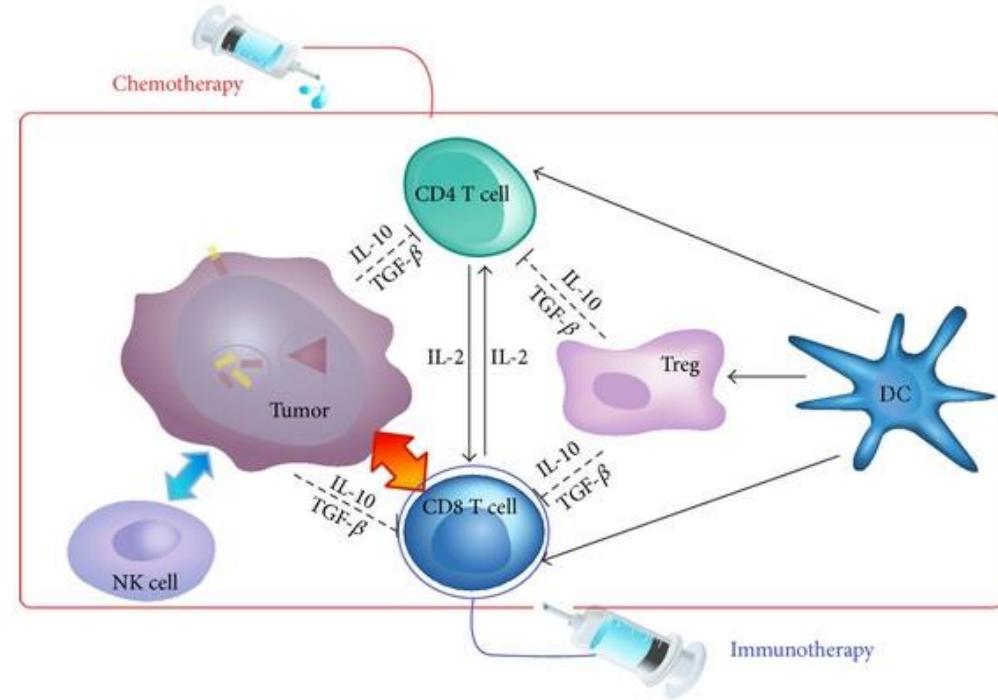


Dynamics of the Kirschner- Panetta Model

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About the model

- Created by Denise Kirschner and John Carl Panetta.
- Published in Journal of Mathematical Biology (1998).
- Wanted a model for short and long-term tumor behavior.
- Rich dynamics for exploring tumor oscillation, relapse, and elimination.



Kim, Kwang Su, Cho, Giphil, Jung, Il Hyo, Optimal Treatment Strategy for a Tumor Model under Immune Suppression, *Computational and Mathematical Methods in Medicine*, 2014, 206287, 13 pages, 2014.
<https://doi.org/10.1155/2014/206287>

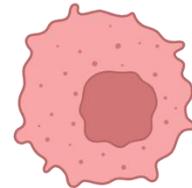
Defining the model

$$\frac{dT}{dt} = r_2(T)T - \frac{\alpha ET}{g_2 + T}$$

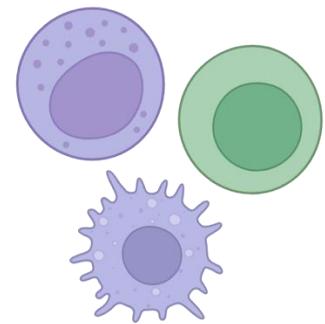
$$\frac{dE}{dt} = cT - \mu_2 E + \frac{p_1 EI_{L2}}{g_1 + I_{L2}} + s_1$$

$$\frac{dI_{L2}}{dt} = \frac{p_2 ET}{g_3 + T} - \mu_3 I_{L2} + s_2$$

Tumor Cells

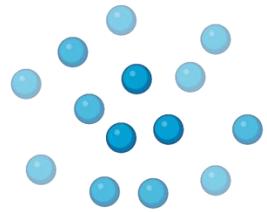


Effector Cells



NK Cells,
T Cells,
Macrophages

**Interleukin-2
(IL-2)**



Non-dimensionalization

- The model is non-dimensionalized for computation.
- This handles large changes more smoothly.

$$\frac{dy}{d\tau} = r_2(1 - by)y - \frac{\alpha xy}{g_2 + y}$$

$$\frac{dx}{d\tau} = cy - \mu_2 x + \frac{p_1 xz}{g_1 + z} + s_1$$

$$\frac{dz}{d\tau} = \frac{p_2 xy}{g_3 + y} - \mu_3 z + s_2$$

Constants in the model

Parameter	Value	Unit	Description
c	$0 \leq c \leq 0.05$	day^{-1}	The antigenicity of tumour; larger values of c represent well recognized antigens
μ_2	3.00×10^{-2}	day^{-1}	Multiplicative inverse of the natural lifespan for effector cells
p_1	1.245×10^{-1}	day^{-1}	Proliferation rate of effector cells estimated by using experimental data (see Kuznetsov et al. 1994)
g_1	2.00×10^7	IU. L^{-1}	Threshold for proliferation of effector cells stimulated by IL-2
s_1		cell. day^{-1}	External source of effector cells
r_2	1.80×10^{-1}	day^{-1}	The logistic growth rate of tumour cells in the absence of an immune response
b	1.00×10^{-9}	cell^{-1}	Multiplicative inverse of the tumour's carrying capacity
a	1.00	day^{-1}	Immune system's strength to eliminate cancer cells
g_2	1.00×10^5	cell	Threshold for cancer removal
p_2	5.00	$\text{IU. L}^{-1}. \text{cell}^{-1}. \text{day}^{-1}$	production rate of IL-2
g_3	1.00×10^3	cell	Threshold for production of IL-2 due to the interaction between cancer cells and effector cells
μ_3	1.00×10^1	day^{-1}	Multiplicative inverse of the lifespan for IL-2
s_2		$\text{IU. L}^{-1}. \text{day}^{-1}$	External source of IL-2

Hamiltonian of the model

Functional:

$$F(s_1, s_2) = \int_0^{t_f} Ax(t) - By(t) - C_1 s_1(t) - C_2 s_2(t) + Dx(t)z(t) - \gamma z(t)^2 dt$$

Lagrangian:

$$\mathcal{L} = -Ax(t) + By(t) + C_1 s_1(t) + C_2 s_2(t) - Dx(t)z(t) + \gamma z(t)^2$$

Hamiltonian:

$$\mathcal{H}(x, y, z, \lambda_1, \lambda_2, \lambda_3) = -\mathcal{L} + \lambda_1 \frac{dx}{dt} + \lambda_2 \frac{dy}{dt} + \lambda_3 \frac{dz}{dt}$$

Functional integrand components

Functional:

$$F(s_1, s_2) = \int_0^{t_f} Ax(t) - By(t) - C_1 s_1(t) - C_2 s_2(t) + Dx(t)z(t) - \gamma z(t)^2 dt$$

Ax(t): Immune cell population.

By(t): Tumor cell population.

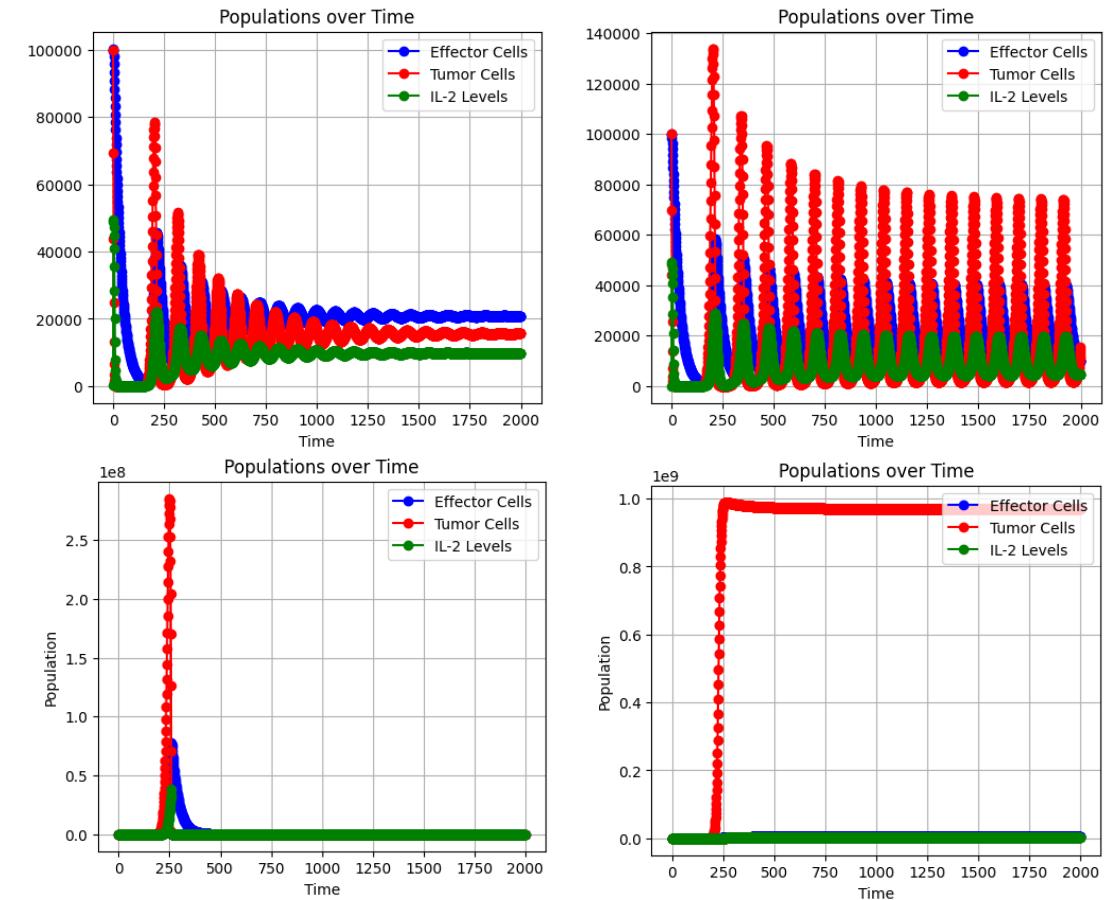
C₁s₁(t): Immune cell treatment dose, C₂s₂(t): IL – 2 treatment dose.

*Dx(t)z(t): Immune cell population * IL – 2, IL – 2 is crucial to immune cell activation, and immune cell presence is essential for IL – 2 treatment effectiveness. This term reflects this dependent relationship.*

γz(t)²: IL – 2 dosage is also comes with toxicity to some extent. Too high dosage can lead to immune cell exhaustion, serious auto – immune reaction, or a phenomenon known as cytokine storm. This term reflects these detrimental effects on the system.

Model behavior without treatment

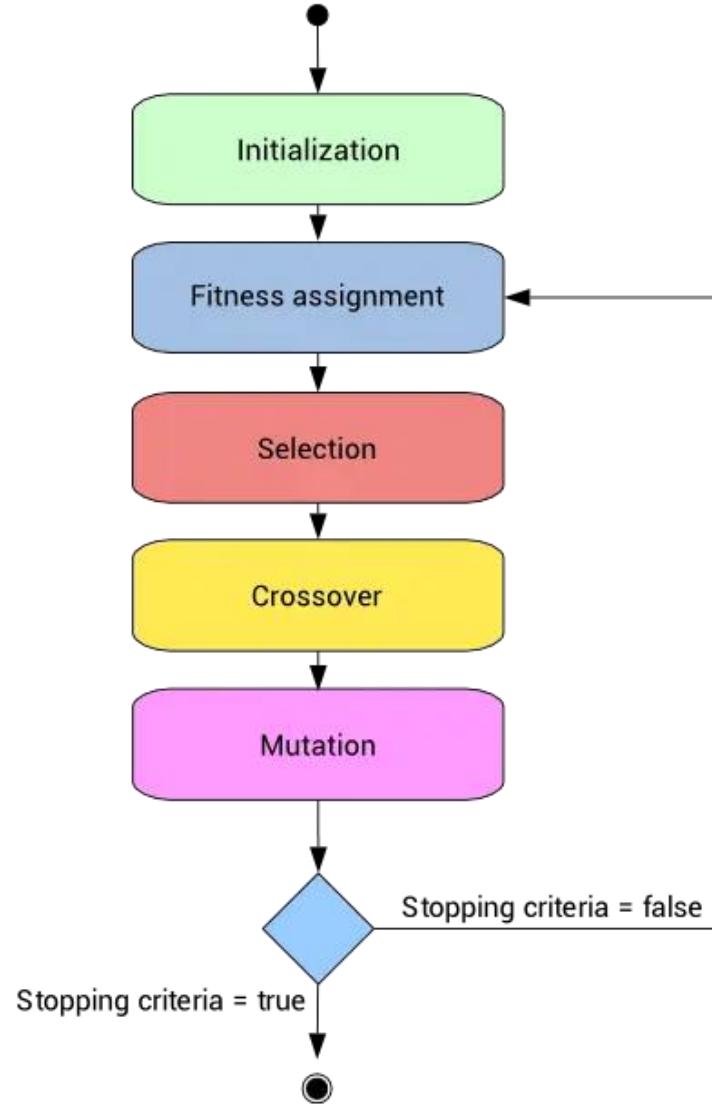
- Effector cell dose, $s_1 = 0$
- IL-2 dose, $s_2 = 0$
- Changing antigenicity (c) can lead to oscillatory population dynamics.



$$c = 0.04, 0.0297, 8e-3, 8e-5$$

Genetic Algorithm (GA)

- Shown to help control chaotic systems
- Fitness function defines how ‘good’ the GA’s solution is.
- GA defines s_1 and s_2 doses.



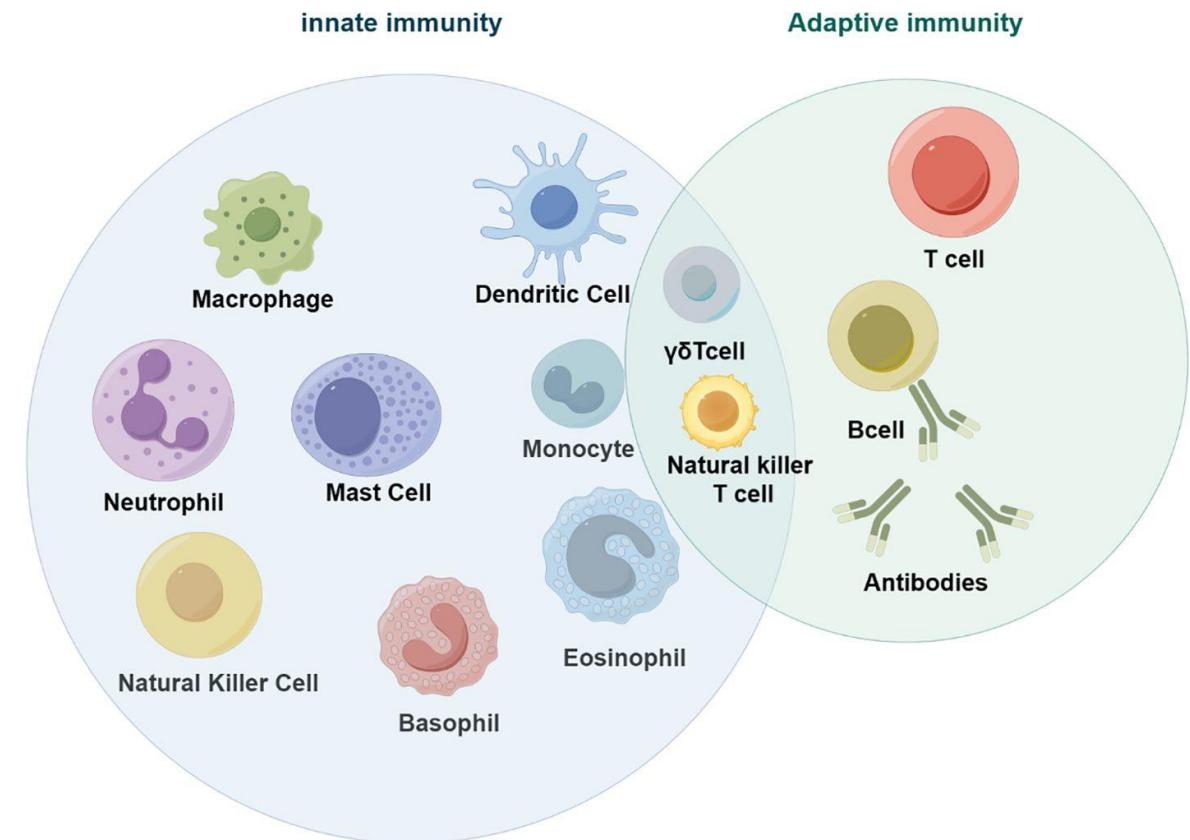
GA: Fitness function

- Can tune for biological relevance.
- Similar but not the same as the integrand of the functional.

```
def fitness_func(ga_instance, solution, solution_idx):  
    """  
    Fitness function maximizing E and minimizing T.  
    """  
    t = ga_instance.environment['t']  
    x = ga_instance.environment['x']  
    y = ga_instance.environment['y']  
    z = ga_instance.environment['z']  
  
    # Extract genes from solution  
    genes1 = solution[0:4]  
    genes2 = solution[4:8]  
  
    # Calculate s_1 and s_2 based on genes and current state  
    s_1 = genes1[0]*x + genes1[1]*y + genes1[2]*z + genes1[3]  
    s_2 = genes2[0]*x + genes2[1]*y + genes2[2]*z + genes2[3]  
  
    # Restrict negative input  
    s_1 = max(0, s_1)  
    s_2 = max(0, s_2)  
  
    E_input = s_1 # Effector cell input from GA  
    IL_input = s_2 # IL-2 input from GA  
  
    t_step = 1 # time step for prediction  
  
    # Calculate derivatives using non-dimensional KP model equations  
    x_pred = x + dx_dt(t=t, x=x, y=y, z=z,  
                         c=0.02, mu_2=0.03, p_1=0.1245, g_1=2e4, s_1=E_input) * t_step  
    y_pred = y + dy_dt(t=t, y=y, x=x, z=z,  
                         r_2=0.18, b=1e-5, alpha=0.002, g_2=1e5) * t_step  
    z_pred = z + dz_dt(t=t, z=z, x=x, y=y,  
                         p_2=5e-7, g_3=1e4, mu_3=10, s_2=IL_input) * t_step  
  
    # Define fitness as maximizing dE_dt and minimizing dT_dt.  
    # Penalize over dose of IL-2.  
  
    a1, a2 = 0.1, 0.1 # weights for immunotherapy components  
    b1, b2, b3, b4 = 0.1, 0.1, 0.1, 0.1 # weights for toxicity components  
  
    immunotherapy = a1*(x_pred) - a2*(y_pred) # Reward high effector cells and low tumor cells  
    toxicity = b1*s_2 + b2*s_1 + b3*(x_pred*s_2) + b4*(z_pred)**2 # Penalty for IL-2 overdose  
  
    c1, c2 = 1.0, 3.0  
  
    fitness = c1*immunotherapy - c2*toxicity # final fitness
```

Immunology and cancer

- Immune cell exhaustion
- Autoimmune response
- Cytokine storm
- Mutational burden
- Immunotherapy



Repository

<https://github.com/jrosgaard/Workspace>

The screenshot shows a GitHub repository page with a dark theme. At the top, there are links for 'README' (highlighted with a red underline) and 'License'. On the right side, there is a pencil icon for editing. The main title of the repository is 'Kirschner Panetta Tumor immunotherapy model' in large white font. Below the title, a description states: 'This project aimed to use a genetic algorithm to perform closed-loop optimization of immunotherapy dosing in the Kirschner-Panetta model.' At the bottom, another description reads: 'Using the slider notebook, you are able to adjust the antigenicity (c) for both the model without treatment and with the GA-optimized treatment regime.' The background of the page is dark, and the text is white or light gray.

References

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