HW08 - Enthalpy & Fossil Fuels

(NOTE: the links in this pdf do not work. The links are active when inside of canvas.)

You might need to grab some data from <u>here</u> for the bond energy problems. Stuck on bomb calorimeters? Here's a video:<u>Thermodynamics - Calorimetry Pt II - Bomb</u> <u>Calorimeter Example</u>

Still feel like you aren't fully there with the conceptual part of calorimetry? Here's a video: <u>Thermodynamics - Calorimetry - Part I</u>

1

A 1.00 g sample of n-hexane (C₆H₁₄) undergoes complete combustion with excess O₂ in a bomb calorimeter. The temperature of the 1815 g of water surrounding the bomb rises from 26.15°C to 29.97°C. The heat capacity of the hardware component of the calorimeter (everything that is not water) is 5068 J/°C. What is thæhange in energyfor the combustion of n-C₆H₁₄? One mole of n-C₆H₁₄ is 86.1 g. The specific heat of water is 4.184 J/g°C.

- O -6.33 x 10⁴ kJ/mol
- O -4.40 x 10³ kJ/mol
- O -4.16 x 10³ kJ/mol
- O -5.25 x 10³ kJ/mol

2

Fill in the blanks to receive credit for each part of this question.

An unknown fuel distilled in a refinery (molar mass 64.0 g/mol) is combusted in a bomb calorimeter holding 991 mL water. When 0.182 grams of the fuel source is combusted in the bomb calorimeter, the temperature of the surroundings raises from 25.0 °C to 27.2 °C. The heat capacity for the hardware component is 2.260 kJ/ °C. The heat capacity of water is 4.184 J/ g °C.

In a bomb calorimeter, the thermometer is in the
The combustion of the fuel that we are measuring here is
· The enthalpy of this reaction is equal to
kJ. The enthalpy per gram of this reaction is about
kJ/g. The enthalpy per mole of this reaction isclosest to
kJ/mol.
iii system iii surroundings iii exothermic iii (no way to tell)
iii endothermic iii -26.8 iii -8.37 iii -22.1 iii -8.73 iii -14.1
iii-8.36 iii-1210 iii-2.56 iii-22.1 iii-77.4 iii-4956
Image:

3

Calculate the change in enthalpy of the following reaction in kJ/mol using bond energy data:

 $CIF + CO \rightarrow COCIF$

4 6 point

Using the bond energy data provided, calculate Δ H for the following reaction: H₂(g) + Cl₂(g) \rightarrow 2HCl (g)

H2(8)	$+ Cl_2(g) \rightarrow 2HCl(g)$
Bond	Bond Energy (kJ/mol)
H-H	436
CI-CI	242
H-CI	432

- O 186 kJ/mol
- O -186 kJ/mol
- O -246 kJ/mol
- O 246 kJ/mol



Estimate the change in enthalpy of the following reaction using bond energy data: $N_2H_4(g) + H_2(g) \rightarrow 2NH_3(g)$

- O 850 kJ/mol
- O 1241 kJ/mol
- O -183 kJ/mol
- O -1469 kJ/mol

6		

What is the value of heat flow for the combustion of hydrogen in kJ/g ? ΔH^* for this process is -286 kJ/mol.

- -572 kJ/g
 572 kJ/g
- -71.5 kJ/g
 -286 kJ/g
- O -143 kJ/g

7 5 point

This is a question that requires you to be completely precise and accurate. The numeric answer to this is exact based on the numbers that you have to use. So the answer is a large integer (4 digits to be exact) and I need you to be EXACTLY right on this. If you follow the steps that I showed in class on 11-16-2021, you should be able to do this easily. QUESTION: What is the heat of combustion (it is a positive value because it is the heat given off - or released from the combustion) of exactly one mole of heptane? You answer has to be exactly right and in kJ.

8 6 points

Which of the following is the most efficient fuel based on its combustion enthalpy per gram?

- O wood
- O coal
- O octane
- O methane
- O hydrogen

9 6 p

What is the more efficient method to break a high molar mass fraction from a crude oil refinery down to a specific fuel?

- O reforming
- O thermal cracking
- O fractional distillation
- O catalytic cracking

10 6 pc

An octane isomer can be made into a more efficient fuel by adding branching through the process of...

- O catalytic cracking
- O catalytic reforming
- O thermal cracking
- O fractional distillation

11 4 poin

If you want to calculate the heat flow involving a temperature change, which equation will you use?

- $O \quad q = mC_s \Delta T$
- $\bigcirc q = m\Delta H$
- $\bigcirc q = mC$
- O Σn bonds breaking -Σn bonds forming
- $O \quad q = 2(m C_s \Delta T)$

4 p	
	w want to calculate the heat flow involving a phase change, which equation will you
use?	
0	$q = m\Delta H_{trans}$
0	q = mC
Ο	$q = 2(m - \zeta_{S}\Delta T)$
Ο	$q = mC_3\Delta T$
0	Σn bonds breaking -Σn bonds forming
4 p	
D	Designate the sign of the heat flow (+ or -) for each of the following physical changes:
Va	'aporization:
	Fusion: Freezing:
	Sublimation:
	Subimation.
	points
	at is the heat required to completely melt a 11.33 g sample of silicon (Si, molar mass = 19 g/mol) solid that is already at its melting point? ΔH_{fus} = 50.2 kJ/mol. Answer in units
	J and round to <u>one decimal place</u> .
5 p	
	t 1 of 4) Draw the heating curve for the process of heating 14.0 g pure ice from -18.0
	o 84 °C and use it to answer the next four questions. at is the heat required to heat the ice to 0 °C? Answer in joules to the nearest whole
num	
	t 2 of 4) What is the heat required to fully melt the ice at 0 °C? Answer in joules to the rest whole number.
5 p	
	t 3 of 4) What is the heat required to heat the water from 0 °C to 84 °C? Answer in
	es to the nearest whole number.
	t 4 of 4) What is the total heat applied during this process? Answer in kilojoules (!) to
uree	e significant figures.
5 p	
	specific heat for liquid argon and gaseous argon is 25.0 J/mol ^o C and 20.8 J/mol ^o C,
respe	ectively. The enthalpy of vaporization of argon is 6506 J/mol. How much energy is
requi	ired to convert 1 mole of liquid Ar from 5 °C below its boiling point to 1 mole of ous Ar at 5 °C above its boiling point?
\bigcirc	6631 J
~	
0	6735 J
	6610 J

- O 229 J
- O 125 J