PURIFICATION PHASES OF ZINC SULPHATE SOLUTION OBTAINED FROM NEUTRAL LEACHING PROCESS OF ZINC CALCINE

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ABSTRACT: The aim of this paper is based on analyze of the zinc sulphate solution obtained from neutral leaching process of zinc calcine in "Trepca" and its purification from impurities. This solution besides zinc contains also other impurities such as (Cu, Cd, Co, Ni, As, Ge, Sb, Se, Te, Ag etc.), which are with different concentrations. The sulphate solution can released from metal impurities based on the fact that they have higher electro - chemical potential than zinc and, as such, during the refining process they separate and precipitate in metallic form. Treatment of solution with appropriate technological methods technologically is very important in continuation of the electrolysis process and is accompanied by economic effects, because of non-ferrous and rare metals benefit. In this paper are described the phases of the purification process including acidic purification, cooling, needed materials and equipments for the refining process. Also, special attention is dedicated to the factors that have influence in the process development such as concentration, temperature, time, compressed air, etc.

Key Words: Zinc, Sulphate solution, Purification process, Impurities,

INTRODUCTION

The zinc industry is one of the most important nonferrous industries in world, based in the demand for zinc which is in increase stage, therefore this effects in rapid development of its industry, and his influence in economy in general is very obvious. Zinc sulfide ores are the main source of zinc metal in the world, however, some zinc is produced from oxide-carbonate ores and different secondary resources such as zinc ash, zinc dross, flue dusts of electric arc furnace, leach residues, etc. Most of the zinc smelting production processes, which includes roasting, leaching, purification and electrolysis, are hydrometallurgical. For the zinc electrolysis process, the purity of the used zinc sulphate solution is an imperative. In the roast-leach process, very pure zinc sulphate solution must be prepared for zinc electrowinning, because the zinc ion is less noble than the hydrogen ion. The purity of electrolyte guarantees both obtaining high-grade cathode zinc deposit and reducing the amount of energy consumed for its procurement which considerably decreases the cost of zinc. The purity of the zinc sulphate solution, which is fed for electrolysis, is of great importance since the purity and structure of cathode zinc depend on it, as well as the coefficient of current utilization, the power consumption, the voltage of the tanks and other indices, which determine the zinc cost price (Boyanov et al., 2004). In the most common hydrometallurgical zinc process, ZnO-rich calcine is first produced from sulphide or oxide-carbonate concentrates and then leached with hot sulphuric acid solution. After liquid/solid separation, the pregnant solution is purified and electrowon for metallic zinc production (Turan et.al., 2004).

Therefore, the purification process is one of the most important unit processes in zinc hydrometallurgy. In hydrometallurgical zinc production, the purity of incoming solution subject to electrolysis is of great importance.

The aim of this paper is description of the phases of purification process of zinc sulphate solution gained from neutral leaching process in Trepca Company, including acidic purification and cooling along with specific equipment and materials. This paper as well described impurities which are contents
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of solution, and their negative effects in next process such is electrolysis process of zinc. Even a small amount of these metallic ions of impurities in the solution may hinder zinc deposition during the electrolysis process. The primary objective of the zinc sulphate solution purification process is to obtain an electrolyte solution that is suitable for zinc electrolysis. Since the zinc powder as material is expensive, its consumption should be minimized during the purification process. In previously published literatures, there have been studies related this issue (Fosnachtet al., 1983; Nyman et al., 1992). Most research has focused on experiments for determining the optimum purification conditions, such as solution temperature, zinc dust particle size, the quantity of zinc dust added, activators concentration and reaction time (Van Der Pas et al., 1996; Moghaddam et al., 2006).

Previously studies has been interested related this issue and published literatures regarding this topic (Ha M K, et.al., 2001; Li et.al., 2012). The membrane filter press (MFP), which is a common machine on solid-liquid separation, has advantages of low cost, high solid content and outstanding efficiency that has been widely used in various industries. The use of a MFP is found to be completely feasible and effective to leach and recover zinc from leaching residues of zinc calcine (Ru et al., 2015). Thereby, improving the technology of zinc production at the stages of solution purification and electrowinning of zinc is relevant. The negative influence of impurities can be reflected both in the decline of the current efficiency - it leads to an increase of the electricity consumption, and in the decrease of the cathode zinc quality due to the increased content of impurities (Sethurajan et al., 2017).

The metal impurities, which are more precious than zinc, can influence the purity of the cathode sediment through co-sedimentation and some of the impurities (cobalt, nickel and copper) facilitate the reverse dissolution of zinc (Muresan et al., 1996) by forming microgalvanic cells with it (Karoleva V., 1986). The harmful effect of these impurities is increased at higher temperatures and acidity of electrolyte, as well as at the simultaneous presence of Sb and Ge, Co and Cu and other elements in the solution. Table 1 shown in which form are metals present in the original zinc concentrate of Trepca company, respectively their mineralogical composition in Zn concentrate in percentage.

<table>
<thead>
<tr>
<th>Minerals</th>
<th>Total %</th>
<th>Zn %</th>
<th>Pb %</th>
<th>Cu %</th>
<th>Cd %</th>
<th>Fe %</th>
<th>CaO %</th>
<th>MgO %</th>
<th>SiO%</th>
<th>S %</th>
<th>Other %</th>
</tr>
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<tbody>
<tr>
<td>ZnS</td>
<td>72.14</td>
<td>48.44</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>23.70</td>
<td></td>
<td></td>
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<tr>
<td>PbS</td>
<td>2.56</td>
<td>2.22</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.34</td>
<td></td>
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</tr>
<tr>
<td>CuFeS₂</td>
<td>1.44</td>
<td>0.50</td>
<td>0.44</td>
<td>0.50</td>
<td></td>
<td></td>
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<tr>
<td>CdS</td>
<td>0.29</td>
<td>0.23</td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td>0.60</td>
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<tr>
<td>Fe₃S₄</td>
<td>0.67</td>
<td>0.31</td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td>0.36</td>
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<tr>
<td>FeS</td>
<td>15.99</td>
<td>10.17</td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td>5.82</td>
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<tr>
<td>CaO</td>
<td>0.46</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.46</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>MgO</td>
<td>0.26</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>0.26</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>SiO₂</td>
<td>2.12</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>2.12</td>
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<td></td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>4.07</td>
<td></td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>4.07</td>
</tr>
<tr>
<td>Total</td>
<td>100.00</td>
<td>48.44</td>
<td>2.22</td>
<td>0.50</td>
<td>0.23</td>
<td>10.92</td>
<td>0.46</td>
<td>0.26</td>
<td>2.12</td>
<td>30.78</td>
<td>4.07</td>
</tr>
</tbody>
</table>

**Table 1.** – The mineralogical composition of Zn concentrate

**BASICS OF PURIFICATION PROCESS OF ZINC SULPHATE SOLUTION**

The purification of zinc sulphate solution from the impurities (metals) is done by separation and cementation and their recovery in metallic forms is overviewed based on the fact that these metals have higher electro-chemical potentials than zinc. The operation factors that have the most profound effect on the process such as impurity concentration, temperature, time, etc. are analyzed, too. The presence of impurities in a zinc electrolyte solution is problematic for the electrowinning process. Impurities act as
catalysts for hydrogen evolution, which leads to a large drop in current efficiency. Therefore, adequate purification of the zinc electrolyte before electrowinning is essential (Dreher et al., 2001).

Purification of zinc sulphate solution process shown that this process play important role related the technological, economical and environmental advantages in industry. This process is dependant of numerous process parameters that need to be controlled carefully for future process such is electrolysis of zinc. A proper technique control can make more efficient this process in the industrial practice in all its steps by properly controlling of all process parameters including the quality of solution, adding of reagents, the speed, duration and temperature of the process (Friedrich et al., 2002).

Depending on the concentration and properties of impurities in the zinc sulphate various chemical and electrochemical refining methods can be used for their recuperation. One of the most known methods is the precipitation or cementation process in which metals of different electrochemical potential are separated by cementation (Kazanbaev et al., 2006). Cementation is the simplest and oldest hydrometallurgical process. It has been used in extractive metallurgy to recover valuable metals as well as to remove unwanted impurities (Pietrzyk, et al., 2017; Fugleberg, 2014).

Impurities in the zinc sulphate solutions, depending on their concentration and some common properties during the separation process, can be classified into the following groups:

I. Fe, Al, As, Sb, Ge, In, Ga, Si
II. Cu, Cd, Co, Ni, Ta
III. Cl, Fl,
IV. K, Na, Mg, Mn

The zinc sulphate solution is considered to have a high refined degree if it is a very clear solution and has the following characteristics (Purification process, 1982):

- solid substances <0.5 g/l,
- Fe²⁺ <5 mg/l,
- total Fe <15 mg/l,
- Sb + As <1 mg/l and
- pH = 5–5.5 (Purification process, 1982)

Various methods such as hydrolyse, cementation, catalyst addition etc. can be used to selectively recover the impurities in these groups. These methods have been subject of several research articles and patents (Nelson et al., 2000), but their application in the industrial practice has still difficulties especially in the selection of optimum factors that influence these processes. Trepca company, as per design capacity, can produce about 90,000 t/year zinc sulphate solutions obtained as a by-product of neutral leaching of zinc calcine (Purification Process, 1982). The overall technological flowsheet of zinc production in the Trepca smelter is described in Figure 1.
Pyrometallurgical and hydrometallurgical routes or their combination can be employed for treating secondary materials. The hydrometallurgical processes are regarded as more eco-friendly for treating such materials having a low zinc content (Ha M K, et al., 2001).

In the hydrometallurgical process, ores containing zinc are pretreated in hot conditions and then dissolved using sulfuric acids into a zinc sulphate solution. This solution is then purified from various impurity metals such as cobalt, nickel, cadmium, antimony etc., by cementation (Lew, 1994). These metal ions must be removed as they would disturb the following electrolysis of zinc. Cobalt and related metals such as nickel cause damage to the cathode used in the electrolysis process as well as contributing to re-dissolution of zinc (Boyanov et al., 2004).

In the hydrometallurgical zinc production process, metallic zinc is produced by reducing it from aqueous solution. The essential part of the process is the purification of zinc sulphate solution, where all elements nobler than zinc are removed by deposition from the solution. Controller performance is highly dependent on the reliability of information which is received from measuring devices and from reference measurements. Measured process data is inherently inaccurate and these measurement errors can be caused by random noise, deteriorating, malfunctioning or wrongly calibrated instruments. The correlation analysis used here applies conversely in process control. The results have been exploited in
the process control of solution purification. In the case of the cobalt removal process, it was necessary to get the continuous solution purification process working reliably and with a high enough deposition efficiency (Näsi, 2007). As zinc consumption is gradually increasing all the time in the world, many zinc plants are looking for possible expansions. Their preliminary profitability calculations can show that a possible total expansion is not economical, when they are looking to expand the whole roasting/acid plant – leaching – solution purification – electrowinning – casting route, as the possible market price for the sulphuric acid is low or the sulphuric acid market is too far away. Based on these facts, their economical study often indicates that implementing a zinc concentrate direct leaching option could be more favourable (Svens, 2012).

Due to the relatively low concentration limits, the preparatory purification is of great importance (Han et al., 2016). Cementation is widely used in hydrometallurgical zinc refining to purify zinc sulphate solutions from impurities. The purification is achieved through electrochemical reactions causing the impurity metals to precipitate on the surface of zinc dust. These reactions are commonly known as cementation reactions. A cementation reaction can be described as electrochemical precipitation where the most noble metal ions in a solution are reduced to solid metal through deposition on the surface of a less noble metal which is oxidized (Sethurajan et al., 2017). In the case of zinc purification, the impurity metals are the nobler metals that are to be reduced while zinc, often in the form of zinc dust, is the less noble metal providing the electrons (Lew, 1994; Free, 2013).

There are many companies over the world that employed more or less similar methods or some other variations within the purification process of zinc sulphate solution obtained from neutral leaching process of zinc calcine.

**PURIFICATION PHASES OF SULPHATE SOLUTION OBTAINED FROM THE LEACHING PROCESS OF ZINC CALCINE**

Based on the analysis presented in the technical documentation of smelter we have these average concentrations of impurities in sulphate solution obtained during the leaching process are given in Table 2.

<table>
<thead>
<tr>
<th>Elements</th>
<th>Concentration (mg/l)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cu</td>
<td>0.2-1.2</td>
</tr>
<tr>
<td>Cd</td>
<td>0.2-1.2</td>
</tr>
<tr>
<td>Co</td>
<td>1.0-10</td>
</tr>
<tr>
<td>Ni</td>
<td>20-50</td>
</tr>
<tr>
<td>As</td>
<td>0.1-0.5</td>
</tr>
<tr>
<td>Ge</td>
<td>0.1-0.5</td>
</tr>
<tr>
<td>Sb</td>
<td>0.1-0.5</td>
</tr>
<tr>
<td>Se</td>
<td>0.1-0.5</td>
</tr>
<tr>
<td>Te</td>
<td>0.1-0.5</td>
</tr>
<tr>
<td>Ag</td>
<td>0.1-0.5</td>
</tr>
</tbody>
</table>

The purpose of the purification of the sulphate solution obtained from leaching process of zinc calcine, which is the subject of this study, is the process with aim to release the sulphate solutions from impurities and produce a highly refined solution in order that the subsequent process of electrolysis, achieves the best possible quality of zinc and make an efficient use of electricity. The zinc sulphate solution is considered to have a high refined degree if it is a very clear solution and has the following characteristics: solid substances <0.5 g/l, Fe^{2+} <5 mg/l, total Fe <15 mg/l, Sb + As <1 mg/l and pH = 5–5.5.
Refining of the zinc sulphate in Trepca smelter is carried out in two phases: cold and hot. The refined solution is subsequently mixed with acid, cooled and sent for electrolysis. The residues of the process are treated to increase the recovery coefficient of cadmium and zinc.

**Cold Refining**

The cold refining is the first phase of the purification process, that enable selective multi-tank sedimentation of impurities such as Cu, Cd, Tl and part of Ni (approximately 12 mg/l). This is carried out through the added zinc powder. The amount of zinc powder needed for the process, is calculated according to the following equation:

\[
Zn(g/l) = 0.5(g/l) + 1.2Cu(g/l) + 1.5Cd(g/l)
\]  

(1)

The zinc powder is used not only in the first tank but also and in the second tank, in order to precipitate any Cd amount which may have accidentally penetrated. Generally, in the cold refining stage the precipitation of Cu and 95% of Cd is achieved as well as part of As, Sb, Tl, Ge and Ni. Duration time of reaction is an important factor in this phase and normally is about 45 mins but longer times especially longer than an hour and a half can cause oxidization of Cd which can pass again in the solution. To increase the speed of filtration the solution needs to be heated in the last tank. Precipitation of the Co and especially of Ni, is done successfully by adding of As and Sb. However if the content of As and Sb in the neutral solution is high - an indication that As and Sb were not removed at the right degree during neutral leaching - Co and Ni can partly precipitate thus increasing the undesirable quantities of Co and Ni in CdSO₄ solution. The filtrate is sent to hot refining stage and the sedimentation is sent for further processing of Cu and Cd (Purification Process, 1982).

**Hot Refining**

Elements which have not precipitated in the first phase such as Ni, Co, Sb, Ge as well and other impurities can be removed during this second phase. This is done by adding zinc powder and antimony oxide (Sb₂O₃) and increasing the temperature up to 90 °C. The amount of zinc powder needed for process is calculated according to the following equation:

\[
Zn(g/l) = 1(g/l) + 100Co(g/l) + 50Ni(g/l)
\]  

(2)

Zinc powder particles must be smaller than 75 µm, in order to speed up their solubility and to avoid using excessive amount of zinc powder. Sb₂O₃ is used as addition because along with the added zinc metal powder it creates a catalyst that accelerates the solubility of zinc and the precipitation of Co and Ni. However, the amount of Sb₂O₃ added to the solution should be limited because a larger than necessary amount causes reversed dissolution of Co precipitate. Furthermore the excess of Sb does not fully precipitate and this has an impact on the process of electrolysis.

In this stage Pb is also added into the solution in the form of oxide or sulphate in order to:

- reduce the repetitive tendency of dissolution of Co precipitate,
- reduce the formation of basic zinc sulphate,
- improve the conditions to remove the Tl.

The solution should be kept in the reservoir a few hours, then empties as necessary because if the solution stays for longer periods, impurities such as Co dissolve and Zn gets oxidized which makes the
filtration process difficult. Normally the duration of chemical reaction in this phase is 3 hours and the temperature 90–95 °C. The solution is filtered and the filtrate is checked for quality. The process is considered cleaned if the filtrate contains approximately 0.15 mg/l Co, and 0.01 mg/l Sb. After filtration when the temperature is about 80 °C, H₂SO₄ electrolyte is added into the cleaned solution until it reaches a concentration of 2–3 g/l free H₂SO₄. This is done in order to avoid the closing of the pipes. The solution is then cooled to near 30 °C and sent to electrolysis. The process continues with re-pulping of Co, Ni, Cu and Cd, in order to increase the yield of Zn. Filtering residues are then sent for processing of Co/Ni (Purification Process, 1982).

The process of refining of sulphate solutions and recovering of valuable materials from it is a complex process whose normal running and its efficiency depends on numerous process parameters. These parameters in the Trepca process include the content of solid substances, Fe²⁺, Fe total and Sb + As concentration of pH of solution, added amount and the size of particles of zinc powder during cold and hot refining, processing time and temperature, the speed of filtration, lead quantity, mixing with the electrolyte.

CONCLUSION

Purification of zinc sulphate solution process shown that this process play important role related the technological, economical and environmental advantages in industry. This process is dependant of numerous process parameters that need to be controlled carefully for future process such is electrolysis of zinc. A proper technique control can make more efficient this process in the industrial practice in all its steps by properly controlling of all process parameters including the quality of solution, adding of reagents, the speed, duration and temperature of the process. This purification way the of sulphate solution can maximize the economic benefits by efficiently separating of valuable metals that are present in the solution including Cu, Cd, Co, Ni, As, Ge, Sb, Se and Te. Based on the above data, it is evident that the selective recovery of these metals from these solutions represents a significant economic added value, and from the environmental point of view this eliminates the need of unnecessary and sometimes risky land filling with valuable metals. In this point of view, in order to revive and improve operation practices as per design capacity in the Trepca Company, a new project needs to be undertaken, in order to increase efficiency of the process through proper control and optimization. The drawbacks of the conventional zinc solution purification process are an economical and technical incentive to search for ways to improve the performance of the purification systems.

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