

Electronic Supplementary Information for:

Use of new cross-linking method to obtain semi-IPN membranes with phosphonic acid groups for a PEMFC application

Etienne Labalme^a, Ghislain David^a, Julien Souquet^b, Pierrick Buvat^b, Janick Bigarre^b

Equipe Ingénierie et Architecture Macromoléculaires, Institut Charles Gerhardt UMR CNRS 5253, Ecole Nationale Supérieure de Chimie de Montpellier, 8 rue de l'Ecole Normale, 34296 Montpellier Cedex 5, France

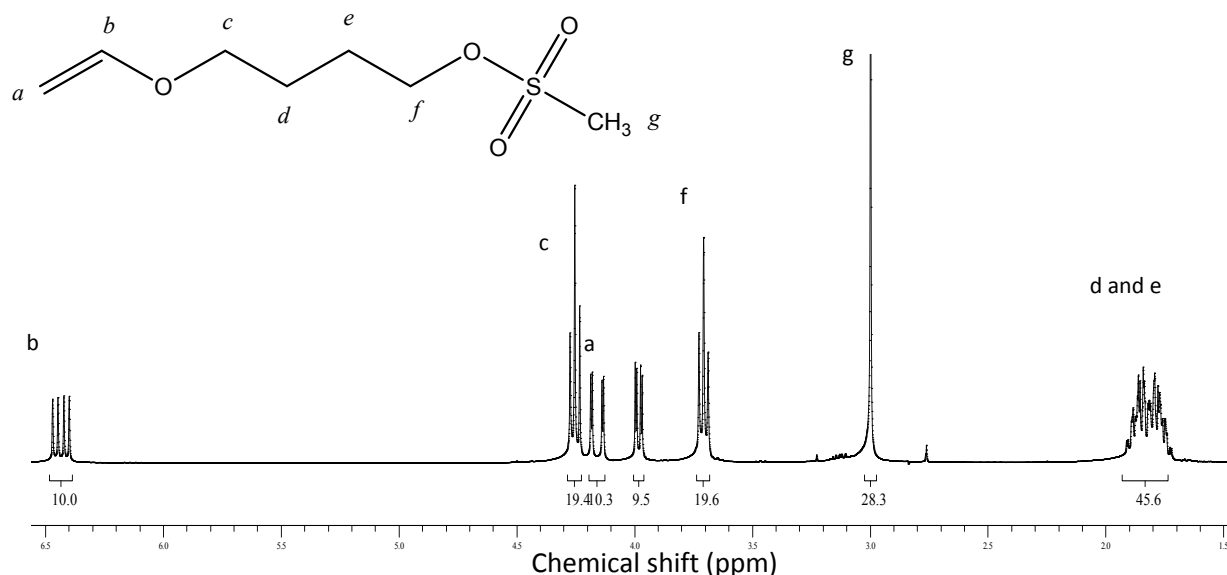


Figure S1 : ¹H NMR spectrum of BVEMs (realized in CDCl₃)

In Figure S1, after purification, the presence of the signal centered at 3ppm representative of the -CH₃ group (g) of the mesylate function can be observed. In addition, we observe a light shielding of the signal representative of the -CH₂- group (f) (from 3.8 to 3.7) in α position of the mesylate group due to the delocalization of the electron cloud of oxygen.

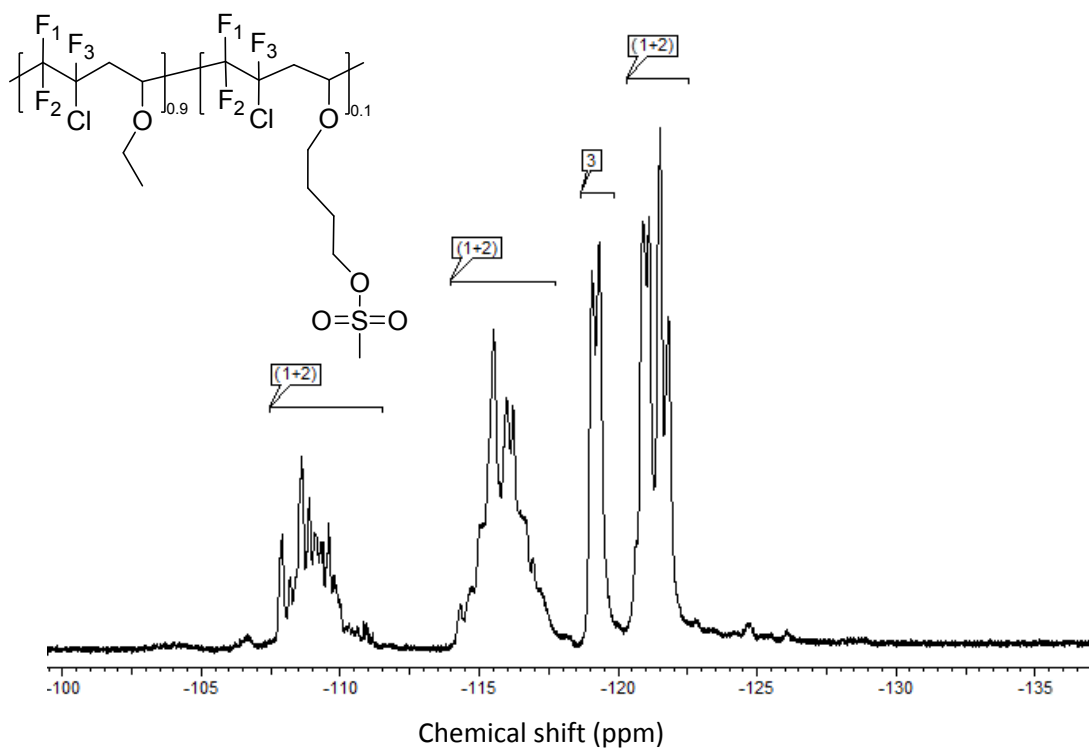


Figure S2: ^{19}F NMR Spectrum of poly[(CTFE-*alt*-EVE)_{0.9}-*co*-(CTFE-*alt*-VBMS)_{0.1}] in CDCl_3

The ^{19}F NMR (Figure S2) spectrum confirms the waiting structure. Indeed, we can observe the presence of the four signals significant of the fluorine atom of the main polymer chain. Concerning the $-\text{CF}_2-$ groups, the large signals range between -108 to -112, -115 to -119, and -120.7 to -123 ppm and for the $-\text{CFCl}-$ the signal is centered at -120 ppm.