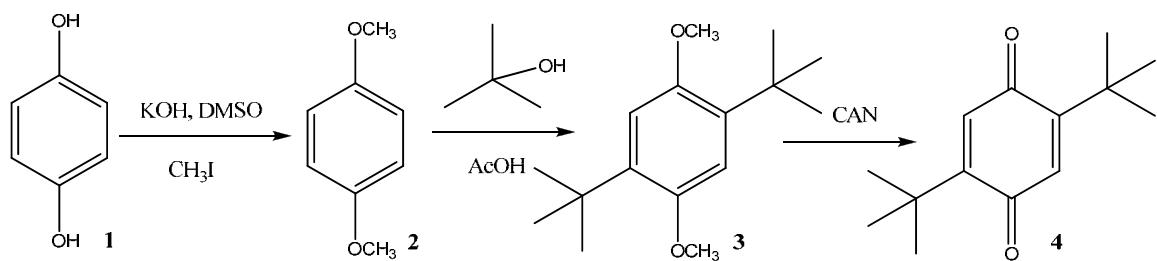


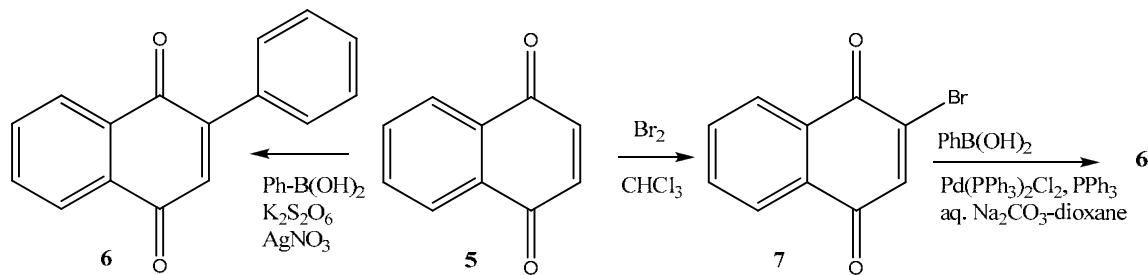
**Synthesis and electrochemical redox properties of arylated *p*-benzoquinones,
naphthoquinones and alkylamidoalkyl-*p*-benzoquinones**

Mariam al Azani, Mazen al Sulaibi, Thies Thiemann, Miguel Ángel Montiel, Carlos
Sánchez Sánchez and Jesus Iniesta

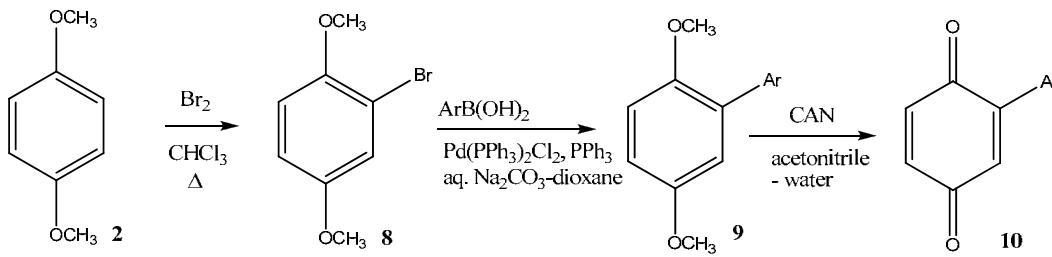
Corresponding authors: thiesthiemann@yahoo.de and jesus.iniesta@ua.es



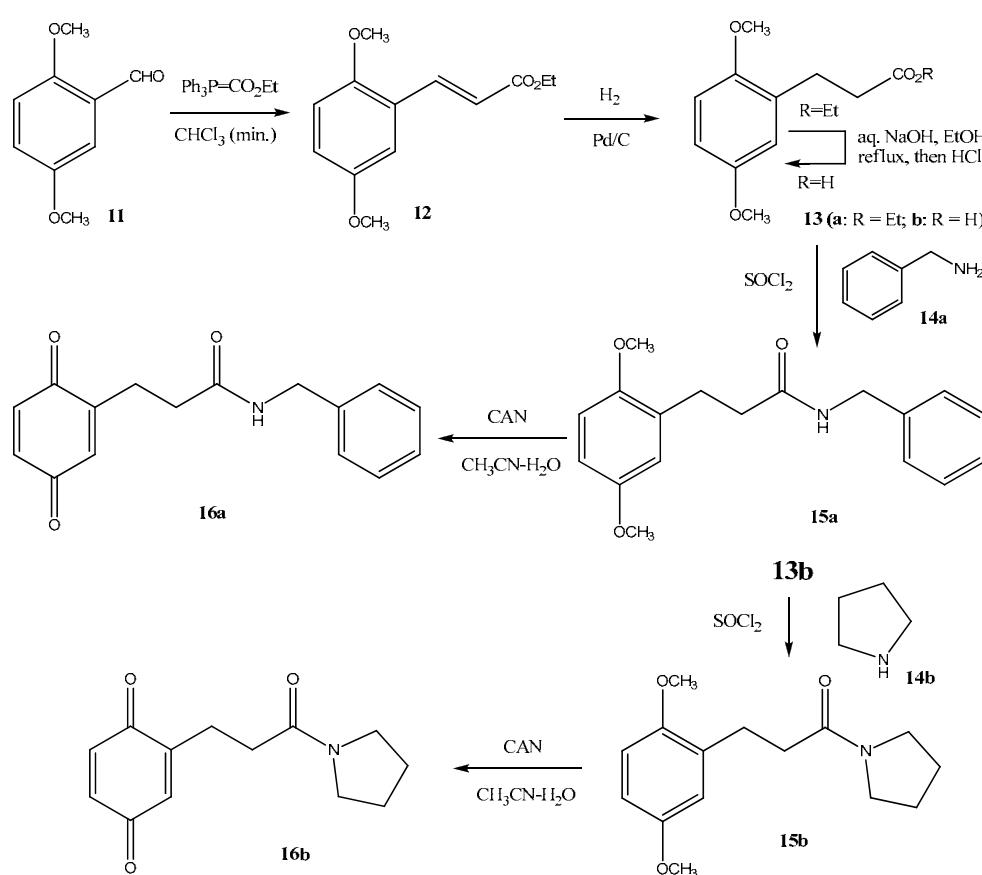
Scheme 1 Preparation of 2,5-*tert*-butyl-*p*-benzoquinone (**4**)



Scheme 2 Preparation of phenylnaphthoquinone **6**



Scheme 3 Preparation of 2-arylbenzo-*p*-quinones **10**



Scheme 4 Preparation of 2-amidoethyl-*p*-benzoquinones **16**

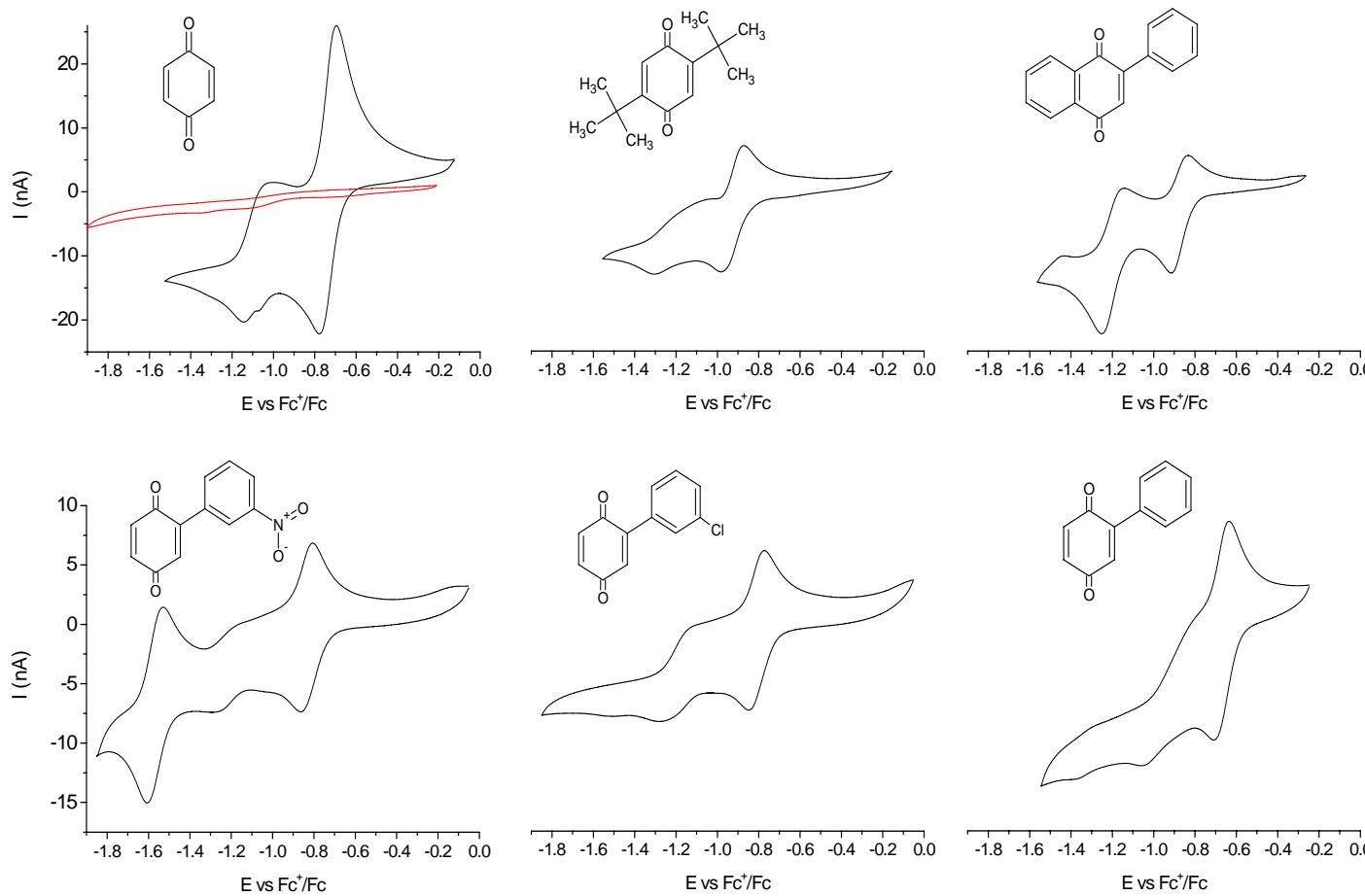


Figure 1. Cyclic voltammetry of 5 mM of benzoquinone (A), phenylbenzoquinone (B), 3-chlorophenylbenzoquinone (C), 3-nitrophenylbenzoquinone (D), 2,5-*tert*-Butyl-p-benzoquinone (E) and phenylnaphthoquinone (F) in the ionic liquid $[\text{bmim}]\text{[BF}_4^-$. 100 μm diameter Au microelectrode. Neat $[\text{bmim}]\text{[BF}_4^-$ (red dashed line). Scan rate 100 mV/s. Third scan recorded.

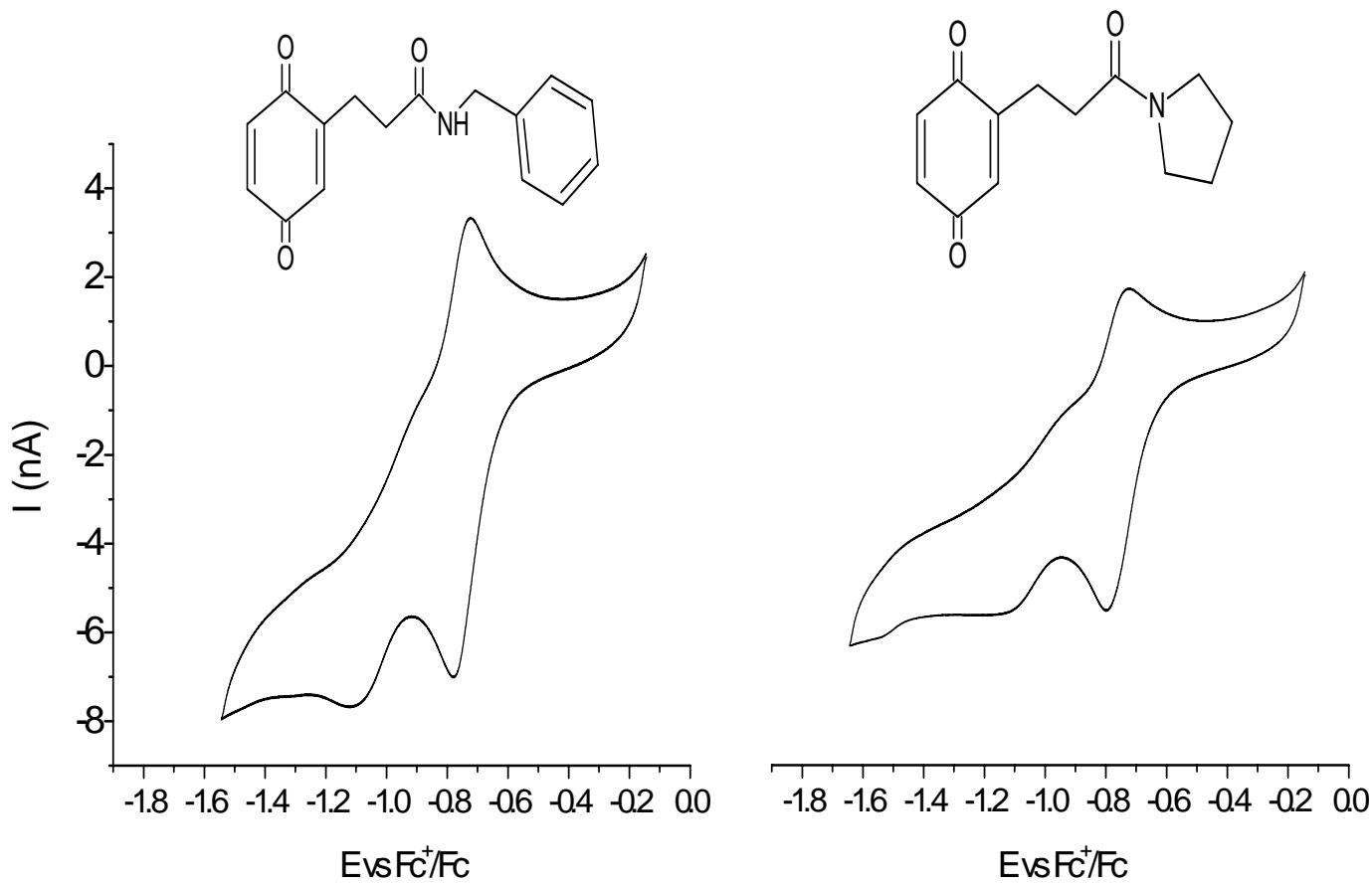


Figure 2. Cyclic voltammetry of 5 mM of *N*-Benzyl 3-(1,4-dimethoxyphen-2-yl)propionamide (**16a**) (A), and *N*-Benzyl carboxamidoethyl-*p*-benzoquinone (**16b**) (B) in the ionic liquid $[\text{bmim}][\text{BF}_4^-]$. 100 μm diameter Au microelectrode. Scan rate 100 mV/s. Third scan recorded.