

**Year of publication 2018**

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Coscoy S, Baiz S, Octon J, Rhoné B, Perquis L, Tseng Q, Amblard F, Semetey V. (2018 Oct 16)  
**Microtopographies control the development of basal protrusions in epithelial sheets**

*Biointerphases* : 13 : 041003 : [DOI : 10.1116/1.5024601](https://doi.org/10.1116/1.5024601)

**Summary**

Venzac B, Madoun R, Benarab T, Monnier S, Cayrac F, Myram S, Leconte L, Amblard F, Viovy JL, Descroix S, Coscoy S (2018 Oct 16)

**Engineering small tubes with changes in diameter for the study of kidney cell organization**

*Biomicrofluidics* : 12 : 024114 : [DOI : 10.1063/1.5025027](https://doi.org/10.1063/1.5025027)

**Summary**

Blanch-Mercader C., Yashunsky V., Garcia S., Duclos G., Giomi L., Silberzan P. (2018 Oct 9)

**Turbulent dynamics of epithelial cell cultures**

*Phys. Rev. Lett.* : 120 : 208001 : [DOI : 10.1103/PhysRevLett.120.208101](https://doi.org/10.1103/PhysRevLett.120.208101)

**Summary**

We investigate the large length and long time scales collective flows and structural rearrangements within in vitro human bronchial epithelial cell (HBEC) cultures. Activity-driven collective flows result in ensembles of vortices randomly positioned in space. By analyzing a large population of vortices, we show that their area follows an exponential law with a constant mean value and their rotational frequency is size independent, both being characteristic features of the chaotic dynamics of active nematic suspensions. Indeed, we find that HBECs self-organize in nematic domains of several cell lengths. Nematic defects are found at the interface between domains with a total number that remains constant due to the dynamical balance of nucleation and annihilation events. The mean velocity fields in the vicinity of defects are well described by a hydrodynamic theory of extensile active nematics.

Duclos G., Blanch-Mercader C., Yashunsky V., Salbreux G., Joanny J.-F., Prost J., Silberzan P. (2018 Oct 3)

**Spontaneous shear flow in confined cellular nematics**

*Nature Physics* : [DOI : 10.1038/s41567-018-0099-7](https://doi.org/10.1038/s41567-018-0099-7)

**Summary**

In embryonic development or tumour evolution, cells often migrate collectively within confining tracks defined by their microenvironment<sup>1,2</sup>. In some of these situations, the displacements within a cell strand are antiparallel<sup>3</sup>, giving rise to shear flows. However, the mechanisms underlying these spontaneous flows remain poorly understood. Here, we show that an ensemble of spindle-shaped cells plated in a well-defined stripe spontaneously develops a shear flow whose characteristics depend on the width of the stripe. On wide stripes, the cells self-organize in a nematic phase with a director at a well-defined angle with the stripe's direction, and develop a shear flow close to the stripe's edges. However, on stripes narrower than a critical width, the cells perfectly align with the stripe's direction and the net flow vanishes. A hydrodynamic active gel theory provides an understanding of these observations and identifies the transition between the non-flowing phase oriented along the stripe and the tilted phase exhibiting shear flow as a Fréedericksz transition driven by the activity of the cells. This physical theory is grounded in the active nature of the cells and based on symmetries and conservation laws, providing a generic mechanism to interpret in vivo antiparallel cell displacements.