Influence of ion size on the stability of the smectic phase of ionic liquid crystals

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General information: All reagents were purchased from commercial sources (Sigma-Aldrich, Merck and Lancaster) and were used without further purification. Solvents used as reaction media were purchased from local sources and were used after distillation. Reactions were monitored using commercially available, pre-coated thin-layer chromatography (TLC) plates (Merck, silica gel 60 F$_{254}$, 0.25 mm) and compounds were visualized under ultraviolet light (254 nm) and by staining with p-anisaldehyde or iodine. $^1$H and $^{13}$C NMR spectra were recorded on a Bruker Advance 200 and 400 instrument operating at 200 MHz ($^1$H), 400 MHz ($^1$H) and 400 MHz ($^{13}$C). Chemical shifts (δ) are quoted in ppm and referenced to internal TMS (δ 0.00 for $^1$H NMR), DMSO-d$_6$ (δ 2.50 for $^1$H NMR & 39.5 for $^{13}$C NMR) or CDCl$_3$ (δ 77.0 for $^{13}$C NMR); coupling constants (J) are quoted in Hz. Melting points were determined on a Buchi instrument and were uncorrected.

Experimental Section

Synthesis of 1-alkyl-3-methylpyridinium halides:

[$C_n$mpy]X ($n = 12, 14, 16; X = Cl, Br, I), 1-dodecyl-3-methylpyridinium halide, 1-tetradecyl-3-methylpyridinium halide, and 1-hexadecyl-3-methylpyridinium halide were synthesised as shown in Scheme 1.

\[
\text{Me} \quad + \quad RX \quad \xrightarrow{100 \, ^\circ \text{C}} \quad \text{Me} \\
R = \text{C}_{12}\text{H}_{25}, \text{C}_{14}\text{H}_{29}, \text{C}_{16}\text{H}_{33}, \quad X = \text{Cl, Br, I}
\]

Scheme 1

[$C_n$mPy]X were synthesized by the reaction of 0.1 mole equivalents of 3-methylpyridine with an excess of the appropriate haloalkane (1.1 mole equivalents) at 100 $^\circ$C for 24 hours. The excess haloalkane allows for the reaction to be stirred without additional solvents. The progress of the reaction was monitored by TLC. After completion of the reaction, the reaction mixture was evaporated under rotavapor to remove traces of starting material to obtain brownish viscous oil. To this was added 50 ml x 3 pet ether, stirred at room temperature for 1h and decanted to obtain ionic
liquid. This was further purified by adding 50 ml x 3 pet ether + ethyl acetate (1:0.5/1:1), stirred at room temperature for 1h and decanted to obtain pure ionic liquid.

**Synthesis of ionic liquid 1-alkyl-3-methylpyridinium tribromide**

\[ [C_{n}mPy]Br_3 \ (n = 12, 14 \& 16), 1\text{-dodecyl-3-methylpyridinium tribromide, 1-tetradecyl-3-methylpyridinium tribromide, and 1-hexadecyl-3-methylpyridinium tribromide were synthesized as shown in Scheme 2.}

\[
\begin{align*}
\text{N} & \quad \text{Me} \\
\text{R} & \quad \text{Br} \\
\uparrow & \\
\text{Br}_2 & \text{0}^0 \text{C-RT} & \text{Br}_3 \\
\end{align*}
\]

\( R = C_{12}H_{25}, C_{14}H_{29}, C_{16}H_{33} \)  

**Scheme 2**

In a fume cupboard, molecular bromine (0.038 mol) was added drop wise over 15 min to the corresponding 1-alkyl-3-methylpyridinium bromide \([C_{n}mPy]Br\) (0.045 mol) with stirring and cooling in a ice bath. After removal of ice bath, the reaction was continued under stirring at room temperature for 2h. The reaction was monitored by TLC. After completion of reaction, the reaction mixture was evaporated under reduced pressure over 5 h at 60 °C on rotavapor to afford a deep red liquid in quantitative yields (95-98%).

**Synthesis of ionic liquid 1-alkyl-3-methylpyridinium triiodide**

\[ [C_{n}mPy]I_3 \ (n = 12, 14, 16), 1\text{-dodecyl-3-methylpyridinium triiodide, 1-tetradecyl-3-methylpyridinium triiodide, and 1-hexadecyl-3-methylpyridinium triiodide were synthesized as shown in Scheme 3.}

\[
\begin{align*}
\text{N} & \quad \text{Me} \\
\text{R} & \quad \text{I} \\
\uparrow & \\
\text{I}_2/\text{CHCl}_3 & \text{0}^0 \text{C-RT} & \text{I}_3 \\
\end{align*}
\]

\( R = C_{12}H_{25}, C_{14}H_{29}, C_{16}H_{33} \)  

**Scheme 3**

Molecular iodine (0.072 mol) was added drop wise over 15 min to the corresponding 1-alkyl-3-methylpyridinium iodide \([C_{n}mPy]I\) (0.036 mol), dissolved in 50ml
chloroform, with stirring and cooling in an ice bath. The reaction was continued for 2h in an ice bath and then left to attain room temperature and continued at room temperature for additional 10h. The reaction was monitored by TLC. After completion of reaction, the reaction mixture was evaporated under reduced pressure over 5 h at 60 °C on rotavapor to afford a deep red liquid. This was further purified by giving wash with pet ether, followed by (50ml x 3) pet ether + ethyl acetate (90:10) mixture to afford red oil in quantitative yields (95-98%).

**Synthesis of ionic liquid 1-alkyl-3-methylpyridinium dichloroiodate**

![Scheme 4](image)

[R = C12H25, C14H29, C16H33]

[CnmPy]X ($n = 12, 14, 16; X = ICl_2$), 1-dodecyl-3-methylpyridinium dichloroiodate, 1-tetradecyl-3-methylpyridinium dichloroiodate, and 1-hexadecyl-3-methylpyridinium dichloroiodate were synthesized as shown in Scheme 4.

A black solution of ICl (19.39 mmol) in dichloromethane (35 ml), was added drop wise to an ice cold solution of 1-alkyl-3-methylpyridinium chloride (16.16 mmol) in water (16 ml) under stirring and then left to attain room temperature. After the reaction mixture was stirred for 1 hour at room temperature, the dichloromethane layer was separated and dried with sodium sulphate and then evaporated under vacuum to afford dark reddish brown ionic liquid 1-alkyl-3-methylpyridinium dichloroiodate in quantitative yields (95-98%). This ionic liquid was stable and stored in dark at 10 °C (in refrigerator) for several months without any change in color, loss of reactivity and degradation (checked by $^1$H NMR).

The NMR ($^1$H and $^{13}$C) data of compounds(a–p) are given as:
1. 1-dodecyl-3-methylpyridinium chloride\(^1\) (a): Dark red viscous semi-solid.

\[ \text{H NMR (200 MHz, CDCl}_3 \delta/\text{ppm): 0.82 (t, 3H, J=6.19 Hz), 1.21 (m, 19H), 2.03 (m, 3H), 2.62 (s, 3H), 4.89 (t, 2H, J=5.56 Hz), 8.01 (s, IH), 8.23 (d, IH, J=7.7 Hz), 9.20 (s, 2H).} \]

\[ \text{C NMR (125 MHz, CDCl}_3, \delta/\text{ppm): 14.01, 18.62, 22.55, 26.01, 29.03, 29.20, 29.28, 29.43, 29.48, 31.77, 31.80, 61.82, 127.86, 139.45, 142.32, 144.27, 145.59.} \]

2. 1-dodecyl-3-methylpyridinium bromide\(^2\) (b): Dark red viscous semi-solid.

\[ \text{H NMR (200 MHz, CDCl}_3 \delta/\text{ppm): 0.82 (t, 3H, 5.75 Hz), 1.22 (m, 18H), 2.16-1.96 (m, 2H), 2.64 (s, 3H), 4.96 (t, 2H, 6.48 Hz), 7.96 (t, 1H, 7.71 Hz), 8.22 (d, IH, 7.96 Hz), 9.19 (d, IH, 5.83 Hz), 9.33 (s, 1H).} \]

\[ \text{C NMR (125 MHz, CDCl}_3, \delta/\text{ppm): 14.01, 18.63, 22.55, 25.98, 28.97, 29.20, 29.25, 29.40, 29.46, 31.77, 31.85, 61.79, 127.77, 139.52, 142.18, 144.38, 145.52.} \]

3. 1-dodecyl-3-methylpyridinium iodide\(^3\) (c): Dark red liquid. (Give integration in NMR)

\[ \text{H NMR (200 MHz, CDCl}_3 \delta/\text{ppm): 0.82 (t, 3H, 5.75 Hz), 1.22 (m, 18H), 2.16-1.96 (sextet, 2H), 2.64 (s, 3H), 4.82 (t, 2H, 6.07 Hz), 7.96 (t, 1H, 7.71 s Hz), 8.26 (d, IH, 7.96 Hz), 9.08 (d, IH, 5.83 Hz), 9.28 (s, 1H).} \]

\[ \text{C NMR (125 MHz, CDCl}_3, \delta/\text{ppm): 13.98, 18.67, 22.52, 25.90, 28.92, 29.17, 29.21, 29.37, 29.43, 31.71, 31.73, 61.71, 127.83, 139.57, 141.83, 144.15, 145.83.} \]

4. 1-tetradecyl-3-methylpyridinium chloride\(^4\) (d): Orange semi-solid.

\[ \text{H NMR (200 MHz, CDCl}_3 \delta/\text{ppm): 0.82 (t, 3H, J=6.06 Hz), 1.22 (s, 18H), 2.16-1.96 (sextet, 2H), 2.29 (2H), 2.62 (s, 3H), 4.90 (t, 2H, J=7.33 Hz), 7.96 (t, 1H, J=6.07 Hz), 8.24 (d, IH, J=7.83 Hz), 9.30 (d, IH, J=5.69 Hz), 9.40 (s, 1H).} \]

\[ \text{C NMR (125 MHz, CDCl}_3, \delta/\text{ppm): 145.29, 144.58, 142.39, 139.43, 127.74, 61.67, 31.88, 31.72, 29.49, 29.45, 29.42, 29.36, 29.20, 29.17, 145.29, 144.58, 142.39, 139.43, 127.74, 61.67, 31.88, 31.72, 29.49, 29.45, 29.42, 29.36, 29.20, 29.17.} \]
28.95, 25.96, 22.49, 18.56, 13.95.

5. 1-tetradecyl-3-methylpyridinium bromide² (e): White solid, M.P. 78-79 °C.

\[
\text{H NMR (200 MHz, CDCl}_3\text{, }\delta/\text{ppm): 0.86 (t, 3H, J=6.07 Hz), 1.22 (s, 22H), 1.99 (t, 2H), 2.64 (s, 3H), 4.89 (t, 2H, J=7.45 Hz), 7.96 (t, 1H, J=6.19), 8.23 (d, IH, J=7.96 Hz), 9.22 (d, IH, J=5.81 Hz), 9.36 (s, 1H).}
\]

\[
\text{C NMR (125 MHz, CDCl}_3\text{): 14.03, 18.65, 22.58, 26.00, 29.00, 29.25, 29.27, 29.43, 29.50, 29.54, 29.57, 31.81, 31.88, 61.80, 127.75, 139.53, 142.18, 144.40, 145.53.}
\]

6. 1-tetradecyl-3-methylpyridinium iodide (f): Dark brown solid, M.P. 44-45 °C.

\[
\text{H NMR (200 MHz, CDCl}_3\text{, }\delta/\text{ppm): 0.86 (t, 3H, J=6.06 Hz), 1.23 (m, 22H), 1.99 (t, 2H), 2.03 (sextet, 2H), 2.65 (s, 3H), 4.87 (t, 2H, J=7.45 Hz), 7.96 (dd, 1H, J=6.06 Hz), 8.23 (d, IH, J=7.96 Hz), 9.07 (d, IH, J=6.06 Hz), 9.25 (s, 1H).}
\]

\[
\text{C NMR (125 MHz, CDCl}_3\text{, }\delta/\text{ppm): 13.97, 18.64, 22.51, 25.89, 28.91, 29.17, 29.20, 29.37, 29.44, 29.47, 29.50, 31.71, 31.73, 61.71, 127.80, 139.53, 141.82, 144.14, 145.80.}
\]

7. 1-hexadecyl-3-methylpyridinium chloride⁴ (g): White creamish solid, M.P. 68-69 °C.

\[
\text{H NMR (200 MHz, CDCl}_3\text{, }\delta/\text{ppm): 0.88 (t, 3H, J=6.69 Hz), 1.22 (m, 26H), 1.98 (m, 2H), 2.62 (s, 2H), 4.91 (t, 2H, J=7.20 Hz), 7.96 (t, 1H), 8.23 (d, IH, J=7.58 Hz), 9.35 (d, IH, J=5.43 Hz), 9.42 (s, 1H).}
\]

\[
\text{C NMR (125 MHz, CDCl}_3\text{, }\delta/\text{ppm): 14.05, 18.67, 22.61, 26.06, 29.04, 29.29, 29.46, 29.54, 29.58, 29.62, 31.84, 31.93, 61.89, 127.75, 139.52, 142.45, 144.49, 146.35.}
\]

8. 1-hexadecyl-3-methylpyridinium bromide⁵ (h): White solid, M.P. 49-50 °C.

\[
\text{H NMR (200 MHz, CDCl}_3\text{, }\delta/\text{ppm): 0.85 (t, 3H, J=5.94 Hz), 1.23 (m, 27H), 1.99 (t, 2H), 1.90 (m, 4H), 2.65 (s, 3H), 4.93 (t, 2H, J=7.33 Hz), 7.96 (t, 1H, J=6.19 Hz), 8.22 (d, IH, J=8.02 Hz), 9.22 (d, IH, J=6.06 Hz), 9.38 (s, 1H).}
\]

\[
\text{C NMR (125 MHz, CDCl}_3\text{, }\delta/\text{ppm): 14.03, 18.65, 22.60, 26.01, 26.04, 29.04, 29.29, 29.46, 29.54, 29.58, 29.62, 31.84, 31.93, 61.89, 127.75, 139.52, 142.45, 144.49, 146.35.}
\]
9. 1-hexadecyl-3-methylpyridinium iodide\textsuperscript{(i)}: Pale yellow solid, M.P. 58-59 °C.

\[ \text{H NMR (200 MHz, CDCl}_3 \text{ δ/ppm): 0.83 (t, 3H, J=6.69 Hz), 1.23 (m, 26H), 1.81 (m, 2H), 2.03 (m, 2H), 2.65 (s, 3H), 4.91 (t, 2H, J=7.33 Hz), 8.04 (t, 1H, J=6.57 Hz), 8.25 (d, 1H, J=7.96 Hz), 9.07 (d, 1H, J=5.94), 9.26 (s, 1H), Hz).} \]

\[ \text{C NMR (125 MHz, CDCl}_3 \text{ δ/ppm): 14.05, 18.71, 22.61, 25.97, 28.99, 29.28, 29.45, 29.54, 29.57, 29.61, 31.78, 31.83, 62.45, 127.77, 139.59, 141.87, 144.28, 145.80.} \]

10. 1-dodecyl-3-methylpyridinium tribromide (j): Dark red liquid.

\[ \text{H NMR (200 MHz, CDCl}_3 \text{ δ/ppm): 0.88 (s, 2H), 1.26 (m, 18H), 2.15 (m, 2H), 2.76 (s, 3H), 4.78 (s, 2H), 8.04 (s, 1H), 8.35 (s, 1H), 8.80 (s, 2H).} \]

\[ \text{C NMR (125 MHz, CDCl}_3 \text{ δ/ppm): 14.01, 20.69, 22.51, 26.67, 29.18, 29.28, 29.43, 29.52, 31.73, 32.48, 34.08, 64.19, 129.26, 140.02, 143.20, 145.75, 147.07.} \]

11. 1-tetradecyl-3-methylpyridinium tribromide (k): Red semi-solid.

\[ \text{H NMR (200 MHz, CDCl}_3 \text{ δ/ppm): 0.85 (t, 3H, J=6.87 Hz), 1.22 (m, 19H), 1.36, (4H, m), 2.06 (quintet, 2H), 2.57 (s, 3H), 4.74 (t, 2H, J=7.63 Hz), 8.03 (t, 1H), 8.31 (d, 1H, J=6.10 Hz), 8.82 (d, 1H, J=6.34 Hz), 8.85 (s, 1H).} \]

\[ \text{C NMR (125 MHz, CDCl}_3 \text{ δ/ppm): 14.03, 19.07, 22.58, 26.16, 26.96, 29.25, 29.28, 29.31, 29.52, 29.55, 29.58, 31.71, 31.81, 62.64, 128.10, 140.22, 141.69, 143.75, 145.08.} \]

12. 1-hexadecyl-3-methylpyridinium tribromide (k): White solid, M.P. 58-59 °C.

\[ \text{H NMR (200 MHz, CDCl}_3 \text{ δ/ppm): 0.89 (t, 3H). 1.26 (s, 26H). 2.20} \]
(s, 2H), 2.79 (s, 3H). 4.83 (s, 2H). 8.04 (s, 1H). 8.36 (s, 1H), 8.89 (s, 2H). 13C NMR (125 MHz, CDCl$_3$ δ/ppm): 14.03, 18.65, 22.60, 26.01, 29.01, 29.27, 29.44, 29.52, 29.57, 29.60, 31.83, 31.88, 61.83, 127.73, 139.54, 142.19, 144.42, 145.52.

13. 1-dodecyl-3-methylpyridinium triiodide (l): Dark red liquid.

1H NMR (200 MHz, CDCl$_3$ δ/ppm): 0.87 (t, 3H, J=6.19), 1.25 (s, 18H), 2.15 (sextet, 2H), 2.72 (s, 3H), 4.73 (t, 2H, J=7.58), 8.09 (t, 1H, J=6.82), 8.39 (d, 1H, J=8.21), 8.57 (s, 1H), 8.64 (s, 1H). 13C NMR (125 MHz, CDCl$_3$ δ/ppm): 14.14, 19.68, 22.66, 26.34, 29.01, 29.32, 29.49, 29.58, 31.68, 31.70, 31.87, 63.32, 128.53, 140.69, 141.63, 143.88, 146.60.

14. 1-tetradecyl-3-methylpyridinium triiodide (m): Dark red liquid.

1H NMR (200 MHz, CDCl$_3$ δ/ppm): 0.84 (t, 3H, J=6.57), 1.25 (s, 22H), 2.17 (m, 2H), 2.72 (s, 3H), 4.74 (t, 2H, J=7.71), 8.09 (t, 1H, J=6.57), 8.39 (d, 1H, J=7.63), 8.67 (d, 2H). 13C NMR (125 MHz, CDCl$_3$ δ/ppm): 14.12, 19.72, 22.64, 26.36, 29.02, 29.32, 29.36, 29.51, 29.61, 29.65, 31.71, 31.86, 63.28, 128.57, 140.66, 141.63, 143.87, 146.64.

15. 1-hexadecyl-3-methylpyridinium triiodide (m): Dark brown solid, M.P. 40-41°C.

1H NMR (200 MHz, CDCl$_3$ δ/ppm): 0.87 (s, 3H), 1.25 (s, 26H), 2.12 (s, 3H), 2.73 (s, 3H), 4.64 (t, 3H), 8.03 (t, H), 8.39 (d, 1H), 8.86 (s, 2H). 13C NMR (125 MHz, CDCl$_3$ δ/ppm): 14.11, 19.64, 22.66, 26.35, 29.01, 29.34, 29.51, 29.60, 29.65, 29.68, 31.68, 31.89, 63.34, 128.52, 140.71, 141.62, 143.88, 146.58.

16. 1-dodecyl-3-methylpyridinium dichloroiodate (n): Dark red liquid.

1H NMR (200 MHz, CDCl$_3$ δ/ppm): 0.87 (t, 3H), 1.26 (s, 18H), 2.17-2.04 (m, 2H), 2.74 (s, 3H), 4.67 (s, 2H), 8.06 (s, 1H), 8.38 (s, 1H), 8.65 (s, 2H). 13C NMR (125 MHz, CDCl$_3$, δ/ppm): 14.07,
19.51, 22.61, 26.34, 29.00, 29.25, 29.31, 29.45, 29.53, 31.82, 31.90, 63.35, 128.51, 140.65, 141.78, 144.27, 146.51.

17. 1-tetradecyl-3-methylpyridinium dichloroiodate (o): Dark red liquid.

\[
\begin{align*}
\text{H NMR} \ (200 \text{ MHz, CDCl}_3 \ \delta/\text{ppm}) & : 0.87 \ (t, \ 3\ H, J=6.06 \ Hz), \ 1.25 \ (m,\ 24H), \ 2.09 \ (t, \ 2H), \ 2.72 \ (s, \ 3H), \ 4.67 \ (t, \ 2H, J=7.20 \ Hz), \ 8.04 \ (t, J=6.32 \ Hz, \ 1H), \ 8.34 \ (d, \ IH, J=7.83 \ Hz), \ 8.70 \ (t, \ 2H). \\
\text{C NMR} \ (125 \text{ MHz, CDCl}_3, \ \delta/\text{ppm}) & : 14.04, \ 19.19, \ 22.58, \ 26.23, \ 28.95, \ 29.25, \ 29.28, \ 29.43, \ 29.55, \ 29.58, \ 31.74, \ 31.81, \ 62.89, \ 128.26, \ 140.55, \ 141.49, \ 143.90, \ 146.31.
\end{align*}
\]

15. 1-hexadecyl-3-methylpyridinium dichloroiodate (p): Pale yellow solid, M.P. 58-59 °C.

\[
\begin{align*}
\text{H NMR} \ (200 \text{ MHz, CDCl}_3 \ \delta/\text{ppm}) & : 0.83 \ (t, \ 3\ H, J=6.87 \ Hz), \ 1.19 \ (m, \ 24 \ H), \ 1.99 \ (\text{quintet, } 2H), \ 2.20 \ (m, \ 3H), \ 2.62 \ (s, \ 3H), \ 4.88 \ (t, \ 2H, J=7.33 \ Hz), \ 8.01 \ (\text{dd, } J=7.78 \ Hz, \ 1H), \ 8.23 \ (d, \ IH, J=8.24 \ Hz), \ 9.19 \ (d, \ IH, J=5.95 \ Hz), \ 9.34 \ (s, \ 1H). \\
\text{C NMR} \ (125 \text{ MHz, CDCl}_3, \ \delta/\text{ppm}) & : 14.05, \ 20.03, \ 22.60, \ 26.54, \ 29.12, \ 29.27, \ 29.37, \ 29.50, \ 29.57, \ 29.62, \ 31.83, \ 32.23, \ 64.13, \ 128.84, \ 140.67, \ 142.51, \ 145.12, \ 146.69.
\end{align*}
\]
1. 1-Dodecyl-3-methylpyridinium chloride (a)

![1H NMR of 1-dodecyl-3-methylpyridinium chloride](image)

1H NMR of 1-dodecyl-3-methylpyridinium chloride

![13C NMR of 1-dodecyl-3-methylpyridinium chloride](image)

13C NMR of 1-dodecyl-3-methylpyridinium chloride

2. 1-dodecyl-3-methylpyridinium bromide (b)
3. 1-dodecyl-3-methylpyridinium iodide (c)

\[ \text{H NMR of 1-dodecyl-3-methylpyridinium bromide} \]

\[ \text{13C NMR of 1-dodecyl-3-methylpyridinium bromide} \]
4. 1-tetradecyl-3-methylpyridinium chloride (d)

$^1$H NMR of 1-dodecyl-3-methylpyridinium iodide

$^{13}$C NMR of 1-dodecyl-3-methylpyridinium iodide
5. 1-tetradecyl-3-methylpyridinium bromide (e)

1H NMR of 1-tetradecyl-3-methylpyridinium chloride

13C NMR of 1-tetradecyl-3-methylpyridinium chloride
6. 1-tetradecyl-3-methylpyridinium iodide (f)

1H NMR of 1-tetradecyl-3-methylpyridinium bromide

13C NMR of 1-tetradecyl-3-methylpyridinium bromide
7. 1-hexadecyl-3-methylpyridinium chloride (g)
8. 1-hexadecyl-3-methylpyridinium bromide (h)
9. 1-hexadecyl-3-methylpyridinium iodide (i)
1H NMR of 1-hexadecyl-3-methylpyridinium iodide

13C NMR of 1-hexadecyl-3-methylpyridinium iodide

10. 1-dodecyl-3-methylpyridinium tribromide (j)
11. 1-tetradecyl-3-methylpyridinium tribromide (k)
1H NMR of 1-tetradecyl-3-methylpyridinium tribromide

13C NMR of 1-tetradecyl-3-methylpyridinium tribromide

12. 1-hexadecyl-3-methylpyridinium tribromide (k)
1H NMR of 1-hexadecyl-3-methylpyridinium tribromide

13C NMR of 1-hexadecyl-3-methylpyridinium tribromide

13. 1-dodecyl-3-methylpyridinium triiodide (I)
14. 1-tetradecyl-3-methylpyridinium triiodide (m)
15. 1-hexadecyl-3-methylpyridinium triiodide (m)
$^1$H NMR of 1-hexadecyl-3-methylpyridinium triiodide

$^{13}$C NMR of 1-hexadecyl-3-methylpyridinium triiodide
17. 1-tetradecyl-3-methylpyridinium dichloroiodate (o)

$^1$H NMR of 1-tetradecyl-3-methylpyridinium dichloroiodate

$^{13}$C NMR of 1-tetradecyl-3-methylpyridinium dichloroiodate

15. 1-hexadecyl-3-methylpyridinium dichloroiodate (p)
$^{1}H$ NMR of 1-hexadecyl-3-methylpyridinium dichloroiodate

$^{13}C$ NMR of 1-hexadecyl-3-methylpyridinium dichloroiodate

References:


