

A review of methods for removal of contaminants in used lubricating oil

Abstract

Management and disposing of used lubricating oil (ULO) poses deleterious effects to air, land and water pollution. These contaminants not only causes environmental problems, they also have bio-accumulation effects on living organism, reduces the lifespan of inhabitants as a result of the spread of diseases, poisoning and fouling of catalyst as well as corrode processing equipment. Removal of contaminants in used lubricating oil is a major step to avoid pollution as discussed thoroughly by many researchers in literature. In addition to curbing pollution, another advantage is converting waste to wealth. This review paper presents insight into various methods for removal of contaminants in used lubricating oil. The advantages and drawbacks of each method were earmarked for further study.

Keyword: *Used lubricating oil, Contaminants. Removal Methods, Treatment*

Introduction

Lubricating oils (LOs) are conventionally obtained from crude oil. Chemical composition of LOs consists on average of about 80–90% base oil and about 10–20% chemical additives and other compounds (Rincón et al., 2005). Lubricating oils mainly helps in reducing friction, dust, corrosion, protection against wear and tear and provision of heat transfer medium in various equipment or machineries. (Shri et al., 2014). During operation time, LOs deteriorate, as well as their additives, and its physical and chemical properties become unsuitable for further use (Tsai, 2011). In the process of usage of lube oil, temperature build up occurs which breaks down the oil and weakens its properties which include pour point, flash point, specific gravity, viscosity etc. (Udonne and Bakare, 2013). These renders the oil unsuitable for regular usage as results of contaminants in the lube oil such as water, wear metals, carbon residue, ash content, gums, varnishes etc. Chemical changes in the oil occurred due to thermal degradation and oxidation. Europe represents 19% of total worldwide market volume of lubricants, consuming around 6.8 million tons in 2015 (Kupareva et al., 2013).

Used lubricating oils (ULOs) are classified as hazardous wastes, and constitute a serious pollution problem not only for the environment, but also for human health due to the presence of harmful contaminants, such as heavy metals, polychlorinated biphenyls (PCBs) and polycyclic aromatic hydrocarbons (PAHs) (Kanokkantapong et al., 2009). Poor management and careless disposal of used lube oil can affect the environment negatively (Lam and Chase, 2012 and Lam et al., 2016). Scientists have reported that in some geographical region e.g. West Africa, the dispersion of the air pollutants could travel at a speed of 10-12 m/s (Emetere, 2017). The implication of this report is that air pollution from burning of waste lubricant is not localized to the source of pollution but could travel with time to other locations. For example, it was recently reported that black soot covered a metropolitan city of Port Harcourt while remote sources were at the suburb settlement (about 22 km away from the city) (Temitayo et al., 2018).

42 On the other hand, ULOs can be considered as valuable resources, in the sense that it is
43 possible to recover energy or profitable materials for further use (Guerin, 2008). The best
44 environmental options, for the management of used lube oils follow the 'waste hierarchy' by
45 recycling, recovering and then disposing. Used lube oils can be used as an alternative fuel in
46 a variety of engine configurations and other applications. Its gross calorific value is greater
47 than 42.9 MJ/kg (Ketlogetswe, C., 1998). The principal objective of any waste management
48 plan is to ensure safe, efficient and economical collection, transportation, treatment and
49 disposal of waste and as well as satisfactory operation for current and foreseeable future
50 scenarios (Stoll and Gupta, 1997). The treatment of used lube oil is important due to; it
51 requires less energy and cost compared to conventional refining of crude oil, it helps in
52 improving air quality, land and water pollution in the environment. The most preferred option
53 by experts is the reuse of the used lube oil generated by consumers (Jafari and Hassanpour,
54 2015). In this paper, a thorough review on various removal and treatment methods for used
55 lube oil would be considered starting from conventional to the most current methods and their
56 limitations; further developments of these fields were also touched. In addition,
57 environmentally friendly and affordable solvent extraction and adsorbents would be
58 developed as a means of removing contaminants in used lube oil.

59 **2. Conventional Methods**

60 The conventional methods of removal of contaminants in used lube oil either requires a high
61 cost technology such as vacuum distillation or the use of toxic materials such as sulphuric
62 acid. These methods also produce contaminating by-products which have highly sulphur
63 levels, especially in the Kurdistan region/Iraq (Hamawand et al., 2013).

64 **2.1. Acid-clay**

65 Assessment of different removal of contaminants processes in used lube oils revealed that
66 acid-clay process had the highest environmental risk and lowest cost. The method involve
67 treatment of used oil with acid and clay (Udonne and Bakare, 2013; Hamawand et al., 2013;
68 Abu-Elella et al., 2015). They all used the clay as an adsorbent to remove the odour and dark
69 colour. What makes acid-clay method unique from others are; its simple method, affordable
70 capital investment, low operating cost and does not need skilled operators (Giovanna et al.,
71 2012; Nwachukwu et al., 2012 and Isah, A.G., 2013).

72 However, this method has many disadvantages; it also produces large quantity of pollutants,
73 is unable to treat modern multigrade oils and it's difficult to remove asphaltic impurities
74 (Fox, M.F., 2007). To reduce these hazardous contaminants from this method; the acid
75 treatment stage of the process can be done under the atmospheric pressure to remove the
76 acidic products, oxidized polar compounds, suspended particles and additives (Falah and
77 Hussein, 2011). Princewill and Sunday, (2010) observed that high recovery rate of treated
78 lube oil from used lube oil depend largely on the source of the used lube oil, pre-treatment
79 mechanisms, extent of contamination and the grade of the acid used. He also showed that the
80 volume of the adsorbent (clay) used could affect the rate at which contaminants are removed
81 and the percentage of recovery of the method.

82 In Abu-Elella et al., (2015) worked on used motor oil. He treated used motor oil with
83 phosphoric acid, sulphuric acid, methanoic acid and acetic acid. He observed that methanoic
84 acid, sulphuric acid and acetic acid have great changes on the kinematic viscosity while

85 phosphoric acid is not affected by used lube oil. He therefore concluded that treatment with
86 acetic acid showed better results than formic acid-clay.

87 **2.2 Solvent Extraction**

88 This method has replaced acid-clay treatment as the preferred method for improving the
89 oxidative stability and viscosity as well as temperature characteristics of the base oils. Base
90 oils obtained from Solvent Extraction are of good quality and contains less amounts of
91 contaminants. In contrast to acid-clay treatment, it operates at higher pressures, requires
92 skilled operating system and qualified personnel. (AERCO 1995). The solvent selectively
93 dissolves the undesired aromatic components (the extract), leaving the desired saturated
94 components, especially alkanes, as a separate phase (the raffinate). (Rincon et al., 2005).

95 Different types of solvents have been used for solvent extraction such as 2-propanol, 1-
96 butanol, methyl ethyl ketone (MEK), ethanol, toluene, acetone, propane etc. (Quang et al.,
97 1974) and (Rincon et al., 2003) used propane as a solvent. He found out that the propane was
98 capable of dissolving paraffinic or waxy material and intermediately dissolved oxygenated
99 material. Asphaltenes which contain heavy condensed aromatic compounds and particulate
100 matter are insoluble in liquid propane. These properties make propane ideal for recycling the
101 used engine oil, but there are many other issues that have to be considered. Propane is
102 hazardous and flammable therefore this process is regarded as hazardous method.

103 Katiyar and Husain (2010); Sterpu et al., (2012) and Hassan et al.,(2012) found out that
104 methyl ethyl ketone has the highest performance due to its low oil percentage losses and high
105 sludge removal while Hussein et al., (2014) and Aremu et al., (2015) found out extraction
106 using butan-1-ol solvent produces the highest sludge removal rate. (Rincon et al., 2005) and
107 Oladimeji et al., 2018) used a composite solvent of methyl ethyl ketone and 2-propanol the
108 oil resulting from this process is comparable to that produced by acid-clay method, its cost
109 was high.

110 Solvent extraction in general involves solvent losses and highly operating maintenance.
111 Also, it occurs at pressures higher than 10 atm and requires high pressure sealing systems
112 which makes solvent extraction plants expensive to construct, operate and the method also
113 produces remarkable amounts of hazardous by-products. (Quang et al., 1974); (Rincon et al.,
114 2003) and Hamawand et al., (2013).

115 Mineral Oil Raffinerie Dollbergen (MRD) solvent extraction process using N-methyl-2-
116 pyrrolidone. The applied oil re-refining process is based on a patent held by AVISTA OIL.
117 (Pöhler et al., 2004) The 'Enhanced Selective Refining' process uses solvent N-methyl-2-
118 pyrrolidone (NMP), which is commonly used in the petroleum refining industry. NMP is a
119 powerful, aprotic solvent with low volatility, which shows selective affinity for unsaturated
120 hydrocarbons, aromatics, and sulphur compounds. Due to its relative non-reactivity and high
121 selectivity, NMP finds wide applicability as an aromatic extraction solvent in lube oil re-
122 refining. The advantages of NMP over other solvents are the non-toxic nature and high
123 solvent power, absence of azeotropes formation with hydrocarbons, the ease of recovery from
124 solutes and its high selectivity for aromatic hydrocarbons. Being a selective solvent for
125 aromatic hydrocarbons and PAH, NMP can be used for the re-refining of waste oils with
126 lower sludge, carbonaceous particles and polymer contents, such as waste insulating,

127 hydraulic and other similar industrial oils.(Lukic, J et al., 2005). The MRD solvent extraction
128 process uses the liquid–liquid extraction principle.

129 The average base oil yield within the process is about 91 %.(Schiessler, N et al., 2007). The
130 base oils produced have high quality (Kupareva et al., 2013). The process is characterized by
131 optimized operating conditions which allow elimination of toxic polyaromatic compounds
132 from the re-refined base oil and preservation of the synthetic base oils like polyalphaolefin
133 (PAO) or hydrocracked oils, which are increasingly present in used oils. However, this
134 method need skilled personnel, proper disposal and management of it waste.

135 **2.3 Vacuum distillation**

136 Extensive research work have been done on vacuum distillation on used oil by the following
137 Martins, J.P, (1997); Shakirullah et al.,2006; Bridjanian and Sattarian (2006); Emam and
138 Shoaib, (2012); Hamawand et al. (2013) and Kannan et al., (2014). In this method, used lube
139 oil collected is heated at a temperature of 120°C to remove the water added to the oil during
140 combustion. Then the dehydrated oil is subjected to vacuum distilled at a temperature of
141 240°C and pressure 20 mmHg. This results to the production of light fuel oil at a temperature
142 of 140°C (the light fuel oil can be used as fuel source for heating) and lubricating oil at
143 240°C. The lubricating oil vapour is condensed and sent for next stage. (Kannan et al., 2014).
144 The advantages of vacuum distillation process over atmospheric pressure distillation are:
145 columns can be operated at lower temperatures; more economical to separate high boiling
146 point components under vacuum distillation; avoid degradation of properties of some species
147 at high temperatures therefore thermally sensitive substances can be processed easily.

148 However, the remaining oil generated at this temperature (240°C) contains the dirt, degraded
149 additives, metal wear parts and combustion products like carbon and is collected as residue.
150 The residue is in the form similar to that of tar, which can be used as a construction material
151 for example road and bitumen production. (Giovanna et al., 2003). The disadvantage of this
152 method is the high investment cost and/or the use of toxic materials such as sulphuric acid.
153 (Havemann, 1978 and Puerto-Ferre, & Kajdas, 1994).

154 **2.4 Hydrogenation**

155 To avoid formation of harmful products and environmental issues based on above methods,
156 some modern processes have been used and the best one is hydrotreating. (Bridjanian and
157 Sattarin, 2006).This method follows vacuum distillation. In this process, the distillate from
158 vacuum distillation is hydrotreated at high pressure and temperature in the presence of
159 catalyst for the purpose of removing chlorine, sulphur, nitrogen and organic components. The
160 treated hydrocarbons resulted in products of improved odour, chemical properties and colour.
161 (Temitayo et al., 2018).

162 Another important aspect of this method is that, this process has many advantages: Produces
163 of high Viscosity Index lube oil with well oxidation resistance and a good stable colour and
164 yet having low or no discards. At the same time, it consumes bad quality feed. In addition to
165 that, this method has advantage that all of its hydrocarbon products have good applications
166 and product recovery is high with no (or very low) disposals. Other hydrocarbon products
167 are: In oil refinery the light –cuts can be used as fuel in the plant itself. Gas oil may be
168 consumed after being mixed with heating gas oil and the distillation residue can be blended

169 with bitumen and consumed as the paving asphalt, because it upgrades a lot its rheological
170 properties. Also, it can be used as a concentrated anti-corrosion liquid coating, for vehicles
171 frames. (Hassan A. D, 2014).

172 The disadvantage of this method is that the residue resulting from the process is of high
173 boiling range of hydrocarbon product fractionated into neutral oil products with varying
174 viscosities which can also be used to blend lube oil (Basel Convention, 2002).

175 **2.5 Membrane Technology**

176 Membrane technology is another method for removal of contaminants of used lubricating
177 oils. In this process, three types of polymer hollow fibre membranes [polyethersulphone
178 (PES), polyvinylidene fluoride (PVDF), and polyacrylonitrile (PAN)] (Lam et al., 2016) were
179 used for recycling the used engine oils. The process is carried out at 40⁰C and 0.1 MPa
180 pressure. The process is a continuous operation as it removes metal and particles and dusts
181 from used lube oil and improves the recovered oils liquidity and flash point. (Dang, C.S,
182 (1997) and Hamawand et al., (2013).

183 Despite the above mentioned advantages, the expensive membranes may get damaged and
184 fouled by large particles with time. (Dang, C.S, (1997) and Hamawand et al., (2013).

185 **2.6 Catalytic Process**

186 For example, Hylube process from Germany. This process allows production of mainly base
187 oils. The Hylube process is a proprietary process developed by Universal Oil Products (UOP)
188 for the catalytic processing of used lube oils into re-refined lube base stocks for re-blending
189 into saleable lube base oils (Kalnes et al., 2006). This is the first re-refining process in which
190 as received used oil is processed, without any pre-treatment, in a pressurized hydrogen
191 environment. A typical HyLube process feedstock consists of a blend of used lube oils
192 containing high concentrations of particulate matter such as iron and spent additive
193 contaminants such as zinc, phosphorous, and calcium (Chari K.R, 2012).

194 The Hylube unit operates with reactor section pressures of 60–80 bar and reactor
195 temperatures in the range 300–350°C (Kalnes T.N and Schuppel A, 2007). The Hylube
196 process achieves more than 85% of lube oil recovery from the lube boiling range
197 hydrocarbon in the feedstock (Kupareva et al., 2013). Besides the advantages of these
198 process, this method is very expensive. This method requires high level personnel due to high
199 temperature and pressure operations.

200 **3. Combined Technologies/methods**

201 These are advance methods that combines two or more generic methods in its process. Due to
202 the complex nature of removal of contaminants in used lube oils, using a single method may
203 not give you the desired standard emission controlled process. Therefore some companies
204 have developed specific processes for treatment and removal of contaminants in used lube
205 oils (Basel Convention, 2002; Brinkman, 2010 and Kupareva et al., 2013), these methods
206 require sophisticated technologies, equipment and processes. Some of these complex
207 processes are briefly discussed below;

208 **3.1 Vaxon process**

209 This process contains chemical treatment, vacuum distillation and solvent refining units. The
210 advantage of the Vaxon process is the special vacuum distillation, where the cracking of oil is
211 strongly decreased. (Chari, K.R, 2012).

212 The chemical final stage does not, however allow the production of high quality base oils;
213 although in Spain the Catalonia refinery produces base stocks accepted by an original
214 equipment manufacturer (OEM). In connection with this fact, the lube distillate obtained
215 from the Vaxon process (Denmark) or North Refining (Netherlands) are precursors for the
216 Avista Oil base. (Kupareva et al., 2013).

217 **3.2 CEP process**

218 This process combines thin film evaporation and hydroprocessing. The used oil is chemically
219 pre-treated to avoid precipitation of contaminants which can cause corrosion and fouling of
220 the equipment. The pre-treating step is carried out at temperatures within 80–170°C. The
221 chemical treatment compound comprises sodium hydroxide, which is added in a sufficient
222 amount to give a pH about 6.5 or higher. (Magnabosco L.M and Rondeau W.A, 1993).

223 Heavy materials such as residues, metals, additive degradation products, etc. are passed to a
224 heavy asphalt flux stream. The distillate is hydropurified at high temperature (315°C) and
225 pressure (90 bar) in a catalytic fixed bed reactor. (Merchaoui et al., 1994) This process
226 removes nitrogen, sulphur, chlorine and oxygenated organic components. In the final stage of
227 the process, three hydrotreating (Hydrofinishing) reactors are used in series to reduce sulfur
228 to less than 300 ppm and to increase the amount of saturated compounds to over 95%, in
229 order to meet the key specifications for API Group II base oil. The final step in this process is
230 vacuum distillation to separate the hydrotreated base oil into multiple viscosity cuts in the
231 fractionator. The yield of base oils is about 70%. (Kupareva et al., 2013).

232 **3.3 Ecohuile process**

233 The re-refining process was based on vacuum distillation and acid-clay treatment steps until
234 the end of 2000. (Audibert, F., 2006). Clay adsorption was banned on 1 January 2001 and the
235 plant was modified and upgraded to the Sotulub process. (Sotulub re-refining process. 2005).
236 Moreover, the addition of injection facilities of so-called Antipoll-additive (1–3 wt% of pure
237 sodium hydroxide) has been provided and has allowed solving the following basic problems:

- 238 • corrosion of dehydration column and cracking column top section due to the organic acidity
239 of the used oil;
- 240 • plugging of equipment and piping due to polymer formation in the cracking section;
- 241 • high losses of base oil in the oily clay due to the high consumption of clay.

242 The Sotulub process (Merchaoui, M, H et al., 1994) is based on treatment of the used oil with
243 an alkali additive called Antipoll and high vacuum distillation. The used oil is pre-heated to
244 about 160°C and mixed with a small amount of Antipoll-additive, which decreases equipment
245 fouling. This process, allows a final product to be obtained with acceptable quality without
246 any additional finishing stage. Oil obtained is additionally fractionated to obtain various base
247 oil cuts. The process provides base oils with a yield of 82–92 %.(Sotulub re-refining process.
248 2005).

249 **3.4 Cyclon process**

250 This process combines the technology of vacuum distillation and hydrofinishing.
251 (Havemann, 1997). The process licence belongs to Kinetic Technology International (KTI).
252 (Kajdas, C, 2000). In this process, used oils taken from storage tanks are dewatered and the
253 light hydrocarbons are removed by distillation. The heavier fraction is sent to high vacuum
254 distillation, where the majority of base oil components are evaporated from the heavy
255 residue. The oils in the residues are extracted with propane in the de-asphalting unit and sent
256 to the hydroprocessing unit where the other oils are processed. Then they are treated with
257 hydrogen and fractionated based on the desired base oil features. The re-refined base oil
258 products obtained are of high quality due to the hydrogenation. (Schiessler, N. et al., 2007
259 and Tsalavoutas, S. et al., 2002)

260 **3.5 STP method**

261 This is another advance method that combines vacuum distillation and hydrofinishing process
262 (Basel Convention, 2002). It produces less harmful pollutants therefore its environmentally
263 friendly (Kupareva et al., 2013). This method involves dehydration, vacuum distillation,
264 separation of the lubricating fraction and hydrofinishing of base oil separation from the
265 residue.

266 **3.6 Interline process**

267 Interline proposes a process based on propane de-asphalting at ambient temperature and
268 under a pressure that facilitates separation in the liquid phase. The lubricating oil yield
269 declared for the Interline process is 79 %.(Monier V and Labouze E, (2001); European IPPC
270 Bureau, Spain (2003). and Aramburu J.A, (2003). The extraction process removes the
271 majority of additives. The process is interesting from the economics point of view because it
272 eliminates thin film distillation and the need for hydrogenation. Both investment and
273 maintenance costs are low.

274 The drawbacks of the Interline process are that the feed should not contain polychlorinated
275 biphenyls (PCBs), and its chlorine content should be below 1000 ppm, since this process has
276 no final hydrofinishing step.

277 **3.7 Propak thermal cracking process**

278 The Propak process consists of screening and dewatering sections, followed by a thermal
279 cracking section, a separation or distillation depending on the product state desired and
280 finally purification and stabilization stages. In certain plant configurations, a heavy boiling
281 fraction is recycled back to the fired process heater. Gasoil in the liquid state is led to the
282 stabilization section from distillation.

283 This technology is characterized by a large operational and product flexibility. Process
284 operating conditions (temperature, pressure, residence time) can be varied to produce a
285 desired product such as heavy fuel oil, gasoil or base oil. (Kupareva et al., 2013)

286 **4.0 Current Technologies for Used Oil Re-refining**

287 Used lube oil normally tends to have a high concentration of potentially harmful pollutant
288 materials and heavy metals which could be dangerous to both living and non-living things on
289 the earth. Used lube oil may cause damage to the environment when dumped into the ground
290 or into water streams including sewers. This may result in ground water and soil

291 contamination. (Hopmans, 1974). Therefore, development of environmentally safe,
292 sustainable and cost-effective solution is required for recycling of used lubricant. (Stehlik,
293 2009).

294 Nowadays due to different treatment and finishing methods, there are currently available
295 many new technologies,(Bridjanian, H and Sattarin, M., 2006) such as pyrolytic distillation
296 method (PDM), pyrolysis process (PP), thin film evaporation (TFE), including combined
297 TFE and clay finishing, TFE and solvent finishing, TFE and hydrofinishing, thermal de-
298 asphaltting (TDA), TDA and clay finishing, TDA and hydrofinishing etc. In addition,
299 environmentally friendly and affordable solvent extraction and adsorbents are being
300 developed as a means of removing contaminants in used lube oil. Some of the current
301 methods are briefly discussed below;

302 From the research conducted by Arpal et al., (2010), a fuel named as diesel-like fuel (DLF)
303 was produced by applying pyrolytic distillation method. Filtration of the waste engine oil
304 sample was done using a quantitative filter. Three additives known as Na_2CO_3 , zeolite and
305 CaO were blended with the purified oil at different ratios and were exposed to thermal and
306 pyrolytic treatment to convert them into a diesel-like fuel. Conclusively, effects of DLF on
307 the oil properties shows a closer range to that of diesel fuel. (Temitayo et al., 2018)

308 Also, Pyrolysis process (PP) has been used as an alternative means of effective conversion of
309 used lubricants to a refined one (Lam et al., (2016); and Manasomboonphan and Junyapoon,
310 (2012). Lam et al., (2016), describe pyrolysis as a thermal process that heats and decomposes
311 substance at high temperature ($300\text{-}1000^\circ\text{C}$) in an inert environment without oxygen.
312 Pyrolysis process is not yet widespread but it has been receiving much attentions nowadays
313 due to its potential to produce energy-dense products from materials. Examples of pyrolysis
314 process includes Microwave Pyrolysis Process (MPP) and Conventional Pyrolysis Process
315 (CPP). The MPP is a thermo-chemical process applied to waste to wealth process of electrical
316 power input of 7.5kW at a flow rate of 5kg/h. (Temitayo et al., 2018).

317 Thin film evaporation technology includes a rotating mechanism inside the evaporator vessel
318 which creates high turbulence and thereby reduces the residence time of feed-stock oil in the
319 evaporator. This is done in order to reduce coking, which is caused by cracking of the
320 hydrocarbons due to impurities in the used oil. Cracking starts to occur when the temperature
321 of the feedstock oil rises above 300°C .

322 However, any coking which does occur will foul the rotating mechanism and other
323 mechanisms such as tube-type heat exchangers are often found in thin film evaporators.
324 Solvent extraction processes are widely applied to remove asphaltic and resinous
325 components.

326 Liquid propane is by far the most frequently used solvent for de-asphalted residues to make
327 lubricant bright stock, whereas liquid butane or pentane produces lower grade de-asphalted
328 oils more suitable for feeding to fuel-upgrading units. The liquid propane is kept close to its
329 critical point and, under these conditions, raising the temperature increases selectivity. A
330 temperature gradient is set up in the extraction tower to facilitate separation. Solvent-to-oil
331 ratios are kept high because this enhances rejection of asphalt from the propane/oil phase.
332 Counter-current extraction takes place in a tall extraction tower. Typical operating conditions
333 can be found in the work by (Mortier and Fox, 2010)

334 Recent studies showed that propane can be replaced by an alcohol–ketone mixture, which
335 reduces coking and fouling problems during distillation. (Kamal and Khan, 2009 & Durrani
336 et al., 2010). The solvent chosen should meet the following requirements: maximum
337 solubility for the oils and minimum solubility for additives and carbonaceous matter; ability
338 to be recovered by distillation. New plant units increasingly use N-methylpyrrolidone
339 because it has the lowest toxicity and can be used at lower solvent/oil ratios, saving energy.
340 Independent of the contacting method used, the end result is two product streams. The
341 raffinate stream is mainly extracted oil containing a limited amount of solvent, while the
342 extract stream is a mixture of solvent and aromatic components. The streams are handled
343 separately during solvent recovery and the recovered solvent streams are recombined and
344 recycled within the plant.

345 However, solvent recovery is an energy-intensive part of the solvent extraction process. For
346 several years, catalytic hydrotreatment stood out as the modern and successful refining
347 treatment from the point of view of the yield and quality of the finished products.
348 Hydroprocessing is more often applied as a final step in the re-refining process in order to
349 correct problems such as poor colour, oxidation or thermal stability, demulsification and
350 electrical insulating properties. . (Kupareva et al., 2013)

351 In hydrofinishing, used oil and hydrogen are pre-heated and then oil allowed to trickle
352 downwards through a reactor filled with catalyst particles where hydrogenation reactions take
353 place. The oil product is separated from the gaseous phase and then stripped to remove traces
354 of dissolved gases or water. Typical reactor operating conditions for hydrofinishing can be
355 found (Mortier and Fox, 2010).

356 The following reactions can be operative: hydrorefining reactions with the objective of
357 removing heteroelements and to hydrogenate olefinic and aromatic compounds, and
358 hydroconversion reactions aiming at modifying the structure of hydrocarbons by cracking
359 and isomerization. (Audibert, 2006).

360 Hydrotreatment catalysts are made of an active phase constituted by molybdenum or tungsten
361 sulfides as well as by cobalt or nickel on oxide carriers. Generally applied combinations are
362 Co-Mo, Ni-Mo, and Ni-W for the active phase and high surface area γ -alumina (transition
363 alumina) carrier. The metal content, expressed as oxides can reach 12–15 wt. % for Mo and
364 3–5 wt. % for Co or Ni. Co-Mo catalysts are preferentially used for hydrodesulphurization
365 and Ni- Mo for hydrogenation and hydrodenitrogenation. Ni-W catalysts are applied for low-
366 sulphur feeds. The most-used carriers are alumina and alumina-silica, the latter being
367 characterized by a higher cracking activity. (Audibert, 2006).

368 The currently applied catalysts in re-refining are modified in order to improve the product
369 base oil quality and to decrease the coke formation, however, their composition is typically
370 not disclosed in an open literature. The technologies applying hydroprocesses require
371 relatively high investments compared with others. However, depending on the technology
372 adopted, the total cost might be lower than in solvent extraction process due to the high
373 operating costs to make up for the solvent losses. On the other hand, solvent extraction and
374 chemical treatment processes do not require catalyst regeneration. Moreover, it is not
375 necessary to establish a hydrogen gas supply facility in these methods which in addition
376 reduces a risk concerning operation safety. (Kupareva et al., 2013)

377 5.0 Conclusion

378 Currently applied technologies can be compared in terms of their operating and capital costs,
379 quality of feedstock and products obtained. These advance combine technology processes
380 and/or methods are mainly found in developed countries but not available in developing
381 countries. These methods when applied generates reduced concentrations of pollutant but
382 require complex and expensive equipment which are rarely found in developing countries.
383 Under increasing environmental pressure of the conventional treatment method such as acid-
384 clay treatment, which was the first oil regeneration process used, it was substituted in the
385 majority of European countries with new technologies based on solvent extraction, pyrolysis,
386 membrane etc. The modern technologies based on solvent extraction, pyrolysis, membrane
387 etc. are environmentally controllable but their operating and capital costs are high, low yields
388 and requires highly skilled personnel (compared to conventional method) is the major
389 drawback. Also, the challenge of cost reduction resulting from the vacuum distillation and
390 hydroprocessing technique. The combined treatment methods have shown remarkable well
391 with high treatment efficiency, environmentally friendly. However, the problem of high cost
392 and season skilled operating personnel remains a major gap in used lube oil treatment.
393 Therefore, there is the need to developed viable, efficient, environmentally friendly,
394 affordable treatment and high yield technique such as solvent extraction coupled with
395 adsorption process to remove contaminants in used lube oil.

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