Electronic Supplementary Material (ESI) for Journal of Materials Chemistry B. This journal is © The Royal Society of Chemistry 2017

#### **Supporting Informations**

Synthesis of  $\alpha$ -amino-lipophosphonates as cationic lipids or co-lipids for DNA transfection in dendritic cells

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Chantal Pichon<sup>2</sup>, Paul-Alain Jaffrès<sup>1</sup> and Patrick Midoux<sup>2</sup>

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Supplementary information (SI)

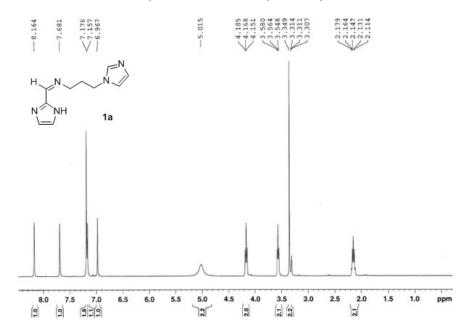
SI; S-1: <sup>1</sup>H, <sup>31</sup>P, & <sup>13</sup>C NMR spectra

SI; S-2: Evaluation of the stability of compounds 5a in acidic media

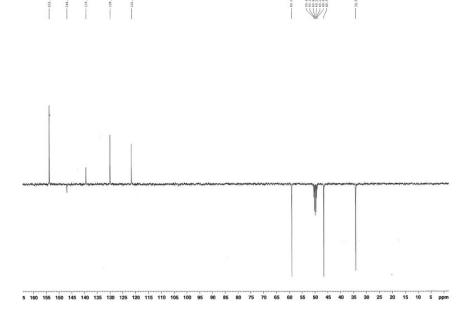
SI; S-3: Evaluation of the stability of compounds 5b in acidic media

SI; S-1: <sup>1</sup>H, <sup>31</sup>P, & <sup>13</sup>C NMR spectra of new compounds

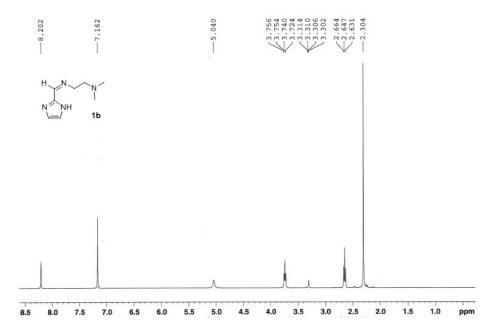
 $^{1}\text{H}$  NMR (400 MHz, CD $_{3}\text{OD})$  of compound 1a.



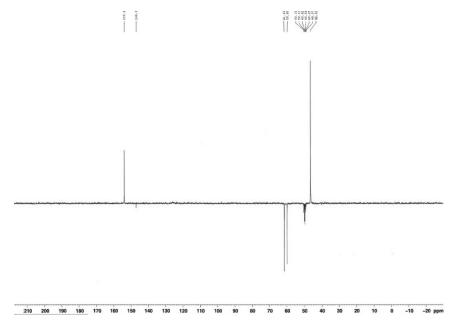
 $^{13}\text{C NMR}$  (125 MHz,  $\text{CD}_3\text{OD})$  of compound 1a.



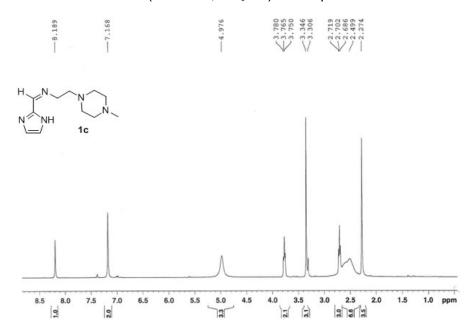
 $^{1}\text{H}$  NMR (400 MHz, CD<sub>3</sub>OD) of compound 1b.



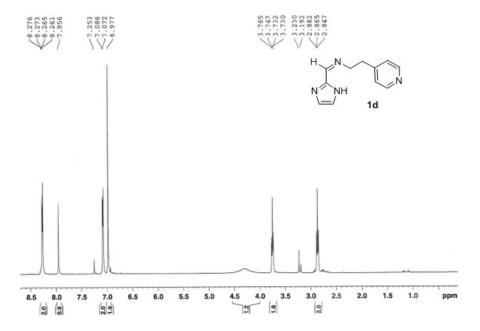
<sup>13</sup>C NMR (125 MHz, CD<sub>3</sub>OD) of compound **1b.** 



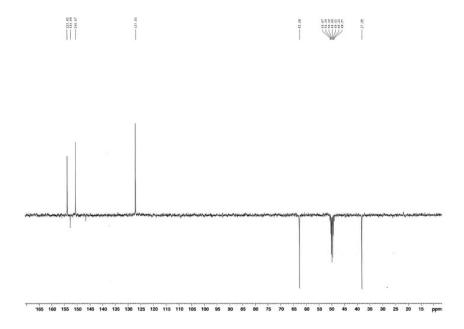
#### <sup>1</sup>H NMR (400 MHz, CD<sub>3</sub>OD) of compound 1c.



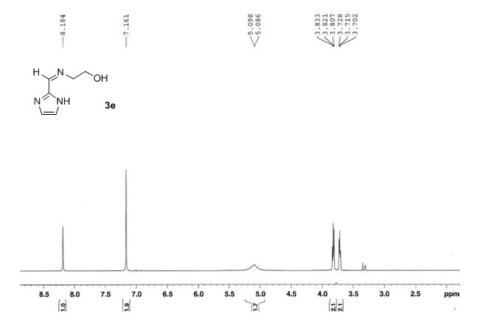
#### <sup>1</sup>H NMR (400 MHz, CD<sub>3</sub>OD) of compound 1d.



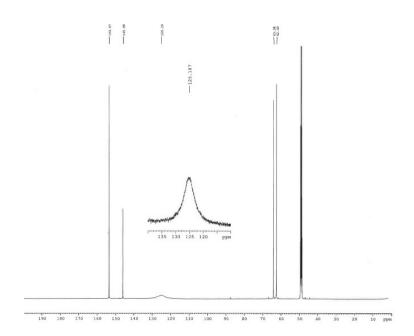
### $^{13}\text{C}$ NMR (125 MHz, $\text{CD}_3\text{OD})$ of compound 1d.



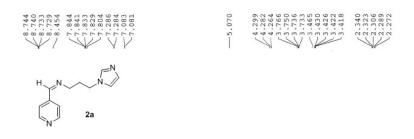
### $^{1}H$ NMR (400 MHz, CD<sub>3</sub>OD) of compound 1e.

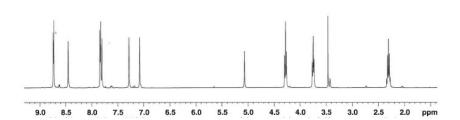


### $^{13}\text{C NMR}$ (125 MHz, CDCl<sub>3</sub>) of compound 3e.

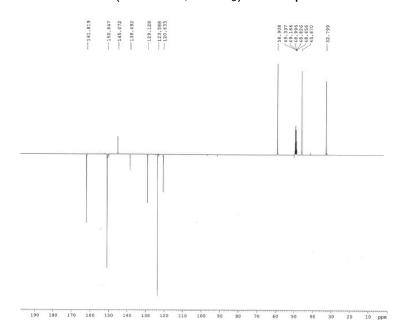


<sup>1</sup>H NMR (400 MHz, CD<sub>3</sub>OD) of compound 2a.



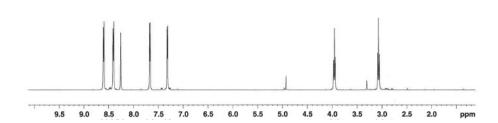


### $^{13}\text{C NMR}$ (125 MHz, CDCl $_3$ ) of compound 2a.

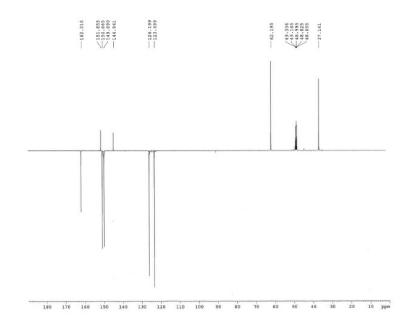


#### <sup>1</sup>H NMR (400 MHz, CD<sub>3</sub>OD) of compound **2b**.

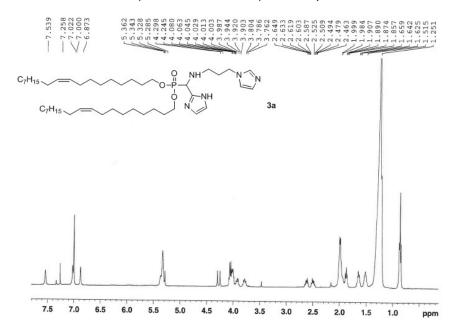




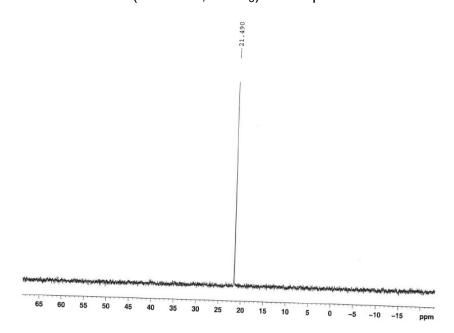
### $^{13}$ C NMR (125 MHz, CDCl<sub>3</sub>) of compound **2b.**



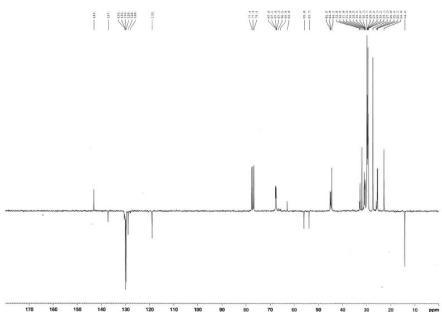
#### <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) of compound **3a.**



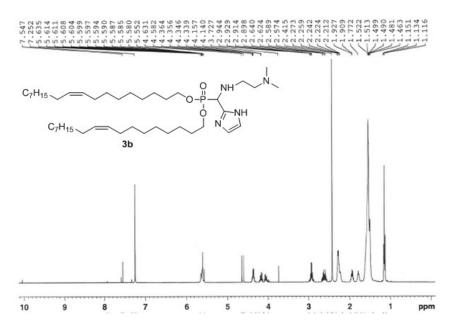
## $^{31}\textbf{P}$ NMR (162 MHz, CDCl $_{3})$ of compound 3a.



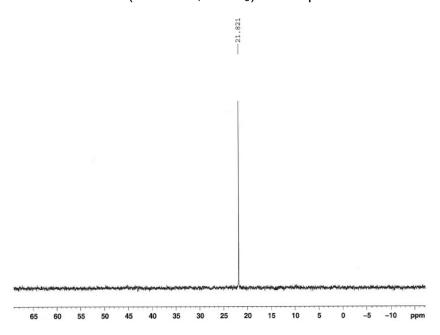
# $^{13}\text{C NMR}$ (125 MHz, CDCl3) of compound 3a.



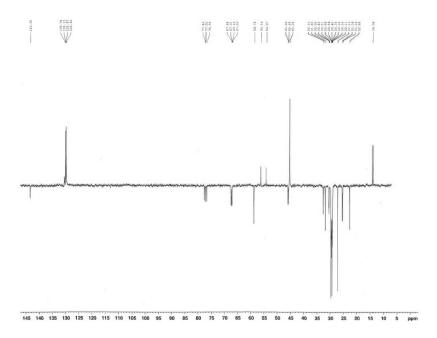
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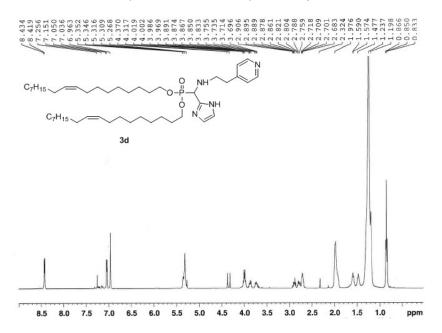
 $^{31}P$  NMR (162 MHz, CDCl<sub>3</sub>) of compound 3b.



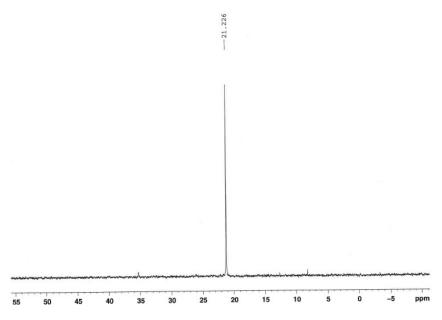
### $^{13}$ C NMR (125 MHz, CDCl<sub>3</sub>) of compound 3b.



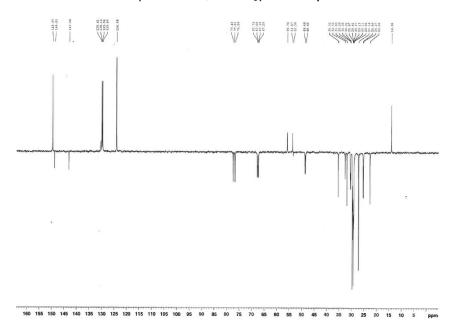
#### <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) of compound **3d.**



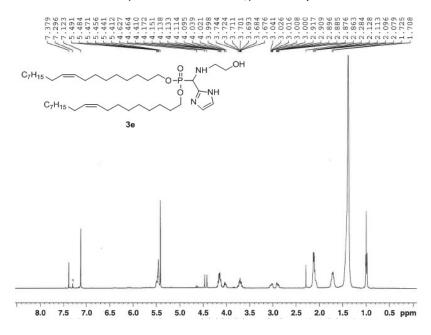
## $^{31}P$ NMR (162 MHz, CDCl<sub>3</sub>) of compound 3d.



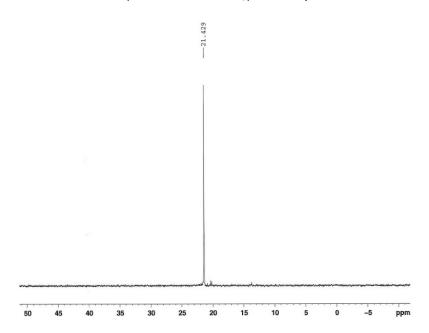
<sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) of compound **3d.** 



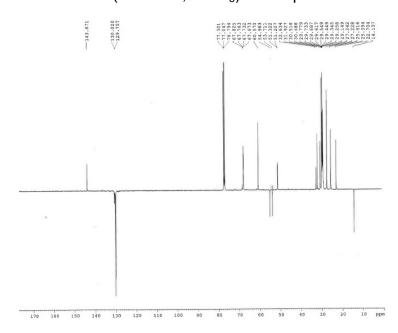
### $^{1}H$ NMR (400 MHz, CDCl<sub>3</sub>) of compound 3e.



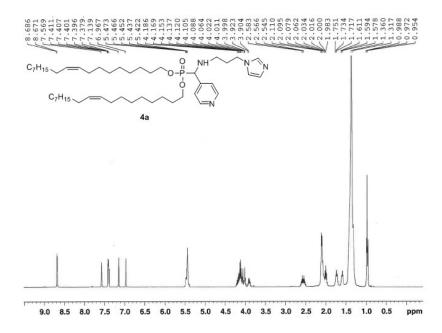
 $^{31}P$  NMR (162 MHz, CDCl<sub>3</sub>) of compound 3e.



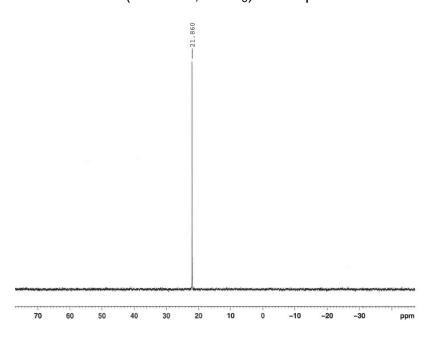
## $^{13}\text{C NMR}$ (125 MHz, CDCl $_3$ ) of compound 3e.



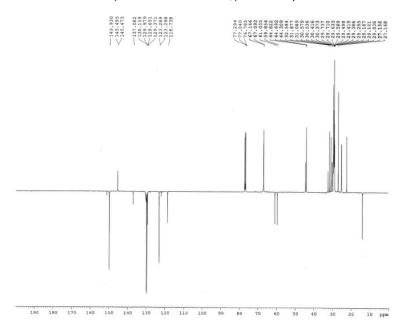
#### <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) of compound **4a.**



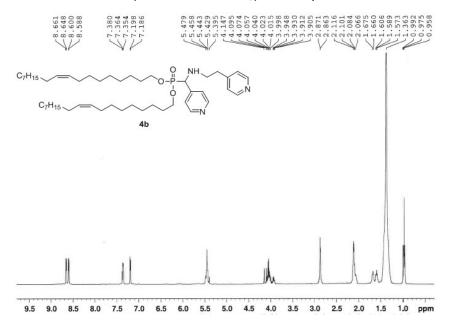
 $^{31}P$  NMR (162 MHz, CDCl<sub>3</sub>) of compound 4a.



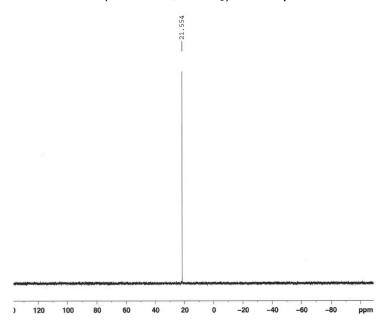
<sup>13</sup>C NMR (125 MHz, CDCl<sub>3</sub>) of compound **4a**.



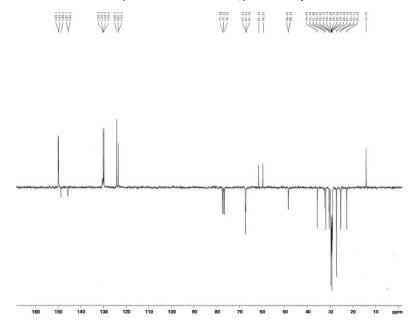
#### <sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) of compound 4b.



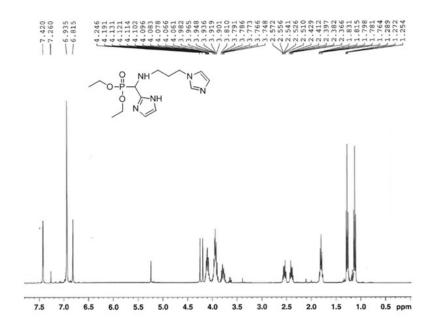
<sup>31</sup>P NMR (162 MHz, CDCl<sub>3</sub>) of compound 4b.



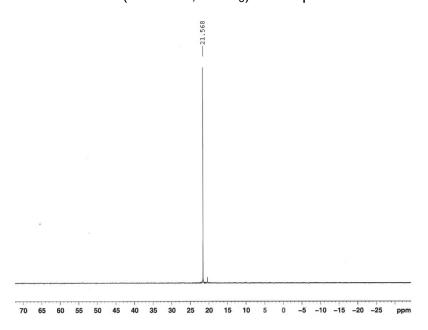
## $^{13}$ C NMR (125 MHz, CDCl<sub>3</sub>) of compound 4b.



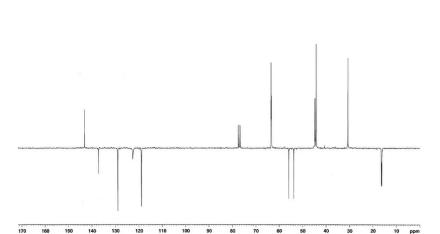
<sup>1</sup>H NMR (400 MHz, CDCl<sub>3</sub>) of compound 5a.



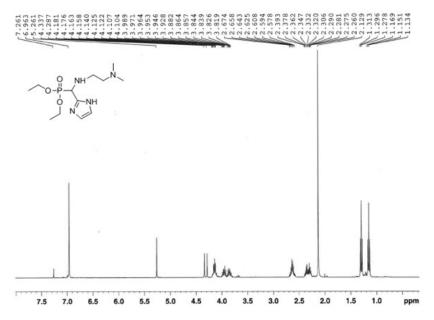
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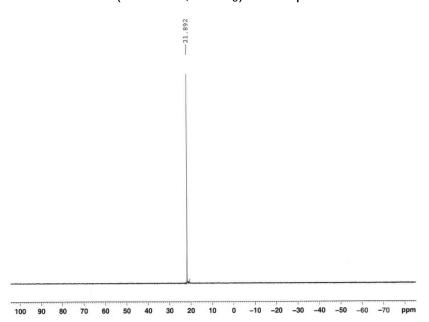
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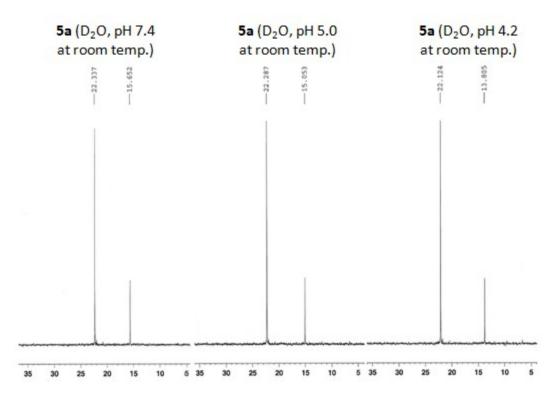


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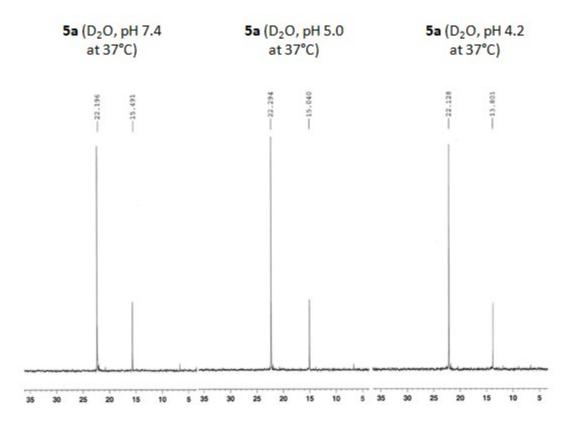


#### SI; S-2: Evaluation of the stability of compounds 5a in acidic media

1- At room temperature at t<sub>0</sub> (in acetate buffer, D<sub>2</sub>O as internal probe)

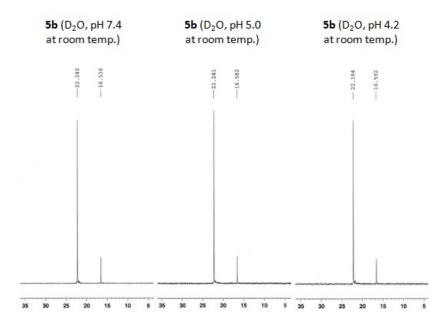


2- At room temperature after 5h at 37°C (in acetate buffer, D<sub>2</sub>O as internal probe)



#### SI; S-3: Evaluation of the stability of compounds 5b in acidic media

1- At room temperature at t<sub>0</sub> (in acetate buffer, D<sub>2</sub>O as internal probe)



2- At room temperature after 5h at 37°C (in acetate buffer, D<sub>2</sub>O as internal probe)

