open science

# The mathematical analysis of a syntrophic relationship between two microbial species in a chemostat 

Tewfik Sari, Miled El Hajji, Jérôme Harmand

## - To cite this version:

Tewfik Sari, Miled El Hajji, Jérôme Harmand. The mathematical analysis of a syntrophic relationship between two microbial species in a chemostat. Mathematical Biosciences and Engineering, 2012, 9 (3), pp.627-645. 10.3934/mbe.2012.9.627 . hal-00487189v2

## HAL Id: hal-00487189 https://hal.science/hal-00487189v2

Submitted on 29 Jul 2011 (v2), last revised 7 Mar 2024 (v3)

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

# ANALYSIS OF A MATHEMATICAL MODEL OF SYNTROPHIC BACTERIA IN A CHEMOSTAT 

TEWFIK SARI, MILED EL-HAJJI, JÉRÔME HARMAND


#### Abstract

A mathematical model involving a syntrophic relationship between two populations of bacteria in a continuous culture is proposed. A detailed qualitative analysis is carried out. The local and global stability analysis of the equilibria are performed. We demonstrate, under general assumptions of monotonicity, relevant from an applied point of view, the asymptotic stability of the positive equilibrium point which corresponds to the coexistence of the two bacteria. A syntrophic relationship in the anaerobic digestion process is proposed as a real candidate for this model.


## 1. Introduction

A synthrophic relationship between two organisms refers to a situation where the species exhibit mutualistism but where, at the opposite of what happens in a purely symbiotic relationship, one of the species can grow without the other. Such a situation can be mathematically formalized as follows. Assume that a first species denoted $X_{1}$ grows on a substrate $S_{1}$ forming an intermediate product $S_{2}$. This intermediate product is required by a second species $X_{2}$ to grow. The limiting substrate of the second bacteria being the product of the first bioreaction, the second bacteria cannot grow if the first one is not present.

Such interactions are quite common in nature: it is why a number of models have already been proposed in the literature. Katsuyama et al. [9], proposed a model involving two mutualistic species for describing pesticide degradation, while a more general case is considered by Kreikenbohm and Bohl [10]. Since mutualism involves generally species interacting through intermediate products, other studies consider mutualistic relationships in food webs. For instance, Bratbak and Thingstad [2], or more recently, Aota and Nakajima [1] considered the mutualism between phytoplankton and bacteria through the carbon excretion by the phytoplankton. A model studied by Freedman et al. [8] was proposed to explain the observed coexistence of such species. However, in the previous studies the models are very specific. In particular, the mathematical analyses of the models are realized for specific growth rates that are explicitely given (in most cases as Monod functions).

To extend the study of mutualism to more general systems, we have recently considered more general assumptions notably with respect to the growth rate functions considered in the models in using qualitative hypotheses, cf. [5] . Furthermore, it was assumed that the species $X_{1}$ may be inhibited by the product $S_{2}$ that it produces itself while the species $X_{2}$ was simply limited by $S_{2}$. An example of such interactions was given by the anaerobic digestion in which mutualistic relationships allow certain classes of bacteria to coexist. A mutualistic relation has been also considered in [4]. See [6] for another model of coexistence in the chemostat.

[^0]In the actual paper, following [7], we revisit the model proposed in [5] in considering two main changes which significantly further extend the range of practical situations covered by the model. First, we assume that there is some $S_{2}$ in the influent. In other terms, the limiting substrate $S_{2}$ on which the species $X_{2}$ grows is not only produced by the species $X_{1}$ but is also available even if the species $X_{1}$ is not present. The second modification of the model is that the second species is supposed to be inhibited by an excess of $S_{1}$, the limiting substrate on which the first species grows. To illustrate the usefulness of such extensions of the original model by El Hajji et al. [5], the biological interpretation of these hypotheses within the context of the anaerobic process is given in the appendix.

The paper is organized as follows. In Section 2, we propose a modified system of four differential equations from the original model in [5]. The positive equilibria are determined and their local and global stability properties are established. In the case when the system has a unique positive equilibrium, the global asymptotic stability results are demonstrated through the Dulac's criterion that rules out the possibility of the existence of periodic solutions for the reduced planar system, the Poincaré-Bendixon Theorem and the Butler-McGehee Lemma. Hence, in this case, for every positive initial conditions, the solutions converge to the positive equilibrium point which corresponds to the coexistence of the two bacterial species as observed in real processes. Simulations are presented in Section 4, an example of a syntrophic relationship is given in Section 6 as a candidate for this model.

## 2. Mathematical model

Let $S_{1}, X_{1}, S_{2}$ and $X_{2}$ denote, respectively, the concentrations of the substrate, the first bacteria, the intermediate product, and the second bacteria present in the reactor at time $t$. We neglect all species-specific death rates and take into account the dilution rate only. Hence our model is described by the following system of ordinary differential equations :

$$
\left\{\begin{align*}
\dot{S}_{1} & =D\left(S_{1}^{i n}-S_{1}\right)-k_{3} \mu_{1}\left(S_{1}, S_{2}\right) X_{1}  \tag{1}\\
\dot{X}_{1} & =\mu_{1}\left(S_{1}, S_{2}\right) X_{1}-D X_{1} \\
\dot{S}_{2} & =D\left(S_{2}^{i n}-S_{2}\right)-k_{2} \mu_{2}\left(S_{1}, S_{2}\right) X_{2}+k_{1} \mu_{1}\left(S_{1}, S_{2}\right) X_{1} \\
\dot{X}_{2} & =\mu_{2}\left(S_{1}, S_{2}\right) X_{2}-D X_{2}
\end{align*}\right.
$$

Where $S_{1}^{i n}>0$ denotes the input concentration of substrate, $S_{2}^{i n}>0$ denotes the input concentration of the intermediate product and $D>0$ is the dilution rate.

Assume that the functional response of each species $\mu_{1}, \mu_{2}: \mathbb{R}_{+}^{2} \rightarrow \mathbb{R}_{+}$satisfies :
A1: $\mu_{1}, \mu_{2}: \mathbb{R}_{+}^{2} \rightarrow \mathbb{R}_{+}$, of class $\mathcal{C}^{1}$,
A2: $\mu_{1}\left(0, S_{2}\right)=0, \quad \mu_{2}\left(S_{1}, 0\right)=0, \quad \forall\left(S_{1}, S_{2}\right) \in \mathbb{R}_{+}^{2}$,
A3: $\frac{\partial \mu_{1}}{\partial S_{1}}\left(S_{1}, S_{2}\right)>0, \quad \frac{\partial \mu_{1}}{\partial S_{2}}\left(S_{1}, S_{2}\right)<0, \quad \forall\left(S_{1}, S_{2}\right) \in \mathbb{R}_{+}^{2}$,
A4: $\frac{\partial \mu_{2}}{\partial S_{1}}\left(S_{1}, S_{2}\right)<0, \quad \frac{\partial \mu_{2}}{\partial S_{2}}\left(S_{1}, S_{2}\right)>0, \quad \forall\left(S_{1}, S_{2}\right) \in \mathbb{R}_{+}^{2}$.
Hypothesis A2 expresses that no growth can take place for species $X_{1}$ without the substrate $S_{1}$ and that the intermediate product $S_{2}$ is obligate for the growth of species $X_{2}$. Hypothesis A3 means that the growth of species $X_{1}$ increases with the substrate $S_{1}$ and it is inhibited by the intermediate product $S_{2}$ that it produces. Hypothesis A4 means that the growth of species $X_{2}$ increases with intermediate product $S_{2}$ produced by species $X_{1}$ while it is inhibited by the
substrate $S_{1}$. Note that there is a syntrophic relationship between the two species.
We first scale system (1) using the following change of variables and notations :

$$
s_{1}=\frac{k_{1}}{k_{3}} S_{1}, \quad x_{1}=k_{1} X_{1}, \quad s_{2}=S_{2}, \quad x_{2}=k_{2} X_{2}, \quad s_{1}^{i n}=\frac{k_{1}}{k_{3}} S_{1}^{i n}, \quad s_{2}^{i n}=S_{2}^{i n}
$$

The dimensionless equations thus obtained are :

$$
\left\{\begin{align*}
\dot{s}_{1} & =D\left(s_{1}^{i n}-s_{1}\right)-f_{1}\left(s_{1}, s_{2}\right) x_{1}  \tag{2}\\
\dot{x}_{1} & =f_{1}\left(s_{1}, s_{2}\right) x_{1}-D x_{1} \\
\dot{s}_{2} & =D\left(s_{2}^{i n}-s_{2}\right)-f_{2}\left(s_{1}, s_{2}\right) x_{2}+f_{1}\left(s_{1}, s_{2}\right) x_{1} \\
\dot{x}_{2} & =f_{2}\left(s_{1}, s_{2}\right) x_{2}-D x_{2}
\end{align*}\right.
$$

Where the functions $f_{1}, f_{2}: \mathbb{R}_{+}^{2} \rightarrow \mathbb{R}_{+}$are defined by

$$
f_{1}\left(s_{1}, s_{2}\right)=\mu_{1}\left(\frac{k_{3}}{k_{1}} s_{1}, s_{2}\right) \quad \text { and } \quad f_{2}\left(s_{1}, s_{2}\right)=\mu_{2}\left(\frac{k_{3}}{k_{1}} s_{1}, s_{2}\right)
$$

Hypotheses A1-A4 satisfied by the functions $\mu_{1}$ and $\mu_{2}$ translate in the following assumptions of the functions $f_{1}$ and $f_{2}$ :

H1: $f_{1}, f_{2}: \mathbb{R}_{+}^{2} \rightarrow \mathbb{R}_{+}$, of class $\mathcal{C}^{1}$,
H2: $f_{1}\left(0, s_{2}\right)=0, \quad f_{2}\left(s_{1}, 0\right)=0, \quad \forall\left(s_{1}, s_{2}\right) \in \mathbb{R}_{+}^{2}$,
H3: $\frac{\partial f_{1}}{\partial s_{1}}\left(s_{1}, s_{2}\right)>0, \quad \frac{\partial f_{1}}{\partial s_{2}}\left(s_{1}, s_{2}\right)<0, \quad \forall\left(s_{1}, s_{2}\right) \in \mathbb{R}_{+}^{2}$,
H4: $\frac{\partial f_{2}}{\partial s_{1}}\left(s_{1}, s_{2}\right)<0, \quad \frac{\partial f_{2}}{\partial s_{2}}\left(s_{1}, s_{2}\right)>0, \quad \forall\left(s_{1}, s_{2}\right) \in \mathbb{R}_{+}^{2}$.
$\mathbb{R}_{+}^{4}$, the closed non-negative cone in $\mathbb{R}^{4}$, is positively invariant under the solution map of system (2). More precisely

Proposition 1. For every initial condition in $\mathbb{R}_{+}^{4}$, the solution of system (2) has positive components and is positively bounded and thus is defined for every positive $t$. The set

$$
\Omega=\left\{\left(s_{1}, x_{1}, s_{2}, x_{2}\right) \in \mathbb{R}_{+}^{4}: s_{1}+x_{1}=s_{1}^{i n}, \quad s_{2}+x_{2}=x_{1}+s_{2}^{i n}\right\}
$$

is a positive invariant attractor of all solutions of system (2).
Proof. The invariance of $\mathbb{R}_{+}^{4}$ is guaranteed by the fact that :
i. $s_{1}=0 \Rightarrow \dot{s}_{1}=D s_{1}^{i n}>0$,
ii. $s_{2}=0 \Rightarrow \dot{s}_{2}=D s_{2}^{i n}+f_{1}\left(s_{1}, 0\right) x_{1}>0$,
iii. $x_{i}=0 \Rightarrow \dot{x}_{i}=0$ for $i=1,2$.

Next we have to prove that the solution is bounded. Let $z_{1}=s_{1}+x_{1}$, then $\dot{z}_{1}=-D\left(z_{1}-s_{1}^{i n}\right)$ from which one deduces :

$$
\begin{equation*}
s_{1}(t)+x_{1}(t)=s_{1}^{i n}+\left(s_{1}(0)+x_{1}(0)-s_{1}^{i n}\right) e^{-D t} \tag{3}
\end{equation*}
$$

Thus $s_{1}(t)$ and $x_{1}(t)$ are positively bounded. Let $z_{2}=s_{2}+x_{2}-x_{1}$, then $\dot{z}_{2}=-D\left(z_{2}-s_{2}^{i n}\right)$ from which one deduces:

$$
\begin{equation*}
s_{2}(t)+x_{2}(t)-x_{1}(t)=s_{2}^{i n}+\left(s_{2}(0)+x_{2}(0)-x_{1}(0)-s_{2}^{i n}\right) e^{-D t} \tag{4}
\end{equation*}
$$

Thus $s_{2}(t)$ and $x_{2}(t)$ are positively bounded. Hence, the solution is defined for all positive $t$. From (3) and (4) we deduce that the set $\Omega$ is an invariant set which is an attractor.

## 3. Restriction on the plane

The solutions of system (2) are exponentially convergent towards the set $\Omega$ and we are interested in the asymptotic behavior of these solutions. It is enough to restrict the study of the asymptotic behaviour of system (2) to $\Omega$. In fact, thanks to Thieme's results [12], the asymptotic behaviour of the solutions of the restriction of (2) on $\Omega$ will be informative for the complete system, see Section 5 . In this section we study the following reduced system which is simply the projection on the plane $\left(x_{1}, x_{2}\right)$, of the restriction of system (2) on $\Omega$.

$$
\left\{\begin{array}{l}
\dot{x}_{1}=\left[\Phi_{1}\left(x_{1}, x_{2}\right)-D\right] x_{1}  \tag{5}\\
\dot{x}_{2}=\left[\Phi_{2}\left(x_{1}, x_{2}\right)-D\right] x_{2}
\end{array}\right.
$$

where

$$
\Phi_{1}\left(x_{1}, x_{2}\right)=f_{1}\left(s_{1}^{i n}-x_{1}, s_{2}^{i n}+x_{1}-x_{2}\right), \quad \Phi_{2}\left(x_{1}, x_{2}\right)=f_{2}\left(s_{1}^{i n}-x_{1}, s_{2}^{i n}+x_{1}-x_{2}\right)
$$

Thus, for (5) the state-vector $\left(x_{1}, x_{2}\right)$ belongs to the following subset of the plane, see Fig. 1:

$$
\mathcal{S}=\left\{\left(x_{1}, x_{2}\right) \in \mathbb{R}_{+}^{2}: 0<x_{1} \leq s_{1}^{i n}, 0<x_{2} \leq x_{1}+s_{2}^{i n}\right\}
$$



Figure 1. The set $\mathcal{S}$

The point $F^{0}=(0,0)$ is an equilibrium of (5). Besides this equilibrium point the system can have the following three types of equilibrium points.

- Boundary equilibria $F^{1}=\left(\bar{x}_{1}, 0\right)$, where $x_{1}=\bar{x}_{1}$ is a solution, if it exists, of equation

$$
\begin{equation*}
\Phi_{1}\left(x_{1}, 0\right)=D \tag{6}
\end{equation*}
$$

- Boundary equilibria $F^{2}=\left(0, \tilde{x}_{2}\right)$, where $x_{2}=\tilde{x}_{2}$ is a solution, if it exists, of equation

$$
\begin{equation*}
\Phi_{2}\left(0, x_{2}\right)=D, \tag{7}
\end{equation*}
$$

- Positive equilibria $F^{*}=\left(x_{1}^{*}, x_{2}^{*}\right)$, where $x_{1}=x_{1}^{*}, x_{2}=x_{2}^{*}$ is a solution, if it exists, of the system of equations

$$
\left\{\begin{array}{l}
\Phi_{1}\left(x_{1}, x_{2}\right)=D  \tag{8}\\
\Phi_{2}\left(x_{1}, x_{2}\right)=D .
\end{array}\right.
$$

We use the following notations

$$
D_{1}=f_{1}\left(s_{1}^{i n}, s_{2}^{i n}\right), \quad D_{2}=f_{2}\left(s_{1}^{i n}, s_{2}^{i n}\right)
$$

The mapping $x_{1} \mapsto \Phi_{1}\left(x_{1}, 0\right)$ is decreasing, and the mapping $x_{1} \mapsto \Phi_{2}\left(x_{1}, 0\right)$ is increasing. If $D_{1}>D_{2}$, there exists a unique real number $\xi_{1}$ satisfying $\Phi_{1}\left(\xi_{1}, 0\right)=\Phi_{2}\left(\xi_{1}, 0\right)$, since

$$
\Phi_{1}(0,0)=D_{1}>D_{2}=\Phi_{2}(0,0), \text { and } \Phi_{1}\left(s_{1}^{i n}, 0\right)=0<\Phi_{2}\left(s_{1}^{i n}, 0\right)
$$

We denote by $\left.D_{3} \in\right] D_{2}, D_{1}$ [ the unique real number (see Figure 2, right) such that:

$$
\Phi_{1}\left(\xi_{1}, 0\right)=\Phi_{2}\left(\xi_{1}, 0\right)=D_{3}
$$



Figure 2. Existence and uniqueness of $\bar{x}_{1}$. On the left, the case $D_{1}<D_{2}$ : $\Phi_{2}\left(\bar{x}_{1}, 0\right)>D$ for all $D<D_{1}$. On the right, the case $D_{1}>D_{2}: \Phi_{2}\left(\bar{x}_{1}, 0\right)>D$ if and only if $D<D_{3}$.

The mapping $x_{2} \mapsto \Phi_{1}\left(0, x_{2}\right)$ is increasing, and the mapping $x_{2} \mapsto \Phi_{2}\left(0, x_{2}\right)$ is decreasing. Hence, if $D_{1}<D_{2}$, there exists a unique real number $\xi_{2}$ satisfying $\Phi_{1}\left(0, \xi_{2}\right)=\Phi_{2}\left(0, \xi_{2}\right)$, since

$$
\Phi_{2}(0,0)=D_{2}>D_{1}=\Phi_{1}(0,0), \text { and } \Phi_{2}\left(0, s_{2}^{i n}\right)=0<\Phi_{1}\left(0, s_{2}^{i n}\right)
$$

We denote by $\left.D_{4} \in\right] D_{1}, D_{2}$ [ the unique real number (see Figure 3, right) such that:

$$
\Phi_{1}\left(0, \xi_{2}\right)=\Phi_{2}\left(0, \xi_{2}\right)=D_{4}
$$

The nature of the trivial equilibrium point $F^{0}$ is given in the following lemma.


Figure 3. Existence and uniqueness of $\tilde{x}_{2}$. On the left, the case $D_{2}<D_{1}$ : $\Phi_{1}\left(0, \tilde{x}_{2}\right)>D$ for all $D<D_{2}$. On the right, the case $D_{2}>D_{1}: \Phi_{1}\left(0, \tilde{x}_{2}\right)>D$ if and only if $D<D_{4}$.

Lemma 1. If $D>\max \left(D_{1}, D_{2}\right)$ then $F^{0}$ is a stable node. If $\min \left(D_{1}, D_{2}\right)<D<\max \left(D_{1}, D_{2}\right)$ then $F^{0}$ is a saddle point. If $D<\min \left(D_{1}, D_{2}\right)$ then $F^{0}$ is an unstable node.

Proof. The Jacobian matrix $J$ of (5), at point $\left(x_{1}, x_{2}\right)$, is given by:

$$
J=\left[\begin{array}{cc}
-\frac{\partial f_{1}}{\partial s_{1}} x_{1}+\frac{\partial f_{1}}{\partial s_{2}} x_{1}+f_{1}-D & -\frac{\partial f_{1}}{\partial s_{2}} x_{1} \\
-\frac{\partial f_{2}}{\partial s_{1}} x_{2}+\frac{\partial f_{2}}{\partial s_{2}} x_{2} & -\frac{\partial f_{2}}{\partial s_{2}} x_{2}+f_{2}-D
\end{array}\right]
$$

where the functions are evaluated at $\left(s_{1}^{i n}-x_{1}, s_{2}^{i n}+x_{1}-x_{2}\right)$. The Jacobian matrix at $F^{0}$ is given by:

$$
J^{0}=\left[\begin{array}{cc}
f_{1}\left(s_{1}^{i n}, s_{2}^{i n}\right)-D & 0 \\
0 & f_{2}\left(s_{1}^{i n}, s_{2}^{i n}\right)-D
\end{array}\right]
$$

The eigenvalues are $D_{1}-D$ and $D_{2}-D$. Thus, if $D>\max \left(D_{1}, D_{2}\right)$ then $F^{0}$ is a stable node. It is an unstable node if $D<\min \left(D_{1}, D_{2}\right)$. It is a saddle point if $\min \left(D_{1}, D_{2}\right)<D<$ $\max \left(D_{1}, D_{2}\right)$.

The conditions of existence of the boundary equilibria $F^{1}$ and $F^{2}$, and their nature, are stated in the following lemmas.

Lemma 2. An equilibrium $F^{1}=\left(\bar{x}_{1}, 0\right)$ exists if and only if $D<D_{1}$. If it exists then it the unique equilibrium on the positive $x_{1}$ semi-axis. If $D_{1}<D_{2}$ then $F^{1}$ is a saddle point for all $D<D_{1}$. If $D_{2}<D_{1}$, then $F^{1}$ is a saddle point for all $0<D<D_{3}$ and a stable node for all $D_{3}<D<D_{1}$.

Proof. An equilibrium $F^{1}=\left(\bar{x}_{1}, 0\right)$ exists if and only if $\left.x_{1}=\bar{x}_{1} \in\right] 0, s_{1}^{i n}[$ is a solution of (6). Let $\psi_{1}\left(x_{1}\right)=\Phi_{1}\left(x_{1}, 0\right)$. Then

$$
\psi_{1}^{\prime}\left(x_{1}\right)=-\frac{\partial f_{1}}{\partial s_{1}}\left(s_{1}^{i n}-x_{1}, s_{2}^{i n}+x_{1}\right)+\frac{\partial f_{1}}{\partial s_{2}}\left(s_{1}^{i n}-x_{1}, s_{2}^{i n}+x_{1}\right) .
$$

By assumption H3, $\psi_{1}^{\prime}\left(x_{1}\right)<0$. Since $\psi_{1}(0)=D_{1}$, and $\psi_{1}\left(s_{1}^{i n}\right)=0$, equation (6) admits a solution in the interval ] $0, s_{1}^{i n}$ [ if and only if $D<D_{1}$. If this condition is satisfied then (6) admits a unique solution since the function $\psi_{1}($.$) is decreasing, see Figure 2. The Jacobian matrix at$ $F^{1}$ is given by:

$$
J^{1}=\left[\begin{array}{cc}
-\frac{\partial f_{1}}{\partial s_{1}} \bar{x}_{1}+\frac{\partial f_{1}}{\partial s_{2}} \bar{x}_{1} & -\frac{\partial f_{1}}{\partial s_{2}} \bar{x}_{1} \\
0 & f_{2}-D
\end{array}\right]
$$

where the functions are evaluated at $\left(s_{1}^{i n}-\bar{x}_{1}, s_{2}^{i n}+\bar{x}_{1}\right)$. The eigenvalues are

$$
f_{2}\left(s_{1}^{i n}-\bar{x}_{1}, s_{2}^{i n}+\bar{x}_{1}\right)-D=\Phi_{2}\left(\bar{x}_{1}, 0\right)-D, \text { and }-\frac{\partial f_{1}}{\partial s_{1}} \bar{x}_{1}+\frac{\partial f_{1}}{\partial s_{2}} \bar{x}_{1}<0
$$

Thus $F^{1}$ is a saddle point if $\Phi_{2}\left(\bar{x}_{1}, 0\right)>D$. If $D_{1}<D_{2}$, this condition is satisfied for all $D<D_{1}$. If $D_{2}<D_{1}$, it is statisfied for all $0<D<D_{3}$, see Figure 2. $F^{1}$ is a stable node if $D_{3}<D<D_{1}$ and $D_{2}<D_{1}$.

Lemma 3. An equilibrium $F^{2}=\left(0, \tilde{x}_{2}\right)$ exists if and only if $D<D_{2}$. If it exists then it the unique equilibrium on the positive $x_{2}$ semi-axis. If $D_{2}<D_{1}$ then $F^{2}$ is a saddle point for all $D<D_{2}$. If $D_{1}<D_{2}$, then $F^{2}$ is a saddle point for all $0<D<D_{4}$ and a stable node for all $D_{4}<D<D_{2}$.

Proof. An equilibrium $F^{2}=\left(0, \tilde{x}_{2}\right)$ exists if and only if $\left.x_{2}=\tilde{x}_{2} \in\right] 0, s_{2}^{\text {in }}$ [ is a solution of (7). Let $\psi_{2}\left(x_{2}\right)=\Phi_{2}\left(0, x_{2}\right)$. Then

$$
\psi_{2}^{\prime}\left(x_{2}\right)=-\frac{\partial f_{1}}{\partial s_{2}}\left(s_{1}^{i n}, s_{2}^{i n}-\tilde{x}_{2}\right)
$$

By assumption H4, $\psi_{2}^{\prime}\left(x_{2}\right)<0$. Since $\psi_{2}(0)=D_{2}$, and $\psi_{2}\left(s_{2}^{i n}\right)=0$, equation (7) admits a solution in the interval ] $0, s_{2}^{i n}$ [ if and only if $D<D_{2}$. If this condition is satisfied then (7) admits a unique solution since the function $\psi_{2}($.$) is decreasing, see Figure 3. The Jacobian matrix at$ $F^{2}$ is given by:

$$
J^{2}=\left[\begin{array}{cc}
f_{1}-D & 0 \\
-\frac{\partial f_{2}}{\partial s_{1}} \tilde{x}_{2}+\frac{\partial f_{2}}{\partial s_{2}} \tilde{x}_{2} & -\frac{\partial f_{2}}{\partial s_{2}} \tilde{x}_{2}
\end{array}\right]
$$

where the functions are evaluated at $\left(s_{1}^{i n}, s_{2}^{i n}-\tilde{x}_{2}\right)$. The eigenvalues are

$$
f_{1}\left(s_{1}^{i n}, s_{2}^{i n}-\tilde{x}_{2}\right)-D=\Phi_{1}\left(0, \tilde{x}_{2}\right)-D, \text { and }-\frac{\partial f_{2}}{\partial s_{2}} \tilde{x}_{2}<0 .
$$

Thus $F^{2}$ is a saddle point if $\Phi_{1}\left(0, \tilde{x}_{2}\right)>D$. If $D_{2}<D_{1}$, this condition is satisfied for all $D<D_{2}$. If $D_{1}<D_{2}$, it is statisfied for all $0<D<D_{4}$, see Figure 3. $F^{2}$ is a stable node if $D_{4}<D<D_{2}$ and $D_{1}<D_{2}$.

Let us discuss now the conditions of existence of positive equilibria $F^{*}$, and their number. An equilibrium $F^{*}=\left(x_{1}^{*}, x_{2}^{*}\right)$ exists if and only if $x_{1}=x_{1}^{*}, x_{2}=x_{2}^{*}$ is a solution of (8) lying in $\mathcal{S}$. One has

$$
\frac{\partial \Phi_{1}}{\partial x_{2}}=-\frac{\partial f_{1}}{\partial s_{2}}\left(s_{1}^{i n}-x_{1}, s_{2}^{i n}+x_{1}-x_{2}\right)
$$

By assumption H3, this partial derivative is positive. Hence, equation $\Phi_{1}\left(x_{1}, x_{2}\right)=D$ defines a function $x_{2}=F_{1}\left(x_{1}\right)$ such that $F_{1}\left(\bar{x}_{1}\right)=0$ when $D<D_{1}$. Recall that $x_{1}=\bar{x}_{1}$ is the solution of (6) which, according to Lemma 2 exists and is unique, if and only if $D<D_{1}$. One has

$$
F_{1}^{\prime}\left(x_{1}\right)=-\frac{\frac{\partial \Phi_{1}}{\partial x_{1}}\left(x_{1}, F_{1}\left(x_{1}\right)\right)}{\frac{\partial \Phi_{1}}{\partial x_{2}}\left(x_{1}, F_{1}\left(x_{1}\right)\right)}=\frac{-\frac{\partial f_{1}}{\partial s_{1}}+\frac{\partial f_{1}}{\partial s_{2}}}{\frac{\partial f_{1}}{\partial s_{2}}}=1-\frac{\frac{\partial f_{1}}{\partial s_{1}}}{\frac{\partial f_{1}}{\partial s_{2}}}>1 .
$$

Hence the function $F_{1}$ is increasing. Since $\Phi_{1}\left(s_{1}^{i n}, 0\right)=0$, the graph $\Gamma_{1}$ of $F_{1}$ has no intersection with the right boundary of the domain $\mathcal{S}$, defined by $x_{1}=s_{1}^{i n}$. This graph separates $\mathcal{S}$ in two regions denoted as the left and right sides of $\Gamma_{1}$, see Figure 4. One has also

$$
\frac{\partial \Phi_{2}}{\partial x_{2}}=-\frac{\partial f_{2}}{\partial s_{2}}\left(s_{1}^{i n}-x_{1}, s_{2}^{i n}+x_{1}-x_{2}\right)
$$

By assumption H3, this partial derivative is positive. Hence, equation $\Phi_{2}\left(x_{1}, x_{2}\right)=D$ defines a function $x_{2}=F_{2}\left(x_{1}\right)$ such that $F_{2}(0)=\tilde{x}_{2}$ when $D<D_{2}$. Recall that $x_{2}=\tilde{x}_{2}$ is the solution of (7) which, according to Lemma 3 exists and is unique, if and only if $D<D_{2}$. One has

$$
F_{2}^{\prime}\left(x_{1}\right)=-\frac{\frac{\partial \Phi_{2}}{\partial x_{1}}\left(x_{1}, F_{2}\left(x_{1}\right)\right)}{\frac{\partial \Phi_{2}}{\partial x_{2}}\left(x_{1}, F_{2}\left(x_{1}\right)\right)}=\frac{-\frac{\partial f_{2}}{\partial s_{1}}+\frac{\partial f_{2}}{\partial s_{2}}}{\frac{\partial f_{2}}{\partial s_{2}}}=1-\frac{\frac{\partial f_{2}}{\partial s_{1}}}{\frac{\partial f_{2}}{\partial s_{2}}}>1
$$

Hence the function $F_{2}$ is increasing. Since $\Phi_{2}\left(x_{1}, s_{2}^{i n}+x_{1}\right)=0$, the graph $\Gamma_{2}$ of $F_{2}$ has no intersection with the top boundary of the domain $\mathcal{S}$, defined by $x_{2}=s_{2}^{\text {in }}+x_{1}$. Thus the point at the very right of $\Gamma_{2}$ lies necessarily on the right boundary of $\mathcal{S}$, defined by $x_{1}=s_{1}^{i n}$. Hence it lies on the right side of $\Gamma_{1}$, see Figure 4.

The graphs $\Gamma_{1}$ and $\Gamma_{2}$ can intersect or not, see Figures 4,5 and 6. If they intersect at some point $F^{*}=\left(x_{1}^{*}, x_{2}^{*}\right)$ then $F^{*}$ is a positive equilibbrium. If the point $A$ at the very left of $\Gamma_{2}$


Figure 4. On the left, the left and right sides of $\Gamma_{1}$. On the center, the point $A$ at the very left of $\Gamma_{2}$ lies on left side of $\Gamma_{1}$ : there are generically an odd number of intersections (3 in this example). On the right, the point $A$ at the very left of $\Gamma_{2}$ lies on right side of $\Gamma_{1}$ : there are generically an even number of intersections (2 in this example).
lies on left side of $\Gamma_{1}$ then $\Gamma_{1}$ and $\Gamma_{2}$ intersect in at least one point $F^{*}=\left(x_{1}^{*}, x_{2}^{*}\right)$. They can have multiple intersections. Generically they have an odd number of intersections (see Figure 4, center). If the point $A$ at the very left of $\Gamma_{2}$ lies on right side of $\Gamma_{1}$ then $\Gamma_{1}$ and $\Gamma_{2}$ can intersect or not. Generically they have an even number of intersections (see Figure 4, right). The nature of a positive equilibrium $F^{*}$ is stated in the following lemmas.
Lemma 4. If an equilibrium $F^{*}=\left(x_{1}^{*}, x_{2}^{*}\right)$ exists then it is a stable node if $F_{1}^{\prime}\left(x_{1}^{*}\right)>F_{2}^{\prime}\left(x_{1}^{*}\right)$. It is a saddle point if the opposite inequality is satisfied.
Proof. The Jacobian matrix at $F^{*}$ is given by:

$$
J^{*}=\left[\begin{array}{cc}
-\frac{\partial f_{1}}{\partial s_{1}} x_{1}^{*}+\frac{\partial f_{1}}{\partial s_{2}} x_{1}^{*} & -\frac{\partial f_{1}}{\partial s_{2}} x_{1}^{*} \\
-\frac{\partial f_{2}}{\partial s_{1}} x_{2}^{*}+\frac{\partial f_{2}}{\partial s_{2}} x_{2}^{*} & -\frac{\partial f_{2}}{\partial s_{2}} x_{2}^{*}
\end{array}\right]
$$

where the derivatives are evaluated at $\left(s_{1}^{i n}-x_{1}^{*}, s_{2}^{i n}+x_{1}^{*}-x_{2}^{*}\right)$. Notice that

$$
\operatorname{tr}\left(J^{*}\right)=-\frac{\partial f_{1}}{\partial s_{1}} x_{1}^{*}+\frac{\partial f_{1}}{\partial s_{2}} x_{1}^{*}-\frac{\partial f_{2}}{\partial s_{2}} x_{2}^{*}<0
$$

and

$$
\operatorname{det}\left(J^{*}\right)=x_{1}^{*} x_{2}^{*}\left[\frac{\partial f_{1}}{\partial s_{1}} \frac{\partial f_{2}}{\partial s_{2}}-\frac{\partial f_{1}}{\partial s_{2}} \frac{\partial f_{2}}{\partial s_{1}}\right]=x_{1}^{*} x_{2}^{*} \frac{\partial f_{1}}{\partial s_{2}} \frac{\partial f_{2}}{\partial s_{2}}\left[F_{2}^{\prime}\left(x_{1}^{*}\right)-F_{1}^{\prime}\left(x_{1}^{*}\right)\right] .
$$

By Assumptions H3 and H4, the product of the partial derivatives is negative. Therefore, the determinant is positive if $F_{1}^{\prime}\left(x_{1}^{*}\right)>F_{2}^{\prime}\left(x_{1}^{*}\right)$ and negative if the opposite inequality is satisfied. Hence the equilibrium $F^{*}=\left(x_{1}^{*}, x_{2}^{*}\right)$ is a stable node if $F_{1}^{\prime}\left(x_{1}^{*}\right)>F_{2}^{\prime}\left(x_{1}^{*}\right)$. It is a saddle point if the opposite inequality is satisfied.

The number of equilibria of (5) and their nature are summarized in the next theorem.
Theorem 1. (1) If $D<\min \left(D_{1}, D_{2}\right)$ then (5) admits the trivial equilibrium $F^{0}$ which is an unstable node, the boundary equilibria $F^{1}$ and $F^{2}$ which are saddle points, and at least one positive equilibrium $F^{*}$. If $F^{*}$ is the unique positive equilibrium then it is a stable node. Generically, the system has an odd number of positive equilibria which are
alternatively stable nodes and saddle points, the one at the very left of these positive equilibria is a stable node.
(2) If $\min \left(D_{1}, D_{2}\right)<D<\max \left(D_{1}, D_{2}\right)$, four subcases must be distinguished
(a) If $D_{1}<D_{2}$ and $D_{1}<D<D_{4}$ then (5) admits the trivial and boundary equilibria $F^{0}$ and $F^{2}$, which are saddle points and at least one positive equilibrium $F^{*}$. If $F^{*}$ is the unique positive equilibrium then it is a stable node. Generically, the system has an odd number of positive equilibria which are alternatively stable nodes and saddle points, the one at the very left of these positive equilibria is a stable node.
(b) If $D_{1}<D_{2}$ and $D_{4}<D<D_{2}$ then (5) admits the trivial equilibrium $F^{0}$, which is a saddle point, and the boundary equilibrium $F^{2}$, which is a stable node. Generically, the system can have an even number of positive equilibria which are alternatively saddle points and stable nodes, the one at the very left of these positive equilibria is a saddle point.
(c) If $D_{2}<D_{1}$ and $D_{2}<D<D_{3}$ then (5) admits the trivial and boundary equilibria $F^{0}$ and $F^{1}$ which are saddle points and at least one positive equilibrium $F^{*}$. If $F^{*}$ is the unique positive equilibrium then it is a stable node. Generically, the system has an odd number of positive equilibria which are alternatively stable nodes and saddle points, the one at the very left of these positive equilibria is a stable node.
(d) If $D_{2}<D_{1}$ and $D_{3}<D<D_{1}$ then (5) admits the trivial equilibrium $F^{0}$, which is a saddle point, and the boundary equilibrium $F^{1}$, which is a stable node. Generically, the system can have an even number of positive equilibria which are alternatively saddle points and stable nodes, the one at the very left of these positive equilibria is a saddle point.
(3) If $D>\max \left(D_{1}, D_{2}\right)$ then (5) admits the trivial equilibrium $F^{0}$ which is a stable node. Generically, the system can have an even number of positive equilibria which are alternatively saddle points and stable nodes, the one at the very left of these positive equilibria is a saddle point.

## 4. Growth functions of Monod type

In this section we consider growth functions $f_{1}$ and $f_{2}$ of the following form

$$
\begin{equation*}
f_{1}\left(s_{1}, s_{2}\right)=\frac{m_{1} s_{1}}{\left(K_{1}+s_{1}\right)\left(L_{1}+s_{2}\right)}, \quad f_{2}\left(s_{1}, s_{2}\right)=\frac{m_{2} s_{1}}{\left(K_{2}+s_{1}\right)\left(L_{2}+s_{2}\right)} \tag{9}
\end{equation*}
$$

Such functions are simply the product of a Monod function in $s_{1}$ by a decreasing functions of $s_{2}$. Such functions are currently used in biotechnology when the growth of a functional species is limited by a substrate while inhibited by another one. Such situations are common in water treatment technology like in the denitrification (limited by the nitrate and inhibited by the dissolved oxygen) or in the anoxic or anaerobic hydrolysis (limited by the slowly biodegradable substrates while inhibited by an excess of oxygen) processes which are modeled this way (cf. [13]).

One can readily check that (9) satisfy Assumptions H1 to H4. By straighforward calculations one has

$$
\begin{aligned}
F_{1}\left(x_{1}\right) & =\frac{-D x_{1}^{2}+\left[m_{1}+D\left(K_{1}-L_{1}+s_{1}^{i n}-s_{2}^{i n}\right)\right]-m_{1} s_{1}^{i n}+D\left(K_{1}+s_{1}^{i n}\right)\left(L_{1}+s_{2}^{i n}\right)}{D\left(K_{1}+s_{1}^{i n}-x_{1}\right)} \\
F_{2}\left(x_{1}\right) & =\frac{D x_{1}^{2}+\left[m_{2}+D\left(L_{2}-K_{2}+s_{2}^{i n}-s_{1}^{i n}\right)\right]+m_{2} s_{2}^{i n}-D\left(K_{2}+s_{1}^{i n}\right)\left(L_{2}+s_{2}^{i n}\right)}{m_{2}-D\left(K_{2}+s_{1}^{i n}-x_{1}\right)}
\end{aligned}
$$

Hence equation $F_{1}\left(x_{1}\right)=F_{2}\left(x_{1}\right)$ giving the abscissa of positive equilibria is an algebraic equation of degree 2. Thus, it cannot have more than two solutions. Hence, the situation depicted on the
center of Figure 4, of three positive equilibria, is excluded. However, the situation depicted in the right of Figure 4, with two positive equilibria can occur. For instance, consider the following


Figure 5. Relative positions of the isocline $\dot{x}_{1}=0$ (in red) and $\dot{x}_{2}=0$ (in green).
values of the parameters

$$
\begin{equation*}
m_{1}=8, \quad m_{2}=4, \quad K_{1}=L_{2}=1, \quad L_{1}=K_{2}=2, \quad s_{1}^{i n}=s_{2}^{i n}=3 \tag{10}
\end{equation*}
$$

Then

$$
D_{1}=6 / 5, \quad D_{3}=8 / 9, \quad D_{2}=3 / 5
$$

There is another bifurcation value, $D=1$ which correspond to the case when the graphs $\Gamma_{1}$ and $\Gamma_{2}$ are tangent, see Figure 6. For this example five cases can occur, see Figure 5:

Proposition 2. Consider system (5) where $f_{1}$ and $f_{2}$ are given by (9) with parameters (10). Then
(1) when $D<3 / 5$, the system has four equilibria, $F^{0}$ which is an unstable node, $F^{1}$ and $F^{2}$, which are saddle points and $F^{*}$, which is a stable node. This is case (1) of Theorem 1, with a unique positive equilibrium.
(2) when $3 / 5<D<8 / 9$, the system has three equilibria, $F^{0}$ and $F^{1}$, which are saddle points and $F^{*}$, which is a stable node. This is case (2.c) of Theorem 1, with a unique positive equilibrium.
(3) when $8 / 9<D<1$, the system has four equilibria, $F^{0}$ and $F_{1}^{*}$, which are saddle points and $F^{1}$ and $F_{2}^{*}$, which are stable nodes. This is case (2.d) of Theorem 1, with two positive equilibria.
(4) when $1<D<6 / 5$, the system has two equilibria, $F^{0}$, which is a saddle point and $F^{1}$ which is a stable node. This is case (2.d) of Theorem 1, with no positive equilibrium.
(5) when $D>6 / 5$, the system has one equilibrium, $F^{0}$, which is a stable node. This is case (3) of Theorem 1, with no positive equilibrium.

In the case when $8 / 9<D<1$ a bistability phenomenon occurs. According to the initial condition, both species can coexist at equilibrium $F_{2}^{*}$, or species $x_{2}$ goes to extinction at equilibrium $F^{1}$. This phenomenon is illustarted numerically with $D=0.95$ in Figure 7.

For the following values of the parameters

$$
\begin{equation*}
m_{1}=8, \quad m_{2}=7, \quad K_{1}=K_{2}=L_{2}=1, \quad L_{1}=3 / 2, \quad s_{1}^{i n}=s_{2}^{i n}=3 \tag{11}
\end{equation*}
$$

the bifurcational values are $D_{1}=4 / 3$ and $D_{2}=21 / 16$. If $D>\max \left(D_{1}, D_{2}\right)$, for instance for $D=3 / 2$, one obtains a bistability phenomenon corresponding to case (3) of Theorem 1, with two positive equilibria. According to the initial condition, both species can coexist at equilibrium $F_{2}^{*}$,


Figure 6. The non hyperbolic cases. When $D=6 / 5, F^{0}$ and $F^{2}$ coalesce. When $D=8 / 9, F_{1}^{*}$ and $F^{1}$ coalesce (saddle node bifurcation). When $D=1$, $F_{1}^{*}$ and $F_{2}^{*}$ coalesce (saddle node bifurcation). When $D=6 / 5, F^{0}$ and $F^{2}$ coalesce.


Figure 7. Numerical solutions in the bistability case $D=0.95$ and parameters values (10). On the left, the separatrix (in green) of the saddle point $F_{1}^{*}$ separate the domain $\mathcal{S}$ in two region which are the basins of attraction of the boundary equilibrium point $F^{1}$ and the positive equilibrium point $F_{2}^{*}$. On the center, the phase portrait. On the right, the isoclines.
or both species go to extinction at equilibrium $F^{0}$. This phenomenon is illustarted numerically in Figure 8.

## 5. Global analysis

Let us establish first that (5) admits no periodic orbit nor polycycle inside $\mathcal{S}$
Theorem 2. There are no periodic orbits nor polycycles inside $\mathcal{S}$.
Proof. . Consider a trajectory of (5) belonging to $\mathcal{S}$. Let us transform the system (5) through the change of variables $\xi_{1}=\ln \left(x_{1}\right), \xi_{2}=\ln \left(x_{2}\right)$. Then one obtains the following system :

$$
\left\{\begin{array}{l}
\dot{\xi_{1}}=h_{1}\left(\xi_{1}, \xi_{2}\right):=f_{1}\left(s_{1}^{i n}-e^{\xi_{1}}, s_{2}^{i n}+e^{\xi_{1}}-e^{\xi_{2}}\right)-D,  \tag{12}\\
\dot{\xi_{2}}=h_{1}\left(\xi_{1}, \xi_{2}\right):=f_{2}\left(s_{1}^{i n}-e^{\xi_{1}}, s_{2}^{i n}+e^{\xi_{1}}-e^{\xi_{2}}\right)-D .
\end{array}\right.
$$

We have

$$
\frac{\partial h_{1}}{\partial \xi_{1}}+\frac{\partial h_{2}}{\partial \xi_{2}}=-e^{\xi_{1}} \frac{\partial f_{1}}{\partial s_{1}}+e^{\xi_{1}} \frac{\partial f_{1}}{\partial s_{2}}-e^{\xi_{2}} \frac{\partial f_{2}}{\partial s_{2}}<0 .
$$





Figure 8. Numerical solutions in the bistability case $D=1.5$ and parameters values (11). On the left, the separatrix (in green) of the saddle point $F_{1}^{*}$ separate the domain $\mathcal{S}$ in two region which are the basins of attraction of the boundary equilibrium point $F^{0}$ and the positive equilibrium point $F_{2}^{*}$. On the center, the phase portrait. On the right, the isoclines.

From Dulac criterion [11], we deduce that the system (12) has no periodic trajectory. Hence (5) has no periodic orbit in $\mathcal{S}$.

Theorem 3. Assume that system (5) has at most one positive quilibrium $F^{*}$, then for every initial condition in $\mathcal{S}$, the trajectories of system (5) converge asymptotically to :

- $F^{*}$ if $D<\min \left(D_{1}, D_{2}\right)$.
- $F^{*}$ if $D_{1}<D_{2}$ and $D_{1}<D<D_{4}$
- $F^{2}$ if $D_{1}<D_{2}$ and $D_{4}<D<D_{2}$.
- $F^{*}$ if $D_{2}<D_{1}$ and $D_{2}<D<D_{3}$.
- $F^{1}$ if $D_{2}<D_{1}$ and $D_{3}<D<D_{1}$.
- $F^{0}$ if $\max \left(D_{1}, D_{4}\right)<D$.

Proof. We restrict the proof to the situation where $D<\min \left(D_{1}, D_{2}\right)$. The other cases can be done similarly. Let $x_{1}(0)>0, x_{2}(0)>0$ and $\omega$ the $\omega$-limit set of $\left(x_{1}(0), x_{2}(0)\right) . \omega$ is an invariant compact set and $\omega \subset \overline{\mathcal{S}}$. Assume that $\omega$ contains a point $M$ on the $x_{1} x_{2}$ axis :

- $M$ can't be $F^{0}$ because $F^{0}$ is an unstable node and can't be a part of the $\omega$-limit set of $\left(x_{1}(0), x_{2}(0)\right)$,
- If $\left.M \in] \bar{x}_{1}, s_{1}^{i n}\right] \times\{0\}$ (respectively $\left.\left.M \in\{0\} \times\right] \bar{x}_{2}, s_{2}^{i n}\right]$ ). As $\omega$ is invariant then $\gamma(M) \subset$ $\omega$ which is impossible because $\omega$ is bounded and $\gamma(M)=] \bar{x}_{1},+\infty[\times\{0\}$ (respectively $\gamma(M)=\{0\} \times] \bar{x}_{2},+\infty[)$,
- If $M \in] 0, \bar{x}_{1}[\times\{0\}$ (respectively $M \in\{0\}, \times] 0, \bar{x}_{2}[$ ). $\omega$ contains $\gamma(M)=] 0, \bar{x}_{1}[\times\{0\}$ (respectively $\gamma(M)=\{0\} \times] 0, \bar{x}_{2}[$ ). As $\omega$ is a compact, then it contains the adherence of $\gamma(M),\left[0, \bar{x}_{1}\right] \times\{0\}$ (respectively $\{0\} \times\left[0, \bar{x}_{2}\right]$ ). In particular, $\omega$ contains $F^{0}$ which is impossible,
- If $M=F^{1}$ (respectively $M=F^{2}$ ). $\omega$ is not reduced to $F^{1}$ (respectively to $F^{2}$ ). By Butler-McGehee theorem, $\omega$ contains a point $P$ of $(0,+\infty) \times\{0\}$ other that $F^{1}$ (respectively of $\{0\} \times(0,+\infty)$ other that $\left.F^{2}\right)$ which is impossible.
Finally, the $\omega$-limit set don't contain any point on the $x_{1} x_{2}$ axis. System (5) has no periodic orbit inside $\mathcal{S}$. Using the Poincar-Bendixon Theorem [11], $F^{*}$ is a globally asymptotically stable equilibrium point for system (5).

Theorem 4. Assume that system (5) has at most one positive quilibrium $F^{*}$, then for every initial condition in $\mathbb{R}_{+}^{4}$, the trajectories of system (2) converge asymptotically to:

- $E^{*}$ if $D<\min \left(D_{1}, D_{4}\right)$.
- $E^{*}$ if $D_{1}<D_{2}$ and $D_{1}<D<D_{4}$.
- $E^{2}$ if $D_{1}<D_{2}$ and $D_{4}<D<D_{2}$.
- $E^{*}$ if $D_{2}<D_{1}$ and $D_{2}<D<D_{3}$.
- $E^{1}$ if $D_{2}<D_{1}$ and $D_{3}<D<D_{1}$.
- $E^{0}$ if $\max \left(D_{1}, D_{2}\right)<D$.

Proof. Let $\left(s_{1}(t), x_{1}(t), s_{2}(t), x_{2}(t)\right)$ be a solution of (2). From (3) and (4) we deduce that

$$
s_{1}(t)=s_{1}^{i n}-x_{1}(t)+K_{1} e^{-D t} \quad \text { and } \quad s_{2}(t)=s_{2}^{i n}+x_{1}(t)-x_{2}(t)+K_{2} e^{-D t}
$$

where $K_{1}=s_{1}(0)+x_{1}(0)-s_{1}^{i n}$ and $K_{2}=s_{2}(0)+x_{2}(0)-x_{1}(0)-s_{2}^{i n}$. Hence $\left(x_{1}(t), x_{2}(t)\right)$ is a solution of the nonautonomous system of two differential equations :

$$
\left\{\begin{align*}
\dot{x}_{1} & =\left[f_{1}\left(s_{1}^{i n}-x_{1}+K_{1} e^{-D t}, s_{2}^{i n}+x_{1}-x_{2}+K_{2} e^{-D t}\right)-D\right] x_{1}  \tag{13}\\
\dot{x}_{2} & =\left[f_{2}\left(s_{1}^{i n}-x_{1}+K_{1} e^{-D t}, s_{2}^{i n}+x_{1}-x_{2}+K_{2} e^{-D t}\right)-D\right] x_{2}
\end{align*}\right.
$$

This is an asymptotically autonomous differential system which converge to the autonomous system (5). The set $\Omega$ is attractor of all trajectories in $\mathbb{R}_{+}^{4}$ and the phase portrait of system reduced to $\Omega(5)$ contains only locally stable nodes, unstable nodes, saddle points and no trajectory joining two saddle points. Thus we can apply Thiemes's results [12] and conclude that the asymptotic behaviour of the solution of the complete system (13) is the same that the asymptotic behaviour described for the reduced system (5) and the main result is then deduced.

## 6. The anaerobic digestion process : An example of a synthrophic relationship

"Methane fermentation" or "anaerobic digestion" is a process that converts organic matter into a gaseous mixture mainly composed of methane and carbon dioxide $\left(\mathrm{CH}_{4}\right.$ and $\left.\mathrm{CO}_{2}\right)$ through the action of a complex bacterial ecosystem (cf. Fig.9). It is often used for the treatment of concentrated wastewaters or to stabilize the excess sludge produced in wastewater treatment plants into more stable products. There is also considerable interest in plant-biomass-fed digesters, since the produced methane can be valorized as a source of energy. It is usually considered that a number of metabolic groups of bacteria are involved sequencially.

One specific characteristic of the anaerobic process is that within such groups, there exists populations exhibiting obligatory mutualistic relationships. Such a syntrophic relationship is necessary for the biological reactions to be thermodynamically possible. In the first steps of the reactions (called "acidogenesis"), some hydrogen is produced. In El Hajji et al.[5], this production of hydrogen at this reaction step was neglected (compare Fig. 9 with Fig. 1 of [5]). This hypothesis constitue the first novelty with respect to [5]. It is to be noticed that an excess of hydrogen in the medium inhibits the growth of another bacterial group called "acetogenic bacteria". Their association with $\mathrm{H}_{2}$ consuming bacteria is thus necessary for the second step of the reaction to be fulfilled. Such a syntrophic relationship has been pointed out in a number of experimental works (cf. for instance the seminal work by [3]). Let us consider the subsystem of the anaerobic system where the VFA (for Volatile Fatty Acids) are transformed into $H_{2}$, $\mathrm{CH}_{4}$ and CO 2 . We can formalize the corresponding biological reactions as a first bacterial consortium $X_{1}$ (the acetogens) transforming $S_{1}$ (the VFA) into $S_{2}$ (the hydrogen) and acetate (cf. Fig.9). Then, a second species $X_{2}$ (the hydrogenotrophic-methanogenic bacteria) grows on $S_{2}$. In practice, acetogens are inhibited by an excess of hydrogen and methanogens by an excess of VFA. Thus, it is further assumed that $X_{1}$ is inhibitied by $S_{2}$ and $X_{2}$ by $S_{1}$. The last inhibition relationship constiute the second novelty with respect to [5]. This situation is precisely the one considered within the model (1).


Considered reactional part


Figure 9. Anaerobic fermentation process

We have proposed a mathematical model involving a syntrophic relationship of two bacteria. It results from this analysis that, under general and natural assumptions of monotonicity on the functional responses, the stable asymptotic coexistence of the two bacteria is possible.

## Acknowledgements

The authors acknowledge Inra and Inria for financial support.

## References

[1] Y. Aota and H. Nakajima, Mutualistic relationships between phytoplankton and bacteria caused by carbon excretion from phytoplankton, Ecological research, 16 (2001), 289-299.
[2] G. Bratbak and T.F. Thingstad, Phytoplankton-bacteria interactions: an apparent paradox? Analysis of a model system with both copetition and commensalism, Ecological research, 25 (1985), 23-30.
[3] M.P. Bryant, E.A. Wolin, M.J. Wolin and R.S. Wolfe, Methanobacillus omelianskii, a symbiotic association of two species of bacteria, Arch. Microbiol. 59 (1967), no. 1, 20-31.
[4] M. El Hajji, J. Harmand, H. Chaker and C. Lobry, Association between competition and obligate mutualism in a chemostat, J. Biol. Dynamics, 3 (2009), no. 6, 635-647.
[5] M. El Hajji, F. Mazenc and J. Harmand, A mathematical study of a syntrophic relationship of a model of anaerobic digestion process, Math. Biosci. Eng. 7 (2010), no. 3, 641-656.
[6] M. El Hajji and A. Rapaport, Practical coexistence of two species in the chemostat - A slow-fast characterization, Math. Biosci. 218 (2009), no. 1, 33-39.
[7] M. El Hajji, T. Sari, J. Harmand, Analyse d'un relation syntrophique: cas d'un chemostat, Proceedings of the 5th conference on Trends in Applied Mathematics in Tunisia, Algeria, Morocco, Sousse, 23-26 Avril 2011, Tunisia, M. Hassine and M. Moakher (Editors), Centre de Publication Universitaire (2011) , 451-456.
[8] H.I. Freedman, R. Kumar, A.K. Easton and M. Singh, Mathematical models of predator mutualists, Canadian Appl. Math. Quart. 9 (2001), 99-111.
[9] C. Katsuyama, S. Nakaoka, Y. Takeuchi, K. Tago, M. Hayatsu and K. Kato, A mathematical model of syntrophic cocultures in the chemostat, J. Theor. Biol. 256 (2009), 644-654.
[10] R. Kreikenbohm and E. Bohl, A mathematical model of syntrophic cocultures in the chemostat, FEMS Microbiol. Ecol. 38 (1986), 131-140.
[11] H.L. Smith and P. Waltman, The theory of the chemostat, Dynamics of microbial competition, Cambridge Studies in Mathematical Biology, Cambridge University Press, 1995.
[12] H.R. Thieme, Convergence results and a Poincaré-Bendixson trichotomy for asymptotically autonomous differential equations, J. Math. Biol. 30 (1992), no. 7, 755-763.
[13] The IWA Task Group on Mathematical Modelling for Design and Operation of Biological Wastewater Treatment, Activated Sludge Models ASM1, ASM2, ASM2d and ASM3, Scientific and Technical Report No.9, (2000), IWA publishing, 130 pages.

Tewfik Sari, Universite de Haute Alsace, LMiA, 4 rue des Frères Lumière, 68093 Mulhouse, \& InraInria Modemic research team, UMR Mistea, SupaAgro, 2 place Viala, 34060 Montpellier, France.

E-mail address: Tewfik.Sari@uha.fr
Miled EL Hajui, ISSAT, Universit de Kairouan 3000 Avenue Beit el hikma, Kairouan Tunisie E-mail address: miled.elhajji@enit.rnu.tn
Jérôme Harmand, LBE-INRA, UR050, Avenue des Étangs, 11100 Narbonne, France \& Inra-Inria Modemic research team, UMR Mistea, SupaAgro, 2 place Viala, 34060 Montpellier, France.

E-mail address: harmand@supagro.inra.fr


[^0]:    Date: July 29, 2011.
    2000 Mathematics Subject Classification. 92A15, 92A17, 34C15, 34C35, 34E18.
    Key words and phrases. Syntrophic relationship; Mathematical modelling; Coexistence; Asymptotic stability; Anaerobic digestion.

