PHA 5127 (Fall, 2008) Case Study #4

- Q1. For the following situations, indicate whether the drug is: filtered, reabsorbed, actively secreted, or reabsorbed through transporters (Assume GFR is 130 mL min⁻¹, urine flow is 1.5 ml min⁻¹)
 - a). A drug with fu = 0.02 and a $Cl_{REN} = 20$ mL min⁻¹
 - b). A drug with fu = 0.40 and a $Cl_{REN} = 52$ mL min⁻¹
 - c). A drug with fu = 0.60 and a Cl_{REN} = 0.9 mL min⁻¹
 - d). A drug with fu = 1.0 and a $Cl_{REN} = 0.3$ mL min⁻¹
- Q2. A male patient is 5 ft 10 inches tall, 40 years old, and weights 80 kg. His serum creatinine is 1.5 mg/dl. Please estimate his GFR.
- Q3. Mark each of the following statements True or False.
 - T F The maximum value of renal clearance can not excess the glomerula filtration rate.
 - T F The renal clearance of a drug (as determined by filtration and reabsorbtion) always depends on the tissue binding of the drug.
 - T F Drinking a lot of water (urine flow is doubled) will increase significantly the renal clearance of aminoglycocsides.
 - T F For an acidic drug with a pka of 1.0, adjustment of the urine pH within physiological ranges will significantly change the renal clearance.
 - T F To determine the clearance of a drug, one needs to know whether the drug is a one or two compartment drug.
 - T F Since creatinine is endogenous and predominantly eliminated by kidney, its clearance is a good estimation of renal active secretion.
 - T F The larger the volume of distribution, the smaller the AUC of a given drug.
- Q4. Define the term linear pharmacokinetics.