

PHA 5127

FALL 2013

Case Study 2

True (A) or False (B)

- 1) A lipophilic drug cannot have a volume of distribution that is smaller than V_T (T/F)
- 2) The pK_a of an acidic drug that shows perfusion limited distribution into tissues is likely to be small. (T/F)
- 3) The same dose of a drug is given orally either as a solution or in form of a slow dissolving crystal suspension. The solution will show higher maximum concentrations in plasma. (T/F)

Assume no active transport.

- 4) T/ F Compared to fat, the liver is likely to have a higher rate of uptake for small lipophilic drugs due to its higher blood flow rate.
- 5) T /F The rate with which hydrophilic compounds will move across well-built membranes will depend on the concentration gradient between total drug in plasma and total drug in tissue.
- 6) T /F Permeability limited distribution is generally seen for small, lipophilic drugs
- 7) T/F Free drug concentrations are always the same in plasma and tissues, when the distribution occurs instantaneously
- 8) A drug (lipophilic, unionized, low molecular weight) is showing in average a pronounced binding to plasma proteins of 99%. Between-subject variability of protein binding is pronounced. 1000mg of the drug is given as IV bolus injection to two patients. Patient 1 has a much stronger plasma protein binding for the drug (99.995%) than the second patient (99.99%). The tissue binding is the same in both patients and is equal to 90%

Based on the given information please indicate whether **patient 1** will have a larger (↑), smaller (↓) or identical (↔) value compared to patient 2 for

- Total initial plasma drug concentration (C_0)
- Free initial plasma drug concentration (free C_0)
- f_u
- V_d

A) $C_0 \uparrow$, free $C_0 \uparrow$, $f_u \downarrow$, $V_d \downarrow$

B) $C_0 \downarrow$, free $C_0 \leftrightarrow$, $f_u \downarrow$, $V_d \uparrow$

C) $C_0 \uparrow$, free $C_0 \downarrow$, $f_u \downarrow$, $V_d \downarrow$

D) $C_0 \uparrow$, free $C_0 \uparrow$, $f_u \uparrow$, $V_d \leftrightarrow$

E) None of the above Combinations

Assume a drug is substrate of a specific transport protein. What of the following statements are True (A) or False (B).

9) T/F Transporters do not use energy

10) T/F Transporters only eliminate drug from the body

11) T/F Transporters are only present in liver and kidney

12) T/F Transporters are saturable