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The isomeric structure of pentacoordinate chiral spirophosphoranes in solution by the combined use of NMR experiments and GIAO DFT calculations of NMR parameters†

Fedor M. Polyancev,¹ Kirill E. Metlushka, Dilyara N. Sadkova, Zilya R. Khisametdinova, Olga N. Kataeva, Vladimir A. Alfonsov, Shamil K. Latypov²* and Oleg G. Sinyashin¹

The interplay of NMR experiments and DFT calculations of NMR parameters is a reliable method for determining the relative configurations of pentacoordinate chiral spirophosphoranes bearing two six- or five-membered rings at the phosphorus atom in solution. The major product of the Betti based derivatives corresponds to the isomers with both substituents at chiral carbons being opposite to the P–H proton. The next populated product corresponds to the isomer with different chiralities at carbons. The least populated isomer is one with both substituents being at the same side of the heterocycle as the P–H bond.

Introduction

Chiral phosphorus compounds are widely used in stereoselective synthesis.^{1–5} For example, chiral phosphorus ligands are of importance in the design of catalysts for asymmetric homogeneous catalysis.⁶ Pentacoordinate phosphorus compounds have drawn special attention for their role as intermediates in some biological processes (*e.g.* enzymatic phosphoryl transfer reactions,^{7–29} hydrolysis/formation of DNA, RNA, cyclic AMP).^{30,31}

Their structural and dynamic properties strongly correlate with the reactivity/selectivity of the processes they are involved in. Although the knowledge about the stereochemistry (the absolute and relative configurations) of these compounds is important in understanding these processes, very few structural studies of pentacoordinate phosphorus systems have been published so far.^{32–38}

Pentacoordinate phosphoranes with an asymmetric phosphorus atom and with several additional chiral chelate ligands are especially difficult to analyze, since these chiral centers may generate a variety of stereoisomers and the determination of their fine structure is not straightforward. To this end X-ray single crystal investigations could be helpful, but obtaining

good single crystals is often time consuming and sometimes impossible. This point, coupled with the fact that most chemical reactions and biological processes occur in solution, makes it highly desirable to find an alternative, reliable, and easy way to determine the relative configurations of such hypervalent chiral phosphorus compounds directly in solution.^{33–38}

Progress in NMR correlation techniques and the state-of-the-art density functional theory (DFT) calculations of NMR parameters has emerged in recent years as a very powerful tool for fine structural analysis of organic compounds and biomolecules. There are a number of examples when DFT calculations of NMR ¹H, ¹³C and ¹⁵N parameters in the frame of the GIAO method allowed the establishment of tautomeric, isomeric and conformational structures as well.^{39–53} However applications of similar approaches to analyze phosphorus compounds by DFT calculations of ³¹P NMR parameters are very rare.^{54–56}

The study here shows that the interplay of the NMR experiment and theory is a powerful method for determining the relative configurations of pentacoordinate chiral spirophosphoranes. For the illustration several pentacoordinate spirophosphoranes bearing two six-membered (Betti based derivatives) and five-membered (α -aminocarbonic acid derivatives) rings at the phosphorus atom were used.

Results

Synthesis

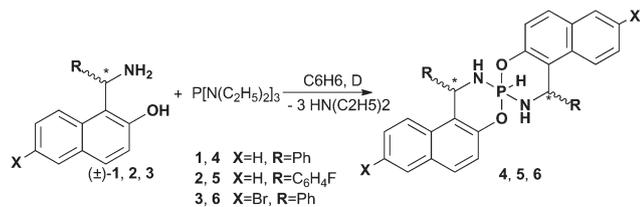
Three chiral elements (two centres and one axis) lead to the four possible combinations. In order to obtain the full set of

A.E. Arbuzov Institute of Organic and Physical Chemistry, Kazan Scientific Center, Russian Academy of Sciences, Arbuzov str. 8, Kazan, 420088, Russian Federation.

E-mail: lsk@iopc.ru

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Scheme 1

stereoisomers we used racemic Betti bases. The synthesis of P-H-spirophosphoranes was carried out by substituting the amino groups in the hexaethyltriimidophosphite by 1-(α -aminobenzyl)-2-naphthols (Scheme 1). For example, in the $^{31}\text{P}\{^1\text{H}\}$ NMR spectra of the reaction mixture of **4** there are two signals with the chemical shifts (CS) characteristic of phosphoranes (-75.4 (65%) and -76.7 (35%) ppm) that may be attributed to the formation of two isomers.

In contrast to the dominant product ($\delta^{31}\text{P} = -75.4$ ppm), the ^1H spectra of which correspond to the high symmetry structure, the second component apparently has no symmetry, and two spiro halves of phosphorane **4** differ in the ^1H NMR spectrum. When diethylamine hydrochloride is used as the catalyst, these two isomers could be obtained in an about 1 : 1 ratio and their isolation in individual form becomes possible

via crystallization. It is interesting that after about 4–6 hours in the NMR spectrum of the individual product that was major in the initial reaction mixture ($\delta^{31}\text{P} = -75.4$ ppm), additional signals appeared which can also be attributed to the structure of high symmetry ($\delta^{31}\text{P} = -80.7$ ppm) with two spiro-connected six-membered heterocycles being equivalent in the ^1H NMR spectra.

In a similar manner for **5** the formation of two isomers was observed which were separated and analyzed (Fig. 1a and b). In this case the ^{19}F NMR is particularly useful and allows distinguishing symmetrical (one singlet, $\delta^{19}\text{F} = -116$ ppm) and non-symmetrical (two singlet's of equal intensity, $\delta^{19}\text{F} = -115.8$ and $\delta^{19}\text{F} = -116.6$ ppm) isomers. The monitoring of the major isomer **5a** via the ^1H and ^{31}P NMR showed that after some time an additional product appeared (10%) in solution ($\delta^{19}\text{F} = -115.5$ and $\delta^{31}\text{P} = -81.0$ ppm) which presumably corresponds to another symmetrical isomer (Fig. 1c).

For compound **6** very similar spectra and their evolution were observed (ESI[†]).

Structure determination

Chemical structure and assignment. A variety of NMR correlations allows one to prove the chemical structure of two main stereoisomers practically directly. For example, first, starting

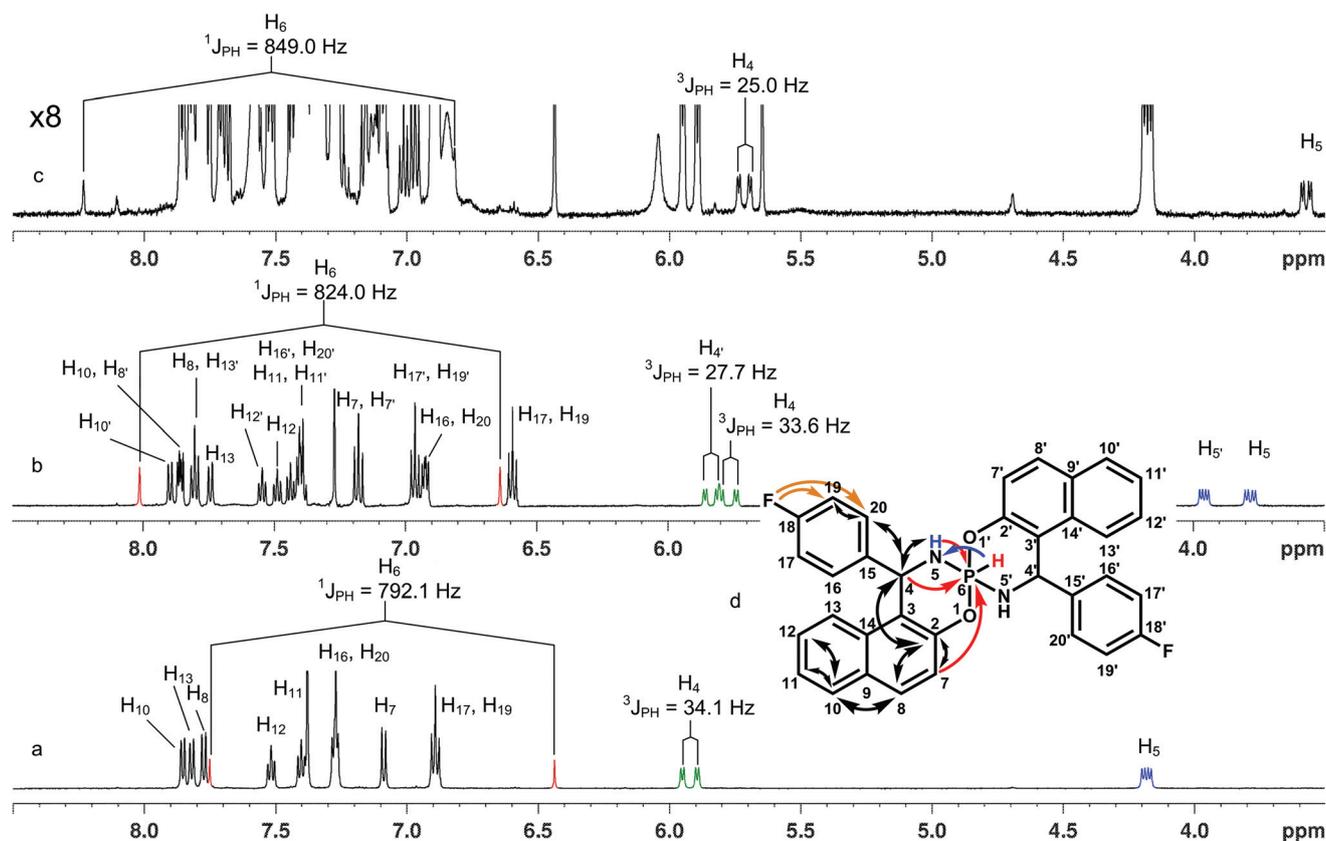


Fig. 1 ^1H spectra of **5** in CDCl_3 at 303 K. (a) Major product (fresh, **5a**); (c) the same product after 24 h; (b) second product; (d) structure of compound **5a** with principal NMR correlations: ^1H - ^1H COSY and ^1H - ^{13}C HMBC (black arrows), ^1H - ^{31}P HMBC (red arrows), ^1H - ^{15}N HMBC (blue arrows), ^{19}F - ^1H HETCOR (orange arrows).



Table 1 Some experimental and calculated NMR parameters^a for compounds 4–6

Comp.	X at C ₁₁	R at C ₄	Isomer	ΔE^b	$\delta^{31}\text{P}$	δH_6	$\delta\text{H}_{16,20}$	$\delta\text{H}_{17,19}$	δH_4	δH_5	$^1J_{\text{PH}}$	$^3J_{\text{PH}_4}$			
4	H	Ph	I	Exp ^c (a)	-75.4	7.08	7.32	7.19	5.95	4.18	788.5	34.1			
				0	-75.2	7.66	7.64	7.41	5.98	3.77	715.4	36.1			
				Exp (b)	-76.7	7.34	7.02, 7.45	6.93, 7.29	5.81, 5.87	3.80, 4.00	823.5	26.3, 33.1			
			II	0.53	-76.4	7.80	7.42, 7.88	7.21, 7.58	5.81, 5.84	3.46, 3.83	742.6	27.6, 35.2			
				Exp (c)	-80.7	7.54	n/d ^d	n/d	5.74	3.58	848.6	25.6			
				1.33	-82.5	7.75	7.87	7.58	5.65	3.36	762.7	25.2			
			5	H	FC ₆ H ₄	I	Exp (a)	-75.2	7.09	7.26	6.88	5.91	4.17	792.1	34.1
							0	-74.4	7.63	7.52	7.04	5.94	3.76	713.9	36.1
							Exp (b)	-76.6	7.32	6.91, 7.39	6.58, 6.95	5.76, 5.82	3.77, 3.95	824.0	27.7, 33.6
II	0.47	-75.7				7.75	7.37, 7.82	6.81, 7.17	5.77, 5.86	3.48, 3.78	738.3	27.6, 34.6			
	Exp (a)	-80.5				7.52	n/d	n/d	5.70	3.56	849.0	25.0			
	1.33	-82.1				7.74	7.82	7.15	5.63	3.33	764.2	24.7			
6	Br	Ph				I	Exp (a)	-76.4	7.08	7.27	7.19	5.89	4.17	790.4	34.1
							0	-75.6	7.62	7.58	7.41	5.92	3.74	718.3	36.0
							Exp (b)	-77.7	7.33	6.96, 7.28	6.96, 7.28	5.74, 5.78	3.77, 3.95	828.0	27.5, 33.7
			II	0.44	-76.5	7.77	7.20, 7.78	7.18, 7.59	5.68, 5.74	3.34, 3.81	742.6	27.6, 35.2			
				Exp (c)	-81.4	7.54	n/d	n/d	5.71	3.58	852.3	25.1			
				0.67	-80.5	7.69	7.78	7.59	5.60	3.25	769.9	24.1			

^a Chemical shifts in ppm, spin–spin couplings in Hz. ^b Relative energy, in kcal mol⁻¹. ^c Experimental data. ^d No data.

from ¹H–¹⁵N HSQC connectivities the NH protons were revealed, then the vicinal H₄ protons were assigned based on COSY data. Next, the ¹H–³¹P HMBC connectivities from the NH and H₄ to phosphorus allowed the determination of the structure of this central moiety (Fig. 1d). Further on, the ¹H–¹³C HSQC/HMBC and ¹H–¹H COSY correlations allow establishing the structures of two aromatic fragments (ESI[†]). In addition, for 5 the ¹⁹F–¹H HETCOR correlations help to establish connectivities up to fluorine atoms. As to the third (minor) isomer due to its low population and extensive overlapping with the signals of the first isomer in the ¹H NMR spectra, only some principal resonances can be identified with confidence (e.g. Fig. 1c). However, the 2D ¹H–³¹P HMBC experiments (ESI[†]) allow one to unambiguously reveal its signal in the ³¹P spectra and also the PH, NH and H₄ proton resonances (e.g. for 5c Fig. 1c).

Isomeric structure. The analysis of the NMR parameters of diastereomers of the phosphoranes 4–6 reveals some general particularities – namely, there are notable differences in some CSs and spin–spin couplings (SSCs) of the isomers (ESI[†]). The most spectacular are the values of direct ¹J_{HP} SSCs, that vary from ~790 (a-product) up to ~850 (c-product) Hz (Table 1), and also the difference in ³J_{PH₄} SSCs. The ¹H CSs of PH, H₄ and aryl protons and ³¹P CSs are diastereomer dependent in these products as well. Thus, it is likely that NMR parameters, being related to local magnetic environments, could be used in fine structural analysis.

In general, three chiral centers (at P, C₄ and C_{4'}) may generate four diastereomers: $\Delta_{\text{P}}R_{\text{C}}R_{\text{C}}/\Lambda_{\text{P}}S_{\text{C}}S_{\text{C}}$, $\Lambda_{\text{P}}R_{\text{C}}R_{\text{C}}/\Delta_{\text{P}}S_{\text{C}}S_{\text{C}}$, $\Delta_{\text{P}}R_{\text{C}}S_{\text{C}}/\Lambda_{\text{P}}S_{\text{C}}R_{\text{C}}$ and $\Delta_{\text{P}}S_{\text{C}}R_{\text{C}}/\Lambda_{\text{P}}R_{\text{C}}S_{\text{C}}$. However, when both aryls in chiral

centers are the same the last two isomers become equivalent in terms of NMR. Thus three main isomers were considered in details: with both aryl substituents being opposite to the P–H bond ($\Delta_{\text{P}}R_{\text{C}}R_{\text{C}}/\Lambda_{\text{P}}S_{\text{C}}S_{\text{C}}$, **I**), with one aryl opposite to the P–H bond and another one being on the same side of the cycle with the P–H bond ($\Delta_{\text{P}}R_{\text{C}}S_{\text{C}}/\Lambda_{\text{P}}S_{\text{C}}R_{\text{C}}$, **II**), in the third isomer both aryls are at the same side of the cycle with the P–H bond ($\Lambda_{\text{P}}R_{\text{C}}R_{\text{C}}/\Delta_{\text{P}}S_{\text{C}}S_{\text{C}}$, **III**) (Fig. 2).

Qualitative consideration of these structures predicts some stereospecific features. There are different dihedral angles between H₄–C and N–P in isomers **I** and **III**, that may lead to different three bond SSCs. As to the **II**-nd isomer, the geometries of its two spiro-halves are very similar to the halves of the **I**-st and **III**-rd isomers, therefore two sets of signals are expected for its ¹H (¹³C, ¹⁹F) spectra.

Thus, preliminary identification can be based on NOEs and symmetry considerations. Presumably, the “a” (major) products can be ascribed to the **I** isomer because there is no NOE between the *ortho*-aryl and P–H protons. Lack of symmetry in the second product (“b”) may be indicative of the **II**-isomer. As to the co-product (“c”) that appeared after some time in solution of the major isomer (“a”), there are no clear NOEs that can be used due to its low population and the signal overlapping with the main product.

To develop a well-argued relationship between the NMR parameters and the fine isomeric structure, DFT calculations of the NMR parameters were carried out for these structural hypotheses. It is well demonstrated that modern GIAO CS calculations on ¹³C, ¹⁵N and ¹H nuclei achieved experimental accuracy and can be used as an additional tool for chemical structure elucidation. There are a number of examples of application of such calculations on these nuclei to obtain fine information on the isomeric, tautomeric and conformational structure.^{39–53,57–60} At the same time the examples of

[†]The isomer **III** might be produced from isomer $\Delta_{\text{P}}R_{\text{C}}R_{\text{C}}$ ($\Lambda_{\text{P}}S_{\text{C}}S_{\text{C}}$) by the Berry pseudorotation process, so its configuration was assigned as $\Lambda_{\text{P}}R_{\text{C}}R_{\text{C}}$ ($\Delta_{\text{P}}S_{\text{C}}S_{\text{C}}$).



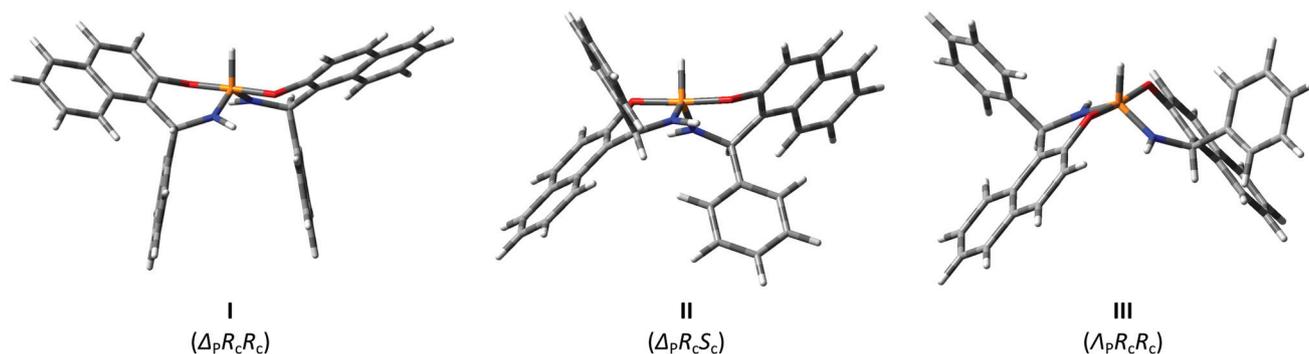


Fig. 2 Energy minimized structures of the main isomers of the title spirophosphorane 4.

application of these approaches on phosphorus are rather sparse.^{54–56}

We used a combination (PBE1PBE/6-311G(2d,2p)//PBE1PBE/6-31+G(d)) that proved to be reliable and effective for ³¹P NMR CS calculations. First, the geometry optimization in the frames of the DFT (PBE1PBE/6-31+G(d)) method predicts that these three main isomers are close in energy (Table 1). Additional calculations with the inclusion of solvent effects (in the frames of the PCM model, Table S1†) do not change notably the energy distribution. In general, these results are in qualitative agreement with experimental findings that all three isomers are present in the reaction mixture after some time.

The results of the CS and SSC calculations (GIAO PBE1PBE/6-311G(2d,2p)) for optimized structures are also in agreement with preliminary conclusions (some essential data are summarized in Table 1). According to the calculations, the PH, H₄ and NH protons and ³¹P CSs should be notably different, particularly, for the I-st and the III-rd isomers. Namely, for the I-isomer the PH proton should resonate at a higher field than for the III-isomer, while for the H₄ and the NH protons the reverse should be observed. Likewise the ³¹P resonance signal of the I-isomer should be at a lower field than in the III-isomer. For the II-isomer, intermediate values are expected (Table 1). Thus, according to the CS data the major (“a”) product should be assigned to the I-isomer for all title compounds. The second non-symmetrical product corresponds to the II-diastereomer while the minor product may be assigned to the III-isomer.

Some SSCs also demonstrate spectacular distinctions for three isomers (Table 1) that correlate well with experimental data. As can be seen the ¹J_{HP} constants are characteristic of these isomers and can be used to assign the isomeric structure. The smallest values (715.4 Hz) are predicted for the I-st isomers while for the III-rd forms these values are notably larger (762.7 Hz). For the II-isomers the PH direct SSCs are expected to have intermediate values (742.6 Hz). The calculations slightly underestimate (by *ca.* 1.1) the experimental data although the correlation is quite good. Thus, the comparison of the calculated *versus* experimental SSCs allows the assignment of the major product to the I-st isomer, the second

product to the II-nd isomer and the least populated product to the III-rd isomer.

The ³J_{HP} between P and H₄ is also very stereospecific (Table 1). First, the calculations predict SSCs to have large absolute values and be essentially different, at least, for the I-st and III-rd isomers. For the II-nd isomers there are two different values for each half but these SSC are close to the corresponding SSCs of the I-st and III-rd isomers, respectively. In spite of the fact that theory slightly underestimates these SSCs, the calculated values and tendencies are in good agreement with the experimentally observed data (Table 1). Thus, the comparison of the experimental *versus* calculated data also allows the assignment that is in accordance with the above conclusion.

These conclusions were supported by solid-state data in some cases. Namely, good crystals suitable for X-ray analysis were grown for the second isomer of racemic 4, which is crystallized as a conglomerate, and for the major product of racemic 5. The results of X-ray single crystal diffraction turned out to be in full agreement with the NMR data. In crystals the major product of 5 is realized in the I form (see Fig. 3) with both aryls directed opposite to the P–H bond. In general, the solid-state geometry is close to the DFT optimized one.

It is interesting that the crystallized product of 4 corresponded to the asymmetrical II-form that is also in agreement with the conclusion based on the NMR data. Thus for the title chiral pentacoordinate spirophosphoranes there are several NMR parameters that can be used to establish the isomeric structure. These relationships are summarized in Fig. 4.

Correlation of NMR parameters with the isomeric structure for other chiral pentacoordinate spirophosphoranes

The interplay of the NMR experiment and theory can be safely used to determine the isomeric structure of chiral pentacoordinate spirophosphoranes with six-membered heterocycles. To this end it is of interest to see if this approach could also be applied to determine the isomeric structure of other similar chiral pentacoordinate spirophosphoranes, *e.g.* with five-membered heterocycles. For example, there are experimental data for a series of chiral pentacoordinate spirophosphoranes obtained from α-amino carbonic acids (Scheme 2).^{33–37} These



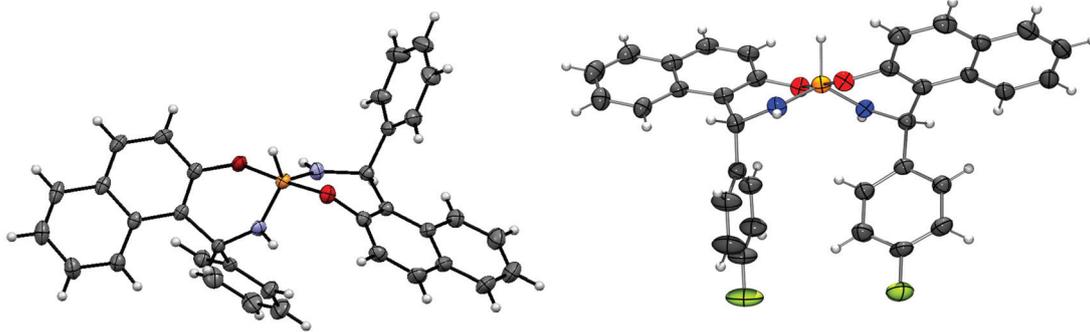


Fig. 3 Molecular geometry of diastereomers 4b (left) and 5a (right) as determined from X-ray single crystal diffraction.

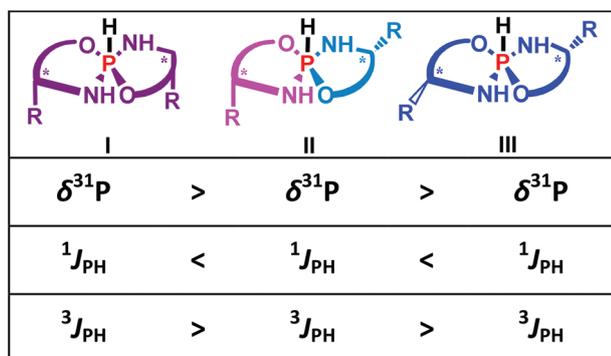
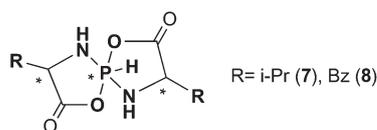


Fig. 4 Schematic representation of three isomers of 4–6 and the main relationships between the NMR parameters and isomeric structure. Magnetically equivalent fragments are colored identically.



Scheme 2

compounds were extensively analyzed by CD and X-ray single diffraction. When enantiomerically pure reagents were being used, the reactions produced only two diastereomers.

A brief survey of available NMR data demonstrates that there are also some specific features of experimental CSs and SSCs characteristic of two diastereomers although somewhat distinct from those observed for the title compounds. First, for five-membered spirophosphoranes the reverse ^{31}P CS distribution is observed, *i.e.* the “I” isomers resonate at a higher field than “III” isomers (some data are included in Table 2).[§] Second, the $^1J_{\text{PH}}$ values are smaller for the “I” isomers than for the “III”-forms. As to the $^3J_{\text{PH}}$ constants they are notably smaller (<13 Hz). Third, there are substantial long range $^4J_{\text{HPH}_4}$

[§] Unfortunately, ^1H CS's data cannot be analyzed in this case as different solvents were used in experiments for different diastereomers.

SSCs (2.4 Hz) in the “III” forms that were not seen in the “I” isomers presumably due to their small values.

These differences are reproduced by theory. According to our calculations performed for the simplest compounds from this series with *i*-Pr (7) or Bz (8) groups at chiral carbons (Table 2), indeed, the “I” isomer has to resonate at a higher field in the ^{31}P NMR spectra than the “III” form.

The calculated SSCs are also qualitatively consistent with the experimental values, although some underestimation is also seen. First, a larger one bond SSC is predicted for the “III” isomers (Table 2) than for the “I” form. According to the calculations, the $^3J_{\text{PH}}$ SSCs should be notably smaller than for the title compounds. Moreover, for the “III”-isomers one may expect a long range four bond SSC between HP and H₄ (2.0 *versus* 0.5 Hz) larger than for the “I” isomer. Thus, several NMR parameters can also be used to determine the relative configuration at chiral carbons in these pentacoordinate spirophosphoranes.

Discussion

Let us consider why some relationships are different in these two series of spirophosphoranes. There are two essential NMR features that distinguish 5-membered spirophosphoranes from 6-membered compounds: (a) $^3J_{\text{PH}}$ – small absolute value for the 5-cycle and large for the 6-cycle; (b) $\delta^{31}\text{P}_\text{I} > \delta^{31}\text{P}_\text{III}$ for 6-membered and $\delta^{31}\text{P}_\text{I} < \delta^{31}\text{P}_\text{III}$ for 5-membered.

To reveal key factors that determine these differences we analyzed the influence of substituents at carbon chiral centers and the nature of the fused aromatic fragments theoretically. To simplify the task and to exclude the specific influence of fused aromatic fragments, the models with the benzo fragment instead of the naphthyl group were analyzed (Fig. 5a and b, Table 3). According to these calculations, the relationships between the NMR parameters and isomeric structure are expected to be essentially similar to the title compounds. Thus, there is no specific effect of the naphthyl moiety on the NMR parameters' distribution in these isomers.

The next step to simplify the model is to check whether there is a specific influence of substituents at chiral carbons (*e.g.* steric interactions). For this purpose we analyzed models



Table 2 Some experimental and calculated NMR parameters^a for compounds 7 and 8

Comp.	R at C ₄	Isomer	ΔE^b	$\delta^{31}\text{P}$	δH_6	δH_4	δH_5	$^1J_{\text{PH}}$	$^3J_{\text{PH}_4}$	$^4J_{\text{H}_6\text{H}_4}$
7	i-Pr	I	Exp ^c (a) ³⁷	-65.0	7.18	3.71	5.76	798.5	13.0	n/d ^d
			0.19	-65.6	7.82	3.73	2.82	697.2	8.8	0.5
		III	Exp (b) ³⁷	-63.7	7.43	3.80	3.47	824.4	n/d	2.4
			0	-62.1	7.79	3.70	2.84	724.0	5.9	2.0
8	Bz	I	Exp (a) ³⁶	-63.0	7.09	n/d	n/d	804.9	n/d	n/d
			0	-68.5	7.95	4.16	3.02	710.3	10.8	0.4
		III	Exp (b) ³⁶	-60.0	5.64	n/d	n/d	810.2	n/d	2.4
			3.58	-64.5	6.62	3.81	2.67	733.6	9.4	2.1

^a Chemical shifts in ppm, spin-spin couplings in Hz. ^b Relative energy, in kcal mol⁻¹. ^c Experimental data. ^d No data.

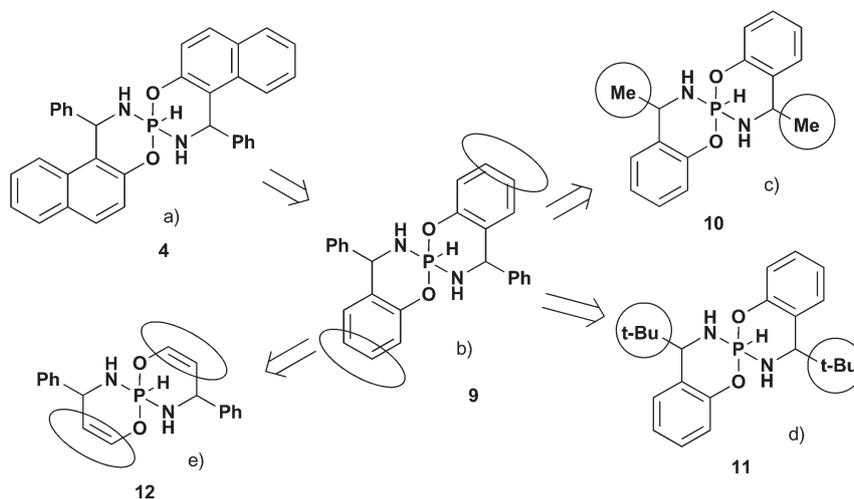


Fig. 5 Step-by-step simplification of the title pentacoordinate spirophosphoranes.

Table 3 Calculated^a NMR parameters for models 9–12

Comp.	R at C ₄	Isomer	ΔE^b	$\delta^{31}\text{P}$	δH_6	δH_4	δH_5	$^1J_{\text{PH}}$	$^3J_{\text{PH}_4}$
9	Ph	I	0	-72.7	7.41	5.11	3.56	717.5	35.4
		III	0.92	-80.6	7.90	4.87	3.24	770.5	20.8
10	Me	I	0	-72.8	7.19	4.04	3.41	707.9	34.5
		III	1.02	-78.5	7.62	3.84	2.78	762.2	25.3
11	t-Bu	I	0	-73.5	7.11	3.77	3.30	687.4	30.1
		III	0.84	-76.7	7.67	3.44	2.74	771.5	21.1
12	Ph	I	0	-93.8	7.71	4.86	3.45	729.1	21.2
		III	0.30	-88.1	7.30	5.28	2.60	826.4	0.6

^a Chemical shifts in ppm, spin-spin couplings in Hz. ^b Relative energy, in kcal mol⁻¹.

with small (Me, 10) and large (*t*-Bu, 11) alkyl groups, as well as aromatic (Ph, 9) fragments at chiral centers (Fig. 5c and d). According to calculations (Table 3), the main tendencies are also the same as for the title compounds. That is there is no specific influence of the R at the chiral carbon atoms on the $\delta^{31}\text{P}$ and $^3J_{\text{PH}}$ difference in the main isomers.

In order to reveal the impact of the fused aromatic unit on NMR parameters, the benzo fragment was replaced by a double bond (12, Fig. 4e) in the next step. In this case the reverse distribution of ^{31}P CSs for the two main isomers and notably smaller $^3J_{\text{PH}}$ SSCs are expected. This is very similar to the trends found in 5-membered compounds.



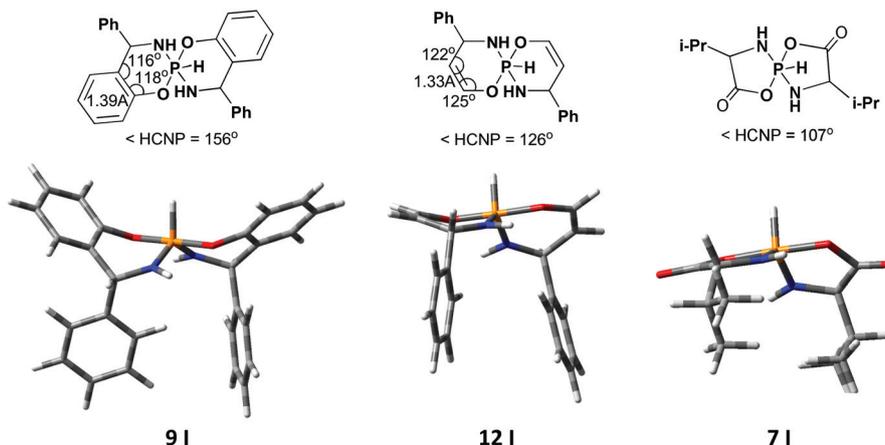


Fig. 6 Some geometrical parameters and calculated 3D structures for different 5- and 6-membered pentacoordinate spirophosphoranes.

Thus, in both series the nature of the substituent at the chiral center is not principal. The nature of the fused aromatic fragment is also not crucial. But NMR parameters are considerably different for compounds with the aromatic system and those without the aromatic system. In this case, it is the geometry of the heterocycle which plays the crucial role: while in 6-membered spirophosphoranes with the fused aromatic system the heterocycle adopts boat conformation, in 5- and 6-membered compounds without the aromatic fragment the heterocycle is essentially flat (or envelope) (Fig. 6). Therefore, in the first case the H_4-C_4-N-P dihedral angle is about 156° and results in a large $^3J_{PH}$ value, in the latter cases these angles are *ca.* 126° and 107° , thus small $^3J_{PH}$ should be observed. The same difference in the conformation leads to a different distribution of ^{31}P CSs in the two main isomers.

Conclusions

The major product of the Betti based derivatives corresponds to the isomers with both substituents at chiral carbons being opposite to the P–H proton. The next populated product corresponds to the isomer with different chiralities at carbons. The least populated isomer is one with both substituents being at the same side of the heterocycle as the P–H bond.

For such pentacoordinate chiral spirophosphoranes bearing two six-membered rings at the phosphorus atom the distinct NMR features, which allow establishing the isomeric structure, can be formulated as follows: the $\Delta_P R_C S_C / \Lambda_P R_C S_C$ isomer has two nonequivalent sets of signals in 1H (^{13}C) NMR; the $\Delta_P R_C R_C / \Lambda_P S_C S_C$ isomer resonances at a lower field in ^{31}P NMR, the PH proton resonances at a higher field, $^1J_{PH}$ is smaller, $^3J_{PH}$ is larger than similar parameters for the $\Delta_P S_C S_C / \Lambda_P R_C R_C$ isomer.

For similar spirophosphoranes bearing two five-membered rings at the phosphorus atom the NMR indicators are slightly different, in full agreement with the theory.

This study shows that the interplay of NMR experiments and DFT calculations of NMR parameters is a powerful and reliable method for determining the relative configurations of the pentacoordinate P–H-spirophosphoranes in solution.

Experimental section

NMR spectroscopy

All NMR experiments were performed with 600, 500 and 400 MHz (600.1, 500.1 and 400.1 MHz for 1H NMR; 150.9, 125.8 and 100.6 MHz for ^{13}C NMR; 242.9, 202.5 and 162.0 MHz for ^{31}P NMR, 60.81 MHz for ^{15}N NMR, 376.5 for ^{19}F respectively) spectrometers equipped with a 5 mm diameter probehead and a pulsed gradient unit capable of producing magnetic field pulse gradients in the z-direction of $53.5 G cm^{-1}$. For 1H - ^{13}C correlations the HSQC experiment is optimized for $J = 145$ Hz. For 1H - ^{13}C long range correlations the HMBC experiment is optimized for $J = 8$ Hz. For 1H - ^{31}P long range correlations the HMBC experiment is optimized for $J = 8$ Hz. For 1H - ^{15}N correlations the HSQC experiment is optimized for $J = 90$ Hz. For 1H - ^{15}N long range correlations the HMBC experiment is optimized for $J = 6$ Hz. For ^{19}F - 1H long range correlations the HETCOR experiment is optimized for $J = 3$ Hz. DOSY experiments were performed with ledbpgp2s, using a stimulated echo sequence and two spoil gradients. NOE experiments were performed with 1D DPFGNOE techniques. CSs (δ in ppm) were referenced to the solvent $CDCl_3$ ($\delta = 7.27$ ppm for 1H and 77.0 ppm for ^{13}C NMR) and to external H_3PO_4 (0.0 ppm) for ^{31}P NMR spectra, to external CH_3CN (235.5 ppm) for ^{15}N NMR spectra, to external C_6F_6 (-164.9 ppm) for ^{19}F NMR spectra.

Calculations

The quantum chemical calculations were performed using the Gaussian 03 software package.⁶¹ Full geometry optimizations have been carried out within the framework of the DFT (PBE1PBE) method using 6-31+G(d) basis sets. ^{31}P CSs were



calculated at the PBE1PBE/6-311G(2d,2p) level of theory.⁵⁶ ³¹P CSs were referred to H₃PO₄ calculated at the same level of theory. The linear scaling procedure was applied.⁵⁶ ¹H CSs were referred to TMS.

Synthesis of 1-(α -aminobenzyl)-2-naphthols (1–3)

Compounds 1–3 were synthesized in several steps. First, the corresponding 1,3-diaryl-2,3-dihydro-1*H*-naphth[1,2-*e*][1,3]oxazines were obtained.^{62–64} which were then hydrolyzed by trifluoroacetic acid in the presence of water.⁶⁵ The free bases were isolated by treating the resulting trifluoroacetates with sodium carbonate.⁶⁶

Synthesis of P–H-spirophosphoranes (4–6)

Spirophosphoranes (4–6) were synthesized by the reaction of the Betti bases and hexaethyltriimidophosphite: (1) without the addition of the catalyst (method A, only 4) resulting in the predominant formation of the ($\Delta_P R_C R_C / \Delta_P S_C S_C$)-isomer; (2) with the addition of diethylamine hydrochloride (5 mol%) as the catalyst (method B), which allows obtaining ($\Delta_P R_C R_C / \Delta_P S_C S_C$)- and ($\Delta_P R_C S_C / \Delta_P S_C R_C \equiv \Delta_P S_C R_C / \Delta_P R_C S_C$)-isomers in a ratio close to 1 : 1 by influencing the rate of P–Z bond cleavage (Z = O, N).⁶⁷

Method A: 1-(α -Aminobenzyl)-2-naphthol (1) (4 mmol) was dissolved in dry benzene (12 mL) under heating. Then the solution of hexaethyltriimidophosphite (2 mmol) in dry benzene (3 mL) was added and the reaction mixture was kept at reflux under argon for 8 hours. After that the solvent was removed and the residue was analyzed by ¹H and ³¹P NMR spectroscopy for the diastereomers' (4a and 4b) ratio determination (($\Delta_P R_C R_C / \Delta_P S_C S_C$) : ($\Delta_P R_C S_C / \Delta_P S_C R_C$) = 65 : 35). Pure ($\Delta_P R_C R_C / \Delta_P S_C S_C$)-P–H-phosphorane (4a) was obtained by recrystallization from benzene.

Method B: 1-(α -Aminobenzyl)-2-naphthols (1–3) (4 mmol) were dissolved in dry benzene (12 mL) under heating. Then dry diethylamine hydrochloride (0.1 mmol, 5 mol%) was added followed by the addition of hexaethyltriimidophosphite solution (2 mmol) in dry benzene (3 mL). The resulting reaction mixture was kept at reflux under argon for 5 hours. After that the solvent was removed and the residue was analyzed by ¹H and ³¹P NMR spectroscopy for the diastereomers' (a and b) ratio determination (($\Delta_P R_C R_C / \Delta_P S_C S_C$) : ($\Delta_P R_C S_C / \Delta_P S_C R_C$) = 1 : 1.1 for 4, 1 : 1 for 5, 1 : 0.9 for 6). Pure ($\Delta_P R_C R_C / \Delta_P S_C S_C$)-4a–6a and ($\Delta_P R_C S_C / \Delta_P S_C R_C$)-4b–6b diastereomers of P–H-phosphoranes were obtained by fractional crystallization from benzene.

Physical data

1-[Amino-(4-fluorophenyl)methyl]-2-naphthol trifluoroacetate (2-CF₃COOH). Yield: 82.2%; m.p. 150–151 °C. δ_H (400.1 MHz; DMSO-*d*₆; TMS) 6.30 (1 H, s, PhCHN), 7.19–8.06 (10 H, m, Ph), 8.80 (3 H, br, NH₃⁺). IR (KBr): ν = 1516 (COO[−]), 1586, 1601 (C=C_{naphth}), 1665 (COO[−]), 3066, 3288 cm^{−1} (NH₃⁺); elemental analysis calcd (%) for C₁₉H₁₅F₄NO₃: C 59.85, H 3.96, N 3.67; found: C 59.68, H 3.77, N 3.55.

1-(α -Aminobenzyl)-6-bromo-2-naphthol trifluoroacetate (3-CF₃COOH). Yield: 84.0%; m.p. 162–164 °C. δ_H (400.1 MHz; DMSO-*d*₆; TMS) 6.29 (1 H, s, PhCHN), 7.30–8.16 (10 H, m, Ph),

8.79 (3 H, br, NH₃⁺). IR (KBr): ν = 1503 (COO[−]), 1582, 1598 (C=C_{naphth}), 1664 (COO[−]), 3066, 3254 cm^{−1} (NH₃⁺); elemental analysis calcd (%) for C₁₉H₁₅BrF₃NO₃: C 51.60, H 3.42, Br 18.07, N 3.17; found: C 51.78, H 3.57, Br 17.89, N 3.35.

1-(α -Aminobenzyl)-6-bromo-2-naphthol (3). Yield: 95.9%; m.p. 137–138 °C. δ_H (500.1 MHz; CDCl₃; TMS) 6.13 (1 H, s, PhCHN), 7.19–7.89 (10 H, m, Ph). IR (KBr): ν = 1589, 1613 (C=C_{naphth}), 3292, 3362 cm^{−1} (NH₂); elemental analysis calcd (%) for C₁₇H₁₄BrNO: C 62.21, H 4.30, Br 24.35, N 4.27; found: C 62.04, H 4.47, Br 24.16, N 4.35.

($\Delta_P R_C R_C / \Delta_P S_C S_C$)-6a-Hydro-5,14-diphenyldinaphtho[2,1-*b*,2,1-*h*]-1,7,5,11,6-dioxadiazaphosphospiro[5.5]undecadiene-7a,16a (4a). (0.54 g, 51.6%, method A); (0.28 g, 26.5%, method B); m.p. 180–182 °C. δ_H (600.1 MHz; CDCl₃; TMS) 7.86 (2 H, d, ³J_{HH} 8.7, H₁₀, H_{10'}), 7.83 (2 H, d, ³J_{HH} 8.7, H₁₂, H_{12'}), 7.74 (2 H, d, ³J_{HH} 8.8, H₈, H_{8'}), 7.49 (2 H, dd, ³J_{HH} 8.7, ³J_{HH} 7.5, H₁₃, H_{13'}), 7.37 (2 H, dd, ³J_{HH} 8.7, ³J_{HH} 7.5, H₁₁, H_{11'}), 7.32 (4 H, dd, ³J_{HH} 8.7, ³J_{HH} 7.3, ⁴J_{HH} 1.8, H₁₆, H_{16'}, H₂₀, H_{20'}), 7.19 (6 H, m, H₁₇–H₁₉, H_{17'}–H_{19'}), 7.08 (1 H, d, ¹J_{PH} 788.5, H₆), 7.07 (2 H, d, ³J_{HH} 8.8, H₇, H_{7'}), 5.95 (2 H, dd, ³J_{PH} 34.1, ³J_{HH} 6.9, H₄, H_{4'}), 4.18 (2 H, dd, ²J_{PH} 14.2, ³J_{HH} 6.9, H₅, H_{5'}). δ_C (150.9 MHz; CDCl₃; TMS) 150.7 (d, ²J_{PC} 6.0, C₂, C_{2'}), 143.3 (s, C₁₅, C_{15'}), 130.4 (s, C₁₄, C_{14'}), 129.4 (s, C₉, C_{9'}), 128.9 (s, C₈, C_{8'}), 128.5 (s, C₁₂, C_{12'}), 128.1 (s, C₁₇, C_{17'}, C₁₉, C_{19'}), 126.7–126.5 (br, C₁₃, C_{13'}, C₁₆, C_{16'}, C₁₈, C_{18'}, C₂₀, C_{20'}), 124.8 (d, ³J_{PC} 8.5, C₃, C_{3'}), 123.5 (s, C₁₁, C_{11'}), 121.6 (s, C₁₀, C_{10'}), 121.2 (d, ³J_{PC} 2.1, C₇, C_{7'}), 53.6 (s, C₄, C_{4'}). δ_P (242.9 MHz; CDCl₃; H₃PO₄) −75.4 (s, P₆). δ_N (60.8 MHz; CDCl₃; NH₃) 60.4 (d, ¹J_{PN} 45.0, N₅, N_{5'}). IR (KBr): ν = 1193 (POC), 1586, 1620 (C=C_{naphth}), 2354 (PH), 3414 cm^{−1} (NH); elemental analysis calcd (%) for C₃₄H₂₇N₂O₂P: C 77.55, H 5.17, N 5.32, P 5.88; found: C 77.35, H 5.01, N 5.49, P 6.02.

($\Delta_P R_C S_C / \Delta_P S_C R_C$)-6a-Hydro-5,14-diphenyldinaphtho[2,1-*b*,2,1-*h*]-1,7,5,11,6-dioxadiazaphosphospiro[5.5]undecadiene-7a,16a (4b). (0.3 g, 28.4%, method B); m.p. 149–150 °C. δ_H (600.1 MHz; CDCl₃; TMS) 7.88 (1 H, d, ³J_{HH} 8.2, H₁₀), 7.85 (1 H, d, ³J_{HH} 8.4, H_{10'}), 7.84 (1 H, d, ³J_{HH} 8.5, H₈), 7.83 (1 H, d, ³J_{HH} 8.2, H₁₃), 7.79 (1 H, d, ³J_{HH} 8.7, H_{8'}), 7.78 (1 H, d, ³J_{HH} 8.4, H_{13'}), 7.52 (1 H, t, ³J_{HH} 7.5, H₁₂), 7.48 (1 H, t, ³J_{HH} 7.5, H_{12'}), 7.45 (2 H, d, ³J_{HH} 8.0, H₁₆, H₂₀), 7.41 (1 H, t, ³J_{HH} 7.5, H₁₁), 7.38 (1 H, t, ³J_{HH} 7.5, H_{11'}), 7.34 (1 H, d, ¹J_{PH} 823.5, H₆), 7.29 (2 H, t, ³J_{HH} 7.5, H₁₇, H₁₉), 7.22 (2 H, d, ³J_{HH} 7.5, H₁₈), 7.19 (1 H, d, ³J_{HH} 8.5, H₇), 7.16 (1 H, d, ³J_{HH} 8.7, H_{7'}), 7.02 (2 H, d, ³J_{HH} 7.3, H₁₆, H₂₀), 6.99 (1 H, d, ³J_{HH} 7.3, H₁₈), 6.93 (1 H, t, ³J_{HH} 7.3, H₁₇, H₁₉), 5.87 (1 H, dd, ³J_{PH} 26.3, ³J_{HH} 6.7, H₄), 5.81 (1 H, dd, ³J_{PH} 33.1, ³J_{HH} 7.1, H₄), 4.00 (1 H, dd, ²J_{PH} 13.8, ³J_{HH} 6.7, H₅), 3.80 (1 H, dd, ²J_{PH} 16.2, ³J_{HH} 7.1, H₅). δ_C (150.9 MHz; CDCl₃; TMS) 151.7 (d, ²J_{PC} 5.9, C₂), 151.6 (d, ²J_{PC} 10.7, C_{2'}), 144.7 (s, C₁₅'), 143.9 (s, C₁₅), 131.5 (s, C₉, C_{9'}), 129.9 (s, C₁₃), 129.2 (s, C₁₉'), 129.7 (s, C₁₃'), 128.6 (s, C₁₉), 127.8 (s, C₁₈'), 127.4 (s, C₁₂, C_{12'}), 127.3 (br, C₁₈, C₂₀'), 126.9 (s, C₂₀), 124.3 (s, C₁₁), 124.2 (s, C_{11'}), 122.4 (s, C₈), 122.3 (s, C_{8'}), 122.1 (s, C₇), 121.0 (s, C₇'), 54.7 (s, C₄'), 53.8 (s, C₄). δ_P (242.9 MHz; CDCl₃; H₃PO₄) −76.7 (s, P₆). δ_N (60.8 MHz; CDCl₃; NH₃) 61.9 (d, ¹J_{PN} 30.0, N₅), 59.9 (d, ¹J_{PN} 38.0, N_{5'}). IR (KBr): ν = 1194 (POC), 1589, 1623 (C=C_{naphth}), 2358 (PH), 3416 cm^{−1} (NH); elemental analysis



calcd (%) for C₃₄H₂₇N₂O₂P: C 77.55, H 5.17, N 5.32, P 5.88; found: C 77.38, H 4.98, N 5.42, P 6.05.

($\Delta_{\text{P}}\text{S}_c\text{S}_c/\text{A}_p\text{R}_c\text{R}_c$)-6a-Hydro-5,14-diphenyldinaphtho[2,1-*b*,2,1-*h*]-1,7,5,11,6-dioxadiazaphosphospiro[5.5]undecadiene-7a,16a (4c). M.p. 180–182 °C. δ_{H} (600.1 MHz; CDCl₃; TMS) 7.54 (1 H, d, ¹J_{PH} 848.6, H₆), 7.14 (2 H, d, ³J_{HH} 8.6, H₇), 5.74 (2 H, dd, ³J_{PH} 25.6, ³J_{HH} 6.1, H₄), 3.58 (2 H, dd, ²J_{PH} 17.3, ³J_{HH} 5.6, H₅). δ_{P} (242.9 MHz; CDCl₃; H₃PO₄) –80.7 (s, P₆). δ_{N} (60.8 MHz; CDCl₃; NH₃) 64.0 (d, ¹J_{PN} 38.0, N₅, N₅). Individual substance could not be isolated.

($\Delta_{\text{P}}\text{R}_c\text{R}_c/\text{A}_p\text{S}_c\text{S}_c$)-6a-Hydro-5,14-di(*p*-fluorophenyl)dinaphtho[2,1-*b*,2,1-*h*]-1,7,5,11,6-dioxadiazaphosphospiro[5.5]undecadiene-7a,16a (5a). (0.34 g, 30.1%, method B); m.p. 171–173 °C. δ_{H} (600.1 MHz; CDCl₃; TMS) 7.84 (2 H, d, ³J_{HH} 8.3, H₁₀, H₁₀), 7.81 (2 H, d, ³J_{HH} 8.3, H₁₃, H₁₃), 7.76 (2 H, d, ³J_{HH} 8.7, H₈, H₈), 7.51 (2 H, t, ³J_{HH} 8.3, H₁₂, H₁₂), 7.39 (2 H, t, ³J_{HH} 8.3, H₁₁, H₁₁), 7.26 (4 H, dd, ³J_{HH} 8.6, ⁴J_{FH} 5.7, H₁₆, H₂₀, H₁₆, H₂₀), 7.09 (1 H, d, ¹J_{PH} 792.1, H₆), 7.08 (2 H, d, ³J_{HH} 8.7, H₇, H₇), 6.88 (4 H, t, ³J_{HH} 8.7, H₁₇, H₁₉, H₁₇, H₁₉), 5.91 (2 H, dd, ³J_{PH} 34.1, ³J_{HH} 6.9, H₄, H₄), 4.17 (2 H, dd, ²J_{PH} 14.2, ³J_{HH} 6.9, H₅, H₅). δ_{C} (150.9 MHz; CDCl₃; TMS) 161.9 (d, ¹J_{CF} 245.3, C₁₈, C₁₈), 150.8 (d, ²J_{PC} 5.8, C₂, C₂), 139.4 (d, ⁴J_{CF} 2.0, C₁₅, C₁₅), 130.5 (s, C₁₄, C₁₄), 129.7 (s, C₉, C₉), 129.4 (s, C₈, C₈), 128.8 (s, C₁₃, C₁₃), 128.5–128.3 (br, C₁₆, C₁₆, C₂₀, C₂₀), 126.9 (s, C₁₂, C₁₂), 124.7 (d, ³J_{PC} 8.2, C₃, C₃), 123.8 (s, C₁₁, C₁₁), 121.5 (s, C₁₀, C₁₀), 121.4 (s, C₇, C₇), 115.1 (d, ²J_{CF} 20.0, C₁₇, C₁₇, C₁₉, C₁₉), 53.2 (s, C₄, C₄). δ_{P} (242.9 MHz; CDCl₃; H₃PO₄) –75.2 (s, P₆). δ_{N} (60.8 MHz; CDCl₃; NH₃) 61.0 (d, ¹J_{PN} 58.0, N₅, N₅). δ_{F} (376.5 MHz; CDCl₃; C₆F₆) –116.0 (s, F₁₈, F₁₈). IR (KBr): ν = 1195 (POC), 1588, 1623 (C=C_{naphth}), 2364 (PH), 3420 cm^{–1} (NH); elemental analysis calcd (%) for C₃₄H₂₅F₂N₂O₂P: C 72.59, H 4.48, N 4.98, P 5.51; found: C 72.66, H 4.31, N 4.87, P 5.32.

($\Delta_{\text{P}}\text{R}_c\text{S}_c/\text{A}_p\text{S}_c\text{R}_c$)-6a-Hydro-5,14-di(*p*-fluorophenyl)dinaphtho[2,1-*b*,2,1-*h*]-1,7,5,11,6-dioxadiazaphosphospiro[5.5]undecadiene-7a,16a (5b). (0.28 g, 24.8%, method B); m.p. 141–143 °C. δ_{H} (600.1 MHz; CDCl₃; TMS) 7.89 (1 H, d, ³J_{HH} 8.0, H₁₀), 7.85 (1 H, d, ³J_{HH} 8.0, H₁₀), 7.84 (1 H, d, ³J_{HH} 8.8, H₈), 7.80 (1 H, d, ³J_{HH} 7.7, H₈), 7.78 (1 H, d, ³J_{HH} 8.1, H₁₃), 7.73 (1 H, d, ³J_{HH} 8.5, H₁₃), 7.54 (1 H, t, ³J_{HH} 7.2, H₁₂), 7.48 (1 H, t, ³J_{HH} 7.5, H₁₂), 7.43 (1 H, t, ³J_{HH} 7.5, H₁₁), 7.39 (3 H, m, H₁₁, H₁₆, H₂₀), 7.32 (1 H, d, ¹J_{PH} 824.0, H₆), 7.18 (1 H, d, ³J_{HH} 8.9, H₇), 7.16 (1 H, d, ³J_{HH} 9.2, H₇), 6.95 (2 H, dd, ³J_{HH} 9.2, H₁₇, H₁₉), 6.91 (2 H, dd, ³J_{HH} 8.8, ⁴J_{FH} 5.4, H₁₆, H₂₀), 6.58 (1 H, dd, ³J_{HH} 8.8, H₁₇, H₁₉), 5.82 (1 H, dd, ²J_{PH} 27.7, ³J_{HH} 6.8, H₄), 5.76 (1 H, dd, ²J_{PH} 33.6, ³J_{HH} 7.6, H₄), 3.95 (1 H, dd, ³J_{PH} 14.0, ³J_{HH} 6.8, H₅), 3.77 (1 H, dd, ²J_{PH} 16.8, ³J_{HH} 7.2, H₅). δ_{C} (150.9 MHz; CDCl₃; TMS) 162.0 (d, ¹J_{CF} 247.3, C₁₈), 161.6 (d, ¹J_{CF} 244.8, C₁₈), 151.1 (d, ²J_{PC} 9.9, C₂), 151.0 (d, ²J_{PC} 14.8, C₂), 139.8 (d, ³J_{PC} 3.0, C₁₅), 139.1 (d, ³J_{PC} 3.4, C₁₅), 130.7 (s, C₁₄), 130.6 (s, C₁₄), 129.8 (d, ⁴J_{PC} 10.0, C₈), 129.4 (d, ⁴J_{PC} 23.6, C₈), 128.9 (s, C₁₀), 128.8 (s, C₁₀), 128.3 (d, ³J_{FC} 8.5, C₁₆, C₂₀), 128.0 (d, ³J_{FC} 8.2, C₁₆, C₂₀), 127.1 (s, C₁₂), 126.9 (s, C₁₂), 124.8 (d, ³J_{PC} 9.1, C₃), 123.9 (s, C₁₁), 123.7 (s, C₁₁), 122.9 (d, ³J_{PC} 10.9, C₃), 121.6 (s, C₁₃), 121.5 (s, C₁₃), 121.4 (d, ³J_{PC} 2.7, C₇), 120.4 (d, ³J_{PC} 1.3, C₇), 115.4 (d, ²J_{FC} 21.5, C₁₇, C₁₉), 114.7 (d, ²J_{PC} 21.3, C₁₇, C₁₉), 53.3 (s, C₄), 52.6 (s, C₄).

δ_{P} (242.9 MHz; CDCl₃; H₃PO₄) –76.6 (s, P₆). δ_{N} (60.8 MHz; CDCl₃; NH₃) 62.4 (d, ¹J_{PN} 30.0, N₅), 60.2 (d, ¹J_{PN} 45.0, N₅). δ_{F} (376.5 MHz; CDCl₃; C₆F₆) –115.8 (s, F₁₈), –116.6 (s, F₁₈). IR (KBr): ν = 1191 (POC), 1586, 1624 (C=C_{naphth}), 2358 (PH), 3422 cm^{–1} (NH); elemental analysis calcd (%) for C₃₄H₂₅F₂N₂O₂P: C 72.59, H 4.48, N 4.98, P 5.51; found: C 72.74, H 4.59, N 4.83, P 5.41.

($\Delta_{\text{P}}\text{S}_c\text{S}_c/\text{A}_p\text{R}_c\text{R}_c$)-6a-Hydro-5,14-di(*p*-fluorophenyl)dinaphtho[2,1-*b*,2,1-*h*]-1,7,5,11,6-dioxadiazaphosphospiro[5.5]undecadiene-7a,16a (5c). M.p. 171–173 °C. δ_{H} (600.1 MHz; CDCl₃; TMS) 7.52 (1 H, d, ¹J_{PH} 849.0, H₆), 5.70 (2 H, dd, ³J_{PH} 25.0, ³J_{HH} 6.8, H₄, H₄), 3.56 (2 H, dd, ²J_{PH} 17.6, ³J_{HH} 5.7, H₅, H₅). δ_{P} (242.9 MHz; CDCl₃; H₃PO₄) –80.5 (s, P₆). δ_{N} (60.8 MHz; CDCl₃; NH₃) 64.2 (d, ¹J_{PN} 45.0, N₅, N₅). δ_{F} (376.5 MHz; CDCl₃; C₆F₆) –115.5 (s, F₁₈, F₁₈). Individual substance could not be isolated.

($\Delta_{\text{P}}\text{R}_c\text{R}_c/\text{A}_p\text{S}_c\text{S}_c$)-6a-Hydro-2,11-dibromo-5,14-diphenyldinaphtho[2,1-*b*,2,1-*h*]-1,7,5,11,6-dioxadiazaphosphospiro[5.5]undecadiene-7a,16a (6a). (0.45 g, 32.8%, method B); m.p. 189–192 °C. δ_{H} (600.1 MHz; CDCl₃; TMS) 7.98 (2 H, br, H₁₀, H₁₀), 7.71 (2 H, d, ³J_{HH} 9.1, H₁₃, H₁₃), 7.65 (2 H, d, ³J_{HH} 8.7, H₈, H₈), 7.55 (2 H, t, ³J_{HH} 9.1, H₁₂, H₁₂), 7.27 (4 H, br, H₁₆, H₂₀, H₁₆, H₂₀), 7.19 (6 H, m, H₁₇, H₁₈, H₁₉, H₁₇, H₁₈, H₁₉), 7.08 (1 H, d, ¹J_{PH} 790.4, H₆), 7.09 (2 H, d, ³J_{HH} 8.7, H₇, H₇), 5.89 (2 H, dd, ³J_{PH} 34.1, ³J_{HH} 6.7, H₄, H₄), 4.17 (2 H, dd, ²J_{PH} 14.0, ³J_{HH} 6.7, H₅, H₅). δ_{C} (150.9 MHz; CDCl₃; TMS) 151.7 (s, C₂, C₂), 143.5 (s, C₁₅, C₁₅), 131.1 (s, C₁₄, C₁₄), 131.0 (s, C₁₀, C₁₀), 130.5 (s, C₁₂, C₁₂), 129.4 (s, C₉, C₉), 128.9 (s, C₁₇, C₁₉, C₁₇, C₁₉), 128.7 (s, C₈, C₈), 127.1–126.9 (br, C₁₆, C₁₈, C₂₀, C₁₆, C₁₈, C₂₀), 125.4 (d, ³J_{PC} 8.6, C₃, C₃), 124.0 (s, C₁₃, C₁₃), 123.0 (s, C₇, C₇), 117.9 (s, C₁₁, C₁₁), 52.2 (s, C₄, C₄). δ_{P} (242.9 MHz; CDCl₃; H₃PO₄) –76.4 (s, P₆). δ_{N} (60.8 MHz; CDCl₃; NH₃) 60.4 (d, ¹J_{PN} 35.0, N₅, N₅). IR (KBr): ν = 1199 (POC), 1587, 1619 (C=C_{naphth}), 2353 (PH), 3419 cm^{–1} (NH); elemental analysis calcd (%) for C₃₄H₂₅Br₂N₂O₂P: C 59.67, H 3.68, Br 23.35, N 4.09, P 4.53; found: C 59.86, H 3.49, Br 23.54, N 3.89, P 4.48.

($\Delta_{\text{P}}\text{R}_c\text{S}_c/\text{A}_p\text{S}_c\text{R}_c$)-6a-Hydro-2,11-dibromo-5,14-diphenyldinaphtho[2,1-*b*,2,1-*h*]-1,7,5,11,6-dioxadiazaphosphospiro[5.5]undecadiene-7a,16a (6b). (0.3 g, 21.8%, method B); m.p. 156–158 °C. δ_{H} (600.1 MHz; CDCl₃; TMS) 8.02 (1 H, d, ⁴J_{HH} 1.7, H₁₀), 7.99 (1 H, d, ⁴J_{HH} 1.9, H₁₀), 7.73 (1 H, d, ³J_{HH} 8.7, H₈), 7.67 (2 H, d, ³J_{HH} 9.1, H₈, H₁₃), 7.63 (1 H, d, ³J_{HH} 9.1, H₁₃), 7.56 (1 H, dd, ³J_{HH} 9.1, ⁴J_{HH} 1.7, H₁₂), 7.52 (1 H, dd, ³J_{HH} 9.1, ⁴J_{HH} 1.9, H₁₂), 7.33 (1 H, d, ¹J_{PH} 828.0, H₆), 7.28 (4 H, m, H₁₆, H₁₇, H₁₉, H₂₀), 7.23 (1 H, m, H₁₈), 7.19 (1 H, d, ³J_{HH} 8.7, H₇), 7.16 (1 H, d, ³J_{HH} 9.1, H₇), 7.01 (1 H, m, H₁₈), 6.96 (4 H, br, H₁₆, H₁₇, H₁₉, H₂₀), 5.78 (1 H, dd, ²J_{PH} 27.5, ³J_{HH} 6.9, H₄), 5.74 (1 H, dd, ²J_{PH} 33.7, ³J_{HH} 7.3, H₄), 3.95 (1 H, dd, ³J_{PH} 13.8, ³J_{HH} 6.5, H₅), 3.77 (1 H, dd, ²J_{PH} 16.3, ³J_{HH} 7.3, H₅). δ_{C} (150.9 MHz; CDCl₃; TMS) 151.8 (d, ²J_{PC} 16.2, C₂), 151.7 (d, ²J_{PC} 20.3, C₂), 144.0 (s, C₁₅), 143.4 (s, C₁₅), 131.3 (s, C₉), 131.2 (s, C₁₀), 131.1 (s, C₁₀), 130.9 (s, C₉), 130.5 (s, C₁₂), 130.4 (s, C₁₂), 129.8 (s, C₁₄), 129.7 (s, C₁₄), 129.1 (br, C₈, C₁₇, C₁₉), 128.7 (s, C₈), 128.5 (s, C₁₇, C₁₉), 127.8 (s, C₁₈), 127.2 (s, C₁₈), 127.0 (s, C₁₆, C₂₀), 126.6 (s, C₁₆, C₂₀), 125.2 (d, ³J_{PC} 7.7, C₃), 124.0 (s, C₁₃), 123.9 (s, C₁₃), 123.5 (d, ³J_{PC} 10.6, C₃), 123.4 (d, ³J_{PC} 2.7, C₇), 123.0 (br, C₇), 117.9 (s, C₁₁), 117.7 (s, C₁₁),



54.5 (s, C_{4'}), 53.6 (s, C₄). δ_{P} (242.94 MHz; CDCl₃; H₃PO₄) -77.7 (s, P₆). δ_{N} (60.8 MHz; CDCl₃; NH₃) 61.9 (d, $^1J_{\text{PN}}$ 30.0, N₅), 59.9 (d, $^1J_{\text{PN}}$ 40.0, N₅). IR (KBr): ν = 1191 (POC), 1586, 1617 (C=C_{naphth}), 2370 (PH), 3422 cm⁻¹ (NH); elemental analysis calcd (%) for C₃₄H₂₅Br₂N₂O₂P: C 59.67, H 3.68, Br 23.35, N 4.09, P 4.53; found: C 59.75, H 3.57, Br 23.12, N 4.27, P 4.65.

($\Delta_{\text{P}}\text{S}_{\text{c}}\text{S}_{\text{c}}/\text{A}_{\text{P}}\text{R}_{\text{c}}\text{R}_{\text{c}}$)-6a-Hydro-2,11-dibromo-5,14-diphenyldi-naphtho[2,1-*b*,2,1-*h*]-1,7,5,11,6-dioxadiazaphosphaspiro[5.5]undecadiene-7a,16a (6c). M.p. 189–192 °C. δ_{H} (600.1 MHz; CDCl₃; TMS) 7.54 (1 H, d, $^1J_{\text{PH}}$ 852.3, H₆), 5.71 (2 H, dd, $^3J_{\text{PH}}$ 25.1, $^3J_{\text{HH}}$ 6.9, H₄, H_{4'}), 3.58 (2 H, dd, $^2J_{\text{PH}}$ 18.1, $^3J_{\text{HH}}$ 5.1, H₅, H_{5'}). δ_{P} (242.9 MHz; CDCl₃; H₃PO₄) -81.4 (s, P₆). Individual substance could not be isolated.

X-ray structure determination

Data sets for single crystals of **4b** and **5a** were collected on Bruker AXS Kappa APEX and Smart Apex diffractometers respectively with graphite-monochromated Mo K α radiation (λ = 0.71073 Å) using complete sphere mode. Programs used: data collection APEX2,⁶⁸ data reduction 'SAINT'⁶⁹ absorption correction SADABS version 2.10,⁷⁰ structure solution SHELXS97,⁷¹ structure refinement by full-matrix least-squares against F^2 using SHELXL-97⁷¹, hydrogen atoms calculated and refined as riding atoms, except the hydrogens attached to phosphorus and nitrogen atoms which were located from a difference Fourier map and refined isotropically. The crystals contain solvent channels, modeled by disordered ethanol and acetone molecules. Crystals **5b** contain a solvate benzene molecule, which is disordered and was refined with geometry and vibrational parameter constraints.

Crystal data for 4b

Formula C₃₄H₂₇N₂O₂P, crystal size 0.39 × 0.39 × 0.15 mm³, M = 526.54, orthorhombic, space group $P2_12_12_1$, a = 8.6029(2), b = 15.2183(3), c = 20.4343(4) Å, V = 2675.3(1) Å³, Z = 4, ρ_{c} = 1.307 g cm⁻³, μ = 0.138 mm⁻¹, T = 150(2) K, θ range = 1.67 to 31.17°, reflections collected: 49 141; independent: 8492 (R_{int} = 0.0354) and 7309 observed reflections [$I \geq 2\sigma(I)$], 364 refined parameters, R_1 = 0.0406, wR_2 = 0.0941 [$I > 2\sigma(I)$]; flack parameter -0.03(2), maximal residual electron density: 0.400/-0.229 e Å⁻³.

Crystal data for 5a

Formula C₃₇H₂₈F₂N₂O₂P, crystal size 0.24 × 0.17 × 0.12 mm³, M = 601.58, monoclinic, space group $C2/c$, a = 38.86(2), b = 8.799(5), c = 17.269(9) Å, β = 95.512(7)°, V = 5878(6) Å³, Z = 8, ρ_{c} = 1.360 g cm⁻³, μ = 0.144 mm⁻¹, T = 293(2) K, θ range = 2.11 to 26.60°, reflections collected: 28 391; independent: 6119 (R_{int} = 0.1310) and 1891 observed reflections [$I \geq 2\sigma(I)$], 406 refined parameters, R_1 = 0.0440, wR_2 = 0.0786 [$I > 2\sigma(I)$]; maximal residual electron density: 0.185/-0.193 e Å⁻³. Due to positional disorder of the solvate benzene molecule the ratio of the observed to collected reflections is low ~31%.

CCDC 1542633 and 1542634 contain the supplementary crystallographic data for this paper.

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