Concentration of Molybdenum Oxides and Salts in a Supercritical Water Medium

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Abstract: The results of an experimental study of the oxidation of industrial 5% water runoff of the propylene epoxidation process carried out under supercritical fluid conditions (T = 673-873 K, P = 25 MPa) on a continuous plant using oxygen as the oxidant are presented. The Chemical Oxygen Demand (COD), pH and mass spectra of the initial sample and reaction products were determined. The regularities of the efficiency of water runoff oxidation on the temperature and duration of the reaction are established. The elemental composition of the solid sediment is determined.

Key words: Molybdenum-containing water runoff, supercritical fluid conditions, oxidation, chemical oxygen demand, solid sediment, COD

INTRODUCTION

Nowadays, the share of chemical and petrochemical industries is growing rapidly as a result of which the question of the environmental component of these industries arises (Gumerov et al., 2012; Gayazova et al., 2013). The main problem of any chemical production is waste disposal (Kim and Kim, 2011; Chin et al., 1997; Vostrikov et al., 2017). If inert and low-hazard industrial wastes are mostly disposed of by landfill, then for toxic waste it is first necessary to carry out thermal neutralization or physico-chemical neutralization (Yang et al., 2017; Xu et al., 2015; Cao et al., 2011). One of the wastes, subjected to thermal neutralization is the waste of the process of hydroperoxide epoxidation of propylene for the joint production of styrene and propylene oxide which is industrially implemented at PTSC Nizhnekamsk Refeklkhim. A distinctive feature of this process is the use of molybdenum compounds soluble in the reaction mass which are used as a catalyst. During the reaction, the catalytic complex is destroyed and removed together with the reaction products in the form of a heavy fraction. After washing, the spent alkaline waste that is formed makes it difficult to separate the waste by distillation or rectification due to the presence of sodium salts and residual alkali which are emulsifiers. As a result, after thermal neutralization, the molybdenum complex is irretrievably lost with waste.

In the literature there is information on the extraction of molybdenum from alkaline waste. The research by Kirpichnikov et al. (1986) describes the return of still bottoms in the process for recycling molybdenum but due to the high corrosivity of the residues, it is not industrially implemented. The researchers carried out the isolation of molybdenum with organic peroxides by treating them with aqueous solutions of alkaline agents (Kaplan, 1970). But in this method there are large losses of target products, soluble in water and pollution of the aqueous phase which complicates the regeneration of the catalyst. A known method of molybdenum extraction with aqueous solutions of acetic, formic, propionic and butyric acids (Askins, 1893). However, as much as possible with this method, no more than 93% of the molybdic product can be extracted. In addition, the high volumetric ratio of the still bottom residue to the extractant, the need for temperature control, the duration of the process complicates the technology of catalyst extraction.

To solve the problem of the molybdenum extraction, the present research considers the possibility of using Supercritical Fluid (SCF) technologies. There are at least three process options. The first option is based on the use of the SCF extraction process. The second method involves the reaction of supercritical oxidation of waste in the aqueous medium. And the third option is based on the combined use of SCF extraction and Supercritical Water Oxidation (SCWO). As for SCF extraction, the researchers of this study have previously been able to obtain results on the extraction of molybdenum and the oxidizability of organic waste contained in the water run-off of the olefin epoxidation process (Kayumov et al., 2012).

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The present research is devoted to the study of the second solution to the problem. The use of water in the SCF state (t-374°C, p-22.1 MPa) allows you to dissolve any organic compounds containing difficult-to-wash substances as well as complex mixtures of organic substances. There are researches on the disposal of water run-off by the SCWO method (Psarov, 2006; Veriansyan and Jae-Duck, 2007).

It should also be noted that in most cases Hydrogen peroxide (H₂O₂) is used as an oxidant which is not profitable from the point of view of the process economic costs. The use of oxygen contained in the air is a promising solution and of course is also an urgent task of this study (Shin et al., 2009; Gong et al., 2008; Leusbrock et al., 2008; Lee et al., 2006).

**MATERIALS AND METHODS**

The studied runoff was the industrial water runoff of styrene and propylene oxide production process at PJSC Nizheukamskneftekhim of the following composition:

- Ethylbenzene C₆H₅CH₂CH₃-2.5% wt.
- Acetophenone CH₃COC₆H₄-1% wt.
- Methylphenylcarbinol C₆H₅CHOHCH₃-6.5% wt.
- Phenol C₆H₅OH-2.5% wt.
- Propylene glycol C₃H₈O₂-12% wt.
- Molybdenum Mo (pure metal)-0.2% wt.
- Water H₂O-40% wt.

As part of the research to develop the process of waste utilization in the SCF conditions of the reaction mixture, an original experimental continuous plant was used (Fig. 1) (Gumerov et al., 2017).

A distinctive feature of this plant is the use of a high-frequency induction heater (13). The advantage of induction heating is that the heat is completely transferred to the reaction mixture while achieving a high heating rate, increasing its uniformity. High concentration and precise localization of the energy of the electromagnetic field provides a short cycle and high performance of the chemical process.

The degree of contamination of wastewater and the oxidation reaction product was determined by the values of COD, obtained on the photometric analyzer “Expert-003-COD” with a thermal reactor for 26 samples in

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**Fig. 1:** Schematic diagram of the SCWO continuous plant: 1) High-pressure pump; 2) Capacity for the initial run-off, air; 3) Air compressor; 4) Valve; 5) Receiver; 6) Manometer; 7) Current isolating element; 8) Step-down transformer; 9) Heat exchanger for heating the feed mixture; 10) Temperature sensor (thermocouple); 11) Reactor; 12) Sediment collection chamber; 13) High frequency induction heater HF-30; 14) Fridge; 15) Wastewater collection tank and 16) Gas flowmeter.
accordance with GOST R 52708-2007 (Aetov et al., 2016). The essence of the COD measurement method was to treat water samples with sulfuric acid and potassium dichromate at a given temperature in the presence of silver sulfate (oxidation catalyst) and mercuric (II) sulfate, used to reduce the effect of chlorides. The COD value in a given concentration range was determined by measuring the optical density of the test solution at a given wavelength of 430 or 605 nm (depending on the measurement range) using the calibration dependence of the optical density of the solution on the COD value. The oxidation efficiency X was determined by Eq. 1:

\[
X = 1 - \frac{\text{COD}_{\text{in}}}{\text{COD}_{\text{oa}}}
\]  

(1)

Where:
- \(\text{COD}_{\text{oa}}\) = Chemical Oxygen Demand for the oxidized water runoff (mgO\(_2\)/L)
- \(\text{COD}_{\text{in}}\) = Chemical Oxygen Demand for the initial flow (mgO\(_2\)/L)

The excess oxygen used in the reaction was calculated using the following Eq. 2:

\[
O_2(\%) = \frac{[O_2]_{i} - [O_2]_{a}}{[O_2]_{i}} \times 100
\]

(2)

Where:
- \([O_2]_{i}\) = The actual concentration of oxygen supplied to the reactor (mmol/L)
- \([O_2]_{a}\) = The stoichiometric oxygen concentration based on the COD of waste water (mmol/L)

The residence time in the flow mode (\(\tau\)) was determined by Eq. 3:

\[
\tau = \frac{V}{Q_1 + Q_2} \times \frac{V_0}{V_i} \times 60
\]

(3)

Where:
- \(V\) = Reactor Volume (mL)
- \(V_0\) and \(V_i\) = The specific volumes of the initial runoff at room temperature, atmospheric pressure and under the reaction conditions (m\(^3\)/kg), respectively
- \(Q_1\) = Feed rate of the initial water run-off (mL/min)
- \(Q_2\) = Air flow rate (mL/min)

The inorganic residue with the target component, molybdenum was analyzed on a portable SI TITAN XRF spectrometer.

**RESULTS AND DISCUSSION**

The oxidation of 5% molybdenum-containing water runoff was carried out with an excess of atmospheric oxygen of 100-400% in the temperature range \(T = 673-873\) K at a pressure \(P = 25\) MPa in the dynamic mode. Figure 2 and 3 present the COD values depending on the process temperature and oxygen excess.

An increase in temperature and an excess of oxidant contribute to a greater oxidation of the waste. The minimum COD value equal to 713 mgO\(_2\)/L is achieved at the maximum temperature (\(T = 873\) K) and 400% excess oxygen. In this case, the estimated residence time of the mixture in the reactor is 230 sec. Obtaining such a result is comparable and even exceeds the COD values for process water (~ 1000 mgO\(_2\)/L) which indicates almost complete oxidability and recycling of industrial waste.

Table 1 shows the data on the efficiency of oxidation of the initial water runoff according to the results of the influence of the process temperature and excess oxygen.

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Fig. 2: Dependence of COD values (mgO\(_2\)/L) on the influence of the temperature of the SCWO process at \(P = 25\) MPa with an excess of oxygen: 1-100%, 2-200%, 3-300%, 4-400%

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Table 1: Efficiency of oxidation (X) of the industrial water runoff reaction products

<table>
<thead>
<tr>
<th>Temp (K)</th>
<th>Excess (O₂, %)</th>
<th>Efficiency of oxidation (X)</th>
<th>Excess (O₂, %)</th>
<th>Efficiency of oxidation (X)</th>
</tr>
</thead>
<tbody>
<tr>
<td>673</td>
<td>100</td>
<td>0.63</td>
<td>300</td>
<td>0.75</td>
</tr>
<tr>
<td>698</td>
<td>100</td>
<td>0.70</td>
<td>300</td>
<td>0.78</td>
</tr>
<tr>
<td>723</td>
<td>100</td>
<td>0.74</td>
<td>300</td>
<td>0.80</td>
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<tr>
<td>748</td>
<td>100</td>
<td>0.82</td>
<td>300</td>
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<td>773</td>
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<td>0.86</td>
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<td>0.87</td>
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<td>798</td>
<td>100</td>
<td>0.89</td>
<td>300</td>
<td>0.94</td>
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<td>823</td>
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<td>0.97</td>
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<td>848</td>
<td>100</td>
<td>0.97</td>
<td>300</td>
<td>0.97</td>
</tr>
<tr>
<td>873</td>
<td>100</td>
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<td>300</td>
<td>0.98</td>
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<td>673</td>
<td>200</td>
<td>0.72</td>
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<td>200</td>
<td>0.99</td>
<td>400</td>
<td>0.99</td>
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Table 2: Mass fractions of dry sediment components (given in pure form)

<table>
<thead>
<tr>
<th>Component</th>
<th>Zn</th>
<th>Fe₂O₃</th>
<th>Cr</th>
<th>Mo</th>
<th>Ag</th>
<th>Cu</th>
<th>SiO₂</th>
<th>Ni</th>
<th>Al₂O₃</th>
<th>CaO</th>
</tr>
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<tbody>
<tr>
<td>Content (%)</td>
<td>38.8</td>
<td>8.2</td>
<td>2.9</td>
<td>2.3</td>
<td>1.4</td>
<td>0.9</td>
<td>0.8</td>
<td>0.8</td>
<td>0.6</td>
<td>0.5</td>
</tr>
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</table>

Bold heading and value are significant

Table 3: The content of molybdenum in the liquid fraction

<table>
<thead>
<tr>
<th>Sample</th>
<th>Sample of the water run-off (Bottom fraction)</th>
<th>Sample of the water run-off (upper fraction)</th>
<th>Purified water run-off (with minimum COD)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mo (mg/L)</td>
<td>3626.00</td>
<td>2939.00</td>
<td>11.98</td>
</tr>
</tbody>
</table>

Fig. 3: Dependence of COD values (mgO₂/L) on excess oxygen (P = 25 MPa, T = 673-873 K)

The efficiency of oxidation is inversely proportional to the value of COD. With a decrease in COD, the efficiency of oxidation increases. As a result, 99% conversion is achieved at the maximum studied parameters of the process.

Table 2 shows the results of mass spectrometric analysis of dry sediment. A small amount of molybdenum (2.3%) is explained by a significant dilution of the industrial runoff itself (5% solution).

In addition to the obtained solid precipitate of the target component, some of the unremoved molybdenum complex remained in the liquid fraction which is clearly shown in Table 3.

CONCLUSION

A study of the process of supercritical water oxidation (T = 673-873 K, P = 25 MPa) of molybdenum-containing water run-off was carried out using air oxygen as an oxidant in a continuous plant. A significant decrease in the COD value of the reaction product in comparison with the COD of the initial water runoff has been established. The optimal conditions for the implementation of the reaction in the considered range of process parameters can take the temperature T = 873 K, the reaction time is 230 sec at a pressure of 25 MPa at which the minimum COD value = 713 mgO₂/L is reached. In the obtained samples of the inorganic residue, the presence of molybdenum was established in the amount of 2.3 wt.% which is the target product for subsequent extraction.

ACKNOWLEDGEMENT

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REFERENCES


