

**BMED 2210**

Fall 2015, Midterm II

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Name: ANSWER KEY

This is a closed book exam. Paper notes are allowed as well as calculators. Show ALL work and express numeric answers using the correct number of significant figures. Reference tables are found on the last page of the exam.

Good luck!

Question #1 \_\_\_\_\_ /25

Question #2 \_\_\_\_\_ /25

Question #3 \_\_\_\_\_ /25

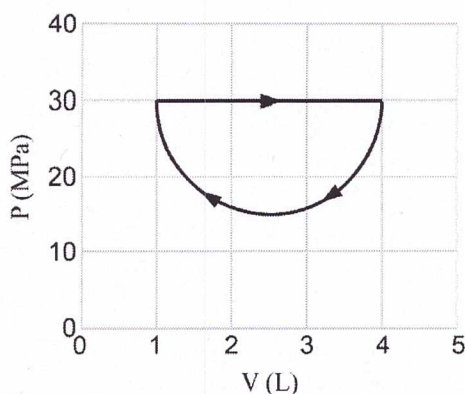
Question #4 \_\_\_\_\_ /25

Total \_\_\_\_\_ /100

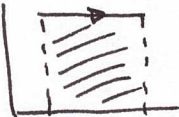


$$\bar{x} = 52.2$$

$$\sigma = 12.2$$

1. [25 pts] Gas within a chamber undergoes the complete cycle shown in the PV diagram below. If the cycle is operated at 20 Hz, calculate the net heat added to the system in one hour.



$W = \int P dV =$  area under curve of PV diagram +5 for recognizing that Work is area under curve

$\Rightarrow W_{\text{cycle}} =$    $+$    $=$    $W_{\text{cycle}} > 0$  (since  $W_1 > W_2$ )

Area of half circle:  $\frac{1}{2} \pi r^2 = \frac{1}{2} \pi \left(\frac{3}{2}\right)^2$

+5 for recognizing area under curve is a half circle (+3 for other functions)  $= \frac{9\pi}{8}$  area units

Each unit square has units of energy given by:

$(10 \text{ MPa})(1 \text{ L}) = (10^7 \text{ Pa})(0.001 \text{ m}^3) = 10^4 \text{ J}$

This means the work done per cycle is:

$\left(\frac{9\pi}{8}\right)(10^4 \text{ J}) = W_{\text{cycle}}$  +5 for correct  $W_{\text{cycle}}$  (+3 for correct idea;  $W = W_1 + W_2$ )

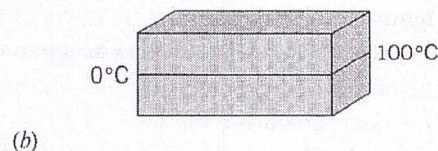
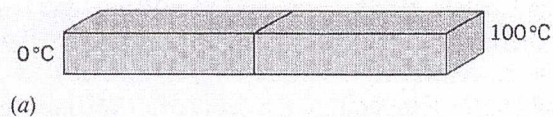
For a cycle we know  $\Delta E_{\text{sys}} = 0 \Rightarrow Q_{\text{cycle}} = W_{\text{cycle}}$  +5 for cons. of energy for closed system

$\Rightarrow$  Total heat added to system in one hour:

$Q_{\text{TOTAL}} = (20 \text{ Hz}) \left(\frac{60 \text{ sec}}{1 \text{ min}}\right) \left(\frac{60 \text{ min}}{1 \text{ hr}}\right) Q_{\text{cycle}}$

$Q_{\text{TOTAL}} = 2.5 \times 10^9 \text{ J}$  +5 for correct answer (+3 for propagating error)

2a. [18 pts] Two identical rectangular rods of metal are welded end to end as shown in (a), and 10 J of heat flows through the rods in 2.0 min. How long would it take 30 J to flow through the rods if they are welded as shown in (b)?



1-D heat flow:

$$\dot{Q} = -\frac{kA}{L} \Delta T$$

For (a) :  $\dot{Q}_a = -\frac{kA}{L} \Delta T = \frac{10 \text{ J}}{2.0 \text{ min}}$

For (b) :  $\dot{Q}_b = -\frac{k(2A)}{(\frac{1}{2}L)} \Delta T = -4 \frac{kA}{L} \Delta T = 4 \dot{Q}_a$

*+5 for recognizing area of (b) is twice that of (a)*

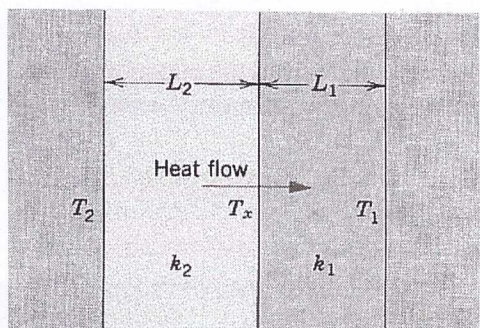
*+5 for recognizing that length of (a) is 2 times that of (b)*

$$\Rightarrow \dot{Q}_b = \frac{30 \text{ J}}{x \text{ min}} = 4 \left( \frac{10 \text{ J}}{2.0 \text{ min}} \right)$$

$$\Rightarrow x = \frac{30 \text{ J}}{20 \text{ J/min}} = \boxed{1.5 \text{ min}}$$

*+8 for correct answer  
(+6 for right expression but wrong math)*

b. [7 pts] The heat that flows across the wet suit of a scuba diver may be modeled as a compound slab where  $T_2$  is the core body temperature,  $L_2$  is the average thickness of skin,  $L_1$  is the thickness of the wet suit, and  $T_1$  is the temperature of the ocean. Circle one equation that correctly expresses the outer surface temperature  $T_x$  of the skin (Hint: use reasoning, not math).



a.  $T_x = \frac{R_1 T_1 + R_2 T_2}{R_1 + R_2}$

c.  $T_x = \frac{R_1 T_1 + R_2 T_2}{R_1 R_2}$

b.  $T_x = \frac{R_1 T_2 + R_2 T_1}{R_1 + R_2}$

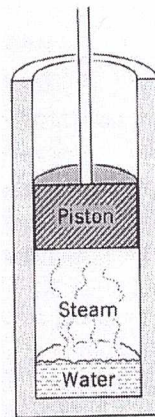
d.  $T_x = \frac{R_1 T_2 + R_2 T_1}{R_1 R_2}$

When  $R_1 = 0$ , then  $T_x = T_1$   
 & when  $R_2 = 0$ , then  $T_x = T_2$

*+3.5 if ~~(a)~~ (b) circled* <sup>3/9</sup> *(c and d doesn't make sense when  $R_1$  or  $R_2 = 0$  since dividing by 0)*  
*+7 if (b) circled*



3. [25 pts total] A cylinder has a well-fitted 2.0-kg metal piston whose cross-sectional area is  $2.0 \text{ cm}^2$ . The cylinder contains water and steam at constant temperature. The piston is observed to fall slowly at a rate of  $0.30 \text{ cm/s}$  because heat flows out of the cylinder through the cylinder walls. As this happens, some steam condenses in the chamber. The density of the steam inside the chamber is  $6.0 \times 10^{-4} \text{ g/cm}^3$  and the atmospheric pressure is  $1.0 \text{ atm}$ .



a. [10 pts] Calculate the rate of condensation of steam (Hint: What properties of the steam, which behaves like an ideal gas, are not changing?).

For the steam, we know that

$$PV = nRT$$

here the pressure is constant (since piston falls slowly, i.e. reversible process  $\Rightarrow P_{\text{int}} = P_{\text{ext}}$ ) and temperature is constant from the problem statement.

$$\Rightarrow \left(\frac{n}{V}\right) = \frac{P}{RT} = \text{constant} \quad \begin{array}{l} +5 \text{ for } \left(\frac{n}{V}\right) = \text{constant} \\ (+3 \text{ for noticing } P \text{ and } T \text{ are constant}) \end{array}$$

$$\text{since } \rho_{\text{steam}} = \frac{m_{\text{steam}}}{V} = \frac{M_{\text{H}_2\text{O}} n_s}{V} = M_{\text{H}_2\text{O}} \left(\frac{n_s}{V}\right)$$

then the  $\rho_{\text{steam}}$  inside chamber is constant, during conversion to water.

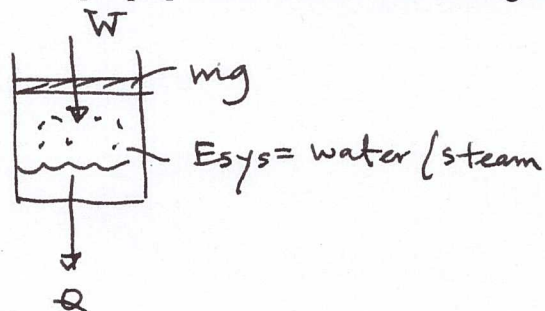
$$\Rightarrow m_{\text{steam}} = \rho_{\text{steam}} V$$

$$\Rightarrow \frac{dm_s}{dt} = \frac{d}{dt}(\rho_s V) = \rho_s A \frac{dx}{dt} \quad \begin{array}{l} \text{cross section of piston} \\ +3 \text{ for expression of } \frac{dm_s}{dt} \\ \sim \text{speed at which piston falls} \end{array}$$

$$\frac{dm_s}{dt} = -(6.0 \times 10^{-4} \text{ g/cm}^3)(2.0 \text{ cm}^2)(0.30 \text{ cm/s})$$

$$= \boxed{-3.6 \times 10^{-4} \frac{\text{g steam}}{\text{s}}}$$

a. [15 pts] At what rate is heat leaving the chamber?



cons. of energy

+5 for energy balance

$$\Delta \dot{E}_{\text{sys}} = \dot{Q} - \dot{W}$$

constant pressure work

heat flowing out of system

internal energy of steam / water

$$\begin{aligned} \Delta \dot{E}_{\text{sys}} &= (\text{heat of condensation}) \left( \frac{dm}{dt} \right) \\ &= (2256 \text{ kJ/kg}) (-3.6 \times 10^{-4} \text{ g/s}) \left( \frac{1 \text{ kg}}{1000 \text{ g}} \right) \\ &= \boxed{-8.1 \times 10^{-4} \text{ kJ/s}} \end{aligned}$$

+2 for  $\Delta \dot{E}_{\text{sys}}$

$$\dot{W} = -P A \frac{dx}{dt} \quad (\text{volume contracting})$$

$$= - \left( 1 \text{ atm} + \frac{mg}{A} \right) (A) \frac{dx}{dt}$$

$$= - \left( 1.01 \times 10^5 \text{ Pa} + \frac{(2 \text{ kg})(9.8 \text{ m/s}^2)}{(2 \text{ cm}^2) \left( \frac{1 \text{ m}}{100 \text{ cm}} \right)^2} \right) \left( 2 \text{ cm}^2 \left( \frac{1 \text{ m}}{100 \text{ cm}} \right)^2 \right) \left( \frac{0.003 \text{ m}}{\text{s}} \right)$$

$$= - (1.99 \times 10^5 \text{ Pa}) (6 \times 10^{-7} \text{ m}^3/\text{s})$$

$$= \boxed{-0.12 \text{ J/s}}$$

+6 for  $\dot{W}$   
(+4 for correct approach but wrong values)

$$\dot{Q} = \Delta \dot{E}_{\text{sys}} + \dot{W} = (-8.1 \times 10^{-4} \text{ kJ/s}) - 0.12 \text{ J/s}$$

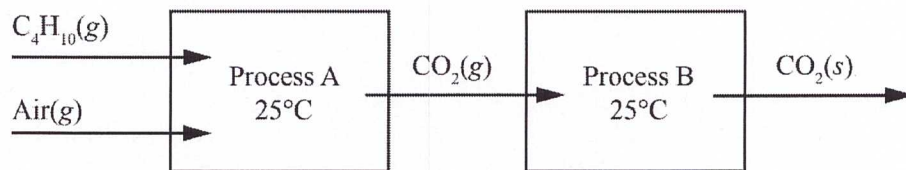
$$= \boxed{-0.93 \text{ J/s}}$$

(i.e. vast majority of heat flow out is used to lower piston)

+2 for  $\dot{Q}$



4. [25 pts total] An industrial two-step process to make dry ice starts with the combustion of *n*-Butane with dry air (79.0% nitrogen and 21.0% oxygen by volume) to form  $\text{CO}_2(\text{g})$  in Process A, which is then converted to  $\text{CO}_2(\text{s})$  in a second Process B. *n*-Butane is fed at 10 kg per second and dry air at 200.0 kg per second, and *n*-Butane flows out at Process A at 1.2 kg per second. The standard heat of formation of dry ice is  $-427.4 \text{ kJ/mol}$ . Both processes are conducted at  $25^\circ\text{C}$ .



a. [15 pts] Find the reaction rate  $R$  for Process A.

First balance stoichiometry for the combustion process in A:  
 $\text{C}_4\text{H}_{10}(\text{g}) + \frac{13}{2}\text{O}_2(\text{g}) \longrightarrow 4\text{CO}_2(\text{g}) + 5\text{H}_2\text{O}(\text{e})$  +5 for correct stoichiometry

Find limiting reagent:

$$\min \left\{ \frac{\dot{n}_{i,s}}{\sigma_s} \right\} \Rightarrow \dot{n}_{\text{butane}} = \left( \frac{10,000 \text{ g}}{\text{s}} \right) \left( \frac{1 \text{ mol}}{58 \text{ g/mol}} \right) = \underline{172.4 \frac{\text{mol}}{\text{s}}}$$

for  $\text{O}_2(\text{g})$ , need  $w_{\text{O}_2}$  +5 for calculating  $w_{\text{O}_2}$

$$\Rightarrow w_{\text{O}_2} = \frac{(0.21)(32 \text{ g/mol})}{(0.21)(32 \text{ g/mol}) + (0.79)(28 \text{ g/mol})} = 0.233$$

$$\Rightarrow \dot{n}_{\text{O}_2} = w_{\text{O}_2} \dot{m}_{\text{air}} / M_{\text{O}_2} = (0.233) \left( \frac{200,000 \text{ kg}}{\text{s}} \right) \left( \frac{1 \text{ mol O}_2}{32 \text{ g/mol}} \right)$$

$$= 1.46 \times 10^3 \text{ mol/s}$$

$$\Rightarrow \left\{ \frac{\dot{n}_{i,\text{O}_2}}{\sigma_{\text{O}_2}} \right\} = \frac{1.46 \times 10^3 \frac{\text{mol}}{\text{s}}}{13/2} = 224$$

$$\left\{ \frac{\dot{n}_{i,\text{butane}}}{\sigma_{\text{butane}}} \right\} = \frac{172.4 \text{ mol/s}}{1} = 172.4 \Rightarrow \text{butane limiting reagent.}$$

$$\Rightarrow R = \frac{\dot{n}_{i,\text{butane}} - \dot{n}_{e,\text{butane}}}{-\sigma_{\text{butane}}} = \frac{(172.4 \text{ mol/s}) - (1.2 \text{ kg}) / (58 \text{ g/mol})}{-(-1)}$$

$$\boxed{R = 151.7 \text{ mol/s}}$$

+5 for R

(+3 for correct expression)  
but wrong values

b. [10 pts] Determine the heat released into the environment every hour for the entire process.

Process A:

$$\Delta H_{\text{rxn}} = \Delta \hat{H}_{\text{C, butane}}^{\circ} R$$

$$= (-2878.52 \text{ kJ/mol}) (151.7 \text{ mol/s})$$

$$= \boxed{-4.4 \times 10^5 \text{ kJ/s}}$$

+3 for  $\Delta H_{\text{rxn}}$  (Process A)  
(+1.5 for incorrect values  
but correct approach)

Process B:

$$\Delta \hat{H}_{\text{rxn}} = \Delta \hat{H}_{\text{f CO}_2(\text{s})}^{\circ} - \Delta \hat{H}_{\text{f CO}_2(\text{g})}^{\circ}$$

$$= (-427.4 \text{ kJ/mol}) - (-393.51 \text{ kJ/mol})$$

$$= -33.89 \text{ kJ/mol CO}_2(\text{g})$$

$$\Rightarrow \Delta H_{\text{rxn}} = (4 \text{ mol CO}_2) (-33.89 \text{ kJ/mol}) \dot{n}_{\text{CO}_2(\text{g})}$$

$$= (4) (-33.89 \text{ kJ/mol}) (151.7 \text{ mol/s})$$

$$= \boxed{-2.1 \times 10^4 \text{ kJ/s}}$$

+3 for  $\Delta H_{\text{rxn}}$  (process B)  
(+1.5 for correct approach)

$$\Delta H_{\text{rxn total}} = -4.4 \times 10^5 - 2.1 \times 10^4 = \boxed{-4.6 \times 10^5 \text{ kJ/s}}$$

+2 for  $\Delta H_{\text{rxn}} = \Delta H_A + \Delta H_B$

$\Rightarrow$  Every hour:

$$\left( -4.6 \times 10^5 \frac{\text{kJ}}{\text{s}} \right) \left( \frac{60 \text{ s}}{1 \text{ min}} \right) \left( \frac{60 \text{ min}}{1 \text{ hr}} \right) = \boxed{-1.7 \times 10^9 \text{ J/hr}}$$

+2 for final answer

END OF EXAM



## Reference Tables

### Factors for Unit Conversion

Quantity	Equivalent Values
Mass	$1 \text{ kg} = 1000 \text{ g} = 0.001 \text{ metric ton} = 2.20462 \text{ lb}_m = 35.27392 \text{ oz}$ $1 \text{ lb}_m = 16 \text{ oz} = 5 \times 10^{-4} \text{ ton} = 453.593 \text{ g} = 0.453593 \text{ kg}$
Length	$1 \text{ m} = 100 \text{ cm} = 1000 \text{ mm} = 10^6 \text{ microns } (\mu\text{m})$ $= 39.37 \text{ in} = 3.2808 \text{ ft} = 1.0936 \text{ yd} = 0.0006214 \text{ mile}$ $1 \text{ ft} = 12 \text{ in} = 1/3 \text{ yd} = 0.3048 \text{ m} = 30.48 \text{ cm}$
Volume	$1 \text{ m}^3 = 1000 \text{ L} = 10^6 \text{ cm}^3 = 10^6 \text{ mL}$ $= 35.3145 \text{ ft}^3 = 220.83 \text{ imperial gallons} = 264.17 \text{ gal}$ $= 1056.68 \text{ qt}$ $1 \text{ ft}^3 = 1728 \text{ in}^3 = 7.4805 \text{ gal} = 0.028317 \text{ m}^3 = 28.317 \text{ L}$ $= 28,317 \text{ cm}^3$
Force	$1 \text{ N} = 1 \text{ kg} \cdot \text{m/s}^2 = 10^5 \text{ dynes} = 10^5 \text{ g} \cdot \text{cm/s}^2 = 0.22481 \text{ lb}_f$ $1 \text{ lb}_f = 32.174 \text{ lb}_m \cdot \text{ft/s}^2 = 4.4482 \text{ N} = 4.4482 \times 10^5 \text{ dynes}$
Pressure	$1 \text{ atm} = 1.01325 \times 10^5 \text{ N/m}^2 (\text{Pa}) = 101.325 \text{ kPa} = 1.01325 \text{ bar}$ $= 1.01325 \times 10^6 \text{ dynes/cm}^2$ $= 760 \text{ mmHg at } 0^\circ\text{C (torr)} = 10.333 \text{ m H}_2\text{O at } 4^\circ\text{C}$ $= 14.696 \text{ lb}_f/\text{in}^2 (\text{psi}) = 33.9 \text{ ft H}_2\text{O at } 4^\circ\text{C}$ $= 29.921 \text{ in Hg at } 0^\circ\text{C}$
Energy	$1 \text{ J} = 1 \text{ N} \cdot \text{m} = 10^7 \text{ ergs} = 10^7 \text{ dyne} \cdot \text{cm}$ $= 2.778 \times 10^{-7} \text{ kW} \cdot \text{hr} = 0.23901 \text{ cal}$ $= 0.7376 \text{ ft} \cdot \text{lb}_f = 9.486 \times 10^{-4} \text{ Btu}$
Power	$1 \text{ W} = 1 \text{ J/s} = 0.23901 \text{ cal/s} = 0.7376 \text{ ft} \cdot \text{lb}_f/\text{s}$ $= 9.486 \times 10^{-4} \text{ Btu/s} = 1.341 \times 10^{-3} \text{ hp}$

Example: The factor to convert grams to  $\text{lb}_m$  is  $\left(\frac{2.20462 \text{ lb}_m}{1000 \text{ g}}\right)$ .

																Noble Gases	
IA	IIA	IIIB	IVB	VB	VIB	VII	VIII	VIII	IB	IIB	IIIA	IVA	VA	VIA	VIIA		
1 H 1.00794																2 He 4.00260	
3 Li 6.941	4 Be 9.01218											5 B 10.81	6 C 12.011	7 N 14.0067	8 O 15.9994	9 F 18.9984	10 Ne 20.1797
11 Na 22.98977	12 Mg 24.305											13 Al 26.98154	14 Si 28.0855	15 P 30.9738	16 S 32.066	17 Cl 35.4527	18 Ar 39.948
19 K 39.0983	20 Ca 40.078	21 Sc 44.9559	22 Ti 47.88	23 V 50.9415	24 Cr 51.996	25 Mn 54.9380	26 Fe 55.847	27 Co 58.9332	28 Ni 58.69	29 Cu 63.546	30 Zn 65.39	31 Ga 69.72	32 Ge 72.59	33 As 74.9216	34 Se 78.96	35 Br 79.904	36 Kr 83.80
37 Rb 85.4678	38 Sr 87.62	39 Y 88.9059	40 Zr 91.224	41 Nb 92.9064	42 Mo 95.94	43 Tc (98)	44 Ru 101.07	45 Rh 102.9055	46 Pd 106.42	47 Ag 107.8682	48 Cd 112.41	49 In 114.82	50 Sn 118.710	51 Sb 121.75	52 Te 127.60	53 I 126.9045	54 Xe 131.29
55 Cs 132.9054	56 Ba 137.33	57* La 138.9055	72 Hf 178.49	73 Ta 180.9479	74 W 183.85	75 Re 186.207	76 Os 190.2	77 Ir 192.22	78 Pt 195.08	79 Au 196.9665	80 Hg 200.59	81 Tl 204.383	82 Pb 207.2	83 Bi 208.9804	84 Po (209)	85 At (210)	86 Rn (~222)
87 Fr (223)	88 Ra 226.0254	89† Ac 227.0278	104 Unq (261)	105 Unp (262)	106 Unh (263)												



Compound	Formula	Mol. wt.	State	$\Delta\hat{H}_f^\circ$ (kJ/g mol)	$\Delta\hat{H}_c^\circ$ (kJ/g mol)
n-Butane	C <sub>4</sub> H <sub>10</sub>	58.12	l	-147.6	-2855.6
Isobutane	C <sub>4</sub> H <sub>10</sub>	58.12	g	-124.73	-2878.52
			l	-158.5	-2849.0
1-Butene	C <sub>4</sub> H <sub>8</sub>	56.104	g	-134.5	-2868.8
			g	1.172	-2718.58
Calcium arsenate	Ca <sub>3</sub> (AsO <sub>4</sub> ) <sub>2</sub>	398.06	c	-3330.5	
Calcium carbide	CaC <sub>2</sub>	64.10	c	-62.7	
Calcium carbonate	CaCO <sub>3</sub>	100.09	c	-1206.9	
Calcium chloride	CaCl <sub>2</sub>	110.99	c	-794.9	
Calcium cyanamide	CaCN <sub>2</sub>	80.11	c	-352	
Calcium hydroxide	Ca(OH) <sub>2</sub>	74.10	c	-986.56	
Calcium oxide	CaO	56.08	c	-635.6	
Calcium phosphate	Ca <sub>3</sub> (PO <sub>4</sub> ) <sub>2</sub>	310.19	c	-4137.6	
Calcium silicate	CaSiO <sub>3</sub>	116.17	c	-1584	
Calcium sulfate	CaSO <sub>4</sub>	136.15	c	-1432.7	
Calcium sulfate (gypsum)	CaSO <sub>4</sub> · 2H <sub>2</sub> O	172.18	aq	-1450.5	
Carbon	C	12.01	c	-2021.1	
Carbon dioxide	CO <sub>2</sub>	44.01	Graphite (β)	0	-393.51
			g	-393.51	
Carbon disulfide	CS <sub>2</sub>	76.14	l	-412.92	
			l	87.86	-1075.2

TABLE 2 SOME HEATS OF TRANSFORMATION

Substance <sup>a</sup>	Melting Point (K)	Heat of Fusion (kJ/kg)	Boiling Point (K)	Heat of Vaporization (kJ/kg)
Hydrogen	14.0	58.6	20.3	452
Oxygen	54.8	13.8	90.2	213
Mercury	234	11.3	630	296
Water	273	333	373	2256
Lead	601	24.7	2013	858
Silver	1235	105	2485	2336
Copper	1356	205	2840	4730