Fluids and Structures Interaction and Modeling

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Flow induced microstructure of emulsions

Sergio Caserta and Stefano Guido





Università di Napoli Federico II

Scuola Politecnica delle Scienze di Base Dipartimento di Ingegneria Chimica, dei Materiali e della Produzione industriale Laboratory of Chemical Engineering @ µ-scale



Drop deformation & Breakup







Non-Newtonian

Collision & Coalescence



Shear Banding



Shear flow cell





Multilamellar Vesicles J. Rheol., 2003, 2007, 2016 PRL 2006, 2008, 2012 Soft Matter 2011, 2013 Langmuir, 2012, 2013

Cell dynamic evolution as Active Bio Soft Matter

Physiological and Pathological Processes





Single Cell Migration Assays





Persistent Random Walk

$$\langle d^2(t)\rangle = 4\mu [t - P(1 - e^{-t/P})]$$

- μ = Random Motility Coefficient
- **P** = Persistence Time

Dickinson & Tranquillo, AIChE J., 1993 Uhlenbeck & Ornstein, Physical R., 1930







n_i= number of Square Displacement for each cell M= number of cells





Wound Healing Assay



Wound Healing Assay



Wound edge speed

Random motility coefficient

$$s = \sqrt{\frac{4 \mu \ln(2)}{t_d}} \qquad \qquad \mu = \frac{s^2 td}{4 \ln(2)}$$

Ascione et al., Chemical Engineering Science, 2017 Ascione et al., Experimental Cell Research, 2017















△ Inner region







	Vρ² (μm/min)²	Vθ ² (degree/min) ²
<u>Outer</u> region	0.0086 ± 0.0050	0.027 ± 0.016
Inner region	0.0075 ± 0.0039	0.019 ± 0.013







M.B.A. Djamgoz et al, Philosophical Transactions of the Royal Society B, 2014





V. Cristini et al., Clinical Cancer Research, 2005 H.B. Frieboes, Cancer Research, 2006

G. Mehta et al., J. Controlled Release, 2012





Matematical model + numerical simulations

$$\Phi\left(\frac{\partial \Phi_{i}}{\partial t} + \nabla \cdot (\mathbf{u}_{i} \Phi_{i})\right) = -\nabla \cdot \mathbf{J}_{i,j} + S_{i};$$

$$(1-1,2, ...)$$
host tissue a
$$S_{1} = \lambda_{M,1} \frac{n}{n_{V}} \Phi_{1} - \lambda_{A,1} \Phi_{1} - \lambda_{N} \mathcal{H}[n_{N} - n] \Phi_{1} - \lambda_{TR} f_{rand} \Phi_{1},$$

$$S_{2} = \lambda_{TR} f_{rand} \Phi_{1} + \lambda_{M,2} \frac{n}{n_{V}} \Phi_{2} - \lambda_{A,2} \Phi_{2} - \lambda_{N} \mathcal{H}[n_{N} - n] \Phi_{2}.$$

$$\mathbf{u}_{i} = -k(\Phi_{i}) \left(\nabla p - \sum_{l} \gamma_{l} \frac{\delta E}{\delta \Phi_{l}} \nabla \Phi_{l}\right) + \chi_{n}(\Phi_{i}, n) \nabla \frac{n}{n_{V}} + \chi_{h}(\Phi_{1})$$

$$0 = D \nabla^{2} n/n_{V} + \nu(1 - n/n_{V}) (1 - p/p_{V})^{+} \delta_{C} - \eta_{i} \Phi_{i}(n/n_{V})$$

(i = 1, 2, ...) Tumor viable species, dead tumor, host tissue and interstitial fluid

- **Φ**: Concentration
- J: Cell flux
- **u**: Cell velocity
- S: Source of cells
- **λ:** Necrosis/Mitosis/Mutation rates
- **n**: substrate concentration

Cristini et al., Clinical Cancer Research 2005 Bearer et al., Cancer Research 2009







Diffusional Instability model

Cristini, Caserta et al., Clinical Cancer Research 2005



Cancer Growth and Invasion: Experiments

 \mathbf{Q}

С













• The dynamic evolution of both isolated and collective cells can be efficiently investigated *in vitro* by Time-lapse Microscopy and image analysis

• A transport phenomena bioengineering approach can be efficiently used to describe the dynamic evolution of cell populations

• Cell dynamic behavior is driven by concentration gradients



... Thank you ...









- S. Caserta
 - - F. Ascione

Main Partners:







V. Cristini

D. Vignjevic

L. Maiuri









DIPARTIMENTO DI INGEGNERIA CHIMICA, DEI MATERIALI E DELLA PRODUZIONE INDUSTRIALE UNIVERSITA' DI NAPOLI FEDERICO II



LABORATORY OF CHEMICAL **ENGINEERING** *(a)* **INTERFACE**



G. Tomaiuolo V. Villella









R. Liuzzi

L. Sicignano C. Caiazza

R.I. Castaldo A. Prato





We are looking forward to welcoming you in Sorrento - Naples





• Sorrento, April 17-20, 2018



Abstract deadline: <u>18th November 2017</u>

www.rheology-esr.org/aerc2018

