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Authors

Pietro Mascheroni¹, Cinzia Stigliano², Melania Carfagna³, Daniela P. Boso¹, Luigi Preziosi³, Paolo Decuzzi⁴, Bernhard A. Schrefler^{1,2}

Title

Predicting the Growth of Glioblastoma Multiforme Spheroids using a Multiphase Porous Media Model

Affiliations

¹ Dipartimento di Ingegneria Civile Edile ed Ambientale, Università di Padova, Via Marzolo 9, 35131 Padova, Italy

² Nanomedicine Department, Houston Methodist Research Institute, 6670 Bertner Street, 77030 Houston, TX

³ Dipartimento di Scienze Matematiche, Politecnico di Torino, Corso Duca degli Abruzzi 24, 10129 Torino, Italy

⁴ Department of Drug Design and Development, Fondazione Istituto Italiano di Tecnologia, 16163 Genova, Italy

Corresponding Author:

Pietro Mascheroni

Email: pietro.mascheroni@dicea.unipd.it

Phone: +39 049 826 5358

Abstract

Tumor spheroids constitute an effective *in vitro* tool to investigate the avascular stage of tumor growth. These three dimensional cell aggregates reproduce the nutrient and proliferation gradients found in the early stages of cancer and can be grown with a strict control of their environmental conditions. In the last years, new experimental techniques have been developed to determine the effect of mechanical stress on the growth of tumor spheroids. These studies report a reduction in cell proliferation as a function of increasingly applied stress on the surface

of the spheroids. This work presents a specialization for tumor spheroid growth of a previous more general multiphase model. The equations of the model are derived in the framework of porous media theory and constitutive relations for the mass transfer terms and the stress are formulated on the basis of experimental observations. A set of experiments is performed, investigating the growth of U-87MG spheroids both freely growing in the culture medium and subjected to an external mechanical pressure induced by a Dextran solution. The growth curves of the model are compared to the experimental data, with good agreement for both the experimental settings. A new mathematical law regulating the inhibitory effect of mechanical compression on cancer cell proliferation is presented at the end of the paper. This new law is validated against experimental data and provides better results compared to other expressions in literature.

Keywords

Mathematical modeling; Cancer growth; U-87MG spheroids; Mechanical compression of spheroids; Growth inhibition

1. Introduction

Cancer is a complex disease involving primarily uncontrolled cell proliferation and migration to distant regions of the body (Longo et al. 2011). From the second half of the last century the scientific community has become more and more aware of the difficulties that arise when treating this illness. Nowadays it is clear that a combined effort from all the physical sciences is necessary to advance our understanding of the disease and promote the discovery of new cures (Michor et al. 2011; Leder et al. 2014). The pioneering works of Greenspan and coworkers (Greenspan 1976) paved the way for the development of mathematical models that could investigate the basic principles underlying cancer progression and predict the outcome of therapies. Most continuum models, as the one presented in this work, deal with the *avascular phase* of tumor growth.

During this stage of cancer progression, a small cluster of cancer cells arise in a healthy tissue due to mutations that alter their biochemical pathways. This small region of abnormal cells grows at the expense of the host counterpart, nourished by oxygen and nutrients that diffuse from the vasculature nearby (Jain 1988; Grantab et al. 2006). At a certain point the external nutrients are not enough to sustain the expansion of the growing mass, leading to the formation of cell proliferation gradients starting from the outer regions of the tumor. As time passes by, cancer cells at the center of the tumor experience severe hypoxia and critical conditions that lead to the death and consequent necrosis of some of them. Finally a steady state arises, where cell proliferation at the tumor border balances cell death at the tumor center (Folkman and Hochberg 1973). The subsequent stage of cancer progression is termed the *vascular phase*, where tumor cells recruit new blood vessels from the host vasculature through tumor angiogenesis. In this second stage the cancer resumes its previous growth and eventually enters the last stage of the illness, the *metastatic phase*, where malignant cells evade the tumor area to form metastases at distant regions of the body.

Since the study of the first stage can be performed in a more controlled experimental setting, a large set of literature is devoted to the analysis of avascular tumor growth *in vitro*. Experiments are usually carried out on tumor spheroids, three dimensional aggregates of cancer cells that grow in an approximately spherical shape (Sutherland et al. 1971; Kim et al. 2010; Vinci et al.

2012; Mikhail et al. 2013). The investigations on tumor spheroids allow evaluating the extent of the gradients of nutrients and cell proliferation and, more recently, the action of a mechanical stress exerted on the cell aggregates. Helmlinger and coworkers (Helmlinger et al. 1997) grow tumor spheroids in gels with varying stiffness, and report a decrease in proliferation for stiffer gels. Another example of spheroids grown in gels with varying stiffness is found in (Kaufman et al. 2005). There, cell proliferation and motility are investigated for different concentrations of collagen in the gel matrix. In that work, a positive correlation between increasing concentrations of collagen and cell invasion is reported, followed by an opposite effect on the growth of the spheroids. The original work of Helmlinger is extended in (Cheng et al. 2009), where the authors perform similar experiments and introduce new tools to quantify spheroid deformation and variations in cell proliferation and apoptosis. Another set of experiments is presented in (Desmaison et al. 2013), where tumor spheroids are subjected to asymmetric stress fields by the use of microstructured substrates. Finally, Montel and Delarue in two subsequent papers (Montel et al. 2011; Delarue et al. 2014) apply mechanical forces on the surface of tumor spheroids through the osmotic effect of Dextran solutions with different concentrations. All these studies report a decrease of tumor cell proliferation as a consequence of the applied stress, even though they are carried out via different experimental configurations.

The earliest continuum models applied to spheroid growth focus on the dependence of cell proliferation on nutrients and other biochemical factors, as reported in the comprehensive reviews of (Preziosi and Tosin 2009a; Lowengrub et al. 2010). They are based on mass balance laws for cells and advection-reaction-diffusion equations for nutrient evolution. Later models include more components and the mechanical interaction between them. For these cases, which are usually defined in the framework of mixture theory or porous media theory, momentum balance equations and constitutive relations are needed for describing the mechanical response of each component (Ambrosi and Mollica 2004; Preziosi and Tosin 2009a; Ehlers et al. 2009; Giverso et al. 2015). Among the first, (Chaplain et al. 2006; Galle et al. 2009) incorporated the effect of mechanical stress on cell proliferation obtaining results matching experimental observations. There followed several papers, where this example of

mechanotransduction is investigated with similar approaches ([Kim et al. 2011](#); [Ciarletta et al. 2013a](#); [Ciarletta et al. 2013b](#); [Mpekris et al. 2015](#)).

This work arises as an extension of the modeling framework presented in ([Sciumè et al. 2013b](#)).

The model is specialized for tumor spheroids and the solution procedure is simplified, as all the new equations are formulated in spatial coordinates with no need of a reference configuration.

A set of experiments is carried out on spheroid cultures to validate the equations. The comparison with the experiments is performed both with spheroids growing freely in the culture medium and subjected to increasing mechanical loads, giving in each case a good match. The experiments suggested the presence of a “master curve”, a common growth curve underlying the spheroid growth dynamics. Finally, a new constitutive relation is proposed to describe the inhibitory effect of the stress on cell proliferation, with a better performance in terms of matching the experimental results when compared to the existing laws in literature.

The rest of the paper is organized as follows. Section 2 introduces the mathematical framework, based on the Thermodynamically Constrained Averaging Theory (TCAT), and the differences with the original model. In the last part of that section, the equations are specialized to the case of tumor spheroids. Section 3 presents the experimental setup for the two culturing conditions. Finally, results from the simulations are presented in Section 4 and the discussion follows in Section 5.

2. A biphasic tumor model

This mathematical model is developed in the framework of the porous media theory, and the governing equations are derived through the TCAT ([Gray and Miller 2005](#)), ([Gray and Miller 2014](#)). We start by defining the problem in terms of microscopic relations amongst the constituents. The TCAT approach is used to transform these microscopic laws into mathematically and physically consistent macroscale relationships, which describe the tumor at the tissue scale. By doing so, the complexity due to the high spatial variability at the microscale is overcome and equations for average quantities describing the tumor behavior are formulated directly. The closed form of the problem is finally obtained by introducing constitutive relations into the macroscale conservation equations. Detailed information about the mathematical

model and its derivation are found in previous work of the authors (Sciumè et al. 2013b; Sciumè et al. 2013a; Sciumè et al. 2014). In this work we describe the behavior of the following constituents, or phases: (i) the tumor cells (TCs), which partition into living (LTCs) and necrotic (NTCs) cells, and (ii) the interstitial fluid (IF) (Fig. 1). The extracellular matrix (ECM) is considered together with the tumor cells and the union of the two entities constitutes the solid skeleton of the tumor. The interstitial fluid phase flows through the pores of this solid matrix, carrying nutrients, growth factors and waste products. Cell proliferation is related to nutrient concentration, whereby cells stop to proliferate and, after some time, undergo necrosis and lysis, if subjected to low levels of nutrients or high levels of mechanical stress. In the following equations, t will denote the union of the tumor cells and ECM and l the interstitial fluid.

2.1 The governing equations

The solid portion of the tumor is modeled as a porous solid with porosity, and its volume fraction is defined as ϕ . Hence, the interstitial fluid occupies the rest of the volume and the sum of all the volume fractions has to be unity:

$$\phi + \phi_l = 1 \quad (12)$$

We write the governing equations for the tumor volume fraction ϕ , the interstitial fluid pressure p_l , the nutrient mass fraction ϕ_n and the necrotic mass fraction ϕ_{nt} . These equations are obtained from the general form of the mass and momentum balance equations of phases and species, according to the TCAT derivation.

The mass balance equations for the phases are written as:

$$\rho_t \frac{d\phi}{dt} + \nabla \cdot (\rho_t \mathbf{v}_t) = \sigma_t \quad (34)$$

$$\rho_l \frac{d\phi_l}{dt} + \nabla \cdot (\rho_l \mathbf{v}_l) = \sigma_l \quad (56)$$

where ρ_t is the density and \mathbf{v}_t the velocity of phase t . The terms σ_t and σ_l account for the inter-phase exchange of mass related to cell growth and cell lysis respectively.

The tumor phase t is composed of two subpopulations, namely necrotic and living cells. The necrotic portion is described by the mass fraction ϕ_{nt} , so that the mass fraction of living cells is ϕ_{lt} . We assume that there is no diffusion for the necrotic and living species, and that necrotic cells

exchange mass with the interstitial fluid through the lysis term. The mass fraction of necrotic tumor cells (ϕ_n) and living tumor cells (ϕ_l) are thus given by:

$$\phi_n = \frac{m_n}{m_n + m_l} \quad (78)$$

$$\phi_l = \frac{m_l}{m_n + m_l} \quad (910)$$

where γ represents an intra-phase exchange term accounting for the rate of death of living tumor cells.

The evolution of the mass fraction of oxygen (ϕ_o), the unique nutrient considered here, follows the equation:

$$\frac{\partial \phi_o}{\partial t} + \nabla \cdot (\mathbf{v}_o \phi_o) = \frac{D_o}{\tau} \nabla^2 \phi_o - \gamma_o \phi_o \quad (1112)$$

where D_o is the diffusion coefficient of oxygen in the extracellular space and γ_o is a mass exchange term accounting for nutrient consumption by tumor cells metabolism and growth. **Note that the mass exchange term in equation 12 is included in the reaction term of equations 4 and 6, since the latter is related to the exchange of mass and nutrients between the two phases (for more details see (Sciumè et al. 2013b) and the references therein).** Following porous media theory (Lewis and Schrefler 1998; Pinder and Gray 2008), the mechanical stress exerted on the solid phase is described through the effective stress tensor $\boldsymbol{\sigma}^s$, given by:

$$\boldsymbol{\sigma}^s = -p \mathbf{I} + \boldsymbol{\sigma} - p \mathbf{I} \quad (1314)$$

where \mathbf{I} is the unit tensor, $\boldsymbol{\sigma}$ the total stress tensor in the tissue, p is the fluid pressure in the interstitial fluid and α is Biot's coefficient defined by:

$$\alpha = \frac{K - K_s}{K} \quad (1516)$$

with K bulk modulus of the unsaturated skeleton and K_s bulk modulus of the solid phase. The relative velocity of the interstitial fluid phase is given by a Darcy type equation obtained by TCAT as:

$$\mathbf{v}_o = -\frac{\kappa}{\mu} \nabla p \quad (1718)$$

where κ is the intrinsic permeability of the solid matrix and μ the dynamic viscosity of the interstitial fluid. The equations of state for the phases can be approximated as:

$$1920 \text{ * MERGEFORMAT ()}$$

$$2122 \text{ * MERGEFORMAT ()}$$

here and are the liquid and solid compressibility respectively and is the normal stress at the solid surface averaged over the solid surface. Considering 20, equation 4 can be written as:

$$2324 \text{ * MERGEFORMAT ()}$$

Following the same steps for equation 6 it is possible to obtain:

$$2526 \text{ * MERGEFORMAT ()}$$

and summing 24 and 26 gives:

$$2728 \text{ * MERGEFORMAT ()}$$

where the gradients of the densities have been neglected and the constrained in 2 has been exploited. Inserting 18 and substituting in 28 leads to:

$$2930 \text{ * MERGEFORMAT ()}$$

2.2 The constitutive relationship for the stress

In order to close the system of equations it is necessary to define a constitutive relation for the stress in the tumor phase. A series of experiments based on single-cell force spectroscopy (Baumgartner et al. 2000; Puech et al. 2005; Helenius et al. 2008; Friedrichs et al. 2013) suggests the following phenomenological description at the microscale. When cells are well separated from each other, they do not experience any interaction. **As soon as the distance between two cells is below a certain threshold they start attracting each other and, once cells are in contact, an adhesive force builds up if they tend to be pulled apart. If the two cells are further pushed together, a repulsive force is observed. This repulsive force tends to high values as cells become more and more packed.** Note that, in the context of porous media theory, the volume fraction of tumor cells can be chosen as a surrogate for cell distance. In mathematical terms, this can be written as a pseudo-potential law (Byrne and Preziosi 2003) (see (Preziosi

and Tosin 2009b) for qualitative analyses on this kind of nonlinear systems) that describes the stress in the tumor tissue:

$$3132 \backslash * \text{MERGEFORMAT} ()$$

In Fig. 2, a schematic of cell behavior is provided depicting the behavior of the cellular system for different volume fractions. Note that with this description the ensemble of tumor cells and ECM behaves like an elastic fluid and the effective stress follows the relation:

$$3334 \backslash * \text{MERGEFORMAT} ()$$

with defined in equation 32 and positive in compression.

2.3 The mass transfer relations

The reaction terms in equation 4 represent tumor cells growth and lysis respectively. In particular, the first term is related to cell proliferation and depends on the exchange of mass between the interstitial fluid and the living portion of the tumor. Its form is given by:

$$3536 \backslash * \text{MERGEFORMAT} ()$$

where the coefficient accounts for the nutrient uptake and for the mass of interstitial fluid that becomes tumor due to cell growth. The function accounts for the effect of nutrient level on cell growth, while describes the inhibition of cell growth due to the mechanical stress exerted on the cells. Finally, the factor accounts for the volume fraction of viable tumor cells (i.e. only viable cells can proliferate).

The second reaction term in equation 4 accounts for cell lysis in the necrotic cell population (NTCs). Its form is given by:

$$3738 \backslash * \text{MERGEFORMAT} ()$$

here, takes into account the degradation of cellular membranes and the mass conversion into interstitial fluid. Since is the volume fraction of necrotic cells, the lysis term is active only on the dead portion of the tumor tissue.

The rate of death of tumor cells in equation 8 is described by the relation:

$$3940 \backslash * \text{MERGEFORMAT} ()$$

where the parameter μ regulates the rate of cell death. The function ϕ describes cell death by lack of nutrient. By doing so, cell death is considered to solely depend on nutrient concentration. Note that equation 40 can be readily modified to include also other effects (i.e. drugs or mechanical pressure).

During the growth of the tumor, nutrients are subtracted from the interstitial fluid, so that the mass exchange term in equation 12 takes the form:

$$4142 \backslash * \text{MERGEFORMAT} ()$$

This expression is validated experimentally in (Casciari et al. 1992a; Casciari et al. 1992b) and takes into account the dependence of nutrient consumption on the local level of nutrient in the tissue. The two oxygen uptake parameters μ_0 and $\mu_{0.5}$ describe respectively the order of magnitude of oxygen uptake in the tumor and the oxygen mass fraction at which oxygen consumption is reduced by half. The functions ϕ , ψ and χ are derived from phenomenological arguments and are selected to be similar to the available literature on the topic (Byrne and Preziosi 2003; Roose et al. 2007; Wise et al. 2008; Kim et al. 2011; Preziosi and Vitale 2011; Ambrosi et al. 2012; Sciumè et al. 2013b; Mpekris et al. 2015). In particular, the following set is assumed:

$$4344 \backslash * \text{MERGEFORMAT} ()$$

$$4546 \backslash * \text{MERGEFORMAT} ()$$

$$4748 \backslash * \text{MERGEFORMAT} ()$$

here $\mu_{0.5}$ is the oxygen threshold value below which cell growth is inhibited, the constant μ_0 is the environmental mass fraction of oxygen and the Macaulay brackets $\langle \cdot \rangle$ indicate the positive value of their argument. Since μ within the spheroid can only be equal or smaller than μ_0 , it follows that the brackets will return a number between one (1) and zero (0) (Sciumè et al. 2013b; Sciumè et al. 2014). The constants μ_0 and $\mu_{0.5}$ (with $\mu_0 > \mu_{0.5}$) account for the action of mechanical stress on cell proliferation, modeling the inhibitory effect of compression on tumor cells duplication (Helmlinger et al. 1997; Cheng et al. 2009; Montel et al. 2012). Note that the expression for μ is different from the usual forms assumed in literature. However, as it will be shown in the next sections, this relation is able to better describe the experimental results.

2.4 The tumor spheroid case

The mathematical model presented above is extended in view of the comparison with the experiments. In this work the focus is on modeling tumor spheroids, which are an aggregate of tumor cells approximately of spherical shape. As a starting point, we adapt the equations of the more general model for spherical symmetry. If the constituents are assumed incompressible and the densities of the phases are supposed to be equal (ρ), equation 28 becomes:

$$\rho \frac{d\mathbf{v}}{dt} = \nabla \cdot \mathbf{T} + \mathbf{f} \quad (49)$$

In spherical symmetry, equation 50 reads:

$$\rho \frac{dv_r}{dt} = \frac{1}{r^2} \frac{d}{dr} (r^2 T_{rr}) - \frac{2}{r} T_{\theta\theta} + f_r \quad (50)$$

which, by symmetry, gives:

$$T_{\theta\theta} = T_{\phi\phi} = T_{\psi\psi} \quad (51)$$

Substituting the new relation in equation 18 gives the expression for the phases velocities as:

$$\mathbf{v} = \frac{1}{\rho} \nabla p \quad (52)$$

From the point of view of the motion of the phases, since the hydraulic permeability is a positive constant, 56 implies that the interstitial fluid is directed opposite to the pressure gradient while tumor cells move along it. When the interstitial fluid pressure is higher in the tumor center, the interstitial fluid flows towards the boundary of the tumor whereas tumor cells move in the direction of the tumor center as observed experimentally in (Dorie et al. 1982; Jain and Stylianopoulos 2010) and already discussed in (Byrne and Preziosi 2003). This mechanism enables the recirculation of interstitial fluid in the tumor tissue: tumor cells in the inner regions of the tumor become necrotic due to nutrient deprivation and turn into interstitial fluid after lysis. This fluid flows towards tumor periphery and can be employed by proliferating cells. From the equilibrium equation for the total stresses (\mathbf{T}) and equations 14 and 34 it follows the important relation:

$$\nabla p = -\nabla \cdot \mathbf{T} \quad (53)$$

The final system of equations for spherical symmetry and with the new expressions for the velocities reads:

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6162* MERGEFORMAT ()

6364* MERGEFORMAT ()

The growth of the spheroid is modeled as a free-boundary problem, and the interface constituted by the tumor cells moves with velocity :

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with R being the radius of the spheroid. To close the differential problem in 60 - 64 it is necessary to define a set of boundary and initial conditions. In particular, no-flow boundary conditions are assumed at the spheroid center, while Dirichlet boundary conditions are assumed on the tumor external surface:

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6970* MERGEFORMAT ()

Note that the first condition in 70 implies prescribing an external stress on the tumor surface, since from 32 and 34 we have:

7172* MERGEFORMAT ()

In the case of a stress-free growing spheroid the external volume fraction satisfies:

7374* MERGEFORMAT ()

Finally, the following initial conditions are assumed throughout the domain:

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3. Materials and methods

3.1 Cell culture and spheroids formation

Human multiforme glioblastoma U-87 MG cells (ATCC) are grown at 37 °C at 5% CO₂ in EMEM (HyClone) supplemented with 50 U/mL penicillin, 50 µg/mL streptomycin and 10% FBS (v/v). Multicellular U-87 MG spheroids are prepared by the liquid overlay method ([Sutherland et al.](#)

1981; Carlsson and Yuhas 1984). Briefly, serum free EMEM medium with 2% (w/v) agar is prepared and sterilized; 50 μ L of the agar solution is added to the bottom of each well of the 96-well plates to prevent cell adhesion onto the well surface. Plates are allowed to cool down before use. U-87 MG cells are counted and then seeded at different densities: 1000, 5000 and 10000 cells/well. Plates are centrifuged for 5 min at 1000 x g. Spheroid diameter is measured every 2 days using Nikon Eclipse Ti microscope (Nikon) with NIS-Element software. The culture medium is replaced with fresh medium every 3 days.

3.2 Cell viability test with Dextran solutions and spheroid compression experiments

For the compression experiments, Dextran is added to the culture medium to exert mechanical stress on the spheroids as reported in (Montel et al. 2011; Montel et al. 2012; Delarue et al. 2014). Briefly, cell culture medium is mixed with the purified Dextran () and the resulting solution is placed at 37°C to obtain full solubilization. To test the effect of the Dextran solutions on U-87 MG cell viability, XTT assay is performed (Stigliano et al. 2015). In particular, 10000 cells are plated in each well of 96-well plate. After 24h, the medium is substituted with different concentrations of Dextran medium (20g/l, 55g/l and 80g/l). After 72h of incubation, the XTT assay is performed according to the **manufacturer's protocol**. The same solutions of Dextran at different concentrations are prepared to test the effect of different mechanical pressures on the surface of the spheroids. In particular, 5000 U-87 MG cells are seeded in 48-well plates (day 0), as reported before, and the Dextran medium is added after spheroid formation (day 3) at a concentration of 20g/l to exert 1kPa, 55g/l to exert 5kPa and 80g/l to exert 10kPa.

The stress acting on the spheroids is estimated as in (Montel et al. 2011; Montel et al. 2012; Delarue et al. 2014) via the following mathematical expression:

where P is the external mechanical pressure (Pa), and C is the concentration of Dextran (% w/w). This expression was originally derived to describe the osmotic pressure of Dextran solutions in colloidal systems (Bonnet-Gonnet et al. 1994) and then validated to hold also for the spheroid compression experiments (Montel et al. 2012).

Spheroid diameter measurement and medium replacement follow the same procedures described above.

4. Results

4.1 Evolution of the phases

The mathematical framework presented above is applied to analyze the growth of a multicellular tumor spheroid *in vitro*. In particular, the growth of the tumor mass is investigated, including necrotic tumor cells and the consumption of nutrient (oxygen) over time. The geometry of the problem and the boundary conditions are described in [Fig. 3](#). At the boundary B1 the TC volume fraction (ϕ), the oxygen mass fraction (ω) and the necrotic cells mass fraction (ϕ_n) are fixed over time. At the symmetry boundaries B2 zero fluxes for all phases are imposed. At the beginning of the simulation the value of the oxygen and of the other variables are fixed, as indicated in the figure. All other governing parameters are listed in [Table 1](#). Numerical results for the evolution of the three principal variables of the model are shown in [Fig. 4](#). In [Fig. 4.a](#), the volume fraction of tumor cells in the system is plotted for different times over the spheroid radius. Initially, the tumor spheroid is composed only by living tumor cells and interstitial fluid. After a few days, necrotic tumor cells appear in the center of the spheroid. Living tumor cells are restricted to the outer portion of the tumor, where there is still enough nutrient to support their growth. At 25 days, necrotic cells occupy the main portion of the spheroid, constituting the “necrotic core”, while the proliferating portion of the tumor is further reduced. Since tumor growth is described by equations 36 and 44, only tumor cells that experience an oxygen level over the critical threshold are allowed to proliferate. This means that the living tumor cells that can actually proliferate are distributed only over a small portion of the radius near the external boundary, called the “proliferative rim”. The remaining portion of living cells are non-proliferating cells that can resume proliferation after an increase in the level of nutrient. Note that the volume fraction of interstitial fluid is approximately constant over time, apart for a small increase at the spheroid center and reduction at the spheroid periphery. These are due, respectively, to the lysis term active at the tumor center and to the fluid consumption induced by cell growth at the boundary.

The evolution of the oxygen mass fraction, the nutrient species considered here, is shown in [Fig. 4.b](#). Each line is plotted over the spheroid radius, every 5 days from the first day. As the spheroid grows, gradients of oxygen concentration develop from the tumor boundary to the center of the spheroid. After a few days the oxygen mass fraction reaches a plateau at the center of the spheroid, with a value below the critical threshold. This can be explained by considering that necrotic cells do not consume oxygen and therefore a gradient is not observed. Moreover, the oxygen consumption is proportional to the amount of oxygen available through equation 42, and this contributes to the shape of the curves.

The evolution of the necrotic mass fraction is presented in the graph of [Fig. 4.c](#). The necrotic portion of the tumor develops from the center towards the boundary. As the interface between living tumor cells and necrotic tumor cells is diffused, the separation between the two is smooth and the model can account for perinecrotic regions.

4.2 Spheroids grown with different initial cell seeding number

The computational model is validated against data from tumor spheroid cultures. U87-MG cells, a human glioblastoma cell line, are cultured with a standard protocol (see materials and methods). The experimental set up is shown in [Fig. 5](#). The bottom of standard cell culture wells is covered with agarose, in order to prevent tumor cells adhesion. Cells are seeded at different initial numbers (1000, 5000, 10000) and rapidly form spheroids suspended in standard culture medium. The evolution of the spheroid radii are then recorded over time via optical microscopy and the resulting growth curves are plotted in [Fig. 6.a](#). It is possible to distinguish between the first stages of growth, characterized by an exponential/linear behavior, followed by a phase of growth saturation where the radius reaches a steady value.

After recording the curves, a series of simulations is run to reproduce the experimental data. The growth curves corresponding to the best fit of the experimental data are shown in [Fig. 6.b](#). The governing parameters are taken from literature when they are available and we use the same order of magnitude of the experimental values when they refer to other cell species. [Table 1](#) lists the parameters used in this study together with the ones determined by the fit of the curves. There is a good agreement with the experimental data, for all the three different

initial cell seeding number. The model captures the growth dynamics both in the first fast-growing phase and in the later phase of growth saturation. Note that the same parameters are used for fitting all the three curves and only the initial radii of the spheroids are changed, showing a good quality of the fit. This experiment is useful for validating the model and displays a second remarkable result. The different initial seeding of tumor cells affects the initial radius of the tumor spheroid, which increases from $\approx 100 \mu\text{m}$ to almost $190 \mu\text{m}$. The data show that, although being constituted by a larger initial number of tumor cells, bigger spheroids reach the same final radius of smaller ones. This result agrees with what is reported in literature about the existence of a steady radius for growing spheroids ([Sutherland et al. 1971](#); [Folkman and Hochberg 1973](#); [Carlsson 1977](#); [Freyer and Sutherland 1986](#)), which in our case takes approximately the value of $475 \mu\text{m}$.

Interestingly, the model reproduces the same behavior of the experiments with the steady state being reached after 25 days from cell seeding. As the spheroids grow freely in the culture medium, the only mechanism to stop cell proliferation is given by equation 44 in the model where it is assumed that cell mitosis and necrosis depend on the local level of nutrient. Therefore, in this modeling framework the hypothesis of nutrient deprivation is sufficient to explain the phenomenon of growth saturation and the existence of an asymptotic radius for the spheroid.

As a further remark about these results, in [Fig. 6.c](#) the previous curves are shifted of the proper amount referring to the different initial condition. It is good to see that they coalesce to a single “master curve”. Spheroids grown from a different initial radius follow the same curve, showing that there exist a common dynamics regulating the growth of these cellular aggregates.

4.2 Compression experiments

The application of a constant mechanical stress on the surface of the growing spheroids is investigated in this series of experiments. Addition of Dextran to the cell culture medium produces an osmotic pressure on the outermost layer of cells located on the spheroid surface. The osmotic pressure acts as a network stress directed to decrease the volume occupied by the spheroid ([Montel et al. 2011](#); [Montel et al. 2012](#); [Delarue et al. 2014](#)). This compressive force

can be calibrated experimentally as in ([Bonnet-Gonnet et al. 1994](#); [Bouchoux et al. 2009](#)), where an empirical law is given to relate the concentration of Dextran in the solution to the external pressure exerted on the surface of the spheroid. Three pressure conditions are explored following the approach described in the Materials and Methods section, namely 1kPa, 5kPa and 10kPa, plus a control experiment with no external pressure. Cell viability is checked as reported in Section 3.2, where it is shown that the addition of Dextran does not alter cell death or growth. The results are presented in [Fig. 7](#), where the growth of the spheroids is followed for 18 days after the addition of Dextran. [Fig. 7.a](#) shows optical images of sample spheroids referring to the control and to the most compressed condition for different time instants. Starting from a similar initial radius (about 200 μm) the two spheroids reach considerable different volumes at 18 days, with the compressed spheroid growing slowly compared to the stress free case. The growing curves for the other external applied pressures are collected in [Fig. 7.b](#). The larger value of the radius is reached for the spheroids grown in the absence of any external compression. When Dextran is added to the medium and the osmotic stress builds up, both the growth rate and the final diameter decrease. If the applied pressure is released the growth of the aggregates resumes, indicating that the effect of the stress is reversible ([Online Resource 1](#)), as shown for the first time in ([Helmlinger et al. 1997](#)). The effect of the external pressure on the growth of the spheroid is included in the model through equation 46, which describes the inhibition of cell proliferation due to the applied mechanical stress on the tumor cells. The most common mathematical expressions for the inhibiting function reported in literature are based on a linear or inversely proportional assumptions ([Roose et al. 2003](#); [Byrne and Preziosi 2003](#); [Kim et al. 2011](#); [Mpekris et al. 2015](#)). In [Fig. 8](#) these forms are tested against the experimental data, together with an exponential relation and a Michaelis–Menten-like expression. The data in [Table 2](#) represent the values of the parameters that provide the best fits to the experimental curves. The linear relationship, applied in [Fig. 8.a](#), underestimates the inhibition effect for low compressions, leading to larger values of the radius for the 1kPa and 5kPa cases. At the same time, for larger compressions the linear relationship gives overestimates for the inhibition, resulting in smaller radii for the 10kPa curve than experimentally measured. The exponential relationship is applied in [Fig. 8.b](#), where it is possible

to observe an improvement for high compressions but a similar underestimation for the 1kPa and 5kPa curves. Another improvement can be seen in [Fig. 8.c](#), referring to the inversely proportional expression. In this case there is a good agreement with the experimental data for all the curves, except for the 1kPa case. The best results are obtained with the Michaelis-Menten-like law, represented in [Fig. 8.d](#). All the different compression levels are well described by the simulations, together with the final radii reached by the spheroids.

4.3 Effect of the growth inhibition parameters

The effect of an external stress acting on the cellular component of the spheroid can be evaluated through a parametric study on the growth inhibition parameters and of equation 46. The growth curves obtained by varying one of the two parameters and keeping the other fixed for the case of an external pressure of 5kPa are presented in [Fig. 9](#). The two solid lines represent reference values assumed by the parameters. In particular, “zero” indicates the curve obtained by setting both and to zero, while “fit” is the curve obtained with the values in [Table 2](#). The effect of different values for is shown in [Fig. 9.a](#). The arrow points in the direction of increasing, while for the value in [Table 2](#) is selected for all the simulations. From the top curve to the bottom one we consider values for that are, respectively, -50%, -25%, +25%, +50% and +75% of the reference value in [Table 2](#). As increases, tumor cells sense more the effect of external stresses on their proliferation. This results in smaller final radii of the spheroids and in slower growth rates. If the value of is sufficiently high then the tumor starts to shrink and an equilibrium radius no longer exists. The same investigation, this time for , is reported in [Fig. 9.b](#). In this case the arrow points in the opposite direction, indicating that an increase in leads to an increase of the final radius. This difference can be easily explained by considering that in equation 46 is directly proportional to and inversely proportional to respectively. However, it is possible to note that the effect of varying is less pronounced on the growth curve, since the reference value in [Table 2](#) is much smaller than the pressures in the compression experiments. These results indicate collectively a double effect of the external environment in limiting the growth of the tumor. Tumor growth may be hindered by nutrient deprivation but also by external stresses exerted by regions close to the tumor.

5. Discussion

In the present study a recent model for tumor growth has been extended to describe the evolution of tumor spheroids. A set of experiments is carried out and the resulting data are compared to numerical predictions. The experimental growth curves validate the model equations both for the free-growth case, where the cells are cultured in three dimensions in standard culture medium, and for the mechanically compressed setup, where the spheroids are subjected to an external pressure. In addition to providing the data for the validation, the first series of experiments highlights the existence of a “master curve” (Fig. 6.c). This common growth trend supports the hypothesis of describing the cell aggregates as a dynamical system, which behavior can be predicted, at least as a first approximation, by the laws of mechanics. The results of the model in terms of tumor volume fraction, oxygen mass fraction and necrotic mass fraction are reported and appear in line with the results of the original model. The second set of experiments about spheroids compression extends the work in (Delarue et al. 2014) by adding another cell species (U87-MG) to their study. The observed evolution curves are similar to their findings and confirm the hypothesis of an inhibitory effect of external stress on cancer cell proliferation. Regarding the description of this phenomenon, the experimental data are exploited to design a constitutive relationship that performs better, compared to the existing laws, in describing the evolution of the system.

Several simplifying assumptions are considered in the work and the model is certainly open to further improvements. In particular, here only one nutrient species, namely oxygen, diffuses in the interstitial fluid and regulates the proliferation of tumor cells. Although the action of other chemicals is implicitly included in the mass transfer term in 36, modeling additional nutrients, growth and necrosis factors could provide supplementary insights in the evolution of the tumor system (Chauhan and Jain 2013; Jain et al. 2014). Another point that should be addressed is the choice of the constitutive relations used to close the differential system. Most of these laws, as it happens frequently in literature, are derived from phenomenological arguments and deserve more experimental work to be linked to the biology of what they are describing. **In particular, here is assumed that the compression of the spheroids induces inhibition of cell growth,**

without modifying the apoptosis rate of the cells. This hypothesis is still a matter of debate in literature (see for example (Montel et al. 2012) and comments therein), and it is adopted here to account for the experimental observations of Delarue et al. (2014) where the experimental setup is similar to the one in this work. A systematic comparison of different compression modalities may improve the understanding of this phenomenon and the design of more accurate constitutive laws.

Finally, here it is considered a very simple mechanical description of the tumor ensemble, function of the volume fraction of the tumor cells. This assumption provides a great simplification of the mechanical equations and describes sufficiently accurately the data, however it does not take into account several phenomena related to the stress experienced by the cells inside the tumor tissue. For example, viscous effects existing at smaller time-scales than cell proliferation are neglected, as well as cellular adhesion bonds breakage and formation during the development of the tumor mass (Ambrosi and Preziosi 2009; Preziosi and Vitale 2011).

Further experiments will be considered in the future, that will provide better estimates for the model parameters and new data in terms of quantities that should be compared to the output of the model equations. A part of the future experimental work will be also devoted to the biochemical understanding of the growth inhibition process due to mechanical stress. Even if some work is already present in literature (Delarue et al. 2014; Fernández-Sánchez et al. 2015), many details remain obscure as well as a proper implementation of the phenomena in the growth equations. A better description of the interactions between the tumor and its external microenvironment (biochemical and mechanical) should offer valuable insights for understanding the progression of the disease and designing new therapeutic treatments.

References

- Ambrosi D, Mollica F (2004) The role of stress in the growth of a multicell spheroid. *J Math Biol* 48:477–499.
- Ambrosi D, Preziosi L (2009) Cell adhesion mechanisms and stress relaxation in the mechanics of tumours. *Biomech Model Mechanobiol* 8:397–413.
- Ambrosi D, Preziosi L, Vitale G (2012) The interplay between stress and growth in solid tumors. *Mech Res Commun* 42:87–91.
- Baumgartner W, Hinterdorfer P, Ness W, et al (2000) Cadherin interaction probed by atomic force microscopy. *Proc Natl Acad Sci U S A* 97:4005–4010.
- Bonnet-Gonnet C, Belloni L, Cabane B (1994) Osmotic pressure of latex dispersions. *Langmuir* 10:4012–4021.
- Bouchoux A, Cayemite PE, Jardin J, et al (2009) Casein micelle dispersions under osmotic stress. *Biophys J* 96:693–706.
- Byrne H, Preziosi L (2003) Modelling solid tumour growth using the theory of mixtures. *Math Med Biol* 20:341–366.
- Carlsson J (1977) A proliferation gradient in three-dimensional colonies of cultured human glioma cells. *Int J Cancer* 20:129–136.
- Carlsson J, Yuhas JM (1984) Liquid-overlay culture of cellular spheroids. *Recent Results Cancer Res* 95:1–23.
- Casciari JJ, Sotirchos S V, Sutherland RM (1992a) Mathematical modelling of microenvironment and growth in EMT6/Ro multicellular tumour spheroids. *Cell Prolif* 25:1–22.
- Casciari JJ, Sotirchos S V, Sutherland RM (1992b) Variations in tumor cell growth rates and metabolism with oxygen concentration, glucose concentration, and extracellular pH. *J Cell Physiol* 151:386–94.
- Chaplain MAJ, Graziano L, Preziosi L (2006) Mathematical modelling of the loss of tissue compression responsiveness and its role in solid tumour development. *Math Med Biol* 23:197–229.
- Chauhan VP, Jain RK (2013) Strategies for advancing cancer nanomedicine. *Nat Mater* 12:958–62.
- Cheng G, Tse J, Jain RK, Munn LL (2009) Micro-environmental mechanical stress controls tumor spheroid size and morphology by suppressing proliferation and inducing apoptosis in cancer cells. *PLoS One* 4:e4632.
- Ciarletta P, Ambrosi D, Maugin GA, Preziosi L (2013a) Mechano-transduction in tumour growth modelling. *Eur Phys J E* 36:23.

- Ciarletta P, Preziosi L, Maugin G a. (2013b) Mechanobiology of interfacial growth. *J Mech Phys Solids* 61:852–872.
- Delarue M, Montel F, Vignjevic D, et al (2014) Compressive stress inhibits proliferation in tumor spheroids through a volume limitation. *Biophys J* 107:1821–1828.
- Desmaison A, Frongia C, Grenier K, et al (2013) Mechanical stress impairs mitosis progression in multi-cellular tumor spheroids. *PLoS One* 8:e80447.
- Dorie MJ, Kallman RF, Rapacchietta DF, et al (1982) Migration and internalization of cells and polystyrene microsphere in tumor cell spheroids. *Exp Cell Res* 141:201–209.
- Ehlers W, Markert B, Rohrlé O (2009) Computational continuum biomechanics with application to swelling media and growth phenomena. *GAMM-Mitteilungen* 32:135–156. doi: 10.1002/gamm.200910013
- Fernández-Sánchez ME, Barbier S, Whitehead J, et al (2015) Mechanical induction of the tumorigenic β -catenin pathway by tumour growth pressure. *Nature* 523:92–95.
- Folkman J, Hochberg M (1973) Self-regulation of growth in three dimensions. *J Exp Med* 138:745–753.
- Freyer JP, Sutherland RM (1986) Regulation of growth saturation and development of necrosis in EMT6/Ro multicellular spheroids by the glucose and oxygen supply. *Cancer Res* 46:3504–3512.
- Friedrichs J, Legate KR, Schubert R, et al (2013) A practical guide to quantify cell adhesion using single-cell force spectroscopy. *Methods* 60:169–78.
- Galle J, Preziosi L, Tosin a. (2009) Contact inhibition of growth described using a multiphase model and an individual cell based model. *Appl Math Lett* 22:1483–1490.
- Giverso C, Scianna M, Grillo A (2015) Growing avascular tumours as elasto-plastic bodies by the theory of evolving natural configurations. *Mech Res Commun* 68:31–39.
- Grantab R, Sivananthan S, Tannock IF (2006) The penetration of anticancer drugs through tumor tissue as a function of cellular adhesion and packing density of tumor cells. *Cancer Res* 66:1033–9.
- Gray WG, Miller CT (2005) Thermodynamically constrained averaging theory approach for modeling flow and transport phenomena in porous medium systems: 1. Motivation and overview. *Adv Water Resour* 28:161–180.
- Gray WG, Miller CT (2014) Introduction to the thermodynamically constrained averaging theory for porous medium systems, 1st edn. Springer International Publishing
- Greenspan HP (1976) On the growth and stability of cell cultures and solid tumors. *J Theor Biol* 56:229–42.

- Helenius J, Heisenberg C-P, Gaub HE, Muller DJ (2008) Single-cell force spectroscopy. *J Cell Sci* 121:1785–91.
- Helmlinger G, Netti PA, Lichtenbeld HC, et al (1997) Solid stress inhibits the growth of multicellular tumor spheroids. *Nat Biotechnol* 15:778–783.
- Jain RK (1988) Determinants of tumor blood flow: A review. *Cancer Res.* 48:2641–2658.
- Jain RK, Martin JD, Stylianopoulos T (2014) The role of mechanical forces in tumor growth and therapy. *Annu Rev Biomed Eng* 16:321–46.
- Jain RK, Stylianopoulos T (2010) Delivering nanomedicine to solid tumors. *Nat Rev Clin Oncol* 7:653–64.
- Kaufman LJ, Brangwynne CP, Kasza KE, et al (2005) Glioma expansion in Collagen I matrices: analyzing collagen concentration-dependent growth and motility patterns. *Biophys J* 89:635–650.
- Kim T-H, Mount CW, Gombotz WR, Pun SH (2010) The delivery of doxorubicin to 3-D multicellular spheroids and tumors in a murine xenograft model using tumor-penetrating triblock polymeric micelles. *Biomaterials* 31:7386–97.
- Kim Y, Stolarska M a., Othmer HG (2011) The role of the microenvironment in tumor growth and invasion. *Prog Biophys Mol Biol* 106:353–379.
- Leder K, Pitter K, Laplant Q, et al (2014) Mathematical modeling of PDGF-driven glioblastoma reveals optimized radiation dosing schedules. *Cell* 156:603–16.
- Lewis RW, Schrefler BA (1998) The finite element method in the static and dynamic deformation and consolidation of porous media. John Wiley
- Longo D, Fauci A, Kasper D, Hauser S (2011) Harrison’s principles of internal medicine. McGraw-Hill Professional
- Lowengrub JS, Frieboes HB, Jin F, et al (2010) Nonlinear modelling of cancer: bridging the gap between cells and tumours. *Nonlinearity* 23:R1–R91.
- Michor F, Liphardt J, Ferrari M, Widom J (2011) What does physics have to do with cancer? *Nat Rev Cancer* 11:657–70.
- Mikhail AS, Eetezadi S, Allen C (2013) Multicellular tumor spheroids for evaluation of cytotoxicity and tumor growth inhibitory effects of nanomedicines in vitro: A comparison of Docetaxel-loaded block copolymer micelles and Taxotere®. *PLoS One* 8:e62630.

- Montel F, Delarue M, Elgeti J, et al (2011) Stress clamp experiments on multicellular tumor spheroids. *Phys Rev Lett* 107:188102.
- Montel F, Delarue M, Elgeti J, et al (2012) Isotropic stress reduces cell proliferation in tumor spheroids. *New J Phys* 14:055008.
- Mpekris F, Angeli S, Pirentis AP, Stylianopoulos T (2015) Stress-mediated progression of solid tumors: effect of mechanical stress on tissue oxygenation, cancer cell proliferation, and drug delivery. *Biomech Model Mechanobiol* 14:1391–1402.
- Pinder GF, Gray WG (2008) *Essentials of multiphase flow in porous media*, 1st edn. Wiley-Interscience
- Preziosi L, Tosin A (2009a) Multiphase and multiscale trends in cancer modelling. *Math Model Nat Phenom* 4:1–11.
- Preziosi L, Tosin A (2009b) Multiphase modelling of tumour growth and extracellular matrix interaction: Mathematical tools and applications. *J Math Biol* 58:625–656.
- Preziosi L, Vitale G (2011) A multiphase model of tumor and tissue growth including cell adhesion and plastic reorganization. *Math Model Methods Appl Sci* 21:1901–1932.
- Puech P-H, Taubenberger A, Ulrich F, et al (2005) Measuring cell adhesion forces of primary gastrulating cells from zebrafish using atomic force microscopy. *J Cell Sci* 118:4199–206.
- Roose T, Chapman SJ, Maini PK (2007) Mathematical models of avascular tumor growth. *SIAM Rev* 49:179–208.
- Roose T, Netti P a., Munn LL, et al (2003) Solid stress generated by spheroid growth estimated using a linear poroelasticity model. *Microvasc Res* 66:204–212.
- Sciumè G, Gray WG, Hussain F, et al (2013a) Three phase flow dynamics in tumor growth. *Comput Mech* 53:465–484.
- Sciumè G, Santagiuliana R, Ferrari M, et al (2014) A tumor growth model with deformable ECM. *Phys Biol* 11:65004.
- Sciumè G, Shelton S, Gray WG, et al (2013b) A multiphase model for three-dimensional tumor growth. *New J Phys* 15:015005.
- Stigliano C, Key J, Ramirez M, et al (2015) Radiolabeled polymeric nanoconstructs loaded with Docetaxel and Curcumin for cancer combinatorial therapy and nuclear imaging. *Adv Funct Mater* 25:3371–3379.
- Sutherland R, Carlsson J, Durand R, Yuh J (1981) Spheroids in Cancer Research. *Cancer Res* 41:2980–2984.

Sutherland RM, McCredie J a, Inch WR (1971) Growth of multicell spheroids in tissue culture as a model of nodular carcinomas. *J Natl Cancer Inst* 46:113–120.

Vinci M, Gowan S, Boxall F, et al (2012) Advances in establishment and analysis of three-dimensional tumor spheroid-based functional assays for target validation and drug evaluation. *BMC Biol* 10:29.

Wise SM, Lowengrub JS, Frieboes HB, Cristini V (2008) Three-dimensional multispecies nonlinear tumor growth-I. Model and numerical method. *J Theor Biol* 253:524–543.

FIGURE CAPTIONS:

Fig. 1 The constituents of the biphasic system

Fig. 2 Scheme of the stress function. The two insets represent forces acting on the cells for different degrees of tumor volume fractions. For high volume fractions, on the right, cells experience a repulsive force. When the volume fractions are low, on the left, cells tend to attract each other

Fig. 3 Scheme for the geometry of a tumor spheroid immersed in a cell culture medium. The initial and boundary conditions for the differential problem are reported on the right

Fig. 4 a Volume fraction of tumor cells, plotted as the difference between the volume fraction inside the spheroid and at the spheroid boundary (here since the spheroids are not compressed). **b** Mass fraction of oxygen. **c** Mass fraction of necrotic tumor cells. All the variables are plotted over the spheroid radius with each line drawn every 5 days from the first

Fig. 5 Optical images of the growth of U-87 MG spheroids from day 5 to day 21. The scalebar is 200 μm and the spheroids are taken from the 10000 seeded cells initial condition. The last image represents a scheme of the culture protocol

Fig. 6 Growth curves recorded from the free growth experiments. **a** Each curve represents a different initial condition in terms of seeded cells. $N \geq 4$ spheroids are considered for each condition. Points are experimental data and error bars are the standard deviations of the measurements. **b** Solid lines are the results of fits with the mathematical model. **c** Growth curve obtained by superimposing the evolution of the radii of spheroids grown at different initial cell seeding numbers

Fig. 7 a Optical images of U-87 MG spheroids grown under the effect of the Dextran solutions. The first row shows the control experiments and the second row a spheroid under the highest compression. The scalebar is $200 \mu\text{m}$ and the initial seeding is 5000 tumor cells. **b** Experimental results for the compression experiments. The points are the experimental data and the error bars represent the standard deviations of the measurements. For each condition $N = 5$ spheroids are considered

Fig. 8 Fit of the experimental data for the compression experiments. Results from the linear **a**, exponential **b**, inversely proportional **c** and Michaelis-Menten-like **d** assumptions for the function in equation 46

Fig. 9 Parametric study for the growth inhibition parameters in equation 46. Effects of **a** and of **b** on the growth of the spheroids