Science and Technology Publishing Inc Environment and Green Technology Available online <u>www.scitecpub.com</u>

ISBN: 978-0-9886890-0-8, 11/25/2012, Page: 1-35

Alternative Liquid Hydrocarbon Fuel Production Experimental Comparative Study for LDPE Waste and LDPE Standard Plastic

¹Moinuddin Sarker*, ²Mohammad Mamunor Rashid

Natural State Research, Inc. Department of Research and Development, 37 Brown House Road (2nd floor), Stamford, CT 06902, USA Phone: 203-406-0675, Fax: 203-4069852

*E-mail: msarker@naturalstateresearch.com

Table of Content	Page No.
1. Introduction	3
2. Material and Method	3
2.1. Materials	3
2.2. Pre-analysis	4
2.3. Process Description	4
3. Result and Discussion	6
3.1. Analytical Procedure	6
3.2. Pre-analysis Discussion	7
3.3. Liquid Fuel Analysis	18
3.4. Solid Residue Analysis	
4. Conclusion	
Acknowledgement	34
References	34

Moinuddin Sarker

1. Introduction

Currently there is a large amount of waste plastics generated in the world. The consumption and production of polymers are increasing worldwide. This increasing amount of polymers wastes generates environmental problems. Utilization of these wastes is important from energetic and political aspects ^[1-4]. The rapid rate of plastic consumption throughout the world has led to the creation of increasing amounts of waste plastics is very small ^[5] and depending on the area of application, the service life of plastic products ranges from 1 to 35 years ^[6]. The weighted average service life of all plastics products are different in various countries based on their life style and economy. Waste plastics can be classified as industrial and municipal waste, according to their origins; these groups have different qualities and properties and are subjected to different management strategies ^[7]. Waste plastics represent a considerable part of municipal wastes; furthermore huge amounts of waste plastic arise as a byproduct of faulty product in industry and agriculture ^[8].

There are several methods of chemical recycling are presently in use, such as direct chemical treatment involving gasification, smelting by blast furnace ^[9] or coke oven ^{[10],} degradation by liquefaction ^{[11],} and gasification [^{12]}. Condensation polymers such as PETE and nylon undergo degradation to produce monomer units ^[13], while from vinyl polymers such as polyolefins, a mixture containing numerous components may be obtained for use as a fuel. Catalytic cracking and reforming facilitate the selective degradation of waste plastics. The use of solid catalysts such as silica-alumina, ZSM-5, Zeolites, and mesoporous, materials has been reported for this purpose ^[14-17]. These materials effectively convert polyolefins into liquid fuel, giving lighter fractions as compared to thermal cracking. Thermal degradation of mixed plastics is currently receiving renewed interest as route for disposal, if the large quantities of plastic wastes collected by different collecting systems. The advantage of thermal degradation of macromolecules in the absence of air (pyrolysis) compared to combustion is a reduction in the volume of product gases by a factor of 5-20 which leads to considerable savings in the gas conditioning equipment. Furthermore, it is possible to obtain valuable hydrocarbon compounds.

The pyrolysis is complicated by the fact that plastics show poor thermal conductivity while the degradation of macromolecules requires large amounts of energy. Other studies in the literature ^[18-23] have investigated the degradation of polyethylene or polypropylene and polystyrene mixtures, and the results of the interactions between these polymers. William and William have analyzed the binary degradation of polystyrene and polypropylene, and it was found that the overall degradation rate of mixture was greater than the combined degradation rates of the individual components. According to the results of the products of polyethylene, polypropylene and polystyrene cracking have the most favourable properties for further energetic application. Linear paraffin content in the gas oil fraction is advantageous because this hydrocarbon structure has the highest cetane rate, but it also has the lowest octane number in the naphtha fraction. Further tests are being performed on how to generate the best quality fuel from these waste plastic. The optimal quality of fuel will represent similar characteristics and chemical properties to commercial fuels. Overall, we have seen that when mixture of waste plastics is processed through the thermal degradation process, the quality and efficiency is achieved.

Keywords: hydrocarbon, liquid fuel, waste plastic, ldpe, thermal, GC/MS, FT-IR

2. Material and Method

2.1. Materials

Low density polyethylene (LDPE) waste plastic collected from Stamford local city grocery store and LDPE waste plastic was shopping bags. Low density polyethylene (LDPE) waste plastic color was varieties type such as transparent, white, black etc. Collected waste plastic comes with foreign raw materials such as food particles, paper,

sand and dust etc. All foreign materials separated by manually and waste plastic washed with liquid detergent. During waste plastic washing period also generating waste water and this waste water keep into separate container for treatment purpose. This process main goal is environmental protection and reduce waste plastic problem from the environment. Waste water treatment process applied only alkali and acidic solution for coagulation and flocculation process. After retreat clean waste water can be use for waste plastic washing purpose. Wash out waste plastic dried into laboratory room temperature and cut into small pieces manually by using cutting scissor. Standard LDPE plastic collected from Sigma Aldrich Company, CAS number is 9002-88-4, lot number is MKBG7350V, Chemical structure is C_2H_4 and catalog number is 428043-1kg. Standard LDPE plastic color is transparent and small pellet size.

2.2. Pre-analysis

Low density polyethylene (LDPE) raw waste plastic and standard LDPE raw plastic was pre-analyzed by using GC/MS, FT-IR, TGA, EA-2400 and ICP. By using GC/MS analysis was showed LDPE waste and standard plastics compound structure identification. FT-IR analysis showed LDPE waste and standard plastic functional group identification and TGA analysis result showed onset temperature which are help to liquefaction temperature determination. By EA- 2400 analysis result showed raw materials carbon, hydrogen and nitrogen percentage. Finally ICP test analysis result showed trace metal present into raw materials which are help to breakdown long chain hydrocarbon to shorter chain hydrocarbon in thermal degradation process. Above all pre-analysis results are described into pre-analysis discussion section.

2.3. Process Description

Low density polyethylene (LDPE) waste plastic and LDPE standard plastic to liquid hydrocarbon fuel production purpose thermal degradation process applied. In the two types experimental process raw sample was used 1 kg of LDPE waste plastic and 1 kg of standard plastic as well. Both experiment performed same parameter and same procedure. Catalyst and vacuum system was not applied in the two experiments. Under Labconco fume hood in present of oxygen and fully close system experiment were performed. Both experiments performed into stainless steel reactor and temperature range was 25- 430 °C. Every experiment temperature was monitored watlow meter and experiment temperature was increase gradually from room temperature to 430°C. For LDPE waste and standard plastic to fuel production set up temperature determine from TGA result based on onset temperature (see **table 4**). 1st experiment was performed with LDPE standard plastic to liquid fuel and 2^{nd} experiment was performed LDPE waste plastic to liquid fuel in the same procedure. 1 kg (1000gm) of standard plastic inserted into reactor chamber and reactor cover was tighten properly, then condensation unit, liquid fuel collection unit, light gas cleaning device, small pump and light gas storage device was set up properly (Seen fig.1). LDPE standard plastic started to melt from 25 °C to 430 °C, subsequently reactor temperature increased gradually. When temperature was increased gradually from 25 °C to 430 °C observed that some vapor started to come out through the condenser unit. Temperature range when 250 °C overcome, found that vapor started to condensed and ultimately turned into drop wise liquid fuel. Because the thermal degradation experimental process first able standard plastic melted, then turn into liquid slurry, then converted to vapor and vapor passed through into condenser unit and at the end liquid fuel produced and collected in the container. In this experiment without vacuum system thermal conversion process was done in presence of moisture and the moistures were not affecting conversion process. Plastic to fuel production process reaction are heterogeneous and strongly endothermic reaction. Plastic hydrocarbon chain bonds are breaking down by heat as well as carbon and moisture produce compounds which is not affecting conversion process. Polymer carbon chain and moisture are reacted each other and possible reaction are mentioned such as $C + H_2O = CO + H_2$, C+ CO₂ =2CO, C+ ¹/₂ O₂= CO, C+O₂ =CO₂, CO+H₂O =CO₂+H₂, C+2H₂= CH₄, CO+ 3H₂= CH₄+H₂O, CO₂+4H₂ = CH_4 +2H₂O. Standard plastic to fuel production period some light fraction gas also generated and those gas mixtures of methane, ethane, propane and butane respectively. Produced light gas was purified by using alkali wash solution

and alkali solution normality was 0.25 (N) and the solution was sodium bicarbonate and in water solvent. Produced light gas was storage into Teflon bag by using small pump.



Figure 1: LDPE waste plastic and standard plastic to fuel production process

Liquid fuel was purified by using RCI purification technology and provided RCI filter purification system used to remove fuel sediment and water portion. From 1 kg standard plastic to liquid fuel production percentage is 80.84%, light gas production percentage is 18.95% and solid black residue percentage is 0.21%. LDPE standard plastic density is 0.78 g/ml. Solid black residue percentage is less because of experimental purposes used 99.99 % pure standard LDPE plastic for liquefaction process to liquid fuel. Standard plastic has low amount of additives content and standard plastic has additives level is parts per billion (ppb). Standard plastic to fuel production mass balance result indicates that from 1 kg of plastic conversion into liquid fuel 808.40 gm, light gas conversion into 189.5 gm and left over residue is 2.1 gm. 1000 gm standard plastic to 1040 ml liquid fuel production input electricity was 6.84 kWh and total time was consumed for whole process 4 hours and 56 minutes. On the other experiment we used waste plastic 1kg (1000gm) and same parameter and same temperature profile. LDPE Waste plastic to fuel production percentage is 80.64%, light gas production percentage is 17.91 % and left over residue percentage is 1.45%. LDPE waste plastic residue percentage is little high because LDPE waste plastic has high percentage

additives. LDPE waste plastic to fuel density is 0.79 g/ml. 1 kg of LDPE waste plastic to 1020 ml fuel production need electricity was 7.25 kWh. LDPE waste plastic to liquid fuel production electricity input was little higher than LDPE standard plastic to liquid fuel. Waste plastic to fuel production total time consumed also higher than standard plastic to fuel production because waste plastic has high amount of metal content and more additives compare with pure standard plastic. Manufacture Company is adding different type of additive for better shape of plastic use. These types of additives we are getting as a solid black residue. Both plastics to liquid fuel production comparison result **table 1** showed for visual understanding. Waste plastic to fuel production technology is patent pending technology and convert all types of waste plastic to fuel for next generation.

Plastics Name	Sample Weight (g.)	Liquid Fuel (g.)	Liquid Fuel (ml)	Sample as Light Gas (g.)	Residue Weight (g.)	Electricity Consumption kWh	Liquid Fuel Yield %	Light Gas Yield %	Solid Residue Yield %
Standard	1000	808.40	1040	189.5	2.1	6.84	80.84	18.95	0.21
LDPE									
LDPE	1000	806.4	1020	179.1	14.5	7.25	80.64	17.91	1.45
Waste									
Plastic									

Table 1: LDPE waste plastic and standard plastic to fuel production percentage data

3. Result and Discussion

3.1. Analytical Procedure

Perkin Elmer TGA pyris-1 was used for raw materials onset and inflection temperature measuring. Helium gas was use for purge and temperature range was used 50-800 °C. Temperature increased range was 20°C/ minute. From this analysis we calculated how much percentage conversion rate from plastic to fuel by using thermal degradation process. TGA analysis gives us liquefaction temperature for plastic and leftover residue percentage. Perkin Elmer FT-IR spectrum 100 was used for two type of sample analysis. 1st pre-analysis of solid raw LDPE standard and LDPE waste plastic and secondly was used for liquid fuel from LDPE standard and waste plastic. Solid sample analysis purpose use ATR system and liquid sample analysis purpose was used NaCl cell system. For liquid sample analysis scan number was 32, resolution was 4 cm⁻¹ and wave range was 4000-400 cm⁻¹. By using FT-IR analysis was giving us wave functional group bend energy which is resemble to calorific value. By using GC/MS analysis was solid hard standard and waste plastic also both liquid fuels. Sold sample was analysis by using pyroprobe and temperature was 1200 °C to sample make volatile for GC column. When liquid fuel was analysis by using GC/MS that time was used auto sample system. Solid and liquid sample analysis purpose was same GC/MS column. Carrier gas was use for sample carrier helium. GC/MS program set up for liquid fuel analysis initial temperature 40 °C and hold for 1 minute, final temperature 325 °C and temperature ramping rate 10 °C per minute. Final temperature hold 15 minutes, equilibration time 0.5 minute and total experiment run time 45.50 minutes. Carrier gas used Helium and Perkin Elmer Elite 5MS capillary column used for GC. Column length 30 m, ID 0.25 mm and DF 0.5 um. Column temperature range was - 60 to 350 °C. MS method set up for mass scan Ion mode EI +, data format Centroid, start mass 35.00, end mass 528, scan time 0.25 sec and inter scan time 0.15 sec. Perkin Elmer EA -2400 used for raw waste plastics CHN percentage analysis. Finally ICP (Induced Couple Plasma) was used for trace metal analysis from raw materials and solid residue.

3.2. Pre-analysis Discussion

Test Method	Trace Metal	Raw LDPE Waste Plastic	Raw LDPE Standard Plastic
Name	Name	(ppm)	(ppb)
ASTM D1976	Silver	<1.0	<1.0
	Aluminum	197.4	99.7
	Boron	2.8	47.1
	Barium	<1.0	42.6
	Calcium	962.6	738.8
	Chromium	<1.0	23.7
	Copper	<1.0	2.2
	Iron	6.0	<1.0
	Potassium	35.4	<50.0
	Lithium	<1.0	<1.0
	Magnesium	25.1	<1.0
	Molybdenum	<1.0	<1.0
	Sodium	45.2	10181.0
	Nickel	<1.0	4.3
	Phosphorus	26.7	<1.0
	Lead	<1.0	<1.0
	Antimony	<1.0	2.5
	Silicon	90.2	<1.0
	Tin	<1.0	<1.0
	Strontium		2.7
	Titanium	2.7	<1.0
	Thallium		<1.0
	Vanadium	<1.0	<1.0
	Zinc	2.6	21.3

Table 2: LDPE waste and standard plastic analysis by ICP

American Standard and Testing Method (ASTM D1976) analysis was performed by ICP (Induced Couple Plasma) of raw waste LDPE plastic numerous metal contents are found in the analysis (table 2). We noticed that LDPE waste and standard plastic some metal content are very high compare to less contents metal in the residue. In raw waste LDPE plastic metal contents are measured in ppm that's Parts per Million and for raw standard LDPE plastic metal contents are measured in ppb that is Parts per Billion. Because of standard plastics is much more pure than waste plastics. LDPE waste plastic has high and low contents metal elements such as Silver <1.0 ppm, Aluminium 197.4 ppm, Boron 2.8 ppm, Barium <1.0 ppm, Calcium 962.6 ppm, Chromium <1.0 ppm, Copper <1.0 ppm, Iron 6.0 ppm, Potassium 35.4 ppm, Lithium <1.0 ppm, Magnesium 25.1 ppm, Molybdenum <1.0 ppm, Sodium 45.2 ppm, Nickel <1.0 ppm, Phosphorus 26.7 ppm, Lead <1.0 ppm, Antimony <1.0 ppm, Silicon 90.2 ppm, Tin <1.0 ppm, Strontium 0 ppm, Titanium 2.7 ppm, Thallium 0 ppm, Vanadium <1.0 ppm and Zinc 2.6 ppm etc. On the other hand Standard Raw LDPE plastic same metal contents are appeared in the analysis but metal quantity contents are making differ of each other. In some case found same quantity but in some case quantity varies on metal to metal including less content and high content. In the analysis of standard LDPE plastic metal quantity contents are found comparative with raw standard LDPE plastic metal contents are following such as Silver <1.0 ppb, Aluminium 99.7 ppb, Boron 47.1 ppb, Barium 42.6 ppb, Calcium 738.8 ppb, Chromium 23.7 ppb, Copper 2.2 ppb, Iron <1.0 ppb, Potassium <50.0 ppb, Lithium <1.0 ppb, Magnesium <1.0 ppb, Molybdenum <1.0 ppb, Sodium 10181.0 ppb, Nickel 4.3 ppb, Phosphorus <1.0 ppb, Lead <1.0 ppb, Antimony 2.5 ppb, Silicon <1.0 ppb, Tin <1.0 ppb, Strontium 2.7 ppb, Titanium <1.0 ppb, Thallium <1.0 ppb, Vanadium <1.0 ppb and Zinc 21.3 ppb etc. In some parameter found that raw standard LDPE plastic and raw waste LDPE plastic metal quantity contents are similar. On the other hand in raw Standard LDPE plastic same metal contents are appeared in the analysis but metal quantity contents are making differ of each other. In some case found same quantity but in some case quantity varies on metal to metal including less content and high content as well. LDPE waste plastic metal content more than standard plastic because when manufacturing company making plastic for consumer that times manufacture company adding extra additives for plastic good shape such as hardness, softness and thickness. Also raw LDPE waste plastics are abandoning in the open nature for long run as well as contaminated with different types of metal as appeared in the analysis.

Test Method Name	Plastics Name	Carbon (C) %	Hydrogen (H) %	Nitrogen (N)%
ASTM D5291_a	Raw LDPE Waste Plastics	85.33	14.31	< 0.30
	Raw LDPE Standard Plastics	86.00	13.68	< 0.30

Table 3: LDPE waste and standard plastic C, H and N percentage by EA-2400

ASTM D5291_a, (American Standard and Testing Method) analysis (table 3) was performed of raw waste LDPE plastic and raw standard LDPE plastic analysis by Elemental Analyzer 2400 (EA-2400) following percentage of Carbon, Hydrogen and Nitrogen contents are appeared. In the analysis of both raw waste and raw standard plastics such as in raw waste LDPE Carbon 85.33%, Hydrogen 14.31% and ultimately Nitrogen % <0.30% as well .On the other hand raw LDPE standard plastics Carbon 86.00%, Hydrogen 13.68% and ultimately Nitrogen <0.30% respectively. In the comparative analysis of carbon, hydrogen and nitrogen contents found that raw standard LDPE plastic has more carbon and hydrogen percentage content than raw waste LDPE plastic and Nitrogen contents are same percentage <0.30% in the both residue as well. Fundamentally raw Standard LDPE plastic is more pure than waste LDPE plastic and waste LDPE plastic are exists in the nature for long run as well as contaminated with metal and other substances. Also during manufacturing different type's additives and dyes are added into the waste LDPE plastics in order to give durable shape and color of waste plastics. Therefore in raw standard LDPE much more pure then raw waste LDPE plastic as aspect of carbon and hydrogen percentage contents.

Table 4: TGA analysis result of LDPE waste plastic and standard LDPE plastic

Sample Name	Sample weight (g.)	Onset temperature (°C)	Inflection point temperature (°C)
LDPE waste plastic	3.13	421.53	457.11
LDPE standard plastic	2.72	434.11	466.57

LDPE waste plastic and LDPE standard plastic was per analysis by using Thermogravimetric analyzer (TGA Pyris-1) for raw sample (**Table 4**) onset temperature measured. TGA analysis purposed helium gas was use as carrier gas and temperature profile was use 50 °C to 800 °C and temperature ramping rate was use 20 °C per minutes. In table 4 analysis result showed LDPE waste plastic onset temperature was 421.53 °C and LDPE standard plastic onset temperature was 434.11 °C from TGA analysis. By using this equipment raw samples onset temperature determined and this temperature was help to setup liquefaction process because before start liquefaction process waste plastic to fuel need to know first temperature profile.

Table 5: FT-IR spectrum of LDPE waste plastic functional group name

Number of	Band Number	Functional	Number of	Band Number	Functional
Peak	(cm ⁻¹)	Group Name	Peak	(cm ⁻¹)	Group Name
1	2916.01	CH_2	5	1377.72	CH_3

Moinuddin Sarker



Figure 2: FT-IR spectrum of LDPE waste plastic

2

3

4

100.0

Perkin Elmer FT-IR analysis of LDPE raw waste plastic (**fig.2 and table 5**) according to their wave number and spectrum band following types of functional groups are appeared in the analysis. In the spectrum field we noticed that higher wave number are emerged in the initial phase and middle index of the spectrum and in higher wave number small and bulky both functional groups are available and in low wave number double bond and single bond functional groups are available such as methane group, cis and trans alkene etc. Hereafter wave number 2916.01 cm⁻¹, functional group is CH₂, 2848.31 cm⁻¹ functional group is C-CH₃ wave number 1645.62 cm⁻¹, functional group is Non-Conjugated, wave number 1462.89 cm⁻¹ and 1377.72 cm⁻¹ functional group is CH₃, and ultimately wave number 729.84 cm⁻¹ and 719.08 functional group is -CH=CH-(cis) as well. Energy values are calculated, using formula is E=hv, Where h=Planks Constant, h =6.626x10⁻³⁴ J, v= Frequency in Hertz (sec⁻¹), Where v=c/ λ , c=Speed of light, where, c=3x10¹⁰ m/s, W=1/ λ , where λ is wave length and W is wave number in cm⁻¹. Therefore the equation E=hv, can substitute by the following equation, E=hcW. According to their wave number several energy values are calculated such as for 2916.01 (cm⁻¹) calculated energy, E=5.79x10⁻²⁰ J, wave number 2848.31 (cm⁻¹), calculated energy, E=3.26x10⁻²⁰ J, wave number 1645.62 (cm⁻¹), calculated ener

9

1462.89 (cm⁻¹), calculated energy, $E=2.90 \times 10^{-20}$ J, wave number 1377.72 (cm⁻¹), calculated energy, $E=2.73 \times 10^{-20}$ J, wave number 729.84 (cm⁻¹), calculated energy, $E=1.44 \times 10^{-20}$ J, Similarly, wave number 719.08 (cm⁻¹) energy, $E=1.42 \times 10^{-20}$ J respectively.



Figure 3: FT-IR spectrum of LDPE standard plastic

Table 6: FT-IR spectrum	of LDPE standard p	lastic functional	group name
-------------------------	--------------------	-------------------	------------

Number of Peak	Band Number (cm ⁻¹)	Functional Group Name	Number of Peak	Band Number (cm ⁻¹)	Functional Group Name
1	2915.53	CH_2	4	1377.28	CH ₃
2	2849.03	CH_2	5	717.51	-CH=CH-(cis)
3	1470.61	CH ₃			

From FT-IR analysis of LDPE raw standard plastic (**fig.3 and table 6**) according to their wave number and spectrum band following types of functional groups are appeared in the analysis. In the spectrum field we noticed that higher wave number are emerged in the initial phase and middle index of the spectrum and in higher wave number small and bulky both functional groups are available and in low wave number double bond and single bond functional groups are available such as methane group, cis and trans alkene etc. Hereafter wave number 2915.53 cm⁻¹,

functional group is CH₂, 2849.03 cm⁻¹ functional group is C-CH₃, wave number 1470.61 cm⁻¹, functional group is CH₃, wave number 1377.28 cm⁻¹ functional group is CH₃, and ultimately wave number 717.51 cm⁻¹ functional group is -CH=CH-(cis) as well. Energy values are calculated, using formula is E=hv, Where h=Planks Constant, h =6.626x10⁻³⁴ J, v= Frequency in Hertz (sec⁻¹), Where v=c/ λ , c=Speed of light, where, c=3x10¹⁰ m/s, W=1/ λ , where λ is wave length and W is wave number in cm⁻¹. Therefore the equation E=hv, can substitute by the following equation, E=hcW. According to their wave number several energy values are calculated such as for 2915.53 (cm⁻¹) calculated energy, E=5.79x10⁻²⁰ J, wave number 2848.03 (cm⁻¹), calculated energy, E=5.65x10⁻²⁰ J, wave number 1470.61 (cm⁻¹), calculated energy, E=2.92x10⁻²⁰ J, wave number 1377.28 (cm⁻¹), calculated energy, E=2.73x10⁻²⁰ J and Similarly, wave number 717.51 (cm⁻¹) energy, E=1.42x10⁻²⁰ J respectively.



Figure 4: GC/MS chromatogram of LDPE waste plastic

Table 7: LDPE waste plastic GC/MS chromatogram compound list

Peak	Retention	Trace	Compound	Compound	Molecular	Probability	NIST
Number	Time	Mass	Name	Formula	Weight	%	Library
	(M)	(m/z)					Number
1	2.18	41	Cyclopropane	C3H6	42	40.3	18854
2	2.25	41	2-Butene, (E)-	C ₄ H ₈	56	16.3	105
3	2.44	42	Cyclopropane, ethyl-	C5H10	70	39.4	114410
4	2.58	67	1,4-Pentadiene	C5H8	68	24.6	114494
5	2.63	66	Cyclopropane, ethylidene-	C5H8	68	10.2	152269
6	2.73	67	Cyclopentene	C ₅ H ₈	68	22.3	19032
7	2.86	67	1,5-Hexadiene	C ₆ H ₁₀	82	30.3	227588
8	2.94	41	Cyclopropane, propyl-	C ₆ H ₁₂	84	26.3	60624

Copyright © 2012 www.scitecpub.com

9	3.54	67	Cyclopentene, 3-methyl-	C ₆ H ₁₀	82	13.6	114408
10	3.68	78	Benzene	C ₆ H ₆	78	73.5	114388
11	3.77	79	Cyclopentene,3- methylene-	C ₆ H ₈	80	19.9	151094
12	3.93	67	Cyclohexene	C6H10	82	19.0	114431
13	4.07	41	1-Heptene	C7H14	98	16.6	19704
14	4.20	43	Heptane	C7H16	100	46.3	61276
15	4.68	83	Cyclohexane, methyl-	C7H14	98	36.1	118503
16	4.99	81	1,4-Hexadiene, 4-methyl-	C7H12	96	7.35	113135
17	5.13	55	1-Octyn-3-ol	C8H14O	126	8.34	113255
18	5.28	67	1,3-Pentadiene, 2,4- dimethyl-	C7H12	96	13.4	114450
19	5.44	79	Cyclopentanepropanol, 2- methylene-	C9H16O	140	19.2	160878
20	5.52	91	Cyclobutene, 2- propenylidene-	С7Н8	92	18.0	29595
21	5.62	81	Cyclopentene, 4,4- dimethyl-	C7H12	96	8.49	38642
22	5.75	82	3,4-Nonadiene	C9H16	124	8.23	54088
23	5.87	67	1,4-Octadiene	C8H14	110	30.5	113431
24	6.09	41	1-Octene	C8H16	112	11.0	191147
25	6.31	43	Hexane, 3-ethyl-	C8H18	114	21.06	113940
26	6.51	67	Bicyclo[5.1.0]octane	C8H14	110	9.46	149566
27	6.65	55	2-Octene	C8H16	112	11.1	118191
28	6.96	67	1,4-Octadiene	C8H14	110	24.5	113431
29	7.17	41	1,9-Nonanediol	C9H20O2	160	11.0	114694
30	7.93	91	Cyclohexanol, 1-ethynyl-, carbamate	C9H13NO2	167	26.0	246016
31	8.44	55	Cyclohexane, cyclopropyl-	C9H16	124	20.9	26670
32	8.53	55	1,8-Nonadiene	C9H16	124	29.8	107523
33	8.81	55	1-Nonene	C9H18	126	10.3	142583
34	9.05	43	Nonane	C9H20	128	16.5	2665
35	9.37	83	Cyclopentane, 1-methyl- 2-(2-propenyl)-, trans-	C9H16	124	43.0	26931
36	10.41	67	Cyclopentene, 1-(3- methylbutyl)-	C ₁₀ H ₁₈	138	10.7	61018
37	10.54	56	5-Dodecene, (E)-	C ₁₂ H ₂₄	168	4.86	61866
38	11.13	56	Nonane, 5-methylene-	C ₁₀ H ₂₀	140	29.4	61927
39	11.47	41	1,9-Decadiene	C ₁₀ H ₁₈	138	32.2	118291
40	11.77	55	1-Decene	C ₁₀ H ₂₀	140	5.50	118883
41	11.99	57	Decane	C ₁₀ H ₂₂	142	27.0	114147
42	12.47	55	Cyclohexane, 1-methyl-2- propyl-	C ₁₀ H ₂₀	140	30.4	114020
43	12.72	55	1,11-Dodecadiene	C ₁₂ H ₂₂	166	11.0	113595
44	13.95	56	Cyclobutane, 1-hexyl-2,3- dimethyl-	C ₁₂ H ₂₄	168	4.65	60877
45	14.38	41	1,10-Undecadiene	C ₁₁ H ₂₀	152	24.1	113574
46	14.66	41	1-Undecanol	C ₁₁ H ₂₄ O	172	5.78	114087
47	14.88	57	Undecane	C ₁₁ H ₂₄	156	28.1	249213

4917.15411,11-Dodecadiene $C_{12}H_{22}$ 16613.45017.40411-Tridecanol $C_{13}H_{28}O$ 2004.515117.6157Dodecane $C_{12}H_{26}$ 17015.75217.95556-Dodecene, (Z)- $C_{12}H_{24}$ 1685.275319.77411,11-Dodecadiene $C_{12}H_{22}$ 1668.035420.00551-Tridecanol $C_{13}H_{28}O$ 2008.105520.1957Tridecane $C_{12}H_{22}$ 18428.7	113595 114368 291499 142611 113595 114368 107767 113609 113612 61052 113925
50 17.40 41 1 -Tridecanol $C_{13}H_{28}O$ 200 4.51 51 17.61 57 Dodecane $C_{12}H_{26}$ 170 15.7 52 17.95 55 6 -Dodecene, (Z)- $C_{12}H_{24}$ 168 5.27 53 19.77 41 $1,11$ -Dodecadiene $C_{12}H_{22}$ 166 8.03 54 20.00 55 1 -Tridecanol $C_{13}H_{28}O$ 200 8.10 55 20.19 57 Tridecane $C_{12}H_{22}$ 184 28.7	114368 291499 142611 113595 114368 107767 113609 113612 61052 113925
51 17.61 57Dodecane $C_{12}H_{26}$ 170 15.7 52 17.95 55 6 -Dodecene, (Z)- $C_{12}H_{24}$ 168 5.27 53 19.77 41 $1,11$ -Dodecadiene $C_{12}H_{22}$ 166 8.03 54 20.00 55 1 -Tridecanol $C_{13}H_{28}O$ 200 8.10 55 20.19 57 Tridecane $C_{12}H_{20}$ 184 28.7	291499 142611 113595 114368 107767 113609 113612 61052 113925
52 17.95 556-Dodecene, (Z)- $C_{12}H_{24}$ 1685.275319.77411,11-Dodecadiene $C_{12}H_{22}$ 1668.035420.00551-Tridecanol $C_{13}H_{28}O$ 2008.105520.1957Tridecane $C_{12}H_{22}$ 18428.7	142611 113595 114368 107767 113609 113612 61052 113925
5319.77411,11-Dodecadiene $C_{12}H_{22}$ 1668.035420.00551-Tridecanol $C_{13}H_{28}O$ 2008.105520.1957Tridecane $C_{12}H_{20}$ 18428.7	113595 114368 107767 113609 113612 61052 113925
54 20.00 55 1-Tridecanol C13H28O 200 8.10 55 20.19 57 Tridecane C12H20 184 28.7	114368 107767 113609 113612 61052 113925
55 20.19 57 Tridecane CraHao 184 29.7	107767 113609 113612 61052 113925
-55 - 20.17 - 57 - 1100000 - 0131128 - 104 - 20.7	113609 113612 61052 113925
56 21.76 56 2-Butyl-1-decene C ₁₄ H ₂₈ 196 24.3	113612 61052 113925
57 22.23 55 1,13-Tetradecadiene C ₁₄ H ₂₆ 194 12.7	61052 113925
58 22.45 41 Cyclotetradecane C ₁₄ H ₂₈ 196 4.46	113925
59 22.62 57 Tetradecane C14H30 198 31.3	
60 24.56 55 1,13-Tetradecadiene C ₁₄ H ₂₆ 194 18.1	113612
61 24.75 55 Cyclopentadecane C ₁₅ H ₃₀ 210 6.52	114876
62 24.91 43 Pentadecane C ₁₅ H ₃₂ 212 22.3	22620
63 26.76 55 11-Hexadecen-1-ol, (Z)- C ₁₆ H ₃₂ O 240 7.82	108369
64 26.93 41 1-Hexadecanol C ₁₆ H ₃₄ O 242 6.55	114116
65 27.07 57 Nonadecane C ₁₉ H ₄₀ 268 25.4	114098
66 27.39 55 1-Hexadecene C ₁₆ H ₃₂ 224 5.03	118882
67 28.84 55 11-Hexadecen-1-ol, (Z)- C ₁₆ H ₃₂ O 240 8.92	108369
68 30.96 83 1-Docosene C ₂₂ H ₄₄ 308 6.54	113878
6931.0757NonadecaneC19H4026817.7	114098
70 32.69 82 11-Hexadecen-1-ol, (Z)- C ₁₆ H ₃₂ O 240 11.1	108369
71 32.83 83 1-Nonadecene C19H38 266 8.38	113626
72 32.93 57 Nonadecane C19H40 268 29.8	114098
7334.48551,19-EicosadieneC20H382788.98	241604
74 34.60 83 1-Docosene C ₂₂ H ₄₄ 308 8.82	113878
75 34.70 71 Nonadecane C ₁₉ H ₄₀ 268 21.8	114098
76 36.19 55 1,19-Eicosadiene C ₂₀ H ₃₈ 278 10.2	241604
77 36.31 83 1-Docosene C ₂₂ H ₄₄ 308 10.2	113878
78 36.39 57 Nonadecane C19H40 268 8.38	114098
79 37.83 55 1,15-Pentadecanediol $C_{15}H_{32}O_2$ 244 10.8	113063
80 37.94 97 1-Eicosanol C ₂₀ H ₄₂ O 298 10.7	113075
81 38.02 71 Heneicosane C ₂₁ H ₄₄ 296 7.46	107569
82 39.40 55 1,15-Pentadecanediol $C_{15}H_{32}O_2$ 244 11.5	113063
83 39.50 83 1-Eicosanol C ₂₀ H ₄₂ O 298 9.71	113075
84 39.56 57 Nonadecane C ₁₉ H ₄₀ 268 7.47	114098
85 40.92 82 1,19-Eicosadiene C ₂₀ H ₃₈ 278 9.30	241604
86 41.00 83 1-Eicosanol C ₂₀ H ₄₂ O 298 12.6	113075
87 41.07 57 Tetracosane $C_{24}H_{50}$ 338 11.0	248196
88 42.44 55 9-Tricosene, (Z)- C ₂₃ H ₄₆ 322 7.69	70967
89 42.51 57 Octacosane C ₂₈ H ₅₈ 394 10.5	149865
90 43.77 82 1,19-Eicosadiene C ₂₀ H ₃₈ 278 8.90	241604
91 45.21 97 1-Eicosanol C ₂₀ H ₄₂ O 298 13.1	113075
92 46.72 83 1-Eicosanol C ₂₀ H ₄₂ O 298 11.4	113075
93 48.34 83 1-Eicosanol C ₂₀ H ₄₂ O 298 13.5	113075

From GC-MS pyroprobe analysis of raw LDPE waste plastics inside of pyroprobe raw solid waste plastics turns into volatile gas with high temperature at 1200 °C and that volatile gas passed through the column to gas chromatography, helium (He) is used as a carrier gas and then sends the volatile gas to the mass spectroscopy and in mass compounds are detected according to the boiling point of individual compound and among those only several compounds are introduced as well elaborated in the analysis (fig.4 and table7). In accordance with the retention time and trace masses numerous different types of hydrocarbon compound and benzene derivatives compounds are appeared in the analysis result index. Many compounds are emerged on the analysis carbon range C_3 to C_{28} . In the initial state of the analysis index according to the retention time such as retention time 2.18 and trace mass 41, compound is single bond Cyclopropane (C_3H_6), retention time 2.25 and trace mass 41 compound is double bond 2-Butene,(E)- (C_4H_8), retention time 2.44 and trace mass 42, compound is single bond Cyclopropane,ethyl-(C_5H_{10}), retention time 2.58 and trace mass 67, compound is 1-4 Pentadiene (C_5H_8), retention time 2.94 and trace mass 41, compound is Cyclopropane, Propyl (C_6H_{12}), retention time 3.54 and trace mass 67, compound is double bond Cyclopentene, 3-methyl- (C_6H_{10}), retention time 3.93 and trace mass 67, compound is Cyclohexane (C_6H_{10}), retention time 4.07 and trace mass 41, compound name is 1-Heptene (C_7H_{14}), retention time 4.20 and trace mass 43, compound is Heptane (C_7H_6), retention time 4.68 and trace mass 83, compound is Cyclohexane, methyl- (C_7H_{14}), retention time 4.99 and trace mass 81, compound is 1,4-Hexadiene, 4-methyl- (C_7H_{12}), retention time 5.28 and trace mass 67, compound is 1,3-Pentadiene, 2,4-dimethyl- (C_7H_{12}), retention time 5.87 and trace mass 67, compound is double bond 1,4-Octadiene (C_8H_{14}), retention time 6.65 and trace mass 55, compound is double bonding 2-Octene (C_8H_{16} , retention time 7.93 and trace mass 91, compound is Cyclohexanol, 1-ethynyl-, carbamate (C9H13NO2) here appearing that oxygen compound are produced because in the reactor during reaction phase oxygen induce from moisture, retention time 8.81 and trace mass 55, compound is 1-Nonene (C_9H_{18}), retention time 9.37 and trace mass 83, compound is Cyclopentane, 1-methyl-2-(2-propenyl)-, trans- (C₉H₁₆), retention time 10.54 and trace mass 56, compound name is double bond 5-Dodecene, (E)- ($C_{11}H_{24}$), retention time 11.99 and trace mass 57, compound is single bond Decane ($C_{10}H_{22}$), retention time 13.95 and trace mass 56, compound is Cyclobutane, 1-hexyl-2,3dimethyl- ($C_{11}H_{24}$), retention time 17.95 and trace mass 55, compound is 6-Dodecene, (Z)- ($C_{12}H_{24}$), retention time 20.19 and trace mass 57, compound is single bond Tridecane (C13H26) etc. As well retention time21.76 and trace mass 56, compound is 2-Butyl-1-decene ($C_{14}H_{28}$), retention time 31.07 and trace mass 57, compound is single bond Nonadecane ($C_{19}H_{40}$), retention time 36.19 and trace mass 55, compound is double bond 1, 19-Eicosadiene ($C_{20}H_{38}$), retention time 37.94 and trace mass 97, compound is alcoholic Eicosanol ($C_{20}H_{42}O$), retention time 39.40 and trace mass 55, compound is 1, 15-Pentadecanediol ($C_{15}H_{32}O_2$) etc. At the last phase of the analysis index retention time 40.92 and trace mass 82, compound is 1, 19-Eicosadiene ($C_{20}H_{38}$), retention time 42.44 and trace mass 55, compound is double bond 9-Tricosene, (Z)- ($C_{23}H_{46}$), retention time 43.77 and trace mass 82, compound is 1,19-Eicosadiene ($C_{20}H_{38}$), retention time 46.72 and trace mass 83, compound is 1-Eicosanol ($C_{20}H_{42}O$) and ultimately retention time 48.34 and trace mass 43 compound is alcoholic 1-eicosanol ($C_{20}H_{42}O$) etc.



Figure 5: GC/MS chromatogram of LDPE standard plastic

Peak	Retention	Trace	Compound	Compound	Molecular	Probability	NIST
Number	Time	Mass	Name	Formula	Weight	%	Library
	(min.)	(m/z)					Number
1	2.21	41	Cyclopropane	C3H6	42	32.7	18854
2	2.28	41	2-Butene, (E)-	C4H8	56	14.0	105
3	2.48	55	Cyclopropane, 1,2-	C5H10	70	17.2	114453
			dimethyl-, trans-				
4	2.61	67	1,3-Pentadiene	C5H8	68	17.3	291890
5	2.76	67	Cyclopentene	C5H8	68	20.8	19032
6	2.90	67	1,5-Hexadiene	C6H10	82	35.2	227588
7	2.97	42	1-Hexene	C6H12	84	25.0	227613
8	3.57	67	Cyclopentene, 1-methyl-	C6H10	82	11.5	231297
9	3.73	78	Benzene	C6H6	78	74.2	221957
10	3.96	67	Cyclohexene	C6H10	82	14.6	114431
11	4.09	56	1-Heptene	C7H14	98	33.5	19704
12	4.23	43	Heptane	C7H16	100	43.3	61276
13	4.49	55	2-Heptene	C7H14	98	12.6	113119
14	4.62	81	Norbornane	C7H12	96	10.6	114371
15	4.66	67	1,4-Heptadiene	C7H12	96	20.0	113639

16	4.72	55	Cyclohexane, methyl-	C7H14	98	27.7	118503
17	5.02	81	Cyclohexene, 4-methyl-	C7H12	96	18.7	125422
18	5.59	91	Toluene	C7H8	92	47.9	61211
19	5.79	67	7-	C9H16	124	7.68	210902
			Methylbicyclo[4.2.0]octa				
20	5.01	4.1	ne	C II	110	165	(2101
20	5.91	41	I,7-Octadiene	C ₈ H ₁₄	110	16.5	62191
21	6.12	55	1-Octene	C ₈ H ₁₆	112	17.1	227923
22	6.34	43	Octane	C ₈ H ₁₈	114	40.2	61242
23	7.00	67	1,4-Octadiene	C ₈ H ₁₄	110	28.3	113431
24	7.28	55	2,/-Octadien-1-ol	С8Н14О	126	10.3	237434
25	7.99	91	Ethylbenzene	C ₈ H ₁₀	106	38.5	114918
26	8.25	91	p-Xylene	C ₈ H ₁₀	106	20.2	150787
27	8.47	55	Cyclopentane, 1-methyl- 2-(2-propenvl)-, trans-	С9Н16	124	24.5	26931
28	8.57	54	1,8-Nonadiene	C9H16	124	31.7	107523
29	8.85	56	1-Nonene	C9H18	126	14.7	107756
30	9.08	43	Nonane	C9H20	128	38.8	228006
31	10.46	67	2-Nonen-1-ol, (Z)-	C9H18O	142	13.2	53342
32	11.53	55	1,9-Decadiene	C10H18	138	21.8	232355
33	11.80	41	1-Decene	C ₁₀ H ₂₀	140	12.8	118883
34	12.04	57	Decane	C ₁₀ H ₂₂	142	41.0	114147
35	13.38	115	Benzene, 1-ethynyl-4-	C9H8	116	22.9	43759
26	14 44		methyl-	C. H.	1.50	26.2	112574
36	14.44	55 41		C11H20	152	36.3	113574
3/	14.70	41	I-Undecene	C11H22	154	5.52	5022
38	14.92	57	Undecane	C11H24	156	29.3	249213
39	15.05	55 41	5-Undecene, (E)-	C11H22	154	4.68	114227
40	17.21	41	1,11-Dodecadiene	C12H22	100	26.6	113595
41	17.45	43	I-Dodecene	C12H24	108	6.65	10/688
42	17.66	5/	Dodecane	С12Н26	1/0	27.4	291499
43	19.84	41	1,13-1 etradecadiene	С14Н26	194	15.5	113612
44	20.06	55	1-Iridecene	С13Н26	182	10.3	10//68
45	20.25	57	Tridecane	С13Н28	184	37.6	114282
46	22.30	41	1,13-Tetradecadiene	C ₁₄ H ₂₆	194	19.2	113612
47	22.51	41	1-Tetradecene	C ₁₄ H ₂₈	196	5.03	69725
48	22.68	57	Tetradecane	C ₁₄ H ₃₀	198	33.5	113925
49	24.63	55	1,13-Tetradecadiene	C ₁₄ H ₂₆	194	8.10	113612
50	24.81	41	1-Pentadecanol	C ₁₅ H ₃₂ O	228	5.53	154949
51	24.97	57	Pentadecane	C ₁₅ H ₃₂	212	20.1	107761
52	26.83	55	11-Hexadecen-1-ol, (Z)-	C ₁₆ H ₃₂ O	240	13.9	108369
53	27.01	41	1-Hexadecene	C ₁₆ H ₃₂	224	8.10	118882
54	27.14	57	Hexadecane	C ₁₆ H ₃₄	226	22.7	114191
55	27.32	57	Z-10-Pentadecen-1-ol	C ₁₅ H ₃₀ O	226	5.51	245485
56	28.91	55	11-Hexadecen-1-ol, (Z)-	C ₁₆ H ₃₂ O	240	6.54	108369
57	29.06	55	E-14-Hexadecenal	C ₁₆ H ₃₀ O	238	8.68	130980
58	29.20	57	Heptadecane	C17H36	240	17.7	107308

59	30.89	55	11-Hexadecen-1-ol, (Z)-	C ₁₆ H ₃₂ O	240	6.44	108369
60	31.03	55	E-15-Heptadecenal	C ₁₇ H ₃₂ O	252	9.30	130979
61	31.14	57	Octadecane	C ₁₈ H ₃₈	254	17.0	57273
62	31.34	57	Oxirane, tetradecyl-	C ₁₆ H ₃₂ O	240	8.07	113147
63	32.78	55	1,19-Eicosadiene	C ₂₀ H ₃₈	278	9.35	241604
64	32.90	55	1-Nonadecene	C19H38	266	12.5	113626
65	33.01	57	Nonadecane	C ₁₉ H ₄₀	268	26.6	114098
66	34.58	55	1,19-Eicosadiene	C ₂₀ H ₃₈	278	15.2	241604
67	34.68	43	1-Nonadecene	C19H38	266	7.49	113626
68	34.79	57	Eicosane	C ₂₀ H ₄₂	282	14.9	149863
69	36.28	55	1,19-Eicosadiene	C ₂₀ H ₃₈	278	18.0	241604
70	36.38	57	1-Docosene	C ₂₂ H ₄₄	308	11.3	113878
71	36.46	57	Heneicosane	C ₂₁ H ₄₄	296	18.7	107569
72	37.92	55	1,19-Eicosadiene	C ₂₀ H ₃₈	278	17.7	241604
73	38.01	55	1-Docosene	C ₂₂ H ₄₄	308	12.9	113878
74	38.10	57	Heneicosane	C ₂₁ H ₄₄	296	10.0	107569
75	39.49	55	1,19-Eicosadiene	C ₂₀ H ₃₈	278	11.8	241604
76	39.58	43	1-Docosene	C ₂₂ H ₄₄	308	8.99	113878
77	41.02	55	1,19-Eicosadiene	C ₂₀ H ₃₈	278	9.60	241604
78	41.08	55	1-Docosene	C ₂₂ H ₄₄	308	10.4	113878
79	42.47	55	1,19-Eicosadiene	C ₂₀ H ₃₈	278	10.4	241604
80	42.53	57	1-Docosene	C ₂₂ H ₄₄	308	9.47	113878
81	43.94	55	1-Eicosanol	C ₂₀ H ₄₂ O	298	7.27	113075
82	45.32	57	1-Eicosanol	C ₂₀ H ₄₂ O	298	21.2	113075
83	46.82	55	1-Eicosanol	C ₂₀ H ₄₂ O	298	20.2	113075

Perkin Elmer GC/MS pyroprobe analysis (Fig.5 and table 8) of raw LDPE standard plastics inside of pyroprobe raw solid standard plastics turns into volatile gas with high temperature at 1200 °C and that volatile gas passed through the column to gas chromatography, helium (He) is used as a carrier gas and then sends the volatile gas to the mass spectroscopy and in mass compounds are detected according to the boiling point of individual compound and among those only several compounds are introduced as well elaborated in the analysis. In accordance with the retention time and trace masses numerous different types of hydrocarbon compound and benzene derivatives compounds are appeared in the analysis result index. Many compounds are emerged on the analysis carbon range C_3 to C_{22} . In the initial state of the analysis index according to the retention time such as retention time 2.21 and trace mass 41, compound is single bond Cyclopropane (C_3H_6), retention time 2.28 and trace mass 41 compound is double bond 2-Butene, (E)- (C_4H_8), retention time 2.48 and trace mass 55, compound is single bond Cyclopropane, 1,2dimethyl-, trans (C_5H_{10}), retention time 2.61 and trace mass 67, compound is 1-3 Pentadiene (C_5H_8), retention time 2.97 and trace mass 42, compound is 1-Hexene (C_6H_{12}), retention time 3.57 and trace mass 67, compound is double bond Cyclopentene, 1-methyl- (C_6H_{10}), retention time 3.96 and trace mass 67, compound is Cyclohexene (C_6H_{10}), retention time 4.09 and trace mass 56, compound name is 1-Heptene (C_7H_{14}), retention time 4.23 and trace mass 43, compound is Heptane (C7H6), retention time 4.66 and trace mass 67, compound is 1,4-Heptadiene (C7H12), retention time 5.02 and trace mass 81, compound is Cyclohexane, 4-methyl- (C_7H_{12}), retention time 5.59 and trace mass 91, compound is Toluene (C_7H_8), retention time 5.79 and trace mass 67, compound is double bond 7-Methylbicyclo[4.2.0]octane (C₉H₁₆), retention time 6.12 and trace mass 55, compound is double bonding 1-Octene (C_8H_{16} , retention time 7.00 and trace mass 67, compound is 1,4-Octadiene (C8H14), retention time 7.18 and trace mass 55, compound is 2,7-Octdoine-1-ol Octadiene (C8H14O), retention time 8.85 and trace mass 55, compound is 1-Nonene (C_9H_{18}), retention time 9.08 and trace mass 43, compound is Nonane (C_9H_{20}), retention time 10.46 and trace mass 67, compound name is 2-Nonene-1-ol (Z)-, ($C_9H_{18}O$), retention time 11.80 and trace mass 41, compound is double bond 1-Decene ($C_{10}H_{20}$), retention time 13.38 and trace mass 115, compound is Benzene, 1-ethynyl-4-methyl- (C_9H_8), retention time 14.92 and trace mass 57, compound is Undecane, ($C_{11}H_{24}$), retention time 17.66 and trace mass 57, compound is single bond Dodecane ($C_{12}H_{26}$) etc. As well retention time22.68 and trace mass 57, compound is Tetradecane ($C_{14}H_{30}$), retention time 31.03 and trace mass 55, compound is alcoholic E-15-Heptadecenal ($C_7H_{32}O$), retention time 36.46 and trace mass 57, compound is Heneicosane ($C_{21}H_{44}$), retention time 37.92 and trace mass 55, compound is 1,19- Eicosandiene ($C_{20}H_{38}$), retention time 39.58 and trace mass 43, compound is 1, 19-Eicosadiene ($C_{20}H_{38}$), retention time 42.47 and trace mass 55, compound is 1, 19-Eicosadiene ($C_{20}H_{38}$), retention time 42.47 and trace mass 55, compound is 1, 19-Eicosandiene ($C_{20}H_{42}O$) and ultimately retention time 46.82 and trace mass 55 compound is alcoholic 1-Eicosanol ($C_{20}H_{42}O$) etc. here appearing that several oxygen compounds are produced because in the reactor during reaction phase oxygen induce from steam and moisture as well.

3.3. Liquid Fuel Analysis



Figure 6: GC/MS Chromatogram of LDPE waste plastic to fuel

Table 9: LDPE waste plastic to fuel GC/MS chromatogram compound list

Peak Number	Retention Time (min.)	Trace Mass (m/z)	Compound Name	Compound Formula	Molecular Weight	Probability %	NIST Library Number
1	1.50	39	Propane	C3H8	44	87.5	18863
2	1.61	43	Butane	C4H10	58	63.5	18940

Copyright © 2012 www.scitecpub.com

3	1.88	42	Cyclopropane, ethyl-	C5H10	70	18.0	250
4	1.91	43	Pentane	C5H12	72	83.1	114462
5	1.95	55	1-Butene, 3-methyl-	C5H10	70	18.1	114463
6	2.02	55	Cyclopropane, 1,2- dimethyl-, cis-	C5H10	70	25.5	19070
7	2.06	67	1,3-Pentadiene	С5Н8	68	21.0	291890
8	2.13	67	1,4-Pentadiene	С5Н8	68	18.2	114494
9	2.25	67	2,3-Diazabicyclo[2.2.1]- hept-2-ene	C5H8N2	96	14.2	142950
10	2.50	41	1-Hexene	C6H12	84	17.3	500
11	2.57	57	Hexane	C6H14	86	73.4	291337
12	2.72	67	3-Hexen-1-ol, (Z)-	C ₆ H ₁₂ O	100	13.4	114154
13	2.84	67	1,3-Hexadiene,c&t	C6H10	82	7.85	231295
14	2.90	56	Cyclopentane, methyl-	C6H12	84	66.5	114428
15	3.00	67	4-Methyl-2-pentyne	C6H10	82	16.6	231299
16	3.14	67	Cyclopentene, 3-methyl-	C6H10	82	14.0	114408
17	3.30	56	Cyclohexane	C6H12	84	24.5	228008
18	3.62	41	Cyclopentane, 1,2- dimethyl-, cis-	C7H14	98	30.3	114027
19	3.74	43	Heptane	C7H16	100	69.0	61276
20	4.06	81	Cyclopentane, 1-methyl- 2-methylene-	C7H12	96	8.69	62523
21	4.16	55	Cyclohexane, methyl-	C7H14	98	58.9	118503
22	4.31	69	Cyclopentane, ethyl-	C7H14	98	40.3	940
23	4.44	67	Norbornane	C7H12	96	9.79	114371
24	4.61	67	1-Ethylcyclopentene	C7H12	96	37.3	114407
25	4.80	91	Toluene	С7Н8	92	24.5	291301
26	4.86	81	Cyclohexene, 4-methyl-	C7H12	96	12.7	125422
27	5.05	79	1,3,6-Heptatriene	C7H10	94	16.3	113127
28	5.15	41	1-Octene	C8H16	112	24.1	1604
29	5.30	41	Octane	C8H18	114	55.3	229407
30	5.39	55	3-Octene, (Z)-	C8H16	112	12.3	113895
31	5.47	95	1-Methyl-2- methylenecyclohexane	C8H14	110	8.30	113437
32	5.65	67	Cyclopentene, 1-(1- methylethyl)-	C8H14	110	8.37	113932
33	5.74	81	Cyclohexane, ethylidene-	C ₈ H ₁₄	110	8.98	1494
34	5.91	41	2,4-Decadien-1-ol	C ₁₀ H ₁₈ O	154	6.65	136415
35	5.98	55	Cyclohexane, ethyl-	C ₈ H ₁₆	112	50.6	113476
36	6.12	67	4-Octyne	C_8H_{14}	110	12.3	118189
37	6.25	41	2,4-Decadien-1-ol	C ₁₀ H ₁₈ O	154	7.79	136415
38	6.55	81	Cyclohexanol, 1-ethynyl-, carbamate	C9H13NO2	167	13.8	313023
39	6.61	67	1-Methyl-2- methylenecyclohexane	C8H14	110	5.15	113437
40	6.87	41	cis-2-Nonene	C9H18	126	10.6	113508
41	6.96	91	Bicyclo[2.1.1]hexan-2-ol, 2-ethenyl-	C ₈ H ₁₂ O	124	32.1	221372
42	7.02	43	Nonane	C9H20	128	34.4	228006

43	7.44	67	Ethylidenecycloheptane	C9H16	124	17.6	113500
44	7.66	55	Cyclopentane, butyl-	C9H18	126	21.8	114172
45	8.12	41	Cyclopentanol, 1-(1- methylene-2-propenyl)-	C9H14O	138	7.53	152742
46	8.24	41	2-Decen-1-ol	C ₁₀ H ₂₀ O	156	12.6	136260
47	8.41	55	E-1,6-Undecadiene	C11H20	152	4.56	245712
48	8.59	41	1-Decene	C ₁₀ H ₂₀	140	25.5	118883
49	8.74	43	Decane	C ₁₀ H ₂₂	142	58.2	114147
50	8.81	55	2-Decene, (Z)-	C ₁₀ H ₂₀	140	15.2	114151
51	9.40	55	Cyclodecane	C ₁₀ H ₂₀	140	6.73	113565
52	9.56	67	Cyclopentene, 1-pentyl-	C10H18	138	10.0	139585
53	9.65	41	Tricyclo[4.2.1.1(2,5)]deca n-3-ol	C ₁₀ H ₁₆ O	152	9.72	191707
54	9.80	41	Cyclohexene, 3-(2- methylpropyl)-	C10H18	138	15.2	27008
55	10.07	41	Carane, 4,5-epoxy-, trans	C ₁₀ H ₁₆ O	152	5.23	156142
56	10.13	41	3-Undecene, (E)-	$C_{11}H_{22}$	154	7.67	60565
57	10.24	41	1-Undecene	$C_{11}H_{22}$	154	6.39	5022
58	10.38	43	Undecane	$C_{11}H_{24}$	156	50.5	114185
59	10.44	41	5-Undecene, (E)-	$C_{11}H_{22}$	154	10.3	114227
60	10.59	41	2-Pentadecyn-1-ol	C ₁₅ H ₂₈ O	224	12.8	36724
61	11.07	41	3-Undecene, (E)-	$C_{11}H_{22}$	154	6.82	60565
62	11.18	67	1-Undecyne	$C_{11}H_{20}$	152	7.22	36306
63	11.40	41	Cyclopentaneacetaldehyd e, 2-formyl-3-methyl-α- methylene-	C ₁₀ H ₁₄ O ₂	166	5.72	57743
64	11.80	41	3-Dodecene, (E)-	C12H24	168	9.23	113960
65	11.92	43	Dodecane	C12H26	170	32.3	291499
66	11.98	41	3-Dodecene, (E)-	C ₁₂ H ₂₄	168	15.8	70642
67	13.15	41	2-Tridecene, (Z)-	C13H26	182	5.66	142613
68	13.27	41	1-Tridecene	C13H26	182	8.29	107768
69	13.39	71	Tridecane	C13H28	184	36.6	114282
70	13.43	41	5-Tridecene, (E)-	C13H26	182	7.54	142619
71	13.58	41	4-Nonene, 5-butyl-	C13H26	182	6.84	34734
72	13.99	41	1,12-Tridecadiene	C ₁₃ H ₂₄	180	7.10	7380
73	14.10	41	1-Nonadecanol	C ₁₉ H ₄₀ O	284	4.35	13666
74	14.64	41	1-Tetradecene	C ₁₄ H ₂₈	196	6.78	34720
75	14.76	57	Tetradecane	C ₁₄ H ₃₀	198	27.1	113925
76	14.80	41	7-Tetradecene	C ₁₄ H ₂₈	196	8.54	70643
77	15.85	41	Z-10-Pentadecen-1-ol	C ₁₅ H ₃₀ O	226	8.55	245485
78	15.95	41	1-Pentadecene	C ₁₅ H ₃₀	210	5.75	69726
79	10.04	57	Pentadecane	C ₁₅ H ₃₂	212	25.3	107761
80	16.08	41	E-2-Hexadecacen-1-ol	C ₁₆ H ₃₂ O	240	13.2	131101
81	16.82	41	E-2-Octadecadecen-1-ol	C ₁₈ H ₃₆ O	268	7.09	131102
82	17.18	41	1-Hexadecene	C ₁₆ H ₃₂	224	9.91	118882
83	17.28	43	Hexadecane	C ₁₆ H ₃₄	226	2.6	114191
84	17.31	55	Hexadecen-1-ol, trans-9-	C ₁₆ H ₃₂ O	240	5.37	141055
85	18.00	41	1-Docosanol	C ₂₂ H ₄₆ O	326	5.95	23377

86	18.35	55	1-Heptadecanol	C ₁₇ H ₃₆ O	256	7.46	113250
87	18.44	43	Heptadecane	C ₁₇ H ₃₆	240	19.9	107308
88	18.47	55	8-Heptadecene	C ₁₇ H ₃₄	238	8.23	113620
89	19.46	83	1-Octadecanol	C ₁₈ H ₃₈ O	270	5.04	221125
90	19.55	57	Octadecane	C ₁₈ H ₃₈	254	18.3	12337
91	19.63	55	1-Eicosanol	C ₂₀ H ₄₂ O	298	7.01	113075
92	20.61	71	Eicosane	C ₂₀ H ₄₂	282	26.1	290513
93	21.52	55	1-Docosene	C ₂₂ H ₄₄	308	7.07	113878
94	21.61	43	Eicosane	C ₂₀ H ₄₂	282	25.3	290513
95	21.97	97	1-Eicosene	C ₂₀ H ₄₀	280	8.35	13488
96	22.56	70	Eicosane	C ₂₀ H ₄₂	282	15.4	290513
97	23.48	99	Eicosane	C ₂₀ H ₄₂	282	14.1	290513
98	23.65	55	1-Eicosanol	C ₂₀ H ₄₂ O	298	8.30	113075
99	24.36	43	Eicosane	C ₂₀ H ₄₂	282	12.3	290513
100	25.21	99	Eicosane	C ₂₀ H ₄₂	282	12.8	290513
101	25.94	55	1-Docosanol	C ₂₂ H ₄₆ O	326	10.4	23377
102	26.03	57	Eicosane	C ₂₀ H ₄₂	282	12.5	290513
103	26.84	57	Heneicosane	C ₂₁ H ₄₄	296	11.8	107569
104	27.64	57	Heneicosane	C ₂₁ H ₄₄	296	11.8	107569
105	28.42	57	Heneicosane	C ₂₁ H ₄₄	296	6.67	107569
106	28.68	57	17-Pentatriacontene	C35H70	490	14.9	233160
107	29.20	57	Hexacosane	C ₂₆ H ₅₄	366	8.06	107147
108	29.99	57	Tetratetracontane	C44H90	618	9.63	23773
109	30.82	57	Heptacosane	C ₂₇ H ₅₆	380	17.6	79427
110	32.86	57	Heptacosane	C ₂₇ H ₅₆	380	15.8	79427

From GC/MS analysis of waste LDPE plastic to fuel (fig.6 and table 9) in accordance with retention time and trace mass indicate various types of compound are present. High intensity compounds are preferred in the analysis. An investigated carbon range in the analyzed plastic is C_3 to C_{44} because large carbon chains are breaking down into small chain resulting in lower carbon range. Sometimes noticed that for high peak intensity compound probability factor is low percentage, where as in low peak intensity compound probability factor is high percentage. Most of the peaks are considered in the analysis and as per their retention time and trace mass maximum peaks are mentioned, in accordance to retention time 1.49 and trace mass 41, derived compound is Cyclopropane (C_3H_6) with probability 67.2%, retention time 1.59 and trace mass 41, compound is 2-Butene, (E)-, (C4H₈) with probability 18.5%, retention time 1.89 and trace mass 43, compound is Pentane (C_5H_{12}) with probability 79.8%, retention time 2.03 and trace mass 67, compound is 1,3-Pentadiene (C5H8) with probability factor 17.4%, retention time 2.96 and trace mass 79, compound is 1,3-Cyclopentadiene, 5-methyl- (C_6H_8) with probability 30.0%, retention time 3.09, trace mass 67, compound is Cyclopentene, 3-methyl- (C_6H_{10}) with probability 12.2%, retention time 3.90 and trace mass 81, compound is 2,3-Dimethyl-1,4-pentadiene, (C_7H_{12}) with probability 7.20%, retention time 4.01 and trace mass 81, compound is Cyclohexene, 3-methyl-, (C_7H_{12}) with probability 9.73%, retention time 4.81 and trace mass 81, compound is Cyclohexene, 3-methyl-, (C₇H₁₂) with probability 8.73%, retention time 5.11 and trace mass 41, compound is 1-Octene (C_8H_{16}) with probability 19.8%, retention time 5.94 and trace mass 83, compound is Cyclohexane, ethyl-, (C_8H_{16}) with probability 56.2%, retention time 6.85 and trace mass 56, compound is trans-7-Methyl-3-octene (C_9H_{18}) with probability 27.5%, retention time 7.00 and trace mass 43, compound is Nonane (C_9H_{20}) with probability 28.8%, retention time 7.85 and trace mass 67, compound is Cyclopentene, 1-butyl- (C_9H_{16}) with probability 42.9% etc. Also in the middle of the analysis index retention time 8.80 and trace mass 41, compound is 2-Decene, (Z)- (C₁₀H₂₀) with probability 16.6%, retention time 10.24 and trace mass 41, compound is 1-Undecene ($C_{11}H_{22}$) with probability 8.04%, retention time 10.58 and trace mass 41, compound is 2-Pentadecyn-1ol, ($C_{15}H_{28}O$) with probability 11.9%, retention time 11.98 and trace mass 41, compound is 3-Dodecene, (E)-(C12H24) with probability 10.2% etc. As well as Retention time 13.59 and trace mass 142, compound is Bicyclo[4.4.1]undeca-1,3,5,7,9-pentaene, $(C_{11}H_{10})$ with probability 29.2%, retention time 14.95 and trace mass 41, compound is Dodecylsuccinic anhydride ($C_{16}H_{28}O_3$) with probability 4.78%, retention time 15.96 and trace mass 41, compound is Cyclopentadecane ($C_{15}H_{30}$) with the probability 7.23% etc. Accordingly retention time 16.10 and trace mass 41, Compound is E-2-Hexadecacen-1-ol ($C_{16}H_{32}O$) with probability 10.9%, retention time 20.55 and trace mass 41, compound is 9-Nonadecene ($C_{19}H_{38}$) with probability 10.7%, retention time 21.65 and trace mass 43, compound is 4-Tetradecene, (E)- $(C_{20}H_{42})$ with probability 41.2% etc. At the end phase of the analysis index high retention time and trace mass such as retention time 22.64 and trace mass 57, compound is Hexadecane ($C_{21}H_{42}$) with probability 31.7%, retention time 23.59 and trace mass 85, compound is Heneicosane ($C_{21}H_{44}$) with probability 10.8%, and ultimately retention time 27.80 and trace mass 57, compound is Octacosane ($C_{28}H_{58}$) with probability 15.9% and retention time 24.53 and trace mass 57 , compound is Octacosane ($C_{28}H_{58}$) with probability 10.7% and ultimately retention time 32.86 and trace mass 57, compound is Heptacosane ($C_{27}H_{56}$) as well. In the earlier phase analysis appearing that in several oxygen compounds are produced because in the reactor during reaction oxygen induced from steam and water. Also noticed in the standard HDPE plastic to fuel including double bond compound as well as benzene and benzene derivatives compounds are available. In the middle of the analysis index also noticed that one or more alcoholic compounds are appeared as well.



Figure 7: GC/MS Chromatogram of LDPE standard plastic to fuel

Table 10: LDPE standard	plastic to fuel GC/	MS chromatogram co	mpound list
		6	

NumberTimeMassNameFormulaWeight%Library(min.)(m/z)Number	Peak Number	Retention Time (min.)	Trace Mass (m/z)	Compound Name	Compound Formula	Molecular Weight	Probability %	NIST Library Number
--	----------------	-----------------------------	------------------------	------------------	---------------------	---------------------	------------------	---------------------------

Copyright © 2012 www.scitecpub.com

1	1		D	0.11	4.4	40.0	100/3
1	1.57	41	Propane	C3H8	44	49.2	18863
2	1.65	43	Isobutane	C4H10	58	52.8	18936
3	1.69	41	1-Propene, 2-methyl-	C4H8	56	34.3	61293
4	1.70	43	Butane	C4H10	58	69.6	18940
5	1.75	41	1-Propene, 2-methyl-	C4H8	56	21.2	61293
6	1.97	42	Cyclopropane, ethyl-	C5H10	70	24.8	19072
7	2.00	43	Pentane	C5H12	72	86.1	114462
8	2.11	55	Cyclopropane, 1,2- dimethyl-, cis-	C5H10	70	24.4	19070
9	2.16	39	1,3-Pentadiene	C5H8	68	19.4	291890
10	2.36	67	2,3-Diazabicyclo[2.2.1]- hept-2-ene	C5H8N2	96	9.55	194998
11	2.42	42	1-Pentene	C5H10	70	8.38	19081
12	2.60	41	Cyclopropane, 1-ethyl-2- methyl-, cis-	C ₆ H ₁₂	84	19.3	113658
13	2.68	41	Hexane	C6H14	86	79.7	61280
14	2.74	55	3-Hexene, (Z)-	C6H12	84	41.8	114381
15	2.83	67	1,3-Butadiene, 2-ethyl-	C6H10	82	12.5	118159
16	2.96	67	trans-1,4-Hexadiene	C6H10	82	8.78	113648
17	3.01	56	Cyclopentane, methyl-	C6H12	84	63.0	114428
18	3.12	67	2,4-Hexadiene, (Z,Z)-	C6H10	82	14.5	113646
19	3.26	67	Cyclopentene, 3-methyl-	C6H10	82	10.6	114408
20	3.42	41	Cyclohexane	C6H12	84	12.3	228008
21	3.74	41	1-Heptene	C7H14	98	16.4	107734
22	3.86	43	Heptane	C7H16	100	69.0	61276
23	3.96	41	3-Heptene	C7H14	98	28.9	113117
24	4.09	81	1,2-Hexadiene, 5-methyl-	C7H12	96	7.97	27634
25	4.30	55	Cyclohexane, methyl-	C7H14	98	55.3	118503
26	4.44	41	Cyclopentane, ethyl-	C7H14	98	34.2	231044
27	4.75	67	1-Ethylcyclopentene	C7H12	96	38.8	114407
28	5.01	81	Cyclohexene, 3-methyl-	C7H12	96	8.09	19639
29	5.30	41	1-Octene	C8H16	112	25.3	1604
30	5.45	43	Octane	C8H18	114	38.6	229407
31	5.55	55	3-Octene, (Z)-	C8H16	112	11.1	113895
32	5.97	67	1-Methyl-2- methylenecyclohexane	C ₈ H ₁₄	110	29.8	113437
33	6.08	41	3-Decyn-2-ol	C ₁₀ H ₁₈ O	154	8.71	53449
34	6.14	83	Cyclohexane, ethyl-	C ₈ H ₁₆	112	55.1	113476
35	6.29	67	3-Octyne	C8H14	110	11.1	118185
36	6.72	81	Cyclohexene, 1,2- dimethyl-	C ₈ H ₁₄	110	8.39	113912
37	7.04	41	cis-2-Nonene	C9H18	126	11.4	113508
38	7.19	43	Nonane	C9H20	128	30.6	228006
39	7.27	55	4-Nonene	C9H18	126	14.2	113904
40	7.61	67	Ethylidenecycloheptane	C9H16	124	19.2	113500
41	7.83	55	Cyclopentane, butyl-	C9H18	126	12.5	114172
42	8.05	67	Cyclopentene, 1-butyl-	C9H16	124	35.9	113491
43	8.42	56	2-Methyl-1-nonene	C ₁₀ H ₂₀	140	11.0	113561

44	8.65	56	trans-3-Decene	C ₁₀ H ₂₀	140	6.74	113881
45	8.77	41	1-Decene	C ₁₀ H ₂₀	140	23.1	118883
46	8.91	57	Decane	C ₁₀ H ₂₂	142	49.0	114147
47	8.99	55	2-Decene, (Z)-	C ₁₀ H ₂₀	140	13.8	114151
48	9.57	55	2,4-Undecadien-1-ol	C ₁₁ H ₂₀ O	168	8.58	136410
49	9.73	67	Cyclopentene, 1-pentyl-	C ₁₀ H ₁₈	138	7.10	139585
50	10.30	56	4-Undecene, (Z)-	C ₁₁ H ₂₂	154	6.33	142600
51	10.42	41	1-Undecene	C ₁₁ H ₂₂	154	5.48	34717
52	10.56	43	Undecane	C ₁₁ H ₂₄	156	50.8	114185
53	10.62	55	5-Undecene, (E)-	C ₁₁ H ₂₂	154	11.9	114227
54	10.76	41	2-Pentadecyn-1-ol	C ₁₅ H ₂₈ O	224	20.1	36724
55	11.24	41	2,4-Pentadien-1-ol, 3- pentyl-, (2Z)-	C ₁₀ H ₁₈ O	154	6.44	142197
56	11.86	41	3-Dodecene, (E)-	C ₁₂ H ₂₄	168	4.00	70642
57	12.11	57	Dodecane	C ₁₂ H ₂₆	170	34.2	291499
58	12.16	41	3-Dodecene, (E)-	C ₁₂ H ₂₄	168	15.3	70642
59	13.33	41	9-Octadecenal	C ₁₈ H ₃₄ O	266	5.16	35819
60	13.44	41	1-Tridecene	C ₁₃ H ₂₆	182	7.48	107768
61	13.56	57	Tridecane	C ₁₃ H ₂₈	184	38.8	114282
62	13.61	55	5-Tridecene, (E)-	C ₁₃ H ₂₆	182	7.68	142619
63	14.28	41	1-Nonadecanol	C19H40O	284	6.59	13666
64	14.71	41	Z-10-Pentadecen-1-ol	C ₁₅ H ₃₀ O	226	7.23	245485
65	14.82	41	3-Tetradecene, (Z)-	$C_{14}H_{28}$	196	4.41	62806
66	14.93	85	Tetradecane	C14H30	198	35.1	113925
67	14.97	55	3-Tetradecene, (E)-	$C_{14}H_{28}$	196	5.61	139981
68	15.12	55	7-Tetradecene	$C_{14}H_{28}$	196	6.70	70643
69	16.02	41	4-Tridecene, (Z)-	C ₁₃ H ₂₆	182	7.93	142617
70	16.12	41	1-Pentadecene	C ₁₅ H ₃₀	210	5.59	69726
71	16.23	57	Pentadecane	C ₁₅ H ₃₂	212	22.5	107761
72	16.41	41	E-2-Hexadecacen-1-ol	C ₁₆ H ₃₂ O	240	5.01	131101
73	16.99	41	1-Nonadecanol	C19H40O	284	11.9	13666
74	17.35	41	1-Hexadecene	C ₁₆ H ₃₂	224	5.81	118882
75	17.45	71	Hexadecane	C ₁₆ H ₃₄	226	36.6	114191
76	17.63	41	2-Methyl-E-7-hexadecene	C17H34	238	3.91	130870
77	18.18	41	1-Eicosanol	C ₂₀ H ₄₂ O	298	6.05	23222
78	18.43	41	E-2-Octadecadecen-1-ol	C ₁₈ H ₃₆ O	268	9.95	131102
79	18.51	41	1-Nonadecene	C19H38	266	5.70	113626
80	18.60	71	Heptadecane	C ₁₇ H ₃₆	240	29.8	107308
81	18.78	41	2-Methyl-E-7-hexadecene	C17H34	238	9.78	130870
82	19.62	41	1-Docosene	C ₂₂ H44	308	5.60	113878
83	19.70	57	Octadecane	C18H38	254	19.9	57273
84	20.67	41	9-Nonadecene	C19H38	266	10.1	113627
85	20.75	71	Eicosane	C ₂₀ H ₄₂	282	14.0	290513
86	21.67	41	1-Docosene	C ₂₂ H ₄₄	308	7.16	113878
87	21.74	57	Eicosane	C ₂₀ H ₄₂	282	25.4	290513
88	21.88	55	1-Eicosene	C ₂₀ H ₄₀	280	10.7	13488
89	22.63	43	10-Heneicosene (c,t)	C ₂₁ H ₄₂	294	8.05	113073

90	22.70	43	Heneicosane	C ₂₁ H ₄₄	296	26.9	107569
91	22.86	55	1-Heneicosanol	C ₂₁ H ₄₄ O	312	7.56	233005
92	23.55	43	1-Docosene	C ₂₂ H ₄₄	308	9.68	113878
93	23.61	71	Heneicosane	C ₂₁ H ₄₄	296	12.5	107569
94	23.79	55	1-Docosanol	C ₂₂ H ₄₆ O	326	12.7	23377
95	24.49	43	Heneicosane	C ₂₁ H ₄₄	296	11.0	107569
96	25.35	57	Tetracosane	C ₂₄ H ₅₀	338	9.46	248196
97	26.18	57	Heptacosane	C ₂₇ H ₅₆	380	7.68	150574
98	26.98	57	Octacosane	C ₂₈ H ₅₈	394	10.2	134306
99	27.78	57	Octacosane	C ₂₈ H ₅₈	394	12.7	134306
100	28.57	57	Heptacosane	C ₂₇ H ₅₆	380	20.1	79427

GC/MS analysis of solid standard LDPE plastic to fuel (Fig.7 and table 10) in accordance with retention time and trace mass indicate various types of compound are present. High intensity compounds are preferred in the analysis. An investigated carbon range in the analyzed plastic is C_3 to C_{28} because large carbon chains are breaking down into small chain resulting in lower carbon range. Sometimes noticed that for high peak intensity compound probability factor is low percentage, where as in low peak intensity compound probability factor is high percentage. Most of the peaks are considered in the analysis and as per their retention time and trace mass maximum peaks are mentioned, in accordance to retention time 1.57 and trace mass 41, derived compound is Propane (C_3H_6) with probability 49.2%, retention time 1.65 and trace mass 43, compound is Isobutane, (C4H10) with probability 52.8%, retention time 1.69 and trace mass 41, compound is 1-Propene, 2-Methyl- (C_4H_8) with probability 34.3%, retention time 1.70 and trace mass 43, compound is Butane (C_4H_{10}) with probability factor 69.6%, retention time 2.00 and trace mass 43, compound is Pentane, (C_5H_{12}) with probability 86.1%, retention time 2.11, trace mass 55, compound is Cyclopropane, 1,2-dimethyl-, cis- (C_5H_{10}) with probability 24.4%, retention time 2.16 and trace mass 39, compound is 1,3-Pentadiene, 2-ethyl-, (C_5H_8) with probability 19.4%, retention time 3.01 and trace mass 56, compound is Cyclopentane, methyl-, (C_6H_{12}) with probability 63%, retention time 3.74 and trace mass 41, compound is 1-Heptene (C_7H_{14}) with probability 16.4%, retention time 3.86 and trace mass 43, compound is Heptane (C_7H_{16}) with probability 69.0%, retention time 4.09 and trace mass 81, compound is 1,2-Hexadiene, 5-methyl- (C_7H_{12}) with probability 7.97%, retention time 4.75 and trace mass 67, compound is 1-Ethylcyclopentene (C_7H_{12}) with probability 38.8%, retention time 5.97 and trace mass 67, compound is 1-Methyl-2-methylenecyclohexane (C_8H_{14}) with probability 29.8%, retention time 6.72 and trace mass 81, compound is Cyclohexene, 1,2-dimethyl- (C_8H_{14}) with probability 8.39% etc. Also in the middle of the analysis index retention time 6.72 and trace mass 81, compound is Cyclohexene, 1,2-dimethyl- (C_8H_{14}) with probability 8.39%, retention time 7.61 and trace mass 67, compound is Ethylidenecycloheptane (C_9H_{16}) with probability 19.2%, retention time 7.83 and trace mass 55, compound is Cyclopentane, butyl- (C_9H_{18}) with probability 12.5%, retention time 8.99 and trace mass 55, compound is 2-Decene, $(Z) - (C_{10}H_{20})$ with probability 13.8% etc. As well in the middle phase of the analysis index retention time 9.57 and trace mass 55, compound is 2,4-Undecadien-1-ol (C11H20O) with probability 8.58%, retention time 10.76 and trace mass 41, compound is 2-Pentadecyn-1-ol ($C_{15}H_{28}O$) with probability 20.10%, retention time 12.11 and trace mass 57, compound is Dodecane ($C_{12}H_{26}$) with the probability 34.2% accordingly. Also at the end phase of the analysis retention time 13.61 and trace mass 55, Compound is 5-Tridecene, (E)- ($C_{13}H_{26}$) with probability 9.82%, retention time 39.58 and trace mass 57, compound is Heneicosane ($C_{21}H_{44}$) with probability 7.68%, retention time 14.28 and trace mass 41, compound is Nonadecanol, $(C_{19}H_{40}O)$ with probability 6.59%, retention time 17.45 and trace mass 71, compound is Hexadecane ($C_{16}H_{34}$) with probability 36.6%, retention time 19.62 and trace mass 41, compound is 1-Docosene ($C_{22}H_{44}$) with probability 5.60%, and retention time 27.78 and trace mass 57, compound is Octacosane $(C_{28}H_{58})$ with probability 12.7% and ultimately retention time 28.57 and trace mass 57, compound is Heptacosane (C₂₇H₅₆) with probability 20.1% as well. In the analysis appearing that in several oxygen compounds are produced because in the reactor during reaction oxygen induced from steam and water. Also noticed in the standard LDPE plastics including single and double bond compound as well as aliphatic and aromatic derivatives compounds are also available.



Figure 8: FT-IR spectrum of LDPE waste plastic to fuel

Table 11: FT-IR	spectrum of LDPE	waste plastic	to fuel	functional	group)
-----------------	------------------	---------------	---------	------------	-------	---

Number of	Band Number	Functional	Number of	Band Number	Functional
Peak	(cm ⁻¹)	Group Name	Peak	(cm ⁻¹)	Group Name
1	3618.53	Free OH	13	1302.59	
2	3078.00	H Bonded NH	14	1137.52	
3	2925.00	C-CH ₃	15	1074.91	
4	2731.20	C-CH ₃	16	992.06	-CH=CH ₂
5	2671.28	C-CH ₃	17	965.00	-CH=CH-(trans)
6	2332.37		18	909.42	-CH=CH ₂
7	2027.66		19	887.93	$C=CH_2$
8	1821.59	Non-Conjugated	20	768.10	
9	1722.64	Non-Conjugated	21	721.95	-CH=CH-(cis)

Copyright © 2012 www.scitecpub.com

10	1641.69	Conjugated	22	674.32	-CH=CH-(cis)
11	1460.01	CH ₃	23	634.23	
12	1377.90	CH ₃	24	552.74	

FT-IR analysis of LDPE raw waste plastic to fuel (fig. 8 and table 11) according to their wave number and spectrum band following types of functional groups are appeared in the analysis. In the spectrum field we noticed that higher wave number are emerged in the initial phase and middle index of the spectrum and in higher wave number small and bulky both functional groups are available and in low wave number double bond and single bond functional groups are available such as methane group, cis and trans alkene etc. Hereafter wave number 3618.53 cm ¹, functional group is Free OH, wave number 3078.00 cm⁻¹, functional group is H Bonded NH, wave number 2925.00 cm⁻¹, functional group is C-CH₃, 2731.20 cm⁻¹ functional group is C-CH₃ wave number 1821.59 cm⁻¹, functional group is Non-Conjugated, wave number 1722.64 cm⁻¹, functional group is Non-Conjugated, wave number 1641.69 cm⁻¹, functional group is Conjugated, wave number 1460.01 cm⁻¹ and 1377.90 cm⁻¹ functional group is CH₃, wave number 992.06 cm⁻¹ functional group is -CH=CH₂, wave number 965.00 cm⁻¹, functional group is -CH=CH-(trans), wave number 887.93, functional group is C=CH₂, wave number 721.95 cm⁻¹, functional group is -CH=CH-(cis) and ultimately wave number 674.32 cm⁻¹, functional group is -CH=CH-(cis) as well. For several wave number energy values are calculated, using formula is E=hv, Where h=Planks Constant, h = 6.626×10^{-34} J, v= Frequency in Hertz (sec⁻¹), Where $v=c/\lambda$, c=Speed of light, where, c=3x10¹⁰ m/s, W=1/\lambda, where λ is wave length and W is wave number in cm^{-1} . Therefore the equation E=hv, can substitute by the following equation, E=hcW. According to their wave number several energy values are calculated such as for 2925.00 (cm⁻¹) calculated energy, $E=5.79 \times 10^{-20}$ J, wave number 2731.20 (cm⁻¹), calculated energy, $E=5.42 \times 10^{-20}$ J, wave number 1641.69 (cm⁻¹), calculated energy, E=3.26x10⁻²⁰ J, wave number 1460.01 (cm⁻¹), calculated energy, E=2.90x10⁻²⁰ J, wave number 1377.90 (cm⁻¹), calculated energy, $E=2.73 \times 10^{-20}$ J, wave number 721.95 (cm⁻¹), calculated energy, $E=1.44 \times 10^{-20}$ J, Similarly, wave number 674.32 (cm⁻¹) energy, $E = 1.33 \times 10^{-20}$ J respectively.

Table 12: FT-IR spectrum of LDP	E standard plastic to fuel	functional group
---------------------------------	----------------------------	------------------

Number of Peak	Band Number (cm ⁻¹)	Functional Group Name	Number of Peak	Band Number (cm ⁻¹)	Functional Group Name
1	3617.44	Free OH	13	1302.51	
2	3078.01	H Bonded NH	14	1137.62	
3	2924.69	C-CH ₃	15	1075.15	
4	2731.20	C-CH ₃	16	992.04	-CH=CH ₂
5	2671.06	C-CH ₃	17	964.98	-CH=CH-(trans)
6	2331.20		18	909.44	-CH=CH ₂
7	2027.67		19	887.85	$C=CH_2$
8	1821.60	Non-Conjugated	20	768.10	
9	1722.59	Non-Conjugated	21	721.98	-CH=CH-(cis)
10	1641.71	Conjugated	22	674.34	-CH=CH-(cis)
11	1456.01	CH ₃	23	634.22	
12	1377.88	CH ₃	24	552.93	



Figure 9: FT-IR spectrum of LDPE standard plastic to fuel

FTIR analysis of LDPE raw standard plastic to fuel (**fig. 9 and table12**) according to their wave number and spectrum band following types of functional groups are appeared in the analysis. In the spectrum field we noticed that higher wave number are emerged in the initial phase and middle index of the spectrum and in higher wave number small and bulky both functional groups are available and in low wave number double bond and single bond functional groups are available such as methane group, cis and trans alkene etc. Hereafter wave number 3617.44 cm⁻¹, functional group is Free OH, wave number 3078.01 cm⁻¹, functional group is H Bonded NH, wave number 2924.69 cm⁻¹, functional group is C-CH₃, 2731.20 cm⁻¹ functional group is C-CH₃, wave number 1821.60 cm⁻¹, functional group is Non-Conjugated, wave number 1722.59 cm⁻¹, functional group is Non-Conjugated, wave number 1456.01 cm⁻¹ and 1377.88 cm⁻¹ functional group is CH=CH-(trans), wave number 887.85, functional group is C=CH₂, wave number 721.98 cm⁻¹, functional group is -CH=CH-(cis) and ultimately wave number 674.34 cm⁻¹, functional group is -CH=CH-(cis) as well. For several wave number energy values are calculated, using formula is E=hv, Where h=Planks Constant, h =6.626x10⁻³⁴ J, v= Frequency in Hertz (sec⁻¹), Where v=c/ λ , c=Speed of light, where, c=3x10¹⁰ m/s, W=1/ λ , where λ is wave length and W is wave number in cm⁻¹. Therefore the equation E=hv, can substitute by the following equation, E=hcW.

According to their wave number several energy values are calculated such as for 2924.69 (cm⁻¹) calculated energy, $E=5.79 \times 10^{-20}$ J, wave number 2731.20 (cm⁻¹), calculated energy, $E=5.42 \times 10^{-20}$ J, wave number 1641.71 (cm⁻¹), calculated energy, $E=3.26 \times 10^{-20}$ J, wave number 1456.01 (cm⁻¹), calculated energy, $E=2.90 \times 10^{-20}$ J, wave number 1377.88 (cm⁻¹), calculated energy, $E=2.73 \times 10^{-20}$ J, wave number 721.98 (cm⁻¹), calculated energy, $E=1.44 \times 10^{-20}$ J, Similarly, wave number 674.34 (cm⁻¹) energy, $E=1.33 \times 10^{-20}$ J respectively.

Table 13: LDPE waste plas	stic to fuel ASTM test result
---------------------------	-------------------------------

Method Name	Test Name	Results	Units
ASTM D240	Gross Heat of Combustion	18681	BTU/lb
ASTM D240	Gross Heat of Combustion	123631	BTU/gal
	(Calculated)		-
ASTM D4052	API Gravity @ 60°F	46.5	°API
ASTM D86-07b	IBP Recovery	95.6	°C
ASTM D86-07b	5% Recovery	120.0	°C
ASTM D86-07b	10% Recovery	136.7	°C
ASTM D86-07b	20% Recovery	164.4	°C
ASTM D86-07b	30% Recovery	193.3	°C
ASTM D86-07b	40% Recovery	222.2	°C
ASTM D86-07b	50% Recovery	248.9	°C
ASTM D86-07b	60% Recovery	262.2	°C
ASTM D86-07b	70% Recovery	288.9	°C
ASTM D86-07b	80% Recovery	365.6	°C
ASTM D86-07b	90% Recovery	-	°C
ASTM D86-07b	95% Recovery	_	°Č
ASTM D86-07b	FBP Recovery	371.1	°C
ASTM D86-07b	Recovery	82.0	Vol%
ASTM D86-07b	Residue	18.0	Vol%
ASTM D2500	Cloud point	-20.6	°C
ASTM D2500	Cloud Point	-5.1	°E
ASTM D2500	Pour point	-3.1	°C
ASTM D97	Pour point	-24	о <u>г</u>
ASTM D2286	Freezing Point	-11.2	°C
ASTM D2380	Freezing Point	10.0 61.0	°E
ASTM D2380	Fleezing Point	01.0	Г °С
ASTM D2624	Temperature Electrical Canductivity	24.0	-C
ASTM D2624	Electrical Conductivity	1.0	pS/M
ASIM D5453	Sultur	<1.0	mg/kg
AST M D1500	ASIM Color	<5.0	
ASTM D4176	Appearance: Clean and Bright	-	
ASTM D4176	Free Water Content/Particles	No water	mg/kg
ASTM D4176	Haze Rating	6.0	
ASTM D4176	Special Observation	Darker than usual	
ASTM D4737	Cetane Index (Procedure A)	63.2	
ASTM D5708_MOD	Vanadium	<1.0	ppm
ASTM D5708_MOD	Nickel	<1.0	ppm
ASTM D5708_MOD	Iron	<1.0	ppm, or, mg/Kg
ASTM D482	Ash	< 0.001	Wt%
ASTM D93	Procedure Used	А	
ASTM D93	Corrected Flash Point	Below room temperature	°C
ASTM D4530	Average Micro Method Carbon Residue 10% distillation	0.4	Wt%
ASTM D664	Procedure Used	А	

Moinuddin Sarker

ASTM D664	Acid Number	0.10	mgKOH/gm
ASTM D130	Copper Corrosion @ 50°C	1a	
	(122°F)/3 hrs.		
ASTM D2709	Sediment and Water	< 0.005	Vol%
ASTM D5291	Carbon Content	86.25	Wt%
ASTM D5291	Hydrogen Content	13.69	Wt%
ASTM D5291	Nitrogen Content	< 0.75	Wt%

Low density polyethylene (LDPE) waste plastic to produced fuel was analysis (table 13) by 3rd party Intertek laboratory New Jersey, USA, and all test was performed ASTM test followed such as ASTM D240 Gross Heat of Combustion :18681 BTU/lb, ASTM D240 Gross Heat of Combustion, (Calculated): 123631 BTU/gal, ASTM D4052 API Gravity @ 60°F: 46.5 °API, ASTM D86-07b IBP Recovery: 95.6 °C, ASTM D86-07b 5% Recovery: 120.0 °C, ASTM D86-07b 10% Recovery: 136.7 °C, ASTM D86-07b 20% Recovery: 164.4 °C, ASTM D86-07b 30% Recovery: 193.3 °C, ASTM D86-07b 40% Recovery: 222.2 °C, ASTM D86-07b 50% Recovery: 248.9 °C, ASTM D86-07b 60% Recovery: 262.2 °C, ASTM D86-07b 70% Recovery: 288.9 °C, ASTM D86-07b 80% Recovery: 365.6 °C, ASTM D86-07b 90% Recovery: °C, ASTM D86-07b 95% Recovery: °C, ASTM D86-07b FBP Recovery: 371.1 °C, ASTM D86-07b Recovery: 82.0 Vol%, ASTM D86-07b Residue: 18.0 Vol%, ASTM D2500 Cloud point: -20.6 °C, ASTM D2500 Cloud Point : -5.1 °F, ASTM D97 Pour point: -24 °C, ASTM D97 Pour point: -11.2 °F, ASTM D2386 Freezing Point : 16.0 °C, ASTM D2386 Freezing Point: 61.0 °F, ASTM D2624 Temperature: 24.0 °C, ASTM D2624 Electrical Conductivity: 1.0 pS/M, ASTM D5453 Sulfur: <1.0 mg/kg, ASTM D1500 ASTM Color: <5.0, ASTM D4176 Appearance: Clean and Bright, ASTM D4176 Free Water Content/Particles: No water mg/kg, ASTM D4176 Haze Rating : 6.0, ASTM D4176 Special Observation: Darker than usual, ASTM D4737 Cetane Index (Procedure A): 63.2, ASTM D5708 MOD Vanadium: <1.0 ppm, ASTM D5708 MOD Nickel: <1.0 ppm, ASTM D5708 MOD Iron : <1.0 ppm, or, mg/Kg, ASTM D482 Ash: <0.001 Wt%, ASTM D93 Procedure Used A ASTM D93 Corrected Flash Point : Below room temperature °C, ASTM D4530 Average Micro Method Carbon Residue 10% distillation: 0.4 Wt%, ASTM D664 Procedure Used A ASTM D664 Acid Number: 0.10 mgKOH/gm, ASTM D130 Copper Corrosion @ 50°C (122°F)/3 hrs.: 1a, ASTM D2709 Sediment and Water: <0.005 Vol%, ASTM D5291 Carbon Content: 86.25 Wt%, ASTM D5291 Hydrogen Content : 13.69 Wt%, ASTM D5291 Nitrogen Content: <0.75 Wt%.

Method Name	Test Name	Results	Units
		incourts	
ASTM D240	Gross Heat of Combustion	19551	BTU/lb
ASTM D240	Gross Heat of Combustion	128606	BTU/gal
	(Calculated)		
ASTM D4052	API Gravity @ 60°F	47.6	°API
ASTM D86-07b	IBP Recovery	86.7	°C
ASTM D86-07b	5% Recovery	104.4	°C
ASTM D86-07b	10% Recovery	120.0	°C
ASTM D86-07b	20% Recovery	151.1	°C
ASTM D86-07b	30% Recovery	188.9	°C
ASTM D86-07b	40% Recovery	223.3	°C
ASTM D86-07b	50% Recovery	246.7	°C
ASTM D86-07b	60% Recovery	271.1	°C
ASTM D86-07b	70% Recovery	297.8	°C
ASTM D86-07b	80% Recovery	-	°C
ASTM D86-07b	90% Recovery	-	°C
ASTM D86-07b	95% Recovery	-	°C

ASTM D86-07b	FBP Recovery 364.4		°C
ASTM D86-07b	Recovery	Recovery 75.0	
ASTM D86-07b	Residue	25.0	Vol%
ASTM D2500	Cloud point	-16.6	°C
ASTM D2500	Cloud Point	2.1	°F
ASTM D97	Pour point	-18.0	°C
ASTM D97	Pour point	-0.4	°F
ASTM D2386	Freezing Point	13.0	°C
ASTM D2386	Freezing Point	55.0	°F
ASTM D2624	Temperature	24.0	°C
ASTM D2624	Electrical Conductivity	<1	pS/M
ASTM D5453	Sulfur	1.0	mg/kg
ASTM D1500	ASTM Color	<4.0	
ASTM D4176	Appearance: Clean and Bright	Fail	
ASTM D4176	Free Water Content/Particles		mg/kg
ASTM D4176	Haze Rating		
ASTM D4176	Special Observation	Darker than usual	
ASTM D4737	Cetane Index	64.7	
	(Procedure A)		
ASTM D5708_MOD	Vanadium	<1.0	ppm
ASTM D5708 MOD	Nickel	<1.0	ppm
ASTM D5708 MOD	Iron	<1.0	ppm OR, mg/Kg
ASTM D482	Ash	<1.0	Wt%
ASTM D93	Procedure Used	А	
ASTM D93	Corrected Flash Point	Below room temperature	°C
ASTM D4530	Average Micro Method Carbon	0.2	Wt%
	Residue 10% distillation		
ASTM D664	Procedure Used	А	
ASTM D664	Acid Number	< 0.10	mgKOH/gm
ASTM D130	Copper Corrosion @ 50°C	1a	
	(122°F)/3 hrs.		
ASTM D2709	Sediment and Water	< 0.005	Vol%
ASTM D5291	Carbon Content	86.45	Wt%
ASTM D5291	Hydrogen Content	13.51	Wt%
ASTM D5291	Nitrogen Content	< 0.75	Wt%

LDPE standard plastic to fuel (table 14) test was performed from 3rd party Intertek laboratory in New Jersey, USA and all ASTM test method followed such as ASTM D240 Gross Heat of Combustion: 19551 BTU/lb, ASTM D240 Gross Heat of Combustion (Calculated): 128606 BTU/gal, ASTM D4052 API Gravity @ 60°F: 47.6 °API, ASTM D86-07b IBP Recovery: 86.7 °C, ASTM D86-07b 5% Recovery: 104.4 °C, ASTM D86-07b 10% Recovery: 120.0 °C, ASTM D86-07b 20% Recovery :151.1 °C, ASTM D86-07b 30% Recovery: 188.9 °C, ASTM D86-07b 40% Recovery: 223.3 °C, ASTM D86-07b 50% Recovery: 246.7 °C, ASTM D86-07b 60% Recovery: 271.1 °C, ASTM D86-07b 70% Recovery: 297.8 °C, ASTM D86-07b 80% Recovery: °C, ASTM D86-07b 90% Recovery: °C, ASTM D86-07b 95% Recovery: °C, ASTM D86-07b FBP Recovery: 364.4 °C, ASTM D86-07b Recovery: 75.0 Vol%, ASTM D86-07b Residue: 25.0 Vol%, ASTM D2500b Cloud point: -16.6 °C, ASTM D2500 Cloud Point: 2.1°F, ASTM D97 Pour point: -18.0 °C, ASTM D97 Pour point: -0.4 °F, ASTM D2386 Freezing Point: 13.0 °C, ASTM D2386 Freezing Point: 55.0 °F, ASTM D2624 Temperature: 24.0 °C, ASTM D2624 Electrical Conductivity : <1 pS/M, ASTM D5453 Sulfur: 1.0mg/kg, ASTM D1500 ASTM Color: <4.0, ASTM D4176 Appearance Clean and Bright: Fail, ASTM D4176 Free Water Content/Particles: 0 mg/kg, ASTM D4176Haze Rating: nil, ASTM D4176 Special Observation: Darker than usual, ASTM D4737 Cetane Index (Procedure A): 64.7, ASTM D5708 MOD Vanadium: <1.0 ppm, ASTM D5708 MOD Nickel: <1.0 ppm, ASTM D5708 MOD Iron: <1.0 ppm OR, mg/Kg, ASTM D482 Ash: <1.0 Wt%, ASTM D93 Procedure Used _A ASTM D93 Corrected Flash Point: Below room temperature °C, ASTM D4530 Average Micro Method Carbon Residue 10% distillation: 0.2 Wt%, ASTM D664 Procedure Used _A ASTM D664 Acid Number: <0.10 mgKOH/gm, ASTM D130 Copper Corrosion @ 50°C (122°F)/3 hrs.:1a, ASTM D2709 Sediment and Water: <0.005 Vol%, ASTM D5291 Carbon Content: 86.45Wt%, ASTM D5291 Hydrogen Content:13.51 Wt%, ASTM D5291 Nitrogen Content : <0.75 Wt%.

3.4. Solid Residue Analysis

Table 15: LDPE waste plastic and LDPE standard plastic to residue analysis result by ICP

Test Method	Trace Metal	LDPE Waste Plastic to	LDPE Standard Plastic to
Name	Name	Residue (ppm)	Residue (ppm)
	0.1	-1.0	-1.0
ASIM D1976	Silver	<1.0	<1.0
	Aluminum	/806	799.9
	Arsenic	23.6	<1.0
	Boron	24.3	14.9
	Barium	11.4	74.8
	Beryllium	<1.0	<1.0
	Calcium	1041	2879
	Cadmium	<1.0	<1.0
	Chromium	29.2	51.0
	Copper	82.4	38.7
	Iron	4280	2112
	Potassium	106.7	<1.0
	Lithium	<1.0	<1.0
	Magnesium	221.6	125.0
	Manganese	17.8	24.4
	Sodium	1201	152.9
	Nickel	148.7	246.7
	Lead	10.8	<1.0
	Antimony	<1.0	<1.0
	Selenium	<1.0	<1.0
	Silicon	77.0	33.8
	Tin	274.3	152.7
	Titanium	191.1	736.5
	Vanadium	17.0	<1.0
	Zinc	1617	84.5

LDPE waste plastic and LDPE standard plastic to fuel production period some residue was left over that residue was combination of metal and metal was analysis by ICP and ASTM method was ASTM D1976 and **table 15** showed metal content result for LDPE waste plastic to residue such as Silver <1.0 ppm, Aluminum 7806 ppm, Arsenic 23.6 ppm, Boron 24.3ppm, Barium 11.4 ppm, Beryllium <1.0 ppm, Calcium 1041 ppm, Cadmium <1.0 ppm, Chromium 29.2 ppm, Copper 82.4 ppm, Iron 4280 ppm, Potassium 106.7 ppm, Lithium <1.0 ppm, Magnesium 221.6 ppm, Manganese 17.8 ppm, Sodium 1201 ppm, Nickel 148.7 ppm, Lead 10.8 ppm, Antimony <1.0 ppm, Selenium <1.0 ppm, Silicon 77.0 ppm, Tin 274.3 ppm, Titanium 191.1 ppm, Vanadium 17.0 ppm, Zinc 1617 ppm level was present into LDPE waste plastic to residue. On the other hand pure standard LDPE plastic to residue also metal content present such as Silver <1.0 ppm, Aluminum 799.9 ppm, Arsenic <1.0 ppm, Copper 38.7 ppm, Iron 2112 ppm, Potassium <1.0 ppm, Lithium <1.0 ppm, Calcium 2879 ppm, Cadmium <1.0 ppm, Manganese 24.4 ppm, Sodium 152.9 ppm, Nickel 246.7 ppm, Lead <1.0 ppm, Antimony <1.0 ppm, Silicon 33.8 ppm, Tin 152.7

ppm, Titanium736.5 ppm, Vanadium <1.0 ppm, Zinc 84.5 ppm level present respectively. LDPE waste plastic and LDPE standard plastic metal content was vary because LDPE waste plastic was come from consumer level and LDPE plastic was made for consumer and it was made with different types of additives and additives percentage is almost 3-4% and those additives come out as solid black residue. On the other hand pure standard LDPE plastic made for analysis or made for plastic manufacturing company making plastic with adding dye or chemical for that reason additives percentage is less then manufactured consumer plastic. During waste plastic to fuel production or standard plastic to fuel production metal content was not come out that much what ever come it was negligible because all metal content melting point temperature more than higher from waste plastic to fuel production temperature. Metal content was help to break down long chain hydrocarbon to short chain hydrocarbon because metal content react as catalyst because all types of catalyst are made by metal.

Test Method Name	Plastics Name	Carbon (C) %	Hydrogen (H) %	Nitrogen (N)%
ASTM D5291.a	LDPE Waste Plastic to Residue	81.81	2.15	< 0.30
	LDPE Standard Plastic to Residue	68.72	2.15	< 0.30

Table 16: LDPE waste plastic and LDPE standard plastic to residue C, H and N % by EA-2400

LDPE waste plastic and LDPE standard plastic to residue was analysis by Elemental Analyzer 2400 in CHN mode with followed ASTM method ASTM D529_a. In **table 16** analysis result indicate that LDPE waste plastic to residue carbon percentage is 81.81%, hydrogen percentage is 2.15% and nitrogen percentage is less then <0.30%. On the other hand standard pure LDPE plastic to residue analysis result indicate that carbon percentage is 68.72 %, hydrogen percentage is 2.15% and nitrogen percentage is <0.30%. LDPE waste plastic additives percentage is higher than LDPE standard plastic. Residue analysis result showed also Btu value more than 5000/lb and this residue could be use as substantial coal or road carpeting and road carpeting. Both residue was heated up to more than 800 °C and check there physical properties and found it did not burn it and make more hard like stone. During heating period we notice that some black smoke came out. Residue was combination of different category of metal which was used during plastic to fuel production finished.

4. Conclusion

LDPE waste plastic and LDPE standard plastic to fuel production process was applied with thermal degradation without catalyst in present of oxygen under laboratory fume hood. Both experiments were performed same parameter and same condition. In laboratory batch scale process LDPE waste plastic and LDPE standard plastic to fuel production temperature were uses in same feature. Collected fuel was analyzed by NSR laboratory and 3^{rd} party analysis for fuel properties determination. In this analysis ASTM result indicate that LDPE waste plastic to fuel BTU value is 123631/gallon and on the other hand LDPE standard plastic to fuel BTU value is 128606/ gallon. LDPE standard plastic to fuel GC/MS analysis result indicate that hydrocarbon range is C₃-C₂₈ on the other hand LDPE waste plastic to fuel GC/MS analysis result indicate that hydrocarbon range is C₃-C₄₄. By using this technology could be solve environmental problems as well as energy problems and the thermal process can able to convert all LDPE waste plastic into liquid hydrocarbon fuel for internal combustion engines. Produced fuel could be use as a feed stock refinery or feed for electricity generation power plant. By using the technologies could be reduce some foreign oil dependency and boost up energy sector for near future.

Acknowledgement

The author acknowledges the support of Dr. Karin Kaufman, the founder and sole owner of Natural State Research, Inc (NSR). The authors also acknowledge the valuable contributions NSR laboratory team members during the preparation of this book.

References

[1] I. Hakki Metecan, Ahmet R. Ozkan, Rahim Isler, Jale Yanik, Mehmet Saglam, Mithat Yuksel, Naphtha derived from polyolefins, Fuel, Volume 84, Issue 5, March 2005, Pages 619–628

[2] Karishma Gobin, George Manos, Polymer degradation to fuels over microporous catalysts as a novel tertiary plastic recycling method, Polymer Degradation and Stability, Volume 83, Issue 2, February 2004, Pages 267–279

[3] N. Miskolczi, L. Bartha, G. Deak, B. Jover, Thermal degradation of municipal plastic waste for production of fuel-like hydrocarbons, Polymer Degradation and Stability, Volume 86, Issue 2, November 2004, Pages 357–366

[4] Ayhan Demirbas, Pyrolysis of municipal plastic wastes for recovery of gasoline-range hydrocarbons, Journal of Analytical and Applied Pyrolysis, Volume 72, Issue 1, August 2004, Pages 97–102

[5] Achilias DS, Roupakias C, Magalokonomosa P, Lappas AA, Antonakou EV. Chemical recycling of plastic wastes made from polyethylene (LDPE and HDPE) and polypropylene (PP), Journal of Hazardous Materials (2007); 149:536-542

[6] Muthaa NH, Patel M, Premnath V. Plastics materials flow analysis for India. Resources, Conservation and Recycling (2006); 47:222-244

[7] Buekens AG, Huang H. Catalytic plastics cracking for recovery of gasoline range hydrocarbons from municipal plastic wastes. Resources Conservation and Recycling (1998); 23:163-181

[8] Misklczia N, Barthaa L, Deak G, Jover B. Thermal degradation of municipal plastic waste for the production of fuel like hydrocarbons. Polymer Degradation and Stability (2004); 86:357-366

[9] Asanuma M, Ariyama T. Recycling of waste plastics in blast furnace. J Jpn Inst Energy (2004); 83(4):252 256

[10] Kato K, Fakuda k, Tachibana H. Waste plastics recycling technology using coke ovens. J Jpn Inst Energy (2004); 83(4):248-251.

[11] Steiner C, Kameda O, Oshita T, Sato T. EBARA's fluidized bed gasification: atmospheric 2x225 t/d for shredding residue recycling and two stage pressurized 30 t/d for ammonia synthesis from waste plastics. In: proceedings of 2nd international symposium on feedstock recycle of plastic and other innovative plastics recycling techniques. Ostend, Belgium; 8-11 September, (2002)

[12] Yoshioka T, Gause G, Eger C, Kamisky W, et all. A Pyrolysis of PETE in fluidized bed plant. Polym Degrad Stabil (2004); 86:499-504

[13] Kaminsky W, Schlesselmann B, Simon CM. Thermal degradation of mixed plastic waste to aromatic and gas. Polym Degrad Stabil (1996); 53:189-197

[14] Nigo A, Bhaskar T, Muto A, Sakata Y. Effect of natural and synthetic zeolites for the gasification of polyethylene and polypropylene, In; Proceeding of 3rd international symposium on feedstock recycle of plastics & other innovative plastics recycling techniques. Karlsruhe, Germany; 25-29 September, (2005), p. 395-401

[15] Kim YM, Kim S, Park YK, Kim JM, et all. Catalytic cracking of HDPE over MCM-48. In; proceeding of 3rd international symposium on feedstock recycle of plastics & other innovative plastics recycling techniques. Karlsruhe, Germany; 25-29 September, (2005), p. 333-339

[16] Aguado J, Serrano DP, Miguel GS, Escola JM, et all. Catalytic activity of zeolitic and mesostructured catalysts in the cracking of pure and waste polyolefins, J Anal Appl Pyrol (2007); 78:153-161

[17] Botas JA, Bravo M, Escola JM, Garcia P. Catalytic upgrading of higher 1-alkanes from polyethylene thermal cracking by modified wacker oxidation. J Mater Cycle Waste Manage (2006); 8:122-125

[18] Todd M. Kruse, Seth E. Levine, Hsi-Wu Wong, Eric Duoss, Andrew H. Lebovitz, John M. Torkelson, Linda J. Broadbelt, Binary mixture pyrolysis of polypropylene and polystyrene: A modeling and experimental study, Journal of Analytical and Applied Pyrolysis, Volume 73, Issue 2, June 2005, Pages 342–354

[19] C.G. Jung, A. Fontana in: J. Scheirs W. Kaminsky (Eds.) Feedstock Recycling and Pyrolysis of waste plastics, Wiley (2006), p. 252 (Chapter 10)

[20] Paul T. Williams* and Elizabeth A. Williams, Interaction of Plastics in Mixed-Plastics Pyrolysis, Energy & Fuels, 1999, 13 (1), pp 188–196, DOI: 10.1021/ef980163x

[21] Jong-Ryeol Kim, Jik-Hyun Yoon, Dae-Won Park, Catalytic recycling of the mixture of polypropylene and polystyrene, Polymer Degradation and Stability, Volume 76, Issue 1, 2002, Pages 61–67

[22] Kyong-Hwan Lee, Dae-Hyun Shin, Characteristics of liquid product from the pyrolysis of waste plastic mixture at low and high temperatures: Influence of lapse time of reaction, Waste Management, Volume 27, Issue 2, 2007, Pages 168–176

[23] Kyong-Hwan Lee, Dae-Hyun Shin, Young-Hwa Seo, Liquid-phase catalytic degradation of mixtures of waste high-density polyethylene and polystyrene over spent FCC catalyst. Effect of mixing proportions of reactants, Polymer Degradation and Stability, Volume 84, Issue 1, April 2004, Pages 123–127