



An experimentalist's guide to electrocatalysis: the Shono oxidation



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ABSTRACT

Electrosynthesis is a powerful method to functionalise organic molecules without the need to use chemical reagents or protecting groups, yet it is not widely used in synthesis. In this study, we investigated the Shono oxidation of a tertiary amide (electrochemical functionalisation of a C–H bond adjacent to an amide nitrogen atom), demonstrating the value of performing cyclic voltammetry, varying voltage and charge per mole, selection of electrolyte and electrode material. We demystify the process to demonstrate a simple relationship between oxidation potential, and charge transfer required, which affords a high conversion to the desired alpha-methoxylated product using an undivided experimental cell.

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Introduction

Electrochemistry meets all the criteria of green, sustainable chemistry for organic synthesis¹ and can be used for protecting group-free synthesis, CH activation chemistry and umpolung reaction centres, generating complex reaction products from simple starting materials.² In our laboratory, our interest has focused on the Shono oxidation,³ which has the remarkable ability to transform C–H groups adjacent to tertiary amides via an *N*-acyl iminium intermediate, to C–X bonds, for example, C–O, C–C bonds. In light of the current interest in C–H activation chemistry,⁴ we sought to demonstrate how this seminal reaction can be performed in a standard University laboratory set-up. In our recent review of the Shono reaction⁵ we identified a paucity of literature on how to get the best results from this reaction. This can be off-putting to those organic chemists considering performing reactions using electrochemistry. The goal of this article is to demonstrate how a simple screen of three parameters can increase the likelihood of success in electrocatalysis, and make this exciting, emerging and enabling branch of synthesis more widely used by the organic community.

Results and discussion

We selected a simple tertiary amide **1** that meets the criteria of the Shono oxidation, yet has not been systematically investigated as far as we know in electrocatalysis. Although Ebersson et al. reported⁶ a one-off galvanostatic route to **2** in 1979, no spectra or characterisation was reported. This method required a high surface area graphitic rod (800 cm²) and a stainless steel cathode placed 1 mm apart to reduce resistance. An applied voltage of between 20 and 26 V and a current of 50 A was passed through a solution of **1** in methanol with 0.01 M Bu₄NBF₄ as the electrolyte to afford **2** in a current yield of 63% and material yield of 76%. Clearly, this high voltage and current is not attainable in most chemical laboratories and the choice of reaction conditions was not understood nor justified.

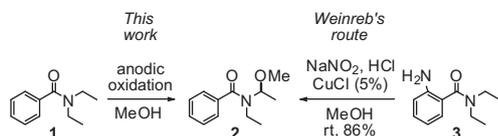
The goal of this study will be to show how model compound **1** can be anodically (electrochemical oxidation) methoxylated to give **2** (Scheme 1) using a simple parameter screen and enabling this sustainable chemistry to be used in synthetic labs, more generally. For comparison, Weinreb and co-workers reported an alternative synthetic route⁷ for the preparation of **2** from **3** using a chemical oxidant and transition metal co-additive (Scheme 1). Of note, Weinreb's route suffered from a dealkylation side reaction due to adventitious water.

Amide **1** was prepared in 85% yield using previously reported chemistry.⁸ Prior to performing electrocatalysis we wished to determine the oxidation potential of **1** in methanol using cyclic voltammetry, however the oxidation peak for **1** to **2** was masked

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Scheme 1. Anodic oxidative methoxylation of **1** to **2** and Weinreb's route to **2** from **3**.

by the oxidation of the solvent beginning at +1.6 V (cyclic voltammograms are located in ESI S1–S4). Therefore, we switched to a solvent with a wider potential window, namely, acetonitrile containing 10% methanol as the co-solvent. Using the relatively cheap electrode material with a large electrode surface area, reticulated vitreous carbon (RVC), as both the working electrode and counter electrode and a silver wire as the reference (see ESI S5 for an example electrode set-up) we obtained typical cyclic voltammograms that exhibited a distinct oxidation peak for **1** which was well resolved prior to the onset of solvent breakdown (Fig. 1).

A scan rate study, as shown in Figure 1b, is a useful voltammetric approach to try and understand the process in hand. In this case, the voltammetry is conducted over a range of scan rates. A plot of peak current (I_p) against scan rate v , and an additional plot of peak current against the square-root of the scan rate allows one to determine if the process is diffusional (since $I_p \sim v^{1/2}$) or adsorbed in nature ($I_p \sim v$) where the plot with the most linear response indicates the dominant process.⁹ In our case, analysis of the data presented in Figure 1 showed a linear response ($I_p/A = 5.16 \times 10^{-4} A/(v/s^{-1})^{1/2} + 4.34 \times 10^{-6} A$; $N = 5$; $R^2 = 0.999$) indicating a diffusional process. Therefore, due to the unique voltammetric signature presented in Figure 1, the appropriate equation for the case of a fully irreversible electron transfer process (not stirred) the Randles–Ševčík equation is as follows:

$$I_p = \pm 0.496(\alpha n')nFAC(FDv/RT)^{1/2}$$

where A is the geometric area of the electrode (cm^2), α is the transfer coefficient (usually assumed to be close to 0.5), n is the total number of electrons transferred per molecule in the electrochemical process, n' is the number of electrons transferred before the rate determining step, F is the Faraday constant, R is the universal gas constant and T is the temperature at which the electrochemical process is performed. This equation clearly shows that temperature,

voltammetric scan rate, the diffusion coefficient and concentration of the analyte under investigation and the electrode area all significantly affect the observed voltammetric signal (I_p). Note that current density (A/m^2) is frequently mentioned in various publications, which from inspection of the units is simply the current (A) divided by the electrode area (m^2), and allows a comparison between experimentalists and represents the current density of the active electrode surface.

In an electrosynthetic experiment, the potential needs to be fixed at a suitable value chosen by the experimentalist. In the literature, we sometimes find cyclic voltammetry exhibits a useful voltammetric signal at +1.5V but then the electrosynthesis is held at an extreme potential of +4.5 V¹⁰ or not using the oxidation potential to enhance selectivity.¹¹ The question is why? In addition, how was this value deduced and what are the implications? If we return to our exemplar Shono electrosynthesis, a key experiment to undertake is a blank voltammogram (Fig. 1a). This is required to understand the exact solvent window (the point in which there is no solvent breakdown) and to ensure that the voltammetric peak of interest is not located in this region, since in addition to the main desired electrochemical process underway; the degradation of the solvent will also occur thus convoluting the electrochemical and electrosynthetic processes (see later). Note the second peak in the voltammogram presented in Figure 1b was due to the electrochemical oxidation of the electrolyte as was evident from comparison of the blank voltammogram (Fig. 1a). With the knowledge that **1** cleanly oxidises at +1.65 V based on the cyclic voltammetry shown in Figure 1b with no interference from solvent degradation, we considered the effect of increasing the potential at which the electrosynthesis was conducted.

Once the voltammetric potential has been carefully chosen, the electrosynthetic reaction (also known as bulk electrolysis) is conducted by holding the reaction at the chosen potential. A common approach is to use chronocoulometry where the total charge (Q) that passes during the time following a potential step is measured as a function of time. Q can be obtained by integrating the current during the applied potential step. In order for the electrosynthetic reaction to go to completion, the amount of charge (C) passed is given by:

$$Q = (m_A/RMM)nF$$

where Q is the charge required to drive the reaction to completion, m_A is the mass (g) of the electroactive analyte, RMM is the relative molecular mass (g mol^{-1}), F is the Faraday constant and n is the

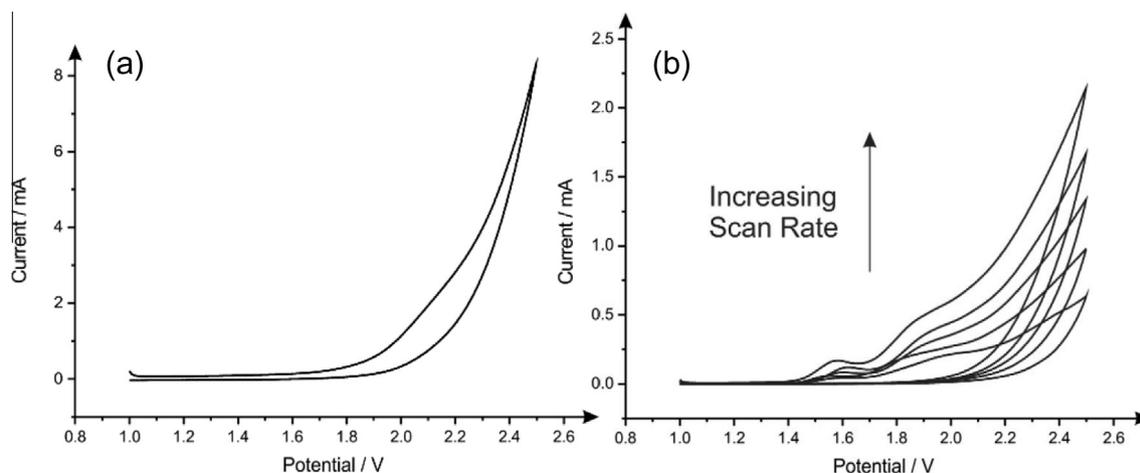


Figure 1. (a): Cyclic voltammetric profile for the blank in MeCN/MeOH (10:1) with 0.47 M TBAP as electrolyte at rates 5 mV/s, 10 mV/s, 25 mV/s, 50 mV/s, 100 mV/s and 250 mV/s at 0 °C. (b): Cyclic voltammograms for **1** in MeCN/MeOH (10:1) with 0.47 M TBAP as electrolyte and 4.7 mM of compound **1** at scan rates of 5 mV/s, 10 mV/s, 25 mV/s, 50 mV/s, 100 mV/s and 250 mV/s at 0 °C.

total number of electrons passed in the electrochemical reaction. Once the amount of charge (C/mol) is known, the chronocoulometry experiment can be performed. Through monitoring the charge, one can determine how long the reaction will take to go to completion. In order to decrease this time, large surface area electrodes and mechanical stirring of the solution are typically employed.

Table 1 demonstrates the percentage conversions obtained for **1** to **2** using a range of voltages (+1.5 to +2.4 V) around the known oxidation potential (+1.65 V). In all cases the temperature was fixed at 0 °C, 4 F/mole was used and identical surface area electrodes were used. All reactions were performed on the same scale and upon reaction completion, the solvent was evaporated and the residue was dissolved in 0.60 mL *d*₆-DMSO (with TMS as the 0.0 ppm internal reference).¹²

It was found that as the voltage of the electro-synthesis reaction increased from +1.5 V to +2.4 V a clear maxima was observed at +1.9 V. However, discoloration of the electrodes (from black to grey and then blue) suggested over reaction and deposition on the electrode surface at this potential which would require replacement of the electrode materials for every reaction, limiting the green potential of this method. As the potential (V) was increased, the potential approaches and becomes outside of the potential window at which point solvent decomposition/degradation occurs. In this instance, the electro-synthetic process becomes complicated by solvent degradation products which affect the efficiency of the formation of the desired product **2**.

In conclusion, based on modification of the voltage parameter, as shown above (**Table 1**) it is pertinent to hold the potential for the electro-synthetic reaction at +1.9 V. Therefore, a simple formula based on cyclic voltammetry can be proposed for this example: Potentiostat voltage = CV oxidation potential (E/V) + a suitably applied over potential voltage (E/V). However, for the following experiments we decided to use +1.8 V for the following variation of the *F*/mole to reduce electrode attrition and reduce the severity of solvent degradation.

The Shono two-electron process should only necessitate the use of 2 *F*/mole, however the use of an excess, often 4 *F*/mole is used. We next investigated whether the reaction can indeed be performed at a lower charge (**Table 2**).

It was evident from **Table 2** that 4 *F*/mole was optimal to achieve the highest conversion. The exact reason as to why this experimental value was twice that required theoretically (see **Scheme 2** for the formation of the intermediate *N*-acyliminium ion **4**) is not fully understood but could possibly be due to the concomitant oxidation of solvent breakdown and/or over-reaction; the exact reason will be considered in future reactions. **Scheme 2** highlights this issue.

Table 1
Variation of potential at a fixed *F*/mole on the percentage conversion of **1** to **2**

Entry	Voltage (V)	<i>F</i> /mole	Time (h)	Normalised % conversion to 2 ^a
1	+1.5	4	2.6	0
2	+1.6	4	2.0	26
3	+1.7	4	3.2	93
3	+1.8	4	2.2	95
4	+1.9	4	2.3	100
5	+2.0	4	1.7	99
6	+2.1	4	0.9	77
7	+2.2	4	1.4	75
8	+2.4	4	0.2	n.d. ^b

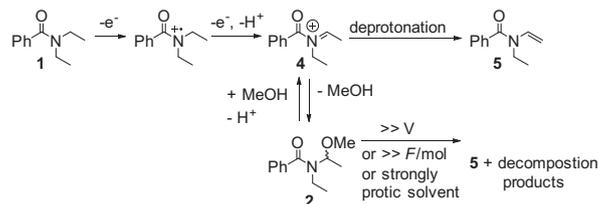
^a The maximal integration of the diagnostic proton at 4.78 ppm is ca. 0.75H due to the existence of rotamers around the amide bond accentuated by unsymmetrical methoxylation of one of the ethyl chains.

^b New aromatic protons were observed in the ¹H NMR spectrum suggesting formation of enamide **5** amongst other side products; n.d. = not determined.

Table 2
The effect of varying *F*/mole. Potential held at +1.8 V in all cases

Entry	<i>F</i> /mole	Time (min)	Normalised % conversion to 2
1	2	17	29
2	3	23	36
3	4	36	95
4	5	43	n.d. ^a

^a Decomposition was observed in the ¹H NMR spectrum, indicative of over-reaction to **5** amongst other side products; n.d. = not determined.



Scheme 2. Postulated mechanism to form **2** via *N*-acyliminium intermediate **4** and possible routes to the side-reaction product, enamide **5** amongst others.

Removal of a single electron from the lone pair of the amide nitrogen atom generates an unstable radical cation. Removal of the second electron via the concomitant expulsion of a proton generates the *N*-acyliminium ion **4** which can be intercepted by methanol to yield **2**. As expected in an acidic environment it is possible for **2** to revert to **4** via an E₂ mechanism. *N*-acyliminium **4** may also deprotonate to enamide **5** via either **2**, **4** or **5** to give a variety of other decomposition products. It was noted that the presence of compounds other than amide **1** or the desired oxidation product **2** could be influenced via increasing the protic strength of the solvent and/or electrolyte (see later), increased charge or the over-voltage applied to the reaction system.

We next considered whether the choice of electrolyte influenced the outcome of the reaction (**Table 3**). Interestingly, the originally selected electrolyte, tetrabutylammonium perchlorate (TBAP),¹³ afforded the highest conversion and yield despite using the same conditions of charge and voltage. Of note, it was found that changing both the electrolyte and solvent resulted in a quantitative conversion to enamide **5** (comparison with Ref. 14) in 100% MeOH. Enamide **5** was not isolated due to instability on silica gel chromatography.

At this point, it was considered whether the perchlorate counter ion accelerated or caused a background chemical oxidation event independent of the electrical voltage applied. It could be seen both from entry 1 in **Table 1** and from a control experiment of **1** in TBAP at 0 °C, that this caused no detectable change to **2**. Therefore, we next sought to optimise the concentration of TBAP employed (**Table 4**).

It was found, that in all cases, conversion of **1** to **2** was achieved but halving the concentration resulted in doubling the time

Table 3
Changing the electrolyte based on the optimal +1.8 V and 4 *F*/mole conditions using MeCN/MeOH; (10:1) unless otherwise stated to convert **1** to **2**

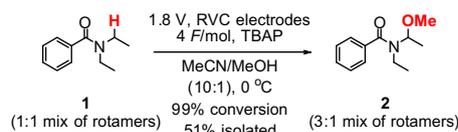
Entry	Electrolyte	Time (min)	Normalised % conversion to 2
1	Bu ₄ NClO ₄ (TBAP)	36	99%
2	Me ₄ NClO ₄ (TMAP)	—	Not soluble
3a	Et ₄ NOTs	60	20%
3b	Et ₄ NOTs ^a	218	0% (99% 5)
4	NaBF ₄	—	Limited solubility
5	NaOSO ₂ Ph	—	Not soluble
6	Bu ₄ NI (TBAI)	—	Not soluble

^a Using 100% MeOH as solvent.

Table 4

Effect of changing the concentration of TBAP using the optimised V and F conditions for the conversion of **1** to **2**

Entry	Electrolyte concentration (M)	Charge (C)	Time (min)
1	0.47	11	36
2	0.24	11.25	67
3	0.12	3.3	188

**Scheme 3.** Optimised electrosynthesis of **2**.**Table 5**

Re-interpretation of the rotamers produced by alpha-methoxylation of **1** to **2** as measured by ¹H NMR spectroscopy

Original assignment ⁷ of 2 in CDCl ₃	Our assignment of 2 in CDCl ₃
7.38–7.27 (m, 5H)	7.36–7.28 (m, 5H)
4.82 and 3.65 (rotamers, 2 bs, 1H)	4.78 (s, 0.75H)
—	3.58 (s, 0.75H)
3.35–3.26 (m, 2H)	3.32–3.26 (m, 1.5H)
3.04 (br s, 3H)	3.00 (s, 2.5H)
1.40–1.38 (d, J = 6.0 Hz, 3H)	1.35 (d, J = 6.0 Hz, 3H)
1.32–1.27 (m, 3H)	1.28–1.23 (m, 3H)

required. Entry 3 was stopped early due to the excessive time required and a spike in the observed resistivity. It appears due to the fact that high levels of resistance encountered in organic media necessitate the use of a high concentration of electrolyte to achieve the reaction in a reasonable time frame.

Taken together, the optimal conditions¹⁵ of electrolyte choice, electrolyte concentration, voltage and Faradays used yielded a preparative conversion of **1** to **2** on a reasonable timescale, without resorting to very large currents and voltages that are only achievable in specialised labs.

The previously optimised conditions were used to convert **1** to **2** in 99% conversion relative to the reference and after extraction and purification by silica gel column chromatography afforded **2** (51% yield) without the need to use transition metals, chemical oxidants or harsh reaction conditions (Scheme 3).

During the course of our analysis of the ¹H NMR spectra associated with **2** and in comparison to Weinreb's spectral data,⁷ a different rotameric ratio of **2** was observed ranging from a 1:1 mix⁷ (generated via a 1,5-H atom radical transfer route) to a ca. 3:1 mixture of rotamers using the alpha-methoxylation route (Table 5).

Conclusions

In conclusion, we have demonstrated that from a rapid cyclic voltammogram measurement, an electrochemical experiment can be undertaken which generates quantitative conversion and isolable amounts of the desired product. This reaction mitigates the use of chemical oxidants and transition group metals to achieve the selective reaction of a C–H bond to give a C–O bond using 'traceless' electrons. This work highlights the various factors that the electrochemical chemist needs to take into consideration. The selection and nature of electrolyte and counter ion are currently under study in our laboratory and will be reported in due course. Together, these results should encourage those wishing to quickly determine whether an electrochemical reaction may work in their organic synthesis and not be a method of last resort. Simply tuning the oxidation potential allows the dial-up of compounds **2**

or **5** from amide **1**. It was also found that increased charge or potential led to degradation products from over-reaction, solvent breakdown and tautomerisation of intermediate **4**.

Acknowledgements

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Supplementary data

Supplementary data (spectroscopic and analytical data including ¹H and ¹³C NMR spectra and cyclic voltammetry of all compounds) associated with this article can be found, in the online version, at <http://dx.doi.org/10.1016/j.tetlet.2015.10.090>.

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- It was envisioned to monitor the diagnostic 4.78 ppm peak for **2** using CDCl₃ but due to overlap of the CHCl₃ un-deuterated reference proton with the phenyl ring in **1** and **2**, *d*₆-DMSO was used for all the other reactions. The data obtained for all the reactions compared favourably either in CDCl₃ or *d*₆-DMSO (ESI S6).
- (a) *Caution*: Tetrabutylammonium perchlorate (TBAP) may intensify fire and is an oxidiser (EU hazard code: H272). Although the safety data sheet for this perchlorate indicates incompatibility with organic materials, sensible precautions allow for the safe use of this material on a preparative scale. For instance, heating may cause an explosion, therefore reactions and work-up are performed at room temperature or below; (b) We thank reviewer 1 for suggesting a control experiment with TBAPF₄ (as in Ref. 6) as a safer electrolyte, however due to limited solubility in acetonitrile a reproducible comparison was not achievable; (c) *Organic Electrochemistry*: 4th ed., *Revised and Expanded*, Lund, H.; Hamerich, O., Eds.; Marcel Dekker: New York, 2001.
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- Representative experimental procedure for the electrochemical synthesis of **1** to generate *N*-ethyl-*N*-(1-methoxyethyl)benzamide **2**: To an undivided glass cell (20 cm³) equipped with a magnetic stirrer along with a rectangular reticulated vitreous carbon (RVC) anode (11 cm²) and rectangular RVC cathode (11 cm²), arranged opposite to one another at a distance of 3.0 mm with a silver wire reference electrode placed 1.0 mm from the working electrode was added **1** (20 mg, 4.73 mM) in acetonitrile (21.6 mL), methanol (2.2 mL) and tetrabutylammonium perchlorate (3.84 g, 0.47 M). The electrolysis was carried out with stirring at a potential of 1.8 V (producing a current density of 7.3 μA/mm² at 0 °C) until a charge of 4.0 F mol⁻¹ had passed. An average current of 5.05 mA passed through the electrodes. The solvent was evaporated under

reduced pressure, and the product was isolated by column chromatography (ethyl acetate/petroleum ether, 10:90 to 15:85) to afford the title compound **2** as a yellow oil, (9.3 mg, 51%). $\nu_{\text{max}}/\text{cm}^{-1}$ 2963, 2913, 2859, 1763, 1738, 1646 and 1550; ^1H NMR (400 MHz, CDCl_3) δ = 7.36–7.28 (m, 5H), 4.78 (s, 0.75H),

3.58 (s, 0.75 H), 3.32–3.26 (m, 1.5H), 3.00 (s, 2.5H), 1.35 (d, J = 6.0 Hz, 3H), 1.28–1.23 (m, 3H); ^{13}C NMR (100 MHz, CDCl_3) δ = 171.9, 136.9, 129.5, 128.6, 126.7, 86.5, 54.7, 34.3, 20.2, 14.4; m/z (ESI) 208 (M+H); Hi-Res LC–MS (ESI) m/z calcd for $\text{C}_{12}\text{H}_{18}\text{NO}_2$ (M+H) 208.1338, found 208.1325.