

Type I Questions: Choose the one most appropriate answer.

- _____ 1. The following is NOT a ligand-gated channel receptor:
- a) GABA_A
 - b) muscarinic acetylcholine receptor**
 - c) serotonin (5-HT₃) receptor
 - d) AMPA receptor
 - e) nicotinic acetylcholine receptor
- _____ 2. Nuclear receptor coactivators:
- a) are not necessary for nuclear receptor activity
 - b) bind to the enhancer elements
 - c) are ubiquitous
 - d) have histone acetyltransferase activity**
 - e) have histone deacetylase activity
- _____ 3. Which statement is NOT true for nuclear receptor enhancer elements:
- a) they are often palindromic in nature
 - b) they are sometimes direct repeats of a sequence
 - c) the retinoic acid receptor response element (enhancer sequence that binds retinoic acid receptors) is identical for all retinoic acid responsive genes**
 - d) the enhancer sequence that binds the estrogen receptor is based on a consensus
 - e) can be found far away from the promoter
- _____ 4. The following statement about the insulin receptor tyrosine kinase is NOT true:
- a) phosphorylates other insulin receptors
 - b) catalyzes transphosphorylation
 - c) can phosphorylate ras**
 - d) can phosphorylate certain other kinases
 - e) can phosphorylate proteins that other receptor tyrosine kinases do not
- _____ 5. Which of the following statements is FALSE
- a) Absorption of a weakly basic drug is likely to occur more readily from alkaline intestine than from the acidic stomach
 - b) Alkalinization of the urine from pH 6 to pH 7 would help to eliminate amphetamine, a basic drug (pK_a=8.5)**
 - c) A weak acid (pK_a=3.5) is more lipid-soluble at pH 2 than at pH 3
 - d) Very few drugs cross cell membranes by way of endocytosis

- e) Uncharged molecules cross cell membranes (by diffusion) more readily than charged molecules

_____ 6. Which of the following is responsible for the step of translocation of dopamine into synaptic vesicles in the presynaptic neuron:

- a) **Vesicular monoamine transporter (VMAT)**
- b) Dopamine transporter
- c) Monoamine oxidase (MAO)
- d) Metabolism by catechol-O-methyltransferase (COMT)
- e) Aromatic L-amino acid transporter

Type II Questions: Questions in this section are to be answered by using the key given below

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- a) If only 1, 2 and 3 are correct
 - b) If only 1 and 3 are correct
 - c) If only 2 and 4 are correct
 - d) If only 4 is correct
 - e) If all are correct

_____ 1. The following statement(s) about acetaminophen is true:

- 1. **it belongs to the para-aminophenol group of compounds**
- 2. its major benefit is its long half-life compared with ASA
- 3. **major toxicity can result from nucleophilic attack on liver cell proteins by a toxic intermediate**
- 4. it is a potent anti-inflammatory

_____ 2. Side effects of aspirin can include:

- 1. **stomach irritation**
- 2. **blood clots**
- 3. **prevention of blood coagulation**
- 4. hypersensitivity to acetaminophen

_____3. Acetylcholine is released from which of the following tissues?

1. **postganglionic parasympathetic neurons innervating intestinal smooth muscle**
2. **postganglionic sympathetic neurons innervating sweat glands**
3. **preganglionic neurons innervating sympathetic ganglia**
4. adrenal medulla

_____4. Acetylsalicylic acid can inhibit both COX-1 and COX-2 because

1. it can fit into the COX-2 side pocket
2. it undergoes extensive first pass metabolism
3. it is more potent than most Coxibs
4. **it acetylates a serine residue in both enzymes**

_____5. Norepinephrine:

1. Is the main neurotransmitter at nicotinic receptors
2. Is the main neurotransmitter at muscarinic receptors
3. Can be ingested orally
4. **Interacts widely with alpha and beta receptors**

_____6. The following statements about Ritalin are true:

1. **Can increase stimulatory impulses in the brain to increase the “signal to noise ratio” of nerve transmission**
2. **Can increase blood pressure**
3. **Can increase heart rate**
4. Has a better affinity for beta receptors than alpha receptors.

_____7. A drug is eliminated by renal filtration and the elimination is a first-order elimination process. The elimination half-life of the drug will depend on the following:

1. volume of distribution
2. extent of plasma protein binding
3. elimination rate constant
4. **plasma concentration**