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#### Hemodynamic Therapy of Middle Cerebral Artery Vasospasm Guided by a Multiphase Model of Oxygen Transport

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## **Middle Cerebral Artery**





#### Rhoton AL: Neurosurgery 2002; 51[Suppl 1]:53-120

## M1 Segment of the MCA



Rhoton AL: Neurosurgery 2002; 51[Suppl 1]:53-120



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Rhoton AL: Neurosurgery 2002; 51[Suppl 1]:53-120

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#### **Cerebral Vasospasm**





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# **Mainstay of Therapy**

- 'Triple H' therapy
  - Hypertension increase flow through pressure
  - Hemodilution increase flow through viscosity
  - Hypervolemia supports goals of above two
- Potential problem
  - Hemodilution decreases oxygen content

$$DO_2 = CO_2 \cdot \dot{Q}$$

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- Blood viscosity is a non-linear, complex function of hematocrit (fraction of blood volume occupied by red cells, normal 0.45, target 0.30)
- Oxygen content is a linear function of hemoglobin concentration (and saturation, here assumed = 1)
- Blood is a non-Newtonian fluid
- The MCA geometry may be additional confounding variable





## **M1 Segment Geometry**





#### **Example Stenotic Geometries**





## **Unstructured Mesh**





## **Experimental Values of Blood Viscosity**





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## **Carreau-Yasuda Viscosity Model**

$$\eta = m \left[ 1 + \left( \lambda \dot{\gamma} \right)^2 \right]^{\frac{n-1}{2}}$$

$$m = 122.28\varepsilon_{rbc}^{3} - 51.213\varepsilon_{rbc}^{2} + 16.30\varepsilon_{rbc} + 1$$
$$n = 0.8092\varepsilon_{rbc}^{3} - 0.8246\varepsilon_{rbc}^{3} - 0.3503\varepsilon_{rbc} + 1$$

Jung J: J Biomech 2006; 39:2064-73



## **Volume-Averaged Blood Density**



#### $\rho_{blood} = \varepsilon_{rbc} \rho_{rbc} + \varepsilon_{plasma} \rho_{plasma}$



## **Single Phase Governing Equations**

$$\rho(\mathbf{u} \cdot \nabla \mathbf{u}) = -\nabla p \mathbf{I} + \nabla \left( \eta \left[ \nabla \mathbf{u} + \left( \nabla \mathbf{u} \right)^T \right] \right) + \mathbf{F}$$
$$\nabla \mathbf{u} = 0$$

COMSOL Incompressible Navier-Stokes Application Mode (ChemEng)



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## **Two Phase Mixture Model (Eulerian-Eularian)**

$$\rho(\mathbf{u} \cdot \nabla \mathbf{u}) = \nabla p - \nabla \left(\rho \theta_d \rho_d / \rho \left(1 - \theta_d \rho_d / \rho\right) \mathbf{u}_{slip} \mathbf{u}_{slip}\right) + \nabla \left(\eta \left[\nabla \mathbf{u} + (\nabla \mathbf{u})^T\right]\right) + \rho \mathbf{g} + \mathbf{F} \left(\rho_c - \rho_d\right) \left[\nabla \left(\theta_d \left(1 - \theta_d \rho_d / \rho\right) \mathbf{u}_{slip}\right) + m_{dc} \rho_d\right] + \rho_c \left(\nabla \mathbf{u}\right) = 0 \nabla \left[\theta_d \mathbf{u} + \theta_d \left(1 - \theta_d \rho_d / \rho\right) \mathbf{u}_{slip}\right] = m_{dc} / \rho_d$$

COMSOL Multiphase Mixture Model Application Mode (ChemEng) Liquid dispersed and continuous phases



## Two Phase Slip Model (Schiller-Naumann)

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$$\frac{3}{4}C_{d} \frac{\rho_{c} \left| \mathbf{u}_{slip} \right| \mathbf{u}_{slip}}{d_{d}} = -\frac{\rho - \rho_{d}}{\rho \nabla p}$$

$$C_{d} = \begin{vmatrix} \frac{24}{\text{Re}_{p}} \left[ 1 + 0.15 \,\text{Re}_{p}^{0.687} \right] & \text{Re}_{p} < 1000 \\ 0.44 & \text{Re}_{p} \ge 1000 \end{vmatrix}$$

$$\text{Re}_{p} = \frac{d_{d}\rho_{c} \left| \mathbf{u}_{slip} \right|}{\eta}$$

Mixture viscosity model same as for single phase flow (Carreau-Yasuda viscosity model based on hematocrit)



## **Calculation of Oxygen Delivery**

 $DO_2 = \int \mathbf{u} \cdot CO_2$   $CO_2 = hct \cdot MCHC \cdot 1.34 \text{ mL/g}$ MCHC = 33 g/dL

hct = hematocrit (single phase) or *phid* (two phase)

MCHC = mean corpuscular hemoglobin concentration



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# Sample Visualizations, 0.5 Stenosis







#### **Single Phase vs Two Phase Results**





## **Effect of Hematocrit with Mild Stenosis**





#### **Moderate to Severe Stenoses**





#### **Very Severe Stenoses**







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#### **Stenosis 0.9**





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## Conclusions



- In this model, single phase and two phase approaches yielded comparable results
- In mild to critical stenoses, where therapy may impact outcome, hemodilution may worsen oxygen delivery and contribute to ischemia
- In very severe stenoses (0.9 and above), hemodilution may improve oxygen delivery, but blood flow is so low (3 orders of magnitude lower) that it would not likely have any clinical impact
- These results assume no change in downstream impedance to blood flow, but are likely still clinically generalizable to the problem at hand