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Regioselective addition of DDQ on a quinoid ring: an entry into chiral zwitterionic bridging ligands

Gabriel Canard,^{*a} Zhongrui Chen,^a Aprilliyan Suryaningtias,^a Marion Jean,^b Nicolas Vanthuyne,^b Michel Giorgi,^c Christian Roussel^b and Olivier Siri^{*a}

The regioselective insertion of DDQ into a C-H bond of the $6\pi + 6\pi$ electron zwitterionic benzoquinonemonoimines **4a-c** results in the formation of the novel chiral C-substituted quinoid ligands **11a-c**. These Michael adducts feature a preserved zwitterionic form and a quaternary stereogenic carbon center as evidenced by the single crystal X-ray structure of the derivative **11a**. The ECD spectra, optical rotations and racemization barriers of the two enantiomers of **11a** were measured subsequently to their separation by a preparative chiral HPLC. Because the racemization of **11a** is stopped in basic media, compounds **11a-c** give a new entry in chiral zwitterionic bridging ligands.

Introduction

The 2,5-dihydroxy-1,4-benzoquinone **1** (Figure 1) featuring a [O,O,O,O] donor set has been extensively used as a bridging ligand to build polymetallic complexes where the effective intermetallic communication can lead, for example, to spin-spin couplings or mixed-valence states.¹ During the last decades, several new classes of bridging ligands were developed by replacing from one to four of the "O" groups of **1** by isoelectronic "NR" ones producing the "amino/imino" quinoid ligands **2-6** (Figure 1).² These replacements can tune the physico-chemical properties of the resulting metallic complexes by introducing unsymmetrical donor sets and/or by selecting the steric and electronic characteristics of the R substituents.² Interestingly, when two amino fragments are introduced on the quinoid skeleton in **3** and **4**, two different electronic distributions are obtained. Indeed, if the quinoidal structure is retained in the structure of **3** having a [N,O,N,O] donor set, the [N,N,O,O] coordinating system of **4** induces an intramolecular proton transfer producing a zwitterion being more stable than its corresponding canonical form (Figure 1).³

This zwitterionic form gives rise to specific physico-chemical properties such as a permanent electric dipole that can guide the self-assembly of **4** derivatives on surfaces.⁴ The zwitterion **4** is also, as the other members of **1-6** series, an efficient coordinating ligand and was incorporated into metallic complexes⁵ that were applied in catalysis,⁶ optical recording,⁷ electrochromism⁸ or vapochromism.⁹ On the other hand, this particular electronic distribution of **4** gives rise to two different kinds of reactivity towards organic and electrophilic reagents (Figure 2).

Fig 1. Stepwise replacement of the "O" groups of **1** by isoelectronic "NR" ones in bridging quinoid ligands **2-6**.

Fig 2. Previously reported reactions of **4** with electrophilic organic reagents.

The addition of a strong acid (HCl) or of a strong alkylating agent ($(R'_3O)BF_4$) on **4** produces the localization of the upper 6 π electron system through the formation of a single O-R' bond ($R' = H, Me$ or Et) in the positively charged derivatives **7** and **8** (Figure 2).¹⁰ On the other hand, the reaction between **4** and the organic electron acceptors tetracyanoethylene (TCNE) or 7,7',8,8'-tetracyanoquinodimethane (TCNQ) yielded the formation of the unexpected adducts **9** and **10** where a regioselective carbon-carbon bond was formed between the two starting components and was preserving the zwitterionic form thanks to an intramolecular proton transfer (Figure 2).¹¹ This insertion of TCNE or TCNQ into the central upper C-H bond was shown to bring from slight to important changes in the coordination and self-assembling properties of **9** and **10** derivatives compared to those of **4**.¹²

The present work illustrates how the addition of 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) in the upper C-H bond adjacent to the $C \equiv O$ bonds of **4** give rise to novel chiral zwitterionic ligands. If the single reported chiral derivatization of **4** was produced by the introduction of chiral amines on its lower N-C-C-C-N part,^{12e} the present one concerns the upper O-C-C-C-O moiety and follows a Michael-type addition. This condensation is accompanied by i) the preservation of the zwitterionic form thanks to an intramolecular proton transfer and ii) the creation of a chiral quaternary carbon center which allowed the separation and characterization of each enantiomer of one **4**-DDQ adduct.

Results and discussion

Organic synthesis.

The reaction between DDQ and the $6\pi+6\pi$ zwitterions **4a-c** at room temperature in a 1:1 ratio results in the progressive formation of orange-red precipitate of the adducts **11a-c** (Figure 3). A first proof of the insertion of DDQ in the C-H bond of **4a-c** was given by the corresponding ^1H NMR and ^{13}C NMR spectra recorded in DMSO- d_6 . For example, the two $C(sp^2)$ -H signals of **4a** located at $\delta = 4.95$ and $\delta = 5.47$ ppm were replaced by a single downfield shifted singlet at $\delta = 5.59$ ppm in the spectrum of **11a** (see the supporting information). If the disappearance of the downfield shifted ^1H NMR signal of H_a could result from the expected C-C oxidative coupling between two molecules of **4a**,¹³ this hypothesis was excluded by the presence of 16 signals in the ^{13}C NMR spectrum of **11a** whose structure was confirmed by high resolution mass-spectrometry and single crystal X-ray diffraction analyses (see below).

Single crystal X-ray diffraction.

The slow cooling of a concentrated toluene solution of **11a** afforded diffracting single crystals suitable for an X-ray structure analysis (Figure 4). Details of the structure determination are given in the experimental part and in Table S1 (see the supporting information). As observed previously with TCNE and TNCQ, the molecular structure of **11a** revealed that a single C-C bond with a length of 1.535(9) Å has been regioselectively formed between DDQ and the zwitterion **4a**.

Fig 3. Reactions of **4a-c** with DDQ.

This insertion is occurring in the C-H bond adjacent to the two C $\equiv$ O bonds of **4a** which retains its zwitterionic form characterized by an equalization of the C-C and C-O bond lengths of its upper O-C-C-C-O moiety as well as the equal C-C and C-N bond lengths of its lower N-C-C-C-N moiety. The newly formed C-C bond involves one of the two DDQ carbon atoms bearing a cyano group as already observed when such addition occurred between DDQ and various nucleophilic carbon atoms of, for example, silyl enol ethers,¹⁴ enamines¹⁵ or indoles.¹⁶ If the C=O bond of the DDQ moiety adjacent to the quaternary carbon atom remains a double bond with a length of 1.203(8) Å, the formation of the zwitterion **11a** is resulting from a proton transfer from the two starting components confirmed by the new C-OH single bond length of 1.334(8) Å. The crystal packing of the racemic **11a** revealed that each enantiomer of **11a** is interacting with the other one thanks to a π - π interaction producing a “tête-bêche” arrangement (Figure 4b).

Fig 4. ORTEP views of the crystal structure of **11a** showing: a) one enantiomer of **11a**; b) the π - π stacking interaction occurring between the two enantiomers of **11a**.

Table 1 Chiral HPLC data ((*S,S*)-Whelk-O1) and specific optical rotations of the two enantiomers of **11a**.

		t(-) (min) ^a	k(-) ^a	t(+) (min) ^a	k(-) ^a	α	Rs	$[\alpha]_{546}^{25}$ ^b	ee
Analytical separation	1 st eluted enantiomer	5.47	0.85						
	2 nd eluted enantiomer			7.73	1.62	1.90	4.02		
Preparative separation	1 st collected fractions	5.48		8.10				+ 650 ($\pm$ 50) ^c	> 99%
	2 nd collected fractions	5.65		7.61				- 650 ($\pm$ 50) ^d	> 96%

^a Detected CD sign at 254 nm. ^b Recorded in methanol. ^c with c = 0.16 mg.cm⁻³. ^d with c = 0.15 mg.cm⁻³.

Chiral HPLC separation and chirality analysis.

The DDQ insertion in the C-H bond of **4a-c** results in the formation of a quaternary stereogenic carbon atom in the corresponding adducts **11a-c**. Consequently, an analytical chiral HPLC analysis of **11a** was conducted in order to see if its two enantiomers could be separated and therefore characterized. After screening different chiral stationary phases and solvents, a so called (*S,S*)-Whelk-O1 column was chosen with a mobile phase combining *n*-heptane, dichloromethane and ethanol containing 0.1% of trifluoroacetic acid (TFA) in an overall 10/10/80 ratio. The addition of an organic acid was required to ensure the elution of the highly polar **11a**. These experimental conditions afforded a good enantioselectivity factor α (1.90) and a high resolution *R*_s (4.02) for the enantiomeric separation of **11a** (Table 1 and Figure 5). Nevertheless, we observed the enantiomerization of each separated enantiomer in the acidic mobile phase during the storage of their collected fractions. A subsequent analysis in the acidic mobile phase gave a corresponding enantiomerization barrier of 96.2 kJ.mol⁻¹ and a half-time value of 65 minutes at 25°C (see the experimental part and the supporting information).

If this racemization process is quite fast in acidic medium, it is completely stopped when a concentrated ethanol solution of sodium carbonate is added during the collection of the separated fractions.

Fig 5. Chromatograms of the chiral HPLC analysis of the racemic **11a**: UV and Circular Dichroism detection at 254 nm on (*S,S*)-Whelk-O1 using heptane / ethanol and tfa / dichloromethane (1/8/1) as a mobile phase and at 1 mL.min⁻¹.

Therefore, using this strategy and the same experimental conditions (stationary and mobile phases), a chiral separation was conducted on a preparative scale giving up to 25 mg of each enantiomer of **11a** with an optical purity exceeding 96% (Table 1). The two enantiomers of **11a** give an identical UV-visible spectrum (recorded in the neutralized mobile phase) featuring three intense absorption bands located at ca. 220, 340 and 350 nm that were previously attributed to $\pi \rightarrow \pi^*$ transitions of the zwitterionic part (Figure 6).¹⁷ On the other hand, the fourth band at ca. 270 nm could be due to the reduced DDQ moiety.¹⁸ The ECD spectrum of the second eluted enantiomer features multiple Cotton effects and is, as expected, the mirror-image of the ECD spectrum recorded for the first eluted one (Figure 6). The same symmetry is observed when measuring the values of the optical rotations in methanol at 546 nm which reaches up to + 650° for the first eluted enantiomer (Table 1).

Fig 6. UV-visible (top) and ECD spectra (below) of the first (green solid line) and the second (red dotted line) eluted enantiomers of **11a** (recorded in methanol).

Racemization mechanism

If the addition of DDQ on **4a-c** is extending the number of such zwitterionic C-substituted derivatives, it is particularly appealing for its concomitant and unprecedented introduction of a chiral centre appended on the upper O-C-C-C-O moiety. This functionalization would be complementary to the single previously reported chiral derivatization of **4** that was performed thanks to the introduction of chiral amines on its lower N-C-C-C-N part.^{4e} Considering that the use of **11a-c** as chiral ligands requires experimental conditions that prevent its racemization, two preliminary experiments were conducted to give first insights of the corresponding racemization process.

The first one was the determination of the racemization barrier that showed a first-order kinetic when using an acidic solution (see the supporting information). If the Michael addition is often used to create chiral centres, it can compete with its reversible dissociation in a Michael–retro-Michael tandem process.¹⁹ This process was found to be responsible of the racemization of **11a** because the ¹H NMR spectrum of an equimolar mixture of **11a** and **4c** (in DMSO-d₆) produces in less than 20 minutes the appearance of traces of the cross coupling products **4a** and **11c** (Figure 7 and Figure S2 in the SI). Within 18 hours,

Chemical reaction scheme showing the equilibrium between two pairs of chemical structures. The structures are labeled 4c, 11a, 4a, and 11c. The structures are shown in equilibrium with each other, indicated by a double-headed arrow.

= 180.3, 169.1, 155.2, 154.3, 142.9, 135.0, 115.5, 115.1, 106.3, 87.7, 81.9, 46.9, 30.0 ppm; UV-Vis (DMSO): λ_{max} ($\epsilon \times 10^{-3}$) = 344 (25.0), 356 (25.9), 478 nm (1.4); HRMS (ESI-TOF): calcd. for $\text{C}_{16}\text{H}_{11}\text{N}_4\text{O}_4\text{Cl}_2^+$ ($[\text{M}+\text{H}]^+$) 393.0152, found 393.0151.

7c: To a solution of **4c** (40 mg, 0.177 mmol, 1 equiv.) in MeOH (4 mL) was added DDQ (41 mg, 0.181 mmol, 1.02 equiv.). The mixture was stirred at room temperature overnight. The resulting precipitate was filtered off, washed successively with MeOH and CH_2Cl_2 before it was dried under vacuum affording the desired product **11c** as an orange powder (36 mg, 0.079 mmol, 45%); ^1H NMR (400 MHz, DMSO- d_6 , 294 K): δ = 9.23 (brs, 2H), 5.73 (s, 1H), 3.62 (m, 4H), 3.48 ppm (m, 4H); ^{13}C NMR (100 MHz, DMSO- d_6 , 294 K): δ = 180.4, 168.9, 155.0, 154.6, 143.4, 134.9, 115.6, 115.4, 106.3, 87.1, 82.5, 58.7, 47.0, 45.7 ppm; UV-Vis (DMSO): λ_{max} ($\epsilon \times 10^{-3}$) = 347 (25.3), 358 (25.5), 491 nm (1.7); HRMS (ESI-TOF): calcd for $\text{C}_{18}\text{H}_{15}\text{N}_4\text{O}_6\text{Cl}_2^+$ ($[\text{M}+\text{H}]^+$) 453.0363, found 453.0363.

Chiral HPLC

The chiral HPLC analyses and separations were performed on Agilent 1260 Infinity unit (pump G1311C, autosampler G1329B, DAD G1365D). A Jasco CD-2095 circular dichroism detector is implemented for the chiroptical detection. The sign given by the on-line circular dichroism at 254 nm is the sign of the compound in the solvent used for the chromatographic separation. The preparative chiral HPLC separation was performed using an Agilent 1260 Infinity unit (pump G1311C, autosampler G1329B, DAD G1365D and fraction collector G1364C). The (*S,S*)-Whelk-O1 analytical (250 x 4.6 mm) and preparative columns (250 x 10 mm, 5 μm) are purchased from Regis Technologies (Morton Grove, USA) and are mainly composed of (3*R*,4*S*)-4-(3,5-dinitrobenzamido)-3-[3-(dimethylsilyloxy)propyl]-1,2,3,4-tetrahydrophenanthrene molecules (selectors) bonded onto silica. HPLC grade Heptane, ethanol and dichloromethane, were degassed and filtered on a 0.45 μm millipore membrane before use. Retention times R_t (in minutes), retention factors $k_i = (R_t - R_{t0})/R_{t0}$, enantioselectivity factors $\alpha = k_2/k_1$ and resolution $R_s = 1.18 (R_{t2} - R_{t1}) / (w_1 + w_2)$ are given. R_{t0} is determined by the injection of tri-tertio-butyl benzene and w_i is the peak width at half-height.

The analysis of the **11a** enantiomer's racemization in the acidic mobile phase was performed starting from a solution of ca. 0.5 mg of the second eluted enantiomer dissolved in 1 mL of the mobile phase and thermostated at 25 °C. Every 12 minutes, 10 μL of this solution was injected on the chiral phase giving a decreasing percentage of the second eluted enantiomer that was monitored.

Single crystal X-ray diffraction

The intensity data for compound **11a** were collected on a Rigaku Oxford Diffraction SuperNova diffractometer using $\text{CuK}\alpha$ radiation ($\lambda = 1.54184 \text{ \AA}$). Data collection, cell refinement and data reduction were performed with CrysAlisPro (Rigaku Oxford Diffraction). The structure was solved with shelXT²⁰ and shelXL²⁰ was used for full matrix least squares refinement. The crystal appeared to be twinned and both domains were refined together using the HKLF5 shelXL instruction to final values of BASF equal to 0.553(2) and 0.447(2) respectively. One aminobutyl was found to be disordered and was refined on two sites of equal occupation 0.5. The hydrogen atoms were all found experimentally except those of the disordered moiety that were introduced at geometrical positions. Their Uiso parameters were fixed to 1.2Ueq(parent atom) for the aromatics and CH_2 carbons and the amine and to 1.5Ueq(parent atom) for the methyls and OH group. CCDC-1572929 (**11a**) contains the supplementary crystallographic data. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif.

Conclusions

The addition of the electron accepting DDQ on the benzoquinonemonoimine **4a-c** results in a C-C bond formation occurring regioselectively on the O-C-C-O part of the zwitterionic reactant. This Michael type addition produces the chiral adducts **11a-c** which retain a zwitterionic form thanks to an intramolecular proton transfer. The optically pure enantiomers of these new quinoidal bridging ligands can be separated and characterized as long as a basic medium is used to prevent their acid-catalysed racemizations that are probably following a retro-Michael process. This unprecedented chiral functionalization of the zwitterionic **4** offers a new entry in chiral zwitterionic quinoidal bridging and coordinating ligands that could be used, for example, in the design of chiral polymetallic structures or redox-active complexes build previously on the parent compound **1** and its derivatives.²¹

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