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# Broadband Dielectric Spectroscopy (BDS) investigation of molecular relaxations in durum wheat dough at low temperatures and their relationship with rheological properties

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#### ABSTRACT

Broadband Dielectric Spectroscopy (BDS) was used to study the dielectric relaxation processes of semolina doughs. The dielectric properties were analyzed as a function of water content, and, additionally, the effects of NaCl presence and semolina characteristics were investigated. The dough was prepared using three different varieties of semolina. BDS measurements were conducted using a custom-made Rheo-dielectric tool composed of a Broadband Dielectric/Impedance Spectrometer connected to a strain-controlled rheometer. The temperature range investigated was from -135 °C to 25 °C, with a step of 5 °C, while the frequency range was  $10^{-1} - 10^7$  Hz. Dielectric spectroscopy turned out to be a valuable technique for dough characterization. It is capable to distinguish the carbohydrates contribution and the different interactions between water and dough components. Moreover, this unique combination allows assessing correlations between rheological and dielectric properties, like the compliance of dough as a function of the relaxation processes and the influence of semolina components.

## 1. Introduction

Durum wheat is the most appropriate raw material for making pasta; it is one of the hardest varieties of wheat, the milling of which produces coarse particles called semolina (Sicignano, Di Monaco, Masi, & Cavella, 2015). Although common wheat is preferred to durum one in the bread-making, in the last years the production and consumption of bread made of durum wheat have been increased (Mefleh et al., 2019). This is due to the several advantages that this kind of bread presents compared to common bread (e.g. yellowish colour, appreciated characteristic taste and odour, fine uniform crumb structure, higher content of proteins, and a longer shelf-life) (Ficco et al., 2017; Pandino, Mattiolo, Lombardo, Lombardo, & Mauromicale, 2020).

The quality of semolina doughs is strongly influenced by the interactions among the ingredients in the presence of water and salt (Fanari, Carboni, Grosso, & Desogus, 2020; Torbica, Mocko Blažek, Belović, & JanićHajnal, 2019). Several studies concern the influence of ingredients and semolina variety on the rheological properties of doughs and, in the most recent literature, new insight into the dough microstructure is presented (Bonilla, Erturk, & Kokini, 2020; Bonilla, Erturk, Schaber, & Kokini, 2020). The role of gluten in conditioning the texture and rheological properties is well known, and due to the formation of a viscoelastic network during hydration and mixing (Bonilla, Schaber, Bhunia, & Kokini, 2019). However, also starch has a relevant influence on the rheological properties of dough. Indeed, even if it has no significant network-forming properties, it is embedded into the gluten network, it has a large volume fraction and interacts, to a certain extent, with the surrounding matrix (Brandner, Becker, & Jekle, 2019). For example, it was found that the viscosity increases with the damaged starch content due to the higher amount of absorbed water (Barak, Mudgil, & Khatkar, 2014). Even the particle size showed effects on viscoelastic properties of durum wheat doughs (Edwards, Dexter, & Scanlon, 2002). On the other side, semolina doughs have been less studied concerning their dielectric properties.

Broadband Dielectric Spectroscopy (BDS) is a non-invasive technique useful to characterize the distribution and influence of the main

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polar components (electrical dipoles within the molecules) which are present in the system (Kang, Lu, Cheng, & Jin, 2015), e.g. water. This technique has great importance in food and in the biomaterials systems field since it is able to discriminate between the free and the various coexisting bond water states in the material and relates the different mobility to the structural and textural changes (Li, Zhang, & Yang, 2019; Nielsen et al., 2016). Various non-equilibrium states and the corresponding transitions, resulting from molecular movements (translation and rotation) and flow dynamics, could be observed during the processing and storage of bakery products (Wang & Zhou, 2017). The rates of these changes are dependent on molecular mobility, product formulation, water activity, storage condition, processing parameters, etc. (Slade & Levine, 1995). Despite the potentiality of this spectroscopic technique, the knowledge and understanding of the dielectric properties of food components are so far quite limited. An important parameter that can be determined using dielectric spectroscopy is the glass transition temperature (Tg), a concept originally developed and applied for synthetic polymers. Meanwhile, it has been applied to many food products to predict their stability (Balasubramanian, Devi, Singh, D Bosco, & Mohite, 2016). Regarding dough, in the literature, we can find few studies related to dough Tg determination. In several publications (Akbarian, Koocheki, Mohebbi, & Milani, 2016; Ding et al., 2015; Matuda, Pessôa Filho, & Tadini, 2011; Meziani et al., 2011), a relaxation process was detected at around -40 °C, through differential scanning calorimetry (DSC), and it was associated with the dough glass transition. However, one should remark that dough and foods are mixtures of different components that have a spatial heterogeneity at longer length scales (e.g. > 5 nm), consequently, it is not possible to define a single glass transition temperature, but rather a temperature range, the width of which is controlled by the dynamic and spatial heterogeneity of the system (Böhmer et al., 1998; Heuer, Wilhelm, Zimmermann, & Spiess, 1995; Tracht et al., 1998). This fact generates the necessity to investigate better the relaxation processes in the dough since its dielectric properties are strongly related to its microstructure, mostly consisting of the so-called gluten network (Miller & Hoseney, 2008). Also starch has a relevant role in determining the microstructural and rheological characteristics of durum wheat dough, but it is often underestimated compared to that of gluten. Indeed, especially when semolina is taken into account, the surface characteristics of starch granules may affect the protein-starch interaction and the viscoelastic behaviour of the dough (Edwards et al., 2002). Dielectric spectroscopy has shown the presence of several molecular relaxations in starch. Butler and Cameron (Butler & Cameron, 2000) reported a relaxation in the range from -120 to -90 °C (denoted with ' $\gamma_1$ '), obeying the Arrhenius law, as reflected by single activation energy, and another relaxation at slightly lower temperatures (denoted with ' $\gamma_2$ '), which was observed only in starch with low water contents. Roudaut et al. (Roudaut, Maglione, Van Dusschoten, & Le Meste, 1999) also observed a secondary relaxation, with apparent activation energy ( $E_a$ ) of about 50 kJ/mol, between -60 and -40 °C (depending on the frequency), which was related to the carbohydrate ingredients (sucrose and starch) of bread. Also non-starch polysaccharides, notwithstanding they represent only about 2-2.5% of the flour mass (Migliori & Gabriele, 2010), have a great influence on dough properties (Bibi, Villain, Guillaume, Sorli, & Gontard, 2016). Ross (Roos, 1993) reported various glass transition temperatures of mono- and di-saccharides, and alditols, in the range between -40 and -70 °C, at which several works previously reported associated relaxation processes. Laaksonen and Roos (Laaksonen & Roos, 2001a), using dielectric analysis at 0.1 Hz, found a Tg of -33 °C and two relaxation processes, one  $\beta$  relaxation at -85 °C and one  $\gamma$  process at -148 °C. In another publication (Laaksonen & Roos, 2000), the same authors noticed that these temperatures increase as frequency increases and they reported four relaxation processes: one first-order phase transition associated with ice melting ( $E_q = 266.4 \text{ kJ/mol}$ ), another one with a glass transition ( $E_a = 97.1 \text{ kJ/mol}$ ), one single  $\beta$  relaxation ( $E_a = 63.0 \text{ kJ/mol}$ ) and one single  $\gamma$  process ( $E_a = 31.5 \text{ kJ/mol}$ ). The results were obtained through

dielectric spectroscopy, which revealed much more sensitive in observing thermal transitions with respect to Differential Scanning Calorimetry (DSC) and Dynamic Mechanical Analysis (DMA). However, even through DMA, at frequencies of 1, 5 and 10 Hz, Ribotta and Le Bail (Ribotta & Le Bail, 2007) found a relaxation process at around -7 °C, corresponding to melting of ice, an  $\alpha$ -relaxation (glass transition) at around -35 °C and a  $\beta$ -relaxation at around -78 °C, while Laaksonen and Roos (Laaksonen & Roos, 2001b) reported a  $\beta$  relaxation at about -80 °C and a  $\gamma$  process in the temperature range from -100 °C to -136 °C at 2.5, 5 and 10 Hz.

Despite starch being contained in high amounts in baked doughs, the main component that influences the dielectric properties is water due to its high mobility, large dipole moment, and ability to act as a plasticizer. Goedeken et al. (Goedeken, Tong, & Virtanen, 1997) reported that the dielectric properties are related not only to the water content but also to the interaction forces between the other components of the material and the dissolved water molecules. According to this, any influence of the starch kind, or crystallinity, is overshadowed by the one of bound water dipoles rotational motion (Roudaut, Simatos, Champion, Contreras-Lopez, & Le Meste, 2004). Water plasticizing properties are well known and, due to this characteristic, moisture content of about 30% wt. Is sufficient to reduce the Tg of many carbohydrates and proteins, including starch and gluten, up to about -10 °C (Levine & Slade, 1990). Moreover, water influences the temperature effect on the dough dielectric properties. These results highlight the need for testing samples with a known and controlled water content. Water dielectric relaxations can be observed at high frequencies higher than 100 MHz (Popov, Ishai, Khamzin, & Feldman, 2016) if the local motion of the water is not substantially constrained. However, the vibration of water atoms or bonds and the local reorientation of small groups of atoms are still possible at lower frequencies, even in the glassy state (Chan, Pathmanathan, & Johari, 1986), and these movements facilitate the structural relaxations (Perera, 2002).

Furthermore, the addition of salt, specifically NaCl, affects the dielectric properties of dough. In more detail, the salt presence significantly increases the dielectric loss ( $\varepsilon$ ") because it generates an additional conductivity due to the ion migration (Fanari, Muntoni, Dachena, Carta, & Desogus, 2020). Indeed, the number of dissolved ions is the most important factor affecting the absolute values of the dielectric loss (Kim & Cornillon, 2001). The ion concentration directly influences water mobility since only the water molecules having sufficient mobility can participate in dissolving salts. In addition, other publications (Laaksonen, Kuuva, Jouppila, & Roos, 2002; Laaksonen & Roos, 2001a, 2003) showed that the presence of NaCl significantly decreases the dough T<sub>g</sub> but also the temperature of the other relaxation processes, if the same mobility is considered.

The objective of the present study is to investigate the effect of ingredients (water and NaCl) and of the semolina composition on the dielectric relaxations occurring at low (sub-zero) temperatures in the dough. In particular, at these temperatures, according to the previous considerations, the dielectric response could be informative on carbohydrates and thus on their role in the definition of the rheological properties of dough, as gluten is expected to be not detectable in such conditions, and the water mobility is reduced as much as possible. Therefore, measuring and understanding the phase transitions and relaxation processes occurring in doughs at sub-zero temperatures, and finding possible correlations with rheological parameters, could be useful for understanding the role of carbohydrates on the mechanical properties of semolina doughs. This new knowledge could find application in the pasta and baking manufacturing, where optimisation of the product texture is crucial. So a more and more precise definition of the rheological characteristics of doughs is necessary. Even the manufacturing of frozen doughs could benefit from this information, which is useful for temperature optimisation to limit changes in the material. Moreover, information about the carbohydrate effects on the rheological properties of foods has the potential to be useful also in other

food sectors, e.g. in gluten-free productions, where the texture reveals to be even more critical.

## 2. Materials and methods

## 2.1. Sample preparation

For each sample, 300 g of semolina (three different varieties were used), distilled water, and sea salt, in different amounts, were kneaded using a measuring mixer type 350 (Brabender® GmbH, Duisburg, Germany). The mixing time was determined as the one required to reach the dough maximum strength point, as observed in previous investigations (Fanari et al., 2019; Fanari, Desogus, Scano, Carboni, & Grosso, 2020), and it was found to be 3-5 min, depending on the sample. The rotational mixing speed was set to 20 rpm to reduce the effect of structural breaks in the dough network. Three different types of semolina were investigated. Two are of non-commercial monovarietal species cultivated in the "S. Michele" experimental farm of the Agricultural Research Agency of Sardinia (AGRIS) in Ussana (Italy), resulting from the milling of Karalis and Cappelli grains. The third kind is a commercial blend acquired in the German retail stores (Gold Puder-Hartweizen Grieß, Aurora Mühlen GmbH, Hamburg, Germany). It should be remarked that their properties are quite different in terms of protein and gluten content and gluten index, as can be seen in Table 1. Gluten index (GI) is a parameter providing information on both gluten quality and gluten quantity, and it expresses the weight percentage of the wet gluten remaining on a sieve after automatic washing with a salt solution and centrifugation (Oikonomou, Bakalis, Rahman, & Krokida, 2015). The semolina protein content was determined by the nitrogen combustion method (ISO 16634-2:2016, 2016) using a Leco FP528 nitrogen analyzer (LECO, Stockport, UK). The gluten content and the gluten index (GI) of semolina were determined following the ICC standard method No. 158 (ICC, 1995) by using the Glutomatic 2200 system (Perten Instruments AB, Huddinge, Sweden). Table 2 reports the mass percentual composition of the samples, with respect to the semolina weight, in terms of water and salt content.

## 2.2. Dielectric characterization

For the dielectric measurements, the sample was placed between two custom-made invar-steel parallel plate electrodes of 25 mm diameter, mounted on an ARES-G2 strain-controlled rheometer (TA Instruments, New Castle, USA). The initial gap in the measurement was 1 mm between the plates. The distance was adjusted during the experiment to maintain the normal force at around 1 N. This is important because, during the freezing and defrosting processes, the sample changes its volume. Thus, the spacing during the experiments resulted from 0.8 to 1.2 mm. The experimental setup is shown in Fig. 1. As soon as the sample loading was completed, the dough was quickly cooled to -135 °C using the rheometer temperature control system. This operation was conducted in the fastest way to avoid the sample's drying, limit the growth of frozen water crystals, and keep the structure and properties of starch unmodified (Yang et al., 2021). After the thermal stabilization, the dielectric analysis was carried out step by step by raising the sample temperature by 5  $\,^\circ\text{C}$  for each step, from -135 to +25  $\,^\circ\text{C}.$  For each temperature step, after thermal stabilization was reached, a dielectric measurement was performed.

#### Table 1

Semolina varieties characteristics (means  $\pm$  STD); values with different letters in the same column are significantly different (p-value  $\leq$  0.05).

	Proteins (%)	Gluten (%)	Gluten Index (%)
Cappelli (CAP) Karalis (KAR) Commercial (COM)	$\begin{array}{c} 14.0 \pm 0.17 \ ^{a} \\ 11.0 \pm 0.05 \ ^{b} \\ 12.0 \pm 0.12 \ ^{c} \end{array}$	$\begin{array}{c} 12.3 \pm 0.30 \; ^{a} \\ 7.0 \pm 0.34 \; ^{b} \\ 8.3 \pm 0.27 \; ^{c} \end{array}$	$\begin{array}{c} 16.5 \pm 2.35 \\ 98.4 \pm 0.79 \\ 90.3 \pm 2.12 \\ ^{\rm c}\end{array}$

Table 2

Composition of the dough reported as wt.% based on semolina weight.

Sample	Semolina variety	Water (wt.%)	Salt (wt.%)
CAP40	Cappelli	40	0
CAP50	Cappelli	50	0
CAP60	Cappelli	60	0
KAR40	Karalis	40	0
KAR50	Karalis	50	0
KAR60	Karalis	60	0
COM40	Commercial	40	0
COM50	Commercial	50	0
COM60	Commercial	60	0
CAP50S	Cappelli	50	1.5
KAR50S	Karalis	50	1.5
COM50S	Commercial	50	1.5

Dielectric measurements were carried out by a Novocontrol Technologies Alpha High-Resolution Broadband Dielectric/Impedance Spectrometer (Montabaur, Germany), within the frequency (f) range  $10^{-1}$  -  $10^7$  Hz, connected with the strain-controlled rheometer parallel plates in a custom-made experimental apparatus already described in previous works (Hyun, Höfl, Kahle, & Wilhelm, 2009; Meins, Dingenouts, Kübel, & Wilhelm, 2012). The signal voltage to generate the electric field was set to 0.5 V. Then, phase and state transitions of wheat dough samples were studied by monitoring the imaginary part of the complex permittivity ( $\varepsilon$ "), the dielectric parameter tan( $\delta$ ) (tan( $\delta$ ) =  $\epsilon''/\epsilon'$ ) and the real part of the conductivity  $\sigma'$  as a function of the temperature ( $\epsilon'$  is the real part of the permittivity). The frequency dependent  $\varepsilon''$  is particularly important since its value is proportional to the conductance and can be related to the energy required to align dipoles or allow the ions movement so that it can be linked to the relaxation processes. To clearly see these relaxation processes, notwithstanding the Ohmic conduction influence, due to the presence of ions, in the cases of samples without salt, it is possible to apply the Kramers-Kronig rule reported in Equation (1) (Wübbenhorst & Van Turnhout, 2002). This relationship, which is a function of the angular frequency  $\omega$ , defines an imaginary value  $\varepsilon_{der}''$ , which approximately equals the one of the Ohmic-conduction-free dielectric loss ( $\varepsilon''_{rel}$ ) for rather broad peaks, like those of the  $\alpha$ -transition or the secondary relaxations (Steeman & van Turnhout, 1994).

$$\varepsilon_{der}^{\prime\prime} = -\frac{\pi}{2} \frac{\partial \varepsilon^{\prime}(\omega)}{\partial [\ln \omega]} \tag{1}$$

A comparison between measured values of  $\varepsilon''$  and the calculated values of  $\varepsilon''_{der}$  for a COM50 sample is shown in Fig. 2a. The derivative method (Equation (1)) emphasizes the  $\varepsilon''$  peaks, in correspondence to the relaxation frequencies, making their individuation and analysis easier.

The relaxation frequency was then calculated as the peak frequency of  $\varepsilon''_{der}$ . To do this, the dependence of  $\varepsilon''_{der}$  on frequency was firstly modeled by using the Havriliak-Negami (HN) relationship (Havriliak & Negami, 1967; Volkov, Koposov, Perfil'ev, & Tyagunin, 2018):

$$\varepsilon^{"} = \Delta \varepsilon \left[ 1 + 2 \cdot (\omega \cdot \tau_{HN})^{\alpha} \cdot \cos\left(\frac{\pi \alpha}{2}\right) + (\omega \cdot \tau_{HN})^{2\alpha} \right]^{\left(-\beta^{\prime}_{2}\right)} \sin(\beta \varphi) \right]$$
(2)

The parameters  $\alpha$  and  $\beta$  are the fractional HN shape parameters (corresponding to high-frequency skewness and breadth, respectively);  $\Delta \varepsilon$  is the relaxation strength, that is the difference between the static (low frequency) permittivity and the permittivity at the high-frequency limit ( $\varepsilon_{s}$ - $\varepsilon_{\infty}$ );  $\varphi$  is a parameter obtainable by Equation (3);  $\tau_{HN}$  is the relaxation time associated with the characteristic frequency  $\omega_{HN}$ , which is related to the frequency of maximum loss ( $\omega_{max}$ ) by Equation (4).

$$\varphi = \arctan \frac{\sin(\alpha \pi/2)}{\left(\omega \tau_{HN}\right)^{-\alpha} + \cos(\alpha \pi/2)}$$
(3)



Fig. 1. Set-up (a) and (b) a scheme of the unique rheology-dielectric spectroscopy combination (Hyun et al., 2009).



**Fig. 2.** a) Comparison between  $\varepsilon''$  (black) and  $\varepsilon''_{der}$  (green) values at -50 °C for COM50 sample; it shows that the derivative method (Equation (1)) enhances the peaks in  $\varepsilon''$ . b) Example of fitting with Havriliak-Negami relationship for COM50. c)  $\varepsilon''_{der}$  as a function of frequency (in Hz) at different temperatures as indicated. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

& Kremer, 2003):

The parameters calculated by fitting Equation (2) (an example of the fitting is shown in Fig. 2b) were used in Equation (4) to calculate the frequency corresponding to the maximum of the  $\varepsilon''$  function (Schönhals

$$\omega_{max} = \omega_{HN} \left[ \frac{\left( \sin \frac{\alpha \beta \pi}{2 + 2\beta} \right)}{\left( \sin \frac{\alpha \pi}{2 + 2\beta} \right)} \right]^{-1/\alpha}$$
(4)

Relaxations can be represented on mobility maps which are displayed as the logarithm of the relaxation frequency as a function of the reciprocal values of the absolute temperature (Arrhenius activation). This can be used to characterize the different motions taking place in a material at a given temperature and frequency range. Then, the apparent activation energy of relaxations was evaluated by Equation (5), describing an Arrhenius type activation behaviour (Iacob, Yoo, & Runt, 2018):

$$\omega_{\beta}(T) = \omega_0 \cdot e^{-\frac{La}{RT}} \tag{5}$$

where  $\omega_{\beta}$  is the relaxation angular frequency,  $E_a$  is the apparent activation energy; *R* is the gas constant, *T* is the absolute temperature, and  $\omega_0$  is a constant, having the meaning of a maximum possible angular relaxation frequency at infinite temperatures. The sensitivity analysis concerning frequency allows the discrimination between relaxation processes and structural changes such as melting or crystallization. So, the values of the activation energies of the samples were calculated for each of them to enhance the difference between the relaxation processes occurring and relate them to the difference in the samples. From Fig. 2c it is possible to observe how the frequency peaks in  $\varepsilon''_{der}$  are moving to lower frequencies as temperature decreases.

## 2.3. Rheological characterization

The rheological experiments were performed using the same device, already described in the previous subsection 2.2, equipped with standard 25 mm plates and a Peltier-based system for temperature control. Immediately after the kneading process, a piece of dough was loaded on the rheometer, compressed to a gap of 2 mm, and then left to rest for 15 min to allow the material relaxation, as suggested in the literature (Phan-Thien & Safari-Ardi, 1998). To prevent evaporation and, consequently, drying of the sample, a layer of silicon oil was applied to its edge. The temperature in the rheometer was kept constant at 25 °C. Then, creep tests were performed applying a 50 Pa constant stress to the sample to stay in the linear viscoelastic region, according to the values of yield stress limit (70-100 Pa) determined through amplitude sweeps previously performed. This value also agrees with the values used to perform creep experiments on dough in other studies found in the literature (Döring, Nuber, Stukenborg, Jekle, & Becker, 2015; Mironeasa & Codină, 2019; Moreira, Chenlo, Torres, & Rama, 2014). The stress was applied for 150 s. Each test for each sample was repeated three times, and the mean value was considered in the present study. The compliance J, which is the ratio between deformation and applied stress, was measured against time to evaluate the rheological properties of the samples. The dependence of J on time is, in general, described in terms of mechanical models composed of basic elements. One of the most used is the Burgers model, which works as a Kelvin and a Maxwell model placed in series. Mathematically, it can be written as (Mainardi & Spada, 2011):

$$J(t) = J_0 + J_m \left( 1 - e^{-\frac{t}{t}} \right) + \frac{t}{\eta_0}$$
(6)

The model parameters are the steady-state viscosity  $\eta_0$ , the instantaneous and delayed compliance, respectively  $J_0$  and  $J_m$ , and the delay time  $\tau$  that indicates how long the material takes to settle to deformation. Many applications of this model to bread dough characterization can be found in the literature (Beck, Jekle, & Becker, 2012; Mironeasa & Codină, 2019; Sun, Koksel, Nickerson, & Scanlon, 2020).

## 3. Results

## 3.1. Dielectric characterization

The  $\varepsilon''$  spectra were analyzed, and the relaxation peaks were detected in terms of peak frequency. All the spectra at temperatures higher than 0 °C present a deflection in the  $\varepsilon''$  values, erroneously attributable to the  $\alpha$ -segmental relaxation process. This peak is, instead, related to the conductivity relaxation and at around 0 °C is significantly affected by water crystallization. The effect of this process can be observed from Fig. 3, where tan( $\delta$ ) and  $\sigma'$  for samples with 50% of water and no salt (CAP50, COM50, and KAR50) at different temperatures are reported as an example. The static conductivity  $\sigma_0$ , detectable from the plateau in  $\sigma'$ , sensibly decreases when the temperature goes to 0 °C and sub-zero values. Consequently, also the corresponding peak in tan( $\delta$ ) is decisively moving at lower frequencies. In this range of temperatures also the  $\alpha$ -segmental relaxation process is screened by the dc conductivity and water crystallization peak.

For each tested temperature, together with  $tan(\delta)$  and  $\sigma_0$ , the relaxation frequencies are reported as a function of inverse temperature in Fig. 4a for CAP50, KAR50, and COM50 samples. It is possible to see that only two relaxation processes are revealed. Both relaxations occur at temperatures lower than the temperatures characteristic of the crystallization process. One process at very low temperatures, classifiable as  $\gamma$  relaxation, and another one at higher temperatures, attributable to a  $\beta$ relaxation, can be found. Their trend suggests a single motion with Arrhenius type behaviour of the relaxation frequency with respect to the temperature. Thus, for each sample, the relaxation frequency data as a function of temperature were fitted using Equation (5) (Arrhenius dependence law). Fig. 4 shows, as an example, the regressions for CAP50, KAR50, and COM50, while the mobility map and respective regressions for the other samples are shown in Figure A1 reported in Appendix A. Additionally, in Table 3, the calculated values of activation energy for  $\beta$  ( $\beta$   $E_a$ ) and  $\gamma$  ( $\gamma$   $E_a$ ) relaxations, for all the samples, and the 95% confidence intervals, are reported. Table 4 reports the values of the statistical parameters of the regressions. The computed statistics ( $R^2$  and MSE) suggest that the exponential Arrhenius dependence reveals to be a simple, though effective, method to describe the relaxation frequency as a function of temperature, with a value of Adj-R<sup>2</sup> higher than 0.99 in most cases. Moreover, the only presence of an Arrhenius dependence advises that there are no contributions of a multimode glass transition, so it is not possible to calculate any Tg since the glass transitions usually show a different behaviour, following a Vogel-Fulcher-Tammann type temperature dependence (Iacob et al., 2018).

The trend of  $tan(\delta)$  as a function of the temperature for a fixed frequency (100 Hz) is shown in Fig. 5 for three samples chosen as an example, and for the other samples in Figures A2-A3 (Appendix A). The experimentally determined  $tan(\delta)$  values in the temperature range investigated show three main peaks for all the samples. The correspondent temperature and  $tan(\delta)$  height values of these peaks are reported in Table 5 for water crystallization, ( $W_CP$ ),  $\beta$  ( $\beta P$ ), and  $\gamma$  ( $\gamma P$ ) processes, respectively. The dielectric excitation frequency of 100 Hz was chosen as it allows to clearly see all the three characteristic peaks identified (Laaksonen et al., 2002). Indeed, at lower frequencies, peaks are not clearly detectable, especially the ones at a higher temperature. The first peak in  $tan(\delta)$  is located at about 0 °C, and it is likely related to water crystallization and not to an  $\alpha$  relaxation process. This theory is supported by visual inspection of Fig. 4 and A1, where conductivity and tan(\delta) relaxation peaks are reported as a function of inverse temperature. In fact, a steep fall in correspondence of the water freezing point (0 °C) is indeed observed. Another peak is visible in the range between -60 and -40 °C, which probably corresponds to the  $\beta$  process observed



**Fig. 3.** Tan( $\delta$ ) and  $\sigma'$  as a function of frequency (logarithmic scale) for (a) CAP50, (b) KAR50 and (c) COM50. The values are reported at different temperatures: 25 °C (blue), 10 °C (magenta), 0 °C (red), -10 °C (black). The black line follows the tan( $\delta$ ) peak moving while the green line is representative of the  $\sigma$ 0 drop. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

in the mobility maps. The other peak which was identified is precisely located at -105 °C for all the samples and could be due to the  $\gamma$  relaxation previously detected in the mobility maps.

The influence of salt is appreciable when comparing the samples data without salt (COM50, KAR50, and CAP50) and with salt (COM50S, KAR50S, and CAP50S). Focusing on the data reported in Fig. 6 and Figure A1, an additional relaxation process is observed when salt is added for both COM and KAR samples. This additional relaxation (named  $\gamma_2$ ) follows an Arrhenius type dependence on the temperature and has activation energy ( $\gamma_2 E_a$ ) in the range of 80–90 kJ/mol. In the CAP50S sample, no additional relaxations were detected, probably because they are hidden by more dominant ones. In effect, the CAP samples are the only ones that are not significantly affected in any of the  $tan(\delta)$  peaks. About the  $tan(\delta)$  peak temperatures, a decrease of 20 °C in the  $\beta$  relaxation peak temperature can be observed for all the samples. while the other peaks show a shift in the order of  $\pm 5-10$  °C. Moreover, for the KAR50S sample, as shown in Fig. 6,  $tan(\delta)$  shows a very sharp additional peak (denoted with "i" in Fig. 6), at about -40 °C, due to the additional relaxation process observed in the mobility map, that probably is hidden by the higher  $\beta$  relaxation peak for CAP50S and COM50S, as it is observable in figure A3 (Appendix A).

## 3.2. Rheological characterization

Results are presented in Fig. 7, where compliance as a function of time for each semolina variety is shown. Results pertaining to CAP, KAR, and COM samples with different water amounts and salt addition are reported in Fig. 7a, 7b, and 7c, respectively. The main outcomes of this analysis are a weak influence of the water amount on the rheological properties for the CAP samples and, on the contrary, a high sensibility to water for the COM samples. On the other hand, KAR samples show intermediate rheological properties. Moreover, the salt addition has a mildly negative effect on the dough deformability, increasing the compliance values of KAR and COM samples while, in the case of CAP

samples, it produces a considerable decrement of compliance, and, as a consequence, an increase in the strength of the dough network. Fig. 7d, instead, reports the semolina variety influence on the rheological properties. It is noticeable that the CAP50 sample presents the lowest compliance. CAP results to be the strongest semolina variety, followed by KAR, and COM turned out to be the weakest, with higher deformability.

Non-linear regression was accomplished on the compliance vs time data in order to calculate the Burgers model (Eq. (6)) parameters. Their values for each sample are reported in Table 6.

## 3.3. Statistical analysis

Pearson correlation coefficients were computed to evaluate correlations between rheological parameters and dielectric ones. Also correlation coefficients between these parameters and water content, protein amount, gluten amount, and gluten index were evaluated. Finally, statistical significance for all these correlations was evaluated through the p-value estimation. The results are reported in Table 7.

#### 4. Discussion

#### 4.1. Water crystallization

The water crystallization process was detected between 0 and -10  $^{\circ}$ C. This process provides information about the water state in the system.

The height of the tan( $\delta$ ) peak shows a very weak and negative correlation with the total water content (r = -0.18). On the other hand, it shows a good and significant correlation with the semolina protein and gluten amount (r > 0.76). This means that the dielectric response is mostly due to the interactions between water molecules and the structural components of the dough, in other words to the bound water.



**Fig. 4.** Mobility map ( $\omega$  in rad/s) and Arrhenius model regression with  $E_a$  values for (a) CAP50, (b) KAR50 and (c) COM50; the samples show two relaxation processes,  $\gamma$  with  $\gamma E_a$  around 30–35 kJ/mol and  $\beta$  with  $\beta E_a$  around 60–70 kJ/mol. Triangles represent tan( $\delta$ ) while green stars  $\sigma_0$  values. The red dashed line highlights the crystallization temperature, in correspondence of which a drop in tan( $\delta$ ) is observable. (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

#### Table 3

Activation energy  $(E_a)$  and 95% confidence interval for dough samples with different water amounts.

Sample	$\beta E_a$ (kJ/mol)	$\gamma E_a$ (kJ/mol)	$\gamma_2 E_a$ (kJ/mol)
COM40	$72.5\pm2.8$	$33.7\pm2.2$	_
COM50	$59.3 \pm 3.5$	$30.3\pm1.1$	-
COM60	$73.6\pm2.2$	$31.5\pm0.9$	-
KAR40	$74.5\pm5.7$	$40.4 \pm 1.0$	-
KAR50	$57.8 \pm 4.2$	$35.6 \pm 1.6$	-
KAR60	$62.8\pm3.4$	$31.8\pm1.5$	-
CAP40	$60.0\pm 6.9$	$36.0\pm1.6$	-
CAP50	$57.7\pm4.0$	$31.8\pm1.4$	-
CAP60	$52.4\pm4.1$	$31.8 \pm 1.3$	-
COM50S	$57.1\pm5.8$	$38.6 \pm 1.0$	$87.9 \pm 3.5$
KAR50S	$63.6\pm6.1$	$39.5\pm1.4$	$83.3 \pm 1.8$
CAP50S	$\textbf{66.9} \pm \textbf{5.2}$	$41.1 \pm 1.2$	-

Table 4				
Statistical	parameters	of the	e regressior	15

Sample	$\beta E_a$		$\gamma E_a$		$\gamma_2 E_a$	$\gamma_2 E_a$		
	RMSE	Adj-R <sup>2</sup>	RMSE	Adj-R <sup>2</sup>	RMSE	Adj-R <sup>2</sup>		
COM40	0.056	0.995	0.051	0.994	-	-		
COM50	0.089	0.992	0.058	0.996	_	-		
COM60	0.047	0.998	0.059	0.997	_	-		
KAR40	0.068	0.994	0.044	0.998	_	-		
KAR50	0.160	0.984	0.035	0.997	_	-		
KAR60	0.143	0.990	0.039	0.997	_	-		
CAP40	0.283	0.942	0.061	0.996	_	-		
CAP50	0.078	0.992	0.050	0.996	_	-		
CAP60	0.054	0.992	0.055	0.996	_	-		
COM50S	0.079	0.988	0.044	0.999	0.025	0.999		
KAR50S	0.047	0.994	0.039	0.998	0.024	0.999		
CAP50S	0.090	0.991	0.035	0.999	-	-		

## 4.2. $\beta$ relaxation

The  $\beta$  relaxation occurred in the temperature range between  $-40 \,^{\circ}\text{C}$ and  $-70 \,^{\circ}\text{C}$  and shows activation energy of about 50–75 kJ/mol. Occurrence of  $\beta$  relaxation phenomena was already observed, at similar temperatures and with similar activation energy values, also through DSC in low molecular weight carbohydrates (Roos, 1993) and through DMA in frozen wheat doughs (Laaksonen & Roos, 2000; 2001b).

The most likely explanation of this process is that it corresponds to the glass transition and is linked to a modification in the mobility of large portions of the amylose and amylopectin molecules. Indeed, these latter systems show similar behaviour, in terms of temperature range and values of activation energy for the relaxation, as reported in the literature at the same frequency of 100 Hz (Butler & Cameron, 2000). Other studies (Roudaut, Maglione, Van Dusschoten, & Le Meste, 1999; Scandola, Ceccorulli, & Pizzoli, 1991) related this relaxation also to dextran or pullulan. This leads to the more general explanation of the rotational motion of the hydroxyl groups present on the  $C_2$ ,  $C_3$ , or  $C_6$  of the glucosidic units. It is also further confirmed by specific studies concerning the dielectric relaxations of glucose (Chan et al., 1986).

Possible connections between gluten sub-units and  $\beta$  relaxation can almost certainly be excluded, mainly based on previous studies, which revealed that gluten does not show any relaxation in the same temperature range with respect to the present work, but only at higher temperatures, with a peak at 27 °C (Roudaut, Maglione, & Le Meste, 1999).

The height of  $\tan(\delta)$  peak is significantly and highly correlated with the semolina protein and gluten amount (r > 0.71), and also with  $W_CP$  (r = 0.75), while the correlation with the water content presents a low and negative coefficient (r = -0.30). The positive and significant correlation with  $W_CP$  suggests that the two processes are largely explainable in the same way, i.e. with the bound water (also the other correlations are similar to those of the water crystallization). Since gluten is not detected, one can assume that this bound water involves carbohydrate molecules.



Fig. 5. Dielectric parameter  $tan(\delta)$  as a function of temperature for CAP50, KAR50, and COM50 samples at 100 Hz (a) and enlargement of the  $\beta$  and  $\gamma$  relaxations zone (b).

## Table 5

Dielectric tan( $\delta$ ) peak values of water crystallization (W<sub>C</sub>P),  $\beta$  relaxation ( $\beta$ P) and  $\gamma$  relaxation ( $\gamma$ P), and corresponding temperature for all the samples, measured at 100 Hz.

	W <sub>C</sub> P		βΡ		γP	γP		
Sample	Tan(δ)	Temp. [°C]	Tan (δ)	Temp. [°C]	Tan (δ)	Temp. [°C]		
COM40 COM50 COM60 KAR40 KAR50 KAR60 CAP40 CAP50 CAP60 COM50S	19.076 19.532 2.464 4.918 5.953 8.061 21.257 21.951 22.099 1.298	-10 -5 0 -5 -5 0 0 -5 -10	$\begin{array}{c} 1.170\\ 0.886\\ 0.308\\ 0.412\\ 0.356\\ 0.473\\ 1.142\\ 1.006\\ 0.949\\ 0.104 \end{array}$	$ \begin{array}{r} -60 \\ -45 \\ -55 \\ -50 \\ -50 \\ -40 \\ -40 \\ -40 \\ -40 \\ -65 \\ \end{array} $	0.465 0.586 0.302 0.410 0.496 0.588 0.472 0.552 0.609 0.287	$\begin{array}{c} -100 \\ -105 \\ -115 \\ -105 \\ -105 \\ -105 \\ -105 \\ -105 \\ -105 \\ -105 \\ -105 \\ -105 \end{array}$		
KAR50S CAP50S	9.368 22.383	$^{-5}_{-10}$	0.302 1.018	-70 -60	0.449 0.541	$-100 \\ -100$		

As activation energy only depends on the energy status of the target groups, the low and negative correlation (r = -0.31) observed with the water amount suggests that water can partially change the energy status of the target group (hydroxyl), and this could be explained with the possibility of interaction with water molecules (through hydrogen bonds). There is no clear explanation of the weak and negative correlation with the protein and gluten amount (r < -0.36) unless it is considered that, in wheat flour, the amount of damaged starch is usually negatively well correlated with the gluten amount (Barak, Mudgil, & Khatkar, 2013). Considering that the damaged starch is the main responsible of the water absorption, a hypothesis could be the greater difficulty in the rotation of the hydroxyl groups due to the lower water

relative availability with respect to the starch.

In the case of samples with added salt, the 20 °C temperature shifting in the  $\beta$  relaxation dielectric tan( $\delta$ ) peak at 100 Hz, that was observed for all the samples, could be linked to the NaCl ability to reduce the transition temperatures of the unfrozen solute, as stated in a previous work (Laaksonen & Roos, 2001a).

## 4.3. $\gamma$ relaxation

The  $\gamma$  relaxation occurred in the temperature range between -115 °C and -100 °C and shows activation energy of about 30-40 kJ/mol. The peak temperature corresponds to that shown in the literature (-112 °C) for wheat doughs at the same frequency of 100 Hz, and the same correspondence was found for the activation energy (31.5 kJ/mol) (Laaksonen & Roos, 2000). From this, the possibility that also gluten influences the  $\gamma$  relaxation process cannot be completely excluded, nor previous studies clearly assessing this aspect were found. However, other researchers (Butler & Cameron, 2000), testing only starch and not doughs, gave an explanation of this relaxation process, claiming that it is likely due to localized chain motions in starch molecules, probably enabled by the motion of the water molecules which are hydrogen-bonded to the amylose and amylopectin molecules, and that small-amplitude oscillations of the glucose rings about the  $(1-4)-\alpha$  and  $(1-6)-\alpha$  linkages are the most likely candidates. The similarity between the dielectric response in this work and ours makes the possibility of a gluten influence on the  $\gamma$  relaxation process very unlikely.

The height of the tan( $\delta$ ) peak presents the weakest correlation with the semolina protein and gluten amount (r > 0.32), and also with the water amount (r = 0.18). Correlation with  $W_CP$  is high (r = 0.71), significant, and similar to the one between  $\beta P$  and  $W_CP$ . However, the correlation between  $\gamma P$  and  $\beta P$  is much lower (r = 0.38) and not significant. This could be supported by the fact that the two processes are



Fig. 6. Tan( $\delta$ ) as a function of temperature for samples KAR50 and KAR50S at 100 Hz (a) and enlargement of the  $\beta$ ,  $\gamma$ , and i relaxations zone (b).



Fig. 7. Compliance against time comparisons among (a) CAP40 (blue), CAP50 (black), CAP60 (red) and CAP50S (green); (b) KAR40 (blue), KAR50 (black), KAR60 (red) and KAR50S (green); c) COM40 (blue), COM50 (black), COM60 (red) and COM50S (green); (d) CAP50 (black), KAR50 (blue) and COM50S (green); d) caP50 (black), KAR50 (blue) and COM50 (red). Data were acquired by creep tests applying a 50 Pa stress at 25 °C and modeled with Burgers model (black lines represent the model regression). (For interpretation of the references to colour in this figure legend, the reader is referred to the Web version of this article.)

#### Table 6

Parameters of the Burgers model estimated for the different samples.

Sample	J <sub>0</sub> (Pa <sup>-1</sup> )	$J_m$ (Pa <sup>-1</sup> )	τ (s)	$\eta_0$ (Pa • s)	Adj-R2
KAR40	$1.35 \cdot 10^{-5}$	$1.26 \cdot 10^{-5}$	27.01	$7.35 \cdot 10^{6}$	0.9997
KAR50	$3.65 \cdot 10^{-5}$	$2.72 \cdot 10^{-5}$	21.20	$3.04 \cdot 10^{6}$	0.9997
KAR60	$1.14 \cdot 10^{-4}$	$8.20 \cdot 10^{-5}$	22.91	$1.27 \cdot 10^{6}$	0.9995
KAR50S	$3.13 \cdot 10^{-5}$	$2.16 \cdot 10^{-5}$	24.20	$3.95 \cdot 10^{6}$	0.9997
CAP40	$1.01 \cdot 10^{-5}$	$9.32 \cdot 10^{-6}$	26.61	$8.95 \cdot 10^{6}$	0.9997
CAP50	$3.10 \cdot 10^{-5}$	$3.11 \cdot 10^{-5}$	24.79	$3.45 \cdot 10^{6}$	0.9995
CAP60	$9.42 \cdot 10^{-5}$	$8.18 \cdot 10^{-5}$	23.63	$1.14 \cdot 10^{6}$	0.9996
CAP50S	$2.26 \cdot 10^{-5}$	$1.88 \cdot 10^{-5}$	26.04	$4.86 \cdot 10^{6}$	0.9997
COM40	$1.53 \cdot 10^{-5}$	$1.43 \cdot 10^{-5}$	27.01	$6.47 \cdot 10^{6}$	0.9997
COM50	$4.42 \cdot 10^{-5}$	$3.05 \cdot 10^{-5}$	24.71	$3.24 \cdot 10^{6}$	0.9995
COM60	$1.20 \cdot 10^{-4}$	$8.63 \cdot 10^{-5}$	22.91	$1.20 \cdot 10^{6}$	0.9995
COM50S	$5.01 \cdot 10^{-5}$	$3.84 \cdot 10^{-5}$	23.45	$2.22 \cdot 10^{6}$	0.9996

related to the motions of different molecular structures. As  $W_C P$  is likely related to bound water and strongly correlated with  $\gamma P$ , this could confirm that both  $\beta$  and  $\gamma$  relaxations are related to different waterbound molecular structures.

 $\gamma E_a$  is substantially not correlated with the protein and gluten content (r < 0.12). The moderate and negative correlation (r = -0.47) with the water amount could mean, similarly to the case of  $\beta$  relaxation, that water molecules can, at least in part, make the target group oscillation easier by interacting with it.

An additional relaxation process in COM and KAR samples (which have the highest gluten index), when salt was added, occurred at slightly lower temperatures and with consistently higher activation energy (80–90 kJ/mol). According to the literature (Butler & Cameron, 2000), this additional phenomenon is likely due to a regime of lower hydration with respect to the main relaxation, which prevails when the hydration increases, making the secondary relaxation undetectable. This is compatible with the presence of salt, which competes for water molecules with the starch.

## Table 7

Correlation among the rheological, dielectric and composition parameters (<sup>a</sup> p-value  $\leq 0.01$ ; <sup>b</sup> 0.01 < p-value  $\leq 0.05$ ; <sup>c</sup> 0.05 < p-value  $\leq 0.10$  (WA:added water amount, PRO: protein content, GLU: gluten content, GI: gluten index,  $\beta E_a$ :  $\beta$  relaxation activation energy,  $\gamma E_a$ :  $\gamma$  relaxation activation energy,  $W_CP$ : water crystallization tan( $\delta$ ) peak height,  $\beta P$ :  $\beta$  relaxation tan( $\delta$ ) peak height,  $\gamma P$ :  $\gamma$  relaxation tan( $\delta$ ) peak height,  $J_0$ :instantaneous compliance,  $J_m$ : delayed compliance,  $\tau$ : delay time and  $\eta_0$ : steady-state viscosity).

	WA	PRO	GLU	GI	$\beta E_a$	γ E <sub>a</sub>	W <sub>C</sub> P	βP	γP	$J_0$	J <sub>m</sub>	τ	$\eta_0$
WA	1												
PRO	0	1											
GLU	0	0.9954 <sup>a</sup>	1										
GI	0	$-0.9705^{a}$	$-0.9891^{a}$	1									
β Ea	-0.3098	-0.3613	-0.3773	0.3951	1								
γ Εα	-0.4734	-0.1181	-0.0849	0.0324	0.2589	1							
$W_C P$	-0.1824	0.7722 <sup>a</sup>	0.7764 <sup>a</sup>	$-0.7690^{a}$	-0.2937	-0.2323	1						
$\beta P$	-0.2964	0.7177 <sup>a</sup>	$0.7350^{a}$	$-0.7483^{a}$	0.0070	-0.0300	0.7526 <sup>a</sup>	1					
$\gamma P$	0.1764	0.3197	0.3597	-0.4147	-0.4383	-0.3697	0.7133 <sup>a</sup>	0.3839	1				
$J_0$	0.9131 <sup>a</sup>	-0.1320	-0.1453	0.1631	-0.0942	-0.5684 <sup>c</sup>	-0.3273	-0.3099	0.0051	1			
$J_m$	0.9166 <sup>a</sup>	-0.0308	-0.0414	0.0572	-0.1569	$-0.5870^{b}$	-0.2461	-0.2171	0.05856	0.9883 <sup>a</sup>	1		
τ	$-0.7460^{a}$	0.3278	0.3210	-0.3047	0.4488	0.3301	0.4860	0.5408 <sup>c</sup>	0.0303	-0.6478 <sup>b</sup>	$-0.6203^{b}$	1	
$\eta_0$	$-0.9287^{a}$	0.1501	0.1649	-0.1848	0.3619	0.4687	0.3015	0.4588	-0.0727	$-0.8249^{a}$	$-0.8261^{a}$	0.8233 <sup>a</sup>	1

## 4.4. Relationship between dielectric and rheological response

Looking at correlation coefficients in Table 7, the first considerations should be done about the relationships between the Burgers model parameters.  $J_0$  and  $J_m$  are very strongly and significantly correlated (r = 0.99), meaning that they are representative of, and influenced by, the same conditions in the dough.  $\eta_0$  is significantly correlated with all the other Burgers parameters, even if a little more slightly ( $r \in [-0.83, -0.82]$ , while  $\tau$  shows weaker, but still significant correlations with  $J_0$  and  $J_m$  ( $r \in [-0.65, -0.62]$ ).

All the Burgers parameters are significantly correlated with the water content, with r = 0.91-0.92 for  $J_0$  and  $J_m$ , -0.92 for  $\eta_0$ , and -0.75 for  $\tau$ . However, no one of them is significantly correlated to protein and gluten amount, nor to the gluten index, and the correlation coefficients are in general low, especially for  $J_0$ ,  $J_m$  and  $\eta_0$ . This could lead to the erroneous conclusion that the semolina composition does not affect the creep response of the dough and that the water amount is the only parameter that influences the rheological behaviour. Actually, the composition alone cannot completely explain the rheological behaviour, but not even the water amount alone, because it is well known that the result depends on the complex interactions among all the dough components.

In effect, looking at the correlations between the Burgers parameters and the dielectric ones, the latter reveal to be much more informative about the effects of the molecular dynamics in the dough on its rheological response.  $J_0$  and  $J_m$  present a significant, negative, and stronger correlation with  $\gamma E_a$  ( $r \in [-0.58, -0.56]$ ) when compared to the semolina composition parameters. Correlations with  $W_CP$  and  $\beta P$  are weaker if compared to the previous one (respectively  $r \in [-0.33, -0.25]$ and  $r \in [-0.31, -0.22]$ ). However, they show to be higher than those with the semolina composition parameters.  $J_0$  and  $J_m$  are not correlated with  $\beta E_a$  ( $r \in [-0.16, -0.09]$ ), nor with  $\gamma P$  ( $r \in [-0.06, -0.01]$ ).  $\tau$  is significantly, even if not very strongly, correlated with  $\beta P$  (r = 0.54).  $\tau$ also presents a moderate correlation value (r = 0.49) with  $W_CP$ , and weaker correlations with  $\beta E_a$  and  $\gamma E_a$  (r = 0.45 and r = 0.33, respectively). No correlation was found between  $\tau$  and  $\gamma P$  (r = 0.03).  $\eta_0$  is the only Burgers parameter which does not present significant correlations with dielectric parameters. However, medium correlation coefficients were found with respect to  $\beta E_a$  and  $\gamma E_a$  (r = 0.36 and r = 0.47, respectively), to  $W_CP$  and  $\beta P$  (r = 0.30 and r = 0.46, respectively). No correlation was found between  $\eta_0$  and  $\gamma P$  (r = – 0.07).

In summary,  $\gamma E_a$  could give the most significant information on  $J_0$ and  $J_m$ ,  $\beta P$  is the most significant dielectric parameter with respect to  $\tau$ , while no dielectric parameter is suitable to give significant information on  $\eta_0$ .  $\gamma P$  revealed to be substantially uncorrelated with respect to all Burgers parameters, thus not informative on the dough rheological behaviour.

Looking at the supposed explanations of the dielectric relaxations, the starch hydration, at the glucose ring level, could be responsible, to a certain extent, of the dough compliance, as far as the water molecules are able to make the molecular movements easier, regardless the actual amount of bound water. On the other hand, the amount of hydroxyl groups in the glucosidic units of starch can act as a retardant of the dough deformation, as an increasing amount of them makes the rheological delay time increase. A similar effect is likely produced on the dough viscosity, even if the correlation with the dielectric parameters is not statistically significant.

#### 5. Conclusions

This work aimed to investigate the dielectric properties of semolina doughs and to find connections with their rheological properties. The goal was to develop a characterization technique able to give information on the process conditions of industrial bakery products and pasta concerning their quality properties in manufacturing and freezing processes.

It was observed that semolina dough displays three molecular relaxation phenomena: the water crystallization process,  $\beta$  and  $\gamma$  relaxations.

Water crystallization occurred in the temperature range between 0 and -10  $^{\circ}$ C. This process showed to be mostly related to the bound water in the system and to the interactions that it establishes with other molecules in the dough.

The  $\beta$  relaxation occurred in the temperature range between -40 °C and -70 °C and showed activation energy of about 50–75 kJ/mol. This process is probably associated with relaxations in glucosidic units, due to the rotational motion of the hydroxyl groups present on carbon atoms (C<sub>2</sub>, C<sub>3</sub>, or C<sub>6</sub>).

The  $\gamma$  relaxation occurred in the temperature range between -115 °C and -100 °C and showed activation energy of about 30–40 kJ/mol. This process is likely due to localized chain motions in starch molecules, probably small amplitude oscillations of the glucose rings about the (1–4)- $\alpha$  and (1–6)- $\alpha$  linkages enabled by hydrogen bound water.

When salt is added to the dough, it is possible to observe an additional relaxation process at slightly lower temperatures and with consistently higher activation energy (80–90 kJ/mol) than the  $\gamma$  one. This additional phenomenon is likely due to a regime of lower hydration compatible with the presence of salt. For the same reason, NaCl is also able to reduce the transition temperatures of the unfrozen solute, producing a temperature shifting in the  $\beta$  relaxation.

Concerning the rheological aspects, the Burgers parameters presented a strong correlation with the water amount, while they were very slightly correlated with the protein and gluten amount. On the other hand, much higher correlations were found between Burgers and most of the dielectric parameters, obtained at sub-zero temperatures, revealing that the latter can be very informative in explaining, to a certain extent, the rheological properties of the dough. Finally, no relationship between the  $\gamma$  relaxation tan( $\delta$ ) peak height and the dough compliance was found in this study.

## CRediT authorship contribution statement

Fabio Fanari: Methodology, Software, Investigation, Data curation, Writing – original draft, Visualization. Ciprian Iacob: Methodology, Validation, Data curation. Gianluca Carboni: Resources, Investigation. Francesco Desogus: Conceptualization, Methodology, Writing – review & editing, Funding acquisition. Massimiliano Grosso: Conceptualization, Software, Formal analysis, Project administration. Manfred Wilhelm: Resources, Supervision.

### Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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## Appendix A



**Fig. A.1.** mobility map and Arrhenius model regression with  $E_a$  values for samples with 40 wt% amount of water CAP40 (a), KAR40 (b), COM40 (c), 60 wt% amount of water CAP60 (d), KAR60 (e), COM60 (f), 50 wt% amount of water and 1.5 wt% amount of salt CAP50S (g), KAR50S (h), COM50S (i) samples. Triangles represent tan( $\delta$ ) while green stars  $\sigma_0$  values. The red dashed line highlights the crystallization temperature, in correspondence of which a drop in tan( $\delta$ ) is observable



**Fig. A.2.**  $tan(\delta)$  as a function of temperature (a) and enlargement of the  $\beta$  and  $\gamma$  relaxations zone (b) for CAP40, CAP50, and CAP60 samples;  $tan(\delta)$  as a function of temperature (c) and enlargement of the  $\beta$  and  $\gamma$  relaxations zone (d) for KAR40, KAR50, and KAR60 samples;  $tan(\delta)$  as a function of temperature (e) and enlargement of the  $\beta$  and  $\gamma$  relaxations zone (f) for COM40, COM50, and COM60 samples. All data reported were measured at the frequency of 100 Hz



Fig. A.3.  $tan(\delta)$  as a function of temperature (a) and enlargement of the  $\beta$  and  $\gamma$  relaxations zone (b) for CAP50 and CAP50S samples;  $tan(\delta)$  as a function of temperature (c) and enlargement of the  $\beta$  and  $\gamma$  relaxations zone (d) for COM50 and COM50S samples. All data reported were measured at the frequency of 100 Hz

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