

A New Formulation of Immiscible Compressible Two-Phase Flow in Porous Media Via the Concept of Global Pressure

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Scaling Up and Modeling for Transport and Flow in Porous Media

Outline

Two-phase immiscible, compressible flow equations

Fractional flow formulation

New global formulation

Simplified global formulation

Numerical comparison of the coefficients

Conclusion

Compressible two-phase flow

We consider two-phase isothermal, compressible, immiscible flow through heterogeneous porous medium. For example, water and gas.

Assumptions:

- ▶ Incompressible fluid: water, $\rho_w = \text{const.}$
- ▶ Compressible fluid: gas, $\rho_g = c_g p_g$.
- ▶ Viscosities μ_w and μ_g are constant.
- ▶ No mass exchange between the phases;
- ▶ The temperature is constant;

Note that the assumptions on the form of mass densities are not essential.

Independent variables: water saturation S_w and gas pressure p_g
($\rho_w = p_g - p_c(S_w)$, $S_g = 1 - S_w$).

Flow equations

Mass conservation: for $\alpha \in \{w, g\}$,

$$\Phi \frac{\partial}{\partial t} (\rho_\alpha S_\alpha) + \operatorname{div}(\rho_\alpha \mathbf{q}_\alpha) = 0,$$

The Darcy-Muscat law: for $\alpha \in \{w, g\}$,

$$\mathbf{q}_\alpha = -\mathbb{K} \frac{k r_\alpha(S_\alpha)}{\mu_\alpha} (\nabla p_\alpha - \rho_\alpha \mathbf{g}),$$

Capillary law:

$$p_c(S_w) = p_g - p_w,$$

No void space.

$$S_w + S_g = 1.$$

Fractional flow formulation

Goal:

- Reformulate flow equations in a form giving less tight coupling between the two differential equations, allowing a sort of IMPES (implicit in pressure and explicit in saturation) numerical treatment.

There are two approaches:

1. Introduce **total velocity**: $\mathbf{Q}_t = \mathbf{q}_w + \mathbf{q}_g$: leads to non-conservative form of the equations
2. Introduce **total flow**: $\mathbf{Q}_t = \rho_w \mathbf{q}_w + \rho_g \mathbf{q}_g$: leads to conservative form of the equations.

We work with **total flow**.

Fractional flow formulation: equations

Total flow:

$$\mathbf{Q}_t = -\lambda(S_w, p_g)\mathbb{K}(\nabla p_g - f_w(S_w, p_g)\nabla p_c(S_w) - \bar{\rho}(S_w, p_g)\mathbf{g}),$$

Total mass conservation:

$$\Phi \frac{\partial}{\partial t}(S_w \rho_w + (1 - S_w) \rho_g(p_g)) + \operatorname{div}(\mathbf{Q}_t) = 0,$$

Water mass conservation:

$$\Phi \rho_w \frac{\partial S_w}{\partial t} + \operatorname{div}(f_w(S_w, p_g)\mathbf{Q}_t + \mathbb{K}\mathbf{g}b_g(S_w, p_g)) = \operatorname{div}(\mathbb{K}a(S_w, p_g)\nabla S_w).$$

Fractional flow formulation: coefficients

phase mobilities $\lambda_w(S_w) = \frac{kr_w(S_w)}{\mu_w}, \quad \lambda_g(S_w) = \frac{kr_g(S_w)}{\mu_g},$

total mobility $\lambda(S_w, p_g) = \rho_w \lambda_w(S_w) + \rho_g(p_g) \lambda_g(S_w),$

water fractional flow $f_w(S_w, p_g) = \frac{\rho_w \lambda_w(S_w)}{\lambda(S_w, p_g)},$

mean density $\bar{\rho}(S_w, p_g) = \frac{\lambda_w(S_w) \rho_w^2 + \lambda_g(S_w) \rho_g(p_g)^2}{\lambda(S_w, p_g)},$

"gravity" coeff. $b_g(S_w, p_g) = \rho_w \rho_g(p_g) \frac{\lambda_w(S_w) \lambda_g(S_w)}{\lambda(S_w, p_g)} (\rho_w - \rho_g(p_g)),$

"diffusivity" coeff. $a(S_w, p_g) = -\rho_w \rho_g(p_g) \frac{\lambda_w(S_w) \lambda_g(S_w)}{\lambda(S_w, p_g)} p'_c(S_w).$

Decoupling of the system

In total flow we want to eliminate saturation gradient:

$$\mathbf{Q}_t = -\lambda(S_w, p_g) \mathbb{K} (\nabla p_g - f_w(S_w, p_g) p'_c(S_w) \nabla S_w - \bar{\rho}(S_w, p_g) \mathbf{g}),$$

- ▶ Idea: introduce a new pressure-like variable that will eliminate ∇S_w term (CHAVENT-JAFFRÉE: MATHEMATICAL MODELS AND FINITE ELEMENTS FOR RESERVOIR SIMULATION)
- ▶ Find a new pressure variable p , called **global pressure**, and a function $\omega(S_w, p)$ such that:

$$\nabla p_g - f_w(S_w, p_g) p'_c(S_w) \nabla S_w = \omega(S_w, p) \nabla p \quad (1)$$

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New global formulation

We introduce unknown function π such that

$$p_g = \pi(S_w, p),$$

where p is *global pressure* to be defined. Then (1) reads

$$\nabla p_g = \omega(S_w, p) \nabla p + f_w(S_w, \pi(S_w, p)) p'_c(S_w) \nabla S_w,$$

or,

$$\begin{aligned} & \frac{\partial \pi}{\partial S_w}(S_w, p) \nabla S_w + \frac{\partial \pi}{\partial p}(S_w, p) \nabla p \\ &= \omega(S_w, p) \nabla p + f_w(S_w, \pi(S_w, p)) p'_c(S_w) \nabla S_w. \end{aligned}$$

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New global formulation

Since p and S_w are independent variables we must have:

$$\frac{\partial \pi}{\partial S_w}(S_w, p) = f_w(S_w, \pi(S_w, p))p'_c(S_w) \quad (2)$$

$$\frac{\partial \pi}{\partial p}(S_w, p) = \omega(S_w, p). \quad (3)$$

Conclusion:

- To calculate $\pi(S_w, p)$ solve the Cauchy problem:

$$\begin{cases} \frac{d\pi(S, p)}{dS} = \frac{\rho_w \lambda_w(S) p'_c(S)}{\rho_w \lambda_w(S) + c_g \lambda_g(S) \pi(S, p)}, & 0 < S < 1 \\ \pi(1, p) = p. \end{cases}$$

- Get $\omega(S_w, p)$ from (3).

New global formulation: Remarks

- It is more natural to replace saturation S_w with capillary pressure $u = p_c(S_w)$ as an independent variable. Then:

$$\begin{cases} \frac{d\hat{\pi}(u, p)}{du} = \frac{\rho_w \hat{\lambda}_w(u)}{\rho_w \hat{\lambda}_w(u) + c_g \hat{\lambda}_g(u) \hat{\pi}(u, p)}, & u > 0 \\ \hat{\pi}(0, p) = p. \end{cases}$$

and $\pi(S_w, p) = \hat{\pi}(p_c(S_w), p)$ [Hat denotes the change of variables.]

- ω is strictly positive and less than 1:

$$\omega(S_w, p) = \exp \left(- \int_0^{p_c(S_w)} \frac{c_g \rho_w \hat{\lambda}_w(u) \hat{\lambda}_g(u)}{(\rho_w \hat{\lambda}_w(u) + c_g \hat{\lambda}_g(u) \hat{\pi}(u, p))^2} du \right),$$

- $p \leq \pi(S_w, p) \leq p + p_c(S_w)$ and therefore $p_w \leq p \leq p_g$.

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- $p \leq \pi(S_w, p) \leq p + p_c(S_w)$ and therefore $p_w \leq p \leq p_g$.

New global formulation: Flow equations

Total flow:

$$\mathbf{Q}_t = -\lambda^n(S_w, p)\mathbb{K}(\omega(S_w, p)\nabla p - \bar{\rho}^n(S_w, p)\mathbf{g}).$$

where the superscript *n* stands for new.

Total mass conservation:

$$\Phi \frac{\partial}{\partial t}(S_w \rho_w + c_g(1 - S_w)\pi(S_w, p)) + \operatorname{div} \mathbf{Q}_t = 0.$$

Water mass conservation:

$$\Phi \rho_w \frac{\partial S_w}{\partial t} + \operatorname{div}(f_w^n(S_w, p)\mathbf{Q}_t + \mathbb{K}\mathbf{g}b_g^n(S_w, p)) = \operatorname{div}(\mathbb{K}a^n(S_w, p)\nabla S_w).$$

New global formulation: Coefficients

New coefficients are obtained from the **old ones** by replacing gas pressure p_g by $\pi(S_w, p)$:

total mobility $\lambda^n(S_w, p) = \lambda(S_w, \pi(S_w, p)),$

water fractional flow $f_w^n(S_w, p) = f_w(S_w, \pi(S_w, p)),$

mean density $\bar{\rho}^n(S_w, p) = \bar{\rho}(S_w, \pi(S_w, p)),$

"gravity" coeff. $b_g^n(S_w, p) = b_g(S_w, \pi(S_w, p)),$

"diffusivity" coeff. $a^n(S_w, p) = a(S_w, \pi(S_w, p)).$

Simplified global formulation

The problem “Find a *mean pressure* p and a function $\omega(S_w, p)$ such that”:

$$\nabla p_g - f_w(S_w, p_g) p'_c(S_w) \nabla S_w = \omega(S_w, p) \nabla p \quad (4)$$

can be solved by introducing a **simplification**:

- Gas density $\rho(p_g)$ can be replaced by $\rho(p)$ without introducing a significant error. Consequently, fractional flow function $f_w(S_w, p_g)$ can be replaced by $f_w(S_w, p)$.

Then, (4) reduces to:

$$\nabla p_g = \omega(S_w, p) \nabla p + f_w(S_w, p) p'_c(S_w) \nabla S_w.$$

Simplified global formulation

Now we can write solution:

$$p_g = p + \int_1^{S_w} f_w(s, p) p'_c(s) ds, \quad (5)$$

$$\omega(S_w, p) = 1 + \frac{\partial}{\partial p} \int_1^{S_w} f_w(s, p) p'_c(s) ds. \quad (6)$$

Note that (5) is **nonlinear equation** w.r.t. p to be solved.

Simplified global formulation: equations

The system written in unknowns p and S_w .

Total flow:

$$\mathbf{Q}_t = -\lambda(S_w, p)\mathbb{K}(\omega(S_w, p)\nabla p - \bar{\rho}(S_w, p)\mathbf{g}),$$

Total mass conservation:

$$\Phi \frac{\partial}{\partial t}(S_w \rho_w + c_g(1 - S_w)p) + \operatorname{div} \mathbf{Q}_t = 0,$$

Water mass conservation:

$$\Phi \rho_w \frac{\partial S_w}{\partial t} + \operatorname{div}(f_w(S_w, p)\mathbf{Q}_t + \mathbb{K}\mathbf{g}b_g(S_w, p)) = \operatorname{div}(\mathbb{K}a(S_w, p)\nabla S_w).$$

Note that the equations are the same, only the coefficients are different.

Simplified global formulation: Applicability

- ▶ Global pressure is well defined for all $p_g \geq 0$ and $S_w \in (0, 1]$ such that $p_w = p_g - p_c(S_w) \geq 0$;
- ▶ $p_w \leq p \leq p_g$;
- ▶ $\omega(S_w, p) > 0$ holds only if certain additional condition is fulfilled. For example: $p \geq p_{min} > 0$ and

$$\forall S_w \in (0, 1], \quad \int_{S_w}^1 \frac{c_g \rho_w \lambda_w(s) \lambda_g(s)}{(\rho_w \lambda_w(s) + c_g \lambda_g(s) p_{min})^2} |p'_c(s)| \, ds < 1.$$

Numerical comparison

The difference in the coefficients of new and simplified models depends on the difference between $\pi(S_w, p)$ and p .

$$0 \leq \pi(S_w, p) - p \leq \int_0^{p_c(S_w)} \frac{\rho_w \hat{\lambda}_w(u)}{\rho_w \hat{\lambda}_w(u) + c_g p \hat{\lambda}_g(u)} du$$

Usually permeability functions depend on dimensionless variable $v = u/P_r$, where P_r is some pressure constant (entry pressure in *Brooks-Corey* or *van Genuchten* models). Then we have:

$$0 \leq \pi(S_w, p) - p \leq P_r \int_0^{+\infty} \frac{\rho_w \hat{\lambda}_w(v)}{\rho_w \hat{\lambda}_w(v) + c_g p \hat{\lambda}_g(v)} dv$$

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Numerical comparison

As an example we show the difference in coefficients:

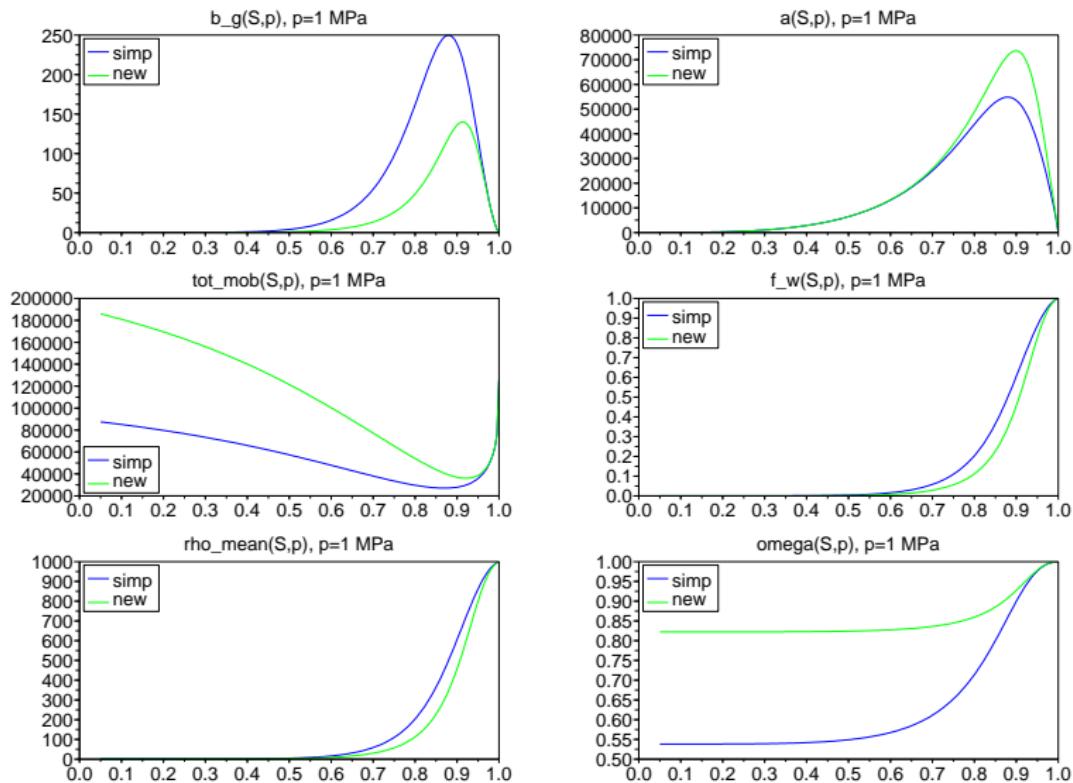
$$\begin{aligned}
 & \lambda(S_w, p) \quad \text{and} \quad \lambda(S_w, \pi(S_w, p)), \\
 & f_w(S_w, p) \quad \text{and} \quad f_w(S_w, \pi(S_w, p)), \\
 & \bar{\rho}(S_w, p) \quad \text{and} \quad \bar{\rho}(S_w, \pi(S_w, p)), \\
 & b_g(S_w, p) \quad \text{and} \quad b_g(S_w, \pi(S_w, p)), \\
 & a(S_w, p) \quad \text{and} \quad a(S_w, \pi(S_w, p)), \\
 & \omega^{simp}(S_w, p) \quad \text{and} \quad \omega^{new}(S_w, p),
 \end{aligned}$$

for van Genuchten functions with parameters (Couple test case):

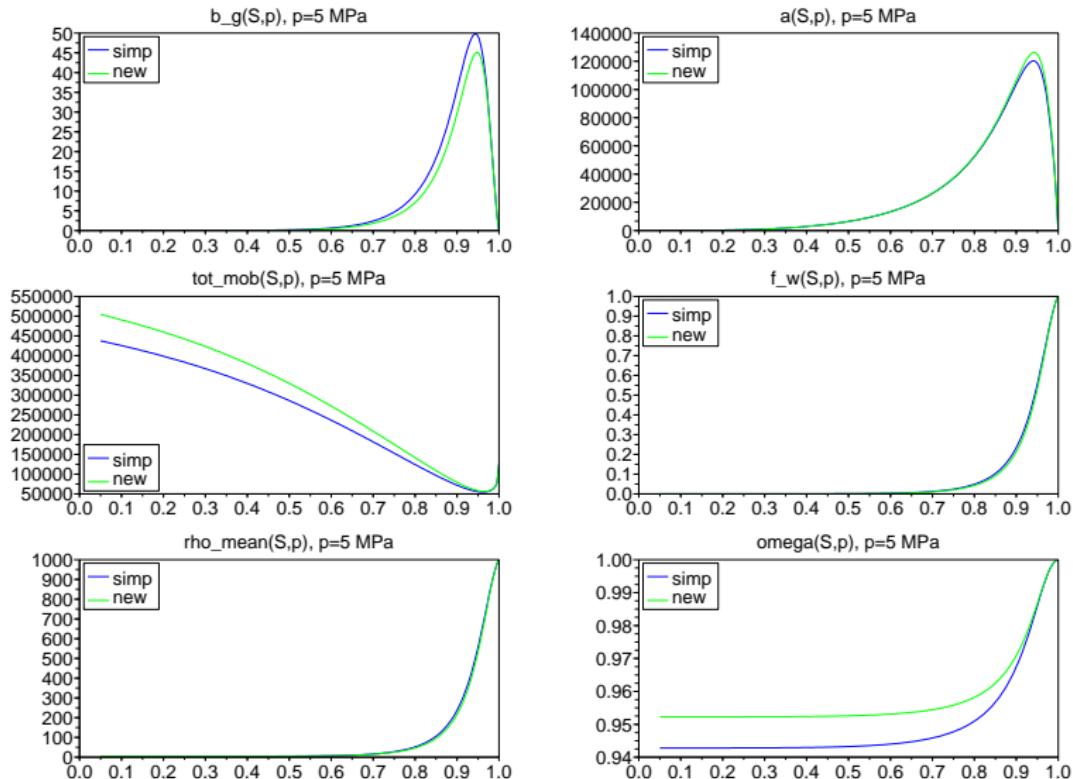
$$\begin{aligned}
 n &= 1.54, \quad Pr = 2 \text{ MPa}, \quad \mu_w = 7.98 \cdot 10^{-3} \text{ Pas}, \quad \mu_g = 9 \cdot 10^{-6} \text{ Pas}, \\
 \rho_w &= 10^3 \text{ kg/m}^3 \text{ and } c_g = 0.808.
 \end{aligned}$$

Notation: **new** = new model; **simpl** = simplified model.

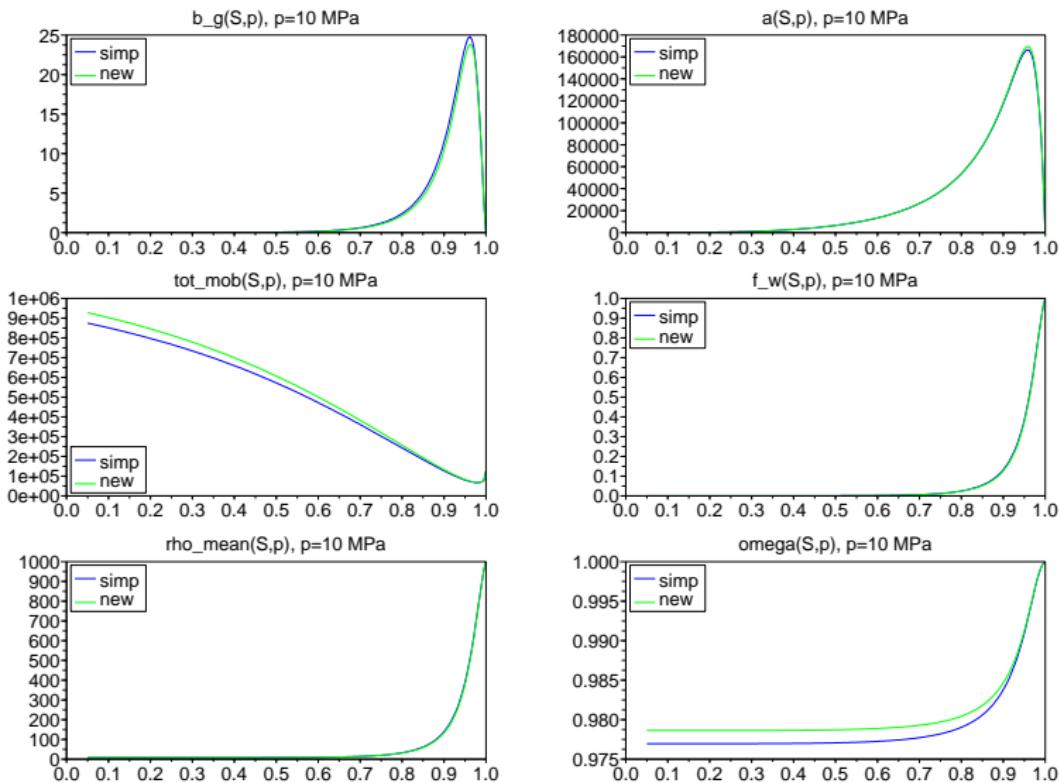
Coefficients at global pressure of 1 MPa



Coefficients at global pressure of 5 MPa



Coefficients at global pressure of 10 MPa



Conclusion

- ▶ In regimes with large global pressures compared to typical capillary pressure the difference in the coefficients is small.
- ▶ For small global pressures and relatively large capillary pressure the difference between new and simplified coefficients becomes significant.
- ▶ Replacing

$$\frac{\partial}{\partial t}(c_g(1 - S_w)\pi(S_w, p)) \quad \text{by} \quad \frac{\partial}{\partial t}(c_g(1 - S_w)p)$$

can have large influence on mass conservation even in for small capillary pressure and elevated global pressure.

Work in progress

- ▶ An extension to multiphase, multicomponent models is straightforward and it is at present in the course of study.
- ▶ The new formulation is well adapted for the mathematical analysis of the model. At present we study existence etc.
- ▶ The discretization of the model by a vertex-centered finite volume scheme is currently studied.

Reference: B. Amaziane, M. Jurak: *A new formulation of immiscible compressible two-phase flow in porous media*, C. R. A. S. Mécanique 336 (2008) 600-605.