



Real-time mass-spectrometric screening of droplet-scale electrochemical reactions

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Real-time mass-spectrometric screening of droplet-scale electrochemical reactions

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A droplet-scale, real-time electrochemical reaction screening platform based on droplet spray ionization mass spectrometry (DSI-MS) has been developed. The N,N-dimethylaniline (DMA) radical cation with a half-life of microseconds was readily detected by MS during the electrooxidation of DMA, and the subsequent reactions followed in real time for minutes.

Determining transient intermediates generated during an electrochemical process is of prime importance to mechanistic studies of electrochemical reactions. In many cases, one-electron¹ or multiple-electrons losses² commonly form a very unstable radical cation. The transient reaction time of such a cation is difficult to detect. Traditional techniques that have been developed for these studies include cyclic voltammetry (CV)³, spectroelectrochemistry⁴, and scanning electrochemical microscopy (SECM)⁵⁻⁷. However, they do not provide direct chemical identification. Combining mass spectrometry (MS) with electrochemistry (EC) is appealing⁸⁻¹² owing to its capabilities for structural characterization, and its high specificity, sensitivity, and speed.

An effective interface between electrochemistry and mass spectrometry for quickly transferring electrochemically generated species from solution to the gas phase for MS analysis is still a challenge. Since EC-MS coupling was described by Bruchkenstein *et al.* in 1971¹³, various ionization methods have been applied for the online coupling of EC-MS, including thermospray¹⁴, fast atom bombardment (FAB)¹⁵, electrospray ionization (ESI)¹⁶, probe electrospray ionization¹⁷, nano desorption electrospray ionization (nanoDESI)¹⁸, and differential electrochemical mass spectrometry (DEMS)^{19, 20}. A sample transfer capillary is commonly used in these EC-MS

couplings to allow electrolytic solution flowing from electrochemical cells to ionization sources. This leads the response times ranged from 0.1 to a few seconds.

Taking advantage of direct sampling, desorption electrospray ionization (DESI)^{21, 22} and easy ambient sonicspray ionization (EASI)²³ reduce the response time to the order of the millisecond scale. In 2015 Zare and coworkers achieved detection of electrogenerated short-lived intermediates by utilizing DESI to directly sample the surface of a rotating waterwheel setup²⁴⁻²⁶ as well as a porous carbon tape with two setups (grooved inclined plane and flat plane)²⁷. In another study by Ran *et al.*²⁸, a hybrid ultramicroelectrode was coupled directly with a mass spectrometer and provided *in situ* information about an electrochemical reaction. However, such techniques require rather complex fabrication. The development of simple and effective screening platform for electrogenerated species is highly desirable and would facilitate mechanistic studies of electrochemical reactions.

The goals of the present work are 1) to develop a droplet-scale, real-time electrochemical reaction screening platform for capturing and characterizing the transient intermediates formed from an electrochemical reaction and 2) to obtain long-time monitoring of electrochemical reaction mechanisms. This setup takes inspiration from the coupling of droplet spray ionization mass spectrometry (DSI-MS)²⁹ with an innovative electrochemical controlling system.

As depicted in Fig. 1, the design employs a glass slip corner positioned in front of the MS inlet, and this slip corner can be both the reservoir for the electrolyte and the spray ionization source. A three-electrode system consisting of a platinum-wire working electrode (WE), a coiled platinum counter electrode (CE), and an Ag/AgCl reference electrode (RE) is mounted on the slip corner. A potentiostat (see Fig. S1, ESI[†]) is used to apply a potential difference across the three electrodes to trigger both the electrochemical reaction and a spray of charged droplets. The electrodes function both as the electrooxidation controller and the high voltage input. The applied voltages are $E + \Delta E$ and E for the WE and CE, respectively. E is the high voltage for the charged spray and ΔE

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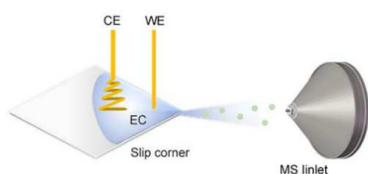
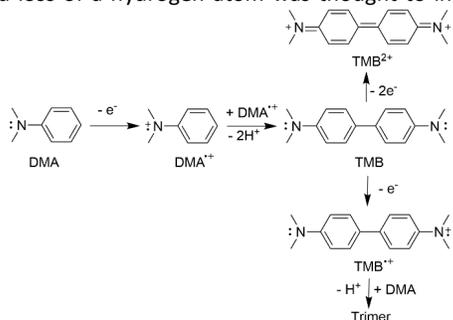


Fig. 1. Schematic representation of the droplet-scale electrochemical reaction screening setup. The reference electrode is not shown. These electrodes function both as the electrode for electrochemistry and the charging source for droplet spray.

is the electrooxidation potential. In addition, the voltage of RE is $E + 0.223$ V and has no influence on the electrochemical reactions. No electrochemical oxidation or reduction is constrained to take place when the WE and CE are held at equal high voltage ($\Delta E = 0$) (see Fig. S2, ESI[†]), indicating that the electrode setup works properly and has no influence on the final detection results.

Electrooxidation of aromatic amines usually gives very unstable radical cations^{1, 30}. One of such examples that has been extensively studied is *N,N*-dimethylaniline (DMA). The DMA radical cation (DMA^{•+}) has a lifetime on the order of microsecond²⁴, which is too short to be detected by conventional methods such as CV³ and spectroscopy³¹. The well-known mechanism for electrooxidation of DMA (Scheme 1) begins with the loss of one electron, leading to the unstable DMA^{•+} cation³². Two DMA^{•+} units rapidly react to yield the dimer, *N,N,N',N'*-tetramethylbenzidine (TMB)^{14, 24}. The electrooxidation then proceeds by loss of one electron from TMB forming TMB^{•+} for the further generation of the trimer in the presence of excess DMA. Loss of two electrons from TMB also occurs during this reaction, which generates the quinoid TMB²⁺¹⁴.

A DMA solution of 0.24 mM was prepared in acetonitrile containing 0.1 mM lithium triflate. When 20 μ L of electrolyte solution was loaded onto the glass slip corner and no potential difference was applied across the electrodes ($E = 4.5$ kV; $\Delta E = 0$ V), the protonated DMA was observed at m/z 122.08 (Fig. 2a). Applying an oxidation potential of 2.0 V across the electrodes ($E = 4.5$ kV; $\Delta E = 2$ V) resulted in the formation of an ion at m/z 121.08, which we ascribe to the DMA radical cation intermediate (Fig. 2b). Upon CID (see Fig. S3, ESI[†]), the observed loss of a hydrogen atom was thought to induce the



Scheme 1. Proposed electrochemical oxidation of DMA.

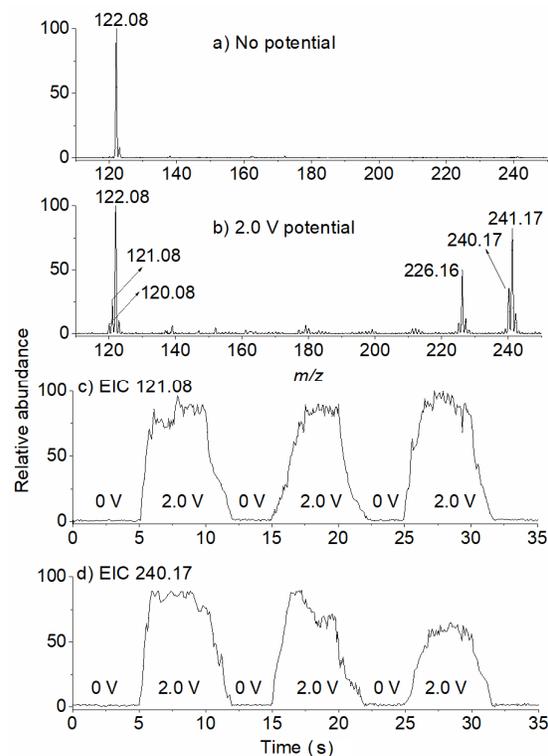


Fig. 2. Full positive-ion-mode mass spectra of DMA with (a) 0.0 V and (b) 2.0 V applied across the electrodes. EIC for the (c) 121.08 peak and (d) 240.17 peak as a function of the applied potential, which was varied between 0.0 V and 2.0 V in 5 s intervals.

formation of a resonance stabilized aromatic imine²⁴. Signals were clearly evident at m/z 241.17 and 240.17, corresponding to the protonated TMB and the radical cation of TMB. Peaks at m/z 226.16 were formed from the loss of CH_2 from the TMB^{•+}. DMA^{•+} and TMB^{•+} were both formed via electrochemical oxidation of DMA, as will be shown below.

The signal intensity of DMA^{•+} increased greatly when an oxidizing potential was applied across the electrodes relative to when no oxidizing potential was applied, as shown in the extracted ion chromatogram (EIC) in Fig. 2c. The MS signal of TMB^{•+} also markedly observed when an oxidizing potential was applied across the electrodes (Fig. 2d). Obviously, the signals of DMA^{•+} and TMB^{•+} were reproducibly observed when an oxidation potential of 2 V was applied compared to that of no oxidation potential (Fig. 2c, 2d). Over time, no significant decrease of DMA^{•+} was observed (Fig. 2c). This may be related to many factors such as short oxidation time and sufficient reactants in solution. As shown in Fig. 2b and 3a, the peak abundances of DMA^{•+} and TMB^{•+} increased over time, indicating that these species were continually formed during the electrooxidation of DMA. This result offers direct and strong evidences for the formation of these species by DMA electrooxidation.

Beyond obtaining real-time information for electrochemical reactions at the initial stage, this droplet-scale electrochemistry screening approach is also applicable to longer real-time monitoring of electrochemical reactions,

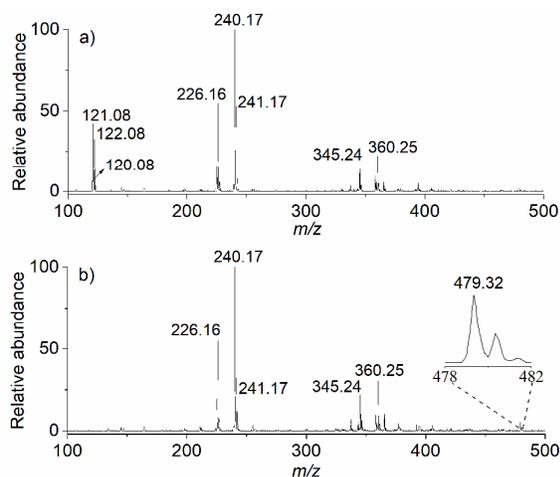


Fig. 3. Full positive-ion-mode mass spectrum of DMA electrooxidation. The mass spectra displayed at the reaction times of (a) 20 s and (b) 90 s.

limited by volatility. Fig. 3 shows monitoring by this method for the duration of minutes. The peak at m/z 120.08 was attributed to the quinoid TMB^{2+} . The protonated trimer (m/z 360.25) was also detected over time (Fig. 2 and 3). A comparison of the mass spectra in Fig. 3a and 3b shows that the peak at m/z 479.32 (protonated tetramer) was unambiguously observed (Fig. 3b inset), although it was detected at low abundance. The trimer and tetramer were also observed even when the DMA was 0.024 mM (Fig. S6), although the relative abundances showed an extent of decrease compared to the concentration of 0.24 mM (Fig. 3b, S6). However, this contrasted with the spectrum using nESI (10 μM DMA) where no signals of trimer and tetramer were detected³³. Besides the concentration, this may be also related to the product transfer efficiency from electrode surface to the spray tip for ionization. No structural information about the trimer and tetramer is available at present. However, this is the first ambient ionization mass spectrometry evidence for the observation of quinoid TMB^{2+} , trimer, and tetramer during the electrooxidation of DMA. DMA electrochemical oxidation as a function of different electrode configurations (wire electrodes, sheet electrodes) has also been studied, and the results are presented in Fig. S4 (ESI[†]).

The electrooxidation of DMA with concentrations ranging from 0.024 to 2.4 mM was also investigated. As shown in Fig. S5 (ESI[†]), peaks of DMA^{2+} were unambiguously observed at different DMA concentrations. In the present work, the great majority of DMA was converted to DMA^{2+} as indicated by higher signal intensities of TMB and TMB^{2+} than that of the protonated DMA. In this regard, we assumed a full conversion of DMA to DMA^{2+} to calculate the half-life of DMA^{2+} , although a complete conversion is impossible. Using the reported rate constant of $2.5 \times 10^8 \text{ M}^{-1} \text{ s}^{-16}$, the half-life of observed DMA^{2+} was calculated to range from 1.7 to 170 μs , depending on the initial starting concentration (see Table S1, ESI[†]). The obtained lifetime cannot be determined by conventional methods such as SECM⁶ and CV³, and is comparable with the results determined by DESI coupling with a waterwheel setup²⁶.

Compared with similar EC-MS methods^{14, 17, 26, 28}, our design has the following merits: (1) This method is simple, easy to fabricate, and can be compatible with a wide range of analysis needs. (2) By using a slip corner as the reservoir for the electrolyte as well as the spray substrate, studies of transient electrogenerated intermediates and size-dependent electrochemical reaction in a droplet is possible. (3) Because the electrolyte droplet on the corner has a capacity of 5 μL up to 60 μL , the maximum spray time can reach about 12 min, limited by solvent evaporation. In this sense, the typical solvent consumption rate is approximate 5 $\mu\text{L}/\text{min}$. Therefore, it achieves a long real-time monitoring on the scale of minutes that offers access to reaction progress and kinetics and to multiple-step reaction monitoring. (4) Replacing the slip corner with a sheet electrode can achieve high electrode surface sampling efficiency, thus facilitating investigation of the chemical composition of an electrode–electrolyte interface³⁴.

In conclusion, by coupling an innovative electrochemistry controlling system with DSI-MS, we have established a droplet-scale real-time electrochemical reaction screening platform. The short-lived DMA radical cation and products (TMB^{2+} , quinoid TMB^{2+} , trimer, and tetramer) resulting from the electrooxidation of DMA were successfully captured and analyzed by MS. These results indicate that isolation of electrogenerated intermediates with half-lives of milliseconds is assessable by employing a slip corner as the reservoir for both the electrolyte and spray ionization source. This platform was also compatible with a wide range of analytical needs such as different concentrations and electrode configurations. Because of the duration of the electrolyte droplet on the corner, the reaction progress on the scale of minutes was readily obtained. All these capabilities bring opportunities to perform and monitor electrochemical reaction studies *in situ*, real-time, and a long-time scale as well. Moreover, this platform has desirable features for studying more complex chemical and biological events in droplets.

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Conflicts of interest

There are no conflicts to declare.

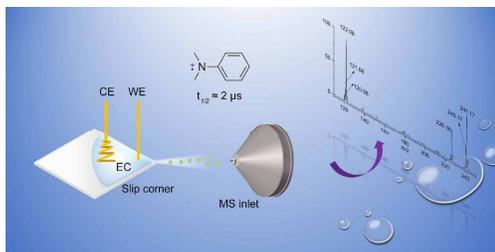
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A real-time electrochemistry-mass spectrometry coupling for detecting the short-lived DMA⁺⁺ and observing DMA reaction propagation is reported.