Present investigation evaluates the LDPE (low-density polyethylene) biodegradation efficiency of polymer degrading bacteria along with UV, nitric acid and surfactant treatments. In current scenario LDPE contamination reported as dominant pollutant in terrestrial and aquatic ecosystem due to its expulsion from commercial and domestic practices. Biodegradation serve as an innovative and effective approach to waste management as compared to land filling and burning processes. The outcomes of UV, nitric acid and surfactant treatments on polymer degradation in addition to bacterial treatment were determined by SEM, FT-IR and electrical conductivity analysis.

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Specifications Table

| Subject area                  | Microbiology, Ecology, Biodegradation |
|-------------------------------|---------------------------------------|
| More specific subject area    | Outcomes of UV, nitric acid and surfactant treatments on biodegradability of low-density polyethylene samples |
| Type of data                  | Tables, Figures, Text file            |

* Corresponding author.

E-mail address: vinaymohanpathak@gmail.com (V.M. Pathak).
How data was acquired
Exploitation of UV, nitric acid and surfactant treatments along with bacterial strains; SEM, FT-IR and electrical conductivity of polymer film was analyzed;

Data format
Analyzed

Experimental factors
Role of physical and chemical treatments on LDPE biodegradation

Experimental features
The relationship between the physical, chemical and biological treatments

Data accessibility
The data are available with this article

Value of the data

- This data could be used as systematic tool for increasing polymer degradation.
- This data will also help in developing the specific and appropriate approach for polymer degradation in a sustainable manner.
- This data represented the impact of physical and chemical treatments on the LDPE biodegradation.

1. Data

The dataset of this article described the consequence of physical and chemical treatments, which include UV, nitric acid and surfactant treatments in LDPE degradation in addition to polymer degrading bacterial strains (*Bacillus subtilis* V8, *Paracoccus aminophilus* B1 4-, *Pseudomonas putida* C 2 5, *Pseudomonas aeruginosa* V1 and *Acinetobacter calcoaceticus* V4). The Figs. 1–3 show the scanning electron microscopy (SEM) micrographs of UV, nitric acid and surfactant treated biodegraded polymer

![Fig. 1.](image-url)

*Fig. 1.* Scanning electron microscopy (SEM) micrographs of UV treated biodegraded polymer samples A- control (without UV treatment), B- *Bacillus subtilis* V8 + UV treated polymer, C- *Paracoccus aminophilus* B1 4- + UV treated polymer, D- *Pseudomonas putida* C 2 5 + UV treated polymer, E- *Pseudomonas aeruginosa* V1 + UV treated polymer, F- *Acinetobacter calcoaceticus* V4 + UV treated polymer after incubation.
Fig. 2. Scanning electron microscopy (SEM) micrographs of nitric acid treated biodegraded polymer samples A- control (without nitric acid treatment), B- *Bacillus subtilis* V8+ nitric acid treated polymer, C- *Paracoccus aminophilus* B1 4+ nitric acid treated polymer, D- *Pseudomonas putida* C 2 5+ nitric acid treated polymer, E- *Pseudomonas aeruginosa* V1+ nitric acid treated polymer, F- *Acinetobacter calcoaceticus* V4+ nitric acid treated polymer after incubation.

Fig. 3. Scanning electron microscopy (SEM) micrographs of surfactant treated biodegraded polymer samples A- control (without surfactant treatment), B- *Bacillus subtilis* V8+ surfactant treated polymer, C- *Paracoccus aminophilus* B1 4+ surfactant treated polymer, D- *Pseudomonas putida* C 2 5+ surfactant treated polymer, E- *Pseudomonas aeruginosa* V1+ surfactant treated polymer, F- *Acinetobacter calcoaceticus* V4+ surfactant treated polymer after incubation.
samples, the Figs. 4–6 show the FT-IR characteristics of UV, nitric acid and surfactant treated biodegraded polymer samples with control. Figs. 7–9 show the electrical conductivity property of UV, nitric acid and surfactant treated biodegraded polymer samples. Electrical conductivity examination is an advanced approach for analyzing the electrical properties of polymer. The examination of electrical conductivity is used for determining the consequences for bacterial growth on polymer surface [3,4,12,17,22]. Bacterial treatment helps in developing electrical conductivity due to bacterial growth on polymer surface and the bacterial presences on it. Such techniques serve as the promising way to analyze the alteration in pure and treated polymer samples. These techniques are facilitated to examine the reproducibility of degradations.

2. Experimental design, materials and methods

The consequences of UV, nitric acid and surfactant treatment on polymer degradation in addition to bacterial treatment were determined by SEM, FT-IR and electrical conductivity analysis. Treated polymer samples were exploited in biodegradation experiment to increase the biodegradation ability of isolated bacterial strains. The 365 nm wave length UV light was used to irradiate polymer strips. Polymer (LDPE) strips of 2 cm diameter (weight 0.247–0.408 g) were placed into a UV box (14 by
Fig. 4. (continued)
26 cm) at a distance of 3 cm from the light source for 8 weeks. UV treated polymer strips were aseptically transferred to mineral broth medium and subjected to biodegradation by isolated bacterial cultures [10,13,14]. In nitric acid treatment strips were treated with nitric acid (99.0%) at 80 °C for 6 days. After nitric acid treatment polymer strips were aseptically transferred to mineral broth medium and subjected to biodegradation by isolated bacterial cultures [2,7,14,21]. The growth of microorganisms effected in the presence of surfactant. Tween 80 (nonionic surfactant) was added at concentration of 0.05% v/v to the media to test the effect of these substances on bacterial attachment to polymer and LDPE degradation by bacterial isolates compared with control sample [1,6,7,9,19,20]. At the end of incubation samples were recovered from culture media and analyzed for degradation by using SEM, FT-IR and electrical conductivity examination [3–5,8,11,12,15,16,18].

Fig. 5. Graph A-E for bacterial treatments with control (untreated LDPE). FT-IR spectra in A for treatment (Bacillus subtilis V8 + nitric acid treated LDPE), B for treatment (Paracoccus aminophilus B1 4- + nitric acid treated LDPE), C for treatment (Pseudomonas putida C 25 + nitric acid treated LDPE), D for treatment (Pseudomonas aeruginosa V1 + nitric acid treated LDPE) and E for treatment (Acinetobacter calcoaceticus V4 + nitric acid treated LDPE).
Fig. 5. (continued)

FT-IR spectra of treatment (Pseudomonas putida C25+ nitric acid treated LDPE) and control

FT-IR spectra of treatment (Pseudomonas aeruginosa V1 + nitric acid treated LDPE) and control

FT-IR spectra of treatment (Acinetobacter calcoaceticus V4+ nitric acid treated LDPE) and control
Fig. 6. Graph A-E for bacterial treatments with control (untreated LDPE). FT-IR spectra in A for treatment (*Bacillus subtilis* V8 + SiO$_2$ + LDPE), B for treatment (*Paracoccus aminophilus* B1 4- + SiO$_2$ + LDPE), C for treatment (*Pseudomonas putida* C2 5 + SiO$_2$ + LDPE), D for treatment (*Pseudomonas aeruginosa* V1 + SiO$_2$ + LDPE) and E for treatment (*Acinetobacter calcoaceticus* V4 + SiO$_2$ + LDPE).
Fig. 6. (continued)

FT-IR spectra of treatment (*Pseudomonas putida C 2 5*) and control

FT-IR spectra of treatment (*Pseudomonas aeruginosa V1*) and control

FT-IR spectra of treatment (*Acinetobacter calcoaceticus V4*) and control
Fig. 7. Electrical properties of UV treated biodegraded polymer samples, graph A for control sample (without treatment), B- Bacillus subtilis V8+ UV treated LDPE, C- Paracoccus aminophilus B1 4+ UV treated LDPE, D- Pseudomonas putida C 25+ UV treated LDPE, E- Pseudomonas aeruginosa V1+ UV treated LDPE and F- Acinetobacter calcoaceticus V4+ UV treated LDPE polymer samples.
Fig. 7. (continued)
Fig. 7. (continued)
Fig. 8. Electrical properties of nitric acid treated biodegraded polymer samples, graph A for control sample (without treatment), B- *Bacillus subtilis* V8+ nitric acid treated LDPE, C- *Paracoccus aminophilus* B1 4+ nitric acid treated LDPE, D- *Pseudomonas putida* C 2 5+ nitric acid treated LDPE, E- *Pseudomonas aeruginosa* V1+ nitric acid treated LDPE and F- *Acinetobacter calcoaceticus* V4+ nitric acid treated LDPE polymer samples.
Fig. 8. (continued)
Fig. 8. (continued)
Fig. 9. Electrical properties of surfactant treated biodegraded polymer samples, graph A for control sample (without treatment), B- *Bacillus subtilis* V8+ surfactant treated LDPE, C- *Paracoccus aminophilus* B1 4+ surfactant treated LDPE, D- *Pseudomonas putida* C 2 5+ surfactant treated LDPE, E- *Pseudomonas aeruginosa* V1+ surfactant treated LDPE and F- *Acinetobacter calcoaceticus* V4+ surfactant treated LDPE polymer samples.
Fig. 9. (continued)
Fig. 9. (continued)
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