Detection of mixed-culture growth in the total biomass data by wavelet transforms

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ABSTRACT

We have shown elsewhere that the presence of mixed-culture growth of microbial species in fermentation processes can be detected with high accuracy by employing the wavelet transform. This is achieved because the crosses in the different growth processes contributing to the total biomass signal appear as singularities that are very well evidenced through their singularity cones in the wavelet transform; however, we used very simple two-species cases.

In this work, we extend the wavelet method to a more complicated illustrative fermentation case of three microbial species for which we employ several wavelets of different number of vanishing moments in order to eliminate possible numerical artifacts. Working in this way allows filtering in a more precise way the numerical values of the Hölder exponents; therefore, we were able to determine the characteristic Hölder exponents for the corresponding crossing singularities of the microbial growth processes and their stability logarithmic scale ranges up to the first decimal in the value of the characteristic exponents. Since calibrating the mixed microbial growth by means of their Hölder exponents could have potential industrial applications, the dependence of the Hölder exponents on the kinetic and physical parameters of the growth models remains as a future experimental task.

Keywords: mixed-culture growth, total biomass, wavelets, Hölder exponents.

RESUMEN

Hemos mostrado en un trabajo anterior que la presencia de crecimiento mixto de poblaciones en procesos de fermentación puede ser inferida con gran precisión por medio del análisis ondeleta. Esto fue realizado debido a que los cruces en los diferentes procesos de crecimiento que contribuyen en la señal de biomasa total surgen como singularidades bien delineadas en los conos de singularidad en la transformada ondeleta; sin embargo, usamos casos muy simples de dos especies.

En este trabajo extendemos el estudio de tipo ondeleta a un caso ilustrativo de fermentación más complicado de tres poblaciones microbianas, en el cual se usaron varios tipos de ondeletas de diferente número de momentos de desvanecimiento para evitar los posibles artefactos numéricos, esto nos permitió filtrar de manera más precisa los valores numéricos de los exponentes de Hölder; por consiguiente, pudimos determinar los exponentes característicos de Hölder para las singularidades de cruce correspondientes a los procesos de crecimiento microbiano así como los rangos estables del logaritmo de la escala hasta el primer decimal en el valor de los exponentes característicos. Debido a que la calibración del crecimiento mixto microbiano por medio de los exponentes de Hölder podría tener aplicaciones industriales potenciales, la dependencia de los exponentes de Hölder en función de los parámetros cinéticos de los modelos de crecimiento queda como una futura tarea experimental.

1. Introduction

The growth of microbial species in media containing two or more substrates limiting the growth is of considerable biotechnological and

bioengineering interest [1]. The mixed growth of microorganisms occurs in many industrial processes. An important class of such processes is that of the traditional food and beverage fermentation in which either ambient

inoculums with selected microorganisms or microorganisms are used. The presence of different microbial species and substrates is a dominant factor for the quality and quantity of the final product. Biomass measurements are among the most important measurements in any cell culture process and detecting the presence or absence of the mixed culture growing processes is a demanding issue. In general terms, the biomass measurements can by divided in continuous and on-line, and discrete and off-line. The continuous biomass acquisition signals and reliable on-line methods yield valuable knowledge on the status of the process and can facilitate the process monitoring and control. Several methods have been developed for the continuous measurement of biomass. Traditional optical methods such as absorbance measurements are probably the easiest ways and are the most commonly used. Additionally, capacitance has also been used to estimate the biomass concentration in a continuous and online way, see for instance the web pages of the Fogale and Applikon companies: http://www.fogale.fr/biotech/pages/yeasts-2.php and http://www.applikonbio.com/applikonbio/c8-1.htm, respectively.

Our main goal is to show that it is possible to infer the mixed culture microbial growth based only on the knowledge of total biomass data, without employing complicated techniques. The alternative procedure that we emphasized recently and it is here shortly presented is to apply the wavelet method to the fermentation data with the purpose of detecting singularities in the growth curves (signals). In this case, one can consider the mixed culture growth curves as more or less regular signals containing crosses due to the difference in growth of the different species. In the wavelet literature, one can find fundamental papers in which it is shown that the wavelet techniques provide an efficient tool for detecting singularities. see for example Mallat and Hwang [2].

2. Microbial kinetics

The fermentation processes on which we focus here evolve in stirred batch tank reactors. In such a case, the biomass converts the substrate in additional biomass and products. The dynamical model of the bioreactor for biomass production is given by the following ordinary differential equations:

$$\frac{dx_{1,i}}{dt} = \mu_i(x_{2,i})x_{1,i},
\frac{dx_{2,i}}{dt} = -\mu_i(x_{2,i})\frac{x_{1,i}}{Y_i},$$
(1)

where $x_{1,i}$ are the biomass concentrations, $x_{2,i}$ the concentrations of the substrate, Y_i is the yield factor defined as the ratio of produced biomass per unit of time and the amount of consumed substrate per unit of time for each species, $\mu_i(x_{2,i})$ are the growth rates of each species and the sub index i represents the number of species and substrates involved in the fermentation process. The growth rates relate the changes in the biomass concentrations with the (negative) changes of the substrate concentrations per unit of time. We will use the formulas of Monod and Haldane, respectively:

1.- For the substrate saturation model (the Monod equation):

$$\mu_i(x_{2,i}) = \frac{\mu_{\max_i}(x_{2,i})}{K_{1,i} + x_{2,i}} , \qquad (2)$$

where is the saturation (or Monod) constant.

2.- For the substrate inhibition model (the Haldane equation):

$$\mu_i(x_{2,i}) = \frac{\mu_{\max_i}(x_{2,i})}{K_{1_i} + x_{2,i} + K_{2_i} x_{2,i}^2},$$
(3)

where, in addition to the saturation constant, there is an inhibition constant K_{2_i} . In both models, μ_{\max_i} is the maximal specific growth rate; for simplicity, this parameter has been taken as unity in the illustrative case presented in the following lines. The value of K_{1_i} expresses the affinity of the biomass for the substrate. The Monod growth kinetics can be considered as a special case of the substrate inhibition kinetics for $K_{2_i} = 0$, i.e., when

the inhibition terms vanish. The total biomass signal (TBS) is naturally given by

$$y(t) = \sum_{i=1}^{m} x_{1,i},$$
 (4)

where *m* is the total number of microbial species inside the bioreactor. In our previous work [3], only the simple cases of two species and two substrates have been considered. In Section 3, we will present a more complicated case, namely the case of three species, i.e., m = 3, with identical initial conditions for all species and substrates.

3. Wavelet analysis

In the context of signal theory, the total biomass signal y(t) is projected onto a wavelet basis through the correlation of the signal with an integral kernel which is a scaled and translated version of an analyzing wavelet

$$T_{\psi}[y](a,b) = \frac{1}{\sqrt{a}} \int_{-\infty}^{+\infty} \psi\left(\frac{t-b}{a}\right) y(t) dt, \qquad (5)$$

where ψ is the analyzing wavelet. The values of (5) are called the wavelet coefficients of signal y(t). In general, a signal can be disentangled into three parts: (a) a mostly smooth continuous structure that can be represented by low-order (piecewise) polynomials, (b) a discontinuous part intrinsic to the particular nature of the signal, and (c) a noise part which can also be quite discontinuous.

In general, for any function f(t) we can define its

moments as $M_k = \int_{-\infty}^{+\infty} t^k f(t) dt$, for k a

nonnegative integer. In the case of wavelets, an analyzing wavelet ψ has p vanishing moments if and only if it satisfies

$$\int_{-\infty}^{+\infty} t^k \psi(t) dt = 0, \text{ for all } 0 \le k < p.$$
 (6)

This property means that the analyzing wavelet ψ is orthogonal to any polynomial having degree up to p-1. The closer a signal is to being orthogonal to the analyzing wavelet, the smaller the resulting wavelet coefficient. Thus, if the analyzed signal can be approximated by a low-order polynomial over the support of the wavelet, then the wavelet is nearly orthogonal to the signal and the resulting wavelet coefficients will be very small (resulting from the approximation error). The smaller the wavelet support, the greater the precision in detecting discontinuities and the shorter the processing time. Moreover, the support of a wavelet is directly proportional to its number of vanishing moments. We will display results presenting the same total biomass signal analyzed with wavelets of the Gaussian family for different vanishing moments, from two to four. In addition, as in our previous work [3], we use the wavelet transform modulus maxima procedure introduced by Mallat and Hwang [2] that provides a precise location of the singularity and a clear cut measure of its Hölder exponent (HE) from the scaling of the wavelet transforms along the so-called maxima lines on which the transforms reaches local maxima with respect to the position coordinate. For more details, the reader is directed to the works [2] and [3].

4. A case study: two Haldane and one Monod species

Depending on the saturation and inhibition parameters and the types of fermentative growth, the crosses in the case of three growing species can occur at well-separated moments. For example, the total biomass signal may have two separated cones of influence in its wavelet transform. Such a case for two Haldane populations and one of Monod type is displayed in the figures of this work for different analyzing wavelets. Processing the same signal with several analyzing wavelets is recommendable when a better identification of the HEs is sought and helps to get rid of all sorts of artifacts. We used the second, third, and fourth derivatives of the Gaussian function as analyzing wavelets. It is well known that these Gaussian derivative wavelets have the same number of vanishing moments as the order of the derivative. In each case the range of scales with the same HE up to the first decimal digit in all three continuous wavelet transforms have been found. The same type of analysis, namely using wavelets of different vanishing moments to sense a given signal, have been used by Arneodo et al. [4] in a research of the fractal properties of DNA genome sequences. The argument for working in this way is that the common features showing up in all wavelet transforms belong with much higher confidence to the signal itself and it is not a numerical artifact caused by the integral transform. Following this idea, in the case of three microbial populations presented here, we can conclude that the HE is 1.6 for the first crossing singularity and 1.3 for the second one. This is so because these values are preserved in sufficiently long scale ranges although not for the entire scale interval that we considered.

The first crossing singularity

The results concerning the first encountered singularity are displayed in Figures 1, 2, and 3. From the plots (c) in each of the figures, we notice that we are able to get a stable first decimal digit of the HE if the low scales in the calculation of the wavelet transform are eliminated roughly proportional with the order of the Gaussian derivative employed.



Figure 1. (a) Total biomass signal (TBS) for three microbial species with different growth rates: two are of Haldane type with $K_{1_1} = 0.02$ g/l and $K_{2_1} = 0.04$ g/l, and $K_{1_2} = 0.03$ g/l and $K_{2_2} = 0.9$ g/l, respectively and the third one is of Monod type with $K_1 = 0.09$ g/l. (b) The singularity analysis has been performed with the second Gaussian derivative wavelet. (c) The HE α is shown in two different scale ranges.



Figure 2. The same case as in Figure 1 but with the singularity analysis performed with the third Gaussian derivative wavelet.



Figure 3. The same case as in previous figures but with the singularity analysis performed with the fourth Gaussian derivative wavelet.

•The second crossing singularity

The results concerning the second singularity are displayed in Figures 4, 5, and 6. For this singularity we could not assess a similar trend for the low

scales. However we identified the scale ranges on which we got a stable first decimal digit of the exponent for the three Gaussian derivatives, as one can see from the right plots (c).



Figure 4. Wavelet analysis performed for the second crossing singularity with the second derivative of the Gaussian as wavelet.



Figure 5. Wavelet analysis performed for the second crossing singularity with the third derivative of the Gaussian as wavelet.



Figure 6. Wavelet analysis performed for the second crossing singularity with the fourth derivative of the Gaussian as wavelet.

•The plateau singularity

The plateau singularity is less interesting because it indicates only the end of the fermentation process. The only useful fact is its detection and we noticed that it is indeed detected with any of the employed wavelets.

We collect the results of our illustrative example in Table 1. One can notice that we can determine R2 as the range of scales where the HE is the same up to the first decimal digit independently of the number of vanishing moments of the used wavelet. In the case of the first crossing, HE is constant over two log-scales (5-6), whereas for the second crossing HE stays constant over three log-scales (3-5). These can be considered as the scales over which the microbial growing process evolves towards two independent singularities. Moreover, for the first crossing singularity HE = 1.6, whereas for the second one HE = 1.3. According to the concept of Hölder regularity, the singularity of the second cross is stronger than the singularity of the first cross since it has a smaller HE. It is worth noting that we selected ranges R2 such that the HEs are obtained in the peaked region of the cones that do not overlap. In this way, we are more confident that the numerical values of the HEs do not express the combined effect of the two crossing singularities.

	Hölder Exponents				
Wavelet	First crossing		Second crossing		Plateau
	singularity		singularity		singularity
	R1f	R2f	R1s	R2s	Detected?
G2	1.5220	1.6028	1.1600	1.3200	Yes
G3	1.8100	1.6357	1.2440	1.3730	Yes
G4	2.2320	1.6533	1.2429	1.3514	Yes

Table 1. Hölder exponents obtained in two different ranges of scales R1 and R2 with the wavelet functions corresponding to the second (G2), third (G3) and fourth (G4) derivatives of the Gaussian function.

5. Conclusions

The presence of mixed culture growth in fermentation processes can be inferred starting from the data of total biomass processed with the wavelet transform acting as detector of the crossing growth singularities.

• The most relevant quantity provided by the wavelet analysis is the HE of the singularities that we determined here through the illustrative example of three microbial species contributing to the fermentation.

• In order to have a real technological application of the wavelet techniques in this area, it is important to perform calibrations of the mixedculture growth in terms of their HEs.

 Elimination of possible numerical artifacts is possible by employing wavelets of different vanishing moments for analyzing the same set of For this goal, we used the Gaussian data. derivative wavelets, which have the same number of vanishing moments as the order of the derivative. It is known from the wavelet theory that working with analyzing wavelets with p vanishing moments provide information on the regularity of the (p-1)th derivative of the analyzed signal. Since the majority of the numerical values of the HEs that we obtained are between one and two, it is clear that the fermentation signal under investigation has singularities in the first derivative. At the same time, this explains why we did not employ the first Gaussian derivative wavelet since it has less vanishing moments than the numerical value of the HEs.

• This work demonstrates that the usage of several wavelets with different vanishing moments together with well-selected scale ranges is helpful in filtering the true values of the HEs in the case of fermentation processes with isolated singularities. On the other hand, when one deals with signals with many close singularities the usage of the multifractal techniques must be considered [6].

• It would be very interesting to use other recent techniques to measure the fractality of the total biomass signals. We mention the DFA (Detrended Fluctuation Analysis) [5] and the MF-DFA (Multifractal-DFA) [6]. These methods are based on determining the scaling properties of the fluctuating moment of order q (q=2 holds for the case of DFA), as a function of the size of the samples and when the data contain noises and trends that are unknown from the point of view of their origin and their specific forms.

In addition, one can consider biological growths whose parameters depend on specific physical parameters, such as temperature, which is common for long-term bacterial activities [7]. Once the signal depends on the temperature, the HE depends as well. It would be highly interesting to examine the temperature dependence of the HE as implied by the temperature dependence of the total biomass signal.

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Authors' Biographies



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Haret C. Rosu was born in 1952 in Bucharest, Romania. He studied physics at the University of Bucharest and obtained the PhD degree in nuclear physics in 1987 during his period as researcher in the campus Magurele-Bucharest, where the main National Physics Laboratory of Romania is located. He staved there for the decade 1980-1990. After 3 years (1990-1993) as visiting professor at the Abdus Salam International Center for Theoretical Physics in Italy, he worked for 8 years as associate professor at the Institute of Physics of the Universidad de Guanajuato (IFUG), in Leon, Mexico. Since January 2002 he has been a full professor at the Instituto Potosino de Investigacion Cientifica y Tecnologica (IPICyT) in San Luis Potosi, Mexico. He was awarded the 2006 Potosi Prize for his academic achievements by IPICyT and he has been a SNI (National System of Researchers) member since 1994, being Level III since 2009. Dr. Rosu has published 125 research papers, most of them since he came to Mexico, and is known for his results in supersymmetric quantum mechanics as well as other areas of theoretical physics and applied sciences. He has also published 6 book chapters and has many contributed papers in proceedings of workshops. His graduate courses on quantum and classical mechanics are freely available on the Internet at the arXiv and other sites. He supervised four PhD students who now are on academic positions in Mexican universities.



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Vrani Ibarra-Junquera received a biochemical engineering degree in 2000, a master's degree in chemical engineering in 2003 and a Ph.D. degree in dynamical system and control in 2006 (at IPICyT). He is currently professor in the Faculty of Chemical Sciences at the Universidad de Colima in the area of Bioprocess Engineering. In the last years his research interest has focused on the monitoring and control of batch fermentations processes and application of bioreactors as alternative for fruit processing to converting it into value added products. Dr. Ibarra-Junquera is currently member of the National System of Researchers, level 1.