

Mechanistic understanding of the effect of initial microstructure of low-alloy carbon steel on its CO₂-corrosion resistance in simulated produced water

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CO₂ corrosion of mild steel and how the precipitation of the corrosion products could affect the protectiveness of the underlying material have been extensively investigated during the last decades. In CO₂ corrosion condition, one almost always encounters the formation of corrosion scale including carbonates phases as well as several kinds of (hydro-) oxides. Initial microstructure of material could highly influence the mechanisms responsible for CO₂ corrosion as well as the resistance of the material against further deterioration. However, not many studies have been performed on understanding the influence of initial microstructure on corrosion behavior in CO₂ sweet conditions on a real scenario i.e. formation water chemistry with electrolytes containing Ca²⁺ ions.

The detailed investigations in this study are focused on understanding the effect of heat-treatment induced microstructural features on CO₂ corrosion behavior and scaling of the low-alloy steels. For this purpose, L80-1Cr material was subjected to several heat treatments (Heating to austenitizing temperature and then cooled in different medias) to obtain different initial microstructures. The heat-treated samples were then electrochemically exposed to CO₂ saturated simulated formation water chemistry. Electrochemical and corrosion behavior of the steels were evaluated using DC polarization and AC impedance techniques. Microstructure, surface chemistry, and surface morphology before and after corrosion experiments were investigated using Scanning Electron Microscopy (SEM), X-ray Diffraction (XRD). Moreover, depth-resolved phase identification of the corrosion scales were applied using synchrotron X-ray diffraction to determine the sequence and extension of corrosion products across the depth.