The surface of a cell has a receptor where ligands become attached.

- **A.** For antigen-detecting receptors like the T Cell Receptor, these ligands come in two types: agonists and antagonists. Assume that molecules arrive according to a Poisson process with rate λ . Among these molecules, a proportion α are agonists (so $0 \le \alpha \le 1$) and the rest are antagonists. Antagonists remain attached for an exponentially-distributed duration with parameter μ_1 , while agonists remain attached for an exponentially-distributed duration with parameter μ_2 . An arriving molecule only becomes attached if the receptor is free of other molecules.
 - What is the percentage of time the receptor is occupied by an agonist? By an antagonist? Free?
- **B.** For this part, assume there is only one type of ligand. The ligand binds to the receptor at rate λ and unbinds at rate μ . Once the ligand is bound, the receptor initiates an intracellular signal at rate k, but it can only do this while a ligand is bound.
 - (a) Write a 3-state continuous-time Markov transition matrix to describe this process.
 - (b) Suppose that at time t = 0, there is a ligand bound to the receptor. How long on average until a signal is transduced?
 - (c) Suppose that the receptor's kinetics are controlled by a control variable, so that $\lambda = c\lambda^*$ and $\mu = c\mu^*$. Again suppose the system starts with a ligand bound. Assume receptor kinetics have been slowed to approximately zero speed, so c = 0, i.e., $\lambda = \mu = 0$. In other words, the ligand is very unlikely to ever unbind.
 - i. According to your intuition, what is the mean time until a signal is transduced?
 - ii. Set $\lambda = \mu = 0$ in the transition matrix. What is the mean time until a signal is transduced? Does this agree with your intuition?
 - iii. Set $\lambda = \mu = 0$ in your answer from Part (b). What is the mean time until a signal is transduced? Does this agree with your intuition?