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► **To cite this version:**

Olivier Lafforgue, Isabelle Seyssiecq, Sébastien Poncet, Julien Favier. Rheological properties of synthetic mucus for airway clearance. *Journal of Biomedical Materials Research Part A*, 2018, 106 (2), pp.386 - 396. 10.1002/jbm.a.36251 . hal-01678912

HAL Id: hal-01678912

<https://hal.science/hal-01678912v1>

Submitted on 4 May 2018

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3 **Rheological properties of synthetic mucus for airway clearance**

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14 ABSTRACT: In this work, a complete rheological characterization of bronchial mucus
15 simulants based on the composition proposed by Zahm et al. [1] is presented. Dynamic Small
16 Amplitude Oscillatory Shear (SAOS) experiments, Steady State (SS) flow measurements and
17 three Intervals Thixotropy Tests (3ITT), are carried out to investigate the global rheological
18 complexities of simulants (viscoelasticity, viscoplasticity, shear - thinning and thixotropy) as
19 a function of scleroglucan concentrations (0.5 to 2wt%) and under temperatures of 20 and 37
20 °C. SAOS measurements show that the limit of the linear viscoelastic range as well as the
21 elasticity both increase with increasing scleroglucan concentrations. Depending on the
22 sollicitation frequency, the 0.5wt% gel response is either liquid-like or solid-like, whereas
23 more concentrated gels show a solid-like response over the whole frequency range. The
24 temperature dependence of gels response is negligible in the 20-37°C range. The Herschel-
25 Bulkley (HB) model is chosen to fit the SS flow curve of simulants. The evolution of HB
26 parameters versus polymer concentration show that both shear-thinning and viscoplasticity
27 increase with increasing concentrations. 3ITTs allow calculation of recovery thixotropic times
28 after shearings at 100s⁻¹ or 1.6s⁻¹. Empiric correlations are proposed to quantify the effect of
29 polymer concentration on rheological parameters of mucus simulants.

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KEYWORDS: synthetic bronchial mucus; viscoelasticity, viscoplasticity, shear - thinning,
thixotropy

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3 **INTRODUCTION**

4 A large number of fluids in human body, such as blood or mucus, produced by different
5 organs, are known to exhibit complex non Newtonian rheological properties under
6 physiological states. When transported in the airways as a result of cough or cilia beating,
7 bronchial mucus is characterized by a non constant, shear rate and time dependent viscosity,
8 in both normal and pathological conditions. Bronchial mucus is mainly composed of water
9 (90-95%), mucins (2-5%), lipids (1-2%), salts (1%), 0.02% of DNA and other molecules such
10 as cells debris [2]. Mucins are high molecular weight glycoproteins insuring a structural
11 protection function. The entangled and cross-linked network of its branched chains forms a
12 3D matrix spanning the mucus gel layer [3]. As a consequence of this complex internal
13 structure, bronchial mucus is a non Newtonian fluid displaying all the possible rheological
14 complexities such as viscoplasticity, shear-thinning, viscoelasticity and thixotropy. All these
15 properties directly affect the way mucus flows and, as a consequence, the vital clearance
16 function of the mucus layer coating the airways. The mechanism of mucus clearance can be
17 described by the following two steps:

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36 Step 1: inhaled particles or pathogens are trapped inside the mucus gel where enzymes and
37 antibodies can biochemically disrupt them.

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42 Step 2 : mucus is mainly transported by the mucociliary mechanism or by cough towards the
43 pharynx where it is either expectorated or digested.

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It is however known that, under certain disease conditions such as Cystic Fibrosis (CF), the
clearance function is affected by modifications of the mucus composition and consequently,
of its viscosity. Due to a lack of hydration (in the case of CF), mucus can indeed become very
thick and difficulties may arise to properly evacuate this fluid from the airways where it can
accumulate and become more easily infected. A good understanding of mucus rheology is

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3 thus of prime importance in order to develop new care solutions for patients suffering from
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5 CF or other chronic respiratory diseases. Among possible care solutions, clearance helping
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7 devices, based on different technologies, have been developed during the last decades. These
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9 small devices can be used by patients at home, on a daily basis to increase the volume of
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11 mucus expectoration and limit the need for respiratory physiotherapy. As an example, a newly
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13 developed apparatus known as the Simeox®, imposes an oscillatory air depression to the air
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15 flow during the exhalation phase of the patient. Based on the thixotropic and shear-thinning
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17 nature of mucus, such a solicitation induces a decrease of its viscosity and stimulates its
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19 expectoration. More insight into the rheology of respiratory mucus is needed to further
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21 improve the efficiency of such clearance helping devices. Although there is a large number of
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23 studies devoted to the rheological characterization of certain types of mucus, the results are
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25 still difficult to interpret, due to the use of different rheological techniques, but also due to the
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27 time evolution (aging) of such biological materials [4]. Furthermore, variations in the method
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29 used to collect samples (contamination issues), together with the natural variability
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31 (depending on the patient, the pathology, the occurrence of an infection...) of this complex
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33 biological fluid lead to important discrepancies in the existing literature, concerning the
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35 results on mucus rheological characterization [5]. Numerous previous works devoted to the
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37 study of mucus rheology (either synthetic or native mucus from different organs), have only
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39 described part of its rheological properties. For instance, many works have used dynamic
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41 oscillatory shear measurements to characterize mucus rheology [6 ; 7 ; 8 ; 9; 10 ; 11 ; 12].
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43 Under small deformations (SAOS) these measurements mostly reflect the properties of mucus
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45 under its native, unperturbed state and are useful to describe the linear viscoelastic response of
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47 mucus. On the contrary, in other works, the authors have made the choice to characterize
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49 mucus rheology using only continuous shear experiments [1 ; 13 ; 14; 15]. In these cases, the
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51 measured properties mostly reflect the flow behavior of mucus and can be used to investigate
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3 its viscosity under physiological shearing rates prevailing in human lungs during normal
4 functioning or during temporary events such as cough. Even in studies where both dynamic
5 oscillatory and shear flow measurements were carried out, the thixotropic nature of mucus
6 was not accounted for [5; 16 ; 17 ; 18 ; 19; 20 ; 21], or at least not on a quantitative point of
7 view [22 ; 23 ; 24]. As a consequence, a complete and intrinsically consistent characterization
8 campaign is still missing in the open literature. As the use of real mucus implies strong issues
9 related to available quantities, and rises questions about the impact of the collection method
10 on the fluid composition, the choice made in this work is to use mucus simulants. In this
11 context, this work proposes a rheological characterization of mucus simulants at different
12 active polymer concentrations (0.5 to 2%), under a temperature of 20 or 37°C allowing to
13 cover the range of air physiological temperature along the airways and using a broad range of
14 available rheological tests (SAOS, controlled shear stress SS flow tests and 3ITT). In an
15 attempt to quantify the measured properties, empirical equations are used to represent the
16 evolution of different rheological parameters as a function of the active polymer
17 concentration.

38 **MATERIALS AND METHODS**

39 **Preparation of mucus simulants**

40 The composition and preparation of polymeric synthetic solutions used to mimic human
41 bronchial mucus are described in Zahm *et al.* [1]. To account for the natural variability of real
42 mucus from one patient to another but also depending on health conditions, gels with different
43 scleroglucan (Actigum™) concentrations ranging from 0.5 to 2wt% were prepared. Mucus
44 simulants are aqueous solutions mainly composed of two types of polymers, Viscogum™ FA
45 (Cargill™) and Actigum™ CS 6 (Cargill™). The polymers used in this work were kindly
46 provided by the Laserson compagny (Etampes, France). Viscogum™ FA is a galactomannan

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3 gum extracted from locust beans and Actigum™ CS 6 is a scleroglucan (branched
4 homopolysaccharide). It consists in a glucose chain branched every three units by an
5 additional glucose forming a three dimensional (triple helix) structure. Sodium chloride
6 (99.8+% NaCl) and di-sodium tetraborate 10aq (99.5+% Na₂B₄O₇·10H₂O) were purchased
7 from Chem-Lab NV (Zedelgem, Belgium). Distilled water used in all preparations was
8 produced using a 2012 distillator (GFL, France). Mucus simulants solutions were prepared
9 within glass bottles filled with 200 mg of distilled water. Then, 0.9wt% of NaCl, 0.5wt% of
10 Viscogum™ FA and a chosen fraction (ranging from 0.5wt% to 2wt%) of Actigum™ CS 6
11 were successively added into the solution under magnetic stirring (Ikamag® RET) at room
12 temperature. The mixture was kept under agitation for 48h at room temperature. After this
13 time period, a mass corresponding to 4mL of di-sodium tetraborate at 0.02M was added. This
14 addition induces the cross-linking of the polymeric chains, building a 3D gel matrix that
15 mimicks the mucin network responsible for the internal structure of real mucus. The agitation
16 is kept for a few more hours before storing the final mixture at 4°C. Before performing the
17 measurements, the solution is fractionnated into several 30mL plastic vials and then allowed
18 to recover at room temperature. Such mucus simulants were found to mimic accurately the
19 main properties of bronchial mucus in the case of different pathologies.

20 21 22 **Rheological measurements**

23 24 *Rheometers and measuring systems*

25 Rheological measurements were performed using two controlled-stress rheometers, the AR
26 550 and the DHR-2 (TA Instruments) equipped with a measuring system consisting of a 2°
27 stainless steel cone (40 mm or 50 mm in diameter). The temperature was controlled by a
28 Peltier plate. A wet steel lid or a thin silicone oil layer insuring a water saturated atmosphere
29 around the sample was used as a dehydration preventing solution.

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3 ***Sample loading***
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5 A small amount of gel was loaded onto the Peltier plate by gently pouring it from the vial in
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7 order to minimize shear history effects. The geometry was then lowered down to the
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9 corresponding gap plus a few micrometers and the excess of fluid was removed on the edges,
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11 the exact gap value was then set. The desired temperature (20 °C or 37°C) was also set before
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13 the tests began.
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17 ***Measurements protocols***
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19 Viscoelastic properties of the simulants were investigated through a series of dynamic shear
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21 experiments (Small Amplitude Oscillatory Shear: SAOS). The results were interpreted based
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23 on the evolution of the elastic and viscous moduli (G' , G'') and the loss angle (δ) as a function
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25 of the sinusoidal input. The stress dependency of the different gels (0.5wt% to 2wt% in active
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27 polymer) response was first measured via stress amplitude sweeps at constant frequency
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29 ($\frac{1}{2\pi}$ Hz). This is a classical test carried out in order to determine, for each solution, the limit
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31 of the Linear ViscoElastic (LVE) range. The frequency dependency (in a maximum range of
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33 10^{-5} to 100 Hz) of the different gels response was also measured at constant stress amplitude,
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35 within the LVE range according to the stress amplitude sweep results. The temperature
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37 dependency of simulants rheological properties was finally investigated by comparing stress
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39 amplitude sweeps obtained for a given sample at either 20°C (ambient air temperature) or
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41 37°C (physiological temperature in the lower airways). However, to fully characterize the
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43 behavior of a mucus layer in response to in vivo solicitations such as cough or air flows
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45 artificially induced by clearance helping devices, the rheological measurements have to be
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47 performed far beyond the LVE range. Rotational controlled shear stress flow tests were then
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49 used to determine the rheological properties of the mucus simulants under shear flow
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51 conditions. In order to quantify the viscoplastic and the shear-thinning effects independently
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3 of the thixotropic ones, steady state rheograms were recorded. Such a steady state curve is
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5 obtained by applying a given shear stress until the corresponding shear rate reaches a constant
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7 value. Steady state (SS) flow curves of the different simulants were modeled using a 3
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9 parameters Herschel-Bulkley (HB) model. This model accounts for the fluid viscoplasticity
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11 via the yield stress value (τ_{0HB}) and for its shear - thinning behavior via the flow (n) and
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13 consistency (K) indexes values. The thixotropy of the more concentrated mucus simulant was
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15 separately quantified using three Intervals Thixotropy Tests (3ITT). A 3ITT test consists in a
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17 stepwise change of stress or strain rate to successively monitor the initial structure, then its
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19 breaking up and finally its recovery. More precisely, the 3ITTs applied here can be described
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21 as follows:
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26 First interval: the sample is submitted to very low shear conditions ($\dot{\gamma} = 0.029s^{-1}$). This
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28 interval gives a reference for the fluid structure "at rest" or at least under very low shear.
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32 Second interval: higher shear conditions are imposed by applying a constant shear stress or
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34 shear rate to disrupt the internal structure until a steady state is reached (depending on the
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36 chosen stress or strain rate value). In our case, a constant shear rate value of either $1.6s^{-1}$ or
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38 $100s^{-1}$ was applied during step 2, in order to submit the sample to shearing conditions
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40 representative of either normal shearing conditions in the airways, or during peculiar events
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42 such as cough.
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46 Third interval: the sample is allowed to recover under very low shear conditions again
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48 ($\dot{\gamma} = 0.029s^{-1}$). A time scale characterizing a given state of regeneration (chosen here at 90%
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50 and 100% of recovery based on the initial structure measured during step 1.) can then be
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52 calculated.
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RESULTS

SAOS tests

Stress amplitude and frequency sweep tests conducted at a constant temperature of 20°C, on simulant gels at different Actigum™ concentrations, all display the same feature. An example is given in figure 1 for a 1.5wt% Actigum™ gel (figure 1 (a) stress amplitude sweep, (b) frequency sweep). A value of the elastic modulus G' above the viscous modulus G'' implies that the elastic behavior dominates the viscous one and indicates a solid-like or gel-like behavior, while $G'' > G'$ indicates a liquid-like behavior. In figure 1 (a), the stress amplitude value for which the transition from one behavior to the other is observed, is denoted by τ_f (flow point value corresponding to the crossover of moduli $G' = G''$, $\tan(\delta) = 1$) and will be discussed hereafter. As a consequence, in the case of figure 1.(b) for which $\tau = 1 \text{ Pa} < \tau_f$, the mucus simulant shows a gel-like behavior for all frequencies ranging from 10^{-3} to 10 Hz. In the case of figure 1.(a), under a constant frequency of $\frac{1}{2\pi} \text{ Hz}$, the two domains are successively observed, with a solid-like behavior for $\tau < \tau_f$, then liquid-like behavior for $\tau > \tau_f$.

Stress amplitude sweep tests

The stress amplitude dependency of the different gels response was recorded via stress amplitude sweeps at a constant frequency ($\frac{1}{2\pi} \text{ Hz}$) for all the concentrations in active polymer (0.5wt% to 2wt%). The results obtained are displayed in figure 2.

For all polymer concentrations, a plateau region for G' and G'' moduli defines the LVE Range at the preset frequency. A yield stress value (τ_y) limits the LVE range and is determined from the end of the moduli plateau. Since the G' curve often deviates first from the plateau, the G'

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3 function is commonly chosen to determine τ_y . Here, a 10% deviation from the plateau has
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5 arbitrarily been set for τ_y calculations . The yield stress is the stress limit below which no
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7 significant change of the internal structure occurs. For $\tau < \tau_y$ the sample is displaying
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9 reversible viscoelastic behavior. On the contrary, above τ_y , the measurements no longer
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11 reflect the structure at rest due to early signs of stress induced microstructural evolutions.
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13 Based on the value of the yield stress and of the corresponding critical deformation γ_c , one can
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15 also calculate the volumetric energy of cohesion of the 3D network E_c in $J.m^{-3}$ (eq. (1)) [25].
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17 This energy of cohesion can be used in a quantitative manner as a measure of the extent of
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19 intermolecular and intramolecular interactions of the polymeric internal structure [26].
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$$E_c = \frac{1}{2} \tau_y \cdot \gamma_c \quad (1)$$

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28 In the case of the mucus simulants tested here, it can be seen in figure 2 that above the critical
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30 yield stress, G' decreases while G'' shows an overshoot before decreasing. This kind of
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32 behavior has been observed on lots of gels by Mezger [27] and is classified as a "Type III
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34 behavior or weak strain overshoot" by Hyun et al [28]. The flow point (τ_f) is identified as the
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36 stress value for which a moduli crossover ($G'=G''$, $\tan(\delta)=1$) is observed. This flow point
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38 corresponds to the stress above which the material becomes more viscous than elastic due to
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40 critical microstructural breakdowns. Finally the stress corresponding to the maximum of G''
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42 (peak overshoot) is denote by τ_{peak} and is, for all concentrations, almost superimposed to the
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44 flow point stress (except for the 0.5wt% gel for which the G'' overshoot does not appear). The
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46 range between τ_y and τ_f is sometimes referred as the "yield zone" or the "yield / flow
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48 transition range" [27]. In this range, despite the predominance of elastic behavior ($G' > G''$),
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50 irreversible deformations might locally have already taken place.
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3 In the case of mucus simulants studied here, G' and G'' moduli plateau values, as well as
4 yield, flow and peak stresses and volumetric cohesive energy, all show increasing values
5 according to the Actigum™ concentration as reported in table 1. The evolution with the
6 Actigum™ concentration of the different transition stresses (τ_y , τ_f and τ_{peak}) as well as the
7 volumetric energy of cohesion (E_c) and the plateau moduli values are all displayed in figure 3.
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14 For quantification purposes, yield, flow and peak stresses variations with Actigum™
15 concentration can be described by power law empirical relationships reported hereafter
16 (equations 2 & 3). As previously mentioned, flow and peak stresses are almost superimposed,
17 which makes sense since they both refer to the actual beginning of macroscopic flow of the
18 gel ($G'' > G'$ and friction starts decreasing). As a consequence a single empirical correlation is
19 needed to account for the effect of polymer concentration on both flow and peak stresses.
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28 Equation 2 gives the quantitative evolution of the yield stress (τ_y in Pa) as a function of
29 Actigum™ concentration (C in wt%):
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$$\tau_y = 1.54 C^{2.31} \quad R^2 = 0.981 \quad (2)$$

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37 Equation 3 accounts for the Actigum™ dependency of both τ_f and τ_{peak} :
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$$\tau_f \text{ \& } \tau_{peak} = 4.66 C^{1.68} \quad R^2 = 0.990 \quad (3)$$

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44 These evolutions are qualitatively compared to the evolution of the yield stress calculated
45 from the HB modelling of the SS flow curve (τ_{0HB}) (see discussion part). Concerning G' and
46 G'' plateau values (in Pa), they increase linearly (Figure 3(b)) with the active polymer
47 concentration (in wt%) with an average slope of 10 for G'' and 49 for G'. The evolution of the
48 volumetric energy of cohesion (E_c in $J.m^{-3}$) of the 3D gel internal network versus the
49 Actigum™ concentration (C in wt%) is also a power law as reported in equation 4:
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$$E_c = 0.076 C^{1.95} \quad R^2 = 0.968 \quad (4)$$

Temperature dependence of the LVE range

The temperature dependence of the LVE range of mucus simulants was measured by comparing stress amplitude sweeps obtained at either 20°C (ambient temperature) or 37°C (physiological temperature in the lower airways). This temperature range was chosen in accordance to the possible range of variation for the air at different stages of respiratory airways. Figure 4 presents the variations of elastic modulus and viscous modulus versus the stress amplitude for a constant frequency ($\frac{1}{2\pi}$ Hz) and a given Actigum™ concentration (0.75 wt%) at either 20°C or 37°C. Each curve is the average of 3 successive measurements. Measurements were also performed with simulants at different polymer concentrations and all display qualitatively identical results.

As it is observed in figure 4, the LVE range of the mucus simulant gel as well as the different stresses (τ_y , τ_f , τ_{peak}) characterizing the solid - liquid transition zone show very little dependence on the temperature in the 20-37°C range. The curves obtained at either 20 or 37°C are almost superimposed and the observed differences are of the same order of magnitude than reproducibility. Concerning the gels studied here, it can be concluded that the effect of temperature in the 20-37°C range on mucus simulants rheological properties is not significant compared to reproducibility.

Frequency sweep tests

Since the frequency is the inverse of a time, frequency sweep tests are usually performed in order to investigate the behavior of a substance as a function of the sollicitation characteristic time under small deformations. Short-term behavior of the sample is then simulated at high frequencies, whereas long-term behavior is displayed under low frequencies. The main

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3 concern under very low frequencies is the conservation of the sample during the important
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5 timelag needed to obtain these measuring points. So in these cases, a special care has to be
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7 taken to control samples dehydration. The frequency dependence of the mucus simulants
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9 response was measured by oscillating stress at increasing frequencies (maximum range: 10^{-5} to
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11 100 Hz) and a constant stress amplitude corresponding to a small deformation, i.e. within the
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13 LVE range ($\tau < \tau_y$) as measured at $\frac{1}{2\pi}$ Hz (see table 1.). Figure 5 gathers the evolutions of
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15 elastic modulus (a), viscous modulus (b) and loss angle (c) as a function of the applied
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17 frequency for the different Actigum™ concentrations.

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23 Since working at very low frequencies implies a long experiment duration and, as a
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25 consequence, a more important risk of sample dehydration, all the simulants have been
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27 studied in the common range 10^{-2} to 10 Hz. only in the case of the more diluted and the more
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29 concentrated gels, the frequency range was enlarged to 10^{-4} - 10 Hz and 10^{-5} - 10 Hz to
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31 investigate the possible occurrence of a low frequency moduli crossover. Over the whole range
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33 of frequencies, simulants with Actigum™ concentrations ranging from 0.75wt% to 2wt%,
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35 show a gel-like behavior characterized by an almost parallel low increase of G' and G'' (in
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37 log-log scale) with G' values 2 to 7 times higher than G'' . Such a behavior is characteristic of a
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39 soft gel in which intermolecular forces are mostly responsible for the 3D internal network
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41 [27]. Therefore elastic behavior dominates the viscous one over the entire frequency range
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43 attesting for the gel stability. G' and G'' only show a slight frequency dependence displaying
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45 an average slope in log-log representation of 0.1 in the case of G' and 0.04 in the case of G''
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47 (see discussion part for interpretation).
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52 The less concentrated simulant (0.5 wt%) displays a different behavior. G' and G'' show a
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54 more important frequency dependence, especially under low frequencies due to a more
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3 flexible network. A moduli crossover (characterized by $\tan(\delta) = 1$ in figure 5 (c)) is also
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5 observed at low frequencies ($< 2 \cdot 10^{-3}$ Hz) for which $G'' > G'$. Such a behavior corresponds to
6
7 a liquid-like behavior and can be observed when a deformation (even very small) is applied at
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9 a sufficiently low rate (i.e. during a long time). On the contrary, an identical deformation
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11 applied at higher frequencies induces a solid-like response. The typical example usually given
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13 in rheology books to illustrate this behavior is the one of a silicone ball. Such a ball left at rest
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15 in a beaker for a long time will finally take the form of the beaker (viscous flow under low
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17 frequency or long time sollicitation), while the same ball thrown onto a wall will bounce,
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19 displaying an elastic behavior under high frequency or small time sollicitation. It is worth
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21 noting that the inverse of the frequency for the moduli crossover is sometimes used as a time
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23 characterizing the elasticity of the gel [29]. However, in this case, it would be delicate to
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25 interpret this value since complex internal structures composed of a large number of
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27 components with different lengths, flexibilities or mobilities, such as mucus simulants, are
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29 most likely to display viscoelastic properties governed by the superposition of several
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31 relaxation modes [2]. The low frequency crossover of moduli is not experimentally reached in
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33 the case of more concentrated simulants (0.75wt% to 2wt%), for the range of tested
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35 frequencies. However, in figure 5 (c), $\tan(\delta)$ increases when the frequency decreases, so that a
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37 moduli crossover would likely appear under sufficiently low frequency at a decreasing
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39 frequency value, as the concentration increases, traducing an increased elasticity for more
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41 concentrated mucus simulants.
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48 **Steady-state (SS) flow test**

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50 Steady state (SS) shear flow tests were performed on three successive samples at each
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52 Actigum™ concentration. The response to the imposed stress (τ) is recorded in terms of
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54 steady state strain rate ($\dot{\gamma}$). The rheograms ($\tau(\dot{\gamma})$ curves) drawn as the average of the three
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3 measurements performed on each Actigum™ concentration displayed a similar shape for all
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5 polymer concentrations. Figure 6 shows an example of such a rheogram obtained in the case
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7 of a 2 wt% simulant.
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11 A yield zone with a solid-like behavior ($\dot{\gamma} \rightarrow 0$ grey zone in figure 6) is first observed until
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13 the progressive departure from a quasi vertical slope to the flow zone showing a shear-
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15 thinning behavior (concave rheogram beyond the yield stress value) is reached. The yield
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17 zone is the manifestation of the yielding behavior of the gel structure that has already been
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19 characterized using SAOS stress amplitude sweeps tests. In SS flow mode, the transition
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21 between solid and liquid-like behaviors is defined by a stress value quoted τ_{0HB} . This value is
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23 also referred in the literature as a yield stress but gives different values compared to the yield
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25 stress measured in SAOS mode (see discussion part). This is why a different notation (τ_{0HB}) is
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27 used to described this yield stress deduced from the fitting of SS flow experimental data by a
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29 3 parameters Herschel-Bulkley model (equation 5) allowing to account for both the SS yield
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31 stress (viscoplastic behavior) and shear-thinning behavior.
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$$\tau = \tau_{0HB} + K\dot{\gamma}^n \quad (5)$$

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40 where τ_{0HB} (Pa) is the SS flow yield stress, K (Pa.sⁿ) and n (-) are the consistency and flow
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42 indexes respectively deduced from the HB modelling. It is worth noting that, as SS flow
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44 curves (as well as the HB model) do not account for the time dependent flow behavior
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46 (thixotropy), the observed yield stress τ_{0HB} is not an inherent property of the material since the
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48 structural strength of the gel is shear history and thus time dependent.
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52 The non Newtonian behavior observed in figure 6, is the direct consequence, in the case of
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54 polymeric substances, of a shear-induced spatial structure transformation [30]. The HB model
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56 parameters obtained for the different polymer concentrations are reported in table 2 and
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3 account for the characterization of the steady-state flow behavior of simulants. The evolution
4 of the different HB parameters with the Actigum™ concentration is drawn in figure 7 and
5 clearly shows that both viscoplasticity (increase in τ_{0HB} value) and shear-thinning behavior
6 (increase in K, decrease in n) increase as the active polymer concentration increases. In the
7 case of the mucus simulants studied here, an attempt has been made to propose a
8 quantification of the different measured rheological properties. Concerning data deduced from
9 SS flow experiments, the evolution of HB parameters with the active polymer concentration
10 (C in wt%) can be quantified by the following empirical relationships (equations 6 to 8).
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$$21 \quad \tau_{0HB} = 8.24 C^{1.34} \quad R^2 = 0.99 \quad (6)$$

$$22 \quad K = 0.315 C^{3.36} \quad R^2 = 0.974 \quad (7)$$

$$23 \quad n = 1 - 0.346 C \quad R^2 = 0.963 \quad (8)$$

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Equation 6 accounts for the increase in viscoplasticity, whereas equations 7 and 8 account for
the increase in shear-thinning behavior, as the active polymer concentration increases.

Three Interval Thixotropy Tests

3ITT tests were performed on the different simulant gels in order to investigate the thixotropic
nature of mucus simulants independently of their SS flow behavior. Thixotropy is
characteristic of complex internal structure systems and is linked to slow time evolutions in
rheological properties due to either restructuring at rest or destructuring initiated by
deformation [30]. Figure 8 represents a first 3ITT performed on a 2 wt% simulant. The second
step imposes a shear flow under an effective shear rate of 1.6 s^{-1} corresponding to the order of
magnitude of the shearing of a mucus layer in the tracheobronchial tree. Indeed, if the
physiological maximum value for shear rate, can reached $10^2\text{-}10^4 \text{ s}^{-1}$ (for example during

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3 coughing), physiological rates in the normal lung are of the order of magnitude of 0.1 to 1 s^{-1} .
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5 [4 ; 8 ; 14]. The three successive steps are plotted as apparent viscosity versus time.
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8 After a shearing under 1.6 s^{-1} during step 2, one can observe in figure 8 that the thixotropic
9 regeneration curve (step 3) allows the calculation of a thixotropic recovery time
10 characterizing a given state of regeneration chosen here at 90% and 100% of recovery, based
11 on the initial structure measured during step 1. The thixotropic recovery time is, in this case of
12 2.7s for 90% and 75s for 100% of recovery.
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20 Figure 9 gives the results of a second test also performed on a 2 wt% simulant. This time, the
21 second step corresponds to a shear flow under a larger shear rate (100 s^{-1}) matching the order
22 of magnitude for the shearing of a mucus layer submitted to cough.
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27 After a shearing at 100 s^{-1} during step 2, one can observe in figure 9 that the thixotropic
28 regeneration curve (step 3) gives a 90% recovery time of 17s and a 100% recovery time of
29 917s, which are logically higher than the recovery times needed after the shearing under 1.6 s^{-1}
30 observed in figure 8.
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38 **DISCUSSION**

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40 During stress amplitude SAOS experiments (figure 2) a G'' overshoot has been observed with
41 mucus simulants. According to Hyun et al [28-31] or Mezger [27], such an overshoot is
42 generally obtained in the case of cross-linked polymers or gel-like internal structures
43 (existence of a 3D network). The overshoot is directly linked to the progressive collapse of
44 the 3D network. During the initiation of flow, at first the friction increases due to spatial
45 rearrangements of some free elements (for example relative motion of end pieces of chains).
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47 These irreversible motions induce an increase in dissipated energy (G'' increases). Then the
48 breakdown of the internal superstructure occurs (at the τ_{peak} value) and the dissipated energy
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3 starts decreasing as the fluid actually flows. In the case of mucus simulants used in this work,
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5 the overshoot behavior shows a strong dependency on the Actigum™ concentration. On the
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7 one hand, the τ_{peak} value increases as the simulant concentration increases (figure 3(a) and eq.
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9 3 for quantification). This is not surprising since more concentrated simulants have a stronger
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11 internal network that will need a higher energy input to collapse. On the other hand, the peak
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13 height is also increasing with the Actigum™ concentration (figure 2(b)). The overshoot
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15 amplitude is directly linked to the amount of friction occurring during the initiation of flow due
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17 to spatial rearrangements of free elements. A more concentrated sample is thus expected to
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19 dissipate more friction energy as a result of its higher network density (more cross-linking and
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21 chains entanglements). A G'' overshoot during stress amplitude sweeps was rarely mentioned
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23 in the literature on the rheological characterization of mucus. In the work of Bastholm [9], a
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25 G'' overshoot seems to occur with some of the tested cervical mucus samples, without any
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27 comment concerning this phenomenon. Celli *et al.* [5], performing a SAOS stress amplitude
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29 sweep on a sane gastric mucus sample (compared to a *Helicobacter pylori* contaminated one),
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31 also observed a "weak strain overshoot" on G'' and referred to the work of Huyn *et al.* [28] for
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33 interpretation. Concerning now the internal network volumetric energy of cohesion E_c values,
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35 we can refer to works by Mori *et al.* [32] or Niraula *et al.* [26] for comparison purposes. Based
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37 on SAOS characterizations on respectively biological suspensions or stabilized oil in water
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39 (O/W) emulsions both exhibiting a 3D cohesive network, they calculated the volumetric
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41 cohesive energy of the material internal network to account for the degree of interaction and
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43 thus of flocculation between droplets in the case of stabilized O/W emulsions [26] and
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45 between bioflocs in the case of activated sludge suspensions [32]. The mucus simulants
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47 studied here are not flocculated suspensions nor liquid-liquid emulsions but aqueous
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49 dispersions of macromolecules. However since a 3D network exists both due to entanglement
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51 and cross-linking of the macromolecules the concept of volumetric cohesive energy can be
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3 useful to quantitatively account for the degree of interaction of the subsequent network. One
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5 can observe that the order of magnitude of E_c for the ActigumTM based mucus simulants (0.02
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7 to 0.36 J.m^{-3}) is intermediate between E_c values calculated for biological activated sludge
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9 suspensions [32] (0.2 to 1.2 J.m^{-3}) and E_c values of stabilized O/W emulsions ($0.24 \cdot 10^{-3}$ to
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11 $2.95 \cdot 10^{-3} \text{ J.m}^{-3}$). It is also worth noting that E_c values showed important dependence upon the
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13 network macromolecular or suspended bioflocs concentration (power law (eq. 4) in this work,
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15 exponential law in [32]). The influence of the polymer concentration on stress amplitude
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17 SAOS measurements displayed in figure 3 for synthetic mucus, has been discussed in some
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19 previous works. As an example, Riley et al. [7] showed that an increase in polymer (Carbopol
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21 934P) concentration induced an increase in both G' and G'' measured in the LVE range at a
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23 constant (5 Hz) frequency. Shah et al. [19] also observed working with mucus simulants that
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25 both G' and G'' (either measured at 1 or 100 rad/s) increased with the coagulant concentration
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27 (0.5, 1.5 and 3%). Finally, Hamel & Fiegl [10] measured an identical increase with cross-
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29 linked polymer concentration of G' and G'' moduli, performing on synthetic mucus strain
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31 sweeps at a constant frequency. Nevertheless, in all of these previous studies, the observed
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33 variations were not quantified. An interesting point that can however be pointed out is the fact
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35 that the achieved order of magnitude of G' and G'' (in the LVE range) with the mucus
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37 simulants tested in this work correspond to those usually observed with sane or pathologic
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39 native respiratory mucus. For instance, CF mucus can reach various viscoelasticity levels as a
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41 function of many factors. In the work of Dawson et al. [18] CF mucus displays viscoelastic
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43 properties, in terms of moduli plateau values, close to those displayed by the 2wt%
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45 ActigumTM gel while in the research of Yuan et al. [11], the measured behavior is close to the
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47 one measured with the 0.75wt% gel. Concerning sane mucus, moduli plateau values close to
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49 those recorded with the 0.5wt% solution have been observed [11]. Concerning the effect of
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51 temperature on the LVE range limit, identical results to those shown in figure 4 have been
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3 described for mucus gels with similar composition, in the same temperature range (20, 32 or
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5 36°C) [33]. A very limited influence of temperature on simulants rheological properties
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7 measured both via stress amplitude sweeps and flow curves was recorded by the authors.
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9 Taylor et al. [6], who studied the rheological characterization of mucin - alginate gels, also
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11 reported no significant differences between the gels behaviors (frequency sweep tests) over a
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13 10 to 60°C temperature range. The rheological properties of such aqueous polyoside solutions
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15 are known to be related to the conformation taken by the macromolecules (ViscogumTM &
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17 ActigumTM). Mucus simulants studied here are composed of a constant amount of
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19 ViscogumTM (galactomannan chains that are cross-linked in the presence of sodium
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21 tetraborate) and various proportions of ActigumTM (extracellular polyoside) that can adopt
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23 either a rigid helicoidal conformation or a random entangled conformation, as a function
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25 notably of temperature [29]. From the measurements displayed in figure 4, it can be
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27 concluded that ActigumTM molecules are likely to present the rigid helicoidal conformation in
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29 simulants tested here, leading to the observed gel-like behavior ($G' > G''$) and also that this
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31 conformation remains when the temperature is raised to values such as 37°C.
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37 During frequency SAOS sweeps, we obtained results (figure 5) that are, for 0.75 to 2wt%
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39 gels, qualitatively comparable to those observed by Coussot et Grossiord [29] on a 0.5wt%
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41 xanthan gum aqueous solution at 25°C. Xanthan gum is an extracellular polyoside close to
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43 ActigumTM in terms of macromolecular composition and structure (glucose chain branched
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45 every 2 units by other oside functions forming a 3D rigid double helix structure), exhibiting
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47 very similar rheological properties. The gel-like behavior of aqueous xanthan solutions over
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49 the whole frequency range is known to be linked to the helicoidal rigid conformation adopted
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51 by polyoside molecules under moderate temperatures or high ionic strength, impairing their
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53 ability to move under small sollicitations. In the case of ActigumTM aqueous solutions,
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3 Actigum™ macromolecules are also known to adopt a rigid triple helix conformation (except
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5 under high temperatures or low ionic strength) that is responsible for the gel-like response
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7 measured over the entire range of tested frequencies in the case of 0.75wt% to 2wt%
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9 Actigum™ gels. Kocevar-Nared et al. [23] also made observations qualitatively comparable
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11 to figure 5 in the case of rehydrated dried crude porcine gastric mucus. Varying the mucin
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13 concentration of rehydrated mucus, they observed that more concentrated samples displayed
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15 no moduli crossover (in the 0.1 - 100 Hz explored range) but a power law slight frequency
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17 dependence for both moduli whereas, more diluted sample displayed a stronger frequency
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19 dependence for both moduli whereas, more diluted sample displayed a stronger frequency
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21 dependence with a crossover point moving towards lower frequencies as the mucin
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23 concentration is increased and as the structure changes to gel-like. In the case of the 0.5wt%
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25 gel studied here, the stronger dependence to the frequency can be linked to the increase in
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27 relative importance of Viscogum™ (flexible entangled network) compared to Actigum™
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29 (rigid network) in comparison to gels with higher Actigum™ concentrations.
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32 During SS flow measurements, the concentration dependent yield stress and shear-thinning
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34 nature of mucus simulants has been observed (figure 6) and quantify (equations 6 to 8).
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36 Identical observations have already been reported but only qualitatively in the case of mucus
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38 simulants and native mucus based on measurements of their rheological flow behavior. The
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40 increase in viscoplasticity (yield stress) with the active polymer concentration has already
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42 been observed in the SAOS stress amplitude tests section of the paper. Yield stresses
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44 measured in flow mode τ_{0HB} show higher values than yield stresses measured in SAOS τ_y .
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46 This is not surprising since τ_y corresponds to the end of the LVE range characterized by early
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48 motions of small elements of the complex internal polymeric structure (at the microscopic
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50 level), while τ_{0HB} corresponds to the proper beginning of flow measured at a macroscopic
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52 scale (existence of a finite shear rate in the measuring gap). It is also worth noting that both τ_y
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3 and τ_{0HB} values are dependent on the calculation method (chosen % of deviation from the
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6 plateau for τ_y and chosen limit value for $\dot{\gamma} \rightarrow 0$ in the case of τ_{0HB}). As a consequence, both
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8 values should only be considered as order of magnitudes or for comparison purposes of data
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10 obtained using the same calculation method. As far as the evolution of yield stresses with the
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12 active concentration is concerned, Malkin et al. [34] reported the case of cysteine / Ag based
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14 colloidal gels for which the yield stress deduced from flow measurements decreases as the
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16 dilution factor of the initial gel increases, showing the decrease in strength of the rigid
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18 structure as the colloidal concentration decreases. If we now focus on the observed shear-
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20 thinning properties of mucus simulants, Puchelle et al. [13] also described the shear-thinning
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22 nature of native and lyophilized pathological bronchial mucus but also of simulants composed
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24 of polyisobutylene solutions (3 and 6% in decalin). Measuring the SS flow curve of these
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26 gels, they obtained shear-thinning indexes (Ostwald de Waele or power law) matching the
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28 range observed in table 2 with our mucus simulants and varying from 0.44 to 0.64. Banerjee
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30 et al. [14] also used a power law to describe the apparent viscosity of shear-thinning mucus
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32 simulants (tragacanth gum) mixed with different surfactants to evaluate their ability to reduce
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34 viscosity. They obtained viscosity reduction ratios varying from 1.5 to 7.2 depending on the
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36 type of surfactant under physiological shear rates (0.1 to 1 s⁻¹ in the tracheobronchial tree).
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38 The shear-thinning behavior of mammalian lung mucus is also cited by Vasquez et al. [16]
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40 but not quantified. It is also the case of Boegh et al. [21] who tried to propose a bio-similar
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42 mucus composition displaying rheological properties (notably in terms of shear-thinning)
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44 close to the composition of a mucus isolated from cultured cells. Finally, Tomaiuolo et al.
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46 [24] also described the shear-thinning nature of different native CF mucus samples without
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48 proposing any quantification of this property. More generally, the increase in shear-thinning
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50 behavior with the polymer concentration, is classical for aqueous polymer solutions. It
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3 corresponds to an increase in shear-thinning capacity for solutions with a higher
4 macromolecular content due to a concentration enhanced reduction of the flow resistance
5 following the alignment and orientation of polymeric chains in the flow direction as the shear
6 rate increases [17 ; 23 ; 34]. Shah et al. [19] compared viscosity curves of respiratory mucus
7 simulants (0.5 to 3% in polyox resin coagulant). However, they only discussed their results in
8 terms of viscosity increase with the coagulant concentration and did not account for the shear-
9 thinning nature of the studied fluids, nor for its evolution with the polymer concentration.
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Finally, 3ITTs performed on a 2wt% simulant gel (figures 8 & 9) allow measuring thixotropic recovery times. If the thixotropic nature of native or synthetic mucus has been reported in some of the previous works devoted to mucus rheological characterization [1 ; 22 ; 23 ; 24], it has often not been quantified. Nielsen et al. [22] but also Tomaiuolo et al. [24], described qualitatively the thixotropic nature of native samples observing an hysteresis loop between viscosity curves successively measured at increasing or decreasing shear rates. Kocevar et al. [23] recorded with porcine gastric mucins solutions at concentrations varying from 10 to 60% the apparent viscosity versus time under a constant shear rate of 50 s^{-1} . They concluded depending on the appearance of a viscosity decrease versus time, that gels above 30% are thixotropic while lower concentrated one showed no thixotropy. Zahm et al. [1] performed identical measurements on mucus simulants and calculated a thixotropic index based on the stress evolution with time under a 1.6 s^{-1} shear rate. The thixotropic index was calculated by the ratio of the initial stress value to the plateau stress value and gave values between 1 (no thixotropy) and 1.8 depending on the samples. However, based on our experiments, since two

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3 different rheometers have been tested here, it seems that the initial stress overshoot height
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5 may be dependent on the response time of the rheometer. Indeed, the more precise DHR-2
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7 (TA Instruments) rheometer gave on the same sample a higher stress overshoot than the
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9 AR500 rheometer (TA Instruments). As a consequence, thixotropic indexes should not be
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11 compared when measured with different apparatus. To the authors' knowledge, 3ITTs results
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13 on mucus or mucus simulants have never been published. For comparison purposes however,
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15 we can refer to the recent work of Toker et al. [30] describing the results of 3ITTs performed
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17 on a mayonnaise or to the work of Mezger [35] in which identical 3ITTs are compared for
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19 two different paints. For a 100 s^{-1} step 2 in both studies, Toker et al. [30] measured a 100%
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21 recovery time varying of 56 to 432 s for a mayonnaise depending on the temperature, while
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23 Mezger [35] measured a 100% recovery time of 60 s or 300 s for the two paint samples
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25 (temperature not given). These results indicate that mucus simulants tested here show a
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27 slower thixotropic recovery compared to paints or mayonnaises (often used as "model"
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29 thixotropic fluids) investigated in these previous works. To conclude with thixotropy, it is
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31 obvious that the thixotropic nature of mucus simulants can give rise to much more studies. In
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33 particular, it would be interesting to perform an important set of 3ITTs with step 2 covering
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35 intermediate shear rate values between 1 s^{-1} and 100 s^{-1} . Modelling these experiments, a time
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37 dependent rheological model for mucus simulants at each active polymer concentration, could
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39 then be deduced. Such a time dependent model could be useful to account for the whole
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41 rheological complexities of these gels and could for instance allow to simulate their behavior
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43 in a model trachea, when submitted to various air pressure signals, such as signals imposed by
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45 clearance helping devices. This will be the purpose of a future work.

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52 In conclusion, in this work, a complete and quantified rheological characterization has been
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54 performed on mucus simulants at different active polymer concentrations (0.5wt% to 2wt% in
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3 ActigumTM). Stress amplitude SAOS sweeps first allowed determining the LVE range of these
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5 gels at 20°C. To account for the end of the LVE range and the subsequent transition towards
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7 flow, different transition stresses (namely, yield τ_y , flow τ_f and peak τ_{peak} stresses) were
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9 described. The increase of these transition stresses with ActigumTM concentration showed a
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11 power law dependence (eq. 2 & 3). Based on the yield stress value and the corresponding
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13 critical deformation, a cohesive volumetric energy ($J.m^{-3}$) that also followed an active
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15 concentration increasing power law (eq. 4) was calculated. Elastic and viscous moduli plateau
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17 values indicated, for all gel concentrations, stability ($G' > G''$), as well as a linear increase
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19 with the ActigumTM concentration. The temperature dependence of the LVE range was not
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21 significant in the range between 20 and 37 °C. Frequency SAOS sweeps displayed, only in
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23 the case of the lowest active polymer concentration, a low frequency moduli cross-over,
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25 showing the liquid-like behavior of this gel under very slow sollicitation and its solid-like
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27 behavior under relatively rapid sollicitation. More concentrated gels displayed a solid-like
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29 behavior over the whole range of tested frequencies. However, based on the $\tan(\delta)$ evolution
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31 one can anticipate a moduli cross-over under lower frequencies as the ActigumTM
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33 concentration increases. In flow mode, to separate viscoplasticity and shear-thinning from
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35 thixotropy, steady state flow curves were recorded and the HB model was used to quantify
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37 simulants SS viscoplasticity (via a yield stress measured in flow mode τ_{0HB}) and shear-
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39 thinning property (via the consistency (K) and flow indexes (n)). The evolutions of these
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41 parameters with the ActigumTM concentration displayed a power law increase for τ_{0HB} and K
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43 (eq. 6 & 7) and a linear decrease for n (eq. 8) indicating an enhanced viscoplasticity and
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45 shear-thinning ability for more concentrated gels. Finally the thixotropy of a 2wt% gel was
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47 tested performing two 3ITTs, submitting the sample to shear rates representative of normal
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49 shearing in the lungs ($1.6 s^{-1}$) or special events such as cough ($100 s^{-1}$). To quantify thixotropy
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51 it was proposed to evaluate a recovery time based on either 90% or 100% of recovery of the
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3 initial structure. Concerning this latter property, further investigations are necessary to fully
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5 characterize the thixotropic nature of mucus simulants, in particular to propose a time
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7 dependent rheological. This is clearly an interesting prospect for future works.
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10 11 **ACKNOWLEDGMENTS**

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14 The authors are indebted to the PhysioAssist Co. and the Association Nationale de la
15
16 Recherche et de la Technologie for the CIFRE grant 2014-1287 (O. Lafforgue PhD thesis)
17
18 and to the Laserson Co. for kindly providing the Actigum™ and Viscogum™ polymers. S.
19
20 Poncet acknowledges also the Canada Foundation for Innovation (John R. Evans Leaders
21
22 Fund n°34582) and the Natural Sciences and Engineering Research Council of Canada
23
24 through the Discovery Grant (RGPIN-2015-06512) for their financial support.
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4 **FIGURE LEGENDS**
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7 **Figure 1:** SAOS sweeps for a 1.5wt% sample. (a) Stress sweep ($\frac{1}{2\pi}$ Hz) (b) Frequency
8 sweep (1Pa).
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13 **Figure 2 :** Evolution of G' (a), G'' (b), tan(δ) (c) vs stress amplitude for different
14 ActigumTM concentrations
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18 **Figure 3:** (a) Limit stresses for LVE range & E_c (b) G' & G'' plateau values vs. ActigumTM
19 concentration
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23 **Figure 4:** Evolution of G' and G'' vs. stress amplitude at either 20°C or 37°C for a 0.75wt%
24 gel at $\frac{1}{2\pi}$ Hz
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30 **Figure 5:** Evolution of G' (a), G'' (b) and tan(δ) (c) vs frequency for different
31 ActigumTM concentrations
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35 **Figure 6:** SS rheogram of a 2wt% simulant
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39 **Figure 7:** HB model parameters vs. ActigumTM concentration
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42 **Figure 8:** 3ITT step 1 at 0.029 s⁻¹, step 2 at 1.6s⁻¹, step 3 at 0.029 s⁻¹
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45 **Figure 9:** 3ITT step 1 at 0.029 s⁻¹, step 2 at 100 s⁻¹, step 3 at 0.029 s⁻¹
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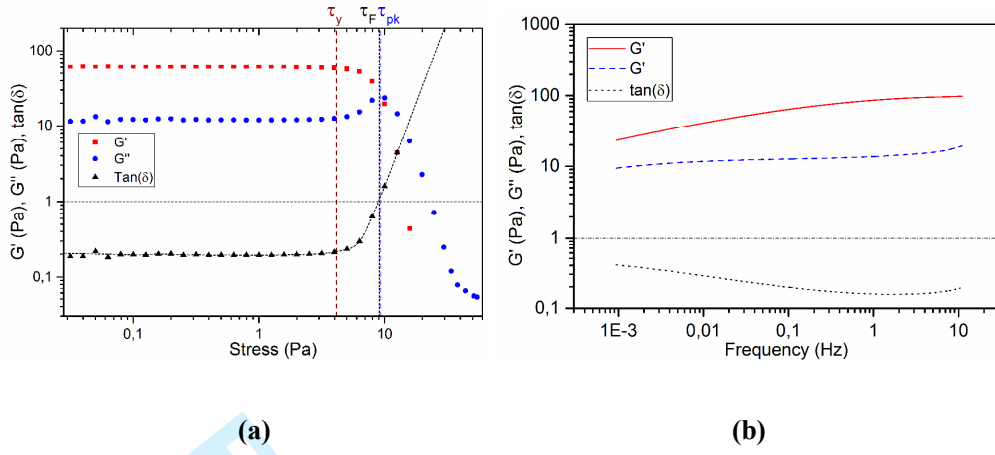


Figure 1.

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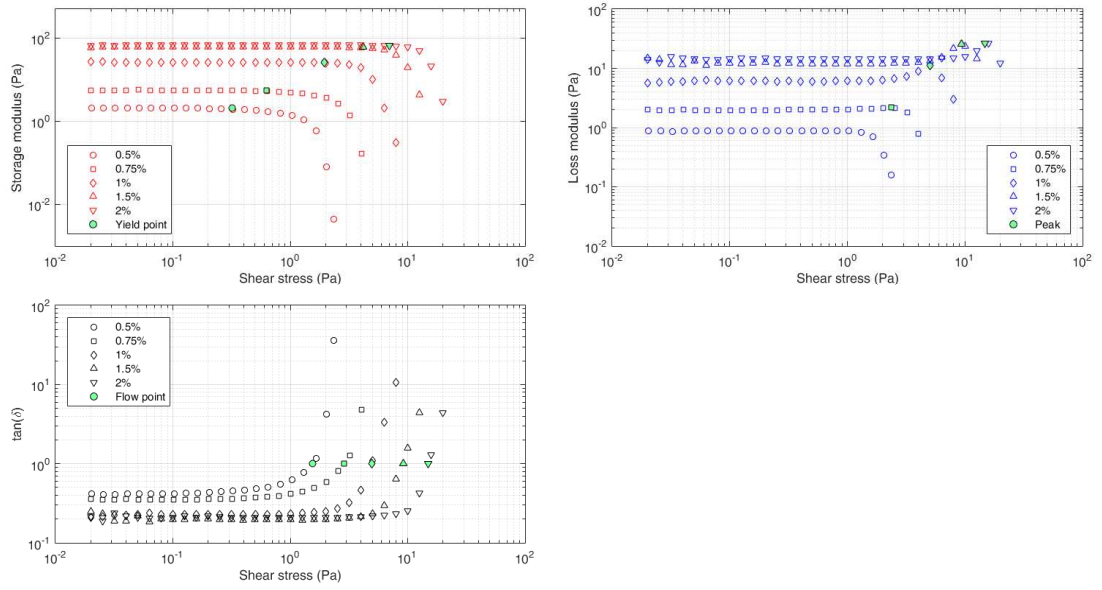


Figure 2

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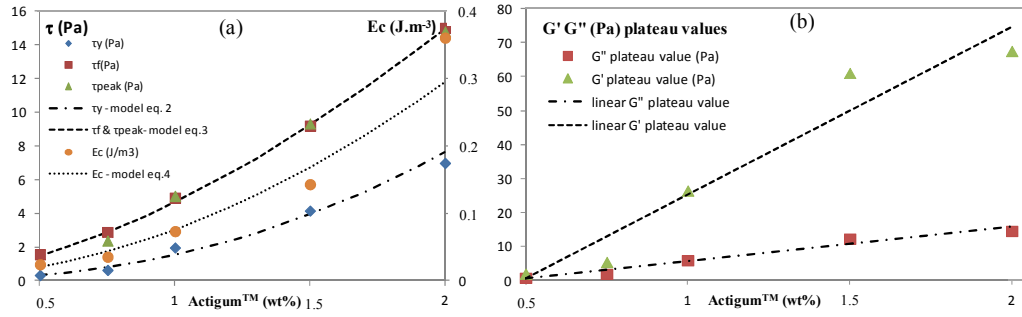


Figure 3

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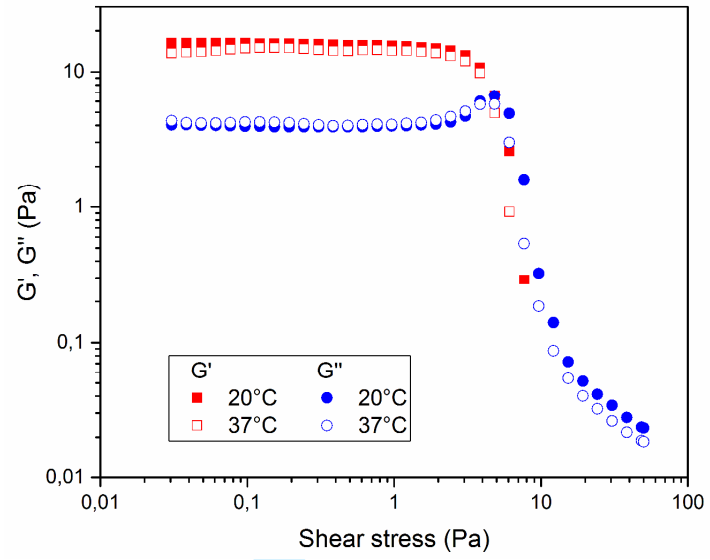


Figure 4

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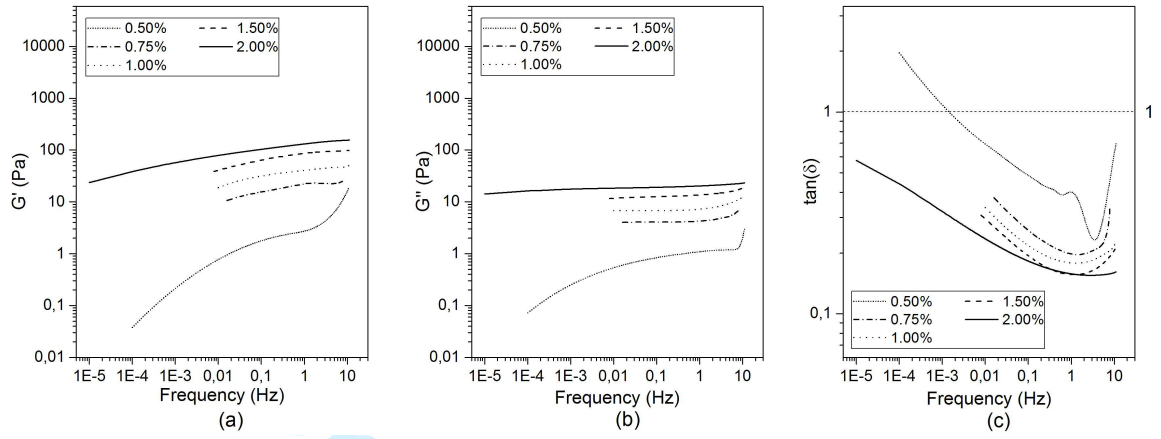


Figure 5

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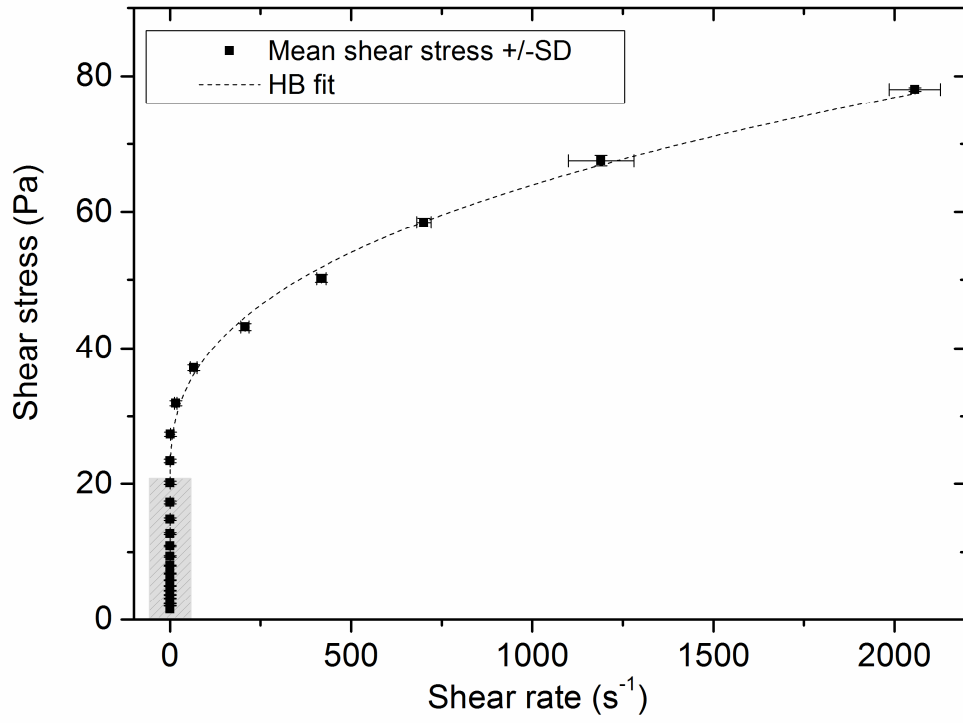


Figure 6: SS rheogram of a 2wt% simulant

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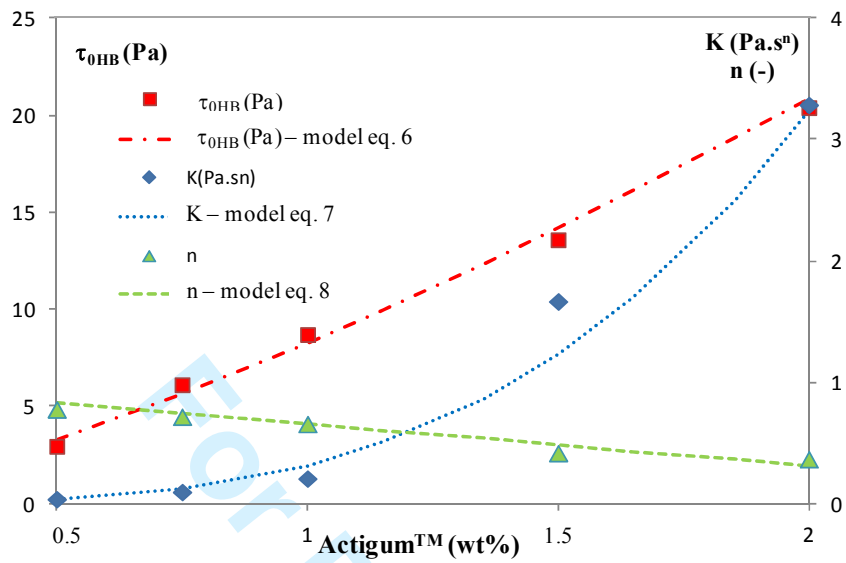


Figure 7

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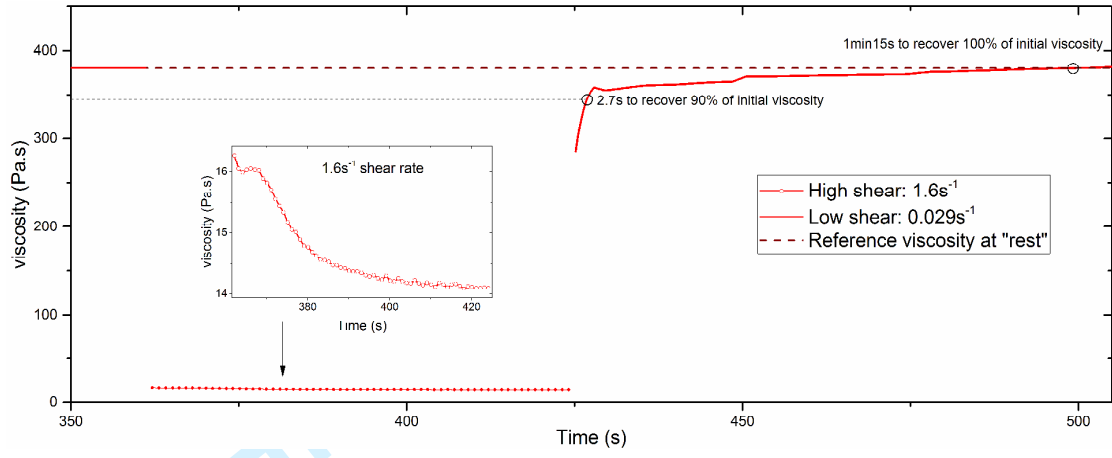


Figure 8

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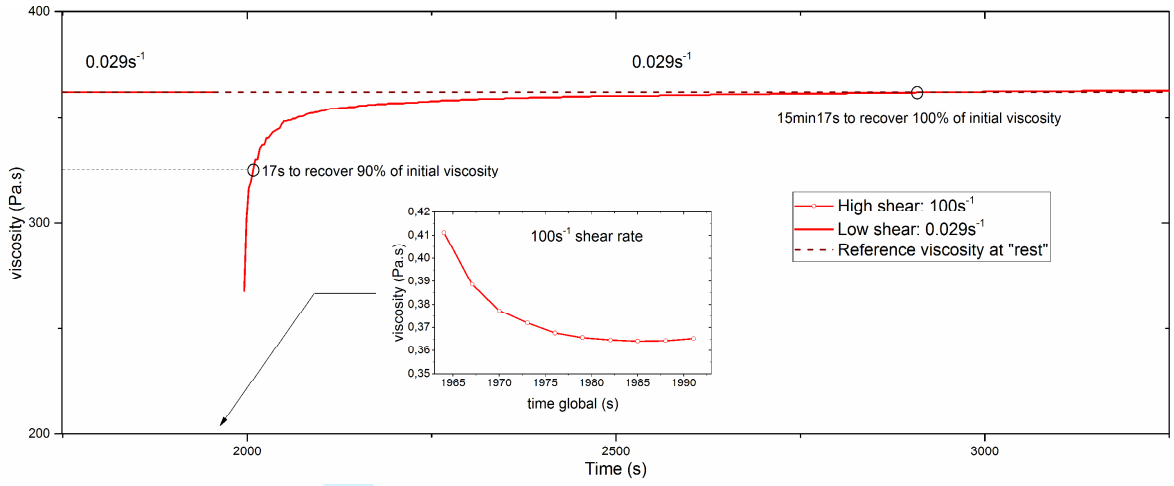


Figure 9

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Table 1. Evolution of limit stresses for LVE range, E_c and moduli plateau values vs active polymer concentrations.

[Actigum TM](wt%)	τ_y (Pa)	τ_f (Pa)	τ_{peak} (Pa)	E_c (J.m ⁻³)	G' (Pa)	G'' (Pa)
0.50	0.32	1.55	-	0.024	2.07	0.88
0.75	0.63	2.88	2.37	0.035	5.52	2.02
1	1.96	4.92	5.02	0.073	26.56	6.06
1.50	4.15	9.20	9.32	0.143	61.18	12.34
2.00	6.98	15.00	14.78	0.360	67.64	14.66

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Table 2. Evolution of the HB model parameters vs. active polymer concentrations.

[Actigum TM](wt%)	τ_{0HB} (Pa)	$K(\text{Pa}\cdot\text{s}^n)$	n
0.50	2.98	0.04	0.78
0.75	6.15	0.1	0.72
1	8.74	0.21	0.66
1.50	13.62	1.67	0.42
2.00	20.43	3.29	0.37

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