Research Article

Volatile Organic Compounds of Anchote Tuber and Leaf Extracted Using Simultaneous Steam Distillation and Solvent Extraction

Yenenesh Ayalew 1, Dargie Tsegay Berhe 1, Nigusse Retta 2, Gulelat Desse 3, Ali Mohammed 4, and Kyong Su Kim 5

1Dilla University, College of Agriculture & Natural Resources, Dilla, Ethiopia
2Addis Ababa University, College of Natural Sciences, Addis Ababa, Ethiopia
3Botswana University, Gaborone, Botswana
4Jimma University, College of Agriculture and Veterinary Medicine, Jimma, Ethiopia
5Chosun University, Department of Food and Nutrition, Gwangju 501-759, Republic of Korea

Correspondence should be addressed to Yenenesh Ayalew; yenenesh.ayalew29@gmail.com

Received 14 March 2022; Revised 28 July 2022; Accepted 18 August 2022; Published 13 September 2022

Academic Editor: Mitsuru Yoshida

Copyright © 2022 Yenenesh Ayalew et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Anchote (Coccinia abyssinica (Lam.) (Cogn)) is an endemic and potentially valuable crop of Ethiopia principally categorized under root and tuber crops, and its newly growing leaves along with the tendrils are also used as nutritious vegetable served after being cooked. Leaf and tuber powders were extracted for the first time to identify volatile organic compounds by simultaneous steam distillation and solvent extraction (SDE) and characterized using gas chromatography mass spectrometry (GC-MS). VOCs having an area percentage above 0.5% were used for identification analysis. From the results, thirty volatile flavor compounds from leaves and fifteen from tubers were identified with the total fraction yield of 770.57 mg/kg and 4536.91 mg/kg, respectively, and from the 30 compounds identified from leaf 16 were distinguished in each of the tested accessions. Ethyl acetate 90.47% (697.13 mg/kg) was detected in a higher amount exhibiting >1% peak area. The rest 6.03% (46.46 mg/kg) were minor quantities (<1%) of the total (770.57 mg/kg) volatile flavor fraction. Among the 15 identified compounds in the tuber, ethyl acetate was the only major compound that accounted together for 99.15% (4498.33 mg/kg) being reported in minor quantities (<1%). The SDE extraction and GC-MS analysis of anchote leaves and tubers successfully identified various volatile flavor compounds, which indicates that anchote was found to be a potential source of volatile flavor compounds that can be used as a food flavoring agent and in folk medicines. Thus, this study confirms that anchote leaf and tuber can be used for more specific and valuable applications in food and medicine industries.

1. Introduction

More than 1,000 low molecular weight (Mr) organic compounds are emitted from floral and vegetative parts of many plant species [1]. The release of volatiles from many vegetative organs has antimicrobial or antitherbivore activity and so could also act to protect valuable reproductive parts of plants from enemies [2, 3].

Volatile organic compounds (VOCs) have great importance in basic and applied research. Plant parts release VOCs into the atmosphere and the soil as a defense mechanism against herbivores and pathogens or as an attraction of pollinators and seed dispersers. In some plants, released VOCs may also act as wound sealers [4].

Flavor, taste, and sensorial quality of food, in which the aroma is formed by a complex group of chemical substances,
are influenced by VOCs. Over 90% of the natural emission of VOCs depends on plant species, and aroma results from the interplay of these emitted VOCs [4, 5]. Identification of VOCs for fragrance, pharmacologically active ingredients, and for flavor compounds is important for potential multipurpose functional use [6, 7]. VOCs formed biochemically during ripening of root bulbs, stem bark, fruits, and other vegetables [8], and among them, only a limited number are important for the characteristic aroma of the food, which are called key odorants [9].

The widely used simultaneous distillation extraction (SDE) technique is the most traditional extraction method, versatile, and relatively simple; however, the high temperature applied during the process, as well as long extraction time, might lead to artifact formation. In certain cases, SDE can provide bigger peaks of thermally stable compounds. SDE is widely applied for food products [10].

Anchote (Coccinia abyssinica (Lam.) (Cogn.)) is one of the important endemic crops principally grown for its edible tubers throughout the south and southwestern parts of Ethiopia. Moreover, its newly grown leaves along with tendrils are served as green vegetable after cooking. It has been used as a traditional medicine for a long period of time to treat various illnesses [11, 12]. Traditional use of anchote is for a number of medicinal reasons including fast mending of broken bones due to its good content of calcium [10] and for making lactating mothers healthier and stronger as well as to treat gonorrhea, tuberculosis, and tumor cancer [13]. Knowing the volatile flavor of anchote would be essential in discovering the key compounds responsible for the aroma and health effects. Therefore, the objective of the present study was to determine the active components and their contribution towards the overall aroma impact and medicinal values of anchote leaves and tubers.

2. Materials and Methods

2.1. Reagents. All reagents used in this experiment were purchased from Sigma Co. (St. Louis, USA) and Fisher Scientific (Waltham, Massachusetts, USA). The organic solvents: diethyl ether and n-pentane were redistilled using a wire spiral packed double distilling apparatus (Normschlif Geratebau, Wertheim, Germany). Besides, Milli-Q water was generated through a water purification system (Millipore Corporation, Bedford, USA), and the anhydrous Na2SO4 was used for dehydration of organic solvents after burning overnight at 650°C in a furnace (F 6000, Barnstead Thermolyne Co., IA, USA) and allowed to cool down in desiccators.

2.2. Extraction of Volatile Organic Compounds. Three anchote accessions, “223090,” “220563,” and “230566,” were selected for volatile flavor profile studies. Three healthy tubers from each accession were harvested from Debre Zeit Agricultural Research Center (DZARC) experimental field in June 2014, washed, peeled, and sliced to small pieces and mixed thoroughly in order to prepare 400 g of samples. Similarly, 200 g bunches of newly growing tips of leaves were first cleaned and chopped into small pieces. Both tuber and leaf samples were oven dried at 105°C to a constant weight in a paper bag by a hot air oven (DHG-9055A, Memment, Germany) set at 105°C. The oven dried anchote leaf and tuber samples were then milled using an electrical miller (FW 100, Yusung Industrial Ltd, China) to fine powder to pass a mesh size of 0.425 mm. Finally, the dried and powdered samples were packed in paper bags and sealed in an airtight polyethylene bag and labeled before storing in a refrigerator set at 4°C for further analysis.

Anchote leaf and tuber samples (30 g) were homogenized in a multimixer blender (MR 350CA, Braun, Spain) and mixed with 500 ml of distilled water which were adjusted to pH 7 using 1 N NaOH and 1 N HCl. After adjusting the pH, 10 ml of n-butyl benzene (110 ppm in n-pentane) was added as an internal standard. The resultant slurry was used for extraction of volatile organic compounds with 100 ml redistilled n-pentane: diethyl ether (1:1, v/v) mixture, using a simultaneous distillation extraction apparatus under atmospheric pressure according to MacLeod et al. [14] for three hrs. The solvent containing the compound extract was dehydrated for 12 hrs using 10 g anhydrous Na2SO4 and then concentrated to approximately 1.5 ml using the Vigeux column (250 ml, Normschlif, Wertheim, Germany). This extract was further concentrated to 0.5 ml under a gentle stream of N2 gas and used for gas chromatography mass spectrometry (GC-MS) analysis (Figure 1).

2.3. Establishment of Retention Index of N-Alkane. The standard value of retention index (RI) was determined by n-alkane mixture (C7 ~ C22). 1 μl mixture of n-alkanes was analyzed to determine the retention time (RT) by GC-MS analysis following the same conditions as mentioned in Table 1. The RI of each peak was established by basic program that substituted the RT of peak of n-alkane confirmed by GC-MS chromatogram.

Retention index was used as a parameter for checking of a solute from the chromatogram by comparing the retention time of both alkanes that appeared above and below the solute.

\[
RI_i = 100Z + 100 \left( \frac{\log VR (i) - \log VR (Z)}{\log VR (Z + 1) - \log VR (Z)} \right),
\]

where RIi is the retention index of compound i, VR(0), VR(Z), and VR(Z+1) are the retention time of standard alkanes (alkanes eluted before and after the substance of interest) which bracket the substance of interest, and factor Z contains the number of carbon eluted, e.g., Z + 1 and Z + 2.

By definition, the retention time of an alkane is the value as a multiple of the carbon number that the compound has to be unrelated with the column solid phase, the temperature of separation, and the requirements of other chromatography. Therefore, n-alkane was indicated as standard index for CH4 (RI = 100), C2H6 = (RI = 200), ..., CnH2n+2 (RI = 100n), and even anything in analysis column.

2.4. Chromatographic Analysis of Volatile Flavor Compounds. Chromatographic analysis was carried out using GC-MS (Model QP-2010, Shimadzu Co., Kyoto,
Japan) in electron impact ionization (EI) mode. The ionization voltage was 70 eV, and the temperatures of the ion source and injector were 230°C and 250°C, respectively. The mass spectrometer was scanned from 50 to 400 m/z. The separation was done by a capillary column, DB-WAX (60 m length × 0.25 mm diameter, 0.25 μm film thickness, Agilent J&W, USA). The program of oven temperature was initially started at 40°C (isothermal for 3 min), which was ramped to 180°C (isothermal for 5 min) at 2°C/min. Subsequently, it increases to 200°C (isothermal for 10 min) at 4°C/min and to 220°C (isothermal for 5 min) at 5°C/min. Finally, it reaches to 250°C (isothermal for 10 min) at 5°C/min. Helium was used as the carrier gas at a flow rate of 1 ml/min, and the sample injector volume was 1 μL using 1 : 100 split ratio.

2.5. Identification and Quantification of Volatile Flavor Compounds. Mass spectra of volatile organic compounds were identified with the aid of our own mass spectral data and those contained within the Wiley 7, NIST 05, and FFNSC 2.0 spectral libraries of the GCMS instrument. In addition, by the comparison of retention indices to the reference data [6, 15–28]. To calculate the relative response factor (RRF), the following formula was used:

\[
RRF = \frac{\text{Peak Area C/Conc. C}}{\text{Peak Area A/Conc. A}},
\]

where peak area C is the peak area of each component in the standard sample, Conc. C is the concentration of each component in the standard sample, peak area A is the peak area of the internal standard in the standard sample, and Conc. A is the concentration of the internal standard in the standard sample.

The quantitative analysis was carried out with the help of peak area percent of the internal standard (n-butylbenzene) using the formula [29]:

Table 1: Retention time of n-alkane mixture for GC-MS retention index.

| Alkanes | Retention time | Alkanes | Retention time |
|---------|----------------|---------|----------------|
| C_7     | 8.405          | C_15    | 62.875         |
| C_8     | 13.095         | C_16    | 68.777         |
| C_9     | 19.775         | C_17    | 74.637         |
| C_{10}  | 27.423         | C_{18}  | 81.002         |
| C_{11}  | 35.188         | C_{19}  | 86.341         |
| C_{12}  | 42.689         | C_{20}  | 92.825         |
| C_{13}  | 49.808         | C_{21}  | 98.567         |
| C_{14}  | 56.528         | C_{22}  | 104.059        |

Figure 1: Schematic diagram for analysis of volatile flavor compounds of anchote leaf and tuber samples.
Component content (mg/kg) = \( (C \times 1000 \text{ g}) / (A \times \text{RRF} \times B_p) \),

where \( A \) is the peak area of the internal standard in the anchote sample, \( B_p \) is the amount of sample, and \( C \) is the peak area of each component in the anchote sample.

3. Results and Discussions

3.1. Volatile Compounds of Anchote Leaf. The identified volatile components, percentage of their relative peak area, and concentrations are shown in Table 2. VOCs having an area percentage above 0.5% were used for identification analysis. In anchote leaf, 30 compounds were identified, among which 16 compounds were distinguished in each of the tested accessions. These compounds are ethyl acetate, acetoin, 1,1-diethoxyethene, 3-methyl-1-butanol (E)-2-hexenal, (Z)-3-hexen-1-ol, benzaldehyde, benzyl-alcohol, phenylacetalddehyde, phenethyl-alcohol, butyrophenone, 2-methoxy-4-vinylphenol, phytone, methyl-palmitate, dibutyl-phthalate, and methyl-linolenate. Ethyl acetate 90.47% (697.13 mg/kg) was detected in a higher amount followed by phenylacetalddehyde 1.88% (14.51 mg/kg) and (E)-hex-2-enal 1.62% (12.47 mg/kg), by exhibiting >1% peak area. They accounted together (93.97% (724.11 mg/kg)). The rest (6.03%) (46.46 mg/kg) were minor quantities (<1%) of the total (770.57 mg/kg) volatile flavor fraction. The VOCs identified in anchote leaves have various applications. A monoterpane compound linalool was presented in very low concentration in accession "220563." It is one of the major volatile components of several aromatic species used in foodstuffs as food additives and pharmacologically to cure a variety of ailments, being a sedative effect inducer, glutamatergic neurons inhibitor, anti-inflammatory, anticarcinogenic, and antiseptic [30]. Furfural has an aroma of almonds and is one of the components found in vanilla. Furfural has a wide variety of uses such as for flavoring food, as herbicide and fungicide, and affects yeast survival and biochemical enzyme activities [6]. Nonanal, 1-hexanol, (Z)-3-hexenol, linalool, and benzaldehyde were considered as important contributors to the aroma of fresh plum fruit [31]. 3-Methyl-1 butanol is a main ingredient in the production of banana oil, an ester found in nature and produced as flavoring in industry. (Z)-3-Hexen-1-ol is an essential aroma compound used in fruit and vegetable flavors as well as in perfumes. 1-Hexanol was used in perfume industry. Benzyl alcohol is produced naturally by many plants and commonly found in fruits and teas. It is also found in a variety of essential oils [32]. Hexanol occurs naturally after hydrolysis or enzymatic reduction reactions and was used in the flavor industry to produce fruity flavors. Benzaldehyde has a characteristic almond-like odor and is the primary component of bitter almond oil. Benzaldehyde is commonly employed to confer almond flavor to foods and scented products and sometimes used in cosmetics products [33]. Acetol or 1,1-diethoxyethene is a major flavoring component of distilled beverages, especially malt whisky. Phenylacetalddehyde is used as an ingredient in fragrances as well as in flavored cigarettes and beverages; its aroma is described as honey-like, sweet, rose, green, and grassy [34]. The aroma of ethyl acetate contributes towards the general perception of fruitiness. Dihydroactinidiolide has a sweet, tea-like odor and was used as a fragrance. Acetoin is used as a food flavoring in baked goods and as a fragrance. At very low concentrations, indole has a flowery smell and is a constituent of many flower scents such as orange blossoms and perfumes. 2-Methoxy-4-vinylphenol is an aromatic substance used as a flavoring agent and is known as one of the compounds responsible for the natural aroma of buckwheat [35]. Caryophyllene oxide, which is an oxygenated terpenoid, is also well known for its preservative characteristics in foods, drugs, and cosmetics. It has a significant central and peripheral analgesic, along with anti-inflammatory activity [36].

From the identified carbonyl compounds, phenylacetaldheyde accounted the highest amount 14.51 mg/kg (74.35%) followed by 1,1-diethoxyethene 2.78 mg/kg (14.27%), while acetoin and nonanal were quantified as 1.32 mg/kg (6.77%) and 0.90 mg/kg (4.60%), respectively. The alcohol group constituted 1.51% of the identified volatile compounds that were specified as 3-methyl-1-butanol (0.46%), [Z]-3-hexen-1-ol (0.61%), 1-hexanol (0.05%), benzyl alcohol (0.32%), 4-nonanol (0.04%), and linalool (0.04%). The remaining four functional groups such as alkanes, hydrocarbons, ketone, and terpene were detected at levels lower than 0.4%. Besides the identified functional groups, nine other volatile compounds were with no identified functional groups and categorized as miscellaneous. Among the miscellaneous (E)-2-hexenal (1.62%) which occupied the major position with >1%, the remaining compounds in content order were as follows: hexadecanoic acid <n->, phenethyl alcohol, methyl linolenate, butyrophenone, 2-methoxy-4-vinylphenol, phytone, (-)-caryophyllene oxide, and indole.

3.2. Volatile Compounds of Anchote Tuber. The identified VOCs in three accessions of anchote tuber that are listed according to their elution order on DB-WAX column with their number of concentrations are shown in Table 3. Fifteen volatile compounds were identified in anchote tubers from three accessions. Among the 15 identified compounds, ethyl acetate was the only major compound that accounted together for 99.15% (4498.33 mg/kg) of the total volatile flavor fraction (4536.91 mg/kg), and 0.85% (38.58 mg/kg) were reported in minor quantities (<1%).

Hexanal content is directly related to oxidative off-flavors, and the compound is easily detected because of its low odor threshold (5 ppb). Propyl acetate is commonly used in fragrances and as a flavor additive. Propyl acetate is synthesized via alcohol or acetic acid having a clear, volatile, and mobile liquid with a characteristic odor reminiscent of acetone and pears and was commonly used in fragrances and as a flavor additive. Typical odors of pyrazine compounds are responsible for the nut-like and peanut butter-like flavors, which are found in roasted barley, coffee, potato chips, and cocoa. 2,3,5-Trimethylpyrazine is categorized as cosmetic, flavor, and fragrance agents, and tetramethylpyrazine is an inhibitor of phosphodiesterase, which has been widely used for the treatment of cardiovascular diseases. Tetramethylpyrazine has also significant vascular protective
properties which have been used widely for the treatment of ischemic neural disorders and cardiovascular diseases [37]. Butyrophenones are widely used drugs for the treatment of psychose and are frequently encountered in forensic chemistry and clinical toxicology. Dibutyl phthalate is an artifact extracted from plastic which often is present in extracted samples from fruit [38, 39]. It is also identified from buckwheat honey [40].

The identified volatile organic compounds in anchote leaves so far belong to the chemical classes of alcohol (7), aldehydes (3), alkanes (2), carbonyl compound (4) esters (3), hydrocarbons (1), ketones (1), terpenes (1), and miscellaneous (8) (Table 4). Esters were dominant with the highest proportion of relative peak area (91.34%) of the emitted volatile organic compounds. Ethyl acetate accounted 99.05% among the ester content (703.84 mg/kg), whereas the remaining percentage was shared by methyl palmitate (0.25%) and dibutyl phthalate (0.71%). Carbonyl compounds were the second major group, accounting 2.53% of the relative peak area. Anchote tubers so far belong to the chemical classes of alcohol (1), aldehydes (1), alkanes (1), carbonyl compounds (3), esters (2), heterocyclic compounds (1), unknown (1), and miscellaneous (5) that are present in Table 4. The order of concentration for the identified functional groups is as follows: esters > carbonyl compounds > alcohols > alkanes > aldehydes > heterocyclic compounds.

| No. | Compound name          | MF | Area% | 223090 mg/kg | RI | Area% | 220563 mg/kg | RI | Area% | 230566 mg/kg | RI |
|-----|------------------------|----|-------|--------------|----|-------|--------------|----|-------|--------------|----|
| 1   | Ethyl acetate C\(_{4}H_{8}O_{2}\) | 4.52 | 19.58 | 606           | 66.58 | 353.49 | 824           | 4.13 | 18.17 | 606           | 66.58 |
| 2   | Ethyl acetate C\(_{4}H_{8}O_{2}\) | 0.10 | 0.45 | 899           |      | —     |              |      | —     |              |     |
| 3   | Ethyl acetate C\(_{4}H_{8}O_{2}\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |
| 4   | Acetoin C\(_{4}H_{8}O_{2}\) | 0.10 | 0.44 | 710           | 0.11 | 0.53 | 710           | 0.08 | 0.35 | 710           |      |
| 5   | 1,1-Diethoxyethane C\(_{4}H_{8}O_{2}\) | 0.19 | 0.81 | 727.22        | 0.17 | 0.87 | 727           | 0.25 | 1.10 | 727           |      |
| 6   | 3-Methyl-1-butanol C\(_{8}H_{12}O\) | 0.28 | 1.19 | 736           | 0.19 | 0.98 | 736           | 0.31 | 1.36 | 736           |      |
| 7   | Hexanal C\(_{8}H_{12}O\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |
| 8   | Furfural C\(_{6}H_{12}O\) | 0.18 | 0.79 | 837           | 0.15 | 0.78 | 837           |      | —     |              | —   |
| 9   | (E)-2-Hexenal C\(_{8}H_{10}O\) | 0.98 | 4.25 | 856           | 0.79 | 3.96 | 857           | 0.97 | 4.26 | 856           |      |
| 10  | (Z)-3-Hexen-1-ol C\(_{8}H_{10}O\) | 0.46 | 1.98 | 858           | 0.27 | 1.38 | 859           | 0.31 | 1.35 | 858           |      |
| 11  | Ethyl benzene (EB) C\(_{10}H_{12}O\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |
| 12  | 1-Hexanol C\(_{6}H_{12}O\) | 0.08 | 0.35 | 872           | —   | —     |              | —   | —     |              | —   |
| 13  | Benzaldehyde C\(_{7}H_{12}O\) | 0.11 | 0.47 | 965           | 0.10 | 0.53 | 965           | 0.11 | 0.50 | 965           |      |
| 14  | Benzyl alcohol C\(_{7}H_{8}O\) | 0.10 | 0.41 | 1036          | 0.13 | 0.68 | 1036          | 0.31 | 1.37 | 1036          |      |
| 15  | Phenylacetaldehyde C\(_{10}H_{16}O\) | 0.60 | 2.60 | 1047          | 1.49 | 7.49 | 1047          | 0.77 | 3.40 | 1047          |      |
| 16  | Phenylacetaldehyde C\(_{10}H_{16}O\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |
| 17  | Butylbenzene C\(_{14}H_{14}O\) | 7.70 | 33.33 | 1059         | 6.61 | 33.33 | 1059         | 7.57 | 33.33 | 1059         |      |
| 18  | 4-Nonanol C\(_{9}H_{20}O\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |
| 19  | Linalool C\(_{10}H_{16}O\) | 0.08 | 0.35 | 1105          | —   | —     |              | —   | —     |              | —   |
| 20  | Nonanal C\(_{10}H_{18}O\) | 0.11 | 0.49 | 1113          | 0.34 | 1.71 | 1113          | 0.35 | 1.54 | 1113          |      |
| 21  | Phenyl alcohol C\(_{10}H_{16}O\) | 0.10 | 0.40 | 1255          | 0.16 | 0.79 | 1255          | 0.30 | 1.25 | 1255          |      |
| 22  | Butyrophenone C\(_{10}H_{16}O\) | 0.08 | 0.34 | 1292          |      | —     |              | —   | —     |              | —   |
| 23  | Indole C\(_{10}H_{14}N\) | 0.12 | 0.60 | 1588          | —   | —     |              | —   | —     |              | —   |
| 24  | 2-Methoxy-4-vinylphenol C\(_{10}H_{12}O\) | 0.07 | 0.30 | 1310          | 0.14 | 0.70 | 1310          | 0.11 | 0.47 | 1310          |      |
| 25  | Beta-ionone C\(_{13}H_{20}O\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |
| 26  | Dihydroactinidiolide C\(_{11}H_{12}O_{2}\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |
| 27  | (-)-Caryophyllene oxide C\(_{15}H_{24}O\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |
| 28  | Phytone C\(_{14}H_{24}O\) | 0.05 | 0.23 | 1841          | 0.12 | 0.59 | 1840          | 0.09 | 0.39 | 1799          |      |
| 29  | Methyl palmitate C\(_{19}H_{32}O_{2}\) | 0.10 | 0.43 | 1922          | 0.18 | 0.93 | 1922          | 0.09 | 0.38 | 1900          |      |
| 30  | Dibutyl phthalate C\(_{22}H_{46}O_{2}\) | 0.28 | 1.21 | 1950          | 0.41 | 2.08 | 1950          | 0.38 | 1.69 | 1899          |      |
| 31  | Hexadecanoic acid C\(_{20}H_{40}O_{2}\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |
| 32  | Methyl linolenate C\(_{19}H_{32}O_{2}\) | 0.19 | 0.81 | 2097          | 0.23 | 1.18 | 2096          | 0.29 | 1.28 | 1999          |      |
| 33  | Docosane C\(_{22}H_{46}\) | —   | —     | —             | —   | —     |              | —   | —     |              | —   |

*Molecular formula, * retention index, and * internal standard.
Most esters have a fruity and floral flavor and may contribute to the aroma and flavor. Carbonyl compounds are widely found in food products, such as fried foods and beverages, and it is caused by the oxidation of fatty acids and higher alcohols, Strecker degradation, aldol condensation, or Maillard reactions [41]. 2-Pentanol was the only compound belonging to the alcohol group constituting 0.22% that is lower than 1% considered as a minor compound.

Aldehydes are particularly important in relation to flavor alterations and from a toxicological perspective. A particular property of the aldehyde volatile oils is their insect repellent activity due to very strong scent [6].

| No. | Compound name        | MF          | Area% | mg/kg  | RI   | Area% | mg/kg  | RI   |
|-----|----------------------|-------------|-------|--------|------|-------|--------|------|
| 1   | 2-Pentanol           | C₅H₁₂O     | 0.20  | 8.09   | 701  | 0.19  | 1.98   | 701  |
| 2   | Heptane              | C₇H₁₄       | —     | —      | —    | —     | 0.14   | 0.76 |
| 3   | Acetoin              | C₅H₁₀O₂    | 0.12  | 4.62   | 710  | 0.12  | 1.26   | 710  |
| 4   | Pyrrole              | C₆H₁₂N     | 0.14  | 0.76   | 699  |
| 5   | 1,1-Diethoxyethane   | C₆H₁₄O₂    | 0.13  | 5.15   | 727  | 0.25  | 2.53   | 727  |
| 6   | Hexanal              | C₆H₁₂O     | 0.05  | 0.26   | 750  |
| 7   | Ethyl acetate        | C₄H₈O₂     | 0.08  | 3.10   | 976  | 0.22  | 2.30   | 899  |
| 8   | Benzaldehyde         | C₇H₈O     | 0.10  | 0.54   | 965  |
| 9   | 2-Pentyl furan       | C₅H₁₀O₂    | 0.10  | 0.56   | 991  |
| 10  | n-Propyl acetate     | C₅H₁₀O₂    | —     | —      | —    |
| 11  | 2,3,5-Trimethyl pyrazine | C₆H₉N₂ | —     | —      | —    | 0.06  | 0.30   | 1001 |
| 12  | Tetramethyl pyrazine  | C₈H₁₂N₂   | —     | —      | —    | 0.38  | 2.11   | 1084 |
| 13  | Butyrophenone        | C₁₀H₁₂O    | 0.03  | 0.24   | 3.97 |
| 14  | Dibutyl phthalate    | C₁₀H₁₂O₄   | —     | —      | —    |

**Table 1: Volatile flavor compounds identified in anchote tuber.**

| No. | Compound name        | MF          | Area% | mg/kg  | RI   | Area% | mg/kg  | RI   |
|-----|----------------------|-------------|-------|--------|------|-------|--------|------|
| 1   | 2-Pentanol           | C₅H₁₂O     | 0.20  | 8.09   | 701  | 0.19  | 1.98   | 701  |
| 2   | Heptane              | C₇H₁₄       | —     | —      | —    | —     | 0.14   | 0.76 |
| 3   | Acetoin              | C₅H₁₀O₂    | 0.12  | 4.62   | 710  | 0.12  | 1.26   | 710  |
| 4   | Pyrrole              | C₆H₁₂N     | 0.14  | 0.76   | 699  |
| 5   | 1,1-Diethoxyethane   | C₆H₁₄O₂    | 0.13  | 5.15   | 727  | 0.25  | 2.53   | 727  |
| 6   | Hexanal              | C₆H₁₂O     | 0.05  | 0.26   | 750  |
| 7   | Ethyl acetate        | C₄H₈O₂     | 0.08  | 3.10   | 976  | 0.22  | 2.30   | 899  |
| 8   | Benzaldehyde         | C₇H₈O     | 0.10  | 0.54   | 965  |
| 9   | 2-Pentyl furan       | C₅H₁₀O₂    | 0.10  | 0.56   | 991  |
| 10  | n-Propyl acetate     | C₅H₁₀O₂    | —     | —      | —    |
| 11  | 2,3,5-Trimethyl pyrazine | C₆H₉N₂ | —     | —      | —    | 0.06  | 0.30   | 1001 |
| 12  | Tetramethyl pyrazine  | C₈H₁₂N₂   | —     | —      | —    | 0.38  | 2.11   | 1084 |
| 13  | Butyrophenone        | C₁₀H₁₂O    | 0.03  | 0.24   | —    | 0.24  | —      | —    |
| 14  | Dibutyl phthalate    | C₁₀H₁₂O₄   | —     | —      | —    |

**Table 4: Content of functional groups in identified volatile components from anchote leaves and tubers.**
followed by carbonyl compounds and alcohols in both leaf and tuber samples.

4. Conclusion
The study of volatile organic compounds in the leaf and tuber parts of the three anchote accessions showed that there are several bioactive volatile components present, which could be isolated and used for various purposes. In general, this study confirms the potential of anchote for various biochemical applications in foods and in folk medicine. Therefore, further studies on the extraction and structure elucidation of various important VOCs from different parts of anchote are essential to promote effective utilization of underexploited genetic resources for more specific and valuable applications in the field of human health and food industries. Since the VOC extraction of anchote tuber and leaf was done for only three accessions, it was difficult to identify several new VOCs using SDE extraction and GC-MS analysis.

Data Availability
The data that support the findings of this study are available upon request to the corresponding author.

Disclosure
The full dissertation has previously been presented [41] as a preprint.

Conflicts of Interest
The authors declare that there is no conflict of interest.

Authors’ Contributions
The authors confirm contribution to the paper as follows: study conception and design were contributed by Y.A.G., A.M., G.D., and N.R. Data collection was contributed by Y.A.G. and D.T.B. Analysis and interpretation of results were contributed by Y.A.G. and D.T.B. Draft manuscript preparation was contributed by Y.A.G. and D.T.B. All authors reviewed the results and approved the final version of the manuscript.

Acknowledgments
This work is financially supported by the Addis Ababa University, Ethiopia.

References
[1] N. Dudareva, E. Pichersky, and J. Gershenzon, “Biochemistry of plant volatiles,” Update on Biochemistry of Plant Volatiles, vol. 135, no. 4, pp. 1893–1902, 2004.
[2] M. Friedman, P. R. Henika, and R. E. Mandrell, “Bactericidal activities of plant essential oils and some of their isolated constituents against Campylobacter jejuni, Escherichia coli, Listeria monocytogenes, and Salmonella enterica,” Journal of Food Protection, vol. 65, no. 10, pp. 1545–1560, 2002.
[3] K. A. Hammer, C. F. Carson, and T. V. Riley, “Antifungal activity of the components of Melaleuca alternifolia (tea tree) oil,” Journal of Applied Microbiology, vol. 95, no. 4, pp. 853–860, 2003.
[4] M. E. Maffe, J. Gertsch, and G. Appendino, “Plant volatiles: production, function and pharmacology,” Natural Product Reports, vol. 28, no. 8, pp. 1359–1380, 2011.
[5] R.-S. Liu, G. H. Jin, D. R. Xiao, H. M. Li, F. W. Bai, and Y. J. Tang, “Screening of the key volatile organic compounds of _Tuber melanosporum_ fermentation by aroma sensory evaluation combination with principle component analysis,” Scientific Reports, vol. 5, no. 1, p. 17954, 2015.
[6] R. Gyawali and K.-S. Kim, “Bioactive volatile compounds of three medicinal plants from Nepal,” Kathmandu University Journal of Science, Engineering and Technology, vol. 8, no. 1, pp. 51–62, 2012.
[7] J. Panten and H. Surburg, “Flavors and fragrances, 1. general aspects,” Ullmann’s Encyclopedia of Industrial Chemistry, vol. 15, pp. 1–9, 2000.
[8] N. Khan, Metals analysis, biochemical properties and irradiation Effects on volatile flavor profile of major aromatic spices from pakistan, Chosun University, Gwangju, Korea, 2014.
[9] A. Plotto, C. A. Margaria, K. L. Goodner, R. Goodrich, and E. A. Baldwin, "Odour and flavour thresholds for key aroma components in an orange juice matrix: terpenes and aldehydes," Flavour and Fragrance Journal, vol. 19, no. 6, pp. 491–498, 2004.
[10] M. N. Wieczorek, M. Majcher, and H. Jelč, “Comparison of three extraction techniques for the determination of volatile flavor components in broccoli,” Food, vol. 9, no. 4, p. 398, 2020.
[11] B. Endashaw, “Study on actual situation of medicinal plants in Ethiopia,” Japan Association for International Collaboration of Agriculture and Forestry, vol. 2, pp. 1–9, 2007.
[12] G. Afera and D. Haile, "Yield and nutrient concentration of Anchote [Coccinia abyssinica (Lam.) Cogn.] affected by harvesting dates and in-situ storage," African Journal of Crop Science, vol. 3, no. 5, pp. 156–161, 2015.
[13] H. Fekadu, F. Beyene, and G. Desse, “Effect of traditional processing methods on nutritional composition and anti-nutritional factors of anchote (Coccinia Abyssinica (Lam.) Cogn) tubers grown in Western Ethiopia,” Journal of Food Processing & Technology, vol. 4, no. 7, p. 249, 2013.
[14] A. J. MacLeod and N. M. Pieris, “Volatile flavor components of wood apple (Feronia limonia) and a processed product,” Journal of Agricultural and Food Chemistry, vol. 29, no. 1, pp. 49–53, 1981.
[15] A. Sayaslan, O. K. Chung, P. A. Seib, and L. M. Seitz, “Volatile compounds in five starches,” Cereal Chemistry Journal, vol. 77, pp. 248–253, 2000.
[16] L. Castro-Vázquez, M. S. Pérez-Coello, and M. D. Cabedoio, “Analysis of volatile compounds of rosemary honey. Comparison of different extraction techniques,” Chromatographia, vol. 57, no. 3–4, pp. 227–233, 2003.
[17] D. S. Garrutti, M. R. B. Franco, M. A. A. P. Da Silva, N. S. Janzantti, and G. L. Alves, “Evaluation of volatile flavour compounds from cashew apple (Anacardium occidentale L) juice by the osme gas chromatography/olfactometry technique,” Journal of the Science of Food and Agriculture, vol. 83, no. 14, pp. 1455–1462, 2003.
[18] M. Zhu, E. Li, and H. He, “Determination of volatile chemical constituents in tea by simultaneous distillation extraction, vacuum hydrodistillation and thermal desorption,” *Chromatography*, vol. 68, no. 7-8, pp. 603–610, 2008.

[19] R. Gyawali and K. Kim, “Volatile organic compounds of medicinal values from Nepalese *Acorus calamus* L.,” *Kathmandu University Journal of Science, Engineering and Technology*, vol. 5, pp. 51–65, 2009.

[20] H. Y. Seo, S. L. Shim, K. Y. Ryu et al., “Analysis of volatile compounds and enantiomeric separation of chiral compounds of dried sancho (*Zanthoxylium schinifolium*)”, *Food Science and Biotechnology*, vol. 18, pp. 18–24, 2009.

[21] S. L. Shim, I. M. Hwang, K. Y. Ryu et al., “Effect of γ-irradiation on the volatile compounds of medicinal herb, *Paoniae Radix*,” *Radiation Physics and Chemistry*, vol. 78, no. 7-8, pp. 665–669, 2009.

[22] H. Cheng, “Volatile flavor compounds in yogurt: a review,” *Critical Reviews in Food Science and Nutrition*, vol. 50, no. 10, pp. 938–950, 2010.

[23] I. Jerković, D. Kovačević, D. Šubarić, Z. Marijanović, K. Mastanjević, and K. Suman, “Authentication study of volatile flavour compounds composition in Slavonian traditional dry fermented salami "kulen,” *Food Chemistry*, vol. 119, no. 2, pp. 813–822, 2010.

[24] D. Jeon, S. Lee, J. Kim et al., “Comparison of volatile components among fresh and semi-fermented leaf of zukro tea,” *Journal of the Korean Tea Society*, vol. 18, pp. 28–36, 2012.

[25] H. Song, S. Shim, S. Lee, D. Kim, and K. Kim, “Analysis of volatile organic compounds of ‘Fuji’ apples following electron beam irradiation and storage,” *Radiation Physics and Chemistry*, vol. 81, no. 8, pp. 1084–1087, 2012.

[26] I. Jerković, D. Gašo-Sokač, H. Pavlović et al., “Volatile organic compounds from *Centaurom erythraea* rafín (Croatia) and the antimicrobial potential of its essential oil,” *Molecules*, vol. 17, no. 2, pp. 2058–2072, 2012.

[27] L. F. Cuevas-glory, J. A. Pino, L. S. Santiago, and E. Sauri-Duch, “A review of volatile analytical methods for determining the botanical origin of honey,” *Food Chemistry*, vol. 103, no. 3, pp. 1032–1043, 2007.

[28] M. D’Auria and R. Racioppi, “The effect of drying of the composition of volatile organic compounds in *Rosmarinus officinalis*, *Laurus nobilis*, *Salvia officinalis* and *Thymus serpyllum*. A HS-SPME-GC-MS Study,” *Journal of Essential Oil Bearing Plants*, vol. 18, no. 5, pp. 1209–1223, 2015.

[29] Y. Ayalew, “Nutritional and phytochemical evaluation of anchote (*Coccinia abyssinica*) (Lam.) (Cogn.) accessions to promote its contribution for food security and medicinal use,” Addis Ababa, Ethiopia, 2016http://etd.aau.edu.et/bitstream/handle/123456789/18164/Yenenesh%20Ayalew%20%202016.pdf?isAllowed=y&sequence=1.

[30] A. T. Peana, P. S. D’Aquila, F. Panin, G. Serra, P. Pippia, and M. D. L. Moretti, “Anti-inflammatory activity of linalool and linalyl acetate constituents of essential oils,” *Phytomedicine*, vol. 9, no. 8, pp. 721–726, 2002.

[31] J. A. Pino and C. E. Quijano, “Study of the volatile compounds from plum (*Prunus domestica* L. cv. Horvin) and estimation of their contribution to the fruit aroma,” *Food Science and Technology*, vol. 32, pp. 76–83, 2012.

[32] M. J. O’Neil, *The merck index: an encyclopedia of chemicals, drugs, and biologicals*, RSC Publishing, 2013.

[33] A. Andersen, “Final report on the safety assessment of benzaldehyde,” *International Journal of Toxicology*, vol. 25, 1_supplement, pp. 11–27, 2006.

[34] C. Kohlpainger, M. Schulte, F. Jürgen, P. Lappe, W. Jürgen, and G. Frey, “Aldehydes, aliphatic,” in *Ullmann’s encyclopedia of industrial chemistry*, 2008.

[35] D. Janež, D. Kantar, S. Kreft, and H. Prosen, “Identification of buckwheat (*Fagopyrum esculentum* Moench) aroma compounds with GC-MS,” *Food Chemistry*, vol. 112, no. 1, pp. 120–124, 2009.

[36] M. J. Chavan, P. S. Wakte, and D. B. Shinde, “Analgesic and anti-inflammatory activity of Caryophyllene oxide from *Annona squamosa* L. bark,” *Phytomedicine*, vol. 17, no. 2, pp. 149–151, 2010.

[37] Y. Jiang, C. Liu, W. Chen, H. Wang, C. Wang, and N. Lin, “Tetramethylypyrazine enhances vascularization and prevents osteonecrosis in steroid-treated rats,” *BioMed Research International*, vol. 2015, Article ID 315850, 12 pages, 2015.

[38] T. Wolsk and K. Tambor, “Identification of honey volatile components by solid phase microextraction (SPME) and gas chromatography / mass spectrometry (GC/MS),” *Journal of Apicultural Science*, vol. 50, pp. 115–126, 2006.

[39] V. M. Osorio and Z. L. Cardeal, “Analytical methods to assess carbonyl compounds in foods and beverages,” *Journal of the Brazilian Chemical Society*, vol. 24, pp. 1711–1718, 2013.

[40] A. Bianco, A. Venditti, S. Foddai, C. Toniolo, and M. Nicoletti, “A new problem. Contamination of botanicals by phthalates. Rapid detection tests,” *Natural Product Research*, vol. 28, no. 2, pp. 134–137, 2014.

[41] A. Venditti, “What is and what should never be: artifacts, improbable phytochemicals, contaminants and natural products,” *Natural Product Research*, vol. 34, no. 7, pp. 1014–1031, 2020.