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► To cite this version:

V. Smanio, Marion Fregonese, Jean Kittel, T. Cassagne, F. Ropital, et al.. Contribution of acoustic emission to the understanding of sulfide stress cracking of low alloy steels. Corrosion Science, 2011, 53 (12), pp.3942-3949. 10.1016/j.corsci.2011.07.041 . hal-01813683

HAL Id: hal-01813683 https://hal.science/hal-01813683

Submitted on 20 Dec 2019

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Contribution of acoustic emission to the understanding of Sulfide Stress Cracking of low alloy steels

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ABSTRACT

The Acoustic Emission technique was applied to standard tests devoted to evaluate sulfide stress cracking susceptibility of steels for oil and gas industry. The mapping of the density of AE signals vs. their location on the specimen gauge length as a function of time allowed early detection of cracking, and gave meaningful information on incubation times and propagation rates. Sulfide stress cracking initiation was correlated with the presence of critical surface defects. A mechanism involving plastic strain and/or metal dissolution was proposed to account for crack propagation.

INTRODUCTION

Hydrogen embrittlement (HE) of steels in H_2S media is a major issue in the oil and gas industry. This type of damage can lead to the failure of important production equipment. Several kinds of damage may be caused by H_2S in contact with low alloy steels [1]. One of the main cracking processes is Hydrogen Induced Cracking (HIC), which is caused by the accumulation of hydrogen at internal traps, such as non-metallic inclusions, regions of anomalous microstructure produced by the segregation of impurities and alloying elements, and other weak interfaces (e.g. the boundary between pearlitic and ferritic phases). When hydrogen concentration reaches a critical value at the defect, recombination to gaseous hydrogen occurs, with the corresponding build-up of high pressure gas leading to internal cracking. Under conditions of applied or residual tensile stresses, HIC cracks can become aligned in the through thickness direction resulting in 'ladder like' crack arrays referred to as Stress Oriented Hydrogen Induced Cracking (SOHIC). This phenomenon is usually associated with low-strength welded ferritic pipe and pressure vessel steels.

The present work is focused on a third kind of wet H₂S cracking mode: Sulfide Stress Cracking (SSC). SSC leads to early failure of the metal due to the combined effect of both hydrogen penetration and tensile stress (residual and/or applied). Crack initiation of SSC always occurs on the surface. High strength metallic materials and hard welded zones are prone to SSC. Although the mechanisms proposed to account for HIC and SOHIC are generally well accepted, the SSC mechanism is still controversial, with several models proposed in the literature. A model based on the reduction of cohesive forces induced by hydrogen was introduced by Troiano [2] and further developed by Oriani [3]. Hydrogen insertion in solid solution would decrease the forces required to fracture the metal along the crystal lattice i.e. decrease cohesion forces of the lattice and the energy necessary to induce cleavage, in particular at defects and interfaces. Other models are based on hydrogen induced local plasticity. Beachem [4] proposed that hydrogen absorbed at the crack tip enhances dislocation glide and therefore plastic strain processes leading to metal fracture. In the hydrogen enhanced localized plasticity (HELP) model, hydrogen increases dislocation mobility. Dislocations can then pile-up more easily, resulting in a distribution of microscopic highly strained areas surrounded by less ductile areas. Fracture is then micro-ductile, with a brittle appearance [5]. Similarly, Lynch [6] proposed the weakening of interatomic bonds by hydrogen adsorption at the crack tip. Localized plastic strain would then lead to the formation of microvoids ahead of the crack tip. Consequently, crack propagation would occur by microvoid coalescence.

In H₂S environment, Crolet [7] proposed a high energy process where proton absorption into the metal induces a dilatation of the crystalline lattice and a local increase of triaxial stress within the first surface layers. The applied stress would then induce SSC initiation and propagation. Moreover, proton absorption within the metal would be enhanced by H₂S itself via the formation of adsorbed HS⁻ on the metallic surface. On the other hand, in the presence of H₂S, a scale of iron sulfide is formed on the surface of the steel. The first stage of iron sulfide formation consists in the nucleation and growth of mackinawite that can be considered as stoechiometric FeS [8]. In longer term exposures, a multi-stage film growth mechanism is involved where mackinawite forms and ruptures. Then, depending on the exposure time, the temperature, the H₂S and the chloride contents, the pyrrhotite and pyrite forms of iron sulfide can grow. Mackinawite was demonstrated to be non protective towards corrosion [9][10] or even suspected to promote corrosion, in incorporating FeSH^+_{ads} ion as: $\text{FeSH}^+_{ads} \rightarrow \text{FeS} + \text{H}^+$. FeSH^+_{ads} formation results from sequential chemisorption followed by anodic discharge: Fe + $\text{H}_2\text{S} \rightarrow \text{FeSH}^-_{ads} + \text{H}^+$ then $\text{FeSH}^-_{ads} \rightarrow \text{FeSH}^+_{ads} + 2\text{e}^-$. [11]. FeSH^+_{ads} can also suffer hydrolysis liberating Fe^{2+} : $\text{FeSH}^+_{ads} + \text{H}^+ \rightarrow \text{Fe}^{2+} + \text{H}_2\text{S}$.

H₂S induced damage evaluation usually consists of ex situ examination of fractured specimens after standard tests. However, this methodology is often insufficient to identify and deeply study the different steps of crack propagation, particularly in the case of SSC. Therefore, the use of an in-situ evaluation method appears to be an interesting development. Among the possible in-situ methods, acoustic emission (AE) exhibits interesting potentialities for corrosion study and monitoring. Indeed, the detection possibility was established and good correlations were obtained between specific AE parameters and the amplitude of corrosion damage in various cases: stress corrosion cracking [12-15], abrasion or erosion corrosion [16], pitting corrosion [17-24], crevice corrosion [25-27], exfoliation corrosion [28], uniform "acidic" corrosion [17][29-30], and also to monitor cracking in sour media [31-36]. Weng et al. [31] found a correlation between HIC damage evaluated by Crack Length Ratio (CLR) [37] and the AE cumulative energy level. Cayard et al. [32] and also Gingell et al. [33] showed that during sour exposure tests under stress, a higher amount of cumulative AE energy was recorded for the experiments leading to fracture compared to non-failure tests. This critical energy build-up associated with specimen failure remains dependent on the steel tested and on the applied stress level. They also showed that AE could be used to discriminate between the initiation and propagation stages of SOHIC.

As far as AE detection of SSC is concerned, most of the published works [34-36] are based on monitoring during standard SSC tests described in NACE TM0177-96 [38]. From these studies, AE appears to be a powerful tool for monitoring H₂S cracking. Some correlations between the AE energy and the extent of cracking were found. However, for most of the past studies, the cracking process was not always clearly identified or documented, especially for SOHIC and SSC, which were not systematically differentiated. Furthermore, the AE signals associated to the cracking process itself were not discriminated. Considering this last point, a step forward was recently reported by the authors [39-40]. Indeed, the AE contributions from H₂ evolution due to acid corrosion, FeS layer build-up, and cracking, all constituting AE sources, were separated by identifying the acoustic signature of each process. This discrimination was performed by isolating physically the active sources through specific

electrochemical polarization procedures. The characteristics of the corresponding AE signals were identified. The most discriminating representation was obtained by plotting absolute energy of the signals as a function of their duration (Figure 1). On this correlation chart, it was possible to separate AE events related to HIC from those related to the corrosion reactions, allowing a better analysis of cracking associated signals. However, this discriminating analysis demonstrated that SSC development led to the emission of AE signals that could not be directly identified on the same energy-duration correlation chart [39]. For that particular process of HE, a specific analysis giving the evolution of the density of AE signals vs. their location and the time of test was necessary.

In that context, the aim of the present work is to validate this new discriminating AE analysis of SSC, and to establish its performance for better understanding and evaluation of this particular H₂S induced cracking process.

EXPERIMENTAL PROCEDURE

Material

This study was conducted on a C110 steel grade usually used as well tubing material. The chemical composition of the steel was analyzed by optical emission spectroscopy (OES), except for carbon and sulfur, analyzed by a chemical method (Table 1). The C110 grade contains alloying elements, such as chromium and molybdenum, providing a good resistance to hydrogen embrittlement. Tensile properties of the steel in the longitudinal direction are given in Table 2. The microstructure of C110 grade is a tempered martensite with macrosegregation. This microstructure is usually resistant to HIC [41]. Two surface states were studied: i/ mechanical polishing as recommended in NACE TM0177-96 standard (final surface roughness below 0.81 μ m) [38] and ii/ electrochemical polishing in 42 %vol. phosphoric acid and 34 %vol. sulphuric acid at 1A for 30 min. In this latter case, diameter of the as-machined samples was reduced by 0.8 %.

Experimental set up

The experimental set up (Figure 2) was based on the standard tensile test described in NACE TM0177-96 (method A) [38]. Constant load was applied with the proof ring. NACE standard specimens were adapted to allow positioning two AE sensors at the bottom of the specimen. The geometry of the gauge section of the standard specimen was unchanged (length = 25.4

mm, diameter = 6.35 mm). During the test, only this section was immersed in the corrosive solution. Tests were conducted in buffered solution according to EFC 16 [42] (50 mg/L NaCl and 4 mg/L CH₃COONa in distilled water) and under 1 bar H₂S. De-aeration of the solution was performed by nitrogen bubbling ensuring less than 15 ppb by weight dissolved oxygen before H₂S introduction. The corrosive fluid (sour media) was circulated between the test cell and a separate glass vessel, in which H₂S bubbling was maintained. This procedure avoided any interfering AE noise detection (e.g. gas bubbling in the test cell would result in AE) and to maintain constant electrochemical conditions in the cell during the test (easier pH control). The pH was measured with a pH meter located in this glass vessel and buffered every day by HCl or NaOH additions.

AE instrumentation consisted of a transducer, a preamplifier and an acquisition device (MISTRAS with AEWIN software from Physical Acoustic Corp.). The gain of the preamplifier was set to 40 dB. A band pass filter with frequency range from 100 to 400 kHz was used. In this study, the threshold was set to 28 dB. The transducers were Nano 30 type from PAC (piezo-electric disks). For each detected AE signal, several acoustic parameters were calculated and recorded, e.g. event amplitude, duration and energy. For each AE signal, the location of source on the specimen was determined by measuring the delay of event detection time between the two sensors. Spatial and temporal post-filtering treatments were also made on the AE data. Only the AE signals located on the gauge section and detected before failure are taken into account for the analysis. The analysis of AE data was first performed according to previous works [39], allowing identifying three typical sources: i/ HIC, ii/ H₂ evolution at the steel surface, iii/ FeS build-up (see Figure 1). In a second configuration, another procedure was applied by using source location. Because the two sensors were placed at either extremity of the specimen, it was possible to determine the position of the source for each AE signal detected on both sensors. This was achieved by comparing the times at which an identical AE signal was detected on each sensor. Then, knowing the distance between both sensors, the position of sources is obtained. Using AE source position as additional parameter, the analysis of AE data consisted in mapping the density of AE signals as a function of position on the gauge section of the specimen and as a function of the time during the test.

RESULTS

Table 3 summarises the different sets of experimental conditions and the observed times to failure. Condition 1 consists of H₂S exposure without applied load. Therefore, no SSC is expected. The goal of this test is to validate that the tested steel is not sensitive to HIC. Conditions 2 and 3 consist of 90 %YS strained specimens exposed to H₂S, with two distinct surface preparations. Corresponding AE results, plotted on absolute energy vs duration correlation charts are shown on Figures 3a, 3b and 3c for conditions 1, 2 and 3 respectively. According to previous results [39], analysis of AE data for condition 1 (Figure 3a) suggests that the only active AE sources are hydrogen evolution and the FeS layer. Both metallographic examination and ultrasonic analysis of the specimens confirmed the absence of HIC phenomena. For condition 2 (Figure 3b), when a constant load is applied on an electropolished specimen, no failure was obtained after 800 h testing and no crack was detected by metallographic examination. AE results are in agreement with these observations: according to previous results [39], only populations of AE signals associated with H₂ and FeS are identified on the correlation chart. For condition 3 (Figure 3c), all of the mechanically polished specimens failed. Times to failure varied between 64 and 200 hours (Table 4). Figure 4a shows a typical fracture surface observed by SEM. Two areas can be identified. Fracture seems to be initiated on the surface of the sample perpendicular to the loading direction (area 1). A brittle type transgranular fracture is observed in area 1, (Figure 5) while microvoids typical of a ductile failure are identified only in area 2 (Figure 6). A profile of the fracture surface along the line drawn on Figure 4a can be extracted from a 3D reconstruction of the surface using a 3D microscope (Figure 4b). Area 1 which is brittle-like corresponds to a flat surface perpendicular to the direction of applied load. Area 2, identified as ductile, corresponds to the 45° region. This profile is typical of a SSC rupture: initial hydrogen assisted crack corresponds to the 90° brittle area. Then, as the cross-sectional area of the loaded part is reduced, the applied stress reaches the UTS and the sample fails by 45° ductile fracture. Corresponding AE results are plotted in the energy vs. duration plane on Figure 3c. In this correlation chart, we cannot distinguish any new population of signal which would correspond to SSC. Either there is no specific acoustic emission related to SSC, or the corresponding signals do not differ from H₂ and FeS signals by energy and duration characteristics. This result is in agreement with previous results providing evidence that SSC signals can not be discriminated from H₂ and FeS signals on absolute energy-duration correlation charts [39]. Therefore, other approaches were tested to extract AE information related to SSC. The best results were obtained by mapping the density of AE signals as a function of position on the gauge section of the specimen and of the time of the test (Figure 7). The density of AE signals is quite homogeneous on the whole gauge length, except in well defined zones where it is higher for a certain period of time. The direct reading of this figure indicates that an AE activity well above the background level was detected at some location and after a certain time. More remarkably, such intense AE activity was always found at the position corresponding to the final failure. Surface examination was also performed at other areas where high AE activity was found, and crack initiation was detected (Figure 8). It could therefore be concluded that these high density signals could be associated with SSC initiation, as previously demonstrated [39].

In order to obtain time and energy information on AE signals associated with SSC, characteristics of AE signals located on the final rupture area (zone 1 on Figure 9) are compared with those of signals located just above the final rupture area, where neither cracking nor increase in AE signals density were detected (zone 2 on Figure 9). Both zones have a surface area of 60 mm². For each zone, both number and energy of AE signals are plotted versus time (Figure 10). During the first part of the test, signals recorded on both areas (zone 1 and zone 2) exhibit the same evolution. From the start of the test to 370 ks, the same cumulative AE activity is recorded for both areas of the specimen. However, from 370 ks until final fracture, the number and energy of AE signals increase drastically on the area of the specimen corresponding to the final fracture (zone 1). At the same time, AE activity of the other area continued to increase at the same rate (zone 2). At the end of the test, 2.5×10^4 signals have been recorded on zone 2 with a cumulative energy of 5 10^5 aJ, with a constant increase, compared to 10^5 signals and 8.2 x 10^5 aJ on zone 1, where the final rupture is located, with a sharp increase of both parameters when SSC initiates. Comparing the evolution of signals recorded on both areas, it can be assessed that a signal related to hydrogen or FeS layer evolution (the only signal type that is detected on zone 2) carries an average energy of 20 aJ, whereas a signal associated to SSC development (detected on zone 1, together with H₂ and FeS layer evolution signals after SSC initiation) carries an average energy that is 5 times lower ((8.2-5):(1-0.25), see Figure 10). SSC propagation is therefore associated with a sharp increase in AE signals density, but SSC associated AE signals have low energies.

From the representation of the location of the density of detected AE signals on the gauge length of the specimen versus the time of test (Figure 7), it is also possible to define the SSC incubation time as the time before signals density increases locally, together with the propagation time of the cracks until final failure. On the other hand, the observation of the brittle areas on the fracture surfaces of the failed specimens (Figure 5) allows measuring the SSC depth and therefore calculating the SSC propagation rate. SSC propagation rates that can be evaluated with this methodology exhibit good reproducibility: $2 \times 10^{-8} \pm 0.5 \times 10^{-8}$ m.s⁻¹. From the representation of the location of the density of detected AE signals on the gauge length of the specimen versus time (Figure 7), it is also possible to define the SSC incubation time as the time before signal density increases locally, together with the propagation time of the cracks until final failure.

DISCUSSION

The present results demonstrate that the steel grade studied here is not prone to HIC when tested without external stress. Metallographic examination, as well as ultrasonic scans showed the absence of internal cracks, in agreement with AE signals analysis: no HIC associated AE signals were detected in any experiments. Both the tempered martensite microstructure and chemical composition of C110 grade can account for its good HIC resistance [41]. When C110 specimens were tested under 90 %YS applied load at pH 3.5 under 1 bar H₂S, most experiments led to SSC fracture. Acoustic Emission analysis consisted first to apply a methodology that gave excellent results in HIC analysis [39-40]. For all SSC experiments, no AE data typical of HIC were detected. This is a clear confirmation that SSC and HIC proceed by different mechanisms. However, another type of analysis of AE data proved the ability of the AE technique to detect differences between failed and non failed experiments. Also, the representation of the density of detected AE signals on the gauge length of the specimen versus the time of test (Figure 7) allowed locating the initiation sites and the final failure zone tens of hours before failure. Early detection of SSC by AE was therefore confirmed to be possible. More detailed analysis of AE signals associated with SSC revealed that most of them seem to have a low energy compared to H₂, FeS and HIC AE signals [40]. When HIC propagation occurred, high AE energy jumps associated with decohesion of interfaces exhibiting cleavage features were observed. From the present AE results, it was therefore confirmed that SSC does not initiate as sub-surface HIC, which is now well recognized. Surface preparation of the specimen had an impact on SSC. Electropolishing provided a higher resistance to SSC vs. mechanical polishing performed as recommended by NACE TM0177-96 [38]. Indeed, for all electropolished specimens, no fracture was observed after more than 720 h. testing. Furthermore, no evidence of SSC initiation was detected by acoustic emission, and no crack was observed at the surface of the specimen by optical microscope

observation. On the other hand, all mechanically polished specimens failed. Fracture surface analysis confirmed that cracking occurred through a SSC rather than a SOHIC mechanism. Therefore, surface preparation appears to be a first order factor for SSC initiation.

SSC initiation:

At the beginning of exposure tests, homogeneous AE data was recorded on the whole specimen surface. This AE activity could be linked with a randomly distributed localised corrosion. Indeed, cross section observations of specimen areas exhibiting a high AE signal density revealed the presence of pits prior to crack propagation (Figure 11). After a certain exposure time, higher AE activity at specific locations at the steel surface was detected. This was associated with crack initiation. The time before such local increase of AE activity was detected was scattered between replicate experiments. This suggests that initiation is not linked with the accumulation of a certain level of absorbed hydrogen. it is more likely that initiation starts at surface defects. Indeed, initiation can occur after more than 100 hours testing, whereas estimations of hydrogen charging kinetics have shown that a HIC standard massive specimen (X65 grade) is saturated with H₂ in about 10 hours in similar experimental conditions [43]. On the contrary to HIC initiation, hydrogen charging time of the material is therefore not the limiting factor for SSC initiation. This explains the importance of surface preparation for SSC specimens. Cernoky et al. [44] demonstrated the importance of specimen surface preparation when tested in acidic H₂S media under constant load. The steel specimens exhibited an increasing resistance to SSC when chemically polished compared to mechanical polishing, itself compared to as-machined surface state. According to Cernoky [44], machining was responsible for the existence of a sub-surface strain-hardened zone, at the origin of SSC susceptibility. Another author [45] also observed that cracking hardly initiated on smooth surfaces in the presence of hydrogen. According to this author, a localized corrosion enhancing pitting would promote crack initiation. Indeed, mackinawite scale growth involves the formation of FeSH ads which can generate a porous sulfide layer and increase the localized corrosion of the steel [11]. It was reported elsewhere that the formation of such a porous sulfide deposit could retain the pit electrolyte within the pits and propagate pit growth [46]. Moreover, immersion of steel in H₂S containing solutions leads to the formation of a rough surface [11]. In the present case, pitting initiation could be enhanced by the presence of H₂S, as reported for martensitic stainless steel [47], or the presence of a relatively rough surface could constitute occluded cells and play the part of initiating pits in which acidic electrolyte can be retained.

SSC propagation:

In the present work, the SSC propagation step was defined to start when a significant local increase of AE signals density was detected. Therefore, the duration of crack propagation could easily be calculated, considering the length of the brittle zone of the fracture surface. From the above results, the corresponding duration of SSC propagation was about 30-35 hours in the present experimental conditions. AE data that were detected in the crack area several hours before the final fracture was detected. They exhibited low energy ($< 10^2$ aJ) and low duration (< 1500 µs), as compared to HIC AE data [39-40]. This could be an indication that SSC propagation involves local plastic strain [4-6], and/or dissolution on the microscale, which is known to be hardly detectable by AE [13][48]. This dissolution mechanism could also account for the difficulties to discriminate SSC AE signals from the signals related to H_2 and FeS, since these two last corrosion processes are also closely linked to anodic dissolution. Evaluation of crack propagation rates was obtained from AE data analysis and SEM observations of fracture surfaces. Although times to failure appeared to be scattered (Table 4), propagation rates lie in the range of 2 x $10^{-8} \pm 0.5$ x 10^{-8} m.s⁻¹ for all tested specimens. In the literature, SSC propagation rate values have already been evaluated through DCB tests. Ernst et al. measured propagation rates ranging from 10^{-7} to 10^{-8} m.s⁻¹ on a C110 steel grade in sour media [49]. Szklarz et al. obtained a 10⁻⁹ m.s⁻¹ propagation rate in the same conditions for a T95 grade steel [50]. The average propagation rates obtained in the present work from AE are therefore in good agreement with literature results. Yet, it is worth noting that during constant load tests, crack propagation is probably not uniform, as suggested by AE results. Indeed, local high AE signal density tended to decrease until failure (Figures 7 and 10), which could indicate a change in the propagation rate. Interrupted tests before failure would then be helpful to obtain a better understanding of the details of the cracking process. The time when the test has to be interrupted could be determined by AE monitoring.

CONCLUSIONS

This work was aimed at investigating the different propagation stages of SSC in a low alloy steel in the presence of wet H_2S , by the use of an acoustic emission technique. The first conclusions of this study concern the acoustic emission technique and its performance for the detection of H_2S assisted cracking:

1. As expected, AE characteristics of SSC exhibits different features from HIC: HIC is associated with high energy AE signals, whereas an emission of low energy signals accompanies SSC propagation.

2. A representation of the density of AE signals *vs.* their location on the gauge length of the specimen and the time of test allows obtaining reproducible, quantitative information on the incubation-initiation time and on the propagation rate.

Initiation and propagation mechanisms were analysed the light of the AE data recorded during SSC experiments.

3. Initiation of cracks is closely linked to surface preparation of the material. It depends on the presence of defects and/or strain-hardened zone at the surface of the metal. SSC initiation is not related to hydrogen saturation of the steel, but to the formation of a critical pit, associated with a surface defect.

4. The low energy of AE signals indicates that SSC propagation is not the result of subsurface HIC. It is suggested that SSC propagation is the result of a cracking mechanism involving hydrogen embrittlement associated with local plastic strain and/or metal dissolution.

ACKNOWLEDGEMENTS

The authors would like to thank M. Boinet (Euro-Physical Acoustics), for his help in the analysis of AE data, and G. Parrain for his technical support. H. Marchebois (Vallourec) is also acknowledged for helpful discussions on H₂S cracking mechanisms, as well as M. Taylor from PennState University and J.L. Crolet.

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Tables

Table 1. Chemical composition (wt %) of C110 steel.

С	Mn	Si	Р	S	Cr	Ni	Mo	Cu	Nb
0,309	0,394	0,343	0,015	0,002	0,964	0,037	0,834	0,018	0,033

Table 2. Mechanical tensile properties of C110 steel (tests conducted according to EN10002-1 standard) (YS: Yield Strength, UTS: Ultimate Tensile Strength, El: Elongation)

YS(MPa)	UTS (MPa)	El%
798	887	20

Table 3. Experimental conditions and corresponding times to failure.

Experimental Conditions	рН	pH ₂ S (bar)	Applied load (%YS)	Surface preparation	Time to failure
1	3,5	1	0	Mechanical polishing	No failure No HIC cracks
2	3,5	1	90	Chemical polishing	No failure (800 h)
3	3,5	1	90	Mechanical polishing	[64-200h]

Table 4. Cracks incubation time, propagation time and propagation rates

Test #	Time to failure (h)	Incubation time (h)	Propagation time
			(h)
1	132	99	33
2	140	104	36
3	90	60	30
4	113	85	28
5	200	121	79
6	64	5	59
7	116	83	34

Figure 1. Correlation chart representing absolute energy as a function of signals duration for a test conducted in the EFC 16 solution at pH 4.5 under 1bar H_2S , on a X65 SwS specimen [33].

Figure 2. Experimental device.

Figure 3. Correlation chart representing absolute energy as a function of signals duration for a test conducted in the EFC 16 solution at pH 3.5 under 1bar H_2S (a) on a C110 specimen without applied load – (b) with an applied stress of 90% YS on a C110 chemically polished specimen – (c) with an applied stress of 90% YS on a C110 standard polished specimen

Figure 4. (a) Fracture surface of C110 steel tested in the EFC 16 solution at pH 3.5 under 1bar H_2S with an applied stress of 90% YS – (b) Profile of the fracture surface (red line on figure 5a).

Figure 5. Detail of the brittle area (1) of C110 fracture surface.

Figure 6. Detail of the ductile area (2) of C110 fracture surface.

Figure 7. Density representation of the location of all the AE signals in function of time (C110 steel, pH 3.5, 1 bar H₂S, σ =90% YS).

Figure 8. SEM observation of the area identified by a circle on the figure 10.

Figure 9. Amplitude of AE signals *vs.* their location on the gauge length of the specimen: identification of AE signals located on the final rupture area of the gauge length (zone 1 - x between 6.7 and 7.0 cm reported from the position of the sensor located at the bottom of the specimen) and AE signals located just above the final rupture area, where no crack neither no increase in AE signals density were detected (zone 2 - x between 6.3 and 6.6 cm).

Figure 10. (a) Number and (b) cumulative energy of AE signals *vs.* time of test for zones 1 and 2.

Figure 11. Cross section observations of high AE density areas.





























