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Application of Multivariate Analysis to the Effects of Additives on Chemical and Sensory Quality of Stored Coffee Brew

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The aim of this work was to obtain a black coffee brew to be consumed hot by extension of its shelf life, by addition of additives. Four pH-regulator agents (sodium and potassium carbonates and bicarbonates), one pH regulator and antioxidant (sodium citrate), three antioxidants [sodium ascorbate, ethylenediaminetetracetic acid (EDTA), and sodium sulfite], and lactoserum were tested by sensory analysis. Sodium carbonate and bicarbonate were selected for a study of the physicochemical (soluble and volatile compounds related to the sensory properties) and sensorial quality of coffee brew stored for 90 days at 4 °C. Although both additives extended the shelf life of the coffee brew up to 60 days, sodium carbonate was the chosen additive because it was the most useful in limiting the pH decrease and perception of sourness, which are some of the main factors involved in the rejection of stored coffee brews, and it better maintained the aroma and taste/flavor. Moreover, the application of multivariate analysis facilitated first the description of the global changes of the coffee brews with or without additives throughout the storage using principal component analysis and second the obtainment of a simple equation only with pH and caffeic acid parameters to discriminate the three types of coffee brews and simplify the analytical process, by means of the stepwise discriminant analysis.

KEYWORDS: Coffee; coffee brews; additives; volatile profile; sensory analysis; storage; multivariate analysis; aroma

INTRODUCTION

It is well-known that the storage of coffee brews leads to deterioration of their sensory characteristics (1-3). This quality loss is generally accompanied by sourness development, partially detectable by a pH decrease, even at refrigeration temperatures (2, 4). This is of importance when aiming for a storage stable packed coffee brew. Although these types of coffee beverages are very popular in some countries, such as Japan, their sensory quality is lower than that of freshly prepared coffee brews (5). This could probably be one of the reasons for the lower success of this type of coffee drink in Western countries, where the traditional image of coffee as a freshly brewed beverage is still deeply rooted. Even so, ready-to-drink coffee beverages have reached great acceptability among certain populations because they are inexpensive and storable, providing affordable alternatives to freshly brewed coffee. However, there is still the need to obtain a stable, good quality black coffee brew to be consumed hot.

Several patents proposed the addition of acid-neutralizing, such as carbonates, hydroxides, etc., antioxidants and other

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additives to avoid, or at least reduce, the chemical and sensory evolution of coffee brews during storage, particularly the increase in acidity (6-8). However, an aroma loss and salty taste were observed (7). Moreover, most of the patents where additives were used in coffee were focused on milk-coffee beverages, cappuccino type coffee, soluble coffee, or cold coffee beverages.

Taking into account the fact that patents show limited detail, to the best of our knowledge, there are no detailed studies dealing with the effect of additives on the changes of coffee chemical compounds and the sensory quality of stored coffee brews. For these reasons, the main aim of this work was to contribute to the knowledge of the coffee brew changes during storage by using additives to obtain a black coffee to be consumed hot by extension of the shelf life of the coffee brew obtained (4, 9). Both physicochemical (soluble and volatile compounds) and sensorial qualities were evaluated. Moreover, multivariate statistical analyses were applied as practical tools to know the global patterns of the coffee brew samples during storage by means of principal component analysis (PCA) and to obtain a simple equation to discriminate the coffee brews

 Table 1. Gradient Solvent System and Flow Used in the Method for the Determination of 5-CQA

time (min)	dilution (acetonitrile:water)	flow (mL/min)
0	12.0:88.0	1.000
5	7.5:92.5	1.600
10	8.0:92.0	1.600
15	25.0:75.0	1.600
20	12.0:88.0	1.100

with or without additives and to simplify the analytical process by means of stepwise discriminant analysis (SDA).

MATERIALS AND METHODS

Coffee. Vacuum-packed Colombian Arabica ground roasted coffee (2.25% water content, $L^* = 19.57 \pm 0.09$) was provided by a local factory. The L^* value was analyzed by means of a tristimulus colorimeter (Chromameter-2 CR-200, Minolta, Osaka, Japan) using the D65 illuminant. The instrument was standardized against a white tile before sample measurements. Ground roasted coffee was spread out in an 1 cm Petri plate, and the L^* value was measured in triplicate and on the CIELab scale.

Chemicals and Reagents. The methanol used was of spectrophotometric grade from Panreac (Barcelona, Spain). Acetonitrile, supragradient high-performance liquid chromatography (HPLC) grade, was provided by Scharlau (Barcelona, Spain). Pure reference standards of caffeine, pentoxyfilline, 5-caffeoylquinic acid (5-CQA), caffeic acid, ferulic acid, 4-vinylguaiacol, propanal, hexanal, 2-ethyl-6-methylpyrazine, and acetic acid were obtained from Sigma-Aldrich (Steinheim, Germany); acetaldehyde, 2-methylpropanal, 2-methylbutanal, 3-methylbutanal, 2-propanone, 2-butanone, 2,3-butanedione, 2,3-pentanedione, 2-ethylpyrazine, 2-ethyl-3,5-dimethylpyrazine, and guaiacol (2-methoxyphenol) were purchased from Acros Organics (Springfield, NJ).

Sodium ascorbate, sodium sulfite, sodium citrate, and lactoserum were provided by ANVISA (Madrid, Spain). Sodium carbonate, sodium bicarbonate, potassium carbonate, potassium bicarbonate, and ethylenediaminetetracetic acid (EDTA) were purchased from Panreac.

Coffee Brew Samples. The ground coffee packages were opened immediately before the preparation of the coffee brew to avoid aroma losses. Coffee brews were prepared from 90 g of ground roasted coffee for a water volume of 1 L, using a French press coffeemaker. The extraction time was 3 min, and the water temperature was 90 ± 2 °C (pH 7.0). Each additive was added immediately after coffee brew extraction in a laminar flow cabin. A reference coffee brew without any additive was prepared. Sterilized glass flasks were filled up to the top (330 mL) with fresh coffee brews in a laminar flow cabin, to ensure aseptic conditions and avoid the microbiological contamination of the samples. Afterward, coffee brews were stored at 4 °C until their analysis. This experiment was made in duplicate.

Microbiological Analysis. Aerobic mesophilic flora were analyzed by colony count technique at 30 °C (ISO 4833:2003). Enumeration of molds and yeasts was made by colony count technique at 25 °C (ISO 7954:1987). These analyses were performed monthly.

pH. The measure was obtained with a Crison Basic 20 pH meter.

Caffeine. Extract preparation, cleanup, and HPLC analysis were performed following the method described by Maeztu et al. (10). HPLC analysis was achieved with an analytical HPLC unit model 1100 (Agilent Technologies, Palo Alto, CA), equipped with a binary pump and an automated sample injector. A reversed-phase Hypersil-ODS (5 μ m particle size, 250 mm × 4.6 mm) column was used. The mobile phase was acetonitrile/milliQ water (15:85) in isocratic conditions at a constant flow rate of 2.0 mL/min at 36 °C. Detection was accomplished with a diode array detector, and chromatograms were recorded at 280 nm.

5-CQA. A 500 μ L amount of the coffee brew was diluted up to 50 mL with milliQ water. 5-CQA HPLC analysis was carried out with the same equipment described above. Conditions of the used gradient solvent system and flow are shown in **Table 1**. The wavelength of detection was 325 nm.

Caffeic Acid, Ferulic Acid, and 4-Vinylguaiacol. The extraction, cleanup, and HPLC analysis of these three compounds were performed simultaneously, according to the method developed by Álvarez-Vidaurre et al. (*11*). The HPLC analysis was carried out with the same equipment described above. The chromatographic separation was achieved at 25 °C by using a complex gradient solvent system with acetonitrile/milliQ water adjusted to pH 2.5 with a phosphoric acid solution (*4*). The wavelengths of detection were 314 nm for caffeic acid, 325 nm for ferulic acid, and 210 nm for 4-vinylguaiacol.

Volatile Compound Analysis. The profiles of volatile compounds were obtained with the method described by Sanz et al. (*12*), adapted to coffee brew by Maeztu et al. (*13*), and using static headspace–gas chromatography–mass spectrometry (SH-GC-MS).

After the flask was opened, 6 mL of a homogenized coffee brew was introduced into a 10 mL vial, which was immediately sealed with a silicone rubber Teflon cap. Each vial was equilibrated at 40 °C for 60 min in the SH sampler (model 7694, Agilent Technologies). Each vial was pressurized with carrier gas for 12 s, and 3 mL of the coffee headspace sample was injected into an HP-Wax glass capillary column (60 m × 0.25 mm × 0.5 μ m film thickness) in an HP 6890 gas chromatograph (Agilent Technologies). The injector temperature was 180 °C, and the carrier gas was helium (1 mL/min linear speed). The oven temperature was maintained at 40 °C for 6 min and then raised at 3 °C/min to 190 °C. Mass spectrometry analysis was performed with a Hewlett-Packard mass selective detector model 5973 (Agilent Technologies) operating in the electron impact ionization mode (70 eV), with a scan range of 33–300 amu. The ion source temperature was set at 230 °C. Each sample was analyzed in triplicate.

Identification and Quantification of the Volatile Compounds. The volatile compounds were identified by comparing their mass spectra with those of the pure reference compounds and also by comparison of their Kovats indices with those of standard compounds. The Kovats indices were calculated according to the method of Tranchant (14). Peak areas were measured by calculation of each volatile total area based on integration of a single ion. The quantification ion of each volatile compound is given in **Table 5**.

Sensory Descriptive Analysis. Twenty judges were recruited among members of the Nutrition, Food Science, Physiology, and Toxicology Department at the University of Navarra. Selection and training were carried out as described by Maeztu et al. (*10, 13*) to have a 10-member panel. Retraining and sensory standards were described by Pérez-Martínez et al. (*4, 9*). A scorecard with the most frequently perceived sensory attributes was developed during training. Two lines for "other" aromas and flavors were added. All of the descriptors were rated on 11-point scales from "none" (0) to "very high" (*10*).

Each coffee brew sample was heated in a microwave oven at 90 \pm 2 °C immediately before tasting and served monadically in a white porcelain coffee cup. The order of presentation was randomized among sessions. A freshly prepared coffee brew was evaluated first, as a reference and to avoid first impressions. All evaluations were conducted in isolated sensory booths illuminated with white light in the sensory laboratory under standardized conditions by UNE 87-004-79 (*15*). Rinse water was provided between samples. After the individual evaluation of each sample, results were discussed in order to find new other sensory attributes that could be developed in the coffee brew during the study and to establish the shelf life by consensus.

Statistical Analysis. Each parameter was analyzed in triplicate. Results are shown as means \pm standard deviations. A two-way analysis of variance (ANOVA) was performed to establish the impact of both the additive addition (sodium carbonate and bicarbonate) and the storage time on several physicochemical and aroma parameters of coffee brew samples (**Table 4**). When interactions were significant, a one-way ANOVA was applied. A *T*-Tukey test was applied as a test a posteriori with a level of significance of 95%.

Correlations among variables were assessed by means of the Pearson correlation test. PCA, based on the Pearson correlation matrix, was applied to the data. Principal components (PC) with eigenvalues higher than 1 were selected. SDA was applied to obtain a simple equation by which the coffee brew samples could be classified. Wilk's Lambda stepwise method was used. The criteria were 0.05 for maximum significance of F to enter and 0.10 minimum significance of F to

Table 2. Sensory Analysis of the Coffee Brews with Additives (75 ppm)^a

reference	sodium carbonate	potassium carbonate	sodium bicarbonate	potassium bicarbonate	sodium citrate	sodium ascorbate	EDTA	sodium sulfite	lactoserum
				aroma					
8 ± 1	7 ± 1	7 ± 1	8 ± 1	6 ± 1	7 ± 0	7 ± 1	7 ± 1	6 ± 1	7 ± 1
8 ± 1	7 ± 1	6 ± 1	8 ± 0	6 ± 1	6 ± 1	7 ± 1	7 ± 0	6 ± 0	7 ± 2
			tas	ste/flavor					
3 ± 1	3 ± 1	5 ± 1	3 ± 1	4 ± 1	6 ± 1	3 ± 1	4 ± 2	5 ± 1	0 ± 1
8 ± 1	7 ± 1	7 ± 1	7 ± 1	5 ± 1	5 ± 0	7 ± 1	7 ± 1	7 ± 1	7 ± 0
0 ± 0	1 ± 1	1 ± 0	1 ± 1	1 ± 1	4 ± 1	0 ± 1	1 ± 1	3 ± 0	1 ± 1
1 ± 1	2 ± 1	4 ± 1	2 ± 0	4 ± 1	3 ± 0	2 ± 1	2 ± 1	5 ± 1	2 ± 1
5 ± 1	4 ± 1	4 ± 0	5 ± 1	5 ± 0	4 ± 1	5 ± 1	5 ± 0	1 ± 0	4 ± 1
0 ± 0	1 ± 1	3 ± 1	1 ± 0	4 ± 1	5 ± 1	0 ± 0	1 ± 1	4 ± 1	1 ± 1
0 ± 0	1 ± 0	0 ± 1	0 ± 0	0 ± 0	0 ± 0	0 ± 0	0 ± 0	1 ± 0	0 ± 1
0 ± 0	1 ± 1	0 ± 1	0 ± 0	0 ± 0	0 ± 0	0 ± 1	0 ± 0	0 ± 0	0 ± 0
	$\begin{array}{c} \text{reference} \\ 8 \pm 1 \\ 8 \pm 1 \\ 0 \pm 0 \\ 1 \pm 1 \\ 5 \pm 1 \\ 0 \pm 0 \\ 1 \pm 0 \\ 0 \pm 0 \\ 0 \pm 0 \\ 0 \pm 0 \end{array}$	$\begin{array}{c c} & sodium \\ \hline reference & carbonate \\ \hline 8 \pm 1 & 7 \pm 1 \\ 8 \pm 1 & 7 \pm 1 \\ \hline 3 \pm 1 & 3 \pm 1 \\ 8 \pm 1 & 7 \pm 1 \\ \hline 0 \pm 0 & 1 \pm 1 \\ 1 \pm 1 & 2 \pm 1 \\ 5 \pm 1 & 4 \pm 1 \\ 0 \pm 0 & 1 \pm 1 \\ \hline \end{array}$	$\begin{array}{c ccccccccccccccccccccccccccccccccccc$						

^a All values are shown as means \pm standard deviations.

Tabl	e 3.	Sensory	Analysis	of	the	Coffee	Brews	with	Additives	during	8	Days	a
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			storage	time (days)	
	fresh coffee (control)	0	1	4	6 8	
	aroma					
aroma intensity	9-10					
sodium carbonate		7 ± 1	7 ± 1	7 ± 1	6 ± 1 $6\pm$	1
sodium bicarbonate		7 ± 1	6 ± 0	7 ± 1	$6\pm1.7\pm$	1
sodium ascorbate		6 ± 1	7 ± 1.6	3 ± 1	$5\pm0.6\pm$	1
lactoserum		8 ± 1	7 ± 1	7 ± 2	$7\pm1.6\pm$	0
aroma freshness	9—10					
sodium carbonate		7 ± 1	7 ± 1	7 ± 1	$6\pm1.6\pm$	1
sodium bicarbonate		7 ± 1	6 ± 1.6	5±1	$5\pm0.7\pm$	1
sodium ascorbate		7 ± 1	7 ± 1.6	5 ± 0	$5 \pm 1.6 \pm$	1
lactoserum		7 ± 1	7 ± 0	1 ± 1	$7\pm1.5\pm$	0
	taste/flav	or				
bitterness	0-1					
sodium carbonate		2 ± 0	2 ± 02	2 ± 1	$3\pm12\pm$	0
sodium bicarbonate		3 ± 1	3 ± 12	2 ± 0	$3\pm0.2\pm$	0
sodium ascorbate		2 ± 1	1 ± 0	1 ± 0	$2 \pm 0.2 \pm$	0
lactoserum	0 10	2 ± 1	3 ± 1	1 ± 0	$1\pm0.3\pm$	1
acidity	8-10	7 . 1	0 1 4 /		C O C	4
sodium carbonate		/±1	8±10) ±		: -
sodium picarbonate		り±0 フ 1	/±1/	/ ± 1		: -
lactocorum		/±1	0±10 6±20		$0 \pm 1 0 \pm 7 \pm 1 4 \pm 1$: I . 1
acioserum	0	0 ± 1	0 ± 2 0	JTI	1 1 1 4 1	
sodium carbonate	0	0 + 0	0 + 0 0	n + n	0 + 0.1 +	0
sodium bicarbonate		0 ± 0 0 + 0	0 ± 0	0 + 0	1+10+	0
sodium ascorbate		0 ± 0 0 ± 1	0 ± 0	2 + 1	1+0.1+	1
lactoserum		0 + 0	$1 + 0^{-1}$	1 + 0	2+13+	1
astringency	0-1	• - •		•		•
sodium carbonate		1 ± 0	0 ± 1^{-1}	1 ± 1	$1\pm10\pm$	0
sodium bicarbonate		1 ± 1	1 ± 0	1 ± 1	0 ± 1 $0\pm$	1
sodium ascorbate		2 ± 1	0 ± 1 ·	1 ± 0	1 ± 0 $0\pm$	1
lactoserum		1 ± 0	1 ± 0	1 ± 1	$1\pm0.1\pm$	1
persistence	9-10					
sodium carbonate		6 ± 1	6±26	3 ± 1	$4\pm0~5\pm$	1
sodium bicarbonate		4 ± 1	5±1 5	5 ± 1	$5\pm1~5\pm$	1
sodium ascorbate		5 ± 1	4±16	3 ± 1	5 ± 0 $4\pm$	1
lactoserum		5 ± 1	3±1 8	5 ± 0	5 ± 1 $3\pm$	1
aftertaste	0					
sodium carbonate		0 ± 1	1 ± 0	l ± 1	2 ± 1 $3\pm$	1
sodium bicarbonate		1 ± 1	2 ± 1	1 ± 0	$1\pm0.0\pm$	1
sodium ascorbate		1 ± 0	2 ± 1	1 ± 0	$2\pm11\pm$	1
lactoserum		2 ± 1	4 ± 12	2±1	$1\pm14\pm$	1

 a All values are shown as means \pm standard deviations.

remove. All statistical analyses were performed using the SPSS v.15.0 software package.

RESULTS AND DISCUSSION

Selection of the Additives. Previous studies on the Colombian Arabica coffee brews showed that staling is mainly due to

Table	4.	Two-Way	ANOVA	Results	of	Coffee	pН	and	Chemical
Compo	our	nds ^a							

	additive	effect	storage	time effect	(additive \times st	torage time)
	F	р	F	p	F	р
pН	399.48	***	295.34	***	15.68	***
caffeine	0.97	NS	6.18	***	1.17	NS
5-CQA	18.29	***	67.60	***	4.83	***
caffeic acid	14.29	***	77.32	***	2.79	**
ferulic acid	1.50	NS	23.52	***	1.77	NS
4-vinylguaiacol	4.64	*	17.31	***	1.42	NS
acetaldehyde	72.70	***	437.04	***	2.68	**
propanal	26.08	***	22.99	***	2.87	**
2-methylpropanal	30.85	***	28.00	***	0.64	NS
2-propanone	13.26	***	34.57	***	2.10	*
2-butanone	20.76	***	23.00	***	2.22	*
2-methylbutanal	51.57	***	33.98	***	1.45	NS
3-methylbutanal	17.34	***	2.77	*	1.05	NS
2,3-butanedione	5.42	**	10.39	***	5.27	***
2,3-pentanedione	6.48	**	28.21	***	4.07	***
hexanal	5.34	**	10.05	***	4.16	***
acetic acid	0.14	NS	27.83	***	0.51	NS

^{*a*} *p*: NS, nonsignificant (p > 0.05); *significant (p < 0.05); **very significant (p < 0.01); and ***highly significant (p < 0.001).

the development of sourness and other nontypical coffee taste/ flavors (rancidity, aftertaste) and loss of aroma, and it is faster in the presence of oxygen (4, 9). For these reasons, pH-regulator and antioxidant agents were previously selected to extend the shelf life of coffee brew. A preliminary study on the sensory effects of additives to coffee brew was made taking into account that sensory properties are crucial for the coffee quality. Four pH-regulator agents (sodium and potassium carbonates and bicarbonates), one pH-regulator and antioxidant (sodium citrate), three antioxidants (sodium ascorbate, EDTA, and sodium sulfite), and lactoserum were tested by sensory analysis, and the results are shown in **Table 2**.

Carbonates and bicarbonates are used to reduce acidity in beverages, including coffee type. Also, these chemical compounds together with polymers are foam-making agents very useful for cappuccino type or milk–coffee beverages. Seventyfive ppm of sodium or potassium carbonates or bicarbonates was added to coffee. Sodium carbonate or bicarbonate showed no influence on aroma and taste/flavor of coffee when they were compared with a reference coffee brew (without additives). However, the addition of potassium salts increased not only the bitterness effect, which is very well-known, but also astringency and aftertaste and slightly decreased freshness aroma, diminishing Colombian coffee quality. Consequently, potassium carbonate and bicarbonate were rejected.

Table 5. Effect of the Sodium Carbonate and Bicarbonate on the Aroma Impact Compounds (Area \times 10⁻³) of Coffee Brews Throughout Storage at 4 $^{\circ}$ C^a

							storage time (day	s)			
QI ^b	KIc		0	З	7	10	15	20	30	60	06
135	1455	2-ethyl-3,5-dimethylpyrazine									
		reference	ND	ND	ND	DN	DN	ND	ND	DN	DN
		NaHCO ₃	QN	DN	ND	DN	QN	ND	DN	DN	ND
		Na ₂ CO ₃	DN	DN	DN	ND	DN	DN	DN	DN	ND
						acids					
45	1480	acetic acid									
		reference 1	338 ± 8 a	512 ± 12 ab	620 ± 68 ab	596 ± 40 ab	514 ± 82 ab	675 ± 33 b	$908\pm156~{ m c}$	$1090\pm507~{ m c}$	1472 ± 473 d
		NaHCO ₃ 1	343 ± 67 a	$494 \pm 98 ab$	542 ± 36 ab	558 ± 20 ab	573 ± 142 ab	750 土 118 b	$1003\pm162~{ m c}$	$990\pm412~{ m c}$	1394 ± 32 d
		Na ₂ CO ₃ 1	$343\pm40\mathrm{a}$	592 ± 40 ab	$663\pm30~ m ab$	587 ± 31 ab	567 ± 23 ab	$672\pm40~{ m b}$	$1032\pm27~{ m c}$	$1003\pm298~{ m c}$	1161 ± 18 d
					phenol	lic compounds					
109	1864	guaiacol									
		reference	QN	QN	QN	QN	QN	QN	QN	DN	QN
		NaHCO ₃	ND	ND	ND	DN	DN	ND	ND	DN	DN
		Na ₂ CO ₃	ND	ND	DN	DN	ND	DN	ND	ND	DN
^a All val	lues are shov	wn as means \pm standard deviation:	s. In each row and	column, different lett	ters indicate statistic.	ally significant differ	ences ($p < 0.05$) thro.	ughout the time. ND,	not detected. The nu	umber in the name in	the sample indicate
- 0104+ +04+	intoi on onim	action with the stores time and th	o odditino oddition	Difforont numbers	o the name in the o	nomala indianta atati	otionly, cianificant diff	- U U U U	hronish off though	to the addition of	dition b OI ion 100
.nal lijere	Was no inter-	action with the storage unite and un	le additive addition.	. UITERENT FIULTIVELS	In the name in the s	sample indicate stat	sucally significant un	erences $(p < 0.00)$	urougnout the time u	ine to the auditive at	idition. 7 Qi, iuti use
for the diff	antification of	the compound c KI Kovete index	· roloniated for the	A vehice vehicles of the							

Table 5. Continued



Figure 1. Effect of the sodium carbonate and bicarbonate on the pH of coffee brews throughout storage at 4 °C.

Citric acid is a pH-regulator and reducing agent commonly used in beverages and many other foods because of its safety. In this study, to avoid the increase of protons, citric acid was added as a sodium salt. However, sourness, and also bitterness and aftertaste, were the highest, and freshness aroma and the typical acidity of Colombian coffee were the lowest in comparison to the reference coffee. For this reason, sodium citrate was also rejected.

Ascorbic acid acts as a potent water-soluble antioxidant by scavenging free radicals such as hydroxyl, peroxyl, and hydroperoxyl radicals (16). Ascorbic acid was added to coffee brew as sodium salt, and a sensory profile similar to the reference coffee was observed.

EDTA is a transition metal chelator. Therefore, it inhibits the formation of hydroxyl radicals from hydrogen peroxide (17). The addition of EDTA to coffee increased the bitterness of the Colombian coffee brew and consequently was rejected.

Sodium sulfite is a potent oxygen scavenger under neutral and basic conditions. Although it is broadly applied for shelf life extension in the food industry, its application is progressively limited because its allergenicity. The addition of sodium sulfite to coffee decreased the aroma and increased the undesirable taste/flavors (bitterness, sourness, astringency, and aftertaste). Therefore, this additive was also rejected.

Taking into account that the addition of milk to coffee and tea is very common to reduce bitterness and astringency and that milk has an almost neutral pH, lactoserum was also used in the preliminary study. This milk component was preselected because the aim of this study is focused on black coffee. The addition of lactoserum to coffee only decreased the bitterness but did not modify any other sensory attributes of the Colombian coffee brew. For this reason, this additive was selected for the next step.

The effect of a higher concentration (100 ppm) of the preselected additives (sodium carbonate, bicarbonate and ascorbate, and lactoserum) was also studied. However, the results of the sensory analysis (data not shown) revealed that the use of a higher dose had a negative influence on the coffee brews. Therefore, the concentration of 75 ppm was used in the subsequent studies.

The next step for the selection of the additives was to carry out a short-term study (8 days) with coffee brews prepared with sodium carbonate, sodium bicarbonate, sodium ascorbate, and lactoserum at a concentration of 75 ppm. As in the previous experiments, sensory analysis was the selection test. A recent coffee brew without additives was used as a reference coffee,



Figure 2. Effect of the sodium carbonate and bicarbonate on the soluble compounds.

namely, fresh coffee, because the final goal of this study was to obtain a stored coffee brew similar to fresh coffee. The results of the sensory analysis are shown in Table 3. Both aroma intensity and freshness kept quite stable throughout the study, even though a tendency to decrease with time was observed. The relative aroma stability might be due to the absence of oxygen. As Charles-Bernard and co-workers observed, this absence had a higher stabilizing effect on the volatile thiols, some of them related to coffee freshness aroma, than antioxidants such as sodium ascorbate (18). Rancid burnt and/or spicy aromas and flavors were not perceived in any of the studied coffee brews along the storage time. Bitterness and astringency maintained low scores, with small variations, throughout the time. In contrast, typical coffee acidity and persistence showed a tendency to decrease whereas sourness and aftertaste tended to increase. These changes were more intense in coffee brews with lactoserum. Moreover, sourness was perceived in the fourth day in coffee brews with sodium ascorbate. Therefore, sodium carbonate and bicarbonate were selected for the long-term study.

Influence of Sodium Carbonate and Bicarbonate on the Coffee Brew Stability. Coffee brews with sodium carbonate or sodium bicarbonate (75 ppm) and a reference coffee brew (without additives), aseptically bottled without headspace and stored at 4 °C for 90 days, were analyzed. The microbiological analysis of the coffee brews during the long-term study showed a colony count number lower than 1 cfu/mL both for mesophilic flora and for molds and yeasts. Therefore, the aseptic handling of the sample preparation and the bottling storage were effective to avoid the microbiological contamination of the coffee brews.

pH and soluble and volatile compounds related to the sensory

properties of coffee brews were studied. A two-way ANOVA was performed to establish the impact of the additive and the storage time on the pH and the chemical compounds of coffee brews (**Table 4**). In most cases, significant interaction between the additive and the storage time has been observed. Those compounds that have no significant interaction effect were significantly affected by both factors, except caffeine, ferulic, and acetic acids, which were not significantly affected by the additives. Moreover, F values corresponding to the storage time were higher than the F values of the additives for all soluble compounds and most of the volatiles, showing greater importance of the storage time effect, which was in detail described in previous works (4, 9).

The effect of the sodium carbonate and bicarbonate on the pH of coffee brews throughout storage at 4 °C is shown in Figure 1. As pH-regulator agents, both additives significantly suppressed the reduction in pH over storage time and increased the pH of coffee brews. At initial time, there were significantly higher pH of the sodium carbonate (5.04) and bicarbonate (5.02)coffee brews than the reference (coffee brew without additives, 4.97). Sodium carbonate coffee brew pH decreased the least. Although both additive coffee brews did not reach a pH lower than 4.8 considered as the limit of the acceptance by some authors (19, 20), the sodium bicarbonate coffee brew pH decreased faster than the carbonate brew. In fact, at 7 days, pH was not significantly different than reference coffee one. However, after 20 days, the pH decrease of bicarbonate coffee brew was progressively slower in comparison to the reference and similar to carbonate coffee brew. This different behavior may be attributed to the hydrogen cation of the bicarbonate that partially contributed to the pH decrease.



Figure 3. Effect of the sodium carbonate and bicarbonate on the sensory profile of coffee brews at 0, 30, 60, and 90 days.

Figure 2 shows the effect of the sodium carbonate and bicarbonate on the soluble compounds of coffee brews throughout storage at 4 °C. The addition of these pH-regulator agents did not influence the changes induced by the storage time (4). Only a higher, but not statistically significant, decrease of 5-CQA in sodium carbonate coffee brew could be observed. This slightly lower amount of 5-CQA could be due to the influence of a higher pH on a lower hydrolysis of chlorogenic acid lactones formed during coffee roasting (21), a higher isomerization to 3-CQA and 4-CQA, or a lower release of chlorogenic acids from noncovalently linked polymeric skeletons, such as melanoidins (22), but not to decomposition to caffeic and quinic acids because the former was also lower, but not significantly, in the carbonate coffee brew.

Coffee aroma is one of the most appreciated characteristics of coffee brews, and its loss is one of the consequences of staling. For this reason, the influence of the sodium carbonate and bicarbonate on the most frequently reported coffee aroma impacts compounds (13, 23-29). One sulfur compound, six aldehydes, four ketones, three pyrazines, one acid, and one phenolic compound were analyzed, and the results are shown in **Table 5**.

Neither methanethiol, a sulfur compound responsible for freshness aroma in ground roasted coffee (30) and in espresso coffee (13), nor guaiacol (2-methoxyphenol), responsible for phenolic and spicy aromas (24) and phenolic and burnt flavors (26), was present at detectable levels in any coffee brew throughout storage. Two ethylpyrazine, 2-ethyl-6-methylpyrazine and 2-ethyl-3,5-dimethylpyrazine, associated with roasty and earthy/musty flavors in ground roasted and brewed coffees (23, 24) and with flowery and fruity aromas of coffee

brews in the case of 2-ethyl-6-methylpyrazine (29) were not detected. Similar results were also observed in coffee brews stored in the presence of air (9).

The most abundant volatile compounds, 3-methylbutanal, 2-methylbutanal, and 2-methylpropanal (Strecker aldehydes), did not show significant interaction in the two-way ANOVA (**Table 4**), but they were significantly affected by both additive and storage time. The addition of sodium carbonate and bicarbonate to coffee brew induced to a lower amount of these volatiles but also for most of the other aldehydes and ketones. Moreover, the absence of oxygen induced less change over time in comparison with results previously reported for coffee brews stored with air headspace, which should lead to better maintenance of coffee aroma (9).

Acetic acid, which has no significant interaction effect in the two-way ANOVA (table 4), was significantly affected only by storage time. The absence/presence of oxygen seems not to have influence in the increase of this volatile because similar results were observed in coffee brews stored with air at the same temperature (4 °C) (9). However, the storage temperature was critical.

Finally, the influence of the sodium carbonate and bicarbonate on the sensory quality of coffee brews throughout storage at 4 °C is shown in **Figure 3**. Although the additives decreased the original acidity of the Colombian coffee brews, the acidity score was adequately high and the other sensory attributes were hardly affected. With time, the typical acidity of the Colombian coffee brews was maintained up to 30 days in sodium carbonate coffee brew, whereas it decreased in the others because the increase of sourness unbalanced the global acidity. Even though the perception of sourness and other nontypical coffee taste and



Figure 4. PCA of the coffee brews throughout storage at 4 $^{\circ}$ C: (a) parameter loadings and (b) sample scores.

flavors, such as aftertaste and astringency, was established at 20 days shelf life for reference coffee brew [confirming the shelf life established in Pérez-Martínez et al. (4, 9)] and 60 days for both coffee brews with sodium carbonate or bicarbonate, the addition of carbonate better maintained the aroma and taste/ flavor of coffee brews.

PCA. PCA is a method that aims to recognize patterns in multivariate data sets or to reduce the dimensionality of a data set obtaining linear combinations of original variables called PCs. In this paper, taking into account the high number of physicochemical and sensorial parameters and points of analysis, this method appeared to be very useful to describe the global changes of the coffee brews with or without additives throughout the storage at 4 °C. Five PCs with eigenvalues higher than 1 were selected by PCA. PC1 and PC2 explained 70.2% of the total variance. Figure 4 shows bidimensional plots of PC1 and PC2 parameter loadings and sample scores. PC1, which explained 53.2% of the total variance, is mainly characterized by sensory attributes, pH, and most of the coffee aroma compounds. PC2, which explained 17.0% of the total variance, is mainly characterized by soluble compounds (caffeic acid, 5-CQA, and caffeine) and the rest of the volatiles.

As can be seen, when the storage time was increased, coffee brews were moved on the left half of the graphic from the top to the bottom due to coffee aroma decrease; however, the typical attributes of Colombian coffee were maintained. On the loss of coffee quality, the products moved to the right half of the graphic because of the decrease of pH, the increase of acetic acid, and the presence of sourness and other nontypical coffee taste and flavors, such as rancidity. Moreover, although all of the coffee brews show a global pattern very similar, the reference coffee brew (without additives) is placed on the right of the coffee



Figure 5. Discriminant scores and centroid values of the coffee brew samples.

brews with additives mainly because of the lower pH and good sensory attributes even during the first days, and sodium carbonate coffee brew is on the left because this coffee brew maintains the coffee quality longer.

Discriminant Analysis (DA). DA is the best-known and most often used supervised classification method in which knowledge of the grouping structure is used to develop rules that predict the group that a new object belongs to. SDA was applied to obtain a simple equation by which the coffee brew samples could be classified. When SDA was applied to all physico-chemical parameters, two discriminant functions (DFs) were obtained. The DF1 that explained 96% of the total variance is shown as follows:

 $y = 32.250 \times pH - 2.830 \times caffeic acid +$ 0.003×2 -propanone - 0.001 × 2,3-butanodione + 0.003 × acetic acid - 158.306

DF1 allowed the classification of the coffee samples into their respective group with a success rate of 75.3%. However, very different parameters participated in the obtained functions. Some of them, such as pH, were measured by simple methods, but others were measured by HPLC (soluble compounds) or HS-GC-MS (volatiles). Consequently, from the analytical point of view, this equation only partially contributed to simplify the analyses to differentiate the three types of coffee brews. For this reason, and because the changes in acidity and taste/flavor-related compounds are the most relevant in coffee brews, a new SDA was applied only to pH and soluble compounds. Two DFs, using only pH and caffeic acid, were obtained. DF1, which explained 100.0% of the total variance, was

 $y = 18.985 \times pH - 3.427 \times caffeic acid - 87.204$

Figure 5 shows the sample results for DF1 and DF2 and the centroids scores. DF1 allowed the classification of the coffee samples into their respective group with a success rate of 81.5%. The sodium carbonate and bicarbonate coffee samples stored

for 90 days were misclassified and included in the group of reference coffee brews, maybe because they overpassed the shelf life, and consequently, they lost the coffee quality.

In summary, among all of the tested additives, both sodium carbonate and sodium bicarbonate were the most effective to keep the coffee brew quality longer. In fact, a shelf life of 60 days was proposed for these coffee brews, in comparison with the 20 days shelf life established for a coffee brew without additives (4). However, sodium carbonate was the chosen additive because it is the most useful to reduce the pH decrease and the appearance of sourness, which are some of the main factors involved in the rejection of stored coffee brews and better maintains the aroma and taste/flavor. Moreover, the application of multivariate analysis facilitated, first, the description of the global changes of the coffee brews with or without additives throughout the storage at 4 °C using the PCA and, second, the obtainment of a simple equation with pH and caffeic acid parameters to discriminate the three types of coffee brews and to simplify the analytical process by means of the SDA.

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