STUDY OF LIQUID FRACTION SEPARATING IN PEROLISIS OF BIOLOGICAL ORGANIC WASTE

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Abstract. Nowadays, the need for clean drinking water is increasing year by year. There are a lot of adsorbents in water treatment, but in obtaining adsorbents, bicoals are one of the most popular methods as adsorbents with a high purification index and a cheap price. Liquid fractions, which were retained as condensates of the mixed gases released during the pyrolysis process, were studied. Our liquid fraction differs from other liquid fractions by the presence of combustible and fuel additives in a small amount. It is recommended to prevent the emission of waste gases into the atmosphere and to use the obtained secondary raw materials as secondary raw materials for paint and antifreeze production enterprises.

Keywords: perolysis, biochar, secondary raw materials, liquid fraction, partial pressure, mixture.

INTRODUCTION: Today, as the number of industrial enterprises increases, the needs of people are also increasing. Year by year, due to the construction of industrial enterprises, new lands are being developed. During the operation of industrial enterprises, the mixing of gas and liquid wastes into the atmosphere, soil, underground and surface water is increasing year by year. Disposal of industrial and domestic wastes causes anthropogenic problems if they are not controlled by separation of gaseous and liquid toxic substances. Based on the world experience, one of the most popular ways to prevent this is to use adsorbents, i.e. biochars. Biochars are obtained from agriculture, forestry, waste. At the same time, it has the ability to adsorb and clean the liquid and gas wastes coming out of the industrial enterprise, and it is distinguished by its low cost.

- Analysis of literature on the topic (Literature review). Humanity has used porous carbon materials (adsorbents) for many centuries. In the 18th century, coal's ability to purify various liquids and absorb some gases was discovered [1]. Until the beginning of the 20th century, carbon sorbents (mainly obtained from wood, bone, agricultural waste) were mainly used for liquid purification in the food industry and winemaking [1].

Today, biochars are secondary raw materials with various carbon content. For example: coconut and citrus fruit waste, wood, cellulose, [2-3] lignite and hard coal, [4] liquid and gaseous hydrocarbons, synthetic [5] polymers, agricultural and o Pyrolysis-activated adsorbents are obtained from vegetable waste [6], waste containing organic substances, bitumen, asphalt, used car tires, waste agricultural and industrial rubber, synthetic polymers.

Activated carbon carbons are a product of the processing of carbon-containing materials, characterized by a highly developed porous surface and high efficiency of soda processing. It is used in industry: it is highly effective in air and gas purification, waste water purification, technical and drinking water absorption, and industrial water purification of wastewater with medium toxic concentration [7].

- **RESEARCH METHODOLOGY** (Research Methodology). In the autumn season, one of the main traffic streets in Olmozor district is crowded with cars, and one of the non-stop traffic

streets is the perennial plant on the side of the intersection of Sagbon small village) Pinus eldarica - pine cones (i.e. spruce 'toys') we collected the spilled pine fir cones and washed it with technical water to remove dust and waste substances from the fir cones. Spilled spruce cone waste mainly contains fiber, fatty esters, phenolic compounds and gasoline vapors, and toxic gases settle in the composition of leaves and spruce cones during the filtering of gases and dust from the atmosphere. Spruce cones are used to reduce the percentage of ash content by washing the denatured substances contained in the washing target (Fig. 1). After we dry the waste of washed spruce cones under the sun for half a day, we put them in a pyrolysis furnace and start burning them in airless conditions (Fig. 2). Starting at 50 °S, the initial carbon was converted to bio-char in 300 minutes at a high temperature of 450-500 °S in an oxygen-free environment (Fig. 2). After more than 40 minutes during the process, the mixed gases coming out of our perolisis furnace are condensed and held in drops using a water cooler. It should be noted that when the temperature reaches 450-500 °S in 240 minutes, 700 ml of liquid fraction is released from 1,800 kg of mature leaves.



Figure 1. Spilled Christmas tree toys.



Figure 2. Our biochar inside our pyrolysis furnace.

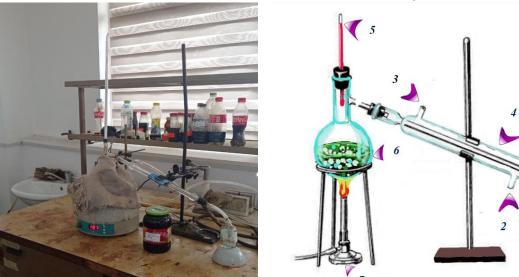


Fig. 3 Separation of volatile substances using the distillation method: collecting tank, 2- inlet of water cooler, 3- outlet of water cooler, 4- water cooler, 5-Thermometer, 6- process of driving liquid fraction, 7- tube heater

Evaporate the obtained liquid fraction at atmospheric pressure. Depending on the conditions, there are three types of driving: 1) driving at atmospheric pressure; 2) drive under low pressure, i.e. in a vacuum; 3) driving with water vapor. Using the first method, substances are boiled and brought to a vapor state. Then the vapors turn into liquid in the cooler. During driving, the amount of substances that boil at low temperatures is greater in the vapor phase than in the liquid phase. When such vapors condense, a fractional liquid with a new composition that boils at a low temperature is formed. A typical driving apparatus consists of a round flask with a side tube (Virs flask), a thermometer, a cooler, a flask, and a collecting flask. The boiling liquid vapor passes into the cooler and condenses (Fig. 3). Therefore, depending on the position of the side tube, the flask is selected according to the boiling temperature of the liquid. If the boiling point of the propellant is high, the side tube of the flask should be bent downwards.

If the boiling temperature is low, the side of the flask should be raised above the neck of the flask. The formula for determining the molecular mass of the substance and the partial pressure of water at the boiling temperature of the mixture using the following formula:

$$\frac{W_A}{W_{H20}} = \frac{P_A - M_A}{P_{H20} \cdot 18,02}$$

In this $\frac{W_A}{W_{H2O}}$ mass of matter and water, P_A , P_{H2O} the substance at the boiling point of the mixture and partial pressures of water, MA is the molecular mass of the substance

Any round-bottom flask with a dephlegmator can be used for driving. A mixture of two substances is divided into fractions depending on the boiling temperature. Two different fractions are usually collected during the first run:

In order to capture volatile chemicals in its composition, we collect the liquid fraction with a temperature of up to 95 oC using a distillation apparatus, and we also collect the curd residue. We can collect chemicals mixed with water up to a temperature of 110 oC with the help of a mechanical device, and we can also collect curd residue.

Using a flask heater, pour 250 ml of our liquid fraction into a heat-resistant 500 ml flask, install the first hand in the flask using a three-hole glass adapter, the second hand put it in the refrigerator, and the third hand 200 °S we install a temperature-resistant thermometer. We collect the volatile chemical substances released during the process in separate storage tanks depending on the temperature at which they are released.) we collect the liquid extracted from the process of perolysis of pine cones of Pinus eldarica at temperatures of 95 °S and 105 °S with the help of a thermometer. In 290 minutes of the process, we separated the liquid fractions and put them in a collecting container, and we sent the liquid fraction to infrared spectroscopy for inspection, and the solid part to a temperature of 120 °S was given for chromatography analysis.

– ANALYSIS AND RESULTS (Analysis and results). Two different liquid fractions obtained, i.e. volatile chemicals obtained at a temperature of 95 °S and liquid fraction obtained at a temperature of up to 120 °S, were checked using the mass selective "5977 AMSD" DRUGS_SKAN.A1 M method orcal of 5% in the "Agileht technoogies 7890NNetwork GC system" chromato-mass spectrometry. with a length of 30 m covered with phenylmethylsiloxane. using a capillary column, the temperature of the injector is 280 oS, and the temperature of the thermostat rises from 150 °S to 289 °S in 1 min. between 10 °S, the analysis time is 29.32 minutes. When we examine our liquid fraction using liquid chromatography, we can see in Table 1 below:

le erator quire strum mple I sc In	r : NH d : 27 ent : Name: fo :		00:33	using AcqMethod	l Drugs_SCAN_A G	_4_auto.M		
al Nur	mber: 7							
Abuno	lance	5.450			TIC: 53_1.D\d	ata.ms		
	3000000							
	2500000							
i i i	2000000							
kernind	1500000							
	1000000		6.678					
	500000	3.875 3.643 4.222.987 5.504 295 4.753 11 388 4.593 5.76	261 7.581	11.890 11.35(2.33	15.573 15.967 0			

N⁰	time taken for	Increase in	Chemical substance %	Minimum
	checking	temperature °S	and name from the	Accuracy
	(minutes).	over time	liquid fraction	Quality
				Percentage %
1	3:30	180	1,44% 3 Metil	
			siklogeksen	76 %
2	3:38	184	0,88% 1 Metil	
			siklogepten	58 %
3	3:50	187	4,46% 3 pentanon metil	53 %
4	3:56	189	9,24%	95 %
			2 furankarboksaldegid	
5	3:59	189	8,95 fenol	95 %
6	4:22	192	3,72%	
			2furankarboksilik	64 %
			kislota, tetrahidro furil	
			metil efir	
7	4:30	195	1,44%	
			2siklopenten, 2gidroksi-	76 %
			3metil	
8	4:45	196	2,90%	
			2 siklopenten, 3 dimetil	87 %
9	4:52	198	4,35%	98 %
			Fenol,2 metil	
10	5:5	201	5,53%	95 %
			kreozol,fenol,3-metil	

11	5:30	206	27,87%	97 %
			Fenol, 2- metoksi	
12	5:55	208	0,88%	84 %
			Fenol,3-dimetil	
13	6:26	212	0,93%	94 %
			Fenol,4-dimetil	
14	6:45	215	11,31%	97 %
			kreozol	
15	7:55	227	2,74	93 %
			fenol,4-etil,2-metoksi	
16			0,93%	
	11:30	235	Ftalik kislota, izobutil	80 %
			propil efir	
17			303%	
			2- benzendikarboksilik	95 %
	11:55	238	kislota,2- metilpropil	
			efir	
18	12:30	245	4,75%	99 %
			fenobarbital	
19	15:45	255	3,23%	99 %
			bis (2-etilheksil) ftalat	
20			1,42%	
			1,2-	
	15:59	259	siklogeksandikarboksilik	35 %
			kislota, siklobutil oktil	
			efiri	

In Table 1, the composition of our liquid fraction obtained up to 95 °S in the "Agileht technoogies 7890NNetwork GC system" chromato-mass spectrometry.

As can be seen in this table, substances with high concentrations of chemicals are: **CH3**, **C6H10O**, **C2H6O**, **C6H5OCH3**, **C9H6O5**, **C7H17N**, **C7H8O**, **C5H10O** organic binders are widely used in the varnish industry as an organic additive solution or solvent.

These organic substances disperse into the environment in volatile gas or liquid form, these substances are characterized by their volatility in open air. If the atmosphere, soil, underground and surface waters are poisoned very quickly.

At the same time, it is considered a chemical poison that affects human health through the respiratory tract, redness or tearing of the eyes, burns under the skin and on the skin, or as an allergen and death.

Our obtained organic chemicals are recommended for use in the food and paint industries.

rator : NHM	2024\Januar V Jan 2024 GCMS		using Ac	qMethod	Drugs_S	CAN_A G_	_4_auto	. м				
: Info : l Number: 9												
Abundance					TIC	53_3.D\da	ata.ms					
9000000	5.450	0										
8000000												
7000000	100.0											
6000000	in i a f e											
5000000												
4000000	3.851	6.678										
3000000	4.5730265											
2000000	4.963 4.222											
1000000	3.666 .504 4.7555.8	7.512	9.597		•	15.550						
lime>	4.00 6.	.00 8.00	10.00	12.00	14.00	16.00	18.00	20.00	22.00	24.00	26.00	28.00

№	time taken for	Increase in	Chemical substance %	Minimum
	checking	temperature °S	and name from the	Accuracy
	(minutes).	over time	liquid fraction	Quality
				Percentage %
1	3:20	184	3,70 %	
			2-butanon, 3-dimetil	72 %
2	3:38	186	4,90%	96 %
			2-siklopenten, 3-metil	
3	3:50	189	14,57%	
			fenol	95 %
4			4,87%	
	4:22	192	2-propenik kislota, 2-	64 %
			metil,(tetrahydro-2-	
			furanil) metal efir	
5	4:30	190	9,16%	
			2-siklopenten-1-bir, 2-	96 %
			gidroksi-3-metil	
6	4:35	193	1,42%	
			2-siklopenten-1-bir, 3-	90 %
			dimetil	
7	4:40	196	4,31%	
			fenol, 2-metil	98 %
8	5:20	202	10,56%	96 %
			p-kresol	
9	5:30	205	21,86%	

			fenol, 2-metoksi	97 %
10	5:45	207	2,11%	
			2-siklopenten-1-bir, 3-	97 %
			etil-2-gidroksi	
11	6:19	212	1,76%	
			fenol, 5-dimetil	96 %
12	6:30	215	1,71%	
			fenol, 3-etil	90 %
13	6:45	217	10,0%	
			kreozol	98 %
14	6:55	219	3,40%	
			katexol	96 %
15	7:30	225	3,27%	
			fenol, 4-etil, 2-metoksi	95 %
16	9:30	245	1,20%	
			1-(4-metiltiofenil), 2-	64 %
			propanon gomovanil	
			kislotasi	
17	15:35	255	1,20%	
			bis (2-etilheksil) ftalat	%

 Table 2 shows the composition of our liquid fraction obtained up to 110 °S in the "Agileht technoogies 7890NNetwork GC system" chromato-mass spectrometry.

As can be seen from this table, it was determined that there are organic chemicals with a high concentration in the liquid state. Our liquid separated from water at a temperature above 110 °S contains: C₆H₁₂, C₆H₆O, CH₃C₆H₄OH, C₆H₅OCH₃, C₇H₈O, C₁₆H₁₂O₃, these substances are distinguished by the category of danger, causing damage to the human nervous system, burning and flashing. It is recommended to use these organic substances as fuel or as a colorant in industry.

CONCLUSION (Conclusion). In this article, when we studied the technology of obtaining biochar by pyrolysis of organic-based waste, we studied the measures to prevent the poisoning of flora, water and soil by flying into the environment as a poisonous gas. In many cases, very little attention is paid to the retention of mixed gases during the perolysis process. The composition of the liquid fraction released during the perolysis process is not studied as a secondary chemical substance. Our next article focuses on obtaining liquid fuel from organic solid waste.

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