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BIOGAS Installation for Processing of ORGANIC BIOMASS

Shakhriyor Khamudillayevich Ergashev, Botir Egamberdiyevich Khayriddinov, Tulkin Amirovich Fayziyev

Assistant of the Department of "Thermal Power Engineering", Karshi engineering-economics institute, Karshi, Uzbekistan.

Doctor of Technical Sciences, Professor of the Department of "Thermal Power Engineering", Karshi engineeringeconomics institute, Karshi, Uzbekistan.

Candidate of technical sciences, Associate Professor of the Department of "Thermal Power Engineering", Karshi engineering-economics institute, Karshi, Uzbekistan.

ABSTRACT: In article are brought results heat physical features of the products of combustion biogas, got from biogas of the installation, built in subsidiary facilities under LLC Muborakneftgaz. To the analyzing energy efficiency of biogas formation, anaerobic processing requires certain temperature conditions and technological processes, preferably close to achieve the optimum process. The article shown factors determining the widespread adoption in the republic of technology of anaerobic decomposition of organic substances to produce biogas.

KEYWORDS: products of combustion, technological processes, degree of the soiling the air, heat of combustion, three atomic gases, factor of the excess of the air, dynamic viscosity, factor heat-conducting.

I.INTRODUCTION

The strategic directions of energy development in the Republic of Uzbekistan provide for the widespread use of non-traditional energy sources, including the energy of organic animal biomass. Calculations show that when processing organic biomass into biological gas, 4.2 times more energy can be produced annually than is produced at power plants in the Republic of Uzbekistan. Closely related to the problem of waste management is another - increasingly exacerbating - environmental protection, which also requires intensive and rational processing of organic biomass.

The use of renewable energy in the world is becoming increasingly important due to the fact that traditional sources of energy (coal, oil, natural gas) are limited, and their use for the production of heat and electricity causes great harm to the environment. In this regard, solar energy is becoming increasingly important, which can be used to produce environmentally friendly heat and electric energy [1,2,3,4].

The sun is a giant source of "clean" energy, not polluting the environment. Efficient use of solar energy can significantly reduce the consumption of natural resources. Climatic and weather conditions in the south of Uzbekistan create wide opportunities for the efficient use of solar energy in the Kashkakdarya region [2,3.4,5].

To achieve maximum efficiency of biogas formation, anaerobic processing requires certain temperature conditions and technological processes, preferably close to achieve the optimum process [6,7].

In the Republic, the main energy fuel is natural gas, in particular biogas, obtained from anaerobic decomposition of organic substances.

In the Republic, the most promising is the use of anaerobic decomposition of organic substances with biogas production.



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II. INSTALLATION DESCRIPTION

Biomass processing under anaerobic conditions is carried out in special, pressurized digesters (Fig. 1). Under the influence of methane-forming bacteria in an oxygen-free environment at a temperature of 30 ... 55 ° C, the biomass is fermented in the reactor with the formation of combustible gas - biogas, which is used for technological and domestic needs. About 1 mt of manure with 90% humidity can produce about 220 m³ of biogas with a calorific value of 28 ... 35 MJ/m³. The remains of the fermentation mass in the digesters are a highly digestible liquid highly concentrated organic fertilizer (humus) easily digestible by plants and devoid of pathogens and weed seeds [8,9].

The main elements of biogas plants are a digester (fermentation reactor) (2) and a gas holder (biogas storage tank) (3). The productivity and economic efficiency of the entire installation depends on the design of the digester.

Biogas plants do not require special expensive equipment. The payback period of these plants is 2 ... 4 years.



Fig. 1. Schematic diagram of a biogas plant for individual solar heating at muborakneftgaz llc:

1-organic waste; 2-digester; 3-gas holder; 4-solid waste; 5-boiler; 6-engine electric generator; 7-gas stove; 8-heating batteries; 9 and 10-lighting and electrical appliances; 11-heated digester.

The main factors determining the widespread adoption in the republic of technology of anaerobic decomposition of organic substances to produce biogas are:

1) high technological readiness and economic profitability; technological and operational simplicity of biogas technology;

2) for biogas technology, raw materials are available almost everywhere (organic industrial, agricultural and household waste, manure, etc.);

3) climatic conditions provide the maximum yield of commercial biogas; winter temperature conditions allow mesophilic fermentation to be ensured (at 30 ... 40 $^{\circ}$ C) at minimum biogas costs for heating in digesters, and in the summer time - heating costs disappear;

4) wide opportunities for the integrated use of biogas plants in conjunction with solar plants.

III.METHODS

Biogas usually contains 75 ... 81% methane. Other components include ethane, propane, butane (1.8 ... 6.2%) [8,9]. Heavier hydrocarbons are of great value as chemical raw materials. The composition of gaseous fuel is set in volume fractions and is written in the following form [6,10,11].

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$$\sum_{i=1}^{i=m} C_n H_{2n+2} + \sum_{i=1}^{i=m} C_n H_{2n} + H_2 + CO + H_2 S + O_2 + N_2 + CO_2 = 100\% .$$
(1)

Thermotechnical calculations are usually carried out for the dry composition of gaseous fuels. The calorific value of dry gas is determined by the volumetric composition in percent (%) and the known calorific value of the components.

Calorific value of gaseous fuel, kJ/m³.

$$Q_{in} = 358CH_4 + 638C_2H_6 + 913C_3H_8 + 1187C_4H_{10} + + 1461C_5H_{12} + 126,5CO + 107,5H_2 + 234H_2S;$$
(2)

the highest

inferior

$$Q_{th} = 398CH_4 + 698C_2H_6 + 993C_3H_8 + 1282C_4H_{10} + 1572C_5H_{12} + 126,5CO + 127,5H_2 + 254H_2S.$$
(2a)

With the known physicochemical composition of natural gas, the highest Q_{th} and lower Q_{in} of the calorific value, the thermal equivalent of natural gas $E = Q_{in}/29300$, are determined. For the resulting biogas is: $Q_{in} = 37600$... 39100 kJ/m³; $Q_{th} = 41800$... 44700 kJ/m³; E = 1.28 ... 1.34. As can be seen from the above values, biogas has a high calorific value. This is due to the fact that biogas has an increased content of heavy hydrocarbons. When these hydrocarbons are extracted (as chemical raw materials), the calorific value of biogas drops by 4 ... 8%.

For thermotechnical calculations, the value of the lower heat of combustion of the fuel is taken. The theoretical amount of air (m^3/m^3) required for the complete combustion of gas is determined by the formula [10.11,17,18,19,20]. $V_{air} = 0.0476 [0.5CO+0.5H_2+2CH_4+$

$$+1,5H_2S + \sum_{i=1}^{i=m} (m+n/4)C_mH_n - O_2].$$
(3)

For biogas is $V_{air} = 10, 4...11, 15 \text{ m}^3/\text{m}^3$.

In real conditions, air for burning fuel is supplied in a larger quantity compared to its theoretical amount V_{air} . The ratio of the actual amount of air V_{aair} supplied to the combustion device to the theoretically necessary V_{tair} is called the coefficient of excess air:

$$\alpha = V_{aair} / V_{air} \,. \tag{4}$$

The coefficient of excess air α depends on the design features of the combustion device, the type and method of burning fuel. Values of α vary in the range of $\alpha = 1.02 \dots 1.5$. The results of theoretical and experimental studies are shown in Fig. 2.

The products of complete combustion of fuel at $\alpha = 1$ contain: dry (non-condensing in the combustion device) triatomic gases R₂O (CO₂ and SO₂); H₂O - water vapor obtained by burning hydrogen; N₂ - fuel nitrogen and nitrogen in the theoretically necessary amount of air.

In addition, the composition of the products of fuel combustion includes water vapor resulting from the evaporation of moisture in the fuel, steam introduced into the furnace with moist air. For $\alpha > 1$, excess air (additional oxygen and nitrogen) will be present in the combustion products.

The biogas combustion products from LLC Muborakneftegaz do not contain sulfur dioxide SO_2 and therefore, $R_2O = CO_2$ is assumed.

The content of combustion products is presented: in percentage and share

$$RO_2 + N_2 + H_2O = 100\%$$
; $X_1 + X_2 + X_3 = 1$;
 $X_1 = N_2/100$; $X_2 = RO_2/100$; $X_3 = H_2O/100$. (5)

in volumetric content

$$V_{cp} = V_{RO} + V_N + V_{HO} \,. \tag{6}$$

Theoretical amount of nitrogen, consisting of air nitrogen and fuel nitrogen

$$V_N = 0,79 V_{air} + 0,8N_2/100; V_N = 8,5...8,86 \text{ m}^3/\text{m}^3.$$
 (7)



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The volume of dry triatomic gases



Fig. 2. The dependence of the volume of biogas obtained on the duration of fermentation of manure of small cattle.

Theoretical volume of water vapor generated by the combustion of hydrogen and contained in the air

$$V_{HO} = 0,01[H_2S + H_2 + 2CH_4 + \sum_{i=1}^{i=m} \frac{n}{2} C_m H_n + 0,124x_2] + 0,0161V_{air};$$
(9)

$$V_{HO} = 2,255...2,349 \text{ m}^3/\text{m}^3.$$
 (9a)

Volumetric content of fuel combustion products according to the formula (6)

$$V_{cp} = 11,862...12,357 \text{ m}^3/\text{m}^3.$$
 (10)

The percentage of components of the products of combustion of fuel

١

$$N_2 = 100 \times X_1 = 68,92...71,79\%; \quad RO_2 = 100 \times X_2 = 8,83...9,2\%;$$
$$H_2O = 100 \times X_3 = 19,01\%. \tag{11}$$

The theoretical fuel temperature is determined based on equations of the energy balance of the combustion process of 1 m^3 of fuel [10,11]

$$Q_{in} \frac{100 - q_3 - q_4 - q_6}{100 - q_4} + Q_6 = \sum_{i=1}^{i=3} (V_i C_i)(T_m - T_o);$$
(12)
$$\sum_{i=3}^{i=3} (V_i C_i) = V_{RO} C_{RO} + V_N C_N + V_{HO} C_{HO};$$
(12a)

where q_3 is the heat loss from chemical underburning,%; q_4 — heat loss from mechanical incompleteness of combustion,%; q_6 - heat loss with physical heat of slag, %; Q_h - heat introduced with hot air heated within the combustion device, kJ/m^3 ;

 C_{RO} , C_N , C_{HO} - specific volumetric heat capacity of the components of the products of fuel combustion, J/(m^3K).



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For gaseous fuels, the components of the heat balance q_3 , q_4 and q_6 are taken $q_3 = q_4 = q_6 = 0$. Thus, the combustion temperature of gaseous fuel in accordance with formula (12) will be determined by the expression:

$$T_m = (Q_{in} + Q_h) / \left[\sum_{i=1}^{l=3} (V_i C_i)\right] + T_o.$$
(13)

Molecular mass

$$M_{cp} = M_{RO} X_2 + M_N X_1 + M_{HO} X_3.$$
(14)

The values of M_{RO} , M_N , M_{HO} are shown in table 1.

$$M_{cp} = 27,571 \text{ kg/kmol.}$$
 (14a)

Mass heat capacity of combustion products

$$C_{cp} = M_{RO} X_2 C_{RO} / M_{cp} + M_N X_1 C_N / M_{cp} + M_{HO} X_3 C_{HO} / M_{cp} .$$
(15)

Values of C_{RO}, C_N, C_{HO} are given in table 1.

The density of dry combustion products, kg/m³

$$\rho_{cp}^{d} = 0,01[1,96\ CO_{2}+1,52\ H_{2}S+1,25\ N_{2}+1,43O_{2}+1,25CO+0,0899H_{2}+$$
$$+\Sigma(0,536m+0,045n)C_{m}H_{n}]; \quad \rho_{cp}^{d} = 0,84...0,875\ \text{kg/m}^{3}.$$
(16)

Density of wet fuel combustion products

$$\rho_{cp}^{w} = \frac{\rho_{cp}^{d} + \frac{x_{c}}{1000}}{1 + \frac{x_{c}}{804}}; \quad \rho_{cp}^{w} = 0.835...87 \text{ kg/m}^{3}.$$
(17)

Density of combustion products with temperature

$$\rho_{dcp=\frac{P_0M_{cp}}{R_0T_i}} \tag{18}$$

The thermal conductivity of the combustion products - gas mixture is determined by the formula [11]

$$\lambda_{cp} = \lambda_1 X_1 + \lambda_2 X_2 + \lambda_3 X_3 ; \qquad \lambda_{cp} = 0.0218 \text{ W/(m K)}.$$
 (19)

The dynamic viscosity of the combustion products is determined by Wilk [11]

$$\mu_{cp} = \frac{\mu_1}{1 + \frac{x_2}{x_1} \Phi_{1,2} + \frac{x_3}{x_1} \Phi_{1,3}} + \frac{\mu_2}{1 + \frac{x_1}{x_2} \Phi_{2,1} + \frac{x_3}{x_2} \Phi_{2,3}} + \frac{\mu_3}{1 + \frac{x_1}{x_3} \Phi_{3,1} + \frac{x_2}{x_3} \Phi_{3,2}}; \quad (20)$$

where $\Phi_{1,2} \dots \Phi_{3,2}$ are viscosity functions:

$$\Phi_{I,2} = \frac{\left[1 + (\mu_1 / \mu_2)^{1/2} (M_2 / M_1)^{1/4}\right]^2}{2\sqrt{2}(1 + M_1 / M_2)^{1/2}}; \ \Phi_{I,3} = \frac{\left[1 + (\mu_1 / \mu_3)^{1/2} (M_3 / M_1)^{1/4}\right]^2}{2\sqrt{2}(1 + M_1 / M_3)^{1/2}};$$

$$\Phi_{2,I} = \frac{\left[1 + (\mu_2 / \mu_1)^{1/2} (M_1 / M_2)^{1/4}\right]^2}{2\sqrt{2}(1 + M_2 / M_1)^{1/2}}; \ \Phi_{2,3} = \frac{\left[1 + (\mu_2 / \mu_3)^{1/2} (M_3 / M_2)^{1/4}\right]^2}{2\sqrt{2}(1 + M_2 / M_3)^{1/2}};$$



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$$\Phi_{3,l} = \frac{\left[1 + (\mu_3 / \mu_1)^{1/2} (M_1 / M_3)^{1/4}\right]^2}{2\sqrt{2} (1 + M_3 / M_1)^{1/2}}; \quad \Phi_{3,2} = \frac{\left[1 + (\mu_3 / \mu_1)^{1/2} (M_2 / M_3)^{1/4}\right]^2}{2\sqrt{2} (1 + M_3 / M_2)^{1/2}}.$$
 (20a)

According to formulas (20), (20a):

$$\mu_{cp} = 1,466 \times 10^{-5} \,\mathrm{Pa} \,\mathrm{c}. \tag{20b}$$

With the known values of C_{cp} , λcp , ρ_{cp} and μ_{cp} , the kinematic viscosity coefficients v_{cp} (m²/s) and thermal diffusivity a_{cp} (m^2/s) are determined, the number

Prandtl Pr_{cp}

where

$$v_{cp} = \mu_{cp} / \rho_{cp}; \quad a_{cp} = \lambda_{cp} / (\rho_{cp} C_{cp}); \quad Pr_{cp} = \mu_{cp} C_{cp} / \lambda_{cp}.$$
 (21)

The dependence of thermal conductivity λ_{cp} and dynamic viscosity μ_{cp} of combustion products on temperature is determined by the formula Sutherland [11.12]

$$\lambda_{cp} = \lambda_{cpo} A_i \; ; \; \mu_{cp} = \mu_{cpo} A_i \; ; \tag{22}$$

$$A_{i} = \frac{273,15 + B_{i}}{T_{oi} + B_{i}} \left(\frac{T_{oi}}{273,15}\right)^{3/2}; \quad T_{oi} = t_{oi} + 273,15.$$
(22a)

Bi values are given in table 1.

Thus, with known values of the thermophysical parameters of the products of fuel combustion, the heat of combustion in accordance with formula (13) is:

$$T_t = 1915...1995 \text{ K} \text{ or } t_t = 1642...1722 \,^{\circ}\text{C.}$$
 (23)

Table 1

Physical parameters of air and combustion products at t = 0 ⁰C CO_2 N_2 H_2O Parameter Air Carbon dioxide Nitrogen Water vapor M_i , kg/kmol 28,95 28 44 18 $100 \times X_i,\%$ 71,79 9,2 19,01 C_{pi} , kDj/(kg K) 0,9956+1,024 +0,8654 +1,833 ++0,00009299t+0,00008855t+0,0002443t+0,0003111t λ_{oi} , W/(m K) 0,02438 0,0243 0,0137 0,0162 $\mu_{oi} \times 10^5$, Pa c 1,7198 0.88 1,66 1,37 122 113 239.7 961 B_i

At combustion temperatures tt> 1650 °C, nitrogen oxides are formed (NO, NO₂; N₂O₄; N₂O₅), which, when combined with moisture and air, form nitric acid [12,13]. The formation of NO and NO₂ accelerates with increasing temperature,



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pressure and excess air. Reducing the formation of NO, NO₂ and their emissions into the atmosphere can be ensured by observing the condition $\alpha = 1$ and lowering the fuel combustion temperature $t_t < 1650$ °C. The best way to reduce the combustion temperature is to dilute the fuel with flue gas, i.e. recirculation use of fuel combustion products. Recirculation of flue gases during the combustion of natural gas can reduce the emission of NO and NO₂ by 60% [13.14,15].

CONCLUSION

The results of the thermophysical characteristics of biogas combustion products obtained from a biogas plant built in a subsidiary farm at Muborakneftegaz LLC are used to study the heat balances of furnace devices and boiler plants, to simulate heat and mass transfer of heat processes using the heat of fuel combustion products. Legend:

ai is the coefficient of thermal diffusivity, m^2 / s ; Ci — specific heat, J/(kg K); Mi - molecular weight, kg / kmol; Po - barometric pressure, Pa; Ro is the universal gas constant, J/(K kmol); t_i is the temperature, ° C; T_i is the absolute temperature, K; x_m - moisture content, kg/kg; Xi - volume fractions of the components of the coolant mixture; λ_i is the coefficient of thermal conductivity, W/(m K); μ_i is the dynamic viscosity coefficient, Pa s; v_i is the kinematic viscosity coefficient, m^2/s ; ρ_i is the density, kg/m³; Pr_i - Prandtl similarity criteria.

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