

The vessel not taken: Understanding disproportionate blood flow

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A new direct numerical simulation which predicts the flow of RBCs through the body's networks of capillaries. Credit: Peter Balogh



Each time a blood vessel splits into smaller vessels, red blood cells (RBCs) are presented with the same decision: Take the left capillary or the right. While one might think RBCs would divide evenly at every fork in the road, it is known that at some junctures, RBCs seem to prefer one vessel over the other. One new computer model looks to determine why RBCs behave this way, untangling one of the biggest mysteries in our vascular system

A pair of researchers from Rutgers University has demonstrated a new direct numerical simulation in *Physics of Fluids*, which predicts the flow of RBCs through the body's networks of capillaries. By constructing a <u>network</u> of virtual capillaries, the team found that not only can the flow through so-called mother vessels sometimes become skewed, leading to an uneven distribution of RBCs in daughter vessels, but these junctions also switch between an even and uneven flow over time.

"This is the biological problem that has immense significance in healthy states and disease states," said Prosenjit Bagchi, one of the authors of the paper. "These phenomena have been known for centuries, but in terms of high-fidelity computational modeling, there has not been much."

Bagchi likens the partitioning of <u>blood</u> cells at bifurcations along <u>blood</u> <u>vessels</u> to cars in traffic where, sometimes, detours happen. Anything from an injury, to a blocked capillary, to a tumor creating new vessels to feed itself can lead to a blood vessel falling out of use. "When it comes to blood vessels, we use or lose it," Bagchi said. "If a city sees that nobody is driving on a particular road, they will stop keeping up with it and might discard it. Microvascular networks are constantly changing their architecture, even in the aging process."

A prominent method to model red blood cell partitioning, called plasma skimming, renders <u>red blood cells</u> as infinitesimally tiny dots that move through a blood vessel. Bagchi's group was surprised to learn how much



this technique underpredicts the partitioning behavior and leads to highly nonuniform distribution of <u>cells</u> in a network. By considering the effect of the cell size, known as cell screening, the team modelled flow with much less heterogeneity in cell distribution.

The group's work casts new light on a longstanding assumption that blood vessel bifurcations either distribute RBCs proportionately or disproportionately with respect to flow. Instead, their findings revealed that vessels can switch between even and uneven partitioning, based on factors including upstream mechanisms that shift RBCs to one side of the mother <u>vessel</u>, bunching up of RBCs at bifurcation points, or changes in flow resistance in the daughter vessels.

Bagchi said he hopes his findings and model will prove to be a helpful tool for researchers looking to better understand blood <u>flow</u> in microvascular networks. In the future, his team is looking towards other particles, including how drug particles are distributed, to accurately predict their transport through capillary networks.

More information: Peter Balogh et al, Analysis of red blood cell partitioning at bifurcations in simulated microvascular networks, *Physics of Fluids* (2018). DOI: 10.1063/1.5024783

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