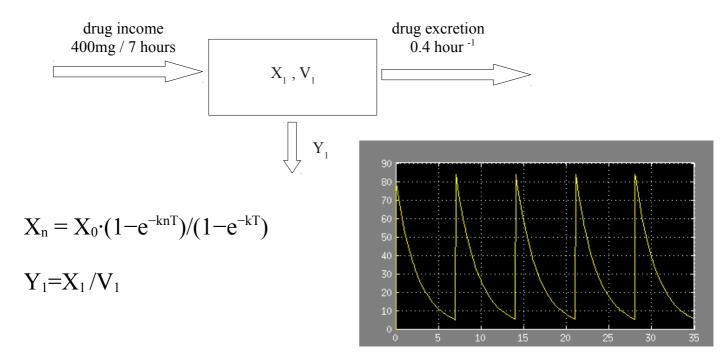
Pharmacokinetics model

Case 1:



Symbol	Importance	Value	Units	
X	ammount of drug	init value: 0	mg	Var
Y	concentration of drug	init value: 0	mg/liters	Variab
V	volume of the system	5	liters	q
K_1	income of the drug	400	mg/time ⁻¹	Param
K _e	rate of drug excretion	0.4	time ⁻¹	n

Case 2:

$$X1' = (-k12-k13)*X1+k21*X2+k31*X3$$

$$X2' = k12*X1-k21*X2$$

$$X3' = k13*X1-k31*X3$$

$$X1' = -0.9*X1+0.2*X2+0.7*X3$$

$$X2' = 0.6*X1-0.2*X2$$

$$X3' = 0.3*X1-0.7*X3$$

$$C = \begin{bmatrix} -0.9 & 0.2 & 0.7 \\ 0.6 & -1 & 0 \\ 0.3 & 0 & -0.7 \end{bmatrix}$$

$$D = \begin{bmatrix} 1 \\ 0 \\ 0 \end{bmatrix}$$

$$D = \begin{bmatrix} 0 \\ 0 \end{bmatrix}$$

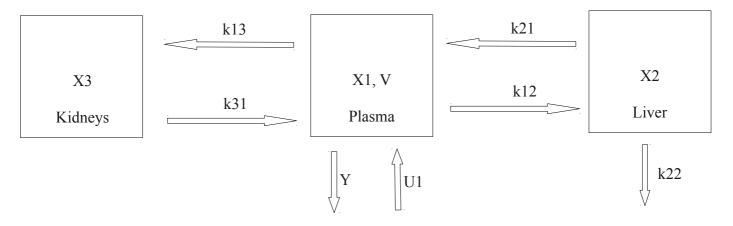


Table of variables

X1	ammount of drug in plasma	init value: 0	mg
Y	concentration of drug in plasma	init value: 0	mg/liters
X2	ammount of drug in kidneys	init value: 0	mg
Х3	ammount of drug in intestine	init value: 0	mg

Table of parameters of the system

Symbol	Importance	Value	Units
V	volume of blood in the system	5	liters
k12	drug transfer from plasma to kidneys	0.6	time ⁻¹
k21	drug transfer from kidneys to plasma	0.2	time ⁻¹
k13	drug transfer from plasma to kidneys	0.3	time ⁻¹
k31	drug transfer from intestine to plasma	0.7	time ⁻¹
k22	excretion rate from the kidneys	0.8	time ⁻¹
U1	drug income	different	mg/time ⁻¹

Figure 2: output of all 3 types of simulation.

