

Investigation of jarosite process tailing waste by means of raman and infrared spectroscopy

Untersuchung des Abfallerzes des Jarosit-Prozesses durch Raman- und Infrarot-Spektroskopie

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The aim of this work was to identify and characterize three samples of jarosite process tailing waste in Mitrovica, Kosovo, using Raman and Fourier transform infrared (FTIR) spectroscopy. The identification is made based on the assignment of bands in the Raman and FTIR spectra. Both Raman and FTIR spectra show the fundamental stretching and bending vibration mode of SO_4^{2-} , OH^- and NH_4^+ groups.

The results obtained by the means of Raman and infrared spectroscopy are compared with cited reference data in order to sum the analysis of vibrational spectra.

Keywords: Jarosite / Waste / Raman / Infrared / Spectroscopy

Schlüsselwörter: Jarosit / Abfall / Raman / Infrarot / Spektroskopie

1 Introduction

Worldwide, jarosite process is the most widely used techniques in the zinc industry. During this process Fe (III) compound of the type $\text{MFe}_3(\text{SO}_4)_2(\text{OH})_6$, where M is usually monovalent Na^+ , K^+ , NH_4^+ , H_3O^+ , Ag^+ or divalent cation such as Pb^{2+} is precipitating, and a huge amount of jarosite waste is released [1]. This jarosite waste contains high concentrations of toxic metals that may become a major environmental and ecological problem. This issue became a dramatic in Mitrovica (Kosovo) were 35 ha of the town has been covered by zinc smelter waste, situated very close to the residential area. Raman and infrared vibrational spectroscopy techniques are regarded as powerful techniques to characterise the minerals, natural jarosite and the jarosite released from the zinc smelter process. The recent study presents the means of these techniques to characterise the jarosite waste in Kosovo.

In general, the methods of the vibrational spectroscopy provide information about stretching and bending vibration that are coupled with variations in interatomic distances and angles. These methods can be used for identification of compounds as well. Raman and Infrared (FTIR) spectroscopy as complementary techniques are suitable for analysis of compounds of jarosite

mineral groups, either natural or synthetic, published in numerous studies [1–10]. Methods are sensitive to vibrational bonding modes of most functional groups in jarosite molecules (OH^- , H_2O , H_3O^+ , SO_4^{2-} and Fe-O) that are particularly suitable for analysis of sulphates [2].

Sulphate anion in aqueous solution has T_d symmetry which is characterized with four Raman bands originating from: symmetric (ν_1) and asymmetric (ν_3) stretching modes and symmetric (ν_2) and asymmetric (ν_4) bending modes [1, 3]. However, symmetry of sulphate anion in crystal structures is reduced (C_{3v}) leading to splitting of these modes. In accordance with the theory group in the spectra of jarosite six basic sulphate bands should occur [1, 3, 4]. Raman and FTIR spectra, except for the determination of group of minerals, can be used for identification of minerals and type of interlayer cation (M^+) as well. In his study, Sasaki et al. reported about the relationship between the type of cation (M^+) and the position of some bands, as a consequence of changes in c unit cell parameter (c parameter increases with increasing of ionic radius) [1]. They have shown that changes in ionic radius of cation M^+ reflects on the position of the vibrational modes of sulphate anions $\nu_1(\text{SO}_4^{2-})$ and $\nu_3(\text{SO}_4^{2-})$, whose wavenumber decreases with increasing parameter c . $\nu_2(\text{SO}_4^{2-})$ and $\nu_4(\text{SO}_4^{2-})$ bending modes are independent of the parameter c [1]. The same authors cite similar dependence for bands position of three Fe-O vibrational modes, which decreases with increasing of parameter c . For Fe-O bands this effect can be used only in Raman spectra, because in FTIR spectra Fe-O bands are present below 600 cm^{-1} that is out of detection limit of most standard FTIR detector devices [1, 3]. Other authors, that studied the stability of the synthetic Na-, K- and NH_4 -jarosite, emphasized that the highest sensitivity to different interlayer cations have positions of $\nu(\text{OH})$ bands (e. g. 3392 cm^{-1} in Natrojarosite and 3434 cm^{-1} in Ammonium jarosite). Identification of jarosite, based on the bands wavenumber

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can be useless if the sample contains a mixture of jarosite with different cations [5].

Besides the bands wavenumber, type of jarosite can be identified using the specific bands that are present only in certain types of mineral e.g. Ammonium jarosite. In the FTIR spectrum of Ammonium jarosite characteristic bands should appear at $\sim 1420\text{ cm}^{-1}$ (H-N-H bending vibrations), ~ 3340 and 3220 cm^{-1} (vibrations of ammonium ion) [1, 5, 6]. Raman spectrum bands of ammonium ion can be visible as a broad peak at 3210 cm^{-1} and a weak peak at 1664 cm^{-1} , were they are assigned as internal vibrations [5].

2 Experimental

2.1 Sampling

In the present study, the waste released during zinc hydrometallurgical process known as jarosite process, was obtained from Trepca zinc smelter tailing waste in Mitrovica Industrial Park, situated in northern Kosovo. Within the sampling framework three samples of 20 cm, 1 m and 2 m depth were collected. Samples were air-dried and sieved through 2 mm size sieve. Raman and infrared spectroscopy were used to evaluate the characteristics of Trepca jarosite that is present in open tailing dams in Mitrovica Industrial Park. Fig. 1 (a, b and c) shows the jarosite of Trepca zinc smelter, Mitrovica, Kosovo.

2.2 Raman spectroscopy

Raman analyses were performed using Bruker SENTERRA Dispersive Raman spectrometer equipped with Olympus microscope. Before analysis samples are transferred on the microscope glass slide and the surface is aligned with other glass slides which were removed prior to analysis. All samples were analyzed at randomly selected points and under the same conditions – Nd: YAG laser operating at 532 nm, nominal laser power 20 mW, excitation power 2 mW, magnification 506, aperture $25 \times 1000\text{ }\mu\text{m}$, spectral resolution $9\text{--}15\text{ cm}^{-1}$, grating 400a, 20 scans, 5 s integration time. Spectra were recorded in frequency range $10\text{--}4000\text{ cm}^{-1}$, and as a detector, a Peltier-cooled charge coupled device (CCD) camera was used. Spectra manipulation such as baseline correction (CRC Concave Rubberband Correction) and automatic determination of peak wavenumbers (Peak Picking) was performed using OPUS 7.0 software.

2.3 Infrared spectroscopy

FTIR spectra were recorded on Bruker Tensor 27 FTIR spectrometer with ATR technique of the diamond cell. Spectra were collected in frequency range $600\text{--}4000\text{ cm}^{-1}$ with 10 scans and spectral resolution 4 cm^{-1} . In the present study we present the results of Raman and Infrared spectroscopy.

3 Results and discussion

FTIR spectra of samples 1, 2 and 3 in the wavenumbers range of $600\text{--}4000\text{ cm}^{-1}$ and Raman spectra in the range of $80\text{--}4000\text{ cm}^{-1}$



a) Sample 1



b) Sample 2



c) Sample 3

Fig. 1. Jarosite of Trepca Zinc Smelter: a) Sample 1, b) Sample 2 and c) Sample 3.

are shown in Fig. 2 and 3. Peak position and assignments are listed in Table 1. Spectra are compared with each other and bands are assigned and interpreted in accordance with the cited literature data [1–10].

FTIR spectra of samples 1, 2 and 3 do not significantly differ, Fig. 2. In spectra, at the range of $\sim 700\text{--}1300\text{ cm}^{-1}$ dominate bad

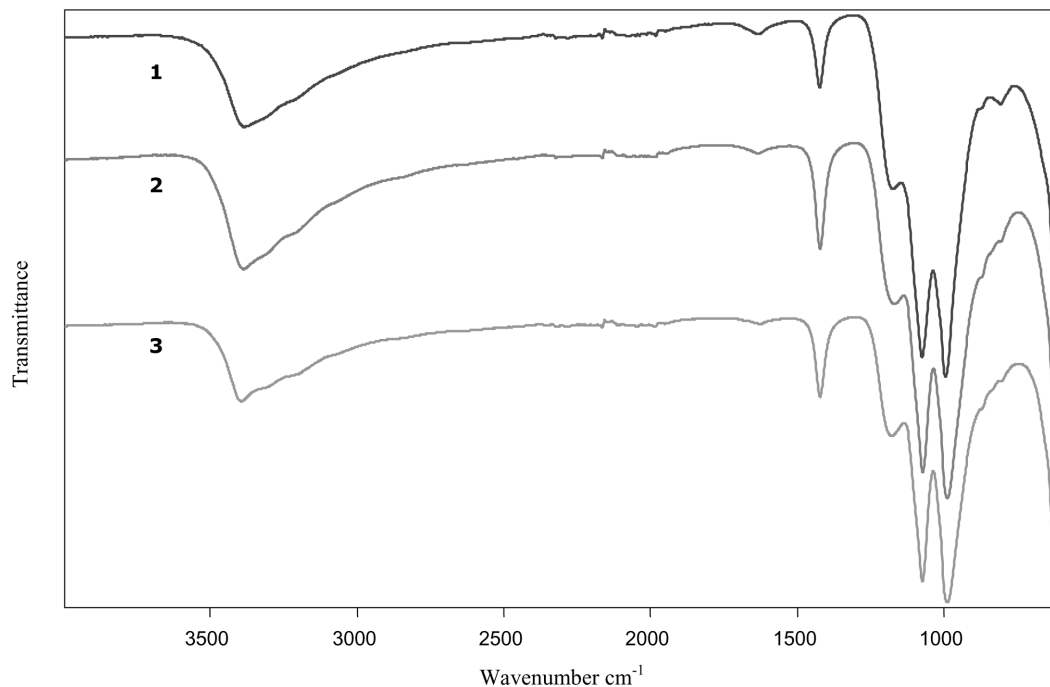


Fig. 2. FTIR spectra of Sample 1, 2 and 3 in frequency range 600–4000 cm^{-1} .

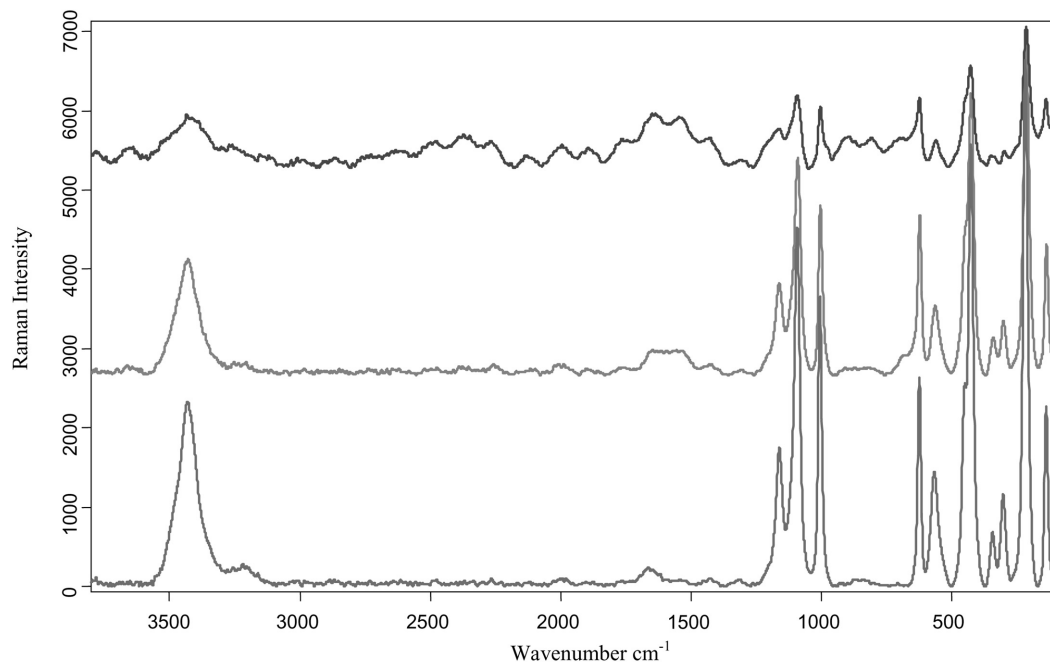


Fig. 3. Raman spectra of Sample 1, 2 and 3 in frequency range 80–3600 cm^{-1} .

resolved bands and a broad peak in the range of higher wavenumbers (2800–3600 cm^{-1}).

According to the group theory, in O-H and N-H stretching region, two bands that originate from O-H stretching vibration are expected to appear. In our spectra appears only one intense but poorly resolved band in range of 3394–3386 cm^{-1} that is

assigned as asymmetric stretching vibrations of hydroxyl, $\nu(\text{OH})$ [6]. On the side of the lower frequencies of these bands can be observed two small shoulders at ~ 3311 and ~ 3213 cm^{-1} , Fig. 4. Music *et al.* and Serna *et al.* reported a band at 3220 cm^{-1} that is assigned to the stretching mode of N-H bonds, $\nu(\text{NH}_4)$, of Ammonium jarosite [7, 8]. In that regard our band at ~ 3213 cm^{-1}

Table 1. Frequencies and assignments of infrared and Raman spectra.

Infrared				Raman			
1	2	3	Assignment	1	2	3	Assignment
3386b	3388b	3394b	$\nu(\text{OH})$	3434b	3388b	3394b	$\nu(\text{OH})$
3311sh	3311sh	3311sh	$\nu(\text{NH}_4)$	~3240wb	~3240wb	~3240wb	$\nu(\text{NH}_4)$
3213sh	3213sh	3209sh	$\nu(\text{NH}_4)$	~1663vw	~1663vw	~1663vw	$\nu(\text{NH}_4)$
1633w	1633w	1629w	$\delta(\text{H}_2\text{O})$	1163w	1162w	1162w	$\nu_3(\text{SO}_4^{2-})$
1423m	1421m	1421m	$\nu(\text{NH}_4)$	1093s	1091s	1093s	$\nu_3(\text{SO}_4^{2-})$
1176w	1171w	1178w	$\nu_3(\text{SO}_4^{2-})$	1003s	1004s	1006s	$\nu_1(\text{SO}_4^{2-})$
1073s	1071s	1072s	$\nu_3(\text{SO}_4^{2-})$	621m	623m	625m	$\nu_4(\text{SO}_4^{2-})$
994s	988s	988s	$\nu_1(\text{SO}_4^{2-})/\text{L}(\text{OH})$	562w	564w	568w	$\delta(\text{Fe-O})/\gamma(\text{OH})$
664vw	664sh	664sh	$\nu_4(\text{SO}_4^{2-})$	450sh	448sh	450vw	$\nu_2(\text{SO}_4^{2-})$
627m	625m	625m	$\nu_4(\text{SO}_4^{2-})$	426s	426s	427s	$\nu_2(\text{SO}_4^{2-})/\nu(\text{Fe-O})$
				343w	343w	345w	$\nu(\text{Fe-O})$
				302w	302w	304w	$\nu(\text{Fe-O})$
				214vs	215vs	219vs	$\nu(\text{Fe-O})/\text{LV}$
				135m	137m	138m	LV

b-broad, m-medium, sh-shoulder, s-strong, v-very strong, w-weak, LV-lattice vibration, L-librational

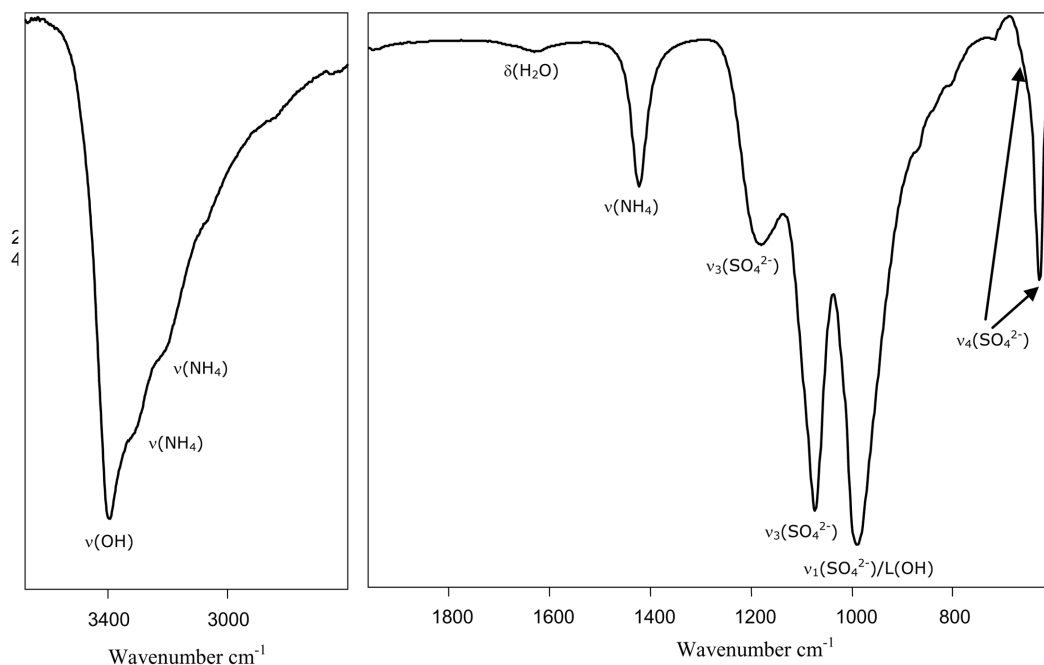


Fig. 4. FTIR spectra of Sample 3 in frequency range 3100–3600 cm^{-1} (left) and frequency range 600–2000 cm^{-1} (right).

can be assigned as $\nu(\text{NH}_4)$ [7, 8]. The same authors also mentioned that the band at 3340 cm^{-1} was assigned as $\nu(\text{NH}_4)$. In our spectra it could possibly be a band at ~3311 cm^{-1} . This band is less visible in spectrum 1 than in spectra 2 and 3.

A more detailed view of the spectra fingerprint area is shown in the Fig. 4. In this part of spectra is present very weak band with maximum of ~1630 cm^{-1} , which is attributed to the bending of surface adsorbed water, $\delta(\text{H}_2\text{O})$ [4, 9]. Medium band at ~1421 cm^{-1} is attributed to bending mode of N-H, $\nu(\text{NH}_4)$ that appears only in Ammonium jarosite. This confirms the assignment of bands at ~3311 and 3213 cm^{-1} [10]. In the spectrum of Ammonium jarosite, there should appear another band at 1655 cm^{-1} , but in our study it was not observed [7, 8]. The weak band, the

center of which depends on the sample is in the range 1171–1178 cm^{-1} , as well as the strong band at ~1072 cm^{-1} are assigned to asymmetric stretching mode of bonds in sulphate anion, $\nu_3(\text{SO}_4^{2-})$. The position of these bands should indicate the type of cation M^+ . As above stated, the Ammonium jarosite is present in the analyzed samples 1, 2 and 3. However, the position of $\nu_3(\text{SO}_4^{2-})$ band that occurs in the range of 1171–1178 cm^{-1} shows deviation from literature values (expected at 1190–1200 cm^{-1}) and actually it indicates the presence of Na^+ or K^+ cations in the structure [1, 6–8]. There is an uncertainty concerning the assignment of strong band at 994 cm^{-1} (Sample 1) and 988 cm^{-1} (Samples 2 and 3). Several authors attributed this band to symmetric stretching vibration of sulphate, $\nu_1(\text{SO}_4^{2-})$. Unlike them, Frost et

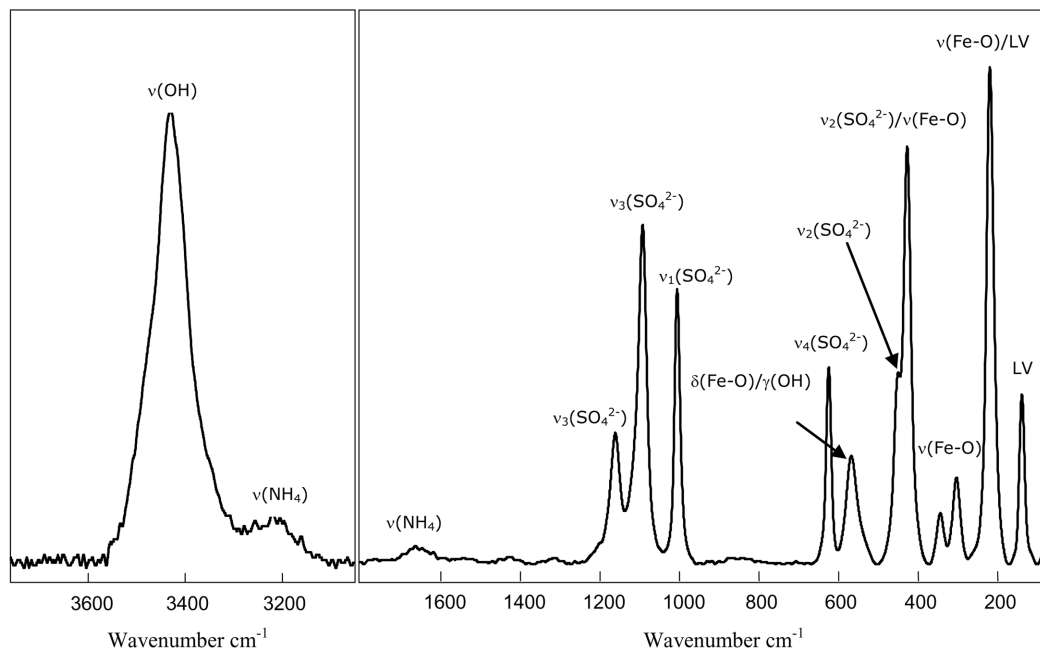


Fig. 5. Raman spectra of Sample 3 in frequency range 3000–3800 cm^{-1} (left) and frequency range 80–1800 cm^{-1} (right).

al. [6] emphasizes that the presence of such strong sulphate peak in FTIR spectrum is unusual, therefore they assigned it to the OH librational mode [1, 6, 8]. According to Frost *et al.* [6] sharp peak of medium intensity at 625 cm^{-1} is in a good agreement with the position of asymmetric sulphate group bending peak ($\nu_4(\text{SO}_4^{2-})$) in Ammonium jarosite, where in the other types of jarosite is close to the position of that peak [3–5, 7, 9, 10]. In fact, this sharp peak doubles with the equivalent peak at 664 cm^{-1} which is poorly resolved in spectrum 1, while in spectra 2 and 3 it appears in a shoulder shape. There are some additional bands at ~ 870 (sh), ~ 807 (vw) and ~ 718 (vw) cm^{-1} that cannot be assigned to jarosite molecules. Therefore, they are considered as impurities present in the study material.

Sampling depth did not impact FTIR spectra of samples 1, 2 and 3, while Raman spectra show certain differences, Fig. 3. The biggest difference can be seen in sample 1 spectrum, where smaller intensity of peaks is observed, peak resolving is worse, and the base line exhibits higher noise. The differences that are stated above are most probably a result of the presence of amorphous impurities in sample 1. In this regard, by examining the spectra one in sample 1 we can conclude that sample 1 gives the same bands as samples 2 and 3, even in similar ratios.

In all Raman spectra there are at least 5 bands that can be assigned to sulphate vibrational modes, Fig. 5. Two equivalent bands, one with low intensity at a range of 1162 cm^{-1} and the other band with strong intensity at a range of 1087–1093 cm^{-1} , are assigned to the asymmetric stretching vibration of sulphate $\nu_3(\text{SO}_4^{2-})$. The band at 1162 cm^{-1} is very close to the wavenumber of Ammonium jarosite spectra according to cited references [1, 5, 6]. In addition there is a band at 1006 cm^{-1} that in previous works was assigned to the symmetrical stretching mode $\nu_1(\text{SO}_4^{2-})$ of the Ammonium jarosite molecule. Two remaining sulphate bands, $\nu_4(\text{SO}_4^{2-})$ and $\nu_2(\text{SO}_4^{2-})$, stands at the range of 625 and 450 cm^{-1} . The last band is poorly resolved. It is visible as

a shoulder on sharp and strong peak of 427 cm^{-1} in spectra 1 and 2, while in spectrum 3 it forms a tiny peak on the same band. In attribution of 427 cm^{-1} the strong peak is questionable. Sasaki *et al.* [1] assigned a band to the stretching vibrations of Fe-O (at 434 cm^{-1}), while the majority of other authors considered such attribution slightly possible and assigned it to the symmetric sulphate bending mode ($\nu_2(\text{SO}_4^{2-})$). As it was predicted by decreasing symmetry of sulphate crystal structure, there are 6 sulphate bands in Raman spectra [1, 3, 5, 6, 10].

Remaining peaks in the spectrum are the results of Fe-O vibration, and provide peaks at 343, 302 and 215 cm^{-1} . The first two bands have low intensity and are not fully resolved, while the last one is the most intensive ones in all spectra. Some authors assigned the 215 cm^{-1} band together with the 138 cm^{-1} band to crystal lattice vibration of the Ammonium jarosite [5, 6]. In accordance with cited references, the band at 568 cm^{-1} is not assigned. The references provide two possibilities, where the bending vibration of $\delta(\text{Fe-O})$ is on a plane or deformation, and the other bending vibration of OH is out of a plane $\gamma(\text{OH})$ [1, 3–6]. Chioa *et al.* [5] assigned that band to the liberation mode of OH functional group, L (OH).

There is a strong band in the high wavenumber region, in the range of 3428–3434 cm^{-1} that is attributed to the $\nu(\text{OH})$ [3, 5, 10], Fig. 5. At low wavenumber region there is a broad and low intensity band, around 3240 cm^{-1} that represents inner vibrations of ammonium ion together with the low intensity band around 1663 cm^{-1} [5].

4 Conclusions

Raman and infrared spectroscopy proves to be very useful tool for the identification and characterization of the jarosite process tailing waste. The results obtained confirm the presence of

Ammonium jarosite in three samples of Trepca tailing waste, Kosovo.

The presence of impurities in jarosite waste is detected by FTIR spectroscopy, at the range of 994 cm^{-1} for sample 1 and 988 cm^{-1} for samples 2 and 3, while the Raman spectroscopy indicates the presence of amorphous impurities in the sample 1.

The present study has also shown that the depth does not impact the FTIR spectra of samples 1, 2 and 3, while the Raman spectra show certain differences. In fact, the biggest difference is shown in the spectrum of sample 1, where small intensity of peaks is observed. The Raman spectra showed 5 bands that are assigned to sulphate vibrational modes. Additionally, there is a strong band in the range of 3428–3434 cm^{-1} that is attributed to the $\nu(\text{OH})$. From this study we can conclude that the Raman and FTIR spectral techniques found all fundamental of vibrational modes of sulphate anion, identified the type of jarosite that is present in the study material together with other impurities detected. In order to resolve the uncertainty of Raman and FTIR techniques further measurement by X-ray diffraction (XRD), SEM, as well as AAS and ICP-AES for determination of trace metals are in process.

5 References

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