Influence of Structural Heterogeneity of High-Strength OCTG Tubes on Sulfide Corrosion Cracking Resistance †

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Abstract: High-strength oil country tubular goods (OCTG) like C110, according to standard API 5CT (yield strength at least 758 MPa), are subject to requirements in terms of mechanical and corrosion properties. In this work, we studied the influence of seamless tubes microstructure with a 177.8 mm diameter and 10.36 mm wall thickness of class С110 high-strength steel to sulfide stress corrosion cracking (SSC) and sulfide stress corrosion cracking with low strain rates (SSRT). Tubes were obtained from continuous billets by screw piercing with preliminary quenching and tempering. It was established that cracking during the tests always begins from the inner surface of the tube. Rough segregation bands were found on the inner tube surface, which occupies about a third of the thickness. It is shown that the SSRT assessment technique allows to estimate the threshold value of the resistance.

Keywords: high-strength OCTG; sulfide corrosion cracking; sulfide stress corrosion tests with low strain rates (SSRT); microstructure; banding; segregation

1. Introduction
During the development of deep oil and gas fields, casing is influenced not only by the aggressive environment of the well (H2S and CO2), but high pressures and temperatures [1–4]. If low-alloyed steel pipes are exposed to these environments, hydrogen cracking can occur, causing corrosion cracks in the hydrogen sulfide environment (SSC) [5]. Casing of high-strength groups in a corrosion-resistant design is used for those conditions, for example C110, according to ANSI/API 5CT (yield strength at least 758 MPa) [6,7].

Extreme operating conditions require high quality of steels and determine the requirements for the level of alloying and purity for detrimental impurities, as well as for the chemical and structural homogeneity of the finished pipes. Therefore, to produce these pipes, it is important to choose the optimal chemical composition of steel and consider that the formation of the final structure is influenced by all redistributions from smelting and casting [8] to rolling [9] and final heat treatment [10].

At satisfactory mechanical properties, determined under static tension, local weak points arise due to non-metallic inclusions and segregations of chemical elements, which can lead to SSC in a corrosive environment (Figure 1, [11]). Near a surface defect (non-metallic inclusions, solid structural components), pitting may occur [12,13], hydrogen accumulates, which is then adsorbed, diffuses into the steel and leads to its embrittlement [11–13]. If segregations and defects are developed into the depth of the pipe, then further opening and development of a corrosion crack occurs along them [11].
Due to the large number of potential causes of steel fracture during tests on SSC, a detailed study of each specific case is required to determine the causes of test failures and to develop technological recommendations to minimize them.

In this work, we studied the influence of seamless tubes microstructure with a 177.8 mm diameter and 10.36 mm wall thickness of class С110 high-strength steel 0.3C-Cr-Mn-Mo+0.15(V+Nb+Ti) to sulfide stress corrosion cracking (SSC) and sulfide stress corrosion cracking with low strain rates (SSRT) [14]. Tubes were obtained from continuous billets by screw piercing with preliminary quenching and tempering.

2. Materials and Research Methods

Samples were tested for resistance to SSC according to NACE TM0177 [15] and API 5CT [7] requirements for strength group C110—exposure in 100% hydrogen sulfide for 720 hours at a load of 644 MPa.

As additional studies, the samples were tested at various loads to determine the threshold stress: at 50%, 62.5%, and 75% of the minimum standard yield strength (MSYS) of 758 MPa. All samples, except those tested at 75% load, showed resistance to SSC (Figure 2).

The study of the corrosion resistance by the SSRT method was carried out using a universal machine UM-10T with an installed reduction gearbox, which allows stretching at speeds from $8.7 \times 10^{-8}$ s$^{-1}$ to $8.7 \times 10^{-4}$ s$^{-1}$ [14].

Mechanical properties were determined using a universal testing machine UM-10T using standard cylindrical fivefold samples. The macro-hardness of the pipe under study was determined on a transverse template using a TK-2M hardness machine by the Rockwell method.

Longitudinal specimens were cut from the pipe and from the tested samples, which were then ground and polished. Then etching with 4% nitric acid. Metallographic studies were performed using a Reichter light optical microscope equipped with a Thixomet automatic image analyzer. The microhardness of individual structural components was determined using a microhardness tester according to the Vickers method with loads from 1 g to 180 g.
The analysis of the dispersed structure and determination of the local chemical composition were studied using a Zeiss Supra scanning electron microscope equipped with an attachment for X-ray spectral analysis.

3. Results

3.1. Mechanical Properties

Testing was made on three cylindrical samples taken from different OCTG quadrants in accordance with GOST 1497 [16]. The test results are presented in Table 1. It presents the yield strength (YS), tensile strength (TS), elongation and hardness (HRC) of the studied samples.

Table 1. Mechanical properties of the OCTG samples.

| Sample | YS, MPa | TS, MPa | Elongation, % | YS/TS | Hardness, HRC |
|--------|---------|---------|---------------|-------|---------------|
| 1      | 795     | 878     | 18.7          | 0.90  | 26.0          |
| 2      | 785     | 875     | 19.1          | 0.89  | 26.5          |
| 3      | 800     | 880     | 20.1          | 0.90  | 26.0          |
| GOST 31446 | 758–828 | >793 | 18.0          | 0.86  | <30.0         |

According to the results of mechanical tests, the material of the OCTG pipe sample corresponds to the strength class C110 (ANSI/API 5CT).

3.2. Metallographic Analysis

The fracture surface of the samples after testing on SSC at 85% of the yield strength was studied using SEM (Figure 3).

Figure 3. Fracture surface of the specimen after testing on SSC 85% yield strength ($Y_{S,min}$).

Crack nucleation on the samples always occurred from side marked in red, followed by a ductile rupture zone (yellow) and a break zone. The fracture structure in Figure 3a has a brittle fracture. The structure of Figure 3b has numerous ductile fracture pits, which
is typical for crack growth zones, the structure of Figure 3c is similar to the previous one, but it has multiple secondary cracks, which corresponds to the break fracture zone.

For metallographic studies, longitudinal thin sections (along the dashed line in Figure 3) were cut out from the samples after testing on the SSC so that the fracture surface fell into the thin section. A panoramic view of the structure of the reverse part of the sample is shown in Figure 4.

![Panoramic image, ×50]

![Panoramic image, ×200]

Figure 4. Fracture surface of the specimen after testing on SSC.

Dark segregation bands were found on the sample, occupying about a third of the surface on the side of the inner pipe surface. It is from this surface that the destruction of samples always begins during tests on the SSC. Moreover, the secondary cracks located perpendicular to the axis of the specimen, according to the stress state diagram during testing, as they develop and meet the segregation band, unfold parallel to the specimen axis.

The structure of OCTG samples is composed of tempered martensite and sorbite (Figure 5a). An inhomogeneity was also found along the pipe wall thickness in the form of segregation bands located at the inner surface and occupying about a third of the thickness (Figure 5b).

![Outer surface of OCTG, ×1000]

![Inner surface of OCTG, ×1000]

Figure 5. OCTG structure at the inner and outer surfaces.

This crude structural heterogeneity results in different properties across the pipe, as measured by hardness. Closer to the outer surface of the pipe, the hardness is lower than
that of the central part but is within the permissible divergence (about 30 HV). It is important to note that closer to the inner surface, there is an unevenness in hardness caused by the presence of segregation bands. The hardness between the structural inhomogeneities differs by 40–50 HV$_{0.18}$, and in the bands, themselves reaches 325 HV$_{0.18}$. Figure 6 shows the structure of the material in dark (left) and bright (right) bands.

![Figure 6. OCTG structure in dark (left) and bright (right) bands.](image)

The structure in the segregation bands is more dispersed and contains many small non-metallic inclusions. According to the results of the chemical analysis of the dark and light bands presented in Table 2 and Figure 7, it was determined that in the segregation areas there is an increased content of molybdenum, chromium, and silicon, inherited from the zonal segregations in the central part of the continuous casting.

| Element | Dark Band Spectrum 1 | Bright Band Spectrum 2 |
|---------|----------------------|------------------------|
| Si      | 0.42                 | 0.32                   |
| Cr      | 1.29                 | 0.96                   |
| Mn      | 0.88                 | 0.93                   |
| Fe      | 95.99                | 97.07                  |
| Mo      | 1.42                 | 0.71                   |
Figure 7. Chemical composition in dark and bright bands.

Figure 8 and Table 3 compare the morphology and composition of carbides in the bright and dark bands. In the bright bands, the carbides are larger, in the dark ones, they are more dispersed. The carbides in dark band have a higher molybdenum content.

Table 3. Carbides composition, wt. %.

| Element | Dark Band | Bright Band |
|---------|-----------|-------------|
|         | Spectrum1 | Spectrum2   | Spectrum3 | Spectrum1 | Spectrum2 |
| Cr      | 3.18      | 3.41        | 1.55      | 3.79      | 6.80      |
| Mn      | 1.77      | 1.98        | 0.96      | 2.34      | 4.35      |
| Mo      | 3.02      | 2.01        | –         | 2.27      | 3.23      |
| Fe      | 92.03     | 92.14       | 96.39     | 91.18     | 84.24     |
| Ti      | –         | –           | –         | –         | 1.00      |

Moreover, crude niobium carbides were found in the dark bands (Table 4 and Figure 9). Judging by their faceted shape (Figure 5b), they were formed at the end of solidification and confirm the liquation nature of the banding found. Similar inclusions are located in the segregation bands in the area of secondary cracks, where the samples failed after SSC. Mostly, these inclusions are niobium carbides, and they are formed during rolling and heat treatment. Coarse niobium and titanium carbonitrides are formed at the end of the solidification of steel.
Figure 9. Nb-Ti Carbides in dark and bright bands.

Table 4. Carbides composition, wt.%

| Element | Spectrum1 | Spectrum2 | Spectrum3 |
|---------|-----------|-----------|-----------|
| Ti      | 1.12      | 1.12      | –         |
| Fe      | 5.76      | 7.31      | 95.13     |
| Nb      | 93.12     | 91.57     | –         |
| Si      | –         | –         | 0.45      |
| Cr      | –         | –         | 0.91      |

3.3. SSRT Testing

SSRT tests were carried out at a tensile speed of $10^{-6}$ s$^{-1}$, on standard specimens with a working part of 25.4 mm, diameter 6.35 mm. According to the test results, the samples were cracked at a load of 700, 780 and 790 MPa. Thus, the experimental results do not confirm the tests carried out according to NACE TM 0177 method A, where the breaking stress was less than 644 MPa. The reason for this result may be the uneven localization of segregation bands over the body of the OCTG, which leads a difference in conditions between the tested samples. Further testing and research are required to investigate these discrepancies and are currently in progress.

4. Conclusions

The reasons for the initiation of cracks on specimens of high-strength steel (strength group C110) after SSC testing are determined. The main reason for the destruction of samples is the presence of segregation banding (dark and bright bands of different degrees of tempering), localized on one of the sides of the sample, along which destruction occurs. Further investigation of the samples from the pipe showed that the banding is localized at the inner surface of the pipe. The dark bands have increased hardness in comparison with bright ones (320 and 280 HV, respectively), and an increased content of chromium, molybdenum and niobium. In addition to dispersed niobium, molybdenum and chromium carbides formed during rolling and heat treatment, coarse niobium and titanium carbonitrides were found in the bands, which were formed in the solidifying metal due to liquation phenomena. Further testing and research with SSRT method is required to investigate these discrepancies and are currently in progress.

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