

# Bis(diphenylphosphorothioyl) trisulfide

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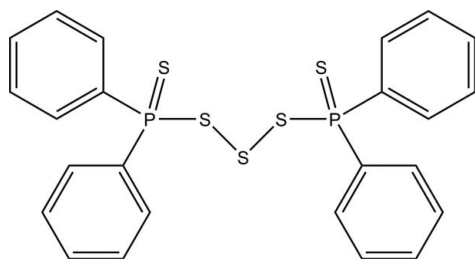
Received 14 July 2008; accepted 29 July 2008

 Key indicators: single-crystal X-ray study;  $T = 297$  K; mean  $\sigma(\text{C}-\text{C}) = 0.005$  Å;  $R$  factor = 0.048;  $wR$  factor = 0.107; data-to-parameter ratio = 18.5.

In the title compound,  $\text{C}_{24}\text{H}_{20}\text{P}_2\text{S}_5$ , the P atoms are arranged *trans* with respect to the  $\text{S}_3$  group and the  $\text{S}=\text{P}-\text{S}-\text{S}$  systems have *cisoid* geometry, with an average  $\text{S}-\text{P}-\text{S}-\text{S}$  torsion angle of  $-56.7^\circ$ . The dihedral angles between the two phenyl rings attached to the P atoms are  $87.33$  (12) and  $75.67$  (10) $^\circ$ . In the crystal structure, the molecules are linked into chains running parallel to the  $a$  axis by weak intermolecular  $\text{C}-\text{H}\cdots\text{S}$  hydrogen bonds. Centrosymmetrically related chains are further connected by  $\pi-\pi$  stacking interactions, with a centroid-to-centroid distance of  $3.795$  (5) Å.

## Related literature

For related literature, see: Deleanu *et al.* (2002); Drake *et al.* (2001*a,b*); Gallacher & Pinkerton (1992*a,b*, 1993); Kulcsar *et al.* (2005); Newton *et al.* (1993); Silvestru *et al.* (1994*a,b*); Buranda *et al.* (1991); Fest & Schmidt (1982); Knopik *et al.* (1993); Lawton (1970); McCleverty *et al.* (1983); Molyneux (1967); Perlikowska *et al.* (2004); Potrzebowski *et al.* (1991, 1994); Tiekink (2001); Tkachev *et al.* (1976); Zhang *et al.* (2004); Emsley (1994); Yadav *et al.* (1989).



## Experimental

### Crystal data

 $\text{C}_{24}\text{H}_{20}\text{P}_2\text{S}_5$   
 $M_r = 530.64$   
 Triclinic,  $P\bar{1}$   
 $a = 9.2287$  (8) Å  
 $b = 11.5476$  (10) Å

 $c = 12.9728$  (12) Å  
 $\alpha = 92.690$  (2) $^\circ$   
 $\beta = 105.287$  (2) $^\circ$   
 $\gamma = 106.124$  (2) $^\circ$   
 $V = 1270.3$  (2) Å<sup>3</sup>
 $Z = 2$   
 Mo  $K\alpha$  radiation  
 $\mu = 0.59$  mm<sup>-1</sup>
 $T = 297$  (2) K  
 $0.35 \times 0.27 \times 0.21$  mm

### Data collection

 Bruker SMART APEX  
 diffractometer  
 Absorption correction: multi-scan  
 (SADABS; Bruker, 2000)  
 $T_{\min} = 0.819$ ,  $T_{\max} = 0.885$ 

 13712 measured reflections  
 5178 independent reflections  
 4536 reflections with  $I > 2\sigma(I)$   
 $R_{\text{int}} = 0.029$ 

### Refinement

 $R[F^2 > 2\sigma(F^2)] = 0.047$   
 $wR(F^2) = 0.106$   
 $S = 1.13$   
 5178 reflections

 280 parameters  
 H-atom parameters constrained  
 $\Delta\rho_{\max} = 0.40$  e Å<sup>-3</sup>  
 $\Delta\rho_{\min} = -0.19$  e Å<sup>-3</sup>

**Table 1**  
 Hydrogen-bond geometry (Å,  $^\circ$ ).

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
$\text{C17}-\text{H17}\cdots\text{S5}^i$	0.93	2.94	3.737 (3)	145

 Symmetry code: (i)  $x + 1, y, z$ .

Data collection: SMART (Bruker, 2000); cell refinement: SAINT-Plus (Bruker, 2001); data reduction: SAINT-Plus; program(s) used to solve structure: SHELXS97 (Sheldrick, 2008); program(s) used to refine structure: SHELXL97 (Sheldrick, 2008); molecular graphics: DIAMOND (Brandenburg & Putz, 2006); software used to prepare material for publication: publCIF (Westrip, 2008).

This work was supported by the Romanian Ministry of Education and Research (CNCSIS grant No. 12/1456/2007). We thank the National Center for X-ray Diffraction (Babes-Bolyai University, Cluj-Napoca) for support in the solid-state structure determination.

Supplementary data and figures for this paper are available from the IUCr electronic archives (Reference: RZ2237).

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**supplementary materials**

*Acta Cryst.* (2008). E64, o1683-o1684 [ doi:10.1107/S1600536808024070 ]

## Bis(diphenylphosphorothioyl) trisulfide

M. Kulcsar, A. Silvestru and M. Vonas

### Comment

Our interest was focused for long time on studies concerning reactions between bis(diorganothiophosphoryl)disulfides and diorganodichalcogenides of type  $R_2E_2$  ( $E = \text{Se}, \text{Te}$ ) in order to obtain new organochalcogen compounds with diorganodithiophosphorus ligands (Newton *et al.*, 1993; Silvestru *et al.*, 1994*a,b*; Drake *et al.*, 2001*a,b*; Kulcsar *et al.*, 2005; Deleanu *et al.*, 2002). Bis(diorganothiophosphoryl)disulfides attracted much interest also due to their potential applications as pesticides (Fest & Schmidt, 1982), additives in motor oils (Molyneux, 1967) and in the rubber vulcanization (McCleverty *et al.*, 1983). Both alkyl and aryl substituted derivatives of type  $[R_2P(S)S]_2$  were already structurally characterized, *e.g.*  $R = \text{OPr-i}$  (Lawton, 1970; Tkachev *et al.*, 1976; Tiekink, 2001; Zhang *et al.*, 2004),  $R = \text{OMe}, \text{OBu-t}$  (Potrzebowski *et al.*, 1991), cyclohexyl (Buranda *et al.*, 1991), Me, Pr-i (Gallacher & Pinkerton, 1992*b*), Ph (Gallacher & Pinkerton, 1993), OPh (Gallacher & Pinkerton, 1993; Knopik *et al.*, 1993), menthoxy (Perlikowska *et al.*, 2004),  $R_2 = \text{OCMe}_2\text{—CMe}_2\text{O}$  (Yadav *et al.*, 1989),  $\text{OCMe}_2\text{—CH}_2\text{O}$  (Potrzebowski *et al.*, 1994). A search of the Cambridge Structure Database revealed that only for one trisulfide,  $[\text{Et}_2\text{P(S)S}]_2\text{S}$ , the X-ray crystal structure was determined (Gallacher & Pinkerton, 1992*a*). Here we report about the phenyl substituted analog,  $[\text{Ph}_2\text{P(S)S}]_2\text{S}$ .

The chalcogen atoms S2 and S3 are doubly bonded to phosphorus [ $\text{S2}=\text{P1} = 1.9351(9)$ ;  $\text{S3}=\text{P2} = 1.9303(9)$  Å], while the  $\text{P1—S1}$  [ $2.1171(9)$  Å] and  $\text{P2—S4}$  [ $2.1282(9)$  Å] distances correspond to single P—S bonds (*cf.*  $[\text{Ph}_2\text{P(S)S}]_2$ :  $\text{P}=\text{S} = 1.930(1)$ ,  $\text{P—S} = 2.139(1)$  Å; Gallacher & Pinkerton, 1993). The sulfur—sulfur distances within the  $\text{S}_3$  group are not significantly different [ $\text{S5—S4} = 2.0407(10)$ ,  $\text{S5—S1} = 2.0440(10)$  Å], corresponding to a S—S single bond. These values are similar to those found in bis(diorganothiophosphoryl)disulfides or in  $[\text{Et}_2\text{P(S)S}]_2\text{S}$ . The  $\text{SPS}_3\text{PS}$  skeleton of the title compound adopts a twisted zigzag chain structure (Fig. 1), with the torsion angles  $\text{S2—P1—S1—S5} = -56.30(5)^\circ$ ,  $\text{P1—S1—S5—S4} = 96.84(4)^\circ$ ,  $\text{S1—S5—S4—P2} = 87.42(5)^\circ$  and  $\text{S5—S4—P2—S3} = -57.13(5)^\circ$ . Although apparently the conformation of the  $\text{SPS}_3\text{PS}$  skeleton is similar to that of the previously reported ethyl derivative, some differences should be noted. In the title compound the phosphorus atoms are *trans* with respect of the central  $\text{S}_3$  group [ $\text{P1—S1}\cdots\text{S5—P2} = 171.4^\circ$ ], as are in the related  $[\text{Et}_2\text{P(S)S}]_2\text{S}$  compound (the torsion angle between corresponding atoms is  $159.8^\circ$ ). In both cases the central S atom and the terminal S atoms, respectively, are placed on opposite sides of the best plane described by the remaining atoms of the skeleton. However, the  $\text{S}=\text{P}\cdots\text{P}=\text{S}$  torsion angle is  $138.8^\circ$ , but only  $-89.4^\circ$  in the ethyl derivative. Moreover, the  $\text{S}=\text{P—S—S}$  system has a *cisoid* geometry [ $\text{S2—P1—S1—S5} = -56.30(5)^\circ$ ,  $\text{S5—S4—P2—S3} = -57.13(5)^\circ$ ], while it has a *transoid* geometry in  $[\text{Et}_2\text{P(S)S}]_2\text{S}$  (average  $\text{S}=\text{P—S—S}$  torsion angle  $179.6^\circ$ ; Gallacher & Pinkerton, 1992*a*). The  $\text{S—P—S}$  angles [ $\text{S2—P1—S1} = 113.77(4)^\circ$  and  $\text{S3—P2—S4} = 114.34(4)^\circ$ ] are consistent with a *cisoid* geometry, similar with that found for  $[\text{Ph}_2\text{P(S)S}]_2$  [ $114.44(4)^\circ$ ; Gallacher & Pinkerton, 1993], but much larger than in the *transoid* derivatives [ $(\text{PhO})_2\text{P(S)S}]_2$  [ $108.39(7)^\circ$ ; Gallacher & Pinkerton, 1993] and  $[\text{Et}_2\text{P(S)S}]_2\text{S}$  [av.  $103.7^\circ$ ; Gallacher & Pinkerton, 1992*a*]. The dihedral angles formed by the plane of the phenyl rings attached to the P1 and P2 atoms are  $87.33(12)$   $75.67(10)^\circ$  respectively. The crystal structure is stabilized by weak intermolecular hydrogen bonding interactions (Emsley, 1994) between the central sulfur atom and an aromatic proton of a neighbouring molecule (Table 1) forming chains parallel to the *a* axis (Fig. 2). Weak intermolecular  $\text{S}\cdots\text{H}$  contacts were observed in  $[(\text{PhO})_2\text{P(S)S}]_2$  [ $2.954(1)$  Å], but they are ab-

## supplementary materials

sent in  $[\text{Et}_2\text{P}(\text{S})\text{S}]_2\text{S}$  or  $[\text{Ph}_2\text{P}(\text{S})\text{S}]_2$ . Centrosymmetrically related chains are further connected by  $\pi$ - $\pi$  stacking interactions involving the C13–C18 phenyl rings, with centroid-to-centroid distance of 3.795 (5) Å.

### Experimental

The title compound was isolated as a by-product during recrystallization of  $\text{PhSeS}_2\text{PPh}_2$  obtained in the reaction between  $[\text{Ph}_2\text{P}(\text{S})\text{S}]_2$  and  $\text{Ph}_2\text{Se}_2$ .

### Refinement

All C-bound H atoms were placed in calculated positions (C–H = 0.93–0.97 Å) and treated using a riding model with  $U_{\text{iso}} = 1.2U_{\text{eq}}(\text{C})$ .

### Figures

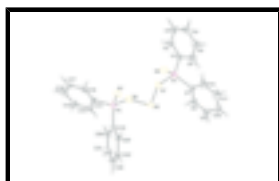


Fig. 1. A view of the title compound, with the atomic numbering scheme, showing displacement ellipsoids at the 50% probability level. H atoms are drawn as spheres of arbitrary radii.



Fig. 2. View of the S...H intermolecular interactions (dashed lines) in the title compound. Only H involved in hydrogen bonding interactions are shown. Symmetry codes: (i)  $1 + x, y, z$ , (ii)  $-1 + x, y, z$ .

### Bis(diphenylphosphorothioyl) trisulfide

#### Crystal data

$\text{C}_{24}\text{H}_{20}\text{P}_2\text{S}_5$

$M_r = 530.64$

Triclinic,  $P\bar{1}$

Hall symbol: -P 1

$a = 9.2287$  (8) Å

$b = 11.5476$  (10) Å

$c = 12.9728$  (12) Å

$\alpha = 92.690$  (2)°

$\beta = 105.287$  (2)°

$\gamma = 106.124$  (2)°

$V = 1270.3$  (2) Å<sup>3</sup>

$Z = 2$

$F_{000} = 548$

$D_x = 1.387$  Mg m<sup>-3</sup>

Mo  $K\alpha$  radiation

$\lambda = 0.71073$  Å

Cell parameters from 4229 reflections

$\theta = 2.4$ – $25.3$ °

$\mu = 0.59$  mm<sup>-1</sup>

$T = 297$  (2) K

Block, yellow

$0.35 \times 0.27 \times 0.21$  mm

*Data collection*

Bruker SMART APEX diffractometer	5178 independent reflections
Radiation source: fine-focus sealed tube	4536 reflections with $I > 2\sigma(I)$
Monochromator: graphite	$R_{\text{int}} = 0.029$
$T = 297(2)$ K	$\theta_{\text{max}} = 26.4^\circ$
$\varphi$ and $\omega$ scans	$\theta_{\text{min}} = 1.6^\circ$
Absorption correction: multi-scan (SADABS; Bruker, 2000)	$h = -11 \rightarrow 11$
$T_{\text{min}} = 0.819$ , $T_{\text{max}} = 0.886$	$k = -14 \rightarrow 14$
13712 measured reflections	$l = -16 \rightarrow 16$

*Refinement*

Refinement on $F^2$	Secondary atom site location: difference Fourier map
Least-squares matrix: full	Hydrogen site location: inferred from neighbouring sites
$R[F^2 > 2\sigma(F^2)] = 0.047$	H-atom parameters constrained
$wR(F^2) = 0.106$	$w = 1/[\sigma^2(F_o^2) + (0.0406P)^2 + 0.4205P]$
$S = 1.13$	where $P = (F_o^2 + 2F_c^2)/3$
5178 reflections	$(\Delta/\sigma)_{\text{max}} = 0.001$
280 parameters	$\Delta\rho_{\text{max}} = 0.40 \text{ e } \text{\AA}^{-3}$
Primary atom site location: structure-invariant direct methods	$\Delta\rho_{\text{min}} = -0.19 \text{ e } \text{\AA}^{-3}$
	Extinction correction: none

*Special details*

**Geometry.** All e.s.d.'s (except the e.s.d. in the dihedral angle between two l.s. planes) are estimated using the full covariance matrix. The cell e.s.d.'s are taken into account individually in the estimation of e.s.d.'s in distances, angles and torsion angles; correlations between e.s.d.'s in cell parameters are only used when they are defined by crystal symmetry. An approximate (isotropic) treatment of cell e.s.d.'s is used for estimating e.s.d.'s involving l.s. planes.

**Refinement.** Refinement of  $F^2$  against ALL reflections. The weighted  $R$ -factor  $wR$  and goodness of fit  $S$  are based on  $F^2$ , conventional  $R$ -factors  $R$  are based on  $F$ , with  $F$  set to zero for negative  $F^2$ . The threshold expression of  $F^2 > \sigma(F^2)$  is used only for calculating  $R$ -factors(gt) *etc.* and is not relevant to the choice of reflections for refinement.  $R$ -factors based on  $F^2$  are statistically about twice as large as those based on  $F$ , and  $R$ -factors based on ALL data will be even larger.

*Fractional atomic coordinates and isotropic or equivalent isotropic displacement parameters ( $\text{\AA}^2$ )*

	$x$	$y$	$z$	$U_{\text{iso}}^*/U_{\text{eq}}$
C1	0.2158 (3)	0.8016 (2)	1.01664 (19)	0.0410 (6)
C2	0.3749 (3)	0.8562 (3)	1.0317 (2)	0.0549 (7)
H2	0.4141	0.8641	0.9724	0.066*
C3	0.4758 (4)	0.8988 (3)	1.1337 (3)	0.0649 (8)
H3	0.5828	0.9348	1.1436	0.078*

## supplementary materials

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C4	0.4171 (4)	0.8878 (3)	1.2203 (3)	0.0689 (9)
H4	0.4848	0.9168	1.2894	0.083*
C5	0.2607 (4)	0.8349 (3)	1.2068 (2)	0.0710 (9)
H5	0.2225	0.8283	1.2665	0.085*
C6	0.1586 (3)	0.7909 (3)	1.1050 (2)	0.0561 (7)
H6	0.0520	0.7543	1.0960	0.067*
C7	-0.1095 (3)	0.7270 (2)	0.8809 (2)	0.0432 (6)
C8	-0.2183 (3)	0.6199 (3)	0.8876 (2)	0.0541 (7)
H8	-0.1877	0.5500	0.8980	0.065*
C9	-0.3720 (4)	0.6158 (4)	0.8789 (3)	0.0721 (10)
H9	-0.4449	0.5435	0.8832	0.087*
C10	-0.4163 (4)	0.7184 (5)	0.8640 (3)	0.0890 (12)
H10	-0.5203	0.7156	0.8568	0.107*
C11	-0.3101 (5)	0.8242 (4)	0.8595 (4)	0.0986 (13)
H11	-0.3412	0.8941	0.8513	0.118*
C12	-0.1564 (4)	0.8300 (3)	0.8671 (3)	0.0735 (9)
H12	-0.0848	0.9030	0.8629	0.088*
C13	0.3127 (3)	0.3345 (2)	0.5892 (2)	0.0398 (5)
C14	0.2557 (3)	0.3370 (2)	0.4795 (2)	0.0479 (6)
H14	0.1519	0.3359	0.4499	0.057*
C15	0.3525 (4)	0.3410 (3)	0.4141 (2)	0.0556 (7)
H15	0.3143	0.3436	0.3406	0.067*
C16	0.5055 (4)	0.3413 (3)	0.4577 (3)	0.0609 (8)
H16	0.5705	0.3430	0.4135	0.073*
C17	0.5620 (3)	0.3390 (3)	0.5657 (3)	0.0619 (8)
H17	0.6657	0.3395	0.5946	0.074*
C18	0.4675 (3)	0.3360 (2)	0.6329 (2)	0.0514 (7)
H18	0.5072	0.3349	0.7065	0.062*
C19	0.0085 (3)	0.2000 (2)	0.61159 (19)	0.0391 (5)
C20	0.0029 (3)	0.0881 (2)	0.6477 (2)	0.0553 (7)
H20	0.0863	0.0805	0.7028	0.066*
C21	-0.1260 (4)	-0.0116 (3)	0.6021 (3)	0.0679 (9)
H21	-0.1291	-0.0866	0.6261	0.082*
C22	-0.2495 (4)	-0.0006 (3)	0.5214 (3)	0.0663 (9)
H22	-0.3370	-0.0679	0.4914	0.080*
C23	-0.2445 (3)	0.1094 (3)	0.4849 (2)	0.0590 (8)
H23	-0.3284	0.1163	0.4297	0.071*
C24	-0.1155 (3)	0.2102 (2)	0.5295 (2)	0.0468 (6)
H24	-0.1123	0.2846	0.5042	0.056*
P1	0.09047 (8)	0.73820 (6)	0.88233 (5)	0.03962 (16)
P2	0.18502 (8)	0.32528 (6)	0.67495 (5)	0.03861 (16)
S1	0.11328 (9)	0.56101 (6)	0.88821 (5)	0.04965 (18)
S2	0.15246 (9)	0.81659 (7)	0.76545 (6)	0.0581 (2)
S3	0.27532 (9)	0.31611 (7)	0.82540 (5)	0.0562 (2)
S4	0.12601 (8)	0.48837 (6)	0.64407 (5)	0.04720 (18)
S5	-0.02304 (8)	0.48041 (6)	0.73690 (6)	0.04856 (18)

Atomic displacement parameters ( $\text{\AA}^2$ )

	$U^{11}$	$U^{22}$	$U^{33}$	$U^{12}$	$U^{13}$	$U^{23}$
C1	0.0431 (14)	0.0406 (13)	0.0377 (13)	0.0128 (11)	0.0096 (11)	0.0023 (10)
C2	0.0462 (15)	0.0608 (18)	0.0556 (17)	0.0146 (13)	0.0134 (13)	0.0027 (14)
C3	0.0443 (16)	0.064 (2)	0.072 (2)	0.0126 (14)	-0.0026 (15)	0.0023 (16)
C4	0.076 (2)	0.0575 (19)	0.0514 (18)	0.0143 (17)	-0.0112 (16)	-0.0003 (14)
C5	0.089 (3)	0.074 (2)	0.0389 (16)	0.0113 (19)	0.0159 (16)	0.0006 (15)
C6	0.0559 (17)	0.0611 (18)	0.0435 (15)	0.0060 (14)	0.0147 (13)	0.0017 (13)
C7	0.0407 (13)	0.0489 (15)	0.0380 (13)	0.0129 (11)	0.0096 (11)	-0.0008 (11)
C8	0.0533 (16)	0.0624 (18)	0.0486 (16)	0.0170 (14)	0.0183 (13)	0.0084 (13)
C9	0.0529 (18)	0.094 (3)	0.061 (2)	0.0040 (18)	0.0249 (16)	-0.0066 (18)
C10	0.055 (2)	0.131 (4)	0.088 (3)	0.040 (2)	0.0229 (19)	-0.009 (3)
C11	0.085 (3)	0.093 (3)	0.137 (4)	0.056 (3)	0.034 (3)	0.007 (3)
C12	0.067 (2)	0.0587 (19)	0.103 (3)	0.0281 (17)	0.0295 (19)	0.0056 (18)
C13	0.0425 (13)	0.0311 (12)	0.0428 (14)	0.0077 (10)	0.0113 (11)	0.0010 (10)
C14	0.0475 (15)	0.0504 (15)	0.0453 (15)	0.0136 (12)	0.0138 (12)	0.0076 (12)
C15	0.0659 (19)	0.0535 (17)	0.0465 (16)	0.0101 (14)	0.0232 (14)	0.0062 (13)
C16	0.0594 (19)	0.0526 (17)	0.072 (2)	0.0048 (14)	0.0353 (17)	-0.0010 (15)
C17	0.0399 (15)	0.0612 (19)	0.078 (2)	0.0090 (13)	0.0144 (15)	-0.0046 (16)
C18	0.0474 (15)	0.0493 (16)	0.0510 (16)	0.0111 (13)	0.0081 (13)	-0.0013 (12)
C19	0.0442 (13)	0.0345 (12)	0.0406 (13)	0.0104 (10)	0.0173 (11)	0.0027 (10)
C20	0.0587 (17)	0.0399 (15)	0.0614 (18)	0.0117 (13)	0.0107 (14)	0.0075 (13)
C21	0.083 (2)	0.0376 (16)	0.072 (2)	0.0061 (15)	0.0157 (18)	0.0063 (14)
C22	0.068 (2)	0.0485 (17)	0.064 (2)	-0.0070 (15)	0.0186 (17)	-0.0096 (15)
C23	0.0486 (16)	0.0630 (19)	0.0515 (17)	0.0052 (14)	0.0052 (13)	-0.0032 (14)
C24	0.0480 (15)	0.0451 (15)	0.0454 (15)	0.0117 (12)	0.0129 (12)	0.0061 (12)
P1	0.0411 (4)	0.0412 (4)	0.0361 (3)	0.0117 (3)	0.0113 (3)	0.0044 (3)
P2	0.0440 (4)	0.0349 (3)	0.0359 (3)	0.0113 (3)	0.0105 (3)	0.0039 (3)
S1	0.0608 (4)	0.0496 (4)	0.0432 (4)	0.0259 (3)	0.0128 (3)	0.0062 (3)
S2	0.0642 (5)	0.0640 (5)	0.0451 (4)	0.0116 (4)	0.0206 (3)	0.0161 (3)
S3	0.0658 (5)	0.0598 (4)	0.0378 (4)	0.0173 (4)	0.0073 (3)	0.0082 (3)
S4	0.0632 (4)	0.0346 (3)	0.0467 (4)	0.0161 (3)	0.0191 (3)	0.0064 (3)
S5	0.0403 (3)	0.0436 (4)	0.0562 (4)	0.0096 (3)	0.0097 (3)	-0.0040 (3)

Geometric parameters ( $\text{\AA}$ ,  $^\circ$ )

C1—C6	1.380 (4)	C14—C15	1.378 (4)
C1—C2	1.384 (4)	C14—H14	0.9300
C1—P1	1.805 (2)	C15—C16	1.375 (4)
C2—C3	1.377 (4)	C15—H15	0.9300
C2—H2	0.9300	C16—C17	1.365 (4)
C3—C4	1.367 (5)	C16—H16	0.9300
C3—H3	0.9300	C17—C18	1.382 (4)
C4—C5	1.361 (5)	C17—H17	0.9300
C4—H4	0.9300	C18—H18	0.9300
C5—C6	1.380 (4)	C19—C24	1.379 (3)
C5—H5	0.9300	C19—C20	1.387 (3)

## supplementary materials

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C6—H6	0.9300	C19—P2	1.816 (2)
C7—C12	1.379 (4)	C20—C21	1.376 (4)
C7—C8	1.383 (4)	C20—H20	0.9300
C7—P1	1.809 (3)	C21—C22	1.369 (4)
C8—C9	1.381 (4)	C21—H21	0.9300
C8—H8	0.9300	C22—C23	1.369 (4)
C9—C10	1.362 (5)	C22—H22	0.9300
C9—H9	0.9300	C23—C24	1.383 (4)
C10—C11	1.353 (6)	C23—H23	0.9300
C10—H10	0.9300	C24—H24	0.9300
C11—C12	1.378 (5)	P1—S2	1.9351 (9)
C11—H11	0.9300	P1—S1	2.1171 (9)
C12—H12	0.9300	P2—S3	1.9303 (9)
C13—C14	1.385 (3)	P2—S4	2.1282 (9)
C13—C18	1.386 (4)	S1—S5	2.0440 (10)
C13—P2	1.808 (2)	S4—S5	2.0407 (10)
C6—C1—C2	119.4 (2)	C14—C15—H15	120.1
C6—C1—P1	121.7 (2)	C17—C16—C15	120.0 (3)
C2—C1—P1	118.9 (2)	C17—C16—H16	120.0
C3—C2—C1	120.5 (3)	C15—C16—H16	120.0
C3—C2—H2	119.8	C16—C17—C18	121.0 (3)
C1—C2—H2	119.8	C16—C17—H17	119.5
C4—C3—C2	119.4 (3)	C18—C17—H17	119.5
C4—C3—H3	120.3	C17—C18—C13	119.1 (3)
C2—C3—H3	120.3	C17—C18—H18	120.4
C5—C4—C3	120.8 (3)	C13—C18—H18	120.4
C5—C4—H4	119.6	C24—C19—C20	119.5 (2)
C3—C4—H4	119.6	C24—C19—P2	123.69 (19)
C4—C5—C6	120.4 (3)	C20—C19—P2	116.8 (2)
C4—C5—H5	119.8	C21—C20—C19	120.1 (3)
C6—C5—H5	119.8	C21—C20—H20	120.0
C1—C6—C5	119.6 (3)	C19—C20—H20	120.0
C1—C6—H6	120.2	C22—C21—C20	120.2 (3)
C5—C6—H6	120.2	C22—C21—H21	119.9
C12—C7—C8	118.9 (3)	C20—C21—H21	119.9
C12—C7—P1	117.5 (2)	C23—C22—C21	120.1 (3)
C8—C7—P1	123.5 (2)	C23—C22—H22	119.9
C9—C8—C7	120.5 (3)	C21—C22—H22	119.9
C9—C8—H8	119.8	C22—C23—C24	120.4 (3)
C7—C8—H8	119.8	C22—C23—H23	119.8
C10—C9—C8	119.6 (3)	C24—C23—H23	119.8
C10—C9—H9	120.2	C19—C24—C23	119.8 (3)
C8—C9—H9	120.2	C19—C24—H24	120.1
C11—C10—C9	120.4 (3)	C23—C24—H24	120.1
C11—C10—H10	119.8	C1—P1—C7	107.49 (11)
C9—C10—H10	119.8	C1—P1—S2	116.30 (9)
C10—C11—C12	120.9 (4)	C7—P1—S2	113.64 (9)
C10—C11—H11	119.5	C1—P1—S1	96.94 (8)
C12—C11—H11	119.5	C7—P1—S1	107.14 (9)

C11—C12—C7	119.6 (3)	S2—P1—S1	113.77 (4)
C11—C12—H12	120.2	C13—P2—C19	106.48 (11)
C7—C12—H12	120.2	C13—P2—S3	116.70 (9)
C14—C13—C18	119.7 (2)	C19—P2—S3	113.09 (8)
C14—C13—P2	120.46 (19)	C13—P2—S4	97.91 (8)
C18—C13—P2	119.8 (2)	C19—P2—S4	106.85 (8)
C15—C14—C13	120.2 (3)	S3—P2—S4	114.34 (4)
C15—C14—H14	119.9	S5—S1—P1	100.37 (4)
C13—C14—H14	119.9	S5—S4—P2	99.52 (4)
C16—C15—C14	119.9 (3)	S4—S5—S1	106.77 (4)
C16—C15—H15	120.1		
C6—C1—C2—C3	0.4 (4)	C6—C1—P1—C7	24.6 (3)
P1—C1—C2—C3	-176.0 (2)	C2—C1—P1—C7	-159.0 (2)
C1—C2—C3—C4	-0.6 (5)	C6—C1—P1—S2	153.3 (2)
C2—C3—C4—C5	0.3 (5)	C2—C1—P1—S2	-30.4 (2)
C3—C4—C5—C6	0.3 (5)	C6—C1—P1—S1	-85.9 (2)
C2—C1—C6—C5	0.1 (4)	C2—C1—P1—S1	90.5 (2)
P1—C1—C6—C5	176.4 (2)	C12—C7—P1—C1	82.6 (2)
C4—C5—C6—C1	-0.4 (5)	C8—C7—P1—C1	-100.7 (2)
C12—C7—C8—C9	1.0 (4)	C12—C7—P1—S2	-47.6 (3)
P1—C7—C8—C9	-175.7 (2)	C8—C7—P1—S2	129.2 (2)
C7—C8—C9—C10	-0.2 (5)	C12—C7—P1—S1	-174.1 (2)
C8—C9—C10—C11	-1.2 (6)	C8—C7—P1—S1	2.6 (2)
C9—C10—C11—C12	1.7 (7)	C14—C13—P2—C19	50.3 (2)
C10—C11—C12—C7	-0.9 (6)	C18—C13—P2—C19	-128.0 (2)
C8—C7—C12—C11	-0.4 (5)	C14—C13—P2—S3	177.66 (17)
P1—C7—C12—C11	176.5 (3)	C18—C13—P2—S3	-0.6 (2)
C18—C13—C14—C15	-0.1 (4)	C14—C13—P2—S4	-59.9 (2)
P2—C13—C14—C15	-178.4 (2)	C18—C13—P2—S4	121.75 (19)
C13—C14—C15—C16	0.7 (4)	C24—C19—P2—C13	-82.9 (2)
C14—C15—C16—C17	-0.8 (4)	C20—C19—P2—C13	95.5 (2)
C15—C16—C17—C18	0.2 (5)	C24—C19—P2—S3	147.60 (19)
C16—C17—C18—C13	0.4 (4)	C20—C19—P2—S3	-34.0 (2)
C14—C13—C18—C17	-0.4 (4)	C24—C19—P2—S4	20.9 (2)
P2—C13—C18—C17	177.9 (2)	C20—C19—P2—S4	-160.63 (19)
C24—C19—C20—C21	-0.3 (4)	C1—P1—S1—S5	-179.08 (9)
P2—C19—C20—C21	-178.8 (2)	C7—P1—S1—S5	70.17 (9)
C19—C20—C21—C22	-0.4 (5)	S2—P1—S1—S5	-56.30 (5)
C20—C21—C22—C23	0.8 (5)	C13—P2—S4—S5	178.76 (8)
C21—C22—C23—C24	-0.5 (5)	C19—P2—S4—S5	68.80 (9)
C20—C19—C24—C23	0.7 (4)	S3—P2—S4—S5	-57.13 (5)
P2—C19—C24—C23	179.1 (2)	P2—S4—S5—S1	87.43 (4)
C22—C23—C24—C19	-0.3 (4)	P1—S1—S5—S4	96.84 (4)

Hydrogen-bond geometry ( $\text{\AA}$ ,  $^\circ$ )

$D-H\cdots A$	$D-H$	$H\cdots A$	$D\cdots A$	$D-H\cdots A$
C17—H17 $\cdots$ S5 <sup>i</sup>	0.93	2.94	3.737 (3)	145

Symmetry codes: (i)  $x+1, y, z$ .

Fig. 1

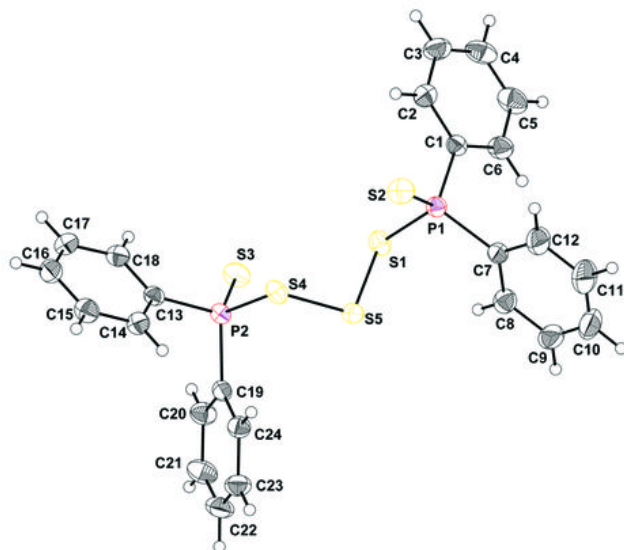


Fig. 2

