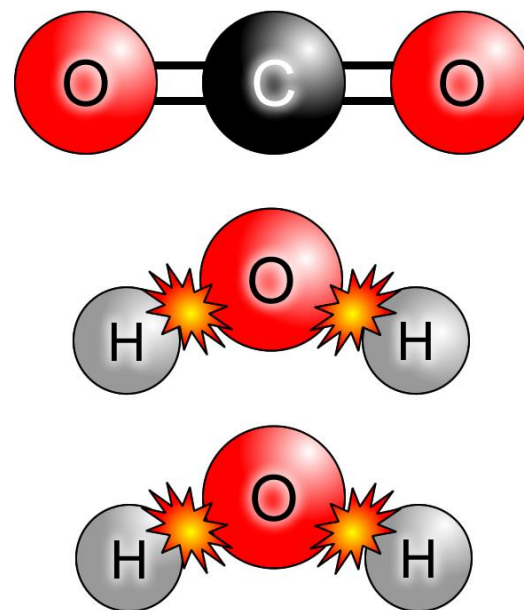
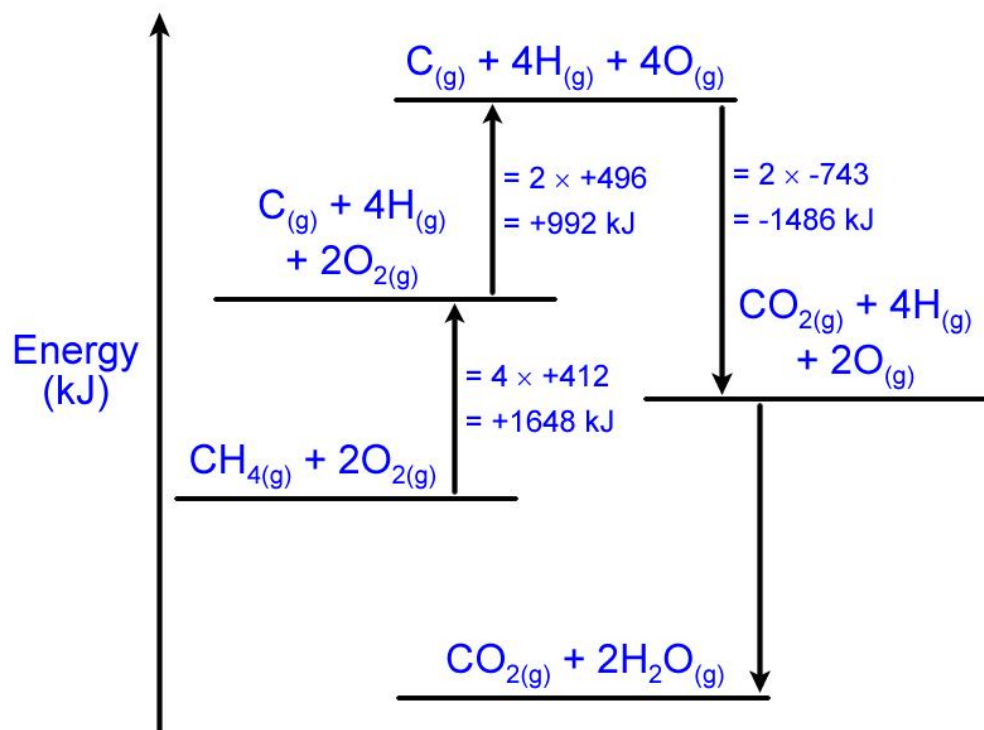
Bond formation is *exothermic*:  $\Delta H$  for this change is *negative*.

## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



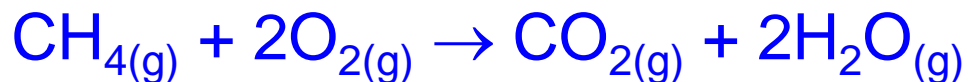
C–H = 412 kJ/mol    O=O = 496 kJ/mol    C=O = 743 kJ/mol    O–H = 463 kJ/mol



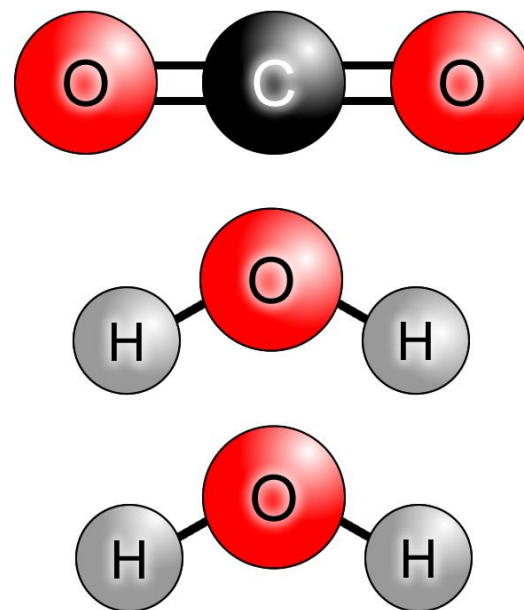
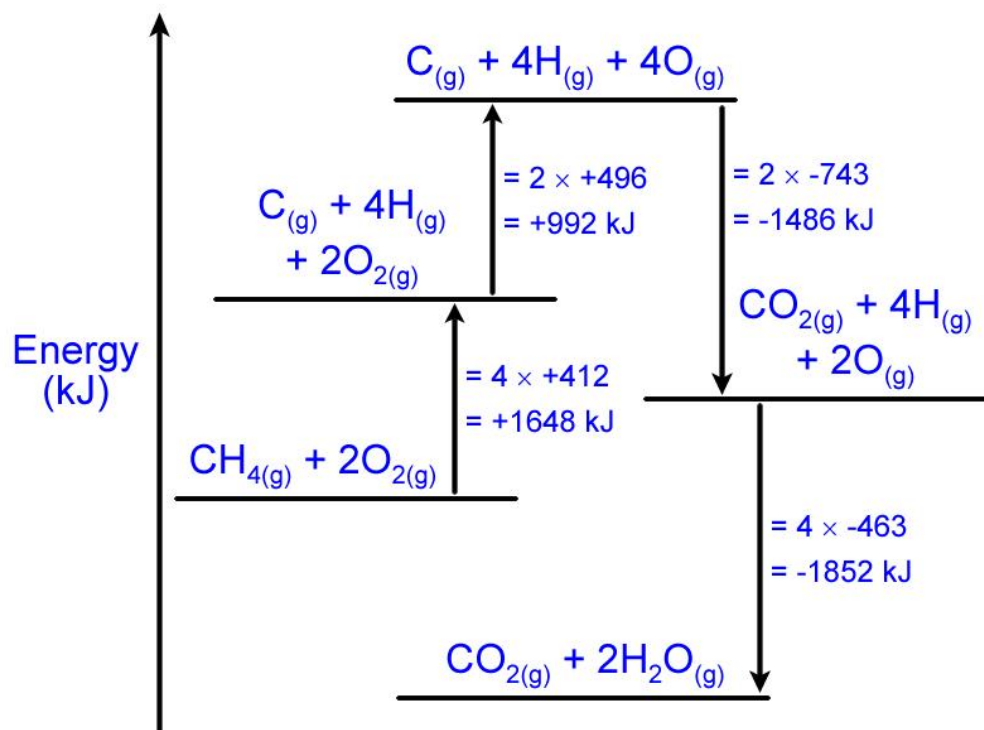
4 × O–H bonds are formed creating 2 ×  $\text{H}_2\text{O}$  molecules.

## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



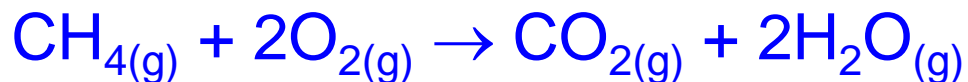
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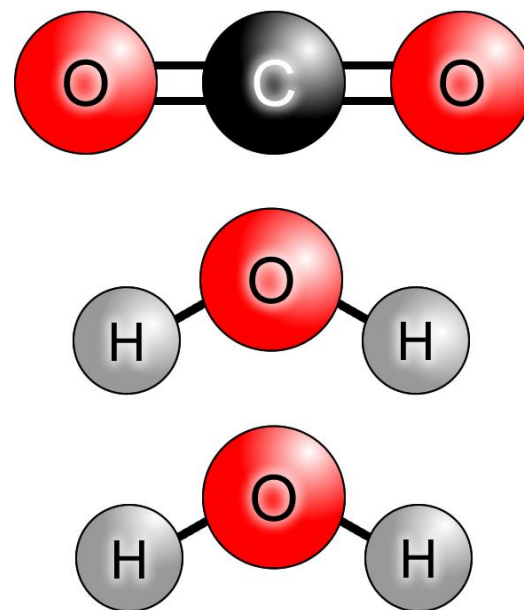
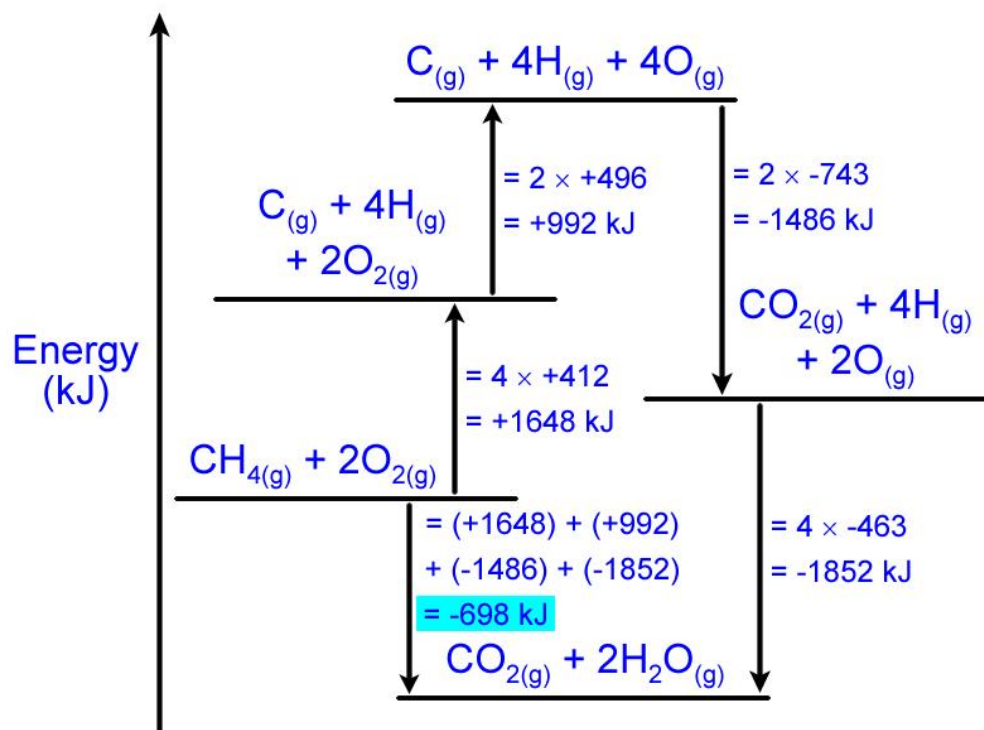
Bond formation is *exothermic*:  $\Delta H$  for this change is *negative*.

## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



C–H = 412 kJ/mol    O=O = 496 kJ/mol    C=O = 743 kJ/mol    O–H = 463 kJ/mol

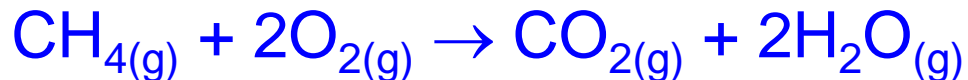


$$(+1648) + (+992) + (-1486) + (-1852) = -698 \text{ kJ}$$

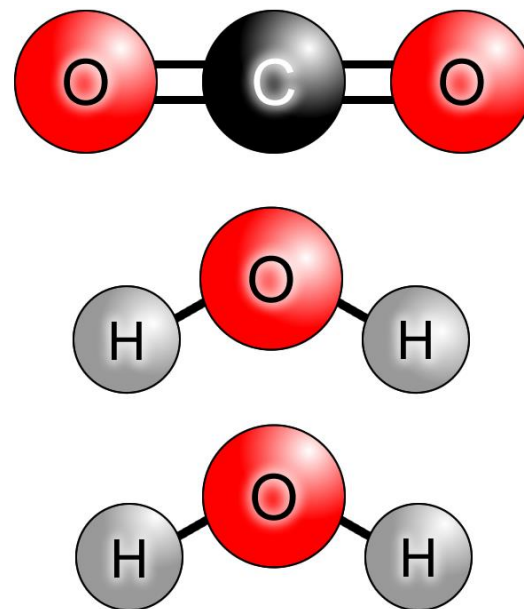
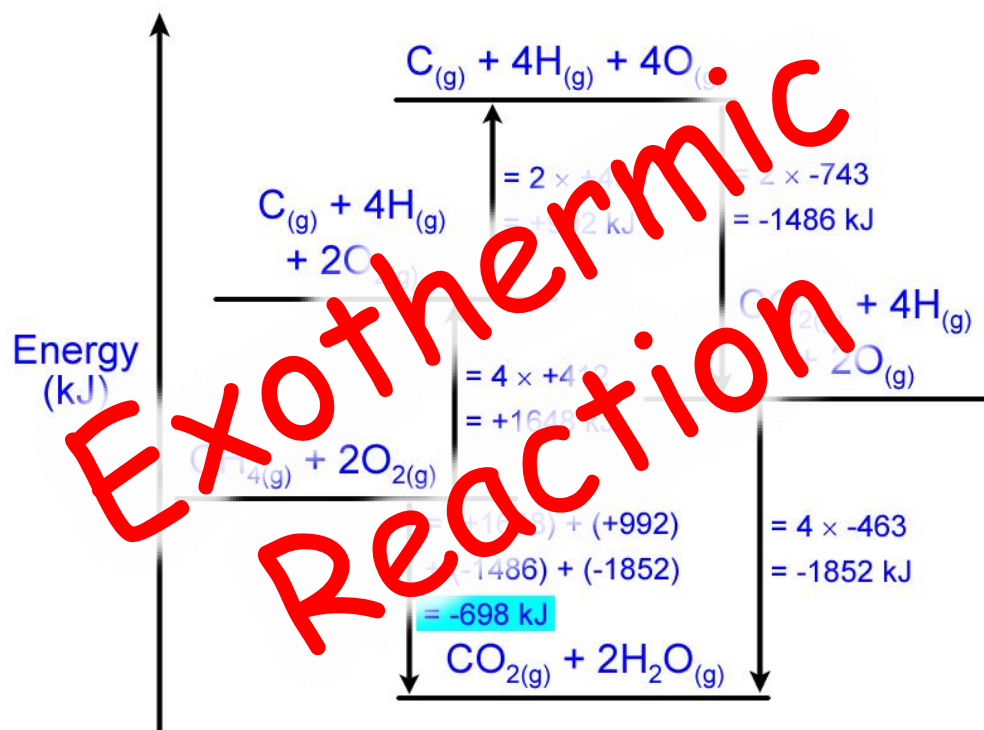


## Energy Change Calculations – Example #3:

The Combustion of Methane forming Carbon Dioxide and Water



C–H = 412 kJ/mol    O=O = 496 kJ/mol    C=O = 743 kJ/mol    O–H = 463 kJ/mol



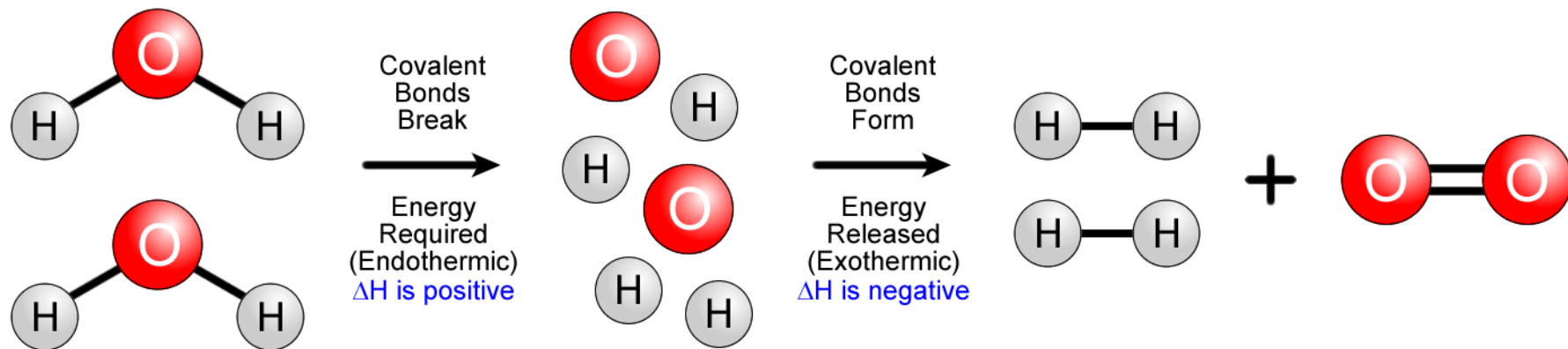
$$(+1648) + (+992) + (-1486) + (-1852) = -698 \text{ kJ}$$

## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



Overview of the Reaction:

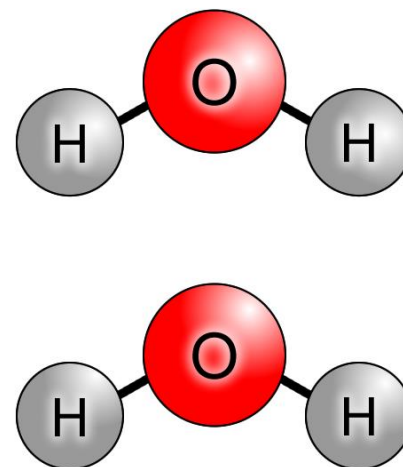
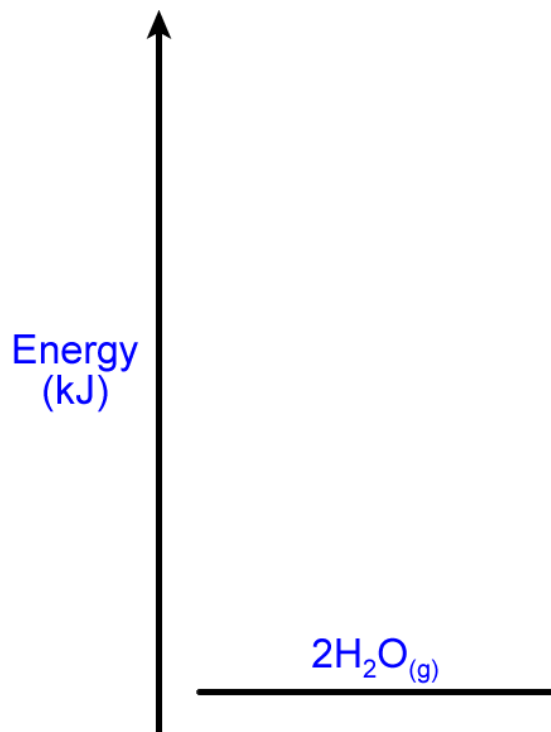


## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



O–H = 463 kJ/mol    O=O = 496 kJ/mol    H–H = 436 kJ/mol

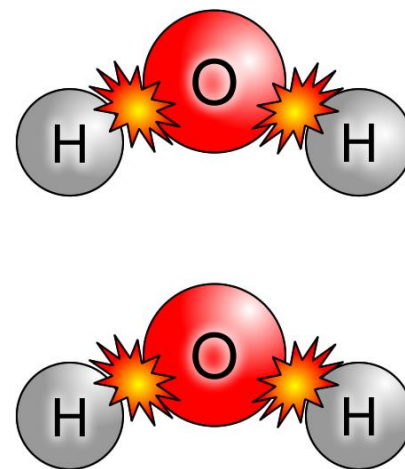
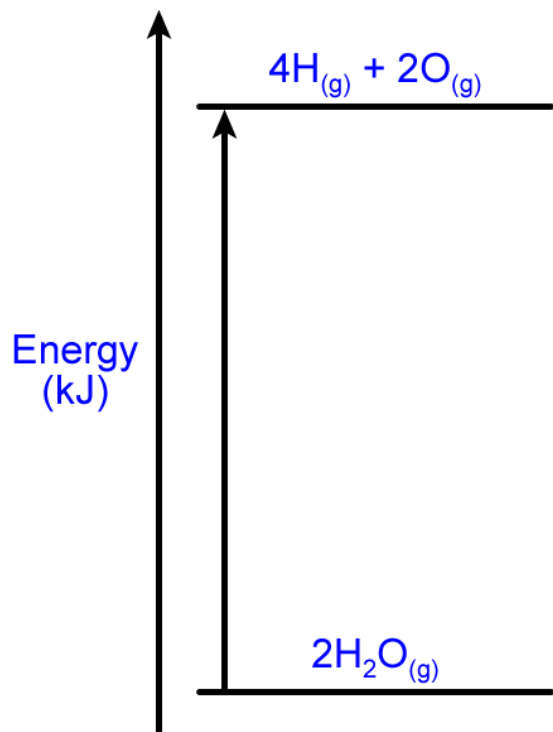


## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



O–H = 463 kJ/mol    O=O = 496 kJ/mol    H–H = 436 kJ/mol



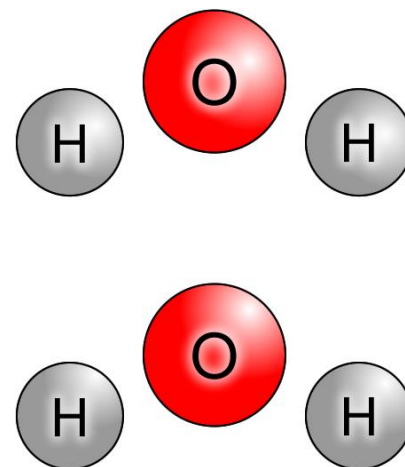
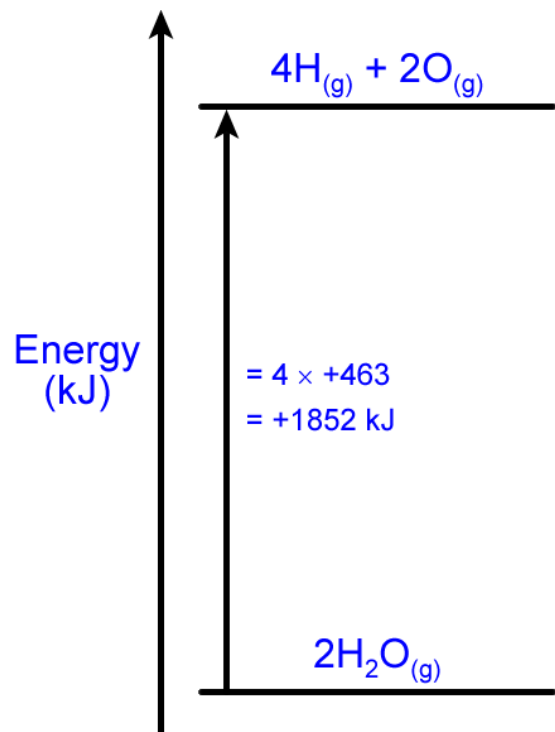
$4 \times \text{O-H}$  bonds are broken forming  $2 \times \text{O}$  and  $4 \times \text{H}$  atoms.

## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



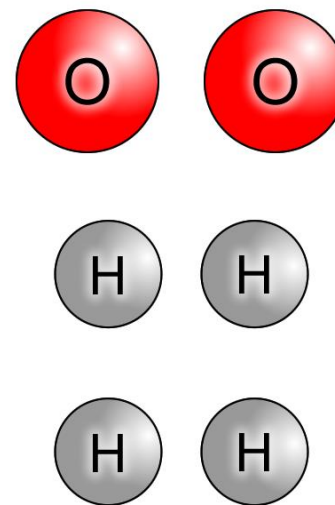
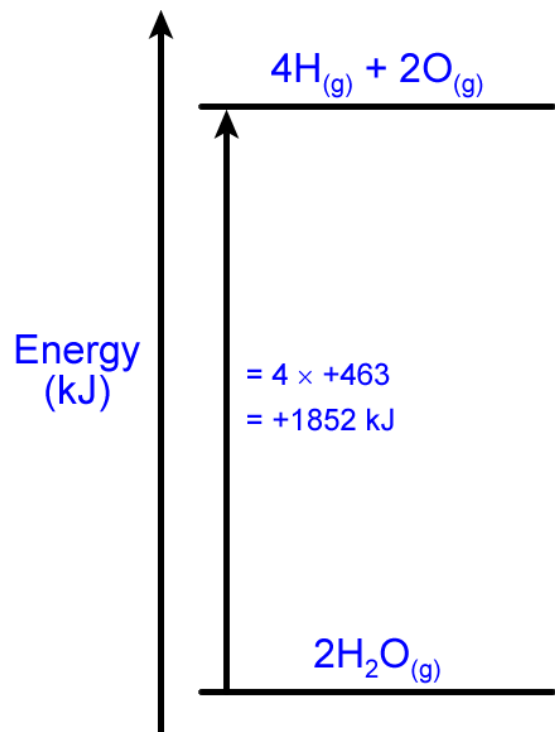
O–H = 463 kJ/mol    O=O = 496 kJ/mol    H–H = 436 kJ/mol



Bond breaking is *endothermic*:  $\Delta H$  for this change is *positive*.

## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



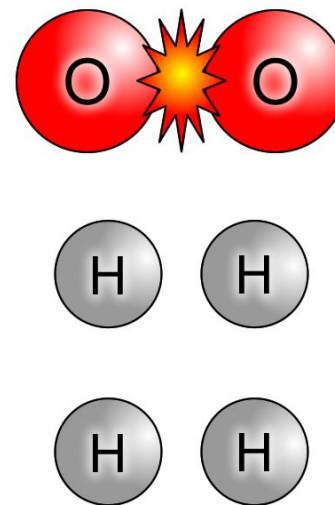
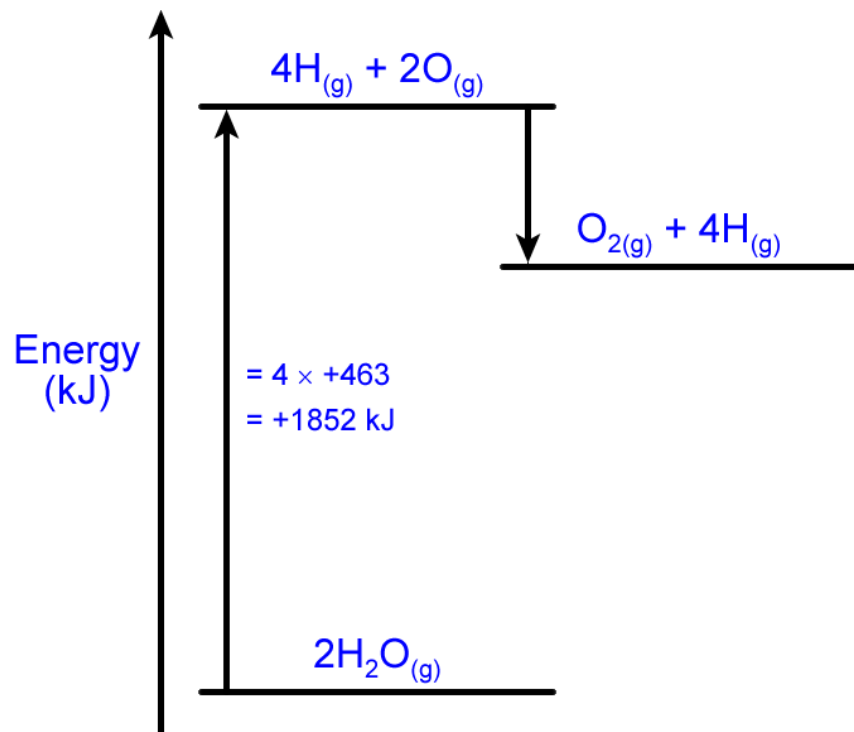
Bond breaking is *endothermic*:  $\Delta\text{H}$  for this change is *positive*.

## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



O–H = 463 kJ/mol    O=O = 496 kJ/mol    H–H = 436 kJ/mol



1  $\times$  O=O bond is formed creating 1  $\times$   $\text{O}_2$  molecule.

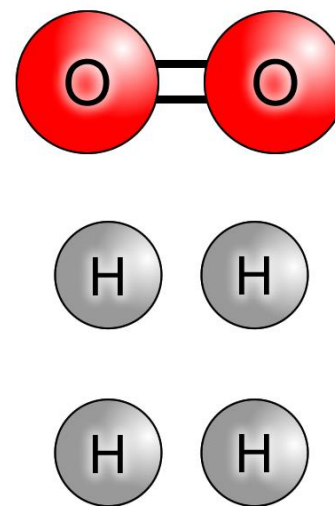
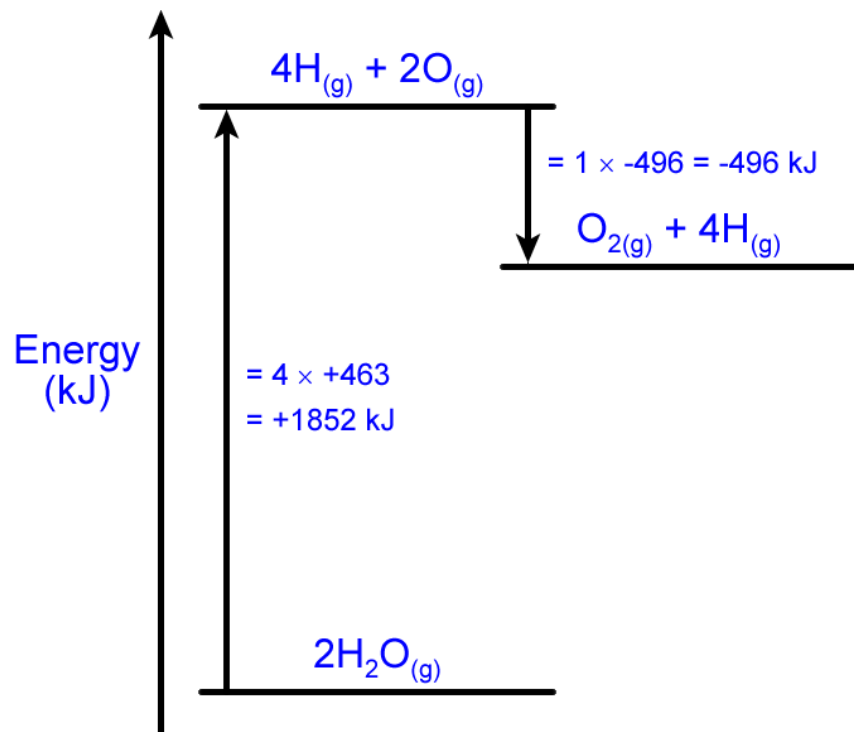


## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



O–H = 463 kJ/mol    O=O = 496 kJ/mol    H–H = 436 kJ/mol



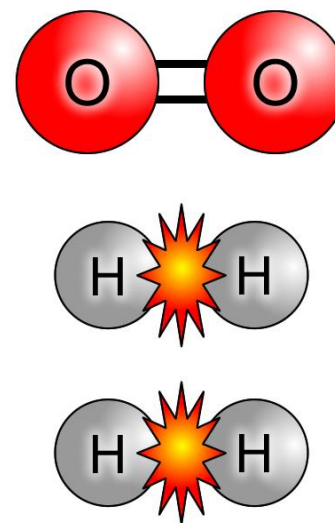
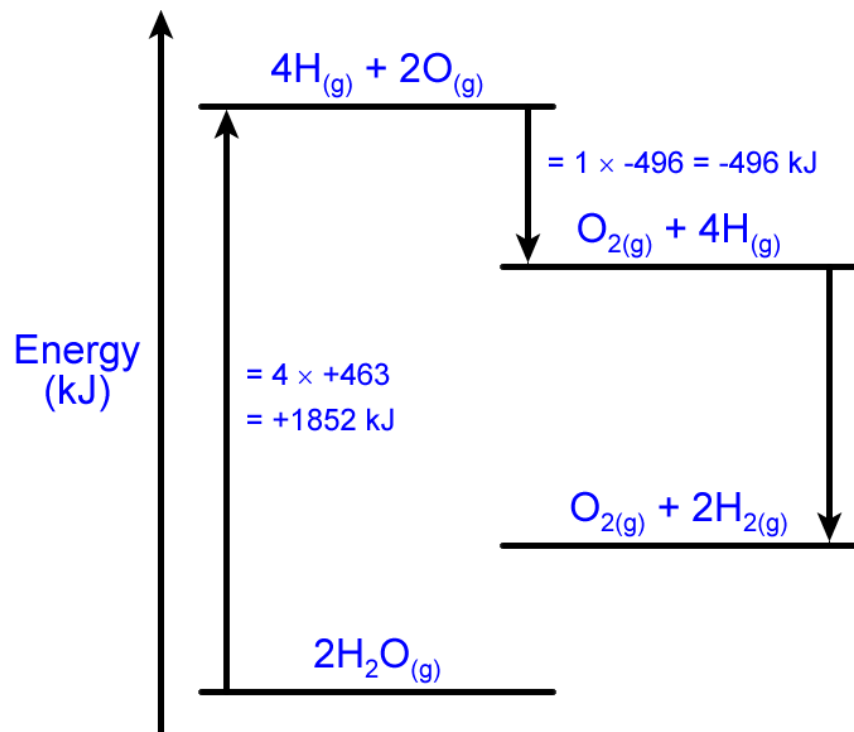
Bond formation is *exothermic*:  $\Delta\text{H}$  for this change is *negative*.

## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



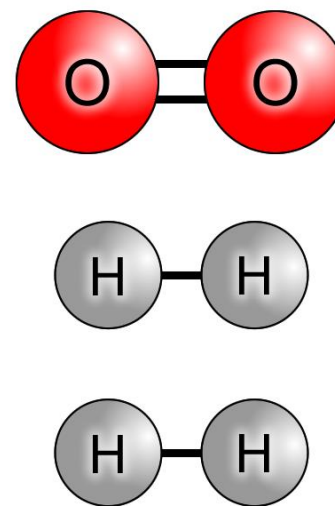
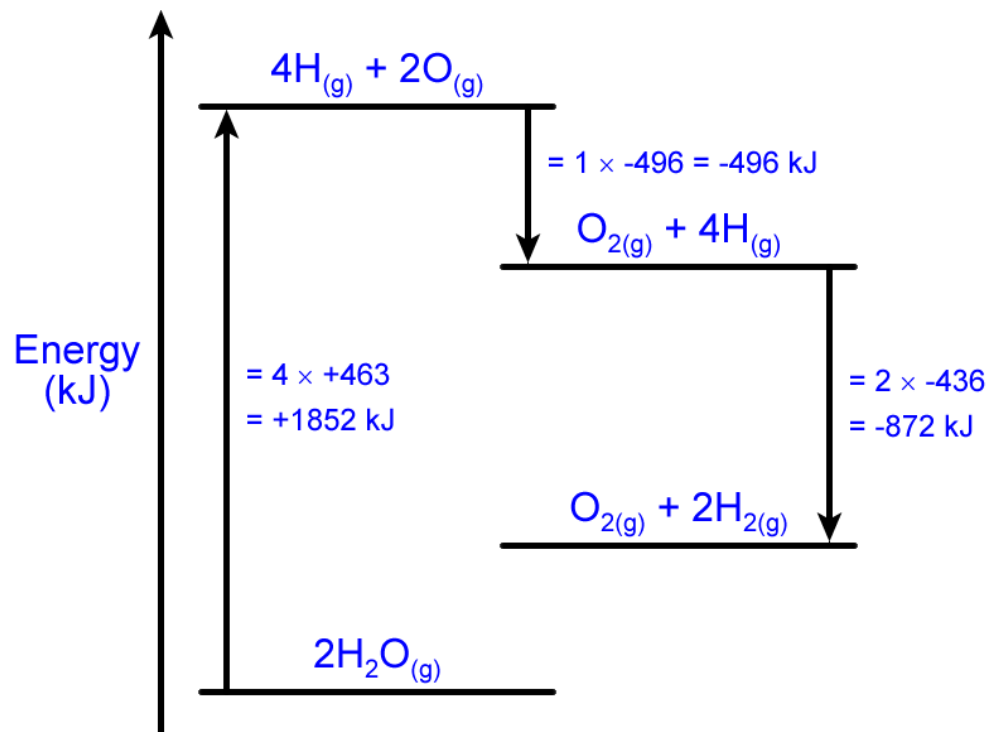
O–H = 463 kJ/mol    O=O = 496 kJ/mol    H–H = 436 kJ/mol



$2 \times \text{H-H}$  bonds are formed creating  $2 \times \text{H}_2$  molecules.

## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



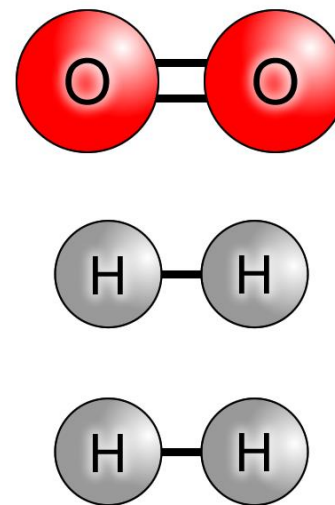
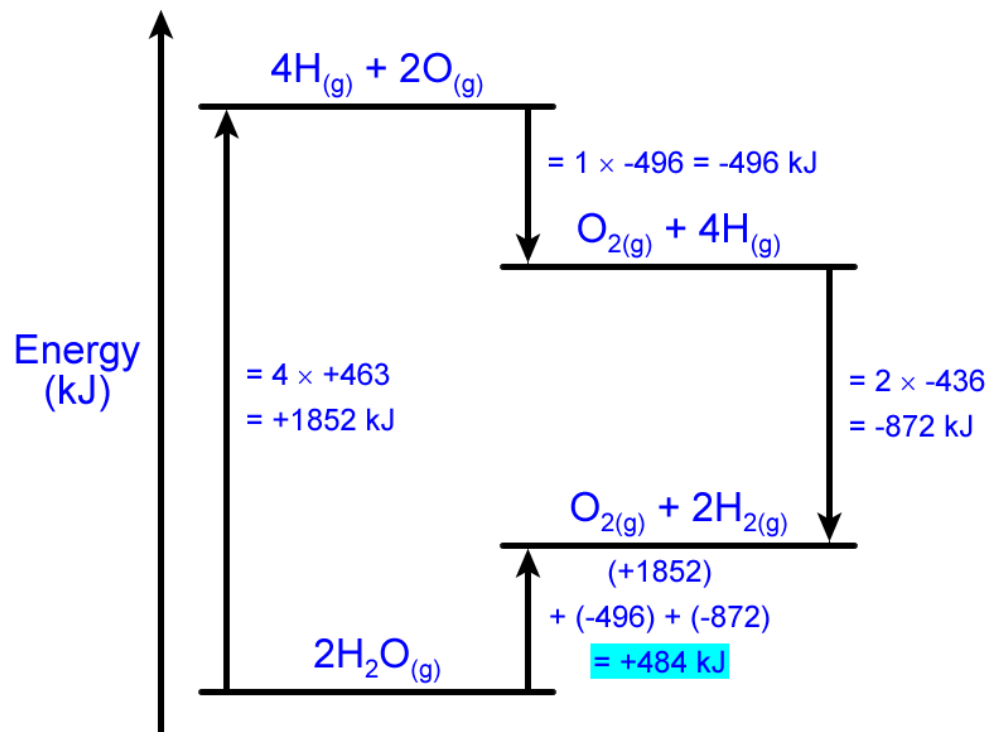
Bond formation is *exothermic*:  $\Delta\text{H}$  for this change is *negative*.

## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



O–H = 463 kJ/mol    O=O = 496 kJ/mol    H–H = 436 kJ/mol



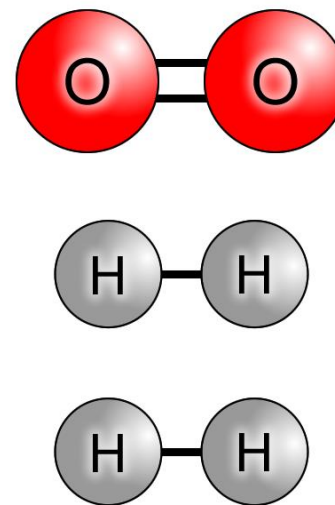
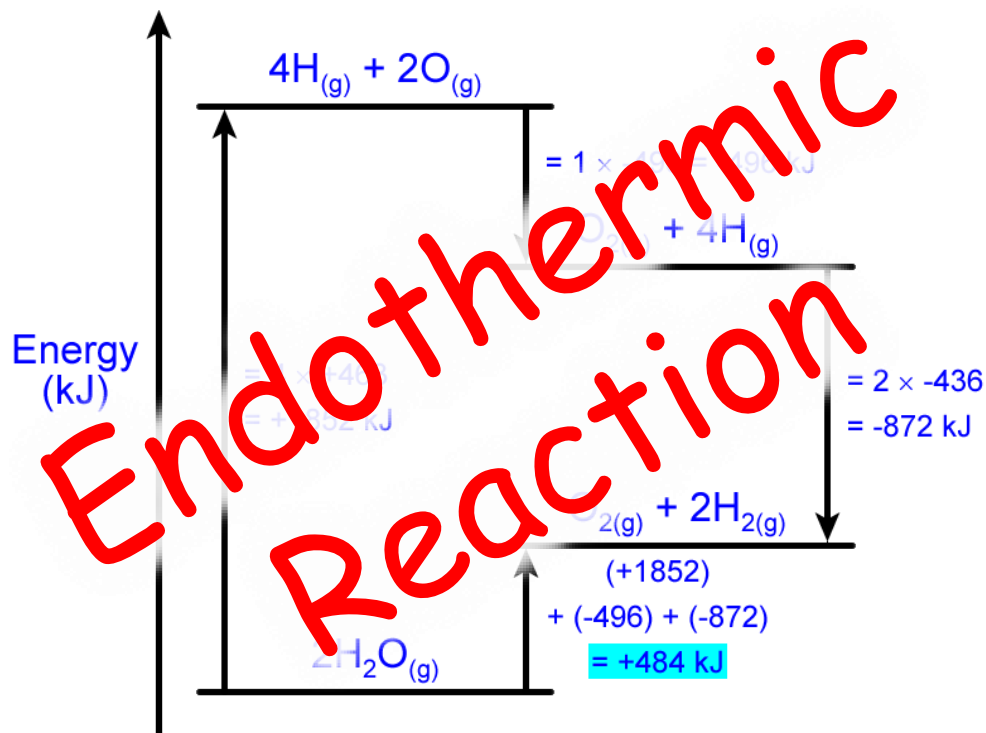
$$(+1852) + (-496) + (-872) = +484 \text{ kJ}$$

## Energy Change Calculations – Example #4:

The Electrolysis of Water to form Hydrogen and Oxygen



O–H = 463 kJ/mol    O=O = 496 kJ/mol    H–H = 436 kJ/mol



$$(+1852) + (-496) + (-872) = +484 \text{ kJ}$$

# Energy from Chemicals



Is a double covalent bond *twice as strong* as a single covalent bond?

# Energy from Chemicals

- Study the bond energies given below:



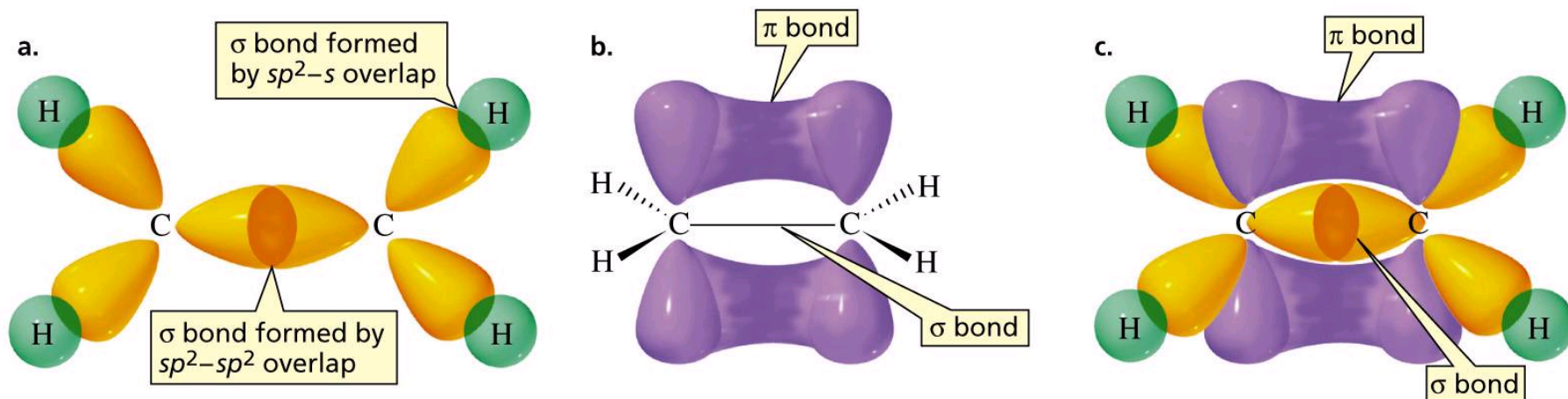
- Is a double covalent bond **twice as strong** as a single covalent bond?
  - For **carbon**,  $2 \times \text{C}-\text{C} = 2 \times 348 = 696 \text{ kJ}$   
 $696 \text{ kJ} \neq 612 \text{ kJ}$   
 $696 \text{ kJ} > 612 \text{ kJ}$
  - For **oxygen**,  $2 \times \text{O}-\text{O} = 2 \times 146 = 292 \text{ kJ}$   
 $292 \text{ kJ} \neq 496 \text{ kJ}$   
 $292 \text{ kJ} < 496 \text{ kJ}$





# Energy from Chemicals

- Why is a double covalent bond not twice as strong as a single covalent bond?



- Because there are different types of covalent bonds called  $\sigma$ -bonds (sigma bonds) and  $\pi$ -bonds (pi bonds).
  - $\sigma$ -bonds and  $\pi$ -bonds have different strengths.

# Energy from Chemicals



The enthalpy change calculations seem to follow the same pattern. Can I automate the calculation in a *spreadsheet*?

# Alkane Combustion Spreadsheet

- Alkanes are **hydrocarbons** with the general formula  $C_nH_{2n+2}$ .
- Alkanes react with oxygen to produce carbon dioxide and water.
  - Design a spreadsheet that will, when the number of carbon atoms in the molecule is keyed in, automatically calculate the energy change for the combustion of one mole of an alkane.

Enthalpy Change of Combustion Spreadsheet for Alkanes - Microsoft Excel non-commercial use

**Spreadsheet to Calculate the Enthalpy Change of Combustion for One Mole of an Alkane**

**Definition:** The enthalpy change of combustion is the energy change produced when one mole of a chemical completely reacts with oxygen under standard conditions.

**Instructions:**

- Choose an alkane by typing a whole number into the cell that is highlighted in blue and press "Return" on your keyboard.
- The spreadsheet will generate the balanced chemical equation for the complete combustion of one mole of the alkane.
- Using average bond energies, the spreadsheet will calculate the enthalpy change of combustion for one mole of the alkane.

**Average Bond Energies:**

C-C = 348 kJ mol<sup>-1</sup>      C-H = 412 kJ mol<sup>-1</sup>      O=O = 496 kJ mol<sup>-1</sup>  
 C=O = 743 kJ mol<sup>-1</sup>      O-H = 463 kJ mol<sup>-1</sup>

**Balanced Chemical Equation:**

$C_2 + H_4 + 2 O_{2(g)} \rightarrow 1 CO_{2(g)} + 2 H_2O_{(l)}$

**Names of the First Ten Straight Chain Alkanes**

CH<sub>4</sub> = Methane  
 C<sub>2</sub>H<sub>6</sub> = Ethane  
 C<sub>3</sub>H<sub>8</sub> = Propane  
 C<sub>4</sub>H<sub>10</sub> = Butane  
 C<sub>5</sub>H<sub>12</sub> = Pentane  
 C<sub>6</sub>H<sub>14</sub> = Hexane  
 C<sub>7</sub>H<sub>16</sub> = Heptane  
 C<sub>8</sub>H<sub>18</sub> = Octane  
 C<sub>9</sub>H<sub>20</sub> = Nonane  
 C<sub>10</sub>H<sub>22</sub> = Decane

**Energy Change During Covalent Bond Breaking:**

0	x	C-C	=	0	x	+348	=	0
4	x	C-H	=	4	x	+412	=	+1648
2	x	O=O	=	2	x	+496	=	+992
							Total =	<b>+2640</b>

**Energy Change During Covalent Bond Formation:**

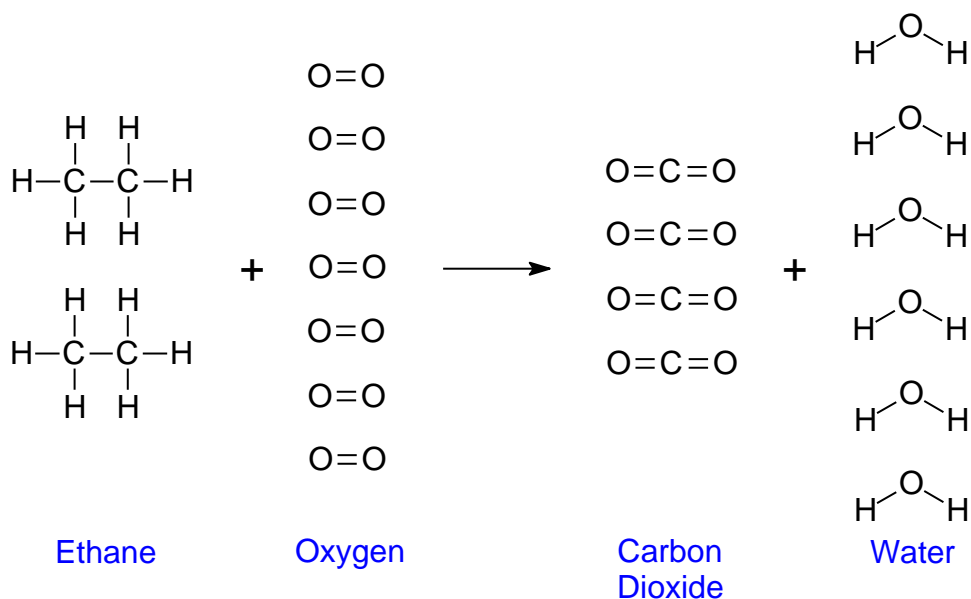
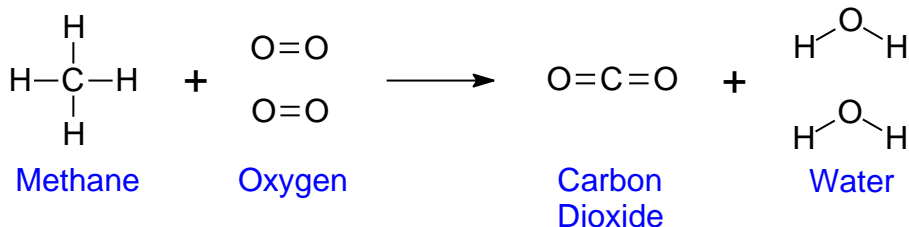
2	x	C=O	=	2	x	-743	=	-1486
4	x	O-H	=	4	x	-463	=	-1852
							Total =	<b>-3338</b>

**Overall Energy Change for the Reaction:**

=	+2640	+	-3338
=	<b>-698</b>	kJ mol <sup>-1</sup>	



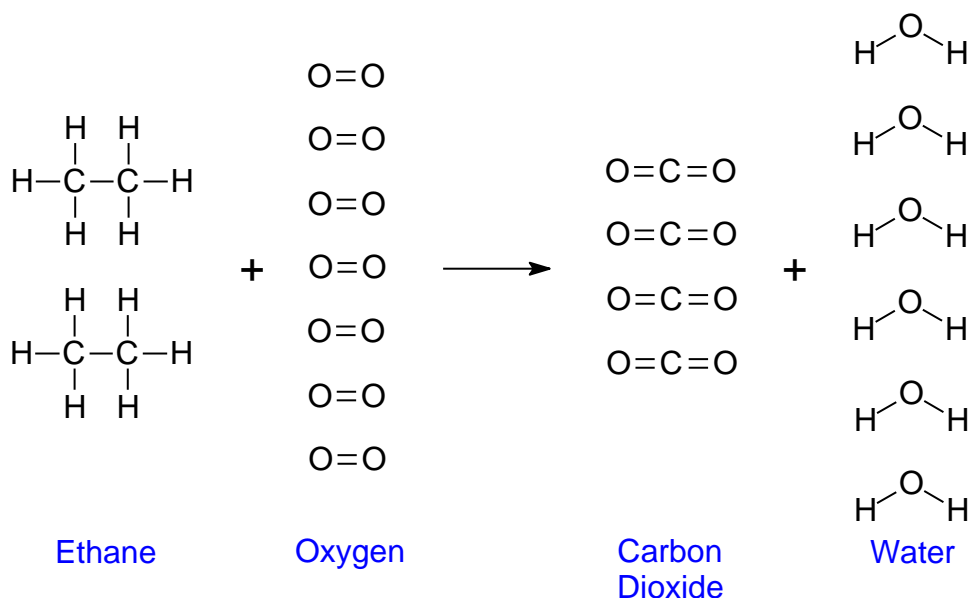
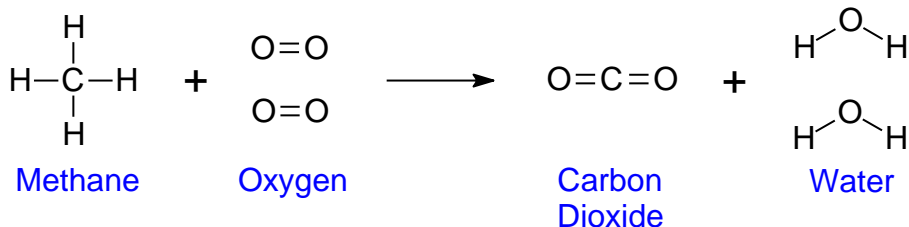
# Alkane Combustion Spreadsheet



- Consider the combustion of **methane** and the combustion of **ethane** shown on the left.
- Let the number of carbon atoms = **n**.
- In relation to **n**, how many C–C bonds break?
- In relation to **n**, how many C–H bonds break?
- In relation to **n**, how many O=O bonds break?
- In relation to **n**, how many C=O bonds form?
- In relation to **n**, how many O–H bonds form?



# Alkane Combustion Spreadsheet



- Consider the combustion of **methane** and the combustion of **ethane** shown on the left.
- Let the number of carbon atoms =  $n$ .
- In relation to  $n$ , how many C–C bonds break?  $n - 1$
- In relation to  $n$ , how many C–H bonds break?  $2n + 2$
- In relation to  $n$ , how many O=O bonds break?  $1.5n + 0.5$
- In relation to  $n$ , how many C=O bonds form?  $2n$
- In relation to  $n$ , how many O–H bonds form?  $2n + 2$



# Alkane Combustion Spreadsheet

- In summary, for an alkane with the general formula  $C_nH_{2n+2}$ :
  - The number of C–C bonds broken =  $n - 1$ .
  - The number of C–H bonds broken =  $2n + 2$ .
- The number of O=O bonds broken =  $1.5n + 0.5$ .
  - The number of C=O bonds formed =  $2n$ .
  - The number of O–H bonds formed =  $2n + 2$ .
  - Average C–C bond energy =  $348 \text{ kJ / mol}$ .
  - Average C–H bond energy =  $411 \text{ kJ / mol}$ .
  - Average O=O bond energy =  $498 \text{ kJ / mol}$ .
  - Average C=O bond energy =  $804 \text{ kJ / mol}$ .
  - Average O–H bond energy =  $463 \text{ kJ / mol}$ .
- Remember, **bond breaking** is endothermic,  $\Delta H$  is positive, while **bond formation** is exothermic,  $\Delta$  is negative.



# Alkane Combustion Spreadsheet

- How do your results compare to the experimental data?

Name of Straight Alkane	Formula of Straight Chain Alkane	Physical State at Room Temperature and Pressure	Standard Molar Enthalpy Change of Combustion at 298 K / kJ/mol
Methane	CH <sub>4</sub>	Gas	-890
Ethane	C <sub>2</sub> H <sub>6</sub>	Gas	-1560
Propane	C <sub>3</sub> H <sub>8</sub>	Gas	-2220
Butane	C <sub>4</sub> H <sub>10</sub>	Gas	-2877
Pentane	C <sub>5</sub> H <sub>12</sub>	Liquid	-3509
Hexane	C <sub>6</sub> H <sub>14</sub>	Liquid	-4163

- Identify any possible sources of error in the calculations.





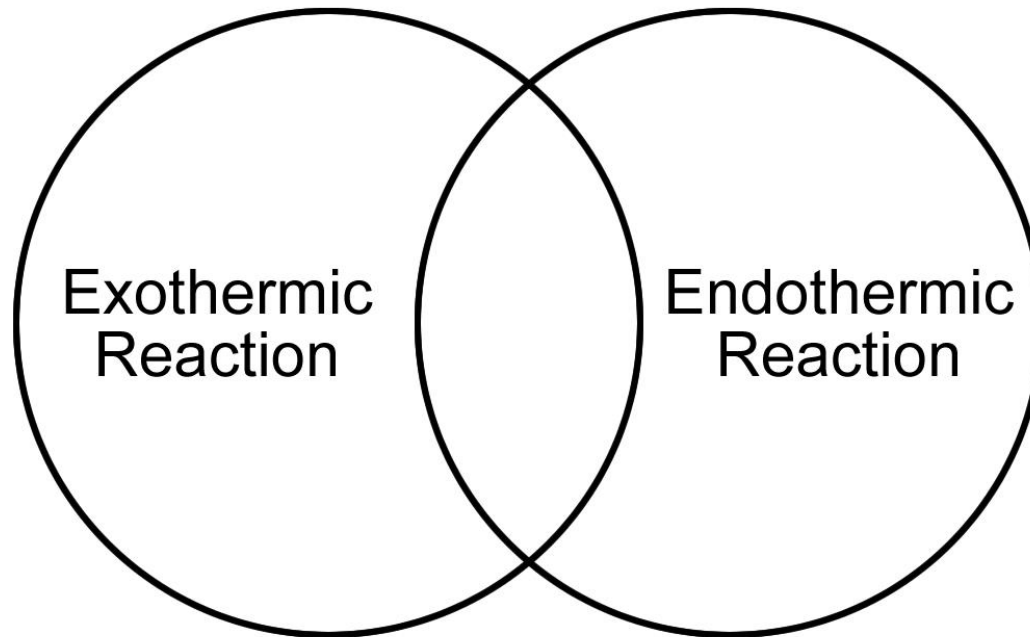
# Energy from Chemicals



Compare and contrast  
*exothermic* and  
*endothermic* reactions.  
In what ways are they  
*similar*? In what ways  
are they *different*?

# Higher Order Thinking Skills

## Compare and Contrast Exothermic and Endothermic Reactions

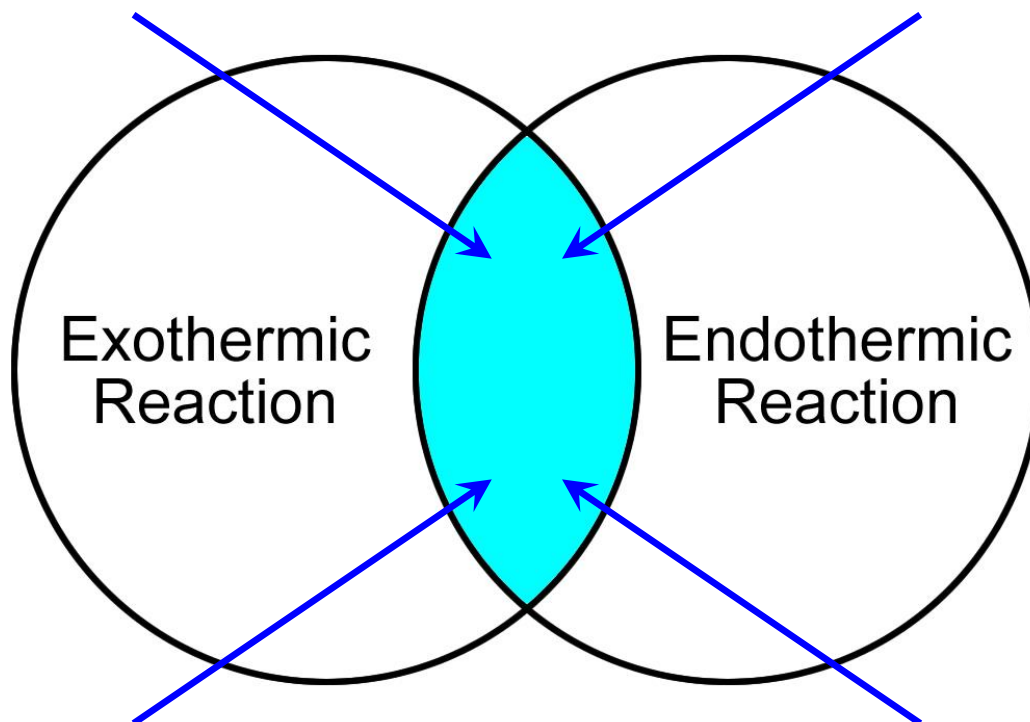


# Higher Order Thinking Skills

## Compare and Contrast Exothermic and Endothermic Reactions

- Reaction involves bond formation.

- Reaction involves bond breaking.

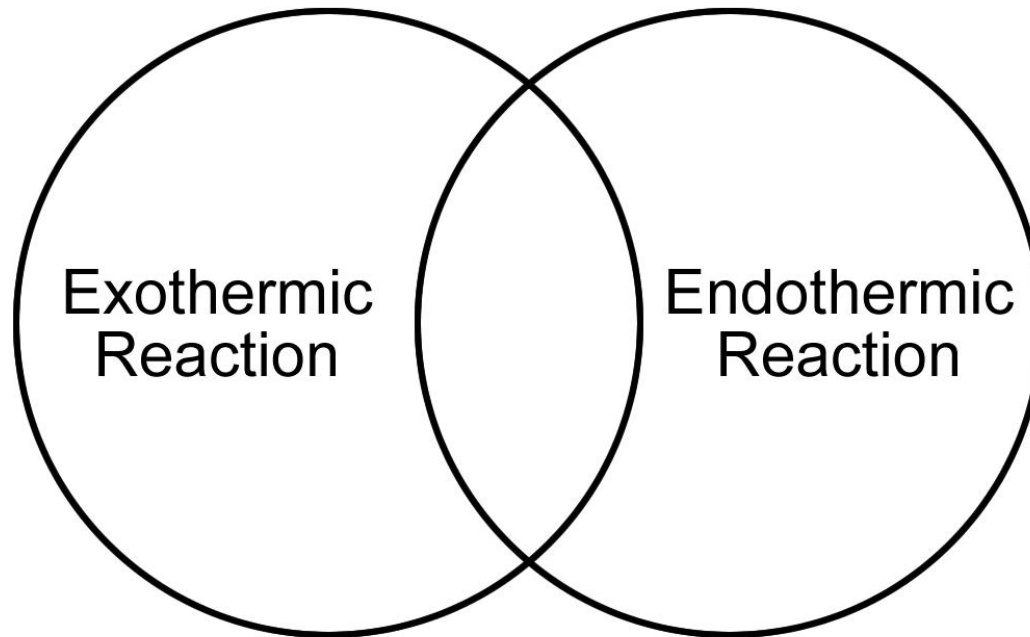


- Reaction has an activation energy.

- A catalyst will lower the activation energy.

# Higher Order Thinking Skills

## Compare and Contrast Exothermic and Endothermic Reactions



# Higher Order Thinking Skills

## Compare and Contrast Exothermic and Endothermic Reactions

- Energy absorbed to break bonds is **less** than the energy released when bonds form.

- Energy absorbed to break bonds is **greater** than the energy released when bonds form.

- Energy of reactants is **greater** than energy of products.

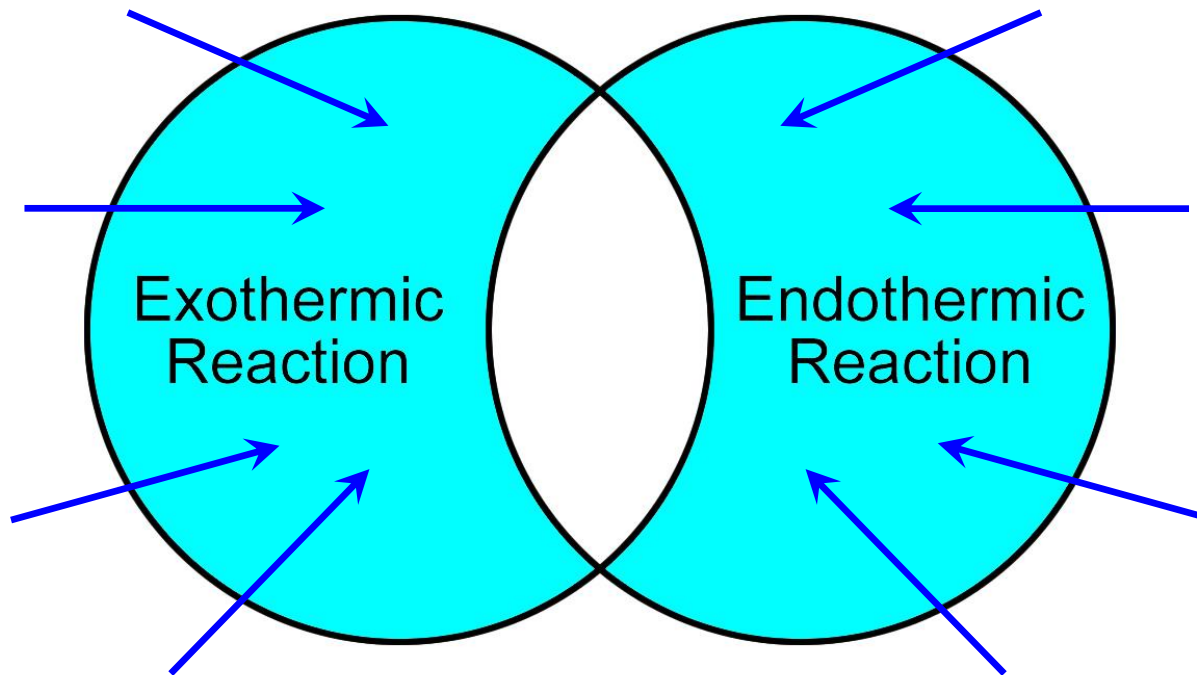
- Energy of reactants is **less** than energy of products.

- $\Delta H$  is **negative**.

- $\Delta H$  is **positive**.

- Energy **released into** the surroundings (temperature increases).

- Energy **absorbed from** the surroundings (temperature decreases).



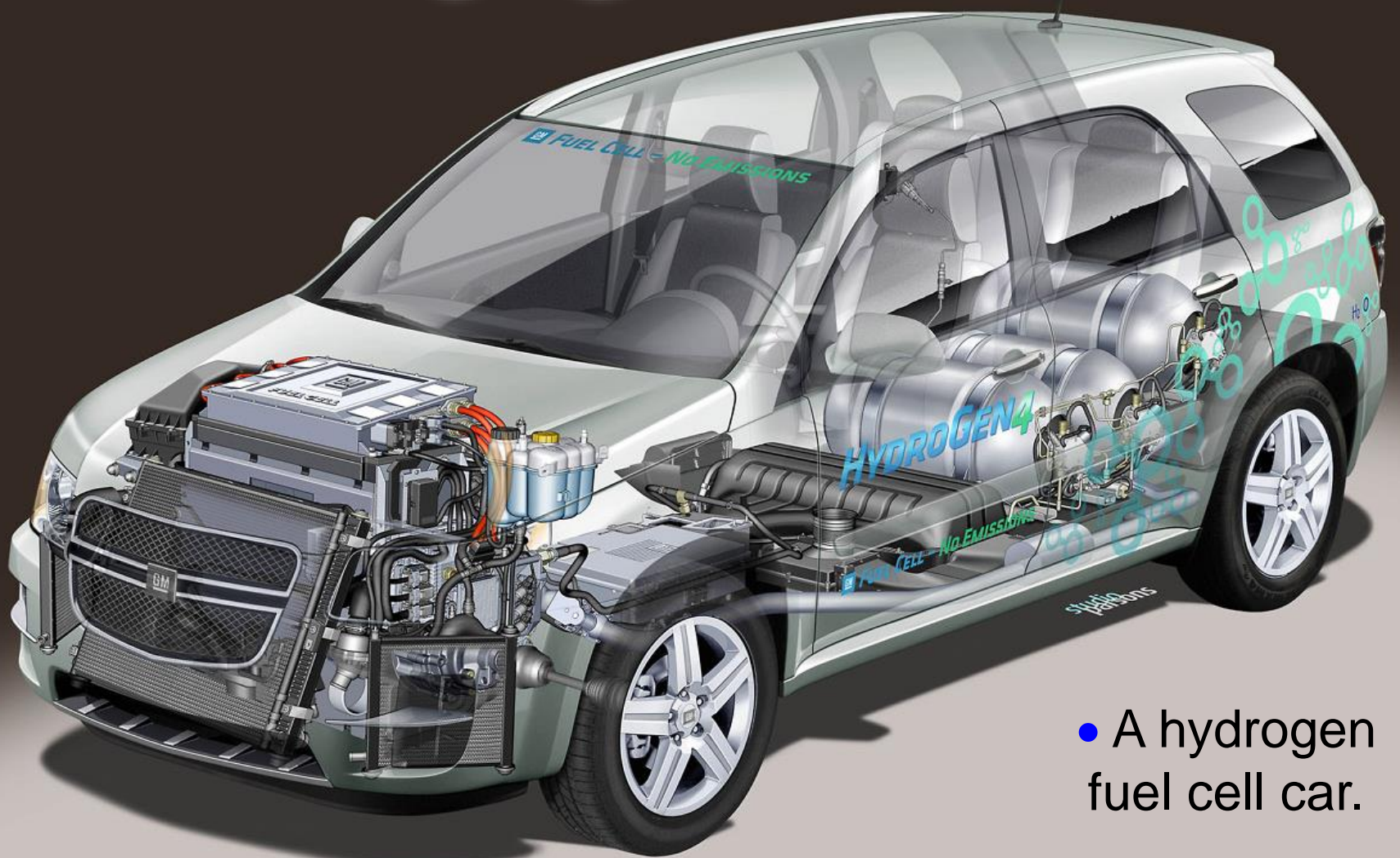
# Energy from Chemicals



*Hydrogen fuel cells* generate electricity without causing any pollution. How do they work?



# Hydrogen Fuel Cells



- A hydrogen fuel cell car.



# Hydrogen Fuel Cells

- The Toyota Mirai – the world's first commercial hydrogen fuel cell car – went on sale in Japan on 15<sup>th</sup> December 2014.



# Hydrogen Fuel Cells

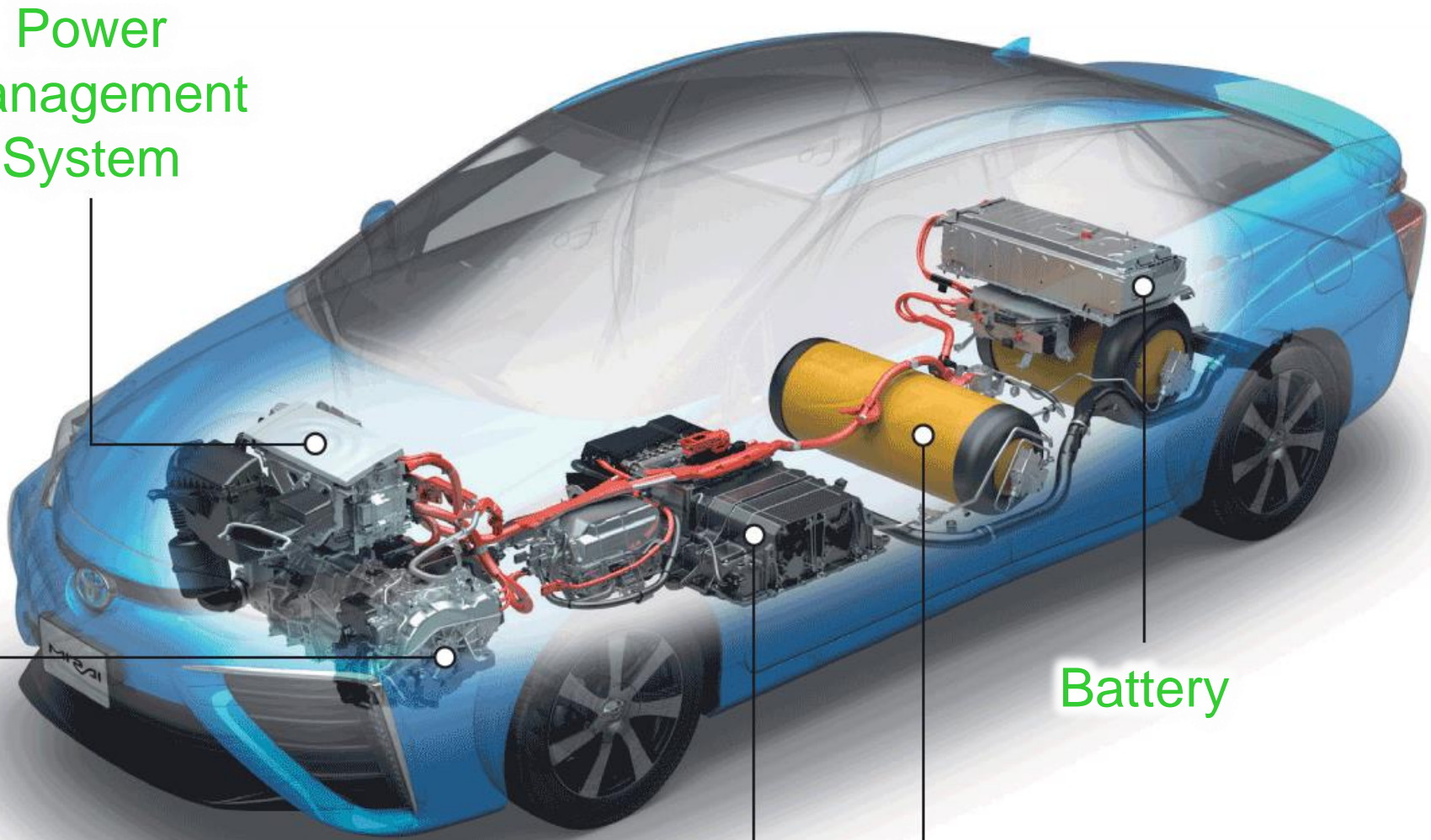
Power  
Management  
System

Battery

Electric Motor

$H_2$  (g) Fuel  
Cell

$H_2$  (g)  
Tank

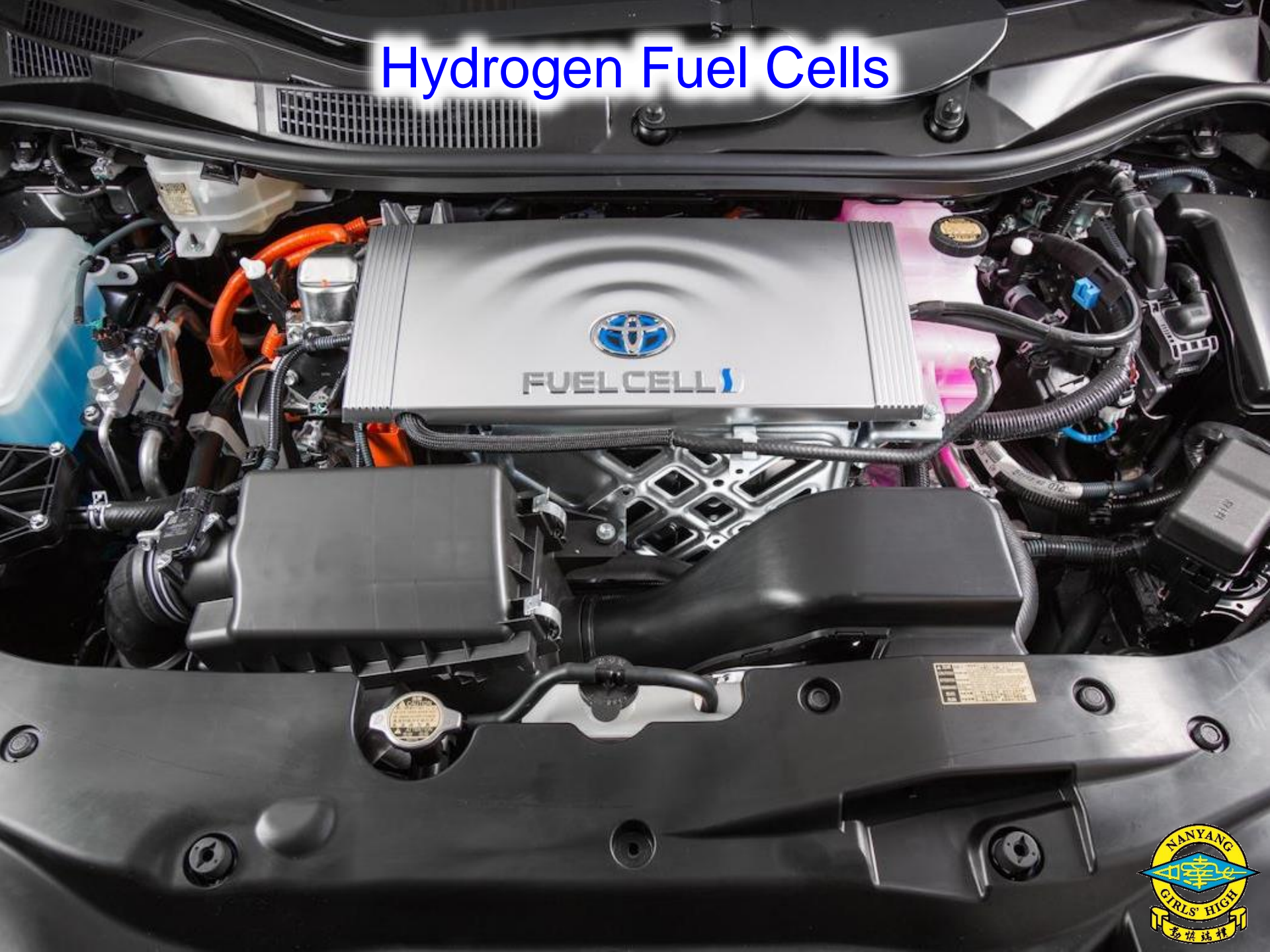




# Hydrogen Fuel Cells



# Hydrogen Fuel Cells



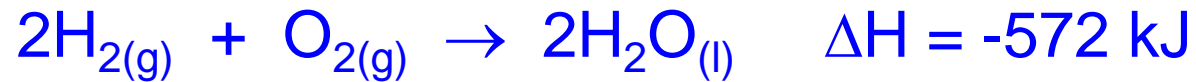


# Hydrogen Fuel Cells



# Hydrogen Fuel Cells

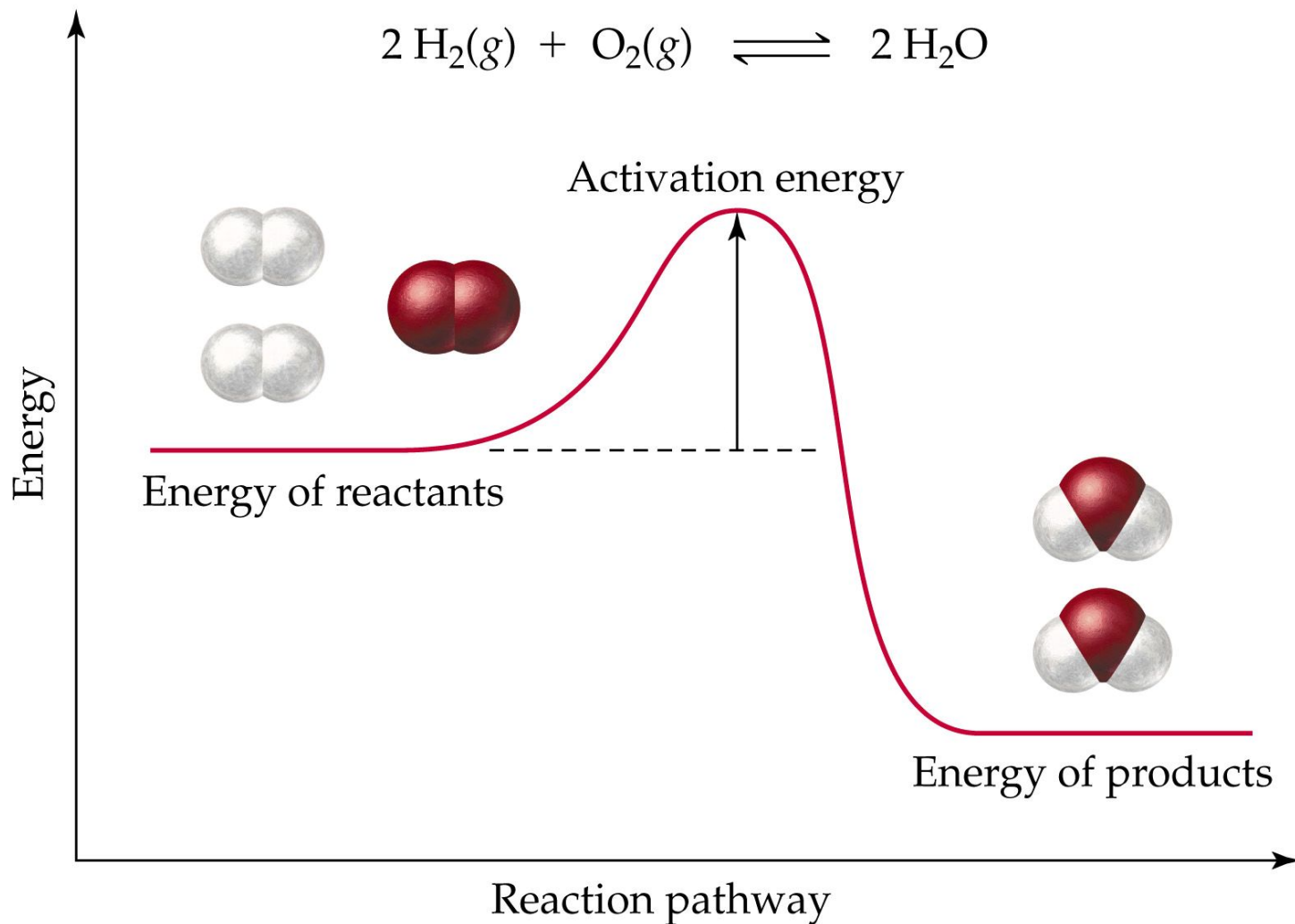
- When hydrogen reacts with oxygen, an exothermic reaction takes place that produces water as the only reaction product.



- In a hydrogen fuel cell, the hydrogen reacts with oxygen to produce water and **energy in the form of electricity**. The electrical energy can be used to do useful work, such as driving an electric motor to move a car.



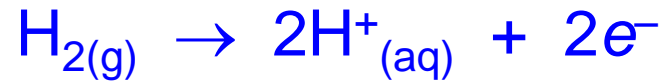
# Hydrogen Fuel Cells





# Hydrogen Fuel Cells

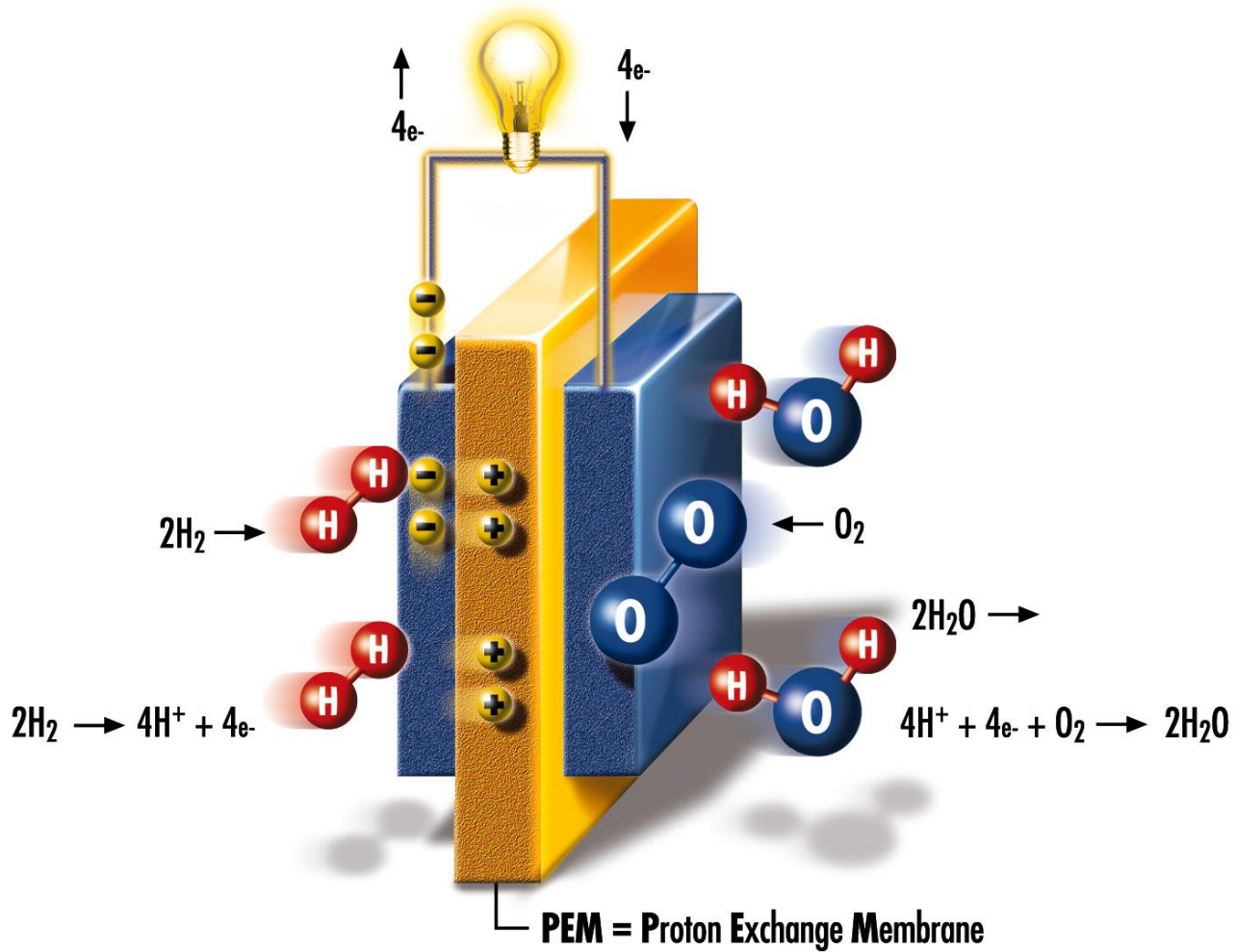
- In the hydrogen fuel cell, a catalyst causes molecular hydrogen to break-down into hydrogen ions and electrons.



- The electrons pass through metal wires in an external circuit where they are made to do useful work, such as drive an electric motor. The hydrogen ions pass through a special membrane.
- On the other side of the membrane, the hydrogen ions, electrons and oxygen combine together to form water.

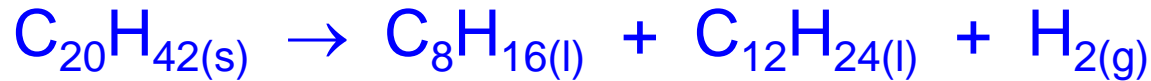


# Hydrogen Fuel Cells



# Hydrogen Fuel Cells

- Hydrogen required for the hydrogen fuel cell is obtained by *cracking* long-chain hydrocarbons from crude oil, e.g.



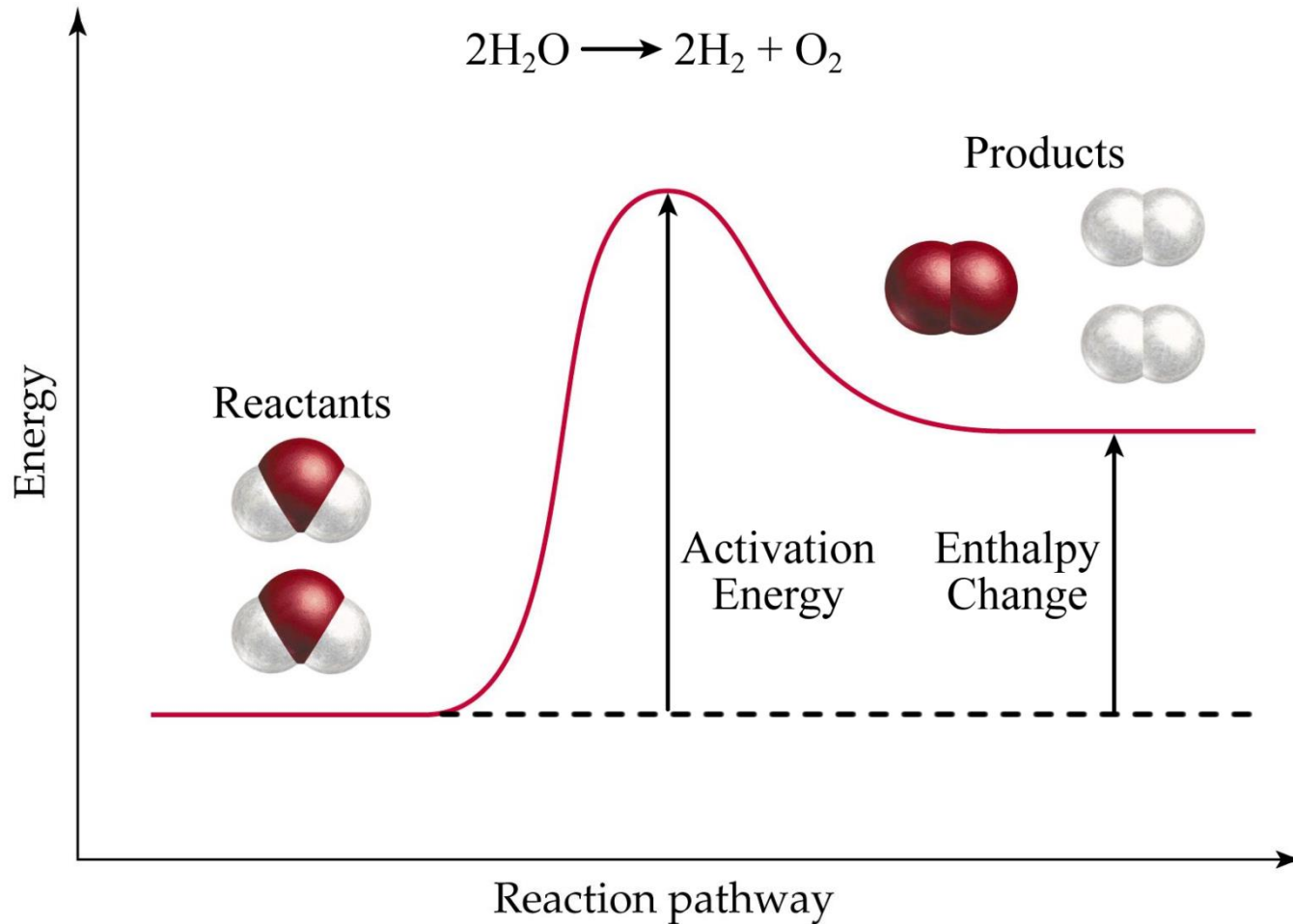
- Hydrogen can also be obtained from the *electrolysis* of water (the decomposition of water by electricity):



- Remember, the hydrogen that is produced by cracking long-chain hydrocarbons is also used in the manufacture of *ammonia* –  $\text{NH}_3$ .
- The hydrogen must be stored with the hydrogen fuel cell. The large scale storage of hydrogen is hazardous because it is a highly flammable gas.



# Hydrogen Fuel Cells



- The electrolysis of water is an *endothermic* process.

# Hydrogen Fuel Cells

- Oxygen required for the hydrogen fuel cell is taken directly from the Earth's atmosphere, which is 21% oxygen.



# Energy from Chemicals



Why is the hydrogen fuel cell considered to be a *clean* source of energy?

# Energy from Chemicals



Why is the hydrogen fuel cell considered to be a *clean* source of energy?

- The only chemical product of the reaction is *water*. Water is not considered to be a pollutant. Compare this to the chemicals that are produced by the internal combustion engine.





# Energy from Chemicals



Is the hydrogen fuel cell really such a *clean* source of energy?

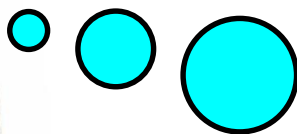
# Energy from Chemicals



Is the hydrogen fuel cell really such a *clean* source of energy?

- Consider how the raw materials (hydrogen and oxygen) for the fuel cell are obtained. To what extent is atmospheric pollution a possible consequence?

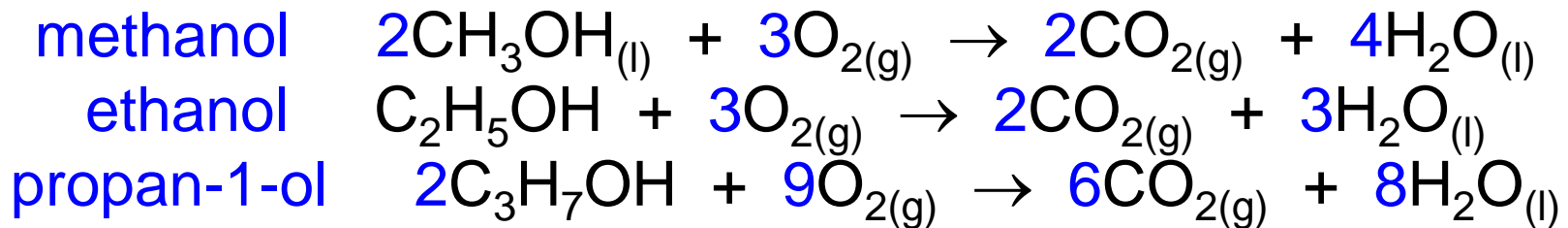
# Energy from Chemicals



How would I *design an experiment* to measure the energy change of a reaction?

# Experimental Design

- The combustion of an alcohol is an exothermic reaction. For example:

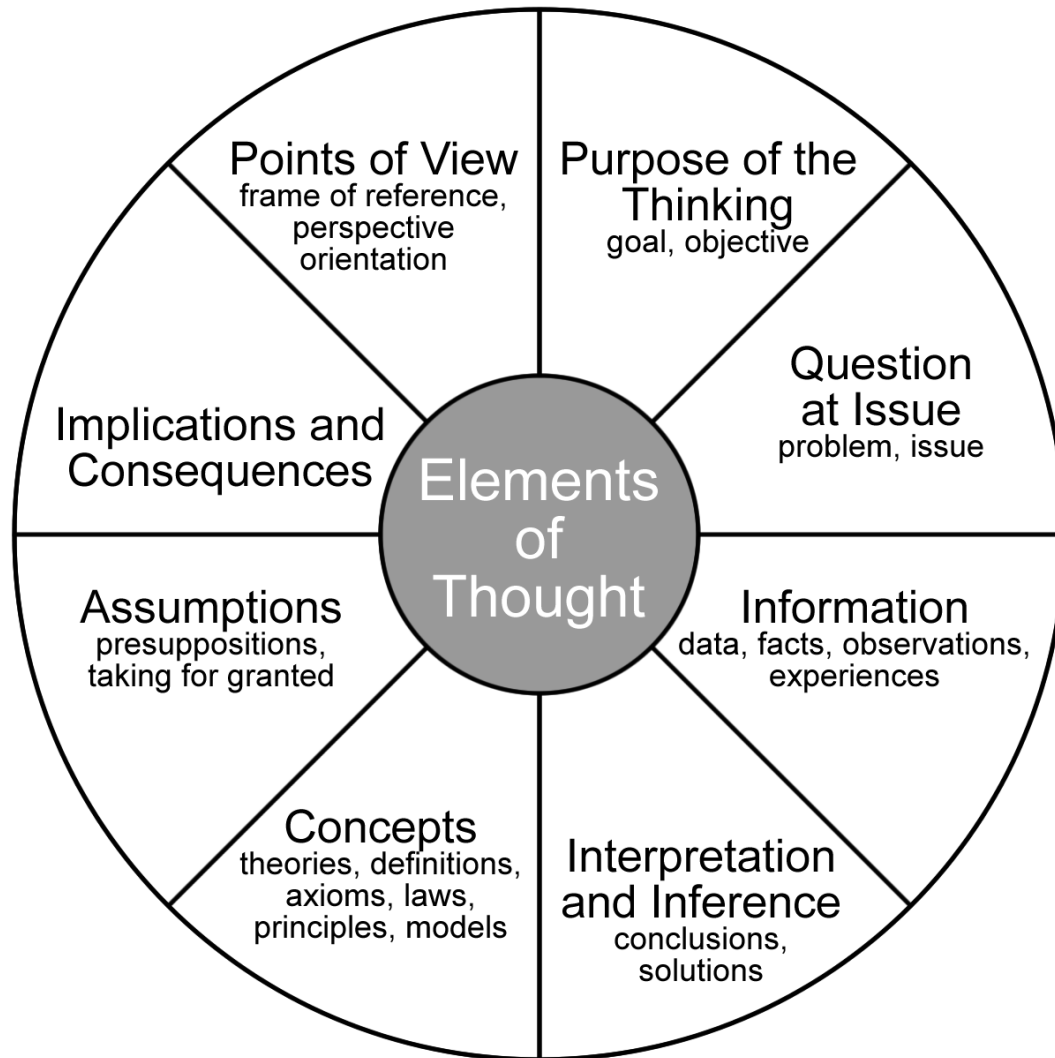


- Design an experiment to determine which alcohol, methanol, ethanol or propan-1-ol, releases the greatest amount of energy when one gram of the alcohol is completely burned in air.



# Experimental Design

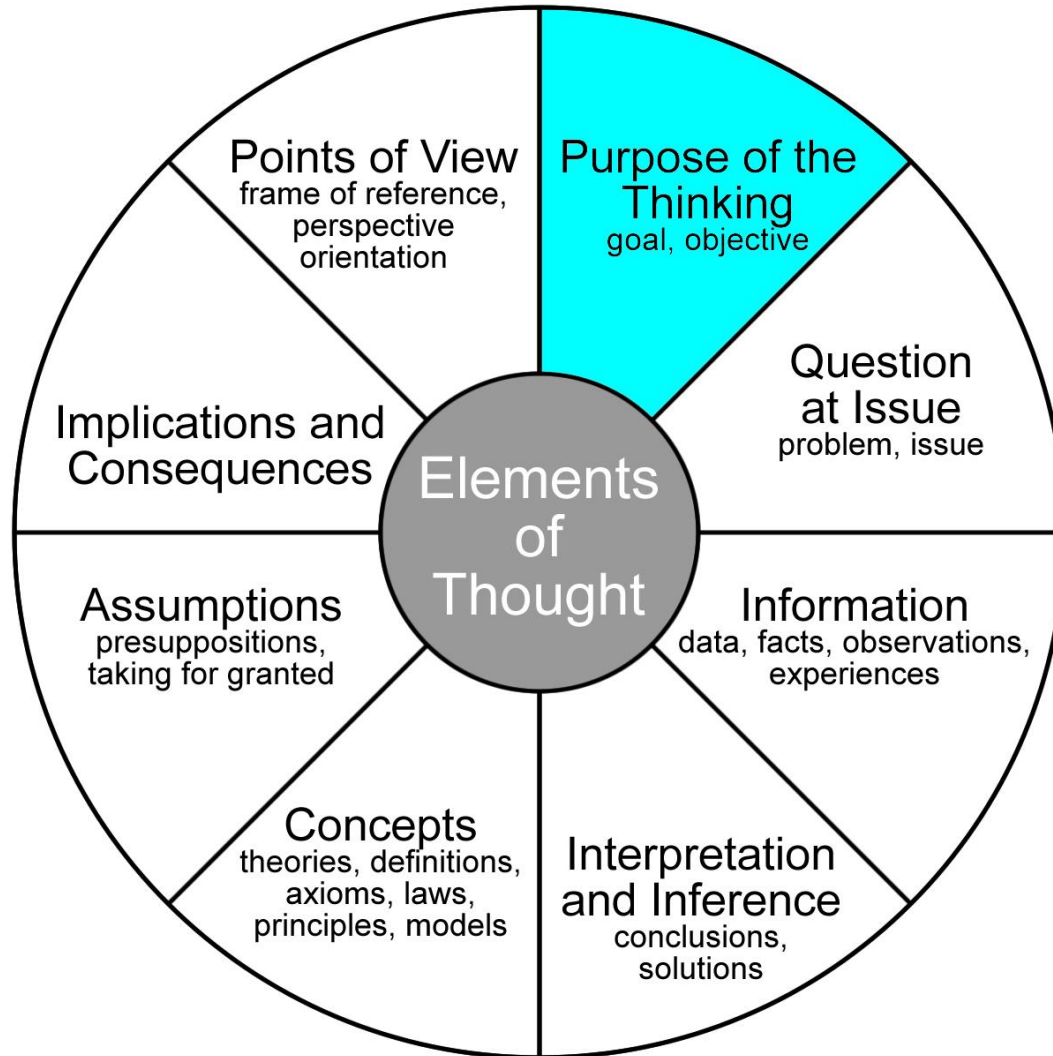
## Paul's Wheel of Reason



- Experiments should be designed using **critical thinking** skills.

# Experimental Design

## Paul's Wheel of Reason

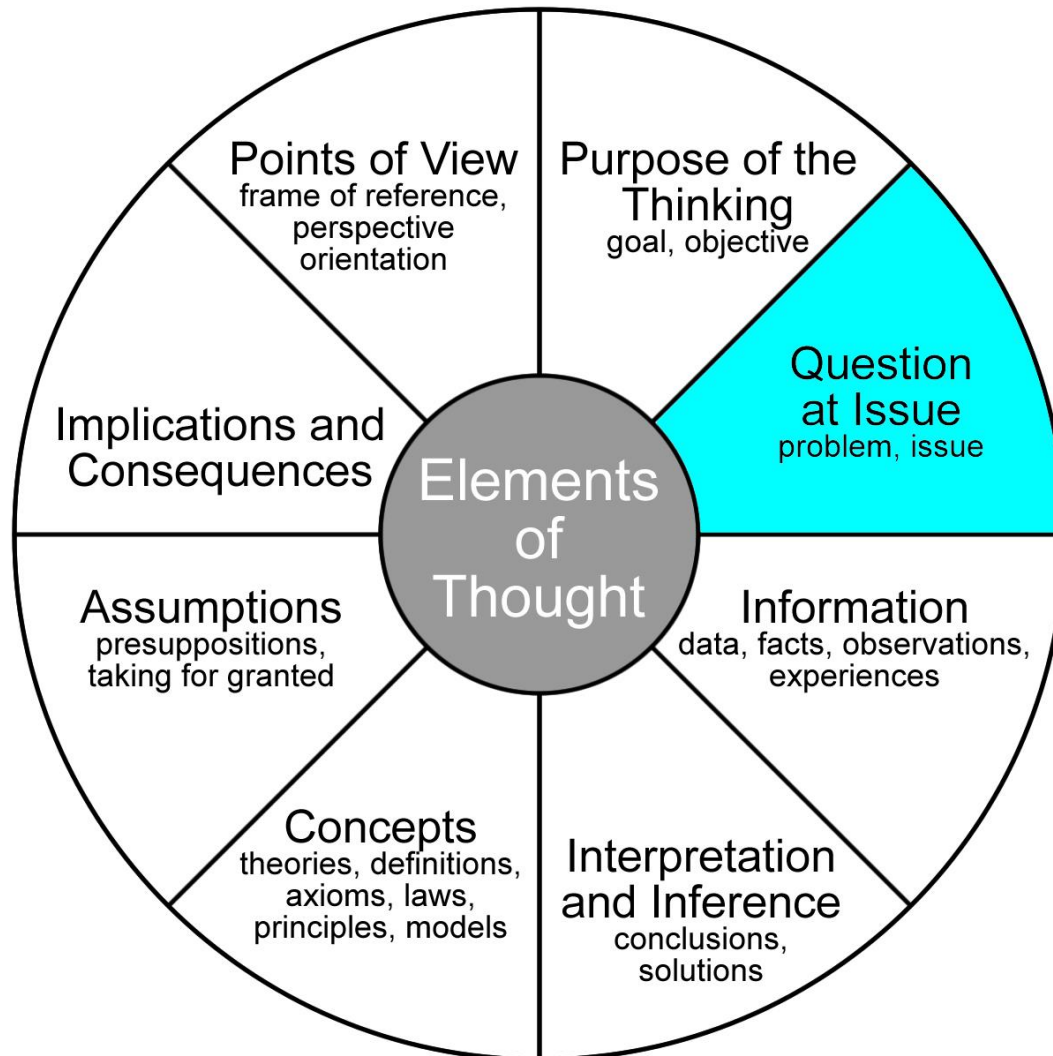


- What is the reason for performing the experiment?
  - What hypothesis is being proved / disproved by performing the experiment?



# Experimental Design

## Paul's Wheel of Reason

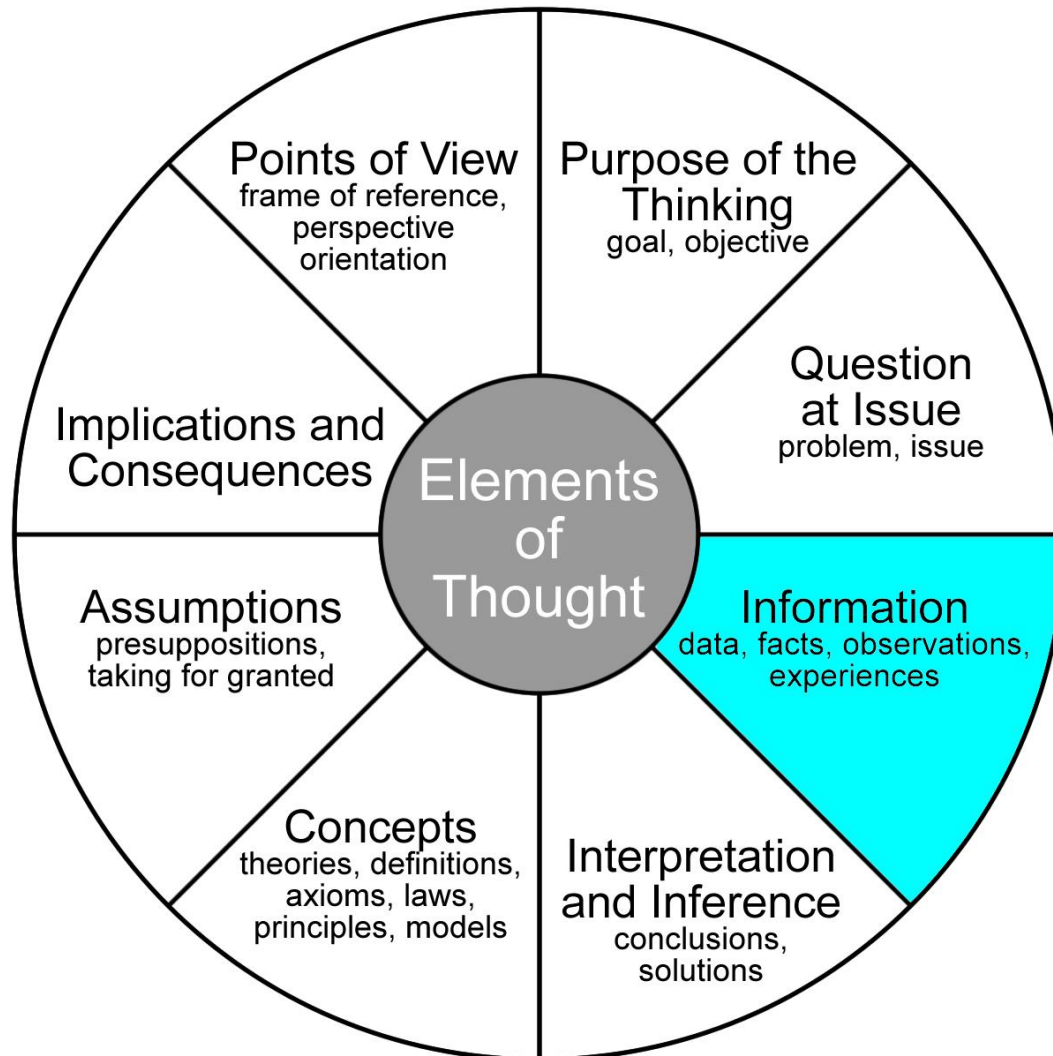


- Clearly state exactly what the experiment is being designed to do.
- What is the problem statement for the experiment?



# Experimental Design

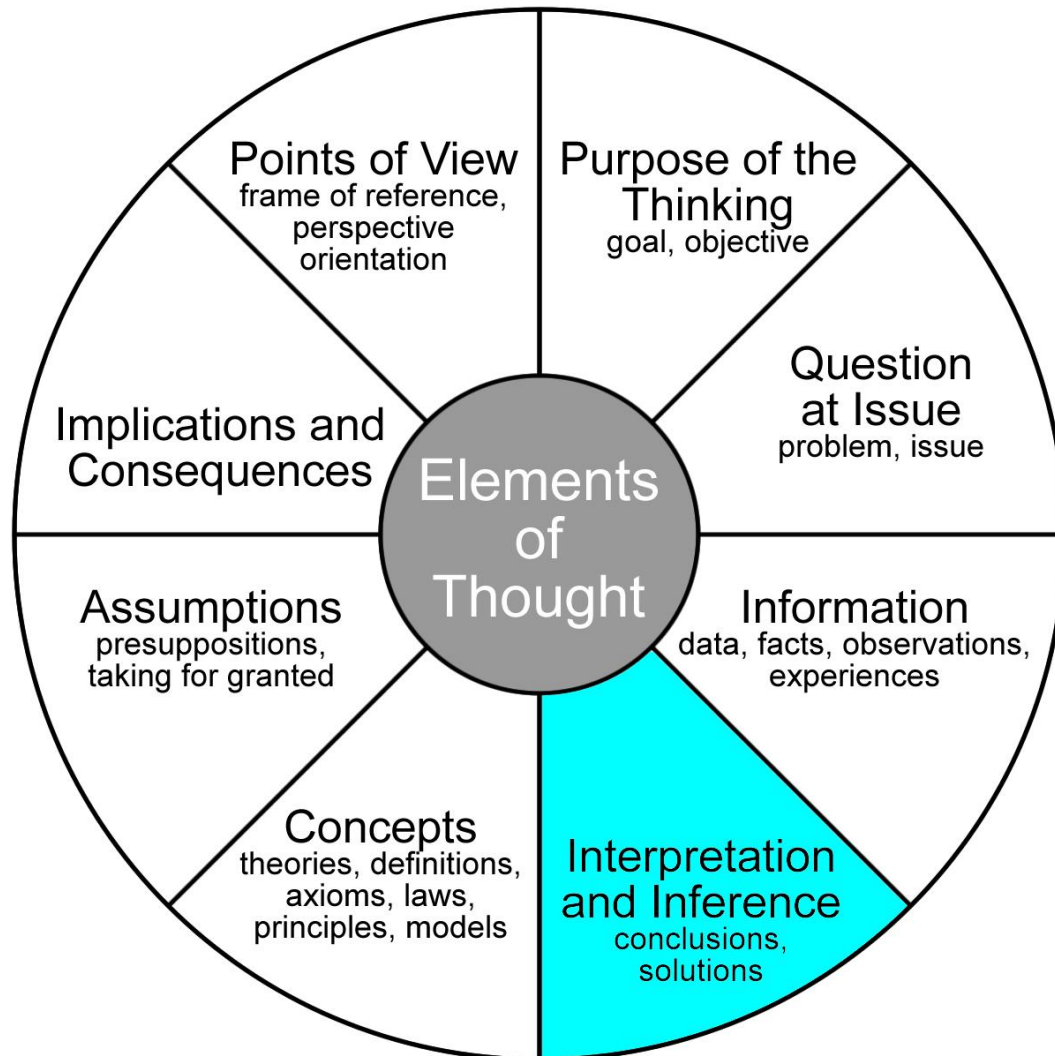
## Paul's Wheel of Reason



- What essential information needs to be collected by the experiment?
  - What observations need to be made?
  - What are the different variables (independent, dependent and control)?

# Experimental Design

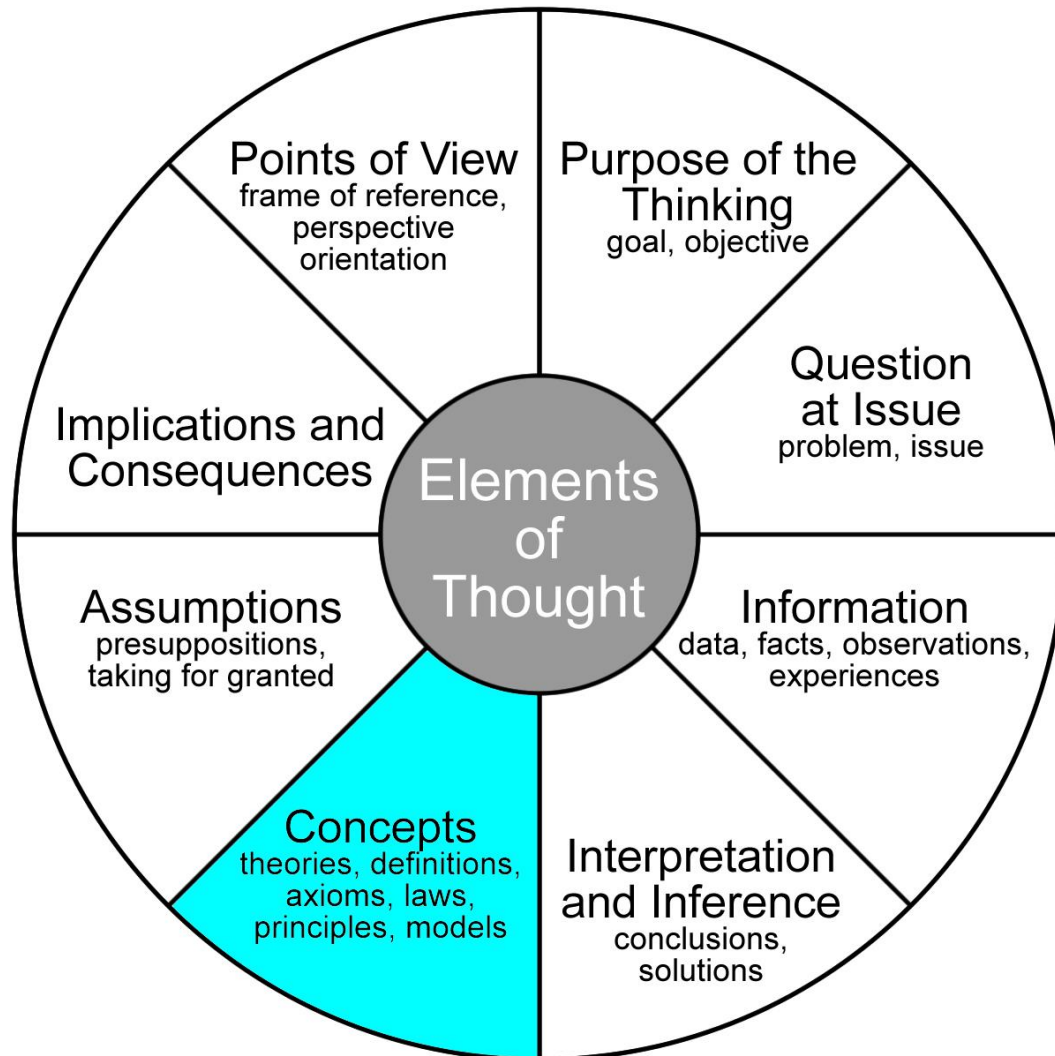
## Paul's Wheel of Reason



- How must the data from the experiment be manipulated and presented?
  - What calculations (if any) need to be done?
- What graphs (if any) need to be plotted?

# Experimental Design

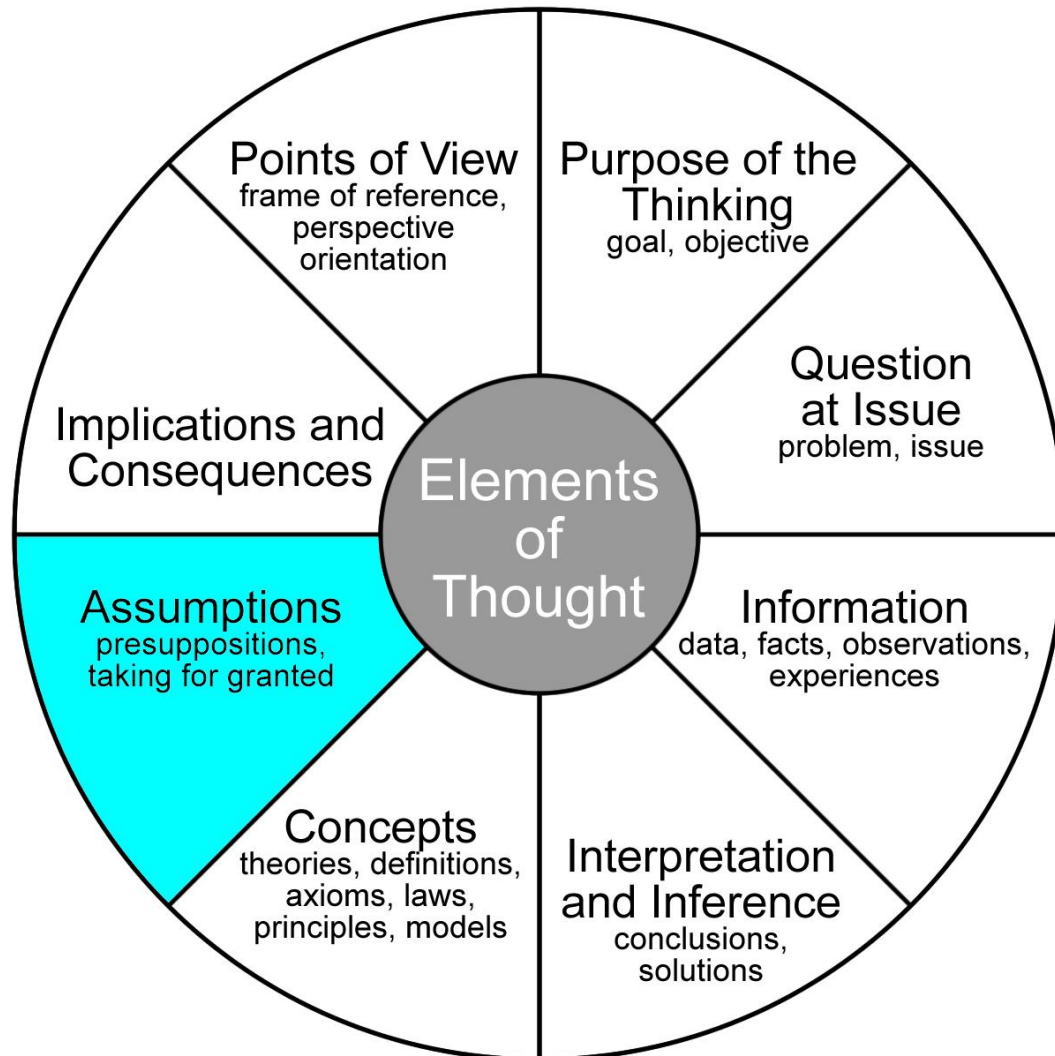
## Paul's Wheel of Reason



- What essential concept / law does the design of the experiment rely on? For example, the concept of energy and the law that energy can not be created or destroyed, but only converted from one form into another.

# Experimental Design

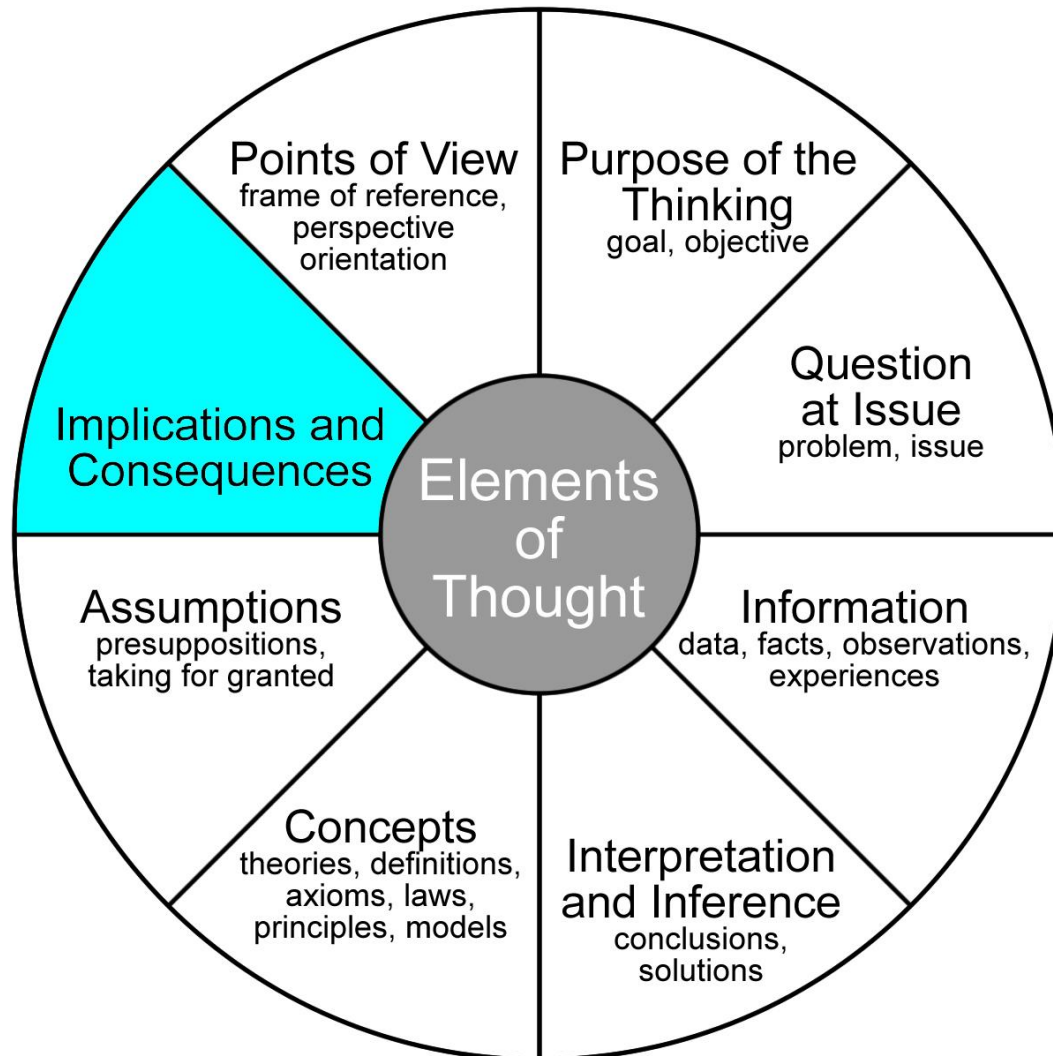
## Paul's Wheel of Reason



- What assumptions are being made when designing the experiment?
- Is it reasonable to make these assumptions?
  - Identifying assumptions can help to identify errors.

# Experimental Design

## Paul's Wheel of Reason

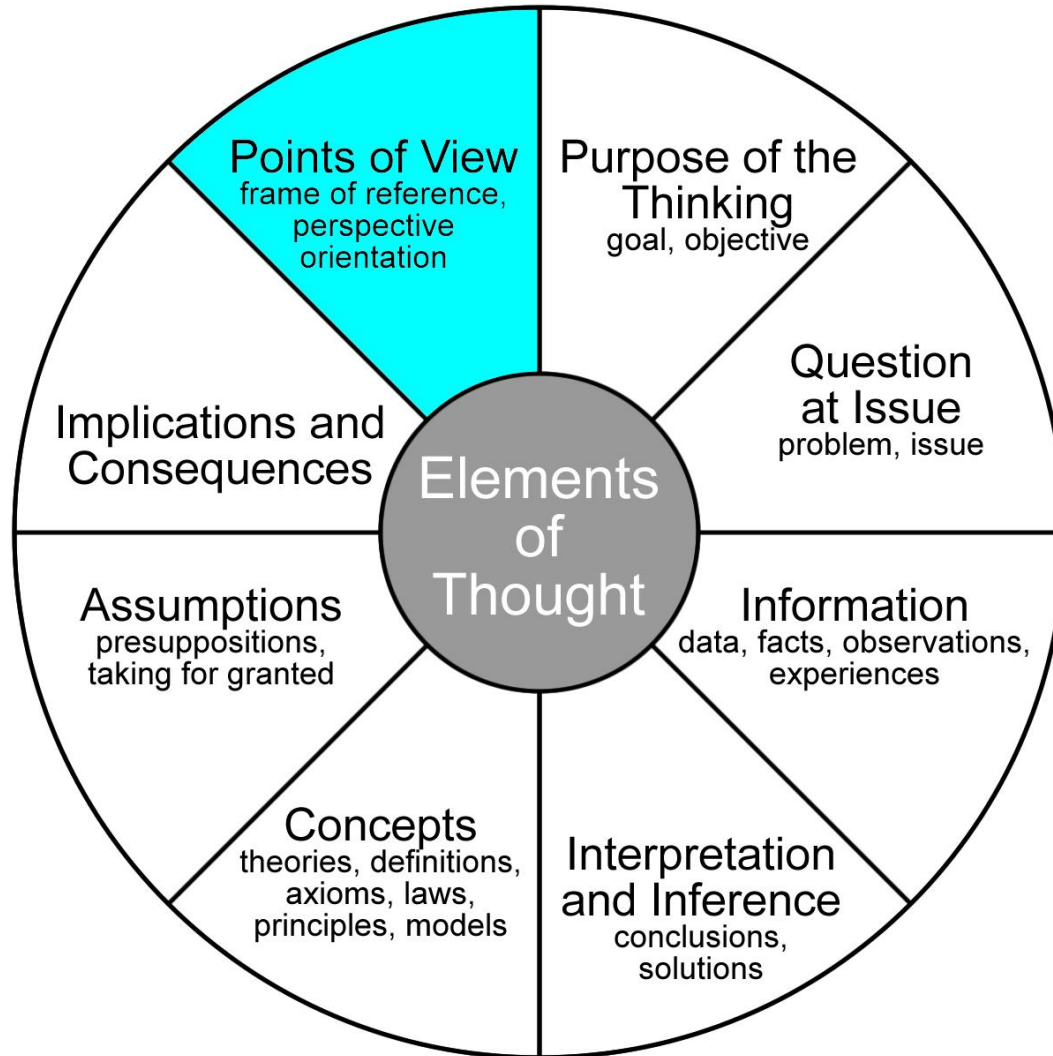


- What are the implications and consequences if the results obtained from the experiment prove / disprove a specific hypothesis?



# Experimental Design

## Paul's Wheel of Reason



- Is there more than just one way of doing this experiment?
- If there is more than one way, which way is the most valid and reliable?

# Experimental Design

- What is your hypothesis or problem statement?
- What are the variables? Which variables must be kept constant?  
What is your independent (input) variable? What is your dependent (output) variable?
- What is the big idea? Provide a general overview of the method.
  - What apparatus is required to perform the experiment?
- How is the apparatus set-up? Provide a clear, labelled diagram.
  - How will you perform the experiment? Provide step-by-step instructions, including how variables will be controlled and what measurements will be taken.
- How will data from the experiment will be recorded? Provide an example results table that includes clear headings and units.
- How will the experimental data be manipulation? What graphs need to be plotted? What calculations need to be performed?
- What errors are associated with the experiment? What effect do these errors have on your results?





# Experimental Design

## Problem Statement

- Does the alcohol containing the greater number of carbon atoms per molecule release more energy when one gram of it is burned?

## Variables

- Mass of water in the copper calorimeter (constant).
  - Mass of alcohol burned during the experiment (independent variable).
- Temperature change of the water in the copper calorimeter (dependent variable).



# Experimental Design

## General Overview of the Method

A known mass of alcohol is burned and used to heat a known mass of water, causing its temperature to increase by a known amount. Using the equation  $E = m \times c \times \Delta T$ , the heat energy absorbed by the water is calculated. It is assumed that the energy used to heat the water originated from combustion of the alcohol, therefore the energy released per unit mass of the alcohol can be calculated.

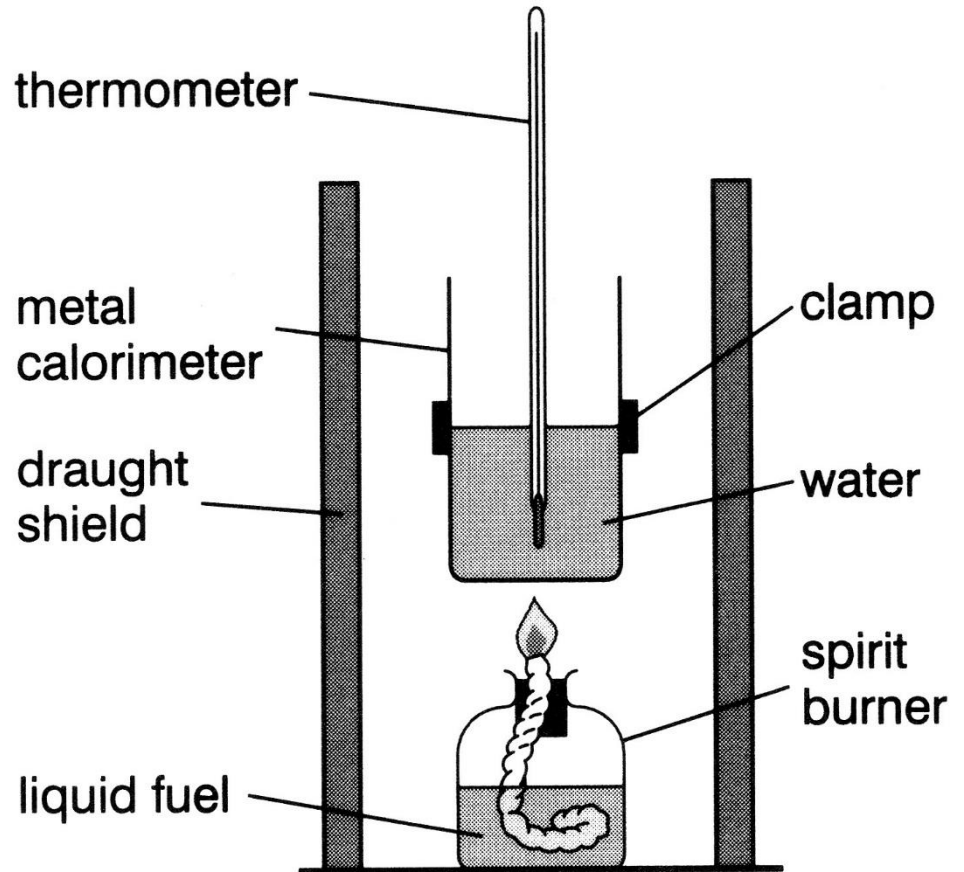


# Experimental Design

## Apparatus and Reagents

- Copper calorimeter
- Digital weighing machine
  - Draught shield
  - Glass rod
  - Matches
- 100 cm<sup>3</sup> Measuring cylinder
- Retort stand and clamp
  - Spirit burner and lid
  - Thermometer
- Methanol
- Ethanol
- Propan-1-ol
- Water

## Diagram



# Experimental Design

## Method

- 1)** Weigh the copper calorimeter on the digital weighing machine. Record the data in the results table.
- 2)** Use the measuring cylinder to pour  $80 \text{ cm}^3$  of water into the copper calorimeter.
- 3)** Re-weigh the copper calorimeter and water and hence calculate the mass of water present in the calorimeter. Record the data in the results table.
- 4)** Support the copper calorimeter in the retort stand and clamp.
- 5)** Place a thermometer in the copper calorimeter and measure the initial temperature of the water. Record the data in the results table.
- 6)** Weigh the spirit burner and lid containing the methanol on the digital weighing machine. Record the data in the results table.



# Experimental Design

## Method

- 7) Place the spirit burner under the copper calorimeter. Adjust the height of the calorimeter so that it is 2 – 3 cm above the spirit burner.
- 8) Remove the lid from the spirit burner and, using the matches, immediately light the spirit burner. Adjust the height of the copper calorimeter once more so that the flame of the burning methanol covers the base of the calorimeter.
- 9) Using the glass rod, gently stir the water in the copper calorimeter. Ensure that no water spills out of the calorimeter.
- 10) When the temperature of the water in the copper calorimeter has increased by 20 °C, place the lid back on the spirit burner to extinguish the flame.



# Experimental Design

## Method

- 11)** Measure the temperature of the water in the copper calorimeter at the end of the experiment. Record the data in the results table.
- 12)** Calculate the change in temperature of the water in the calorimeter. Record the data in the results table.
- 13)** Re-weigh the spirit burner on the digital weighing machine at the end of the experiment. Record the data in the results table.
- 14)** Calculate the mass of methanol burned during the experiment. Record the data in the results table.
- 15)** Repeat the experiment for the other two alcohols, ethanol and propan-1-ol.
- 16)** Repeat the entire experiment at least once more.



# Experimental Design

## Results

	Methanol	Ethanol	Propan-1-ol
Mass of the copper calorimeter / g	X	X	X
Mass of the copper calorimeter + water / g	X	X	X
Mass of water in the copper calorimeter / g	X	X	X
Mass of spirit burner + lid + alcohol at the start of the reaction / g	X	X	X
Mass of spirit burner + lid + alcohol at the end of the reaction / g	X	X	X
Mass of alcohol burned during the reaction / g	X	X	X
Temperature of the water in the copper calorimeter at the start of the reaction / g	X	X	X
Temperature of the water in the copper calorimeter at the end of the reaction / g	X	X	X
Temperature increase of the water in the copper calorimeter during the reaction / g	X	X	X





# Experimental Design

## Calculation

- It is assumed that all of the energy released during combustion of the alcohol is absorbed by the water in the copper calorimeter, *i.e.*

energy produced by combustion of alcohol = energy gained by water

- Energy gained by water =  $m_w \times c \times \Delta T$

where:

$m_w$  = mass of water / g

$c$  = specific heat capacity of water = 4.20 j / g / °C

$\Delta T$  = change in temperature of the water / °C

- Energy released by burning 1 g of alcohol =  $(m_w \times c \times \Delta T) \div m_a$

where:

$m_a$  = mass of alcohol / g



# Experimental Design

## Source of Error

### 1) The alcohol undergoes incomplete combustion.

Incomplete combustion of the alcohol releases less energy than complete combustion. This will reduce the amount by which the water in the calorimeter increases in temperature, resulting in a smaller value for the calculated energy change.

### 2) Heat energy is lost to the surroundings.

Heat energy from combustion of the alcohol is lost to the surroundings rather than being transferred into the water. This will reduce the amount by which the water in the calorimeter increases in temperature, resulting in a smaller value for the calculated energy change.



# Experimental Design

## Source of Error

### 3) Heat energy is absorbed by the copper calorimeter.

Heat energy from combustion of the alcohol is absorbed by the copper calorimeter rather than being transferred into the water.

This will reduce the amount by which the water in the calorimeter increases in temperature, resulting in a smaller value for the calculated energy change.



# Experimental Design

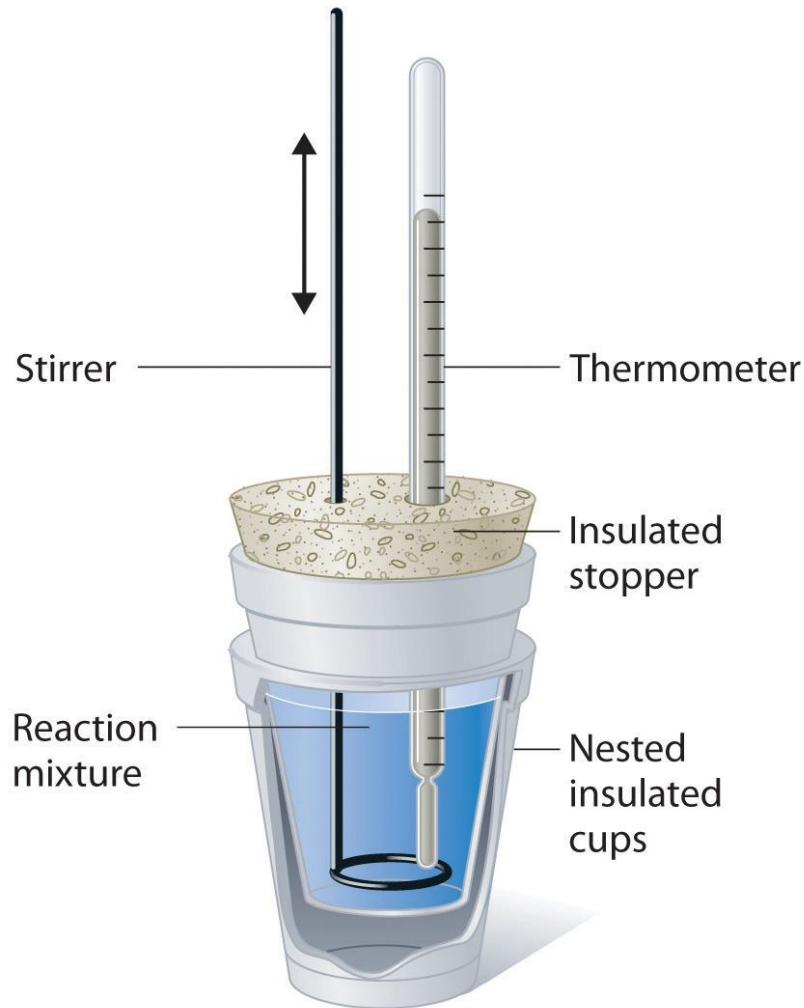
- When ammonium chloride dissolves in water, an endothermic energy change takes place:



- Design an experiment to determine the energy change when one mole of ammonium chloride dissolves completely in water.



# Experimental Design



- The apparatus that could be used for conducting this experiment is shown in the diagram on the left.
- Refer to the diagram to help you design the experiment.

# Energy from Chemicals

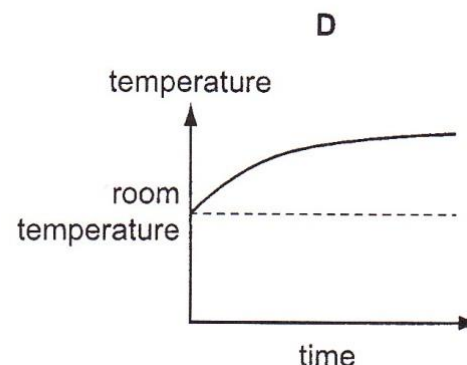
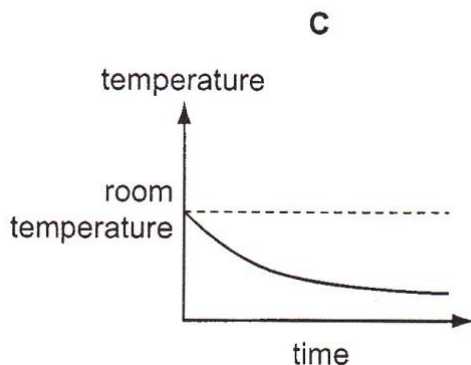
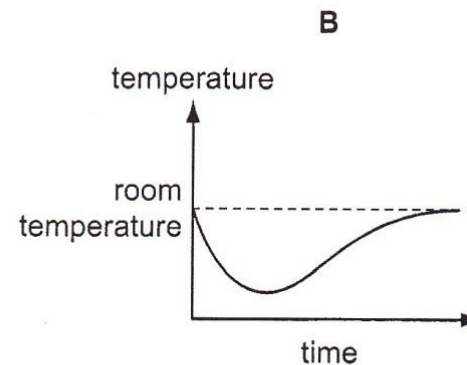
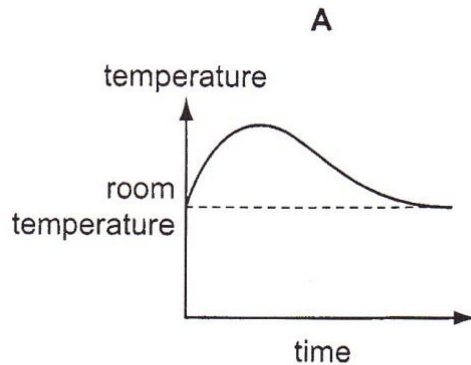


Could I please have  
some questions to  
*test my understanding*  
of the topic Energy  
from Chemicals?

# Assessment for Learning

## Question 1:

- Dissolving ammonium nitrate in water is endothermic. Which graph shows how the temperature alters as the ammonium nitrate is added to water and the solution is left to stand?

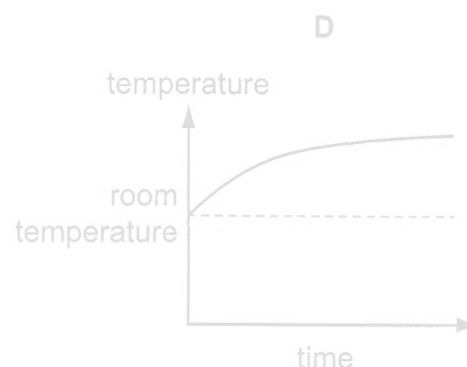
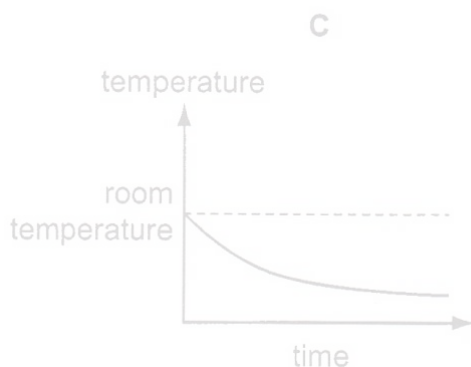
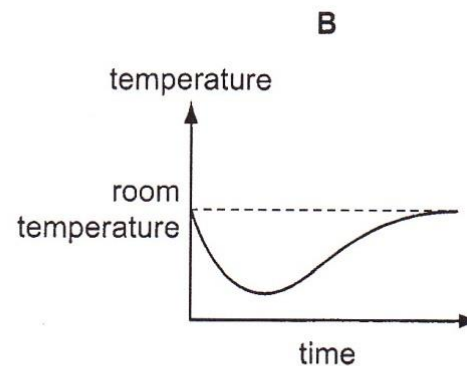
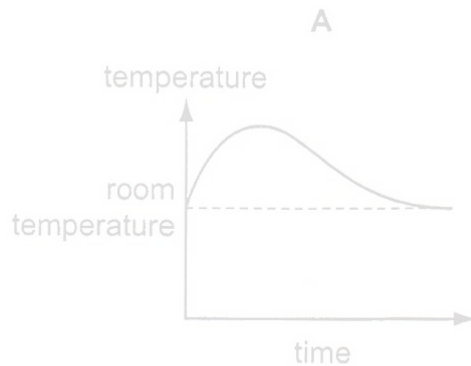




# Assessment for Learning

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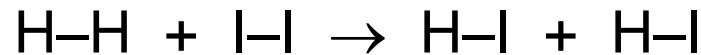
- Dissolving ammonium nitrate in water is endothermic. Which graph shows how the temperature alters as the ammonium nitrate is added to water and the solution is left to stand?



# Assessment for Learning

## Question 2:

- The formation of hydrogen iodide from hydrogen and iodine is an endothermic reaction.



What may be deduced from this information?

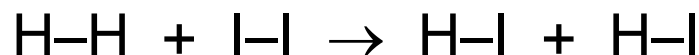
- A** The number of bonds broken is greater than the number of bonds formed.
- B** The formation of H-I bonds absorbs energy.
- C** The products possess less energy than the reactants.
- D** The total energy change in bond formation is less than that in bond breaking.



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# Assessment for Learning

## Question 3:

- The table below shows the energy released by the complete combustion of some compounds used as fuels.

Compound	Formula	$M_r$	$\Delta H$ in kJ / mol
methane	$\text{CH}_4$	16	-880
ethanol	$\text{C}_2\text{H}_5\text{OH}$	46	-1380
propane	$\text{C}_3\text{H}_8$	44	-2200
heptane	$\text{C}_7\text{H}_{16}$	100	-4800

Which fuel produces the most energy when 1 g of the compound is completely burned?

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**C** Methane                      **D** Propane



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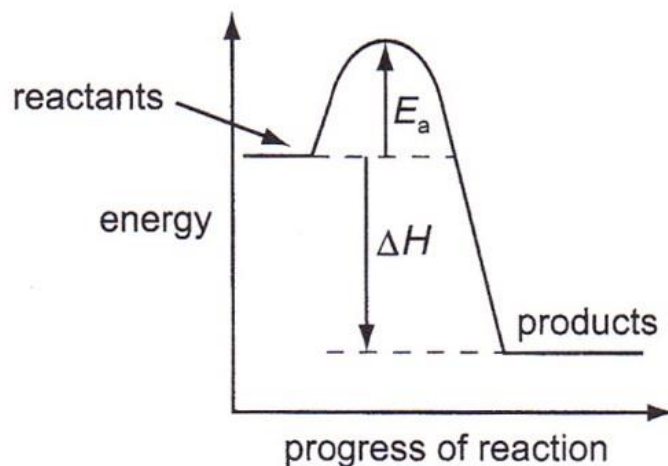
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**D** Propane



# Assessment for Learning

## Question 4:

- The enthalpy diagram shows an uncatalysed, exothermic reaction.



The reaction was repeated in the presence of a catalyst. What effect does the catalyst have on the activation energy,  $E_a$ , and the enthalpy change,  $\Delta H$ ?

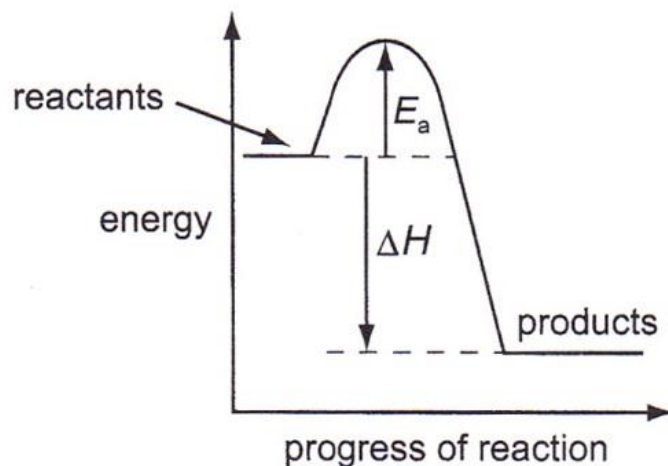
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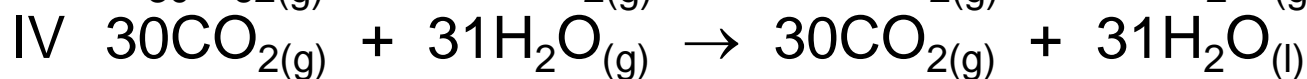
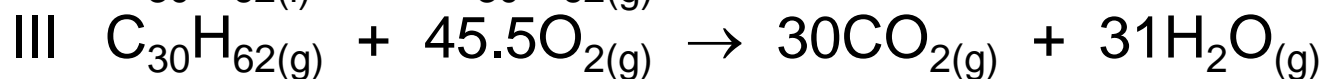
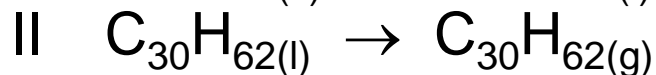
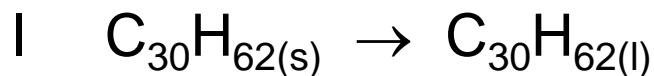




# Assessment for Learning

## Question 5:

The scheme shows four stages, I to IV, in the conversion of solid candlewax,  $C_{30}H_{62}$ , into carbon dioxide and water.



Which stages are exothermic?

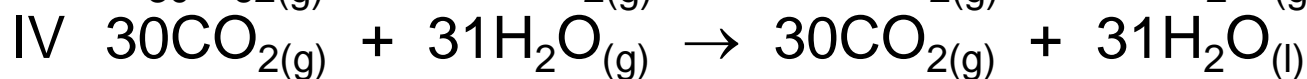
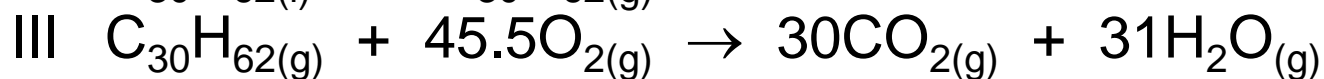
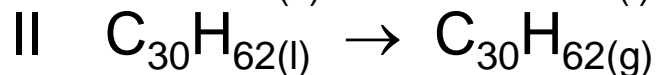
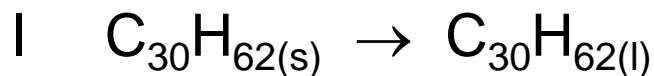
- A** I and II.
- B** II and III.
- C** III and IV.
- D** I and IV.



# Assessment for Learning

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- A I and II.
- B II and III.
- C III and IV.**
- D I and IV.



# Energy from Chemicals



Presentation on  
**Energy from Chemicals**  
by Dr. Chris Slatter

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8<sup>th</sup> February 2016

