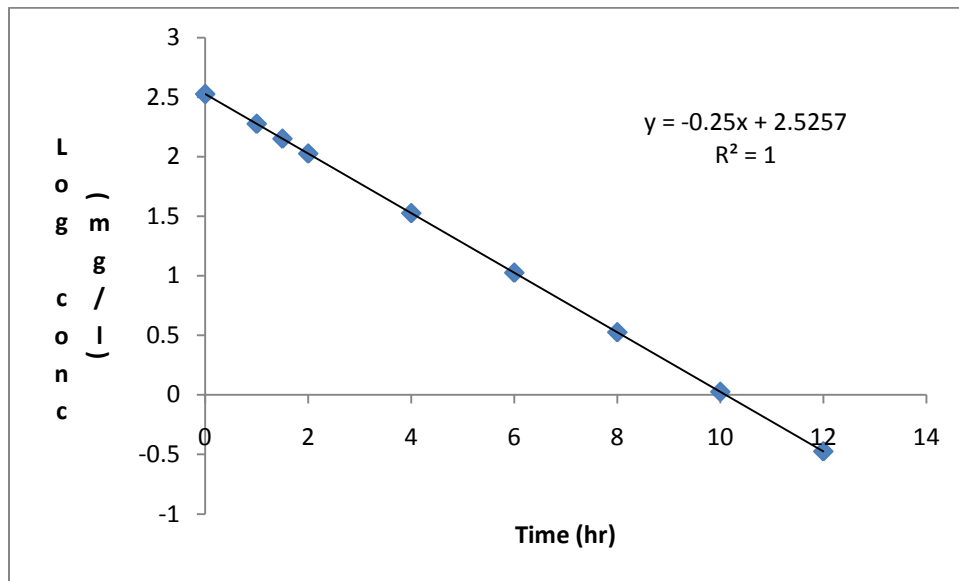


1) The table1 shows the serum concentration profiles of a certain drug in patient X. Please specify the units for the Pk parameters to get full credit.

- a) Determine if the elimination process is a first order or a zero order process. (Plot on a graph paper) [1]
- b) Calculate K_e , the first order elimination rate constant. [1]
- c) Calculate AUC 0-tlast and AUC 0-inf by trapezoidal rule. [2]
- d) Calculate the concentration of the drug X in serum at time 5hr.[1]

time(hr)	conc (mg/l)
0	12.50
1	9.74
1.5	8.59
2	7.58
4	4.60
6	2.79
8	1.69
10	1.03
12	0.62

Solution



a) It is clear from the figure that the \ln conc vs time is a straight line. So the elimination process is a **first order** process.

b) The value of k_e is **0.25 hr⁻¹**

c) The $AUC(0-12hr)$ is **48.16 mg*hr/L**

The $AUC(0-inf)$ is **50.64 mg*hr/L**

d) Using the equation $C = C_0 * \exp(-k_e * t)$

$C_0 = 12.5mg/L$, $k_e = 0.25hr^{-1}$; **$C(5hr) = 3.58mg/L$**

Q2 Derive the equation $k_e = 0.693/t_{1/2}$ for an i.v. bolus one compartment body model. [2]

Solution:

Substitute $C = C_0/2$ in the equation $C = C_0 * \exp(-k_e * t)$ and solve for the relationship between k_e and $t_{1/2}$

True or False

- a) When the change in amount of the drug in the body is related to the amount by the following equation $\frac{dX}{dt} = -K_e * X^0$, where X is the amount of the drug at a given time t , then we say the elimination is a zero order process. (T/F) [1]
- b) Drugs with a low volume of distribution have a narrow therapeutic window. (T/F) [1]
- c) When whole blood is collected in a heparinized tube and then centrifuged, the supernatant that is obtained is plasma. (T/F) [1]