Real Time Imaging of Deuterium in a Duplex Stainless Steel Microstructure by Time-of-Flight SIMS

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For more than one century, hydrogen assisted degradation of metallic microstructures has been identified as origin for severe technical component failures but the mechanisms behind have not yet been completely understood so far. Any in-situ observation of hydrogen transport phenomena in microstructures will provide more details for further elucidation of these degradation mechanisms. A novel experiment is presented which is designed to elucidate the permeation behaviour of deuterium in a microstructure of duplex stainless steel (DSS). A hydrogen permeation cell within a TOF-SIMS instrument enables electrochemical charging with deuterium through the inner surface of the cell made from DSS. The outer surface of the DSS permeation cell exposed to the vacuum has been imaged by TOF-SIMS vs. increasing time of charging with subsequent chemometric treatment of image data. This in-situ experiment showed evidently that deuterium is permeating much faster through the ferrite phase than through the austenite phase. Moreover, a direct proof for deuterium enrichment at the austenite-ferrite interface has been found.

The popularity of Duplex Stainless Steel (DSS) arises from an attractive combination of properties, including high strength and excellent corrosion resistance. Due to these properties, this type of steel is being used in many applications and environments which frequently provide critical conditions for hydrogen absorption in metallic microstructures1. By cathodic charging or during free corrosion of DSS in hydrogen environments retarding recombination, overall hydrogen concentrations of more than 40 at.ppm have frequently been observed and found to be harmful2,3. Dependent on the respective solution and trapping mechanisms, hydrogen taken up from various sources might cause significant degradation of the mechanical properties and consequently might entail hydrogen assisted cracking (HAC) of components, as documented by severe failure cases. Although HAC represents an important issue in industrial applications and attracted enormous research attention for decades, the mechanisms of hydrogen assisted crack initiation and propagation as well as those of hydrogen transportation in the respective microstructures are not yet fully understood4–7. Local in-situ characterization of the hydrogen distribution in steel microstructures represents a key for elucidation of HAC as well as hydrogen transportation in the respective microstructures8,9. One, if not the only, laterally resolving method of chemical analysis for imaging the locally hydrogen distribution and interaction in a specific microstructure is secondary ion mass spectrometry (SIMS)10. Up to the present, all of the SIMS experiments aiming on the elucidation of hydrogen effects were carried out using externally charged samples and dynamic SIMS instruments (for a survey cf. SI). In contrast, real time imaging results have been obtained in the present study by static ToF-SIMS monitoring of the deuterium distribution in a DSS microstructure while continuously charged for the first time (cf. Fig. 1). In the experiments, deuterium that permeated through the steel’s microstructure has been imaged at the outer surface of an electrochemical permeation cell made from DSS itself which is located within the ultra-high vacuum chamber of the instrument (for details see SI). Analysis of that vacuum/steel interface avoids any impact of the electrolyte on the imaged surface. In the permeation experiment, the DSS surface has been investigated with increasing time of charging: First deuterium secondary ions were detected after 28 days of charging, saturation after 37 days. The limited sensitivity of SIMS (secondary ion emission probabilities may be small as $10^{-4}$) leads to an underestimation of the “break-through” time which is the time of charging required for detecting the first intentionally
introduced deuterium on the analyzed surface. To gain a maximum of information from the SIMS image data acquired multivariate data analysis by Principal Component Analysis (PCA) has been performed. By using key co-variations between element and fragment secondary ions the image contrast between features is substantially increased. In comparison to a previous SIMS study of deuterium effusion from the DSS sample reported by the authors\(^1\) the steps beyond the present experimental state are (i) the real time monitoring of the permeation, diffusion and distribution of deuterium in individual phases of the DSS microstructure and (ii) the multivariate treatment of SIMS image data.

The use of the hydrogen isotope deuterium emerges from the requirement to distinguish between hydrogen species diffused through the microstructure of DSS to the imaged surface and hydrogen containing residual gas species adsorbed thereon. Of course, there are differences in diffusibility and solubility for both hydrogen isotopes, but these range within the same order of magnitude. It is well-known that, in principal, deuterium behaves very similar to hydrogen in steel microstructures\(^7,10,12–14\).

Results of the in-situ permeation experiment are displayed in Fig. 2. Figure 2a shows the scores image for the first principle component (PC1) obtained from the positive secondary ion data set comprising matrix related secondary ions. The related loadings plot (cf. Figure S2 in the SI) indicate Ni\(^+\), Mn\(^+\) and Fe\(^+\) as key secondary ions that characterize the variance captured by PC1 obtained from the positive secondary ion data set. Thus the different phases in DSS can clearly be distinguished. The ferrite appears dark and the austenite appears bright. Figure 2b exhibits the PC1 scores image delivered by PCA of the negative secondary ion data set where deuterium related secondary ions contribute to. In this case the related loadings plot (cf. Figure S2 in the SI) indicates D\(^-\) and OD\(^-\) as key (fragment) secondary ions that characterize the variance captured by PC1. By comparison of Fig. 2a,b a preferential deuterium accumulation in the ferrite phase becomes evident. For single grains inside austenite and ferrite phases, intensity variations in Fig. 2b have been observed. This observation is assumed to be due to the dependency of the secondary ion emission probability of deuterium related secondary ions on the respective grain lattice plane exposed to the impinging primary ions.

Based on Fig. 2a,b, a higher permeation of deuterium and faster saturation of the ferrite phase is concluded when compared to the austenite phase. This experimental result is consistent with literature data, where deuterium in the ferrite phase is reported to have an about five orders of magnitude higher diffusivity\(^7,15,16\). By this, the current in-situ measurements provide evidence to previous statements\(^15,17,18\) based on models and experiments that deuterium preferentially diffuses through the ferrite matrix. In accordance to the diffusion coefficient\(^19\), within the time frame of the reported experiment, only a minor amount of deuterium will be diffusing into and trapped in the austenite grains at considerably lower diffusion coefficients.

Reduction of the negative image data matrix used for score images displayed in Fig. 2b by the use of only a mass range 20–50 m/z reveals images shown in Fig. 2c,d. Figure 2c presents an image of the scores of PC1 where significant positive loadings have been found only for m/z = 26, i.e. CN\(^-\) fragment ions (cf. Figure S2 in the SI). Comparison of Fig. 2c with Fig. 2a reveals that these occur in areas corresponding to the austenite phase. Moreover, the comparison of Fig. 2c with Fig. 2b reveals that certain regions in the ferrite phase where a bright region related to CN\(^-\) correlates with a bright area representing high deuterium concentrations.

Figure 2d displays the PC2 scores image belonging to the PC1 image in Fig. 2c. The loadings plot for PC2 shows strong positive loading exclusively at m/z = 43 assigned to CN\(_2\)HD\(^-\) (cf. Figure S2 in the SI) and displays distinct positive scores at grain boundaries within a specific phase and high positive scores at ferrite-austenite
Figure 2. Principle component score images of the ToF-SIMS data sets acquired after 34 days of charging. 
(a) PC1 scores image for identifying phases using the positive ToF-SIMS data set acquired after 34 days of charging DSS with deuterium by using a peak list with secondary ions covering the main alloying elements. The bright regions represent the austenite phase (as proved by the loadings plot in shown Figure S2a). (b) PC1 scores image identifying locations rich in deuterium obtained from the negative SIMS data set acquired after 34 days and using the full peak list. Bright pixel indicate positive scores and correlate with the ferrite phase. (c) PC1 scores image obtained from the same data set but using a different peak list prepared to focus on carbides and
nitrides. This image is identifying locations rich in N and C. Bright pixels indicate positive scores and correlate with the austenite phase. (d) PC2 scores image identifying locations rich in N, C and D (bright pixel means positive score) indicating graded accumulation of deuterium at the ferrite and austenite grain boundaries and at the ferrite/austenite interface (highest scores) as well. All ToF-SIMS image acquisition settings and parameters are given in table S3 in SI.

interfaces. This image provides evidence that regions at the grain boundaries and interfaces contain carbon and nitrogen as well as accumulated deuterium at the same time. Such findings indicate that grain and phase boundaries act as sinks in the microstructure which attract hydrogen or deuterium ions.

The SIMS data sets leading to images displayed in Fig. 2 have been taken from a region of interest (ROI) selected at the beginning of the in-situ permeation experiment which has been analyzed many times before to identify the break-through of deuterium. To rule out any unintended measurement artifacts (e.g. due to beam damage) a fresh ROI was selected for ToF-SIMS imaging after 37 days of charging (shown in Fig. 3). Again, the ferrite phase appears dark and the austenite is bright in Fig. 3a. Also, the significantly higher deuterium concentration in the ferrite grains in comparison to the austenite has been confirmed by Fig. 3b. Similar results to those obtained after break-through of deuterium have been found for the saturated state. Additionally an ion-induced secondary electron image (Fig. 3c) of the duplex microstructure at the new ROI is taken where also phase and even grain boundaries can be distinguished.

Principally the in-situ permeation experiment should enable also a determination of kinetic data as, e.g., overall diffusion coefficients. Unfortunately quantitative evaluation of such values will be limited by the detection limit of the SIMS instrument for deuterium and the tortuous effect occurring in a two phase alloy. However, it has to be expected that most of the deuterium detected at the time of the break-through permeated predominantly through the ferrite phase on paths circumventing the austenitic phase (cf. ref. [15]) characterized by considerably lower diffusivity and higher solubility for deuterium.

It has been demonstrated for a technically relevant multiphase metallic microstructure, here DSS, that imaging ToF-SIMS together with subsequent image data treatment by PCA is a powerful approach for phase identification (positive SIMS mode) and in-situ visualizing the deuterium distribution (negative SIMS mode). The in-situ approach using a deuterium charging cell in the analysis chamber of a ToF-SIMS instrument provided clear experimental evidence for a preferred diffusion of deuterium through the ferrite phase, while ex-situ SIMS studies could only show preferential accumulations in the phases subsequently to charging with hydrogen or deuterium.

The experiment also enables an investigation of interactions of deuterium with grain and phase boundaries, carbon and nitrogen in the DSS microstructure. However, it addresses the diffusion and permeation of deuterium through the microstructure without any external mechanical loading on the specimen. It is anticipated that, under these specific circumstances, deuterium accumulates in the lattice and is trapped at dislocations and already existing twins with much higher solubility and lower diffusivity of hydrogen/deuterium in austenite as compared to ferrite. The authors currently develop an advanced experiment aiming on the realization of an in-situ mechanical loading of deuterium charged specimens to study the degradation process of each phase under these more complex circumstances by imaging ToF-SIMS.

The real time and in-situ imaging ToF-SIMS based approach introduced in this letter widens the perspectives for experiments aiming on deuterium and hydrogen (or even other mobile constituents) behavior in a range of relevant metallic microstructures. It will contribute to a better understanding of hydrogen and deuterium permeation and subsequent cracking in metallic microstructures known to be crucial for the design of engineering components in industrial branches with high impact as petrochemical, off-shore constructions and oilfield applications.

Finally it should be mentioned that the in-situ ToF-SIMS experiment reported here is part of recent community overarching activities to develop new experiments e.g. on the big and flexible sample holder within the analysis chamber of ION TOF’s ToF-SIMS instruments. One example is a microfluidic device for in situ ToF-SIMS analysis of the Liquid/Vacuum interface introduced in ref. 23.

Methods

ToF-SIMS analysis procedure. ToF-SIMS measurements were performed using a ToF-SIMS IV instrument (ION-TOF GmbH, Muenster, Germany). 25 keV Bi\(^+\) primary ions delivered by a liquid metal ion gun (LMIG) were used for analysis. A sputter ion gun providing 3 keV Ar\(^+\) ions for sputtering has been used for cleaning the region of interest. Previously to the measurements an area of \(300 \times 300 \mu m^2\) was sputtered by Ar\(^+\) primary ions for 5 min to remove surface contaminants originating from the cell production process. ToF-SIMS data from the surface of the steel were acquired from 100 × 300 \(\mu m^2\) sample ROI performing the burst alignment mode with a beam diameter of approximately \(\sim 250 \text{nm}\) providing sufficient lateral resolution for imaging the steel’s micro structure. The information depth of the SIMS method equals to the first and second monolayer of the steel surface.

The first step of the experiment was to image the finely polished DSS surface of the cell before charging by ToF-SIMS in positive mode. A representative ROI of 100 × 100 \(\mu m^2\) was selected to identify the austenitic and the ferritic phases by the detection of Cr\(^+\), Mn\(^+\) and Fe\(^+\) secondary ions; first with instrument settings delivering high mass resolution and thereafter with settings for high lateral resolution. Subsequently, negative secondary ion images were acquired from one and the same ROI at increasing times of charging with deuterium over a period up to saturation (37 days).

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PCA and data preprocessing. Image data were treated by PCA (for details see SI). PCA was performed using the Solo+Mia toolbox (v7.5.2, Eigenvector Research Inc.) which were run under MATLAB (v7.9.0.529, MathWorks Inc.). Prior to principal component analysis the raw data of secondary ions were pre-processed by normalization to the total intensity of each respective mass spectrum, Poisson scaling and mean centering finally.

Design and details of the in-situ charging experiment. The results have been obtained by a novel experimental approach, i.e. by mounting a specifically designed cell (Fig. 1) in the ultra-high vacuum (UHV) analysis chamber of the ToF-SIMS instrument. The cell was made using a rolled cylindrical bar of a duplex stainless steel grade 2205 (UNS S31803, EN 1.4462). The material was purchased from ThyssenKrupp and specified.
by Deutsche Edelstahlwerke GmbH according to applicable standards\textsuperscript{26–31} to have a ferrite content of 51%. After rolling, the bar was solution annealed at 1050 °C and water quenched. The cell was fully made from this DSS material and has been machined in a way enabling observation of diffusion perpendicular to the rolling direction. This cell allows continuous electrochemical charging of the duplex stainless steel (DSS) with deuterium from inside and analyzing its outer surface exposed to UHV at the same time. The inner surface of the cell was flat and plan-parallel to its monitored outer surface. In that way a 0.5 mm thick steel membrane has been established for permeation experiments with deuterium. To introduce deuterium into the steel a continuous flow of an electrolyte solution of 0.05 M D\textsubscript{2}SO\textsubscript{4} and 0.01 M NaAsO\textsubscript{2} (recombination poison) in D\textsubscript{2}O was realized through the tubular test sample. This was accomplished by pumping the electrolyte through a flexible polymer pipe. The pipe was lead from outside the UHV chamber to the tubular sample within the chamber by using flexible stainless steel bellows. A Pt wire was acting as counter electrode (anode) in the galvano-static charging process using a current density of 10 mA·cm\textsuperscript{-2}.

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Author Contributions
The project was planned and supervised by W.U. and Th.B. The experimental design was performed by F.S. and Th.W. Data was collected by F.S. analyzed by O.S. and G.H. and discussed by O.S., G.H, W.U. and Th.B. The manuscript was prepared by O.S., G.H, Th.B. and W.U. All authors reviewed the manuscript.

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