This paper presents the results from an experimental study of the thermal method applied for obtaining platinum-group metals (PGMs) by melting used autocatalysts in an 80 kV A plasma reactor, using suitable fluxes and a reducing agent. As the collector of noble metals, grey cast iron was used during the melting. The melt products were the alloy, inert vitrified slag, syngas and fly ash. In the alloy, when melting the catalyst based on cordierite, there was 99.27 % PGMs and 99.00 % in the case of Cr/Ni-strip catalysts. The loss of PGMs in the slag during the melting of cordierite catalysts was 0.34 % and for Cr/Ni catalysts, it was 0.26 %. A chemical analysis of the ash confirmed that in addition to the mechanically removed CaO, SiO₂, MgO, Al₂O₃, it also contained condensed vapours of Fe+PGMs. During the melting of the cordierite catalyst, the ash contained 0.40 % of the refined metals, while 0.74 % was the figure for the CrNi-strip catalyst. The synthesis gas from the process had a very low heating value (0.16 MJ m⁻³ or 0.24 MJ m⁻³).

Keywords: waste, catalyst, platinum-group metals, plasma technology

1 INTRODUCTION

Plasma technology for the processing and disposal of different types of waste is becoming a subject of interest for many waste producers. This technology can be used to process, e.g., municipal waste, ash from thermal power stations, dangerous asbestos-based waste, etc. Plasma technology is also used when processing various kinds of industrial waste. Noble metals can be recovered from the metallic waste containing them. The recoverability of noble metals from slag is, for example, the subject of several papers. Noble metals can also be obtained from slag using chemical-based methods (e.g., leaching). The basis for this is the change of a given phase to the liquid phase. The extraction of Cu, Ag, Au and Pd by leaching in nitric acid is described by references. The yield of silver using this method is up to 87 % and for copper, it is 98 %. The leaching of Cu, Zn and Pb from PCBs using electro-generated chlorine in a hydrochloric acid solution is described in reference.
cars – NOx, CO and unburnt hydrocarbons. This means that the aforesaid pollutants, with the help of the catalytic elements (Pt, Pd and Rh) found on the walls of the catalyst carrier, are reduced or oxidized to less harmful chemical compounds.

Platinum and palladium take the predominant share in the removal of carbon monoxide and hydrocarbons; the rhodium additive is intended to improve the activity of an autocatalyst as reduction reactions lower the content of nitrogen oxide. First, platinum with rhodium catalyses the splitting of nitrogen oxides into nitrogen and oxygen, and then platinum with palladium catalyses the oxidation of carbon monoxide (CO) and unburnt hydrocarbons with free oxygen into CO2 and water.

3 Material, Technology, Experimental Part

The autocatalysts were melted in the 80 kV A plasma reactor with a dependent electric arc. The plasma-forming nitrogen gas was fed into the electric arc through a hollow graphite electrode where it was transformed into a plasma at a temperature of the order of 103 K. At such a temperature, independent of the partial pressure of oxygen, the organic and partly inorganic components of the waste are decomposed into simple compounds according to the chemical reaction in Equation (1):11

\[
C_{x}H_{y}O_{z}Cl_{s}N_{a} = xCO_{g} + yH_{2(g)} + zN_{2(g)} + wH_{2}O(g) + rSO_{2(g)} + uHCl(g)
\]

The non-gaseous components of the waste are melted and two unmixable liquid phases are formed at the bottom of the plasma reactor – metallic alloy and slag. When melting autocatalysts, the gaseous product is a synthesis gas, which contains two combustible gases (hydrogen and carbon monoxide).

The plasma reactor15 was used to melt autocatalysts based on cordierite and autocatalysts based on Cr/Ni strips. A monolithic autocatalyst based on cordierite with a plastic structure (with a circular or elliptical cross-section) is made of ordinary cordierite with isomorphic iron (2FeO·2Al2O3·5SiO2) or ordinary cordierite with isomorphic magnesium (2MgO·2Al2O3·5SiO2). The cordierite is coated with the γ-Al2O3 gel film containing noble metals (platinum, palladium, iridium, rhodium). The average noble-metal amount in a catalyst is from 1.8 g to 2 g per 1 kg of ceramics with a Pt:Rh ratio of 5:1. The housing of the autocatalyst, the closure carrier, is made of stainless steel with a high amount of nickel and chromium (Figure 1).

An autocatalyst on a Cr/Ni strip is made from a chromium nickel strip wound into an S shape, on which there is a coating consisting of a thin film of γ-Al2O3 containing Pd, Pt, Rh (Figure 2). Autocatalysts of this type are also supplied as monolithic carriers (Figure 3) in the form of the plastic of ordinary cordierite with isomorphic Fe or Mg. The monolithic carriers contain a thin film of γ-Al2O3 containing mainly Pd.

Catalysts containing gold, silver, rhenium, rhodium, palladium, iridium or platinum have the waste code 16 08 01, while the ADR waste code is B 1150.

Catalysts made of ordinary cordierite with isomorphic Fe or Mg were ground, before the experiment, to a granularity of < 8 mm. The mix was of a grey colour. Autocatalysts on a Cr/Ni strip were mechanically separated, before the experiment, from the catalytic layer of γ-Al2O3 containing Pd, Pt, Rh with a granularity < 1 mm. The catalysts applied onto a Cr/Ni strip resulted in dust of a yellow-to-cream colour.

Before the experimental tests, the samples were chemically analysed. The largest components of the cordierite-based autocatalyst were oxides SiO2 (37.03 % mass fraction), Al2O3 (33.20 % mass fraction), MgO (9.80 % mass fraction) and Fe2O3 (1.14 % mass fraction). The autocatalyst based on Cr/Ni strips included Al2O3 (55.67 % mass fraction), CeO2 (12.00 % mass fraction), La2O3 (6.87 % mass fraction), SiO2 (2.69 % mass fraction) and ZrO2 (2.59 % mass fraction). In this melting process, the slag-forming elements are considered to be SiO2, Al2O3, Fe2O3 and MgO or CaO. The fractions gained in this way, containing noble metals were subsequently melted in the plasma reactor with an addition of fluxes (quick lime) and the slag had a relatively low melting point (below 1400 °C). At this temperature, the slag was saturated with a compound of Fe or Mg. The monolithic carriers contain a thin film of γ-Al2O3 containing mainly Pd.

Catalysts containing gold, silver, rhenium, rhodium, palladium, iridium or platinum have the waste code 16 08 01, while the ADR waste code is B 1150.

Catalysts made of ordinary cordierite with isomorphic Fe or Mg were ground, before the experiment, to a granularity of < 8 mm. The mix was of a grey colour. Autocatalysts on a Cr/Ni strip were mechanically separated, before the experiment, from the catalytic layer of γ-Al2O3 containing Pd, Pt, Rh with a granularity < 1 mm. The catalysts applied onto a Cr/Ni strip resulted in dust of a yellow-to-cream colour.

Before the experimental tests, the samples were chemically analysed. The largest components of the cordierite-based autocatalyst were oxides SiO2 (37.03 % mass fraction), Al2O3 (33.20 % mass fraction), MgO (9.80 % mass fraction) and Fe2O3 (1.14 % mass fraction). The autocatalyst based on Cr/Ni strips included Al2O3 (55.67 % mass fraction), CeO2 (12.00 % mass fraction), La2O3 (6.87 % mass fraction), SiO2 (2.69 % mass fraction) and ZrO2 (2.59 % mass fraction). In this melting process, the slag-forming elements are considered to be SiO2, Al2O3, Fe2O3 and MgO or CaO. The fractions gained in this way, containing noble metals were subsequently melted in the plasma reactor with an addition of fluxes (quick lime) and the slag had a relatively low melting point (below 1400 °C). At this temperature, the slag was saturated with a compound of Fe or Mg. The monolithic carriers contain a thin film of γ-Al2O3 containing mainly Pd.

Catalysts containing gold, silver, rhenium, rhodium, palladium, iridium or platinum have the waste code 16 08 01, while the ADR waste code is B 1150.

Catalysts made of ordinary cordierite with isomorphic Fe or Mg were ground, before the experiment, to a granularity of < 8 mm. The mix was of a grey colour. Autocatalysts on a Cr/Ni strip were mechanically separated, before the experiment, from the catalytic layer of γ-Al2O3 containing Pd, Pt, Rh with a granularity < 1 mm. The catalysts applied onto a Cr/Ni strip resulted in dust of a yellow-to-cream colour.

Before the experimental tests, the samples were chemically analysed. The largest components of the cordierite-based autocatalyst were oxides SiO2 (37.03 % mass fraction), Al2O3 (33.20 % mass fraction), MgO (9.80 % mass fraction) and Fe2O3 (1.14 % mass fraction). The autocatalyst based on Cr/Ni strips included Al2O3 (55.67 % mass fraction), CeO2 (12.00 % mass fraction), La2O3 (6.87 % mass fraction), SiO2 (2.69 % mass fraction) and ZrO2 (2.59 % mass fraction). In this melting process, the slag-forming elements are considered to be SiO2, Al2O3, Fe2O3 and MgO or CaO. The fractions gained in this way, containing noble metals were subsequently melted in the plasma reactor with an addition of fluxes (quick lime) and the slag had a relatively low melting point (below 1400 °C). At this temperature, the slag was saturated with a compound of Fe or Mg. The monolithic carriers contain a thin film of γ-Al2O3 containing mainly Pd.

Catalysts containing gold, silver, rhenium, rhodium, palladium, iridium or platinum have the waste code 16 08 01, while the ADR waste code is B 1150.
the xCaO·yAl2O3 type or pure aluminium oxide to prevent corrosion of the lining in the plasma reactor with a high Al2O3 content. When melting the autocatalysts, in addition to the flux in the plasma reactor, 15 kg of grey cast iron was added as a collector of noble metals in the form of cast-iron castings (for each melt). The composition of the charge for both autocatalysts is presented in Table 1. The table also shows the duration of the experiment, the average electricity consumption and the consumption of the nitrogen plasma-forming gas.

The amounts of the noble metals of interest and the chemical compositions of the alloys from the melting and gasification process for the two autocatalysts are given in Table 2. The chemical analysis shows that the alloy of grey cast iron and platinum-group metals is a metal collector. The Fe+PGM alloy is the raw material for the hydro-metallurgical production of the noble metals Pt, Pd and Rh.

The chemical composition of the slag from the melting and gasification of these autocatalysts is summarized in Table 3. The table shows that all the elements with a high affinity to oxygen, such as Al, Si, Mg and Ca, were concentrated in the slag.

In the slag, there were also non-sedimentary particles of Fe-containing noble metals, which represent mechanical losses of these metals in the slag (Figure 4). It was therefore necessary to recycle the slag by crushing, grinding and with the subsequent magnetic separation.

The results of the leaching tests for slag confirmed that this slag is inert and non-leachable. This means that no undesirable metals or elements leach from the slag and, therefore, it can be deposited in a landfill as other waste. This slag can also be used in civil engineering as a backfill or drainage material. The slag is also suitable for the production of glass fibres used in the production of heat- and electrical-insulation materials, as well as in the metallurgy of iron and steel.

The gas generated in the melting process was removed from the plasma reactor through a cyclone

<table>
<thead>
<tr>
<th>Catalyster</th>
<th>Slag-forming</th>
<th>Reduction agent</th>
<th>Time of experiment (min)</th>
<th>Consumption of el. energy (kWh)</th>
<th>Consumption of plasma gas (dm3 min–1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>cordierit</td>
<td>313.5</td>
<td>47.1</td>
<td>77.5</td>
<td>1863</td>
<td>77.20</td>
</tr>
<tr>
<td>Cr/Ni strips</td>
<td>90.0</td>
<td>77.5</td>
<td>77.5</td>
<td>1175</td>
<td>73.85</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*volume of gas media at a pressure of 101.325 kPa and temperature of 25 °C

<table>
<thead>
<tr>
<th>Catalyster</th>
<th>Slag-forming</th>
<th>Reduction agent</th>
<th>Time of experiment (min)</th>
<th>Consumption of el. energy (kWh)</th>
<th>Consumption of plasma gas (dm3 min–1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>cordierit</td>
<td>16.50</td>
<td>9.47</td>
<td>22.33</td>
<td>6.18</td>
<td>96.04</td>
</tr>
<tr>
<td>Cr/Ni strips</td>
<td>18.00</td>
<td>5.095</td>
<td>16.74</td>
<td>3.28</td>
<td>97.21</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Catalyster</th>
<th>Slag-forming</th>
<th>Reduction agent</th>
<th>Time of experiment (min)</th>
<th>Consumption of el. energy (kWh)</th>
<th>Consumption of plasma gas (dm3 min–1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>cordierit</td>
<td>353.50</td>
<td>16.77</td>
<td>12.47</td>
<td>30.08</td>
<td>37.85</td>
</tr>
<tr>
<td>Cr/Ni strips</td>
<td>187.30</td>
<td>42.00</td>
<td>0.33</td>
<td>3.86</td>
<td>47.37</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Figure 3: Carrier of the catalyster

Figure 4: Macrophotography of mechanically removed particles of iron with Pt, Pd and Rh in the slag matrix

Table 1: Basic data on the input raw materials for the experiment

Table 2: Masses of alloys and their chemical compositions

Table 3: Masses of slags and their chemical compositions

![Figure 3](image1.png)  
![Figure 4](image2.png)
separator where most of the mechanically removed ash was trapped. The chemical composition of the fly ash captured mainly in the cyclone separator, and subsequently in the other filtration devices, is summarized in Table 4. The chemical analysis shows that the ash (apart from the oxides of CaO, SiO$_2$, MgO, Al$_2$O$_3$ mechanically removed from the batch and the unreacted carbon added to the batch) also contained condensed vapour of Fe. The fly ash can be recycled without any major problems, thus increasing the share of the precious metals extracted from the waste being processed.

<table>
<thead>
<tr>
<th>catalyser</th>
<th>Fly ash (kg)</th>
<th>CaO (v%)</th>
<th>MgO (v%)</th>
<th>SiO$_2$ (v%)</th>
<th>Al$_2$O$_3$ (v%)</th>
<th>C$_{celk}$ (g)</th>
<th>Fe$_{celk}$ (g)</th>
<th>PGM</th>
</tr>
</thead>
<tbody>
<tr>
<td>cordierite</td>
<td>5.7</td>
<td>16.50</td>
<td>6.55</td>
<td>18.39</td>
<td>13.34</td>
<td>2.47</td>
<td>15.13</td>
<td>2.5</td>
</tr>
<tr>
<td>Cr/Ni strips</td>
<td>5.9</td>
<td>28.05</td>
<td>3.05</td>
<td>7.92</td>
<td>12.00</td>
<td>2.96</td>
<td>1.40</td>
<td>3.4</td>
</tr>
</tbody>
</table>

5 DISCUSSION OF THE RESULTS OBTAINED

From the point of view of the material and energy utilization of the waste, the plasma melting and gasification of autocatalysts in the 80 kVA plasma reactor can be evaluated as follows. Besides the alloy and slag, a by-product of the melting of autocatalysts is synthesis gas. It results from the reduction of the catalysts and the high-temperature pyrolytic decomposition of the organic portions of feedstocks. The gas was removed from the plasma reactor via a cyclone separator. Most of the mechanically removed ash was captured here. Subsequently, the synthesis gas was cooled from 1500–1600 °C to 160–170 °C in a hydrocyclone and then microparticles were collected in a sleeve filter. In the next step, the acidic constituents of the synthesis gas were neutralized in a countercurrent absorption column (pH = 9 of the neutralizing solution of NaOH). From the absorption column, the synthesis gas was fed into a system of coolers where, in the first step, it was cooled below the dew point of 60 °C to precipitate the moisture, with which the synthesis gas was saturated after the cooling in the hydrocyclone and the neutralization process. After the cooling, the gas was heated above the dew point and then burnt in a cogeneration unit with a CAPSTONE C65 microturbine.

The concentration of the main synthesis-gas components was continuously measured with a chromatograph during the melting. The results of the measurement of the proportions of H$_2$, CO and CO$_2$ in the gas, resulting from the melting of the cordierite autocatalyst, is shown in Figure 6. The chromatograph also searched for methane but it was not found in the synthesis gas. Of flammable gases, the syngas contained only hydrogen and carbon monoxide. The proportion of H$_2$ fluctuated between 0.79 % and 0.98 % (an average of 0.89 %), and the CO content ranged from 1.04 % to 1.27 % (an average of 1.15 %).

The average concentrations of the major components of the synthesis gas from the continuous chromatographic measurement of CO, CO$_2$, H$_2$ and N$_2$ during the melting of both types of catalysts are summarised in Table 5.

<table>
<thead>
<tr>
<th>catalyser</th>
<th>syngas (m$^3$·h$^{-1}$)</th>
<th>H$_2$ (v%)</th>
<th>CO (v%)</th>
<th>CO$_2$ (v%)</th>
<th>N$_2$ (v%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>cordierite</td>
<td>15.77</td>
<td>0.89</td>
<td>1.14</td>
<td>0.40</td>
<td>97.57</td>
</tr>
<tr>
<td>Cr/Ni strips</td>
<td>14.91</td>
<td>0.60</td>
<td>0.74</td>
<td>0.28</td>
<td>98.38</td>
</tr>
</tbody>
</table>

*volume of gas media at a pressure of 101.325 kPa and temperature of 25 °C

Because of the small proportion of flammable gases, the lower heating value (LHV) of the synthesis gas was very low. When processing the cordierite catalyst, the average LHV was around 0.24 MJ m$^{-3}$, and only 0.16 MJ m$^{-3}$ when melting the Cr/Ni-based catalyst. The energy benefit of the synthesis gas with the given LHV was negligible in the energy-production process. However, it is appropriate to use synthesis gas in the cogeneration unit.

A division of the platinum-group metals obtained from the alloy, slag and fly ash is shown in Table 6. It is
obvious that minimum amounts of these metals passed into the slag and fly ash. For the Cr/Ni-strip-based autocatalyst, it was 1.00 % and only 0.74 % for the cordierite autocatalyst.

<table>
<thead>
<tr>
<th>catalyst</th>
<th>alloy</th>
<th>slag</th>
<th>fly ash</th>
</tr>
</thead>
<tbody>
<tr>
<td>cordierite</td>
<td>99.27</td>
<td>0.33</td>
<td>0.40</td>
</tr>
<tr>
<td>Cr/Ni strips</td>
<td>99.00</td>
<td>0.26</td>
<td>0.74</td>
</tr>
</tbody>
</table>

The diffusion of metals into the plasma-reactor lining did not occur because the entire volume of the liquid phases (metal and slag) was kept in the graphite part of the reactor. This resulted in a minimum loss of the metals in the lining.

Cast iron was used as the collector of the noble metals. It is affordable, has high sorption capabilities with regard to platinum-group metals and magnetic properties that allow mechanically removed metal particles in slag to be separated. This means that the debris from the experimental melts were magnetically separated after grinding and milling, thereby increasing the yield of the metals of interest (Pt, Pd and Rh in Fe).

**6 CONCLUSION**

The liquidation of used autocatalysts in a plasma reactor can clearly reduce the environmental burden. Three major products can be obtained from the melting process: Fe+PGM alloy, slag and synthesis gas. The alloy of grey cast iron and platinum-group metals from the process of melting is the metal collector. The alloy is a raw material suitable for hydro-metallurgical production of noble metals Pt, Pd and Rh.

The slag with non-sedimentary Fe particles, which also contain noble metals, needs to be recycled to reduce the mechanical loss of metals. After crushing and grinding the slag, the iron from the platinum-group metals can be separated using magnets. The slag can also be used in ferrous metallurgy and civil engineering.

The synthesis gas from the melting of both types of autocatalysts has energy parameters that enable its use in the production of heat and electricity in a cogeneration unit. The share of the synthesis gas of the average natural-gas consumption is less than 0.5 %.

An accompanying feature of melting autocatalysts is fly ash in the synthesis gas that, in addition to the CaO, MgO, SiO₂ and Al₂O₃ oxides removed from the batch, also contains condensed vapour of Fe. It is therefore advisable to recycle fly ash, thereby additionally increasing the return of the metals of interest from the process of melting the catalysts.

Based on the results obtained by melting the autocatalysts, it can be stated that this waste can be further processed with the plasma technology into a product of an inert nature and the resulting melt products. This means that the presented elimination of the analysed waste by plasma melting is a waste-free BAT technology.

Previous experimental studies confirm that the prospective areas of use of plasma technologies will be the recycling and disposal of hazardous waste and low-level radioactive materials, and the refining of materials to high purity.

**Acknowledgements**

This research was realised with the support of the KEGA Grant Agency, nos. 003TUKE-4/2016 and SP2017/37-FMMIVŠBUO.

**7 REFERENCES**


