

*Regular article*

## The use of splitting methods in the numerical simulation of reacting flows \*

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Received: 20 July 2002 / Accepted: 23 March 2003

Published online: 23 January 2004 – © Springer-Verlag 2004

Communicated by: M.S. Espedal, A. Quarteroni, A. Sequeira

**Abstract.** The aim of this paper is to study discretizations of convection-diffusion-reaction equations using splitting methods. Estimates for the physical splitting errors and the numerical splitting errors are established. These estimates lead to the selection of optimal sequences and coupling of physical phenomena and adequate use of implicitness and explicitness. Numerical simulations of two chemical industry problems are included.

### 1 Introduction

Numerical simulation of reacting flows is required in all problems coming from the chemical industry, and also in the modelling of certain regions of hypersonic aerodynamics. The added complexity of this kind of problems comes from the fact that often a wide range of time scales is present. This leads to numerical difficulties related to the possible stiffness of reaction terms. Considering the Partial Differential Equations that describe reacting flows, we may split them additively into advective transport, diffusive transport and chemical transformations. We thus obtain submodels corresponding to different physical phenomena, that are easier to solve separately. Recent literature ([2, 4, 6]) has provided several ways of dealing with operator splitting as far as the sequence of these physical phenomena and also the possible coupling between some of them are concerned. Numerical simulations in [0,  $t$ ] pointed out that the coupling of reactions with diffusive transport in a sequence composed by a convection problem in a certain interval [0,  $t/2$ ], a diffusion-reaction problem in [0,  $t$ ] and a final convection problem in [ $t/2$ ,  $t$ ] perform notably better than an uncoupled splitting. In this paper we present a theoretical justification of this numerical evidence by studying the splitting errors and the stability properties of some numerical methods, based on different sequences and couplings of physical phenomena.

\* This work has been supported by Centro de Matemática da Universidade de Coimbra and Project POCTI/35039/MAT/2000

The splitting error has two main contributions: a physical splitting error that would exist even if submodels were solved exactly, and a numerical splitting error, related with the approximation of each submodel and the way these submodels are linked. In Sect. 2 we establish the order of the physical splitting error for different sequences and couplings of the phenomena. In Sect. 3 we show how the stability properties of numerical splitting errors based on the previous sequences and couplings, depend on the alternating use of implicit and explicit methods. Numerical simulations of two chemical industry problems will be presented in Sect. 4. Namely we exhibit results concerning a fixed bed catalytic reactor and a paper industry digester.

### 2 Splitting methods and splitting errors

Let us consider convection-diffusion-reaction equations of type

$$\frac{\partial c}{\partial t}(z, t) = F_1(c, z, t) + F_2(c, z, t) + F_3(c, z, t), \quad (1)$$

where  $c$  denotes a specie concentration and  $F_1(c, z, t)$ ,  $F_2(c, z, t)$ ,  $F_3(c, z, t)$  represent respectively the convection, the diffusion and the reaction terms.

Let us define in  $[0, T]$  the splitting grid  $\{t_s\}$  with  $t_s = s\Delta t$  and  $t_{s+1/2} = t_s + \Delta t/2$  where  $\Delta t$  denotes the splitting step size. We suppose that the concentration  $c$  at  $t = t_s$  is known (at least approximately). The computation of an approximation of  $c$  at time level  $t_{s+1}$  can be obtained using several splitting algorithms. In this paper we analyse from a theoretical viewpoint two functional splitting procedures, suggested in [6], to study an atmospheric pollution problem.

**Functional splitting I** Problem (1) is decomposed in five subproblems respectively (Convection, Diffusion, Reaction, Diffusion, Convection) as follows:

$$\begin{cases} \frac{\partial u}{\partial t}(z, t) = F_1(u, z, t), t \in [t_s, t_{s+1/2}], \\ u(t_s) = c(t_s), \end{cases} \quad (2)$$

$$\begin{cases} \frac{\partial v}{\partial t}(z, t) = F_2(v, z, t), t \in [t_s, t_{s+1/2}], \\ v(t_s) = u(t_{s+1/2}), \end{cases} \quad (3)$$

$$\begin{cases} \frac{\partial w}{\partial t}(z, t) = F_3(w, z, t), t \in [t_s, t_{s+1}], \\ w(t_s) = v(t_{s+1/2}), \end{cases} \quad (4)$$

$$\begin{cases} \frac{\partial v^*}{\partial t}(z, t) = F_2(v^*, z, t), t \in [t_{s+1/2}, t_{s+1}], \\ v^*(t_{s+1/2}) = w(t_{s+1}), \end{cases} \quad (5)$$

$$\begin{cases} \frac{\partial u^*}{\partial t}(z, t) = F_1(u^*, z, t), t \in [t_{s+1/2}, t_{s+1}], \\ u^*(t_{s+1/2}) = v^*(t_{s+1}). \end{cases} \quad (6)$$

The splitting solution I at  $t_{s+1}$  is  $u^*(t_{s+1})$  – the exact solution of (2)–(6) – which represents an approximation of  $c(t_{s+1})$ .

**Functional splitting II** Problem (1) is decomposed in three subproblems (Convection, Diffusion+Reaction, Convection) as described below

$$\begin{cases} \frac{\partial u}{\partial t}(z, t) = F_1(u, z, t), t \in [t_s, t_{s+1/2}], \\ u(t_s) = c(t_s), \end{cases} \quad (7)$$

$$\begin{cases} \frac{\partial v}{\partial t}(z, t) = F_2(v, z, t) + F_3(v, z, t), t \in [t_s, t_{s+1}], \\ v(t_s) = u(t_{s+1/2}), \end{cases} \quad (8)$$

$$\begin{cases} \frac{\partial u^*}{\partial t}(z, t) = F_1(u^*, z, t), t \in [t_{s+1/2}, t_{s+1}], \\ u^*(t_{s+1/2}) = v(t_{s+1}). \end{cases} \quad (9)$$

The splitting solution II at  $t_{s+1}$  is  $u^*(t_{s+1})$  – the exact solution of (7)–(9) – which stands for an approximation of  $c(t_{s+1})$ .

Let us represent by  $c_I$  and  $c_{II}$  respectively the splitting solutions I and II. In the case operators  $F_1, F_2$  are linear and  $F_3$  depends only on  $z$  we can estimate the splitting errors  $\|c - c_I\|_\infty$  and  $\|c - c_{II}\|_\infty$ .

**Proposition 1.** Let  $F_1(c, z, t) = -\alpha \frac{\partial c}{\partial z}(z, t)$ ,  $F_2(c, z, t) = \beta \frac{\partial^2 c}{\partial z^2}(z, t)$ ,  $\beta > 0$ , and  $F_3(c, z, t) = F_3(z)$ , with  $|F_3'| \leq q$ ,  $c(z, 0) = c_0(z)$  and  $\lim_{z \rightarrow \pm\infty} c(z, t) = 0$ , for all  $t$ . Then  $\|c - c_I\|_\infty = O(\Delta t^{3/2})$  and  $\|c - c_{II}\|_\infty = O(\Delta t^2)$

*Proof.* Using Fourier transforms we have for the exact solution of (1)

$$\begin{aligned} c(z, t) &= \frac{1}{\sqrt{\pi}} \int_0^t \int_{\mathbb{R}} F_3\left(z - \alpha\xi + 2\sqrt{\beta\xi}y\right) e^{-y^2} dy d\xi \\ &\quad + \frac{1}{\sqrt{\pi}} \int_{\mathbb{R}} c_0\left(z - \alpha t + 2\sqrt{\beta t}\xi\right) e^{-\xi^2} d\xi. \end{aligned} \quad (10)$$

It is a tedious but straightforward task to establish that the splitting solutions  $c_I$  and  $c_{II}$  are given respectively by

$$\begin{aligned} c_I(z, t) &= \frac{1}{\sqrt{\pi}} \int_{\mathbb{R}} t F_3\left(z - \frac{\alpha t}{2} + 2\sqrt{\frac{\beta t}{2}}y\right) e^{-y^2} dy \\ &\quad + \frac{1}{\sqrt{\pi}} \int_{\mathbb{R}} c_0\left(z - \alpha t + 2\sqrt{\beta t}\xi\right) e^{-\xi^2} d\xi, \end{aligned} \quad (11)$$

$$\begin{aligned} c_{II}(z, t) &= \frac{1}{\sqrt{\pi}} \int_0^t \int_{\mathbb{R}} F_3\left(z - \frac{\alpha t}{2} + 2\sqrt{\beta\xi}y\right) e^{-y^2} dy d\xi \\ &\quad + \frac{1}{\sqrt{\pi}} \int_{\mathbb{R}} c_0\left(z - \alpha t + 2\sqrt{\beta t}\xi\right) e^{-\xi^2} d\xi. \end{aligned} \quad (12)$$

From (10) and (11) we have

$$\begin{aligned} &|c(z, t) - c_I(z, t)| \\ &\leq \frac{q}{\sqrt{\pi}} \left| \int_0^t \int_{\mathbb{R}} \left(\alpha\xi - \frac{\alpha k}{2} + 2\sqrt{\beta\xi}y - \sqrt{\frac{\beta t}{2}}y\right) e^{-y^2} dy d\xi \right| \end{aligned}$$

and considering  $\int_{\mathbb{R}} e^{-y^2} dy = \sqrt{\pi}$ ,  $\int_{\mathbb{R}} e^{-y^2} |y| dy = 1$  we may easily conclude that  $\|c - c_I\|_\infty = O(\Delta t \sqrt{\Delta t})$ .

From (10) and (12) we conclude

$$|c(z, t) - c_{II}(z, t)| \leq \frac{q}{\sqrt{\pi}} \left| \int_0^t \int_{\mathbb{R}} \left(\alpha\xi - \frac{\alpha t}{2}y\right) e^{-y^2} dy d\xi \right|$$

and consequently  $\|c - c_{II}\|_\infty = O(\Delta t^2)$ .  $\square$

We note that the integral terms in (11) and (12), concerning the initial conditions  $c_0(z)$ , introduce no error in  $c_I$  and  $c_{II}$ . Also if  $F_3$  is a linear function of  $z$  then  $c_I = c_{II}$ .

In [6], the authors, while studying an atmospheric problem, point out that splitting II lead to more accurate numerical results. The estimates of Proposition 1 can justify from a theoretical point of view this numerical evidence.

The following example illustrates the estimates of Proposition 1. Let us consider the initial-boundary value problem

$$\begin{aligned} \frac{\partial c}{\partial t}(z, t) &= \gamma_1 \frac{\partial c}{\partial z}(z, t) + \gamma_2 \frac{\partial^2 c}{\partial z^2}(z, t) \\ &\quad + \gamma_3(c(z, t) - c_w(t)) + \gamma_4 r(c) \end{aligned} \quad (13)$$

with initial-boundary conditions

$$\begin{cases} c(z, 0) = f(z), z \in (0, 1), \\ c(0, 0) = c_F(0), \\ \gamma_2 \frac{\partial c}{\partial z}(0, t) = [c(0, t) - c_F(t)]\gamma_1, t > 0, \\ \frac{\partial c}{\partial z}(1, t) = 0, t \geq 0. \end{cases} \quad (14)$$

In (13),  $\gamma_i$ ,  $i = 1, \dots, 5$ , are real constants,  $r(c)$  and  $c_w(t)$  represent known functions. In (14),  $f$  and  $c_F$  are also assumed to be known.

We took in (14),  $r(c) = c^2$ ,  $\gamma_1 = -1$ ,  $\gamma_2 = 10^{-3}$ ,  $\gamma_3 = -0.5$ ,  $\gamma_4 = 0.5$ ,  $c_w(t) = 0.8$  and  $f(z) = 1.11067 + 0.359664z + 0.325977z^2$ .

In Fig. 1 are plotted the exact and numerical splitting solutions for  $t = 0.5$  and  $t = 1$ . We note that splitting II solution is much more accurate than splitting I solution.

In Proposition 1 we consider a convection-diffusion-reaction problem in an unbounded domain. In the case of convection-reaction equations estimates have been established under a boundary condition along  $z = 0$ .

**Proposition 2.** Let  $F_2 = 0$  and  $F_3(c, z, t) = F_3(c)$ , with  $|F_3'| \leq q$ .

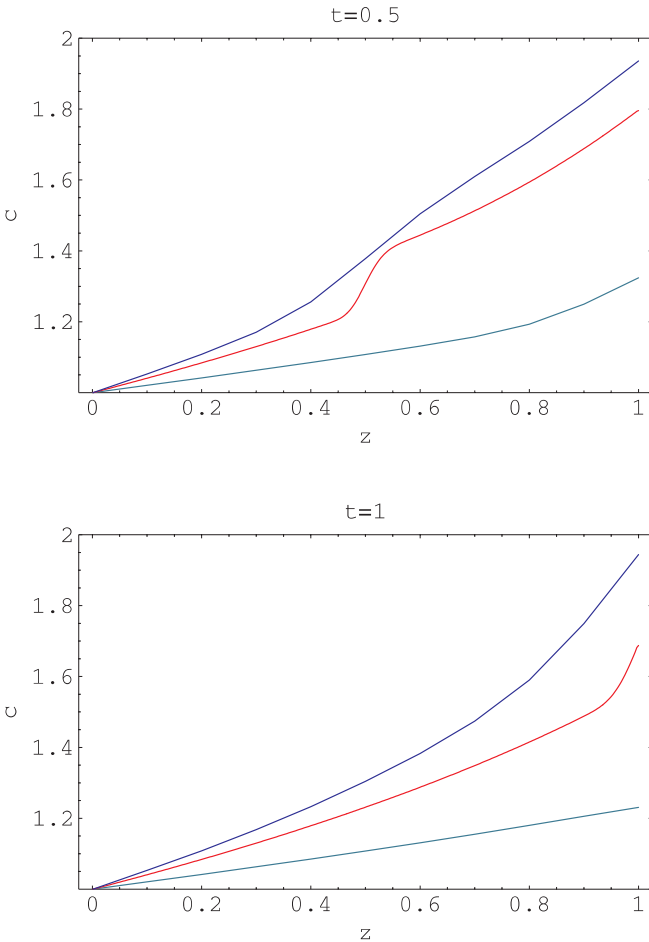
(i) If  $c(z, 0) = c_0(z)$  for  $z \in (0, +\infty)$  and  $c(0, t) = g(t)$  with  $c_0$  and  $g$  enough smooth functions, then:

(a) if  $F_1(c) = \alpha(z, t) \frac{\partial c}{\partial z}$  we have  $c_{II} = c$  for  $z > \alpha t$  and

$$\|c - c_{II}\|_\infty = O(\Delta t) \text{ for } z \leq \alpha t;$$

(b) if  $F_1(c) = \frac{\partial}{\partial z} f(c)$  we have  $\|c - c_{II}\|_\infty = O(\Delta t)$ .

(ii) If  $c(z, 0) = c_0(z)$ ,  $z \in \mathbb{R}$ , then the splitting solution is the exact solution.



**Fig. 1.** Numerical solutions (split I: green; split II: blue) and exact solution (red) for (13)–(14)

*Proof.* We just present the proof of 1(a). The exact solution satisfies

$$\begin{cases} \int_{g(\frac{\alpha t - z}{\alpha})}^{c(z,t)} \frac{dc}{F_3(c)} = \frac{z}{\alpha}, & z \leq \alpha t, \\ \int_{c_0(z - \alpha t)}^{c(z,t)} \frac{dc}{F_3(c)} = t, & z > \alpha t. \end{cases} \quad (15)$$

The solution  $c_{II}$  of (7)–(9) verifies

$$\begin{cases} c(z, t) = g\left(\frac{\alpha t - z}{\alpha}\right), & 0 \leq z \leq \frac{\alpha t}{2}, \\ \int_{g(\frac{\alpha t - z}{\alpha})}^{c(z,t)} \frac{dc}{F_3(c)} = \frac{z}{\alpha}, & \frac{\alpha t}{2} < z \leq \alpha t, \\ \int_{c_0(z - \alpha t)}^{c(z,t)} \frac{dc}{F_3(c)} = t, & z > \alpha t. \end{cases} \quad (16)$$

From (15) and (16) the result follows.  $\square$

We observe that if no boundary conditions are considered in the convection-reaction problem, as in (ii) of Proposition 2, the exact solution is obtained. When a boundary condition is considered, as in (i) of Proposition 2, an error of order  $\Delta t$  propagates in the domain defined by  $z \leq \alpha t$ .

### 3 Numerical splitting methods

#### 3.1 General description of the family of methods

We consider, in what follows, numerical methods obtained by discretizing (7)–(9). In the interval  $[0, T]$  we consider the splitting grid  $\{t_s\}$  and in the space domain we define the grid  $\{z_j\}$  with  $z_j - z_{j-1} = h$ . Let  $U_h^s$ ,  $V_h^s$  and  $U_h^{s*}$  be numerical approximations at  $t_s$  of  $u(t_s)$ ,  $v(t_s)$  and  $u^*(t_s)$ , respectively defined by

$$\begin{cases} U_h^{s+1/2} = F_{1,h}(U_h^s), \\ U_h^s = c_h^s, \end{cases} \quad (17)$$

$$\begin{cases} F_{3,h}^*(V_h^{s+1}) = F_{2,h}(V_h^s) + F_{3,h}(V_h^s), \\ V_h^s = U_h^{s+1/2}, \end{cases} \quad (18)$$

$$\begin{cases} U_h^{s+1*} = F_{1,h}(U_h^{s+1/2*}), \\ U_h^{s+1/2*} = V_h^{s+1}, \end{cases} \quad (19)$$

where  $U_h^{s+1*}$  represents a numerical approximation of  $c_{II}(t_{s+1})$  and consequently of  $c(t_{s+1})$ . In this algorithm  $F_{1,h}$  is a linear operator resulting from the discretization of the convection equation (7). Equation (18) represents a discretization of (8), where we assume that  $F_{2,h}$  is linear and  $F_{3,h}$ ,  $F_{3,h}^*$  are nonlinear operators. These operators take into account the boundary conditions prescribed for (7)–(9).

### 3.2 Stability

We establish in what follows the stability of the methods described by algorithm (17)–(19). Let  $c_h^{s+1}$  and  $\tilde{c}_h^{s+1}$  be two solutions computed using this splitting algorithm. The difference  $c_h^{s+1} - \tilde{c}_h^{s+1}$  satisfies the following equation:

$$\begin{aligned} & c_h^{s+1} - \tilde{c}_h^{s+1} \\ &= F_{1,h} (JF_{3,h}^*)^{-1} (F_{2,h} + JF_{3,h}) F_{1,h} (c_h^s - \tilde{c}_h^s) \end{aligned} \quad (20)$$

where  $JF_{3,h}^*$  is the Jacobian matrix of  $F_{3,h}^*$  computed in  $\sigma V_h^{n+1} + (1 - \sigma) \tilde{V}_h^{n+1}$  for some  $\sigma \in [0, 1]$ . The Jacobian matrix  $JF_{3,h}$  is defined analogously. The following proposition can be easily established:

**Proposition 3.** *If there exists  $S_C \in (0, 1)$  such that*

$$\left\| F_{1,h} (JF_{3,h}^*)^{-1} (F_{2,h} + JF_{3,h}) F_{1,h} \right\| \leq S_C \quad (21)$$

*then the splitting method (17)–(19) is stable.  $\square$*

Let us suppose now that in each step of (17)–(19) we introduce a perturbation  $\varepsilon_i$ , that is

$$\begin{cases} \tilde{U}_h^{s+1/2} = F_{1,h} (\tilde{U}_h^s) + \varepsilon_1, \\ U_h^s = c_h^s, \end{cases} \quad (22)$$

$$\begin{cases} F_{3,h}^* (\tilde{V}_h^{s+1}) = F_{2,h} (\tilde{V}_h^s) + F_{3,h} (\tilde{V}_h^s) + \varepsilon_2, \\ \tilde{V}_h^s = \tilde{U}_h^{s+1/2}, \end{cases} \quad (23)$$

$$\begin{cases} \tilde{U}_h^{s+1*} = F_{1,h} (\tilde{U}_h^{s+1/2*}) + \varepsilon_3, \\ \tilde{U}_h^{s+1/2*} = \tilde{U}_h^{s+1}, \end{cases} \quad (24)$$

Then we can study how these perturbations propagate from time level  $t_s$  to time level  $t_{s+1}$ . Considering that

$$\begin{aligned} c_h^{s+1} - \tilde{c}_h^{s+1} &= F_{1,h} (JF_{3,h}^*)^{-1} ((F_{2,h} + JF_{3,h}) \\ &\quad (F_{1,h} (c_h^s - \tilde{c}_h^s) + \varepsilon_1) + \varepsilon_2) + \varepsilon_3 \end{aligned}$$

we obtain the internal stability factor

$$\varrho_s = F_{1,h} (JF_{3,h}^*)^{-1} ((F_{2,h} + JF_{3,h}) \varepsilon_1 + \varepsilon_2) + \varepsilon_3$$

which satisfies

$$\begin{aligned} \|\varrho_s\| &\leq \left\| F_{1,h} (JF_{3,h}^*)^{-1} (F_{2,h} + JF_{3,h}) \right\| \|\varepsilon_1\| \\ &\quad + \left\| F_{1,h} (JF_{3,h}^*)^{-1} \right\| \|\varepsilon_2\| + \|\varepsilon_3\|. \end{aligned}$$

Attending to the last inequality we have the following proposition.

**Proposition 4.** *If there exists  $I_C \in (0, 1)$  such that*

$$\left\| F_{1,h} (JF_{3,h}^*)^{-1} (F_{2,h} + JF_{3,h}) \right\| \leq I_C, \quad (25)$$

$$\left\| F_{1,h} (JF_{3,h}^*)^{-1} \right\| \leq I_C \quad (26)$$

*then the splitting method is internally stable.  $\square$*

### 3.3 Discretization errors

We recall that  $c(t_s)$  represents the solution of the convection-diffusion-reaction equation at  $t = t_s$ ,  $c_h^s$  a numerical approximation computed using splitting method (17)–(19) and  $c_{II}(t_s)$  the solution computed using the functional splitting (7)–(9). We have

$$\|c(t_s) - c_h^s\|_\infty \leq \|c(t_s) - c_{II}(t_s)\|_\infty + \|c_{II}(t_s) - c_h^s\|_\infty.$$

The first norm in the right hand side has been estimated in Proposition 1. We compute in what follows

$$\|c_{II}(t_s) - c_h^s\|_\infty.$$

Let  $T_{i,h}^{s+1}$ ,  $i = 1, 2, 3$  be the truncation errors at  $t = t_{s+1}$  associated with the discretizations (17), (18) and (19) respectively. By  $e_{i,h}^{s+1}$ ,  $i = 1, 2, 3$ , we denote the global errors defined by  $e_{1,h}^{s+1} = u(t_{s+1}) - U_h^{s+1}$ ,  $e_{2,h}^{s+1} = v(t_{s+1}) - V_h^{s+1}$  and  $e_{3,h}^{s+1} = u^*(t_{s+1}) - U_h^{s+1*}$ . These errors satisfy (22), (23) and (24) with  $\varepsilon_i = T_{i,h}^s$ . Then

$$\begin{aligned} & c_{II}(t_{s+1}) - c_h^{s+1} \\ &= F_{1,h} (JF_{3,h}^*)^{-1} (F_{2,h} + JF_{3,h}) F_{1,h} (c_h^s - \tilde{c}_h^s) \\ &\quad + F_{1,h} (JF_{3,h}^*)^{-1} ((F_{2,h} + JF_{3,h}) T_{1,h}^s + T_{2,h}^s) + T_{3,h}^s. \end{aligned}$$

Therefore, we obtain for the truncation error of the splitting method,  $T_h^s$ , the following equation

$$T_h^s = F_{1,h} (JF_{3,h}^*)^{-1} ((F_{2,h} + JF_{3,h}) T_{1,h}^s + T_{2,h}^s) + T_{3,h}^s.$$

From the previous results and stability Propositions 3 and 4 we may easily establish the next convergence estimate for the discretization error.

**Proposition 5.** *If (21), (25) and (26) are satisfied then*

$$\begin{aligned} & \|c_{II}(t_s + 1) - c_h^{s+1}\| \\ & \leq \frac{1 - S_C^{s+1}}{1 - S_C} (I_C + 1) \max_{i=1,2,3} \max_{\ell=1,\dots,s+1} \|T_{i,h}^\ell\|_\infty. \end{aligned} \quad \square$$

### 3.4 Special families of methods

Let us consider equation (1) with  $z \in [0, 1]$ ,

$$F_1(c) = -\alpha \frac{\partial c}{\partial z}, \quad F_2(c) = \beta \frac{\partial^2 c}{\partial z^2}, \quad F_3(c) = f(c).$$

We discretize (7)–(9) using backward and second order centered differences respectively for first and second order space derivatives. The time derivative is discretized using the  $\theta$ -method which is defined by

$$y^{s+1/2} = y^s + \frac{\Delta t}{2} ((1 - \theta)G(y^{s+1/2}) + \theta G(y^s)),$$

when  $y' = G(y)$  is to be solved.

In this case we have,

$$[F_{1,h}(U_h^s)]_1 = \frac{1 - a\hat{\theta}}{1 + a} (U_h^s)_1,$$

and, for  $i = 2, \dots, N$ ,

$$[F_{1,h}(U_h^s)]_i = \frac{1+\hat{\theta}}{1+a} \sum_{j=1}^{i-1} \left( \frac{a}{1+a} \right)^{i-j} (U_h^s)_j + \frac{1-a\hat{\theta}}{1+a} (U_h^s)_i.$$

For  $F_{2,h}$  we have

$$[F_{2,h}(V_h^s)]_1 = \left( 1 - 2\beta \frac{\Delta t}{h^2} \right) (V_h^s)_1 + \beta \frac{\Delta t}{h^2} (V_h^s)_2;$$

for  $i = 2, \dots, N-1$ ,

$$[F_{2,h}(V_h^s)]_i = \beta \frac{\Delta t}{h^2} (V_h^s)_{i-1} + \left( 1 - 2\beta \frac{\Delta t}{h^2} \right) (V_h^s)_i + \beta \frac{\Delta t}{h^2} (V_h^s)_{i+1};$$

and

$$[F_{2,h}(V_h^s)]_N = \beta \frac{\Delta t}{h^2} (V_h^s)_{N-1} + \left( 1 - 2\beta \frac{\Delta t}{h^2} \right) (V_h^s)_N.$$

Finally, for  $\sigma \in [0, 1]$ ,  $F_{3,h}$  is defined, for  $i = 1, \dots, N$ , by

$$[F_{3,h}(V_h^s)]_i = \Delta t \sigma f((V_h^s)_i),$$

and  $F_{3,h}^*$  is defined, for  $i = 1, \dots, N$ , by

$$[F_{3,h}^*(V_h^{s+1})]_i = (1 - \Delta t(1 - \sigma)) f((V_h^{s+1})_i).$$

Let us denote by  $M[\theta, \sigma]$  the splitting method just described.

It is a simple but tedious task to establish the expressions of  $I_C$  and  $S_C$  for  $M[\theta, \sigma]$ . Let  $\Delta t_0$  be an upper bound for the time step size. Assuming that  $|F_3| \leq q$  we have:

- (i) For  $\theta = 0$  then, for  $\gamma \in (0, 1)$  and for  $\Delta t$  small enough, we have

$$S_C := \gamma^2 \frac{1 + \Delta t_0 \sigma q}{1 - (1 - \sigma) \Delta t_0 q},$$

$$I_S := \gamma \frac{1 + \Delta t_0 \sigma q}{1 - (1 - \sigma) \Delta t_0 q};$$

- (ii) For  $\theta \in (0, 1)$ , there exists  $K(\theta) < 1$  such that, for  $\Delta t$  small enough,

$$S_C := K(\theta)^2 \frac{1 + \Delta t_0 \sigma q}{1 - (1 - \sigma) \Delta t_0 q},$$

$$I_S := K(\theta) \frac{1 + \Delta t_0 \sigma q}{1 - (1 - \sigma) \Delta t_0 q};$$

- (iii) If  $\theta = 1$  then  $I_C$  and  $S_C$  are greater or equal to one.

Attending to the values of  $S_C$  and  $I_C$  we conclude that  $M[0, 1]$  and  $M[0, 0]$  are stable and internally stable. Nevertheless the stability constant of  $M[0, 1]$  is bounded by  $\gamma^2(1 + \Delta t_0 q)$  while  $M[0, 0]$  has a stability constant bounded by  $\gamma/(1 - \Delta t_0 q)$ .

As far as the spatial truncation error is concerned we have, for  $j = 1, 3$ ,  $w = u, u^*$ ,

$$T_{j,h}^{s+1} = \frac{\partial^2 w}{\partial t^2} \frac{\Delta t}{2} \left( \frac{1}{2} - (1 - \theta) \right) + \alpha \frac{h}{2} \frac{\partial^2 w}{\partial x^2} + O(h^2, \Delta t^2),$$

and

$$T_{2,h}^{s+1} = \frac{\Delta t}{2} \left( \frac{\partial^2 w}{\partial t^2} - (1 - \sigma) f'(v(x_i, t_{s+1})) \frac{\partial w}{\partial t} \right) + O(h^2, \Delta t^2),$$

where the partial derivatives are evaluated at some points in  $(x_{i-1}, x_i) \times (t_s, t_{s+1})$ .

From Proposition 5 if  $M[\theta, \sigma]$  is stable and internally stable – with stability constants  $S_C, I_S$  less than one – then the discretization error  $c_{II}(t_s) - c_h^s$  satisfies

$$\|c_{II}(t_{s+1}) - c_h^{s+1}\|_\infty \leq \frac{1 - S_C^{s+1}}{1 - S_C} (I_S + 1) O(h, \Delta t).$$

## 4 Numerical examples

### 4.1 A fixed bed catalytic reactor

This section is concerned with the nonlinear initial boundary value problem (13)–(14) which assumes importance in chemical engineering, for instance in the modeling of fixed bed catalytic reactors ([5]). In this model it is assumed that for a given axial point, the temperature values at a characteristic position of the tube radius are representative of the whole cross section. In (13),  $\gamma_i, i = 1, \dots, 4$ , are real constants, representing the intensity of some mechanisms occurring in the system. The wall temperature  $c_w(t)$  reflects the distributed nature of the interaction between the process and its environment and the term  $r(c)$  is a nonlinear known function standing for a chemical reaction rate. In the initial-boundary condition (14),  $f$  and  $c_F$  are assumed to be known functions.

Computations have been carried with parameters:

$$\begin{cases} \gamma_1 = -1, \gamma_2 = 10^{-3}, \gamma_3 = -0.5, \gamma_4 = 0.5, \\ c_F(0) = 1.1, c_F(t) = 1, t > 0, \\ r(c) = c^2, \\ f(z) = 1.11067 + 0.359664z + 0.325977z^2, z \in (0, 1), \\ c_w(t) = 0.8, t \geq 0. \end{cases}$$

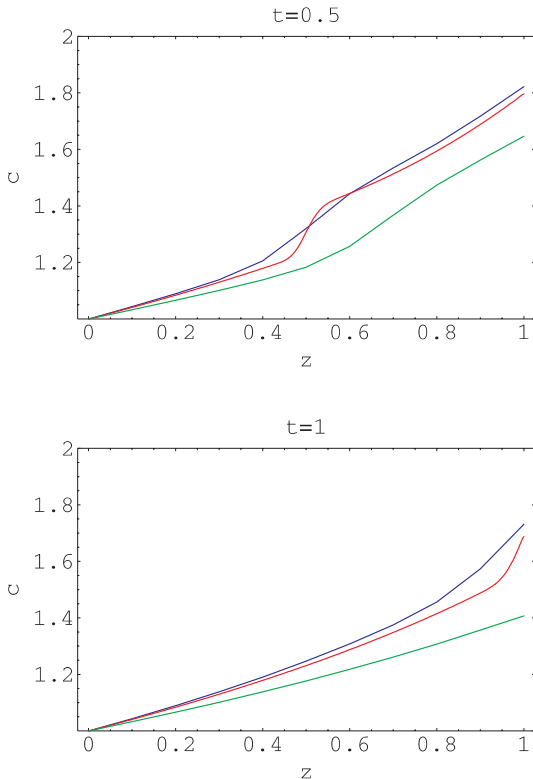
In Fig. 2 we plot the numerical solution computed with  $M[1, 1]$  and  $M[0, 1]$  with  $h = 10^{-2}$  and  $\Delta t = \frac{h}{2}$ .

The behaviour of the numerical approximation obtained using  $M[0, 0]$  is analogous to the one presented by the numerical approximation obtained using  $M[0, 1]$ . However the computational cost of  $M[0, 0]$  is greater than the computational cost of  $M[0, 1]$ .

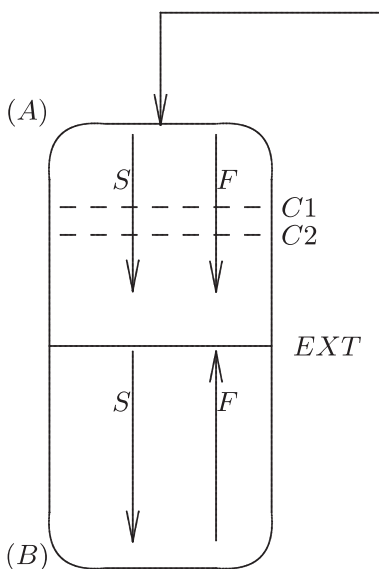
### 4.2 A moving bed reactor used in the paper industry

In this section we consider the system of hyperbolic partial differential equations that describes the behaviour of a moving bed reactor – the digester – used in the paper and pulp industry ([1]).





**Fig. 2.** Numerical approximations ( $M[1,1]$ : green;  $M[0,1]$ : blue) and exact solution (red) for (13)–(14)

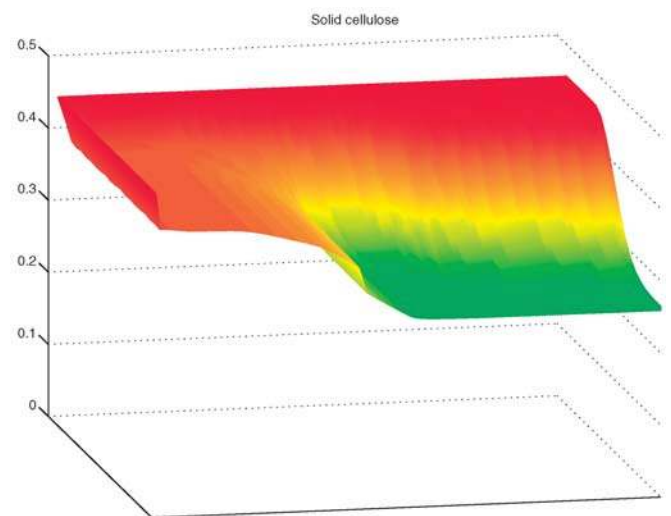


**Fig. 3.** In the first part of the digester the phases flow concurrently while in the second part the flows are countercurrently

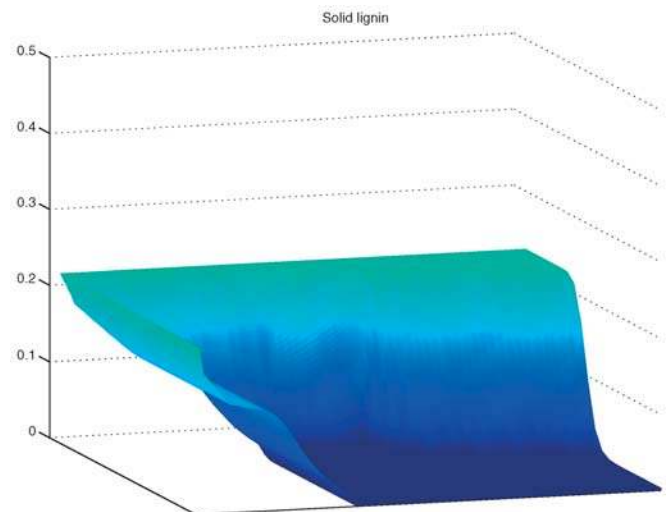
The digester is a heterogeneous reactor with an almost cylindrical shape, where wood chips – solid phase – react with an aqueous solution – free liquid phase – of sodium hydroxide and sodium sulfide, to remove the lignin from the cellulose fibers (Fig. 3). As the wood chips are porous a third phase – entrapped liquid phase – is also considered.

From the physical point of view we can describe the behaviour of a digester as follows: (i) In its upper part the

solid and the liquid phase flow downwards concurrently and at  $z = EXT$  the liquid phase is completely extracted; (ii) In its lower part a free liquid flows upwards countercurrently with the solid phase; (iii) At several levels of the digester's height – the circulations  $C1$  and  $C2$  – the free liquid is extracted, enriched and heated before being reinjected. From an industrial point of view it is important to know the temperature and the concentration of several chemical species – lignin, cellulose, hemicellulose – in the solid phase, free liquid phase and entrapped liquid phase. If  $y_i$  denotes the concentration of some chemical product then  $y_i^F$ ,  $y_i^E$  and  $y_i^S$  denote its concentration in the liquid, entrapped and solid phases respectively. For  $\ell = F, E, S$ ,  $y_i^\ell$ ,  $i = 1, 2, 3$ , represents the concentration of cellulose, hemicellulose and lignin respectively. For  $\ell = F, E$ ,  $i = 4, 5$ ,  $y_i^\ell$  represents the concentration respectively of sodium hydroxide and sodium sulfide. Finally  $y_6^F, y_6^E$  stand for the temperature of the free and the entrapped liquid. The behaviour of PDE's



**Fig. 4.** The evolution in time of the concentration of solid cellulose



**Fig. 5.** The evolution in time of the concentration of solid lignin

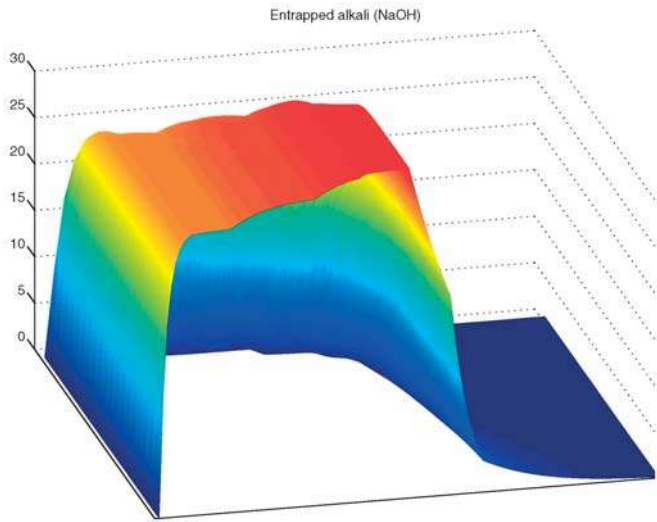


Fig. 6. The evolution in time of the concentration of entrapped alkali

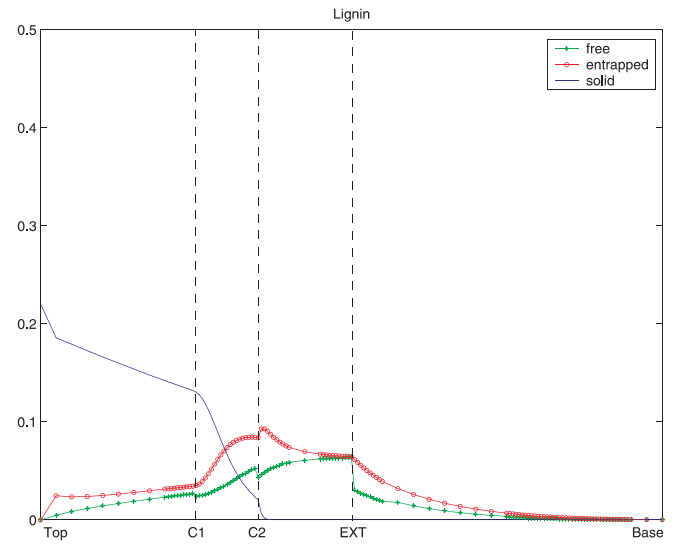


Fig. 9. Numerical stationary lignin solution for (27)

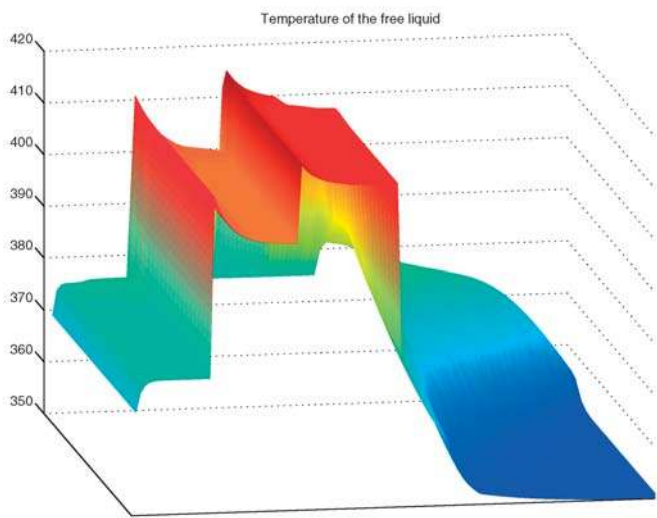


Fig. 7. The evolution in time of the temperature free liquid

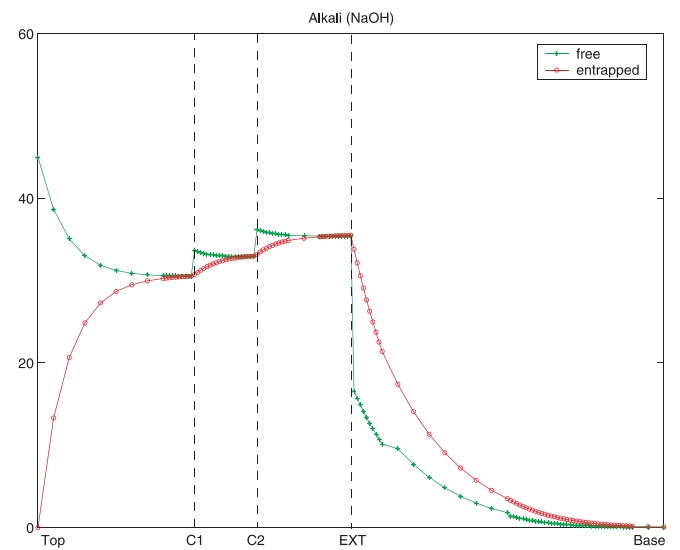


Fig. 10. Numerical stationary alkali solution for (27)

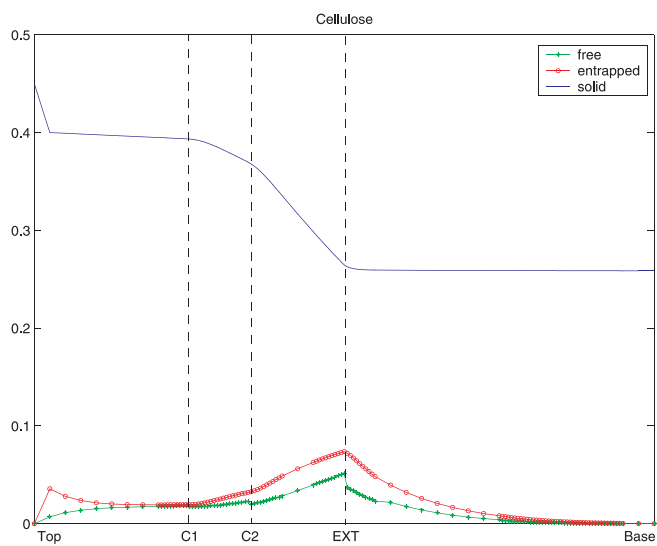


Fig. 8. Numerical stationary cellulose solution for (27)

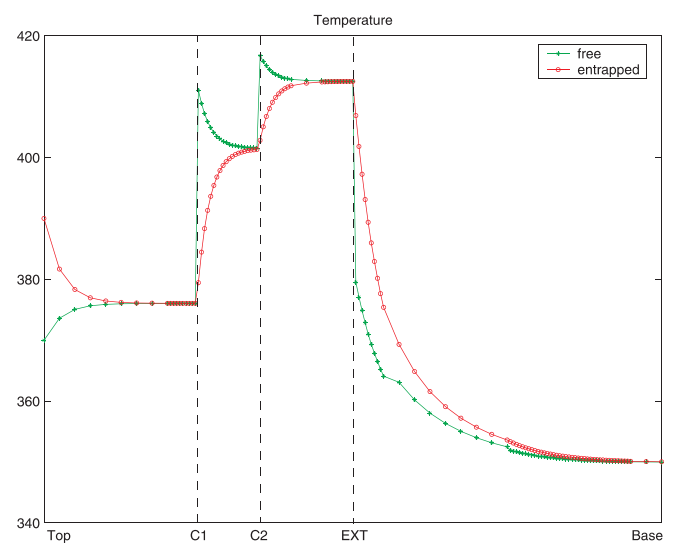


Fig. 11. Numerical stationary temperature solution for (27)

$$\left\{ \begin{array}{l} \xi_1 \frac{\partial y_i^F}{\partial t} = -u_F \frac{\partial y_i^F}{\partial z} + \varrho_1 (y_i^E - y_i^F), \quad i = 1, \dots, 5, \\ \xi_1 \frac{\partial y_6^F}{\partial t} = -u_F \frac{\partial y_6^F}{\partial z} + \varrho_2 (y_6^E - y_6^F), \\ \xi_2 \frac{\partial y_i^E}{\partial t} = -u_S \frac{\partial y_i^E}{\partial z} - R_i \\ \quad - \varrho_3 (y_i^E - y_i^F), \quad i = 1, \dots, 5, \\ \xi_2 \frac{\partial y_6^E}{\partial t} = -u_S \frac{\partial y_6^E}{\partial z} - \varrho_4 (y_6^E - y_6^F), \\ \xi_3 \frac{\partial y_i^S}{\partial t} = -u_S \frac{\partial y_i^S}{\partial z} + R_i, \quad i = 1, \dots, 3, \end{array} \right. \quad (27)$$

where  $\xi_i$ ,  $i = 1, 2, 3$ ,  $\varrho_i$ ,  $i = 1, \dots, 4$ , represent physical constants.

In system (27), the velocity  $u_F$  is positive for  $i = 1, 2, 3$  – the liquid phase flows downwards – and is negative for  $i = 4, 5$ , and  $z \geq EXT$  – because the liquid phase flows upwards countercurrently. The reaction terms  $R_i$  are represented by exponential functions depending on  $y_i^E$  and are experimentally established in [1]. The boundary and initial conditions as well as the values of the parameters used in the model were obtained experimentally and are established in [1].

The evolution in time of the concentration of solid cellulose, solid lignin and entrapped alkali are plotted respectively in Fig. 4, Fig. 5 and Fig. 6. In Fig. 7 the time evolution of the temperature of the free liquid is plotted. The stationary concentrations of cellulose, lignin, alkali and the temperature are plotted respectively in Fig. 8, Fig. 9, Fig. 10 and Fig. 11.

The computations have been made using M[0, 1] with constant step-size and local refinement near the circulation points. The results obtained for the evolution problem present physical evidence. The stationary results agree with the numerical results in [1].

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