In-Situ Observation of Concurrent Oxidation and Mechanical Deformation in Al and Zr

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Metals will naturally form an oxide layer at the surface when exposed to air. The oxide skin is a diffusion barrier that may slow down the corrosion rate significantly. Since structural materials are usually working under external/internal stresses, cracks may be generated inside the oxide layer during deformation, reducing the protectiveness of oxide. Understanding the response of metal surface oxide to concurrent chemical reaction and deformation, is critical for unravelling the stress corrosion cracking (SCC) mechanisms of metals and development of alloys with improved corrosion-resistance. However, direct observation of the physical processes in the oxide layer as well as at the metal-oxide-interface (MOI) is challenging, as they are usually buried by nature [1]. Herein, using an environmental TEM (E-TEM), we report \textit{in-situ} observations of the concurrent oxidation and mechanical deformation in aluminum (Al) and zirconium (Zr). These two metals are interesting systems to compare: Al has an amorphous native oxide at room temperature while Zr develops a crystalline oxide.

For Al, we develop a mechanical break-junction approach enabling direct observation of the environmental deformation of the naturally-formed alumina on Al surface at room temperature. The experimental procedures are illustrated in Fig. 1a-c. We prepare two Al nanotips, and bring them to contact inside an ETEM (Fig. 1a). The nanotips are cold-welded [2] together (Fig. 1b). Afterwards, pure oxygen is injected inside the chamber (Fig.1c). We then pull the nanotips and observe the evolution of the Al oxide layer and MOI simultaneously. We found that native alumina is liquid-like during deformation, and it can remain its integrity without any cracks/spallation (Fig. 1d-e) at moderate strain rate. Also, it can keep matching the deformation of metal even when the metal surface inclinations have changed significantly. When the strain rate is higher, fracture in metal will occur, exposing fresh metal surface. We then visualized the initial oxidation of Al at the metal-gas interface at atomic resolution (Fig. 2a-c). Unlike traditional thin film growth or nanoglass consolidation processes, we observe seamless coalescence of new oxide islands without forming any glass-glass interface or surface grooves (Fig. 2b), indicating greatly accelerated glass kinetics [3] at the surface compared to the bulk.

For Zr, we visualize the high temperature oxidation process when no external force is applied. It is surprising that the oxidation-induced internal stress lead to direct cracks/pores formation in the thin film (fig. 2d), which in turn accelerates the oxidation kinetics by providing new oxygen diffusion shortcuts.

In conclusion, our research reveals the liquid-like superplasticity in thin amorphous alumina during deformation at room temperature. Our findings have important implications for not only understanding of the unique oxidation resistance in Al, but also development of new corrosion-resistant coatings and alloys.
References:


Figure 1. (a−c) Schematic drawing of the experimental procedures. (d−e) Sequential TEM images showing the super-elongation and self-healing process of aluminum oxide when stretched in O₂ environment. The strain of oxide at (e) is about 117%. All scale bars, 5 nm. Reprinted with permission from [4]. Copyright 2018 American Chemical Society.

Figure 2. (a−c) Sequential HRTEM images of the initial oxidation process on aluminum fresh surface. Reprinted with permission from [4]. Copyright 2018 American Chemical Society. (d) HAADF image of the Zr thin film after high temperature oxidation, showing numerous small cracks and pores.