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Synthesis of bio-oil from waste *Trichosanthes cucumerina* seeds: a substitute for conventional fuel

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The present study explores the methodology for the synthesis of bio-oil from waste *trichosanthes cucumerina* seeds by the solvent extraction method. It investigates the yield percentage, concentration of free fatty acids and acid contents in the extracted bio-oil. Effects of size of the crushed seeds, moisture content, extraction time, solvent to seed ratio and extraction temperatures were examined. The non-polar hexane solvent resulted in a higher percentage of oil yield ($28.4 \pm 0.4\%$) for the crushed seed size of 0.21 mm, 6% moisture content, 270 min extraction time, 68 °C temperature and 6:1(ml/g) of solvent to seed ratio. The synthesized bio-oil was characterized using Fourier Transform Infra-Red spectrum and Gas Chromatography–Mass Spectroscopy analysis. The properties of the bio-oil and biodiesel were assessed according to the American Society for Testing and Materials and the Association of Official Analytical Chemists standards. The obtained methyl-ester by trans-esterification process results in the fuel properties closer to the conventional fuel. Thus, *Trichosanthes cucumerina* bio-diesel can be used as a potential substitute.

The increase in the percentage of energy consumption by various sectors makes the fossil sources in deficiency and makes the researchers start to shift to renewable energy sources¹. Among the various alternative sources, straight vegetable oil has identical fuel properties, sulphur-free and bio-degradable, which makes it as the substitute for fossil fuels². The bio-diesel from renewable sources shown positive impacts for better engine performances and cleaner emissions in the automotive sector³. In general, the vegetable oils are classified as edible and non-edible feedstocks. The edible feedstocks namely coconut, olive, palm, peanut, rice bran, soybean and sunflower are the predominant in the production of bio-diesel but leads to risk in food supplies, bio-diversity and increase the cost of the fuel. Therefore, the non-edible feedstocks like cottonseed, jatropha, jojoba, polanga, karanja, linseed, mahua, neem, rubber seed and tobacco were considered as the substitutes^{4,5}.

The bio-origin materials produce a variety of biofuels through the conversion routes of pressing, extraction, chemical processes (hydrolysis and trans-esterification), biochemical processes (fermentation and anaerobic digestion), thermochemical processes (flash pyrolysis, gasification and hydrothermal liquefaction) and direct combustion⁶. The reactive extraction is considered as the most viable technology in the production of bio-diesel compared with other methods. The reactive extraction based soxhlet extractor combines the oil extraction and transesterification processes used for the bio-oil production using various seeds^{7,8}. *Crotalaria juncea* oil, jojoba oil, *zanthoxylum bungeanum* oil, *jatropha* oil, *indigofera colutea* oil, soybean oil, *senna occidentalis* oil, *cassia javanica* oil and palm oil were produced with the soxhlet extractor and reported in the literature^{9,10}.

In the trans-esterification process, the traces of free fatty acids in the oil extracted gets removed and produce the by-products of ester and crude glycerol with the assistance of an acid catalyst, alkali catalyst and purification process^{11–13}. This trans-esterification process is considered as the most feasible and commercially used technique for the conversion of methyl esters and effective reduction of viscosity for the bio-oils¹⁴. The physicochemical properties such as cetane number, heating value, kinematic viscosity, density, flash and fire points of the bio-diesel obtained using trans-esterification processes were assessed with ASTM standard methods¹⁵.

The bio-oil extraction from *jatropha curcas* seeds was studied with the functional parameters of solvent namely extraction temperature, type of solvent used (n-hexane and petroleum ether), solvent-to-seed ratio, extraction time and particle size of the seeds. The n-hexane solvent was seen with the maximum efficiency and 1.3% higher when compared to petroleum ether for the optimum condition¹⁶. The separation of bio-oil from urban waste *putranjiva roxbughii* seeds were carried out with the five different solvents (Diethyl ether, Hexane, Isopropanol, Toluene and Chloroform). The non-polar solvent hexane seen with better results compared to other

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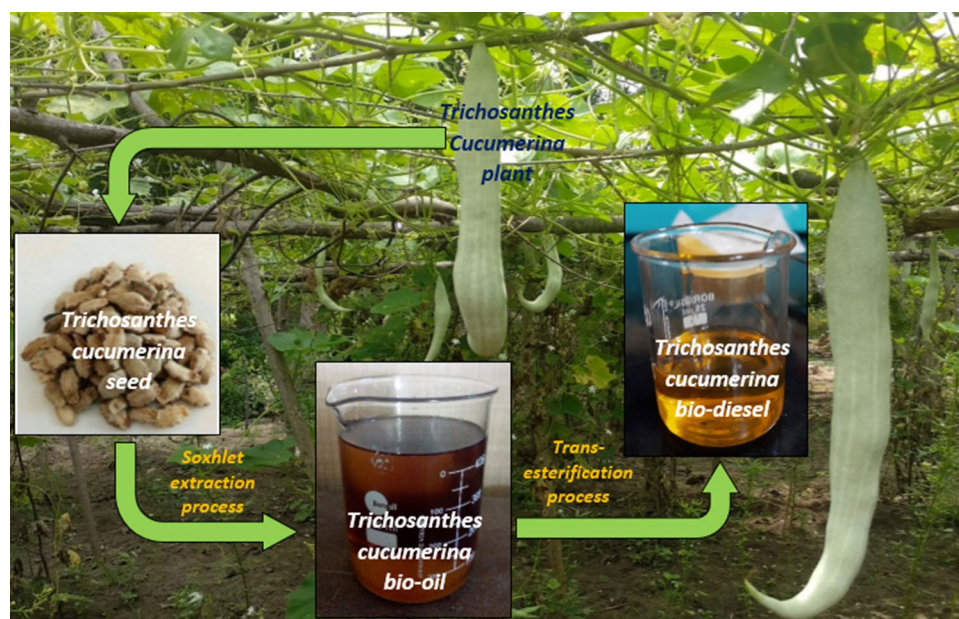


Figure 1. *Trichosanthes cucumerina* fruit, waste seed, TCO and TCB. It presents the image of *Trichosanthes cucumerina* fruit and its collection of waste seed. Also, the extraction of bio-oil and production of bio-diesel.

Solvent parameters	Solvents used				
	Acetone	Methanol	Hexane	Ethanol	Isopropanol
Chemical formula	C ₃ H ₆ O	CH ₃ OH	C ₆ H ₁₄	C ₂ H ₅ OH	C ₃ H ₇ OH
Boiling point (°C)	56.1	64.5	68.7	78.3	82.3
Density (kg/m ³)	784	791.4	655	789.3	786
Refractive index at 20 °C	1.36	1.33	1.375	1.36	1.376
Dielectric Constant at 20 °C	20.7	32.7	1.88	24.5	18.6
Latent heat of vaporization (kJ/kg)	511	1100	335	836	666
Viscosity at 20 °C (mPa-s)	0.33	0.593	0.32	1.19	2.37
Surface tension at 20 °C (mN/m)	25.20	22.70	18.43	22.10	23

Table 1. Properties of different solvents used^{18–21}.

considered solvents and also reported that the higher boiling point of toluene requires more heat to form vapour during the solvent recovery distillation process compared to hexane¹⁷. The non-polar solvents (toluene, chloroform and n-hexane) produced maximum bio-oil yield compared to polar solvents (methanol and 2-propanol) in the extraction of bio-oil from date palm seeds¹⁸.

This research is focussed on the synthesis of bio-oil extraction from the waste *trichosanthes cucumerina* seeds and its fundamental properties were investigated. Further, the trans-esterification process was carried out to remove the free fatty acids in the bio-oil and converted to bio-diesel. The *trichosanthes cucumerina* biodiesel was obtained from trans-esterification processes with effective fuel properties. The presence of functional groups and the free fatty acid content were characterized using GC-MS and FTIR analysis. The properties of *trichosanthes cucumerina* bio-oil and bio-diesel were measured using the AOAC and ASTM standard methods and compared with the conventional fuel.

Materials and methods

***Trichosanthes cucumerina* seed and solvents.** The *trichosanthes cucumerina* is an oil-rich seed plant that comes under the family of curcubitaceae. It originated in Asian countries and referred as snake gourd, viper gourd and snake tomato. Figure 1 shows the *trichosanthes cucumerina* which are 40–120 cm long, pale-green in colour, 0.5–1 kg weight (single fruit) and contains 40–70 seeds. Seeds are received from the local market and segregated for the bio-oil production. Five solvents namely polar protic (methanol, ethanol and isopropanol), dipolar aprotic (acetone) and non-polar (hexane) purchased from SRL Chemicals Pvt Ltd, Mumbai, India. The physical properties of the different solvents are listed in Table 1.

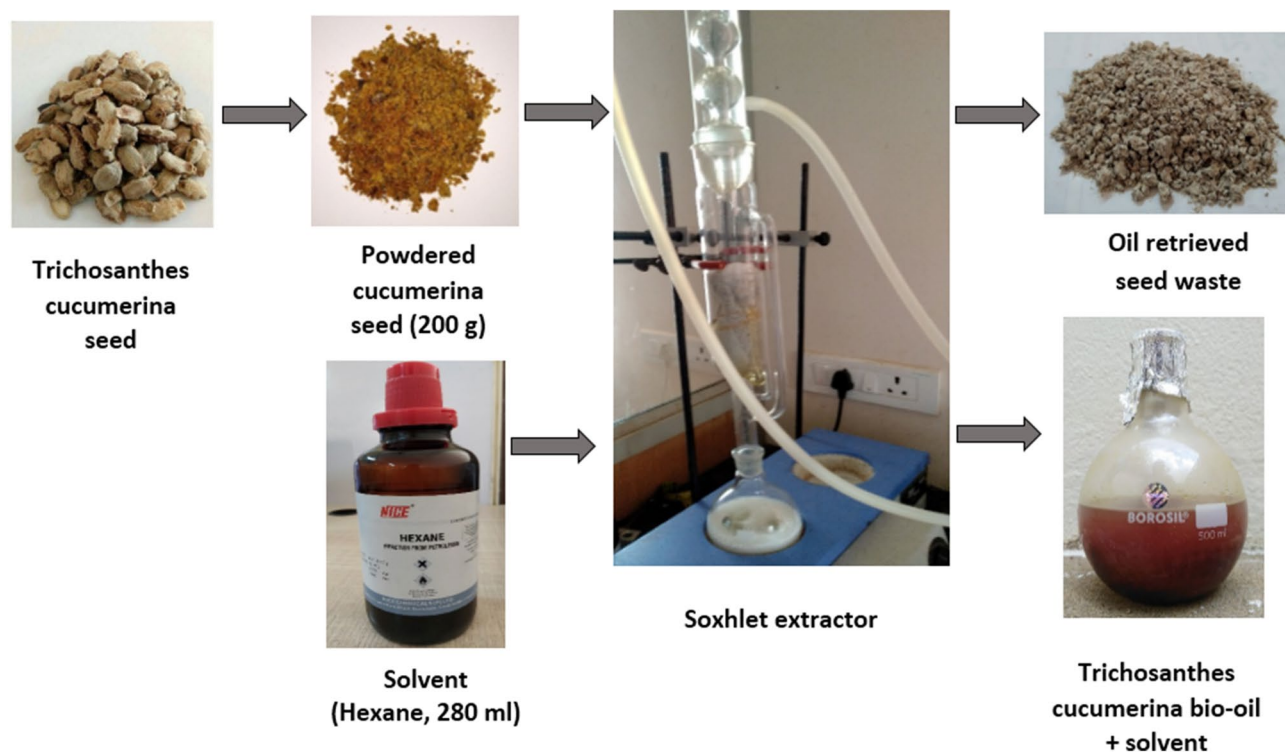


Figure 2. *Trichosanthes cucumerina* bio-oil extraction flow process. The bio-oil extraction from powdered cucumerina seed with hexane using a Soxhlet apparatus produces the oil retrieved seed waste and bio-oil associated with hexane.

Bio-oil extraction process. The segregated seeds were washed with distilled water (in-house) for the removal of impurities. The seeds outer layer were removed and dried in an oven at 65 °C temperature. The dried seeds were crushed in the oilseed crushing machine (SASTRA Deemed University, Tanjore, Tamil Nadu, India) and obtained with 7% of the bio-oil yield from the taken 200 g of *trichosanthes cucumerina* seeds. From the literature, the soxhlet extractor was selected to extract the bio-oil yield^{8,9}. The round bottom flask of 500 ml capacity, filled with 280 ml of hexane and kept in the heating mantle. The powdered cucumerina seeds of 200 g were packed with a satin cloth which was held inside the thimble. A reflux condenser was attached at the top with inlet and outlet ports for cooling water circulation using aquarium motor. The heated solvent vapors were passed over the bio-mass through distillation path and cooled using a condenser. The cold vapor drips back into the chamber, which was emptied by siphoning action. It results in an extracted *trichosanthes cucumerina* bio-oil with solvent and the oil retrieved seed waste is depicted in Fig. 2. In batch distillation process, the obtained bio-oil and solvent mixtures were heated (25 to 85 °C) to evaporate the solvents to separate the bio-oil from the solvent as shown in Fig. 3. The experimental work was performed for three times with each solvent and the average was considered.

Trans-esterification process. The bio-oil leached from *trichosanthes cucumerina* seeds found with FFA content via GC-MS test which needs the trans-esterification process. In the process, the triglycerides of *trichosanthes cucumerina* bio-oil converted into its mono-esters by reacting with alcohols in the presence of NaOH or KOH^{18,22}. The bio-oil prepared from the seeds of *trichosanthes cucumerina* fruit was subjected to the transesterification process using acid-catalyzed esterification, alkali catalyzed esterification and purification stages. Initially, the bio-oil was mixed with methyl alcohol in 16:1 molar ratio then added with 1% of H₂SO₄. The mixture was heated at 60 °C for 45 min which reduced the acid value of bio-oil to less than 4 mg of KOH/g²³. Next, the bio-oil (750 ml) and methyl alcohol (400 ml) were mixed with alkali catalyst (NaOH) and subjected to stirring at a constant speed of 1500 rpm for 30 min duration. The end of the process observed with fatty acid methyl ester, glycerol and the traces of NaOH. Further processing with the addition of HCL and H₂O and followed by the purification process, the traces were removed and obtained with 93.4 ± 0.2% *Trichosanthes Cucumerina* Biodiesel (TCB). The trans-esterification process, as shown in Fig. 4, was carried out at SASTRA Deemed University, Tanjore, Tamil Nadu, India.

Bio-oil yield. The percentage of bio-oil yield was calculated with the formula (1) for all the five different solvents considered in the research (methanol, ethanol, isopropanol, acetone and hexane).

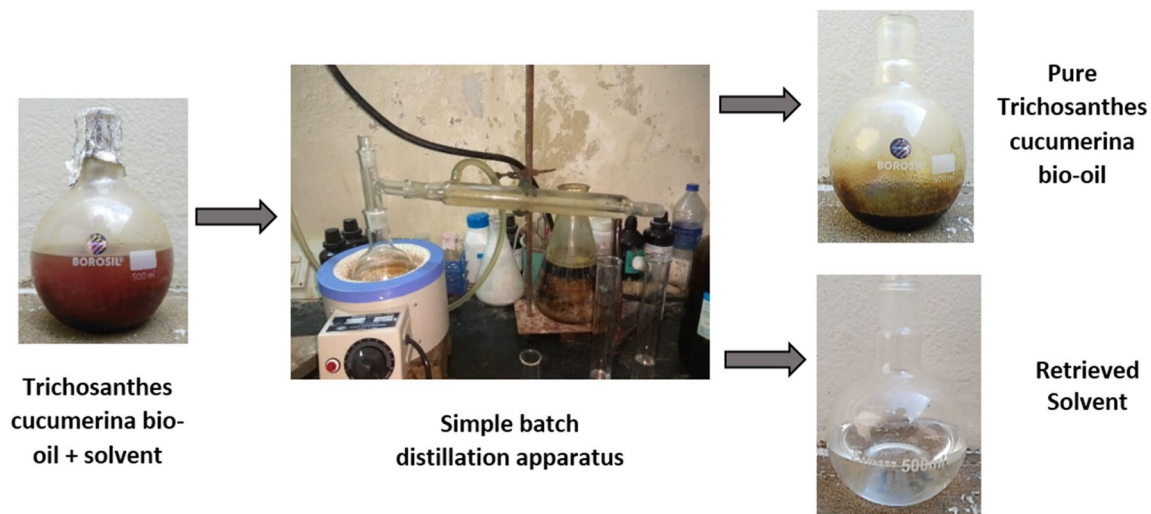


Figure 3. Batch distillation flow process. The mixture of bio-oil with solvent collected from Soxhlet extractor was separated through a separation process. It resulted in around 60% of the solvent, and 28.4% of bio-oil yield.

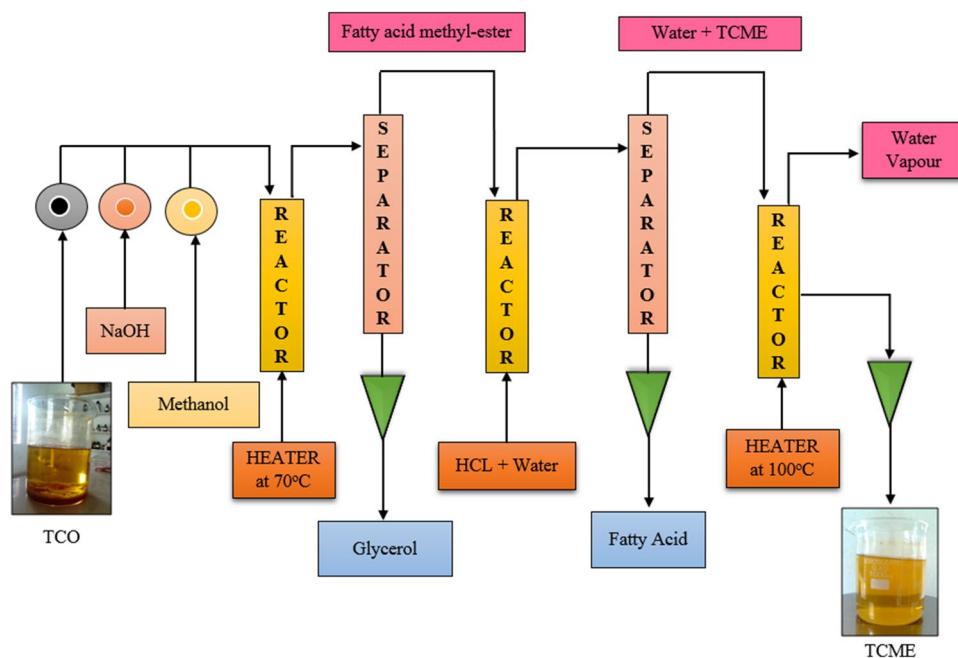


Figure 4. The sequence of trans-esterification process. It is the process to which triglycerides of *Trichosanthes cucumerina* bio-oil converted into their mono-esters by reacting them with alcohols (methanol or ethanol) in the presence of a catalyst (NaOH or KOH).

$$\% \text{ of bio - oil yield} = \frac{\text{Trichosanthes cucumerina bio - oil (gm)}}{\text{Trichosanthes cucumerina seed (gm)}} \times 100 \quad (1)$$

The corresponding percentage differences between the different solvents is shown in Fig. 5. It was evident that hexane resulted in a high percentage of yield was about $28.4 \pm 0.4\%$. It may be attributed due to its low latent heat of vaporization, the minimum value of dielectric constant and surface tension, lower density and viscosity as compared with other solvents are listed in Table 1. The least percentage of $20.54 \pm 0.2\%$ was found with ethanol due to their higher polarity and solubility in water which declined the bio-oil yield. The results obtained in this research were in line with the results of the other researchers^{8,17,20,24-27}, are reported in Table 2.

Results and Discussion

Effect of moisture content. The weight of the seeds with moisture and without moisture was measured using infrared moisture analyzer. For the first set of experiments, *trichosanthes cucumerina* seeds with the moisture content of $1 \pm 0.02\%$ were considered to check the yield of bio-oil with hexane as the solvent and resulted in $20.5 \pm 0.3\%$. The increase in bio-oil yield was observed for the moisture content of seed between $1 \pm 0.02\%$ to

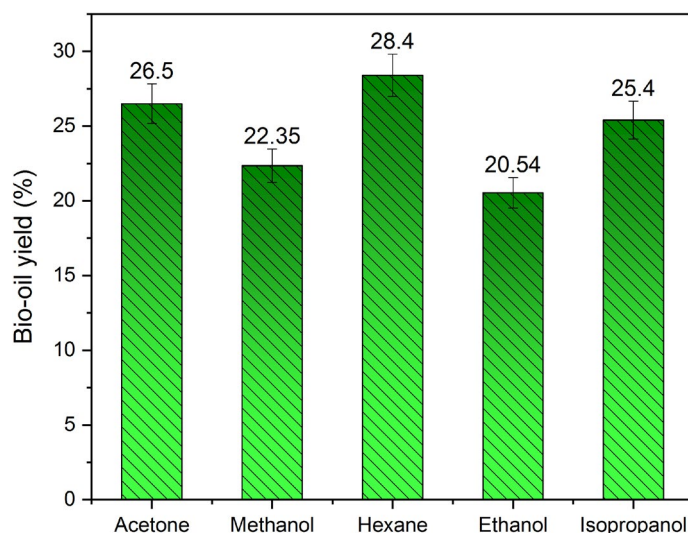


Figure 5. The maximum percentage of bio-oil yield with different solvents. The bar charts represent the comparison of the maximum percentage of bio-oil yield with different solvents used.

Bio-oils	Bio-oil yield (%)								
	Acetone	Methanol	Hexane	Ethanol	Isopropanol	Butanol	Petroleum ether	Chloroform	Toulene
TCO [This study]	26.50 ± 0.2	22.35 ± 0.15	28.40 ± 0.4	20.54 ± 0.2	25.40 ± 0.3	–	–	–	–
ZSO ⁸	–	20.2	26.3	23.4	–	–	25.3	–	–
PRSO ¹⁷	–	–	43.07	–	40.61	–	36.9	25.6	30.5
CGAO ²⁰	–	8.6	11.76	–	9.8	–	–	7.6	9
ULO ²⁴	6.55 ± 0.04	7.51 ± 0.04	8.53 ± 0.04	7 ± 0.05	7.89 ± 0.06	–	–	–	–
SSO ²⁵	19.28	20.3	22.65	–	23.46	–	–	18.72	15.23
CFSO ²⁶	12.5	18	27.03	–	22	–	–	15.2	–
CJSO ²⁷	20.5	18.5	24.4	–	22.3	–	–	16.5	20.6

Table 2. Comparison of *Trichosanthes cucumerina* bio-oil (TCO) yield using different solvents. ZSO *Zanthoxylum bungeanum* seed oil, SSO *Senna occidentalis* seed oil, ULO *Ulva lactuca* oil, CFSO *Cassia fistula* seed oil, CJSO *Cassia javanica* seed oil, PRSO *Putranjiva roxburghii* seed oil, CGAO *Cladophora glomerata* algal oil.

6 ± 0.02%, as shown in Fig. 6. Further increase in moisture content reduced the yield since the penetration of hexane into the *trichosanthes cucumerina* seed, and the higher moisture content functioned as the barrier for bio-oil extraction^{28–30}. The higher percentage of yield (28.01 ± 0.3%) was observed at 6% of moisture content, by keeping other parameters as constant (size of seed as 0.21 mm, extraction time of 270 min, at 68 °C and the solvent-to-seed ratio of 6:1 ml/g hexane). The further increase in the percentage of moisture content shows in the reduction of bio-oil yield. Farsie and Singh³¹ reported that the maximum percentage of bio-oil yield was obtained from sunflower seeds expressed at 6% of moisture content. Muhammad Muhammad Fadhlullah et al.³² determined the effect of moisture content in the bio-oil generation using *Calophyllum inophyllum* L. seeds. It was observed that the yield was increased from 28.87% to 33.39% for the moisture content of the seeds of 0% and 1.2% respectively. In contrast, the increase in moisture content to 20% the yield gets reduced to 15.56%. Orhevba et al.³³ experimented with 6.3, 8.1, 13.2 and 16.6% of moisture content of neem seed kernel and observed with 22.3, 24.86, 21.21 and 15.62% of bio-yield respectively. The maximum percentage of bio-oil yield of 24.86% was observed at the optimum moisture of 8.1% while the least yield of 15.62% was recorded at the highest moisture content of 16.6%. Suganya and Renganathan²⁴ carried out bio-oil extraction from marine macroalgae *Ulva Lactuca* with 12 different solvents. The authors observed the maximum bio-oil yield at 5% of moisture content with hexane as solvent after with 5% the yield gets declined.

Effect of seed size. The extraction of bio-oil was carried out with different sizes of the seed in the range 0.6 ± 0.02 mm to 0.15 ± 0.02 mm were present in Fig. 7. Initially, the seed of size 0.6 ± 0.02 mm was considered for the experimentation with hexane as a solvent which resulted in 20.54 ± 0.3% of bio-oil yield. Further experiments have seen with an increasing trend of yield till 0.21 ± 0.02 mm size of the seed (bio-oil yield as 28.30 ± 0.4%) after

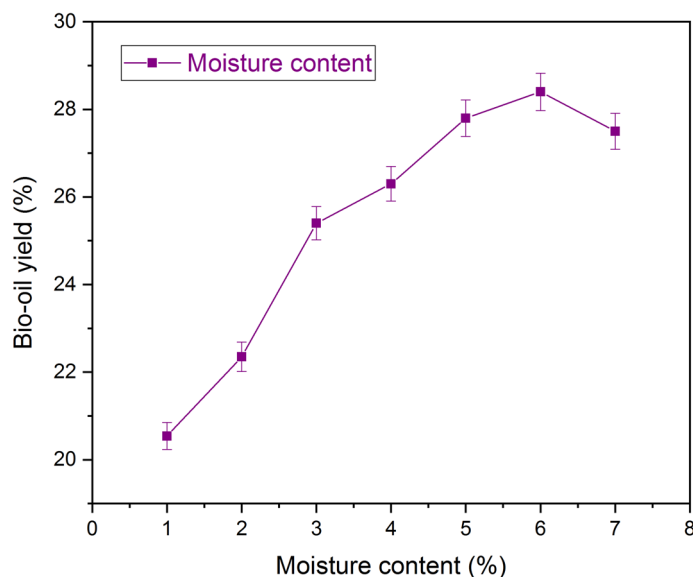


Figure 6. Effect of moisture content on bio-oil yield using hexane. The moisture content (%) of the seed increases with an increase in bio-oil yield with the optimum conditions of 0.21 mm seed size, 270 min extraction time, 68 °C temperature and 6:1(ml/g) of solvent to seed ratio.

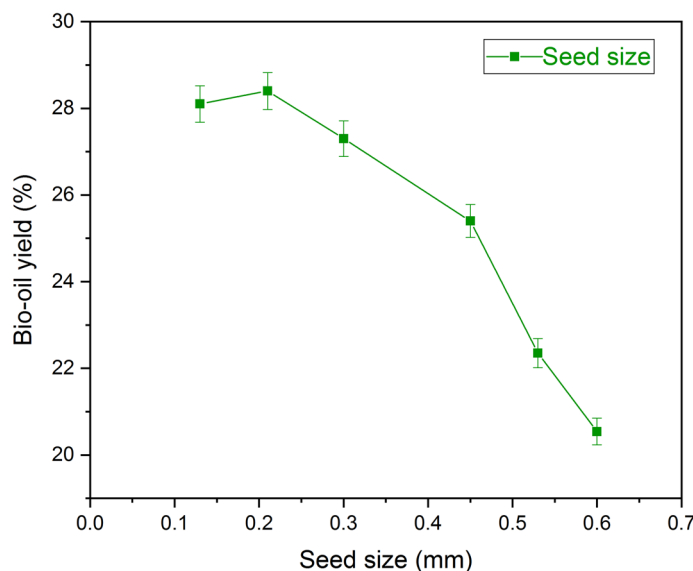


Figure 7. Effect of seed size on bio-oil yield using hexane. The crushed seed size (mm) decreases with an increase in bio-oil yield (optimum conditions: 6% moisture content, 270 min extraction time, 68 °C temperature and 6:1(ml/g) of solvent to seed ratio).

which for the seed size of 0.15 ± 0.02 mm, the bio-oil yield get declined. Thereby the seed size seems to be the next critical parameter for the bio-oil yield extraction. The increase in contact between the solvent and seed, and mass transfer of seed (solid-state) to solvent (liquid state) shows that the decrease in the size of the seeds increased the bio-oil yield³⁴. Qian et al.³⁵ reported that yield from cottonseed gets increased with the reduction of particle size; in contrast, the further reduction has not shown the improvement in the yield extraction. The decrease in seed size would not always increase the yield due to the range limit, which helps in optimizing the yield³². The results obtained in this work were in line with the reported works of literature, the *Ulva Lactuca* reported with a high yield of 10.9% with the seed size of 0.15 mm²⁹ and *Adenanthera pavonina* seen with a high yield of 26.2% for the particle size of 0.25 mm²⁵.

Effect of extraction time. The effect of extraction time at different time intervals of 90,150,210,270 and 330 min for the bio-yield was examined. Figure 8 illustrates that the increase in extraction time increased the

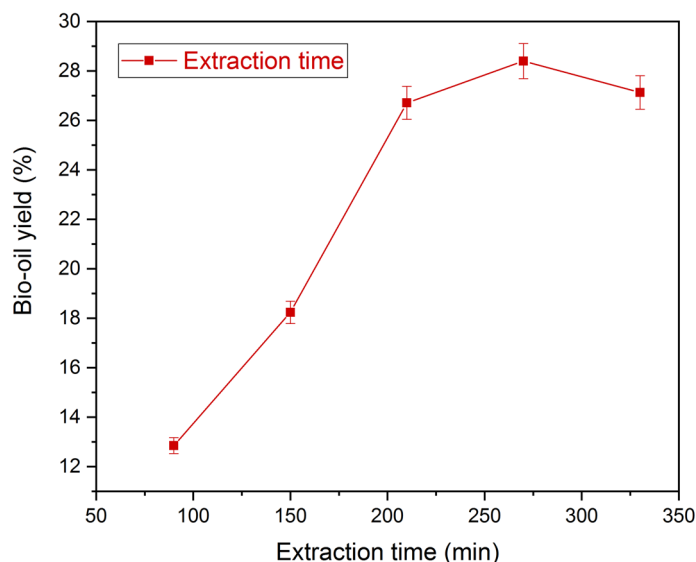


Figure 8. Effect of extraction time on bio-oil yield using hexane as a solvent. The bio-oil extraction from *Trichosanthes cucumerina* seed using hexane with the impact of time, as shown in the graph. The hexane produces maximum bio-oil yield ($28.15 \pm 0.3\%$) as compared with other solvents at an optimum extraction time of 270 min.

Solvent-to-seed ratio (ml/g)	Bio-oil yield (%)	Extraction temperature (°C)	Bio-oil yield (%)
2:1	12.85 ± 0.1	28	18.34 ± 0.15
3:1	20.35 ± 0.2	38	21.05 ± 0.2
4:1	22.35 ± 0.2	48	23.74 ± 0.2
5:1	26.5 ± 0.3	58	26.8 ± 0.3
6:1	28.4 ± 0.4	68	27.81 ± 0.4
7:1	26.2 ± 0.3	78	25.3 ± 0.3

Table 3. Effects of solvent-to-seed (ml/g) ration and temperature (°C) on bio-oil yield extraction.

bio-oil yield from 90 to 270 min, a further increase in extraction time decreased the yield (say at 330 min). The bio-oil extraction time increased gradually, which also increases the yield percentage up to a certain extent, after that the yield reaches a plateau at longer time duration³⁶. Hence the extraction time 270 min was remarked to be the optimum time for further examinations. From the literature, it was observed that the *Senna occidentalis* seeds produced the maximum oil yield 23.46% with the optimum extraction time of 210 min³⁷. Theresa et al.³⁸ investigated the extraction time between 30 to 300 min using *Indigofera colutea* seeds with hexane as the solvent and seen with maximum yield (38.45%) obtained at 210 min, further extension of time decreased the bio-oil yield.

Effect of solvent-to-seed ratio. The effect of solvent-to-seed ratio was also explored by varying the ratio between 2:1 to 7:1, as given in Table 3. The increase in the bio-oil yield percentage was observed until 6:1, a further increase in ratio reduced the yield percentage. The increase in solvent volume significantly improved the yield until the value of the equilibrium reached, a further increase in solvent volume not seen with any improvement²⁹. Suganya and Renganathan²⁴ experimented on marine macroalgae *Ulva Lactuca* with the solvent-to-seed ratio varied from 3:1 to 6:1, which increased the bio-oil yield from 9% to 10.88%. The yield gets reduced for the further increase of solvent volume.

Effect of extraction temperature. The variation of extraction temperature on the bio-oil yield obtained from *Trichosanthes cucumerina* seed with all solvents was conducted over the range of 28 to 78 °C. The increase in bio-oil yield (hexane as solvent) from $18.34 \pm 0.15\%$ to 25.3 ± 0.3 was obtained as given in Table 3. The rise in extraction temperature increases the bio-oil yield, due to the mass transfer rate and solubilization of hexane. The lower viscosity (0.32 mPa s) and surface tension (18.43 mN/m) of hexane which improves the diffusivity and solubilization inside the solid matrix at higher temperature resulted in maximum extraction rate. The solvent dissolution capacity also would increase the bio-oil yield. On the further rise in temperature beyond 68 °C (boiling point of hexane), the bio-oil yield content decreased³⁹. This rise in temperature increased the solvent boil off and reduced the active contact area between solid and liquid phases⁴⁰. The hexane provided the maximum

Fatty acids	Molecular formula	Molecular weight	Retention time	%Peak area
Hexanoic acid	C ₆ H ₁₂ O ₂	116	9.31	0.2581
Nonanoic acid	C ₉ H ₁₈ O ₂	158	11.71	0.0418
Undecanoic acid, ethyl ester	C ₁₃ H ₂₆ O ₂	214	13.85	0.0806
Octanoic Acid	C ₈ H ₁₆ O ₂	144	14.04	2.7896
2-Decenal, (E)-	C ₁₀ H ₁₈ O	154	15.34	0.3680
2-Dodecanone	C ₁₂ H ₂₄ O	184	15.91	0.0403
2H-Pyran-2-one, 6-ethyltetrahydro-	C ₇ H ₁₂ O ₂	128	16.25	0.6635
2,4-Decadienal	C ₁₀ H ₁₆ O	152	16.82	0.3009
2H-Pyran-2-one, tetrahydro-6-propyl-	C ₈ H ₁₄ O ₂	142	22.86	0.2332
Dodecanoic acid	C ₁₂ H ₂₄ O ₂	200	27.67	13.1398
2H-Pyran-2-one, 6-hexyltetrahydro-	C ₁₁ H ₂₀ O ₂	184	32.62	0.1087
Decanoic acid, ethyl ester	C ₁₂ H ₂₄ O ₂	200	34.41	0.0549
Tetradecanoic acid	C ₁₄ H ₂₈ O ₂	228	34.61	7.5717
2-Nonadecanone	C ₁₉ H ₃₈ O	282	37.08	0.0885
Hexadecanoic acid, ethyl ester	C ₁₈ H ₃₆ O ₂	284	38.85	0.0460
n-Hexadecanoic acid	C ₁₆ H ₃₂ O ₂	256	39.12	13.2114
Oleic Acid	C ₁₈ H ₃₄ O ₂	282	42.62	43.2662
Octadecanoic acid	C ₁₈ H ₃₆ O ₂	284	42.84	12.3392
Hexadecanoic acid, 2-hydroxy-1-(hydroxymethyl)ethyl ester	C ₁₉ H ₃₈ O ₄	330	43.56	0.2971
Z,E-3,13-Octadecadien-1-ol	C ₁₈ H ₃₄ O	266	43.81	0.4736
9,12,15-Octadecatrienoic acid, (Z,Z,Z)-	C ₁₈ H ₃₀ O ₂	278	44.81	3.1031
9-Octadecenoic acid (Z)-, 2-hydroxy-1-(hydroxymethyl)ethyl ester	C ₂₁ H ₄₀ O ₄	356	46.39	1.5240

Table 4. Fatty acid composition of *Trichosanthes cucumerina* bio-oil through GC–MS analysis.

amount of oil extraction ($27.81 \pm 0.4\%$) with the optimum temperature of $68\text{ }^{\circ}\text{C}$. It was detected that the solubility of the solvent increased with an increase in the diffusion rate⁴¹. Sayyar et al.¹⁶ considered the role of extraction temperature, the higher percentage of bio-oil yield of 47.3% from *Jatropha curcas* using hexane as the solvent at $68\text{ }^{\circ}\text{C}$. Milan D. Kostic et al.⁴² optimized the extraction of bio-oil from hemp seeds using n-hexane solvent. The results show that the extraction temperature of n-hexane (at $70\text{ }^{\circ}\text{C}$) increased the hemp seed oil yield because of improved oil solubility in the solvent. Karthikeyan Murugesan and Renganathan Sahadevan²⁷ optimized the non-edible bio-oil extraction from *Cassia javanica* seeds using different solvents. The hexane was the best extractor to extract the maximum percentage of bio-oil yield was about 24.4% with the optimum conditions (extraction temperature at $68\text{ }^{\circ}\text{C}$ and extraction time at 3.5 h).

GC–MS Analysis. The GC–MS test analysis (refer Table 4) was performed to measure the various factors of bio-oil, presence of FFA composition and different types of hydrocarbons^{43,44}. The *trichosanthes cucumerina* bio-oil organic compounds were measured by the test method of gas chromatography (Fig. 9), model PerkinElmer Clarus 500 coupled with a mass spectrometer. The Capillary Column Elite-5MS was used for the separation of components in bio-oil fraction for the length of 30 m. The flow rate of carrier gas fixed at 1 mL/min and the GC was operated at 58.3 min for the helium flow rate of 1 mL/min. The temperature ranges from 150 to $280\text{ }^{\circ}\text{C}$ at $10\text{ }^{\circ}\text{C}/\text{min}$ was maintained in the column. A sample volume of 1.0 μL *trichosanthes cucumerina* bio-oil in chloroform was injected through a split mode, with 1:10 split ratio. The MS condition for the mass range was set to 40 to 450 amu with the complete mode of electron ionization with the electron energy of 70 eV. The components in the sample have been identified using Turbomass ver 5.2.0 software and NIST 2005 mass spectral library. From the GC–MS analysis, 22 free fatty acids were found as listed in Table 4. The oleic acid (C₁₈H₃₄O₂) contributed 43.27%, dodecanoic acid (C₁₂H₂₄O₂) with 13.14%, n-Hexadecanoic acid (C₁₆H₃₂O₂) with 13.21%, Octadecanoic acid (C₁₈H₃₆O₂) with 12.3% of the total composition percentage. Among them, 61.14% of unsaturated and 34.09% of saturated fatty acids were present in the *trichosanthes cucumerina* bio-oil.

FTIR analysis. The presence of polymer, organic and inorganic materials were identified through the Fourier Transform Infrared (FTIR) Spectroscopy analyzer. An infrared light source scans the test samples and analyses the chemical properties⁴⁵. This analytical method was used to analyze the chemical bonds and its nature, based on its stretching or bending on exposure to infrared radiation. In this study, the Deuterated Tri Glycine Sulphate (DTGS) detector was used, which works on the variation in the temperature and IR radiation intensity. The FTIR spectra were recorded between 500 and 4000 cm^{-1} in the transmission mode for the *trichosanthes cucumerina* bio-oil, as shown in Fig. 10. The various functional groups which were identified are tabulated in Table 5. The C–H stretching at 2922.59 cm^{-1} , 2855.1 cm^{-1} shows the presence of alkanes and the C=C stretch ensure the presence of the aromatic compounds^{17,46}. The presence of oxygenated functional groups (O–H, C–O and C=O) indicated the high percentage of oxygen content in the bio-oil and inferred the acidic nature. The presence of hydrocarbon groups indicates the potential usage of bio-oil as the alternate source of energy⁴⁷.

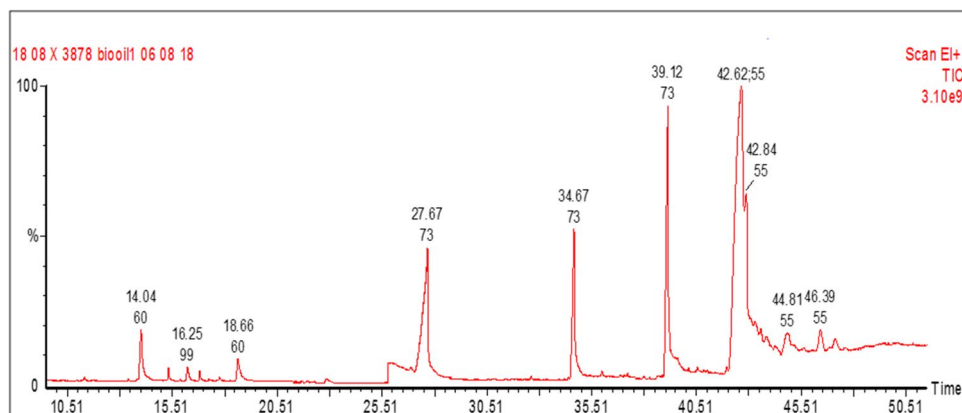


Figure 9. GC-MS Chromatogram of *Trichosanthes cucumerina* bio-oil. The *trichosanthes cucumerina* bio-oil organic compounds were measured by the test method of gas chromatography. It found twenty-two free fatty acids were present in the *trichosanthes cucumerina* bio-oil. It results in the percentage of unsaturated fatty acids as high as saturated fatty acids.

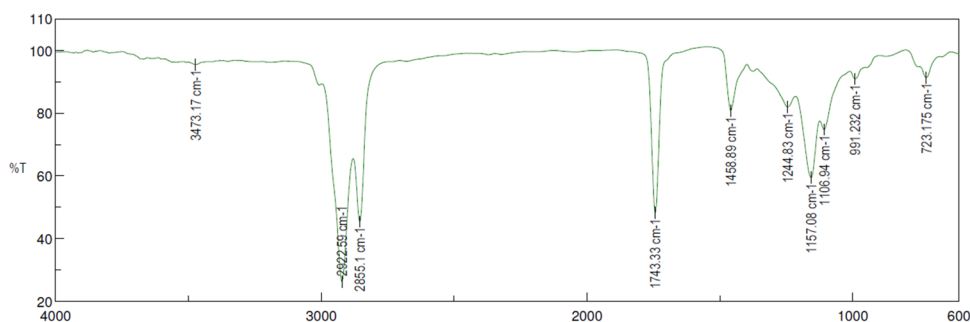


Figure 10. Spectra of *Trichosanthes cucumerina* seed oil according to FTIR analysis. The FTIR spectra were recorded between 500 cm^{-1} and 4000 cm^{-1} in the transmission mode for the *trichosanthes cucumerina* seed oil. The C-H stretching at 2922.59 cm^{-1} and 2855.1 cm^{-1} shows the presence of alkanes. The C=C stretch proves the presence of the aromatic compounds. The presence of oxygenated functional groups (O-H, C-O and C=O) indicates the high percentage of oxygen content in the bio-oil.

Wave Number (cm^{-1})		Functional groups	Compound class	References
Theoretical Range	Experimental value			
3200–3600	3473.17	O–H Stretching	Phenols, Alcohols	48
2810–3000	2922.59	C–H Stretching	Aliphatics–Alkanes	
2810–3000	2855.1	C–H Stretching	Aliphatics–Alkanes	
1715–1755	1743.33	C=O Stretching	Aldehydes and Ketones	49
1400–1550	1458.89	C=C Stretching	Aromatic	50
1210–1260	1244.83	C–O Bending	Ester	51
1110–1210	1157.08	C–OH Stretching	Tertiary Alcohol	47
1080–1160	1106.94	C–O Stretching	Aliphatic ether, Secondary alcohol	52
950–1010	991.232	=C–H Bending	Alkene	47
670–810	723.175	C–H Bending	Aromatic	49

Table 5. Functional groups of *Trichosanthes cucumerina* bio-oil according to FTIR analysis.

Fuel properties. The fuel properties of *Trichosanthes cucumerina* bio-oil (TCO), *Trichosanthes cucumerina* biodiesel (TCB) and Diesel fuel (DF) are tested as per the AOAC and ASTM standards and carried out at Delta Inspection & Research Laboratory, Chennai, Tamil Nadu, India. The properties like kinematic viscosity (ν), fuel density (ρ), heating value, iodine number and saponification value (SV), cetane number, flash and fire points were checked and listed in Table 6.

Fuel properties	Diesel fuel	TCO	TCB	Standard methods
Chemical formula	C ₁₀ H ₂₂	C ₁₆ H ₃₀ O ₂	C ₁₂ H ₂₆ O ₂	Calculation
Molecular weight (g/mol)	142.282	252.487	203.61	Calculation
Kinematic viscosity (cSt)	2.71	42.85	4.26	ASTM D 445
Density (kg/m ³)	836	912	856	ASTM D 4052
Heating value (MJ/kg)	42.5	32.45	38.50	ASTM D 3286
Cetane number (CN)	51	35	44	ASTM D 613
Flash Point (°C)	67	236	158	ASTM D 93
Fire Point (°C)	86	262	169	ASTM D 93
Acid value (mg KOH/g)	0.1	2.16	1.74	ASTM D 664
Sulphur (mg/kg)	29	–	–	ASTM D 2622
Iodine value (gI ₂ /100 g)	38.3	82.4	67.2	AOAC CD1-25
Saponification value (mg KOH/g)	–	121.37	74.3	AOAC CD3-25

Table 6. Comparison of DF, TCO and TCB fuel properties.

Fuel density (ρ) and kinematic viscosity (ν). The fuel density was measured at the reference temperature of 15 °C using hydrometer and found to be 8.33% (TCO) and 6.14% (TCB) higher than diesel. The higher number of unsaturated fatty acid contents in the bio-oil increases the molecular weight. It results in higher fuel density which leads to increased compression ratio, specific fuel consumption and the rate of oxidation. The TCB density was very closer to the value of diesel fuel caused more thermal stability of the biodiesel^{53,54}. The Redwood viscometer was used to measure the viscosity. The TCO obtained 42.75 cSt of kinematic viscosity which was 14.82 times higher than that of diesel which attributed to an increase in the free fatty acid chain, the breakdown of intermolecular forces and adhesion between biofuel molecules⁷. The TCB resulted in decreased kinematic viscosity and 9.05 times lesser than TCO, which improves the fuel characteristics of atomization and vaporization and provides better engine performance⁵⁵.

Other properties. The other properties like fuel heating value, iodine number, saponification value, cetane number can also be referred in Table 6. The fuel heating values were numerically calculated using Dulong's formula⁵⁶. The higher proportion of oxygen content in the bio-oil resulted in lower heat energy (32.45 MJ/kg) as compared to diesel. The heating value of TCB moderately increased with the trans-esterification to 38.50 MJ/kg. It can release a higher amount of heat energy during fuel combustion, which improves engine performance^{57,58}. The percentage of unsaturated FFA present in the bio-oil was measured as the iodine number. The *Trichosanthes cucumerina* bio-oil has an iodine number of 82.4. The degree of unsaturation affects the thermal stability of the fuel and results in carbon deposits⁵⁹. The saponification values were calculated as the amount of potassium hydroxide required for the complete hydrolysis per gram of bio-oil. The saponification value of the *Trichosanthes cucumerina* bio-oil obtained as 121.37 mg KOH/g. In the other reported works, the saponification values of 192 and 212 mg KOH/g were obtained for *Calophyllum inophyllum* L. and *Ulva lactuca* seeds respectively^{24,60} which are higher than the value obtained in this research. The higher saponification values may result in corrosion problem in the diesel engine⁶¹. The cetane number of 51, 35 and 44 was obtained for diesel, TCO and TCB respectively. Higher cetane number of TCB entails shorter ignition delay leads to improved diffusion part of combustion equated with premixed phase^{39,62}.

Conclusion

Five different solvents were used to synthesis the bio-oil from *Trichosanthes cucumerina* seeds. The hexane was found to be the better solvent as compared with other solvents. The maximum bio-oil yield was achieved of 28.4 ± 0.4% at a temperature of 68 °C, 0.21 mm crushed seed size, 6% moisture content, 270 min extraction time, and 6:1 (ml/g) of solvent to seed ratio. The FTIR analysis shows the higher percentage of oxygen content and less sulphur content in the *trichosanthes cucumerina* bio-oil. The GC-MS results are seen with the saturated (43.27%) and unsaturated fatty acids (61.14%). The biodiesel was produced from *trichosanthes cucumerina* bio-oil through the three stages of the trans-esterification process, which produced 93.4 ± 0.2% of biodiesel. The physicochemical properties of the *trichosanthes cucumerina* bio-oil and biodiesel were analyzed using AOAC and ASTM standards. The biodiesel properties obtained (heating value—38.74 MJ/kg, kinematic viscosity—4.26 cSt and cetane number—44) were closer to diesel fuel, and it can be considered in the diesel engine as an efficient alternative source.

Data availability

The datasets generated during and/or analyzed during the current study are available from the corresponding author on reasonable request.

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Author contributions

R. Manimaran as being the corresponding author, was worked on the synthesis of bio-oil, the conversion of biodiesel and testing. K. Murugu Mohan Kumar has guided in the processing and analysis of test results. N. Sathiyarayanan contributed in the English reading and corrections. All authors are thankful to the unknown reviewers for their valuable inputs to improve the quality.

Competing interests

The authors declare no competing interests.

Additional information

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