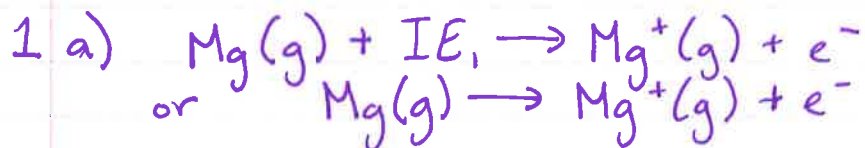


## Practice test 2 Key

- 1) a
- 2) a
- 3) b
- 4) c
- 5) a

- 6) b
- 7) a
- 8) e
- 9) d
- 10) c



b) Each IE takes one  $e^-$ . After the second IE ( $\text{IE}_2$ ) you have removed both valence electrons from Mg.  $\text{IE}_3$  is the energy required to take a 3rd electron from Mg which is from its closed (stable) core.

c) 738 kJ/mol

$$\frac{738 \text{ kJ}}{1 \text{ mol}} \left( \frac{10^3 \text{ J}}{1 \text{ kJ}} \right) \left( \frac{1 \text{ mol}}{6.022 \times 10^{23}} \right) = 1.226 \times 10^{-18} \text{ J}$$

$$E = \frac{h \cdot c}{\lambda} \Rightarrow \lambda = \frac{h \cdot c}{E} = \boxed{1.62 \times 10^{-7} \text{ m}}$$

$$\text{or } \boxed{162 \text{ nm}}$$

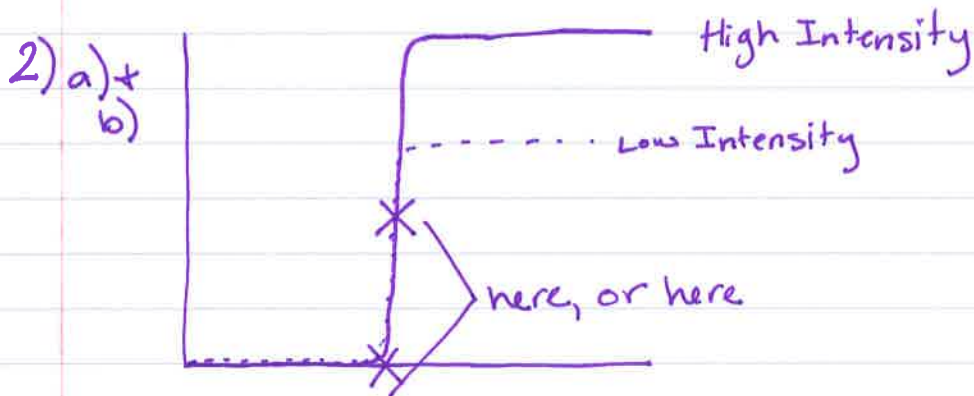
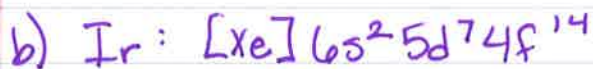
$$2 \quad \frac{1}{\lambda} = R \left( \frac{1}{m^2} - \frac{1}{n^2} \right)$$

$$\frac{1}{4652.84 \text{ nm}} = 1.097 \times 10^{-2} \text{ nm}^{-1} \left( \frac{1}{m^2} - \frac{1}{7^2} \right)$$

$$\frac{1}{m^2} = .04$$

$$\boxed{m = 5}$$

Short Answer

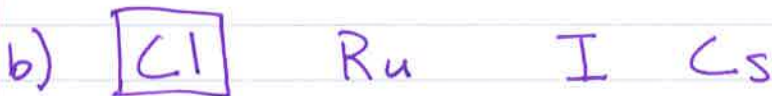


c) The extra energy is converted to kinetic energy in the  $e^-$ . So, the extra energy will go toward the velocity of the  $e^-$  as it is ejected.



these 3 are isoelectronic, so most protons means smallest

Just using trend makes you know Kr is smaller than Rb

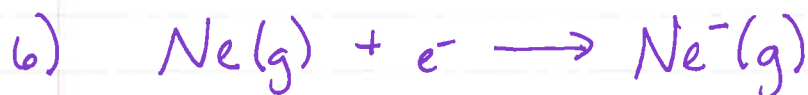


Just trend alone here.

4) Be has a full s-subshell which is stable. This stability causes it to require more energy to remove than ~~the~~ the p-electron of B despite the trend.

5) a)  $\boxed{F}$  s Br Zn

b) Si P  $\boxed{N}$  C



Because you are having to begin filling a new shell in order to add  $1e^-$ .

### \*Naming\*

- diphosphorus pentasulfide
- HF(aq)
- lithium sulfide
- chromium(III) phosphate
- $Mn(C_2H_3O_2)_2$

### \*Problems\*

1)  $23.7g K_3PO_4 \left( \frac{1 mol K_3PO_4}{212.27g K_3PO_4} \right) \left( \frac{3 mol K}{1 mol K_3PO_4} \right) \left( \frac{39.10g K}{1 mol K} \right)$

$= \boxed{13.1g K}$

2)  $12.465 \text{ g CO}_2 \left( \frac{1 \text{ mol CO}_2}{44.01 \text{ g CO}_2} \right) \left( \frac{1 \text{ mol C}}{1 \text{ mol CO}_2} \right) = .2832 \text{ mol C} *$

$7.656 \text{ g H}_2\text{O} \left( \frac{1 \text{ mol H}_2\text{O}}{18.02 \text{ g H}_2\text{O}} \right) \left( \frac{2 \text{ mol H}}{1 \text{ mol H}_2\text{O}} \right) = .8497 \text{ mol H} *$

$.2832 \text{ mol C} \left( \frac{12.01 \text{ g C}}{1 \text{ mol C}} \right) = 3.4016 \text{ g C}$

$.8497 \text{ mol H} \left( \frac{1.01 \text{ g H}}{1 \text{ mol H}} \right) = .858197 \text{ g H}$

$14.300 \text{ g sample} - 3.4016 \text{ g C} - .858197 \text{ g H}$   
 $= 10.04 \text{ g Cl}$

$10.04 \text{ g Cl} \left( \frac{1 \text{ mol Cl}}{35.45 \text{ g Cl}} \right) = .2832 \text{ mol Cl} *$

$\text{C} \frac{.2832}{.2832} \text{ H} \frac{.8497}{.2832} \text{ Cl} \frac{.2832}{.2832}$

