A multiscale model of complex endothelial cell dynamics in early angiogenesis

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Set- up	Specifications	Figures
Betup 1: Individual sprout	$N_I^x = 25, \ N_I^y = 29$ $\mathcal{I}_{init} = \{i = (i_x, i_y)^T : \ i_x = 1, 2, \ i_y = 14, 15\}$ $\mathcal{I}_{VP} = \{i = (1, i_y)^T : \ i_y = 14, 15\}$	For WT cells: Figs 7, 8, 10, 13, 14 and S7-S9 Figs. For mutant cells: Fig 15, S3-S9 Figs.
	$s_{init} = r$ $c_{init} = 0.0, m_{init} = 1.0$	
	$V \in \{0, 2500, 25000\}$, uniform distribution over the lattice, which correspond to 0, 5 and 50 ng/ml VEGF concentration, respectively. The exact value specified in the text.	
	and S1 Appendix, respectively).	
Setup 2: Vessel in VEGF gradient	$N_I^x = 25, \ N_I^y = 29$ $\mathcal{I}_{init} = \{i = (i_x, i_y)^T : \ i_x = 12, 13, \ i_y = 1, \dots, 29\}$	Fig 9
	$L_{VP} = L_{init}$ s_{init} - not specified	
	$c_{init} = c_{max}, m_{init} = 0.0$ $V(i_x, i_y) = 2500 \frac{i_y}{N_I^y}$. This corresponds to a VEGF gradient linearly increasing from 0 to 5 ng/ml along the <i>y</i> -axis.	
	Simulations with this setup are performed with WT cells (see S1 Table).	
etup 3: Cell bead	$N_I^x = 25, \ N_I^y = 29$	Fig 11
	$\mathcal{I}_{init} = \{ i = (i_x, i_y)^T : \ (i_x - 13)^2 + (i_y - 9)^2 \le 5, \ i \in \mathcal{I} \}$	
	$\mathcal{I}_{VP} = \mathcal{I}_{init}$	
	s_{init} - not specified	
	$c_{init} = c_{max}, m_{init} = 0.0$	
	$V(i_x, i_y) = 2500 \frac{i_y}{N_I^y}$. This corresponds to a VEGF gradient linearly increasing from 0 to 5 ng/ml along the <i>y</i> -axis.	
N	Simulations with this setup are performed with WT cells (see S1 Table).	
etup 4: Linear sprout	$N_I^x = 25, \ N_I^y = 2$	Fig 12, Table 3
	$\mathcal{I}_{init} = \{ i = (i_x, i_y)^T : i_x = 1, \dots, 25, i_y = 1, 2 \}$	
	$\mathcal{I}_{VP} = \mathcal{I}_{init}$	
	$s_{init} = r$	
	$c_{init} = 0.0, m_{init} = 1.0$	
	V = 15000, corresponding to a constant uniform distribution of VEGF at the concentration 30 ng/ml (used in [1]).	
Ň	Simulations with this setup are performed with mutant cells (S1 Appendix).	

S4 Table. Setups of simulation experiments. For each setup of numerical simulation we specify the lattice dimensions, N_I^x and N_I^y ; the set of indices corresponding to the vascular plexus, \mathcal{I}_{VP} ; the initial cell nuclei positions, \mathcal{I}_{init} ; the initial polarisation direction, s_{init} ; the initial ECM and BM concentrations, c_{init} and m_{init} , respectively; the VEGF distribution over the lattice, V; and cell line used in simulations.

References

1. Jakobsson L, Franco CA, Bentley K, Collins RT, Ponsioen B, Aspalter IM, et al. Endothelial cells dynamically compete for the tip cell position during angiogenic sprouting. Nature Cell Biology. 2010;12(10):943.