# Yield and Chemical Composition of Essential oil of Piperaceae in one Segment of the Semi deciduous Forest of Paraná State, Brazil, in Seasonal Samplings

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Abstract— Essential oils are composed of a complex mixture of various classes of substances; among them are phenylpropanoid, monoterpenes, and sesquiterpenes, belonging to the secondary metabolism of plants. However, these compounds can be influenced by seasonal factors, among others. The objective of this study is to realize the prospect of obtaining Piperaceae with aromatic potential from a segment of the semideciduous forest of the Atlantic Forest Biome in the northwester region of Paraná State. The aim is to assess the qualitative and quantitative characteristics of its essential oil in the collection during the winter and summer seasons of year 2016. The statistical design was internally randomized in a factorial of 4 (species) x 3 (replicates). The species studied were Piper mosenii C. DC., Piper xylopioides Kunth, Piper diospyrifolium (Kunth) Kunth ex C. DC., and Piper gaudichaudianum Kunth. A total of 78 compounds were identified, 68 in winter and 71 in summer. The species presented a variation in the yield and composition of essential oils, both in winter and summer. The predominant chemical sesquiterpenes followed monoterpenes, with prominence of (E)-caryophyllene, germacrene D, bicyclogermacrene, α-pinene and β-

Keywords— Bioprospecting, genus Piper, protected areas, secondary metabolism, species aromatics.

## I. INTRODUCTION

Brazil has a wide territorial area with a rich biodiversity, which provides a valuable source of plant species. Many of these are little studied and constitute a large biological collection for scientific research. The prospect of finding aromatic species in the forests of the Atlantic Forest biome, with its significant biodiversity, may represent the discovery of new essential oils with a potential for beneficial use, increasing the viability of sustainable management of this ecosystem, which is highly damaged, with a gerat need for conservation.

The plants are inexhaustible sources of natural products, many of them essential oils and secondary metabolites, mainly used in perfumery and cleaning products. They are also a source of active ingredients for the pharmaceutical industry (5; 33; 11). They can also be used in crop protection against pests and diseases, with the advantage of not accumulating in the environment and have a wide spectrum of action, which reduces the risk of developing resistant strains pathogenically (10).

More than 2000 plant species produce essential oils, among them are several representatives of the Piperaceae family, which possess high agronomic and commercial significance, as they are used as condiments, adornments, food, and in popular medicine. Some species of the genus *Piper* are also used in folk medicines, especially in Brazil, and many have proven to be of great significance due to the pharmacological activity and/or production of essential oils in their structure (13).

Piperaceae encompasses approximately 2,000 species allocated in approximately five genera. Just over 500 species are recorded in Brazil, distributed in four genera (30). The *Piper* genus is represented by 290 species and the Atlantic Forest is one of the centers of diversification and endemism of the genus, with about 150 species (14).

The production of essential oils in Brazil is still incipient to meet the demand. However, the national and international market has demonstrated a great interest in new essences; something that this biodiversity has a great potential to meet. In this context, the conservation units are an excellent laboratory for research and prospecting of essential oils. Various floristic studies were carried out at

the Caiuá Ecological Station, North Diamond, Paraná State, however, phytochemical and biological studies with the aromatic native species in this forest formation are scarce, which makes this research highly relevant.

The objective of this study is to evaluate, quantitatively and qualitatively, the essential oil in the seasonal samplings of Piperaceae, which is commonly found in a segment of the semideciduous forest of the Atlantic Forest Biome located in the Northwest of Paraná State.

## II. MATERIAL AND METHODS

The study was conducted in the Caiuá Ecological Station, a Conservation Unit in the state, with an area of 1,427.30 ha, located in the northwestern region of Paraná State, Municipality of North Diamond, with approximate coordinates between 52°49' to 52° 53' W and 22° 34' to 22° 37' S and an altitude that varies from 240 m to 380 m. It belongs to the hydrographic basin of the Lower Paranapanema River, with part of the area occupying the banks of the reservoir of the Rosana Hydroelectric Power Station (UHE Rosana), remnant of the Paranapanema River.

According to the Koeppen climate classification, the northwestern region of Paraná State presents the Cfa type climate - mesothermal, humid, without a dry season, and with hot summers. The average temperature of the coldest month is below 18°C and the average temperature of the hottest month is above 22°C. The average annual rainfall is 1,200 to 1,400 mm, being the quarter (December, January, and February) when it rains. The average annual

temperature is between 21°C and 22°C, being the average of the hottest month (February) 24 to 25°C and the coldest month (July) 17°C to 18°C. The relative humidity of the air average is 75%, (4).

The formation of the majority of the soil in the Caiuá Ecological Station, is represented by soils derived from river sediments in portions adjacent to the Paranapanema River, with a predominance of Red Latosols, Red Argisols, Red-Yellow Argisols, and Quartzarenic Neosols, respectively (7). Its vegetation cover is inserted in the Atlantic Forest biome in the region of the Semideciduous Forest vegetation type, whose ecological concept is conditioned to the dual climate seasonality (15).

The studied species were *Piper mosenii* C. DC., *Piper xylopioides* Kunth, *Piper diospyrifolium* (Kunth) Kunth ex C. DC., and *Piper gaudichaudianum* Kunth species. These occurred commonly in the conservation unit.

The field work consisted of the collection of approximately 1 kg of plant material of each species (leaves and branch terminals) for extraction and quantification, and for determining the moisture content of the samples. The collection and transport of the plant material was prompted by the Environmental Institute of Paraná, under proper environmental authorization, with number 03/2016. The herbarium specimens were transported to the Botanical Museum Hall of Curitiba city where they were herborized and classified (Table 1).

Table 1- Description of species collected during the winter and summer of 2016 in the Caiuá Ecological Station, North Diamond /PR.

Species	Herbarium Latitude		Longitude	Altitude	
	number				
Piper mosenii C. DC	MBM 396409	22°36'8.5"S	52°53 4.7W	288m	
Piper gaudichaudianum Kunth.	MBM 396403	22°36'29.3"S	52°51'58.4"W	277m	
Piper xylopioides Kunth	MBM 396405	22°36'34.4"S	52°51'48.7"W	273m	
Piper diospyrifolium (Kunth)	MBM 396413	22°36'21.7S	52°52'4.8W	268m	
Kunth ex C. DC.					

The identification of species was performed with the aid of specialized bibliographies, comparison of herbarium specimens declared at the herbarium, and consultation with experts on the respective groups of plants of these species.

The oil extraction was done by means of hydrodistillation during four-and-a-half hours in a graduated Clevenger apparatus using 100 g of fresh leaves and one liter of distilled water, with three repetitions.

The essential oils were diluted in hexane at a ratio of 1% and 1.0  $\mu$ L of the solution where it was injected, with a split flow of 1:20, in an Agilent 6890 gas chromatograph coupled with a mass selective detector Agilent 5973N. The injector was maintained at 250°C. The separation of the constituents was obtained in a capillary column HP-5MS (5%-phenyl-95%-dimethyl polysiloxane, 30 m x 0.25 mm x 0.25  $\mu$ m), using helium as a carrier gas (1.0 ml min<sup>-1</sup>). The temperature of the oven was scheduled to be 60°C to 240°C at a rate of 3°C

min<sup>-1</sup>. The detector of mass was operated on electronic ionization mode (70 eV), at a rate of  $3.15~s^{-1}$  sweeps and mass range of  $40~\mu$  to  $450~\mu$ . The transfer line was maintained at  $260^{\circ}$ C, the source of ions at  $230^{\circ}$ C, and the analyser at  $150^{\circ}$ C.

For the quantification, the diluted samples were injected into the chromatograph Agilent 7890A equipped with flame ionization detector (FID), operated at 280°C. The same column and analytical conditions described above were employed, except for the carrier gas used, which was hydrogen, at a flow rate of 1.5 mL min<sup>-1</sup>. The percentage composition was obtained by electronic integration of the signal of the FID by dividing the area of each component by the total area (area %).

The identification of the constituents was obtained by comparison of their mass spectra with those of (32) and (18) and also by their linear retention indices calculated from the injection of a homologous series of hydrocarbons (C7-C26) and compared with data from the literature (1).

The results were submitted to analysis of variance and the means of treatments were compared by the Tukey test at 5% probability, using the software SISVAR (9) and the principal component analysis (PCA) using the program BioEstat v.5.

### III. RESULTS AND DISCUSSION

There was a significant difference in the essential oil content among species and at different seasons of collection (Table 2). The species *P. xylopioides* presented an average yield of oil statistically superior to others, and in the summer this content was higher. *P. mosenii*, *P. gaudichaudianum*, and *P. diospyrifolium* showed similar levels of oil in winter, however, in the summer these species differed in the levels among them, whereas, *P. xylopioides* presented the highest content followed by *P. diospyrifolium* and *P. gaudichaudianum*. *P. mosenii* did not produce oil in the summer contrary to the study of (24) conducted on the coast of Paraná State, where the species showed a low variation in the essential oil content in the winter, spring, and summer seasons.

Table 2 – Averages of the content of essential oils of fresh samples of species collected during the winter and summer of 2016, in the Caiuá Ecological Station, North Diamond, Paraná State, Brazil

Species	Oil content (%)*		
	Winter	Summer	
Piper mosenii C. DC	0.28 b A		
Piper gaudichaudianum Kunth.	0.13 b A	0.05 cB	
Piper xylopioides Kunth	0.94 a B	1.37 a A	
Piper diospyrifolium (Kunth) Kunth ex C. DC.	0.13 b B	0.38 b A	
CV (%)	23.54		

<sup>\*</sup> Medium followed by the same letter in column and capitalized on the line did not differ statistically among themselves by Tukey test at 5% probability.

The species *P. mosenii* and *P. gaudichaudianum* presented an essential oil yield that was higher in winter. As stated, *P. xylopioides* and *P. diospyrifolium* presented higher yield in summer.

The study on *P. gaudichaudianum* performed by (23) in a population in Santa Maria, Rio Grande do Sul State, found an average content of oil from fresh leaves of 0.38%, superior to that found in the present study, but without the effect of seasonality.

The chemical composition of essential oils is generally a characteristic of a given species and from the point of view of the chemical composition it is genetically and epigenetically controlled. The quantity, quality, and concentration of these species are influenced by the environmental components. Among the environmental factors that can be highlighted are, light intensity and photoperiod, the latitude, temperature (minimum and maximum average), soil (chemical and physical

properties), wind, and the availability of water, or even a combination of some of these subfactors and seasonality (26).

Studies conducted by (24) on the *Piper* genus in the Atlantic Forest, on the coast of Paraná, showed the influence of seasonality on the yield and the constituents of essential oils. (16), (27), (29), (17), (21), (20), and (19), also identified the influence of seasonality on the chemical profile of the oils analyzed.

The chemical composition of the essential oil of the studied species identified 78 constituents, corresponding to an average of 90% of chemical compounds of the essential oil, in the identified samples (Table 3). The species *P. diospyrifolium* was the one that presented the highest number of compounds identified, with 55, followed by *P. gaudichaudianum* with 50, *P. xylopioides* with 35, and *P. mosenii* with 33 compounds.

Table 3 - Phytochemical analysis of essential oils of fresh samples of species collected during the winter and summer of 2016 in the Caiuá Ecological Station, North Diamond, Paraná State, Brazil.

Compounds	IR <sup>c</sup>	$\mathbf{I}\mathbf{R}^{t}$	P. mos	P. gau	P. xyl	P. dio
3Z-hexenol	849	850		0.31		
	931	932		5.09	4.14	1.01
α-pinene	931	932		(2.86)	(2.40)	(1.97)
	974	974		6.62	0.25	2.14
β-pinene	<i>71</i> 4	<i>)</i> / <del>-</del>		(3.88)	0.23	(1.46)
6-methyl-5-hepten-2-one	984	981		0.77		
	989	988		0.26	1.22	0.23
myrcene	767	700		0.20	(0.82)	(0.28)
	1004	1002			1.56	
α-phellandrene		1002			(1.16)	
o-cymene	1022	1022			0.13	
limonene	1026	1024	0.10	0.72		7.77
			0.10			(5.22)
	1027	1025			12.50	
β-phellandrene					(8.97)	0.24
(7) 0	1028	1032		0.18	0.89	0.24
(Z)-β-ocimene					(0.37)	(0.18)
(E) 0:	1045	1044			0.33	2.64
(E)-β-ocimene					(0.32)	(1.96)
γ-terpinene	1055	1054				(0.16)
linalool	1100	1095			(0.25)	(0.10)
illialooi	1100	1073			7.52	0.39
δ-elemene	1333	1335	1.27		(7.38)	(0.28)
				0.57		0.20
α-cubebene	1345	1345	0.16	(0.54)		(0.21)
cyclosativene	1366	1369		(0.55)		(0.40)
α-ylangene	1366	1373				(0.27)
gene				3.15		1.95
α-copaene	1370	1374	0.80	(4.33)		(1.88)
0.1	1205	1200			1.91	0.77
β-elemene	1387	1389	2.15	(0.10)	(1.91)	(0.57)
Z-caryophyllene	1399	1408				(0.54)
J 1 J				0.54		0.11
α-gurjunene	1403	1409		(0.44)		(0.14)
(T) : 1 H	1410		16.20	7.40	2.59	20.56
(E)-cariophyllene	1413	1417	16.39	(7.25)	(2.34)	(20.65
β-copaene	1422	1430	0.69			(0.41)
	1 422	1.421		4.94		0.44
β-curjunene	1422	1431		(5.13)		0.44
γ-clemene	1429		0.78			0.27
y-cicilicite	14 <i>2</i> 7	1434	0.76			(0.24)
α-guaiene	1433	1437	0.83			
	1431	1439		1.44	0.30	0.70
aromadendrene	1431	1437		(0.37)	(0.35)	(1.10)
6.9-guaiadiene	1437	1442		0.18		(0.14)

				(0.18)		
				(0.10)	0.40	
trans-muurola-3.5-dieno	1441	1451			(0.35)	
				2.32	0.50	1.66
α-humulene	1446	1452	3.21	(1.36)	(0.50)	(1.70)
		1102		5.33		
geranyl acetate	1451	1453		(5.66)		
geranyracetate				(3.00)	5.49	
allo-aromadendrene	1453	1458	0.13		(4.52)	0.35
		1430		0.28	(4.32)	0.21
cis-cadina-1(6).4-diene	1456	1461	0.23	(0.31)		(0.27)
		1401		5.24	0.12	(0.27)
day as 50 diams	1471	1471				(0.18)
dauca-5.8-diene	1.471	1.475		(6.26)	(0.14)	(1.60)
Trans-cadina-1(6).4-diene	1471	1475				(1.68)
	1474	1478		0.61	0.35	1.55
γ-muurolene		1.70		(0.56)	(0.39)	
germacrene D	1476		30.36	1.41	3.09	10.06
germaerene B	1170	1484	30.30	(1.73)	(1.75)	(6.84)
aristolechene	1477	1487			0.13	
0 1	1.450		0.60	0.13		0.37
β-selinene	1479	1489	0.69	(0.36)		(0.42)
				0.29		` /
δ-selinene	1484	1492	0.19	(0.36)		
Trans-muurola-4(14).5-dieno	1484	1493				0.14
	1484	1493				(0.15)
Epi-cubebol Epi-cubebol	1404	1493				(0.13)
1	1486	1406			0.58	
valencene		1496		4.15	(0.56)	0.42
biciclogermacrene	1490	4.700	13.46	4.15	29.83	8.42
		1500		(4.81)	(33.45)	(8.51)
	1495	1500		4.00	0.77	0.55
α-muurolene				(4.98)	(0.83)	(0.60)
(E.E)-α-farnesene	1499		0.94	0.34		
(2.2) & 1		1505	0.5	(0.48)		
germacrene A	1497	1508			(0.56)	
	1501	1511				0.76
$\delta$ -amorphene	1501	1511				(0.43)
				3.77	1.27	1.57
γ-cadinene	1508	1513	1.56	(4.05)	(1.93)	(1.92)
2 44				5.59	3.69	2.72
δ-cadinene	1518	1522	2.74	(7.12)	(3.87)	(3.22)
				0.57	(0.0.)	(= :==)
trans-cadina-1.4-diene	1526	1533		(0.83)		0.49
trans cadma 1.4 diene				0.31	0.20	0.53
α-cadinene	1532	1537		(0.37)	(0.26)	(0.34)
u-caumenc				0.19	(0.20)	
n aamaan 11 s 1	1543	1539				
α-copaen-11-ol				(0.14)		
,	1535	1544		0.25		
α-calacorene				(0.58)		
selina-3.7(11)-Diene	1536	1545				(0.40)
germacrene B	1548	1559	4.53	0.43		1.31

				(0.21)		(1.27)
(F) naralidal	1561		1.42	6.13	1.83	1.72
(E)-nerolidol	1561	1561	1.42	(7.11)	(1.78)	(0.23)
	1568	1574			4.52	
germacrene D-4-ol	1500	1374			(8.57)	
espatulenol	1569		1.86	3.20		0.77
		1577		(0.82)		(0.78)
caryophyllate oxide	1574	1500	1.60	3.14	(0.90)	2.30
1: 41-1 Di-41-4	1505	1582	0.15	(3.69)		(3.22)
diethyl Phthate	1585	1590	0.15			0.13
globulol	1582	1590		0.10	0.26 (0.22)	(0.25)
viridiflorol	1582		0.45	0.21		
		1592		(0.20)		
.c. 1. 1	1594	1600		0.50		
rosifoliol	1500	1.000	0.45	(0.51)		(0.40)
ledol	1593	1602	0.45			(0.40)
humulene epoxi II	1600	1608	0.16	0.86 (0.51)		(0.14)
1-10-di-Epi-cubenol	1607		0.17	0.67		0.16
1-10-di-Epi-cubenoi	1007	1618	0.17	(0.77)		(0.29)
10-Epi-γ-eudesmol	1612	1622				(0.13)
1-Epi-cubenol	1621		0.93	1.22		0.38
1-Epi-cubellol	1021	1627	0.93	(1.59)		(1.91)
	1623	1630		0.57		
γ-eudesmol	1023	1050		(0.61)		
	1621				1.13	
muurola-4.10(14)-dien-1-β-ol		1630			(0.92)	
F ' 1' 1	1634	1638				1.55
Epi-α-cadinol				0.10		(1.91)
caryophile-4(12). 8(13)-dien-5β-ol	1630	1639	0.24	0.18 (1.20)		
		1039		1.94	2.67	
Epi-α-muurolol	1634	1640	1.88	(2.21)	(3.40)	1.55
		1010		1.89	0.50	
α-muurolol	1639	1644	1.03	(2.17)	(0.68)	(0.34)
	1 < 4 4			0.66	` '	0.20
β-eudesmol	1644	1649		(0.76)		0.39
α-cadinol	1647		3.27	1.26	4.51	1.38
u-caumoi	104/	1652	J.41	(1.31)	(4.86)	(1.46)
	1680	1683		1.88		
Epi-α-bisabolol	1000	1005		(1.38)		
	Monoth	nerpenes (%)	3.03	12.50	21.87	17.50
		· ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	()	(4.44)	(18.75)	(15.22
Oxyge	enated monoth	nerpenes (%)	)		(2.12)	( )
			57.58	45.83	(3.13)	() 57.5(
	Sesqui	terpenes (%)	)	45.83 (55.56)	(53.12)	(58.70
			36.36	35.42	21.88	22.50
Oxyg	enated sesqui	terpenes (%)	)	(35.56)	(25.00)	(26.08
			()	(33.30)	(23.00)	(20.0

	Dhanvilanon ancida (0/)	3.03	2.08	3.13	2.50
	Phenylpropanoids (%)	()	(2.22)	()	()
_	Othors (0/)		4.17		
	Others (%)	()	(2.22)	()	
_	Total compounds identified (%)	94.89	91.66	96.07	80.21
	Total compounds identified (%)	() (89.21)	(96.73)	(81,23)	

IRc= retention index calculated; IRt= retention index of literature; P. mos= *Piper mosenii*; P. gau= *Piper gaudichaudianum*; P. xyl= *Piper xylopioides*; P. dio= *Piper diospyrifolium*. Values between brackets correspond to the composition of the essential oil samples collected in the summer.

The chemical composition of the species *P. mosenii* identified during the winter had an average of 57.58% of sesquiterpenes hydrocarbon, 36.36% of oxygenated sesquiterpenes, and 3.03% of monoterpenes and phenylpropanoids. A majority of the constituents of samples performed was identified as germacrene D (30.36%), (E)-caryophyllene (16.39%), and biciclogermancrene (13.46%), with the total compounds identified as 94.89%. During the summer an yield of oil was not observed for that species.

The chemical composition of essential oils is determined by genetic factors, however, according to (17), other factors may cause significant changes in the production of secondary metabolites. In fact, the secondary metabolites represent a chemical interface between plants and the environment. The stimuli arising from the environment in which the plant is located can redirect the metabolic pathway, causing a biosynthesis of different compounds. Among these factors, we can highlight the interactions between plant/microbial, plant/insect, and plant/plant; age and stage of development, abiotic factors such as brightness, temperature, rainfall, nutrition, time, and time of collection, as well as techniques during harvest and postharvest. It is valid to note that these factors can present correlations between themselves and not act in isolation. They can exercise joint influence on secondary metabolism, which causes a variation in the income and composition of the essential oil analyzed.

The yield of essential oil of the species *P. gaudichaudianum* collected during the winter season showed a proportion of 45.83% of sesquiterpenes hydrocarbon, 35.42% of oxygenated sesquiterpenes, 12.50% of monoterpenes, 2.08% phenilpropanoids, and 4.14% of other components (chain of 8 and 12 carbons), with the total of compounds identified of 91.66%. Samples collected in the summer showed a proportion of 55.56% of sesquiterpenes hydrocarbon, 35.56% of oxygenated sesquiterpenes, 4.44% of monoterpenes, 2.22% phenilpropanoids and for other components (chain of 8 and 12 carbons). The majority of constituents were (E)-caryophyllene (7.40%), β-pinene (6.62%), (E)-

nerolidol (6.13%),  $\delta$ -cadinene (5.59%), geranyl acetate (5.33%), Dauca-5,8-diene (5.24%),  $\alpha$ -pinene (5.09%),  $\beta$ -gurjunene (4.94%), and bicyclogermacrene with 4.15%.

Yet for *P. gaudichaudianum*, it was observed that in the summer there was a reduction in the proportion of monoterpenes (12.50% to 4.44%) and increase of hydrocarbons of sesquiterpenes (45.83% to 55.56%) and the total compounds identified was 89.21%. The main constituents identified were (E)-caryophyllene (7.25%),  $\beta$ -pinene (6.62%),  $\delta$ -cadinene (7.12%), (E)-nerolidol (7.11%), dauca-5,8-diene (6.26%), geranyl acetate (5.66%),  $\beta$ -gurjunene (5.13%), and  $\alpha$ -muurolene with 4.98%

Among the principal constituents, a majoritarian checked in some municipalities of the state of Rio Grande do Sul/Brazil for essential oils from leaves and inflorescences of P. gaudichaudianum, stand out αhumulene (13.3–37.5%), \(\beta\)-caryophyllene (10.4–19.3%), ß-pinene (5.6-7%),E-nerolidol (5.32–22.4%), Ecaryophyllene(8.9%), bicyclogermacrene (7.4%), selinene (3.7-15.7%),α-selinene (8.9-16.6%),alloaromadendrene (7.7%), linalool (4.8%) (3; 25). Already in a seasonal study in the municipality of Atalanta (Santa Catarina State) the constituents were observed to be  $\beta$ -caryophyllene (10.4–12.5%),  $\alpha$ caryophyllene (8.2–10.4%),  $\delta$ -selinene (5.4–6.9%),  $\delta$ cadinene (6.0–7.3%); E-nerolidol (3.0-7.2%), Z- $\beta$ -(5.5-5.6%),δ-cadinene (6.4-7.3%) and guaiene Valencene (4.0-5.6%) (22).

This difference in the chemical composition of the essential oil of the study population in comparison with the other studies carried out with the same *P. gaudichaudianum* extract could occur due to environmental, genetic, and biotic factors. Second, (31) stated, the chemical variability may be the result of the selection pressure of the environment and/or the ecology or characterizing a chemical adjustment to the environmental conditions prevalent.

The phytochemical composition of the essential oil from samples of *P. xylopioides* collected during the winter season showed a proportion of 53.12% of sesquiterpenes hydrocarbon, 21.88% of oxygenated

sesquiterpenes, 21.87% of monoterpenes, and 3.13% of phenylpropanoids, and a total of 96.07% of identified compounds. The majority of the constituents were bicyclogermacrene 29.83%,  $\beta$ -phellandrene 12.50%,  $\delta$ -elemene 7.52%, and alloaromadendrene 5.49%. In the summer we found a proportion of 53.12% of sesquiterpenes hydrocarbon, 25% of oxygenated sesquiterpenes, 18.75% of monoterpenes, and 3.13% of oxygenated monoterpenes, with a total of 96.73% of compounds identified. The proportion of constituents found by the majoritarian was bicyclogermacrene 33.45%,  $\beta$ -phellandrene 8.97%, and germacrene D-4-ol 8.57%.

The samples of *P. diospyrifolium* collected during the winter season showed the proportion of oxygenated monoterpenes of 57.50%, oxygenated sesquiterpenes of 22.50%, monoterpenes of 17.50%, and 2.50% of phenylpropanoids, with a total of 80.21% compounds identified. The majority of the constituents were (E)-

caryophyllene of 20.56%, germacrene D of 10.06%, bicyclogermacrene of 8.42%, and limonene of 7.77%. Already, during the summer season we found a proportion of 58.70% of sesquiterpenes hydrocarbon, 26.08% of oxygenated sesquiterpenes, and 15.22% of monoterpenes hydrocarbons, with a total of 81.23% compounds identified, and the proportion of constituents the majoritarian found was (E)-caryophyllene of 20.65%, germacrene D 6 of 68%, bicyclogermacrene of 8.51%, and limonene of 5.22%.

The predominance of sesquiterpenes in the genus *Piper* was also observed by (8), (6), (28), and (12).

In the winter/2016 station, the dendrogram (Fig. 1) demonstrates the similarity of the chemistry among the *Piper* genus studied, where three distinct groups of Euclidean distances were observed. This can also be explained by the genetic variability among species and populations (2).

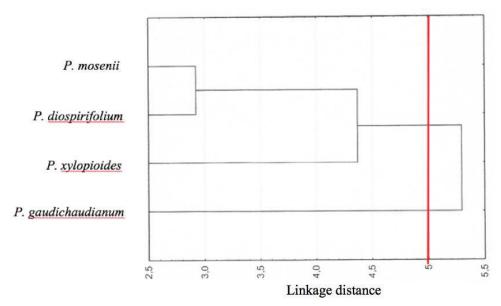


Fig.1: Dendogram of the Piper species showing functions of the chemical compounds in the essential oil of fresh samples collected in winter/2016, using the Euclidean distance.

The first grouping is formed by two Species: P. mosenii and P. diospyrifolium. This cluster is characterized by (E)-caryophyllene (16.39–20.56%), germacrene D, (10.06-30.36%) bicyclogermacrene (8.42-13.46%), γ-cadinene (1.56–1.57%), β-elemene (0.77– 2.15%), δ-cadinene (2.72–2.74%), (E)-nerolidol (1.42– 1.72%), Óx. caryophyllene (1.60–2.30%), epi- $\alpha$ -muurolol (1.55-1.88%), and  $\alpha$ -muurolol (0-1.03%). The cluster II includes the species P. xylopioides, which features (E)caryophyllene (2.59%),germacrene D (3.09%)bicyclogermacrene (29.83%), γ-cadinene (1.27%), βelemene (1.91%), δ-cadinene (1.27%), (E)-nerolidol (1.83%), epi- $\alpha$ -muurolol (2.67%), and  $\alpha$ -muurolol (0.5%). The last group consists of (E)-caryophyllene (7.40%), germacrene D (1.41%) bicyclogermacrene (4.15%),  $\gamma$ -cadinene (3.77%),  $\delta$ -cadinene (5.59%), (E)-nerolidol (6.13%), ox. caryophyllene (3.14%), epi- $\alpha$ -muurolol (1.94%), and  $\alpha$ -muurolol (1.89%) in the species P. gaudichaudianum.

To determine the degree of variations in the phytochemical, a principal component analysis (PCA) was performed using a correlation matrix of all chemical compounds (Table 4 and Fig. 2).

Table 4 – Eigen values and accumulated variance for factors obtained from principal components analysis, based on the chemical composition of the species of Piper, studied on the basis of the composition of its essential oils during the winter season/2016.

Compounds	Factors*				
	1	2	3		
(E)-cariofilene	0.015	-0.152	0.338		
germacrene D	0.521	0.327	-0.274		
biciclogermacrene	-0.019	0.653	-0.098		
γ-cadinene	0.459	-0.135	0.433		
ß-elenene	0.414	-0.137	-0.162		
δ-cadinene	-0.152	0.067	-0.584		
(E)-nerolidiol	-0.348	0.247	0.117		
Óx. cariofileno	-0.124	-0.502	-0.317		
Epi-α-muurolol	-0.223	0.105	0.325		
α-muurolol	0.130	-0.238	0.150		
α-cadinol	-0.347	-0.149	0.058		
Eigenvalues	5.83	3.35	1.82		
% of variance	56.64	33.44	16.55		
Cumulative %	56.64	90.08	100,00		

<sup>\*</sup> significance  $\geq 60\%$ 

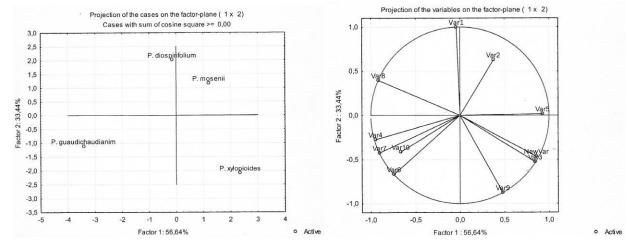


Fig.2 - Principal Component Analysis (PCAs) for the species of Piper based on chemical composition of essential oils of fresh samples collected in the winter season/2016. New Var: (E)-caryophyllene; Var1: germacrene D; Var2: bicyclogermacrene; Var3: γ-cadinene; Var4: β-eleneno; Var5: δ-cadinene; Var6: (E)-nerolidol; Var7: Óx. caryophyllene; Var8: epi-α-muurolol; Var9: α-muurolol; Var10: α-cadinol.

Results obtained by PCA, based on 11 chemical compounds, are shown in Figure 2 and Table 4. The three factors explain 100% of the accumulated variation in the data; the first two factors being considered the most important, as they described 90.08% of the accumulated variance (Table 4). The compounds germacrene D, γ-cadinene, and β-elemene, demonstrate the relevant contributions, with 56.64% of the variation in the principal components (PC 1). Bicyclogermacrene, germacrene D, and (E)-nerolidol are compounds that contributed, by explaining 33.44% of the variance of principal components (PC 2).

For summer 2016, the dendrogram (Fig. 3) is submitted to chemical similarity of the *Piper* genus where

three distinct groups of Euclidean distances were observed. The first grouping is formed only by the species P. mosenii, which showed an absence of the production of oil from this station. The cluster II includes the species P. gaudichaudianum and P. diospyrifolium, which present as constituents (E)-caryophyllene (7.25–20.65%), germacrene D (1.73–6.84%) Bicyclogermacrene (4.81–8.51%),  $\gamma$ -cadinene (7.12–3.22%),  $\beta$ -elemene (0.10–0.57%),  $\delta$ -cadinene (0.37–0.34%), (E)-nerolidol (7.11–0.23%), caryophyllene oxide (3.39–3.22%), Epi- $\alpha$ -muurolol (2.21%) and  $\alpha$ -muurolol (2.17–0.34%) and  $\alpha$ -cadinol (1,31–1,46%).

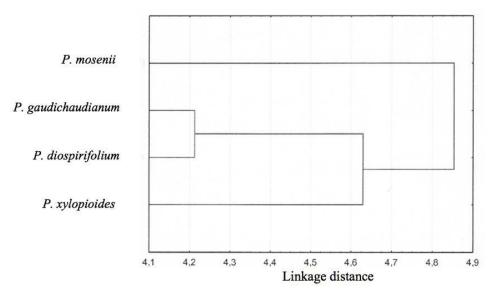


Fig.3 - Dendrogram for the Piper species and function of the chemical compounds of the essential oil of fresh samples collected in the summer/2016, using the Euclidean distance.

The last group consists of (E)-caryophyllene (2.34%), germacrene D (1.75%) bicyclogermacrene (33.45%),  $\gamma$ -cadinene (1.93%),  $\beta$ -elemene (1.91%),  $\delta$ -cadinene

(3.87%), (E)-nerolidol (1.78%), caryophyllene oxide (0.90%), epi- $\alpha$ -muurolol (3.40%),  $\alpha$ -muurolol (0.68%), and  $\alpha$ -cadinol (4.84%) in the species *P. xylopioides*.

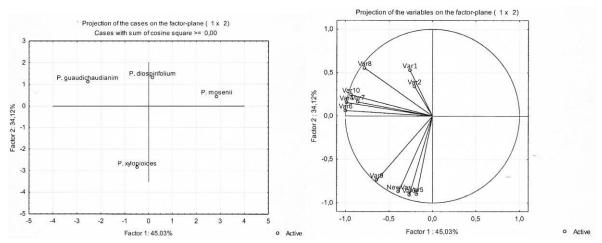


Fig.4 - Principal Component Analysis (PCAs) for the species of Piper based on chemical composition of essential oils of fresh samples collected in the summer season/2006. NewVar: (E)-caryophyllene; Var1: germacrene D; Var2: bicyclogermacrene; Var3: γ-cadinene; Var4: β-elemene; VAR5: δ-cadinene; Var6: (E)-nerolidol; Var7: Óx. cCaryophyllene; Var8: Epi-α-muurolol; Var9: α-muurolol; Var10: α-cadinol.

Results obtained by PCA based on the analyzed species and 11 chemical compounds are presented in Fig. 4 and Table 5. The three factors explain 100% of the accumulated variation in the data; the first two factors being considered most important, because they described 79.15% of accumulated variance. The  $\alpha$ -muurolol

compounds, (E)-nerolidol, and germacrene D, demonstrate the relevant contributions, with 45.03% of the variation for the principal components (PC 1). Epi- $\alpha$ -muurolol and  $\delta$ -cadinene are compounds that contributed by explaining 34.12% of the variance in the principal components (PC 2).

Table 5 – Eigen values and accumulated variance for factors obtained by principal components analysis (PCA) based on the chemical composition of the species of Piper studied, based on the composition of their essential oils during the summer season/2016.

Compounds	Factors*				
	1	2	3		
(E)-cariophyllene					
	0.117	0.093	0.120		
germacrene D	0.272	0.176	-0.470		
biciclogermacrene	-0.534	-0.606	-0.208		
γ -cadinene	0.010	0.071	-0.146		
ß-elenene	0.041	-0.061	-0.392		
$\delta$ -cadinene	-0.586	0.484	-0.281		
(E)-nerolidiol	0.278	-0.110	-0.273		
Óx. cariofilene	0.070	-0.154	-0.587		
Epi-α-muurolol	-0.253	0.520	-0.068		
α-muurolol	0.284	0.184	0.019		
α-cadinol	-0.235	-0.101	0.209		
Eigenvalues	4.953	3.753	2.293		
% of variance	45.03	34.12	20.85		
Cumulative %	45.03	79.15	100.00		

<sup>\*</sup> significance ≥ 60%

The different stations influenced both the quantity and the chemical compounds of the evaluated essential oils of the species of genus *Piper*. With the exception of *P*. Mosenii, compounds α-pinene, (E)-caryophyllene, aromadendrene, α-humulene, germacrene bicyclogermacrene, α-muurolene, γ-cadinene, δ-cadinene, α-cadinene, and (E)-nerolidol were found in other species in both stations of collection. The compounds of  $\alpha$ phellandrene, β-phellandrene, trans-muurola-3,5-diene, valencene, germacrene D-4-ol, and muurola-4,10(14)dien-1-β-ol were found only in the species P. xylopioides in the collection of two seasons.

Differences in chemical constituents can be justified by the regulation of gene expression of the enzymes involved in the biosynthetic route of terpenes. In addition, climatic conditions (collections at different seasons) contributed to the chemical characterization of the essential oil of the species analyzed.

#### IV. CONCLUSION

The results presented here demonstrate that the environmental factor of seasonality interfered with the levels and the average percentage of the chemical constituents of essential oils.

The studied species of genus *Piper* can be distinguished into three groups per workstation, in accordance with the composition of the essential oil of fresh leaves.

The chemical composition of the predominant species evaluated consisted of sesquiterpenes followed by

monoterpenes, with emphasis on (E)-caryophyllene, germacrene D, bicyclogermacrene,  $\alpha$ -pinene and  $\beta$ -pinene,  $\alpha$ -phellandrene,  $\beta$ -phellandrene, trans-muurola-3,5-diene, and Valencene; germacrene D-4-ol and muurola-4,10(14)-dien-1- $\beta$ -ol were found only in the species *P. xylopioides* in the two seasons of collection.

With the exception of *P. mosenii* compounds,  $\alpha$ -pinene, (E)-caryophyllene, aromadendrene,  $\alpha$ -humulene, germacrene D, bicyclogermacrene,  $\alpha$ -muurolene,  $\gamma$ -cadinene,  $\delta$ -cadinene,  $\alpha$ -cadinene, and (E)-nerolidol were found in the other species and in both workstation collections.

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