

Inorg. Chem., 1997, 36(14), 2992-3000, DOI:10.1021/ic960794b

Terms & Conditions

Electronic Supporting Information files are available without a subscription to ACS Web Editions. The American Chemical Society holds a copyright ownership interest in any copyrightable Supporting Information. Files available from the ACS website may be downloaded for personal use only. Users are not otherwise permitted to reproduce, republish, redistribute, or sell any Supporting Information from the ACS website, either in whole or in part, in either machine-readable form or any other form without permission from the American Chemical Society. For permission to reproduce, republish and redistribute this material, requesters must process their own requests via the RightsLink permission system. Information about how to use the RightsLink permission system can be found at http://pubs.acs.org/page/copyright/permission.html



Copyright © 1997 American Chemical Society

Fig.1 pH dependence of the ¹H-NMR resonances of ligand PCTA-[14] recorded at 400 MHz and 25°C.

Fig.2 pH dependence of the ¹H-NMR resonances of ligand PCTA-[13] recorded at 400 MHz and 25° C.

Fig.3 ¹H NMR spectra of **a**) LaPCTA-[14], **b**) LaPCTA-[13], **c**) LaPCTA-[12], recorded at 400 MHz and 50°C, in D₂O at pD = 7. The label w indicates HDO signal, whereas the asterisk indicates CH₃OH signal present as a impurity. *tert*-butyl alcohol was added as an internal reference ($\delta = 1.29$ ppm).

Fig.4 ¹H NMR spectra of a) LuPCTA-[14], b) LuPCTA-[13], c) LuPCTA-[12] recorded at 400 MHz and 50°C. in D₂O at pD = 7. The label w indicates HDO. *tert*-butyl alcohol was added as an internal reference (δ = 1.29 ppm).

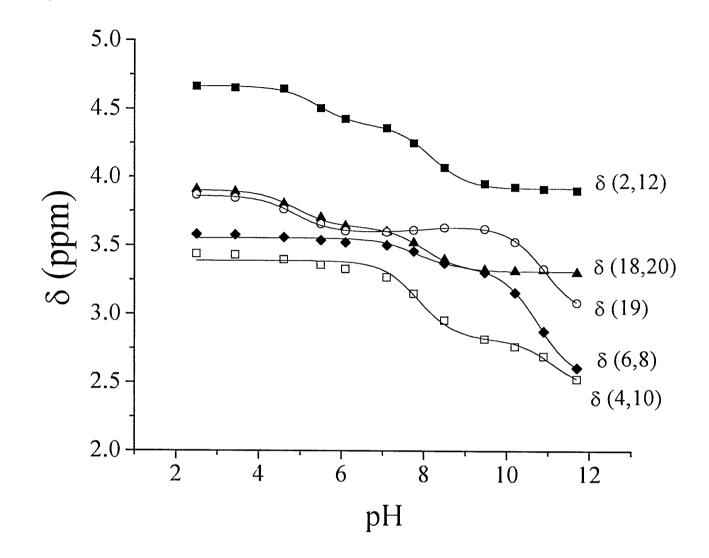
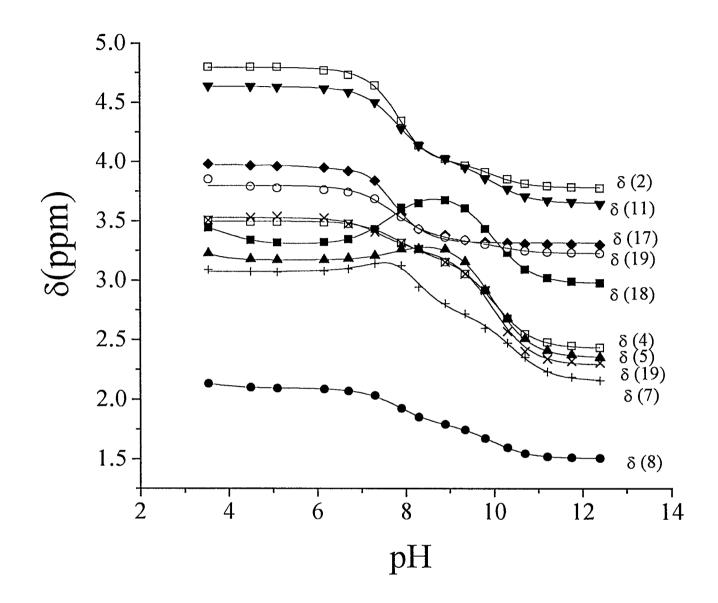


Figure 1



SUPPL. MAT

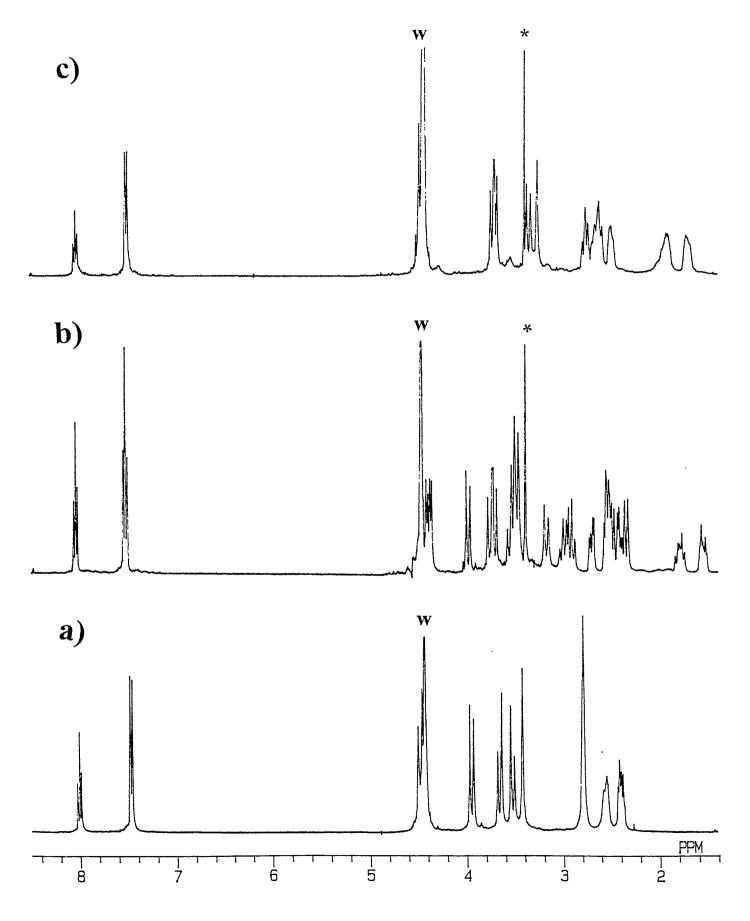


Figure 3

_

SUPPL. MAT

