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WAVELETS IN THE TRANSPORT THEORY OF HETEROGENEOUS REACTING SOLUTES

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In this paper we consider the one-dimensional convection (advection) dispersion equation of the transport theory of heterogeneous reacting solutes in porous media. A wavelet solution is in the framework of multi-resolution analysis.

Рассматривается дисперсионное уравнение одномерной конвекции (адвекции) теории транспорта гетерогенных реагирующих растворенных веществ в пористых средах. Вейвлетное решение получено в рамках мультирезольвентного анализа.

Розглядається дісперсійне рівняння одновимірної конвекції (адвекції) теорії транспорту гетерогенних реагуючих розчинених речовин у пористих середовищах. Вейвлетний розв'язок одержано в рамках мультирезольвентного аналізу.

INTRODUCTION

The transport theory of reacting solutes in a porous medium is considered under the hypotheses that the motion of solute transport is unidirectional, isothermal and devoid of instabilities; it takes place in a heterogeneous porous medium and the content, the density and the viscosity of the water in the medium are constant during the process. Since we are mainly interested in the chemical reaction equations we assume that the physical parameters defining the medium are unaffected by the transport, leaving the size of the pores, their distribution in the solid (and so on) unchanged. The chemical species defining the solid are at rest (while the chemical species defining the solute are mobile) in the medium, so that the mathematical model of the transport solution under chemical reactions (source-free) is the following algebraic-differential system $[6](\Theta, D, Q \text{ constants})$

$$\begin{cases} \Theta \frac{\partial c_i}{\partial t} = Lc_i, \\ L \stackrel{def}{=} D \frac{\partial^2}{\partial x^2} - Q \frac{\partial}{\partial x} \\ f_r \left(c_i, \frac{\partial c_i}{\partial t} \right) = 0, \quad (i = 1, \dots, \#P; r = 1, \dots) , \end{cases}$$
(1)

also called the convection-dispersion system. System (1) is a differential and algebraic system in the unknown #P functions $c_i(x,t)$, #P being the number of tenads [?] involved in the chemical reaction. The algebraic equations (1)₃ express the relationships

among concentrations of reaction participants, to be fulfilled irrespective of the contributions of the individual processes (both chemical and/or physical) influencing the concentrations c_i .

1. ONE DIMENSIONAL SOLUTE TRANSPORT EQUATION

In a one-dimensional domain $\Omega \subset \mathcal{R}$ let x be the coordinate, I a finite interval of time t, $(I \stackrel{def}{=} \{t: 0 < t < T, T < \infty\})$. We consider a sufficiently fast and reversible reaction of heterogeneous type [6],where the medium's original solution consists of two reacting cations M_1, M_3 in equilibrium with a cation exchanger ${}^1 \overline{M}_e$. The displacing solution contains a reacting cation M_1 and a non-reacting anion M_2 . The transport affecting reaction is represented by the chemical reactions (for binary cation exchange)

$$M_1 + \overline{M_3 M_e} \rightleftharpoons M_3 + \overline{M_1 M_e}.$$

There are four tenads in the system, three reacting M_1, M_3, M_e and one chemical non-reacting species M_2 . The basic equations (1) for the concentrations

¹ Dissolved in water and solid phase transport participants are given symbols M and \overline{M} , respectively.

 $c_1, \bar{c}_1, c_2, c_3, \bar{c}_3$ are [6, p. 1237]

$$\begin{cases} \Theta \frac{\partial c_1}{\partial t} + \rho \frac{\partial \bar{c}_1}{\partial t} = Lc_1, \\\\ \Theta \frac{\partial c_2}{\partial t} = Lc_2, \\\\ \Theta \frac{\partial c_3}{\partial t} + \rho \frac{\partial \bar{c}_3}{\partial t} = Lc_3, \\\\ \rho \frac{\partial \bar{c}_1}{\partial t} + \rho \frac{\partial \bar{c}_3}{\partial t} = 0 \\\\ K_{13} = \frac{\bar{c_1}c_3}{c_1\bar{c}_3} \end{cases}$$
(2)

with K_{13} a given constant. The *e* subscript, representing M_e , has been dropped from the \bar{c} functions, for them ρ is the anologous of Θ and represents the porous medium's bulk density (mass of the medium's solids/the medium's volume). In the heterogeneous reactions both the liquid and solid phases are involved in the tenads. From $(2)_4$ we get

$$\bar{c}_1 + \bar{c}_3 = c \tag{3}$$

since the exchange capacity does not vary with x. Then it follows:

$$\Theta \frac{\partial}{\partial t} \left(c_1 + c_3 \right) = L \left(c_1 + c_3 \right)$$

that, with the change of variable $u = c_1 + c_3$, becomes the linear equation

$$\Theta \frac{\partial u}{\partial t} = Lu,$$

whose solution can be expressed in terms of wavelets [2]. Deriving the condition $(2)_5$ with respect to t and taking into account equation (3) we obtain from $(2)_{1,4}$ the following nonlinear system [6]:

$$\left(\Theta + \frac{\rho p_i}{g} \right) \frac{\partial c_i}{\partial t} - \frac{\rho p_i}{g} \frac{\partial c_j}{\partial t} = Lc_i \qquad i, j = 1, 3$$
$$p_i \stackrel{def}{=} K_{13} \left(c - c_i \right), \quad g \stackrel{def}{=} K_{13} c_1 + c_3,$$

from which we obtain the non-linear equation

$$a\frac{\partial g}{\partial t} = gLg, \quad L \stackrel{def}{=} D\frac{\partial^2}{\partial x^2} - Q\frac{\partial}{\partial x}$$

with a suitable constant a. In particular, we restrict ourselves to the following one-dimensional initialboundary problem:

$$\frac{\partial g}{\partial t} = gLg, \quad L \stackrel{def}{=} D \frac{\partial^2}{\partial x^2} - Q \frac{\partial}{\partial x}, \quad (a = 1)$$

$$\begin{cases} u(x,0) = u_B(x), \quad 0 \le x \le 1, \quad t = 0, \\ u(0,t) = 0, \quad x = 0, \quad t > 0, \\ u(1,t) = 0, \quad x = 1, \quad t > 0 \end{cases}$$
(4)

and we assume as the initial function

$$u_B(x) = \begin{cases} 1, & x \in \Lambda_1, \\ -1, & x \in \Lambda_{-1}, \\ 0, & x \notin \Lambda_1 \cup \Lambda_{-1}, \end{cases}$$
(5)
$$\Lambda_1 \stackrel{def}{=} \{ x : 0 < x_0 \le x \le x_1 < 1 \}$$
$$\Lambda_{-1} \stackrel{def}{=} \{ x : 0 < x_1 \le x \le x_2 < 1 \}$$

which corresponds to the realistic case of an impulse function when $|x_2 - x_0| \rightarrow 0$.

2. MULTI-RESOLUTION ANALYSIS IN HAAR BASIS

The Haar scaling function $\Phi_k^n(x) \stackrel{def}{=} 2^{n/2} \Phi(2^n x - k)$ has a compact support on the dyadic interval

$$D_k^n \stackrel{\text{def}}{=} \left[\frac{k}{2^n}, \frac{k+1}{2^n} \right)$$

where its value is 1. The Haar family of wavelets

$$\Psi_{k}^{n}(x) \stackrel{def}{=} \left\{ 2^{\frac{n}{2}} \Psi(2^{n} x - k) \right\}_{k, n \in \mathbb{Z}}, \qquad (6)$$

is a complete orthonormal system for the $L_2(\mathcal{R})$ functions [5]

$$\Psi_{k}^{n}(x) = \begin{cases} 1, & x \in \left[\frac{k}{2^{n}}, \frac{k+1/2}{2^{n}}\right] \\ -1, & x \in \left[\frac{k+1/2}{2^{n}}, \frac{k+1}{2^{n}}\right] \\ 0, & \text{elsewhere }. \end{cases}$$
(7)

Let V_n , $n \in \mathbb{Z}$ be the subspace of $L_2(\mathcal{R})$ defined as the set of the piecewise constant functions f(x) of compact support on D_k^n (*n* fixed) , and W_n the orthogonal subspace such that the axioms of multiresolution (or multiscale) analysis [4, 5] are fulfilled,

$$\begin{cases}
L_2(\mathcal{R}) = \bigoplus_{n \in Z} W_n = V_q \oplus \bigoplus_{j \ge q} W_j, \quad q \in Z \\
V_{n+1} = V_n \oplus W_n,
\end{cases}$$
(8)

being \oplus the direct sum of orthogonal spaces. The set of functions $\{\Psi_k^n\}$ $(n \in \mathbb{Z})$ represents an orthonormal basis for $L_2(\mathcal{R})$.

Fixing the resolution value $N < \infty$, in (8), the $L_2(\mathcal{R})$ space is approximated by $L_2(\mathcal{R}) \cong \bigoplus_{\alpha} W_{\alpha}$,

that is,

$$f(x) \cong \pi^{N+1} f(x) \stackrel{def}{=} \alpha_0^0 + \sum_{n=0}^N \sum_{k=0}^n \beta_k^n \Psi_k^n(x), \quad (9)$$

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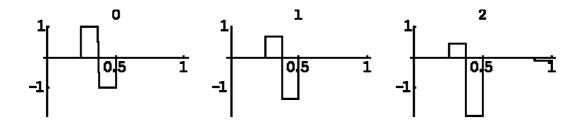


Рис. 1. Haar-wavelet representation of the wave solution

being π^n a projection operator into V_{n+1} , so that $\pi^n: L_2(\mathcal{R}) \to V_{n+1}$.

Choosing a number of dyadic nodes $x_k \stackrel{def}{=} \frac{k}{2^n}, (k = 0, ..., 2^n - 1)$ the dyadic discretization is the operator $\nabla^n : L_2(\mathcal{R}) \to L_2(\mathbb{Z}(2^{-n}))$, with $L_2(\mathbb{Z}(2^{-n})) \subset L_2(\mathcal{R})$ being the set of $L_2(\mathcal{R})$ functions sampled at x_k . The action of ∇^n on f(x) is such that $\nabla^n f(x) = \mathbf{f}^n$ with $\mathbf{f}^n = \{f_0, f_1, ..., f_{2^n-1}\}$ and $\{f_k \stackrel{def}{=} f(x) \mid_{x=x_k}, 0 \le k \le 2^n - 1\}$. The fast Haarwavelet transform \mathcal{H} of \mathbf{f}^N is the linear operator [1, 4]

$$\mathcal{H} : L_2 \left(Z(2^{-n}) \right) \to V_n \mid \mathbf{f}^N \mapsto \mathcal{H}\mathbf{f}^N = = \left\{ \alpha_0^0, \beta_k^n \right\}_{k=0,\dots,n}^{n=0,\dots,2^N-1},$$
(10)

where α_0^0 , β_k^n are the wavelet coefficients.

According to the above definitions, the projection operator $\pi^n : L_2(\mathcal{R}) \to V_{n+1}$ is factorized as $\pi^n = \mathcal{H}\nabla^n$.

A *p*-order Cardinal spline, is a $C^{p-1}([0,1))$ differentiable operator

$$\mathcal{S}^p : L_2(Z(2^{-N})) \to C^{p-2}([0,1)) : \mathbf{f}^N \mapsto s(x) \stackrel{def}{=} \mathcal{S}^p \mathbf{f}^N,$$

such that for the differential operator

$$L: L_2(Z(2^{-N})) \to L_2(Z(2^{-N}))$$

it is

$$L\mathcal{H} = \mathcal{H}L\mathcal{S}^p. \tag{11}$$

There follows that, given the set \mathbf{f}^N and computed the spline of a sufficiently large order, the splinederivative of $\mathcal{H}\mathbf{f}^N$ belongs to the same space of \mathbf{f}^N [1, 4].

According to the above, using splines and wavelets up to the resolution N, the approximate solution of the equation (4) is the vector $\mathbf{u}^N (\in V_{N+1})$, i.e. assuming the Euler formula for the time-derivative

$$\pi^N \frac{\partial u}{\partial t} = \frac{\mathbf{u}^{N+1} - \mathbf{u}^{N+1}}{\Delta t}$$

we have from (4)

$$\mathbf{u}^{N+1} = \mathbf{u}^N + \Delta t L \left(\mathcal{H} \mathbf{u}^N \right)$$

and, according to (11)

$$\mathbf{u}^{N+1} = \left(1 + \Delta t \mathcal{H} L \mathcal{S}^p\right) \mathbf{u}^N. \tag{12}$$

With the boundary condition (5), time step $\Delta t = 0.01$, and assuming in $(4)_1 \ Q = D = 1$ and in $(5)x_0 = \frac{1}{4}, \ x_1 = \frac{3}{8}, \ x_2 = \frac{1}{2}$, after 3 time steps (t = 0.05) we obtain the evolving function of Fig.

- (t = 0.05) we obtain the evolving function of Fig.
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