

Spatio-temporal Scanning Electrochemical Microscopy

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Abstract

Scanning Electrochemical Microscopy (SECM) above all else, is a technique to be used to gather information in the spatial domain about a particular system of interest. Even when a time coordinate is assigned to a spatial scan, it is always assigned to the scan as a whole, without any temporal distinction between the individual data points. These however, could have very different time coordinates, because SECM is relatively slow. Scan times are often measured in minutes, during which a dynamic system change dramatically. The difference between the actual, and the assigned time coordinate could lead to difficulties during evaluation.

This paper proposes mixed domain scanning. During the scan, the time coordinates of each individual data acquisition points are stored as well as the spatial ones. From the data, interesting new, spatiotemporal plots can be assembled, which can be used to study dynamically changing systems, and answer questions like how a species of ion is distributed in the system, and how does that distribution change in time. As an example, consecutive scans were recorded on a pre-defined line above a galvanically corroding magnesium sample, and the distribution of hydrogen ions were monitored in time along that line.

Ide még azt kéne leírni, hogy nagy térbeli ÉS időbeli felbontás az SECM-mel lehetetlen, mert a nagy térbeli felbontáshoz sok pont kell, ami sok időbe telik. Mivel sok időbe telik, csak alacsony frekvenciával készíthetők teljes képek. Ezért néha csak egyes vonalpáasztázásokat alkalmaznak, hogy növeljék az időbeli felbontást - pl. korai korróziós események nyomon követésére - és az EGÉSZ vonalhoz egy időpillanatot rendelnek hozzá. Még ez is torzított képet ad a rendszerről, hiszen a vonal pontjai nem egy időben lettek felvéve. Egy másik megközelítés egy érdekes pont kitüntetése, és a jel követése ezen a ponton, ilyenkor természetesen abszolút nincs térbeli felbontás.

Keywords: scanning electrochemical microscopy, potentiometry, spatiotemporal, mixed domain scanning

1. Introduction

Citation [1].

2. Theory

Eq. 1 describes the transient cell response when the measuring electrode is brought to contact with a solution of different analyte activity.

$$E_{cell}(t) = E_{cell}(\infty) + [E_{cell}(0) - E_{cell}(\infty)]e^{-t/RC} \quad (1)$$

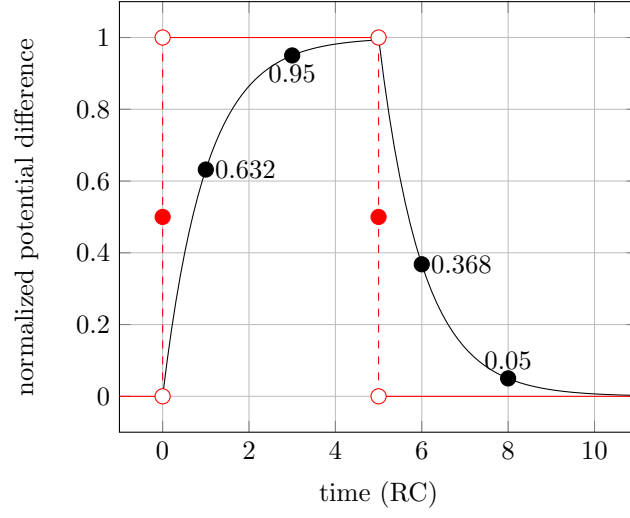


Figure 1: Charging and discharging the series RC circuit. Red: normalized input voltage (V_{in}) to the series RC circuit, two consecutive *Heaviside step functions*, the second one is inverted and shifted $5RC$ to the right. Black: normalized output voltage (V_{out}) of the series RC circuit.

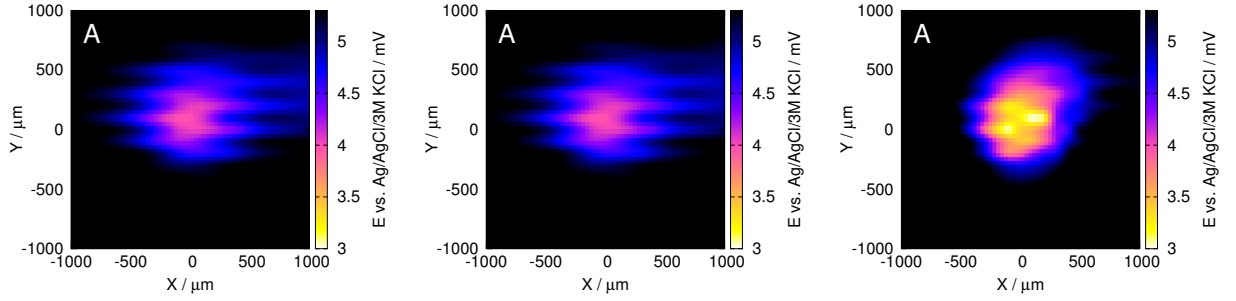


Figure 2: Caption.

Table 1: Comparison of the scanning algorithms.

Algorithm	Number of sampling points	Total scan time (s)	Mean squared error
Meander	441	440	2.75×10^{-2}
Fast comb	441	520	2.07×10^{-2}
Comb	441	881	2.75×10^{-2}
Web	110	109	9.63×10^{-3}
Arc	341	340	2.95×10^{-3}

3. Material and methods

4. Results and discussion

5. Conclusions

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References

- [1] P. J. Eaton, P. West. Atomic force microscopy. Vol. 10. Oxford: Oxford University Press, 2010.

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