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Effect of ohmic heating on the structure and properties of flexible multilayer packaging

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Abstract: Food-packaging-processing interactions define packaging materials' performance properties and product quality. This study evaluated the effect of ohmic heating (OH) processing and different food simulants on the properties of four multilayer flexible packaging materials (PET_{met}/PE , PET_{met}/PP , $PET/A1/PE$, and $PET/A1/PA/PP$). OH treatment was applied to the sealed packages containing the simulants using a voltage gradient of 3.7 V/cm at a frequency of 20 kHz, resulting in a thermal process of at 80 °C for 1 min. The structure and performance of the different packages were then evaluated. The materials did not show changes in chemical groups nor thermal properties. However, the simulant-packaging-processing interaction resulted in changes in crystallinity, morphology, mechanical and barrier properties (water and oxygen), especially for metallized films in contact with acidic food simulants. The results indicate that although OH resulted in changes in packaging materials, these materials can be used under the conditions applied in this study.

Keywords: emerging technology; food packaging; tensile strength; heat sealing strength; barrier properties; food thermal process

1. Introduction

Industrial processes demand around 1/3 of global energy, generating significant potential to reduce and/or manage energy consumption (Napp et al., 2014; Sakr & Liu, 2014). Particularly in the food industry, energy is necessary to increase the temperature of the process, such as generating steam, hot water, heating food, among others. In such processes, heat can be generated directly or indirectly transferring heat to the material, and be produced from fuels, electric energy and others, (Sakr & Liu, 2014). The quality of a product can be determined by the performance of a heating system (Van Der Gotot et al., 2016). However, efficient systems to manufacture products with the required quality and at the lowest cost are necessary.

Thus, the development of new food processes, with the aim of ensuring greater food quality and safety, but also with emphasis on energy efficiency, lower water consumption and reduction of gas and effluent emissions, will reduce food costs production and improve sustainability in the food chain (Fasolin et al., 2019; Misra et al., 2017; Pereira & Vicente, 2010).

Ohmic heating (OH) is a promise technology as it can process food in a short period of time and reduce the energy input (Sakr & Liu, 2014; Yildiz-Turp et al., 2013). Different mechanisms applied during this method reduce the resistance to the heat and mass transfer, compared to conventional methods (Misra et al., 2017; Pereira & Vicente, 2010). OH is defined as a process in which electrical current (usually in alternating mode) passes through conductive materials resulting in internal heat dissipation (Gavahian et al., 2019; Knirsch et al., 2010). The application of the electrical field is made through two electrodes in contact with food, and generally from low to moderate intensity (<1000 V/cm), with a frequency of 50 Hz to 20 kHz (Fasolin et al., 2019). Heating occurs in the form of an internal transformation of energy (from electrical to thermal) within the material, in a very rapid and volumetric heating (Fasolin et al., 2019; Knirsch et al., 2010), and the amount of heat is directly related to the voltage gradient and electrical conductivity of the product (Yildiz-Turp et al., 2013). However, for processing to occur effectively, the materials must contain sufficient water and electrolytes to allow the passage of electrical current (Knirsch et al., 2010).

This technology is bringing a new paradigm to the thermal processing of foods, reducing excessive thermal load (it does not depend on conduction and convection heat transport mechanisms) and benefiting from the non-thermal effects of electric fields (Fasolin et al., 2019). In addition to rapid heating rates, OH is versatile, allowing to process of various food systems, whether particulate or not, which can lead to increased quality of the final product and increasing its added value. OH has numerous applications, including blanching, evaporation, dehydration, fermentation, extraction, sterilization and heating of foods (Makroo et al., 2020).

To minimize the risk of post-process contamination and avoid expensive aseptic packaging lines, packaging before OH treatment can be considered. Developments for processing OH packaged foods are focused on packaging materials with incorporated aluminum electrodes and conductive packaging materials (Kamonpatana, 2018), with main focus on the food itself (and not the packaging performance). A V-shaped PET/Al/PE (PET: polyethylene terephthalate, Al: aluminum foil and PE: polyethylene) packaging with two metal sheets used as electrodes to heat food was developed, however, heating was not uniform inside the packaging. To facilitate uniform heating, the V-shaped packaging has been redesigned to a rectangular shape with flat sides. A temperature distribution study on a 227 g package showed the presence of cold regions on the electrodeless sides due to current channeling through a hotter, more conductive center (Somavat et al., 2012). A uniform temperature distribution was obtained with the use of two heating strips installed on the external side of the materials (Jun & Sastry, 2005, 2007) Using this approach, treatments at 121°C for 6 min and 130°C for 1 min was adequate to inactivate 5 log *Geobacillus stearothermophilus* spores, confirming that sterilization was achieved (Somavat et al., 2012).

On another study, carbon black was incorporated into polypropylene (PP) film due to its conductive properties. The conductive film was connected to the PP film on the left and right sides of the bag to form a rectangular shaped conductive package, liable to be subjected to OH. However, to eliminate the cold zone at the edges and corners of the package, the system required the use of a means to carry the current and channel the electrode current through the package, where the electrical conductivity of the medium was greater than that of the food inside the packaging (Kanogchaipramot et al., 2016). Subsequently, this conductive packaging was used to evaluate the effect of OH in the processing of packaged orange juice. The temperature in the center of the conductive package containing orange juice was 84 °C when a voltage of 300 V was applied for 300 s. A 6 log inactivation of *Escherichia coli* was achieved, indicating that pasteurization of fruit juice using OH through conductive packaging was effective.

Few studies are available in the literature regarding *in-situ* application of OH. In terms of packaging, studies are focused on evaluating the materials' electrical conductivity and heat distribution. To the best of our knowledge, no studies have been carried out that evaluated the effect of OH on the structure and properties of packaging materials, as this technology has great potential for pasteurization and/or sterilization of foods. Still in this context, evaluating the effect of OH on the properties of packaging materials will result in a diagnosis for the application of this technology to process packaged foods. Therefore, the objective of this research was to evaluate the effect of OH processing on the structure and properties of different real packaging materials in the pasteurization range. The innovation lies in the description and understanding of the mechanism of food-packaging-process interaction of this promising emerging technology, paving the application of this technology in the food industry. Finally, for this research, commercial packaging materials widely used for thermos-processed foods and beverages were selected to understand whether these materials are resistant to OH processing.

2. Materials and methods

2.1. Materials

Four multilayer packaging materials (Finepack, São Paulo, Brazil) that are commonly used to package thermos-processed foods were evaluated, as described below:

- PET_{met}/PE $(11/44 \text{ }\mu\text{m})$
- PET_{met}/PP $(12/39 \text{ }\mu\text{m})$
- PET/Al/PE $(12/7/45 \text{ }\mu\text{m})$
- PET/Al/PA/PP $(12/7/15/38 \mu m)$

In which PET: polyethylene terephthalate, met: metallized, PE: polyethylene, PP: polypropylene, Al: aluminum foil and PA: polyamide.

2.2.Packaging sample preparation, model foods and ohmic heating processing

The different samples with dimensions of 80 mm \times 220 mm were filled with 120 mL of two different food simulants / model foods. The rectangular shape of the packaging materials was selected based on the best distribution of electrical potential and, consequently, the uniformity of heating, as described in the literature (Somavat et al., 2012).

The two evaluated food simulants were: 3% (v/v) acetic acid solution and 0.1% sodium chloride solution (w/v), referred to as AA and NaCl, respectively. Thus, food simulants representing acidic and low-acid food products were considered. The food simulants AA and NaCl showed electrical conductivity of 1597 ± 5 µs cm⁻¹ and 1613 ± 45 µs cm⁻¹ at 23 °C, respectively.

The OH was performed using two stainless steel electrodes placed on the lateral ends of the inner part of each package. Each package was then hermetically sealed, allowing the external part of the electrodes to be connected to the power system. Through this apparatus, it was possible to apply the OH effect inside the packaging. The supplied voltage, and consequently the temperature, were controlled through the use of a function generator (1 Hz– 25 MHz and 1–10 V; Agilent 33220A, Penang, Malaysia) connected to an amplifier system (4505 Precision Power Amplifier; Miko-Kings Instruments Ltd, Hong Kong, China). The applied electrical frequency was 20 kHz and an electric field intensity of *ca.* 3.7 V cm−1 was applied until reaching the target treatment temperature of 80 °C. The temperature was measured with a type K thermocouple, placed inside the packaging, in the geometric center of the sample volume. Upon reaching the target treatment temperature, the samples were maintained at 80 °C for 1 min - conditions similar to those applied to pasteurize carrot and apple juices (Abedelmaksoud et al., 2018; Mannozzi et al., 2018, 2019).

For each packaging sample and each food simulant, 10 treatments were carried out. The treatments performed were described as follows: control (non-processed samples), OH-AA and OH-NaCl. The different samples were characterized in terms of morphology, structure, thermal, mechanical and barrier properties, according to the methods described below.

2.3.Characterization of packaging materials

2.3.1. Fourier-transform infrared (FT-IR) spectroscopy

FT-IR analyzes was performed on a Fourier transform infrared spectrometer (Bruker, Alpha II, Billerica, USA) equipped with attenuated total reflectance (ATR). For all samples, the inner layers were directly analyzed with a spectral resolution of 4 cm⁻¹, 24 scans in the range 4000–600 cm⁻¹ (ASTM-E573-01, 2021; ASTM-E1252-98, 2021). The inner layers were selected based on direct contact with the food simulants and also on the configuration of the electrodes, that is, inside the packaging. Three spectra were recorded at different locations on the film for each sample to estimate the average film inhomogeneous potential and measurement variability.

2.3.2. Differential Scanning Calorimetry (DSC)

The thermal behavior of the packaging samples was evaluated using a differential scanning calorimeter (DSC 6000, Perkin Elmer, Waltham, USA). About 5 mg were weighed and placed in aluminum containers. The test temperature was from 40 $^{\circ}$ C to 300 $^{\circ}$ C at a fixed rate of 10 °C/min and pure nitrogen gas flow of 20 mL/min (ASTM-D3418-21, 2021). The melting temperature (T_m) and enthalpy of melting (ΔH) were estimated based on the DSC thermograms. The results were the average of three replications.

2.3.3. X-ray diffraction (XRD)

X-ray diffraction analyzes were performed on an XRD equipment (Bruker D8, Odelzhausen, Germany) with Cu-Kα radiation ($λ = 1.54056$ Å, 40 kV, 40 mA at a scan rate of 0.0333°/s). Measurements were obtained with steps of 0.04° from 10° to 30° (2θ) and an acquisition time of 1.2 s. The samples were cut with dimensions of 4 cm x 3cm and the reading was taken from the internal layer, that is, the layer in direct contact with the food simulants and also according to the configuration of the applied processing, that is, with the electrodes on the inside of the packaging. The percentage of overall crystallinity of the films was estimated by the ratio between the area under the peaks (crystalline regions) and the total area (crystalline and amorphous regions) through deconvolution of the peaks using the free software Fityk. The areas of the curves were determined by numerical integration.

2.3.4. Scanning electron microscopy (SEM)

Morphological analysis of the film surface was performed in an ultra-high resolution Field Emission Gun Scanning Electron Microscopy (FEG-SEM) (NOVA 200 Nano SEM, FEI Company). Topographic images were obtained with a Secondary electron detector at an acceleration voltage of 10 kV. Before morphological analyses samples were covered with a thin film (5 nm) of Au-Pd (80-20 weight %), in a high-resolution sputter coater (208HR Cressington Company), coupled to a MTM-20 Cressington High Resolution Thickness Controller. SEM images were analyzed at 5000× magnification. Similar to FT-IR and XRD analysis, the inner layer was selected to obtain the SEM images.

2.3.5. Mechanical properties

Tensile strength and seal strength tests were carried out at 23 ± 2 °C and 50 ± 5 % RH after conditioning the samples for a period of at least 48 h under the same conditions. Samples measuring 25.4 mm wide were used and tests were carried out in the machine direction (MD) and in the transverse direction (TD) of the material. All tests were performed on a universal testing machine (Instron, 5966-E2, Norwood, USA). Tensile strength was determined using a 100 N load cell with a test speed of 50 mm min-1 with an initial distance of 40 mm for the MD and 100 mm for the TD. (ASTM-D882, 2018). The seal strength was determined using a 1 kN

load cell at a speed of 300 mm min⁻¹ and an initial distance of 10 mm. (ASTM-F88/F88M, 2021). The tests were carried out in five replications.

2.3.6. Barrier properties

2.3.6.1. Water vapor transmission rate

The water vapor transmission rate (WVTR) was determined using a PERMATRAN equipment (3/34 G, MOCON, Minneapolis, USA) with an infrared sensor device. The test was carried out at 38 °C and 90% RH with a permeation area of 5 cm². WVTR tests were carried out from the outer layer to the inner layer (ASTM-F1249, 2020).

2.3.6.2. Oxygen transmission rate

The oxygen transmission rate (OTR) was determined using an OXTRAN equipment (2/22 H, MOCON, Minneapolis, USA). The outer side of the sample was placed in contact with the permeating gas (100% O_2). The test was carried out at 23 °C dry with a permeation area of 5 cm². The reading has been corrected for a permeant gas partial pressure gradient of 1 atm. OTR tests were carried out from the outer layer to the inner layer (ASTM-D3985, 2017).

2.4.Statistical analysis

The results were presented as mean \pm standard deviation and statistically evaluated by analysis of variance (ANOVA) and Tukey's test to compare mean values $(p<0.05)$.

3. Results and discussion

3.1. Fourier-transform infrared (FT-IR) spectroscopy

The FT-IR spectra of the inner layers of the four films show some characteristic absorption bands for polyethylene - Figure 1(a) and (c) - and polypropylene - Figure 1(b) and (d) - films, which are consistent with those reported in the literature (Alaburdaite & Krylova,

2023; Wang & Ajji, 2022). Regarding the FT-IR spectra of the PE films, the characteristic peaks were attributed to the asymmetric and symmetric stretching vibrations of $CH₂$ (2,915 and 2,849) cm^{-1}), respectively, to the C–C stretching vibration (1,646 cm⁻¹), to CH deformation vibrations in CH₂ (1,471 cm⁻¹), to bending vibrations in CH₃ (1,375 cm⁻¹) and to C-C equilibrium vibrations in CH² (717 cm−1) (De Geyter et al., 2008; Kochetov et al., 2017; Marangoni Júnior et al., 2023; Turriziani, Vieira, et al., 2023). For PP films, the characteristic peaks are attributed to asymmetric and symmetric stretching vibrations in CH₃ and CH₂ (2,951 to 2,838 cm⁻¹), asymmetric deformation vibrations of CH₃ or scissor vibrations of CH₂ (1,456 to 1,376 cm⁻¹), to CH₃ symmetric strain vibrations $(1,359 \text{ cm}^{-1})$ and numerous small peaks in the range referring to C–C asymmetric stretching, CH³ asymmetric rocking, C–H vibrations, C–C asymmetric stretching vibrations, and to $CH₂$ oscillation vibrations (1,300 to 700 cm⁻¹) (Alaburdaitė & Krylova, 2023; Morent et al., 2008; Smith, 2021). FT-IR is one of the most common and economical methods for analyzing the chemical structure of materials. Thus, it was used to evaluate whether different food simulants and OH would have any influence on the chemical structure of the polymers that make up the internal layer of the studied packaging materials. No changes were observed in the PE and PP spectra after contact with food simulants and OH in relation to untreated films, suggesting that the applied conditions did not modify the chemical nature of PE and PP.

Figura 1. FT-IR spectra of inner layer of different packaging materials processed by ohmic heating. (a): PET_{met}/PE , (b): PET_{met}/PP , (c): $PET/A1/PE$ and (d): $PET/A1/PA/PP$.

3.2.Differential Scanning Calorimetry (DSC)

Figure 2 and Table 1 show the thermal profiles and melting temperatures (*Tm*) of each polymeric material of the multilayer structures, as well as the total enthalpies of melting (Δ*H*) of the entire multilayer structure before and after contact with the different food simulants followed by OH.

Tm: melting temperature and Δ*H**: total enthalpy of melting.

Values referring to the mean of three repetitions ± standard deviation.

^{a,b,c} means followed by the same letter in the line do not differ at the 95% confidence level ($p <$

0.05).

Figure 2. DSC of different packaging materials processed by ohmic heating. (a): PET_{met}/PE, (b): PET_{met}/PP , (c): $PET/A1/PE$ and (d): $PET/A1/PA/PP$.

For the PET_{met}/PE and $PET/AI/PE$ samples, three melting peaks were observed, with the 1st peak corresponding to LDPE, the 2nd peak corresponding to LLDPE and the 3rd peak corresponding to PET, with T_m values varying between 110.2 °C and 112.3 °C, 117.1 °C to 117.8 ºC and 232.3 ºC to 241.1 ºC, respectively. These ranges are similar to those reported in the literature for LDPE, LLDPE (Benítez et al., 2013; Marangoni Júnior et al., 2023) and PET (Marangoni Júnior, Oliveira, Bócoli, et al., 2020; Marangoni Júnior, Oliveira, Dantas, et al., 2020). For the PET_{met}/PP samples, two melting peaks were observed, where the 1st represents PP and the 2nd represents PET. For the PET/Al/PA/PP samples, 3 melting peaks were found, the 1st referring to PP, the 2nd referring to PA and the 3rd to PET. The PET T_m values were close to those described for the PET_{met}/PE and $PET/A1/PE$ samples. Regarding the T_m of PP and PA, the values were 153.1 ºC to 155.4 ºC and 207.2 ºC to 214.5 ºC, respectively. Values close to

those reported in the literature for PP (Sanetuntikul et al., 2023) and PA (Marangoni Júnior, Oliveira, Dantas, et al., 2020). For the four different multilayer packaging materials, *T^m* and Δ*H* were not significantly influenced by the food simulant in contact with the material and by OH, showing that the materials are thermally stable for the conditions in which they were subjected.

3.3.X-ray diffraction (XRD)

The X-ray diffraction patterns of the different samples before and after OH are shown in Figure 3. For the metallized films (PET_{met}/PE and PET_{met}/PP) a crystalline peak is observed at the angular position of $2\theta = 26^\circ$ - Figures 3 (a) and (b). These peaks are attributed to the semi-crystalline structure of PET (Dasdemir et al., 2012). The peaks at approximately $2\theta = 21^{\circ}$ correspond to PE (Figure 3(a)) (Turriziani, Perez, et al., 2023) and the less intense peaks between $2\theta = 15^{\circ}$ and 21° correspond to PP (Figure 3(b)) (Yetgin, 2019). For the PET/Al/PE sample, the PE crystalline peak is also observed at approximately $2\theta = 21^{\circ}$ (Figure 3(c)) and for the PET/Al/PA/PP sample in addition to the PP peak at $2\theta = 15^{\circ}$ a more intense peak is observed at approximately $2\theta = 24^{\circ}$ referring to PA (Figure 3(d)) (Bhunia et al., 2016; Marangoni Júnior et al., 2023).

The overall crystallinity of PET_{met}/PE films processed by OH was higher compared to the control, suggesting that OH induced crystallization of this material. This increase can be attributed mainly to PET, as it is susceptible to the phenomenon of chemicrystallization, where hydrolytic degradation of the polymer occurs at temperatures above the glass transition of PET (65–75 °C). During this process, the amorphous region undergoes chain scission to produce smaller chain fragments with greater mobility, which aids in crystallization (Bhunia et al., 2016; Sammon et al., 2000). On the other hand, the overall crystallinity of the PET_{met}/PP films was lower after OH, where possibly high temperature processing led to a fragmentation of the crystalline structure that was ordered differently upon cooling. Similar behavior was also observed for multilayer films with PP layer processed by conventional heating and microwaveassisted thermal sterilization (Bhunia et al., 2016; Dhawan et al., 2014). The same behavior was observed for PET/Al/PA/PP films, which may also be the result of exposure of the film's hydrophilic polymer (PA) in an environment with high humidity, resulting in the plasticization of the PA and consequently causing some distortions in the crystalline structure of the polymer (Miri et al., 2009; Parodi et al., 2017). Finally, in PET/Al/PE samples almost no change in crystallinity was observed after OH, with values varying between 20.6% and 21.6%. In this sense, the main hypothesis is that changes must also have occurred in the crystalline rearrangement of this material during high-temperature processing. However, after cooling, this rearrangement returned to its initial conformation, thus minimizing the impact on crystallinity.

Figure 3. X-ray diffraction patterns of different packaging materials processed by ohmic heating. (a): PET_{met}/PE , (b): PET_{met}/PP , (c): $PET/AI/PE$ and (d): $PET/AI/PA/PP$.

Scanning electron microscopy was used to visualize the surface morphologies of the films' inner layers before and after OH, as shown in Figure 4.

Figure 4. Scanning electron microscopy of the inner layers surface of different packaging

and i): PET/Al/PE and (j, k and l): PET/Al/PA/PP. The red arrows indicate changes on the film surface.

The PET_{met}/PE sample showed a rough surface - Figures 4 (a), (b) and (c) - similar to that found for the PET/Al/PE samples - Figures 4 (g), (h) and (i). In relation to the PET_{met}/PP and PET/Al/PA/PP samples, the surfaces were more homogeneous and had a smooth appearance - Figures 4 (d), (e) and (f)) and Figures 4 (j), (k) and (l) - respectively. This difference between the materials is associated with the polymer of the inner layer being different, where PE has greater roughness compared to PP. It is observed that after OH the PETmet/PE sample showed small inclusions that differ from the other samples (Figure 4(b)) and in the PET_{met}/PP sample some holes were detected (Figure 4(e)), which is associated with the food simulant-packaging-processing interaction, mainly considering that the acidic food simulant is considered the most critical simulant for metallized packaging. In relation to films laminated with aluminum foil (Al) treated by OH, it is also possible to observe some inclusions and some typical defects related to bending and/or crushing of the aluminum foil, but without breaking the aluminum layer. The defects caused by processing and interaction with food simulants are not perceptible to the human eye. However, as it is necessary to assess whether they cause any damage to the material barrier that makes the use of these packaging in this application unfeasible. This approach is presented in section 3.6.

3.5.Mechanical properties

The tensile strength (TS) and elongation at break (EB) results in the machine direction (MD) and transverse direction (TD) of the different samples before and after OH are presented in Figure 5. The tests were carried out in two different directions of the material (MD and TD), as in general the mechanical properties are influenced by the orientation of the molecules in the manufacture of the film, that is, they differ in relation to the direction of the material. The TS

values in the MD and TD showed no significant difference between the processed and unprocessed samples, with the exception of the $\text{PET}_{\text{met}}/\text{PE}$ sample which showed a significant reduction in TS in TD (Figure 5(a)). Regarding EB, OH did not result in significant changes in both directions of the material when compared to the control, an exception was for the PET/Al/PA/PP film, which showed a significant reduction in EB in TD (Figure 5 (H)). These small changes in TS and EB may possibly be associated with changes in the degree of crystallinity of the polymers after OH. However, this behavior did not present a trend for all samples and as the changes were punctual and minimal, it can be concluded that OH in the applied conditions did not result in a negative impact on the tensile properties of the materials studied – a positive result.

The multilayer PET_{met}/PE and $PET/AI/PE$ materials presented maximum seal strength values equal to or greater than those of materials composed of PET_{met}/PP and $PET/AI/PA/PP$. this result is attributed to the excellent characteristics sealing of the PE layer. For metallized films (PET_{met}/PE and PET_{met}/PP) OH did not result in significant changes in seal strength, regardless of the food simulant in contact with the material (Figure 6(a) and (b)). However, for the PET/Al/PE film, OH resulted in a significant increase in the maximum seal strength in MD (Figure 6(c)), which may be due to overheating of the film due to of the presence of aluminum foil in the multilayer structure, as highlighted by Kamonpatana (2018) and Kanogchaipramot et al. (2016). Thus, the hypothesis is that this overheating possibly led to an increase in the temperature of the PE film, reaching close to the PE melting temperature (Table 1), favoring the fusion of the sealing layer and consequently resulting in more resistant bonds between the layers interfaces of the sealing layers when cooling the polymer. In principle, this result is not negative. On the other hand, the PET/Al/PA/PP film processed by OH showed a tendency to reduce the maximum seal strength values in MD in relation to the control (Figure 6(d)), possibly due to the increase in pressure inside the packaging. Furthermore, this reduction may be based on the hypothesis that the presence of a polar polymer (PA) in the structure, which may have

favored the absorption of food simulants and consequently had an impact on the sealing layer (Marangoni Júnior et al., 2023). However, no leakage was observed in the packaging sealing region after OH, thus ensuring the integrity of the packaging. It should also be noted that even though there is a reduction in the seal strength of the films, these values are within those recommended for sterilizable bags, as described by (Marangoni Júnior, Oliveira, Dantas, et al., 2020).

Figure 5. Tensile strength (a)(c)(e)(g) and elongation at break (b)(d)(f)(h) of different packaging materials processed by ohmic heating. (a)(b): PET_{met}/PE , (c)(d): PET_{met}/PP , (e)(f):

PET/Al/PE, (g)(h): PET/Al/PA/PP, MD: machine direction and TD: transverse direction. Lowercase letters (MD) and uppercase letters (TD) do not differ at the 95% confidence level (*p*< 0.05).

Figure 6. Maximum seal strength of different packaging materials processed by ohmic heating. (a): PET_{met}/PE , (b): PET_{met}/PP , (c): $PET/A1/PE$, (d): $PET/A1/PA/PP$, MD: machine direction and TD: transverse direction. Lowercase letters (MD) and uppercase letters (TD) do not differ at the 95% confidence level ($p < 0.05$).

3.6.Barrier properties

The WVTR and OTR results of the different films before and after OH are presented in Table 2. The WVTR of the PET_{met}/PE and PET_{met}/PP control metallized films were 1.79 ± 0.23 g_{water}.m⁻².day⁻¹ and 1.52 ± 0.21 g_{water}.m⁻².day⁻¹, respectively. Regarding OTR, the values were

 5.45 ± 0.46 mL(step, m⁻².day⁻¹ and 5.94 ± 0.67 mL(step, m⁻².day⁻¹, respectively. For PET/Al/PE and PET/Al/PA/PP films, the WVTR and OTR were below the quantification limit of the equipment used. These results are similar to those previously reported for metallized and aluminum foil laminated films (Dantas et al., 2022; Marangoni Júnior, Alves, et al., 2020; Marangoni Júnior et al., 2018). The greater barrier to oxygen and water vapor of films laminated with aluminum foil in relation to metallized films is attributed to the excellent barrier properties of the aluminum foil and the absence of microholes in the Al layer, as observed in Figures 4 (g) and (i) .

Packaging material	$WVTR (gwater.m-2.day-1)$ 38 °C/90% RH			OTR $(mL(STP).m-2.day-1)$ $23^{\circ}C$		
	Control	OH-AA	OH-NaCl	Control	OH-AA	OH-NaCl
$PET_{met}/PE*$	$1.79 + 0.23^b$	$5.22 + 1.19^a$	2.77 ± 0.37^b	$5.45 + 0.46^b$	$21.28 + 2.72^a$	$7.46 + 0.83^b$
$PET_{met}/PP*$	$1.52 + 0.21^a$	$3.69 + 0.93^{\circ}$	$3.48 + 1.25^{\circ}$	$5.94 + 0.67^b$	$9.70 + 1.88^a$	$5.21 + 0.49^b$
PET/Al/PE**	$\leq 0.01^{(1)}$	$\leq 0.01^{(1)}$	$\leq 0.01^{(1)}$	$\leq 0.05(1)$	$\leq 0.05(1)$	$\leq 0.05(1)$
PET/Al/PA/PP**	$\leq 0.01^{(1)}$	$\leq 0.01^{(1)}$	$\leq 0.01^{(1)}$	$\leq 0.05(1)$	$\leq 0.05^{(1)}$	$\leq 0.05(1)$

Table 2. WVTR and OTR of different packaging materials processed by ohmic heating.

Values referring to the mean of *three and **two repetitions \pm standard deviation.

 (1) Results below the quantification limit of equipment.

^{a,b,c} means followed by the same letter in the line do not differ at the 95% confidence level ($p <$ 0.05).

It is observed that OH in synergy with the acidic food simulant resulted in a significant increase in the WVTR of the PET_{met}/PE film and the OTR of the PET_{met}/PE and PET_{met}/PP films. These results are attributed to the defects observed in the SEM images (Figure 4) and may also have some association with changes in the overall crystallinity of the films. It should be noted that the acidic food simulant is the most critical simulant in terms of interaction with metallized packaging materials. Added to this, the high process temperature (80 ºC) possibly

favored interactions between the food simulant and the packaging material, which consequently had an impact on the film barrier. However, this behavior does not prevent the use of the material for this application, if an assessment is made regarding the impact that this loss of barrier will have on the stability of acidic foods treated by OH during storage. This is to ensure that the materials do not compromise the product's quality. In relation to PET/Al/PE and PET/Al/PA/PP films, OH and the different simulants did not influence the barrier of the materials, which although presented morphological defects, there was no rupture of the aluminum foil, which guaranteed maintenance excellent barrier properties.

4. Conclusions and future research outlook

The different multilayer packaging materials were processed by OH at pasteurization conditions in contact with different food simulants. The materials proved to be stable in terms of chemical structure and thermal properties. On the other hand, the OH associated with different food simulants resulted in an increase in the overall crystallinity of the PET_{met}/PE film and a reduction in the crystallinity of the PET_{met}/PP and $PET/A1/PA/PP$ films. Added to this, OH resulted in changes in the surface of the films, especially in metallized films in contact with the acidic food simulant. Regarding mechanical properties, OH resulted in a small and punctual reduction in the tensile strength of the PET_{met}/PE film and in the elongation at break of the PET/Al/PA/PP film. Furthermore, there was an increase in the heat-sealing tensile strength of the PET/Al/PE film after OH treatment and a reduction for the PET/Al/PA/PP film. The metallized films in contact with the acidic food simulant followed by OH treatment have an increase in water vapor permeability rate and oxygen permeability rate. Finally, although interactions among food, packaging, and processing have caused modifications to the packaging materials under study, they remain suitable for this application.

Future studies are suggested covering different OH processing conditions, examining other packaging materials and evaluating the impact of post-packaging OH on the quality of different food products, aiming to expand knowledge about food-packaging-processing relationships.

Declaration of 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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CRediT authorship contribution statement

Luís Marangoni Júnior: Conceptualization, Validation, Formal analysis, Methodology, Investigation, Data curation, Formal analysis, Resources, Funding acquisition, Writing – original draft. *Rui M. Rodrigues:* Conceptualization, Validation, Methodology, Formal analysis, Writing - Review & Editing. *Ricardo N. Pereira:* Conceptualization, Validation, Methodology, Formal analysis, Writing - Review & Editing. *Pedro Esteves Duarte* *Augusto:* Formal analysis, Writing - Review & Editing. *Danielle Ito:* Investigation, Methodology, Writing - Review & Editing. *Fábio Teixeira Gomes:* Investigation, Methodology, Writing - Review & Editing. *Marisa Padula:* Conceptualization, Validation, Supervision, Resources, Funding acquisition, Writing – review & editing. *António A. Vicente:* Conceptualization, Validation, Supervision, Resources, Funding acquisition, Writing – review & editing.

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