Ge 212 Problem Set 2

Due Wednesday October 14, 2015

1. In class it was (or will be?) asserted that, for any stable or metastable material, the reversible adiabats are steeper in *P*-V space than the isothermals. Once we learn the second law we will see that this is equivalent to the statement (∂V/∂*P*)S > (∂V/∂*P*)*T*. Use the table of Jacobians (<http://www.asimow.com/jacobiantable.jpg>) and the constraint (coming later) that *Cp* > 0 for any stable or metastable material to prove that this is always true, regardless of the sign of (∂V/∂*T*)*P*. It is possible for (∂*V*/∂*T*)*P* to be zero as well; what happens then?
2. In class we used a stream of red paint being dripped into a pot of green paint, stirred at some steady rate, to show the idea that there exists a relaxation time against which we can judge whether any spontaneous change occurs. Question: according to the definitions given in class, is the case of very slow dripping, where the paint in the pot is always uniform in color, an example of a *reversible* process or a *quasistatic* process? **Explain**.
3. Consider an electric circuit consisting of a capacitor with capacitance *C* (in farads, 1 farad = 1 coulomb/volt) in series with a resistor with resistance *R* (in ohms, 1 ohm = 1 volt/amp, and 1 amp = 1 coulomb/second). Let *V*0 be the voltage to the left of the capacitor, *V*1 the voltage between capacitor and resistor, and *V*2 the voltage to the right of the resistor. So the charge on the capacitor is C(*V*0 – *V*1) and the current across the resistor is (*V1 – V2)*/*R.* When the left and right sides are shorted so that *V*0 = *V*2, this circuit obeys the equation *C*(d(*V0 – V1)*/d*t*) – (*V1 – V0)*/*R* = 0 and its only equilibrium state is *V*1 = *V*0 = *V*2 with no charge on the capacitor. Say instead we connect the ends of this circuit to a variable voltage source so we can apply a difference V2 – V0 and we want to charge up the capacitor *reversibly…*
	1. What does it mean to change voltage reversibly in this case? What is a criterion for assessing whether the change in reversible?
	2. What is the value of the time constant  we should use to evaluate whether we are approaching reversible changes or not?
4. Say we have a sphere of matter with thermal conductivity *k* (in W/m/K), heat capacity *C*p (in J/K/kg), density  (in kg/m3), and radius *R* (in m). Say we want to heat this sphere *reversibly* by changing the surface temperature…
	1. What does it mean to change temperature reversibly in this case? What is a criterion for assessing whether the change in reversible?
	2. What is the characteristic time  that we have to think about to determine whether a temperature change will approach reversibility?
5. Let’s think about component transformations for a little while…Consider the component sets **n** = {CaO, Al2O3, SiO2} and **n’** = {CaSiO3, Al2SiO5, CaAl2Si2O8}. Write the matrix ****i’i that maps a composition in terms of component set **n’** to **n**. In class I asserted that the reverse transformation matrix ****ii’ = {****i’i}–1. Is this true here? If not, what is wrong with this example?
6. *Component transformation in real life*

In the MELTS model, pyroxenes are represented as a seven-component system using these components:

 En Enstatite Mg2Si2O6

 Di Diopside CaMgSi2O6

 Hd Hedenbergite CaFe2+Si2O6

 Jd Jadeite NaAlSi2O6

 Abf Alumino-buffonite\* CaTi0.5Mg0.5AlSiO6

 Bf Buffonite CaTi0.5Mg0.5Fe3+SiO6

 Es Essenite CaFe3+AlSiO6

(\* I am not making this up)

a. How many oxides are present in this model (counting FeO and Fe2O3 separately)?

b. Consider that you are given a composition in terms of moles of the 7 components, which you can write as a vector {En, Di, Hd, Jd, Abf, Bf, Es}. Give me the transformation matrix you would premultiply by this vector in order to obtain this composition in moles of oxides (make sure you specify the order of the oxides in the resulting vector, please!).

c. Now consider the reverse problem, which occurs in real life when you measure a pyroxene composition by electron probe and Mössbauer or by wet chemistry and then need to express it in pyroxene components: given the composition as a vector of moles of oxides, is there a matrix you can multiply by in order to obtain the composition in moles of the 7 components? If not (hint, hint), why not? – please explain this both mathematically and *conceptually*.

d. Assuming you answered ‘no’ to the first part of (c), propose a reasonable method to accomplish the task of converting from an oxide analysis to component moles. Is there a sense in which your method does this task as well as possible?