

**Problem R-12N** (C<sub>22</sub>H<sub>28</sub>ClP<sub>4</sub>Pd). Analyze the <sup>1</sup>H decoupled 32.4 MHz <sup>31</sup>P NMR spectra of a palladium-phosphine complex shown on the next page (Bartsch, R.; Carmichael, D.; Hitchcock, P. B.; Meidine, M. F.; Nixon, J. F.; Sillett, G. J. D. *J. Chem. Soc., Chem. Commun.* **1988**, 1615).

(a) Identify all signals in the low temperature spectrum (-75 °C), and report approximate coupling constants using the form:  $\delta$  \_\_\_\_\_,  $^{\times}J_{1-2}$  =\_\_\_\_ Hz. Use the numberings shown on the structure. For each signal briefly give your reasoning for the assignment.

	$Me_3C \sim P^2 \qquad P^4Et_3$
	$P^1 - Pd - Cl$
	$P^3$ $\downarrow$
p1	Me <sub>3</sub> C
'	$C_{22}H_{28}CIP_4Pd$

$P^2$			

(b) Identify the process which is responsible for the changes in the NMR spectrum at the higher temperatures (-30  $^{\circ}$ C and +50  $^{\circ}$ C). The signal at -122 ppm in the +50  $^{\circ}$ C spectrum is a triplet. Draw a structure or an equation.

(c) What is the proton frequency (MHz) of the spectrometer which was used for these spectra?

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This is the P closest to the two PEt<sub>3</sub> groups, so expect triplet splitting. The dtd (J = 502, 49, 22 Hz) at  $\delta$  -17 is the only signal that shows a triplet, so this must be P<sup>1</sup>, which should also be coupled to both P<sup>2</sup> And P<sup>3</sup>, as observed.

$$P^{2}$$
 $P^{1}$ 
 $P^{1}$ 
 $P^{2}$ 
 $P^{1}$ 
 $P^{2}$ 
 $P^{3}$ 
 $P^{2}$ 
 $P^{3}$ 
 $P^{4}$ 
 $P^{3}$ 
 $P^{5}$ 
 $P^{5$ 

$$\delta$$
 -17,  ${}^{1}J_{\text{P1-P2}}$  = 502 Hz,  ${}^{2}J_{\text{P1-P4/5}}$  = 49 Hz,  ${}^{2}J_{\text{P1-P3}}$  = 20 Hz

 $P^2$  should also show the large  $^1J$  to  $P^1$ , so it must be the dd (J = 500, 45 Hz) at  $\delta$  18.  $P^1$  and  $P^2$  form an  $ABXY_2$  system, would need to do an AB quartet calculation to get accurate chemical shifts

 $P^2$   $\delta$  18,  $^1J_{P1-P2} = 500 \text{ Hz}$ ,  $^2J_{P1-P3} = 40 \text{ Hz}$ 

<sup>3</sup> P<sup>3</sup> is coupled to both P<sup>1</sup> and P<sup>2</sup> (dd, J = 46, 25 Hz), so it has to be the δ 112 signal  $\delta 112, {}^{2}J_{P3-P2} = 40 \text{ Hz}, {}^{2}J_{P3-P1} = 20 \text{ Hz}$ 

This is the signal with double area at  $\delta$  -122, d, J = 48 Hz  $P^4. P^5 \qquad \delta -122, \, ^1J_{P4/5-P1} = 46 \text{ Hz}$ 

(b) Identify the process which is responsible for the changes in the NMR spectrum at the higher temperatures (-30  $^{\circ}$ C and +50  $^{\circ}$ C). The signal at -122 ppm in the +50  $^{\circ}$ C spectrum is a triplet. Draw a structure or an equation.

The Pd migrates back and forth between  $P^1$  and  $P^2$ , so their chemical shifts are averaged, and both  $P^3$  and  $P^4/P^5$  become triplets, equally coupled to both. Since the two coupling constants are fairly close (expect the  $P^4/P^5$  coupling to be (49+0)/2=25, and  $P^3$  coupling to be (40+20)/2=30 Hz in size, the  $P^1/P^2$  signal becomes an approximate quartet.

The exchange is intramolecular, since the coupling between P<sup>1</sup>/P<sup>2</sup> and P<sup>4</sup>/P<sup>5</sup> is maintained in the high temperature spectrum

(c) What is the proton frequency (MHz) of the spectrometer which was used for these spectra?

32.4x(100/40.49) = 80 MHz

