



Graphical Flashcards

Pharmacology

For the Primary FRCA

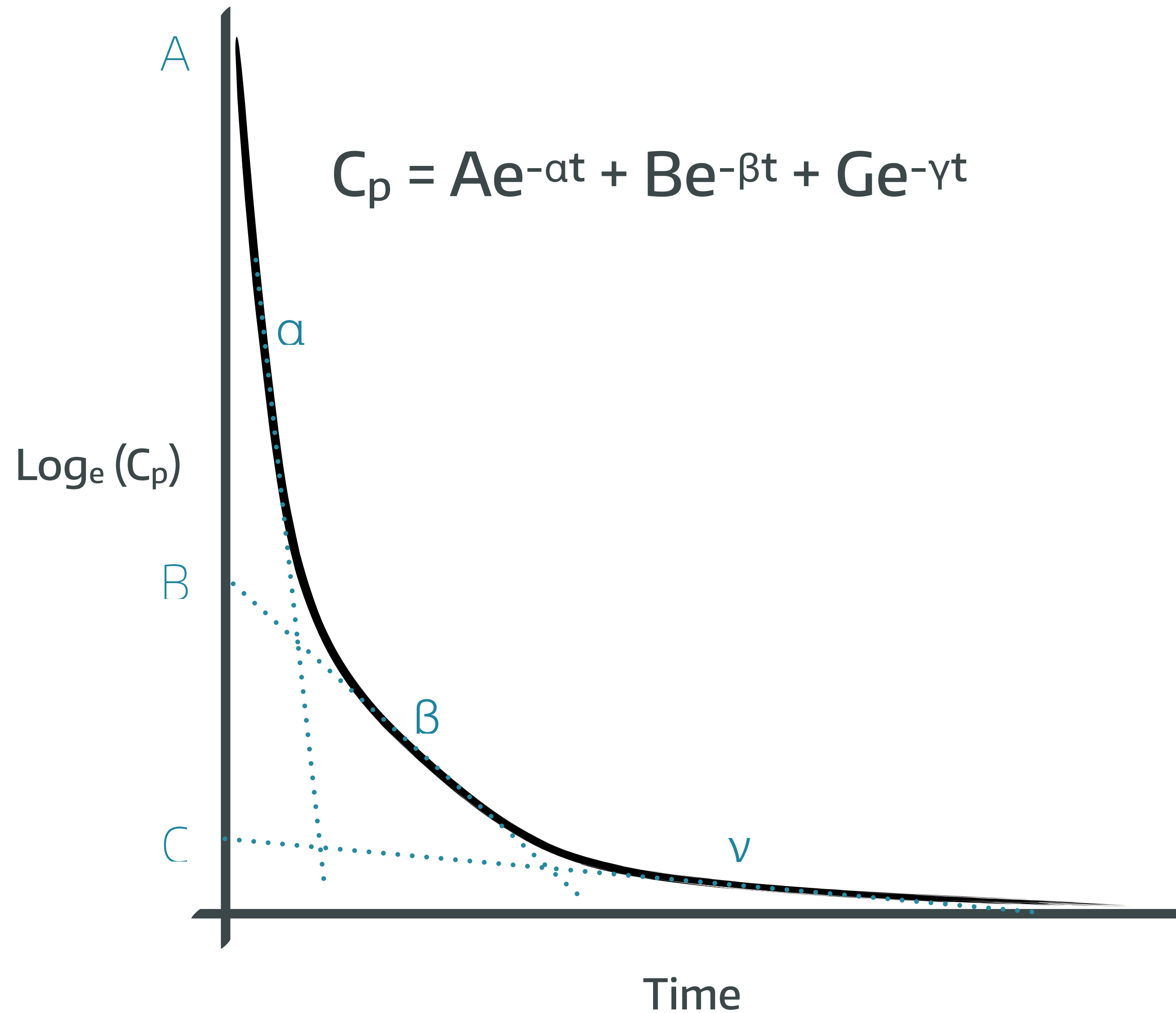


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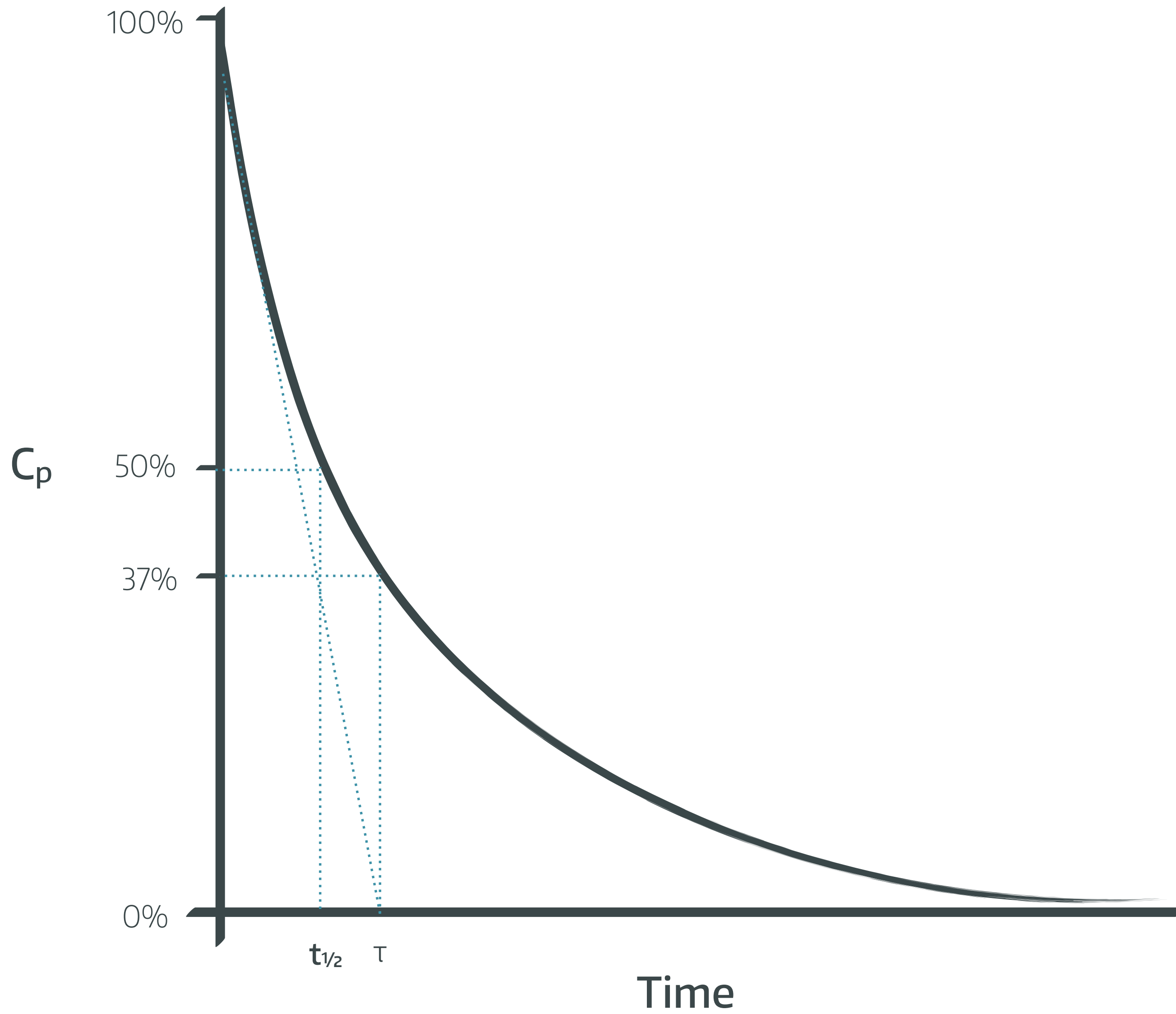
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Tri-exponential Decline in Plasma Concentration of Drug



PHARMACOKINETICS

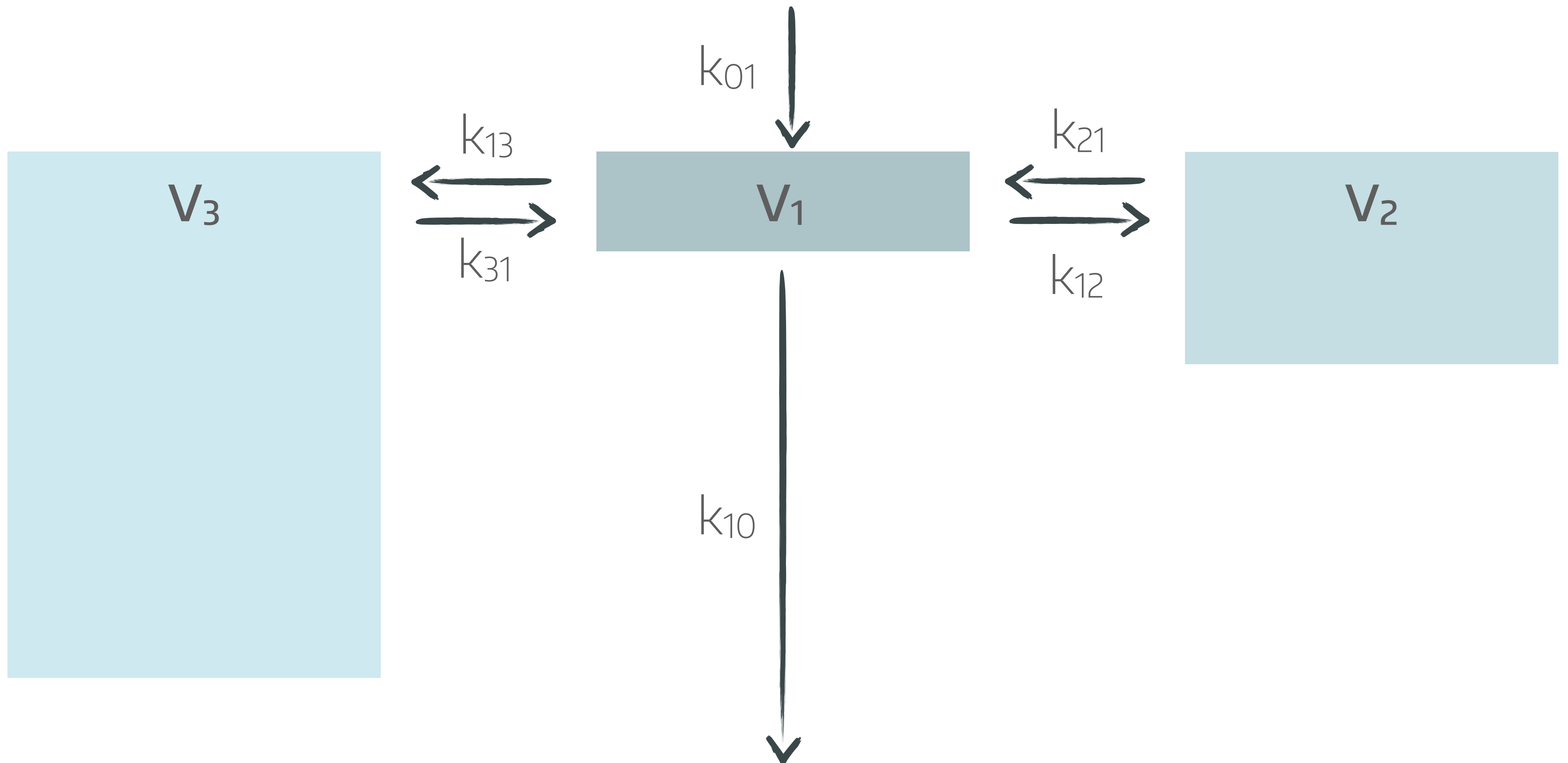


$$t_{1/2} = 0.693 \times \tau$$

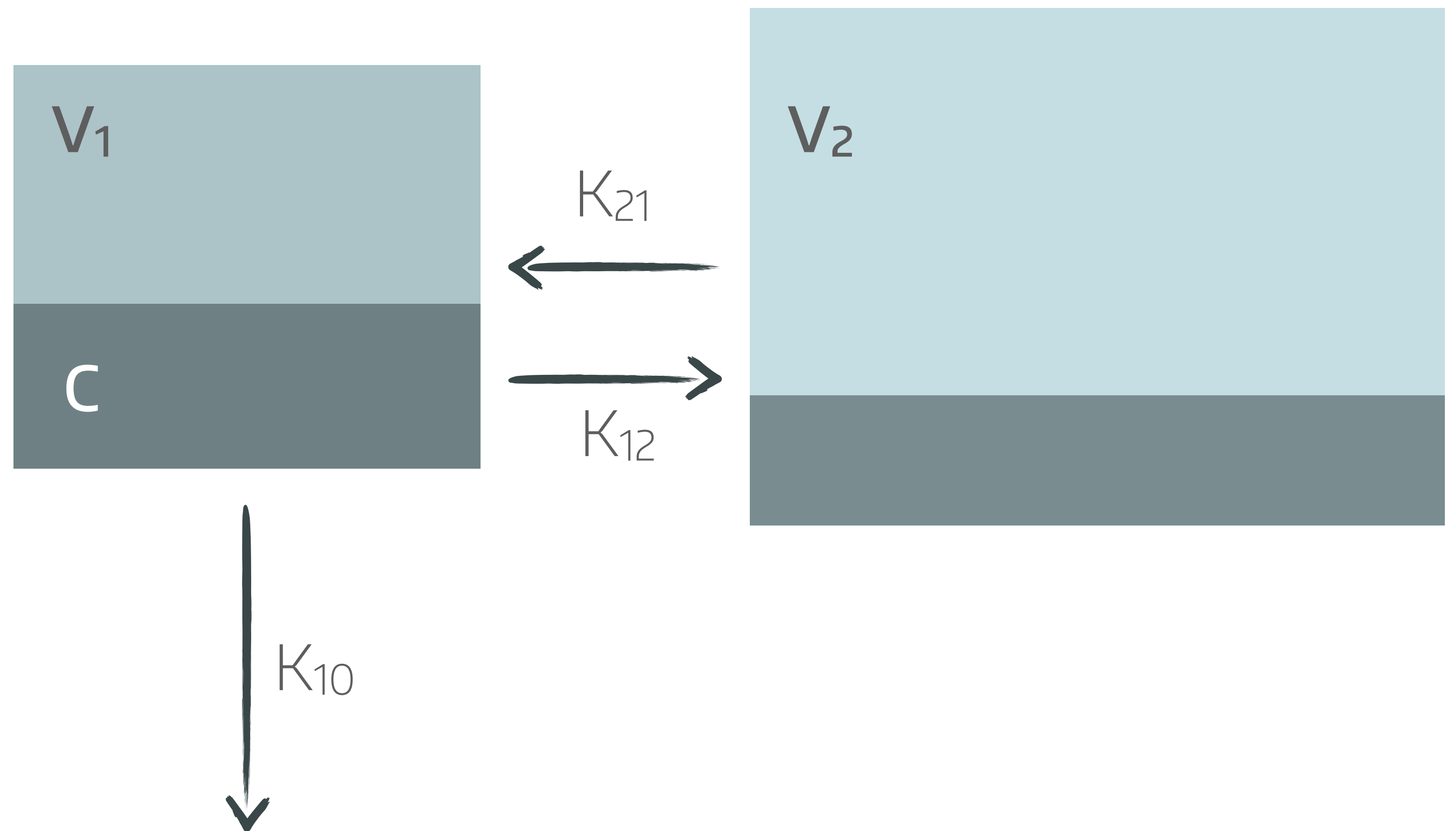
$$k = \tau^{-1}$$

$$C_p = C_0 \cdot e^{-kt}$$

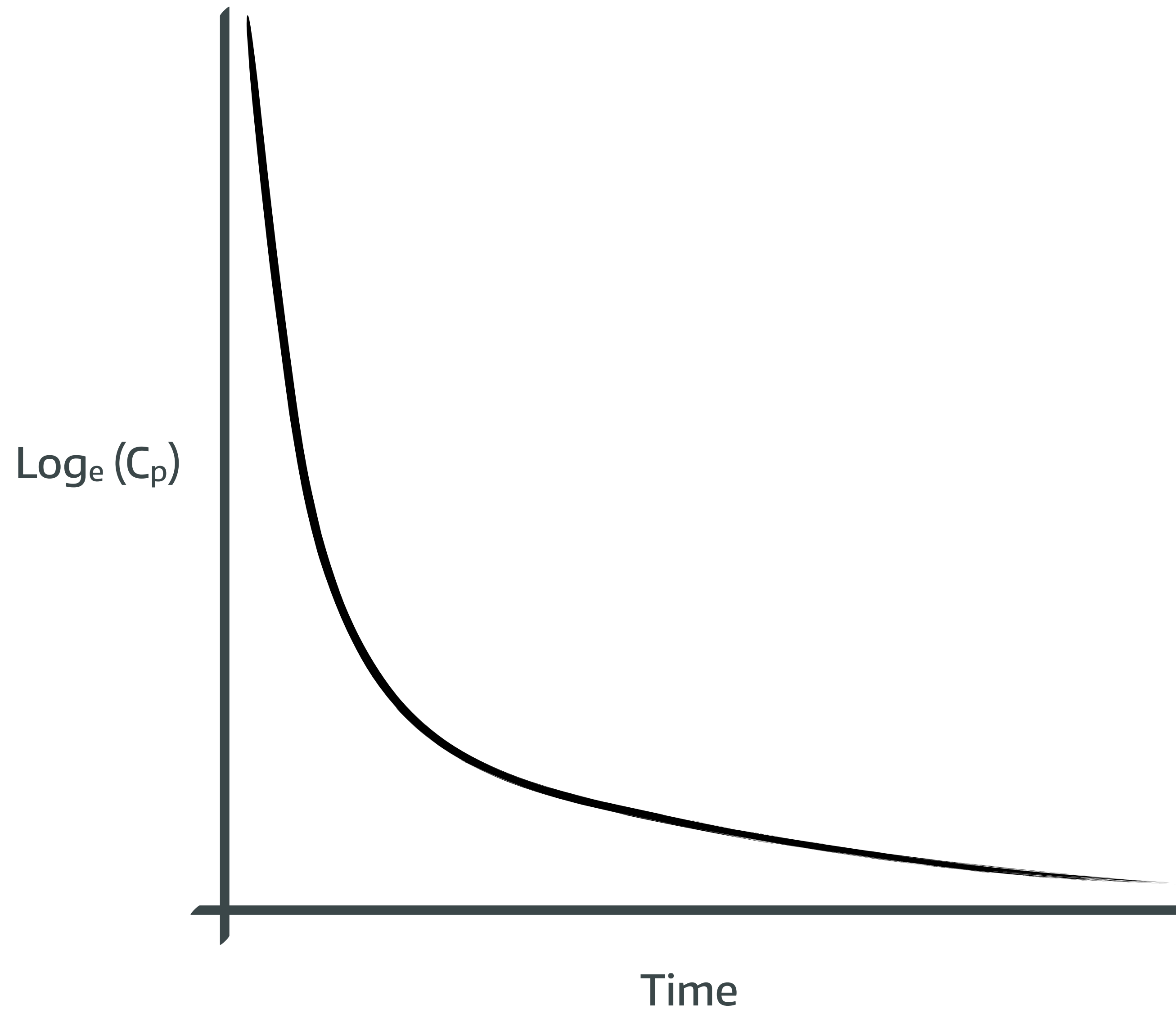
The 3-Compartment Model



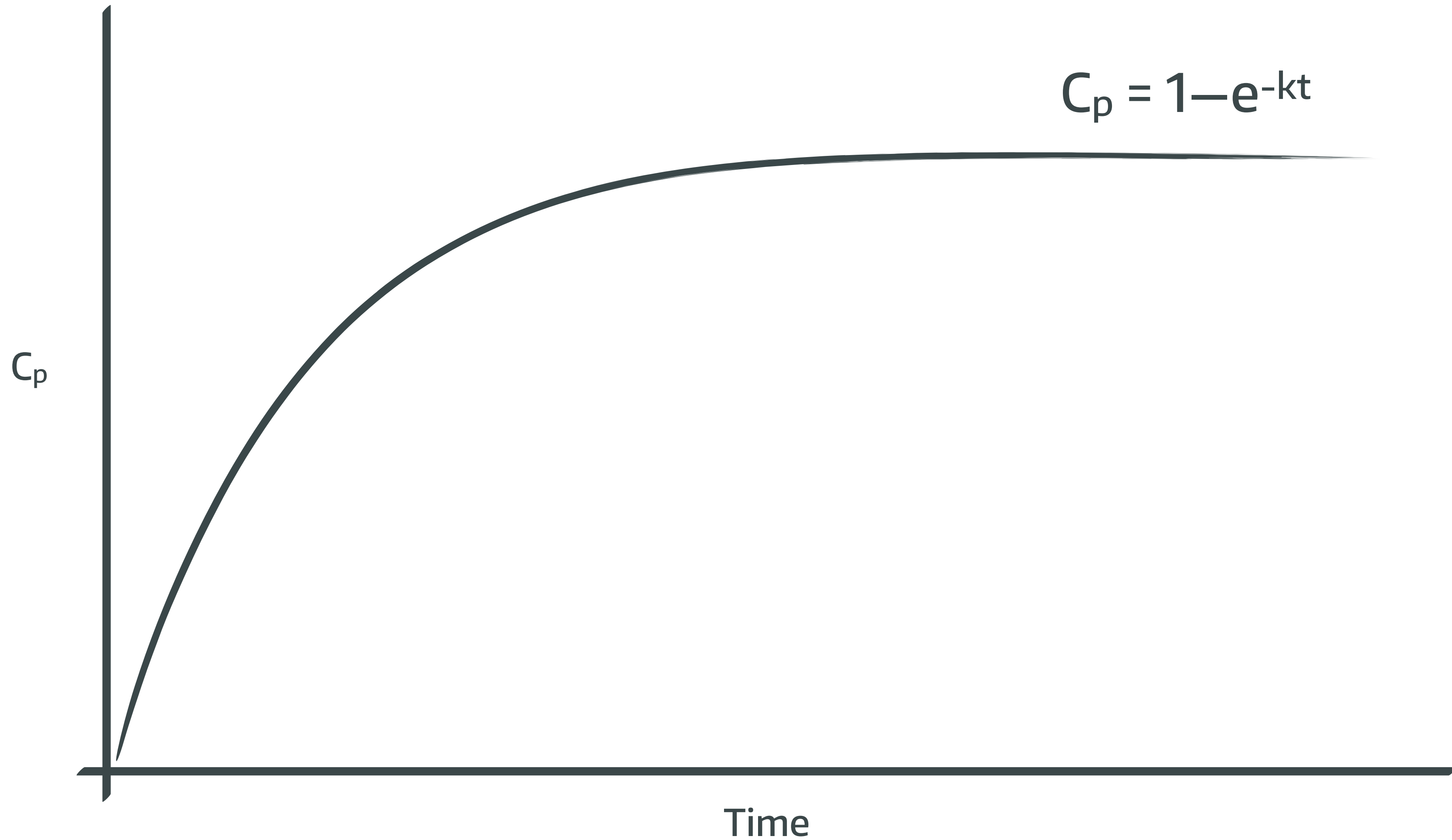
The 2-Compartment Model



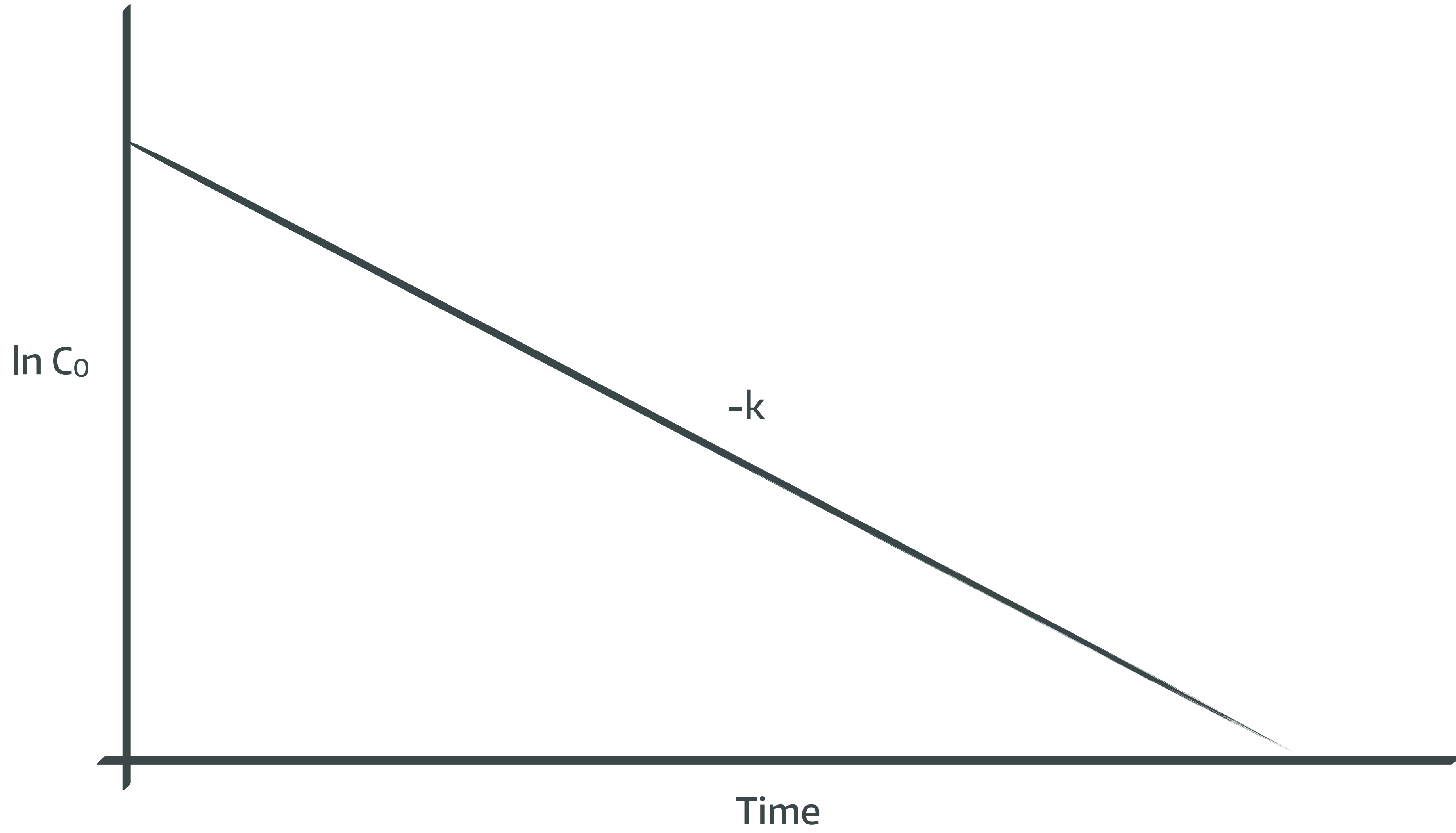
Bi-exponential Decline in Plasma Concentration of Drug



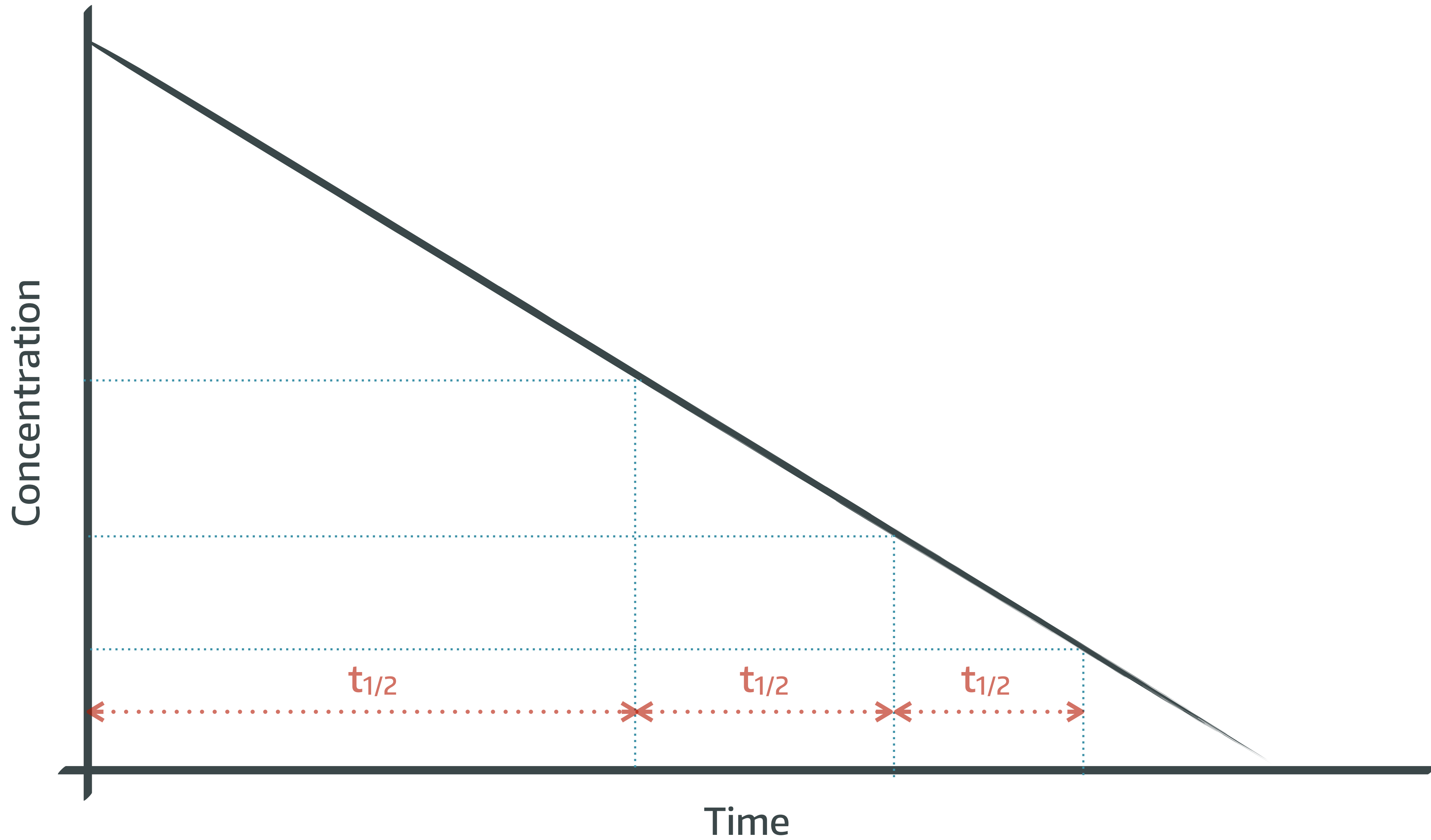
Wash-in Curve for Infusion Building to Steady State



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Zero Order Kinetics

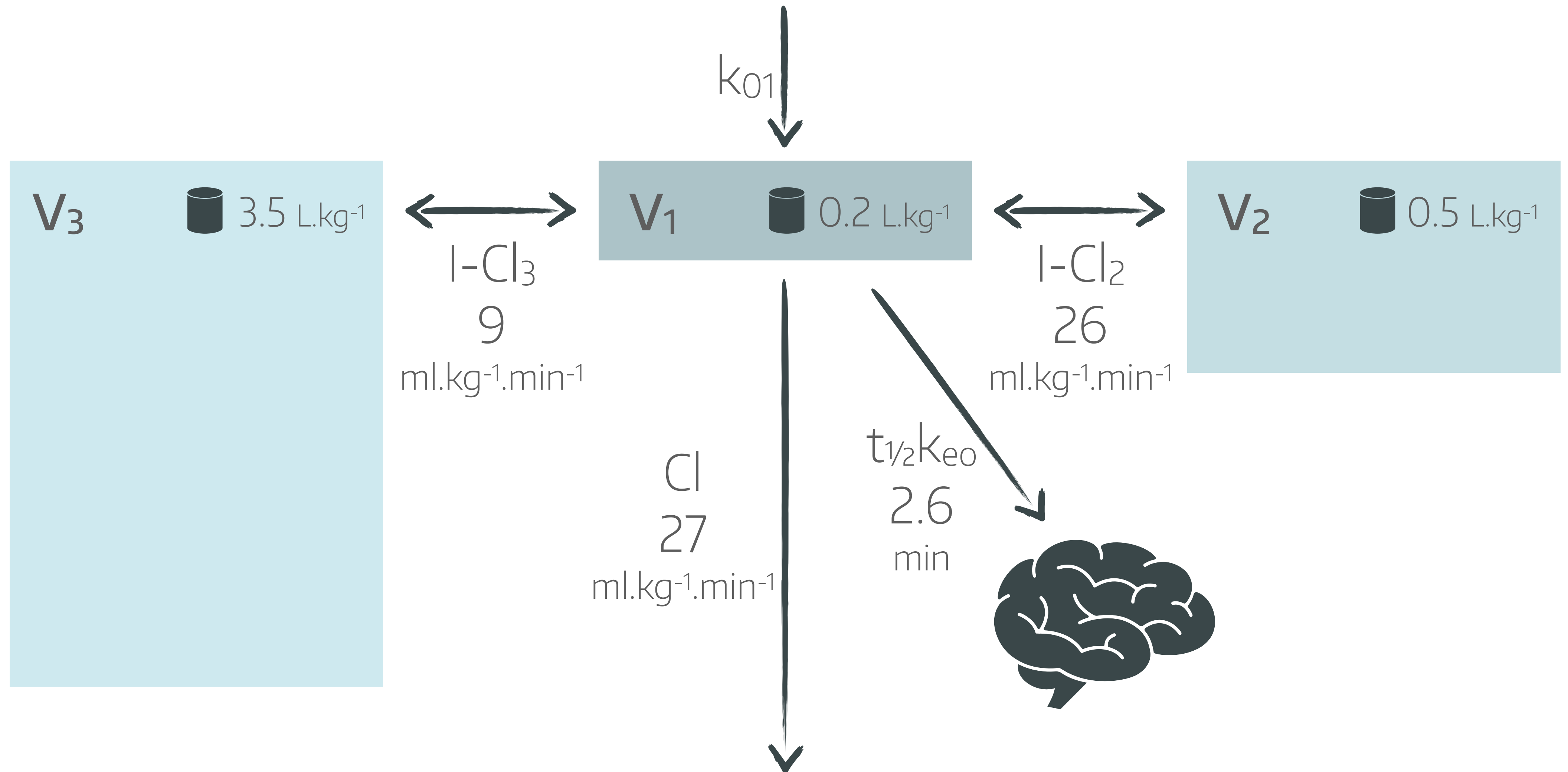


The 1-Compartment Model

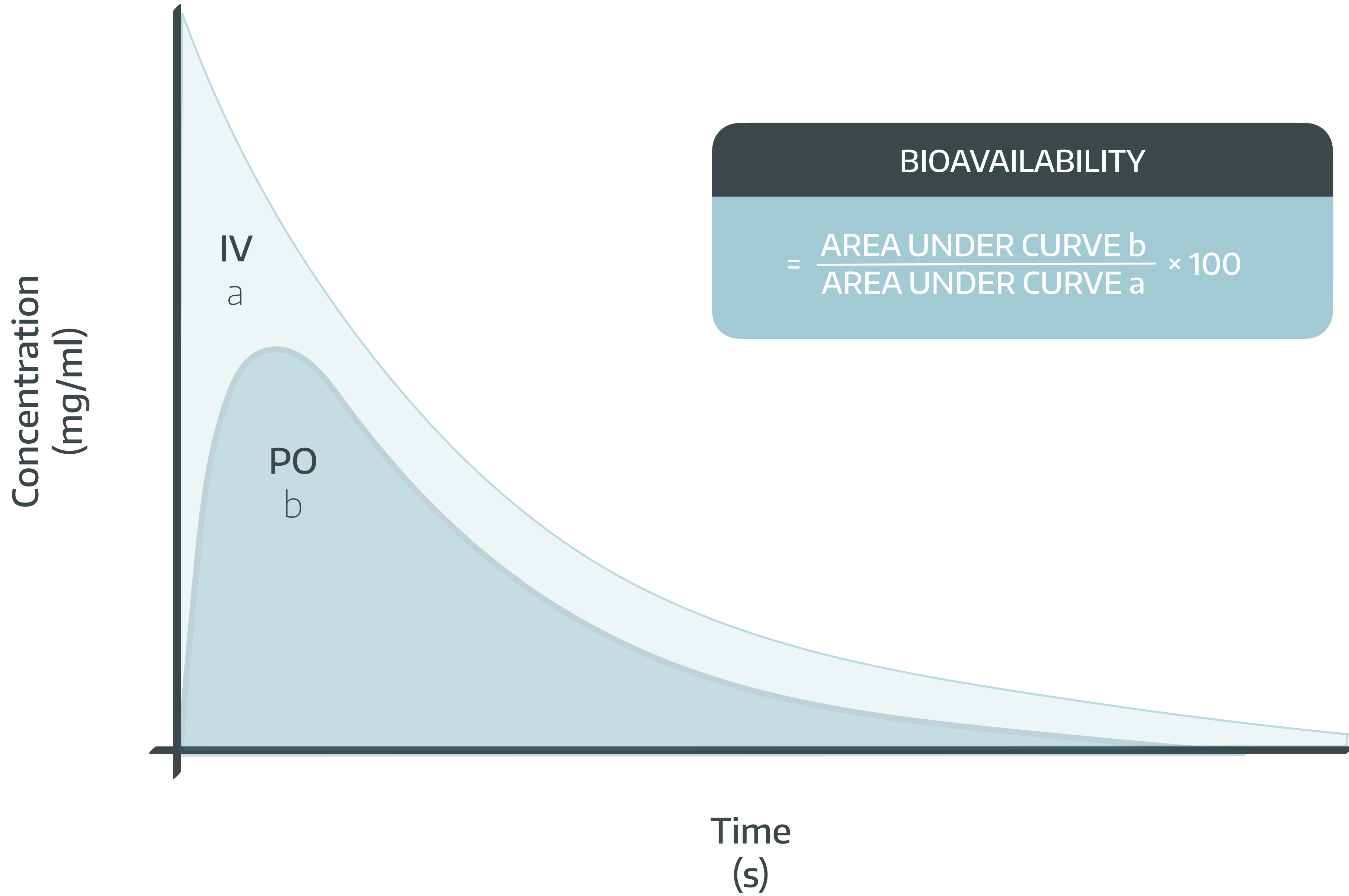


$$V_d = \frac{\text{Dose}}{\text{Concentration}}$$

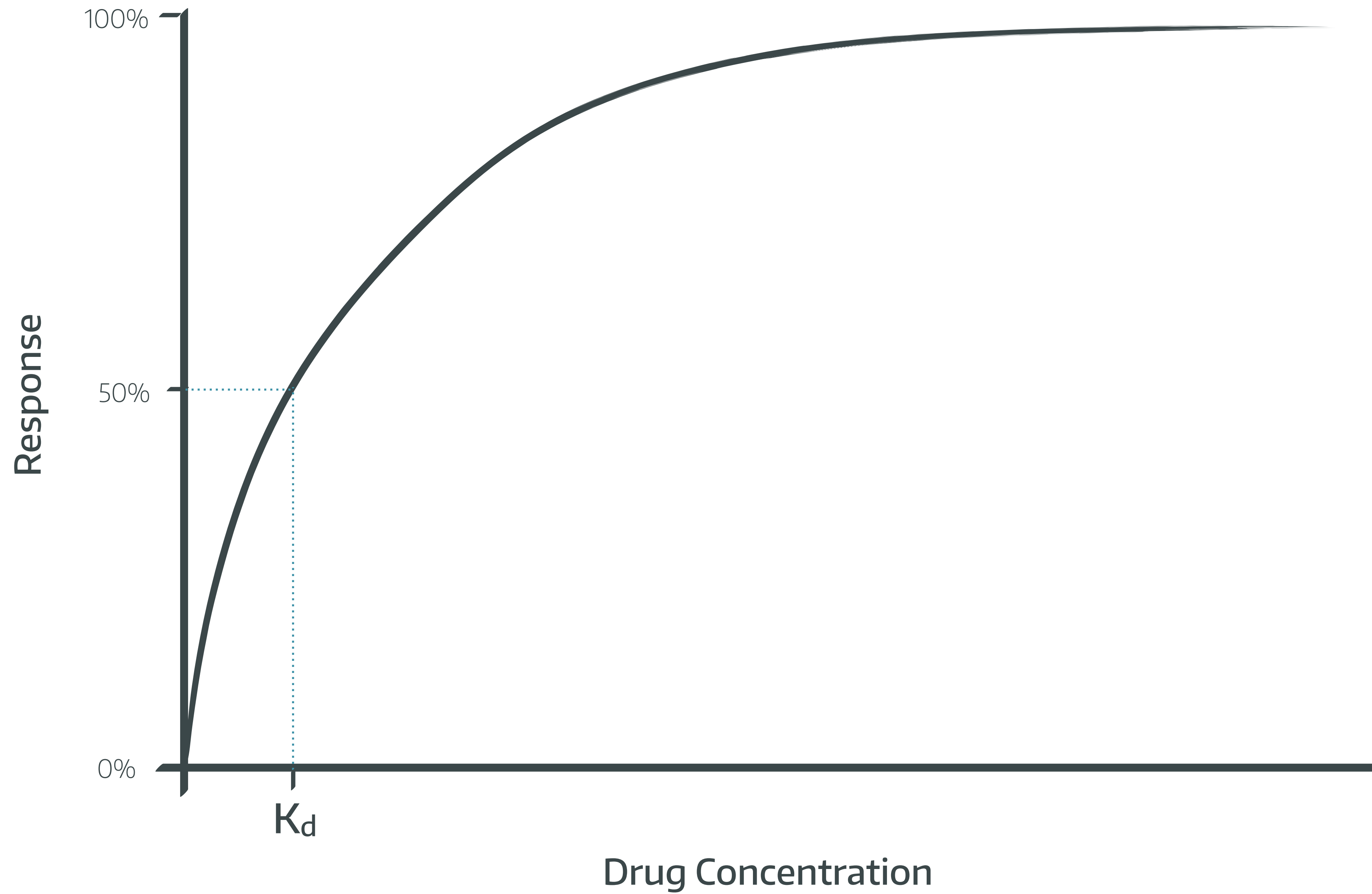
PROPOFOL: The 3-Compartment Model + Effect Site



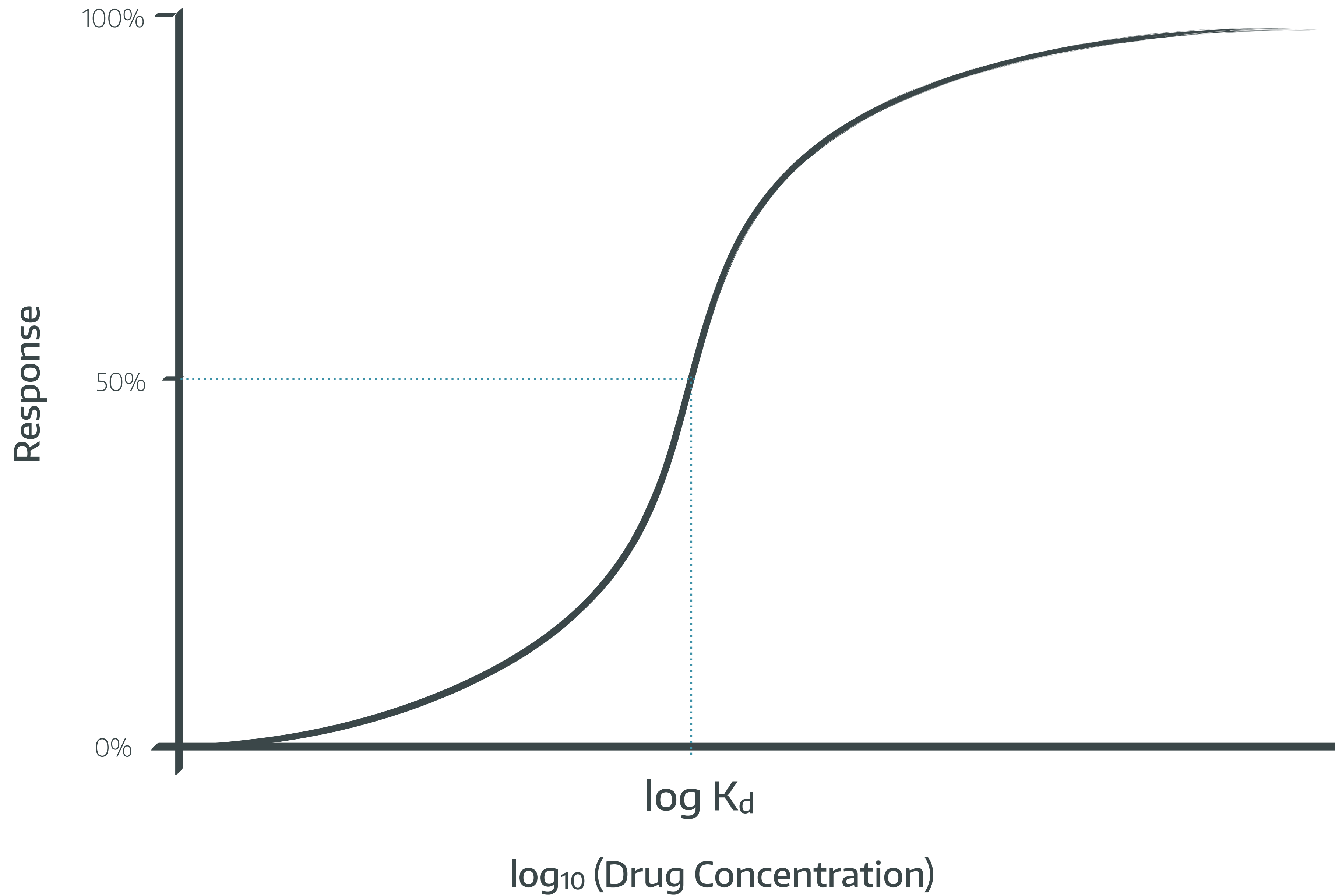
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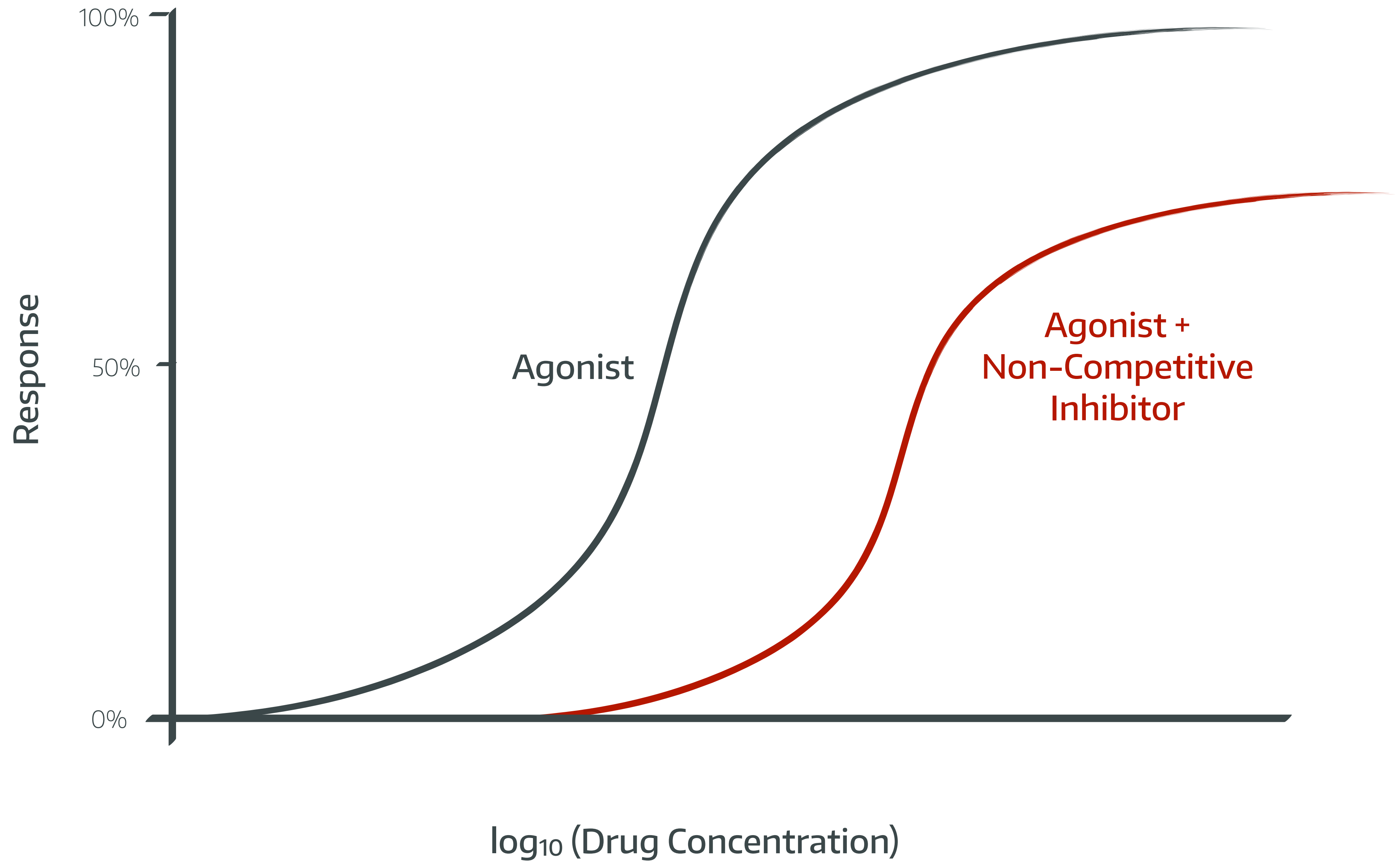
PHARMACODYNAMICS



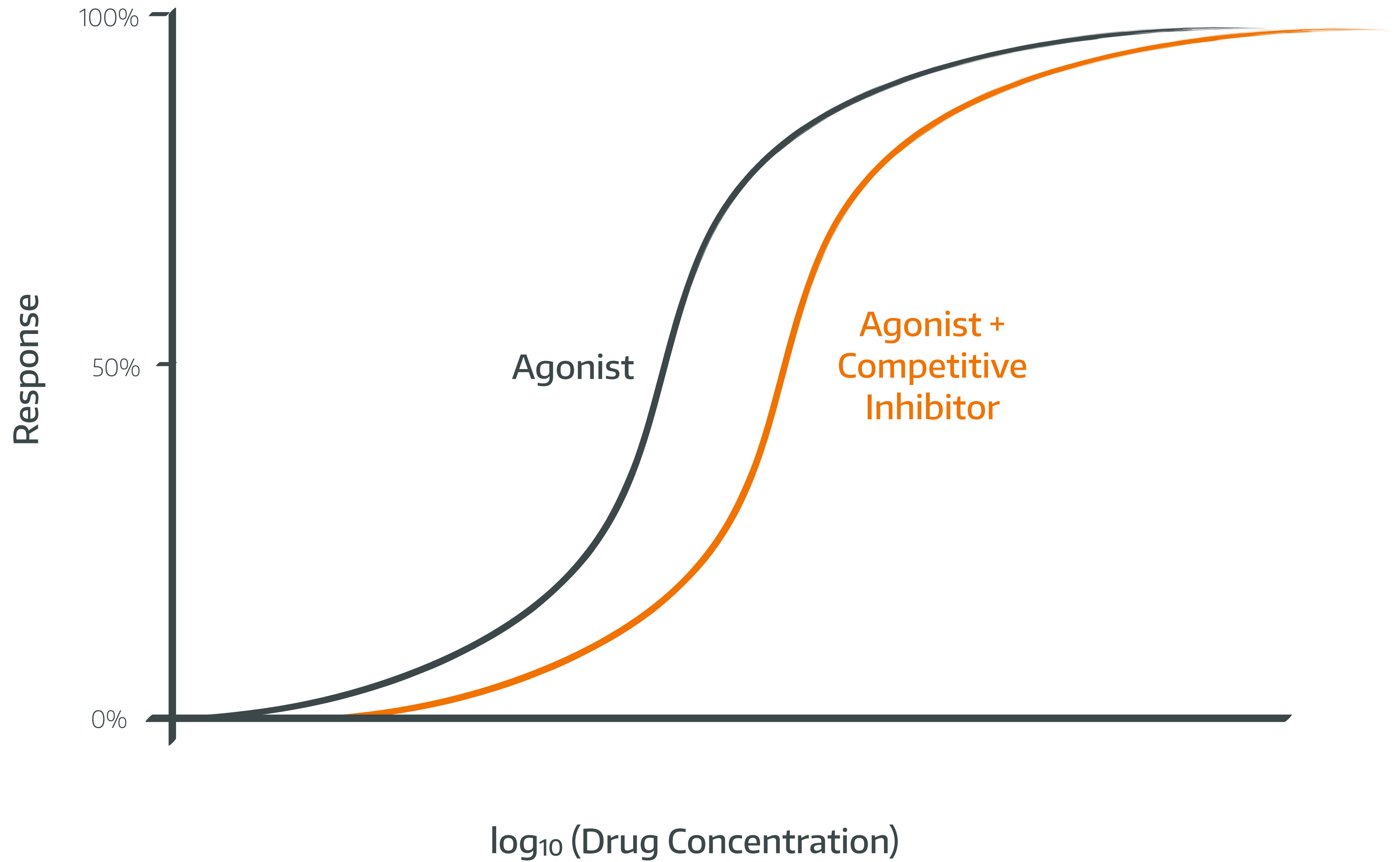
PHARMACODYNAMICS



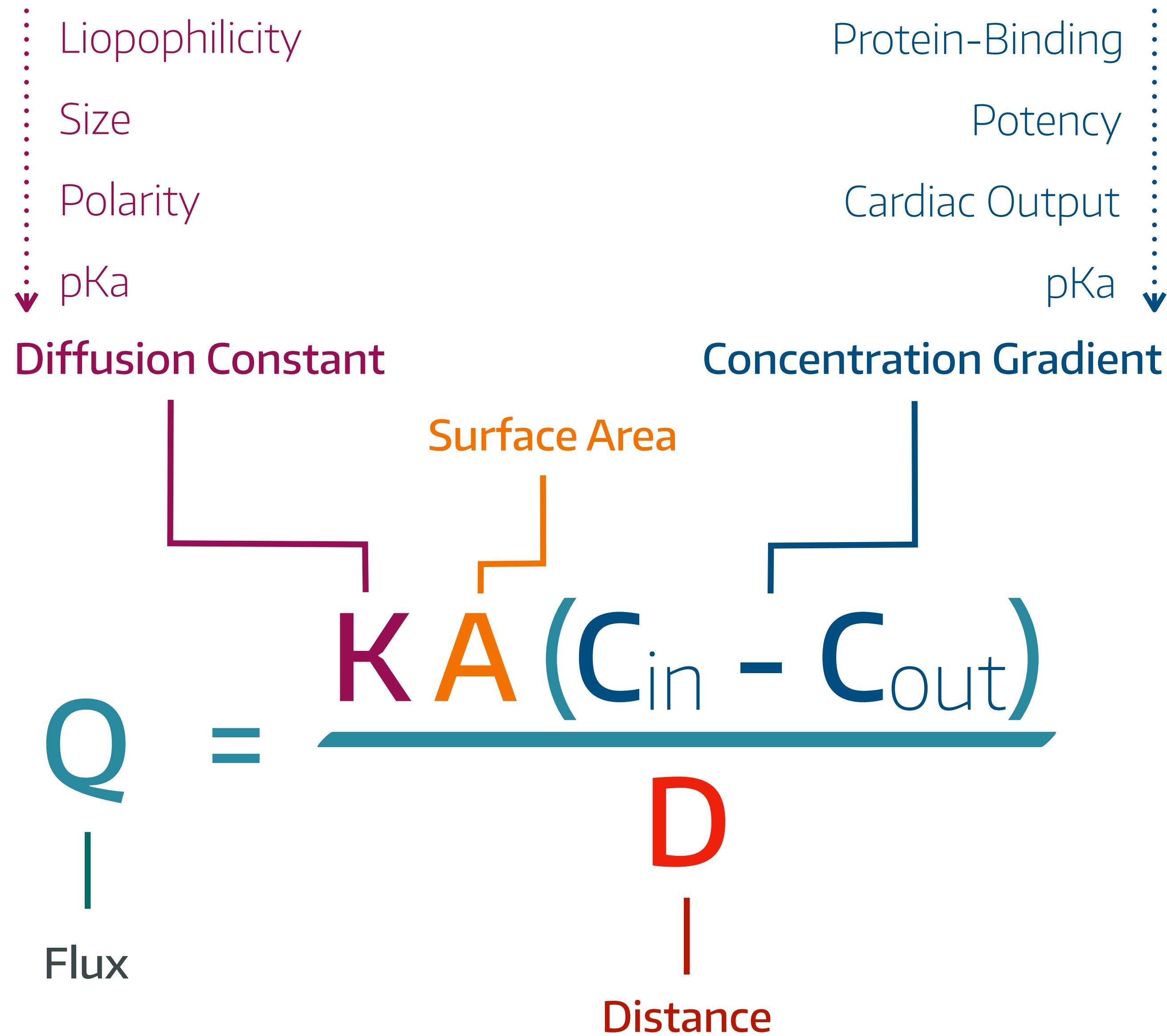
PHARMACODYNAMICS



PHARMACODYNAMICS



DRUGS ACROSS THE PLACENTA



Local Anaesthetics

Foetal Toxicity only occurs if mother toxic
Bupivacaine can ion-trap

Anaesthetic Agents

Thiopentone crosses (no harm).
Inhaled agents cross but excreted rapidly.

Muscle Relaxants

Suxamethonium crosses in very small quantities.
NDMRs do not cross.

Opioids

Beware that Pethidine (and active metabolite Norpethidine) cross and can ion-trap in foetus.

DRUG ELIMINATION

$$C_t = C_0 \cdot e^{-kt}$$

Diagram illustrating the drug elimination equation:

- C_t : Concentration at time t
- C_0 : Concentration at time 0
- k : Rate Constant
- t : Time

The diagram shows the equation $C_t = C_0 \cdot e^{-kt}$ with labels and connecting lines. Below C_t is the label 'Concentration at time t'. Below C_0 is the label 'Concentration at time 0'. Above the k in the exponent is the label 'Rate Constant'. Above the t in the exponent is the label 'Time'. A bracket connects 'Rate Constant' to k . A bracket connects 'Time' to t . There are also vertical lines connecting C_t to its label and C_0 to its label.

$$C_t = C_0 \cdot e^{-kt}$$

$t_{1/2}$ = time taken for C_0 to drop by 50%

∴ substitute 1 for C_0

$$1/2 = 1 \cdot e^{-kt_{1/2}}$$

$$\ln(1/2) = \ln(1) + -kt_{1/2}$$

$$\ln(1/2) = 0 - kt_{1/2}$$

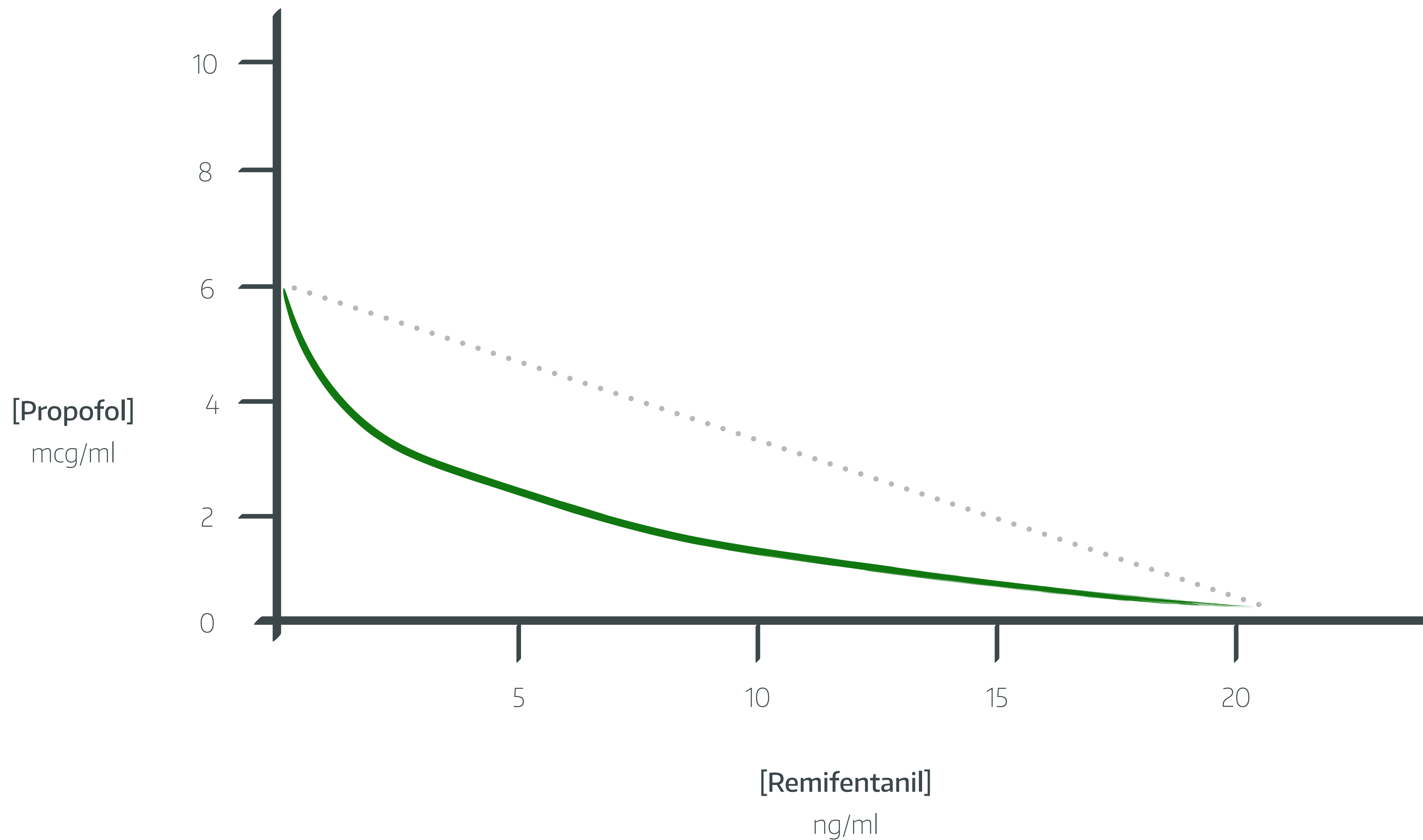
$$kt_{1/2} = \ln(2)$$

$$t_{1/2} = \frac{\ln(2)}{k}$$

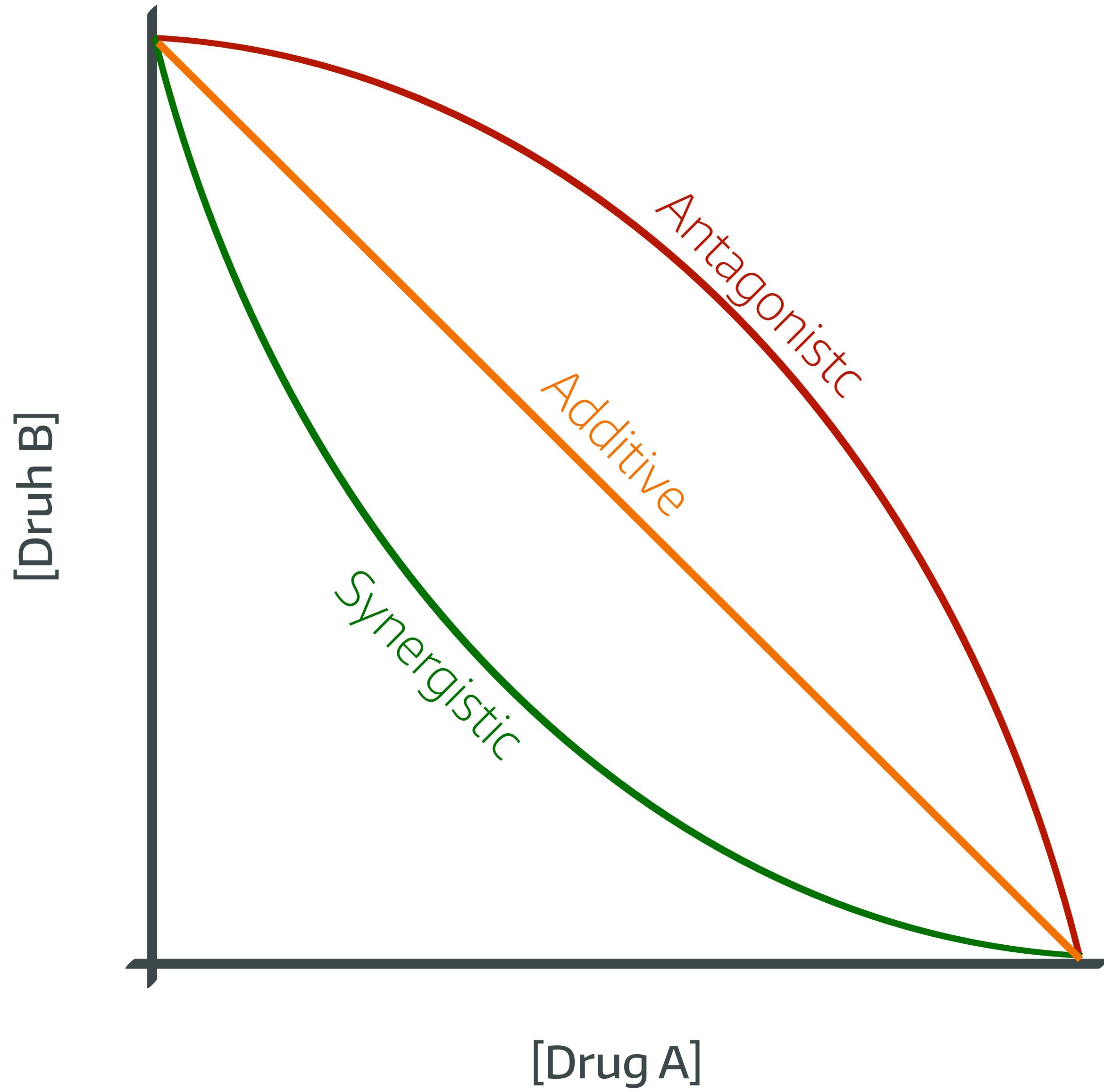
DRUG ELIMINATION

$$\begin{array}{ccccc} \text{Clearance} & & \text{Volume of} & & \text{Elimination Rate} \\ & & \text{Distribution} & & \text{Constant} \\ & & \text{└──┐} & & \text{└──┐} \\ & & V_D & \cdot & k \\ \text{ml.min}^{-1} & = & \text{ml} & \cdot & \text{min}^{-1} \end{array}$$

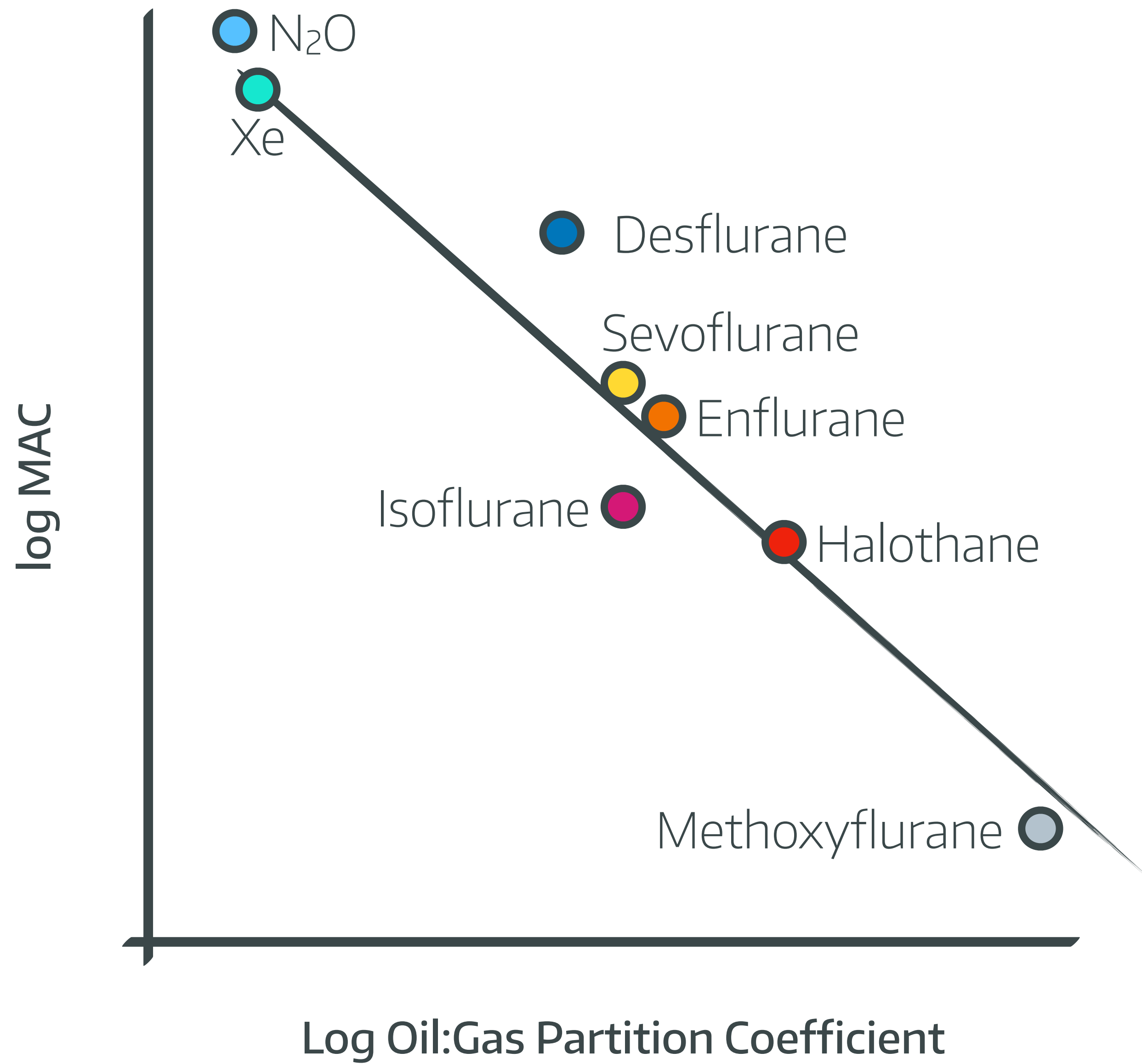
PROPOFOL VS REMIFENTANIL ISOBOLOGRAM



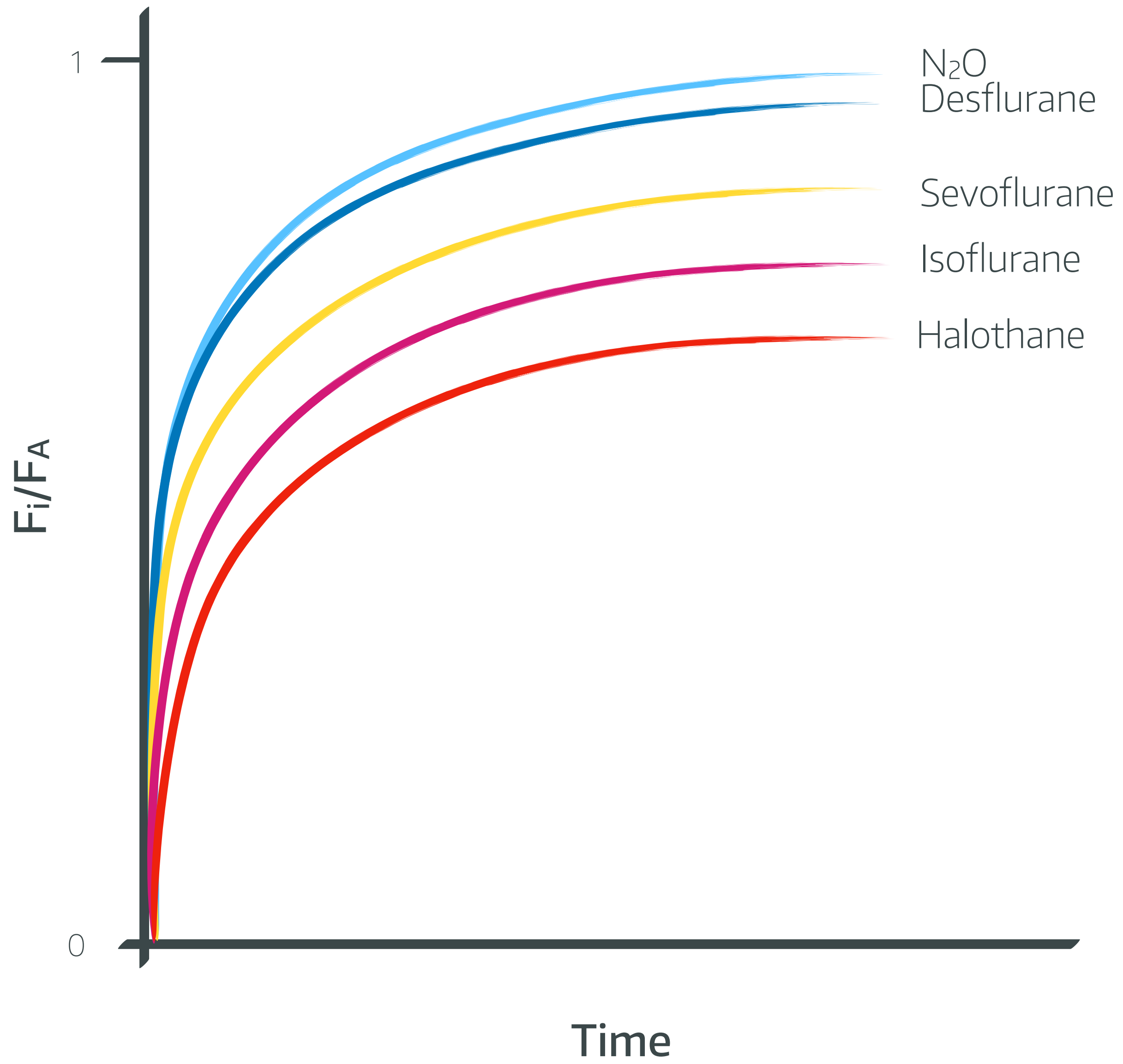
ISOBOLOGRAM



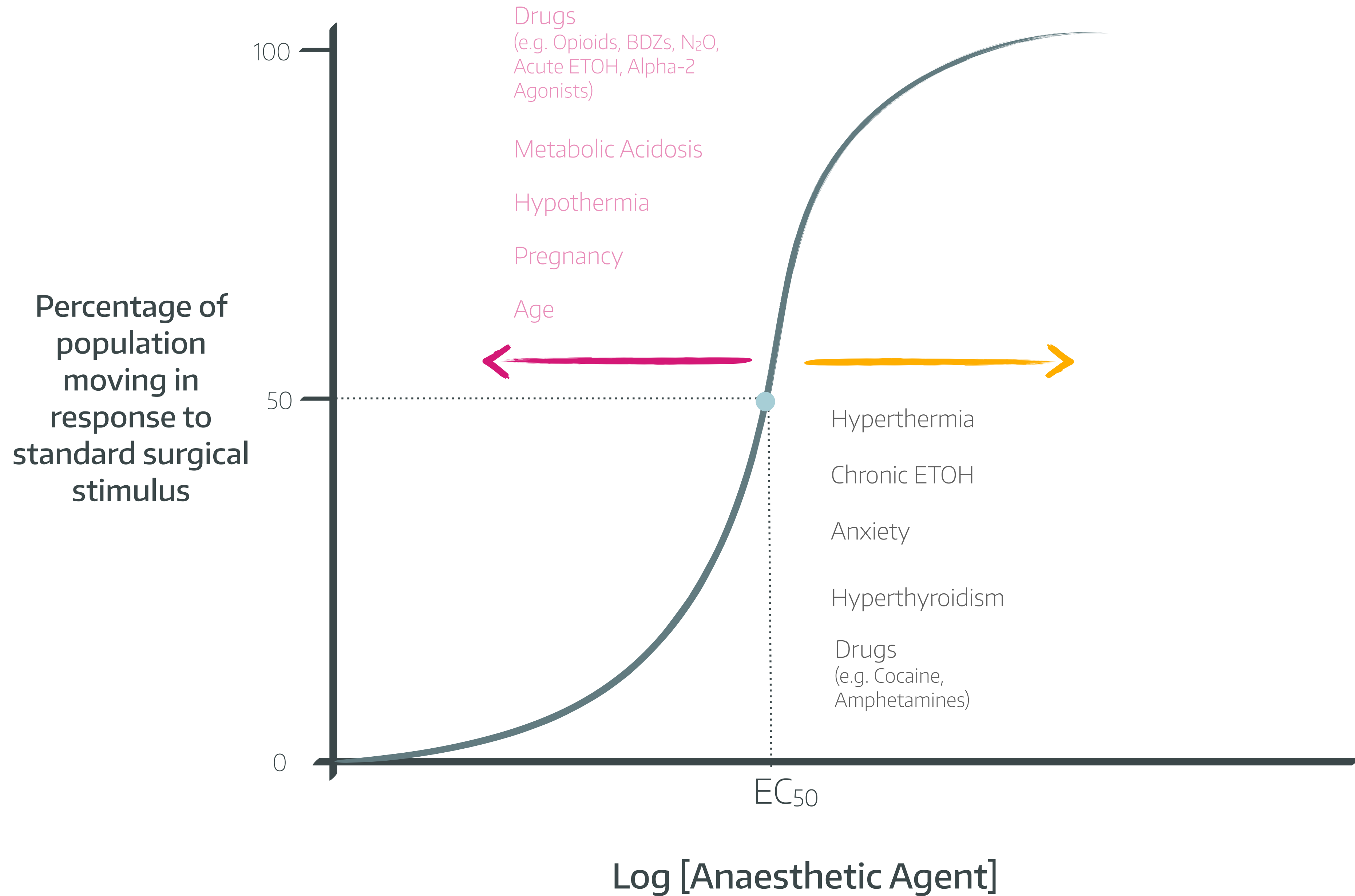
MEYER-OVERTON



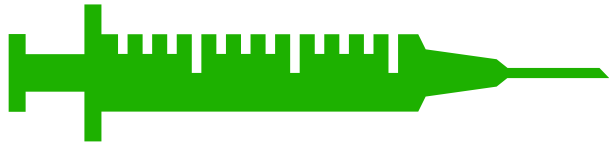
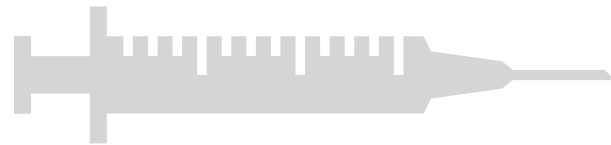
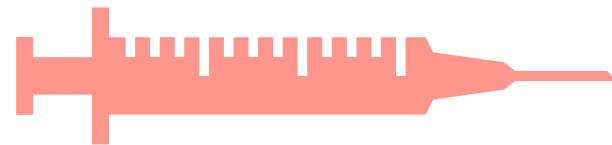
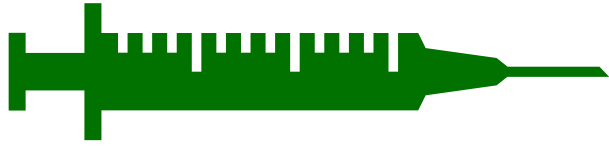
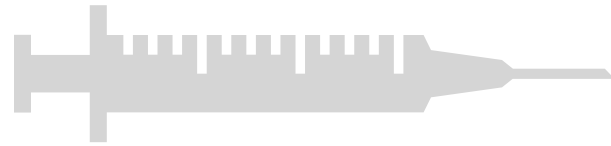
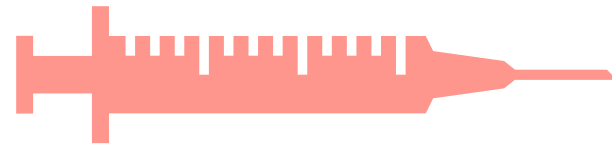
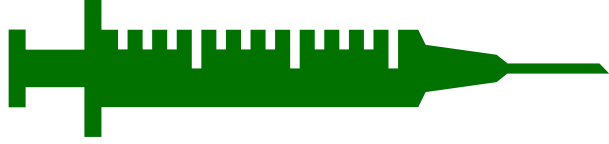
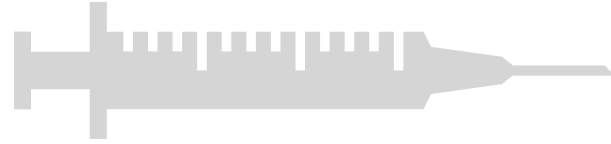
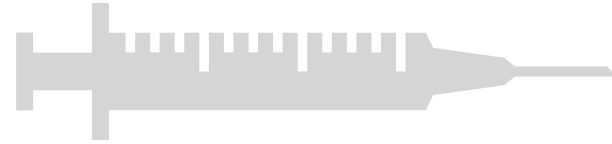
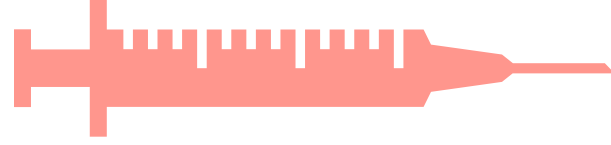
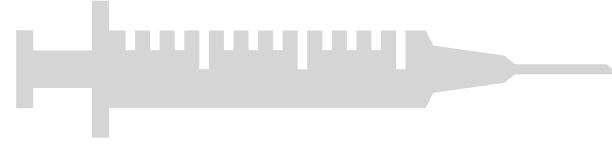
PHARMACOKINETICS



PHARMACOKINETICS



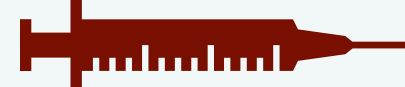
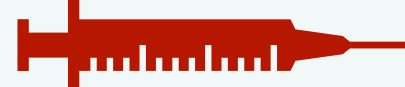
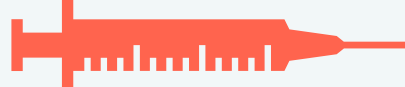
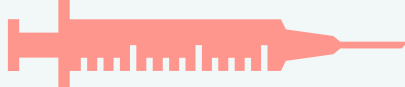
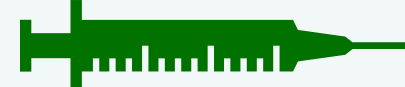
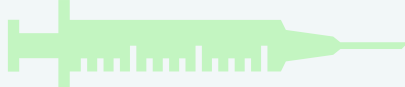
HYPONOTIC AGENTS

	GABA _A Receptors	Glycine Receptors	NMDA Receptors	nACh Receptors
Propofol				
Thiopentone				
Etomidate				
Ketamine				

Key

Strength of Agonistic effect

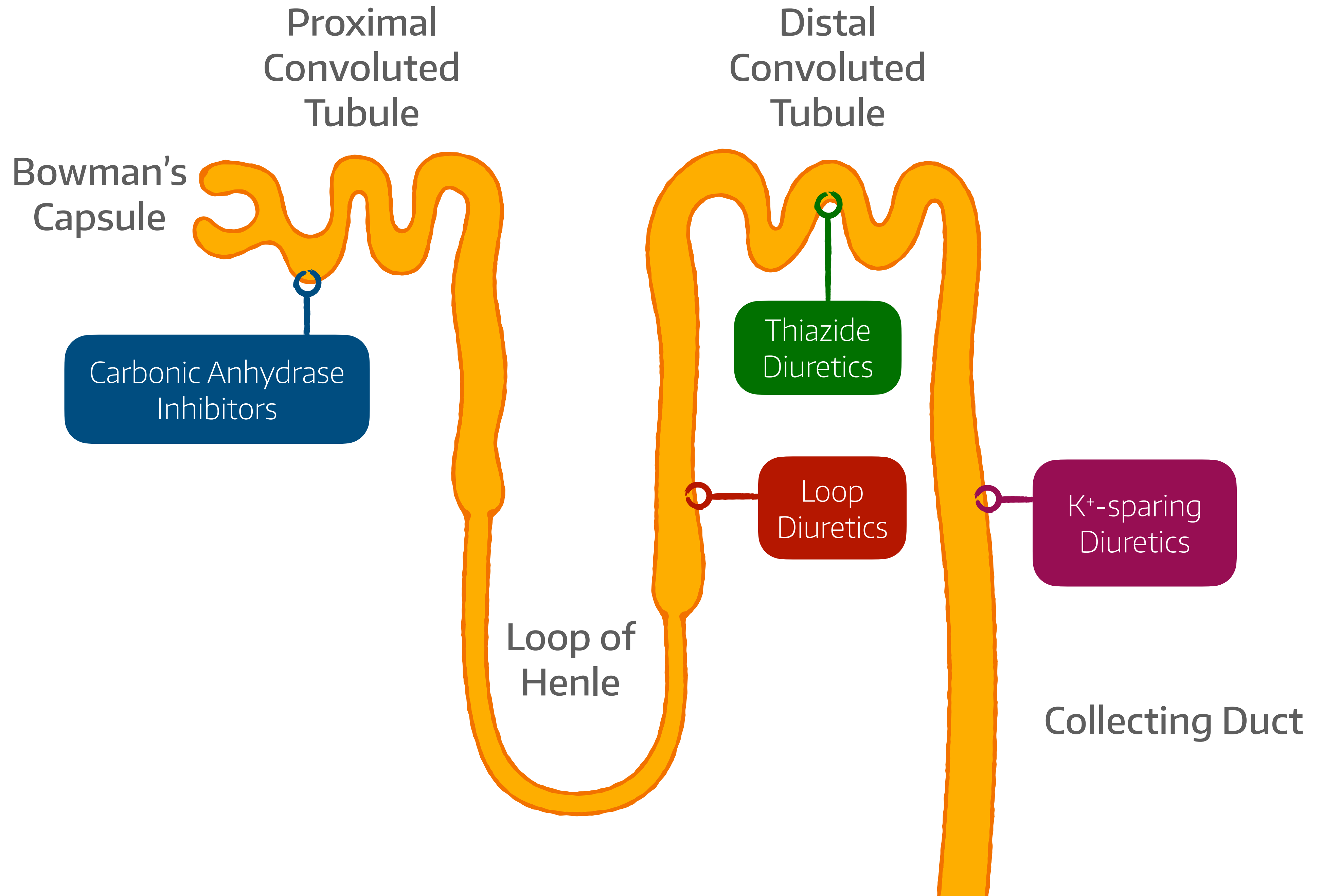
Strength of Antagonistic effect



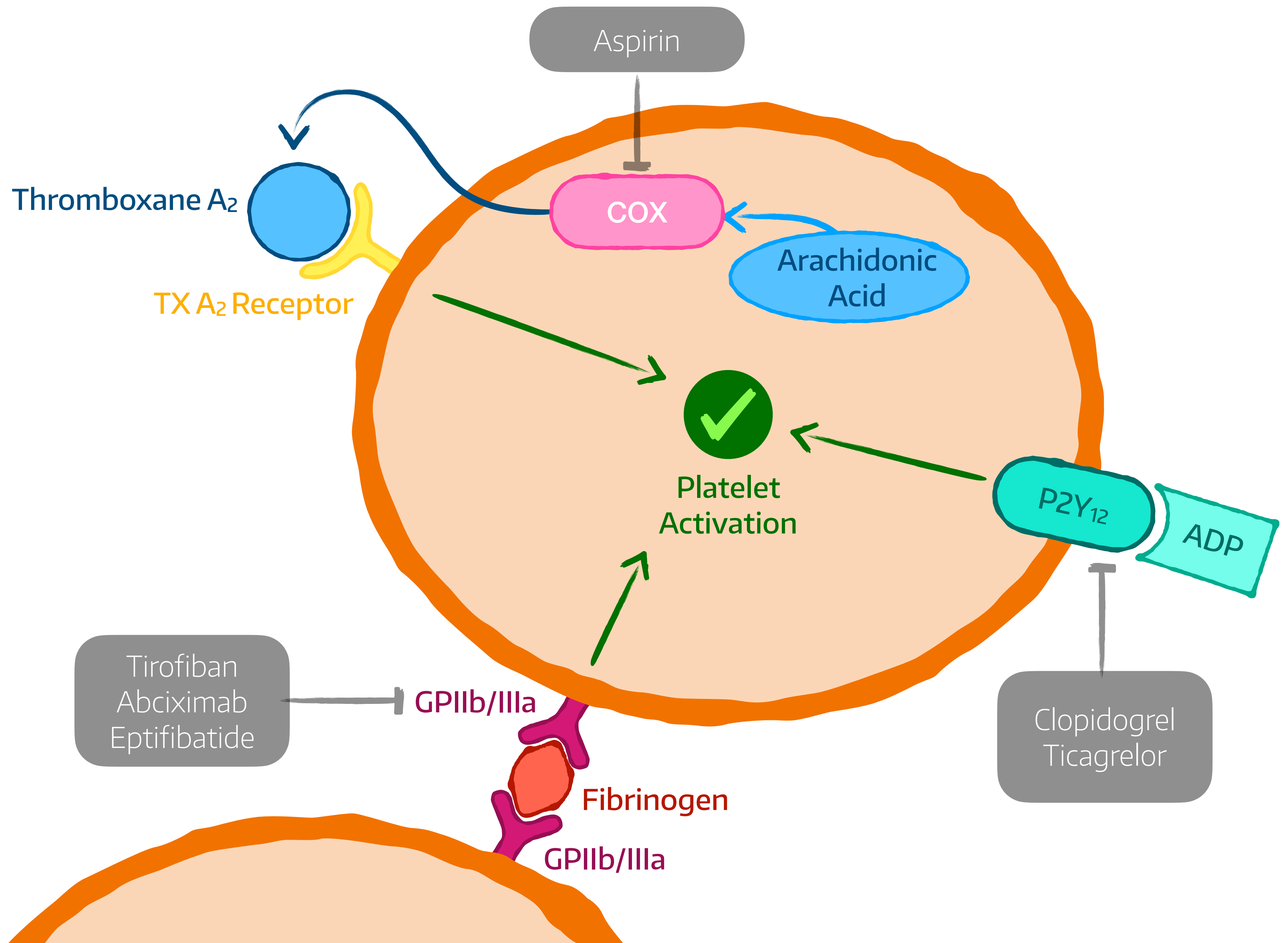
BENZODIAZEPINES

	Equipotent Dose (mg)	Onset (Mins)	Half-life (Hours)	Duration of Effect (Mins)
Chlordiazepoxide	25	60 PO		
Diazepam	10	1 IV 15-30 PO		
Lorazepam	1	2 IV 30-60 PO		
Midazolam	2.5	2-5 IV		

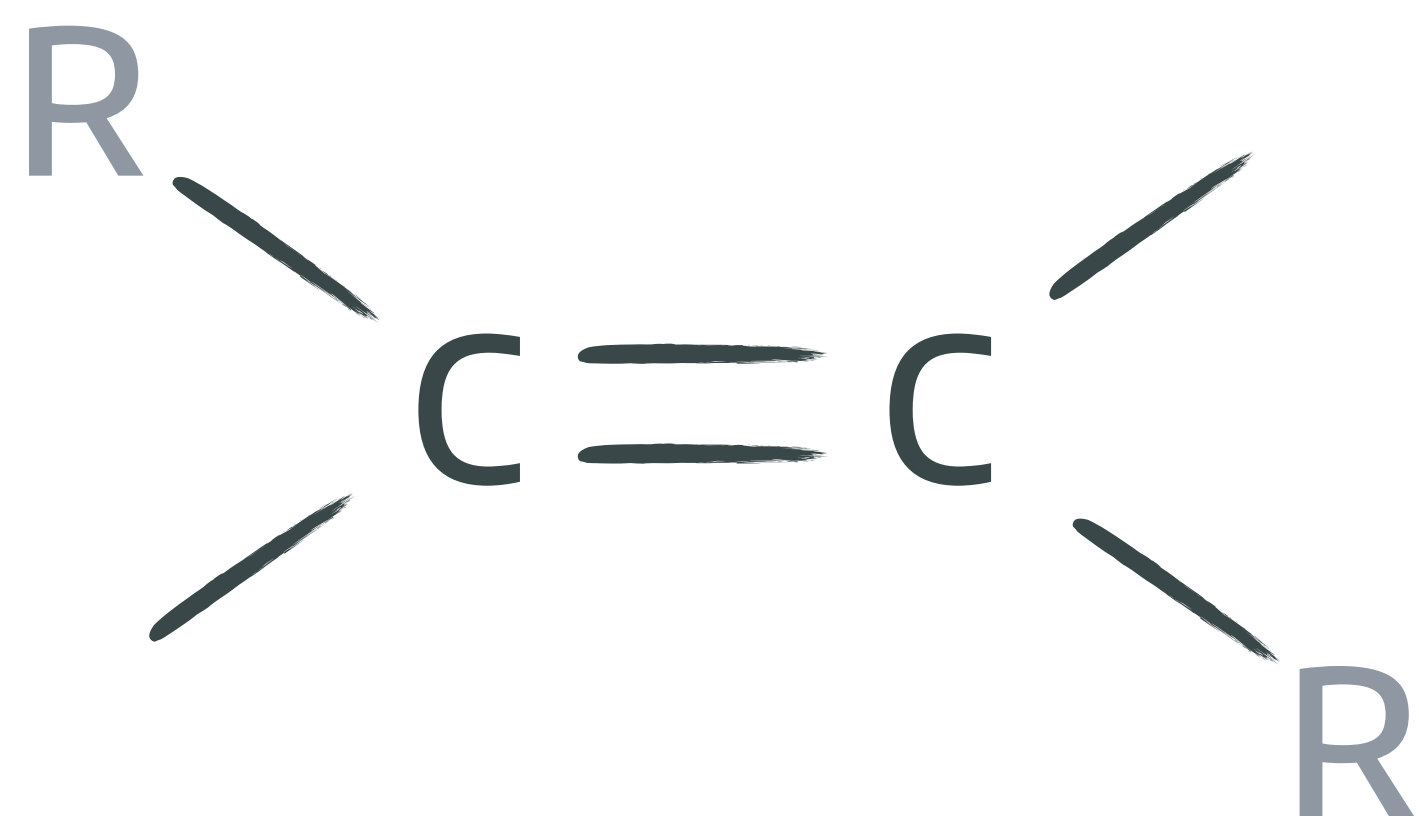
DIURETICS



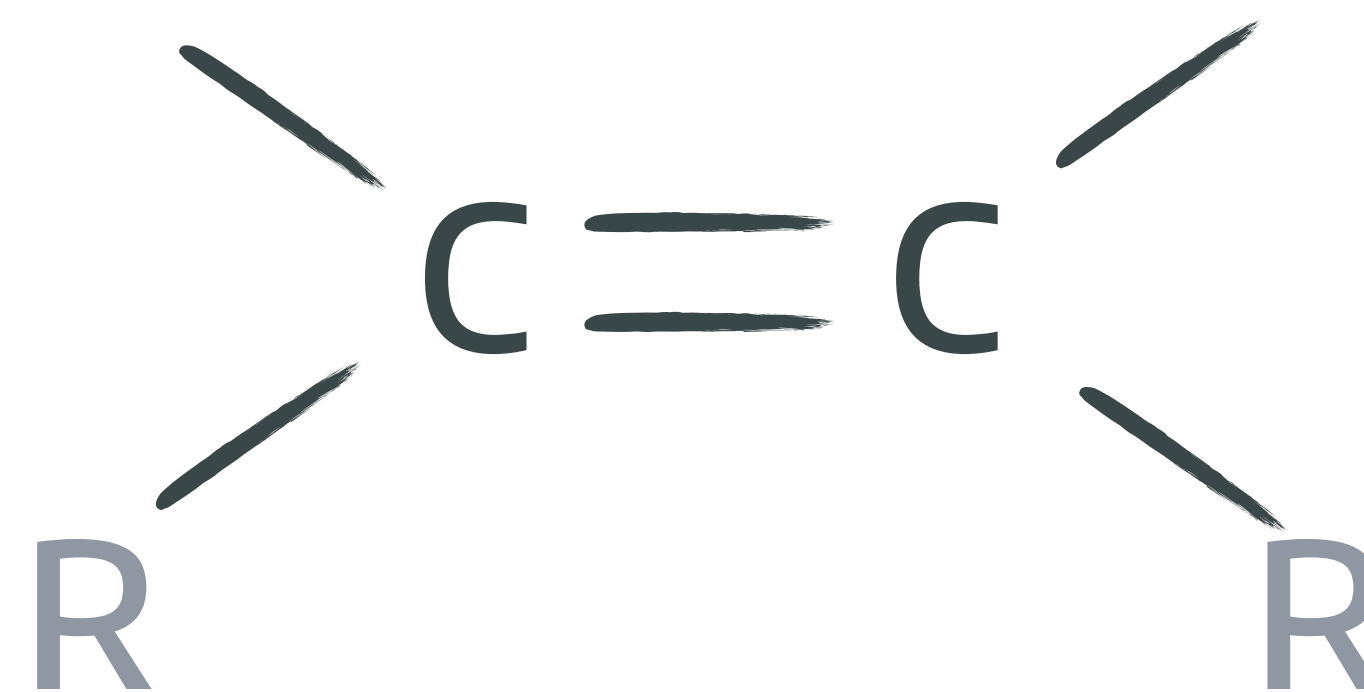
ANTIPLATELET AGENTS



GEOMETRIC ISOMERISM

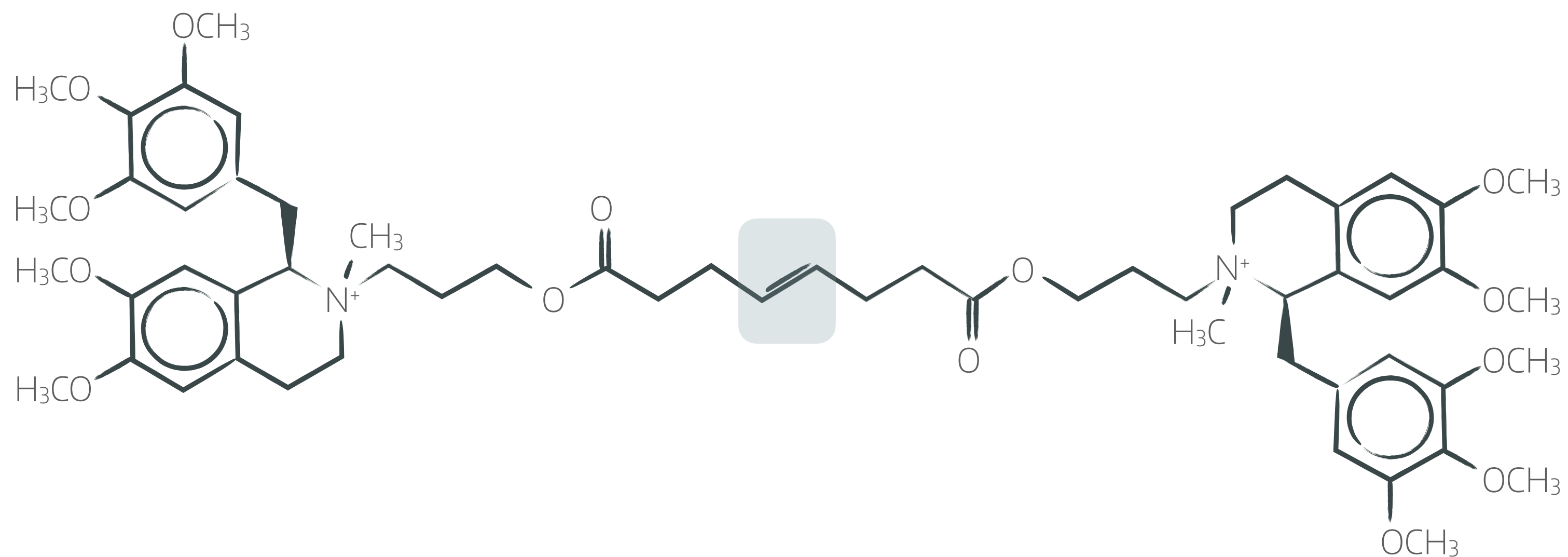


TRANS



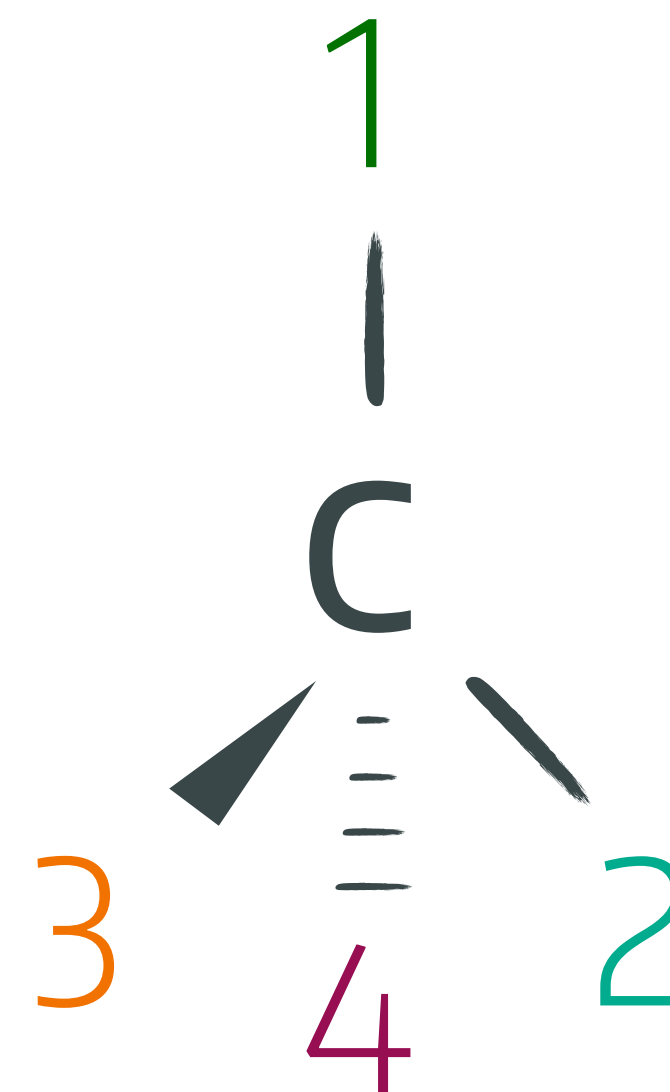
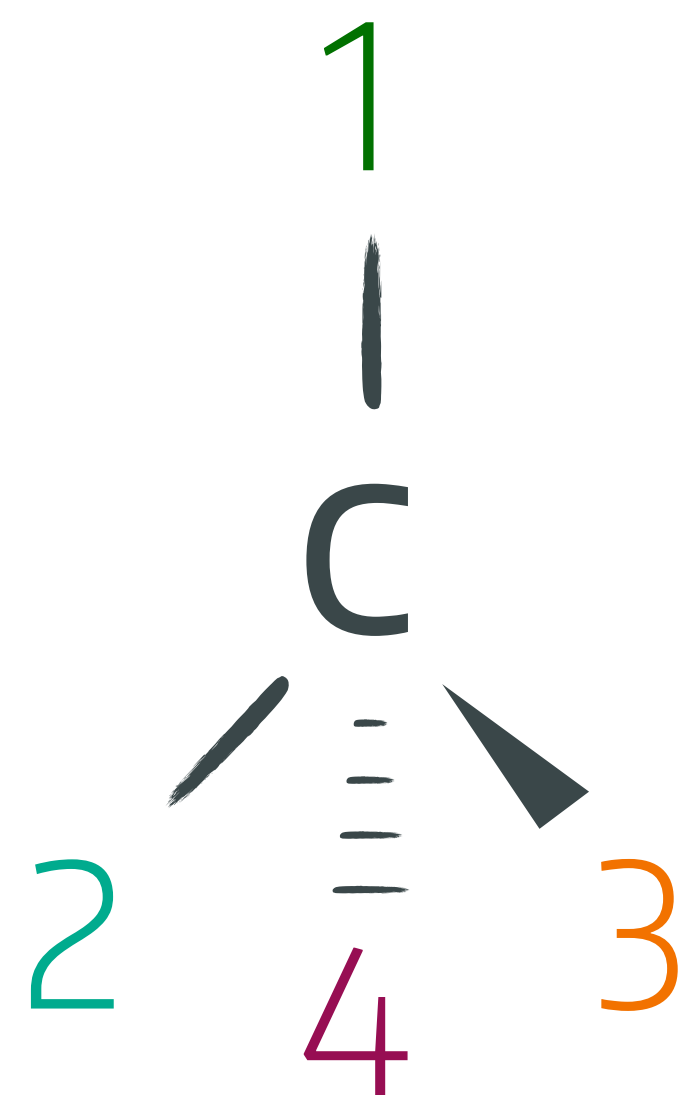
CIS

GEOMETRIC ISOMERISM



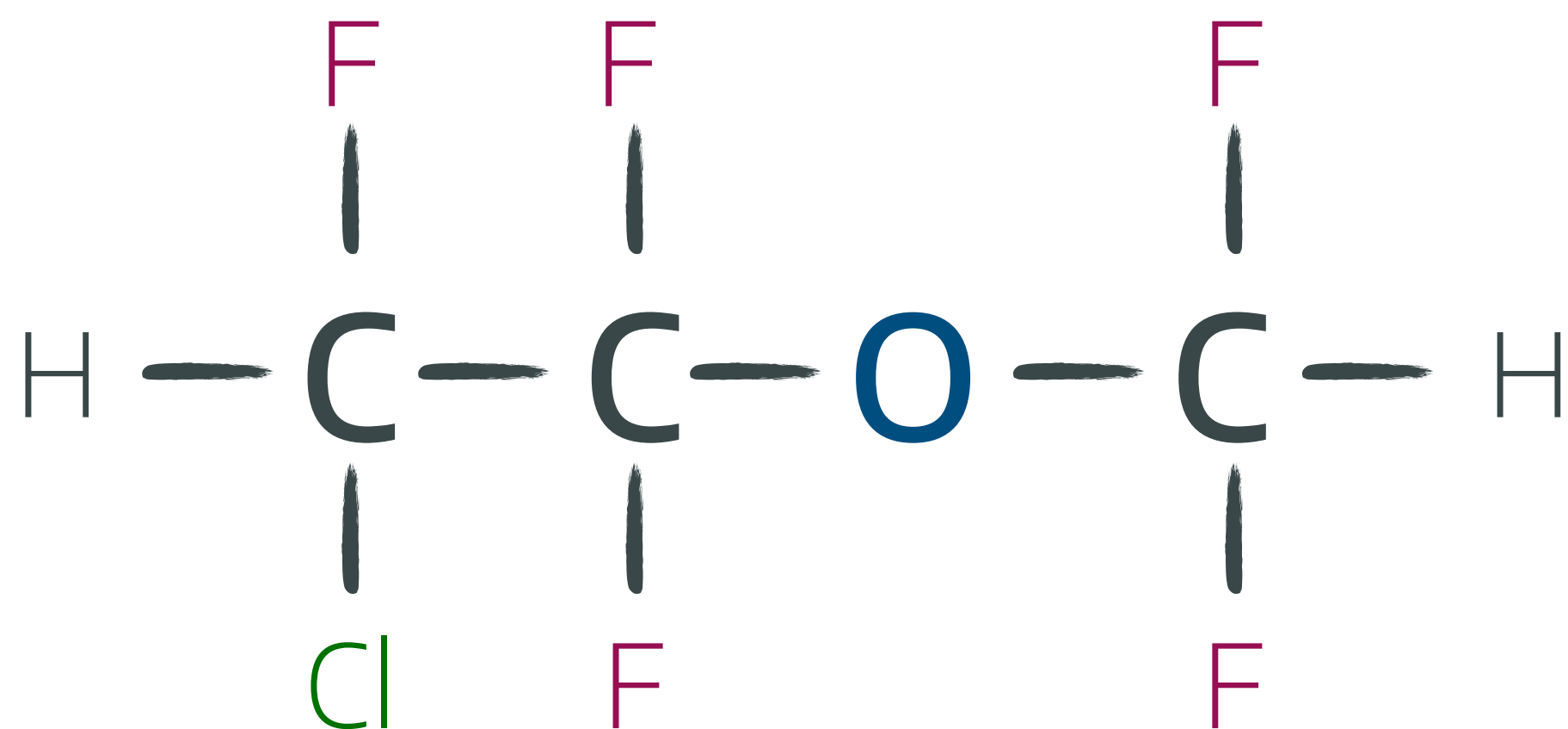
MIVACURIUM

OPTICAL ISOMERISM



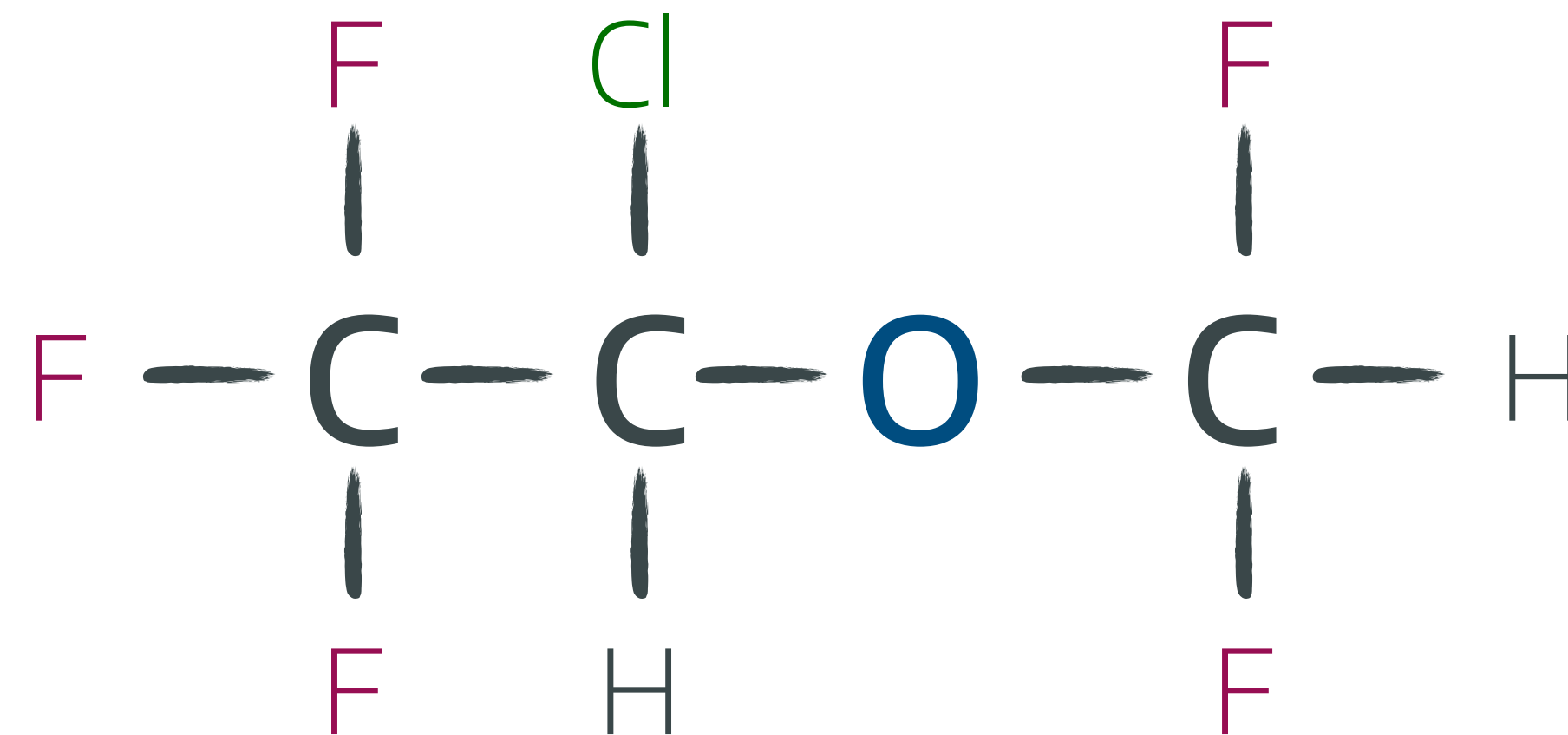
STRUCTURAL ISOMERS

Enflurane



MAC	BP	SVP	B:G
1.68	56	23.1	1.9

Isoflurane

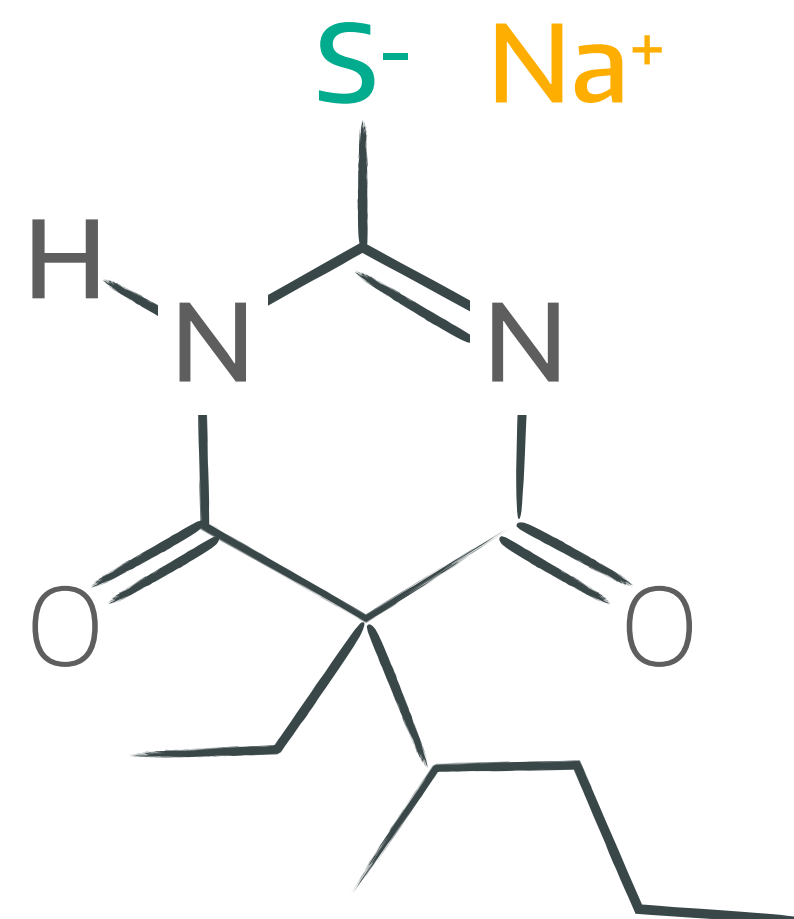


MAC	BP	SVP	B:G
1.15	48	31.5	1.4

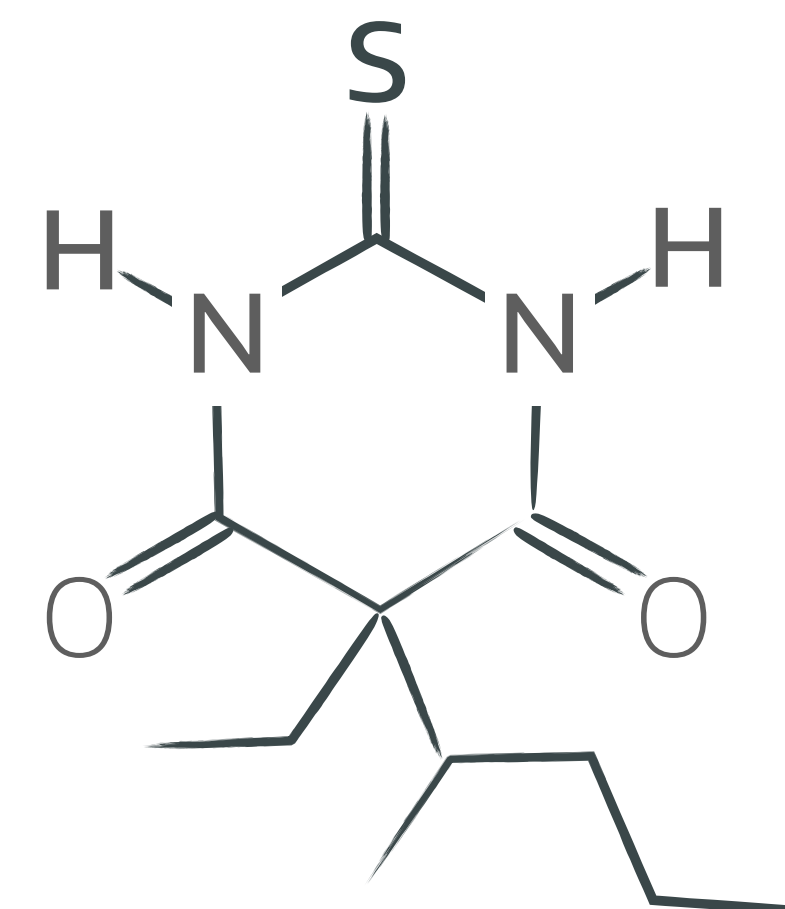
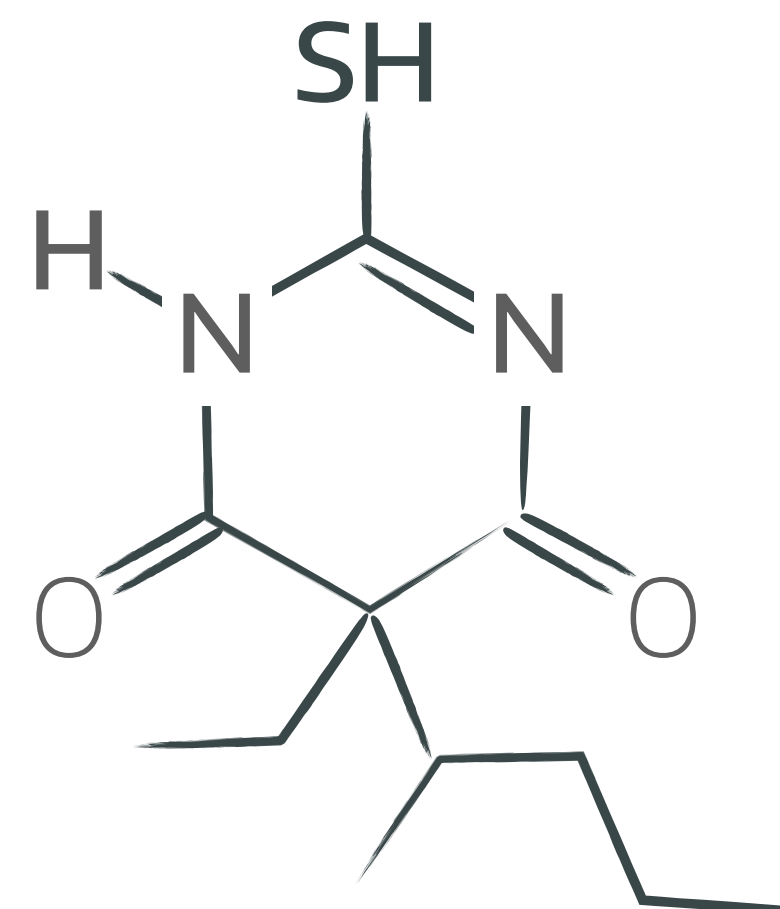
TAUTOMERISM

THIOPENTONE

Ionised

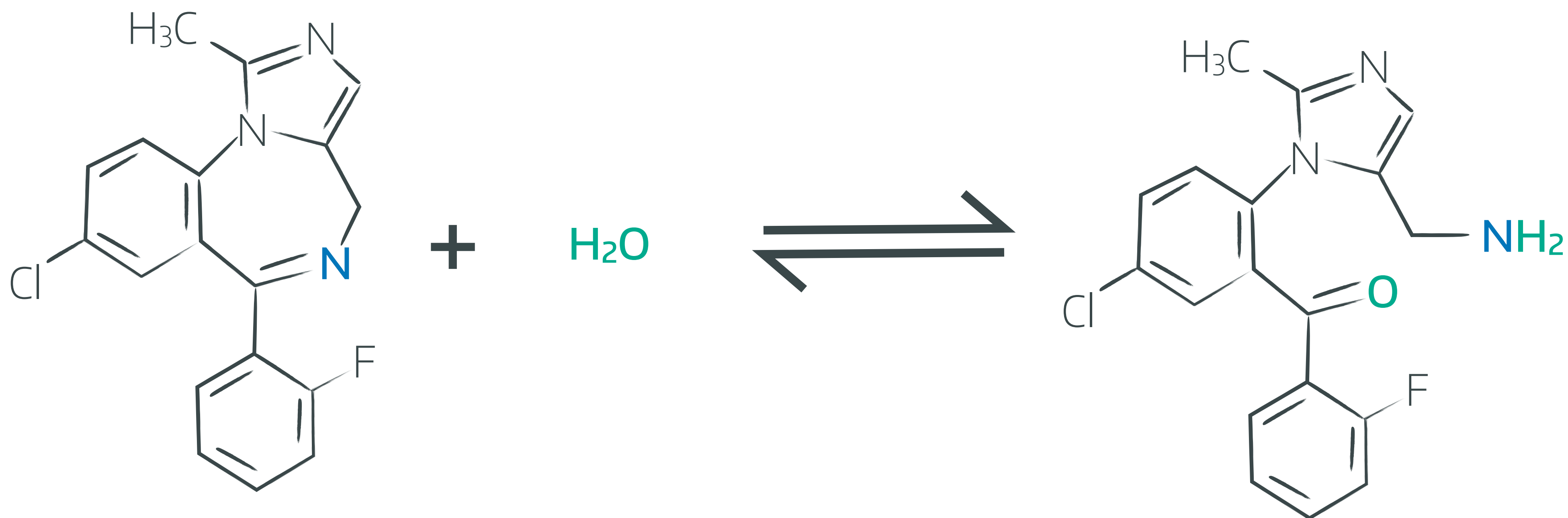


Unionised

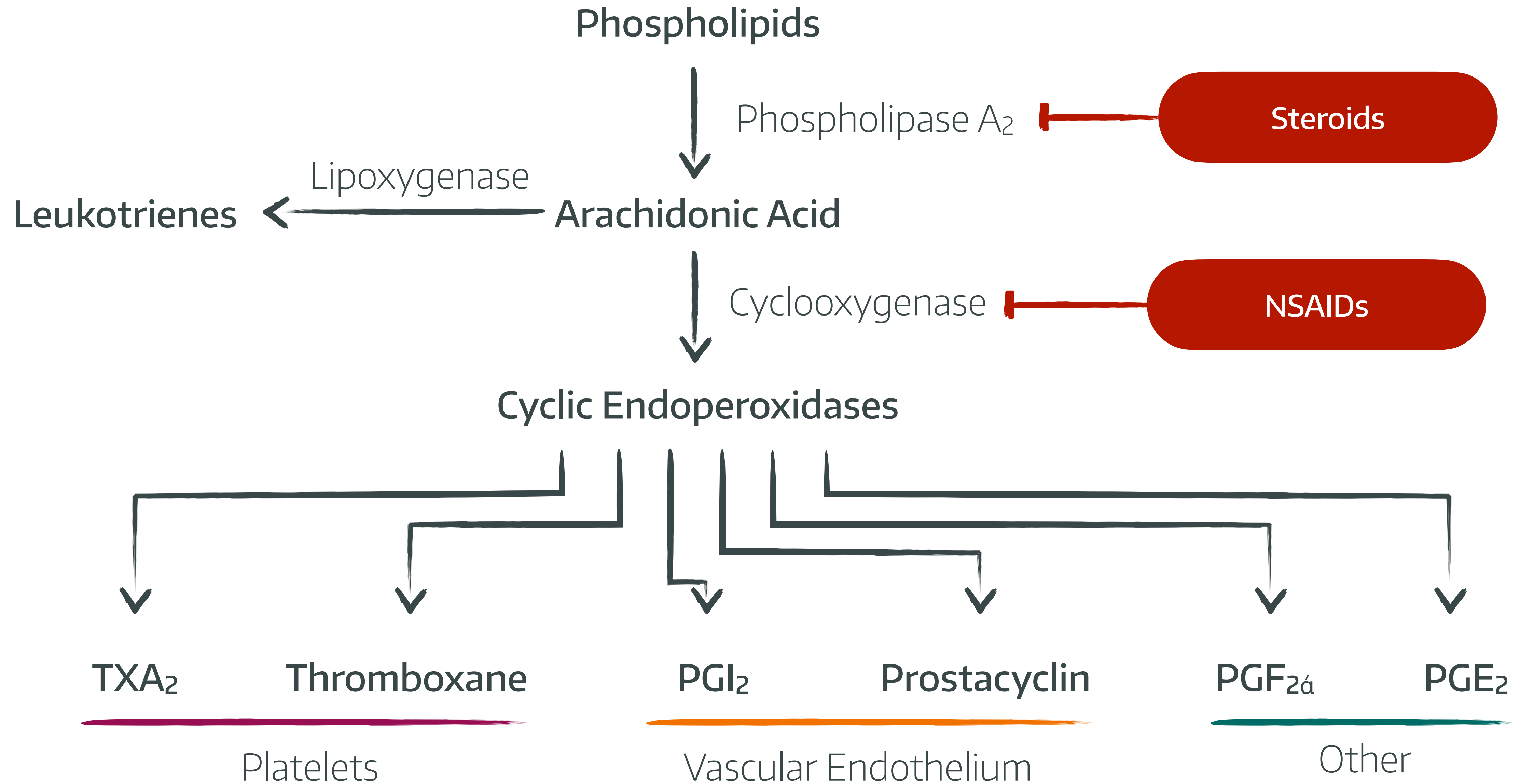


TAUTOMERISM

MIDAZOLAM

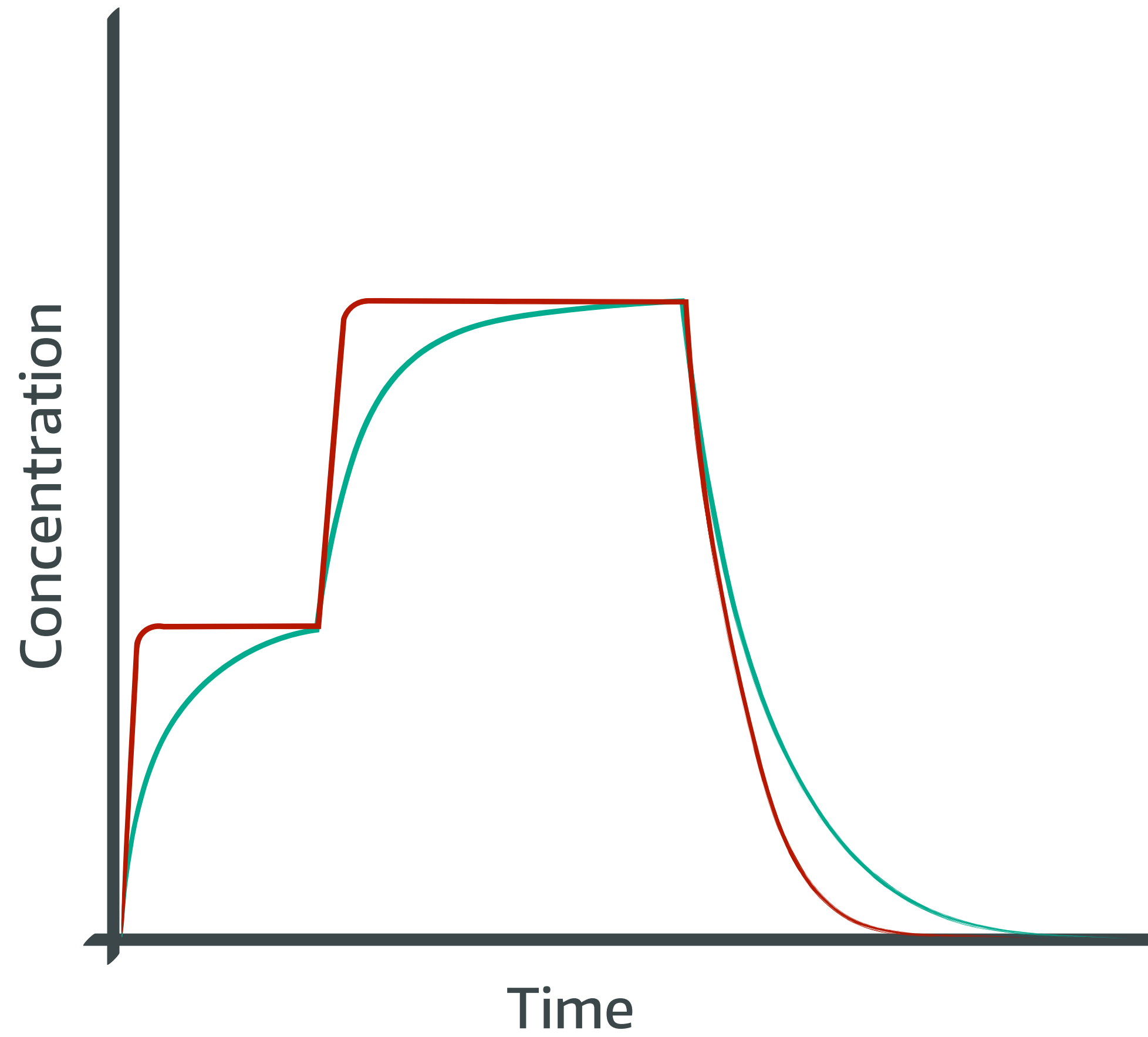


ARACHIDONIC ACID CASCADE

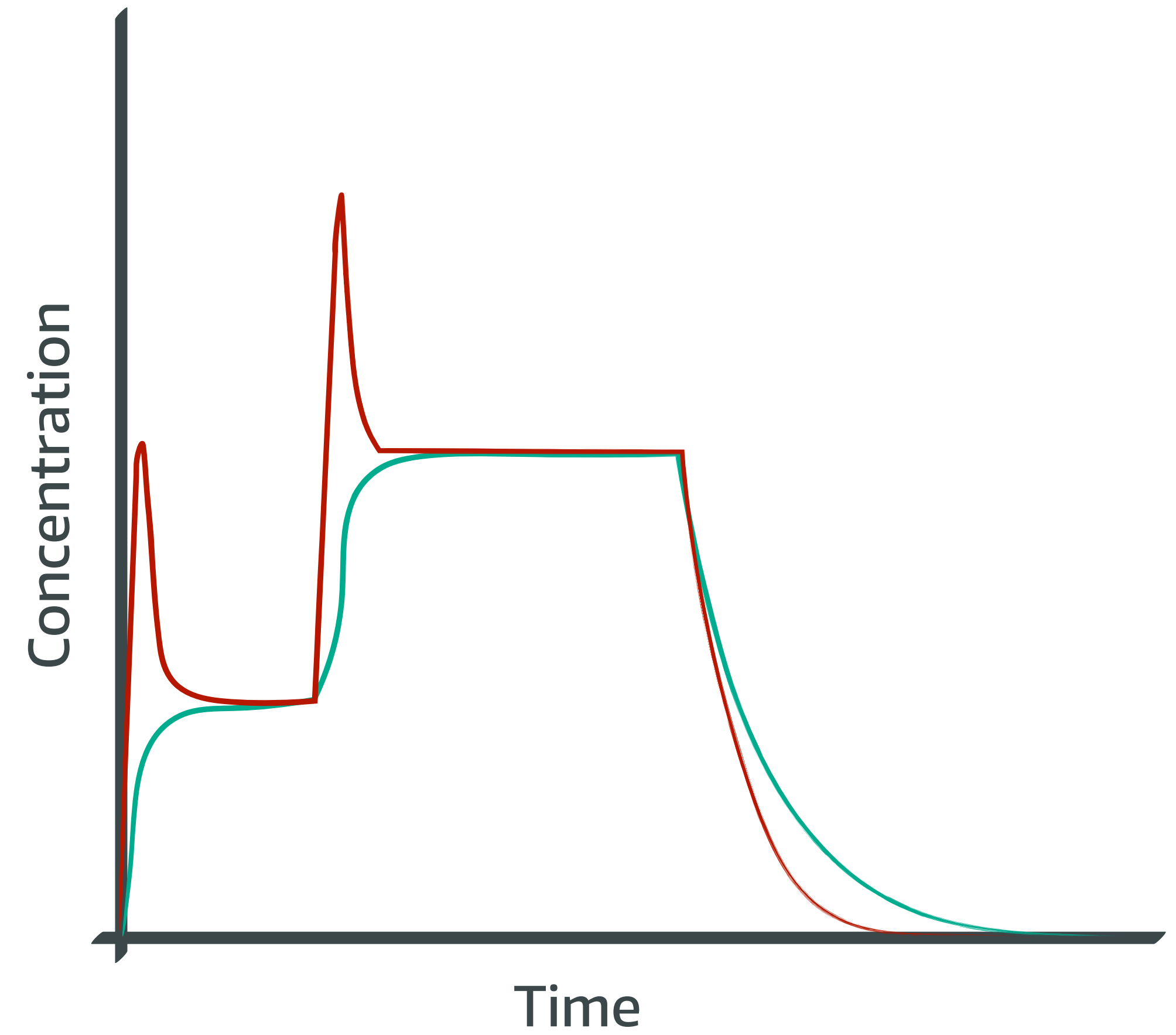


TARGET-CONTROLLED INFUSIONS

Plasma Control

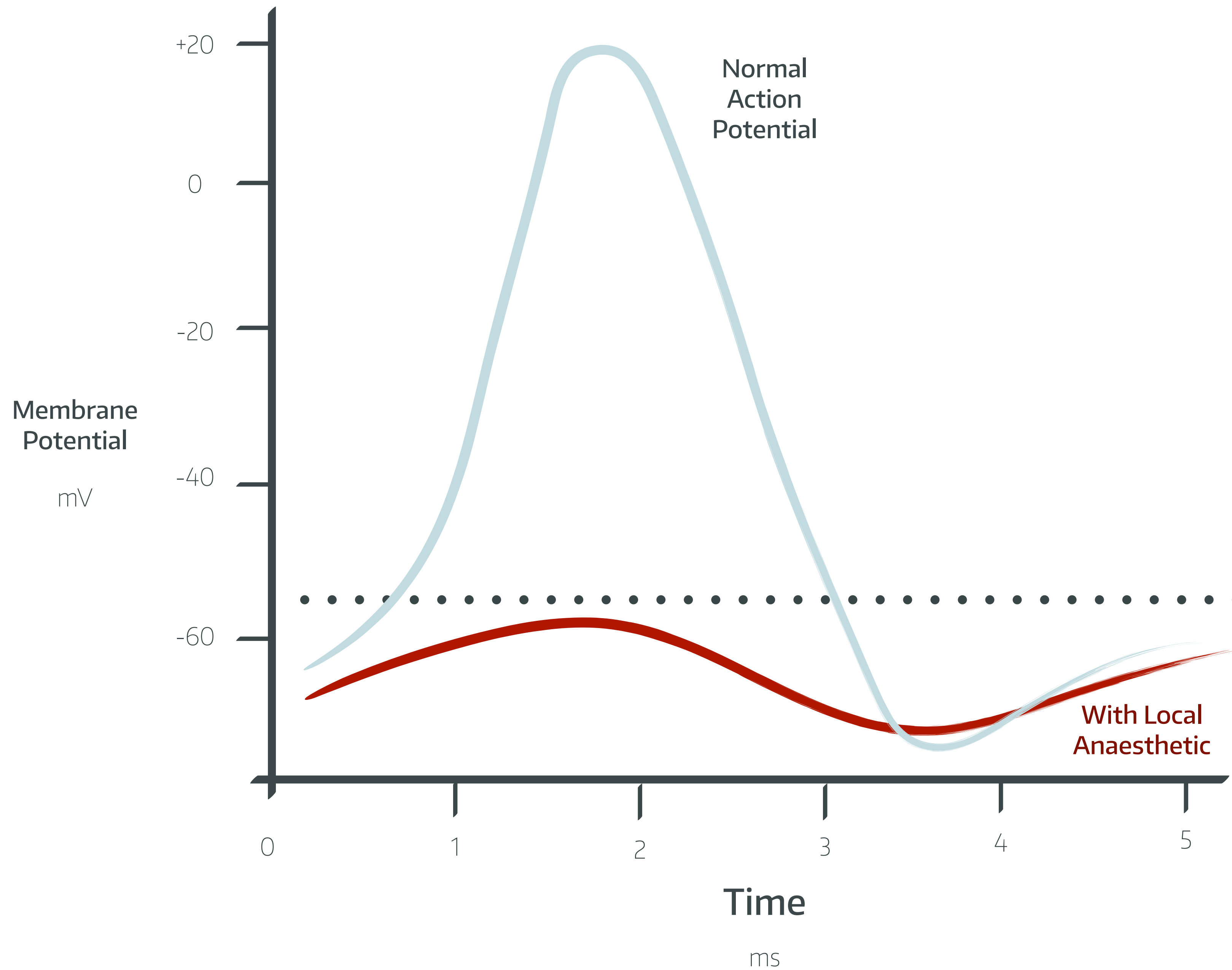


Effect Site Control

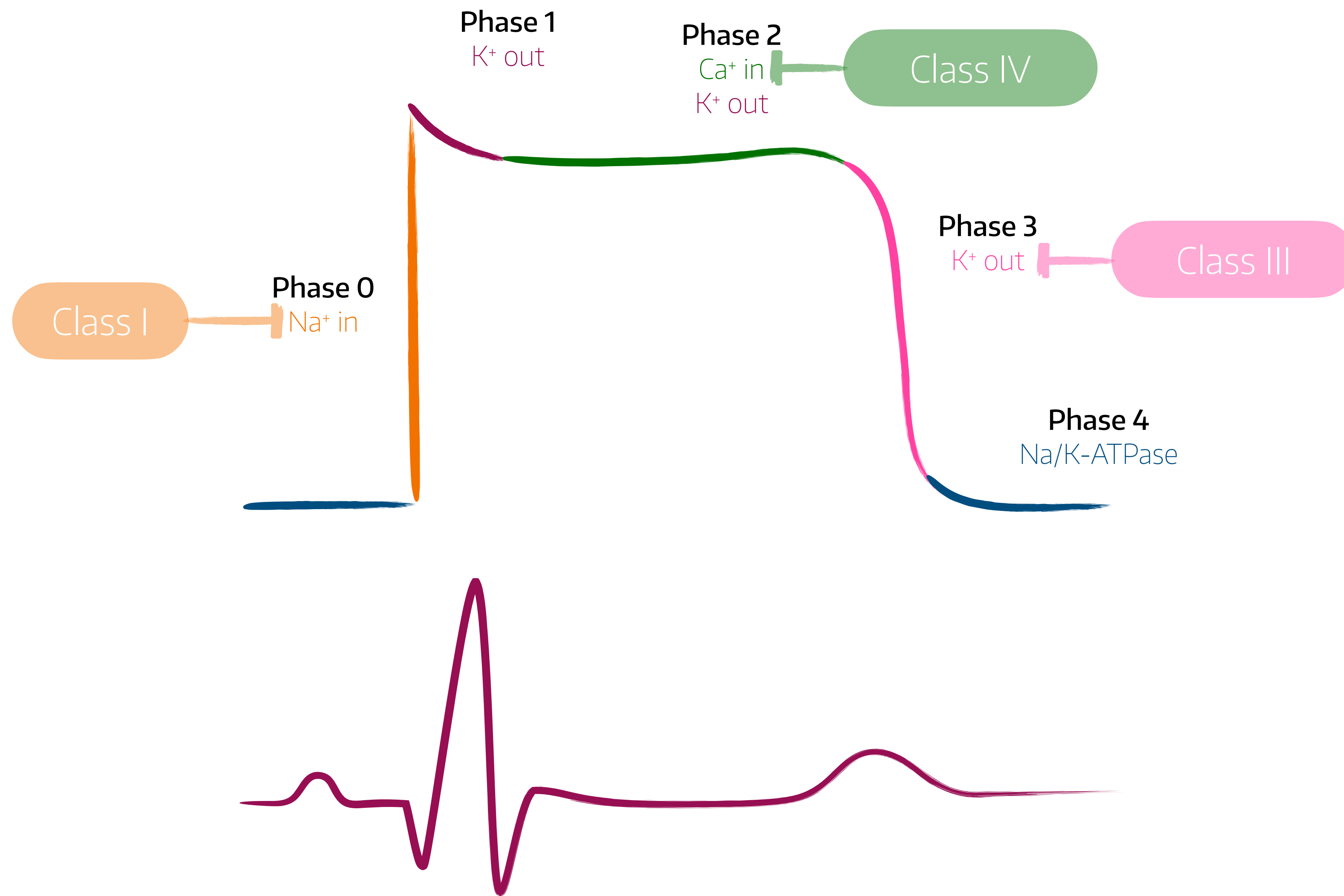


Effect Site
Plasma

LOCAL ANAESTHETICS



ANTIARRHYTHMICS



Indications for Beta-Blockade

Cardiovascular

Angina

Arrhythmia

Hypertension

Heart Failure

Myocardial Infarction

Endocrine

Thyrotoxicosis

Neurological

Migraine Prophylaxis

Anxiety

Glaucoma (topical)

Beta-Blockers

Competitive Antagonists

Selective β_1

Non-Selective

α & β

β_1 & β_2

Bisoprolol

Labetalol

Timolol

Esmolol

Carvedilol

Sotalol

Atenolol

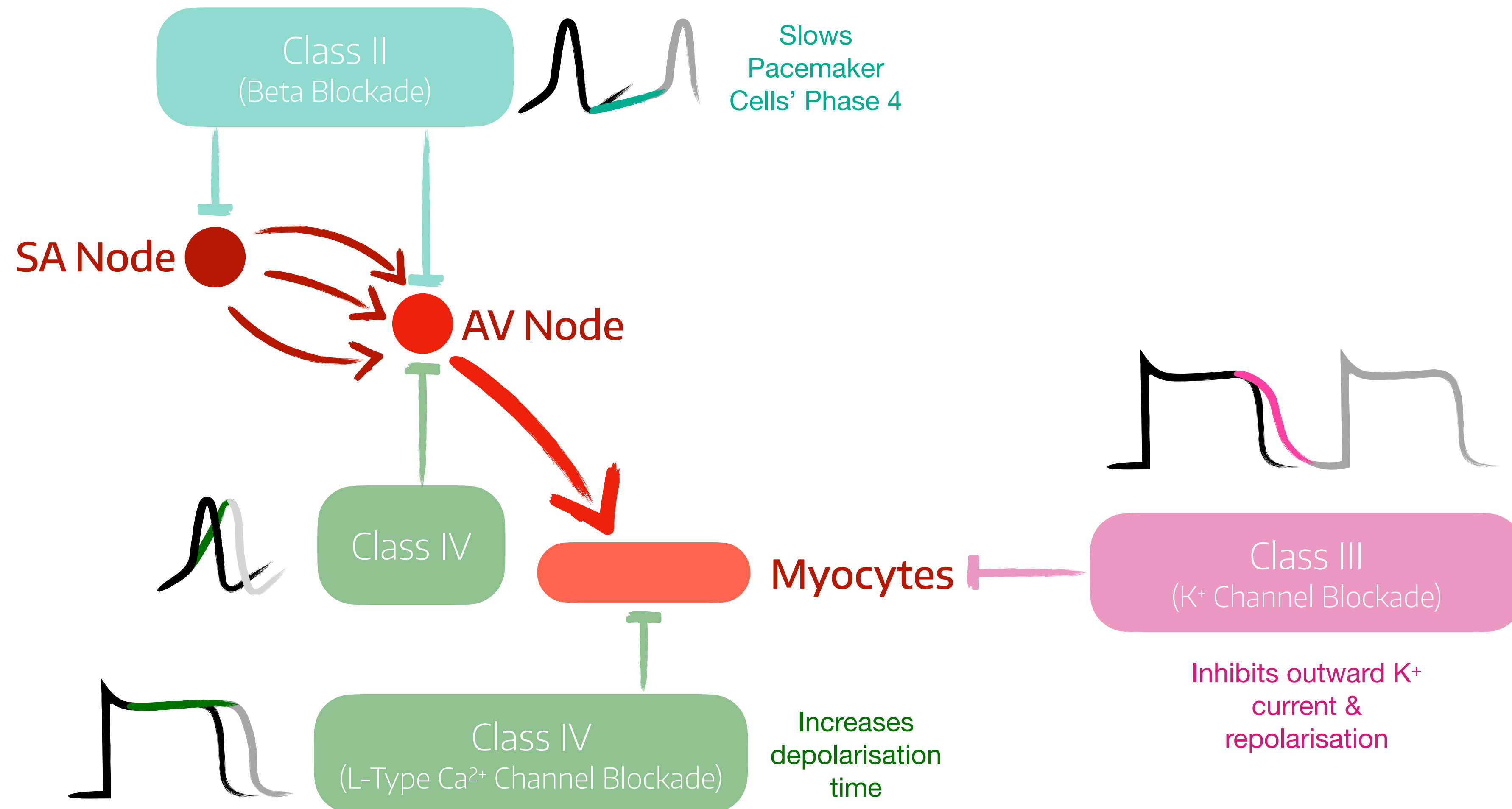
Propranolol

Metoprolol

DRUG ADDITIVES

Buffers	Solvents	Preservatives	Antioxidants
Hydroxide	Water	Benzyl Acohol	Sulphites
	Propylene Glycol		
	Mannitol		
	Soy Bean Oil		

Actions of Amiodarone

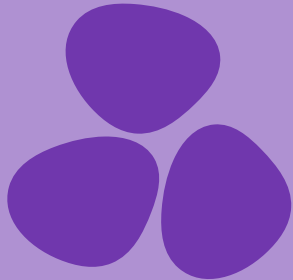


BARIATRIC DRUG DOSING

(Society for Obesity & Bariatric Anaesthesia)

Total Body Weight kg	Lean Body Weight Boer / James / Hume Formulae	Adjusted Body Weight $= 1.4 \times \text{IBW}$
Suxamethonium mg/ml	Propofol (Induction) mg/ml	Propofol (Infusion) mg/ml
LMWHs mg/ml	Thiopentone mg/ml	Neostigmine mg/ml
	(Al)Fentanyl mg/ml	Sugammadex mg/ml
	Morphine mg/ml	Antibiotics mg/ml
	Non-Depolarising NMBDs mg/ml	
	Local Anaesthetics mg/ml	

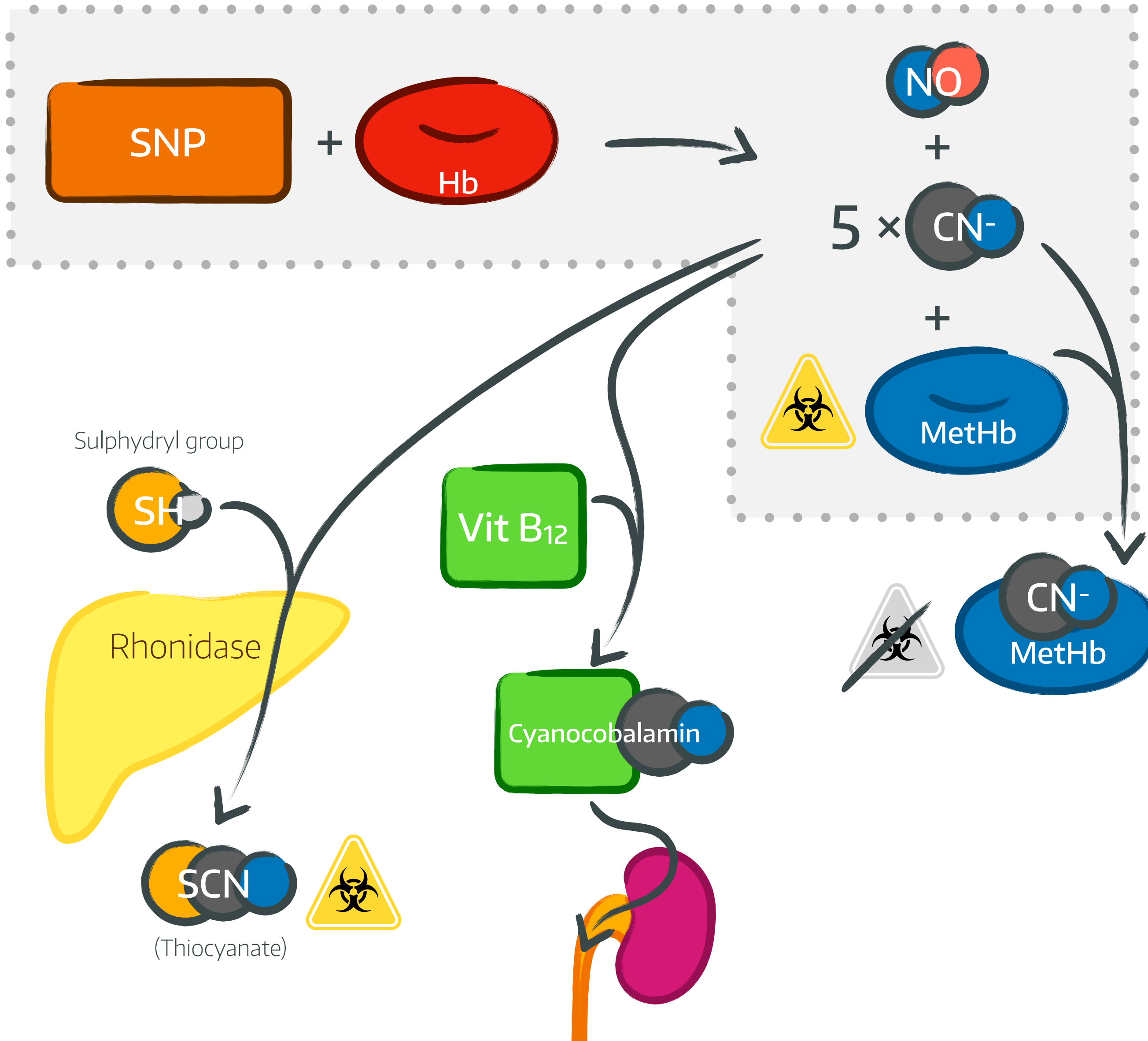
CLASSIFICATION OF BACTERIA

	Gram Positive		Gram Negative	
	Cocci 	Bacilli 	Cocci 	Bacilli 
Aerobic	Staphylococci Streptococci Enterococci	Bacillus	Neisseria Moraxella	Haemophilus Klebsiella Pseudomonas Enterobacter Mycobacteria Campylobacter Legionella Proteus
Anaerobic	Streptococcus viridans	Actinomyces Clostridium Lactobacillus Listeria		Escherichia

THIS IS NOT A COMPREHENSIVE CLASSIFICATION OF ALL BACTERIA, BUT IS A USEFUL TABLE TO BEAR IN MIND WHEN CONSIDERING ANTIBIOTIC THERAPIES.

SODIUM NITROPRUSSIDE

Metabolism

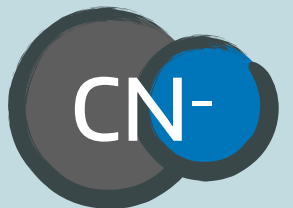


Toxicity



Cyanide

Binds cytochrome oxidase
Impairs aerobic respiration
8micrograms/ml = toxic level



Thiocyanide

100 x less toxic



Treatment

1. Stop SNP
2. Dicobalt Edetate
3. Sodium Thiosulphate (provides sulphhydryl groups)
4. Nitrates (converts oxyhaem to methaem which has higher affinity for CN- than cytochrome oxidase)