## **ABBREVIATIONS**

These are included for reference only - do not try to memorise them!

 $\alpha$  = slope of the component of the plasma concentration / time curve attributable to distribution. Used for predicting C

 $\mathbf{A}$  = the intercept of this line on the Y axis. Used for predicting C,

AUC = area under the plasma concentration / time curve

 $AUC_{0-inf}$  = area under the plasma concentration / time curve extrapolated to infinity

 $AUC_{0.12}$  = area under the plasma concentration / time curve for the first 12 hours

**AUMC** = area under the moment curve. A theoretical concept used for deriving the MRT.

 $\beta$  = slope of the component of the plasma concentration / time curve attributable to elimination. Used for predicting C

 $\mathbf{C}_{_{t}}$   $\mathbf{B}$  = the intercept of this line on the Y axis. Used for predicting  $\mathbf{C}_{_{t}}$ 

**bsa** = body surface area. Corresponds more closely to metabolic rate than weight, especially important with drugs with a low therapeutic ratio. Used for extrapolating doses from big animals to small ones and vice versa.

 $C = C_p$  = plasma concentration of drug. Units usually  $\mu$ g/mL (M rarely used).

Css = Cpss = plasma concentration at a steady state, ie, the amount of drug going in is the same as the amount of drug going out.

**CL** = clearance = the volume of blood cleared of drug per unit time. Units usualy mL/min/kg

 $CL_{\text{systemic}} = CL_{\text{total}} = \text{the sum of } CL_{\text{hepatic}}, CL_{\text{renal}}, \text{ etc}$   $C_{\text{max}} = \text{maximum plasma concentration reached after a}$ 

dose of drug.  $\mathbf{D} = \mathbf{O} = \text{dose}$  or quantity, ie, amount of drug given.

 $\mathbf{F}$  = bioavailability (fraction of dose reaching the systemic circulation).

 $\mathbf{k_a}$  = absorption rate constant

**kel** = elimination rate constant - slope of the plasma concentration / time curve in a single compartment model. Used in deriving the half life and other parameters.

Ln = natural logarithm

 $\lambda z$  = slope of the terminal elimination phase in a multicompartment model (corresponding to  $k_{el}$  in a single compartment model)

**MRT** = mean residence time =  $AUMC_{0-inf}$  /  $AUC_{0-inf}$  Gives some indication of how long a drug persists in the body. nb - covers absorption as well as distribution and elimination.

 $\mathbf{t}_{1/2}$  = half life = the time it takes for drug concentration to fall by half.

 $\mathbf{t}_{1/2\alpha}$  = half life of the distribution phase

 $\mathbf{t}_{1/2\beta}$  = half life of the elimination phase

 $\overrightarrow{Vd}$  = volume of distribution = the volume the drug would occupy if it was evenly distributed at the concentration found in the plasma. Gives some idea of where the drug goes.

 $Vd_c$  = volume of distribution of the central compartment

 $Vd_{ss}$  = volume of distribution at a steady state

 $Vd_{\lambda z} = Vd_{\beta} = Vd_{area} = volume$  of distribution during the terminal elimination phase.