

Essential Oils: Composition, Applications and Analytical Considerations

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Introduction

Essential oils are complex mixtures of volatile organic compounds naturally synthesized by plants as part of their secondary metabolism. They are typically obtained through steam distillation, hydrodistillation, cold expression in the case of citrus peels, or other mechanical extraction processes. The resulting products are generally hydrophobic liquids, partially or completely insoluble in water, characterized by high volatility and a distinctive, intense aroma.

The term *essential* refers to their function as the essence of the plant, meaning the fraction responsible for its characteristic fragrance and biological activity. From a biological perspective, essential oils play multiple roles in plant physiology. They contribute to defense mechanisms against herbivores and pathogenic microorganisms, support protection against environmental stress factors, participate in inter-plant communication processes, and attract pollinators.

Chemical Composition

Essential oils are composed predominantly of terpenoid compounds, along with smaller fractions of phenylpropanoids and other aromatic molecules. Terpenes represent the major class and are constructed from isoprene units, each containing five carbon atoms. Terpenes are classified based on the number of such units: hemiterpenes (1 isoprene unit, 5 carbon atoms), monoterpenes (2 units, 10 carbon atoms), sesquiterpenes (3 units), diterpenes (4 units) up to 8 units.

Monoterpenes are generally the most abundant constituents in essential oils and are responsible for their high volatility. Sesquiterpenes and diterpenes, which have higher molecular weight, contribute to stronger and more persistent aromatic notes. The relative proportion of low and high molecular weight terpenes determines whether a plant secretion behaves as a volatile essential oil or as a semi-solid resin. Essential oils remain liquid at room temperature due to the predominance of low molecular weight volatile compounds. Many terpenoid compounds exhibit documented biological activity, including antimicrobial, antiviral, anti-inflammatory and antioxidant effects. Examples include essential oils derived from pine, lemon, rosemary and tea tree.

Properties and Uses

Essential oils exhibit a broad spectrum of biological activities that are directly related to their chemically

diverse composition, which includes terpenes, phenols, alcohols and other volatile compounds. Their lipophilic nature facilitates interaction with biological membranes, contributing to antimicrobial, antiviral, anti-inflammatory and antioxidant effects, as well as smooth muscle relaxation activity. Beyond their biological activity, essential oils are widely used in cosmetic and personal care formulations, natural preservation systems, environmental fragancing and household sanitation products, due to both their functional properties and characteristic aromas. In topical applications they are generally used in diluted form, since several essential oils may cause irritation or sensitization if applied undiluted, while certain citrus-derived oils may induce photosensitivity because of furanocoumarin content. In aromatherapy, inhalation through diffusion or steam systems stimulates olfactory pathways that can influence emotional and autonomic responses, with reported effects including stress reduction, sleep modulation, mood improvement and mild analgesic or antiemetic support.

Oxidation Mechanisms in Essential Oils

Although essential oils are often associated with antioxidant properties, their chemical composition makes them intrinsically susceptible to oxidative degradation. Unlike vegetable oils, which are lipid matrices composed mainly of triglycerides and unsaturated fatty acids, essential oils are not lipid systems. They are complex mixtures of volatile, low molecular weight compounds, predominantly unsaturated monoterpenes and sesquiterpenes. In vegetable oils, oxidation primarily involves unsaturated fatty acids and follows classical lipid autoxidation pathways, leading to the formation of lipid hydroperoxides and secondary aldehydic compounds. In contrast, oxidation in essential oils mainly concerns terpene molecules, which can readily react with atmospheric oxygen through radical-mediated autoxidation mechanisms. Monoterpenes such as limonene, linalool and α -pinene are particularly prone to oxidation. During storage and exposure to light, heat or oxygen, these compounds can form: terpene hydroperoxides, epoxides, aldehydes and ketones, secondary rearrangement products. For example, limonene can be converted into limonene hydroperoxides, linalool into linalool oxides, and pinene into pinene oxides. These oxidation products are not only responsible for changes in aroma profile, with the development

of off-notes and loss of freshness, but may also increase the sensitization potential of the oil. From a regulatory perspective, oxidized terpenes are of particular concern in the fragrance and cosmetic sectors. Several oxidation products have been identified as potential skin sensitizers and are subject to attention under IFRA standards and cosmetic safety assessments. Therefore, the monitoring of primary oxidation products is not only a matter of organoleptic quality but also of toxicological relevance.

The high volatility and chemical reactivity of terpene-rich matrices further complicate stability control. Oxidation may be accelerated by improper storage conditions, including exposure to oxygen, elevated temperatures and ultraviolet radiation. Packaging in dark glass containers, reduced headspace oxygen and controlled storage temperatures are commonly recommended to limit oxidative degradation.

For these reasons, the evaluation of oxidative stability represents a critical step in essential oil quality control and provides the scientific basis for the determination of analytical parameters such as Peroxide Value.

Official Methods for Chemical Analysis

The key parameters for monitoring essential oil quality are **Peroxide Value** and **Free Fatty Acids Value**, which provide information on oxidative degradation and chemical stability. Industry guidance is supported by organizations such as the **International Fragrance Association (IFRA)** and the **European Federation of Essential Oils**, which define analytical and safety frameworks for commercial applications.

The determination of **Peroxide Value** is commonly performed according to the **IFRA Analytical Method for Peroxide Value**, based on iodometric titration. The method quantifies hydroperoxides formed during the early stages of oxidation and expresses results as **milliequivalents of active oxygen per kilogram (meq O₂/kg)** or alternatively as **millimoles of peroxide per liter (mmol/L)**. The analysis is conducted in an acidic organic medium, typically a glacial acetic acid and isooctane mixture, followed by titration of liberated iodine with standardized sodium thiosulfate. Due to the volatility and reactivity of terpene-rich matrices, strict control of light exposure, reagent standardization and blank correction is required to ensure reproducibility.

The determination of **Free Fatty Acids Value** is performed according to **ISO 1242**, which defines an acid-base titration method for the quantification of free acidic constituents in volatile oils. The sample is dissolved in neutralized alcoholic solvent and

titrated with standardized potassium hydroxide, with results expressed as **mg KOH per gram of sample (mg KOH/g)**. Elevated acidity may indicate hydrolytic or oxidative degradation and poor storage conditions. The method requires careful handling of flammable solvents and alkaline reagents, and attention to evaporation losses due to the volatile nature of essential oils.

Given the high chemical reactivity of terpene-rich systems, minimizing oxidative exposure during storage and processing is essential. At the same time, quality control protocols should aim to reduce sample manipulation and analysis time while ensuring accurate and reliable monitoring of oxidative stability.

An alternative approach to analysis: CDR FoodLab® system

In contrast to conventional iodometric and acid-base titration methods, which require significant solvent volumes, glassware, controlled laboratory conditions and trained personnel, **CDR FoodLab®** provides a rapid and simplified alternative for the determination of key oxidative parameters: **Free Fatty Acids Value** and **Peroxide Value**.

Based on photometric analysis with ready-to-use reagents, the system delivers results within minutes, enabling immediate quality control at raw material receiving, during storage, or directly in production environments. The method requires only microquantities of sample, an important advantage when analyzing essential oils with intense intrinsic coloration, as the minimal sample volume prevents color interference and ensures reliable optical measurement. The absence of toxic organic solvents significantly improves operator safety and reduces environmental impact compared to classical titration techniques. The system does not require highly specialized personnel, making analytical control accessible even to small and medium-sized manufacturing operations. Its versatility allows application to both liquid oils and solid fats, covering a broad range of raw materials.



Comparison of Analytical Methods for Essential Oils

Feature	Traditional Official Methods (Titration)	CDR FoodLab® System
Methodology	Titration Iodometric titration (Peroxide Value) or Acid–base titration ISO 1242 (Free Fatty Acids Value).	Photometric Analysis Spectrophotometric reading.
Reagents & Solvents	Complex & Hazardous Requires toxic organic solvents (e.g., glacial acetic acid, isooctane), flammable solvents, and alkaline reagents (KOH).	Ready-to-use & Safer Pre-filled reagents. No toxic organic solvents are required.
Analysis Time	Lengthy Requires preparation, standardization of reagents, and blank correction.	Rapid Delivers results within minutes.
Sample Volume	Standard Requires significant solvent volumes and standard sample amounts.	Microquantities Requires only a very small amount of sample.
Interference	Risk of Color Interference Intense intrinsic coloration of essential oils can make titration endpoints difficult to see.	No Interference Minimal sample volume prevents color interference, ensuring reliable optical measurement.
Equipment & Skill	Specialized Requires glassware, controlled laboratory conditions, and trained personnel.	Simplified Does not require glassware or highly specialized personnel. Can be used in production environments.
Safety & Environment	Lower Safety Involves handling of hazardous chemicals and flammable solvents.	High Safety Significantly improves operator safety and reduces environmental impact.

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