

Q1) Say true or false

T **F** Clearance defines the amount of drug eliminated from body per unit time

T **F** For any drug, the AUC is affected by its dose and volume of distribution

T **F** As intrinsic clearance describes the liver's innate ability to clear unbound drug from intracellular water via metabolism or biliary excretion, the higher the liver blood flow, the higher will be a drug's clearance

T **F** For a high extraction drug that has predominant hepatic metabolism, the liver blood flow is a rate limiting step in its clearance

T **F** For a low extraction drug, the higher the protein binding the higher will be the clearance

Q2) Predict the changes in Hepatic Clearance ( $CL_H$ ) for the following scenarios

Parameter	Change occurring	Effect on low extraction drug	Effect on High extraction drug
$f_u$	↓	↓	↔
$CL_i$	↑	↑	↔
$Q_h$	↑	↔	↑

For a low extraction drug :

$$CL_H = \frac{Q_H \cdot f_u \cdot CL_{int}}{Q_H + f_u \cdot CL_{int}}$$

$$CL_H = \frac{Q_H \cdot f_u \cdot CL_{int}}{Q_H} = f_u \cdot CL_{int}$$

For a high extraction drug :

$$CL_H = \frac{Q_H \cdot f_u \cdot CL_{int}}{Q_H + f_u \cdot CL_{int}}$$

$$CL_H = Q_H$$

Q3) Predict changes in Oral bioavailability for the following scenarios

Parameter	Change occurring	Effect on low extraction drug	Effect on High extraction drug
$f_u$	↓	↔	↑
$CL_i$	↑	↔	↓
$Q_h$	↑	↔	↑

For a low extraction drug:

$$F = \frac{Q_H}{Q_H + f_u \cdot CL_{int}} \quad F = 1$$

For a high extraction drug:

$$F = \frac{Q_H}{Q_H + f_u \cdot CL_{int}} \quad F = \frac{Q_H}{f_u \cdot CL_{int}}$$

Q4) A patient is to be started on two medications (A and B) administered by IV bolus injections. Blood samples were taken at 1 and 4 hours following the first injections of drug A or B alone in order to determine whether concentrations were in an appropriate range for each drug. See table below for these levels and additional information.

Drug	Dose (mg)	$C_p$ at 1h (mg/L)	$C_p$ at 4h (mg/L)	$E_H$	$f_u$
A	400	1.22	0.76	0.1	0.3
B	1200	0.92	0.51	0.8	0.1

Assume liver blood flow of 90 L/h, where  $E_H$  is the extraction ratio and  $f_u$  is the fraction unbound. Renal elimination and hepatic metabolism are the only clearance pathways for both of these drugs.

- Calculate the  $CL$ (hepatic) for Drug B and  $CL$  (renal) for Drug B
- Also calculate the  $AUC_{0-\infty}$  Drug B

$$K_e = \ln C_2 - \ln C_1 / (t_2 - t_1) = 0.20 \text{ /hr}$$

$$T_{1/2} = 0.693 / 0.2 = 3.45 \text{ hr}$$

$$C_0 = C_t / \exp(-k_e \cdot t)$$

$$C_0 = 1.13 \text{ mg/L}$$

$$V_d = \text{dose} / C_0$$

$$V_d = 1200 / 1.13 = 1062 \text{ L}$$

$$CL (\text{hepatic}) = Q_H * E_H = 90 * 0.8 = 72 \text{ L/hr}$$

$$Cl(\text{tot}) = V_d * K_e = 1062 * 0.20 = 212.4 \text{ L/hr}$$

$$CL (\text{renal}) = Cl (\text{total}) - Cl (\text{hepatic}) = 212.4 - 72 = 140.4 \text{ L/hr}$$

$$AUC_{0-\infty} = \text{dose} / Cl(\text{tot}) = 1200 / 212.4 = 5.65 \text{ mg hr/L}$$

We cannot use just the hepatic clearance to calculate the AUC, because the drug is cleared extra-hepatically as well