Monitoring of the Forced Hydrolysis of FeCl₃ Solutions in the Presence of Sodium Dodecyl Sulphate

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Abstract: Precipitations by the forced hydrolysis of 0.2 M FeCl₃ aqueous solutions between 2 and 72 hours in the presence of 1 % sodium n-dodecyl sulphate (SDS) were investigated. In the absence of SDS a direct phase transformation β-FeOOH → α-Fe₂O₃ via dissolution/recrystallization occurred in the precipitation system. In the presence of SDS small amounts (traces) of α-FeOOH as an intermediate phase precipitated, and with a prolonged time of forced hydrolysis α-FeOOH also transformed to α-Fe₂O₃ via the dissolution/recrystallization mechanism. On the basis of Mössbauer spectra it was concluded that in the presence of SDS the α-Fe₂O₃ phase exhibited a lower degree of crystallinity. During the precipitation process in the presence of SDS the competition between the stability of Fe(III)-dodecyl sulphate on one side and the formation of iron oxide phases on the other also played an important role. Thermal field emission scanning electron microscopy (FE SEM) revealed that the big α-Fe₂O₃ particles possessed a substructure. The elongation of primary α-Fe₂O₃ particles produced in the presence of SDS was noticed. This effect can be assigned to the preferential adsorption of sulphate groups on nuclei and crystallites of FeOOH and α-Fe₂O₃ phases during the forced hydrolysis of FeCl₃ solutions.

Keywords: FeCl₃ hydrolysis, sodium n-dodecyl sulphate, α-Fe₂O₃, α-FeOOH, XRD, Mössbauer, FT-IR, FE SEM.

INTRODUCTION

EMATITE (α-Fe₂O₃) particles have found different applications as pigments, photocatalysts, electrode materials, fine abrasives and cosmetic additives. α-Fe₂O₃ can also be used as an adsorbent for toxic elements or radioisotopes in wastewater treatment. In surface and colloid chemistry these particles are often used as model systems due to their excellent acid-base surface properties. The investigation of colloid stability of α-Fe₂O₃ particles is important to understand the nature of this phenomenon.

The simplest way to produce α-Fe₂O₃ particles is by forced hydrolysis in a boiling 0.01 M FeCl₃ solution,[1] whereas at temperatures below 70 °C akaganèite (β-FeOOH) particles (spindle- or cigar-shaped) will precipitate. The synthesis and characterization of β-FeOOH and its decomposition products in vacuum were investigated.[2] Musić et al.[3] used Mössbauer spectroscopy to investigate the hydrolysis of 0.1 M solutions of Fe(NO₃)₃, FeCl₃, Fe₂(SO₄)₃ or NH₄Fe(SO₄)₂ at 90 °C. A proposal was made concerning the mechanism of formation of the hydroxides and oxides of Fe³⁺ ions in these precipitation systems. It is generally accepted that the phase transformation β-FeOOH → α-Fe₂O₃ in hydrolysing FeCl₃ solutions at elevated temperature is driven by a dissolution/recrystallization mechanism.[4–6] The application of Mössbauer spectroscopy to investigate the precipitation of iron oxides has recently been reviewed by Musić et al.[7]

The nano/microstructure of α-Fe₂O₃ plays a very important role in many of its applications For this reason many researchers investigated conditions for the preparation of α-Fe₂O₃ with different properties. Katsuki and Komarneni[8] used the forced hydrolysis of the FeCl₃ solution.
at hydrothermal conditions to investigate the influence of morphology on the colour of the red pigment (α-Fe₂O₃) in the production of porcelain. Sugimoto et al.[9] noticed remarkably different effects of Cl⁻, OH⁻, SO₄²⁻ or PO₄³⁻ anions on the shape and internal structure of α-Fe₂O₃ particles. Hollow α-Fe₂O₃ spheres were produced by the forced hydrolysis of the FeCl₃ solution containing H₃PW₁₂O₄₀.[10] Mesoporous α-Fe₂O₃ particles were produced by the forced hydrolysis of the FeCl₃ solution containing L-phenylalanine and NaOH.[7] Žic et al.[11] investigated the effect of surfactants on the precipitation of colloidal particles by the forced hydrolysis of the FeCl₃-HCl solution. The hydrothermal transformation β-FeOOH → α-Fe₂O₃ in dense aqueous suspensions, prepared by a partial neutralization of the concentrated FeCl₃ solution with the concentrated NaOH solution, was investigated by Žic et al.[11] Žic et al.[12] also investigated the precipitation of α-Fe₂O₃ from dense β-FeOOH suspensions containing ammonium amidosulphonate. The peanut-type α-Fe₂O₃ particles, as well as particles in the form of double cupolas interconnected with the neck were obtained. These particles showed a substructure. Double cupolas were porous and consisted of linear chains of small α-Fe₂O₃ particles (also interconnected) which were directed from the centre towards the surface of cupolas.

The aim of the present work was to obtain more data about the influence of sodium dodecyl sulphate on the crystallization kinetics, phase transformations and particle morphology in the precipitates produced by the forced hydrolysis of aqueous FeCl₃ solutions. This work is a continuation of our longtime investigations in the precipitation chemistry of iron oxides (group name for hydroxides, oxyhydroxides and oxides of iron). In many cases the phase analyses of solid hydrolytical products of iron ions are a demanding task, specifically if iron oxide phases vary from amorphous to a well-crystalline nature. For this reason three complementary techniques, XRD, ⁵⁷Fe Mössbauer and FT-IR were used in the phase analysis. Generally, it is known that surface active agents act very differently on the precipitation processes in dependence on their characteristics (polar, nonpolar, chemical nature of the organic chain, pH).

**EXPERIMENTAL**

**Sample Preparation**

AnalaR grade FeCl₃·6H₂O was supplied by Kemika. Sodium n-dodecyl sulphate (SDS) was supplied by Sigma Aldrich (Cat. No.: 151-21-3; ACS grade reagent). Twice distilled water was prepared in own laboratory and used in all experiments. A stock solution 2M FeCl₃ was prepared. The concentration of 0.2M FeCl₃ solutions was adjusted in all precipitation systems. Experimental conditions for the preparation of samples are given in Table 1. The yellowish precipitate was formed by adding the FeCl₃ solution to a clear aqueous solution of SDS, thus indicating the formation of Fe(III)-dodecyl sulphate. Thus obtained suspension was homogenized in an ultrasound bath. The precipitation systems were autoclaved at 160 °C using a Teflon-lined, non-stirred pressure vessel manufactured by Parr Instruments (model 4744). The autoclaves were heated between 2 and 72 h in a DX 300 gravity oven (Yamato; temperature uniformity ± 1.9 °C at 100 °C and ± 3 °C at 200 °C). The autoclaving times were corrected for the time needed that the autoclave reaches the predetermained temperature. After a proper autoclaving time the autoclaves were abruptly cooled with cold water. The mother liquor was separated from the precipitate with the ultracentrifuge (Sorvall model Super T21). The separated precipitates were subsequently washed with twice-distilled water and one time with C₂H₅OH, then dried.

**Instrumentation**

XRD patterns were recorded with an APD 2000 powder diffractometer manufactured by ItalStructures (G NR Analytical Instruments Group, Italy). ⁵⁷Fe Mössbauer spectra were recorded at 20°C in the transmission mode, using a standard configuration by WissEl GmbH (Starnberg, Germany). The ⁵⁷Co/Rh Mössbauer source was used. The velocity scale and Mössbauer parameters refer to the metallic α-Fe absorber at 20 °C. Deconvolution of Mössbauer spectra was performed using the MossWin program.

**Table 1.** Experimental conditions for forced hydrolysis of 0.2 M FeCl₃ solutions.

<table>
<thead>
<tr>
<th>Sample</th>
<th>V(FeCl₃, 2M)/mL</th>
<th>SDS[SDS] / g</th>
<th>V(H₂O)/ml</th>
<th>t / °C</th>
<th>Time of ageing / h[η]</th>
</tr>
</thead>
<tbody>
<tr>
<td>R1</td>
<td>4</td>
<td>36</td>
<td>160</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>R2</td>
<td>4</td>
<td>36</td>
<td>160</td>
<td>6</td>
<td></td>
</tr>
<tr>
<td>R3</td>
<td>4</td>
<td>36</td>
<td>160</td>
<td>24</td>
<td></td>
</tr>
<tr>
<td>R4</td>
<td>4</td>
<td>36</td>
<td>160</td>
<td>72</td>
<td></td>
</tr>
<tr>
<td>S1</td>
<td>4</td>
<td>0.4</td>
<td>36</td>
<td>160</td>
<td>2</td>
</tr>
<tr>
<td>S2</td>
<td>4</td>
<td>0.4</td>
<td>36</td>
<td>160</td>
<td>4</td>
</tr>
<tr>
<td>S3</td>
<td>4</td>
<td>0.4</td>
<td>36</td>
<td>160</td>
<td>6</td>
</tr>
<tr>
<td>S4</td>
<td>4</td>
<td>0.4</td>
<td>36</td>
<td>160</td>
<td>10</td>
</tr>
<tr>
<td>S5</td>
<td>4</td>
<td>0.4</td>
<td>36</td>
<td>160</td>
<td>16</td>
</tr>
<tr>
<td>S6</td>
<td>4</td>
<td>0.4</td>
<td>36</td>
<td>160</td>
<td>24</td>
</tr>
<tr>
<td>S7</td>
<td>4</td>
<td>0.4</td>
<td>36</td>
<td>160</td>
<td>72</td>
</tr>
</tbody>
</table>

[SDS] = sodium dodecyl sulphate.
[η] = hour.
FT-IR spectra were recorded at RT with a Perkin-Elmer spectrometer (model 2000). The powders were mixed with KBr, then pressed into tablets using the Carver press.

The samples were also inspected with a thermal field emission scanning electron microscope (FE SEM, model JSM-7000F) manufactured by Jeol Ltd.

RESULTS AND DISCUSSION

The XRD patterns of reference samples R1, R2 and R4 are given in Figure 1 corresponding to single phases of $\beta$-FeOOH (sample R1) or to $\alpha$-Fe$_2$O$_3$ (samples R2 and R4).

The Mössbauer spectra of reference samples R1 to R4 produced in the absence of SDS surfactant are shown in Figure 2. The Mössbauer spectrum of sample R1 shows the superposition of two doublets with quadrupole splittings $\Delta_1 = 0.55$ and $\Delta_2 = 0.99$ mm s$^{-1}$. The parameters of this spectrum can be assigned to $\beta$-FeOOH (Table 2). The spectra of samples R2, R3 and R4 are characterised by one sextet with parameters corresponding to $\alpha$-Fe$_2$O$_3$. The increase in the hyperfine magnetic field (HMF) from 51.2 to 51.5 T (Table 2) with the autoclaving time prolonged from 6 to 72 h can be related to crystalline ordering in $\alpha$-Fe$_2$O$_3$.

Table 2. $^{57}$Fe Mössbauer parameters for reference samples (R) and (S) obtained by precipitation in the presence of SDS.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Line</th>
<th>$\delta /$ mms$^{-1}$</th>
<th>$\Delta$ or $E_q /$ mm s$^{-1}$</th>
<th>HMF / T</th>
<th>$\Gamma /$ mm s$^{-1}$</th>
<th>$A /$%</th>
<th>Identification</th>
</tr>
</thead>
<tbody>
<tr>
<td>R1</td>
<td>Q1</td>
<td>0.38</td>
<td>0.55</td>
<td></td>
<td>0.30</td>
<td>57</td>
<td>$\beta$-FeOOH</td>
</tr>
<tr>
<td></td>
<td>Q2</td>
<td>0.37</td>
<td>0.99</td>
<td></td>
<td>0.33</td>
<td>43</td>
<td></td>
</tr>
<tr>
<td>R2</td>
<td>M</td>
<td>0.37</td>
<td>-0.21</td>
<td>51.2</td>
<td>0.34</td>
<td>100</td>
<td>$\alpha$-Fe$_2$O$_3$</td>
</tr>
<tr>
<td>R3</td>
<td>M</td>
<td>0.36</td>
<td>-0.21</td>
<td>51.5</td>
<td>0.29</td>
<td>100</td>
<td>$\alpha$-Fe$_2$O$_3$</td>
</tr>
<tr>
<td>R4</td>
<td>M</td>
<td>0.37</td>
<td>-0.21</td>
<td>51.5</td>
<td>0.34</td>
<td>100</td>
<td>$\alpha$-Fe$_2$O$_3$</td>
</tr>
<tr>
<td>S1</td>
<td>Q(a)</td>
<td>0.38</td>
<td>0.79</td>
<td></td>
<td>0.23</td>
<td>100</td>
<td>$\beta$-FeOOH</td>
</tr>
<tr>
<td>S2</td>
<td>Q(a)</td>
<td>0.38</td>
<td>0.81</td>
<td></td>
<td>0.48</td>
<td>85</td>
<td>$\beta$-FeOOH</td>
</tr>
<tr>
<td></td>
<td>M1</td>
<td>0.37</td>
<td>-0.26</td>
<td>36.7</td>
<td>1.14</td>
<td>5</td>
<td>$\alpha$-FeOOH</td>
</tr>
<tr>
<td></td>
<td>M2</td>
<td>0.37</td>
<td>-0.21</td>
<td>48.1</td>
<td>0.58</td>
<td>10</td>
<td>$\alpha$-Fe$_2$O$_3$</td>
</tr>
<tr>
<td>S3</td>
<td>Q(a)</td>
<td>0.38</td>
<td>0.75</td>
<td></td>
<td>0.54</td>
<td>39</td>
<td>$\beta$-FeOOH</td>
</tr>
<tr>
<td></td>
<td>M1</td>
<td>0.37</td>
<td>-0.26</td>
<td>37.4</td>
<td>0.97</td>
<td>5</td>
<td>$\alpha$-FeOOH</td>
</tr>
<tr>
<td></td>
<td>M2</td>
<td>0.37</td>
<td>-0.20</td>
<td>48.5</td>
<td>0.31</td>
<td>56</td>
<td>$\alpha$-Fe$_2$O$_3$</td>
</tr>
<tr>
<td>S4</td>
<td>Q</td>
<td>0.37</td>
<td>1.18</td>
<td></td>
<td>0.53</td>
<td>4.7</td>
<td></td>
</tr>
<tr>
<td></td>
<td>M(a)</td>
<td>0.37</td>
<td>-0.21</td>
<td>48.3</td>
<td>0.29</td>
<td>95.3</td>
<td>$\alpha$-Fe$_2$O$_3$</td>
</tr>
<tr>
<td>S5</td>
<td>M(a)</td>
<td>0.37</td>
<td>-0.20</td>
<td>49.2</td>
<td>0.27</td>
<td>100</td>
<td>$\alpha$-Fe$_2$O$_3$</td>
</tr>
<tr>
<td>S6</td>
<td>M(a)</td>
<td>0.37</td>
<td>-0.21</td>
<td>49.8</td>
<td>0.27</td>
<td>100</td>
<td>$\alpha$-Fe$_2$O$_3$</td>
</tr>
<tr>
<td>S7</td>
<td>M</td>
<td>0.37</td>
<td>-0.21</td>
<td>50.7</td>
<td>0.25</td>
<td>100</td>
<td>$\alpha$-Fe$_2$O$_3$</td>
</tr>
</tbody>
</table>

All data are given relative to $\alpha$-Fe standard.

Key: $\delta$ = isomer shift; $\Delta$ or $E_q$ = quadrupole splitting; HMF = hyperfine magnetic field; $\Gamma$ = line width; $A$ = area under the peaks.

Errors: $\delta = \pm 0.01$ mm s$^{-1}$; $\Delta$ or $E_q = \pm 0.01$ mm s$^{-1}$; HMF = $\pm 0.2$ T.

(a) Fitted with distribution of quadrupole doublets or hyperfine magnetic fields.

Samples S4 and S5 could contain traces of $\alpha$-FeOOH, as denoted with arrows (G) in corresponding figures.
Figure 3 shows the FT-IR spectra of reference samples R1 to R4. The FT-IR spectrum of sample R1 can be assigned to β-FeOOH. Sample R1 shows an IR band centred at 848 cm$^{-1}$, two shoulders at 700 and 640 cm$^{-1}$ and two shoulders at 498 and 425 cm$^{-1}$. According to an earlier work\[15,16\] the band at 848 cm$^{-1}$ and the shoulder at 700 cm$^{-1}$ can be assigned to the deformation vibration of OH groups, whereas the intense shoulder at 640 cm$^{-1}$ can be related to the interaction of Fe-OH groups with Cl$^-$ ions.

Generally, β-FeOOH exhibits a hollandite-type crystal structure while the structural channels in β-FeOOH produced from the FeCl$_3$ solution contain water and a small amount of Cl$^-$ ions.\[14\] These Cl$^-$ ions (less than ~ 2 %) in structural channels stabilize the crystal lattice of β-FeOOH and cannot be removed by simple washing. Weckler and Lutz[17] discussed two sets of vibrations at 847 and 820 cm$^{-1}$, and also those at 697 and 644 cm$^{-1}$ in terms of two O–H–Cl hydrogen bonds present in β-FeOOH. The IR spectrum of β-FeOOH was also the subject of discussion by Murad and Bishop.\[18\] The FT-IR spectra of samples R2 to R4 can be assigned to α-Fe$_2$O$_3$. Wang et al.\[19\] tabulated optical parameters for bulk α-Fe$_2$O$_3$ and investigated the influence of the geometrical shape of α-Fe$_2$O$_3$ particles on the corresponding FT-IR spectrum. Generally, the IR spectrum of α-Fe$_2$O$_3$ shows six active vibrations, two $A_{2u}$ ($E_{\parallel}$) and four $E_{u}$ ($E_{\perp}$).

The XRD patterns of selected samples S1, S2, S4, S6 and S7, prepared in the presence of SDS, are shown in Figure 4. The effect of adding SDS to the precipitation systems is clearly visible. The sharpness of XRD lines of sample S1 is decreased in relation to sample R1, which can be related to lower crystallinity and/or decreased crystallite size of β-FeOOH particles. This can also be related to different integral intensities between XRD lines 310 and 211 for β-FeOOH in sample R1 relative to S1 and S2 (Figures 1 and 4). Additionally, XRD patterns of sample S2 showed the presence of very small amounts of goethite (G) and hematite (H). After 10 h of forced hydrolysis a small amount of α-FeOOH (sample S4) was also visible in the corresponding XRD pattern (line 110). XRD patterns of samples S6 and S7 showed the presence of α-Fe$_2$O$_3$ as a single phase.
Figures 5 and 6 show the Mössbauer spectra of samples S1 to S7. Upon 2h of forced hydrolysis of the 0.2 M FeCl₃ solution containing SDS surfactant (sample S1) a central quadrupole doublet was recorded. This spectrum was fitted for one average doublet with $\delta = 0.79$ mm s⁻¹ (Table 2) and could be assigned to $\beta$-FeOOH.[20] The spectrum of sample S2 was fitted as a superposition of the central quadrupole doublet and two sextets. The central quadrupole doublet ($\Delta = 0.81$ mm s⁻¹) could be assigned to $\beta$-FeOOH (85 %), whereas the two sextets of very small intensities with HMF = 36.7 T (5 %) and HMF = 48.1 (10 %) can be assigned to $\alpha$-FeOOH and $\alpha$-Fe₂O₃, respectively. Upon further prolongation of autoclaving time up to 6 h there was a decrease in $\beta$-FeOOH to 39 % and a corresponding increase in $\alpha$-Fe₂O₃ to 56 %, whereas $\alpha$-FeOOH remained at 5 % (sample S3). In sample S4 the $\alpha$-Fe₂O₃ fraction increased to 95.3 %. The same spectrum also showed central quadrupole doublet with $\Delta = 1.18$ mm s⁻¹ (4.7 %). In line with XRD pattern the peak denoted as G in the Mössbauer spectrum of sample S4 could be assigned to $\alpha$-FeOOH. Mössbauer spectrum of sample S7 corresponds to $\alpha$-Fe₂O₃ as a single phase (Figure 6). The HMF values of the $\alpha$-Fe₂O₃ phase increased from 48.1 T (samples S2) to 50.7 (sample S7), thus indicating a much lower degree of $\alpha$-Fe₂O₃ crystallinity in relation to reference samples R2 to R4 (51.2 to 51.5 T).

The effect of SDS is also visible in the FTIR spectra shown in Figure 7. The spectrum of sample S1 can be

![Figure 5. $^{57}$Fe Mössbauer spectra of samples S1 to S6 prepared in the presence of SDS.](image1)

![Figure 6. $^{57}$Fe Mössbauer spectrum of sample S7 prepared in the presence of SDS.](image2)

![Figure 7. FTIR spectra of samples S1 to S7 prepared in the presence of SDS.](image3)
assigned to the β-FeOOH phase. The only difference between samples S1 and S2 is in the relative intensities of the shoulders at 696, 648, and 425 cm⁻¹. Additionally, the IR bands at 1128 cm⁻¹ (sample S1) and 1209, 1133 cm⁻¹ (sample S2) are also visible. Sample S3 shows IR bands that can be assigned to β-FeOOH and two additional IR bands at 570 and 470 cm⁻¹ that can be assigned to α-Fe₂O₃.[21] The IR band at 896 cm⁻¹ could be assigned to a small amount of the α-FeOOH phase, but the second IR band close to 800 cm⁻¹, also typical of α-FeOOH, is not visible due to the overlapping with a broad IR band of β-FeOOH centred at 848 cm⁻¹.[21] Generally, the IR band of α-FeOOH centred at 896 cm⁻¹ is assigned to the in-plane Fe–O–H bending bend (δOH), whereas the out-of-plane bending band (γOH) is typically around 800 cm⁻¹.[21] In the FT-IR spectra of samples produced in the presence of SDS additional bands are also noticed. For example, in the spectrum of sample S3 four IR bands at 1206, 1115, 1036, and 976 cm⁻¹ are well visible. These IR bands can be related to the sulphate group. The ν₃(SO₄) fundamental vibration is split into 3 active IR bands due to the formation of a surface bidentate bridging complex between the sulphate group and surface iron atoms.[22] The presence of an IR band at 977 cm⁻¹ is due to the ν₁(SO₄) vibration. It is generally known that the specific adsorption of oxyanions reaches its maximum at acidic pH values and decreases with an increase in pH values.

Generally, it is also known that the presence of oxyanions, surface active or complexing agents may strongly influence the shape of hematite particles in precipitation processes. For example, Ocaña et al.[23] precipitated spindle-shaped α-Fe₂O₃ particles by a forced hydrolysis of the Fe(ClO₄)₃ solution at 100 °C. The phosphate anions that could not be washed out were discussed as responsible for the formation of spindle-shaped particles. The adsorbed phosphates were visible in the IR spectrum as evidence, with several peaks between 1036 and 934 cm⁻¹. In the present work Mössbauer, XRD and FT-IR measurements showed a direct phase transformation β-FeOOH → α-Fe₂O₃ in reference samples (R) produced by the forced hydrolysis of the 0.2 M FeCl₃ solution. In the presence of SDS (samples S1 to S7) the kinetics of this phase transformation was retarded and in samples produced between 4 and 24 hours of forced hydrolysis a small fraction (traces) of an intermediate phase α-FeOOH were detected. The HMF values of α-Fe₂O₃ particles prepared in the presence of SDS varied from 48.1 to 50.7 T. These HMF values were significantly below 51.75 T measured for well-crystalline α-Fe₂O₃ as reported by Murad and Johnston.[24] It can be concluded that the presence of SDS lowers the crystallinity of precipitated α-Fe₂O₃. Musić et al.[25] investigated the effect of HCl additions on the forced hydrolysis of FeCl₃ solutions. They found out that under certain conditions the forced hydrolysis of the FeCl₃ solution containing only HCl additions can also produce an α-FeOOH phase. Wang et al.[26] investigated the precipitation of α-Fe₂O₃ nanoparticles by the forced hydrolysis of FeCl₃ solutions with no additive presence. In dependence on experimental conditions, primary nanoparticles showed different morphologies. Beside the β-FeOOH → α-Fe₂O₃ phase transformation there was also a direct phase transformation of the amorphous fraction.

![Image](image_url)

**Figure 8.** FE SEM images of (a) sample R1 and (b, c) sample R2.
into α-Fe₂O₃. In the present work the addition of SDS to the precipitation system caused the formation of a small fraction of α-FeOOH as an intermediate phase, which at a pro-
longed time of autoclaving was transformed into α-Fe₂O₃ by way of the dissolution/recrystallization mechanism.

The FE SEM image of sample R1 (Figure 8a) showed the dominant presence of β-FeOOH rods, but star- and X-shaped particles were also noticed. Sample R2 showed micron size α-Fe₂O₃ particles (Figure 8b) which consisted of primary α-Fe₂O₃ nanoparticles (elongated), as evidenced by Figure 8c. α-Fe₂O₃ particles produced upon 72 h of the forced hydrolysis of FeCl₃ solutions at 160 °C (sample R4) are shown in Figure 9a. These particles also showed a substructure (Figure 9b); however, the primary α-Fe₂O₃ particles increased in size. Figure 10 shows the FE SEM images of selected samples prepared in the presence of SDS. The FE SEM image of sample S1 (Figure 10a) shows the aggregates of very fine β-FeOOH needles. In addition to very fine β-FeOOH needles, Figure 10b (sample 10b), also shows peanut particles of α-Fe₂O₃. The substructure of these peanut particles consisting of laterally arrayed subparticles is visible. The FE SEM images of samples S4 and S5, respectively, are shown in Figures 10c,d, demonstrating the predominant presence of big α-Fe₂O₃ particles of different morphol-
gies (spheres, double cupolas with ring and peanut shaped particles). Similar shapes of α-Fe₂O₃ particles are also visible in the FE SEM images of sample S7 (Figure 10e,f).

CONCLUSION

The effect of SDS (1 %) on the kinetics, phase composition and shape of the particles precipitated by the forced hydrolysis of the 0.2 M FeCl₃ solution was investigated. A direct phase transformation β-FeOOH → α-Fe₂O₃ via dis-
solution/recrystallization was present in the absence of SDS. On the other hand, in the presence of SDS this trans-
formation was retarded in relation to reference kinetics. In the presence of SDS a small fraction of α-FeOOH precipi-
tated, which was transformed into the end product α-Fe₂O₃ also via the dissolution / recrystallization mechanism. α-
Fe₂O₃ particles precipitated in the presence of SDS showed lower crystallinity in relation to reference samples, as con-
cluded on the basis of the Mössbauer spectra. This effect was explained by a competition between the stability of Fe(III)-dodecyl sulphate on one side and the formation of iron oxide phases on the other. Precipitated α-Fe₂O₃ particles showed a substructure, i.e., consisted of much smaller primary particles. The effect of SDS on the microstructure of α-Fe₂O₃ particles was noticed. The influence of SDS on the forced hydrolysis of FeCl₃ solutions can be related to the specific adsorption of sulphate groups on the nuclei and crystalites of FeOOH and α-Fe₂O₃ phases. The steric effect of n-alkyl chain cannot be excluded. This work also showed

Figure 9. FE SEM images of (a,b) sample R4.

Figure 10. FE SEM images of (a) sample S1, (b) sample S2, (c) sample S4, (d) sample S5 and (e,f) sample S7 precipitated in the presence of SDS.
how the important is combination of XRD, Mössbauer and FT-IR spectroscopic methods in the analysis of small amounts (traces) of iron oxides produced in the precipitation processes.

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