Discontinuous Galerkin method for two-component miscible liquid-gas porous media flows

Alexandre Ern∗ Igor Mozolevski†

February 26, 2012

Abstract

We consider two-component (typically, water and hydrogen) compressible liquid-gas porous media flows including mass exchange between phases possibly leading to gas-phase (dis)appearance, as motivated by hydrogen production in underground repositories of radioactive waste. Following recent work by Bourgeat, Jurak, and Smaî, we formulate the governing equations in terms of liquid pressure and dissolved hydrogen density as main unknowns, leading mathematically to a nonlinear elliptic-parabolic system of PDEs, in which the equations degenerate when the gas phase disappears. We develop a discontinuous Galerkin method for space discretization, combined with a backward Euler scheme for time discretization and an incomplete Newton method for linearization. Numerical examples deal with gas-phase (dis)appearance, ill-prepared initial conditions, and heterogeneous problem with different rock types.

1 Introduction

Multicomponent multiphase porous media flows are encountered in several applications including petroleum engineering and various hydrology models related, e.g., to agricultural engineering and groundwater remediation. Such flows have received an enhanced attention recently in connection with gas sequestration and the disposal of radioactive waste in underground repositories. This last application actually constitutes the main motivation for the present work. In such repositories, the corrosion of metallic components, and also marginally the radiolysis of water, leads to hydrogen production. An important issue in the design and safety analysis of the underground repository is then to understand and predict the migration of hydrogen through the host rock. A typical model to describe this situation is to consider a two-phase (liquid and gas), two-component (water and hydrogen) flow. During the simulation, the gas phase is generally not present in the whole domain, as hydrogen gradually penetrates into the host rock which is initially saturated. When both phases are present, the hydrogen component is assumed to be in thermodynamic equilibrium between both phases, and in the context of moderate hydrogen concentrations, this equilibrium is described by the linear Henry law for hydrogen dissolution. For simplicity, we assume herein that water does not vaporize, which is a reasonable assumption in the present context. Additionally, the compressibility of the gas phase must be accounted for.

Multicomponent multiphase porous media flows are described by well-established models covered in several monographs, see, e.g., Bear [6], Chavent and Jaffré [10], Helmig [20], Marle [26], and Peaceman [30]. There is a satisfactory mathematical theory for incompressible two-phase flows hinging on the theory of

∗Partially supported by the Groupement MoMaS (PACEN/CNRS, ANDRA, BRGM, CEA, EdF, IRSN), France and by CNPq, Brazil.
∗Université Paris-Est, CERMICS, Ecole des Ponts ParisTech, 77455 Marne la Vallée cedex 2, France (ern@cermics.enpc.fr).
†Departamento de Matemática, Universidade Federal de Santa Catarina, 88040-900 Florianópolis, SC, Brazil (igor.Mozolevski@mtm.ufsc.br).
elliptic-parabolic PDEs, see, e.g., [2, 7, 23, 24, 28]. Recent mathematical results in the compressible case can be found in [3, 25] without mass exchange between phases and in [27, 33, 34] with mass exchange.

The possibility of gas-phase disappearance rises the issue of selecting the main unknowns in the governing equations since, typically, the phase saturations are not appropriate for that purpose. Following [3, 22], we select here the liquid pressure together with the dissolved hydrogen density. This choice is motivated by the fact that the liquid phase is always present in the whole domain, while the dissolved hydrogen density is always well defined, regardless of the presence of a gas phase. As observed in [22], Henry’s law can be used within a formulation with complementary constraints to determine the presence of the gas phase. Here, as in [8, 32], we adopt the simpler approach where the gas-phase saturation is recovered from the main unknowns using the reciprocal function of the capillary pressure extended by zero. An alternative approach to phase disappearance (where both phases can disappear in different parts of the domain) is discussed in [1, 29].

The goal of the present work is to design and evaluate numerically a discontinuous Galerkin (dG) method for two-component compressible liquid-gas porous media flows including mass exchange between phases. The dG method is used for space discretization, in conjunction with a backward Euler scheme for time discretization and an inexact Newton solver for linearization. Of special interest in the numerical evaluation of the dG method are test cases featuring gas-phase (dis)appearance, ill-prepared initial conditions, and heterogeneous problems with different rock types. Introduced forty years ago, dG methods have experienced a vigorous development over the last decade in many fields of engineering. Attractive features offered by dG methods include the possibility of enforcing locally basic conservation principles, the flexibility in designing the mesh and local degrees of freedom, and the ability to enforce interface conditions in the context of multi-domain problems. We refer to [4, 14] for a unified analysis of dG methods and to recent monographs on the subject [21, 31, 11]. DG methods for incompressible two-phase porous media flows without inter-phase mass exchange have been developed, e.g., in [5, 13, 15, 16, 18], but, to our knowledge, this is the first time where compressible flows with inter-phase mass exchange are covered.

This paper is organized as follows. In §2, we present the governing equations and formulate the mathematical model. In §3, we describe the numerical method and, in particular, design the dG method for space discretization. Finally, in §4, we present the numerical results and draw some conclusions.

2 Problem setting

In this section, we first present the governing equations in their basic form. Then, we specify the choice of main unknowns and derive the mathematical model.

2.1 Governing equations

We assume that the porous medium is isothermal and undeformable. We adopt the terminology of our targeted application related to hydrogen production within geological repositories of radioactive waste. The two phases are indicated by a subscript $\alpha \in \{l, g\}$ referring to liquid and gas respectively. We consider two components, indicated by a superscript $\beta \in \{w, h\}$ referring to water and hydrogen respectively.

The mass conservation equation for each component $\beta \in \{w, h\}$ can be written as

$$
\Phi \sum_{\alpha \in \{l, g\}} \partial_t (s_\alpha \rho_\alpha^{\beta}) + \sum_{\alpha \in \{l, g\}} \nabla \cdot (\rho_\alpha^{\beta} \mathbf{q}_\alpha + \mathbf{j}_\alpha^{\beta}) = F_\beta, \quad (1)
$$

where $\Phi$ denotes the porosity, $s_\alpha$ the saturation of phase $\alpha$, $\rho_\alpha^{\beta}$ the density of component $\beta$ in phase $\alpha$, $\mathbf{q}_\alpha$ the volumetric flow rate for phase $\alpha$, $\mathbf{j}_\alpha^{\beta}$ the mass diffusion flux of component $\beta$ in phase $\alpha$, and $F_\beta$ the source term of component $\beta$. The volumetric flow rate of each phase is obtained from the Darcy–Muskat
law in the form

\[ q_\alpha = -K\lambda_\alpha(s_\alpha)\nabla p_\alpha, \quad \alpha \in \{l, g\}, \quad (2) \]

where \( K \) denotes the absolute permeability of the medium (taken to be scalar-valued for simplicity), \( \lambda_\alpha \) the mobility of phase \( \alpha \) (assumed to be a given function of the saturation \( s_\alpha \) such that \( \lambda_\alpha \) vanishes if the phase \( \alpha \) is absent), and \( p_\alpha \) the pressure of phase \( \alpha \). For simplicity, gravity forces are neglected in (2). The phase saturations take values in \([0, 1]\) (in hydrogeology models, they often take values in a subinterval of \([0, 1]\) depending on residual saturations of both phases, cf \S 4) and are such that

\[ s_l + s_g = 1. \quad (3) \]

Furthermore, assuming that the gas phase is the non-wetting phase, the phase pressures are such that

\[ p_g = p_l + \pi(s_g), \quad (4) \]

where \( \pi : [0, 1) \to [\pi(0), +\infty) \) denotes the capillary pressure (assumed to be a given function of the gas saturation). The quantity \( \pi(0) \) denotes the possibly nonzero entry pressure.

Concerning the water component, we assume incompressibility in the liquid phase and we neglect water vaporization. As a result,

\[ \rho_l^w = \rho_l^{\text{std}}, \quad \rho_g^w = 0, \quad (5) \]

where \( \rho_l^{\text{std}} \) denotes the standard water density at the given temperature of the medium. Concerning the hydrogen component, we assume the ideal gas law in the gas-phase and that hydrogen phase changes are in thermodynamic equilibrium as governed by Henry’s law. This leads to

\[ \rho_g^h = C_g p_g, \quad C_g = \frac{M^h}{RT}, \quad (6) \]

and

\[ \rho_l^h = C_h p_g, \quad C_h = HM^h, \quad (7) \]

where \( M^h \) denotes the hydrogen molar mass, \( R \) the universal gas constant, \( T \) the (absolute) temperature, and \( H \) the (temperature-dependent) constant in Henry’s law. Furthermore, the hydrogen diffusion flux in the liquid phase, \( j^h_l \), is evaluated using a Fick-type law in the form

\[ j^h_l = -\Phi s_l D^h_l \nabla \rho^h_l, \quad (8) \]

where \( D^h_l \) denotes the (temperature-dependent) molecular diffusion coefficient of hydrogen in the liquid phase. Since for a two-component system, the diffusion fluxes in the liquid phase satisfy \( \sum_{\beta \in \{w, h\}} j^\beta_l = 0 \), we infer \( j^w_l = -j^h_l \). The use of the Fick-type law (8) using density, and not concentration, gradients and neglecting cross-diffusion effects is reasonable as long as the hydrogen component is dilute in the liquid phase.

To sum up, the above simplifying assumptions allow us to recast the mass conservation equations (1) for both components as

\[ \Phi \partial_t s_l + \nabla \cdot (\rho_l^w q_l + j^h_l) = F^w, \quad (9a) \]

\[ \Phi \partial_t (\rho_l^h s_l + C_g p_g s_g) + \nabla \cdot (\rho_l^h q_l + C_g p_g q_g + j^h_l) = F^h. \quad (9b) \]
2.2 Mathematical model

Owing to the possible disappearance of the gas phase in some parts of the computational domain that are a priori unknown, it is not appropriate to select any of the saturations or the gas pressure as one of the main unknowns of the mathematical model. Indeed, the saturations vanish identically, or are identically equal to one, in those regions where only the liquid phase is present, while the gas pressure is not even defined in those regions. Following the recent ideas of Smaï ([32] (see also Bourgeat, Jurak, and Smaï [8] for a slightly different choice), we select as main unknowns of the mathematical model the liquid pressure \( p_l \) (since the liquid phase is always present throughout the domain) and the dissolved hydrogen density \( \rho_l \). In what follows, we set

\[
y = (y_1, y_2), \quad y_1 := p_l, \quad y_2 := \rho_l.
\]

The mass conservation equations (9) can then be recast into the form

\[
\begin{align*}
\partial_t b_1(y) - \nabla \cdot (A_{11}(y)\nabla y_1 + A_{12}(y)\nabla y_2) &= F_1, \\
\partial_t b_2(y) - \nabla \cdot (A_{21}(y)\nabla y_1 + A_{22}(y)\nabla y_2) &= F_2,
\end{align*}
\]

or, in more compact form, for all \( i \in \{1, 2\} \),

\[
\partial_t b_i(y) - \sum_{j \in \{1, 2\}} \nabla \cdot (A_{ij}(y)\nabla y_j) = F_i,
\]

with source terms \( F_1 := F^w \), \( F_2 := F^h \), and

\[
\begin{align*}
b_1(y) &= -\Phi \rho_l^{\text{std}} s_g(y), \\
b_2(y) &= \Phi a(s_g(y)) y_2, \\
A_{11}(y) &= \rho_l^{\text{std}} K \lambda_l (1 - s_g(y)), \\
A_{12}(y) &= -\Phi (1 - s_g(y)) D_l^h, \\
A_{21}(y) &= y_2 K \lambda_l (1 - s_g(y)), \\
A_{22}(y) &= y_2 K \lambda_q (s_g(y)) \omega C_h^{-1} + \Phi (1 - s_g(y)) D_l^h,
\end{align*}
\]

where \( \omega = \frac{C_h}{\sigma_h} \) and \( a(s) = 1 + (\omega - 1)s \). Finally, the gas saturation is recovered from

\[
s_g(y) = \pi^{-1} \left( \frac{y_2}{\sigma_h} - y_1 \right),
\]

where \( \pi^{-1} : \mathbb{R} \to [0, 1) \) denotes the extension by zero to \( \mathbb{R} \) of the inverse function of the capillary pressure \( \pi \), cf Figure [4]. We observe that \( s_g \) is a continuous function of \( y \); this function is actually differentiable if the van Genuchten model is considered for the capillary pressure, as is often the case in hydrogeology and is actually the case in our numerical experiments presented in §4 below. If the Brooks–Corey model is used instead, the function \( \pi^{-1} \) is not differentiable at the entry pressure \( \pi(0) \), and a semi-smooth version of the linearization procedure described in § 3.1 needs to be considered.

The governing equations (12), supplemented with (13) and (14), are posed in the computational domain \( \Omega \) which we assume to be a bounded, open, polyhedral domain in \( \mathbb{R}^d \), \( d \geq 1 \), with boundary denoted by \( \partial \Omega \) and outward unit normal by \( \mathbf{n} \). Dirichlet and Neumann boundary conditions are enforced at the boundary. Given a partition of \( \partial \Omega \) into \( \partial \Omega_D \cup \partial \Omega_N \), we enforce on \( \partial \Omega_D \) the value \( y = y_D \), and on \( \partial \Omega_N \), we enforce \( \mathbf{n} \cdot \sigma_i(y) = \sigma_i \) for all \( i \in \{1, 2\} \), where the total fluxes \( \sigma_i \) are defined as

\[
\sigma_i(y) = -(A_{i1}(y)\nabla y_1 + A_{i2}(y)\nabla y_2).
\]
It is possible to consider a setting where the partition into Dirichlet and Neumann boundaries depends on the component \( i \in \{1, 2\} \), but this generality is not needed for the test cases envisaged herein. Finally, initial conditions are imposed on \( y \) in the form \( y(x, 0) = y_0 \) for all \( x \in \Omega \).

Equation (11a) is a parabolic equation in \( y_1 \) (since the liquid phase is always present) that degenerates into an elliptic equation if the gas phase disappears. Equation (11b) is a parabolic equation in \( y_2 \) that degenerates into an elliptic equation if there is no dissolved hydrogen. Therefore, (11a) is loosely termed the “pressure equation” and (11b) the “hydrogen equation.” To examine the coupled system, we write it in nondimensional form. Let \( y_1 = p_0 \bar{y}_1 \) and \( y_2 = C_h p_0 \bar{y}_2 \), where \( p_0 \) denotes a reference pressure. Dividing (11a) by \( \varrho \bar{v}_i^{\text{std}} \) and (11b) by \( C_h p_0 \) yields

\[
\begin{align*}
\partial_t \bar{v}_1(\bar{y}) - \nabla \cdot (\varpi_{11}(\bar{y}) \nabla \bar{y}_1 + \varpi_{12}(\bar{y}) \nabla \bar{y}_2) &= F_1, \\
\partial_t \bar{v}_2(\bar{y}) - \nabla \cdot (\varpi_{21}(\bar{y}) \nabla \bar{y}_1 + \varpi_{22}(\bar{y}) \nabla \bar{y}_2) &= F_2,
\end{align*}
\]

with \( F_1 := F^w / \varrho \bar{v}_i^{\text{std}} \), \( F_2 := F^h / (C_h p_0) \), and letting \( \bar{\pi}_y(\bar{y}) = \pi^{-1}(p_0(\bar{y}_2 - \bar{y}_1)) \),

\[
\begin{align*}
\bar{v}_1(\bar{y}) &= -\Phi \bar{\pi}_y(\bar{y}), \\
\bar{v}_2(\bar{y}) &= \Phi a(\bar{\pi}_y(\bar{y})) \bar{y}_2, \\
\varpi_{11}(\bar{y}) &= p_0 K \lambda_1(1 - \bar{\pi}_y(\bar{y})), \\
\varpi_{12}(\bar{y}) &= -(C_h p_0 / \varrho \bar{v}_i^{\text{std}}) \Phi(1 - \bar{\pi}_y(\bar{y})) D_h^i, \\
\varpi_{21}(\bar{y}) &= \bar{y}_2 p_0 K \lambda_2(1 - \bar{\pi}_y(\bar{y})), \\
\varpi_{22}(\bar{y}) &= \bar{y}_2 p_0 K \lambda_2(1 - \bar{\pi}_y(\bar{y})) \omega + \Phi(1 - \bar{\pi}_y(\bar{y})) D_h^i,
\end{align*}
\]

where \( \bar{v}_1(\bar{y}) \) is nondimensional and \( \varpi_{ij}(\bar{y}) \) scales as m²/s. The matrix \( \varpi \) yields a positive definite quadratic form (and hence ellipticity on the space differential operator) if and only if

\[
4\varpi_{11} \varpi_{22} - (\varpi_{11} + \varpi_{12} \varpi_{21})^2 > 0. \tag{18}
\]

For applications related to hydrogen migration in underground repositories, typical values are \( p_0 = 1 \) MPa, \( C_h p_0 = 1.5 \cdot 10^{-2} \) kg/m³, and using the values listed in Table 1, we obtain the estimates (in m²/s) \( \varpi_{11} \approx 5 \cdot 10^{-11}, \varpi_{12} \approx 6.8 \cdot 10^{-15}, \varpi_{21} \approx 5 \cdot 10^{-11}, \varpi_{22} \approx 4.5 \cdot 10^{-10} \) (for \( \varpi_{22} \), only the second term is used for the estimate, but the first term can take much larger values if the gas phase is present). Hence, condition (18) holds true. We observe that \( \varpi_{12} \ll \varpi_{11} \), whence, neglecting \( \varpi_{12} \) (and using only the second term for \( \varpi_{22} \)), we infer from (18) the condition \( p_0 K \lambda_1(1) < 4 \Phi D_h^i \) for ellipticity. In situations where this condition fails (e.g., because \( K \) is too large or \( D_h^i \) too small), a smallness condition on the dissolved hydrogen density can be invoked, with an upper bound typically depending on \( K \) and \( D_h^i \). Finally, an important result under the assumption \( \varpi_{12} \approx 0 \) is that the change of variables (see Små [53, 54])

\[
\bar{y}_1 = \bar{y}_1 + \omega^{-1} e^{\omega \pi_2}, \quad \bar{y}_2 = e^{\omega \pi_2}, \tag{19}
\]
yields a new system in the variables $\{\pi_1, \pi_2\}$ fitting the framework of the Alt–Luckhaus theorem for the existence of weak solutions [2], namely the time-derivative term involves the gradient of a convex potential and the space-derivative terms yield a symmetric positive definite matrix.

### 3 Numerical method

In this section, we first present the time discretization together with the linearization procedure and then describe the dG method for space discretization. We consider the nondimensional form (16); to alleviate the notation, bars are henceforth omitted.

#### 3.1 Time discretization and linearization

Let $\{t_m\}_{0 \leq m \leq M}$ be a partition of time interval $[0, T]$ such that $\ell^0 = 0$ and $\ell^M = T$, the given simulation time, and set $\tau^m = t_m - t_{m-1}$ for $m = 1, \ldots, M$. Time discretization of (12) is achieved using the backward Euler method: Starting from the initial condition $y^0 := y_0$, we seek, for all $m = 1, \ldots, M$, the function $y^m$ such that, for all $i \in \{1, 2\}$,

$$
\frac{1}{\tau^m}(b_i(y^m) - b_i(y^{m-1})) - \sum_{j \in \{1, 2\}} \nabla \cdot (A_{ij}(y^m)\nabla y^m_j) = F_i^m,
$$

where a superscript $m$ on any problem data (e.g., on $F_i$) indicates evaluation at the discrete time $t_m$.

The linearization procedure is based on an incomplete Newton solver. It first involves a fixed-point iteration (indicated by an index $l$) on the coefficients in the diffusive terms, leading to

$$
\frac{1}{\tau^m}(b_i(y^m_{l+1}) - b_i(y^{m-1})) - \sum_{j \in \{1, 2\}} \nabla \cdot (A_{ij}(y^m_l)\nabla y^m_{j,l+1}) = F_i^m.
$$

The fixed-point iteration is initialized with the values at the previous time step, i.e., we take $y^m_0 := y^{m-1}$. The second ingredient in the linearization procedure is a linear approximation of the time-derivative terms,

$$
\frac{1}{\tau^m}(b_i(y^m_{l+1}) - b_i(y^{m-1})) = \frac{1}{\tau^m}(b_i(y^m_{l+1}) - b_i(y^m_l)) + \frac{1}{\tau^m}(b_i(y^m_l) - b_i(y^{m-1}))
\approx \partial_1 b_i(y^m_l) \frac{y^m_{1,l+1} - y^m_{1,l}}{\tau^m} + \partial_2 b_i(y^m_l) \frac{y^m_{2,l+1} - y^m_{2,l}}{\tau^m} + \frac{b_i(y^m_l) - b_i(y^{m-1})}{\tau^m}.
$$

As a result, for all $m = 1, \ldots, M$ (time loop) and for all $l \geq 0$ (fixed-point loop), we seek $y^m_{l+1}$ such that, for all $i \in \{1, 2\}$,

$$
- \sum_{j \in \{1, 2\}} \nabla \cdot (A_{ij}(y^m_l)\nabla y^m_{j,l+1}) + \sum_{j \in \{1, 2\}} \partial_1 b_i(y^m_l) \frac{y^m_{1,l+1} - y^m_{1,l}}{\tau^m} = G_i^m.
$$

Solving for $y^m_{l+1}$ amounts to solving the linear system of PDEs,

$$
- \nabla \cdot (A_{11}(y^m_l)\nabla y^m_{1,l+1} + A_{12}(y^m_l)\nabla y^m_{2,l+1}) + \frac{1}{\tau^m}(\partial_1 b_1(y^m_l)y^m_{1,l+1} + \partial_2 b_1(y^m_l)y^m_{2,l+1}) = G_{1,l},
$$

$$
- \nabla \cdot (A_{21}(y^m_l)\nabla y^m_{1,l+1} + A_{22}(y^m_l)\nabla y^m_{2,l+1}) + \frac{1}{\tau^m}(\partial_1 b_2(y^m_l)y^m_{1,l+1} + \partial_2 b_2(y^m_l)y^m_{2,l+1}) = G_{2,l}.
$$

(20a), (20b)
with the right-hand sides, for all $i \in \{1, 2\}$,

$$G_{i,l}^m = F_i^m + \frac{1}{\tau m} b_i(y^{m-1}) + \frac{1}{\tau m} (\partial_1 b_i(y^{m})y_i^m + \partial_2 b_i(y^{m})y_2^m - b_i(y^{m})).$$

(21)

together with the boundary conditions

$$y_{i+1}^m = y_D^m \quad \text{on } \partial \Omega_D,$$

(22a)

$$n \cdot \sigma_i(y_{i+1}^m) = \sigma_i^m, i \in \{1, 2\}, \text{ on } \partial \Omega_N.$$  

(22b)

### 3.2 Space discretization: dG method

Let $\{T_\delta\}_{\delta > 0}$ be a family of shape-regular meshes of the domain $\Omega$ (possibly containing hanging nodes), where $\delta$ denotes the maximum element diameter in $T_\delta$. We say that the set $F$ is a mesh interface (resp., boundary face) if $F$ has nonzero $(d - 1)$-dimensional measure and if there exist distinct $T^-, T^+ \in T_\delta$ such that $F = \partial T^- \cap \partial T^+$ (resp., if there exists $T \in T_\delta$ such that $F = \partial T \cap \partial \Omega$). Interfaces are collected in the set $\mathcal{F}_\rho^b$, boundary faces in the set $\mathcal{F}_\rho^b$, and we let $\mathcal{F}_\rho := \mathcal{F}_\rho^b \cup \mathcal{F}_\rho^d$. For all $F \in \mathcal{F}_\rho$, $\delta_F$ denotes its diameter. We suppose that the meshes $T_\delta$ are fitted to the partition $\partial \Omega = \partial \Omega_D \cup \partial \Omega_N$; accordingly, the set of boundary faces $\mathcal{F}_\rho^b$ is partitioned as $\mathcal{F}_\rho^b = \mathcal{F}_\rho^b \cup \mathcal{F}_\rho^N$ with obvious notation.

For a scalar- or vector-valued function $v$ that is possibly two-valued at an interface $F = \partial T^- \cap \partial T^+ \in \mathcal{F}_\rho^b$, we define its jump and average at $F$ as

$$[v] = v^+ - v^-, \quad \{v\} = \frac{1}{2}(v^+ + v^-), \quad v^\pm = v|_{T^\pm}$$

and extend these definitions to boundary faces $F = \partial T \cap \partial \Omega \in \mathcal{F}_\rho^b$ by setting $[v] = \{v\} = v|_T$. For all $F = \partial T^- \cap \partial T^+ \in \mathcal{F}_\rho^b$, we define $n_F$ as the unit normal vector to $F$ pointing from $T^-$ toward $T^+$, whereas for $F \in \mathcal{F}_\rho^b$ we set $n_F = n$. The sign arbitrariness in the definition of $n_F$ and of the jump across $F$, for all $F \in \mathcal{F}_\rho^b$, is irrelevant in what follows.

Let $k \geq 1$ be an integer. The dG method is based on the discrete space

$$V^k_\delta := \{v_\delta \in L^2(\Omega); \forall T \in T_\delta, v_\delta|_T \in P_k(T)\},$$

(23)

where $P_k(T)$ denotes the vector space spanned by polynomials of total degree $\leq k$ on $T$. Let $y_\delta \in [V^k_\delta]^2$ and let $i, j \in \{1, 2\}$. The interior penalty dG bilinear form discretizing the differential operator $-\nabla \cdot (A_{ij}(y_\delta)\nabla \cdot)$ can be written, for all $u_\delta, v_\delta \in V^k_\delta$, as

$$a_\delta^{ij}(y_\delta; u_\delta, v_\delta) = \sum_{T \in T_\delta} \int_T A_{ij}(y_\delta) \nabla u_\delta \cdot \nabla v_\delta$$

$$- \sum_{F \in \mathcal{F}_\rho^b \cup \mathcal{F}_\rho^d} \int_F n_F \cdot \{A_{ij}(y_\delta)\nabla u_\delta\} [v]$$

$$- \theta^{ij} \sum_{F \in \mathcal{F}_\rho^b \cup \mathcal{F}_\rho^d} \int_F n_F \cdot \{A_{ij}(y_\delta)\nabla v_\delta\} [u]$$

$$+ \sum_{F \in \mathcal{F}_\rho^b \cup \mathcal{F}_\rho^d} \eta^{ij}_F \frac{\sigma}{\delta_F} \int_F \|v_\delta\|^2 [u],$$

where $\theta^{ij} = 0$ for $i \neq j$ and $\theta^{ii} = 1$ for $i \in \{1, 2\}$ (so that the diagonal blocks of the linear system matrix remain symmetric), while $\eta^{ij} = 0$ for $i \neq j$ and $\eta^{jj} = \|A_{ii}\|_{L^\infty(\Omega)}$ for $i \in \{1, 2\}$ (in the present context, the variations of $A_{ii}$ in $\Omega$ are sufficiently mild to use a global scaling for the penalty parameter). Moreover, the user-dependent parameter $\sigma$ is typically set to 10.
We consider the following dG method: Given $y_{\delta}^{m-1} \in [V_{\delta}^k]^{L_2}$ from the previous time step (or the $L_2$-projection of the initial condition), the fixed-point loop is initialized as $y_{\delta,0}^m = y_{\delta}^{m-1}$, and, for all $l \geq 0$, we seek $y_{\delta,l+1}^m \in [V_{\delta}^k]$ such that, for all $v_\delta, w_\delta \in V_{\delta}^k$,

$$a_3^{11}(y_{\delta,l}^m; y_{1,\delta,l+1}^m, v_\delta) + a_3^{12}(y_{\delta,l}^m; y_{2,\delta,l+1}^m, v_\delta) + \frac{1}{\tau^m} \int_{\Omega} \left( \partial_1 b_1(y_{\delta,l}^m) y_{1,\delta,l+1}^m + \partial_2 b_1(y_{\delta,l}^m) y_{2,\delta,l+1}^m \right) v_\delta$$

$$= \int_{\Omega} G_{1,1}^m v_\delta - \sum_{F \in F_\delta^1} \int_F \sigma_1^m v_\delta$$

$$+ \sum_{F \in F_\delta^1} \int_F \left( \phi_{F}^{11}(y_{\delta,l}^m; v_\delta) y_{1,D}^m + \phi_{F}^{12}(y_{\delta,l}^m; v_\delta) y_{2,D}^m \right),$$

(24)

$$a_3^{21}(y_{\delta,l}^m; y_{1,\delta,l+1}^m, w_\delta) + a_3^{22}(y_{\delta,l}^m; y_{2,\delta,l+1}^m, w_\delta) + \frac{1}{\tau^m} \int_{\Omega} \left( \partial_1 b_2(y_{\delta,l}^m) y_{1,\delta,l+1}^m + \partial_2 b_2(y_{\delta,l}^m) y_{2,\delta,l+1}^m \right) w_\delta$$

$$= \int_{\Omega} G_{2,1}^m w_\delta - \sum_{F \in F_\delta^1} \int_F \sigma_2^m w_\delta$$

$$+ \sum_{F \in F_\delta^1} \int_F \left( \phi_{F}^{21}(y_{\delta,l}^m; w_\delta) y_{1,D}^m + \phi_{F}^{22}(y_{\delta,l}^m; w_\delta) y_{2,D}^m \right),$$

(25)

with $\phi_{F}^{ij}(y_{\delta}; v_\delta) = -\theta^{ij} n_F \cdot A_{ij}(y_{\delta}) \nabla v_{\delta} + \eta_i^{ij} \frac{\partial k}{\partial y} v_{\delta}$, for all $i, j \in \{1, 2\}$. Integrating by parts elementwise the volume term in the bilinear forms $a_3^{ij}$, it is seen that the above formulation weakly enforces the PDEs (11) in each mesh element, the boundary conditions (22) on all boundary faces, and, on all mesh interfaces $F \in F_\delta^1$, the transmission conditions

$$[y_{\delta}] = 0, \quad n_F \cdot [\sigma_i(y_{\delta})] = 0, \quad \forall i \in \{1, 2\},$$

(26)

consistently with the properties satisfied by the exact solution.

**Remark 3.1.** For heterogeneous media consisting of different rocks with contrasting properties, weighted averages can be considered in the formulation of the dG method, as in [7], for linear advection-diffusion equations. Furthermore, we observe that (26) implies that the dissolved hydrogen density $\rho_{H_2}$ is continuous at any interface. Owing to Henry’s law, the gas pressure (if this phase exists on both sides of the interface) is also continuous; hence, since the liquid pressure is continuous (also by (25)), the capillary pressure is also continuous at the interface. However, the gas saturation is in general discontinuous at the interface; this fact is illustrated in the test case of [4,5]. Instead, for two-component porous media flows without inter-phase mass exchange where one of the saturations is used as one of the main unknowns, the penalty strategy in the dG method must be revised to enforce weakly a nonlinear jump condition on the saturation, cf [10].

### 4 Numerical results

In this section, we evaluate numerically the dG method derived in §3 on three one-dimensional test cases, dedicated respectively to gas-phase (dis)appearance, ill-prepared initial conditions, and heterogeneous problems with different rock types. The first two test cases have been proposed within the GNR MoMaS in connection with the Couplex-Gas benchmark proposed by the French National Radioactive Waste Management Agency (ANDRA); reference solutions can be found in [19]. The third test case has been investigated in [9].
We use the van Genuchten model for capillary pressure and the van Genuchten–Mualem model for relative permeability, i.e.,

$$\pi(s_g) = P_h s_{le}(s_g)^{-1/m} - 1 \right)^{1/n},$$  \hspace{1cm} (27a)

$$\lambda_l(s_g) = \frac{1}{\mu_l} s_{le}(s_g)^{1/2} \left( 1 - (1 - s_{le}(s_g)^{1/m})^m \right)^2,$$ \hspace{1cm} (27b)

$$\lambda_g(s_g) = \frac{1}{\mu_g} s_{ge}(s_g)^{1/2} \left( 1 - s_{le}(s)^{1/m} \right)^2,$$ \hspace{1cm} (27c)

where $m = 1 - \frac{1}{n}$, $\mu_l$ and $\mu_g$ are viscosities, while

$$s_{ge}(s_g) = \frac{s_g - s_{gr}}{1 - s_{lr} - s_{gr}}, \quad s_{le}(s_g) = 1 - s_{ge}(s_g),$$ \hspace{1cm} (28)

are the relative gas and liquid saturations and $s_{ge}$, $s_{lr}$ are the residual saturations in the gas and liquid phases respectively. Values for the above physical parameters are specified below for each test case. In all cases, the convergence criterion in the incomplete Newton method uses a tolerance of $10^{-8}$ in the $L^2(\Omega)$-norm, and, in the present one-dimensional setting, a direct solver is used for the linear systems.

### 4.1 Gas-phase (dis)appearance

In this test case, hydrogen is injected into the porous medium $\Omega = (0, 200)$ (m) initially saturated by water. Injection is performed through the left boundary $x = 0$ during the injection time $T\text{inj} = 10^5$ (years), while the simulation time is $T = 10^6$ (years). There are no external sources, i.e., $F^w = F^h = 0$. This example illustrates the potential of the method to simulate gas-phase appearance and disappearance related to hydrogen injection.

<table>
<thead>
<tr>
<th>Porous medium</th>
<th>Fluid characteristics</th>
</tr>
</thead>
<tbody>
<tr>
<td>Param</td>
<td>Value</td>
</tr>
<tr>
<td>$\Phi$</td>
<td>0.15 (-)</td>
</tr>
<tr>
<td>$K$</td>
<td>$5 \times 10^{-20}$ m$^2$</td>
</tr>
<tr>
<td>$P_h$</td>
<td>$2 \times 10^6$ Pa</td>
</tr>
<tr>
<td>$n$</td>
<td>1.49 (-)</td>
</tr>
<tr>
<td>$s_{lr}$</td>
<td>0.4 (-)</td>
</tr>
<tr>
<td>$s_{gr}$</td>
<td>0 (-)</td>
</tr>
</tbody>
</table>

Table 1: Parameter values for the porous medium and fluid characteristics used in test case 1.

The porous medium and fluid characteristics are presented in Table 1. Initial and boundary conditions are given by

$$-\mathbf{n} \cdot \mathbf{\sigma}_1|_{x=0} = 0, \quad -\mathbf{n} \cdot \mathbf{\sigma}_2|_{x=0} = q_{\text{inj}};$$ \hspace{1cm} (29)

$$p_l|_{x=200} = 10^6 \text{ (Pa)}, \quad q^h_{l}|_{x=200} = 0;$$ \hspace{1cm} (30)

$$p_l|_{t=0} = 10^6 \text{ (Pa)}, \quad q^h_{l}|_{t=0} = 0;$$ \hspace{1cm} (31)

where $q_{\text{inj}} = 5.57 \times 10^{-6} \chi_{[0,T_{\text{inj}}]}(t)$ (kg/m$^2$/year) and $\chi_A$ denotes the characteristic function of the set $A$. The ellipticity criterion (18) can be verified at each point in space and in time; the left-hand side of (18) takes values in the range $[1, 8] \times 10^{-4} / (m^2/year)^2$.

We consider the first-order dG space $V_h^1$ ($k = 1$) and use a uniform mesh of 200 elements in $\Omega$. In time, we consider the partition $[0, 1, 2, 3, 5, 7, 10] \times 10^5$ (years) of the time interval $[0, T]$ and use the time steps $\tau = [125, 500, 1000, 5000, 10000, 50000]$ (years) within each time slab.
Figures 2 and 3 present selected results of our simulations at various times. During an initial transient up to \(\approx 2 \times 10^3\) (years), the dissolved hydrogen molar density increases owing to gas injection, while the liquid pressure remains roughly constant, and there is no gas phase. The gas phase appears at the time \(\approx 1.3 \times 10^4\) (years), and the liquid pressure begins to increase until it reaches its maximum at the time \(\approx 10^5\) (years). The gas saturation continues to increase up to the time \(\approx 5 \times 10^5\) (years), while the gas pressure reaches its maximum at \(x = 0\) at the time \(\approx 10^5\) (years) and decreases afterward. When the gas injection is eventually stopped, the liquid pressure starts to decrease (owing to the absence of entering water flux at \(x = 0\)) and reaches its minimum at the time \(\approx 5.6 \times 10^5\) (years). Afterward, the water pressure increases again, the dissolved hydrogen molar density and the gas saturation decrease, while the system has almost reached equilibrium at the final simulation time. All these results agree well with those presented in [19, 32].

### 4.2 Ill-prepared initial condition

In this test case, we consider the domain \(\Omega = (0, 1)\) (m) with zero flux boundary conditions and no external sources, while uniform liquid pressure and discontinuous gas pressure are considered as initial conditions. Owing to the discontinuity of the gas pressure, the system is initially out of the mechanical equilibrium. This example illustrates the potential of the method to simulate the evolution of the system back to equilibrium starting from an ill-prepared initial condition.

![Table 2: Parameter values for the porous medium and fluid characteristics used in test case 2.](image)

<table>
<thead>
<tr>
<th>Porous medium</th>
<th>Fluid characteristics</th>
</tr>
</thead>
<tbody>
<tr>
<td>Param. Value</td>
<td>Param. Value</td>
</tr>
<tr>
<td>(\Phi)</td>
<td>(D_l)</td>
</tr>
<tr>
<td>(K)</td>
<td>(\mu_1)</td>
</tr>
<tr>
<td>(P_r)</td>
<td>(\nu_g)</td>
</tr>
<tr>
<td>(n)</td>
<td>(H(303K))</td>
</tr>
<tr>
<td>(s_{gr})</td>
<td>(M^h)</td>
</tr>
<tr>
<td>(s_{g,r})</td>
<td>(\sigma_1^{ad})</td>
</tr>
</tbody>
</table>

The left-hand side of (13) takes values in the range \([6, 9] \times 10^3\) (m²/year²).

We consider the first-order dG space \(V_h^1\) (\(k = 1\)) and use a uniform mesh of 512 elements in \(\Omega\). The simulation time is \(T = 10^6\) (s). We consider the partition \(T = [0, 10^2, 5 \times 10^2, 10^3, 5 \times 10^3, 10^4, 2.1 \times 10^5, 5 \times 10^5, 10^6]\) (s) of the time interval \([0, T]\) and use 32 uniform time steps within each time slab.

Figures 4 and 5 present selected results of our simulations at various times. The gas phase is initially present in the whole domain. Owing to the initial discontinuity in the gas pressure, the gas saturation and the dissolved hydrogen density are also discontinuous initially. After a short time (10 s), the liquid pressure exhibits a jump of the order of 0.8 MPa. Then, a liquid pressure front propagates and reaches the boundaries at \(\approx 10^2\) seconds. Afterward, up to the time \(\approx 10^4\) seconds, the liquid pressure increases at the left boundary and decreases at the right boundary. At the time \(\approx 10^4\) (s), the difference of left and right boundary values is of the order of 1 MPa, and the liquid pressure still exhibits a sharp gradient at \(x = 0.5\). Next, the difference between boundary values begins to decrease, and the liquid pressure smoothly
converges to its equilibrium position, which it has almost reached at time $10^6$ (s). Furthermore, the initially discontinuous profile of dissolved hydrogen density is smoothed owing to fast hydrogen diffusion in water. Thus, this profile reaches equilibrium much earlier than the liquid pressure, at time $\approx 10^4$ (s). Finally, the gas saturation profile exhibits a very steep front up to the time $\approx 10^3$ (s); then, the shock slowly dissipates, but the profile still exhibits a sizable gradient at time $\approx 10^5$ (s). The gas saturation profile has almost relaxed back to equilibrium at the final simulation time. All these results agree well with those reported in [19].

4.3 Hydrogen injection in heterogeneous medium

In this test case, we consider the heterogeneous porous medium $\Omega = (0, 200)$ (m) composed of two rocks occupying respectively the subdomains $\Omega_1 = (0, 20)$ and $\Omega_2 = (20, 200)$. The porous medium and fluid characteristics are presented in Table 3. The column for the porous medium contains two values, one for each rock type. We observe that the rock occupying the subdomain $\Omega_2$ has a finer texture.

As in test case 1, the initial and boundary conditions are given by (29), except that the gas injection is not stopped at time $T_{\text{inj}}$. This example illustrates the potential of the method to handle heterogeneous porous media. The left-hand side of (18) takes values in the range $[10^{-4}, 1]$ (m$^2$/year)$^2$.

<table>
<thead>
<tr>
<th>Porous medium</th>
<th>Fluid characteristics</th>
</tr>
</thead>
<tbody>
<tr>
<td>Par. Value</td>
<td>Par. Value</td>
</tr>
<tr>
<td>$\Phi$ (0.3, 0.15) (-)</td>
<td>$D^h_1$ 3 $10^{-9}$ m$^2$/s</td>
</tr>
<tr>
<td>$K$ (10$^{-18}$, 5 10$^{-20}$) m$^2$</td>
<td>$\mu_1$ 1 $10^{-3}$ Pa s</td>
</tr>
<tr>
<td>$P_c$ (2 10$^8$, 15 10$^8$) Pa</td>
<td>$\mu_2$ 9 $10^{-6}$ Pa s</td>
</tr>
<tr>
<td>$n$ (1.54, 1.49) (-)</td>
<td>$H(303K)$ 7.65 $10^{-6}$ mol/Pa/m$^3$</td>
</tr>
<tr>
<td>$\rho_0$ (0.01, 0.4) (-)</td>
<td>$M_h$ 2 $10^{-3}$ kg/mol</td>
</tr>
<tr>
<td>$\rho_c$ (0.0) (-)</td>
<td>$\rho_{\text{std}}$ 10$^3$ kg/m$^3$</td>
</tr>
</tbody>
</table>

Table 3: Parameter values for the porous medium and fluid characteristics used in test case 3.

We consider the first-order dG space $V^1_\delta (k = 1)$ and use a uniform mesh of 16 elements in $\Omega_1$ and 144 elements in $\Omega_2$ so that the mesh is fitted to the interface separating $\Omega_1$ and $\Omega_2$. The simulation time is $T = 10^6$ (years). We consider the partition $[0, 6 \cdot 10^4, 7 \cdot 10^4, 10^6]$ (years) of the time interval $[0, T]$ and use the time steps $\tau = [200, 1000, 2000, 20000]$ (years) within each time slab. We recall that the dG method weakly enforces the continuity of the normal component of the total fluxes and that of the liquid pressure and of the dissolved hydrogen density, cf [29].

Figures 6 presents selected results of our simulations at various times. Owing to gas injection, the gas phase appears in the first subdomain at the time $\approx 3 \cdot 10^4$ (years) and reaches the interface separating the two rock types at the time $\approx 4.2 \cdot 10^4$ (years). The saturation at the left of the interface (in the coarser rock) starts to increase while maintaining the jump in the saturation such that capillary pressure continuity is preserved, and the gas phase penetrates into the second (finer) rock. Next, at the time $\approx 1.3 \cdot 10^5$ (years), the liquid pressure starts to decrease, and the system relaxes back to equilibrium, as reflected by the fact that the net difference of total hydrogen flow at inflow and outflow boundaries tends to zero. We stress that the capillary pressure is continuous at the interface at all times. The results are in good agreement with those reported in [31].

4.4 Conclusions

The above numerical results show that the proposed methodology is capable of delivering accurate discrete solutions capturing the correct physical behavior in various complex situations. Future work can aim at assessing the proposed methodology on higher-dimensional test cases and at analyzing the discrete scheme to derive stability and error estimates.

Acknowledgment. The authors are grateful to Farid Smaï (IRSN) for fruitful discussions.
References


Figure 2: Liquid pressure $p_l$ (top line, MPa), dissolved hydrogen molar density $\rho_l^h/M^h$ (second line, mol/m$^3$), and gas saturation $s_g$ (bottom line, %) at times $\{2 \cdot 10^3, 1.4 \cdot 10^4, 10^5, 1.4 \cdot 10^5, 5 \cdot 10^5\}$ (left column, years) and $\{5.1 \cdot 10^5, 5.6 \cdot 10^5, 6.1 \cdot 10^5, 6.6 \cdot 10^5, 10^6\}$ (right column, years).
Figure 3: From top to bottom and from left to right, as a function of time: liquid and gas pressures $p_l$ and $p_g$ at $x = 0$ (MPa), gas saturation $s_g$ at $x = 0$ (%), front position of the gas phase (m), total water flux at $x = 200$ (kg/m²/year), and total hydrogen flux at $x = 200$ (kg/m²/year).
Figure 4: Liquid pressure $p_l$ (top line, MPa), dissolved hydrogen molar density $\rho_{lh}/M_h$ (second line, mol/m$^3$), and gas saturation $s_g$ (bottom line, %) at times $\{10, 10^2, 5 \cdot 10^2, 10^3\}$ (left column, s) and $\{5 \cdot 10^3, 10^4, 5 \cdot 10^4, 10^5\}$ (right column, s).
Figure 5: Liquid pressure $p_l$ (MPa), dissolved hydrogen molar density $\rho_h^l/M_h^l$ (mol/m$^3$), and gas saturation $s_g$ (%) at times $\{2 \cdot 10^5, 5 \cdot 10^5, 10^6\}$ (s).
Figure 6: Liquid pressure $p_l$ (MPa), dissolved hydrogen molar density $\rho_h/M_h$ (mol/m$^3$), gas saturation $s_g$ (%), and capillary pressure $\pi$ (MPa) at times $\{3 \cdot 10^4, 4.2 \cdot 10^4, 1.3 \cdot 10^5, 10^6\}$ (years).