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Questa è la Versione finale referata (Post print/Accepted manuscript) della seguente pubblicazione:

Original Citation:

BIOMASS GROWTH IN UNSATURATED POROUS MEDIA: HYDRAULIC PROPERTIES CHANGES / BORSI I; FARINA A; FASANO A; M. PRIMICERIO. - STAMPA. - (2007), pp. 196-207. [10.1142/9789812709394_0018]

Availability:

This version is available at: 2158/258636 since:

Publisher: World Scientific

Published version: DOI: 10.1142/9789812709394_0018

Terms of use: Open Access

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BIOMASS GROWTH IN UNSATURATED POROUS MEDIA: HYDRAULIC PROPERTIES CHANGES

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We present a model to describe the biomass growth process taking place in an unsaturated porous medium during a bioremediation process. We focus on the so-called column experiment. At the initial time biomass and polluted water is inoculated in the column. The subsequent changes of hydraulic properties are analyzed. We also show some preliminary simulations.

Keywords: Microbial growth; Porous media; Bioremediation.

1. Introduction

The effect of the microbial growth on hydraulic properties of porous media is a topic studied in the framework of many applications, e.g. oil recovery, wastewater treatment, bioremediation, etc. (see [1]).

Studies on flow through porous media in presence of biomass growth are presented in the papers by Rockhold et al. [2–4]. As stated there, additional work is needed for modelling unsaturated condition.

The objective of this study is thus to analyze the flow through a contaminated unsaturated porous medium in presence of biomass growth processes which induce changes in the hydraulic properties of the medium itself.

In this paper we focus only on *anaerobic* processes, namely the model we develop does not account for the O_2 consumption and diffusion.

2. Problem description and physical assumptions

We consider a vertical column (whose high is $L \sim 1 m$) of an unsaturated contaminated soil which represents a "laboratory scale" of a real vadose zone (the so-called "column experiment", see Fig. 1). The physical model is developed considering a 1-D approximation, so that x denotes the vertical





Fig. 1. A schematic of the column experiment.

coordinate of the column, pointing upwards.

At the initial time (t = 0) the saturation degree of the medium corresponds to the steady state. Then we inoculate biomass and (possibly) polluted water through the top surface. Our goal is to model the evolution of the biomass, pollutant concentration and hydraulic properties of the soil as well.

Hereafter we list the most significant physical assumptions (see also [5]):

- A.1 The soil is a homogeneous, unsaturated, rigid porous medium.
- A.2 The liquid phase which shares the empty space with air is composed by water (main component). We shall neglect the liquid density variation.
- A.3 The pollutant is dissolved in water and adsorbed onto the soil grains. Moreover, the dissolved pollutant (below a certain concentration) acts as nutrient for the bacteria (*bio-reduction*), but above a certain threshold may become a toxic agent (see [6]).
- A.4 The biomass is distributed either in water as suspension ("free biomass") or attached on the soil grains ("attached biomass"). In particular, there is no cluster formation in the free biomass. The mass of the single bacterium is known and denoted by \mathfrak{m}_b .
- A.5 The attached biomass forms *porous clusters* so that the liquid can diffuse through them. The clusters porosity is a known constant denoted by ε_b . The pores are saturated at all times. Moreover, the concentration of the pollutant in the "entrapped water" equals the one of the "free water" (see [3], for instance). The number of bacteria that forms the unit mass of the attached biomass is a known constant denoted by N^* , $[N^*] = Kg^{-1}$. Of course, N^* , ε_b and \mathfrak{m}_b depend on the type of bacteria which are present inside the column.
- A.6 We consider the attachment of free biomass on the clusters, but we neglect the inverse process (i.e. we neglect detachment). Indeed the

experiments show that detachment is mainly due to the mechanical action caused by the "fast" water flux, [10].

A.7 The concentration of pollutant and bacteria in the liquid phase is low (few ppm).

Free biomass and attached biomass are responsible for different effects changing hydraulic properties. More precisely:

- E.1 The free biomass causes essentially viscosity and surface tension variations.
- E.2 The attached biomass growth causes medium porosity variations and affects the contact angle.
- E.3 The above variations induce, in turn, changes in the permeability and in the relative saturation of the medium.

3. Notations and basic equations

We introduce the following quantities:

- ε_0 , $[\varepsilon_0] = [-]$, initial porosity of the column (known parameter).
- ε_b , $[\varepsilon_b] = [-]$, clusters porosity (considered a known constant).
- ϕ^f , volume fraction occupied by the liquid and the gaseous (air) phase.
- ϕ^c , volume fraction occupied by the clusters.
- σ , $[\sigma] = [-]$ liquid phase saturation.

•
$$\theta_{lm} = \frac{\text{volume of "mobile" liquid}}{\text{porous medium volume}}, [\theta_{lm}] = [-]. \text{ In particular, } \theta_{lm} = \sigma \phi^f.$$

•
$$\theta_{lb} = \varepsilon_b \phi^c = \frac{1}{\text{porous medium volume}}, \ [\theta_{lb}] = [-].$$

We thus have that

$$\phi^f + \phi^c = \varepsilon_0$$

So, the volume fraction occupied by the liquid (accounting for mobile and stored) is

$$\theta_{lm} + \theta_{lb} = \sigma \phi^f + \varepsilon_b \phi^c = (\sigma - \varepsilon_b) \phi^f + \varepsilon_b \varepsilon_0. \tag{1}$$

The dependent variables which have to be determined are:

- $\sigma(x, t)$, the liquid phase saturation.
- $\phi^c(x,t)$, or alternatively $\phi^f = \varepsilon_0 \phi^c$.
- $N_l(x,t) = \frac{\text{number of free bacteria in the liquid phase}}{1}$ $[N_l] = Kg^{-1}.$ unit mass of free liquid • $w_A(x,t) = \frac{\text{mass of pollutant dissolved in the liquid phase}}{\text{mass of pollutant dissolved in the liquid phase}}$
- unit mass of free liquid

• $w_s(x,t) = \frac{\text{mass of pollutant adsorbed onto soil grains}}{\text{unit mass of solid matrix}}.$

Darcy's law and Richards equation

We define the water pressure P and introduce the *capillary pressure* P_c and the *pressure head* ψ , setting

$$P_c = P_{air} - P = -P, \qquad \psi = \frac{-P}{\rho g} = \frac{P_c}{\rho g},$$

since, as usual, P_{air} has been rescaled to 0. The well-known Darcy's law describes the *specific discharge* q,

$$q = -\frac{\rho g \mathcal{K}_{sat}}{\mu} k_{rel} \left(\frac{\partial \psi}{\partial x} + 1\right),$$

where ρ water density, g gravity acceleration and

- $\mathcal{K}_{sat} = \mathcal{K}_{sat}(\phi^f)$, saturated permeability, $[\mathcal{K}_{sat}] = m^2$. For instance, we mention the Kozeny-Carman formula [11].
- $k_{rel} = k_{rel}(\psi), [k_{rel}] = [-], relative permeability (see [14]).$
- μ is the liquid phase viscosity. We assume the following law (based on experimental observations [2])

$$\mu = \tilde{\mu}(N_l) = \mu_0 \left(1 + b_3 N_l\right), \tag{2}$$

with b_3 empirical parameter and μ_0 water viscosity (i.e. the liquid viscosity in absence of biomass).

Next, the Richards' equation [14] reads as

$$\frac{\partial}{\partial t} \left[\theta_{lm} + \theta_{lb} \right] = -\frac{\partial q}{\partial x}.$$
(3)

Now, introducing the saturation curve $\sigma = \tilde{\sigma}(\psi)$, we have $\theta_{lm} = \phi^f \tilde{\sigma}(\psi)$ so that, exploting (1), the mass conservation (3) rewrites as

$$\frac{\partial}{\partial t} \left[(\sigma - \varepsilon_b) \phi^f \right] = -\frac{\partial q}{\partial x}.$$
(4)

Evolution equation for the attached biomass phase According to the literature (see for instance, [7] $\S 2.3$ and [12]) we set

$$\frac{\partial \phi^c}{\partial t} = \underbrace{(c_1 N^\star) \left[\varepsilon_0 f \left(w_A \varepsilon_b \phi^c\right) - \phi^c\right] \phi^c}_{\text{biomass "bulk growth"}}$$
(5)

$$+\underbrace{\lambda\mathfrak{H}\left(w_{s}-w_{A}\right)N_{l}\theta_{lm}}_{\text{attachment}},$$

where:

- $[c_1] = Kg \, s^{-1}$.
- $w_A \varepsilon_b \phi^c$ is the amount of nutrient available for the attached biomass.
- $\varepsilon_0 f(w_A \varepsilon_b \phi^c)$, is a modified form of the *carrying capacity* (see [13] § 1.2). Actually, ε_0 is the maximum volume fraction allowed for the attached biomass and it is "modulated" by the function $f, 0 \leq f \leq 1$, which accounts of both amount of nutrient and toxic effects (see assumption A.3).
- $\mathfrak{H}(\cdot)$ is the Heaviside function. We notice that the attachment term could be multiplied also by a function of ϕ^c , i.e. an "effective surface" term modeling the so-called *collector* and *collision* (or *sticking*) efficiencies.

Evolution equation for the free biomass

The free biomass is a component of the liquid phase. Therefore, following [8,9], the evolution equation for N_l is

$$\frac{\partial}{\partial t} \left(N_{l} \theta_{lm} \right) = \underbrace{-\frac{\partial}{\partial x} \left[q \, N_{l} \right] + \frac{\partial}{\partial x} \left[a_{L} \left| q \right| \frac{\partial N_{l}}{\partial x} \right]}_{\text{advection/dispersion}} + \underbrace{c_{1} \left[N_{max} f(w_{A}) - N_{l} \right] N_{l} \theta_{lm}}_{\text{free biomass growth}}$$
(6)

 $\underbrace{\lambda \mathfrak{H} \left(w_s - w_A \right) N_l \theta_{lm}}_{\text{attachment}}$

The first term in (6) is the divergence of the advective flux $\mathbf{J}_{adv} = -qN_l$. The second term represents the dispersive flux \mathbf{J}_{disp} , which in 1–D setting has the following form

$$\mathbf{J}_{disp} = -\sigma \phi^f D, \quad D = \frac{a_L |q|}{\sigma \phi^f},$$

with a_L longitudinal dispersion coefficient (see [14]). The "bulk growth" of the free biomass is modeled as in the case of the attached biomass, i.e. by means of a logistic-type dynamics. Here N_{max} is the equilibrium value. We set

$$N_{max} = \mathfrak{n}N^{\star}, \quad 0 < \mathfrak{n} < 1.$$

Equation for the adsorbed pollutant.

We describe the dynamics of w_s by the following equation

$$\frac{\partial w_S}{\partial t} = \underbrace{h_A w_A \varepsilon_b \phi^c(w^\star - w_S)}_{\text{adsorption term}}$$
(7)

$$-\underbrace{h_{DB} N^* \phi^c w_S}_{\text{bio-reduction term}},$$

stipulating, essentially, that only two effects are important: adsorption (desorption) and bio-reduction. In particular,

- $w_A \varepsilon_b \phi^c = \varepsilon_b \theta_{lb}$ is the amount of pollutant dissolved inside the biomass clusters.
- w^* is maximum concentration of pollutant (known parameter) which can be adsorbed by the soil.
- h_A , $[h_A] = s^{-1}$, is the adsorption/desorption rate per unit concentration.
- h_{DB} , $[h_{DB}] = Kg s^{-1}$, bio-reduction rate per unit mass.

Equation for w_A .

The total amount of pollutant (per unit volume of porous medium) dissolved in the free and "entrapped" water is $w_A(\theta_{lm} + \theta_{lb})$, i.e. $w_A(\theta_{lm} + \varepsilon_b \phi^c)$. According to [9], we write for it the following equation

$$\frac{\partial}{\partial t} \left[w_A(\theta_{lm} + \varepsilon_b \phi^b) \right] = -\underbrace{\frac{\partial}{\partial x} \left[q \, w_A \right]}_{A} + \frac{\partial}{\partial x} \left[a_L \left| q \right| \frac{\partial w_A}{\partial x} \right]$$

advection/dispersion term

$$-\underbrace{h_A w_A \varepsilon_b \phi^c(w^* - w_S)}_{\text{adsorption term}}$$
(8)

$$-\underbrace{h_{DB}\left[\theta_{lm}N_{l}+\phi^{c}N^{\star}\right]w_{A}\left(\theta_{lm}+\varepsilon_{b}\phi^{c}\right)}_{\text{bio-reduction term}}.$$

The bio–reduction term in (8) depends on the amount of bacteria which are present either in the liquid and in the clusters.

Summarizing, we have to solve the system of the governing equations (4), (5), (6), (7) and (8), which has to be endowed with initial and boundary conditions.

4. A simplified approach: biofilm and fluid media scaling

We now illustrate the basic idea of an approach to simplify the problem. The key point is to consider a porous medium constituted by a network of capillary tubes distributed uniformly in space. Next, we assume that the attached biomass phase forms a uniform layer (*biofilm*) completely coating the internal surfaces of the capillary tubes.

Focusing now on a single capillary tube, we compare two scenarios: capillary tube partially filled with "pure" water and capillary tube whose walls are coated by the biofilm and partially filled by a liquid whose components are water, bacteria and pollutant.

Denoting by p_c and $p_{c, bio}$ the capillary pressures which refer to the above scenarios, we may write the following Laplace formulas

$$p_c = \frac{2\gamma \cos \alpha}{R}, \quad p_{c,bio} = \frac{2\gamma_{bio} \cos(\alpha_{bio})}{R_{bio}},$$

where R and R_{bio} denote the capillary radii, γ and γ_{bio} are the surface tensions and α , α_{bio} are the contact angles. Therefore,

$$\frac{p_{c,bio}}{p_c} = \frac{\gamma_{bio}}{\gamma} \frac{\cos \alpha_{bio}}{\cos \alpha} \frac{R}{R_{bio}}$$

We now assume that the above formula holds true also for the averaged quantities, i.e.

$$\frac{P_{c,bio}}{P_c} = \frac{\gamma_{bio}}{\gamma} \frac{\cos \alpha_{bio}}{\cos \alpha} \frac{\langle R \rangle}{\langle R_{bio} \rangle},\tag{9}$$

where $P_{c,bio} = \langle p_{c,bio} \rangle$, $P_c = \langle p_c \rangle$, since $\langle \cdot \rangle$ denotes the R.E.V. average. Now, selecting appropriate constitutive equations^a for

$$\frac{\cos \alpha_{bio}}{\cos \alpha}, \frac{\gamma_{bio}}{\gamma} \text{ and } \frac{\langle R \rangle}{\langle R_{bio} \rangle}$$

we can define the parameter $\beta = \beta (\phi^f, N_l)$, (called *scaling factor* for the capillary pressure), such that

$$\frac{P_{c,bio}}{P_c} = \beta \left(\phi^f, N_l \right), \quad \Rightarrow \quad \psi = \beta \left(\phi^f, N_l \right) \psi_0, \tag{10}$$

where $\psi_0 = -\frac{P_c}{\rho g}$ is the pressure head in absence of biomass. Therefore,

$$\theta_{lm} = \tilde{\sigma} \left(\beta \left(\phi^f, N_l \right) \psi_0 \right) \left(\varepsilon_0 - \phi^c \right).$$

7

^aSuch equations strongly depend on the intrisic geometry of the medium, see [5].

Such an approach (often called *fluid media scaling*, see [15]), presents an evident advantage: once the flow problem is specified, we determine ψ_0 using the "classical" Richards' equation, i.e. equation (4) where the term due to porosity changes is absent. As second step we evaluate ψ exploiting (10) and q through (2). Therefore, the mathematical problem is strongly simplified and its numerical solving turns out easier. Of course, the fluid media scaling suffers from an evident drawback: ψ and q do not fulfill the Richards' equation, i.e. mass conservation. Hence, such a property needs to be tested a posteriori.

5. Numerical simulations

In this section we present few numerical simulations we worked out considering the simplified model. A deeper analysis of the results can be found in the forthcoming paper [5]. Our main goals are: (i) to show that the found solution satisfies (within a suitable tolerance) the Richards' equation; (ii) to show that the results obtained agree, at least qualitatively, with the experimental data; (iii) to put in evidence that, in certain circumstances, variation of porosity and hydraulic properties is significant.

5.1. Problem setting

The PDEs system was solved in a 1D domain. Concerning the soil, we used the well-known van-Genuchten and Mualem forms for the saturation and permeability curve, respectively (see [14]). Moreover, a Cozeny-Karman function for the saturated permeability $k_{sat}(\phi^f)$ has been selected (see [11]). Finally, we run the simulation considering $T_{max} = 7 \, day$ as maximum time of the process.

Initial conditions

The initial stage of the experiment is characterized by absence of attached biomass and a given pollutant concentrations. Hence we set

$$w_A(x,0) = w_A^0, \quad w_S(x,0) = w_S^0, \quad N_l(x,0) = 0, \quad \phi^c(x,0) = 0,$$

with $w_A^0 = 0.3$ and $w_S^0 = 0.7$.

Boundary conditions

Following [4] we set on the column bottom, x = 0,

$$\left. \frac{\partial N_l}{\partial x} \right|_{x=0} = 0, \quad \left. \frac{\partial w_A}{\partial x} \right|_{x=0} = 0.$$

On the column top, x = 1 water (with pollutant and bacteria) is inoculated in the medium. We stipulate that pollutant and bacteria concentration in the inflow water are known, hence,

$$N_l(1,t) = N_1(t), \quad w_A(1,t) = w_{A,1}(t).$$

5.2. Simulation results

First, we mention the computed value of the quantity

$$\max_{\substack{(x,t)\in[0,1]\times[0,t_c]}} \left| \frac{\partial(\theta_{lm}+\theta_{lb})}{\partial t} + \frac{\partial q}{\partial x} \right| \sim 3.8 \times 10^{-7}$$

which definitively shows that the computed solution $\psi = \beta \psi_0$ satisfies the Richards' equation.

Moreover, in Fig. 5.2-5.2 we report the plot of the most significant quantities computed during the simulation. All the values are plotted at initial, intermedium and final time step.

Further comments on the obtained results will be reported in the forthcoming work [5].

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Fig. 2. Plot of the scaling factor β at different time values.



Fig. 3. Pressure head ψ at different times.



Fig. 4. Moisture content $(\theta_{lm} + \theta_{lb})$ at different times.





Fig. 5. Relative permeability $k_{rel} \mbox{ at different times}.$