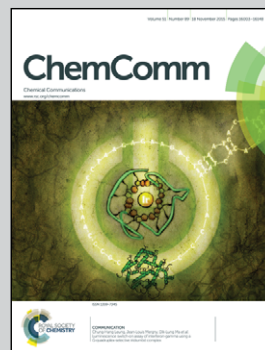


Showcasing research from Andreas Erbe's laboratory,  
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Solvent-starved conditions in confinement cause chemical  
oscillations excited by passage of a cathodic delamination front

The use of Raman spectroscopy to investigate the initial stages  
of a corrosion process at a polymer/metal interface reveals  
oscillations in pH. These oscillations are caused by the limited  
supply of water as solvent and reagent.

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## Solvent-starved conditions in confinement cause chemical oscillations excited by passage of a cathodic delamination front

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**After passage of a delamination front at a polymer/zinc interface, pH oscillations and oscillations in the quantity of corrosion products are observed. The reason for these oscillations is the low quantity of water in the confined reaction volume, water consumption by oxygen reduction, and water regeneration after precipitation of ZnO.**

Confinement enables chemical processes which are not possible in bulk solution. Catalyst particles exhibit different catalytic activities inside carbon nanotubes compared to free particles.<sup>1,2</sup> The inside of other carbon nanostructures,<sup>3–5</sup> and silica networks can be used as “nanoreactors”.<sup>6,7</sup> Spatial confinement was used to synthesize nanosheets.<sup>8,9</sup> The ability to create confinement may have been crucial in the origin of life on the earth.<sup>10</sup> Confinement enables the control of the characteristics of metal (electro)catalysts.<sup>11–13</sup> In O<sub>2</sub> electrochemistry, confinement affects reaction rates.<sup>14,15</sup> Corrosion works differently in confinement, e.g. in “crevice corrosion”,<sup>16</sup> which has recently been monitored in real time.<sup>17</sup> In corrosion protection, polymer coatings are frequently employed. During their deadhesion from the base metal substrate, a confined reaction volume is created, which shall be explored here. The most prominent mechanism of deadhesion of a polymer from steel and galvanised steel is the electrochemical process of cathodic delamination.<sup>18</sup> Inspired from the wealth of precipitation patterns formed in chemical gardens,<sup>19</sup> this work studies the corrosion of Zn in the confinement created after passage of a delamination front between Zn and delaminating poly(vinyl butyral) [PVB] as a model for a weakly protecting polymer coating.

The mechanism of cathodic delamination has been shown to start with the electrochemical O<sub>2</sub> reduction (ORR) at a front, which propagates underneath the coating, with anodic metal dissolution and random formation of corrosion products following after the front.<sup>18,20–22</sup> We will show that oscillations are observed in the quantity of corrosion products formed and

in pH between a delaminating polymer and a Zn substrate after the passage of a cathodic delamination front.

For the experiments, zinc sheets (1 cm × 1 cm × 1.5 mm; purity 99.95%; Goodfellow, Cambridge, UK) were initially mechanically polished using 4000 grit polishing paper and then cleaned ultrasonically in ethanol. The polished zinc sheets were spin coated with 10 weight% solution of PVB in ethanol (Sigma Aldrich) followed by drying in an oven at 70 °C, to obtain ~2–3 μm thick polymer layers. Afterwards, a small defect in the polymer coating was created at the edge of the sample and the sample was placed on a zinc base plate. Then a small reservoir was built around the edge of the sample with the help of a fast drying two component adhesive to place an electrolyte.

To carry out *in situ* and operando Raman measurements, the assembly of the sample mounted on a zinc substrate was introduced into a custom made chamber with a glass window opening to allow spectral acquisition (Fig. 1). To minimize electrolyte evaporation, the humidity was controlled at >90% by mixing humid and dry air with flow controllers. The *in situ* Raman spectra were recorded using a Horiba Jobin Yvon LabRAM confocal Raman microscope with an Argon ion laser source (emission at 514.342 nm) and a power output of 12 mW. The typical spectral acquisition time was in the range of 20 to 30 s. The measurements were performed in backscattering geometry

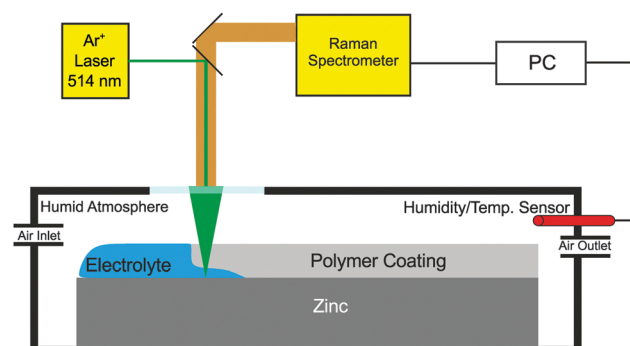


Fig. 1 Schematic view of the setup used for the experiments.

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with a long working distance 50 $\times$ -objective (numerical aperture 0.5) for sample illumination and scattered light collection. The laser spot size on the sample surface was approximately 10  $\mu\text{m}$  and it was focused roughly 2–3 mm away from the edge of the sample. To start the cathodic delamination process, the reservoir was filled with 1 M aqueous KCl solution. During the *in situ* measurements, the laser was focused on the same spot throughout the experiment. (Mapping of the sample surface, though desirable, failed because (a) it is difficult to keep the focus at the polymer/metal interface inside the *in situ* cell during mapping and (b) because during mapping, information on the temporal and spatial dependence are interleaved.)

Raman spectra (examples in Fig. 2) show modes of the forming ZnO corrosion products, of H<sub>2</sub>O, and of the PVB polymer layer. Raman spectra are analysed in the regions containing ZnO modes (wavenumber  $\tilde{\nu}$  360–660  $\text{cm}^{-1}$ ) and water modes (3050–3800  $\text{cm}^{-1}$ ). The ZnO region of the spectra is dominated by an E<sub>2</sub> mode of wurzite ZnO at  $\sim 440 \text{ cm}^{-1}$ , and its LO phonon at 570  $\text{cm}^{-1}$ .<sup>23,24</sup> The LO phonon of single crystalline Wurzite ZnO is inactive in the backscattering geometry used here; its observation in the spectra therefore shows the presence of defects, *e.g.* oxygen or zinc vacancies.<sup>25,26</sup> The peak positions are stable throughout the experiments, except for a shift of the LO peak to higher wavenumbers in the first minutes of the experiment. Consequently, the average environment inside the ZnO lattice is not changing, except in the initial phase of the experiment. The changes affect the environment of defect sites.

Normalised Raman intensities  $I_N(\tilde{\nu})$  shown in Fig. 3a and b have been obtained for each individual spectrum by setting the spectrum's minimum intensity  $I_{\text{min}}$  to zero and the spectrum's maximum intensity  $I_{\text{max}}$  to 1, *i.e.* as

$$I_N(\tilde{\nu}) = \frac{I(\tilde{\nu}) - I_{\text{min}}}{I_{\text{max}} - I_{\text{min}}} \quad (1)$$

In the normalised intensities, oscillations are visible in the intensity of the ZnO modes (Fig. 3a and c). Intensity maxima

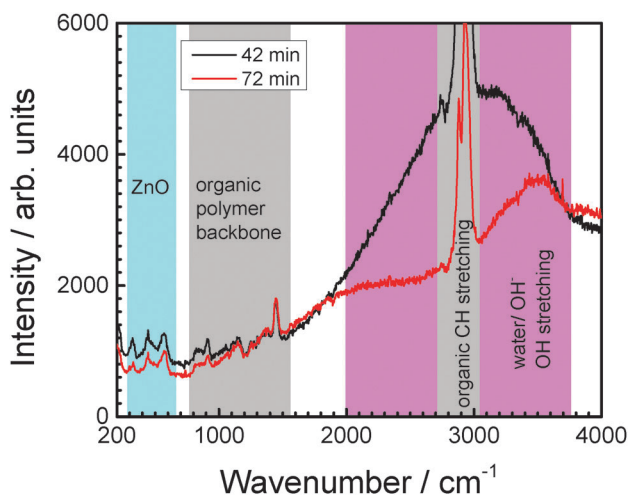


Fig. 2 Examples of *in situ* Raman spectra after two different delamination times as indicated on the graph. Regions with contributions from the different species in the system are labelled.

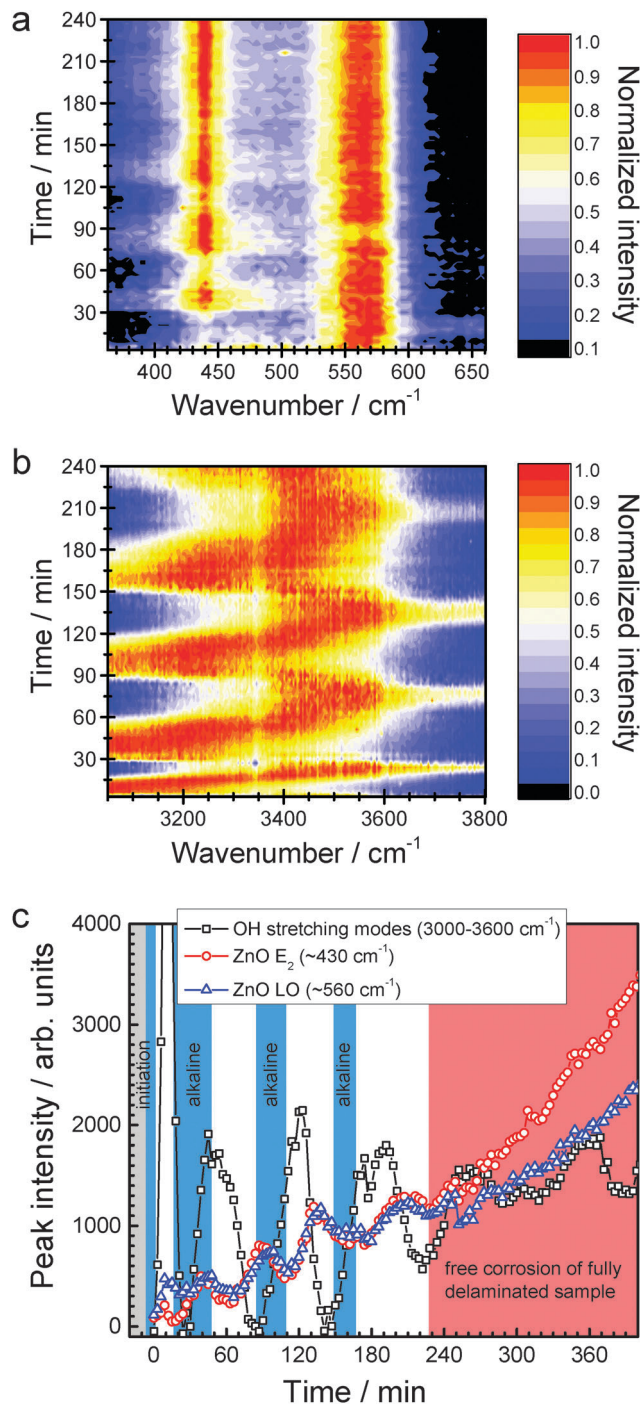


Fig. 3 Normalised Raman spectra in the regions dominated by (a) ZnO phonons and (b) OH stretching modes. Panel (c) shows the time dependence of the intensities at peak maxima. Times where the maximum of the OH stretching modes is below 3350  $\text{cm}^{-1}$  are labelled "alkaline".

and minima are observed in both ZnO modes at the same times, *i.e.* there is no shift between the two curves. Because the observation of the LO phonon in backscattering geometry is closely related to defects,<sup>25,26</sup> this observation excludes the possibility that oscillations between defect-rich and defect-poor ZnO occur. Rather, the amount of the ZnO corrosion product oscillates. On the other hand, the maximum intensity of the OH stretching



modes is observed with a certain delay after the ZnO intensity maxima (Fig. 3c).

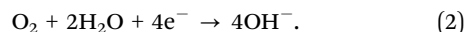
Maxima of the OH stretching modes of H<sub>2</sub>O clearly show oscillations (Fig. 3b), between  $\approx 3170\text{ cm}^{-1}$  during intervals with increasing intensity of the OH stretching modes and  $\approx 3500\text{ cm}^{-1}$  during decreasing OH intensities. A shift of the maximum to lower wavenumbers is indicative of higher OH<sup>-</sup> concentration in water.<sup>27,28</sup> Oscillations in intensity become obvious when plotting the Raman intensity at the maximum of the respective peak (Fig. 3c).

From scanning Kelvin probe investigations, a delamination rate of  $\approx 3\text{ mm h}^{-1}$  has been determined. The oscillations terminate after  $\approx 4\text{ h}$ , which coincides with the time for delamination of the full, 1 cm long coated area.

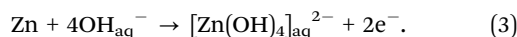
Chemical oscillations are encountered in a number of chemical systems far from equilibrium. Mostly, an autocatalytic step is involved in the reaction, *e.g.* because one reaction product or intermediate blocks or catalyses a follow-up reaction in a multistep process.<sup>29</sup> Chemical oscillations have especially been observed and exploited in heterogeneous multi-phase systems.<sup>30,31</sup> When coupled to oscillating reaction, interesting dynamics can be observed in follow-up processes.<sup>32–36</sup> Oscillations in pH can lead to propagating fronts, when coupling to dynamic aggregation of nanoparticles.<sup>37,38</sup> Even phase formation and dissolution can oscillate.<sup>39</sup>

No oscillations have so far been reported in delamination processes. In electrochemical systems, however, oscillations and pattern formation have been observed, *e.g.* during dissolution of n-type Si.<sup>40</sup> On the example of CO electrooxidation, the mechanisms leading to oscillations involving several micro-electrodes have been studied. With appropriate cooperativity, microscopic oscillations can be transformed into macroscopic patterns.<sup>41</sup> In corrosion processes, local cathodes and local anodes are usually randomly distributed over the surface.<sup>21</sup> Consequently, no oscillations are observed in a typical corrosion process. (Exceptions exist—oscillations in current and potential are observed *e.g.* in corrosion of stainless steels,<sup>42,43</sup> of carbon steels or iron in acids,<sup>44–46</sup> and copper,<sup>47</sup> amongst others, mostly because of active-passive transitions.)

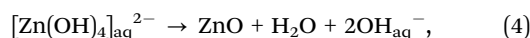
The proposed mechanism which leads to the oscillations observed here is summarised in Fig. 4. At a cathodic delamination front, ORR proceeds under water consumption,<sup>48,49</sup>



The initial increase in the intensities of the OH stretching modes marks the approaching delamination front; the passage of the delamination front coincides with the initial maximum intensity of the OH modes. The quantity of water present between the metal surface and polymer film in the initial stages of delamination is small. Therefore, immediately after passage of the delamination front, the conditions are highly alkaline.<sup>48,49</sup> According to classical kinetics, the rate of the ORR should hence decrease. In the illuminated region ( $\sim 10\text{ }\mu\text{m}$ ), the beginning of ZnO formation is closely associated with this passage of the delamination front. The ORR subsequently consumes water and produces OH<sup>-</sup>. The alkaline conditions thus created lead to dissolution of the initially formed ZnO. After the delamination front has moved on, the surface becomes anodic, featuring metal dissolution. At pH > 12, set by the conditions created as a result of ORR, Zn dissolution occurs according to ref. 50:



An increase in the concentration of soluble Zn species together with a decrease in pH due to the binding of OH<sup>-</sup> inevitably makes the formation of the solid corrosion product ZnO favourable, which forms as



under water release, possibly *via* Zn(OH)<sub>2</sub> as an intermediate.<sup>50</sup> The corresponding Raman spectra show increasing intensities of ZnO and OH modes. ZnO formation results in an increase in the number of available water molecules triggering a restart of the ORR, this time on a ZnO-coated surface, where O<sub>2</sub> was accumulated again. The observed potentials are known from Kelvin probe measurements to permit the reduction of O<sub>2</sub>.<sup>20</sup> The consequence is alkalisation, dissolution of the formed ZnO, hence a decrease of the intensities of the ZnO modes. Alkalisation and consumption of water will stop the ORR again at some point, making the region anodic, with the start of the next cycle. ZnO peak intensities oscillate with an increasing slope of the running average. This increase is consistent with the release of only one water molecule per unit of ZnO formed, while in the ORR, (at least) two water molecules are needed for one O<sub>2</sub> molecule to be reduced, so that full regeneration of initial conditions is impossible. As water is also present in solvation shells of the dissolved species, the species mentioned in reactions 2–4 do not represent all water molecules participating in the process.

When the polymer coating is fully delaminated, the confined reaction volume disintegrates as no interaction prevents water from spreading on the metal. The consequence is an overall excess of water and the standard monotonous growth of corrosion products.

In conclusion, the ORR at a cathodic delamination front excites chemical oscillations, including an oscillating pH, on corroding Zn. The oscillations involve the formation of a solid

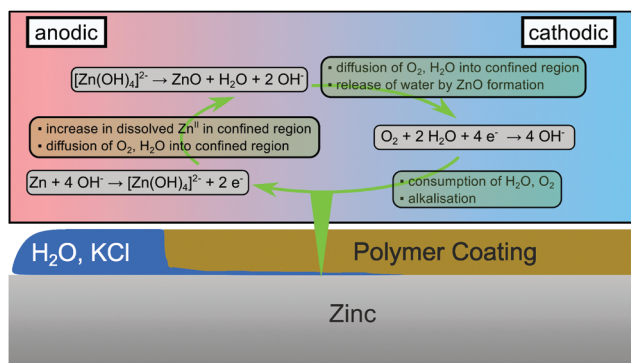


Fig. 4 Scheme of processes occurring consecutively after passage of a cathodic delamination front before complete delamination of the polymer coating.



ZnO phase. Because the solvent and reagent water is present in limited amounts (in analogy to the “crowded” situation in a cell),<sup>51</sup> oscillations are sustained until the sample is fully delaminated. Consequently, after passage of the delamination front, not only the quantity of free water, but also the pH oscillates over a large range. It is unique in the investigated system that the solvent itself takes part in the feedback mechanism leading to oscillations. The observation of oscillations also shows that correlated and not random cathode–anode-structures are present in a delaminating system.<sup>18,21,22</sup> In the field, full delamination is rarely observed, and confinement can be sustained for long times. Modern self-healing concepts are thus only viable if these oscillations are suppressed, or a redox-triggered self-healing mechanism is used.<sup>52</sup>

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