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## Comparative studies on physicochemical properties of virgin coconut oil (VCO) with different coconut oils

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### Abstract

This study aims to compare the physicochemical properties of Virgin Coconut Oil (VCO), Refined, Bleached and Deodorized (RBD) oil and Cold Pressed Oil (CPO, Chekku oil). The physicochemical properties taken for the study includes moisture content, refractive index, saponification value, iodine value, acid value, unsaponifiable matter, polenske value and % free fatty acids. The physicochemical properties of all three samples obtained are in compliance with Asian and Pacific Coconut Community (APCC) standards. Of all the samples, VCO showed 1.3% FFA which was higher when compared, that it should be consumed only to a certain limit for cooking. VCO has very low moisture content of 0.11% where the shelf life of oil can be naturally preserved. The acid value of VCO was 2.7 mg KOH/g which has the highest amount of AV whereas for other samples it was less than 1.0 mg KOH/g. These findings can be used for chemical analysis and adulteration detection.

**Keywords:** Virgin coconut oil (VCO), commercial coconut oil samples, physicochemical properties, APCC standard, free fatty acids (FFA)

### 1. Introduction

Coconut oil is an edible oil that has been consumed in tropical countries for thousands of years (Gopala *et al.*, 2010) <sup>[14]</sup>. Over the last decade, the world production of coconut oil has been raised because of its edible characteristics. In the point of edible nature, there are two type of coconut oil, RBD (refined, bleached, and deodorized) which was used in major and the virgin coconut oil (VCO) where the consumption is comparatively low because of its high cost (Perera *et al.*, 2020) <sup>[20]</sup>. It is largely consumed for both edible and non-edible purposes that includes cooking, bakery, confectionary, pharmaceutical, infant foods, and cosmetics (Dia *et al.*, 2005; Gopala *et al.*, 2010) <sup>[11, 14]</sup>.

Commercial edible RBD coconut oil is derived from the kernel of *Cocos nucifera* L., a tropical plant (Gopala *et al.*, 2010) <sup>[14]</sup> from which dried copra was taken then extracted and the resulting crude oil is processed on an industrial scale by washing, bleaching and deodorization (F.M. Dayrit *et al.*, 2007) <sup>[8]</sup>. Coconut oil contains 92% saturated fatty acids (SFA) (Da Silva Lima & Block, 2019) <sup>[7]</sup> hence it is consumed less (Gopala *et al.*, 2010) <sup>[14]</sup>. However, research on populations that eat diets high in coconut oil have shown no harmful effects on the population's health (Thampan *et al.*, 2009). The major fatty acid of coconut oil is a medium-chain fatty acid (MCFA). MCFA serves as the source of energy and is easy to absorb, metabolize and store in the body (Schönfeld & Wojtczak, 2016) <sup>[24]</sup>. Lipid oxidation is a significant degradation mechanism that occurs in oils, thereby affecting the organoleptic properties of food products coupled with oil (Perera *et al.*, 2020) <sup>[20]</sup>. As coconut oil is composed of more saturated fatty acids, it is more resistant to oxidation and polymerization than the oils with unsaturated fatty acids (Da Silva Lima & Block, 2019) <sup>[7]</sup>. But when the oil is processed, there are certain thermal and chemical treatments that influence the natural quality and composition of fatty acids (Vidya & Seeja, 2016) <sup>[27]</sup>.

Virgin coconut oil (VCO) on the other hand, obtained from fresh and mature coconut kernels (12 months after pollination) (*Cocos nucifera* L.) by mechanical or natural means, with or without heat application, which does not alter the quality of the oil. VCO has not been exposed to chemical refining, deodorization and bleaching. In its natural state, it can be consumed without the need for further processing. Virgin coconut oil consists primarily of peroxidation-resistant medium chain triglycerides.

Colorless, sediment-free virgin coconut oil with a perfect, fresh coconut fragrance. It is free from rancid odor or taste (APCC, 2009; Nevin & Rajamohan, 2004) [3, 18]. The production can be of two methods which includes wet and dry methods (Ghani *et al.*, 2018) [13]. In the dry method, the kernel is dried to remove the moisture from microbial invasion occurring. The dried kernel is pressed mechanically by means of bulls or electric source known as cold pressed oil. In contrast, in the wet method the VCO is extracted from the coconut meat/kernel by either chilling and centrifugation, enzymatic or fermentation to destabilize the emulsion of coconut milk without requiring the process of drying (Raghavendra & Raghavarao, 2010) [22]. The lauric acid content, a medium chain fatty acid was present in high amount in the VCO that shows potential for anti-obesity treatments (Assunção *et al.*, 2009; F.M. Dayrit, 2015) [5, 9]. Due to the presence of higher MCFA they perform some specific functional properties (Savva & Kafatos, 2015; Srivastava *et al.*, 2018) [23, 25] that includes antiviral, antibacterial, antiplaque, antiprotozoal, healing, anti-inflammatory and anti-obesity effects (German & Dillard, 2004) [12]. And VCO also has several benefits for curing some minor illnesses (List, 2016) [16] such as diarrhea, skin inflammations, gastrointestinal problems, minor wounds, injures and swelling (Nevin & Rajamohan, 2006) [19]. The VCO therefore exhibits more benefits in terms of both functional and nutritional aspects than other commercial edible coconut oil. Recently, here grows the interest in the consumption of VCO for cooking and medicinal benefits (Ghani *et al.*, 2018) [13]. Due to its complicated processing steps, virgin coconut oil costs much more than other commercial coconut oils. Hence there is a risk of adulteration in VCO, so it was important to study the quality characteristics of VCO. Therefore, the subject of this paper is to focus on the comparative study of physicochemical properties of commercial edible RBD coconut oil, virgin coconut oil and cold pressed coconut oil.

## 2. Material and Methods

### 2.1 Raw material

The coconut oils samples that include virgin coconut oil (VCO) were purchased from Pavithra Kera product, Palakkad (Kerala), Refined, Bleached and Deodorized coconut oil (RBD) and cold pressed coconut oil (CPO) were purchased from the local market in the city. All solvents and chemicals used were of analytical grade.

### 2.2 Determination of physicochemical properties

The physicochemical properties were carried out for all the three coconut oil samples. All the tests were performed with reference to the Food Safety and Standards Authority of India (FSSAI), American Oil Chemists Society (AOCS), Asian Pacific Coconut Community (APCC) and Indian standards (IS).

#### 2.2.1 Moisture content (MC)

Moisture and volatile are an important factor to determine the oil quality. Moisture content (MC) for the different oils samples were determined based on the American Oil Chemists Society (AOCS) (Firestone *et al.*, 2009) method. About 5.0g of oil sample was weighed in a crucible with lid which was preheated, weighed and dried. It was then heated at 105 °C for about 24 hours until there is no change in the successive observations. The crucible with sample was then placed in the desiccator and allowed to cool to a room

temperature. The crucible with oil sample was then re-weighed. The moisture content were then calculated using the following formula:

$$\text{Moisture content(\%)} = \frac{\text{initial weight} - \text{final weight}}{\text{initial weight}} \times 100\%$$

#### 2.2.2 Refractive index at 40 °C (RI)

The refractive index (RI) of the oil samples were measured using precision Abbes refractometer based on AOAC official method and ISI having a measuring range of refractive index of 1.300-1.700 with the accuracy within  $\pm 0.0002$ . (Ghani *et al.*, 2018) [13]. The refractive index was calculated by sandwiching 1 or 2 drops of oil sample between illuminating and refracting prisms of Abbes refractometer by means of syringe. The sample was illuminated with monochromatic light from sodium vapor lamp and it was allowed to stand for 5min before taking the reading displayed on the screen.

#### 2.2.3 Saponification value (SV)

Coconut oil has a relatively high saponification value due to its high concentration of short and medium chain triglycerides. The saponification value (SV) of the oil samples were determined using the International Union of Pure and Applied Chemistry (IUPAC) method (Rigaudy & Klesney, 1992) and AOAC official method 920.160. Approximately 2.0g of sample was weighed into a clean dried Erlenmeyer flask and 25ml of 0.5N ethanolic KOH was added and the mixture was boiled in a reflux condenser for 60 min. The mixture was then cooled to a room temperature and 1% phenolphthalein solution as an indicator was added to the cooled mixture and subsequently titrated against 0.5N HCl until the color of the mixture changes from pink to colorless. The volume of HCl was recorded and represented as S. Similarly, the same was repeated for the blank, and the volume of HCl was noted as B. The saponification value was calculated using.

$$SV = \frac{(B-S) \text{ ml of HCl} \times 28.05}{\text{weight of sample (g)}}$$

#### 2.2.4 Iodine value (IV)

Iodine value (IV) for the oil samples were determined by using Wijs method (AOCS, 2004). Approximately 4.0g of sample was mixed with 20ml cyclohexane to dissolve the fat content; 25 ml of Wijs solution was then added. The flask was closed, and the solution was shaken for 30 min continuously. Simultaneously, 20ml aqueous KI solution (15% v/v) and 100 ml of water were added to the mixture. It was then titrated with 0.1N Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> until the disappearance of yellow color. Then a few drops of starch solution, turning the solution to blue, were added and the titration continued until the blue color vanished. The volume of Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> consumed was recorded and represented as S. For the analysis, the same was repeated with blank sample and volume of Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub> consumed was recorded as B. The IV was calculated using:

$$IV = \frac{(B-S) \times N \text{ of sodium thiosulphate} \times 12.69}{\text{weight of sample (g)}}$$

#### 2.2.5 Acid value (AV)

An important index of physicochemical property of oil which indicates the quality, age, edibility and suitability of oil for its uses. The acid value (AV) for oil samples were determined by the International Union of Pure and Applied Chemistry (IUPAC) 2.201(1979)/ISO 660:1996). Diethyl ether and ethanol of equal volume (25ml) were mixed together and 1ml

of 1% phenolphthalein indicator solution was added. The mixture was then neutralized with 0.1M potassium hydroxide solution and the oil sample (between 1-10g) was dissolved in the neutralized solvent mixture. The sample mixture was then titrated against 0.1M potassium hydroxide solution with constant shaking until appearance of pink color that persists for about 15seconds is obtained. The acid value was calculated by:

$$AV = \frac{\text{titre value(ml)} \times 5.61}{\text{weight of sample(g)}}$$

### 2.2.6 Unsaponifiable matter

The Unsaponifiable matter was determined based on the (Commission, 1999) [6] AOAC, official method 933.08. It uses the neutralized liquid after the titration for the determination of saponification value. The neutralized liquid was transferred quantitatively into a separating funnel using 50 ml of water for washing the flask. It was repeated for 3 times while still warm with 50ml diethyl ether, all the ether extracts combined into another separating funnel and washed vigorously with 20ml portions of water and the water was discarded. Following that it was also washed with 20 ml portions of aqueous 0.5 M potassium hydroxide. The extract was transferred into the beaker and kept in boiling water bath for evaporation, 2-3 cm<sup>3</sup> acetone was added and heated on water bath. It was further dried to constant weight and then dissolved in 2ml of diethyl ether. 10 ml of neutralized ethanol was added and titrated with 0.1 M alcoholic potassium hydroxide. The unsaponifiable matter was obtained from the equation:

$$\text{Unsaponifiable matter} = \frac{M - 0.028v \times 1000}{W}$$

Where

V= volume of potassium hydroxide used

M= Mass of oil used for saponification value

W= Mass of unsaponifiable matter

### 2.2.7 Polenske value (PV)

Most of the oils/fats didn't require this test to be performed whereas coconut oil and palm kernel oil contains appreciable quantities of caprylic, capric and lauric acid glycerides and therefore requires this test. The polenske value is historically being determined based on the AOAC 925.41. About 5 g of oil/fat was weighed and a mixture of 20 g glycerol and 2 ml of 50% of NaOH (Sodium hydroxide) was added. The whole mixture is heated in a Bunsen burner with constant stirring till saponification is initiated (liquid becomes clear) and if liquid is dark it indicates excess heating where the procedure has to be repeated. Then the mixture was cooled to permit the addition of water and distillation is performed. Then the alcoholic solution of insoluble volatile acids was collected after distillation and 0.25 ml of phenolphthalein indicator was added. Finally, it was then titrated against 0.1 N NaOH until the disappearance of pink color and the same was performed for blank.

$$PV = 10 \times V \times N$$

Where

V= Volume of NaOH required for titration (in ml)

N= Normality of NaOH

### 2.2.8 Free fatty acid (FFA)

The free fatty acid in the oil sample was measured based on the standard Association of Official Agricultural Chemists

(AOAC) method. Approximately 7.0g of different oil samples were mixed with 2 ml of phenolphthalein solution and a few drops of 0.1 M NaOH. Next 50 ml of ethanol was mixed and constantly shaken until a permanent faint pink color was obtained, which was then titrated against 0.25 N NaOH. The volume of NaOH spent was recorded as S and the same was repeated for the blank and the volume was recorded as B. the percentage of free fatty acid was calculated by the formula:

$$\% \text{ FFA} = \frac{(B-S) \text{ ml of NaOH} \times N \times 56}{1.99 \times \text{weight of sample(g)}}$$

Where

N-normality of NaOH.

### 2.3 Statistical analysis

The physicochemical properties of oil samples were performed in the set of 5 trials (n=5) for repeatability and reliability. The result is presented as the average value of the data obtained. The mean, standard deviation and standard errors were analyzed using Toolkit in Ms-Excel with the significance level of 95 percent and the quality characteristics were compared.

### 3. Result and discussion

The physicochemical properties of virgin coconut oil (VCO) were compared with the Refined, Bleached, Deodorized (RBD) coconut oil and cold pressed oil (Chekku oil) and represented in the Table. 1 (Asian Pacific Coconut Community, 2009) [3].

#### 3.1 Moisture content

The moisture content of oil samples was found to be in the range of 0.10%-0.80% (w/w), which is within the value recommended by APCC (<0.5% w/w) except the cold pressed oil. This indicates that the VCO were able to produce with low moisture content 0.1%. On the other hand, RBD oil and CPO contain slightly higher moisture content than VCO which may be due to the keeping quality of oil and initiation of oxidation. Lower moisture content of VCO (Choe & Min, 2006) indicates the increase in shelf life and prevents oxidation and rancidity process.

**Table 1:** Physicochemical properties of coconut oil samples (RBD, CPO and VCO)

Parameters	Coconut oil samples			APCC standard
	RBD	CPO	VCO	
MC %	0.13±0.005	0.7±0.044	0.11±0.008	Max 0.5
RI at 40 °C	1.4485	1.4484	1.4487	1.4480 – 1.4492
SV (mg KOH/g)	252±0.75	257±0.63	259±0.63	Min 250
IV (gI <sub>2</sub> /100g)	9.2±0.074	8.2 ±0.04	7.0±0.06	4.0 – 11.0
AV (mg KOH/g)	0.5±0.05	0.9±0.006	2.7±0.04	Max 4.0
Unsaponifiable matter %	0.16±0.007	0.1±0.049	0.12±0.006	<0.5
PV (meqO <sub>2</sub> /kg)	14.0±0.10	14.5±0.05	14.6±0.06	Min 13.0
FFA %	0.25±0.01	0.42±0.006	1.3±0.05	Max 2.0

RBD-Refined, Bleached and Deodorized coconut oil, CPO-Cold Pressed Oil (Chekku oil), VCO-Virgin coconut oil.

#### 3.2 Refractive index

The RI values of the VCO was 1.4487 and that of RBD oil and CPO was 1.4485 and 1.4484 respectively. Results indicate that the refractive index of VCO was higher when compared with RBD and CPO. The values are within the specifications suggested by APCC set proposed. Most

possibly, the disparity is due to the high FFA and impurity of VCO sample (F. Dayrit *et al.*, 2007) [10]. Therefore, certain compositions of FFA results in higher RI of VCO compared with RBD and CPO. Hence, this parameter can be considered for the adulteration detection of VCO with other vegetable oils.

### 3.3 Saponification value

The SV of VCO was about 259 mg KOH/g fats whereas the codex range was between 248-265 mg KOH/g fats (Codex, 2001). Comparatively VCO has the highest SV. The SV is related to the mean molecular mass of fats and oils and inversely proportional to the chain length of the fatty acids of fats and oils (Gopala Krishna *et al.*, 2010) [14]. Hence, the VCO contains higher content of short chain fatty acids in contrast to RBD and CPO oil.

### 3.4 Iodine value

The samples were found to have IV in the acceptable range compared to the APCC standard (4.0 to 11.0). The lowest IV was noted for the VCO (7.0) whereas the CPO and RBD oil has the IV of 8.2 and 9.2 respectively. Hence, IV indicates the weight percentage of coconut oil related to unsaturated fatty acids that can absorb halogens such as iodine (R *et al.*, 2019) [21]. It was analysed that the VCO has less content of unsaturated fatty acids to bind such halogens but the RBD and CPO had higher IV which was due to the presence of high amount of unsaturated fatty acids. Because of the presence of low amount of unsaturated fatty acids safety limit of consuming VCO should be studied and adulteration detection can be made wise with the iodine value.

### 3.5 Acid value

The AV of CPO and RBD oil was 0.9 mg KOH/g and 0.5 mg KOH/g respectively, much lower than the VCO (2.7 mg KOH/g) and moreover all the samples were within the APCC limit. It can be understood that the behaviour of either lipase activity or light and heat exposure was relatively higher in VCO rather than other samples (Akubugwo *et al.*, 2008; Warra *et al.*, 2011) [2, 28]. The lipase activity and hydrolytic action or oxidation was used to correlate the % FFA in fat or oil.

### 3.6 Unsaponifiable matter

VCO and CPO, which have the highest saponification value of 259 and 257 mg KOH/g, also have the lowest unsaponifiable matter of 0.12 per cent and 0.1 per cent, respectively, which confirms that their properties will be ideally suited for soap production compared to 0.16 per cent in RBD oil and would be better suited for use in cosmetics. This would also refer to those substances that are not saponifiable by alkali hydroxides but soluble in regular fat solvents (Afolayan *et al.*, 2014) [1] and to products of saponification that are soluble in such solvents.

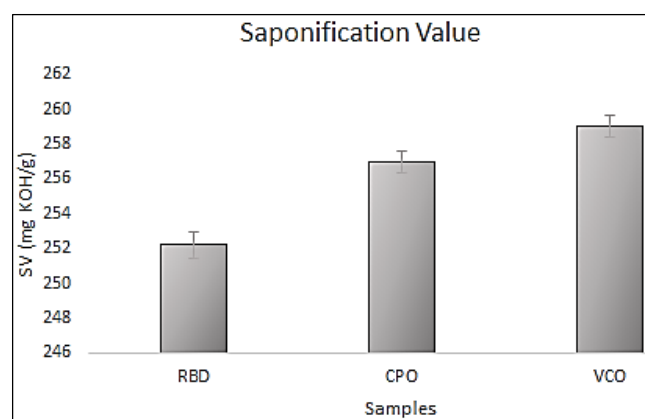
### 3.7 Polenske value

In comparison VCO has slightly high PV of 14.6 m-eqO<sub>2</sub>/kg than CPO and RBD oil having 14.5 m-eqO<sub>2</sub>/kg and 14.0 m-eqO<sub>2</sub>/kg respectively. Results reveal that the VCO has higher quantities of caprylic, capric and lauric acids. These glycerides form the composition of free fatty acids. Therefore, higher PV would indicate the presence of high % FFA. These fatty acids are steam volatile but insoluble in water and gives high polenske number (Nwinuka N.M *et al.*, 2009). It can be

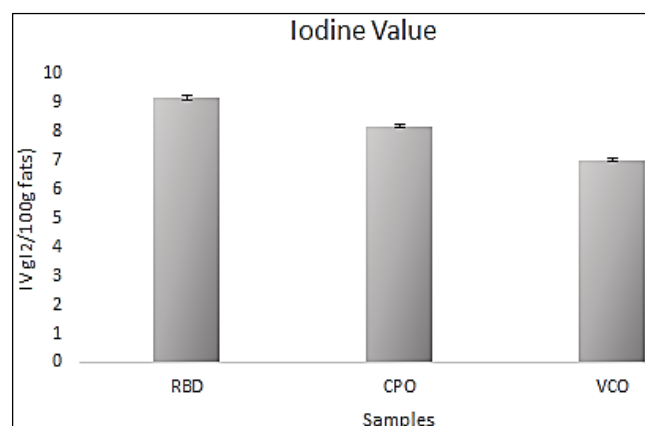
observed that PV is directly related to the free fatty acids present in the sample.

### 3.8 Free fatty acids

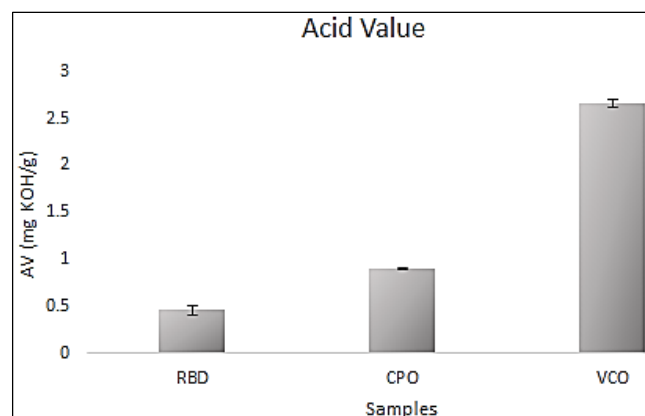
The % FFA of all the samples were in the range of 0.25%-1.3% which was acceptable and within the recommended limit by APCC. However, RBD oil has the lowest % FFA when compared to CPO and VCO. FFA are produced by the hydrolysis of oils and fats and are susceptible to oxidation (Kumar & Krishna, 2015; Marina *et al.*, 2009) [29, 17]. In RBD oil while processing with chemical treatments it may be reduced and regulated without suppressing the quality. FFA can be used as an indicator of taste and aroma of oil. The trend followed to interrupt % FFA from refractive index, acid value and polenske value in oil samples can be used vice versa.



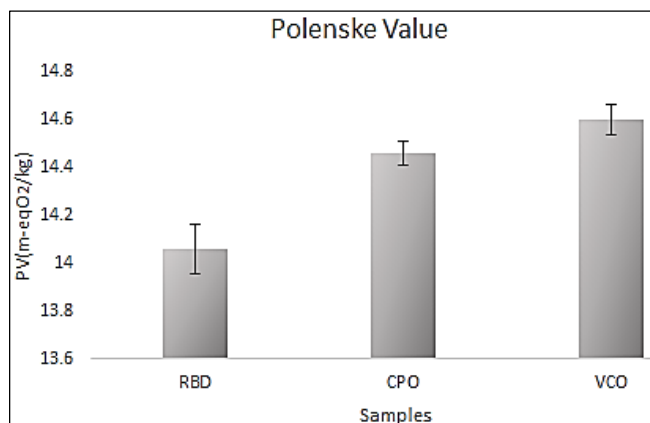
(A)



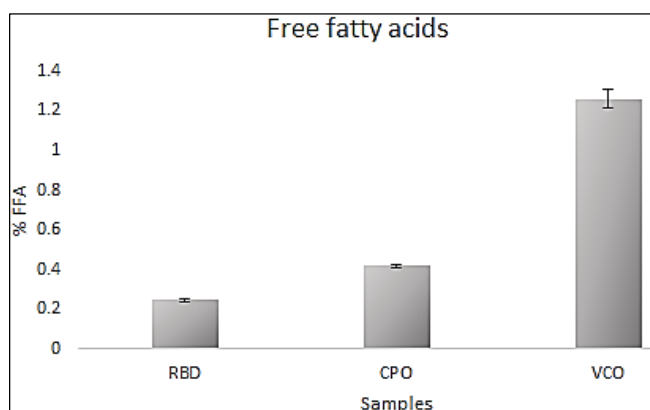
(B)



(C)



(D)



(E)

**Fig 1:** Comparative study on physicochemical properties of VCO with other coconut oil samples (A) Saponification Value, (B) Iodine Value, (C) Acid Value, (D) Polenske Value, (E) Free fatty acids

#### 4. Conclusions

This research findings reveals that most of the physicochemical properties tested are within the acceptable range referred by APCC 2009 [3] (Standard, 2009) [26]. The Iodine value of the VCO was 7.0 gI<sub>2</sub>/100g, which was lower compared to other samples of coconut oil, indicating that the VCO has more oxidation stability and the lower moisture content of VCO could enhance the shelf life of oil. The FFA of VCO was 1.3% which was slightly higher, would mean that consumption for cooking purpose was minimal. VCO showed good results against the RBD coconut oil and CPO which can be used in the cooking and other edible purposes ensuring quality and safety. Therefore, the parameters including SV, AV, IV and % FFA can be used for the adulteration detection of VCO to have significant results.

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