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Solvent-induced selectivity of Williamson etherification in the pursuit of amides resistant against oxidative degradation†

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This article reports two discoveries. (1) 2-Methoxyethanol induces unprecedented selectivity for etherification of 5-hydroxy-2-nitrobenzoic acids without forming undesired esters. (2) Such compounds are precursors for amides showing unusual robustness against oxidative degradation, essential for molecular electrets that transfer strongly oxidizing holes at about -6.4 eV vs. vacuum.

Introduction

Williamson etherification (WE), involving alcohols and alkanes with good leaving groups as starting materials, remains the most broadly used method for the preparation of ethers.¹ WE offers an important means for adding electron-donating groups to aromatic conjugates, essential for preparing electron-rich *p*-conducting organic materials. Despite its immense importance, the lack of selectivity presents challenges for the utility of WE.^{1c}

To harness dipole effects on charge transfer (CT),² we develop bioinspired molecular electrets that are based on polypeptide structures composed of anthranilamide (Aa) residues.³ (Electrets are systems with ordered electric dipole moments.) The amide and hydrogen bonds of the electrets are a source for their intrinsic dipole moments.⁴ The side chains of Aa, R₁ and R₂ (Chart 1a), on the other hand, provide a means for controlling their electronic properties.^{4a,5} Indeed, electric dipoles present key paradigms for controlling CT, which is of tremendous importance for energy conversion and organic electronics, among many other fields.^{2a} Molecular dipoles rectify CT,^{2b,2c,3c,3d} affecting charge separation and charge recombination.^{3c}

Electron-donating side chains afford hole-transfer electrets.^{5d} For long-range CT in organic materials, it is important to

attain a hopping (or incoherent) mechanism, for which the kinetics exhibits negligible distance dependence beyond about 1 nm.⁶ In order to prevent oxidative degradation of electrets mediating such hole hopping, it is crucial for the comprising Aa residues to form stable radical cations, Aa^{•+}.^{5a-c} For attaining such stability, we have determined that: (1) the spin density distribution (SDD) of Aa^{•+} should not extend over its C-terminal amide;^{5b} and (2) the reduction potentials for oxidizing Aa, should not be too large, *i.e.*, $E_{\text{Aa}^{\bullet+}|\text{Aa}} < 1.5$ V vs. SCE, to prevent the inherent oxidative degradation of the amides.^{5b} The latter places a limit on how oxidizing the transferred holes can be. Hole hopping along moieties with as positive $E_{\text{Aa}^{\bullet+}|\text{Aa}}$ as possible ensures the potency of the holes for attaining large open-circuit voltages and for driving chemical transformations. Because placing alkoxy side chains on Aa residues brings the reduction potentials of their radical cations, Aa^{•+}, to the limit of

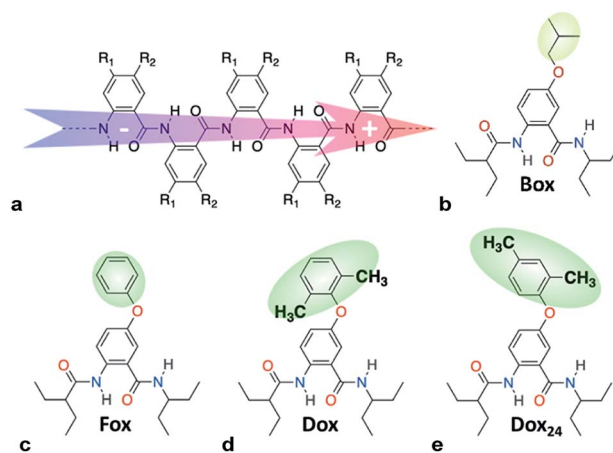


Chart 1 (a) Molecular anthranilamide (Aa) electret with its permanent electric dipole. (b–e) Electret Aa residues with either side chains that are highlighted.

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1.5 V vs. SCE,^{5a-c} such ether conjugates present a key paradigm in the pursuit of organic derivatives that can transduce strongly oxidizing charge carriers.

Established peptide-synthesis procedures, used on aromatic β -amino acids with protected amines, lead to non-reactive cyclic intermediates, preventing the formation of **Aa** derivatives.^{5f,7} Therefore, in the synthesis of **Aa** conjugates, we introduce each residue as its 2-nitrobenzoic acid (**NBA**) analogue.

WE offers a means for synthesizing alkyloxy derivatives of **NBA** from alkyl halides and the corresponding **NBA** hydroxy analogues, **1** (Scheme 1a).^{5a} Because the carboxyl of **1** has a smaller pK_a value than the hydroxyl, both are deprotonated under the basic conditions of WE and can act as nucleophiles, leading to the formation of the esters of the alkoxy **NBA** derivatives, **2** (Scheme 1a).^{5a} This lack of selectivity of WE sets the requirement for an additional step, *i.e.*, hydrolysis, for obtaining the free-acid precursor, **3**, for **Aa** synthesis (Scheme 1a).^{5a}

Herein, we present the discovery of conditions for WE that allow for selective etherification of **1** without forming the undesired esters. Employing 2-methoxyethanol (**2ME**) as a solvent for WE of **1** with 1-iodobutane under microwave irradiation, affords 5-butoxy-2-nitrobenzoic acids, **3**, in quantitative yields ($R = n\text{-C}_4\text{H}_9$, Scheme 1a, Table 1). Reacting **1** with other butyl halides under the same conditions, also results in direct production of **3** without formation of the ester, **2**, or other side products. Reversing the WE paradigm, by taking advantage of the propensity of electron-deficient **NBA** fluorides to undergo nucleophilic aromatic substitution, provides routes to phenoxy **NBA** derivatives (Scheme 1b). As examined with cyclic voltammetry (CV), the electrochemical oxidation of the butoxy **Aa** derivatives barely shows any reversibility at scan rates of about 0.1 V s^{-1} or less. Conversely, some of the phenoxy **Aa** conjugates show not only improved chemical reversibility, but also $E_{\text{Aa}^{\bullet+}|\text{Aa}}$ exceeding 1.5 V vs. SCE by about 200 mV. This result appears surprising because aliphatic and aromatic carboxamides readily undergo irreversible oxidative degradation at about 1.5 V vs.

Table 1 Reaction yields of direct formation of **3** from **1** and X-R^a

-R =	X =		
	I	Br	Cl
-(CH ₂) ₃ CH ₃	97%	34%	15%
-CH ₂ CH(CH ₃) ₂	40%	19%	Trace ^b
-CH(CH ₃)CH ₂ CH ₃	41%	Trace ^b	Trace ^b
-C(CH ₃) ₃	Trace ^b	Trace ^b	N.D. ^c

^a Yields are based on isolated and purified products, **3** (Scheme 1a).

^b Products are not apparent on TLC but are detected using HRMS.

^c Not detected with HRMS.

SCE.⁸ Therefore, these ether **Aa** conjugates have the potential to transduce strongly oxidizing holes making them comparable to a wide range of inorganic CT materials.

Results and discussion

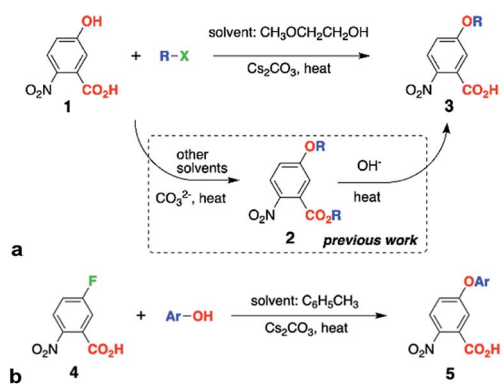
Attaining selectivity for Williamson etherification

Others and we have shown the two-step synthesis of **3**, starting with the preparation of **2** using polar solvent media at about 150 °C (Scheme 1a).^{5a,9} To lower the temperature and considerably shorten the reaction times, we resort to microwave radiation instead of conventional heating. Microwaving mixtures of **1** and various butyl halides dissolved in different solvents that are traditionally used for this reaction, such as *N,N*-dimethylacetamide and *N,N*-dimethylformamide (in the presence of Cs₂CO₃), does not produce even detectable amounts of either **2** or **3**, as confirmed with high-resolution mass spectrometry (HRMS) of the reaction mixtures.

Polar solvents, however, are not usually beneficial for microwave-driven reactions. Medium polarity and hydrogen bonding can, indeed, stabilize transition states and lower activation energies. In such reaction mixtures, however, the polar solvents, rather than the reactants, tend to be the principal absorbers of the microwave energy. Conversely, non-polar solvents with high boiling points ensure that reactants absorb substantial amount of the microwave radiation. This direct transfer of energy to the reactants is a principal advantage that microwave-driven chemical conversions have over reactions under conventional heating.

Using low-polarity solvents, such as toluene and ethyl acetate, allows the formation of only traces of **3**, as detected with HRMS. This minute conversion of **1** to **3** renders it impractical. Medium-polarity aprotic solvents, such as tetrahydrofuran (**THF**) and halogenated alkanes, do not produce even traces of **2** or **3**. This lack of conversion extends even to the neat-reaction conditions, *i.e.*, when the butyl halide reactant is the solvent.

Conversely, when **2ME** is the solvent for the same WE reaction of **1** with butyl halides, we observe the formation solely of the free acid, **3**, as NMR analysis, *i.e.*, NOESY, confirms (see ESI†). Using other alcohols as solvents, such as isopropanol, also affords traces of **3** without **2**, as HRMS reveals. While employing ethylene glycol for WE results in chromatographically detectable amounts of **3**, the minute yields of a few percent



Scheme 1 Etherification of 2-nitrobenzoic acid (**NBA**) derivatives. (a) $-X = -\text{Cl}, -\text{Br}, -\text{I}$; $-R = -(\text{CH}_2)_3\text{CH}_3, -\text{CH}_2\text{CH}(\text{CH}_3)_2, -\text{CH}(\text{CH}_3)\text{C}_2\text{H}_5, -\text{C}(\text{CH}_3)_3$; for direct conversion of **1** to **3**, 1 equiv. **1**, 4 equiv. **RX** and **3** equiv. Cs₂CO₃; microwave heating: $2 \times 30 \text{ s}$, 60 W, temperature set to 130 °C, and ambient-pressure control; or conventional heating: 12 h, pressure tube at 130 °C; (b) $-\text{Ar} = -\text{C}_6\text{H}_5, -2,6\text{-C}_6\text{H}_3(\text{CH}_3)_2$ and $-2,6\text{-C}_6\text{H}_3(\text{CH}_3)_2$; 1 equiv. **4**, 2–4 equiv. **ArOH** and 3–6 equiv. Cs₂CO₃; conventional heating: 12 h, pressure tube at 110 °C.



renders it impractical. Ether solvents, such as THF and 1,2-dimethoxyethane, conversely, do not mediate the conversion of **1** to **3**. While similar to ethylene glycol and 1,2-dimethoxyethane ($R'O-CH_2CH_2-OR''$, R' and $R'' = H$ or CH_3), **2ME** provides a unique reaction environment that under microwave radiation allows for etherification of the phenol OH of **1**, without the side reaction of esterification of the carboxyl. This finding of the **2ME**-mediated selectivity of WE is a key breakthrough of this study, eliminating the need for the second step, *i.e.*, hydrolysis of **2** (Scheme 1a).

As a variation of WE, Purdie–Irvine alkylation, involving Ag_2O as a reagent with high affinity for the formed halide ions, allows for monomethylation of diols.¹⁰ Cs^+ are the only available metal ions in the WE for converting **1** into **3** (Scheme 1a), and they do not have the chelation properties of Ag^+ . Conversely, strongly basic conditions involving treating alcohols with NaH or Na, and excess of these alkoxides when reacting them with chloro- or bromo-acetic acid, produce ethers capped with free carboxylates.^{10c,11} Such conditions keep the media dry and the dryness of the base is important for enhancing the WE yields.¹² In the contrary to the previously reported synthesis of ethers with free carboxylates, we use: (1) a relatively mild base, Cs_2CO_3 , (2) a hydroxyl-carboxyl starting material, **1**, and (3) halides in four-fold excess (Scheme 1a). In addition to its selectivity, all these features represent key advantages of the **2ME**-mediated WE.

The reaction yields of the direct synthesis of **3** from **1** manifest dependence not only on the halide, but also on the alkyl chain. Improving the leaving group, by varying the starting materials from chloride to bromide and iodide, increases the reaction yields (Table 1). Concurrently, branched primary butyls, $X-CH_2CH(CH_3)_2$, show lower yields than the linear ones, $X-(CH_2)_3CH_3$. The yields decrease further for secondary butyls, $X-CH(CH_3)C_2H_5$; and for tertiary, $X-C(CH_3)_3$, we observe only traces of **3** or no product at all (Table 1). These effects of the halides and the alkyl chains on the yields are consistent with the S_N2 mechanism expected for such WE reactions. In addition, the inherent instability of *t*-butyl ethers of electron-deficient phenols reflects the negligible and no yields of **3** when $R = C(CH_3)_3$.

The cleanliness of this WE reaction represents a principal advantage. Reacting **1** with 1-iodobutane produces the *n*-butyl analogue of **3** in quantitative yields (Table 1). For the other butyl halides the isolated-product yields are under 50%. Nevertheless, the reason for these yields was a lack of conversion despite the left over starting material **1**, rather than a production of side products. Increasing the reaction times does not increase the ratio between **3** and **1**, and does not improve the yields.

Transferring this microwave synthetic procedure to pressure-tube settings provides the means for scaling up and for using less expensive conventional heating sources. Heating for 12 hours at 130 °C, *i.e.*, about 5° above the boiling point of **2ME** under normal pressure, results in similar selectivity of the production of **3**. Monitoring the progress of the reaction, using TLC and HRMS, reveals formation of the ester **2**, but with isolated yields that do not exceed 1%. Often, **2** disappears with extended reaction times. This finding suggests that **2ME** has the

perfect proticity resulting in pH of the reaction mixture that ensures the ether formation, but makes the ester unstable. Even if the ester **2** forms, the traces of water that build up with the progress of the reaction can drive its hydrolysis.

Switching the WE pattern allows for pursuing the synthesis of phenoxy **Aa** precursors. Instead of starting with halide derivatives of the side chains (Scheme 1a), we introduce the side chains as the WE nucleophiles (Scheme 1b). Fluorines on electron-deficient aromatic rings are immensely susceptible to nucleophilic aromatic substitution. Hence, introducing the side chain as the corresponding phenolate to fluorinated **NBA**, **4**, allows for making the precursors, **5**, for phenoxy **Aa** residues (Scheme 1b). The reaction yields for the phenoxy and 2,4-dimethylphenoxy analogues of **5**, *i.e.*, the precursors for **Fox** and **Dox₂₄** (Chart 1c and e), are about 90%. The synthesis of the precursor of **Dox**, however, proceeds with yields smaller than 45%, most likely because of the steric hindrance induced by the two methyl groups at the ortho positions of the phenoxy side chain (Chart 1d).

Attempts to employ this nucleophilic aromatic substitution (Scheme 1b) for the synthesis of alkoxy **Aa** derivatives prove unfeasible. While the formation of the alkoxide ions, using metal sodium or strong imide bases, is quite trivial, **4** appears unstable under such extremely basic conditions.

Oxidation reversibility at relatively high potentials

Converting the ether **NBA**s (**3** and **5**, Scheme 1) to their amide analogues capped with alkyls (Chart 1b and e), allows for examining the electrochemical properties of these **Aa** residues.^{5a-c,5e} We select one aliphatic and three aromatic ether derivatives: **Box** (with isobutyl), **Fox** (with phenyl), **Dox** (with 2,6-dimethylphenyl) and **Dox₂₄** (with 2,4-dimethylphenyl) (Chart 1b and e). In **Dox** and **Dox₂₄**, the *ortho* and *para* positions of the electron-donating methyls ensure strong electronic coupling with the **Aa** ring. When both methyls are *ortho* to the oxygen, they can enforce orthogonality between the two aromatic rings and impede rotations around the ether linker, which represent the principal difference between **Dox** and **Dox₂₄** (Chart 1d and e).

At moderate scan rates, **Box** exhibits irreversible oxidation at about 1.5 V vs. SCE for dichloromethane (**DCM**) media (Fig. 1a), which is consistent with the behaviour of other **Aa** residues with alkyloxy side chains.^{5a} Replacing the aliphatic with aromatic ether, methylated at the two *ortho* positions, leads to reversible oxidation at potential of about 1.7 V vs. SCE (*i.e.*, **Box** vs. **Dox**, Fig. 1b and Table 2), which is unprecedented for amides.

At the same scan rates of 0.1 V s⁻¹ and smaller, the other two aromatic-ether residues, **Fox** and **Dox₂₄**, exhibit irreversible oxidation similar to **Box**. Like **Dox**, however, **Fox** and **Dox₂₄** oxidize at more positive potentials than **Box** (Fig. 1a).

Increasing the scan rates beyond 0.5 V s⁻¹ leads to partial oxidation reversibility of **Box**, **Fox** and **Dox₂₄**, as evident from the small cathodic peaks that appear in their voltammograms (Fig. 1c). These findings indicate that the lifetimes of their radical cations are in the order of 1 s, which is sufficiently long for most CT applications. This oxidation reversibility (even if only partial),



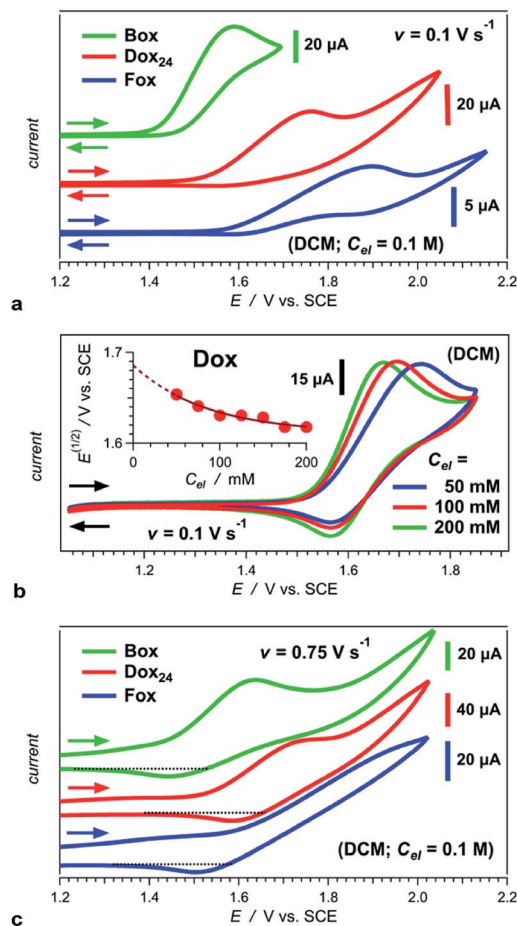


Fig. 1 Cyclic voltammograms of Aa ether residues (Chart 1b and e) for DCM in the presence of an electrolyte, $N(n\text{-C}_4\text{H}_9)_4\text{PF}_6$, with varied concentration, C_{el} . (a) Irreversible oxidation of **Box**, **Fox** and **Dox₂₄** at scan rate, $v = 0.1 \text{ V s}^{-1}$, and $C_{\text{el}} = 100 \text{ mM}$. (b) Oxidation of **Dox** showing reversibility at 0.1 V s^{-1} . Inset: dependence of the half-wave potential on C_{el} and extrapolation of its value for neat DCM, *i.e.*, for $C_{\text{el}} = 0 \text{ M}$. (c) Partial reversibility of the oxidation of **Box**, **Fox** and **Dox₂₄** at 0.75 V s^{-1} .

at potentials as high as 1.7 and 1.8 V vs. SCE, reveals an important effect that ethers have on aromatic amides.

Employing acetonitrile (MeCN) as a solvent yields the same trends as DCM. As expected, the reduction potentials for MeCN are not as positive as those for DCM (Table 2).

The Aa residues with aromatic ethers are harder to oxidize than the one with an alkyloxy group. Adding two methyls to the phenyloxy substituents causes a slight negative shift in the reduction potentials, *i.e.*, **Fox** vs. **Dox** and **Dox₂₄** (Table 2). While the latter agrees with the methyl electron-donating properties, the former appears counter-intuitive assuming that the phenyloxy groups extend the π -conjugation of the residues. Nevertheless, the zero-to-zero energies, ϵ_{00} , of all ether residues are similar, around 3.5 eV (Table 2), regardless the aromatic vs. aliphatic nature of the substituent. It suggests that the resonance effect of the aromatic ethers on the π -conjugation does not necessarily dominate the electronic properties of the residues.

Table 2 Reduction potentials and optical excitation energies of Aa ether residues

	$E_{\text{Aa}^{+/+} \text{Aa}}^{(1/2)}/\text{V vs. SCE}^a$		$\epsilon_{00}/\text{eV}^b$	
	DCM	MeCN	DCM	MeCN
Box	1.56 ± 0.05	1.42 ± 0.07	3.48	3.52
Fox	1.82 ± 0.03	1.57 ± 0.02	3.53	3.56
Dox	1.69 ± 0.02	1.55 ± 0.03	3.52	3.55
Dox₂₄	1.70 ± 0.04	1.56 ± 0.04	3.51	3.55

^a Values for neat solvents, *i.e.*, extrapolated to $C_{\text{el}} = 0 \text{ M}$ (Fig. 1b) from cyclic voltammograms recorded at 0.05 and 0.1 V s^{-1} . For irreversible oxidation, the values of the half-wave potentials at various C_{el} are obtained from the inflection points at the rise of the anodic peaks.¹³

^b Estimated from the wavelengths, λ_{00} , where the normalized absorption and fluorescence spectra cross, $\epsilon_{00} = hv/\lambda_{00}$,^{5c,14} which is a rough approximation for the transition-dipole-moment representation.^{14a}

The Hammett constants and the field Swain–Lupton parameters of phenyloxy groups are more positive than those of alkyloxy substituents.¹⁵ When the conformations suppress the resonance effects, allowing the inductive ones to dominate, the phenyloxy groups are stronger electron-withdrawers than the alkyloxy ones, which is consistent with our findings.

Why do the methyls at the *ortho* positions of the phenyloxy substituent provide extra stability of the radical cations of **Dox**? The sterics of the *ortho*-methyls forces orthogonality between the two aromatic rings of **Dox**, and SDD does not extend over the phenyl side chain. That is, **Dox** and **Box** manifest similar SDD that extends over the N-terminal amide and the ether oxygen (Fig. 2). Conversely, the aromatic rings of **Fox** and **Dox₂₄** are not orthogonal to each other, and the SDD expands over the phenyls of the side chains (Fig. 2).

These contrasts between SDD of **Dox**^{•+} and of the radical cations of the other two aromatic ethers could account for the difference between the reversibility of their oxidation.

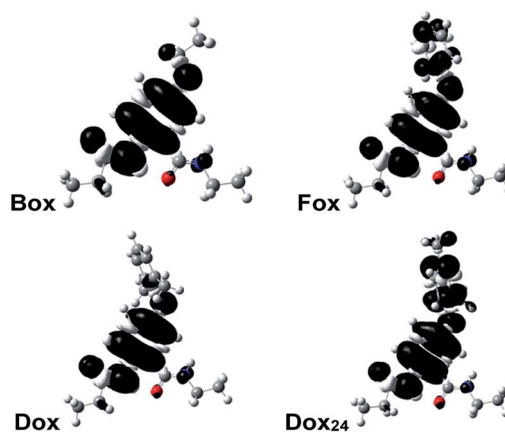


Fig. 2 Spin-density distributions (SDDs) of the radical cations of the Aa ether residues (*anti* conformers) showing the delocalization of the positive charge. For the DFT computations, the alkyls chains are truncated to ethyls.



Conversely, **Dox** shows reversibility at low scan rates, while **Box** does not. Yet, they have quite similar SDDs (Fig. 2).

Two conformers of **Dox** provide orthogonality between its aromatic rings: *i.e.*, with the phenyloxy at *syn vs. anti* orientation in relevance to the C-terminal amide. While the computed energy differences between the *syn*- and *anti*-**Dox** are comparable to the thermal energy, $k_B T$ at $T \approx 20$ °C, the NMR spectra show a single set of peaks for the 2,6-dimethylphenyloxy protons. We ascribe it to an interexchange between the two conformers that is faster than the NMR acquisition timescales. The structural features of **Dox** make the electronic properties of its radical cation similar to those of **Box**^{•+}, rather than **Fox**^{•+} and **Dox**₂₄^{•+}. Unlike **Box**^{•+}, however, **Dox**^{•+} has a polarizable aromatic side chain.

This strategy for decreasing the susceptibility of aromatic amides to oxidative degradation by adding ether substituents is considerably more attractive than an alternative means involving modification of their -CONH- groups. Introducing capping groups with quaternary carbons that connect to the amides of **Aa** residues does not detectably improve the stability of their radical cations. Oligomers of **Aa** residues that oxidize irreversibly, still show irreversible voltammograms.^{5f} Conversely, converting the C-terminal amide, **Aa**-CONH-R, to a tertiary amide, *i.e.*, to **Aa**-CONR₂, induces partial reversibility in the voltammograms of residues that oxidize irreversibly.^{5b} It is consistent with the importance of the proton of the C-terminal amide for the pathways of oxidative degradation. In an **Aa** oligomer (Chart 1a), however, the introduction of such tertiary amides along its back bone would disrupt the hydrogen-bonding pattern that is important for its extended conformation and macrodipole. The C-terminal protons of such oligomers are not involved in the hydrogen bonding network, as NMR studies show.^{3b} Therefore, capping the C-terminus of an **Aa** oligomer with tertiary amide will not disrupt its conformational integrity. Still, computational analysis show that the positive charge of a singly oxidized oligomer, composed of identical **Aa** units, localizes on the N-terminal residue. Therefore, tertiary amide on C-terminus would prove beneficial only if the hole transfer away from the C-terminal residue is slower than its oxidative degradation. In light of this discussion, gaining oxidation reversibility *via* adding ether side chains is a superb strategy because it does not involve disruption of the amides along the backbone that maintain the extended secondary conformation of the **Aa** oligomers.

Conclusions

Ether substituents show important propensity for stabilizing oxidized amides. Ethers are strong enough electron-donating groups to remove the positive charge in the radical cations from the amide groups susceptible to oxidative degradation, but not strong enough to cause drastic negative shifts in reduction potentials. Orthogonality of the aromatic rings of such ether substituents is especially promising for the design of hole-transfer amides. Selectivity for etherification of aromatic hydroxyls in the presence of other nucleophilic groups is important for the synthesis of such charge-transfer conjugates

that, along with their newly discovered electronic properties, opens new doors for organic electronics and energy science.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- (a) A. Williamson, *Philos. Mag. Ser. 3*, 1850, **37**, 350–356; (b) R. Gigg and P. M. Carroll, *Nature*, 1961, **191**, 495–496; (c) M. Sauthier, A. Mortreux and I. Suisse, in *Carbohydrate Chemistry*, ed. A. P. Rauter, T. K. Lindhorst and Y. Queneau, Royal Society of Chemistry, 2014, vol. 40, pp. 73–98; (d) S. Mandal, S. Mandal, S. K. Ghosh, P. Sar, A. Ghosh, R. Saha and B. Saha, *RSC Adv.*, 2016, **6**, 69605–69614.
- (a) J. B. Derr, J. Tamayo, E. M. Espinoza, J. A. Clark and V. I. Vullev, *Can. J. Chem.*, 2018, **96**, 843–858; (b) E. Galoppini and M. A. Fox, *J. Am. Chem. Soc.*, 1996, **118**, 2299–2300; (c) S. Yasutomi, T. Morita, Y. Imanishi and S. Kimura, *Science*, 2004, **304**, 1944–1947; (d) C. Shlizerman, A. Atanassov, I. Berkovich, G. Ashkenasy and N. Ashkenasy, *J. Am. Chem. Soc.*, 2010, **132**, 5070–5076; (e) Y. Chen, J. Viereck, R. Harmer, S. Rangan, R. A. Bartynski and E. Galoppini, *J. Am. Chem. Soc.*, 2020, **142**, 3489–3498.
- (a) V. I. Vullev, *J. Phys. Chem. Lett.*, 2011, **2**, 503–508; (b) B. Xia, D. Bao, S. Upadhyayula, G. Jones and V. I. Vullev, *J. Org. Chem.*, 2013, **78**, 1994–2004; (c) D. Bao, S. Upadhyayula, J. M. Larsen, B. Xia, B. Georgieva, V. Nunez, E. M. Espinoza, J. D. Hartman, M. Wurch, A. Chang, C.-K. Lin, J. Larkin, K. Vasquez, G. J. O. Beran and V. I. Vullev, *J. Am. Chem. Soc.*, 2014, **136**, 12966–12973; (d) M. Krzeszewski, E. M. Espinoza, C. Cervinka, J. B. Derr, J. A. Clark, D. Borchardt, G. J. O. Beran, D. T. Gryko and V. I. Vullev, *Angew. Chem., Int. Ed.*, 2018, **57**, 12365–12369.
- (a) M. K. Ashraf, R. R. Pandey, R. K. Lake, B. Millare, A. A. Gerasimenko, D. Bao and V. I. Vullev, *Biotechnol. Prog.*, 2009, **25**, 915–922; (b) S. Upadhyayula, D. Bao, B. Millare, S. S. Sylvia, K. M. M. Habib, K. Ashraf, A. Ferreira, S. Bishop, R. Bonderer, S. Baqai, X. Jing, M. Penchev, M. Ozkan, C. S. Ozkan, R. K. Lake and V. I. Vullev, *J. Phys. Chem. B*, 2011, **115**, 9473–9490.
- (a) J. M. Larsen, E. M. Espinoza, J. D. Hartman, C.-K. Lin, M. Wurch, P. Maheshwari, R. K. Kaushal, M. J. Marsella, G. J. O. Beran and V. I. Vullev, *Pure Appl. Chem.*, 2015, **87**, 779–792; (b) E. M. Espinoza, J. M. Larsen and V. I. Vullev, *J. Phys. Chem. Lett.*, 2016, **7**, 758–764; (c) J. M. Larsen-Clinton, E. M. Espinoza, M. F. Mayther, J. Clark, C. Tao, D. Bao, C. M. Larino, M. Wurch, S. Lara and V. I. Vullev, *Phys. Chem. Chem. Phys.*, 2017, **19**, 7871–7876; (d) J. M. Larsen,



- E. M. Espinoza and V. I. Vullev, *J. Photonics Energy*, 2015, **5**, 055598; (e) E. M. Espinoza, J. M. Larsen-Clinton, M. Krzeszewski, N. Darabedian, D. T. Gryko and V. I. Vullev, *Pure Appl. Chem.*, 2017, **89**, 1777–1797; (f) K. Skonieczny, E. M. Espinoza, J. B. Derr, M. Morales, J. M. Clinton, B. Xia and V. I. Vullev, *Pure Appl. Chem.*, 2020, **92**, 275–299.
- 6 J. Jortner, M. Bixon, T. Langenbacher and M. E. Michel-Beyerle, *Proc. Natl. Acad. Sci. U. S. A.*, 1998, **95**, 12759–12765.
- 7 Y. Hamuro, S. J. Geib and A. D. Hamilton, *J. Am. Chem. Soc.*, 1996, **118**, 7529–7541.
- 8 (a) J. F. O'Donnell and C. K. Mann, *J. Electroanal. Chem. Interfacial Electrochem.*, 1967, **13**, 157–162; (b) T. Golub and J. Y. Becker, *Org. Biomol. Chem.*, 2012, **10**, 3906–3912.
- 9 (a) A. Andersson, J. B. Baell, P. J. Duggan, J. E. Graham, R. J. Lewis, N. G. Lumsden, C. E. Tranberg, K. L. Tuck and A. Yang, *Bioorg. Med. Chem.*, 2009, **17**, 6659–6670; (b) H. Bertrand, R. Guillot, M.-P. Teulade-Fichou and D. Fichou, *Chem.–Eur. J.*, 2013, **19**, 14654–14664; (c) A. F. Donnell, Y. Zhang, E. M. Stang, D. D. Wei, A. J. Tebben, H. L. Perez, G. M. Schroeder, C. Pan, C. Rao, R. M. Borzilleri, G. D. Vite and S. Gangwar, *Bioorg. Med. Chem. Lett.*, 2017, **27**, 5267–5271.
- 10 (a) T. Purdie and J. C. Irvine, *J. Chem. Soc., Trans.*, 1903, **83**, 1021–1037; (b) A. Bouzide and G. Sauve, *Tetrahedron Lett.*, 1997, **38**, 5945–5948; (c) A. Tuchscherer, D. Schaarschmidt, S. Schulze, M. Hietschold and H. Lang, *Dalton Trans.*, 2012, **41**, 2738–2746.
- 11 O. Zech, M. Kellermeier, S. Thomaier, E. Maurer, R. Klein, C. Schreiner and W. Kunz, *Chem.–Eur. J.*, 2009, **15**, 1341–1345.
- 12 B. Aribat, Y. Le Bigot and A. Gaset, *Tetrahedron*, 1996, **52**, 8245–8256.
- 13 (a) E. M. Espinoza, J. A. Clark, J. Soliman, J. B. Derr, M. Morales and V. I. Vullev, *J. Electrochem. Soc.*, 2019, **166**, H3175–H3187; (b) A. Purc, E. M. Espinoza, R. Nazir, J. J. Romero, K. Skonieczny, A. Jeżewski, J. M. Larsen, D. T. Gryko and V. I. Vullev, *J. Am. Chem. Soc.*, 2016, **138**, 12826–12832; (c) E. M. Espinoza, B. Xia, N. Darabedian, J. M. Larsen, V. Nunez, D. Bao, J. T. Mac, F. Botero, M. Wurch, F. Zhou and V. I. Vullev, *Eur. J. Org. Chem.*, 2016, **2016**, 343–356; (d) D. Bao, S. Ramu, A. Contreras, S. Upadhyayula, J. M. Vasquez, G. Beran and V. I. Vullev, *J. Phys. Chem. B*, 2010, **114**, 14467–14479; (e) D. Bao, B. Millare, W. Xia, B. G. Steyer, A. A. Gerasimenko, A. Ferreira, A. Contreras and V. I. Vullev, *J. Phys. Chem. A*, 2009, **113**, 1259–1267.
- 14 (a) G. Angulo, G. Grampp and A. Rosspeintner, *Spectrochim. Acta, Part A*, 2006, **65A**, 727–731; (b) S. Guo, D. Bao, S. Upadhyayula, W. Wang, A. B. Guvenc, J. R. Kyle, H. Hosseinibay, K. N. Bozhilov, V. I. Vullev, C. S. Ozkan and M. Ozkan, *Adv. Funct. Mater.*, 2013, **23**, 5199–5211.
- 15 (a) C. Hansch, A. Leo and R. W. Taft, *Chem. Rev.*, 1991, **91**, 165–195; (b) C. G. Swain and E. C. Lupton Jr, *J. Am. Chem. Soc.*, 1968, **90**, 4328–4337.

