Monitoring the Shelf-Life of Minimally Processed Fresh-Cut Apple Slices By Physical–Chemical Analysis and Electronic Nose

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Abstract

Fresh-cut apples, in slices or in cubes, are minimally processed products, which are currently collecting a great interest by fruit marketers for their promising diffusion. Their shelf life, from a microbiological point of view, has been fixed about 2 or 3 weeks under refrigeration. However in a few days they undergo biochemical degradations with production of off-flavors and texture breakdown. In this work, the change of aromatic fingerprint of apple slices packaged in air and in a modified atmosphere (with 100% N2) was measured, by using a commercial electronic nose. The obtained data were also compared with sensory evaluation of judge’s panel. Moreover, quality parameters such as total acidity, total soluble solids and firmness were determined at different sampling times (0, 4, 8 and 12 days).

The data show that the electronic nose is able to discriminate between the two different storage conditions applied: the multivariate analysis, Principal Component Analysis, presents clearly differences among the four sampling times when the apple slices are stored in air and in N2.

Our results indicate that the electronic nose can be considered a valid supplementary tool to human sensory panel assessment especially in food quality safety and control and it can be a simple, objective and rapid method to control the food quality during the storage.

Keywords: Fresh-cut apples; Shelf-life; Electronic nose; Sensory analysis; Food storage; Minimally processed fruit

Introduction

Recently the demand for minimally processed fruits and vegetables is increasing. The ready-to-eat fresh-cut products are one of the major growing segments in food markets. The rapid growth is due to the new lifestyles and to the health-consciousness of the consumers [1]. In fact, they request fresh-like processed products with high quality attributes (such as appearance, texture, and flavor) similar to those of the raw products [2] to satisfy the daily needs of antioxidants, minerals and dietary fibers. The ready-to-eat fresh-cut products are slightly processed products, retaining intact the positive characteristics of the fresh fruits and vegetables. Minimal processing has been defined as a combination of procedures, such as washing, peeling, slicing or chopping, not affecting the fresh-like quality and flavor of the food [3]. However, fresh-cut fruits are more challenging to obtain than other processed products for the difficulties in preserving their fresh-like properties during prolonged periods. The tissue integrity of fruits, in fact, can be easily altered during the shelf-life time [4-6].

Fresh-cut apples, in slices or in cubes, are lightly processed products and the fruit marketers have shown a great interest in their development. Although from a microbiological point of view, their shelf-life, in refrigerated conditions, has been fixed to 2 or 3 weeks [7], biochemical degradations, such as enzymatic browning, off-flavors release and texture breakdown, occur in a few days [8]. The addition of chemical additives (preservatives, antioxidants, edible coating, colorants, etc.), although used to prolong the shelf-life [9-11] can be associated with health problems such as allergies and more [12]. On this basis, new techniques to maintain the natural qualities of ready-to-eat fruits without using chemical preservatives were developed. The Nicoli’s group first proposed the use of a modified atmosphere composed by 80% N2 and 20% CO2 to better preserve the apple slice quality [13]. Ever since, many attempts have been done to identify the better atmosphere composition to preserve the fresh fruit quality [14,15].

The fruit quality is not a single and well-defined attribute but comprises many properties or characteristics. In many cases, indicators of fruit maturity, such as color, total soluble solids and titratable acids, may not be sufficient to determine optimal sensory quality [16].

Aroma is one of the most important quality parameter perceived by consumers. Gas chromatography technique is usually employed to identify the fruit aroma, while sensory analysis is used to assess the intensity of the aroma descriptors [17]. Whether the chromatographic techniques require special equipment and dedicated staff, the sensory analysis also presents some drawbacks, such as the duration of the panel training and, sometimes, the dubious objectivity of the results. New technologies have been tested on fruits, aiming for fast and, in some cases, non-destructive volatile detection [18,19]. The electronic

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olfactory systems or electronic noses are the new technologies that can
be used to predict fruit quality [20-23].

In this work we studied the changes of aromatic fingerprint of apple
slices, packaged in atmospheric air and in a modified atmosphere
(100% N<sub>2</sub>), stored at 4°C, for 0, 4, 8 and 12 days by an electronic nose.
A commercial type of electronic nose, EOS<sup>835</sup>, based on metal oxide
semiconductor sensors (MOS), to detect changes in volatile profiles
due to storage conditions and during shelf life was used. Moreover,
classical quality parameters, such as total acidity, total soluble solids,
firmness and sensory profile by trained judges, were also determined.

Materials and Methods

Apples and fruit processing

‘Fuji’ apples were harvested on September 2009 in Caltagirone
(Sicily, Italy) and were transported to the laboratory in a
refrigerator, at 4°C, for 0, 4, 8 and 12 days by an electronic nose.
A judge’s panel was engaged in the sensory analysis. The judges (10
persons) were trained in some preliminary sessions by using different
apple samples, in order to develop a common vocabulary for the
description of the sensorial attributes. Eight descriptors were chosen to
describe the quality of apple’s slices: appearance, browning, flavor,
consistency, juiciness, sweetness, acidity, and pleasantness. Samples
were evaluated by assigning a score between 1 (absence of the
sensation) and 5 (extremely intense), except for the descriptor
‘browning’ where a reverse evaluation (1=maximum; 5=minimum)
was adopted. Water at room temperature was used to rinse the sample
before tasting. No statistical analysis of the data was carried out.

Electronic Nose measurements

An EOS<sup>835</sup> (Sacmi, Italy) was used to detect the aromatic
fingerprint at different storage times. The EOS<sup>835</sup> instrument,
belonging to the so-called "electronic noses", consists of an array of
metallic oxide semiconductor (MOS) resistances installed inside a
patented measuring cell, the "sensors chamber". The interaction with
odor molecules causes variation of the electrical conductivity. The
MOS sensors are doped with different metallic oxides (Table 1) that
react differently to the same odorous molecules, thus generating a set
of signals (olfactory imprint or aromatic fingerprint) characteristic of
each analyzed sample. All measurements were done in triplicate,
cutting 3 g of slices and placing them into 20 ml vials. Each vial was
sealed and conditioned at 40°C for 10 minutes before analysis to
ensure that the aroma reached the vial headspace. After equilibration,
4 ml of headspace was drawn in. The interaction between flavors and
sensor layers causes a modification of the sensor resistance. A typical
measurement profile is given in Figure 1. After each measurement the
sensors are constantly purged with GC-grade air. The difference
between the resistance value at rest, R<sub>0</sub>, and that one induced by
sample volatile compounds, R, is considered as the sensor response.
After each measurement, the electronic nose chamber is flushed again
with pure grade air.

Physical and chemical analyses

The juice soluble solid content (SSC) was determined by an optical
refractometer (Atago Co., Ltd., Japan); results are expressed as °Brix.
Total acidity (TA) was measured by titration with 0.1 N NaOH.
Three replicates were performed and the results expressed as g/L of
citric acid.

Firmness measurements were performed by EFFEGI Texture
Analyzer by determining the maximum force during the penetration
of a 8 mm diameter stainless steel cylinder into the apple slice tissue.
Results were expressed as kgf.

Statistical analysis was carried out using the STATSOFT 6.0
program (Vigonza, Padova, Italy). The statistical differences among
different storage times were evaluated by variance analysis (ANOVA),
and mean separation was performed by using the Tukey’s test.

Sensory evaluation

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Table I: MOS sensor array configuration of the EOS.

Results and Discussion

Physical and chemical parameters

The physical and chemical characteristics of apples processed in Air and in N₂ and stored for 0, 4, 8 and 12 days are reported in Table 2. The SSC and TA values do not show noteworthy changes as function of the storage time both in Air and in N₂.

Penetration measurements performed in both experimental condition (air and N₂) showed significant differences in the apple texture, especially in the sample packaged in Air, as the p value calculated for firmness modifications was p ≤ 0.001. In the case of apple slices packaged in N₂, the p value was equal to 0.05. After 12 days of storage, the texture of apple samples packed in Air was softer than that one of samples stored in N₂, thus indicating that a modified atmosphere was more able to save the sample hardness [24,25]. The maintenance of the texture is considered a positive attribute in processed apples.

Sensory evaluation

Results concerning sensory analysis are shown in the radar plot for the 8 chosen descriptors (Figures 2a and b). ‘Sweetness’, ‘acidity’ and ‘consistency’ are in agreement with analytical data results. Figure 2a illustrates that the scores of global ‘appearance’ for apple slices stored in Air after 8 and 12 days were the same, probably due to the presence of brown and soft parts that influenced the acceptance of the judges. Samples stored in N₂ (Figure 2b) showed the lowest ‘appearance’ score after 12 day packing time. The ‘flavor’ descriptor behavior is different for apple slices stored in Air and in N₂: the same value was attributed to samples packaged in Air for 0-4 days and smaller value was assigned to samples examined after 8-12 days. In the N₂ case, the same values were attributed to apples packaged for 0, 4, 8 days and smaller value for those saved for 12 days.

Electronic Nose data

A PCA correlation matrix was carried out on the electronic nose data (Figure 3). For the apple slices packaged in air, the two first principal components, PC1 and PC2, accounted for 94.07% of the total variance (73.92% and 20.15% respectively) (Figure 3a). Samples were distributed along PC1 according to the storage time. The aroma showed changes from 0 to 12 days of storage. At 0 and 4 days, as well as at 8 and 12 days, the aromatic fingerprint is in agreement with the panel estimation.
In the case of apple slices packaged in nitrogen, the two first principal components, PC1 and PC2, accounted for 94.60% of the total variance (75.88% and 18.72% respectively) (Figure 3b). Also in this set, samples were distributed along PC1 according to the storage time. The behavior of aromatic fingerprint was different from that observed for samples packaged in air. In fact, data suggest that the aroma is better preserved in nitrogen: the aroma of samples stored for 0, 4, and 8 days in N2 was separated in a cluster independent on that obtained in the case of samples stored for 12 days. Also in this case the electronic nose response was similar to value assigned by the panel.

Furthermore, the electronic nose results confirm data previously obtained by penetration measurements: the aroma changes are consequence of firmness loss.

Conclusions

Fresh-cut apple tissues undergoing changes during storage, including production of compounds affecting the fruit flavor. Our data show that changes of the aroma fingerprint detected by electronic nose measurements are in agreement with results obtained by physical-chemical analysis and human evaluation panel.

Furthermore, product storage in modified atmosphere (N2) is able to prolong the shelf-life of the fresh-cut apple slices up to 1 week.

The approach here used can be applied to evaluate the loss of aroma quality of other fresh-cut products, in which enzymatic and microbiological processes are responsible for the decrease of the shelf life.

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