HIGH-TEMPERATURE GASIFICATION OF RDF WASTES AND MELTING OF FLY ASH OBTAINED FROM THE INCINERATION OF MUNICIPAL WASTES

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ABSTRACT. This paper presents innovative solutions for thermal processing of selected components of municipal wastes (so-called RDF wastes) using a low-ionized dependent plasma arc generated by a progressive and promising plasma reactor technology. The application of this process can transform hazardous wastes into inert wastes while significantly reducing the overall volume of wastes. The experimental results presented in this paper show the outputs achieved with thermal disposal of RDF wastes and ash from municipal wastes.

KEYWORDS: plasma, gasification, waste, fly ash, vitrified slag.

1. INTRODUCTION

Currently, about 80% of the world's primary energy production relates directly or indirectly to the combustion of fossil energy carriers, such as coal, oil and natural gas [1, 16]. Combustion of these carriers releases the carbon in the structure of the energy source in the form of CO_2 . When thermal processes are applied, excessive production of CO_2 from fossil fuel combustion increases the concentration of the gas product in the atmosphere. CO_2 contributes significantly to global warming, to the increase in the average year-round temperature in the global context, and to changes in climatic conditions [2]. In recent research and development, priority is being given to new technologies that will be able to re-use the energy potential of various kinds of wastes produced by mankind [3].

Wastes, as such, often contain several valuable raw materials with a relatively high energy and material potential. Recovery of raw materials and energy from wastes is of significant economic and also environmental importance. The elimination of wastes that cannot be recycled by other processes is also of fundamental importance.

This paper investigates selected combustible components of municipal wastes (so-called RDF wastes), which are usually incinerated or deposited in a landfill. Energy recovery in the combustion process produces large amounts of unwanted by-products in the form of slag, fly ash and gaseous emissions [1, 8, 9]. Due to the environmental problems associated with incineration, we are looking for new ways of targeted recovery and disposal of municipal wastes. One way that has been investigated for disposing of wastes is to gasify them and melt them in a plasma reactor.

2. Study area

Technologies using thermal plasma offer a unique ability to transfer extremely high performance from a small volume of plasma to the charge being processed [15]. The energy radiated from a source of thermal energy (a plasma column) and then transferred to batch particles decomposes the organic fraction of the waste, independently from the partial pressure of the oxygen. The part of the waste is reduced into simple gaseous compounds. Their presence in the synthesis gas is characterized by the composition of the batch and the boundary conditions of the gasification or of the high-temperature pyrolysis. Fig. 1 shows a block diagram representing the connection of the experimental laboratory plasma reactor that is used for laboratory investigations of the gasification and melting of wastes. A detailed description of the methodology for gasifying and melting the wastes is described in [1].

Decomposition of a heterogeneous batch of RDF at a temperature above 1000 °C, depending on the amount of available oxygen contained in the mixture, is characterized predominantly by the formation of certain components in the form of carbon monoxide, hydrogen and solid carbon [4]. Reducing the amount of untreated carbon in the system and increasing the amount in the synthesis gas in the form of CO can be achieved by introducing an oxidizing agent into the process. In order to increase the concentration of carbon monoxide and hydrogen in the synthesis gas, it is convenient to use oxidizing agents in the form of oxygen, carbon dioxide and, in some cases, water vapour [1]. The oxidation reactions are in Table 1.

The use of water vapour as an oxidizing agent provides an opportunity to increase the amount of hydrogen in the resulting mixture. However, the

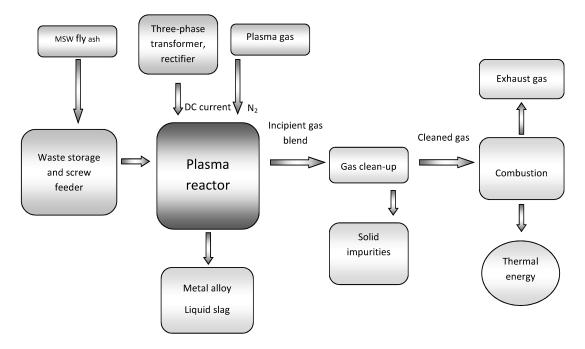


FIGURE 1. Block diagram representing the stages in the lab unit for gasifying and melting the wastes

Reaction	$\Delta H_{298}^0 \ (10^3 \mathrm{J mol}^{-1})$
$2C + O_2 \rightarrow 2CO$	-221.0
$\rm C+O_2\rightarrow \rm CO_2$	-393.5
$C + CO_2 \rightarrow 2CO$	+172.0
$\rm C + \rm H_2O \rightarrow \rm H_2 + \rm CO$	+131.3
$2C + 2H_2O \rightarrow 2CO + 2H_2O$	+90.1

TABLE 1. Oxidation reactions.

water gas reaction is endothermic. The endothermic reaction increases the specific energy consumption, based on the weight (kg) of the processed batch. It is therefore necessary to optimize the ratio of $C/O/H_2O$ in the mixture, and to optimize the assessment of the energy and the economic profitability of the process.

3. RAW MATERIALS USED FOR GASIFICATION AND THE PROPERTIES OF THE GAS

For the purposes of the experimental tests on plasma processing of RDF wastes, we used a sample that was obtained by separating the combustible components of municipal wastes (Fig. 2). This mixture of selected components of municipal waste (plastic, paper, tetrapacks, dendromass, fabrics, etc.) was compacted after separation and stored in bales before it was dispatched for its final use.

The basic physical and chemical properties that characterized the batch (Fig. 2) were as follows: 37.93 wt% of C, 4.63 wt% of H, 5.72 wt% of O, 12.33 wt% of Cl, 0.1 wt% of S, 0.05 wt% of N, 26.34 wt% of ash and 24.90 wt% of moisture. The slag-forming oxides, which form the major part of the



FIGURE 2. Sample of RDF wastes.

ash arising from RDF waste annealing of the samples at $815 \,^{\circ}$ C, are represented in the flowing proportions: $17.1 \, \text{wt\%}$ of SiO₂; $48.63 \, \text{wt\%}$ of CaO; $25.71 \, \text{wt\%}$ of Al₂O₃; $2.48 \, \text{wt\%}$ of MgO.

Given the high alkaline content of the calcium oxide in the ash, an acid flux is the most appropriate flux for reducing the melting point of the slag generated in the process of high-temperature gasification of the RDF waste. Silica flux was therefore used as a fluxing agent. The mixing ratio of the batch and the silica sand (96 wt% of SiO₂) was set at 10 : 1.1 (RDF : flux). This ratio is based on theoretical considerations, on the changes in the composition of the melting point of the net batch and the batch with the addition of the flux. The changes in the composition can be determined on the basis of the ternary diagram of the dominant slag-forming oxides (Fig. 3a). Adding a flux in a defined ratio reduces the melting point of the emerging slag to approximately 1350 °C(Fig. 3b).

Energy recovery of waste materials with a high

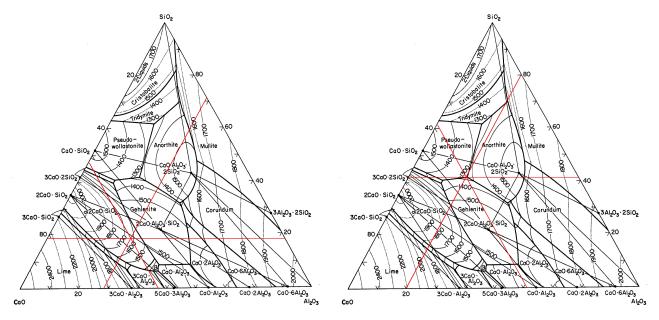


FIGURE 3. Ternary diagram CaO–SiO₂–Al₂O₃ for RDF wastes (a) and for RDF wastes + flux (b) [6].

Operating parameters		1st stage	2nd stage
Gasification temperature	(°C)	1450 - 1500	1450 - 1500
Duration of experiment	(s)	1800	1200
Batch mass	(kg)	2.4	2.4
Dosage speed	$(10^{-3}\mathrm{kgs^{-1}})$	1.333	2
Plasma gas flow (nitrogen)	$(10^{-3} \mathrm{m^3 s^{-1}})$	0.3133	0.1533
Oxygen flow	$(10^{-3} \mathrm{m^3 s^{-1}})$	0.1333	0.1333
Production of synthesis gas based on kg batch*	$(\mathrm{m}^3\mathrm{kg}^{-1})$	1.35	1.803

*parameter based on the nitrogen balance

TABLE 2. Basic operating parameters for high-temperature gasification of RDF wastes.

content of flammable matter further requires a highly efficient batch decomposition process, while ensuring maximum possible production of combustible gaseous components such as CO and H₂ [5]. Based on the given ratio of carbon and oxygen in the mixture of 3.158 : 1.741 (molar ratio), the theoretical total partial oxidation of the carbon atoms will be achieved by bringing a volume of oxygen corresponding to 1.417 mole per 100 g of the batch. This indicates that the process of gasification requires the additions of an oxidizer. The volume of oxidant from an external source in the form of pure oxygen is determined by calculation, taking into account results from previous experiments, with a flow rate of $0.133 \cdot 10^{-3} \, \text{m}^3 \, \text{s}^{-1}$.

4. EXPERIMENT

High-temperature gasification of RDF wastes was examined at 30 kVA on a plasma reactor with a dependent arc. A block diagram is shown in Fig. 1. The gasification process was carried out in the temperature range between 1450 and 1500 $^{\circ}$ C, for two different times of the experiment (two stages), at different speeds and with a different plasma gas flow. The basic parameters describing the gasification process

and the amount of synthesis gas that was produced per 1 kg of the batch are summarized Table 2.

In the first stage of batch gasification a sample of the syngas was taken. Its composition was as follows: 0.82 vol% of CH₄, 39.2 vol% of H₂, 0.04 vol% of O₂, 17.4 vol% of N₂, 3.3 vol% of CO₂, and 39.1 vol% of CO. In the next hour, the plasma reaction chamber of the plasma reactor was tempered to a temperature of about 1400 °C. During this time, it can be expected that the batch in the reservoir will dry up. The dosage batch restarted at a temperature of about 1447 °C. The ratio of the six most dominant gaseous components was as follows: 0.26 vol% of CH_4 , 50.9 vol% of H₂, 0.03 vol% of O₂, 25.4 vol% of N₂, 1.1 vol% of CO₂, and 43.5 vol% of CO. The presence of chlorine, sulfur, heavy metals and other elements that are used in plastics as flame retardants (e.g., bromine) were not analyzed during gasification of the RDF wastes and therefore they do not form a part of the discussion in this paper.

High percentages of the two most dominant flammable components of the synthesis gas (i.e., H_2 and CO), which represent 78–94 vol% of the total composition of the synthesis gas, indicate its potential energy utilization. The high ratio of H_2 and CO

Compound	RDF Poland	RDF Poland	MSW	MSW	MSW	RDF Ostrava	RDF Ostrava
Methane	0.01	0.16	8.59	4.77	2.54	1.84	0.72
Hydrogen	16.3	31.6	44.5	48.9	30.3	45.6	36.74
Oxygen	0.07	0.02	0.16	0.11	0.74	0.65	0
Nitrogen	10.6	8.63	6.03	6.61	15.9	17.2	18.26
CO_2	28.9	11.3	6.6	1.66	2.42	1.46	1.56
CO	44.1	48.3	32.5	37.1	47.3	32.9	42.62
Ethylene (C_2H_4)	0.001	0.004	0.97	0.49	0.52	0.27	0.08
Ethan (C_2H_6)	0.001	0.001	0.055	0.031	0.023	0.013	0.002
Ethyne (C_2H_2)	0.001	0.002	0.42	0.24	0.15	0.044	0.017
Amount of C_3 hydrocarbons	0.01	0.01	0.011	0.004	0.004	0	0
Amount of C_4 hydrocarbons	0.005	0.005	0.0099	0.004	0.005	0	0
Amount of C_5 – C_8 hydrocarbons	0.005	0.005	0.17	0.12	0.09	0.0026	0.006
Heating capacity $(MJ m^{-3})$	7.35	9.59	13.12	12.31	10.71	9.93	_
Estimated volume production $(m^3 kg^{-1})$	1.065	1.453	1.069	0.973	0.404	1.403	1.324

TABLE 3. The composition (in vol%) of the synthesis gas obtained by gasification of RDF wastes and municipal solid wastes (MSW).

in the gas mixture that was produced was also demonstrated in experiments similar to high-temperature gasification of batches with a substantial proportion of combustibles (Table 3). Recovery of the synthesis gas, however, must be preceded by a purification process in which impurities are removed from the gas in the form of particulate matter (PM), and also sulfur and chlorine compounds.

Melting and gasification of wastes with a substantial proportion organic fraction in the plasma reactor produces not only the synthesis gas (which can be used for producing electricity and heat), but also a separate liquid oxidic phase (slag) and also flue dust [6, 11]. The slag produced in the process is a gasification product. Its composition can be modified by adding fluxes, so that the product can be used e.g. in construction. This means that the plasma melting process and gasification of wastes, depending on the type of waste that is treated, is a technology that meets all the requirements for a so-called BAT technology. This technology enables an assessment of the material and energy content of various types of wastes.

The second kind of material that we investigated was fly ash from the incineration of municipal wastes (without separating the unburned part). This type of waste belongs to the category of so-called hazardous wastes. It contains a high concentration of undesirable metals (Hg, Cd, Pb, Zn, As, Sb, Sn, Cu, Ni, Cr, V and Ba and their soluble salts), persistent organic substances, dioxins and furans. For this reason according to Paragraph 21, Section 8 of Act number 223/2001 on the fly ash produced by the incineration of municipal wastes, the ash must be stabilized before it is finally disposed of [1].

So-called "cementing" has become a relatively

widespread technique for stabilizing fly ash. The fly ash is mixed with additives and with liquid reagents, and after the mixture hardens, the pollutants are fixed in the silicate matrix that has formed. The objective of the fly ash stabilization method is to ensure safer storage of the fly ash in landfills with a controlled regime.

A new, very promising way of environmentally friendly disposal of this hazardous waste, with subsequent use of final product, is vitrification in the plasma reactor [7,8,11,13,14]. During the vitrification process, the fly ash is melted together with the flux material at high temperature so as to create an oxidic glassy slag melt. This melt is then granulated or cast into moulds. The glassy nature of this product and its structure prevents leaching of toxic elements and their compounds that are trapped within (Table 4). Oxidic fly ash resulting from the process and consisting essentially of metals with a low boiling point (Hg, Cd, Zn, Pb and the like.), mechanical fly ash, alkali metal (Na, K) and chlorides of metals, is collected in the cleaning circuit of the synthesis gas. This fly ash could have suitable properties for further metallurgical processing [7].

Tests of fly ash vitrification, a product of energy recovery from municipal wastes in incinerators, have proved that the plasma reactor technology is suitable for recovering and disposing of wastes of this type, both from the point of view of the technological process and from the point of view of the environment. The melting experiments showed the inert character of the oxidic vitrified slag that was produced (Table 4). The volume reduction of the processed batch is between 45–65%. In addition, it is expected that the vitrified glass can be recovered.

Measured value		Aqueo	us extract	Landfill Class			
	Sample	Sample	Sample	Sample	SKIO*	SKNNO*	SKNO*
	fly ash 1	vitrified slag 1	fly ash 2	vitrified slag 2	Lea	chability class	
		stag 1		51ag 2	I.	II.	III.
pH		6.53	12.8	6.5	6 - 12	5.5 - 13	4 - 13.5
Arsenic (As)	0.013	< 0.001	< 0.001	< 0.001	0.05	0.2	2.5
Barium (Ba)	4.721	0.018	0.345	0.012	2	10	30
Cadmium (Cd)	< 0.01	< 0.0003	< 0.0003	< 0.002	0.004	0.1	0.5
Total Chrome (Cr)	5.51	0.003	0.318	0.002	0.05	1	7
Cooper (Cu)	0.861	0.019	0.006	0.012	0.2	5	10
Mercury (Hg)	69.6	< 0.0001	0.0002	< 0.0001	0.001	0.02	0.2
Molybdenum (Mo)	< 0.05	< 0.004	0.234	< 0.005	0.05	1	3
Nickel (Ni)	0.468	< 0.002	< 0.002	< 0.01	0.04	1	4
Lead (Pb)	8.612	< 0.005	0.6	< 0.01	0.05	1	5
Antimony (Sb)	0.176	< 0.001	< 0.001	< 0.001	0.006	0.07	0.5
Selenium (Se)	0.095	< 0.001	< 0.001	< 0.001	0.01	0.05	0.7
Zinc (Zn)	37.21	0.012	3.14	0.022	0.4	5	20
Chlorides	37900	< 1	3404	< 2	80^{\dagger}	1500	2500
Fluorides	32.92	< 0.1	3.8	0.03	1	15	50
Sulfates	12400	< 2	1255	< 5	$100^{+,\ddagger}$	2000	5000
Phenol index		< 0.03	< 0.01	< 0.002	0.1	50	100
CRL*	103470	34	9852	38	400	6000	10000
Eco-toxicity	positive	negative	undetermined	negative			

[†]If the determined value of CRL is $400 \,\mathrm{mg}\,\mathrm{l}^{-1}$, it is not necessary to establish values for chlorides and sulfates.

[‡]The value for sulphates may be exceeded, if the determined CRL value does not exceed $600 \text{ mg} \text{ l}^{-1}$.

*Note: CRL – entirely soluble matter; SKIO – landfill with inert waste; SKNNO – landfill with no dangerous matter.

TABLE 4. Results of the maceration test and norms for waste disposal (in $mg l^{-1}$).

5. CONCLUSION

High temperature gasification tests have been carried out on sorted components of municipal wastes (RDF wastes) in order to investigate the possibility of recovering energy from them. If the results of this experiment are confirmed, it will be possible to move away from present-day waste management methods, such as landfilling. This pilot test has confirmed that it is possible to recover energy from the primary product - the synthesis gas - which is a by-product of plasma waste treatment. In addition, this technology offers effective reduction of the volume of RDF. The synthesis gas that is produced is rich in hydrogen and in carbon monoxide. After all undesirable ingredients have been removed, the gases can be used for energy production. The production of a gas mixture rich in hydrogen and carbon monoxide was confirmed for other samples with basic properties similar to the processed RDF wastes.

RDF waste gasification tests in a plasma reactor further confirmed that it is technologically possible to gasify the types of waste with high content of flammable matter. The solid products (glassy slag, alloys, flue dust) considered as by-products of gasification may also, depending on their composition, have some economic and social added value.

The technology presented in this paper could also handle certain types of hazardous wastes. Tests that have already been carried out involving melting the ash in a plasma reactor have shown that the plasma reactor technology, in cooperation with the equipment installed in WtE processes, e.g. incineration, provides an efficient way to dispose of the ash and to acquire an inert intermediate product in the form of a glassy slag. The economic justification for the process is dependent on the relation between the energy consumption for the process itself, the limiting conditions for batch processing, and the amount and the value of the products that can be recovered through the process (glassy vitrified slag, flue dust and synthesis gas). Anticipated future tightening of the limits on the content of pollutants in fly ashes sent for landfill, and also increasing costs for waste landfilling, indicate that the plasma melting process is a very promising new technology for dealing with RDF wastes.

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