SYNTHESIS OF ACRYLATED EPOXIDIZED BIOBASED RESIN FROM GROUNDNUT SEED OIL

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Abstract: In this study, groundnut seed oil was epoxidized in situ using hydrogen peroxide (30%) and formic and acetic acid. The reaction conditions were monitored at a temperature of 70°C, stirring speed of 750 rpm and time of 6 hours. After epoxidation, a further modification was done using acrylic acid in the presence of hydroquinone at a temperature of 120°C. Comparatively, peroxyformic acid performed more effectively than the peroxyacetic acid during epoxidation with an iodine value (26.4 gl/100g oil) and oxirane content(3.27%). FTIR analysis of the raw, epoxidized, and acrylated groundnut seed oil indicates that they were suitably functionalized.

Keywords: Groundnut seed oil, epoxidation, acrylation, formic acid, acetic acid.

INTRODUCTION

The use of petroleum-based feedstock in the manufacture of polymers has experienced some levels of decline in recent years due to the spiralling prices and high rate of depletion of the stocks. Furthermore, the unhealthy effect of polymer wastes on the environment resulting from its non-biodegradability is of great concern. This has inspired researchers to investigate sources of renewable natural materials as an alternative source of monomers for the polymer industry, which could substitute for the petroleum-based monomers for the manufacture of polymers, hence the need to develop biobased resins from plant seed oils [1-4].

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Plant seed oils are essential raw materials in the formation of bio-based thermosets. Different researchers have worked extensively on plant seed oil such as sunflower oil, soybean oil, groundnut seed oil and linseed oil due to their high iodine value and fatty acid profile (oleic, linoleic and linolenic acids) contained [5-7].

Vegetable oils containing unsaturated fatty acids triglycerides can be suitably epoxidized and further modified to biobased resins. These C=C bonds are the reactive sites for the modifications [20]. Studies have revealed that rubber seed oil, soya bean oil, sesame seed oil, melon seed oil and groundnut seed oil containing high unsaturated fatty acid triglycerides can be modified through several chemical processes [8-12].

Wool and Sun [13]. reported on various synthetic pathways by which an epoxidized plant oil can be suitably functionalized using acrylic acid to give acrylated epoxidized triglycerides and maleic anhydride to give malenized triglycerides.

Groundnut is a principal oilseed crop that is cultivated on a large scale throughout the world. It is an annual crop principally for its edible oil and protein-rich kernel seeds, borne in pods that develop and mature below the soil surface [14]. In India, 80% of the groundnut produce is crushed to extract oil and accounts for 36.10% of the total oil production. Groundnut seed contains 44 – 56% oil and 22 – 30% protein on a dry seed and is a rich source of mineral (Phosphorous, calcium, magnesium and potassium) and vitamins E, K and B group [15]. The fatty acid composition consist of Iodine value (g/100 g) = 85; oleic acid (18:1) = 46.8%. linoleic acid (18:2) = 33.4. This composition makes the oil amenable to various chemical modification; it can be successfully epoxidized and further modified [12].

The use of bio-based resins has helped reduce fossil fuel consumption, and they have been utilized in the production of various products, such as inks, paints, coatings, and plastics. Also, it has been used in the production of technical materials for housing (doors, window frames, hot tubs and composite decking), for aerospace (aero craft wings, tails, propellers, and interior), boats, bathtubs, pools, storage tanks and in the manufacture of vehicle parts [10, 16-18]. This paper, therefore, adds value to groundnut seed oil by synthesizing epoxides and acrylates from its oil, which could be used as a substitute to the non-renewable ones made from petrochemical sources that have adverse effects on the environment. Table 1., below represents the fatty acids (oleic, linoleic and linolenic acids) present in groundnut seed oil.

Table 1. Composition of unsaturated fatty acids present in melon seed oil and their molecular weights, [12].

| Fatty acids    | Composition (wt%) |
|---------------|-------------------|
| Oleic acid    | 53.0              |
| Linoleic acid | 32.0              |
| Linolenic acid| -                 |
MATERIAL AND METHODS

Materials

Magnetic heater with a stirrer, Three-necked round bottom flask, Thermometer, Condenser, Feed funnel, Stirring bulb, measuring cylinder, weighing balance, Separation funnel, Rotary evaporator. Groundnut seed oil was used as raw material, Formic acid (85 wt%), Acetic acid (85 wt%), Hydrogen peroxide (30 wt%), Sodium Carbonate, Distilled water, Acrylic acid, Hydroquinone powder, Toluene, Sodium sulphate.

Methods

Epoxidation procedure

(35 g) of groundnut seed oil was measured and placed in the three-necked round bottom flask, 2.43 g of formic acid was added to the flask, and the mixture was stirred continuously for 30 min. Then 16.15 g of 30% aqueous hydrogen peroxide was added drop-wise to the reaction mixture as an oxygen donor while stirring. The mole ratio of the components used was 1:1.5:0.5, \( \text{H}_2\text{O}_2: \text{HCOOH} \). After the complete addition of hydrogen peroxide, the mixture was heated under reflux and maintained at a temperature of 70°C with rapid stirring. The rapid stirring was maintained throughout the experiment to achieve fine dispersion of oil and avoid high peroxide concentration zones that could lead to an explosive mixture. Samples were collected hourly from the set-up for analysis. The collected samples of epoxidized melon seed oil (EMSO) were then washed with a solution of 10 g of sodium carbonate mixed in 200 ml of distilled water to remove the free fatty acids and other unwanted components. The 10 g of \( \text{Na}_2\text{CO}_3 \) was first dissolved in 100 ml of distilled water before the remaining 100 ml was finally added, mixed with the sample and separated using a separation funnel. Subsequent extraction was used to recover the remaining samples after washing.

Synthesis of Acrylated epoxidized groundnut seed oil (AMMSO)

35 g of epoxidized melon and groundnut seed oil was heated at room temperature, acrylic acid (4.89 g) containing hydroquinone (0.013 g of 0.25 wt% acrylic acid) was added after 30 min. The reaction mixture was heated under reflux from 8 to 12 hours under constant stirring. The obtained product, Acrylated epoxidized groundnut seed oil, was then isolated.

Characterization of groundnut seed oil

The pure, epoxidized, and acrylated groundnut seed oil was characterized using Fourier Transform Infrared (FTIR) Spectroscopy Technique to determine surface functional groups present. The FTIR analyses were carried out on the samples using Shimadzu FT-IR-8400S Spectrophotometer with a resolution of 4 cm\(^{-1}\) in the range of 400 - 5000 cm.
Analytical techniques
Iodine value

Wiji's method of the Association of Oil Chemists determined the iodine value of the test oil sample. First, 0.5 g of the sample was poured into a conical flask. Next, 10 ml of carbon tetrachloride was added to the oil and shook to allow the oil to dissolve. Also, 20 ml of Wiji's iodine solution was later added to the mixture, stirred vigorously, stoppered and kept in the dark for 30 minutes. Subsequently, 15 ml of potassium iodide solution followed by 100 ml of distilled water was added. The mixture was titrated against 0.01 N sodium thiosulphate solution. A reagent black was titrated as well.

The iodine value of epoxidized samples was calculated after analysis using the formula in eq. (1).

\[
IV = \frac{(B - S) \times M \times 12.69}{W}
\] ………….. (1)

Where:
IV = Iodine value of samples
S = Volume of Na\(_2\)S\(_2\)O\(_3\) used for sample (ml),
B = Volume of Na\(_2\)S\(_2\)O\(_3\) used for blank (ml),
W = Weight of sample used (g),
M = Molarity of the Na\(_2\)S\(_2\)O\(_3\) used.

Oxirane Oxygen content

The percentage of the oxirane oxygen was determined by the direct method established by using a hydrobromic acid solution in glacial acetic acid. First, the content of oxirane oxygen (O.O.) was calculated according to the consumed amount of the halogen atom.

The Oxirane Oxygen Content of the analyzed samples was calculated using the formula in eq. (2).

\[
OV = \frac{(B - S) \times M \times A_o \times 100}{1000W}
\] ………….. (2)

Where:
S = Volume of NaOH used for sample (ml)
B = Volume of NaOH used for blank (ml)
M = Molarity of the NaOH used
W = Weight of sample used (g)
A\(_o\) = Atomic weight of oxygen
RESULTS AND DISCUSSION

Table 2. Iodine and oxirane values of the epoxidized samples

| Samples                                | Iodine value (g /100 g of oil) | Oxirane value (%) |
|----------------------------------------|-------------------------------|------------------|
| Epoxidised groundnut seed oil with acetic acid | 26.92                        | 2.00             |
| Epoxidised groundnut seed oil with formic acid | 26.4                         | 3.27             |

The iodine value and oxirane oxygen content are essential parameters in the characterization of epoxidized vegetable oils [19]. While the iodine value indicates the remaining unsaturated fatty acid triglyceride present after the epoxidation reaction, the oxirane oxygen content indicates the epoxy groups incorporated in the products. From the results presented in Tab. 2, it was observed that the epoxidized groundnut seed oil with formic acid had better-functionalized samples than the oil epoxidized with acetic acid. With formic acid, the iodine value was obtained as 26.4 g/100 g oil, which is lower than 26.92 g/100 g oil for acetic acid and the oxirane value of 3.27% was obtained for samples where formic acid was used. This is a higher than 2.00% value result obtained for samples where acetic acid was used. Although the oil samples were epoxidized successfully with both formic and acetic acids as used in this study, the results indicated that formic acid gave a better performance than acetic acid for the epoxidation process of groundnut seed oil. This result agrees with the findings of [19].
Discussion

From results of the FT-IR spectra as presented in Fig. 1-5, the presence of carbon-carbon double bonds (C=C) were observed in the untreated groundnut seed oil, which was indicated by the appearance of peaks at 3500 cm\(^{-1}\) and 3000 cm\(^{-1}\) as shown in Fig. 1, this indicates the functionality of the oil. As presented in Fig. 2 in the acetic acid epoxidized groundnut seed oil, the absorption band for the epoxy group was indicated by the single peak at 3527.92 cm\(^{-1}\). For the same oil epoxidized with formic acid, the peak was also indicated at 3506.70 cm\(^{-1}\); this shows a close margin between both carboxylic acids, indicating their suitability in the epoxidation process. Fig. 3 shows that the oils were suitably epoxidized in both cases, and these results were achieved after six hours (6 h) reaction time, stirring speed of 750 rpm and 70°C temperature.
The I.R. spectra for the acrylic acid modified samples revealed acrylic group indicated by the peak at 3489 cm\(^{-1}\) for the acrylated epoxidized groundnut seed oil with acetic acid and 3483 cm\(^{-1}\) for the acrylated oil epoxidized with formic acid, as shown in Fig. 4 and Fig.5, respectively. These peaks were observed to be absent in both the epoxy resin and the pure, unmodified groundnut seed oil, which was an indication that the epoxy groups in the epoxidized oils have been converted to acrylated groups after the acrylation process was carried out on the epoxidized groundnut seed oil, this result is in agreement with the findings of [20].

CONCLUSIONS

The results obtained during this study found that groundnut seed oil is a good starting material for oil epoxy synthesis. Although there may be competition with food needs because of its edibility, increasing pressure on supply, consolidating on this research would upscale its production and increase its diversification in other areas of application. Under the same reaction conditions of temperature 70°C, stirring speed of 750 rpm, and time of 6 hours, peroxyformic acid was observed to be more efficient for in situ epoxidation than peroxyacetic acid.

However, both carboxylic acids gave good results during the process of epoxidation and acrylation of the oil.

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Sažetak: U ovoj studiji ulje semena kikirikija je epoksidovano in situ pomoću vodonik-peroksida (30%) mravlje i sirčetne kiseline. Uslovi reakcije praćeni su na temperaturi od 70 °C, brzini mešanja od 750 o/min i vremenu trajanja od 6 sati. Posle epoksidacije, izvršena je naredna modifikacija upotrebom akrilne kiseline u prisustvu hidrohinona na temperaturi od 120°C. U poređenju sa navedenim tretmanima, peroksiformna kiselina se pokazala efikasnijom od peroksiocetne kiseline tokom epoksidacije sa jodnom vrednošću (26,4 gl/100 g ulja) i oxiran sadržajem (3,27%).
FTIR analiza sirovog, epoksidovanog i akrilovanog ulja semenki kikirikija ukazuje na to da semenke kikirkija funkcionalno odgovaraju ovom procesu.

**Ključne reči:** Ulje semena kikirikija, epoksidacija, akrilacija, mravlja kiselina, sirčetna kiselina.

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