

# Numerical Simulation of Concentration Polarization to Estimate Gypsum and Calcium Carbonate Scaling on Membrane Surfaces



Asunción Santafé-Moros, José M. Gozávez-Zafrilla

ISIRYM. Universitat Politècnica de Valencia, C/ Camino de Vera s/n, 46022 Valencia, Spain



## Aim

To estimate the potential of salt precipitation (*scaling*) in spiral-wound membrane modules as a consequence of the concentration polarization of ionic species.

Scaling indices: LSI for calcium carbonate and gypsum saturation.

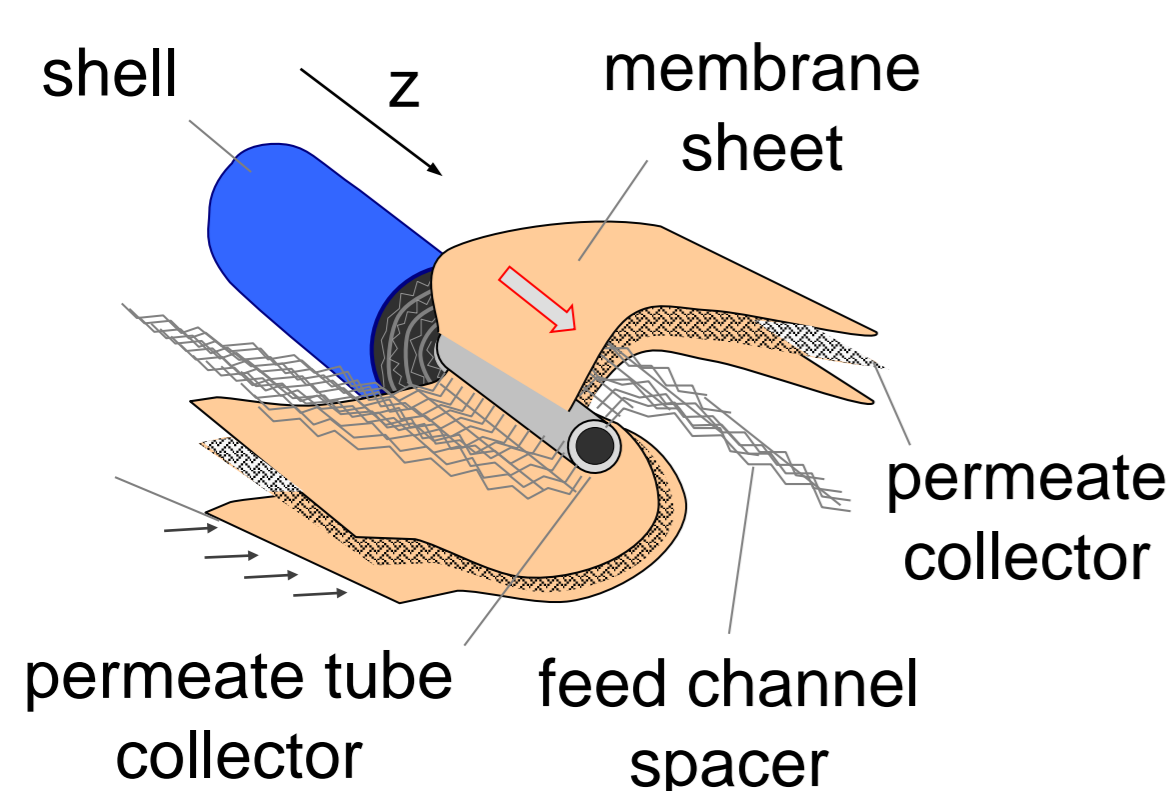


Fig. 1. Structure of spiral-wound module

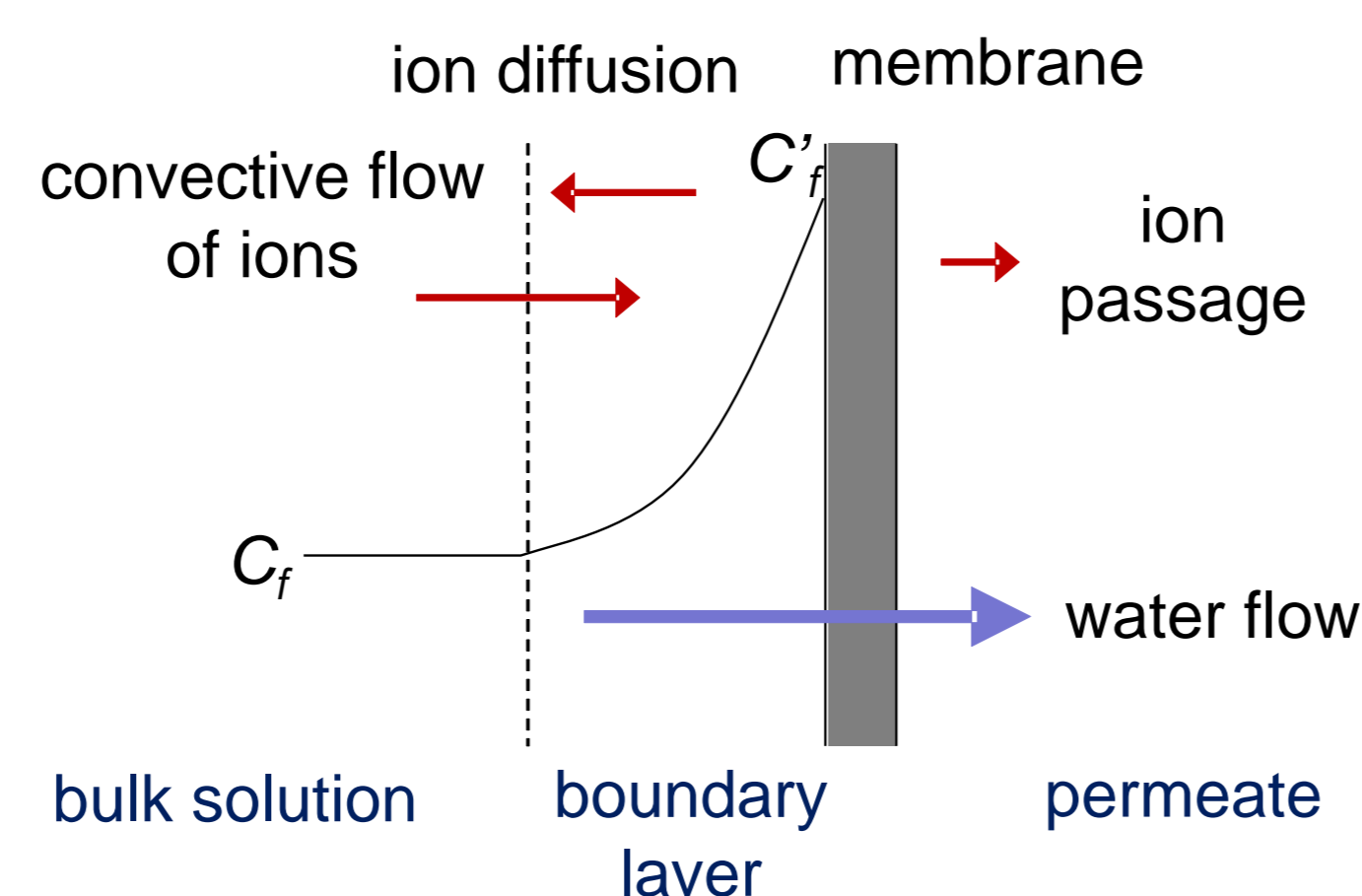


Fig. 2. Concentration polarization

## Cases studied

- Reverse osmosis membrane of permeability:  $A_w = 2,1 \text{ cm}^3 \cdot \text{m}^2 \cdot \text{s}^{-1}$
- Ionic systems for  $\text{CaCO}_3$  scaling (I) and gypsum scaling (II):
  - I)  $[\text{Na}^+] = [\text{Cl}^-] = 20 \text{ mol/m}^3$   $[\text{Ca}^{2+}] = [\text{HCO}_3^-] + [\text{CO}_3^{2-}] = 3 \text{ mol/m}^3$
  - II)  $[\text{Na}^+] = [\text{Cl}^-] = 20 \text{ mol/m}^3$   $[\text{Ca}^{2+}] = [\text{SO}_4^{2-}] = 6 \text{ mol/m}^3$
- Operating conditions:  $\text{pH} = 6.7$ ,  $T = 25 \text{ }^\circ\text{C}$ ,  $P = 5 \text{ bar}$

## Model

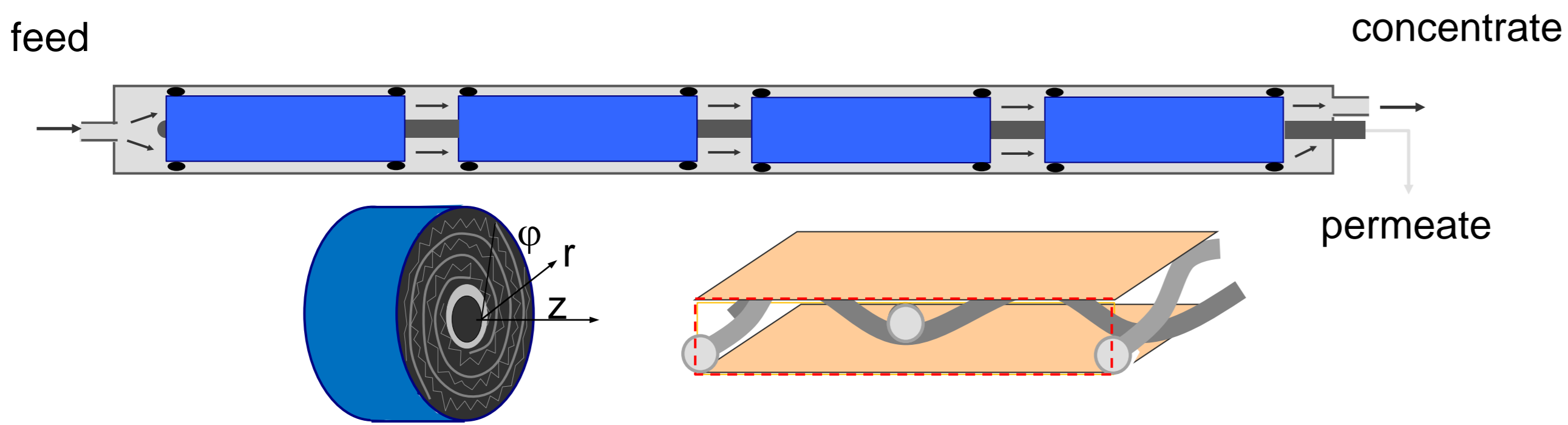


Figure 3. Domain selection

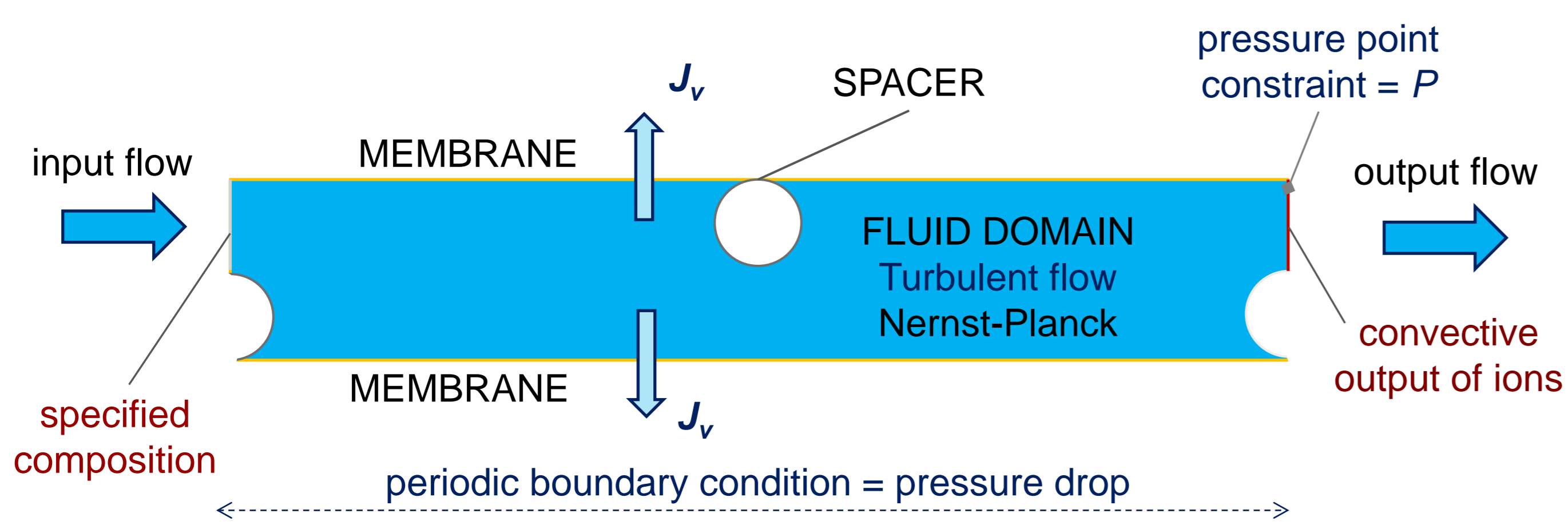


Fig. 4. Boundary specification

### Flow

- Different cross-flow velocity attained by varying pressure drop
- Volumetric flux calculated from phenomenological model:

$$J_v = A_w \cdot (\Delta P - \Delta \Pi)$$

### Ionic transport

- Nernst-Planck mode
- High rejection → 'No flux' condition applied to membranes

## Solving procedure

### Meshing

- Fluid domain: 'finer' free-triangular mesh → cell Reynolds  $< 3$
- Membranes and spacers walls: boundary layer mesh

### Solver steps

1. Solution of hydrodynamics
2. Solution of ionic transport for given hydrodynamics
3. Application of segregated solver to the coupled model

## Conclusions

- The higher concentration in the membrane wall caused by concentration polarization compared to that of the bulk implies a risk of scaling underestimation if the calculation is based on bulk conditions
- The developed model can be useful to set suitable chemical and operating conditions in the concentrate to minimize the risk of scaling

## Results

- The low cross-flow velocity in the areas near the base of the spacer causes the greatest values of LSI (scaling occurs for  $\text{LSI} > 0$ )

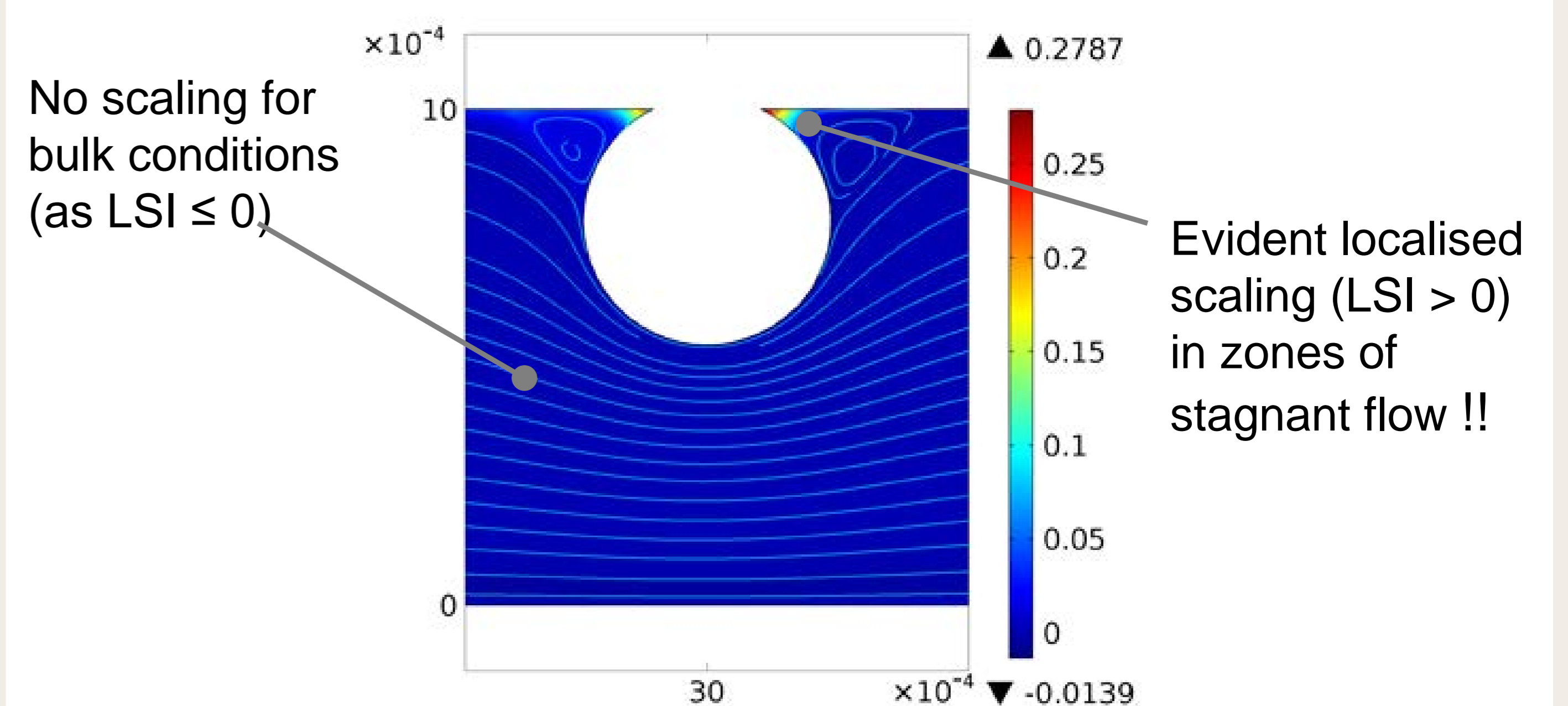


Figure 5. Flow streamlines and mapping of the Langelier Saturation Index (LSI) at cross-flow velocity of 0.5 m/s

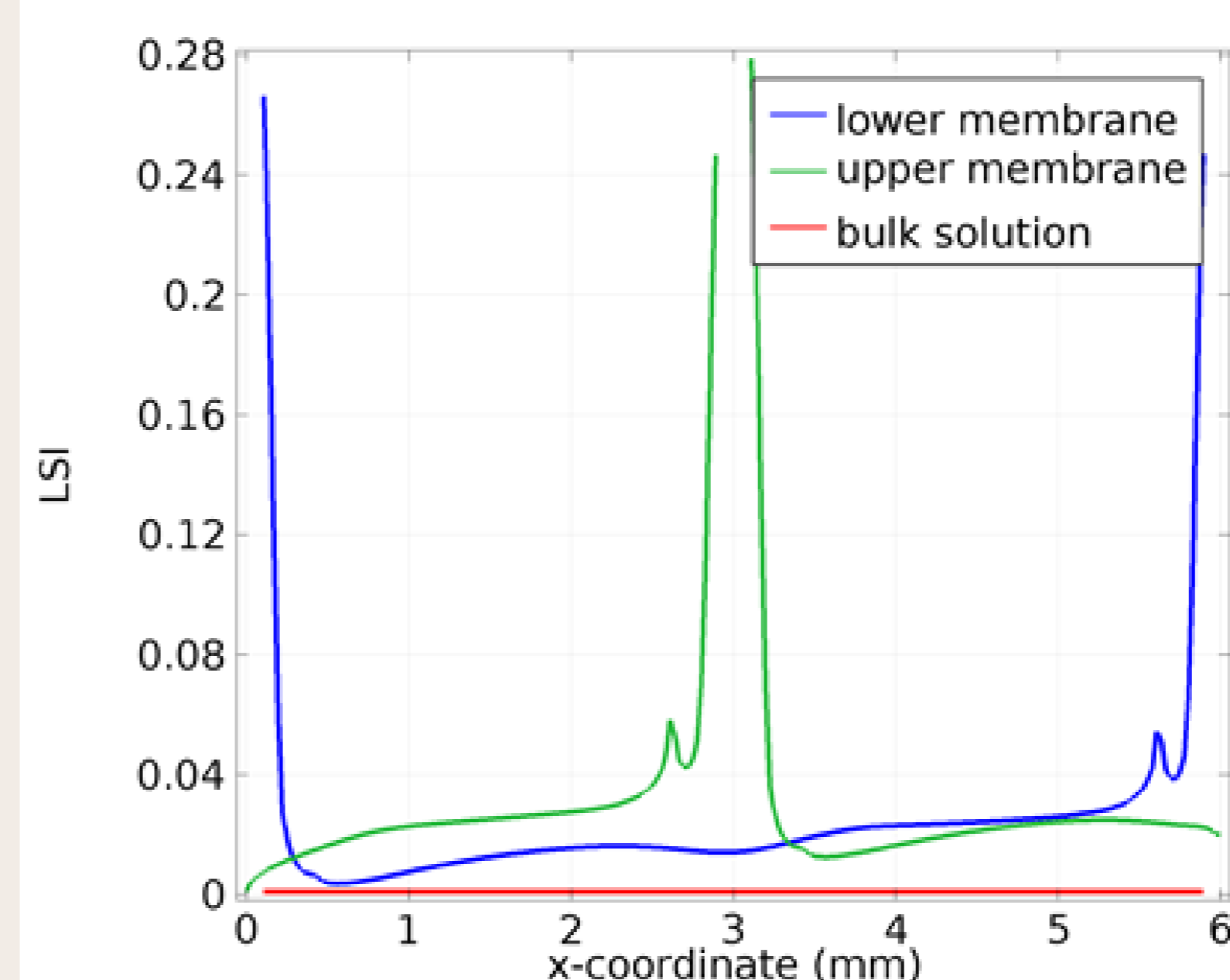


Figure 6. Flow streamlines and LSI mapping at cross-flow velocity of 0.5 m/s

- Fig. 6 shows  $\text{LSI} > 0$  on the whole membrane area, predicting calcium carbonate scaling in the long term
- For gypsum similar results were obtained (not shown)
- Acidification or proper change of operating conditions can correct the scaling.

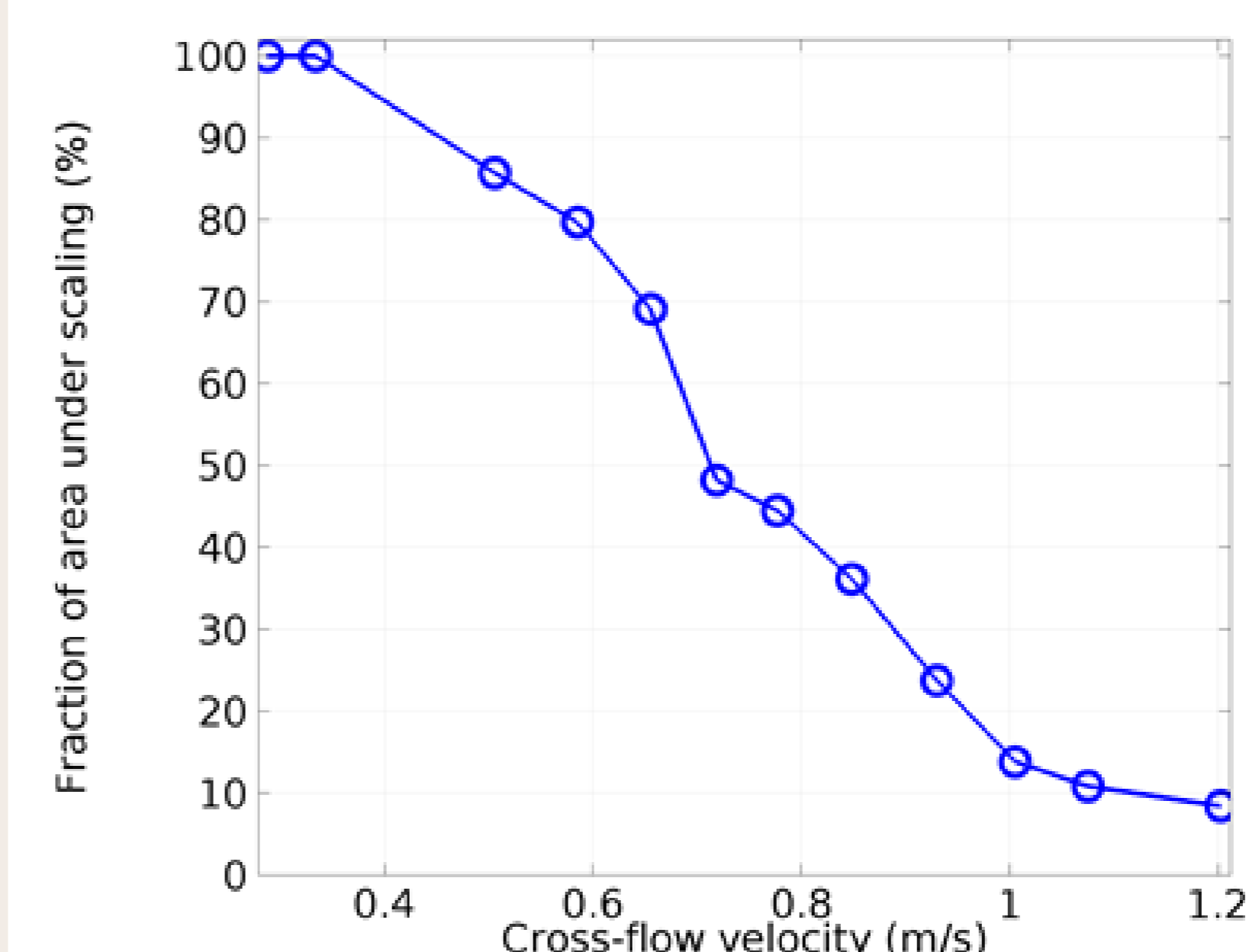


Figure 7. Fraction of membrane area under scaling condition of  $\text{LSI} > 0$  at different cross-flow velocities

- Fig. 7 shows how the membrane area under scaling is reduced as a consequence of the increase in cross-flow velocity.

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