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This is the author's manuscript

Original Citation:

Availability:
This version is available http://hdl.handle.net/2318/40535 since

Published version:
DOI:10.1088/1478-3975/4/4/P01

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PERSPECTIVE

Physical aspects of cancer invasion

Caterina Guiot1,2, Nicola Pugno3, Pier Paolo Delsanto2,4,5 and Thomas S Deisboeck6

1 Department of Neuroscience, University of Torino, Torino, Italy
2 CNISM, Torino, Italy
3 Department of Structural Engineering and Geotechnics, Politecnico of Torino, Torino, Italy
4 Department of Physics, Politecnico of Torino, Torino, Italy
5 Bioinformatics and High Performance Computing Labs, Bioindustry Park of Canavese, Colleretto Giacosa, Italy
6 Complex Biosystems Modeling Laboratory, Harvard-MIT (HST) Athinoula A. Martinos Center for Biomedical Imaging, Massachusetts General Hospital, Charlestown, MA 02129, USA

E-mail: deisboec@helix.mgh.harvard.edu

Received 3 August 2007
Accepted for publication 29 October 2007
Published 17 December 2007
Online at stacks.iop.org/PhysBio/4/P1

Abstract

Invasiveness, one of the hallmarks of tumor progression, represents the tumor’s ability to expand into the host tissue by means of several complex biochemical and biomechanical processes. Since certain aspects of the problem present a striking resemblance with well-known physical mechanisms, such as the mechanical insertion of a solid inclusion in an elastic material specimen (G Eaves 1973 The invasive growth of malignant tumours as a purely mechanical process J. Pathol. 109 233; C Guiot, N Pugno and P P Delsanto 2006 Elastomechanical model of tumor invasion Appl. Phys. Lett. 89 233901) or a water drop impinging on a surface (C Guiot, P P Delsanto and T S Deisboeck 2007 Morphological instability and cancer invasion: a ‘splashing water drop’ analogy Theor. Biol. Med. Model 4 4), we propose here an analogy between these physical processes and a cancer system’s invasive branching into the surrounding tissue. Accounting for its solid and viscous properties, we then arrive, as a unifying model, to an analogy with a granular solid. While our model has been explicitly formulated for multicellular tumor spheroids in vitro, it should also contribute to a better understanding of tumor invasion in vivo.

1. Introduction

Due to the complexity of the mechanisms involved in neoplastic growth, in silico experiments are becoming the tool of choice [4] for the description and interpretation of the observed phenomena, based on models ranging from microscopic to macroscopic. Microscopic models are suitable to describe malignant transformations, which lead to profound alterations at the sub-cellular and cellular levels. Macroscopic models of solid tumor growth may be related to universal scaling laws [5], in analogy with the model of ontogenic growth for all living organisms proposed by G B West and collaborators [6, 7]. A similar quest for universalities is currently being pursued in a completely different scientific context (continuum mechanics and elasticity) [8] and extended to other fields as well [9]. In all cases, a bridging between a microscopic and a macroscopic description is of fundamental importance [10] and can be best achieved by means of intermediate, so-called mesoscopic models (see [11] and references therein).

Tumor invasion involves a variety of biochemical and biomechanical processes that ultimately lead to cell detachment from the primary tumor, attachment to extracellular matrix (ECM), infiltration into adjacent tissue and, in some instances, to dissemination via bloodstream or lymphatic system en route to seeding secondary satellites or
show that, in a nutrient-poor microenvironment, the only
unifying features that should be noted. There are, however, some
complex experimental background. There are, however, some
endogenous substances (integrins, focal adhesion kinases)
produced by the tumor itself [23], how the ECM is remodeled
host' interface is now available. Hence, one can learn how
about the biochemical mechanisms occurring at the 'tumor–
but that growth instabilities are also required at the interface
with the host. In fact, a remarkable amount of information
on matrix homogeneity and
invasive mass (defined as 'podosomes', or adhesive) and the
invasive pattern.

Several theoretical models of tumor invasion have been
proposed, either describing the tumor–host interface as
a traveling wave edge [17], through extracellular matrix
deradation [18], or for specific applications to invasive
gliomas [19]. The role of tensions of homeostasis on malignant
growth has also been investigated at a microscopic level [20].
Recent observations of invasive branching (or fingering) in
multicellular tumor spheroids (MTS) [21, 22] suggest that
invasion cannot be solely caused by tumor cell proliferation,
but that growth instabilities are also required at the interface
with the host. In fact, a remarkable amount of information
about the biochemical mechanisms occurring at the 'tumor–
host' interface is now available. Hence, one can learn how
the local host stroma is affected by the degrading enzymes
produced by the tumor itself [23], how the ECM is remodelled
by endogenous substances (integrins, focal adhesion kinases)
[24] and how all these molecular mechanisms cross-interact
[25]. No simple model can account for such a detailed and
complex experimental background. There are, however, some
unifying features that should be noted.

(1) Short-range processes, i.e. the enzyme cascade is confined
to the cell surface of the invading pseudopodia [26].
(2) Borderline processes have important implications also
for nutrient availability: the appearance of sprouts and
invasive branches affects the tumor's surface-to-volume
ratio. The fractal dimension (and other related parameters,
such as the scaling parameter of the 'West-like' growth
law [2, 27]) are therefore increased with respect to
2/3, which would correspond to diffusion across a
spherical surface. Similar patterns have been observed in
bacterial colonies cultured in more rigid media (i.e. a high
concentration of agar) and poor nutrient concentration
[28].

2. Why an amorphous 'solid–fluid' model?

Two numerical simulation models have been recently proposed
to describe tumor invasion, assuming that the role of the
microenvironment is as crucial as that of the tumor itself.
Both of them are consistent with the well-known existence
of the two main invasive patterns [29]: the 'smooth margin'
invasive mass (defined as 'podosomes', or adhesive) and the
'fingering' (or invadopodia, i.e. protrusive pattern). They also
explain the invasion dependence on the microenvironment
characteristics, i.e. on matrix homogeneity and/or nutrient
availability. In particular, Macklin and Lowengrub [30]
show that, in a nutrient-poor microenvironment, the only
viable invasion strategy for the tumor is to break into small
fragments which penetrate the surrounding tissue. Conversely,
under nutrient-rich conditions, the tumor behavior depends
mostly on the biomechanical property of the medium: a soft,
mechanically responsive medium enhances 'smooth margin'
invasion while, in the opposite situation, long invasive fingers
are produced, whose characteristics largely depend on cell-
to-cell adhesion. Similarly, Anderson et al [31] found that a
homogeneous matrix elicits the smooth margin invasive type,
while a heterogeneous matrix ('bumpy' or 'grainy', depending
on the macro or cell–level scale of the inhomogeneities) favors
fingering. Also, in their model, the tumor oxygenation level,
together with the cell-to-cell adhesion value, impacts the
invasive pattern.

From a physical point of view, while the case of a
homogeneous, elastic medium is somehow trivial (and already
investigated in several papers, see for instance [3]), it is
intriguing to represent the microenvironment as a 'grainy'
medium (see [31]). Also, the mechanically 'unresponsive'
medium considered by the same authors (and promoting tumor
fingering) is more comparable to a viscous fluid.

Here, we attempt to reconcile the solid and the fluid
mechanical models [2, 3], starting from the fact that, in many
ways, an amorphous solid can behave as a fluid and vice
versa. Thus, fluid- and solid-like behaviors often coexist.
For instance, it is well known that the earth's crust, during
asteroid or meteorite impacts, displays a fluid-like behavior.
In particular, the classical bowl-like crater often shows a central
 pinnacle, due not simply to elastic recoil or melting, but as a
consequence of drop-like reflux [32]. Conversely, a fluid under
strong shear stress behaves as a solid, a fact which is used to
smartly increase the protective capability of 'liquid' armors,
i.e. solid armors impregnated by colloidal suspensions [33].
An attempt to treat solids and liquids in a unified manner is
represented by the theory of granular solids (or, conversely,
of viscous fluids). In fact, such systems behave as an amorphous
solid if their grains are confined, but as a viscous fluid if they
are not [34]. Accordingly, the model formulated in the present
paper represents a first step in the direction of treating a tumor
as a granular solid. A granular material is by definition a
conglomerate of discrete macroscopic particles. If they are
non-cohesive, then the forces between them are only repulsive
so that the shape of the material is determined by external
boundaries and gravity. Despite this seeming simplicity, a
granular material behaves differently from any of the other
familiar forms of matter solids, liquids or even gases, and
should therefore be considered as an additional state of matter
[35].

We note that the two main cancer invasive mechanisms,
namely 'smooth margin' invasive mass and 'fingering',
resemble a progressive damage growth in comminuted solids
or a drop splashing in liquids. Thus, even if this analogy is
crude, we believe that considering a granular material can
give better prediction with respect to the used and abused
hypothesis of the continuum. Accordingly, we use quantized
fracture mechanics ([36–38], see appendix A) in order to arrive
at an asymptotic matching between the predictions of the
number of invasive branches for a solid (derived according to
classical linear elastic fracture mechanics [39]) and for a liquid (derived according to the splashing water drop analogy [3], and see appendix B). Note that fracture mechanics have already been used to model tumor invasion, suggesting, despite the biological details, that a tumor invasion resembles a growing inclusion in a fracturable matrix (the biological details are included in this model by considering opptiune material properties) [2].

It is remarkable that a unique, simple relationship between few, measurable physical variables can predict the occurrence of a fingering pattern (the computational details are given in appendices A and B, for the solid and the liquid analogies, respectively). Following equations (A.6) and (B.4), the value $N_f = 1$ separates the case of no branching (hence no tumor invasion) from the one in which invasion takes place. By defining the dimensionless invasion parameter,

$$IP = PR/\sigma,$$

invasive behavior is expected in all cases but for $IP < 1$ (which implies large tumor surface tension, small confining pressure and/or small tumor radius values). According to equation (1), which can be derived with both the ‘mechanistic’ and the ‘viscous fluid’ models, tumor invasion is controlled by three parameters given below.

2.1. Tumor surface tension

Multicellular aggregates of three different malignant astrocytoma cell lines (U-87MG, LN-229 and U-118MG) investigated by Winters et al [40] with surface tensions of about 7, 10 and 16 dyne cm$^{-1}$, respectively, showed a significant inverse correlation between invasiveness and surface tension. Moreover, these authors showed that the anti-edematous therapeutic agent Dexamethason increases tumor surface tension or cohesiveness between cells; accordingly, Dexamethason would have anti-invasive effects as well. Therefore, surface tension can indeed be a predictor of in vitro invasiveness, with a threshold value for $\sigma$ of about 10 dyne cm$^{-1}$.

2.2. Microenvironmental pressure

The mechanical interaction between matrix and tumor has been recently investigated more thoroughly. For instance, Paszek et al [41] observed that stiffer tissues promote malignant behavior. Similarly, the experimental work on NIH3T3 fibroblasts by Georges and Jannmey [42] shows that they keep a roughly spherical shape (suggesting prevalence of cohesive forces) when embedded in a soft polyacrilamide gel, while in a stiff gel they exhibit finger-like patterns (consistent with the preponderance of adhesive forces). We note that the effect of external pressure on the growth of tumor cell colonies has also been studied by Bru and Casero [43], showing that geometrical and dynamical patterns are markedly dependent on the pressure exerted by the surrounding medium. Moreover, pressure can act either as an inhibitor or as an enhancer for tumor cell proliferation, depending on the particular cell line (see e.g. [44]).

2.3. Tumor radius

Finally, Tamaki et al [45] investigated C6 astrocytoma spheroids with different diameters (i.e., 370, 535 and 855 µm on average), which were implanted in collagen type I gels. The authors showed that spheroid size indeed correlated with a larger total invasion distance and increased rate of invasion.

Taken together, any therapeutic strategies that solely or in combination are geared toward reducing tumor burden, diminishing surrounding mechanical pressure and increasing (residual) tumor surface tension, may eventually hold promise in clinics.

3. Conclusions

In this contribution we have reviewed two models, based on analogies with well-known mechanisms of fracture mechanics and fluid dynamics, which have been recently proposed to illustrate some of the features of tumor invasion. In the former, tumors are visualized as amorphous solids, while in the latter as viscous fluids. We have attempted, in this paper, to reconcile the two representations by means of an intermediate one, i.e., the ‘granular’ solid model. Remarkably, the most useful result that we have obtained, i.e. the formula for the so-called invasion parameter, is consistent with both models (and, of course, with the intermediate one). Such a formula identifies the most relevant physical parameters, whose control should be the target of dedicated therapies, e.g. the tumor’s surface tension, its radius and the confining host tissue pressure. Understanding their role could explain why some therapies fail while others prove to be effective in locally controlling tumor expansion. While a reliable, patient-specific assessment of tissue properties poses a formidable challenge, in principle one should be able to predict whether a particular tumor type in a given host organ exhibits finger-like invasion patterns or not. Eventually, a cancer type, organ site and patient-specific invasion parameter, $IP$, may be of significant value for diagnostic purposes, as most of this multicellular behavior occurs well below the current non-invasive imaging resolution limits. However, for any such future iteration, a more realistic description should obviously take into account the heterogeneity of both tumor and microenvironment, which would not only imply regional differences in the $IP$ value but also argue for a dynamic behavior of $IP$. Similarly, the schematic ‘sequence’ suggested in figure 1 would then refer to a single site rather than to the entire tumor, as invasive branching would become desynchronized at numerous sites across the tumor surface.

To conclude, we believe that, although an all-encompassing model of cancer invasion (or, in general, tumor growth) is desirable, for the time being it may be more expedient to use ‘composite’ models in which the different facets of the problem are considered individually, not as alternative but as complementary descriptions. Likewise, for the numerical simulation of neoplastic growth (both for diagnostic and therapeutic purposes), a multilevel approach [46] may be most promising, i.e. one that includes both microscopic and macroscopic scales and its mesoscopic bridging level.
Figure 1. Schematic representation of an invasion sequence or ‘cycle’. (a) Initial condition: tumor (black) in an elastic matrix (gray). (b) Non-invasive phase: the interfacial stress increases in the course of the tumor’s elastic growth and interaction with the matrix. (c) Invasive phase: when the tumor induced stress reaches the matrix strength tolerance threshold, invasion takes place ‘ideally’ reducing the confining stress to zero (due to matrix-degrading enzymes for instance [55]). (d) Final condition: the invasion cycle is concluded and the non-invasive growth phase starts anew (see the text, and [22] for more details).

Acknowledgments

This work has been supported in part by NIH grant CA 113004 (Center for the Development of a Virtual Tumor, CViT (http://www.cvit.org)). Helpful discussions with M Griffa are gratefully acknowledged.

Appendix A. The amorphous solid analogy

In this mechanistic analogy, cracks correspond to the cellular infiltration channels of figure 1(c) and failure is usually assumed to arise, for un-notched specimens, when the stress $\sigma$ reaches the material strength tolerance $\sigma_C$. In notched specimens, it is not the stress $\sigma$ but the stress-intensity factor, $K$, that must reach a critical value $K_C$ for fracture propagation [39]. Thus, $K \equiv \sqrt{\pi \chi \sigma l} = K_C$, i.e. the stress-intensity factor at the tip of a crack of length $l$ loaded by a stress $\sigma$ must be equal to the fracture toughness $K_C$ of the material; $\chi$ is a geometrical factor, e.g., for a crack at the edge of a large medium $\chi \approx 1.12$. Recently, a more powerful criterion (valid both for small and large values of $l$) for predicting the strength of solids has been derived [36–38] by simply removing the assumption of continuum crack propagation. Accordingly, the failure stress is estimated as

$$\sigma_f = \frac{\sigma_C}{\sqrt{1 + \frac{\pi \chi \sqrt{\sigma l}}{K_C}}}$$

(A.1)

which represents an asymptotic matching between the two previously discussed solutions. Further details, such as the number of branches during invasion, can be deduced as follows: let us consider a cylindrical tumor of radius $R$.
embedded in a linear elastic matrix. Take \( N \) cracks of length \( a \), starting at and perpendicular to the interface, equally spaced and thus with an angular period of \( 2\pi/N \). According to fracture mechanics the stress intensity factor at the tip of each crack is

\[
K_I = P \sqrt{\pi a F \left( \frac{a}{R + a}, N \right)},
\]

where \( P \) is the tumor-to-matrix interface pressure and \( F \) is a known function [47]. Note that \( F(0, N) \approx 2.243 \), whereas \( F(1, N) \approx 2/\sqrt{N} \), for the large value of \( N \) (> 10). According to quantized fracture mechanics [36–38], propagation will take place when

\[
[K_{1c}^\text{aq}] = K_{IC},
\]

in which \( \left\langle \bullet \right\rangle \equiv \frac{1}{q} \int_a^{a_{aq}} \bullet \, da \) \( K_{IC} \) represents the material fracture toughness and \( q \) the fracture quantum (related to the microstructure) of the matrix. Note that according to classical fracture mechanics \( q \rightarrow 0 \). By introducing equation (A.2) into equation (A.3) and inverting the latter with respect to \( N \), we obtain the number of branches during tumor invasion:

\[
N_f \approx \frac{4\pi}{P^2(a + q/2)} K_{IC}^2 C,
\]

with

\[
C = \frac{\langle a F^2 N/4 \rangle_{aq} a_{aq}}{a + q/2}.
\]

For large cracks it follows that \( C \approx 1 \). We note that \( q \) can be fixed by imposing the same predictions in the case of small cracks with those derivable according to the "splashing water drop" analogy (see appendix B). Accordingly

\[
q \approx \frac{K_{IC}^2}{2\pi} \sqrt{\frac{R}{\pi a \sigma^3}}
\]

where \( \sigma \) is the surface tension. Thus, for large cracks, \( N_f \approx \frac{4\pi P^2}{K_{IC}^2} \), whereas for small cracks

\[
N_f \approx \frac{P R}{\sigma}.
\]

**Appendix B. The viscous fluid analogy**

In 1898, the naval engineer H J S Hele-Shaw observed that a drop of liquid injected in a more viscous environment would generate an instability which leads to a variable number of "fingers". The macroscopic details of this so-called Hele-Shaw effect depend on the combination of selected fluids and their viscosity. Recently, this effect has been widely studied because of its intrinsic fractality, and the fractal dimensions have been measured for many pairs of fluids. The appearance of the same patterns in multicellular tumor spheroids, MTS, is probably related to the strong viscosity of the commonly used ECM gel MATRIGEL \(^\text{TM}\) (in the order of 10 Pa s, see www.tbmc.it), while the viscosity of the MTS is unfortunately unavailable (but probably lower than in Matrigel). Also other common culture media, such as collagen, edible gelatine and agar can reach large viscosity values, up to 100 Pa s, after sol–gel transition. Apparently, cell membrane viscosity can vary over a wide range of values. For instance, Yu-Qiang et al [48] found for breast cancer cells a very large value (0.021 pN s \( \mu m^{-3} \) corresponding to \( 2.1 \times 10^8 \) Pa s), while Dunham et al [49] obtained for keratinocytes values between 60 and 120 cP (i.e. 0.06–0.12 Pa s). Further investigation in biological tissues is rather cumbersome, due to the need for accurate measurements of their viscosity, but in principle it should be possible to predict whether a particular tumor type in a given tissue or organ would exhibit a finger-like invasive pattern or not.

As observed in [3], there is a striking analogy between MTS invasion and a liquid drop impacting on a solid surface and causing the formation of a fluid ‘crown’ (‘Rayleigh’ or ‘Yarin–Weiss’ capillary instability [50–52]) as shown in figure 2. They seem to share several features, although they may not be easily recognized due to the unfamiliar terminology: e.g., the occurrence of invasive ‘fingering’ corresponds to the secondary jets, the evidence for branch confluence corresponds to hole nucleation near the fluid rim and, finally, the proliferating aggregates emerging within the invasive cell population [53] correspond to the outgoing small drops at the fluid–air interface. Intriguingly, the number of fingers can then be predicted on the basis of the following parameters: the fluid density \( \rho \), the drop radius \( R \), the deceleration \( a \), the fluid viscosity \( \mu \) and the surface tension \( \sigma \) [54]:

\[
N_f = \frac{2\pi R}{\lambda}.
\]

where

\[
\lambda = 2\pi (3\sigma/\rho a)^{1/2}.
\]

Assuming for simplicity a spherical shape and a radius \( R \) at the invasion time,

\[
a = F/m = P S/\rho V = 3P/\rho R,
\]

which, remarkably, yields again equation (A.6):

\[
N_f \approx \sqrt{\frac{P R}{\sigma}}.
\]

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