Palm Oil Mill Effluent Treatment Processes—A Review

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Abstract: Ponding system or land application techniques are widely used at industrial scale to treat palm oil mill effluent (POME) prior to discharge to the environment. POME is considered as one of the major problems that has generated voluminously from the palm oil industries. The main purpose of this article is to organize the scattered available information on various aspects and a wide range of promising current POME treatments including biological microorganisms, physicochemical methods of coagulation, and membrane and thermochemical process. In addition, the integrated system of anaerobic-aerobic bioreactor (IAAB), which has been touted as highly efficient with easy control at acceptable temperature range and shorter treatment time, has potential to be exploited for POME treatment. The main influencing factors for IAAB POME treatment are highlighted as outstanding characteristics for challenges and future prospects.

Keywords: palm oil mill effluent; ponding system; biological treatment; physicochemical treatment; membrane treatment; thermochemical treatment

1. Introduction

Now, palm oil is recognized as high-quality oil used popularly and primarily for cooking in developing countries. Thus, oil palm tree plantations are spreading across Asia, Africa, and Latin America. From the viewpoint of soil and water pollution, palm oil production provides serious problems. Palm oil mill effluent (POME) exists in forms of high solid, oil, and grease and includes biochemical oxygen demand (BOD) and chemical oxygen demand (COD) in richness. These are major concerns in environmental pollutant due to their adverse effects to many forms of life [1]. The direct discharge of POME in the environment is a matter of concern for both toxicological and esthetical reasons. The majority of the palm oil mills use the addition of water during the extraction process of crude palm oil (CPO) from the fresh fruit bunch (FFB). As a result, a considerable amount of wastewater was generated [2] and stored in ponds. In major oil palm-producing nations such as Indonesia and Malaysia, the production of CPO has resulted in an annual production of up to 3 billion pounds of POME. Furthermore, it is estimated that 5–7.5 tonnes of water are required to produce 1 tonne of CPO, with more than 50% of the water ending up as POME [3,4].

POME exists as viscous and brownish liquid containing approximately more than 95–96% of water, together with 0.6–0.7% of oils and 2–4% of suspended solids for the source of fruit debris, and is known as a non-toxic substance [5]. The brownish colour is due to the origins of carotene (8 mg/L), pectin (3400 mg/L), tannin, phenolic (5800 mg/L), and lignin (4700 mg/L), thus making it a nutrient-rich substance [5]. POME is known to be a major problem in every oil palm factory due to its voluminous abundance and disposal problems. The direct discharge of POME on the land results in clogging, water logging of the soil, and kills the vegetation on contact. Discharge of the effluent into waterway causes water depletion, resulting in aquatic pollution [6]. In extreme cases, river water consequently turns brown with an unpleasant smell and viscous characteristic. Mortality of fish and...
other aquatic organism increases, and local people are deprived of the local water sources for domestic uses and fishing. Therefore, the effective techniques for POME treatment are of the utmost significance.

During the past two decades, several biological, physical, and chemical techniques have been reported as treatment methods; few, however, have been accepted by the industries [7]. Amongst the numerous and latest alternative treatment techniques are integrated systems of anaerobic-aerobic bioreactor (IAAB). The procedure of choice gives the best results in POME treatment as IAAB can reduce COD and BOD within 115 days [8]. Recently, numerous approaches have been studied for the development of cheaper and effective treatment techniques. Many non-conventional low-cost methods including utilization of microorganisms were proposed as techniques of POME treatments [9–19].

The present article deals with the technical feasibility of various non-conventional and highly effective treatments for POME. The main aim of this review is to provide a summary of recent topics concerning the optimum treatment output with low COD, BOD, total suspended solid (TSS), and colour with short treatment time. Science Direct were searched (final search February 2021) using a combination of search terminology of palm oil effluent, treatment, and fermentation. Among all mooted POME treatment processes, only a final selection of 124 papers fostering the most effective treatment processes was identified and included as references.

2. Palm Oil Production Process and Oil Trapping Pond System

The majority of the oil palm mill employed a wet process for the oil extraction, utilizing fresh fruit bunch (FFB). No chemicals are added during the oil extraction process from the oil palm fruits, thereby making all wastes generated as non-toxic to the environment. By using United Oil Palm (UOP) Industries Sdn. Bhd., Malaysia as a reference point, the treatment process starting from the FFB and the extent of the properties changes occurring after decomposition at each individual pond until final discharge was presented in Figure 1.

For grading, initially, FFB is retrieved from the farm and transported to the mills prior to processing. As categorized accordingly based on their weight and quality, the grading step is crucial to ensure only good quality of FFB selected and can eliminate the deteriorated FFB. The utilization of damaged palm fruits will render low-quality palm oil. The selected good quality FFB were dropped onto a ramp and transferred into horizontal sterilizer cages.

The sterilizer cages with the FFB were exposed to high-temperature pressurized steam under 40 psi for 75–90 min at 140 °C. There are four objectives of the FFB sterilization: (1) most importantly, to deactivate the enzymes responsible for the formation of free fatty acids; (2) to remove external impurities; (3) to soften and loosen the fruitlets from bunches; and (4) to detach the kernels from the shells prior stripping process. The sterilization process acts as the main contributor to the POME accumulation in the form of sterilizer condensate. Also, the sterilized fruit bunches were fed and lifted into a rotating drum. In the drum stripper, the fruitlets were detached and separated from the bunches. At this stage, the loose fruitlets were softened and digested under steam-heated conditions by a steam jacket or by direct live steam injection. Simultaneously, the high pressure and high temperature broke the oil-bearing cells of the mesocarp and were channelled to a mechanical twin screw-press machine for crude palm oil (CPO) extraction [20,21] following clarification.

It was known that the digested CPO consisted of 35–45% palm oil, 45–55% water, and residual fibrous materials and was pumped into a clarification tank in order to separate the oil from CPO at 90 °C in order to further enrich the oil separation process. Then, the oil was skimmed-off from the top of the tank. The bottom phase of the clarification tank still contained remaining residual oil and again passed through the sludge separator, namely decanter heavy phase (DHP) tank and de-oiling tank. The sludge is the by-product of the clarification, and the purification process is the second source of POME in terms of its polluting strength and quantity. The recovered oil was returned to the clarification tank.
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Figure 1. Palm oil production process and oil trapping pond for POME treatment in Malaysia.

In the depericarping, the nuts were removed by the digestion and pressing processes for depericarp and then the residual fibres transported to the nut cracking machine or plant. The cracked mixture of kernels and shells was then separated in a hydrocyclone and claybaths. The third source of POME was fed from the wash-water of the hydrocyclone. The kernel produced was then stored into kernel bunkers before being transferred to the mill for oil extraction. Shell wastes were piled with the mesocarp fibers in the boiler for power generation. The remaining wastewater was discharged accordingly to their subsequent treatment process. Overall, the production of CPO will generate a by-product of solid wastes such as mesocarp fibres and EFB. These solid wastes still can be utilized for other purposes. Raw POME is an example of liquid waste and is subjected to various stages of pond treatments. Detailed elaboration associated with POME treatment will be discussed in the next sub-topic. In contrast, the de-oiling tank feeds POME waste oil to the trapping pound. The ponding system involved several series of ponds of acidic, cooling, anaerobic, aeration (aerobic), facultative and final polishing ponds.

3. Characteristics of Palm Oil Mill Effluent (POME)

Typically, in the process of oil palm, POME is accumulated from three different sources such as clarification wastewater, sterilizer condensate, and hydro-cyclone wastewater [22]. The general characteristics of each raw POME are presented in Table 1.
Table 1. General characteristics of raw POME.

| Parameters                          | Concentration Range | Current Discharge Limit |
|-------------------------------------|---------------------|-------------------------|
| Chemical oxygen demand (COD)        | 15,000–100,000      | -                       |
| Biochemical oxygen demand (BOD)     | 10,250–43,750       | 100                     |
| Total suspended solids (TSS)         | 5000–54,000         | 400                     |
| Oil and grease                      | 130–18,000          | 50                      |
| Temperature                         | 80–90               | 45                      |
| pH                                  | 3.4–5.2             | 5–9                     |

All values, except pH and temperature, were expressed in mg/L.

The raw POME is a thick brownish colour in the viscous and colloidal suspension, which contains high water contents (95–96%), suspended solids (2–4%), and oils (0.6–0.7%). POME exists as an acidic substance with unpleasant odour, high value of COD, and BOD, which can be concomitantly sensitive to environmental issues [23,24]. Many researchers reported on the POME nutritional composition, as well as the minerals, trace elements, and heavy metals [23,25]. The XRF analysis of POME, sampled in Penang, Malaysia, was obtained below; C 51.0%, O 35.3%, Na 0.0632%, Mg 1.09%, Al 0.215%, Si 0.552%, P 0.429%, S 0.553%, Cl 2.75%, K 6.77%, Ca 1.09%, Mn 0.0243%, Fe 0.141%, and Rb 0.0286%. The quality of the effluent produced was affected by many factors including maturation degree of FFB, different treatment systems, as well as varied batches of FFB and factories system [26]. Hence, in order to control and minimize the pollution impact towards the environment, the Department of Environment (DOE), Malaysia, had enacted Environmental Quality Act (EQA) 1974 and set standard limits for the discharge of POME into the environment [27].

4. POME Treatment Process

4.1. Ponding System or Land Application

Ponding system or land application is the most widely used at the majority of palm oil mills (85%) for effluent treatment, due to cost-effectiveness, low maintenance costs, energy efficiency, system reliability, and simple design [28]. However, this obsolete ponding system is often necessary for long hydraulic retention time (HRT) and large pond area. This is considered to be inefficient in certain ways [29].

The configuration of POME treatment via a pond system is also known as waste stabilization ponds of acidic to final polishing ponds contain an overall HRT of 100–120 days prior to discharge to the environment. The raw POME in the oil-trapping pond was pumped into the acidification pond and retained for about 6 days. Then, the POME was pumped into a cooling pond through a cooling tower and retained for another 7 days. Cooling pond reduced the POME temperature to be 35–38 °C and stabilised the pH prior to the anaerobic stage. While cooling, high initial temperature of POME promotes thermal and anaerobic digestion, mainly via hydrolysis and acidogenesis. Anaerobic treatment ponds were noticed as the most effective method to treat high-strength wastewater by anaerobic decomposition.

The hydrolysis process involved the breakdown of complex polymers such as carbohydrates, proteins, and lipid into respective monomers. For example, sugar, amino acids, and fatty acids were facilitated by the thermo-tolerant microbes. In the acidogenesis process, carbon-containing monomers are fermented into volatile fatty acids (VFA), such as acetic acid (HAc), propionic acid (HPr), butyric acid (HBu), valeric acid (HVa) with trace-non VFA, and lactic acid [23]. Acetogenesis reduced the long chain VFA (HPr, HBu and HVa) from the acidogenesis into hydrogen and carbon dioxide (CO₂). Hydrogen and CO₂ were then consumed by hydrogenotrophic methanogens, while HAc and CO₂ were utilized by acetoclastic methanogens that produced methane gas. The methane gas is considered as the final value-added product for the biogas production. Following degradation of the long chain fatty acids (LCFA) and VFA, anaerobically treated POME became alkaline and existed as blackish brown due to partial degradation of the lignin into the phenolic [24]. Anaerobic treatment stages comprised of four ponding series with total HRT of 54–60 days.
Then, anaerobic POME is further treated using a three series of aeration ponds with the floating aerators for about 20 days before being discharged into the facultative ponds. Facultative ponds consisted of three ponding series, which are very crucial to further reduce the organic content in the effluent, prior to discharging to the waterway, as complied with the Environmental Quality Act (EQA) 1974, the stipulated regulation for crude palm oil treated effluent discharge. Through sedimentation requiring approximately 2 days, final polishing pond clarifies suspended microbes from the aerobically-treated POME. This ponding system has been described by many researchers [30,31] successes for effluent treatment and complies with the disposal standard requirement with some limitation to the minimum disposal standard requirement [32].

The anaerobic ponding system was found to demonstrate great performance in treating POME due to reduction of high-level organic properties in POME [33]. However, 100% decolourization was unable to be achieved using this treatment system [34]. Open ponding system successfully reduced the levels of COD (100–1725 mg/L), BOD (100–610 mg/L) and ammoniacal nitrogen (100–200 mg/L) to a certain range. Nevertheless, the open ponding system requires extensive land area, concomitantly leading to serious environmental problems by the production of methane gas, resulting in ozone layer depletion [35]. Furthermore, the open and closed digesting tanks were commonly exploited after the ponding system, due to its availability on a wide range of volumetric capacity. Although these types of tanks had low capital and operating costs, with shorter HRT (20–25 days) and limited land area requirement [36], they still had a significant amount of harmful biogas (approximately 5.5 kg of CH4 per tonne), which was produced due to the lack of a gas retrieval device in the system [37]. Even though the implementation of CH4 capturing system provides various advantages such as reducing the greenhouse gaseous emission, generating renewable energy, and improving soil quality and FFB production, the system is too costly for the next stage in commercialisation [38].

4.2. Biological Treatment

Although ponding system is popular for POME treatment technique at the industrial scale, large land area (30–45 acres) and long HRT (100–160 days) entail an economical drawback. Thus, there is a need for the development of low-cost and utilization of available materials, which can be used more economically at the industrial scale. Due to the aforementioned problems, research interest has increased regarding the alternative production technique to partially replace the obsolete open ponding system. Especially for POME treatment by biological techniques in the presence of bacteria, fungi, yeast, microalgae, and other microbial cultures, which have been the subjects of many recent researcher.

Table 2 summarizes various biological processes for POME treatments. In contrast to the ponding system or land application, the anaerobic system would be more preferable, due to higher organic conversion efficiency of COD, BOD, TSS, and colour, followed by oil and grease [8]. Nowadays, the anaerobic system is claimed to be unique and beneficial for environmentally and friendly stabilized systems and has the ability to produce an energy source in the form of the methane gas, CH4 [8,20,31]. Emerging from the current anaerobic treatment, some modifications on the anaerobic treatment system have been developed for the POME wastewater by aerobic suspended growth processes, attached growth anaerobic process with immobilised cell bioreactors, anaerobic fluidised bed reactors and anaerobic filters, anaerobic sludge blanket processes using both up-flow anaerobic sludge blanket reactor and anaerobic baffled bioreactor, membrane separation anaerobic treatment process, and integrated anaerobic treatment processes [14,31]. Chan et al. [39] concluded that the organic matter in an anaerobic effluent is not fully stabilized and additional post treatment was necessary in order to remove some ions’ presence in the effluent. Hence, in order to meet the discharge standard, the anaerobic system was coupled with aerobic degradation of the POME.

Referring to Table 2, the highest COD removal (99.60%) was obtained by utilizing the integrated system of the anaerobic-aerobic batch (IAAB) reactors [8]. IAAB enabled to
operate at organic loading rate (OLR) of 7–12 kg COD/m$^3$day with a continuous mixing flowrate of 140 m$^3$/h. A significant amount of CH$_4$ gas (45–68%) was produced during the treatment. So this revealed high potential POME treatment to efficiently utilize the main substrate for the microbial growth and biogas production [8]. The system was capable of reducing the treatment time to less than 115 days, as compared with the current pond treatment system. In another study, the thermophilically mixed culture was able to reduce up to 90.00% of the COD in the POME within only 6 days for working volume of 2.5 L by batch continuous stirred tank reactor [40]. Until now, there have been insufficient scientific reports associated with the utilization of the aerobic post-treatment anaerobically digested POME. However, sequencing batch reactors (SBR) or activated sludge reactors (ASR) were reported to be utilized as post treatment for anaerobic processes. Better performance was achieved at optimum OLR of 1.8–4.2 kg COD/m$^3$day utilizing SBR on the COD (96.00%), BOD (99.00%) and TSS (99.00%) removal efficiency within 22 h [2], simultaneously leading the discharge limits. The controlled dissolved oxygen and pH were controlled throughout the fermentation, thus demonstrating positive results of using the SBR system. Apart from the anaerobic and aerobic digestion, an integrated system of biological and membrane separation treatment was identified as an ultra-sonic membrane assisted system (UMAS) with maximum COD removal efficiency (98.90%) within 7 days for 200 L volume of raw POME [41]. The aerobic treatment under mesophilic condition varied accordingly with efficient removal of the COD, BOD, and TSS, depending on the different temperatures during the process [14,42].

Phytoremediation is another green technology that has received huge interest over the past decades [15]. The application of phytoremediation in the wastewater treatment offers various advantages in terms of cost-effective alternatives, aesthetic appearances, and sustainability concern. The key element is plant selection prior to proceeding with the implementation of remediation techniques. The suitable properties were high nutrients uptake, as well as heavy metals uptake in the wastewater [43]. Phytoremediation of POME using Typha latifolia was effective in reducing the COD concentration up to 97.18% [44]. The utilization of Vettiveria zizanioides was reported to reduce the COD and BOD concentration effectively up to 94% and 90% within 14 days for 45 L POME volume [15]. Water spinach (Ipomoea aquatic Forsk) was another promising agent for successful reduction of the COD and colour of POME at 86.3% and 95.3%, respectively [45]. This was conducted for 15 days with better achievement in decolourising 75% (7.5 L POME + 2.5 L tap water, respectively) [45]. Furthermore, Baihaqi et al. [46] reported the use of Spirogyra sp. to reduce the COD concentration to 63.6% within 20 days for 75 mL L$^{-1}$ of POME. Interestingly, the majority of the phytoremediation techniques had high potential to reduce the nitrate and phosphate content, as well as the wastewater contaminants including chromium [47], lead [48], and textile dyes [49] in highly polluted media, which have been reported on by Chanu and Gupta [48]. Phytoremediation is easy to handle, cost effective, readily available, and environment friendly [50].

In addition to the aquatic plants such as Pistia stratoites [51] and Eichornia crassipes [52], there were a few attempts in cultivating microalgae in the POME such as Chlorella spp. UKM 8 and Chlamydomonas sp. UKM 6 in reducing the COD concentration up to 92% within 21 days for 2 L of POME under different inoculum sizes that ranged from 5–15% v/v [53]. Hariz et al. [54] utilized two stages of treatment for about 25 days by utilizing Scenedesmus sp. UKM9 and Chlorella sp. UKM2 in treating POME, resulting in 71% reduction of COD, 86% reduction of total nitrogen, and 85% reduction of phosphate with 25 days of treatment for 85% v/v of POME. Overall performance of phytoremediation techniques was evaluated for about 14–25 days of treatment period with varied results.

Due to high content of organic and mineral constituents in POME, these properties were an advantage of convenient fermentation processes as a feasible treatment method. POME accumulated lipid, bioplastics synthesis, citric acid, and bioethanol production by various microorganisms [16,55–57]. The 50% (v/v) POME was found to generate the highest biomass growth (8.09 g/L) and lipid accumulation (1.46 g/L), with lipid content
of 18.04% on dry weight basis utilising *Bacillus cereus* via batch fermentation system [16]. The substantial reduction in pollution load specifically in COD and BOD for 50% (**v/v**) POME demonstrating removal efficiencies of 79.35% and 72.65% within 6 days. The *B. cereus* cultivation in POME could be a promising method for attaining higher biomass growth and lipid production in parallel with the bioremediation concept [16]. By a double-step process [55], the breakdown of POME involved acetic acid and propionic acid, thus continuing with the polyhydroxyalkanoates (PHA) synthesis by *Rhodobacter sphaeroides*. Furthermore, the usage of POME hydrolysated a substrate in producing citric acid under controlled parameters and then effectively reduced the COD level more than 80% and saved the treatment costs, simultaneously minimising the current environmental problems [56]. Spontaneous fermentation of bioethanol utilising POME was performed by Wakil et al. [57] for both, with amendments of glucose, corn steep liquor, and sugarcane bagasse, and without amendments. The highest ethanol content was obtained at 18.17 g/L (2.3%) with the amendment, whereas 3.95 g/L ethanol (0.5%) was achieved without the amendments. In the POME fermentation, the optimum condition was of 30 g/L glucose and 30 g/L sugarcane bagasse at pH 8.5 for 12 days, although there were no reported values on the COD, BOD, and TSS throughout the fermentation. To date, the organic content in POME can be completely converted into useful bioethanol, but the process yet requires complicated handling procedures and time-consuming processes in the industrial scale of the optimizing fermentation condition.

Microbial fuel cell (MFC) has gained attention by many researchers in treating POME. In contrast to the obsolete ponding system, MFC was performed at only room temperature with low substrate concentration levels. With the aid of microorganisms, an organic-rich wastewater was used to generate electrical energy, which is considered as cost-effective and sustainable energy source. Even though POME can be transformed into an electricity source utilizing a single chamber of MFC, still the COD and BOD removal efficiencies remains without satisfaction. The observation of low COD and BOD removal was due to the substrate type, treatment parameters, reactor design, and operational costs [58]. Barani Tharan et al. [59] treated both raw POME and diluted POME in a dual chamber of MFC by using polyacrylonitrile carbon felt and a Nafion 117 membrane. The lower COD removal efficiency (45.00%) was obtained in the raw POME, as compared to the diluted POME (70%) after 15 days of the incubation period. However, the power electricity generation increased with the increment of initial COD value, implying that the MFC system with raw POME performed better than the diluted one. A combination of two-staged MFCs coupled with the immobilised biological aerated filter was conducted in replacing the conventional method of anaerobic biological treatment system. This was an advanced system for successful reduction of the COD up to 96.50% within 10 days operation in 2.36 L of working volume [60]. Although MFC is an easy handling and simple system, some of the limitations including the capital costs and the technical parts of the electron transfer performances still remain as major drawbacks.
| No. | Microbes or Biological Seeding                                                                 | COD (mg/L) | BOD (mg/L) | TSS (mg/L) | Colour (mg/L) | Oil (mg/L) | CO₂ Fixation (g CO₂/L/day) | CH₄ Release(%) | Working Volume (L) | HRT (days or h) | Treatment                                                                 |
|-----|-------------------------------------------------------------------------------------------------|------------|------------|------------|---------------|------------|---------------------------|----------------|---------------------|----------------|--------------------------------------------------------------------------|
| 1.  | Current standard discharge limit                                                                | 100        | 100        | 400        | 200           | 50         | -                         | -              | -                   | -              | DOE, 1982                                                                |
| 2.  | Future standard discharge limit                                                                | -          | 20         | 200        | 100           | 5          | -                         | -              | -                   | -              | DOE, 2015                                                                |
| 3.  | Mesophilic-thermophilic bacterial seed                                                           | 99.60%     | 99.00%     | 99.00%     | -             | -          | 45–68                     | -              | 75                  | 115 d          | Integrated anaerobic-aerobic bioreactor (IAAB).                           |
| 4.  | Thermophilic mixed culture                                                                     | 90.00%     | -          | -          | -             | -          | 64                        | 2.5            | 6                   | 6 d             | Anaerobic, Continuous stirred tank reactor (CSTR).                       |
| 5.  | Activated sludge basins                                                                        | 96.00%     | 99.00%     | 99.00%     | -             | -          | -                         | -              | 1.8                 | 22 h           | Ultra-sonic-assisted batch reactor (SBR).                                 |
| 6.  | Anaerobic culture in POME                                                                       | 98.90%     | -          | 93.00%     | -             | -          | -                         | -              | 200                 | 7 d             | Anaerobic expanded granular sludge bed (EGSB) reactor—Pilot scale steel tank (6 m × 16 m) |
| 7.  | Inoculum from open anaerobic ponds of POME                                                      | 94.89%     | -          | -          | -             | -          | 65–70                      | -              | -                   | 9.8 d          | Anaerobic filter (AFR) with immobilized bio-balls with addition of Ni.    |
| 8.  | POME + inoculum from cow manure digester farm                                                   | 92.00%     | -          | -          | -             | -          | 30                        | 5              | 25                  | 25 d           | Anaerobic, Carbon dioxide fixation.                                      |
| 9.  | Microalgae (Chlorella spp. UKM 8 and Chlamydomonas sp. UKM 6)                                  | 92.00%     | -          | -          | -             | -          | -                         | 15.00%         | -                   | 21 d           | Anaerobic mixed microbial consortium for all bacteria–fungi combination (ABFC). |
| 10. | Micrococcus luteus 101PB, Stenotrophomonas maltophilia 102PB, Bacillus cereus 103PB, Providencia vermicola 104PB, Klebsiella pneumonia 105PB, Bacillus subtilis 106PB, Aspergillus fumigatus 107PF, Aspergillus niger 108PF, Aspergillus nomius 109PF, and Meyerozyma guilliermondii 110PF | 91.06%     | 90.23%     | 92.23%     | -             | -          | -                         | -              | 1                   | 50 d           |阿森纳混合微生物群体。为了所有细菌-真菌组合（ABFC）。                                      |
### Table 2. Cont.

| No. | Microbes or Biological Seeding | Concentration, mg/L or Removal Efficiency, % | Process Aspects | Ref. |
|-----|--------------------------------|---------------------------------------------|-----------------|-----|
| 11. | Mixed culture: (Bacillus cereus 103 PB, and Bacillus subtilis 106 PB) | COD (mg/L): 90.64% | Working Volume (L): 0.25 | Aerobic fermentation using single reactor. | [64] |
| 12. | Thermophilic mixed culture | BOD (mg/L): 93.11% | HRT (days): 5 d | Anaerobic digestion | [65] |
| 13. | Fungal strains Emericella nidulans NFCCI 3643 | TSS (mg/L): 90.00% | CH4 Release (%): 0.25 | | [66] |
| 14. | Thermophilic mixed culture | Colour (mg/L): 88.23% | CO2 Fixation (g CO2/L/day): 0.25 | Anaerobic Fermentation (Shake flasks) | [9] |
| 15. | Saccharomyces cerevisiae strain | COD: 86.30% | Working Volume (L): 52 | Anaerobic fluidised bed reactor (AFBR) | [67] |
| 16. | Moringa oleifera extract | TSS: 97.30% | CO2 Fixation (g CO2/L/day): 4.5 | Anaerobic co-digestion. | [68] |
| 17. | Microalgae (Scenedesmus sp. UKM9 and Chlorella sp. UKM2) | BOD (mg/L): 98.30% | HRT (days): 36 d | Integrated 2 stage treatment. | [54] |
| 18. | Activated sludge treatment | TSS (mg/L): 92.24% | CH4 Release (%): 2 | Anaerobic treatment | [69] |
| 19. | Raw POME | COD (mg/L): 71.00% | CH4 Release (%): 2 | Aerobic BioAX treatment, AnaerobicEG bioreactors (Pilot scale bioreactor) | [70] |
| 20. | Thermophilic anaerobic-aerobic treated POME | BOD (mg/L): 88.00% | HRT (days): 11 months | Pectin and chitosan in Jar Test protocol. | [71] |
| 21. | POME + dairy manure | Colour (mg/L): 99.00% | CH4 Release (%): 1 | Continuous stirred tank reactor (CSTR). | [72] |
| 22. | 13 phyla of bacteria (Proteobacteria, Chloroflexi, Firmicutes, Bacteroidetes, and Nitrospirae) | COD (mg/L): 80.76% | CH4 Release (%): 57 | Moving bed biofilm reactor (MBBR). | [73] |
| 23. | Typha latifolia | BOD (mg/L): 88.00% | CH4 Release (%): 59 | Phytoremediation using single plant. | [44] |
| 24. | Vetiveria zizanioides | TSS (mg/L): 90.00% | CH4 Release (%): 45 | Phytoremediation using single plant. | [15] |
Table 2. Cont.

| No. | Microbes or Biological Seeding                                                                 | Concentration, mg/L or Removal Efficiency, % | Process Aspects                                                                 | Ref. |
|-----|------------------------------------------------------------------------------------------------|---------------------------------------------|---------------------------------------------------------------------------------|------|
| 25. | Water spinach *(Ipomoea aquatic Forsk)*                                                        | COD (mg/L) = 86.30% BOD (mg/L) = - TSS (mg/L) = - Colour (mg/L) = 95.30% Oil (mg/L) = - CO₂ Fixation (g CO₂/L/day) = - CH₄ Release (%) = - Working Volume (L) = 10 HRT (days or h) = 15 d | Phytoremediation using single plant.                                              | [45] |
| 26. | *Spirogyra* sp.                                                                                  | COD (mg/L) = 63.60% BOD (mg/L) = - TSS (mg/L) = - Colour (mg/L) = - Oil (mg/L) = - CO₂ Fixation (g CO₂/L/day) = - CH₄ Release (%) = - Working Volume (L) = 0.75 HRT (days or h) = 20 d | Phytoremediation using single plant.                                              | [46] |
| 27. | *Pistia stratoites*                                                                               | COD (mg/L) = 30.00% BOD (mg/L) = 93.00% TSS (mg/L) = - Colour (mg/L) = - Oil (mg/L) = - CO₂ Fixation (g CO₂/L/day) = - CH₄ Release (%) = - Working Volume (L) = 6 HRT (days or h) = 15 d | Phytoremediation using single plant.                                              | [51] |
| 28. | *Eichorniacrassipes*                                                                             | COD (mg/L) = 50.00% BOD (mg/L) = 50.00% TSS (mg/L) = - Colour (mg/L) = - Oil (mg/L) = - CO₂ Fixation (g CO₂/L/day) = - CH₄ Release (%) = - Working Volume (L) = 5 HRT (days or h) = 30 d | Phytoremediation using single plant.                                              | [52] |
| 29. | *Bacillus cereus* MF661883                                                                       | COD (mg/L) = 79.35% BOD (mg/L) = 72.65% TSS (mg/L) = - Colour (mg/L) = - Oil (mg/L) = - CO₂ Fixation (g CO₂/L/day) = - CH₄ Release (%) = - Working Volume (L) = 0.2 HRT (days or h) = 6 d | Batch fermentation mode, lipid accumulation and bioremediation.                  | [16] |
| 30. | *Aspergillus niger* ATCC 9642                                                                    | COD (mg/L) = 63.00% BOD (mg/L) = - TSS (mg/L) = - Colour (mg/L) = - Oil (mg/L) = - CO₂ Fixation (g CO₂/L/day) = - CH₄ Release (%) = - Working Volume (L) = 1 HRT (days or h) = 7 d | Fermentation of POME into citric acid.                                           | [56] |
| 31. | Raw POME and anaerobic sludge                                                                    | COD (mg/L) = 70.00% BOD (mg/L) = - TSS (mg/L) = - Colour (mg/L) = - Oil (mg/L) = - CO₂ Fixation (g CO₂/L/day) = - CH₄ Release (%) = - Working Volume (L) = 0.45 HRT (days or h) = 15 d | Microbial fuel cell-dual chamber                                                   | [59] |
| 32. | Anaerobic digestion and activated sludge from Gaopeidian sewage treatment plant                   | COD (mg/L) = 96.50% BOD (mg/L) = - TSS (mg/L) = - Colour (mg/L) = - Oil (mg/L) = - CO₂ Fixation (g CO₂/L/day) = - CH₄ Release (%) = - Working Volume (L) = 2.36 HRT (days or h) = 10 d | Integrated two-staged MFCs coupled with the immobilised biological aerated filter | [60] |
4.3. Physicochemical Treatment

Although there are various types of biological treatment, certain processes or systems may not deliver the required results with respect to the minimum requirements set by the authority. Physicochemical treatment processes or also known as polishing techniques are viable means to treat POME. Biomass and other waste materials such as fibres and fruits wastes may offer an inexpensive and renewable source for POME treatment. These waste materials have low or no economic value and often present a disposal problem. Therefore, there is a need to valorise these low-cost by-products. Thus, their conversion into sorbent for removing or/and reducing oil, BOD, COD, and Total Suspended Solid (TSS) from POME has an economic value to reduce the cost of waste disposal. Most importantly, these provide a potentially inexpensive alternative treatment technique. A wide variety of sorption materials was known for treatment of biomass and other wastes by using esterified sago bark fibre [17], activated bioadsorbent from oil palm kernel shell [74–76], banana peel [5,77], empty fruit bunch (EFB) [78,79], coconut shell [80,81], oil palm mesocarp fibre [82], rice husk [83] and cocoa peels [84]. Currently, natural fibres are considered as highly potential sources to be utilized as natural sorbents for removing oil from POME with excellent adsorption properties, as well as being chemical-free and highly biodegradable [85].

Table 3 shows the summary of various bioadsorbent materials prepared from biomass, which exhibit high sorption properties. Mohammed and Chong [5] reported that the highest COD (100%), BOD (97%), TSS (100%), and colour removal (96%) were obtained after 30 h treatment by banana peel adsorbent dosage of 30 g/100 mL. A well-known biosorbent material such as industrial-grade alum at 10 g/L dosage was reported in reducing the COD (98%) and TSS (99%) value in 25% v/v POME [78]. An empty fruit bunch-based activated carbon with a dosage of 20 g/L was utilized to treat POME, significantly reducing the COD, BOD, and TSS at 98%, 88%, and 96% within only 30 min [79]. Mohammed [74] successfully removed the POME colour (100%) within 12 h by using oil palm kernel shell (PKS), followed by chemical activation (potassium hydroxide) and microwave treatment. The high feasibility in adsorption method especially in oil removal was also reported using esterified sago bark fiber and sago bark fibre [17]. Those performances were evaluated using adsorption tests particularly for 24 h, resulting in 80.23% and 57.77%, respectively. However, no analysis of COD and BOD was done in this work. Based on the compiled literature, we conclude that until now, there is no further investigations on the POME adsorption scalability.

Alternative physicochemical treatments such as membrane filtration [4,18,86], coagulation [87], coagulation-flocculation [88–90], electrocoagulation [91–93], and fenton-oxidation [94–96] were only capable of migrating the organic pollutants from one form into another. However, those still could not mineralise the organic pollutants [97]. These methods are considered to be costly due to the chemical usage and involved excess-sludge treatment [98]. Worldwide, membrane techniques have been extensively used in POME polishing treatments prior to reducing the concentration of COD, BOD, TSS, colour, and any organic elements. Membrane usage such as nanofiltration, ultrafiltration, and reverse osmosis were capable to increase the removal efficiencies of pollutants [4,18,86]. Ahmad et al. [4] utilized membrane filtration with success to remove more than 99.40% of COD, 98.80% of BOD, with 100% of TSS removal within 4 h of treatment for 200 mL POME. Azmi and Yunus [18] adopted micro filtration membrane (MF) method applying for POME with COD and BOD reduction in 180 mL POME at 90% within 1.5 days. Amat et al. [86] successfully decolourised POME at 97.40–97.90% removal with MF system throughout 4 h for a volume of 15 mL of POME. Several researchers evaluated the removal of pollutants from POME final discharge by using membrane bioreactor (MBR), as classified as a hybrid system of two interdependent processes of biological and membrane filtration [99]. An integrated treatment process for POME final discharge was based on MBR adopted by Ahmed and co-workers [20] for obtaining high quality effluent. However, the major problem when using the membrane is generally caused by pore blocking at the membrane surface, which simultaneously affects the capital cost and maintenance [100].
Coagulation-flocculation process is acknowledged to have a superior performance technology in filtering the suspended solids [89]. More than 90% of colour removal was achieved within 23 min of POME decolourisation using dual-coagulants, ferric chloride-anionic polyacrylamide [30]. The coagulant is basically composed of chemical compounds, as they are easy to handle, cheap, and portray good removal efficiency towards the wastewater application [101]. Another study utilised mango pit as a coagulant and fly ash as an adsorbent in the coagulation-flocculation process [88] and recorded COD and TSS removal at 89% and 96% within 1 h. Through the coagulation process or other tertiary treatment processes, the capability of adsorption process in removing the pollutants is potentially high. Therefore, the combination with other treatment techniques would enhance the performance of POME treatment. However, after treatment of the flocculation process, huge sludge remained as an additional problem.

Electrochemical processes including electrolysis and electrocoagulation techniques have been effectively employed for removing pollutants in various industrial wastewaters [102]. The latter process is claimed to be an effective wastewater treatment technique, as it involves a series of treatments: coagulation, precipitation, and flotation. Electrocoagulation uses aluminium or iron anodes to produce aluminium or iron hydroxide flocs by the hydrolysis reaction of the anodes. The ability in treating POME was studied on the decolourisation through the polishing step [91,92], resulting in almost 100% colour removal being recorded in POME decolourisation utilizing aluminium electrode within 65 min operation. Che Sayuti and Mohd Azoddein [91] successfully reduced the COD from 4900 mg/L to 210 mg/L (95.71%) and TSS from 4000 mg/L to 30 mg/L (99.25%), respectively. The final COD value almost met the DOE requirement of 200 mg/L, whereas TSS was 100 mg/L, completely fitting the requirement. The highest efficiency was obtained at optimum pH 7.44 with electrocoagulation time of 25 min and voltage of 100 V utilising aluminium electrodes. Interestingly, this method was found to be efficient and capable in reducing treatment time compared to the ponding system.

Electro-Fenton system demonstrated better performance to COD removal than the electrocoagulation system, despite both being capable in reducing most of the organic compounds [101]. Fenton-oxidation has become another favourable treatment system in the past decades [103]. For example, a total of 85% COD reduction was achieved under the acidic POME at a pH close to 3. For the decolourisation and organic decomposition [104], Fenton-oxidation process was useful, resulting in a higher COD removal (82%), with more than 90% colour removal in the case of 50 mM H$_2$O$_2$ with Fe$^{2+}$ for the 30 min oxidation process.

In other studies, ultrasonic cavitation technology, coupled with the use of hydrogen peroxide, H$_2$O$_2$, was used as an alternative technique in treating POME [104]. Cavitation is defined as the phenomenon of formation, growth, and collapse of microbubbles in a specific liquid. There are several types of cavitation, described as acoustic, hydrodynamic, optic, and particle. The application of ultrasonic cavitation was widely employed for degradation of textile dyes [105], degradation of chemical contaminants [106], and industrial wastewater treatment [107]. Manickam et al. [108] successfully investigated the effect of hydrogen peroxide in reducing the COD patterns by using the combination of ultrasound/H$_2$O$_2$ (30% w/v), yielding better results compared to ultrasound/H$_2$O$_2$ (6% w/v). An impressive result was obtained at final COD of 27,000 mg/L (almost 50% reduction) for 6 h. Thus, the final COD obtained portrayed the suitability method of chemical/ultrasound combination as a pre-treatment step for POME, as followed by another digestion method. In addition to COD reduction, POME de-colouring was possible from brown to light yellow with this treatment. Thus, it was believed that this technique could be one of the green process engineering techniques for POME treatment [108].

Photocatalytic reaction demonstrates high efficiency in the degradation of organic compounds [109]. TiO$_2$ are the most widely utilized owing to its low cost, low toxicity, high performance, and high chemical stability. The heterogenous photocatalytic system used TiO$_2$ nanoparticles as the photocatalyst, and simultaneously performed better in
degrading the organic pollutants, especially in POME \cite{110,111}. The COD reduction (78\%) was obtained from the initial value of 168 mg/L in POME settling pond \cite{111}. Organic compounds in POME were successfully decomposed within 7 h or reaction by using 0.83 g/L of 20\% wt. Cu/TiO$_2$ loading \cite{112}. Despite their advantages, the photocatalytic system also portrayed several drawbacks such as inefficient catalyst separation and low penetration of light incident during the reaction. Thus, some modifications including cylindrical column photoreactor \cite{109}, double-cylindrical shell photoreactor \cite{113}, and ZSM-5 zeolite support with Fe$^{3+}$ and Ni$^{2+}$ doping \cite{114} could overcome their problems. By introducing steam-high temperature, Cheng et al. \cite{115} and Ng et al. \cite{116} successfully generated novel approaches for catalytic POME steam, reforming into a valuable syngas. In the former, the highest COD and BOD removals were achieved at 93.8\% and 93.7\%, respectively, at 873 K and 51.44\% of H$_2$ selectivity under 90 mL/h POME flow in 20 mL/min of N$_2$ as the carrier gas. In the latter, 99.7\% of COD was reduced over the same catalyst at 1173 K within 1 h of operation. It was apparent that high temperature and partial pressure of POME favoured higher H$_2$ production. Despite the potential of H$_2$ gas production, the energy consumption and net energy in transforming the POME into syngas should be furthermore verified to identify the scale-up feasibility prospect.

4.4. Thermochemical Treatment

Thermochemical treatment was one of the prominent ways to transform the wet biomass into valuable products, as also known by ‘synthesis gas’ with abbreviation of syngas. By using the minimization of total Gibbs free energy method, the thermodynamic simulation is performed to study the effect of reaction temperature on product yield and syngas ratio (H$_2$:CO) \cite{117}. The result evinced the steam reforming all POME organic compounds, which is feasible at 773 K and beyond this point. Bioconversion of the highest H$_2$-rich syngas occurred at 1073 K within the study temperature ranging from 573–1173 K \cite{117}.

The supercritical water gasification (SCWG) was conducted at 653 K with 250 bar prior to obtaining syngas from the dual feed of POME and empty fruit bunch (EFB) \cite{118}. The POME served as the reaction medium, whereas the EFB acted as the main substrate. The pure POME was gasified into the valuable gaseous products yielding about 13 mmol/mL of CO$_2$, 20 mmol/mL of H$_2$, 0.3 mmol/min of CH$_4$, and 0.6 mmol/mL CO, respectively. Evidently POME had high potential substrate in producing valuable and beneficial gas products under SCWG condition.

In spite of the substantially enhanced H$_2$ gas production, the carbon deposition on the catalyst surface may deter the overall process. In addition, in order to efficiently perform this process at a large-scale, the reduction of overall energy consumption to produce syngas should be addressed.
Table 3. Summary of various physicochemical treatments at different dosages with removal efficiency of final discharge POME.

| No. | Bioadsorbent Material | Bioadsorbent Dosage (g/L) | COD (mg/L) | BOD (mg/L) | TSS (mg/L) | Colour (mg/L) | Oil (mg/L) | CH4 Release (%) | Working Volume (L) | HRT (day/h/min) | Treatment | Ref. |
|-----|-----------------------|---------------------------|------------|------------|------------|--------------|------------|----------------|------------------|----------------|-----------|-----|
| 1.  | Standard discharge limit | -                         | 100        | 100        | 400        | 200          | 50         | -              | -                | -              | DOE, 1982  |     |
| 2.  | Future discharge limit  | -                         | -          | -          | -          | -            | -          | -              | -                | -              | DOE, 2015  |     |
| 3.  | Esterified sago bark fibre | 5                         | -          | -          | -          | -            | 80.23%     | -              | 0.1              | 24 h          | Ethyl acetate and calcium oxide. | [17] |
| 4.  | Oil palm kernel shell   | 5 g/100 mL                | -          | -          | -          | 100%         | -          | -              | 0.1              | 14 h          | Potassium hydroxide activation, and microwave. | [74] |
| 5.  | Banana peel             | 30 g/100 mL               | 100%       | 97.00%     | 100%       | 96.00%       | -          | 0.1            | 30 h             | Activated carbon (AC). | [5] |
| 6.  | Industrial grade alum   | 10                        | 98.00%     | -          | 99.00%     | -            | -          | 1.0            | -                | Adsorption-coagulation | [78] |
| 7.  | Empty fruit bunch-based activated carbon | 20                        | 98.00%     | 88.00%     | 96.00%     | -            | -          | 0.1            | 30 min           | Pyrolysis, followed by physical activation. | [79] |
| 8.  | Oil palm kernel shell   | 10                        | 89.00%     | 85.00%     | 80.00%     | 88.00%       | -          | -              | -                | Sodium-potassium hydroxide mixture (NaOH-KOH) activation. | [75] |
| 9.  | Oil palm kernel shell   | -                         | 80.00%     | -          | -          | -            | -          | 0.25           | 24 h             | Steaming | [76] |
| 10. | Coconut shell           | 50                        | 70.00%     | -          | 70.00%     | 70.00%       | -          | -              | -                | Commercial activated carbon | [80] |
| 11. | Oil palm mesocarp fibre | 10                        | 70.00%     | -          | 88.00%     | -            | -          | -              | -                | Consecutive treatments. | [82] |
| 12. | Coconut shell           | 5                         | 61.00%     | -          | 39.00%     | 61.00%       | -          | 0.1            | 48 h             | Acid-washed activated carbon (Kinetic models) | [81] |
| 13. | Fruit peel (banana and orange peels) | 10                        | 52.00%     | 46.00%     | 18.00%     | 57.00%       | -          | -              | -                | 3 days Biochar from pyrolysis of fruit peels. | [77] |
| No. | Bioadsorbent Material                      | Bioadsorbent Dosage (g/L) | COD (mg/L) | BOD (mg/L) | TSS (mg/L) | Colour (mg/L) | Oil (mg/L) | CH₄ Release (%) | Working Volume (L) | HRT (day/h/min) | Treatment                                                                 | Ref. |
|-----|------------------------------------------|---------------------------|------------|------------|------------|---------------|-------------|----------------|-------------------|-----------------|---------------------------------------------------------------------------|------|
| 14  | Palm oil mill effluent (POME)            | -                         | 99.40%     | 98.80%     | 100%       | -             | -           | -              | 500               | 4 h             | Integrated Membrane ultrafiltration + Reverse osmosis                   | [4]  |
| 15  | Palm oil mill effluent (POME)            | -                         | 90.00%     | 90.00%     | -          | -             | -           | -              | 0.18              | 1.5 days        | Membrane ultrafiltration (MF)                                           | [18] |
| 16  | Palm oil mill effluent (POME)            | -                         | -          | -          | 97.90%     | -             | -           | -              | 0.15              | 4 h             | Membrane ultrafiltration (MF) + Nanofiltration (NF)                     | [86] |
| 17  | Rice husk                                | 50                        | 41.00%     | -          | -          | 88.00%        | -           | -              | 10                | 18 h            | Microwave incineration.                                                   | [83] |
| 18  | Cocoa peels                              | 1.5                       | -          | -          | 83.45%     | 80.88%        | -           | -              | 0.05              | 5 h             | Activated carbon with nitric acid (HNO₃)                                 | [84] |
| 19  | Modified cassava peel starch and alum    | 7.5–50                    | -          | -          | 92.75%     | -             | -           | -              | 30 min            | (2 min          | Dual coagulation process                                                  | [87] |
| 20  | Alum                                     | 2.12                      | 59.00%     | -          | -          | -             | -           | -              | 1                 | 30 min          | Coagulation-flocculation                                                 | [88] |
| 21  | Mango pit and fly ash                    | 50                        | 89.00%     | -          | -          | 96.00%        | -           | -              | 0.3               | 1 h             | Coagulation-flocculation                                                 | [89] |
| 22  | Polyacrylamide (C-PAM, A-PAM)            | 300                       | 70.90%     | -          | 96.40%     | -             | -           | -              | 1 min             | Coagulation-flocculation                                                 | [90] |
| 23  | Aluminium Electrode                       | -                         | 95.71%     | -          | 99.25%     | -             | -           | -              | -                 | 25 min          | Electrocoagulation                                                      | [91] |
| 24  | Aluminium Electrode                       | -                         | 75.40%     | -          | -          | 100%          | -           | -              | 0.5               | 65 min          | Electrocoagulation                                                      | [92] |
| 25  | Polyacrylamide                           | 1–3                       | 70.00%     | -          | -          | -             | -           | -              | 3                 | 6 h             | Electrocoagulation                                                      | [93] |
| 26  | Calcium lactate-Polyacrylamide           | 0.5                       | 93.90%     | 88.70%     | -          | >90.00%       | -           | -              | 0.5               | 23 min          | Coagulation-flocculation                                                 | [30] |
| 27  | H₂O₂ and Fe²⁺                             | 1.33–5.67, 1.12–4.48      | 85.00%     | -          | -          | -             | -           | -              | 0.25              | 90 min          | Electron-fenton oxidation                                               | [94] |
Table 3. Cont.

| No. | Bioadsorbent Material | Bioadsorbent Dosage (g/L) | COD (mg/L) | BOD (mg/L) | TSS (mg/L) | Colour (mg/L) | Oil (mg/L) | CH4 Release (%) | Working Volume (L) | HRT (day/h/min) | Process Aspects | Treatment | Ref. |
|-----|-----------------------|---------------------------|------------|------------|------------|--------------|------------|----------------|------------------|-----------------|---------------|-----------|------|
| 28. | H2O2 and Fe2+         | 5.0 mM, 1.0 mM           | 82.00%     | -          | -          | -            | -          | -              | 0.25             | 30 min          | Electron-fenton oxidation | [104]     |
| 29. | H2O2                  | 10                        | 50.00%     | -          | -          | -            | -          | -              | 0.05             | 6 h             | Ultrasonic cavitation coupled with H2O2 | [108]     |
| 30. | TiO2                  | 0.5                       | 78.00%     | -          | -          | -            | -          | -              | 0.01             | 4 h             | UV-TiO2 photocatalytic reaction | [111]     |
| 31. | Cu/TiO2               | 0.83                      | 40.00%     | -          | -          | -            | -          | -              | 0.3              | 7 h             | Cu-TiO2 photocatalytic reaction | [112]     |
| 32. | Ni/Al2O3              | 0.1                       | 93.80%     | 93.70%     | -          | -            | -          | -              | 0.6–3.3          | 40              | Photocatalytic by steam (873 K) | [116]     |
| 33. | Ni/Al2O3              | 0.1                       | 99.70%     | -          | -          | -            | -          | -              | 0.28             | -               | Photocatalytic by steam (1173 K) | [117]     |
| 34. | Palm oil mill effluent| 50                        | 73.08%     | -          | 98.33%     | -            | -          | -              | -                | 6 h             | Activated carbon (AC) and ultrasound (US). | [119]     |
| 35. | Cultures of Aspergillus fumigatus | 10%       | 71.00%     | -          | 71.00%     | -            | -          | 0.01           | 5 days           |                | Different carbon sources—centrifugation. | [120]     |
| 36. | Palm oil mill effluent (POME) | 5–20           | 46.00%     | -          | 61.00%     | 39.00%       | -          | 0.1            | 6 h              |                | Combined magnetic field and commercial activated carbon (adsorption) | [121]     |
4.5. Torrefaction or Hydrothermal Treatment

Conversion of solid waste to high-density briquettes via torrefaction or hydrothermal as a pre-treatment process is touted as the ideal solution to avert accumulation of solid waste, especially in the agro-industry sector. Various agro-industry solid wastes such as woodchips, olive pulp, and olive stones are pre-treated to produce biomass fuel with a heating value (HV) ranging from 16.7–20.3 MJ/kg [122,123]. In lieu of the aforementioned pre-treatment process, the mixture of POME and oil palm trunk (OPT) is simply subjected to being oven dried at 105 °C until constant weight is observed and followed by forming into briquettes [124]. POME and OPT with the ratio 90:10 bode the highest HV of 30.85 MJ/kg [124]. The high HV value obtained by utilizing POME corroborates its potential applications for biomass fuel, as compared to other agro-wastes.

5. Life Cycle Analysis of POME

Life cycle analysis (LCA) is a common method used for assessing environmental impacts associated with new technology designs or processes, herein specifically focused on various POME treatments. The severity of environmental impact upon utilizing the designated POME treatment processes can be quantitatively evaluated by monitoring the parameters such as level of BOD, COD, and CO₂ emissions. Among all mooted POME treatment processes listed in Tables 2 and 3, only the biological treatment utilizing an integrated anaerobic-aerobic bioreactor in the presence of mesophilic-thermophilic bacterial and physicochemical treatment, using banana peel as a bioadsorbent material, can be categorized as the most efficient process. The utilization of agro-waste such as banana peel and EFB as bioadsorbent are deemed to be cost-efficient. The aforementioned waste material has no economic value and is often present as a voluminous solid waste. Thus, valorization of these by-products will aid in reducing the waste disposal costs and creating other value-added products such as bioadsorbents.

6. Conclusions

Overall, among the aforementioned POME treatments, phytoremediation, MFC techniques, and agro-waste-derived bioadsorbents seemed to be more cost-effective, but still were unable to produce satisfactory results in lowering the COD and BOD values. Numerous physicochemical treatment methods of biological processes, bioadsorbents, membranes, and coagulants were employed for treating POME, but no single process can be utilized as stand-alone at the commercial scale, yet. The usage of these adsorbents and chemical coagulants seems uneconomical and impractical. The thermochemical treatment for POME including the integrated system of anaerobic-aerobic bioreactor (IAAB) demonstrated the highest COD removal efficiency and is touted as the ideal treatment system. The judicious utilization of various affordable biomaterials, short treatment time, and scalable and clean technologies will be expected to pave the way towards an efficient POME treatment. These initiatives are in line with United Nations Sustainable Development Goals 7: Affordable and clean energy. Finally, we wish to comment on the sustainable aspect. Most treatment technologies should deem to possess environment-friendly properties to some extent, either partially or, preferably, through elimination of chemical solvent usage. Their many advantageous characteristics of biological treatment over harmful chemical treatment utilization justify their use as an eco-friendly treatment process.

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Abbreviations

| Abbreviation | Description |
|--------------|-------------|
| POME         | Palm oil mill effluent |
| IAA          | Integrated anaerobic-aerobic bioreactors |
| BOD          | Biochemical oxygen demand |
| COD          | Chemical oxygen demand |
| TSS          | Total suspended solids |

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