Recent topics in the modeling and analysis of diffuse interface tumor growth

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NSF-CBMS Conference on THE CAHN-HILLIARD EQUATION:
RECENT ADVANCES AND APPLICATIONS

joint works with Cecilia Cavaterra (Milano) – Hao Wu (Fudan) and Alain Miranville (Poitiers) – Giulio Schimperna (Pavia)





Fondazione Cariplo and Regione Lombardia Grant MEGAsTaR 2016-2019

Outline

- Phase field models for tumor growth
- 2 The model HZO by [A. Hawkins-Daarud, K.-G. van der Zee and J.-T. Oden (2011)]
- 3 Joint work with C. Cavaterra and H. Wu, AMO (online)
- Well-posedness
- 5 Long-term dynamics
- 6 The optimal control problem
- Joint work with A. Miranville and G. Schimperna, JDE (online)
- Well-posedness
- Dissipativity and existence of the attractor
- Perspectives and Open problems

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Setting

Tumors grown in vitro often exhibit "layered" structures:

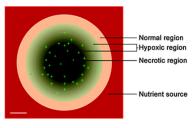


Figure: Zhang et al. Integr. Biol., 2012, 4, 1072–1080. Scale bar $100\mu m = 0.1 mm$

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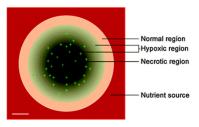


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- sharp interfaces are replaced by narrow transition layers arising due to adhesive forces among the cell species: a diffuse interface separates tumor and healthy cell regions
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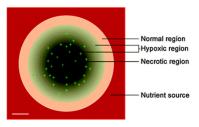


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Regarding modeling of diffuse interface tumor growth we can quote, e.g.,

 Ciarletta, Cristini, Frieboes, Garcke, Hawkins-Daarud, Hilhorst, Lam, Lowengrub, Oden, van der Zee, Wise, also for their numerical simulations → complex changes in tumor morphologies due to the interactions with nutrients or toxic agents and also due to mechanical stresses

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- Frieboes, Jin, Chuang, Wise, Lowengrub, Cristini, Garcke, Lam, Nürnberg, Sitka, for the interaction of multiple tumor cell species described by *multiphase mixture models*

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HZO: the free energy

- $u = \text{tumor cell volume fraction } u \in [0, 1]$
- ullet n= nutrient-rich extracellular water volume fraction $n\in [0,1]$
- $f(u) = \Gamma u^2 (1-u)^2$: a double well
- $\chi(u,n) = -\chi_0 un$: chemotaxis driving the tumor cells toward the oxygen supply

$$E = \int_{\Omega} \left(f(u) + \frac{\epsilon^2}{2} |\nabla u|^2 + \chi(u, n) + \frac{1}{2\delta} n^2 \right) dx. \tag{4}$$

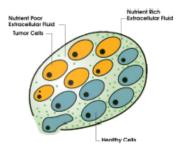


Figure 1. Four-species model: illustration of the four-species mixture. The tumor and healthy cell populations are assumed to have a thin diffuse interface, whereas the nutrient-rich and nutrient-poor extracellular water are segregated by a wide smooth interface.

Diffuse interface models of tumor growth

The plot of the summand $f(u) + \chi(u, n)$

The lowest energy state is when u = 1 and n = 1, when there is a full interaction between the tumor species and the nutrient-rich extracellular water.

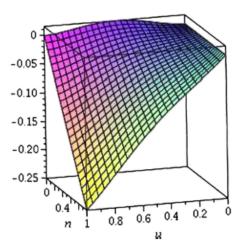


Figure 2. Graph of homogeneous free energy: $f(u) + \chi(u,n)$. ($\Gamma = \chi_0 = 0.25$).

$$u_{t} = \nabla \cdot (M_{u} \nabla \mu_{u}) + \gamma_{u}, \quad \mu_{u} = \partial_{u} E = f'(u) + \partial_{u} \chi(u, n) - \epsilon \Delta u$$

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$$\gamma_u = P(u)(\mu_n - \mu_u), \quad \gamma_n = -\gamma_u, \quad \text{where}$$

$$P(u) = \begin{cases} \delta P_0 u & \text{if } u \geq 0 \\ 0 & \text{elsewhere} \end{cases}$$

being δ a small positive constant and $P_0 \geq 0$.

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Simulations by HZO: the tumor starts growing increasingly more ellipsoidal at first and eventually begins forming buds growing toward the higher levels of nutrient

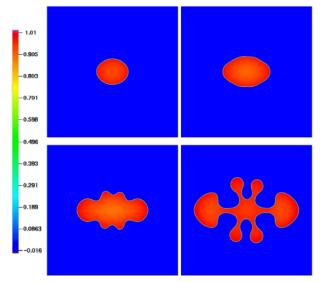


Figure 7. Example simulation: snapshots are shown at t=20,40,60, and 80 of a simulation with $\Gamma=0.045, \epsilon=0.005, \chi_0=0.05, \delta=0.01, P_0=0.1, \hat{M}=200, \text{ and } \hat{D}=1.$

Simulations by HZO: the influence of χ_0 and δ

- When the ratio χ_0/Γ is small, the tumor remains circular $u\sim 0,1$
- When $\chi_0 \sim \Gamma$ the tumor goes into an ellipse
- When χ_0/Γ and χ_0/ϵ are big, u no longer takes on values close to 0 and 1: it begins moving quickly toward the regions with higher nutrients
- Only when χ_0 is large the value of δ makes a difference in simulations

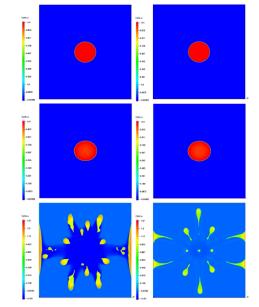
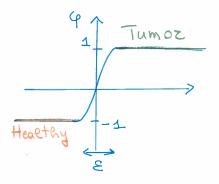


Figure 10. Effects of parameter χ_0 : illustrated here are the effects of different values of χ_0 when $\Gamma=0.045$ and $\epsilon=0.005$ are held constant. In the first row, $\chi_0=0.005$; in the second row, $\chi_0=0.05$; and in the third row, $\chi_0=0.5$. In the first column, $\delta=0.1$; and in the second column, $\delta=0.01$.

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Our notation for the tumor phase parameter $(u=)\phi \in [-1,1]$



The sharp interface S replaced by a (threekness E) thin transition layer $\varphi = -1$ in the Healthy tissue phase $\varphi = \pm 1$ in the Tumoz phase

 In terms of the theoretical analysis most of the recent literature is restricted to the two-phase variant, i.e., to models that only account for the evolution of a tumor surrounded by healthy tissue.

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 control and long-time behavior of solution, have been established in a number of
 papers of a number of authors which include: Agosti, Ciarletta, Colli, Frigeri,
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 - for tumor growth models based on the coupling of Cahn-Hilliard (for the tumor density) and reaction-diffusion (for the nutrient) equations, and
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In this talk we concentrate on two recent results on optimal control and long-time behavior of solution. Long-time dynamics and optimal control

Long-time dynamics and optimal control

- The state system consists of a Cahn-Hilliard type equation for the tumor cell fraction and a reaction-diffusion equation for the nutrient
- The possible medication that serves to eliminate tumor cells is in terms of drugs and is introduced into the system through the nutrient
- In this setting, the control variable acts as an external source in the nutrient equation
- 1 First, we consider the problem of "long-time treatment" under a suitable given source and prove the convergence of any global solution to a single equilibrium as $t \to +\infty$.
- 2 Then we consider the "finite-time treatment" of tumor, which corresponds to an optimal control problem. Here we also allow the objective cost functional to depend on a free time variable, which represents the unknown treatment time to be optimized. We prove the existence of an optimal control and obtain first order necessary optimality conditions for both the drug concentration and the treatment time.

The main modelling idea

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By establishing the Lyapunov stability of certain equilibria of the state system (without external source), we see that ϕ_Ω can be taken as a stable configuration, so that the tumor will not grow again once the finite-time treatment is completed

The state system: Cahn–Hilliard + nutrient model with source terms

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The PDE system is an approximation of the model proposed in [HZO: A. Hawkins-Daarud, K.-G. van der Zee and J.-T. Oden (2011)] in $Q:=\Omega\times(0,T)$:

$$\phi_t - \Delta \mu = P(\phi)(\sigma - \mu), \qquad \mu = -\Delta \phi + F'(\phi)$$
$$\sigma_t - \Delta \sigma = -P(\phi)(\sigma - \mu) + \mathbf{u}$$

subject to initial and boundary conditions

$$\phi|_{t=0}=\phi_0, \quad \sigma|_{t=0}=\sigma_0, \quad \text{in } \Omega\,, \quad \partial_\nu\phi=\partial_\nu\mu=\partial_\nu\sigma=0, \quad \text{on } \partial\Omega\times(0,T)$$

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- The state variables are:
 - ▶ the tumor cell fraction ϕ : $\phi \simeq 1$ (tumorous phase), $\phi \simeq -1$ (healthy tissue phase)
 - **b** the nutrient concentration σ : $\sigma \simeq 1$ and $\sigma \simeq 0$ indicate a nutrient-rich or nutrient-poor extracellular water phase
- F is typically a double-well potential with equal minima at $\phi=\pm 1$
- $P \ge 0$ denotes a suitable regular proliferation function
- The choice of reactive terms is motivated by the linear phenomenological constitutive laws for chemical reactions
- The control variable u serves as an external source in the equation for σ and can be interpreted as a medication 4□ > 4回 > 4 重 > 4 重 > 重 のQで

Energy identity

The system turns out to be thermodynamically consistent. In particular, when u=0 the unknown pair (ϕ,σ) is a dissipative gradient flow for the total free energy:

$$\mathcal{E}(\phi,\sigma) = \int_{\Omega} \left[\frac{1}{2} |\nabla \phi|^2 + F(\phi) \right] dx + \frac{1}{2} \int_{\Omega} \sigma^2 dx.$$

Moreover generally, under the presence of the external source u, we observe that any smooth solution (ϕ, σ) to the problem satisfies the following energy identity:

$$\frac{\mathrm{d}}{\mathrm{d}t}\mathcal{E}(\phi,\sigma) + \int_{\Omega} \left[|\nabla \mu|^2 + |\nabla \sigma|^2 + P(\phi)(\mu - \sigma)^2 \right] \mathrm{d}x = \int_{\Omega} u\sigma \,\mathrm{d}x,$$

which motives the twofold aim of the present contribution.

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- 2. Denoting by $T\in(0,+\infty)$ a fixed maximal time in which the patient is allowed to undergo a medical treatment, we derive necessary optimality conditions for
 - (CP) Minimize the cost functional

$$\mathcal{J}(\phi, \sigma, u, \tau) = \frac{\beta_Q}{2} \int_0^{\tau} \int_{\Omega} |\phi - \phi_Q|^2 dx dt + \frac{\beta_{\Omega}}{2} \int_{\Omega} |\phi(\tau) - \phi_{\Omega}|^2 dx + \frac{\alpha_Q}{2} \int_0^{\tau} \int_{\Omega} |\sigma - \sigma_Q|^2 dx dt + \frac{\beta_S}{2} \int_{\Omega} (1 + \phi(\tau)) dx + \frac{\beta_u}{2} \int_0^{\tau} \int_{\Omega} |u|^2 dx dt + \beta_T \tau$$

subject to the state system and the the control constraint

$$u \in \mathcal{U}_{\mathrm{ad}} := \{u \in L^{\infty}(Q): u_{\min} \leq u \leq u_{\max} \text{ a.e. in } Q\}, \quad \tau \in (0,T)$$

Comments on the cost functional

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- $\tau \in (0, T]$ represents the treatment time of one cycle, i.e., the amount of time the drug is applied to the patient before the period of rest, or the treatment time before surgery, ϕ_Q and σ_Q represent a desired evolution for the tumor cells and for the nutrient, ϕ_Ω stands for desired final distribution of tumor cells
- ullet The first three terms of ${\mathcal J}$ are of standard tracking type and the fourth term of ${\mathcal J}$ measures the size of the tumor at the end of the treatment
- ullet The fifth term penalizes large concentrations of the cytotoxic drugs, and the sixth term of ${\mathcal J}$ penalizes long treatment times



The choice of ϕ_{Ω}

After the treatment, the ideal situation will be either the tumor is ready for surgery or the tumor will be stable for all time without further medication (i.e., u=0). This goal can be realized by making different choices of the target function ϕ_{Ω} in the above optimal control problem (CP).

- For the former case, one can simply take ϕ_{Ω} to be a configuration that is suitable for surgery.
- While for the later case, which is of more interest to us, we want to choose ϕ_Ω as a "stable" configuration of the system, so that the tumor does not grow again once the treatment is complete.

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- While for the later case, which is of more interest to us, we want to choose ϕ_{Ω} as a "stable" configuration of the system, so that the tumor does not grow again once the treatment is complete.

For this purpose, we prove that any local minimizer of the total free energy $\mathcal E$ is Lyapunov stable provided that u=0. As a consequence, these local energy minimizers serve as possible candidates for the target function ϕ_Ω . Then after completing a successful medication, the tumor will remain close to the chosen stable configuration for all time.

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However, for our coupled system this property no longer holds, which brings us new difficulties in analysis.

• Besides, quite different from the Cahn-Hilliard-Oono system (cf. Miranville's lesson) in which the mass $\int_{\Omega} \phi(t) \, \mathrm{d}x$ is not preserved due to possible reactions, here in our case it is not obvious how to control the mass changing rate:

$$\frac{\mathrm{d}}{\mathrm{d}t} \int_{\Omega} \phi \, \mathrm{d}x = \int_{\Omega} P(\phi)(\sigma - \mu) \, \mathrm{d}x.$$

Similar problem happens to the nutrient as well, that is

$$\frac{\mathrm{d}}{\mathrm{d}t}\int_{\Omega}\sigma\,\mathrm{d}x=-\int_{\Omega}P(\phi)(\sigma-\mu)\,\mathrm{d}x+\int_{\Omega}u\,\mathrm{d}x.$$

 The observation that the total mass can be determined by the initial data and the external source:

$$\int_{\Omega} (\phi(t) + \sigma(t)) \, dx = \int_{\Omega} (\phi_0 + \sigma_0) \, dx + \int_0^t \int_{\Omega} u \, dx \, d\tau, \quad \forall t \ge 0$$

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- Based on the above mentioned special structure of the system, by introducing a new version of Łojasiewicz-Simon inequality we are able to prove that every global weak solution (ϕ,σ) of the problem will converge to a certain single equilibrium $(\phi_{\infty},\sigma_{\infty})$ as $t\to +\infty$ and, moreover, we obtain a polynomial decay of the solution.

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- Based on the above mentioned special structure of the system, by introducing a new version of Łojasiewicz-Simon inequality we are able to prove that every global weak solution (ϕ, σ) of the problem will converge to a certain single equilibrium $(\phi_{\infty}, \sigma_{\infty})$ as $t \to +\infty$ and, moreover, we obtain a polynomial decay of the solution.
- Besides, a nontrivial application of the Łojasiewicz-Simon approach further leads to the Lyapunov stability of local minimizers of the free energy \mathcal{E} (we only consider the case u = 0 for the sake of simplicity).

Outline

- Phase field models for tumor growth
- 2 The model HZO by [A. Hawkins-Daarud, K.-G. van der Zee and J.-T. Oden (2011)]
- 3 Joint work with C. Cavaterra and H. Wu, AMO (online)
- Well-posedness
- 5 Long-term dynamics
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- Perspectives and Open problems

Let $\phi_0 \in H^2_N(\Omega) \cap H^3(\Omega)$ and $\sigma_0 \in H^1(\Omega)$ and assume that

- (P1) $P \in C^2(\mathbb{R})$ is nonnegative. There exist $\alpha_1 > 0$ and some $q \in [1,4]$ such that, for all $s \in \mathbb{R}$, $|P'(s)| \le \alpha_1(1+|s|^{q-1})$
- (F1) $F = F_0 + F_1$, with $F_0, F_1 \in C^5(\mathbb{R})$. There exist $\alpha_i > 0$ and $r \in [2, 6)$ such that $|F_1''(s)| \le \alpha_2$, $\alpha_3(1+|s|^{r-2}) \le F_0''(s) \le \alpha_4(1+|s|^{r-2})$, $F(s) \ge \alpha_5|s| \alpha_6 \quad \forall s \in \mathbb{R}$ (U1) For any T > 0, $u \in L^2(0, T; L^2(\Omega))$.

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Theorem (Strong solutions)

(1) For every T > 0, the state system admits a unique strong solution:

$$\begin{split} \|\phi\|_{L^{\infty}(0,T;H^{3}(\Omega))\cap L^{2}(0,T;H^{4}(\Omega))\cap H^{1}(0,T;H^{1}(\Omega))} + \|\mu\|_{L^{\infty}(0,T;H^{1}(\Omega))\cap L^{2}(0,T;H^{2}(\Omega))} \\ + \|\sigma\|_{C([0,T];H^{1}(\Omega))\cap L^{2}(0,T;H^{2}_{N}(\Omega))\cap H^{1}(0,T;L^{2}(\Omega))} \leq K_{1}. \end{split}$$

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(2) Let (ϕ_i, σ_i) be two strong solutions. Then there exists a constant $K_2 > 0$, depending on $\|u_i\|_{L^2(0,T;L^2)}$, Ω , T, $\|\phi_0\|_{H^3}$ and $\|\sigma_0\|_{H^1}$, such that

$$\begin{aligned} \|\phi_{1} - \phi_{2}\|_{L^{\infty}(0,T;H^{1}) \cap L^{2}(0,T;H^{3}) \cap H^{1}(0,T;(H^{1})')} + \|\mu_{1} - \mu_{2}\|_{L^{2}(0,T;H^{1})} \\ + \|\sigma_{1} - \sigma_{2}\|_{C([0,T];H^{1}) \cap L^{2}(0,T;H^{2}) \cap H^{1}(0,T;L^{2})} \leq K_{2}\|u_{1} - u_{2}\|_{L^{2}(0,T;L^{2})}. \end{aligned}$$

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Long-term dynamics

We make the following additional assumptions:

- (P2) P(s) > 0, for all $s \in \mathbb{R}$
- (F2) F(s) is real analytic, for all $s \in \mathbb{R}$
- (U2) $u\in L^1(0,+\infty;L^2(\Omega))\cap L^2(0,+\infty;L^2(\Omega))$ and satisfies the decay condition $\sup_{t\geq 0}(1+t)^{3+\rho}\|u(t)\|_{L^2(\Omega)}<+\infty,\quad \text{for some }\rho>0.$

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Theorem (1. The stationary problem)

For any $\phi_0 \in H^1(\Omega)$, $\sigma_0 \in L^2(\Omega)$, the state system admits a unique global weak solution (ϕ, μ, σ) : $\lim_{t \to +\infty} (\|\phi(t) - \phi_{\infty}\|_{H^{2}(\Omega)} + \|\sigma(t) - \sigma_{\infty}\|_{L^{2}(\Omega)} + \|\mu(t) - \mu_{\infty}\|_{L^{2}(\Omega)}) = 0$, where $(\phi_{\infty}, \mu_{\infty}, \sigma_{\infty})$ satisfies the stationary problem

$$\left\{ \begin{array}{ll} -\Delta\phi_{\infty}+F'(\phi_{\infty})=\mu_{\infty}, & \text{in }\Omega\\ \\ \partial_{\nu}\phi_{\infty}=0, & \text{on }\partial\Omega\\ \\ \int_{\Omega}(\phi_{\infty}+\sigma_{\infty})\,dx=\int_{\Omega}(\phi_{0}+\sigma_{0})\,dx+\int_{0}^{+\infty}\!\!\int_{\Omega}u\,dx\,dt \end{array} \right.$$

with μ_{∞} and σ_{∞} being two constants given by $\sigma_{\infty} = \mu_{\infty} = |\Omega|^{-1} \int_{\Omega} F'(\phi_{\infty}) dx$.

The convergence rate

Theorem (2. Convergence rate)

Moreover, under the same assumptions, the following estimates on convergence rate hold

$$\begin{split} &\|\phi(t) - \phi_{\infty}\|_{H^{1}(\Omega)} + \|\sigma(t) - \sigma_{\infty}\|_{L^{2}(\Omega)} \leq C(1+t)^{-\min\left\{\frac{\theta}{1-2\theta}, \frac{\rho}{2}\right\}}, \quad \forall \, t \geq 0, \\ &\|\mu(t) - \mu_{\infty}\|_{L^{2}(\Omega)} \leq C(1+t)^{-\frac{1}{2}\min\left\{\frac{\theta}{1-2\theta}, \frac{\rho}{2}\right\}}, \quad \forall \, t \geq 0, \end{split}$$

$$\|\mu(t) - \mu_{\infty}\|_{L^{2}(\Omega)} \leq C(1+t)^{-\frac{\epsilon}{2}\min\{\frac{\epsilon}{1-2\theta},\frac{\epsilon}{2}\}}, \quad \forall t \geq 0,$$

where C>0 is a constant depending on $\|\phi_0\|_{H^1(\Omega)}$, $\|\sigma_0\|_{L^2(\Omega)}$, $\|\phi_\infty\|_{H^1(\Omega)}$,

 $\|u\|_{L^1(0,+\infty;L^2(\Omega))}$, $\|u\|_{L^2(0,+\infty;L^2(\Omega))}$ and Ω ; $\theta\in(0,\frac{1}{2})$ is a constant depending on ϕ_{∞} .

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- ullet Then we give a characterization on the ω -limit

$$\omega(\phi_0,\sigma_0) = \{(\phi_\infty,\sigma_\infty) \in (H^2_N(\Omega) \cap H^3(\Omega)) \times H^1(\Omega) : \exists \{t_n\} \nearrow +\infty \text{ such that}$$
$$(\phi(t_n),\sigma(t_n)) \to (\phi_\infty,\sigma_\infty) \text{ in } H^2(\Omega) \times L^2(\Omega)\}.$$

And we have the following result

Theorem (3. The ω -limit)

Assume (P1), (F1), (U2). For any initial datum $(\phi_0, \sigma_0) \in H^1(\Omega) \times L^2(\Omega)$, the associated ω -limit set $\omega(\phi_0, \sigma_0)$ is non-empty. For any element $(\phi_\infty, \sigma_\infty) \in \omega(\phi_0, \sigma_0)$, σ_∞ is a constant and $(\phi_\infty, \sigma_\infty)$ satisfies the stationary problem. Besides, μ_∞ is a constant given by $|\Omega|^{-1} \int_{\Omega} F'(\phi_\infty) dx$ and the following relation holds

$$P(\phi_{\infty})(\sigma_{\infty} - \mu_{\infty}) = 0$$
, a.e. in Ω .

And the positivity of P entails immediately also $\sigma_{\infty} = \mu_{\infty}$.



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• Finally, we prove the convergence of the trajectories and polynomial decay by means of a proper Łojasiewicz–Simon inequality: Given any initial datum $(\phi_0, \sigma_0) \in H^1(\Omega) \times L^2(\Omega)$ and source term u satisfying (U2), we denote by

$$m_{\infty} := |\Omega|^{-1} \left(\int_{\Omega} (\phi_0 + \sigma_0) \, \mathrm{d}x + \int_0^{+\infty} \int_{\Omega} u \, \mathrm{d}x \, \mathrm{d}t \right)$$

the total mass at infinity time. Then we are able to derive the following

Theorem (Łojasiewicz-Simon Inequality)

Let (F1), (F2), (P1), (P2) and (U2) be satisfied. Suppose that $(\phi_{\infty}, \mu_{\infty}, \sigma_{\infty})$ is a solution to the elliptic stationary problem. Then there exist constants $\theta \in (0, \frac{1}{2})$ and $\beta > 0$, depending on ϕ_{∞} , m_{∞} and Ω , such that for any $(\phi, \sigma) \in H^2_N(\Omega) \times H^1(\Omega)$ satisfying

$$\|\phi - \phi_{\infty}\|_{H^{1}(\Omega)} < \beta,$$

$$\int_{\Omega} (\phi + \sigma) dx + m_{u} |\Omega| = \int_{\Omega} (\phi_{\infty} + \sigma_{\infty}) dx = m_{\infty} |\Omega|,$$

where m_u is a certain constant fulfilling $|m_u| \leq |\Omega|^{-\frac{1}{2}} ||u||_{L^1(0,+\infty;L^2(\Omega))}$, then we have

$$\|\mu - \overline{\mu}\|_{(H^1(\Omega))'} + C\|\nabla\sigma\|_{L^2(\Omega)} + C\|\sqrt{P(\phi)}(\mu - \sigma)\|_{L^2(\Omega)} + C|m_u|^{\frac{1}{2}}$$

$$\geq |\mathcal{E}(\phi, \sigma) - \mathcal{E}(\phi_{\infty}, \sigma_{\infty})|^{1-\theta}, \quad \text{where}$$

 $\mu = -\Delta \phi + F'(\phi) \text{ and } C > 0 \text{ depends on } \Omega, \ \phi_{\infty}, \ m_{\infty}, \ \|\phi\|_{H^2(\Omega)}, \ \|\sigma\|_{H^1(\Omega)}, \ \|u\|_{L^1(0,+\infty;L^2(\Omega))}.$

Energy minimizers with u = 0

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Let us now assume u = 0. Then it follows that the total mass of the system is now conserved:

$$\int_{\Omega} (\phi(t) + \sigma(t)) \, \mathrm{d}x = \int_{\Omega} (\phi_0 + \sigma_0) \, \mathrm{d}x, \quad \forall \, t \geq 0.$$

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Let $m \in \mathbb{R}$ be an arbitrary given constant. Set

$$\mathcal{Z}_m = \left\{ (\phi, \sigma) \in H^1(\Omega) \times L^2(\Omega) : \int_{\Omega} (\phi + \sigma) \, \mathrm{d}x = |\Omega| m \right\}.$$

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Any $(\phi^*, \sigma^*) \in \mathcal{Z}_m$ is called

• a local energy minimizer of the total energy

$$\mathcal{E}(\phi, \sigma) = \int_{\Omega} \left[\frac{1}{2} |\nabla \phi|^2 + F(\phi) \right] dx + \frac{1}{2} \int_{\Omega} \sigma^2 dx$$

if there exists a constant $\chi > 0$ such that $\mathcal{E}(\phi^*, \sigma^*) \leq \mathcal{E}(\phi, \sigma)$, for all $(\phi, \sigma) \in \mathcal{Z}_m$ satisfying $\|(\phi - \phi^*, \sigma - \sigma^*)\|_{H^1(\Omega) \times L^2(\Omega)} < \chi$

• If $\chi = +\infty$, then (ϕ^*, σ^*) is called a *global energy minimizer* of $\mathcal{E}(\phi, \sigma)$ in \mathcal{Z}_m .

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We first derive some properties for the critical points of $\mathcal{E}(\phi,\sigma)$ in \mathcal{Z}_m .

$$\begin{cases} -\Delta \phi + F'(\phi) = \mu, & \text{in } \Omega, \\ \partial_{\nu} \phi = 0, & \text{on } \partial \Omega, \\ \int_{\Omega} (\phi + \sigma) \, \mathrm{d}x = |\Omega| m, \end{cases}$$

where μ and σ are constants given by $\sigma = \mu = |\Omega|^{-1} \int_{\Omega} F'(\phi) dx$.

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Theorem (4. Critical points)

Let assumption (F1) be satisfied. Then we have:

$$\left\{ \begin{array}{ll} -\Delta\phi+F'(\phi)=\mu, & \text{ in } \Omega, \\ \partial_{\nu}\phi=0, & \text{ on } \partial\Omega, \\ \int_{\Omega}(\phi+\sigma)\,\mathrm{d}x=|\Omega|m, \end{array} \right.$$

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Theorem (4. Critical points)

Let assumption (F1) be satisfied. Then we have:

(1) If $(\phi^*, \sigma^*) \in H^2_N(\Omega) \times \mathbb{R}$ is a strong solution to the stationary problem above, then (ϕ^*, σ^*) is a critical point of $\mathcal{E}(\phi, \sigma)$ in \mathcal{Z}_m . Conversely, if (ϕ^*, σ^*) is a critical point of $\mathcal{E}(\phi, \sigma)$ in \mathcal{Z}_m , then $\phi^* \in H^2_N(\Omega)$, $\sigma^* \in \mathbb{R}$ satisfy the stationary problem above

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- (2) If (ϕ^*, σ^*) is a local energy minimizer of $\mathcal{E}(\phi, \sigma)$ in \mathcal{Z}_m , then (ϕ^*, σ^*) is a critical point of $\mathcal{E}(\phi, \sigma)$.
- (3) The functional $\mathcal{E}(\phi, \sigma)$ has at least one minimizer $(\phi^*, \sigma^*) \in \mathcal{Z}_m$ such that

$$\mathcal{E}(\phi^*, \sigma^*) = \inf_{(\phi, \sigma) \in \mathcal{Z}_m} \mathcal{E}(\phi, \sigma)$$

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Theorem (5. Lyapunov stability)

Assume that (F1), (F2), (P1), (P2) are satisfied and u = 0. Given $m \in \mathbb{R}$, let (ϕ^*, σ^*) be a local energy minimizer in \mathcal{Z}_m of

$$\mathcal{E}(\phi,\sigma) = \int_{\Omega} \left[\frac{1}{2} |\nabla \phi|^2 + F(\phi) \right] dx + \frac{1}{2} \int_{\Omega} \sigma^2 dx.$$

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Then, for any $\epsilon > 0$, there exists a constant $\eta \in (0,1)$ such that for arbitrary initial datum $(\phi_0,\sigma_0) \in (H^2_N(\Omega)\cap H^3(\Omega)) \times H^1(\Omega))$ satisfying $\int_{\Omega} (\phi_0+\sigma_0)\,\mathrm{d}x = |\Omega|m$ and $\|\phi_0-\phi^*\|_{H^1(\Omega)}+\|\sigma_0-\sigma^*\|_{L^2(\Omega)} \leq \eta$, the state system admits a unique global strong solution (ϕ,σ) such that

$$\|\phi(t)-\phi^*\|_{H^1(\Omega)}+\|\sigma(t)-\sigma^*\|_{L^2(\Omega)}\leq \epsilon, \quad \forall \ t\geq 0.$$

Namely, any local energy minimizer of $\mathcal{E}(\phi, \sigma)$ in \mathcal{Z}_m is locally Lyapunov stable.

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Conclusions on long-term dynamics

- The result on long-time behavior derived in Theorem 1 and 2 can be applied to the global strong solution obtained in Theorem 5
- Although it is still not obvious to identify the asymptotic limit $(\phi_{\infty}, \sigma_{\infty})$, we are able to conclude that $(\phi_{\infty}, \sigma_{\infty})$ also satisfies

$$\|\phi_{\infty} - \phi^*\|_{H^1(\Omega)} + \|\sigma_{\infty} - \sigma^*\|_{L^2(\Omega)} \le \epsilon$$

• In particular, if (ϕ^*, σ^*) is an isolated local energy minimizer then it is locally asymptotic stable

Outline

- Phase field models for tumor growth
- The model HZO by [A. Hawkins-Daarud, K.-G. van der Zee and J.-T. Oden (2011)]
- 3 Joint work with C. Cavaterra and H. Wu, AMO (online)
- Well-posedness
- 5 Long-term dynamics
- 6 The optimal control problem
- Joint work with A. Miranville and G. Schimperna, JDE (online)
- Well-posedness
- Dissipativity and existence of the attractor
- Perspectives and Open problems



Assumptions for the optimal control problem

Hence we can take the target $\phi_{\Omega} = \phi_{\infty}$ in the following:

(CP) Minimize the cost functional

$$\mathcal{J}(\phi, \sigma, u, \tau) = \frac{\beta_Q}{2} \int_0^{\tau} \int_{\Omega} |\phi - \phi_Q|^2 dx dt + \frac{\beta_{\Omega}}{2} \int_{\Omega} |\phi(\tau) - \phi_{\Omega}|^2 dx + \frac{\alpha_Q}{2} \int_0^{\tau} \int_{\Omega} |\sigma - \sigma_Q|^2 dx dt + \frac{\beta_S}{2} \int_{\Omega} (1 + \phi(\tau)) dx + \frac{\beta_u}{2} \int_0^{\tau} \int_{\Omega} |u|^2 dx dt + \beta_T \tau$$

subject to the state system and the the control constraint

$$u \in \mathcal{U}_{ad} := \{ u \in L^{\infty}(Q) : u_{\min} \le u \le u_{\max} \text{ a.e. in } Q \}, \quad \tau \in (0, T),$$

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where $T \in (0, +\infty)$ is a fixed maximal time. We assume:

- (C1) β_Q , β_Ω , β_S , β_u , β_T , α_Q are nonnegative constants but not all zero.
- (C2) $\phi_Q, \, \sigma_Q \in L^2(Q), \, \phi_\Omega, \, \sigma_\Omega \in L^2(\Omega), \, u_{\min}, \, u_{\max} \in L^\infty(Q), \, \text{and} \, u_{\min} \leq u_{\max}, \, \text{a.e. in } Q.$
- (C3) Let \mathcal{U}_R be an open set in $L^2(Q)$: $\mathcal{U}_{\mathrm{ad}} \subset \mathcal{U}_R$ and $\|u\|_{L^2(Q)} \leq R$, for all $u \in \mathcal{U}_R$.

From the well-posedness results it follows that the $\emph{control-to-state operator } \mathcal{S}$

$$u \mapsto \mathcal{S}(u) := (\phi, \mu, \sigma)$$

is well-defined and Lipschitz continuous as a mapping from $\mathcal{U}_R\subset L^2(Q)$ into the following space

$$(L^{\infty}(0,T;(H^{1}(\Omega))')\cap L^{2}(0,T;H^{1}(\Omega)))\times L^{2}(0,T;(H^{1}(\Omega))')\times (L^{\infty}(0,T;(H^{1}(\Omega))')\cap L^{2}(Q)).$$

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The triplet (ϕ, μ, σ) is the unique weak solution to the state system with data (ϕ_0, σ_0, u) over the time interval [0, T]. For convenience, we use the notations $\phi = \mathcal{S}_1(u)$ and $\sigma = \mathcal{S}_3(u)$ for the first and third component of $\mathcal{S}(u)$.

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Theorem (Existence of the optimal control)

Assume that (P1), (F1), (U1) and (C1)–(C3) are satisfied. Let $\phi_0 \in H^2_N(\Omega) \cap H^3(\Omega)$ and $\sigma_0 \in H^1(\Omega)$. Then there exists at least one minimizer $(\phi_*, \sigma_*, u_*, \tau_*)$ to problem (CP). Namely, $\phi_* = \mathcal{S}_1(u_*)$, $\sigma_* = \mathcal{S}_3(u_*)$ satisfy

$$\mathcal{J}(\phi_*, \sigma_*, u_*, \tau_*) = \inf_{\substack{(w,s) \in \mathcal{U}_{\text{ad}} \times [0, T] \\ \text{s.t. } \phi = \mathcal{S}_1(w), \, \sigma = \mathcal{S}_3(w)}} \mathcal{J}(\phi, \sigma, w, s).$$

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$$\begin{split} \partial_t \xi - \Delta \eta &= P'(\phi_*)(\sigma_* - \mu_*) \, \xi + P(\phi_*)(\rho - \eta), \qquad \eta = -\Delta \xi + F''(\phi_*) \, \xi, \\ \partial_t \rho - \Delta \rho &= -P'(\phi_*)(\sigma_* - \mu_*) \, \xi - P(\phi_*)(\rho - \eta) + h \\ \partial_n \xi &= \partial_n \eta = \partial_n \rho = 0, \qquad \xi(0) = \rho(0) = 0. \end{split}$$

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We can apply [Theorems 3.1, 3.2, CGRS] for the well-posedness of the linearized system and the Fréchet differentiability of the control-to-state operator $\mathcal S$ with respect to u.

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$$\begin{split} \mathcal{Y} := & \left(H^1(0, T; (H_N^2(\Omega))') \cap L^{\infty}(0, T; L^2(\Omega)) \cap L^2(0, T; H_N^2(\Omega)) \right) \times L^2(Q) \\ & \times \left(H^1(0, T; L^2(\Omega)) \cap L^2(0, T; H^2(\Omega)) \right). \end{split}$$

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$$\times \left(H^1(0, T; L^2(\Omega)) \cap L^2(0, T; H^2(\Omega)) \right).$$

For any $u_* \in \mathcal{U}_R$, the Fréchet derivative $DS(u_*) \in \mathcal{L}(L^2(Q), \mathcal{Y})$ is defined as follows: for any $h \in L^2(Q)$, $DS(u_*)h = (\xi^h, \eta^h, \rho^h)$, where (ξ^h, η^h, ρ^h) is the unique solution to the linearized system associated with h.

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First order optimality conditions

Define a reduced functional

$$\widetilde{\mathcal{J}}(u,\tau) := \mathcal{J}(S_1(u), S_3(u), u, \tau).$$

Since the control-to-state mapping \mathcal{S} is also Fréchet differentiable into $C^0([0,T];L^2(\Omega))$ with respect to u, then the reduced cost functional $\widetilde{\mathcal{J}}$ is Fréchet differentiable in \mathcal{U}_R .

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Theorem (Existence of solutions to the adjoint system)

Assume (P1), (F1), (U1), (C1)–(C3), $\phi_0 \in H^2_N(\Omega) \cap H^3(\Omega)$, and $\sigma_0 \in H^1(\Omega)$. Then the adjoint system

$$-\partial_t p + \Delta q - F''(\phi_*) q + P'(\phi_*)(\sigma_* - \mu_*)(r - p) = \beta_Q (\phi_* - \phi_Q)$$

$$q - \Delta p + P(\phi_*)(p - r) = 0, \qquad -\partial_t r - \Delta r + P(\phi_*)(r - p) = \alpha_Q(\sigma_* - \sigma_Q)$$

$$\partial_n p = \partial_n q = \partial_n r = 0, \qquad r(\tau_*) = 0, \quad p(\tau_*) = \beta_\Omega (\phi_*(\tau_*) - \phi_\Omega) + \frac{\beta_S}{2}$$

has a unique weak solution (p, q, r) on [0, T]:

$$p \in H^{1}(0, T; (H_{N}^{2}(\Omega))') \cap C^{0}([0, T]; L^{2}(\Omega)) \cap L^{2}(0, T; H_{N}^{2}(\Omega)),$$

$$q \in L^{2}(Q), \qquad r \in H^{1}(0, T; L^{2}(\Omega)) \cap C^{0}([0, T]; H^{1}(\Omega)) \cap L^{2}(0, T; H_{N}^{2}(\Omega)).$$

Necessary optimality conditions

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Theorem (Necessary optimality conditions)

Let $(u_*, \tau_*) \in \mathcal{U}_{ad} \times [0, T]$ denote a minimizer to the optimal control problem (CP) with corresponding state variables $(\phi_*, \mu_*, \sigma_*) = S(u_*)$ and associated adjoint variables (p,q,r), then it holds:

$$\beta_u \int_0^T \! \int_\Omega u_*(u-u_*) \, \mathrm{d}x \, \mathrm{d}t + \int_0^{\tau_*} \! \int_\Omega r(u-u_*) \, \mathrm{d}x \, \mathrm{d}t \geq 0, \quad \forall \, u \in \mathcal{U}_{\mathrm{ad}}.$$

Besides, setting

$$\mathcal{L}(\phi_*, \sigma_*, \tau_*) = \frac{\beta_Q}{2} \int_{\Omega} |\phi_*(\tau_*) - \phi_Q(\tau_*)|^2 dx + \beta_\Omega \int_{\Omega} (\phi_*(\tau_*) - \phi_\Omega) \, \partial_t \phi_*(\tau_*) \, dx$$
$$+ \frac{\alpha_Q}{2} \int_{\Omega} |\sigma_*(\tau_*) - \sigma_Q|^2 \, dx + \frac{\beta_S}{2} \int_{\Omega} \partial_t \phi_*(\tau_*) \, dx + \beta_T$$

we have

$$\mathcal{L}(\phi_*, \sigma_*, au_*) egin{array}{ll} \geq 0, & ext{if } au_* = 0, \ = 0, & ext{if } au_* \in (0, T), \ \leq 0, & ext{if } au_* = T. \end{array}$$

Interpretation of the first condition

Besides, if we extend r by zero to $(\tau_*, T]$, then we can express the variational inequality

$$\beta_u \int_0^T \int_{\Omega} u_*(u-u_*) \, \mathrm{d}x \, \mathrm{d}t + \int_0^{\tau_*} \int_{\Omega} r(u-u_*) \, \mathrm{d}x \, \mathrm{d}t \ge 0, \quad \forall \, u \in \mathcal{U}_{\mathrm{ad}}.$$

as

$$\int_0^T \int_{\Omega} (\beta_u u_* + r)(u - u_*) \, \mathrm{d}x \, \mathrm{d}t \ge 0, \quad \forall \, u \in \mathcal{U}_{\mathrm{ad}},$$

which allows the interpretation that the optimal control u_* is the $L^2(Q)$ -projection of $-\beta_u^{-1}r$ onto the set $\mathcal{U}_{\mathrm{ad}}$ (provided that $\beta_u>0$).

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We consider here the long time dynamics for the following model for tumor growth:

$$\varphi_{t} - \Delta \mu = (\mathcal{P}\sigma - \mathcal{A})h(\varphi),$$

$$\mu = -\Delta \varphi + \Psi'(\varphi),$$

$$\sigma_{t} - \Delta \sigma = -\mathcal{C}\sigma h(\varphi) + B(\sigma_{s} - \sigma),$$

settled in $\Omega \times (0, +\infty)$, and complemented with the Cauchy conditions and with no-flux (i.e., homogeneous Neumann) boundary conditions for all unknowns.

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settled in $\Omega \times (0, +\infty)$, and complemented with the Cauchy conditions and with no-flux (i.e., homogeneous Neumann) boundary conditions for all unknowns.

- Here h(s) is an interpolation function such that h(-1) = 0 and h(1) = 1, and
 - $h(\varphi)\mathcal{P}\sigma$ proliferation of tumor cells proportional to nutrient concentration
 - $h(\varphi)A$ apoptosis of tumor cells
 - $h(\varphi)\mathcal{C}\sigma$ consumption of nutrient by the tumor cells
- The constant σ_s denotes the nutrient concentration in a pre-existing vasculature, and $B(\sigma_s \sigma)$ models the supply of nutrient from the blood vessels if $\sigma_s > \sigma$ and the transport of nutrient away from the domain Ω if $\sigma_s < \sigma$.

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- A regular double-well potential Ψ , e.g., $\Psi(s)=1/4(1-s^2)^2$



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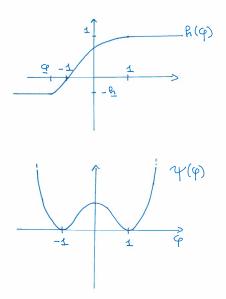
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- that admits the global attractor in a proper phase space.

The main difference with respect to the previous model is that here we do not have the total energy baance we had before. Here we only have

$$\frac{\mathrm{d}}{\mathrm{d}t} \left(\frac{1}{2} \|\nabla \varphi\|^2 + \int_{\Omega} \Psi(\varphi) \, \mathrm{d}x \right) + \|\nabla \mu\|^2 = \int_{\Omega} (\mathcal{P}\sigma - \mathcal{A}) h(\varphi) \mu \, \, \mathrm{d}x.$$

Examples of functions h and Ψ



The basic assumptions on the potential

The configuration potential Ψ lies in $C^{1,1}_{loc}(\mathbb{R})$. Moreover its derivative is decomposed as a sum of a monotone part β and a linear perturbation:

$$\Psi'(r) = \beta(r) - \lambda r, \quad \lambda \ge 0, \ r \in \mathbb{R}.$$

We normalized so that $\beta(0) = 0$ and further β complies with the growth condition

$$\exists c_{\beta} > 0: \ |\beta(r)| \leq c_{\beta}(1 + \Psi(r)) \ \forall r \in \mathbb{R},$$

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In order to prove uniqueness of solutions we also need that there exists c>0 such that

$$|\beta(r) - \beta(s)| \le c|r - s|(1 + |\beta(r)| + |\beta(s)|) \quad \forall r, s \in \mathbb{R}.$$

Note that this is still consistent with asking an at most exponential growth of β .

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Remark

The function $h(\varphi)$ is assumed to satisfy h(-1)=0 and h(1)=1. The simplest situation when this occurs is the "symmetric" case when we have $\underline{h}=0$ and $\underline{\varphi}=-1$. On the other hand we will see in what follows that dissipativity of trajectories may not hold in such a case. This motivates our choice to consider the possibility of having $\underline{h}>0$.

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Remark

We could also take $h(\varphi)=k\varphi+h_0(\varphi)$, where k>0 and h_0 is smooth and uniformly bounded. This situation is somehow simpler because, at least as long as we can guarantee that $\mathcal{P}\sigma-\mathcal{A}>0$, the linear part of h drives some mass dissipation effect in the Cahn-Hilliard type equation $\varphi_t-\Delta\mu=(\mathcal{P}\sigma-\mathcal{A})h(\varphi)$.

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We assume the initial data to satisfy

$$\sigma_0 \in L^{\infty}(\Omega), \qquad 0 \le \sigma_0 \le 1 \text{ a.e. in } \Omega,$$

$$\varphi_0 \in H^1(\Omega), \qquad \Psi(\varphi_0) \in L^1(\Omega).$$

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Theorem (Well-posedness)

Then the tumor-growth model

$$\begin{split} & \varphi_t - \Delta \mu = (\mathcal{P}\sigma - \mathcal{A})h(\varphi), \quad \varphi(0) = \varphi_0, \quad \partial_n \varphi = 0 \text{ on } \partial\Omega, \\ & \mu = -\Delta \varphi + \Psi'(\varphi), \quad \partial_n \mu = 0 \text{ on } \partial\Omega, \\ & \sigma_t - \Delta \sigma = -\mathcal{C}\sigma h(\varphi) + B(\sigma_s - \sigma), \quad \sigma(0) = \sigma_0, \quad \partial_n \sigma = 0 \text{ on } \partial\Omega \end{split}$$

admits one and only one global in time weak solution:

$$\varphi \in H^{1}(0, T; H^{1}(\Omega)') \cap C^{0}([0, T]; H^{1}(\Omega)) \cap L^{2}(0, T; H^{2}(\Omega)),$$

$$\beta(\varphi) \in L^{2}(0, T; L^{2}(\Omega)), \quad \mu \in L^{2}(0, T; H^{1}(\Omega)),$$

$$\sigma \in H^{1}(0, T; H^{1}(\Omega)') \cap C^{0}([0, T]; L^{2}(\Omega)) \cap L^{2}(0, T; H^{1}(\Omega)) \cap L^{\infty}(0, T; L^{\infty}(\Omega));$$

Moreover, for any T>0 there exists $\overline{\sigma}_T\geq 1$ such that

$$0 \le \sigma(t, x) \le \overline{\sigma}_T$$
, for a.e. $(t, x) \in (0, T) \times \Omega$,

where we can take $\overline{\sigma}_T$ independent of time if B - Ch > 0 and $\overline{\sigma}_T = 1$ if h = 0.

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Assumptions for dissipativity

Let the parameters in

$$\varphi_t - \Delta \mu = (\mathcal{P}\sigma - \mathcal{A})h(\varphi),$$

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satisfy (where
$$\mathit{h}(\mathit{r}) \equiv -\underline{\mathit{h}}$$
 for all $\mathit{r} \leq \underline{\varphi} \leq -1$)

(H1)
$$\underline{h} > 0$$
, $B - C\underline{h} > 0$,

(H2)
$$\frac{B\sigma_s}{B-C\underline{h}}<1,$$

$$\label{eq:harmonic} \mbox{(\emph{H3})} \quad \mbox{\mathcal{A}} - \mathcal{P} \frac{\mbox{B} \sigma_{\text{s}}}{\mbox{B} - \mbox{\mathcal{C}} \underline{\mbox{h}}} > 0.$$

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$$\label{eq:B-condition} (\textit{H1}) \quad \underline{\textit{h}} > 0, \qquad \textit{B} - \textit{C}\underline{\textit{h}} > 0,$$

(H2)
$$\frac{B\sigma_s}{B-C\underline{h}} < 1$$
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(H3)
$$A - P \frac{B\sigma_s}{B - C\underline{h}} > 0.$$

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These conditions essentially prescribe \underline{h} to be *strictly positive*, but *small*. Let also β have a superquadratic behavior at infinity, namely

$$\exists \, \kappa_{\beta} > 0, \, C_{\beta} \geq 0, \, p_{\beta} > 2: \, \beta(r) \operatorname{sign} r \geq \kappa_{\beta} |r|^{p_{\beta}} - C_{\beta} \, \, \forall r \in \mathbb{R}.$$



Starting from spatially homogeneous initial data we reduce to the following ODE system:

$$X' + (A - PS)h(X) = 0,$$

$$S' + CSh(X) + B(S - \sigma_s) = 0$$

where X = X(t) and S = S(t) are the spatial mean values of φ and σ .

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$$B\sigma_s - (C+B)S \leq S' \leq B\sigma_s - (B-C\underline{h})S$$

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If $C\underline{h} \geq B$, i.e. H1) ii) does not hold and X(0) << 0, S(0) >> 0 (in such a way that $\mathcal{P}S - \mathcal{A} > 0$), then it follows

$$X' = -(\mathcal{P}S - \mathcal{A})\underline{h} < 0,$$

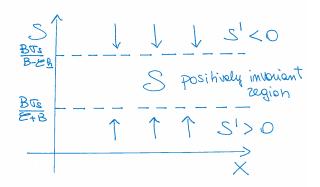
$$S' = B\sigma_s + (\mathcal{C}h - B)S > 0$$

and both |X| and S go increasing forever. Even if we restrict ourselves to $S(0) \le 1$, if X(0) < -1 then the physical constraint $S(t) \in [0,1]$ is not respected.

Assume (H1): $\underline{h} > 0$, $B - C\underline{h} > 0$. Then, the region

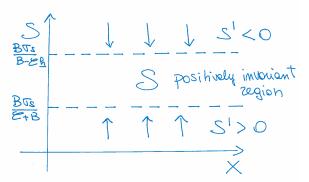
 $\mathcal{S} := \left\{ (X,S): \ \frac{B\sigma_s}{\mathcal{C} + B} \leq S \leq \frac{B\sigma_s}{B - \mathcal{C}\underline{h}} \right\} \text{ is positively invariant for the dynamical process}$

because $B\sigma_s - (C + B)S \le S' \le B\sigma_s - (B - C\underline{h})S$:



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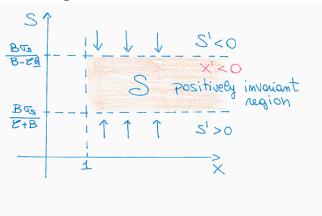
Now, if we want to keep the physical constraint $S(t) \in [0,1]$, we need to assume

(H2):
$$\frac{B\sigma_s}{B-Ch} < 1$$

Let us assume that X(0) > 1, which also implies h(X) = 1. Then, we have:

$$X' = (\mathcal{PS} - \mathcal{A})$$

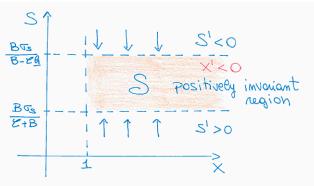
and condition (H3): $\frac{A}{P} > \frac{B\sigma_s}{B-Ch}$ prescribes that in $S \cap \{X > 1\}$ we have X' < 0:



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and condition (H3): $\frac{A}{P} > \frac{B\sigma_s}{B-Ch}$ prescribes that in $S \cap \{X > 1\}$ we have X' < 0:



On the other hand when $\frac{A}{P} \leq \frac{B\sigma_s}{B+C}$, dissipativity cannot hold. Indeed if

 $S(0) \in \left[\frac{B\sigma_s}{C+B}, \frac{B\sigma_s}{B-C\underline{h}}\right]$ and $X(0) \ge 1$, then X(t) is forced to increase forever (X' > 0).

Dissipativity and Attractor

We can define the "energy space"

$$\mathcal{X} := \left\{ (\varphi, \sigma) \in H^1(\Omega) \times L^{\infty}(\Omega) : \ \Psi(\varphi) \in L^1(\Omega) \right\}$$

and we correspondingly introduce the "magnitude" of an element $(arphi,\sigma)\in\mathcal{X}$ as

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Theorem (Dissipativity)

Under the previous compatibility conditions, there exists a positive constant C_0 independent of the initial data and a time T_0 depending only on the \mathcal{X} -magnitude of the initial data such that any weak solution satisfies

$$\|(\varphi(t),\sigma(t))\|_{\mathcal{X}}\leq C_0$$
 for every $t\geq T_0$.

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Theorem (Existence of the Attractor)

Under the previous compatibility conditions the dynamical system generated by weak trajectories on the phase space $\mathcal X$ admits the global attractor $\mathcal A$. More precisely, $\mathcal A$ is a relatively compact subset of $\mathcal X$ which is also bounded in $H^2(\Omega) \times H^1(\Omega)$ and uniformly attracts the trajectories emanating from any bounded set $\mathcal B \subset \mathcal X$.

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- 4. Include a stochastic term in phase-field models for tumor growth representing for example uncertainty of a therapy or random oscillations of the tumor phase (ongoing project with C. Orrieri and L. Scarpa).

Many thanks to all of you for the attention!

Preliminaries

- Def. \mathcal{B}_0 is an absorbing set for a semigroup S(t) on a metric space (X, d_X) iff
 - \triangleright \mathcal{B}_0 is bdd
 - ▶ $\forall B \subset X \text{ bdd } \exists T_B \geq 0 \text{ s.t. } S(t)B \subset \mathcal{B}_0 \quad \forall t \geq T_B.$
- Theorem. Let S(t) be a strongly continuous semigroup on a c.m.s. (X, d_X) . Moreover, if
 - S(t) admits an absorbing set \mathcal{B}_0 ;
 - ▶ $\forall B \subset X \text{ bdd } \exists t_B > 0 \text{ s.t. } \bigcup_{t > t_B} S(t)B \text{ is compact in } X,$

then S(t) admits a *universal attractor* A that is

$$\mathcal{A} = \bigcap_{ au>0} \overline{\bigcup_{t> au} S(t) \mathcal{B}_0}.$$

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With our work we aim to provide a contribution to the theory of free terminal time optimal control in the context of diffuse interface tumor models, where the control is applied in the nutrient equation.

FLRS: A multispecies model with velocities - with Frigeri, Lam, Schimperna

Typical structure of tumors grown in vitro:

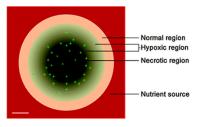


Figure: Zhang et al. Integr. Biol., 2012, 4, 1072–1080. Scale bar $100\mu\mathrm{m}=0:1\mathrm{mm}$

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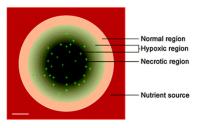


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A continuum thermodynamically consistent model is introduced with the ansatz:

- sharp interfaces are replaced by narrow transition layers arising due to adhesive forces among the cell species: a diffuse interface separates tumor and healthy cell regions
- proliferating and dead tumor cells and healthy cells are present, along with a nutrient (e.g. glucose or oxigene)
- tumor cells are regarded as inertia-less fluids: include the velocity satisfying a Darcy type law with Korteveg term

- $\varphi_p, \varphi_d, \varphi_h \in [0, 1]$: the volume fractions of the cells:
 - $\triangleright \varphi_{p}$: proliferating tumor cell fraction
 - $\triangleright \varphi_d$: dead tumor cell fraction
 - φ_h : healthy cell fraction
- The variables above are naturally constrained by the relation $\varphi_p + \varphi_d + \varphi_h = 1$ hence it suffices to track the evolution of φ_p and φ_d and the vector $\boldsymbol{\varphi} := (\varphi_p, \varphi_d)^\top$ lies in the simplex $\Delta := \{ \boldsymbol{y} \in \mathbb{R}^2 : \ 0 \leq y_1, y_2, \ y_1 + y_2 \leq 1 \} \subset \mathbb{R}^2$

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- n: the nutrient concentration (it was σ before)
- $u:=u_i$, i=1,2,3: the tissue velocity field. We treat the tumor and host cells as inertial-less fluids and assume that the cells are tightly packed and they march together
- q: the cell-to-cell pressure



Letting J_i , $i \in \{p, d, h\}$, denote the mass fluxes for the cells, then the general balance law for the volume fractions reads as

$$\partial_t \varphi_i + \operatorname{div}(\varphi_i \mathbf{u}) = -\operatorname{div} \mathbf{J}_i + \mathbf{S}_i \quad \text{ for } i \in \{p, d, h\}$$

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where $F = F_0 + F_1$ is a multi-well configuration potential, e.g.

$$\begin{split} & \digamma_0(\varphi_{p},\varphi_{d}) := \varphi_{p} \log \varphi_{p} + \varphi_{d} \log \varphi_{d} + (1 - \varphi_{p} - \varphi_{d}) \log (1 - \varphi_{p} - \varphi_{d}) \\ & \digamma_1(\varphi_{p},\varphi_{d}) := \frac{\chi}{2} \left(\varphi_{d} (1 - \varphi_{d}) + \varphi_{p} (1 - \varphi_{p}) + (1 - \varphi_{d} - \varphi_{p}) (\varphi_{d} + \varphi_{p}) \right) \end{split}$$

Letting J_i , $i \in \{p, d, h\}$, denote the mass fluxes for the cells, then the general balance law for the volume fractions reads as

$$\partial_t \varphi_i + \operatorname{div}(\varphi_i \boldsymbol{u}) = -\operatorname{div} \boldsymbol{J}_i + \boldsymbol{S}_i \quad \text{ for } i \in \{p, d, h\}$$

where we set $S_h = 0$, whereas S_p , S_d may depend on n, φ_p and φ_d Assume: the tumor growth process tends to evolve towards (local) minima of the free energy functional of Ginzburg–Landau type:

$$E(\varphi_{p}, \varphi_{d}) := \int_{\Omega} \frac{F(\varphi_{p}, \varphi_{d})}{\int_{\Omega} F(\varphi_{p}, \varphi_{d})} + \frac{1}{2} |\nabla \varphi_{p}|^{2} + \frac{1}{2} |\nabla \varphi_{d}|^{2} dx$$

where $F = F_0 + F_1$ is a multi-well configuration potential, e.g.

$$\begin{split} & F_0(\varphi_p, \varphi_d) := \varphi_p \log \varphi_p + \varphi_d \log \varphi_d + (1 - \varphi_p - \varphi_d) \log (1 - \varphi_p - \varphi_d) \\ & F_1(\varphi_p, \varphi_d) := \frac{\chi}{2} \left(\varphi_d (1 - \varphi_d) + \varphi_p (1 - \varphi_p) + (1 - \varphi_d - \varphi_p) (\varphi_d + \varphi_p) \right) \end{split}$$

The fluxes J_i are defined as follows:

$$J_i = -M_i \nabla \mu_i, \quad \mu_i := \frac{\delta E}{\delta \varphi_i} = -\Delta \varphi_i + F_{,\varphi_i} \quad \text{ for } i = p, d$$

FLRS: the velocity and nutrient evolutions

We set $J_h = -J_p - J_d$, then upon summing up the three mass balances for i = p, d, h, using the fact that $\varphi_p + \varphi_d + \varphi_h = 1$ and $S_h = 0$, we deduce the following relation:

$$\operatorname{div} \boldsymbol{u} = S_p + S_d =: S_t$$

The velocity field u is assumed to fulfill Darcy's law:

$$\mathbf{u} = -\nabla \mathbf{q} - \varphi_{p} \nabla \mu_{p} - \varphi_{d} \nabla \mu_{d}$$

where q denotes the cell-to-cell pressure and the subsequent two terms have the meaning of Korteweg forces

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Since the time scale of nutrient diffusion is much faster (minutes) than the rate of cell proliferation (days), the nutrient is assumed to evolve quasi-statically:

$$0=-\Delta n+\varphi_p n$$

where $\varphi_p n$ models consumption by the proliferating tumor cells



Goal: to study this multispecies model including different mobilities, singular potential and non-Dirichlet b.c.s on the chemical potential. The main problems are:

• we have two different Cahn-Hilliard equations with non-zero right hand sides: $\partial_t \varphi_i - \text{div}(M_i \nabla \mu_i - \varphi_i \mathbf{u}) = S_i$ and if we do not choose the Dirichlet b.c.s on μ_i then we need to estimate the mean values of $\mu_i = -\Delta \varphi_i + F_{,\varphi_i}$ containing a multiwell logarithmic type potential F_0

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- such a relation does not involve directly the singular part F_0 . Hence, the evolution of y_p, y_d are not automatically compatible with the physical constraint and this has to be proved by assuming proper conditions on coefficients and making a careful choice of the boundary conditions
- the choice $(M_i \nabla \mu_i \varphi_i \mathbf{u}) \cdot \mathbf{n} = 0$ seems essential

FLRS: The weak notion of solution

Definition. $(\varphi_p, \varphi_d, \mathbf{u}, q, n)$ is a weak solution to the problem in $(0, T) \times \Omega$ if the previous equations hold, for a.e. $t \in (0, T)$ and for i = p, d, in the following weak sense:

$$\begin{split} &\langle \partial_t \varphi_i, \zeta \rangle + \int_{\Omega} M_i \nabla \mu_i \cdot \nabla \zeta - \varphi_i \boldsymbol{u} \cdot \nabla \zeta \, dx = \int_{\Omega} S_i \zeta \, dx \quad \forall \zeta \in H^1(\Omega), \\ &\int_{\Omega} \mu_i \zeta \, dx = \int_{\Omega} \nabla \varphi_i \cdot \nabla \zeta + \eta_i \zeta + F_{1,\varphi_i}(\varphi_p, \varphi_d) \zeta \, dx \quad \forall \zeta \in H^1(\Omega), \\ &\int_{\Omega} \boldsymbol{u} \cdot \nabla \xi \, dx = -\int_{\Omega} (S_p + S_d) \xi \, dx \quad \forall \xi \in H^1_0(\Omega), \\ &\int_{\Omega} \boldsymbol{u} \cdot \zeta \, dx = \int_{\Omega} -\nabla q \cdot \zeta - \varphi_p \nabla \mu_p \cdot \zeta - \varphi_d \nabla \mu_d \cdot \zeta \, dx \quad \forall \zeta \in (L^2(\Omega))^d, \\ &0 = -\Delta n + \varphi_p n \quad \text{a.e. in } \Omega, \\ &\eta_i = F_{0,\varphi_i}(\varphi_p, \varphi_d) \quad \text{a.e. in } \Omega, \\ &S_p = \Sigma_p(n, \varphi_p, \varphi_d) + m_{pp} \varphi_p + m_{pd} \varphi_d \quad \text{a.e. in } \Omega, \\ &S_d = \Sigma_d(n, \varphi_p, \varphi_d) + m_{dp} \varphi_p + m_{dd} \varphi_d \quad \text{a.e. in } \Omega. \end{split}$$

Moreover, there hold the initial conditions

$$\varphi_p(x,0) = \varphi_{p,0}(x), \quad \varphi_d(x,0) = \varphi_{d,0}(x)$$
 a.e. in Ω ,

where $\langle \cdot, \cdot \rangle$ denotes the duality pairing between $H^1(\Omega)$ and its dual $H^1(\Omega)'$.



Set $\Sigma(n, \varphi_p, \varphi_d) := (\Sigma_p, \Sigma_d)$ and $\underline{\underline{M}} = (m_{ij}), i, j \in \{p, d\}$, the matrix of the coefficients of the mass souces in the Cahn-Hilliard equations: $(S_p, S_d) = \Sigma + \underline{M}(\varphi_p, \varphi_d)^T$

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- \bullet Σ is globally Lipschitz and
- that there exist a closed and sufficiently regular subset Δ_0 contained in the open simplex Δ and constants $K_{p,-}, K_{p,+}, K_{d,-}, K_{d,+} \in \mathbb{R}$, with $K_{p,-} \leq K_{p,+}$ and $K_{d,-} \leq K_{d,+}$, such that $\Sigma(\mathbb{R}^3) \subset [K_{p,-}, K_{p,+}] \times [K_{d,-}, K_{d,+}]$
- for any $\mathbf{x}=(x_p,x_d)\in [K_{p,-},K_{p,+}]\times [K_{d,-},K_{d,+}]$, there holds

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Assumptions on the initial data:

- $\bullet \ \varphi_{p,0}, \varphi_{d,0} \in H^1(\Omega) \text{ with } 0 \leq \varphi_{p,0}, \quad 0 \leq \varphi_{d,0}, \quad \varphi_{p,0} + \varphi_{d,0} \leq 1 \text{ a.e. in } \Omega,$
- the mean values satisfy $(\frac{1}{|\Omega|}\int_{\Omega}\varphi_{p,0}(x)\,dx, \frac{1}{|\Omega|}\int_{\Omega}\varphi_{d,0}(x)\,dx) \in \operatorname{int}\Delta_0$ and $F_0(\varphi_{p,0},\varphi_{d,0}) \in L^1(\Omega)$

FLRS: Examples of mass sources

Examples of mass sources in $\partial_t \varphi_i - \operatorname{div}(M_i \nabla \mu_i - \varphi_i \mathbf{u}) = S_i$ for $i \in \{p, d\}$ complying with the assumptions in the "logarithmic" case are:

$$S_p = \lambda_M g(n) - \lambda_A \varphi_p$$
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for positive constants λ_M , λ_A , λ_L (with $\lambda_M(\lambda_A + \lambda_L) < \lambda_A \lambda_L$, $\lambda_A < 2\lambda_L$) and a bounded positive function g such that $0 < g(s) \le 1$, e.g., $g(s) = \max(n_c, \min(s, 1))$ for some constant $n_c \in (0, 1)$.

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- the growth of the proliferating tumor cells due to nutrient consumption at a constant rate λ_M
- the death of proliferating tumor cells at a constant rate λ_A , which leads to a source term for the necrotic cells
- ullet the lysing/disintegration of necrotic cells at a constant rate λ_L

FLRS: Existence of weak solutions

The main result of S. Frigeri, K.-F. Lam, E. R., G. Schimperna, Comm Math Sci. (2018)

Theorem

For every T>0 here exists at least one weak solution $(\varphi_p,\mu_p,\eta_p,\varphi_d,\mu_d,\eta_d,\mathbf{u},q,n)$ to the multi-species tumor model on [0,T] with the regularity

$$\varphi_{i} \in H^{1}(0, T; H^{1}(\Omega)') \cap L^{\infty}(0, T; H^{1}(\Omega)) \cap L^{2}(0, T; H^{2}(\Omega)),$$
with $0 \leq \varphi_{i} \leq 1$, $\varphi_{p} + \varphi_{d} \leq 1$ a.e. in Q , for $i = p, d$,
$$\mu_{i} \in L^{2}(0, T; H^{1}(\Omega)), \quad \eta_{i} \in L^{2}(Q),$$

$$\mathbf{u} \in L^{2}(Q) \text{ with div } \mathbf{u} \in L^{2}(Q), \quad q \in L^{2}(0, T; H^{1}(\Omega)),$$

$$n \in (1 + L^{2}(0, T; H^{2}(\Omega) \cap H^{1}(\Omega))), \quad 0 \leq n \leq 1 \text{ a.e. in } Q.$$

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$$n \in (1 + L^{2}(0, T; H^{2}(\Omega) \cap H^{1}(\Omega))), \quad 0 \leq n \leq 1 \text{ a.e. in } Q.$$

Notice that the boundary conditions:

$$(M_i \nabla \mu_i - \varphi_i \mathbf{u}) \cdot \mathbf{n} = 0, \quad \partial_{\mathbf{n}} \varphi_i = 0, \quad \mathbf{q} = 0, \quad \mathbf{n} = 1 \text{ on } \Gamma$$

are incorporated in the definition of weak solutions



FLRS: an idea of the proof

1 consider a regularized version of this problem by replacing the singular potential F_0 by its Moreau–Yosida approximation F_ε , and by introducing some suitable truncation functions. The latter choice is due to the fact that F_ε is no longer a singular function, and consequently the uniform boundedness properties $0 \le \varphi_p$, $0 \le \varphi_d$, $\varphi_p + \varphi_d \le 1$ are not expected to hold in the approximation level.

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- 2 to prove existence of a solution to the regularized system a further regularization and a Schauder fixed point argument: only exploits elementary existence and uniqueness results methods for PDEs
- 3 derive the bounds independent of the regularization parameters in order to pass to the limit in the approximation scheme via compactness tools: the main problem is to bound the mean values of φ_i away from the potential bareers

Denoting $\mathbf{y}(t) := ((\varphi_p)_{\Omega}(t), (\varphi_d)_{\Omega}(t)), (\Sigma)_{\Omega} = ((\Sigma_p)_{\Omega}, (\Sigma_d)_{\Omega}),$ then by testing by 1 the mass balances

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Taking
$$t=t_*$$
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$$\frac{d}{dt}\mathbf{y}(t_*)\cdot\mathbf{n}<0.$$

Hence y(t) cannot leave Δ_0 and so there exist positive constants $0 < c_1 < c_2 < 1$:

$$c_1 \leq (\varphi_p)_{\Omega}(t), (\varphi_d)_{\Omega}(t) \leq c_2, \quad c_1 \leq (\varphi_p + \varphi_d)_{\Omega}(t) \leq c_2 \quad \forall t \in [0, T].$$