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# The possible effect of high magnetic fields on the aqueous corrosion behaviour of Eurofer

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## Abstract

In defining the corrosion control requirements for DEMO, the impact of the mixed Eurofer-97/AISI 316 steel system and plant specific effects should be considered throughout, in particular, the effect of the intense magnetic fields present. A substantial amount of data related to corrosion resistance of structural materials is available for industrial applications in fission, but applies to different materials and neutronic conditions. Experimental work is being carried out under the DEMO Breeding Blanket Project of the EUROfusion programme, which will further develop the understanding of irradiation effects. However, there is very limited information regarding magnetic field-assisted corrosion under conditions relevant for the fusion environment readily available in the literature. This work reviews current knowledge and progress in establishing the possible influence of the intense magnetic field on corrosion behaviour of the main structural material, Eurofer-97, in the breeding blanket. To support the relevance of this problem statement, preliminary corrosion experimental results of Eurofer-97 coupons, obtained by using a simple apparatus that allows exposure to a magnetic field intensity of 0.88 T and temperatures up to 80°C in water at atmospheric pressure, are presented as an initial qualitative investigation of possible magnetic field related effects.

Keywords: Eurofer, Corrosion, Magnetic field, DEMO, Breeding Blanket, WCLL

## 1. Introduction

The water coolant circuits of the DEMO fusion power plant will experience extreme conditions that will challenge the coolant-facing structural materials. Namely, high energy neutron irradiation to large cumulative doses, elevated temperature and pressure, the influence of tritium and intense magnetic fields [1]. The effect of a magnetic field on corrosion behaviour is an unusual, and not very well understood, variable. It is of particular importance when a material contains magnetic phases. Prior work has generally been performed at much lower field intensities than those expected for the breeding blanket coolant location. Despite this, results have shown clear effects on corrosion behaviour, with both inhibiting and enhancing effects reported, dependent on the effect on mass transport and corrosion product morphology.

This paper reports initial progress in establishing the possibility of an influence of the intense magnetic fields on the corrosion behaviour of the main structural material in the breeding blanket, Eurofer-97, a reduced activation ferritic-martensitic (RAFM) steel. Three apparent mechanisms by which the high magnetic field could affect corrosion and related processes have been postulated.

*Modification of the adherent corrosion layer:* The Eurofer-97 corrosion product film is expected to comprise a high proportion of magnetite which has a high magnetic moment and so may be structurally

affected by growth within the field, as sub-domains attempt to align to the magnetic field direction. This could credibly lead to increases or decreases in corrosion rate, as the film could foreseeably become denser and more protective, or porous and less passivating. Such observations have been made previously [2], although not for materials comparable to Eurofer-97 or conditions relevant to DEMO coolant circuits.

*Interaction with corrosion product particulates:* It is foreseeable that the high magnetic field will interact with any precipitated or spalled corrosion product particulates with a tangible magnetic moment. The predominance and nature of such particulates is not known and is difficult to predict with accuracy. For instance, PWR primary circuits may accumulate significant inventories of nickel and iron based corrosion products as a result of release from the nickel based alloy steam generator tubes and subsequent mobilization. On the other hand, VVERs which are stainless steel based circuits generally suffer much less from this problem. A generic assumption would be that a high magnetic field will tend to favour deposition of any corrosion product particulates on the surfaces of the coolant channels, as it will effectively increase mass transport rates of the particles to the wall where they may impinge and adhere. Assuming that this effect will be strongest at locations with the highest field would point to enhanced deposition in the breeding blanket, particularly at inboard locations. In addition, it is known that the morphology of precipitation products is also affected by the presence of magnetic fields [3].

*Magnetohydrodynamic (MHD) effects:* The presence of both a high magnetic field and current flows (from corrosion processes) can lead to induced magnetohydrodynamic fluid flow. These are complex phenomena but have been studied through simulation and experiment for various systems [4, 5], including the LiPb system within DEMO [6]. The water coolant circuit differs significantly in that it has a much higher flow rate and a considerably lower conductivity, which would tend to suggest that MHD effects would be lower order. Consider also, however, that the occluded solution within pits or under corrosion scabs could have a much higher ionic concentration and hence local conductivity. In addition to this, though possibly of lower importance, the use of any alkalisng agent would increase the bulk conductivity. The possible enhancement of mass transport rates within an occluded cell (see Fig. 1), where streaming potentials or currents might otherwise be relatively low (diffusion controlled) could be more significant. This mechanism has been observed during artificial pit experiments by Tang & Davenport [7]. Again, the impact is difficult to judge since increased mass transport could enhance corrosion product transport away from an active site and so increase rates, but conversely could also remove the aggressive species and thereby result in pit re-passivation.

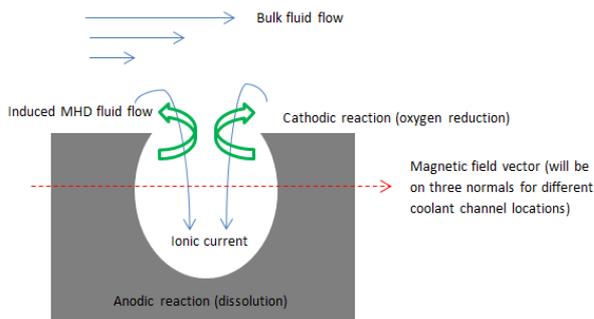


Fig. 1: Representation of possible MHD effect on pitting or crevice corrosion within a high magnetic field.

The following sections of this paper give a summary of the DEMO plant and operating conditions (Section 2), followed by an overview of the relevant literature on corrosion behaviour of materials exposed to high magnetic fields (Section 3), which form the basis of a problem statement for possible impact of magnetic fields on corrosion behavior within the circuit. Notably, it has been found that the literature does not include any information specific to Eurofer-97 and very little of relevance to the DEMO cooling circuit operating conditions. Furthermore, it is understood by the authors that the available corrosion testing facilities which can reproduce conditions close to expected in DEMO cooling circuits are not able to apply intense magnetic fields. Therefore, in order to provide an initial level of information in the identified knowledge gaps on the possible impact of magnetic fields on Eurofer, some preliminary tests have been undertaken to identify qualitatively any effects. The results of these initial tests involving the exposure of Eurofer-97 to water at

comparatively low temperatures in a magnetic field are described in Section 4. These represent an improved level of information specific to this material as well as a providing a basis for comparison to results already reported in the literature and a starting point for any subsequent, more detailed, investigations.

## 2. Plant description

The EU DEMO baseline design defines a reactor of  $\sim 2$  GW thermal power output with a net electric output of  $\sim 500$  MW delivered to the grid in pulses of  $\sim 2$  hours separated by dwell periods of  $\sim 0.5$  hours. The breeding blanket provides the functions of breeding tritium, removing nuclear heat from the plasma, and shielding of the vacuum vessel and other components from the direct neutron flux. The water-cooled lithium-lead (WCLL) blanket design [8, 9] is a set of modular boxes that form the inner wall of the tokamak, through which LiPb flows slowly to generate tritium. A set of coolant pipes manufactured from Eurofer-97 carry water in a radial-toroidal direction to extract the heat deposited (Fig. 2). The First Wall is the front face of each module, cooled independently and consisting of a Eurofer-97 steel plate with water cooling channels. Module dimensions are 1.0 m and 1.4 m in width and 0.55 m and 0.9 m in radial thickness (inboard and outboard respectively).

The WCLL concept utilises a primary coolant of pressurised water with nominal conditions of 15.5 MPa pressure, a reactor inlet temperature of  $295^\circ\text{C}$ , and an outlet temperature of  $328^\circ\text{C}$ . During the dwell period, the primary coolant continues to flow at, or close to, its nominal flow rate, and the average coolant temperature remains the same; that is, the inlet temperature will increase while the outlet temperature decreases to maintain an average temperature of  $311.5^\circ\text{C}$ . The flow velocity of the coolant inside the blanket coolant channels is  $2\text{--}3\text{ m s}^{-1}$ , in the feeding pipes inside the vacuum vessel that provide flow to each module the velocity is around  $7\text{ m s}^{-1}$ , and elsewhere outside of the vacuum the velocity is higher at around  $18\text{ m s}^{-1}$ . The total residence time of the coolant in-vessel where the highest fields are present is of the order of 5-10 seconds.

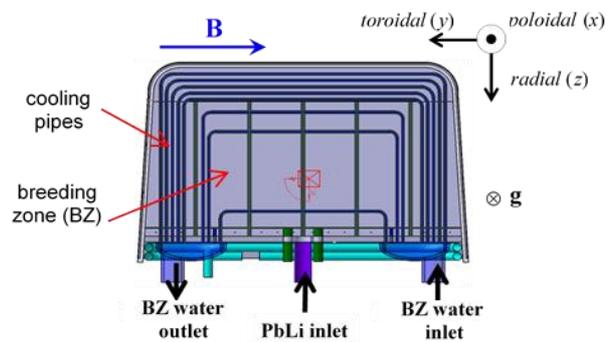


Fig. 2: Radial-toroidal cross-section of an individual WCLL blanket module (adapted from [8]).

The magnetic field seen by the coolant channels varies with position. The toroidal magnetic field varies with a  $1/r$  dependence and the blanket modules have a radial thickness of around 1 m, leading to fields from 9.9 T to 8.6 T across the inboard blanket and from 4.2 T to 3.7 T across the outboard blanket, both in a radial direction.

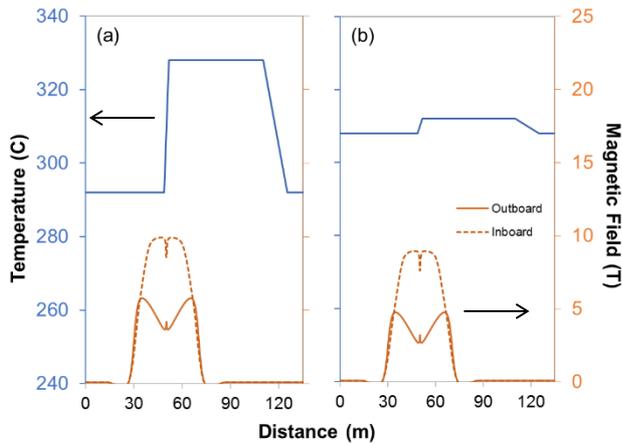


Fig.3: Indicative magnetic field and temperature with distance around the WCLL coolant loop for a notional circulating coolant volume in: (a) pulse and (b) dwell conditions.

Additional contributions to the magnetic field arise from the ferromagnetic structural material itself, and due to the poloidal field generated by the poloidal field coils and the plasma. These add around 0.5 T to the total field. During the dwell period, the toroidal field coils will remain energised, leaving the total field largely unchanged relative to that during the pulse, although the poloidal field component will not be present.

Figure 3 shows illustratively the temperature and field experienced by a notional coolant volume as it flows around the coolant circuit and equatorial midplane blanket modules during pulse and dwell. Coolant enters the vessel via the upper port and, after an initial increase, the magnetic field increases or decreases depending on whether it flows to the inboard or outboard side.

### 3. Literature review corrosion of metallic alloys under influence of magnetic field.

Investigations into the effect of magnetic fields on corrosion have been carried out on a number of different metals and alloys, within different conditions. These are summarised briefly below, although none have yet addressed the impact on aqueous corrosion of Eurofer-97 or comparable steel in high temperature water.

Rućinski *et al.* [10] studied the effect of a magnetic field on the corrosion of AISI 303 stainless steel in a  $\text{FeCl}_3$  solution with the field direction perpendicular to the surface of the metal. It was found that the magnetic field inhibited corrosion, as evident from an increase in repassivation potential, decrease in mass loss, and reduction in pit number. Voltammetric measurements

performed on both Pt and stainless steel electrodes revealed that the cathodic reaction was accelerated by the presence of a magnetic field. This magnetoinhibition was explained as a result of a passive layer formed whilst within the magnetic field with a passivation capacity higher than that formed in the absence of the field.

A number of works have investigated the corrosion of iron and low carbon steel under magnetic fields. Bikul'chuyus *et al.* [2] investigated the corrosion of low-carbon steel in tap water in a permanent 0.05 T field using gravimetric and electrochemical methods over period of 2 hours in circulating water at up to  $70^\circ\text{C}$ . It was found that the presence of a magnetic field decreased the rate of corrosion by 14% on average, regardless of temperature. This was attributed to the presence of the field favouring the formation of an adherent corrosion product film rather than transport away from the surface. Lu *et al.* [11] investigated the anodic polarisation behaviour of iron within 0.5 M sulfuric acid solutions with and without a 0.4 T DC magnetic field. It was found that the presence of an applied magnetic field accelerates the cathodic diffusion process and has little effect on the anodic polarisation. Cathodic limiting current densities were observed to increase when a magnetic field was applied. Further work in 2005 [12] used similar conditions to investigate the effects of a 0.4 T horizontal magnetic field on the electrode processes of iron within sulfuric acid solutions with dichromate. It was found that polarisation resistance significantly decreased (thus decreasing corrosion resistance) in the presence of a magnetic field. Nasher and Shalash [13] measured the magnetic field effect on the activation-controlled dissolution and hydrogen reduction reactions occurring on iron in solutions of up to 5% NaCl. Results revealed an increase in the rate of corrosion for samples placed in the magnetic field.

Sueptitz *et al.* [14] found that the current density of iron in sulphuric acid could be increased or decreased by altering the configuration of the magnetic field to the electrode, due to its effect (via Lorentz force and magnetic field gradient force action) on surface pH during anodic dissolution. Additional work [15] investigating the influence of high gradient magnetic fields on the anodic dissolution of iron in sulfuric acid resulted in observation of a localisation of the material loss within all potential regions of anodic dissolution of iron, with the exception of the passive region. The anodic current density was also affected by the presence of the magnetic field, again attributed to Lorentz force and field gradient effects.

In 2010, Li *et al.* [16] investigated the effect of a magnetic field on pitting corrosion in pure magnesium by stochastic approaches, and identified an increase in the pitting corrosion susceptibility as a result of the magnetic field. It was observed that the field altered the pit initiation mechanism, from the parallel birth and death stochastic model to the parallel birth stochastic model. The mechanism for pit growth was unchanged by the magnetic field, however there was a change in the

growth rate: stable pits were more likely to grow with higher growth rate and develop into larger pits.

The influence of a magnetic field (0.2 T or 0.08 T) on the electrochemical reactions occurring on a copper electrode in 0.5 M HCl by electrical impedance spectroscopy (EIS), electrochemical noise (EN) and potentiodynamic polarisation curve measurements was explored by Garcia-Ochoa *et al.* [17]. Lorentz forces were minimised by aligning the symmetry axis of the magnetic field with the ion flow. It was found that the presence of a magnetic field affected cathodic reactions considerably, with an increase in corrosion rate in the presence of oxygen in acid media, or a decrease in the absence of oxygen. The corrosion potential was shifted towards anodic values in the presence of oxygen, or a cathodic shift in the absence of oxygen. Furthermore, it was found that the hydrogen ion reduction reaction was retarded by the magnetic field, as such, corrosion current was also lowered. Changes to the morphology of the electrodeposits were also observed as a result of the presence of the field. The corrosion of Ag and Cu within nitric acid solutions was found to be partially suppressed by a magnetic field of up to 5 T [18]. The effect was considered to be the result of the removal of the catalyst from the electrode and electrode surface passivation, due to Lorentz force.

A mechanistic investigation into magnetic fields effects was undertaken using artificial pits by Tang and Davenport [7]. This simulated pitting corrosion using one-dimensional artificial pits and two-dimensional pits in metal thin films. It was found that, for ferromagnetic materials (iron, cobalt, nickel), pitting dissolution was affected significantly by the presence of a magnetic field. Upon the application of a parallel magnetic field, pitting dissolution increased. This was the result of local stirring within the artificial pits produced as a result of the Lorentz force and the high field gradient force causing an increase in the diffusion-limited current. In comparison, the application of a perpendicular magnetic field resulted in a reduction in pitting dissolution. This was caused by paramagnetic corrosion products being drawn towards the surface of the metal by the field gradient force, forming a protective layer.

Pondichery *et al.* [19] investigated the effects of a 0.75 T external magnetic field on a range of materials, both non-magnetic and ferromagnetic, within 3.5% NaCl solution. By comparison of the electrochemical behaviour by both potentiodynamic polarisation and corrosion potential vs. time tests, it was found that the magnetic field has a notable effect on both the corrosion rate and the corrosion potential. An increase in corrosion rate and a cathodic shift of corrosion potential was observed for ferromagnetic materials (416 stainless steel, 1018 carbon steel), whereas for non-magnetic materials (304 stainless steel, Ti alloy, pure Zn) no such effect was measured.

Studies have been performed into magneto-hydrodynamic (MHD) effects on corrosion of Eurofer-97 [20] and other materials [21] in flowing LiPb, although

these are considered to have limited relevance to behaviour in water coolant.

Of all the studies outlined, those with most relevance to Eurofer-97 are likely to be those on AISI 416 martensitic steel [19]. The observation that corrosion rates were enhanced for this material is in common with many of the other studies, although this appears to be determined by other application dependent factors, such as geometry and transport processes. Overall these findings point to the likelihood of a magnetic field effect on the corrosion of Eurofer-97 and therefore underline the value of further work. Consequently, initial tests were undertaken in conditions which could be compared to the literature sources above, as a facility which can impose magnetic fields on corrosion specimens under the conditions of interest is not currently available.

## 4. Corrosion Tests on Eurofer-97 within a Magnetic Field

### 4.1 Methodology

A series of preliminary tests were undertaken with the aim of identifying qualitatively whether there is an effect of high magnetic fields on the corrosion of Eurofer-97. A simple apparatus was employed to expose coupons of Eurofer-97 to a high magnetic field at temperatures up to 80°C at atmospheric pressure. Samples were held within a glass tube in thermostatically controlled water within the annulus of a Halbach array of NdFeB magnets (see Fig 4). The array was rated as having an annulus field intensity of 0.88 T, which was confirmed with a magnetic field meter and found to be constant within the annulus. The corrosion coupons were held either within the annulus or well outside it (zero field) with nylon monofilament. In order to avoid heating and possible demagnetisation of the magnet, a cooling jacket comprised of PTFE tubing was placed between the glass tube (outer diameter 27 mm) and the array (inner diameter 30 mm). A perfusor syringe was used to maintain a constant solution level and the solution was deoxygenated by a continuous low flow rate sparge of nitrogen. The top of the tube was filled with stainless steel wool and uppermost ceramic fibre lagging to reduce the ingress of oxygen into the system. Samples of Eurofer-97 were cut into coupons of approximately 5 x 5 x 1 mm, ground to SiC P1200, degreased in isopropanol, washed in demineralised water and dried prior to aqueous exposure. Test solutions were composed of high purity demineralised water in some cases with additions of sodium chloride and citric acid (Analytical grade, Fisher Scientific). The following test conditions were applied, which were deoxygenated in all cases.

- Demineralised water at 80°C for 100 hours
- 100 mg kg<sup>-1</sup> chloride (as sodium chloride) at pH 3.5 at 60°C for 140 hours.
- 1% w/w chloride (as sodium chloride) at pH 2.4 at 22°C for 190 hours

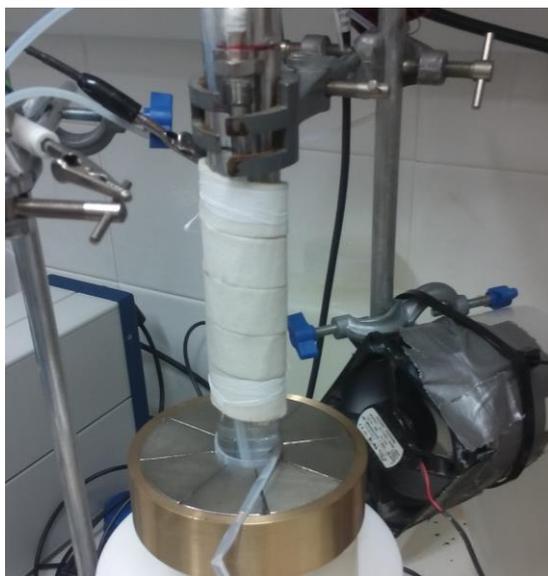


Fig. 4: Apparatus employed for initial trials on Eurofer-97 corrosion within a magnetic field.

The post-exposure samples were analysed by scanning electron microscopy (SEM) and energy dispersive X-ray (EDX) using a TESCAN Mira3 XMH FE-SEM microscope equipped with an X-Max 80 EDS detector from Oxford Instruments and a Zeiss Sigma FE-SEM. The electron beam was operated at 10-30 kV. Surface sectioning by ion beam milling was undertaken with a FEI Helios NanoLab 600 DualBeam FIB-SEM. X-ray diffraction (XRD) spectra were recorded with an X'pert Pro MPD with a Cu K- $\alpha$  source. All spectra were normalised to the Fe(400) reflection and phase identification performed by comparison with standard literature [22, 23]. Secondary-ion mass spectrometry (SIMS) measurements were made using a double focusing magnetic sector instrument VG Ionex IX-70S at VTT. A 5 keV  $O^{2+}$  primary beam with a current of 500 nA and the intensities of the positive secondary ions at mass-to-charge ratios of 48 (Ti), 51 (V), 52 (Cr), 56 (Fe), 60 (Ni) and 181 (W) were measured as a function of time. Negative secondary ions were measured with a 12 keV  $Cs^+$  primary beam using a current of 100 nA. In the case of Cs bombardment secondary ions 16 (O), 35 (Cl), 52 (Cr), 56 (Fe), 68 (CrO) and 72 (FeO) were followed [24].

## 4.2 Results and Discussion

Exposure tests on Eurofer-97 in deoxygenated demineralised water were found to produce only very

slow film formation at temperatures between 60 and 80°C. More aggressive conditions involving the addition of sodium chloride and citric acid enhanced corrosion to an observable extent. Although these conditions were less representative of the expected WCLL chemistry, they could have relevance to localised accumulation of impurities within an occluded crevice or corrosion scale. They are also comparable to those used in literature studies which generally employed elevated levels of aggressive anions and/or acidic matrices [7, 10-19]. At 100 mg kg<sup>-1</sup> chloride and pH 3.5, a mild but visible darkening of the sample surface occurred, with a change in hue, indicative of the formation of a corrosion film. A chloride concentration of 1% w/w at pH 2.4 at 22°C increased the corrosion rate further and led to a matte grey surface. Nonetheless, the changes in mass recorded over the periods studied were too small to allow gravimetric distinction between those within and without the magnetic field. Visual inspection showed some differences in the hue of the corrosion film and in some cases an evident greater darkening of the samples within the field, although other replicates showed less distinct contrast.

An SEM image of the surface of a reference sample prepared by grinding alone without aqueous exposure is shown in Figure 5a. The surface of a sample exposed to demineralised water at 80°C with no magnetic field for 100 hours is shown in Figure 5b and shows no sign of corrosion attack. A slight increase in oxygen is indicative of the formation of a thin protective oxide layer. Secondary electron images of sample surfaces exposed to 100 mg kg<sup>-1</sup> Cl at pH 3.5 for 100 hours at 60 °C, with 0.9 T field and no field are shown in Figures 5c and 5d respectively. Both samples are characterised by a thicker oxide layer than that produced in demineralised water, although this still too thin to allow accurate thickness measurement by FIB-SEM (Figure 5c). The chemical compositions analysed by EDX indicate presence of various forms of iron hydroxide and possibly hematite (Fe<sub>2</sub>O<sub>3</sub>) and magnetite (Fe<sub>3</sub>O<sub>4</sub>). The XRD analyses indicated the likely presence of highly textured magnetite, alongside other unidentified phases, with a higher abundance in the presence of the magnetic field. Additional analyses are required to confirm the dominant corrosion products, but the presence of magnetite is expected due to the deoxygenated exposure conditions.

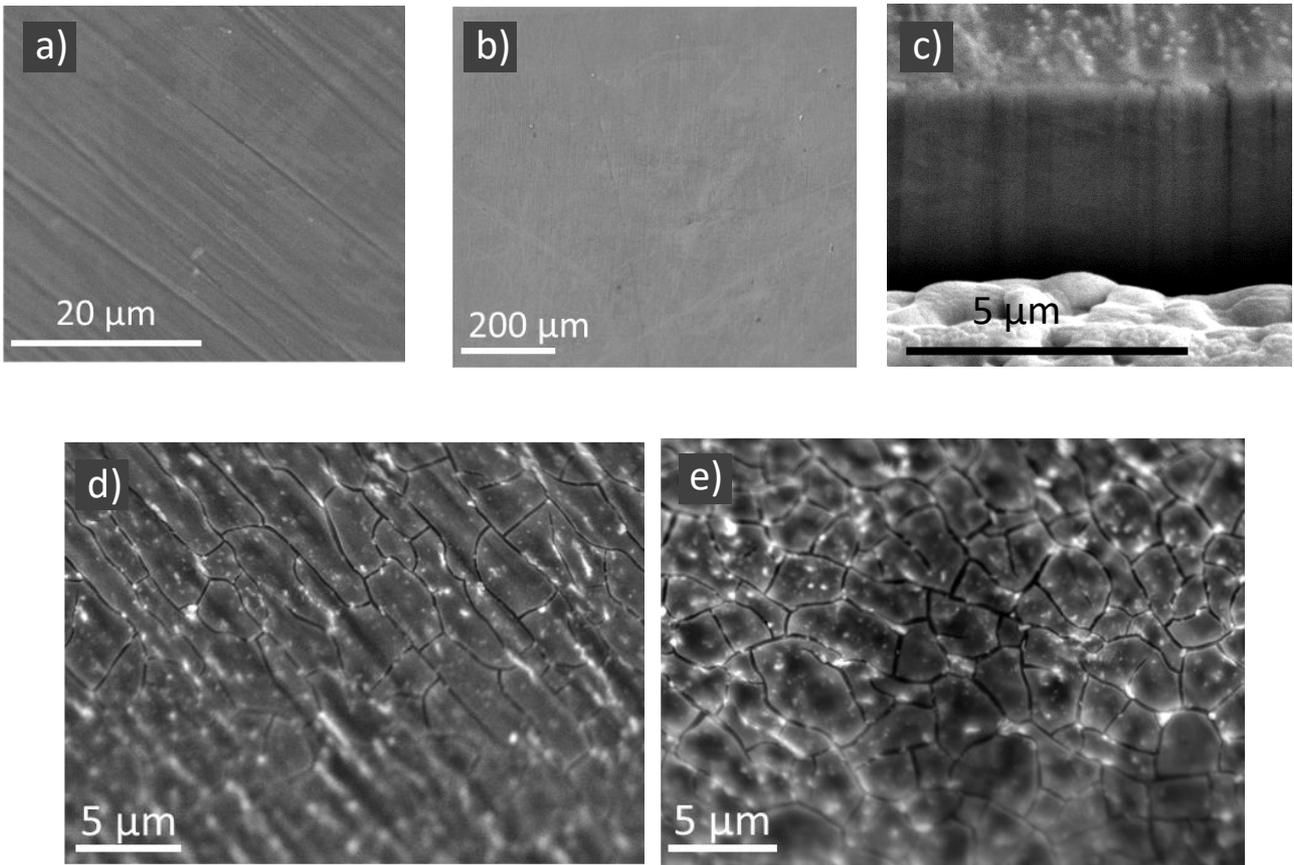


Fig. 5: Scanning electron micrographs of Eurofer-97 coupon surfaces showing: reference sample (a), sample exposed for 100 hours to deoxygenated demineralised water at 80°C without magnetic field (b), FIB-SEM section of corrosion film on sample exposed for 140 hours to deoxygenated pH 3.5 water with 100 mg kg<sup>-1</sup> chloride at 60°C with a 0.9 T magnetic field (c), sample exposed for 140 hours to deoxygenated pH 3.5 water with 100 mg kg<sup>-1</sup> Cl at 60°C, with a 0.9 T magnetic field (d), and without a field (e).

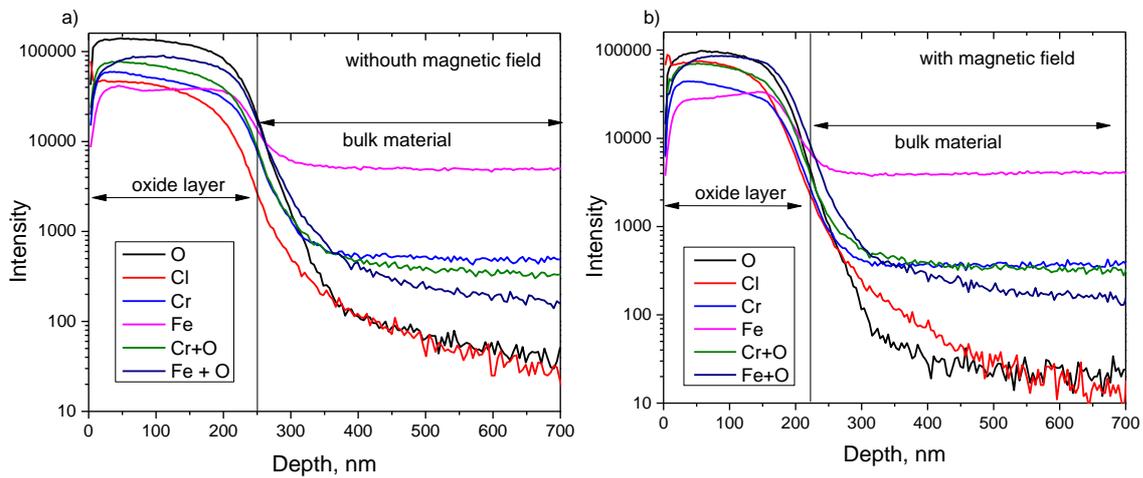


Fig. 6: Results of SIMS analyses of Eurofer-97 sample surface exposed for 140 hours to deoxygenated pH 3.5 water with 100 mg kg<sup>-1</sup> Cl at 60°C without magnetic field (a) and with 0.9 T magnetic field (b).

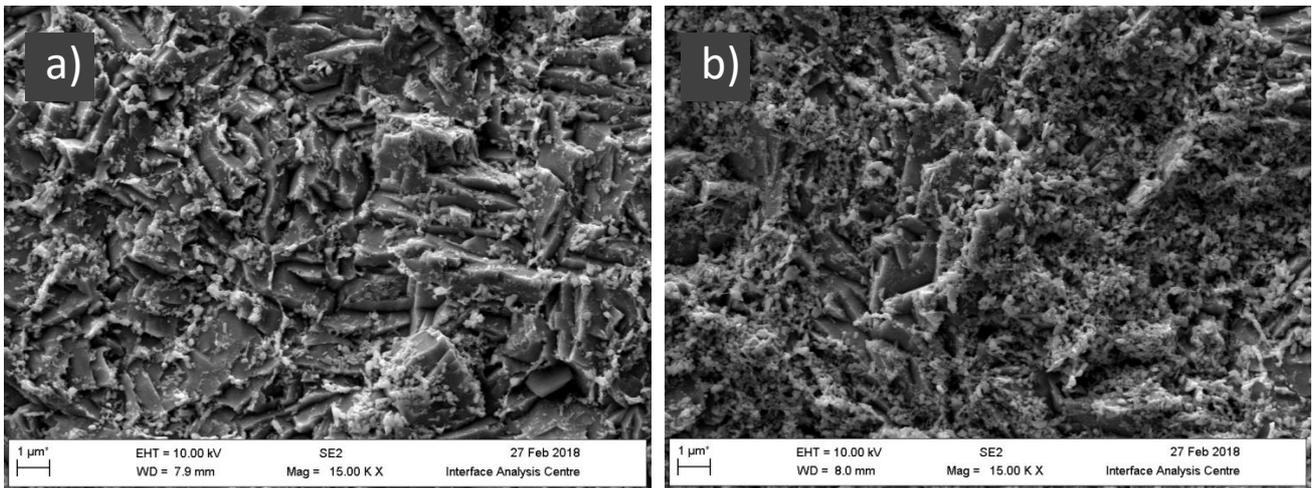


Fig. 7: Scanning electron micrographs of samples exposed for 190 hours to deoxygenated water with 1% w/w chloride at 22°C with a 0.9 T magnetic field (a), and without a field (b).

EDX point analysis of the surface showed enrichment of the corrosion product layer in W up to 1.6 wt.% (control: 1.1 wt.%), Cr up to 12 wt.% (control: 8.8 wt.%), and Ni up to 0.09 wt.% (control: 0.015 wt.%). Higher enrichment is observed in sample exposed in the magnetic field. The cracked pattern observed on the oxide layer is attributed to internal stress accumulation or thermal shock of the corrosion layer. Interestingly the crack pattern of the oxide layer in Figure 5e looks uniform, without any particular orientation. On the contrary, the crack pattern on the sample formed within the magnetic field (Figure 5d) tends to form parallel lines, as if the layer formed compact elongated islands along one direction and less compact zones in between. It might be speculated that the magnetic field affected the oxide film formation due to the development of distinct magnetic domains in the ferrimagnetic magnetite comprising the corrosion film. The ion milled section (Figure 5c) shows that the film formed in these conditions was very thin, clearly well under 0.5  $\mu\text{m}$ .

SIMS spectra taken on samples are shown in Figure 6. Figure 6a shows depth profiles of O, Cl, Cr, and Fe in the corrosion products layer of sample corroded in the solution without applying the magnetic field and Figure 6b shows the depth profiles of the sample corroding under the magnetic field. The horizontal axis represents depths calculated based on the sputtering time, and the vertical axis is the intensity of each element. The depth profiles obtained are relatively similar and suggest an oxide thickness of around 180-250 nm, which is consistent with the FIB-SEM section. However, the oxide layer developed in the magnetic field is approximately 20-30 nm thinner. The spectra obtained by both beams (Cs and O, not presented here) show Cr enrichment across oxide layer in the sample exposed to the magnetic field. A greater enhancement is apparent for the abundance of oxygen in the sample without the field, which would be consistent with a less oxidised corrosion product (notably, magnetite, compared to FeOOH).

A number of surface points were selected at random on samples exposed for 190 hours to deoxygenated water with 1% w/w chloride at 22°C for SEM inspection of the corrosion layer morphology. The film on these samples was significantly more developed and showed a difference in the dominance of corrosion product morphology depending on whether they were exposed in the presence of a magnetic field. The sample exposed in the presence a magnetic field had a greater coverage of semi-octahedral crystallites with a smaller amount of fine granular particles (Figure 7a). The sample exposed in the absence of a field was dominated by the fine granular phase (Figure 7b), with underlying semi-octahedral phase visible in some areas, but not others. The dimensions of the semi-octahedral faces were also larger on the sample exposed to the magnetic field. A tentative assignment based on typical corrosion product morphologies can be made that the semi-octahedral phase is magnetite, with the finer granular phase being lepidocrocite or goethite (both FeOOH). These observations could suggest that the magnetic field stabilised the ferrimagnetic corrosion product phase and thus caused it to cover a greater proportion of the surface area, as well as forming larger crystallites. This corresponds to the preceding observation of changes in the thinner corrosion product layer under somewhat less aggressive chemistry.

These data are currently too sparse to allow definitive analysis, however, they are noted to be broadly consistent with the literature [19]. The modification of similar corrosion product phases by a magnetic field was observed [3], and this has been related effects on corrosion rate, which may either increase or reduce rates [7, 14], depending on whether the corrosion film became more [10, 18] or less [12, 13, 19] passivating.

## 5. Operational Relevance

Although the data presented in the preceding section point to a qualitative effect of high magnetic fields on Eurofer-97, this does not indicate the extent of any enhancement (or indeed diminishment) of corrosion rates. The impact of a change in corrosion product morphology is not clear and even if this results in an initial increase in film formation rate, then as long as the properties of the film formed are not significantly impaired, then this may in fact result in a more rapid surface passivation. The greater concern would be that the modified corrosion layer was more highly porous or defected and so did not effectively passivate the surface from ongoing corrosion.

Should a significant long term increase in corrosion rates associated with high magnetic field be identified, then clarification of the mechanism by which this occurs may allow mitigations to be identified. One of the main mitigations which can be proposed if ongoing work in this area suggests a possible corrosion performance issue is the use of an extended “hot functional test” during the DEMO commissioning phase. This would aim to develop the Eurofer-97 corrosion film fully before the first plasma is produced and in the absence of a magnetic field. The potential for modification of the existing corrosion product when the field is applied would need to be considered, but this approach, alongside other measures, has been effective in ensuring passivation and low corrosion product release in commercial LWR plant.

Changes in corrosion behaviour of alloy surfaces within the irradiated coolant circuit could have an associated impact on uptake and release of activated species which could affect the radiation fields in out of flux circuit locations due to redeposition. The high magnetic field could also have an effect on any circulating corrosion product particulate which is likely to be held up and could have a higher likelihood of deposition on surfaces, with the possibility of “crud-bursts” when the magnetic field is removed at the end of an operational phase.

## 6. Summary and Outlook

A comparison between the expected conditions in the DEMO breeding blanket WCLL water coolant circuit, and the relatively limited available literature on the effect of high magnetic fields on corrosion of steels suggests that some effect can be expected on the performance of the Eurofer-97 coolant-facing structural material. This is due to it being ferromagnetic and the primary corrosion product formed under deoxygenated conditions (magnetite) being ferrimagnetic.

Due to the lack of an identified corrosion testing facility capable of reproducing key aspects of the DEMO WCLL cooling circuit conditions at the same time as applying high magnetic fields, preliminary trials have been undertaken to provide a qualitative indication as the potential for magnetic field-assisted corrosion to occur. These scoping tests involved exposure of Eurofer-97 coupons to a range of temperatures which are

comparatively low compared to the DEMO operating conditions, and with a range of chemistries to allow comparison to literature studies. The test solution was maintained in a deoxygenated state in keeping with the expected DEMO operating conditions, which will also favour formation of magnetite as a corrosion product which is ferrimagnetic. The results of these trials have indicated qualitative differences in corrosion film morphology as a result of the magnetic field. Differences in micro-crack orientations were observed which could be related to the development of magnetic domains in the corrosion product. An increased dominance and enhanced crystal size in one corrosion product phase (tentatively identified as magnetite) was also identified in the presence of the magnetic field. These observations are comparable to results from the literature for broader material classes.

These tests are too limited to provide indications as to the likely extent of any deleterious effect of magnetic fields on the service performance of Eurofer-97, however, the literature points to an increase in corrosion rates of ferromagnetic materials exposed to high magnetic fields. On this basis, the possibility of an effect on Eurofer-97 must be considered further. A key aspect of any further work must be the identification or development of a facility which will allow corrosion tests in the presence of high magnetic fields and temperature, and chemistry conditions relevant to DEMO WCLL cooling circuits.

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