

sp Carbon Chains Surrounded by sp³ Carbon Double Helices: Coordination-Driven Self-Assembly of Wirelike $Pt(C=C)_nPt$ Moieties That Are Spanned by Two $P(CH_2)_mP$ Linkages

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Abstract: Reactions of trans, trans- $(C_6F_5)(p-tol_3P)_2Pt(C\equiv C)_4Pt(Pp-tol_3)_2(C_6F_5)$ and diphosphines $Ar_2P(CH_2)_m$ -PAr₂ yield trans, trans-(C₆F₅)(Ar₂ $\overset{P}{P}$ (CH₂)_mPAr₂)Pt(C=C)₄ $\overset{P}{P}$ t(Ar₂P(CH₂)_mPAr₂)(C₆F₅), in which the platinum atoms are spanned via an sp and two sp³ carbon chains (Ar/m = 3, Ph/14, 87%; 4, p-tol/14, 91%; 5, $p-C_6H_4-t-Bu/14$, 77%; 7, Ph/10, 80%; 8, Ph/11, 80%; 9, Ph/12, 36%; only oligomers form for m > 14). Crystal structures of 3-5 show that the sp³ chains adopt chiral double-helical conformations that shield the sp chain at approximately the van der Waals distance, with both enantiomers in the unit cell. The platinum square planes define angles of 196.6°-189.9° or more than a half twist. Crystal structures of 7–9, which have shorter sp³ chains, exhibit nonhelical conformations. Reaction of the corresponding Pt- $(C \equiv C)_6$ Pt complex and Ph₂P(CH₂)₁₈PPh₂ gives an analogous adduct (27%). The crystal structure shows two independent molecules, one helical and the other not. Low-temperature NMR data suggest that the enantiomeric helical conformations of 3-5 rapidly interconvert in solution. Cyclic voltammograms of 3-5 show more reversible oxidations than model compounds lacking bridging sp³ chains. These are the only double-helical molecules that do not feature bonding interactions between the helix strands, or covalent bonds to templates dispersed throughout the strands, or any type of encoding. The driving force for helix formation is analyzed.

Introduction

Of the many types of helical molecules,^{1,2} double helices have captured the greatest attention.²⁻⁸ The structure elucidation of the first compound recognized as a double helix, DNA,³ has

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been immortalized in books and plays. Since this structural motif intrinsically shields interior functional domains, it represents an astute evolutionary choice for carrying the genetic code. However, we are not aware of any efforts to employ double helices as "protecting groups" for abiological functional arrays.

To our knowledge, all double-helical molecules feature either (1) bonding interactions between the helix strands, such as hydrogen bonds, or (2) covalent bonds to metal templates dispersed throughout the strands. Factors that promote or enforce complementarity, e.g., the programming⁹ or "encoding of specific structural and conformational information",4b are thought to be required. We wondered whether it would be possible to design double-helical molecules that lack these conventional driving forces. In the extreme, might certain substances appear to yield double helices because "they felt like it"?

Toward this end, we sought rodlike building blocks, represented by A in Scheme 1. Such moieties are seeing extensive use in the development of molecular-scale devices.¹⁰ The most

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fundamental and radially compact would be a chain of sp hybridized carbons.¹¹ An atom that supports at least two additional bonds would then be grafted onto each terminus. Transition metals are obvious candidates,¹² as exemplified by the square planar endgroups **B** in Scheme 1. These could anchor two long flexible chains, containing for example polymethylene $((CH_2)_m)$ or polydifluoromethylene $((CF_2)_m)$ domains.

Conformation is then an issue. Will the flexible chains twist about the rod to give double helices as in the case of C (Scheme 1)? Alternatively, will they preferentially adopt conformations with "holes" (D) or "fingers" (E)? Since monocyclic crown ethers and related macrocycles seldom exhibit conformations with holes,¹³ we discounted possibility **D**. However, **E** was viewed as a plausible alternative. Regardless, all of these conformations sterically shield the rodlike segment to varying degrees.

Such assemblies would have potential applications. For example, platinum(II)-capped polyynes $Pt(C \equiv C)_n Pt$ (polyynediyls) with as many as 28 sp carbon atoms have been isolated,¹⁴ corresponding to a metal-metal separation of 3.9 nm. However, all polyynes become increasingly more labile with sp carbon chain length. Since the decomposition pathways are believed to be bimolecular, systems of the type C-E should exhibit

enhanced kinetic stabilities. New length regimes would likely be accessible.

Also, molecules in which unsaturated ligands span two transition metals can often be generated in more than one oxidation state, each of which can undergo interesting chargetransfer phenomena.¹⁵ However, the stabilities of oxidized or reduced species derived from metal-capped polyynes dramatically decrease with sp chain length.¹⁶ No examples with more than eight carbon atoms have been isolated,¹⁷ hampering certain applications of these wirelike species. The flexible chains in C-E should again serve a stabilizing function, which for oddelectron systems would be conceptually analogous to the insulation about household electrical wire. Others have reported complementary approaches to shielding unsaturated assemblies that connect two electroactive groups.18-21

Hence, we set out to investigate whether $Pt(C \equiv C)_n Pt$ moieties could serve as core units for double-helical assemblies C. One of several underlying considerations involved the wellestablished redox chemistry of monoplatinum(II) complexes. In most but not all cases, electrochemical oxidations to platinum-(III) are only partially reversible.²² Thus, it would be simple to qualitatively detect a stabilizing effect of the flexible chains upon the corresponding radical cations.

As summarized in Scheme 2, two synthetic strategies were considered. The first involved reactions of preformed $Pt(C \equiv$ C)_nPt species (F) with $bis(\alpha, \omega$ -diarylphosphino)alkanes, in which the phosphorus atoms are separated by flexible polymethylene chains. This can be viewed as a coordination-driven "self-assembly" process with potential for operationally simple nanodevice synthesis. In terms of precedent, Sünkel had shown that when the chloride-substituted diplatinum ethynediyl complex Cl(Ph₃P)₂PtC=CPt(PPh₃)₂Cl was treated with excess PEt₃, all of the PPh₃ ligands could be replaced.²³

The second strategy involved the synthesis of platinum chloride complexes with ω -functionalized *n*-alkyl monophosphine ligands (G). The sp carbon chain would then be introduced, and the phosphine ligands on opposite platinum termini coupled, generating bridging diphosphines within the metal coordination sphere. In both approaches, there would be the possibility of byproducts in which the diphosphine ligands span trans positions of the same platinum atom, as shown in H'.

In this paper, we report successful syntheses of the target molecules C by the first strategy. This simple route allows rapid access to a number of complexes but also has limitations as described below. In the following paper, we report successful

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applications of the second strategy, using alkene metathesis as the means of ligand coupling.²⁴ This longer method is more reliable, but the overall yields are somewhat lower. Portions of this work have been communicated,²⁵ and additional details are described elsewhere.²⁶

Results

1. Syntheses of Target Molecules: 8 sp and 14 sp³ Carbons. Reactions of the *p*-tolyl-substituted diplatinum octatetraynediyl complex *trans,trans-(p*-tol)(Ph₃P)₂Pt(C \equiv C)₄Pt-(PPh₃)₂(*p*-tol) (1; Scheme 3)^{14b} were investigated first. An NMR tube was charged with 1 and the diphosphine Ph₂P(CH₂)₁₂-PPh₂,^{27,28} which features 12 sp³ carbon atoms. Displacement of the PPh₃ ligands proceeded rapidly below room temperature, generating one platinum-containing product as assayed by ³¹P NMR and TLC. However, when solutions were concentrated, insoluble, presumably oligomeric or polymeric materials formed. Only small portions could be induced to redissolve.

Comparable results were obtained with the longer-chain diphosphine Ph₂P(CH₂)₁₆PPh₂.²⁸ Extensive attempts to crystallize the major products from dilute solutions were unsuccessful. Oligomeric structures are rigorously established for similar Pt-

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insoluble oligomers and/or polymers

 $(C \equiv C)_n$ Pt-derived materials below. Monoplatinum complexes of other types of long-chain chelating diphosphines have also been observed to oligomerize.^{28b}

In an attempt to avoid the generation of oligomers, two strategy modifications were implemented: (a) platinum endgroups with enhanced Lewis acidities were targeted, with the idea that the dissociation of one terminus of the diphosphine ligand in **H** (Scheme 2) would be retarded – a logical initial event in oligomerization; (b) since this might in turn render substitution reactions of **F** (Scheme 2) more difficult, approaches involving functionalized phosphines (**G**) were investigated. In this way, no platinum—phosphorus bond-making or -breaking reactions are required after the initial substitution. As described in the following paper,²⁴ this sequence proved successful with pentafluorophenyl-substituted platinum endgroups and provided the first syntheses of the target molecules. However, an obvious question remained: would modification (a) alone have sufficed?

As shown in Scheme 4, the answer is in many cases yes. A CH₂Cl₂ solution of the diphosphine Ph₂P(CH₂)₁₄PPh₂ (2.6 equiv),²⁷ which features 14 sp³ carbon atoms, was added to a moderately dilute CH₂Cl₂ solution of the pentafluorophenyl-substituted diplatinum complex *trans,trans*-(C₆F₅)(*p*-tol₃P)₂-Pt(C=C)₄Pt(P*p*-tol₃)₂(C₆F₅) (**2**; 0.0025 M).^{14a} Workup gave the target complex *trans,trans*-(C₆F₅)(Ph₂P(CH₂)₁₄PPh₂)Pt-(C=C)₄Pt(Ph₂P(CH₂)₁₄PPh₂)(C₆F₅) (**3**) as an analytically pure yellow powder in 87% yield. The structure was verified crystallographically, as described below. Analogous reactions

crystallographically, as described below. Analogous reactions with similar phenyl-substituted diphosphines, $(p-RC_6H_4)_2P-(CH_2)_{14}P(p-C_6H_4R)_2$ (R = CH₃, *t*-Bu),³⁰ gave the corresponding complexes **4** and **5** in 77–91% yields.

Complexes **3–5**, and all other compounds isolated below, were stable in air as solids for extended periods. They were characterized by IR, NMR (¹H, ¹³C, ³¹P), and UV–visible

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⁽²⁹⁾ A similar reaction of the Pt(C≡C)₈Pt homologue^{14a} of 2 and 10 with Ph₂P(CH₂)₂₈PPh₂ gave exclusively oligomeric products.

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spectroscopy, and mass spectrometry, as summarized in the Experimental Section. The IR and UV-visible spectra were quite similar to those of the precursor 2^{14a} and essentially identical with those of the model compound 6.30 The latter is a



better electronic model for **3**, lacking only bridging sp³ carbon chains. The PtC \equiv C¹³C NMR signals of **3**-**5** (98.6/94.2, 99.2/ 94.1, 99.1/94.4 ppm) differed slightly from those of 2 (100.6/ 96.7) but were nearly identical with those of 6 (100.0/94.2 ppm). The PCH₂CH₂CH₂CH₂¹³C NMR signals were also very close to those of 6 (28.1-28.3/25.5-25.7/30.6-30.7 vs 28.2/25.5/31.3 ppm). The mass spectra exhibited intense molecular ions (100%). DSC data for 3 and 4 showed endotherms at 244 and 213 °C (T_e ; presumably phase transitions) with exotherms beginning at higher temperatures (271 and 260 °C, T_i) that were accompanied by mass loss (TGA data).

2. Crystallography: 8 sp and 14 sp³ Carbons. Complexes 3, 4, and 5 readily crystallized. The first was obtained as both benzene and toluene sesquisolvates (3. (benzene)1.5, 3. (toluene)_{1.5}), and the last, as two different toluene solvates (5. $(toluene)_{5,5}$ and **5**· $(toluene)_2$) or pseudopolymorphs.³³ The crystal structures were solved as summarized in the Supporting Information. Those of $3 \cdot (benzene)_{1.5}$ and $3 \cdot (toluene)_{1.5}$ were very similar. Key metrical parameters are given in Tables 1-3.

The structures of $3 \cdot (toluene)_{1.5}$, 4, and $5 \cdot (toluene)_{5.5}$ are compared in Figure 1 (I-K). In all three compounds, the sp³ carbon chains exhibit a striking double-helical conformation. Helices are intrinsically chiral, and both enantiomers are present in the crystal lattice. With $3 \cdot (benzene)_{1,5}$ and $3 \cdot (toluene)_{1,5}$, the twisting is less "uniform" than that in the other complexes, a feature that can be analyzed via torsion angle patterns (Table 3) as described below. The idealized structures (i.e., as represented in Scheme 4) belong to the point group D_2 .

The bond lengths, bond angles, and slight sp chain curvatures in $3 \cdot (toluene)_{1.5}$, 4, and $5 \cdot (toluene)_{5.5}$ are unexceptional. They are very close to those of 2,14a which lacks bridging diphosphines, as well as higher and lower homologues.14a,25b Considerably greater sp chain curvature has been observed,¹¹ a normal phenomenon that has various measures, such as the average bond angles in Table 1 or the difference between the platinumplatinum distances and the sum of the intervening bond lengths. These parameters and others have been analyzed for all $Pt(C \equiv$ $C)_n$ Pt complexes in detail elsewhere.¹¹

The angles defined by the square planes of the platinum termini are of particular interest. Of the measures summarized in Table 1, we favor the planes defined by the P-Pt-P linkage on one endgroup and the platinum on the other. These emphasize the twist defined by the P-Pt-P linkages and are less effected by various geometric nonidealities. The values in $3 \cdot (\text{benzene})_{1.5}$,

Scheme 4. Reactions of the Pt(C≡C)₄Pt Complex 2 with Bis(α, ω -diarylphosphino)alkanes



 $3 \cdot (toluene)_{1.5}$, 4, and $5 \cdot (toluene)_{5.5}$ range from 189.8° to 196.6°, more than a half twist. This feature is emphasized in the partial structure of 3·(toluene)_{1.5} shown in Figure 2 (L). Helical objects are also characterized by a degree of helicity and a pitch (Table 1).^{1,31} The former is derived by dividing the preceding angles by 360°, and the latter represents the hypothetical distance at which a full turn would be reached.

The crystal structure of precursor 2 has been previously reported,14a and two views with all atoms at van der Waals radii are illustrated in Figure 3 (M, N; parallel and perpendicular to the planes of the C_6F_5 rings). The sp carbon chain is clearly accessible to external reagents. Figure 3 also gives analogous representations of $3 \cdot (toluene)_{1.5}$ (Q, R), as well as structures from which the phenyl rings and hydrogen atoms are omitted (O, P). For reference, comparable views of 4 are depicted (S, **T**); **5**·(toluene)_{5.5} is very similar. These structures nicely illustrate the extensive shielding of the sp chains in 3-5, although **R** and T reveal that a line-of-sight through a protecting rim of peripheral groups remains.

Figure 3 shows that the sp and sp³ carbon chains of $3 \cdot (toluene)_{1,5}$ and 4 entwine with a spacing slightly greater than their van der Waals radii (1.78 and 1.70 Å).32 A cylindrical helix is characterized by a radius, which can be approximated by calculating the average distances of the sp³ carbon atoms from the platinum-platinum vectors. As summarized in Table 1, these span the narrow range from 3.977 to 4.053 Å. A higher value indicates a "looser" helix. Accordingly, there is an inverse correlation with the degree of helicity. The packing diagrams show essentially parallel sp chains, the spacing between which

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Table 1. Key Crystallographic Distances [Å] and Angles [deg] for Helical Complexes

complex	3. (benzene) _{1.5} ^a	3-(toluene) _{1.5}	4	5·(toluene) _{5.5}	11·(CH ₂ Cl ₂) _{0.67} ^b
Pt•••Pt	12.775(8)	12.7653(14)	12.6984(5)	12.8867(3)	17.9897(8)
sum of bond lengths,	12.956	12.914	12.890	12.896	18.023
Pt1 to Pt2					
Pt1-C1	2.003(3)	1.987(4)	1.991(8)	1.988(4)	1.962(13)
C1-C2	1.207(5)	1.215(7)	1.211(9)	1.205(6)	1.207(19)
C2-C3	1.368(5)	1.365(7)	1.345(10)	1.369(6)	1.38(2)
C3-C4	1.207(5)	1.214(7)	1.201(9)	1.206(6)	1.23(2)
C4-C5	1.354(5)	1.360(7)	1.363(10)	1.360(6)	1.37(2)
C5-C6	1.216(5)	1.207(7)	1.198(9)	1.209(6)	1.16(2)
C6-C7	1.368(5)	1.360(7)	1.372(10)	1.356(6)	1.36(2)
C7-C8	1.209(5)	1.212(6)	1.222(9)	1.220(6)	1.22(2)
C8-C9	-	-	-	-	1.34(2)
C9-C10	-	-	-	-	1.21(2)
C10-C11	-	-	-	-	1.41(2)
C11-C12	-	-	-	-	1.176(19)
C8-Pt2 or C12-Pt2	1.994(3)	1.994(4)	1.987(7)	1.983(4)	1.998(15)
Pt1-C1-C2	178.0(3)	175.9(5)	178.8(6)	179.3(4)	179.0(14)
C1-C2-C3	177.5(4)	178.9(6)	176.4(8)	178.3(5)	174.7(18)
C2-C3-C4	177.3(4)	178.4(6)	176.0(8)	179.9(7)	178(2)
C3-C4-C5	176.9(5)	178.2(6)	175.3(8)	178.5(6)	177(2)
C4-C5-C6	178.5(5)	177.0(6)	176.0(8)	177.9(6)	174(2)
C5-C6-C7	176.3(4)	177.3(6)	177.9(8)	178.5(5)	174(2)
C6-C7-C8	174.1(4)	172.8(5)	178.2(8)	179.1(5)	178(2)
C7-C8-C9	-	-	-	-	177(3)
C8-C9-C10	-	-	-	-	180(3)
C9-C10-C11	-	-	-	-	176.9(18)
C7-C8-Pt2 or	171.7(3)	171.6(4)	171.5(6)	179.3(4)	177.8(15)
C11-C12-Pt2					
av $Pt-C_{sp}-C_{sp}$	174.9	173.8	175.2	179.3	178.4
av $C_{sp} - C_{sp} - C_{sp}$	176.8	177.1	176.6	178.7	176.6
av sp/sp ³ distance ^c	4.053	3.977	4.042	3.988	3.868
av π stacking ^d	3.896	3.755	3.772	3.704	3.907
P-Pt-P/Pt angle ^e	196.6	196.5	189.9	193.3	163.2
$Pt+P+P+C_i+C1$ angle ^e	198.1	200.8	188.6	193.5	163.9
helix pitch ^f	23.72	23.66	24.45	24.02	39.76
degree of helicity	54.6%	54.6%	52.7%	53.7%	45.3%
$C_{sp}-C_{sp}^{g}$	8.071	7.974	9.339	11.959	9.028
fractional offset ^h	0.55	0.38	0.49	0.44	0.44

^{*a*} In order to better compare the two similar noncentrosymmetric solvates of **3**, the numbering of the PtC₈Pt segment in **3**·(benzene)_{1.5} has been reversed in this table. ^{*b*} Data for the nonhelical molecule in the unit cell is given in Table 2. ^{*c*} Average distance from every CH₂ group to the Pt–Pt vector. ^{*d*} Distance between midpoints of the C₆F₅ and C₆H₄R rings. ^{*e*} Angle between planes defined by these atoms on each endgroup. ^{*f*} The hypothetical distance at which a full helix turn is reached. ^{*s*} The shortest carbon–carbon distance between parallel sp chains in the crystal lattice. ^{*h*} Of parallel sp chains: see ref 11.

increases from 7.97 to 11.96 Å as the sizes of the p-aryl substituents increases (Table 1).

In contrast to the preceding four crystal structures, that of the pseudopolymorph $5 \cdot (toluene)_2$ exhibited a nonhelical conformation. As shown in Figure 4, the sp³ carbon chains define loops with respect to the sp chain. However, when atoms are viewed at van der Waals radii, there are no "holes" as in the case of **D** (Scheme 1). The average distance of the sp^3 carbon atoms from the platinum-platinum vector (Table 2) is somewhat greater than that in helical 5. (toluene)_{5.5} (4.064 vs 3.988 Å). Interestingly, the crystal density of $5 \cdot (toluene)_2$, a property sometimes used to gauge the relative stabilities of polymorphs,³³ is higher than that of 5·(toluene)_{5.5} (1.333 vs 1.256 Mg/m³); in contrast to the other structures, the lattice features two nonparallel sets of parallel sp chains. In any event, V and W (Figure 4) show that the sp chain in $5 \cdot (toluene)_2$ is only slightly less shielded than those in the helical complexes. Importantly, the structure requires that nonhelical conformations of 5 be populated in solution.

In all of the preceding structures, including that of **2**, aryl/ C_6F_5 /aryl stacking interactions are evident. These are found for

nearly all (C₆F₅)Pt adducts of aryl phosphines,^{14a,25b,34,35} and average distances are summarized in Tables 1 and 2. The origin of this phenomenon, which is general for highly fluorinated arenes, is well understood.³⁶ Although it plays a role in crystal packing, it is not believed to be a determinant of sp³ chain conformation in the title molecules. Note that the shortest average distance is found in nonhelical **5**·(toluene)₂.

3. Syntheses of Target Molecules: Other sp/sp³ Carbon Chain Lengths. Several questions follow from the preceding results: (a) How short an sp³ carbon chain can be introduced? (b) How long an sp³ carbon chain can be introduced? (c) Can analogous chemistry be effected with longer sp carbon chains? When the sp³/sp carbon ratio becomes too small, helical conformations should no longer be possible. However, such species would still possess sp carbon chains that are to various degrees sterically protected. Furthermore, when the sp³ and sp chains have similar numbers of atoms, they are constrained over

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 (35) Shima, T.; Bauer, E. B.; Hampel, F.; Gladysz, J. A. Dalton Trans. 2004, 1012

⁽³⁶⁾ Reichenbächer, K.; Süss, H. I.; Hullinger, J. Chem. Soc. Rev. 2005, 34, 20.

Table 2. Key Crystallographic Distances [Å] and Angles [deg] for Nonhelical Complexes

complex	5·(toluene) ₂ ^a	7·(toluene) ₄	7 • (CHCl ₃) ₂	9-(CH ₂ Cl ₂) _{2.5}	11 · (CH ₂ Cl ₂) _{0.67} ^{a,b}
Pt•••Pt	12.8820(21)	12.9120(3)	12.6417(31)	12.9580(2)	17.6087(13)
sum of bond lengths,	12.907	12.933	12.953	12.980	18.06
Pt1 to Pt2					
Pt1-C1	1.989(7)	2.014(6)	1.998(5)	2.009(4)	1.960(16)
C1-C2	1.220(9)	1.192(8)	1.219(6)	1.205(6)	1.23(2)
C2-C3	1.356(10)	1.366(8)	1.359(7)	1.379(6)	1.36(2)
C3-C4	1.202(10)	1.208(8)	1.224(7)	1.214(7)	1.19(2)
C4-C5	1.377(15)	1.361(9)	1.362(6)	1.365(7)	1.38(3)
C5-C6	-	1.215(8)	1.212(6)	1.222(7)	1.24(2)
C6-C7	-	1.368(8)	1.369(6)	1.365(6)	1.34(4)
C7-C8	-	1.210(8)	1.209(6)	1.212(6)	-
C8-Pt2	-	1.999(6)	2.001(4)	2.009(4)	-
Pt1-C1-C2	175.7(7)	174.0(6)	171.1(4)	176.0(5)	166.7(14)
C1-C2-C3	175.6(8)	178.2(8)	176.4(6)	178.5(6)	170.8(18)
C2-C3-C4	179.4(10)	178.3(7)	177.4(5)	178.9(6)	174.5(18)
C3-C4-C5	178.8(12)	178.5(9)	177.4(5)	179.6(6)	176(2)
C4-C5-C6	-	179.6(9)	177.0(5)	179.2(6)	178(2)
C5-C6-C7	-	178.5(8)	175.0(5)	178.6(5)	178(2)
C6-C7-C8	-	177.5(7)	173.5(5)	178.7(5)	-
C7-C8-Pt2	-	176.7(6)	169.4(4)	175.1(4)	-
av $Pt-C_{sp}-C_{sp}$	175.7	175.4	170.3	175.6	166.7
av $C_{sp} - \dot{C}_{sp} - \dot{C}_{sp}$	177.9	178.4	176.1	178.9	175.5
av sp/sp ³ distance ^c	4.064	3.932	3.989	4.090	4.157
av π stacking ^d	3.633	3.803	3.756	3.766	3.979
P-Pt-P/Pt angle ^e	0.0	0.2	5.1	12.9	0.0
$Pt+P+P+C_i+C1$ angle ^e	0.0	2.7	40.4	18.3	0.0
$C_{sp} - C_{sp}^{f}$	11.433	7.296	7.780	9.738	9.028
fractional offset ^g	0.88	0.41	0.10	0.32	-

^{*a*} This molecule exhibits an inversion center. ^{*b*} Data for the helical molecule in the unit cell is given in Table 1. ^{*c*} Average distance from every CH₂ group to the Pt–Pt vector. ^{*d*} Distance between midpoints of the C₆F₅ and C₆H₄R rings. ^{*e*} Angle between planes defined by these atoms on each endgroup. ^{*f*} The shortest carbon–carbon distance between parallel sp chains in the crystal lattice. ^{*g*} Of parallel sp chains: see ref 11.

Table 5. TOISION ANGLES [DEG] INVOLVING SP [®] CAIDON ALONIS IOLAL ₂ F(CH ₂) ₁₄ FAI ₂ -DIUGEU COL	npiexes
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complex	3·(benzene) _{1.5} ^a	3-(toluene) _{1.5} ^a	4 ^a	5-(toluene) _{5.5} ^a	5·(toluene) ₂
P-C1-C2-C3	-158.8/-80.8	-161.2/85.3	175.2/170.2	178.3/-179.4	177.9/-170.6
C1-C2-C3-C4	66.5/-71.9	64.6/-173.8	54.9/64.3	66.8/64.9	173.5/-68.8
C2-C3-C4-C5	171.4/173.3	173.2/166.5	62.1/62.1	63.9/64.4	$69.2/-89.8^{b}$
C3-C4-C5-C6	-174.9/176.9	179.0/-176.4	174.8/179.5	169.9/174.9	$176.8/-160.4^{b}$
C4-C5-C6-C7	-174.8/166.5	-176.3/170.1	-172.1/179.5	173.1/179.2	$-179.3/-82.1^{b}$
C5-C6-C7-C8	175.8/-172.6	-178.6/173.1	175.2/-177.0	-178.5/-178.7	$-160.9^{b}/148.0^{b}$
C6-C7-C8-C9	66.6/173.6	67.8/61.9	172.3/178.2	170.2/170.7	$-84.6^{b}/-84.6^{b}$
C7-C8-C9-C10	173.4/171.1	171.5/172.1	-175.8/176.2	179.6/178.7	$148.0^{b}/-160.9^{b}$
C8-C9-C10-C11	173.8/68.5	174.1/172.4	176.2/-176.4	178.7/177.7	$-82.1^{b}/-179.3$
C9-C10-C11-C12	173.2/166.0	174.2/177.9	-164.4/179.0	178.0/173.2	$-160.4^{b}/176.8$
C10-C11-C12-C13	58.0/172.0	57.0/165.3	68.3/61.9	65.9/62.7	$-89.8^{b}/69.2$
C11-C12-C13-C14	57.1/74.9	57.1/69.2	-176.1/55.3	63.3/63.4	-68.8/173.5
C12-C13-C14-P'	177.7/-158.6	-179.4/-150.9	169.4/171.6	-179.8/175.4	-170.6/177.9

^{*a*} The sp³ carbon chains in these structures are not equivalent. The torsion angles for both are presented starting from the same platinum atom. ^{*b*} There is considerable vibrational disorder associated with the atoms from which these values are derived.

their entire lengths to be in close proximity, providing a valuable reference state.

As shown in Scheme 4 (bottom), CH_2Cl_2 solutions of the shorter-chain diphosphines $Ph_2P(CH_2)_mPh_2$ ($m = 10, 11, 12)^{27,28}$ were added to CH_2Cl_2 solutions of **2** that were more dilute than those employed above. Workups gave the substitution products **7**, **8**, and **9** in 80%, 80%, and 36% yields, respectively. In the last reaction, which is closely related to the unsuccessful synthesis in Scheme 3, oligomeric byproducts were evident, and a higher dilution was necessary ([**2**] = 0.0005 M). The IR, NMR, UV-visible, MS, and DSC/TGA properties of **7**–**9** were very similar to those of **3**–**5**, including the ¹³C NMR chemical shifts of the sp and sp³ carbon atoms.

Analogous reactions of longer-chain diphosphines Ph₂P- $(CH_2)_mPPh_2$ (m = 16, 18, 32)^{27,28} with highly dilute CH₂Cl₂ solutions of **2** (0.0001, 0.0001, 0.000 075 M) gave only insoluble oligomers (85%, 94%, 61%, respectively).^{26a} The IR spectra of these materials were very similar. Mass spectra did not exhibit any ions for diplatinum complexes.

As shown in Scheme 5 (top), a CH₂Cl₂ solution of the diphosphine Ph₂P(CH₂)₁₈PPh₂ was added to an extremely dilute CH₂Cl₂ solution of the diplatinum dodecahexaynediyl or Pt-(C \equiv C)₆Pt complex **10** (0.000 03 M).^{14a} Workup and crystallization gave the substitution product **11**, which has sp and sp³ chains of 12 and 18 carbon atoms, in 27% yield. Oligomeric material was also isolated in 64% yield. As with **3–9**, the IR





Figure 1. Thermal ellipsoid plots (50% probability level) of the doublehelical $Pt(C \equiv C)_4 Pt$ complexes with hydrogen atoms and solvate molecules omitted: **I**, **3**·(toluene)_{1.5}; **J**, **4**; **K**, **5**·(toluene)_{5.5}.



Figure 2. View of **3**•(toluene)_{1.5} along the platinum—platinum axis with non-*ipso* aryl carbon atoms, hydrogen atoms, and solvate molecules omitted.

and UV-visible properties, and ¹³C NMR chemical shifts of the sp carbon atoms, were similar to those of the unbridged precursor. However, as summarized in Scheme 5 (bottom), reactions of **10** and longer-chain diphosphines gave almost exclusively oligomeric products.²⁹ In the cases of $Ph_2P(CH_2)_{19}$ -



Figure 3. Space filling representations of selected $Pt(C \equiv C)_4Pt$ complexes: **M**, **N**, views of **2**·(toluene) parallel and perpendicular to the planes of the C₆F₅ ligands; **O**, **P**, analogous views of **3**·(toluene)_{1.5} with phenyl rings, hydrogen atoms, and solvate molecules omitted; **Q**, **R**, analogous views of **3**·(toluene)_{1.5} but with phenyl rings and hydrogen atoms; **S**, **T**, analogous views of **4**.

 PPh_2 and $Ph_2P(CH_2)_{20}PPh_2$, mass spectra of the reaction mixtures showed ions for diplatinum complexes, and trace amounts could be extracted from the oligomers.

Experiments were conducted to confirm the oligomeric nature of these products. First, the material derived from **10** and Ph₂P(CH₂)₂₀PPh₂ gave an appropriate microanalysis. Second, it was treated with an excess of the monophosphine PEt₃. Trialkylphosphines are commonly more basic than diarylalkylphosphines. As shown in Scheme 5, substitution occurred to give the known complex *trans*,*trans*-(C₆F₅)(Et₃P)₂Pt(C \equiv C)₆Pt-(PEt₃)₂(C₆F₅) (**12**), which was isolated in 56% yield.^{14a} Analogous additions of PEt₃ to oligomers derived from **2** and Ph₂P(CH₂)_mPPh₂ (m = 16, 18, 32) similarly gave *trans*,*trans*-(C₆F₅)(Et₃P)₂Pt(C \equiv C)₄Pt(PEt₃)₂(C₆F₅).^{14a}

4. Crystallography: Other sp/sp³ Carbon Chain Lengths. Complexes 7, 9, and 11 could also be crystallized. In the case of 7, two pseudopolymorphs, $7 \cdot (toluene)_4$ and $7 \cdot (CHCl_3)_2$, were obtained. Their crystal structures, and those of $9 \cdot (CH_2Cl_2)_{2.5}$ and $11 \cdot (CH_2Cl_2)_{0.67}$, were determined as summarized in the Supporting Information. Portions of the sp³ carbon chains of $7 \cdot (toluene)_4$ and $9 \cdot (CH_2Cl_2)_{2.5}$ were disordered, but this could



Figure 4. Structure of the nonhelical Pt($C \equiv C$)₄Pt complex 5·(toluene)₂ with solvate molecules omitted: U, thermal ellipsoid plot (50% probability level) with hydrogen atoms omitted; V, W, views parallel and perpendicular to the planes of the C₆F₅ ligands with atoms at van der Waals radii.

Scheme 5. Reactions of the $Pt(C \equiv C)_6Pt$ Complex **10** with $Bis(\alpha, \omega$ -diphenylphosphino)alkanes



be resolved and only data for the dominant conformations are presented. Key metrical parameters are listed in Table 2. The structures of $7 \cdot (\text{chuch}_3)_2$, and $9 \cdot (\text{CH}_2\text{Cl}_2)_{2.5}$ are



Figure 5. Thermal ellipsoid plots (50% probability level) of other nonhelical $Pt(C \equiv C)_4Pt$ complexes with hydrogen atoms and solvate molecules omitted: **X**, **7**·(toluene)₄ (dominant conformation); **Y**, **7**·(CHCl₃)₂; **Z**, **9**·(CH₂Cl₂)_{2.5} (dominant conformation).

shown in Figure 5. In all three cases, the angles defined by the P–Pt–P/Pt planes were near 0° (0.2°–12.9°).



Figure 6. Structures of the two independent $Pt(C \equiv C)_{12}Pt$ complexes in crystalline $11 \cdot (CH_2Cl_2)_{0.67}$ with solvate molecules omitted: **AA**, thermal ellipsoid plot (50% probability level) of the double-helical complex with hydrogen atoms omitted; **BB**, **CC**, views parallel and perpendicular to the planes of the C_6F_5 ligands with atoms at van der Waals radii; **DD**, **EE**, **FF**, analogous views of the nonhelical complex.

As would be expected from the shorter sp³ carbon chains, **7**•(toluene)₄ and **7**•(CHCl₃)₂ crystallized in nonhelical conformations. The sp³ chains of the former laterally shielded the sp chain, whereas those of the latter bowed in similar directions, exposing more of the sp chain. The average sp/sp³ distance in the former (3.932 Å) was shorter than that in the double-helical complexes; that in the latter (3.989 Å) was comparable to the double-helical complexes. However, **7**•(CHCl₃)₂ was more dense (1.616 vs 1.468 Mg/m³), likely due in part to the heavier solvate molecule. In **9**•(CH₂Cl₂)_{2.5}, the longer sp³ chains also bowed in similar directions, with a somewhat greater average sp/sp³ distance (4.090 Å).

The dodecahexaynediyl complex $11 \cdot (CH_2Cl_2)_{0.67}$ exhibited two independent molecules in the unit cell (2:1 ratio). As shown in Figure 6, the dominant form showed a double-helical conformation (AA–CC). However, it was less twisted than the others, with an angle of 163.2° between the P–Pt–P/Pt planes and a helix pitch of 39.76 Å. Hence, the sp carbon chain is much less shielded than with the octatetraynediyl complexes in Figures 1–3. The second form was nonhelical (**DD**–**FF**), with an average sp/sp³ distance much greater than that of the helical form (4.157 vs 3.868 Å). Metrical data for the first are given in Table 1, and those for the second are presented in Table 2. From these results, we propose that an sp³/sp carbon ratio of at least 1.50 is required for a helical conformation.

5. Dynamic and Redox Properties. For each of the above molecules that could reasonably be expected to adopt a double-helical conformation, a confirmatory crystal structure has been obtained. Thus, even though **5** and **11** can also crystallize in nonhelical conformations, we thought it likely that helical motifs would be favored in solution and not solely imposed by crystal lattices. Helical molecules often contain diastereotopic groups, such as the PAr_2 substituents in **3–5**. In solution, separate NMR signals are frequently observed.^{4b} However, when the helix can

Table 4. Cyclic Voltammetry Data

complex ^a	E _{p,a} [V]	E _{p,c} [V]	<i>E</i> ° [V]	ΔE [mV]	İ _{c'a}
2^b	1.261	1.143	1.202	118	0.48
3	1.306	1.224	1.265	82	0.78
4	1.259	1.168	1.214	91	0.71
5	1.249	1.156	1.203	93	0.76
7	1.315	1.179	1.247	136	0.10
8	1.326	1.241	1.284	85	0.36
10 ^b	1.467^{b}	1.306	1.387	161	-
11	1.465	1.372	1.418	93	0.35
6	1.301	1.214	1.258	87	0.68

^{*a*} Conditions: $(7-9) \times 10^{-4}$ M, *n*-Bu₄N⁺ BF₄^{-/}CH₂Cl₂ at 22.5 ± 1 °C; Pt working and counter electrodes, potential vs Ag wire pseudoreference; scan rate 100 mV s⁻¹; ferrocene = 0.46 V. ^{*b*}Data from ref 14a.

"untwist" and rewind to an enantiomer, net exchange occurs. When this is rapid on the NMR time scale, only a single signal is observed.

Room-temperature NMR spectra of **3**–**5** showed only a single signal for all potentially diastereotopic atoms or groups. Thus, numerous low-temperature spectra were recorded, some of which are depicted elsewhere.^{26b} However, even at the lowest feasible temperatures, separate signals could not be observed (**3**, CD₂Cl₂ (–40 °C; solubility limit), toluene- d_8 (–90 °C; ¹³C signals *i/p* to P); **4**, THF- d_8 (–120 °C; ¹³C signals *i/p* to P; CH₃ ¹H and ¹³C signals); **5**, THF- d_8 (–120 °C; ¹³C signals *i/p* to P; C(CH₃)₃ ¹H and ¹³C signals). As analyzed further below, we interpret this as indicating a low barrier, ≤12 kcal/mol, for interconverting enantiomeric double helices.

In order to characterize the redox properties of the preceding complexes, cyclic voltammograms were recorded in CH_2Cl_2 . The conditions were identical to those used previously for diplatinum complexes such as 2 and 10.^{14a} Data are summarized in Table 4, and typical traces are compared in Figure 7.

Oxidations, presumably to the corresponding mixed-valent radical cations,²² were always observed, but no reductions took place prior to the solvent-induced limit.³⁷ Although the oxidations were not chemically or electrochemically reversible, the degree of reversibility, as reflected by the $i_{c/a}$ and ΔE values, was greater for the helical complexes. This is readily seen in the pairs of traces in Figure 7. Although the anodic currents for **3** and **4** are less than or equal to those of the nonhelical model compounds **6** and **2**, their cathodic currents are significantly greater, indicative of longer-lived radical cations.

Discussion

1. Syntheses. Our route to the title molecules (Schemes 4, 5) can be viewed as a one-dimensional analogue of the coordination-driven self-assembly processes used by Stang and Fujita to access two- and three-dimensional polypalladium and polyplatinum complexes with other types of rigid spacers.³⁸ However, as noted above, success is capricious. In some cases, oligomerization appears to be competitive; in other cases, the target molecules appear to form but oligomerize upon attempted workup. As described in the following paper,²⁴ many complexes that cannot be accessed are easily prepared by the alternative



Figure 7. Comparison of the cyclic voltammograms of double-helical complexes 3 and 4 with those of model compounds 6 and 2.

route in Scheme 2, using alkene metathesis/hydrogenation sequences. The resulting samples show no tendency to oligomerize.

It is also not obvious why systems with *p*-tolyl-substituted platinum endgroups (**1**, Scheme 3) are more disposed toward oligomerization than those with pentafluorophenyl-substituted end groups (**2**, **10**; Schemes 4, 5). This may reflect subtle differences in the relative rate constants for the initial phosphine substitution step and the rate-determining step for oligomerization. However, deeper insight will require mechanistic investigations beyond the scope of this study. The use of Pp-tol₃ ligands in **2** and **10** as opposed to the PPh₃ ligands in **1** gives somewhat more soluble educts, in which the phosphines remain weakly basic enough to be displaced by the monophosphine PPh₂(*n*-octyl) (leading to **6**)³⁰ and the electronically similar diphosphines Ph₂P(CH₂)_mPPh₂.

In principle, the reactions in Schemes 3-5 could also give isomers of the type **H'** (Scheme 2), which contain *trans*-spanning diphosphine ligands. However, attempts to generate similar monoplatinum species via substitution reactions proceed in very low yields and give much oligomer.³⁴ Authentic samples have been synthesized by another route²⁴ and do not readily equilibrate with the title molecules.

2. Spectroscopic and Redox Properties. The IR and UV– visible spectra of diplatinum polyynediyl complexes *trans,trans*-(Ar')(Ar₃P)₂Pt(C \equiv C)_nPt(PAr₃)₂(Ar') have been extensively analyzed in previous papers.¹⁴ As noted above, the termini-spanning diphosphine ligands in **3–9** and **11** have no detectable effect. Given the magnetic anisotropy and deshielding effects that

⁽³⁷⁾ Reduction of the Pt(C≡C)₈Pt homologue of 2 and 10 can be observed in THF, which has a wider cathodic window. However, species with shorter sp carbon chains exhibit no reduction.

 ^{(38) (}a) Leininger, S.; Olenyuk, B.; Stang, P. J. Chem. Rev. 2000, 100, 853. (b) Fujita, M.; Tominaga, M.; Hori, A.; Therrien, B. Acc. Chem. Res. 2005, 38, 369.





characterize C≡C linkages,39 we had thought that NMR chemical shifts of the sp³ carbon chains might reflect average distances from the sp carbon chain. However, DFT calculations show that when an sp³ carbon atom is moved from 3.5 Å to 5.5 Å away from the midpoint of an octatetrayne in a perpendicular plane, the ¹³C NMR chemical shift varies negligibly.⁴⁰

The thermal stabilities of 3-9 and 11 are also comparable to analogues without termini-spanning diphosphine ligands. However, the redox properties show several interpretable differences. Oxidations of other diplatinum(II) complexes with bridging unsaturated ligands afford mixed valent platinum(II)/ platinum(III) radical cations.²² Furthermore, there is good evidence that platinum(III) species often decompose via solvent interaction with the vacant axial coordination site. Stabilities are enhanced when the axial position is sterically shielded.²² Importantly, the sp³ carbon chains in the title molecules markedly increase axial shielding (compare Q and S vs M in Figure 3).

Consider the cyclic voltammetry data for 3 in Table 4. It undergoes the most reversible oxidation of the $Pt(C \equiv C)_4 Pt$ complexes, as reflected by the $i_{c/a}$ and ΔE values. Complexes 4 and 5 are comparable; their E° values are slightly less positive, indicating thermodynamically more favorable oxidations, consistent with the electron-releasing *p*-alkyl PAr₂ substituents. In going from 3 to 6, the sp³ chains are replaced by *n*-octyl groups, making the sp carbon chains and axial positions on platinum more accessible. The E° values of **3** and **6** are quite close (1.265 vs 1.258 V), but the ΔE and $i_{c/a}$ values (82 vs 87 mV; 0.78 vs 0.68) indicate a loss of reversibility, as highlighted in the overlaid traces in Figure 7 (top).

In going from 4 to 2, the sp³ chains are replaced by p-tolyl substituents. The ΔE and $i_{c/a}$ values (91 vs 118 mV; 0.71 vs 0.48) now indicate a substantial loss of reversibility (see Figure 7, bottom). In 7 and 8, the shorter sp^3 chains do not shield the $Pt(C \equiv C)_4 Pt$ moiety as extensively. Also, some ring strain may be introduced. Accordingly, reversibilities drop dramatically ($i_{c/a}$ 0.10 and 0.36). In accord with MO analyses,⁴¹ complexes with longer sp chains are thermodynamically more difficult to oxidize14,16 and furthermore show much poorer reversibilities. Both trends are evident with the $Pt(C \equiv C)_6 Pt$ complexes 10 and 11. However, the sterically shielded complex again exhibits superior reversibility.

3. Crystal Structures. Of the structural features described above, the conformations of the CH₂CH₂CH₂CH₂ and CH₂CH₂-CH₂PAr₂ moieties are particularly deserving of further analysis. As illustrated in Scheme 6, gauche and anti geometries are possible; the former are chiral. Although there are other types Stahl et al.

of chiral conformational moieties in the title molecules, the helical chirality is derived from these gauche segments. In order for the enantiomers to interconvert, each gauche segment must invert. One would expect the lowest energy pathway to involve "untwisting" to an anti conformation and then continued twisting to a gauche conformation of the opposite chirality. Only in complexes with longer sp³ chains would "belt-tightening" syn transition states be conceivable.

One question is whether there are any "conserved regions" in the $Pt(C \equiv C)_4 Pt$ complexes with $Ar_2P(CH_2)_{14}PAr_2$ bridges. As noted above and reflected by the torsion angles in Table 3, the structures of $3 \cdot (benzene)_{1.5}$ and $3 \cdot (toluene)_{1.5}$ are very similar. One sp³ chain exhibits an identical pattern of four gauche and nine anti segments (angles given on left). The other exhibits three mismatches, with three and four gauche segments, respectively. In neither molecule are the gauche/anti patterns of the two sp³ chains identical or symmetrical about the midpoint. In contrast, the sp³ chains of 5·(toluene)_{5.5} exhibit identical patterns (four gauche, nine anti) that are symmetrical about the midpoint. The sp³ chains in **4** are very similar, with just one anti/gauche mismatch.

All of the CH₂CH₂CH₂PAr₂ segments in these four molecules exhibit anti conformations, which avoid 1,5-synperiplanar interactions⁴² with PAr₂ groups. With a single exception, the immediately adjacent CH2CH2CH2CH2 segments exhibit gauche conformations, representing the onset of helicity. In 4 and 5. (toluene)_{5,5}, the anti CH₂CH₂CH₂CH₂ segments are grouped in the middle of the chain, leading to the motif evident in S in Figure 3. In all of the sp³ chains except one of $3 \cdot (benzene)_{1.5}$, the gauche torsion angles have the same sign, indicating the same helical chirality.

In the nonhelical structure $5 \cdot (toluene)_2$, the sp³ chain must connect two phosphorus atoms that are syn as opposed to anti. Hence, the distance that must be spanned is shorter. Accordingly, the sp³ chains in $5 \cdot (toluene)_2$ are more kinked, with five gauche segments each. Since this structure has a center of symmetry, the chains are identical, but the gauche segments are asymmetrically distributed. In the nonhelical complexes with shorter sp³ chains (Figure 5), the distances that must be spanned per methylene group are greater. All exhibit fewer gauche segments; in the nondisordered structure 7. (CHCl₃), there is only one in each chain (Table s2, Supporting Information).

In both the helical and nonhelical molecules in 11. (CH2-Cl₂)_{0.67}, the CH₂CH₂CH₂PAr₂ and immediately adjacent CH₂-CH₂CH₂CH₂ segments exhibit anti and gauche conformations, respectively. Each sp³ chain features 10-12 anti and 5-7 gauche conformations, the latter of both signs, as tabulated in the Supporting Information. In one sp³ chain of the helical form, the pattern is symmetrical about the midpoint; the other contains two mismatches. There is no discernible pattern in the nonhelical form, in which the chains are symmetry equivalent. Although the angle defined by the P-Pt-P/Pt planes in the helical form (163.2°) is only ca. 15% less than that with 3-5, the much greater platinum-platinum distance results in a much longer helix pitch (39.76 vs 23.66–24.45 Å).

4. Solution Structures and Dynamic Properties. The crystallographic data for the Pt(C≡C)₄Pt complexes with Ar₂P(CH₂)₁₄PAr₂ bridges suggest that they exist as equilibrium

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mixtures of at least several chiral double-helical and formally achiral nonhelical conformations in solution. As analyzed above and below, we believe that the former should dominate. However, we have been unable to establish a chiral ground state, which would feature diastereotopic aryl groups, by lowtemperature NMR. This in turn raises several questions.

First, might the double-helical conformations in Figures 1-3reflect some deep-seated lattice packing preference? For example, diplatinum polyynediyl complexes trans, trans-(X)(R₃P)₂- $Pt(C \equiv C)_n Pt(PR_3)_2(X)$ $(n \ge 3; R = alkyl, Ar; X = Ar', Cl)$ always crystallize with essentially coplanar endgroups,14,25b,43 even though DFT calculations show no electronic driving force.⁴¹ Since a packing driving force would be difficult to prove, we have focused on structural modifications that might raise the barrier for interconverting enantiomeric helical ground states.

Several components to this barrier can be identified. First, as noted above, there are torsional motions involving the CH2-CH₂CH₂PAr₂ and CH₂CH₂CH₂CH₂ segments and other portions of the molecule. These would occur both sequentially and in concert. The activation energy for the conversion of gauche- to anti-butane (see Scheme 6) is ca. 2.7 kcal/mol. If the ratedetermining step featured one such energy maximum in each sp³ chain, the conformation of which would logically be derived from a local minimum higher than the ground state, the barrier should significantly surpass 5.4 kcal/mol. Second, there should be attractive van der Waals and possibly CH/π interactions between the sp and sp³ chains. These should be maximized in helical conformations and diminished in the transition state for the rate-determining step.

Thus, higher barriers would be expected for complexes with longer sp and sp³ chains, which should have greater van der Waals contact surfaces. This possibility is explored in the following paper.²⁴ Another approach would be to decrease the sp chain length, such that endgroup-endgroup interactions destabilize the achiral planar ground state, analogous to the situation in ortho-substituted biaryls. Efforts in this direction have also been initiated.^{25b} Alternatively, functional groups could be introduced on the sp³ chain, which might enhance attractive interactions with the sp chain. Recently, similar complexes with ethereal diphosphines have been reported.44

The gain that might be associated with the first strategy is illustrated by data of Moore. He measured the binding energies of rodlike guests in the inner cavities of single-helical foldamer hosts.⁴⁵ These increased with increasing lengths of the hosts, up to the point where that of the guest molecule was exceeded. Nonetheless, none of these approaches or others⁴⁶ have yet led to a complex for which a chiral ground state can be demonstrated. Hence, the dominant conformation of the title molecules in solution remains an unsettled question.

5. Further Analyses of Helicity. The preceding issues aside, the title molecules are unique in the *absence* of bonding

expected, rates should slow under pressure.

interactions or complementarity between the two helical chains. The sp carbon chain provides a mechanical support, without which the helical conformation would not be energetically competitive in solution or the solid state. Although the comparison is unflattering in the nanotechnology era, our complexes can be viewed as molecular analogues of beanpoles, implements used by the most ancient agrarian societies to template upwardly spiraling vines.

Comparable molecules are challenging to identify. Distant relatives would include biphenyls in which the o/o and o'/o'positions are bridged by sp³ carbon chains. Systems with as many as four methylene groups have been described.⁴⁷ However, the helix strands are splayed outward from the biphenyl axis. Future studies of "carbo-mer" or (C≡C)-expanded analogues⁴⁸ would be worthwhile and may yield structures resembling those in Figures 1-3.

There is an extensive literature of single helical molecules,¹ some of which are particularly relevant to the title molecules. Rebek has demonstrated that when guests with longer $(CH_2)_m$ segments bind within cavitands or container molecules of appropriate diameters and/or lengths, helical conformations are induced.49 This can be viewed as an "external" templating, in contrast to our "internal" beanpole template. Both we and Hirsch have prepared polyynes in which the termini are joined by a single $(CH_2)_m$ tether.^{50,51} However, no structural data are so far available.

Although we could not obtain barriers for the interconversion of the enantiomers of the helical molecules in Figures 1-3, the values would represent a measure of the sp carbon chain protection, analogous to the way insulation for electrical wire is graded. A correlation to the extra activation energy required for attack upon the sp carbon chains, versus model compounds such as 6, would be expected. However, from the standpoint of engineering functional molecular devices, a portfolio of flexible bridges with a gradation of barriers would be optimal.

6. Conclusion

This study has demonstrated the accessibility of an unprecedented new class of double-helical molecules in which two sp³ carbon chains wrap around an sp carbon chain via a onestep coordination-driven self-assembly process. In contrast to other double-helical molecules, there is an absence of bonding interactions or complementarity between the strands. There are, however, synergies between the chains. The sp chain provides a mechanical support, without which helical conformations would not be energetically competitive. The sp3 chain in turn serves as a protecting group, enhancing the stabilities of mixedvalent radical cations. There is an obvious analogy to household insulated wire. Alternative routes to this class of molecules, and a variety of isomers and derivatives, will be detailed in upcoming full papers.24,44b

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^{2006. 39. 11.} (46) Helical structures might have greater inversion barriers in more polar solvents. However, these tend to have higher freezing points, hampering low temperature measurements. Also, since a positive ΔV^{\ddagger} value would be

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Experimental Section

$trans, trans-(C_6F_5)(Ph_2P(CH_2)_{14}PPh_2)Pt(C \equiv C)_4Pt(Ph_2P(CH_2)_{14}PPh_2)-(CH_2)_{14}PPh_2)-(CH_2)_{14}PPh_2)Pt(C \equiv C)_4Pt(Ph_2P(CH_2)_{14}PPh_2)-(CH_2)_{14}PPh_2)Pt(C \equiv C)_4Pt(Ph_2P(CH_2)_{14}PPh_2)-(CH_2)_{14}PPh_2)Pt(C \equiv C)_4Pt(Ph_2P(CH_2)_{14}PPh_2)-(CH_2)_{14}PPh_2)Pt(C \equiv C)_4Pt(Ph_2P(CH_2)_{14}PPh_2)-(CH_2)_{14}PPh_2)Pt(Ph_2P(CH_2)_{14}PPh_2)Pt(Ph_2P(CH_2)_{14}PPh_2)-(CH_2)_{14}PPh_2)Pt(Ph_2P(CH_2)_{14}PPh_2)Pt(Ph_2P(CH_2)_{14}PPh_2)-(CH_2)_{14}PPh_2)Pt(Ph_2P(CH_2)Pt(Ph_2P(CH_2)Pt(Ph_2P(CH_2)Pt(Ph_2P(CH_2)Pt(Ph_2)Pt(Ph_2P(CH_2)Pt(Ph_2P(CH_2)Pt(Ph_2)Pt(Ph_2P(CH_2)Pt(Ph_2P(CH_2)Pt(Ph_2P(CH_2)Pt(Ph_2)Pt(Ph_2P(CH_2)Pt(Ph_2P(CH_$

(C₆F₅) (3). A Schlenk flask was charged with *trans,trans*-(C₆F₅)(*p*-tol₃P)₂Pt(C=C)₄Pt(*Pp*-tol₃)₂(C₆F₅) (2;^{14a} 0.204 g, 0.100 mmol) and CH₂Cl₂ (40 mL). A solution of Ph₂P(CH₂)₁₄PPh₂ (0.150 g, 0.265 mmol)²⁷ in CH₂Cl₂ (20 mL) was passed through a 2 cm silica gel pad directly into the Schlenk flask with stirring. After 2 h, the solvent was removed by oil pump vacuum, and ethanol (30 mL) was added. The yellow solid was collected by filtration, washed with ethanol (2 × 10 mL), and dried by oil pump vacuum to give **3** (0.170 g, 0.087 mmol, 87%), mp ≥ 245 °C dec. Calcd for C₉₆H₉₆F₁₀P₄Pt₂: C, 59.01; H, 4.95. Found: C, 59.48; H, 5.27. DSC: endotherm with *T_i*, 181.6 °C; *T_e*, 244.0 °C; *T_p*, 263.7 °C; *T_c*, 270.3 °C; *T_f*, 270.2 °C; exotherm with *T_i*, 271.0 °C.⁵² TGA: onset of mass loss (*T_i*), 286.0 °C.

NMR (δ , CDCl₃) ¹H 7.40 (m, 16H of 8 Ph), 7.28 (m, 8H of 8 Ph), 7.19 (m, 16H of 8 Ph), 2.68 (m, 8H, PCH₂), 2.15 (m, 8H, PCH₂C*H*₂), 1.63–1.47 (m, 8H, PCH₂CH₂C*H*₂), 1.46–1.22 (m, 32H, remaining CH₂); ¹³C{¹H}^{53,54} 145.6 (dd, ¹*J*_{CF} = 223 Hz, ²*J*_{CF} = 22 Hz, *o* to Pt), 136.3 (dm, ¹*J*_{CF} = 240 Hz, *m/p* to Pt), 132.8 (virtual t, ²*J*_{CP} = 5.7 Hz, *o* to P), 132.0 (virtual t, ¹*J*_{CP} = 27.9 Hz, *i* to P), 130.1 (s, *p* to P), 127.8 (virtual t, ³*J*_{CP} = 5.0 Hz, *m* to P), 98.6 (s, ¹*J*_{CPt} = 960 Hz,⁵⁵ PtC≡), 94.2 (s, ²*J*_{CPt} = 266 Hz,⁵⁵ PtC≡*C*), 63.4 (s, PtC≡C*C*), 57.9 (s, PtC≡ CC≡*C*), 30.6 (virtual t, ³*J*_{CP} = 8.0 Hz, PCH₂CH₂CH₂), 30.0 (s, CH₂), 29.5 (s, CH₂), 29.3 (s, CH₂), 28.1 (virtual t, ¹*J*_{CP} = 18.3 Hz, PCH₂), 28.0 (s, CH₂), 25.6 (s, PCH₂CH₂); ³¹P{¹H} 14.9 (s, ¹*J*_{PPt} = 2582 Hz).⁵⁵ IR (cm⁻¹, powder film): $\nu_{C=C}$ 2148 (m), 2007 (w). UV-vis:⁵⁶ 265 (72 800), 291 (86 000), 318 (109 000), 353 (7200), 379 (4800), 412 (2400). MS:⁵⁷ 1954 (**3**⁺, 100%), 1784 ([**3**-C₆F₅-H₂]⁺, 2%), no other significant peaks >1300.

$trans, trans - (C_6F_5)(p-tol_2P(CH_2)_{14}Pp-tol_2)Pt(C \equiv C)_4Pt(p-tol_2P-tol_2P) + (C \equiv C)_4Pt(p-tol_2P-tol_2P) + (C \equiv C)_4Pt(p-tol_2P) + (C \equiv C)_$

(CH₂)₁₄Pp-tol₂)(C₆F₅) (4). Complex 2 (0.153 g, 0.075 mmol), CH₂Cl₂ (40 mL), and a solution of *p*-tol₂P(CH₂)₁₄Pp-tol₂ (0.120 g, 0.120 mmol)³⁰ in CH₂Cl₂ (20 mL) were combined in a procedure analogous to that for **3**. An identical workup gave **4** as a yellow solid (0.140 g, 0.068 mmol, 91%), mp ≥210 °C dec. Calcd for C₁₀₄H₁₁₂F₁₀P₄Pt₂: C, 60.46; H, 5.46. Found: C, 60.42; H, 5.60. DSC: endotherm with *T*_i, 196.2 °C; *T*_e, 213.2 °C; *T*_p, 242.2 °C; *T*_c, 255.6 °C; *T*_f, 259.3 °C; exotherm with *T*_i, 260.0 °C.⁵² TGA: onset of mass loss (*T*_i), 260.3 °C.

NMR (δ , CDCl₃) ¹H 7.27 (m, 16H, *o* to P), 6.99 (m, 16H, *m* to P), 2.63 (m, 8H, PCH₂), 2.29 (s, 24H, CH₃), 2.10 (m, 8H, PCH₂CH₂), 1.60–1.47 (m, 8H, PCH₂CH₂CH₂), 1.46–1.21 (m, 32H, remaining CH₂); ¹³C{¹H}^{53,54} 145.7 (dd, ¹J_{CF} = 222 Hz, ²J_{CF} = 22 Hz, *o* to Pt), 140.3 (s, *p* to P), 136.3 (dm, ¹J_{CF} = 250 Hz, *m/p* to Pt), 132.8 (virtual t, ²J_{CP} = 6.0 Hz, *o* to P), 128.7 (virtual t, ¹J_{CP} = 29.4 Hz, *i* to P), 128.5 (virtual t, ³J_{CP} = 5.5 Hz, *m* to P), 99.2 (s, PtC=), 94.1 (s, ²J_{CPt} = 269 Hz,

PtC≡*C*), 63.5 (s, PtC≡*CC*), 57.8 (s, PtC≡*CC*≡*C*), 30.7 (virtual t, ${}^{3}J_{CP}$ = 7.8 Hz, PCH₂CH₂CH₂, 30.0 (s, CH₂), 29.6 (s, CH₂), 29.4 (s, CH₂), 28.3 (virtual t, ${}^{1}J_{CP}$ = 18.4 Hz, PCH₂), 28.0 (s, CH₂), 25.5 (s, PCH₂CH₂), 21.2 (s, CH₃); ${}^{31}P{}^{1}H{}$ 13.5 (s, ${}^{1}J_{PPt}$ = 2562 Hz).⁵⁵ IR (cm⁻¹, powder film): $\nu_{C=C}$ 2146 (m), 2003 (w). MS:⁵⁷ 2066 (4⁺, 100%), 1973 ([4-C₆F₅-H₂]⁺, 2%), 1897 ([4-2C₆F₅-H₂]⁺, 4%), no other significant peaks > 850.

trans,trans-(C₆F₅)((p-t-BuC₆H₄)₂P(CH₂)₁₄P(p-C₆H₄-t-Bu)₂)Pt(C=

C)₄Pt((p-t-BuC₆H₄)₂P(CH₂)₁₄P(p-C₆H₄-t-Bu)₂)(C₆F₅) (5). Complex 2

(0.153 g, 0.075 mmol), CH₂Cl₂ (50 mL), and a solution of (*p*-*t*-BuC₆H₄)₂P(CH₂)₁₄P(*p*-C₆H₄-*t*-Bu)₂ (0.149 g, 0.188 mmol)³⁰ in CH₂Cl₂ (94 mL) were combined in a procedure analogous to that for **3**. An identical workup gave **5** as a yellow solid (0.138 g, 0.057 mmol, 77%), mp >210 °C dec. Calcd for C₁₂₈H₁₆₀F₁₀P₄Pt₂: C, 63.99; H, 6.71. Found: C, 64.72; H, 6.92.

NMR (δ , CDCl₃) ¹H 7.30 (m, 16H, *o* to P), 7.17 (m, 16H, *m* to P), 2.61 (m, 8H, PCH₂), 2.20 (m, 8H, PCH₂CH₂), 1.63–1.49 (m, 8H, PCH₂-CH₂CH₂), 1.46–1.25 (m, 32H, remaining CH₂), 1.23 (s, 72H, CH₃); ¹³C{¹H}^{53,54} 153.3 (s, *p* to P), 145.5 (dd, ¹J_{CF} = 224 Hz, ²J_{CF} = 24 Hz, *o* to Pt), 136.0 (dm, ¹J_{CF} = 248 Hz, *m/p* to Pt), 132.6 (virtual t, ²J_{CF} = 5.8 Hz, *o* to P), 128.8 (virtual t, ¹J_{CP} = 28.7 Hz, *i* to P), 124.7 (virtual t, ³J_{CP} = 5.1 Hz, *m* to P), 99.1 (s, PtC=), 94.4 (s, PtC=C), 63.7 (s, PtC=CC), 57.5 (s, PtC=CC=C), 34.7 (s, C(CH₃)₃), 31.1 (s, CH₃), 30.6 (virtual t, ³J_{CP} = 7.9 Hz, PCH₂CH₂CH₂), 30.4 (s, CH₂), 29.9 (s, CH₂), 29.5 (s, CH₂), 28.3 (virtual t, ¹J_{CP} = 18.3 Hz, PCH₂), 27.9 (s, CH₂), 25.7 (s, PCH₂CH₂); ³¹P{¹H} 12.7 (s, ¹J_{PPt} = 2561 Hz).⁵⁵ IR (cm⁻¹, powder film): $\nu_{C=C}$ 2146 (m), 2003 (w). UV-vis:⁵⁶ 292 (114 000), 318 (154 000), 351 (8000), 379 (8000), 410 (4000). MS:⁵⁷ 2402 (**5**⁺, 100%), no other significant peaks >975.

$\textit{trans-trans-}(C_6F_5)(Ph_2P(CH_2)_{10}PPh_2)Pt(C \blacksquare C)_4Pt(Ph_2P(CH_2)_{10}PPh_2)-(CH_2)_{10}PPh_2)Pt(C \blacksquare C)_4Pt(Ph_2P(CH_2)_{10}PPh_2)Pt(C \blacksquare C)_4Pt(Ph_2P(CH_2)_{10}PPh_2)Pt(Ph_2P(CH_2)PPh_2)Pt(Ph_2P(CH_2)_{10}PPh_2)Pt(Ph_2P(CH_2)PPh_2)Pt(Ph_2P(CH_2)Pt(Ph_2P(Ph_2P(Ph_2)PPh_2)Pt(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2)PPh_2)Pt(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph_2P(Ph$

(C₆F₅) (7). Complex 2 (0.153 g, 0.075 mmol), CH₂Cl₂ (75 mL), and a solution of Ph₂P(CH₂)₁₀PPh₂ (0.093 g, 0.180 mmol)^{27,28} in CH₂Cl₂ (180 mL) were combined in a procedure analogous to that for **3**. An identical workup gave **7** as a yellow solid (0.111 g, 0.060 mmol, 80%), mp >200 °C dec. DSC: endotherm with T_{i} , 222.9 °C; T_{e} , 258.1 °C; T_{p} , 271.5 °C; T_{c} , 279.1 °C; T_{f} , 278.9 °C; exotherm with T_{i} , 279.1 °C.⁵² TGA: onset of mass loss (T_{i}), 264 °C.

NMR (δ , CDCl₃) ¹H 7.38 (m, 16H of 8 Ph), 7.29 (m, 8H of 8 Ph), 7.21 (m, 16H of 8 Ph), 2.71 (m, 8H, PCH₂), 2.01 (m, 8H, PCH₂CH₂), 1.58–1.47 (m, 8H, PCH₂CH₂CH₂), 1.40–1.25 (m, 16H, remaining CH₂); ¹³C{¹H}^{53,54} 145.8 (d, ¹J_{CF} = 227 Hz, *o* to Pt), 136.4 (dm, ¹J_{CF} = 239 Hz, *m/p* to Pt), 132.8 (virtual t, ²J_{CP} = 5.8 Hz, *o* to P), 131.9 (virtual t, ¹J_{CF} = 27.5 Hz, *i* to P), 130.3 (s, *p* to P), 127.9 (virtual t, ³J_{CP} = 5.1 Hz, *m* to P), 101.6 (s, PtC=), 94.0 (s, PtC=C), 64.4 (s, PtC=CC), 59.3 (s, PtC=CC=C), 32.2 (virtual t, ³J_{CP} = 7.6 Hz, PCH₂-CH₂CH₂), 30.7 (s, CH₂), 29.9 (s, CH₂), 29.2 (virtual t, ¹J_{CP} = 18.3 Hz, PCH₂), 26.4 (s, PCH₂CH₂); ³¹P{¹H} 14.4 (s, ¹J_{PPt} = 2572 Hz).⁵⁵ IR (cm⁻¹, powder film): $\nu_{C=C}$ 2138 (m), 1996 (w). MS:⁵⁷ 1841 (7⁺, 100%), 1763 ([7-C₆H₅-H]⁺, 4%), no other significant peaks >900.

$\textit{trans,trans-}(C_6F_5)(Ph_2P(CH_2)_{11}PPh_2)Pt(C \equiv C)_4Pt(Ph_2P(CH_2)_{11}PPh_2)-$

(C₆F₅) (8). Complex 2 (0.153 g, 0.075 mmol), CH₂Cl₂ (75 mL), and a solution of Ph₂P(CH₂)₁₁PPh₂ (0.094 g, 0.180 mmol)^{27,28} in CH₂Cl₂ (180 mL) were combined in a procedure analogous to that for **3**. An identical workup gave **8** as a yellow solid (0.113 g, 0.060 mmol, 80%), mp > 220 °C dec. DSC: endotherm with T_i , 188.5 °C; T_e , 225.1 °C; T_p , 244.1 °C; T_c , 251.8 °C; T_i , 254.1 °C; exotherm with T_i , 256 °C.⁵² TGA: onset of mass loss (T_i) 267 °C.

NMR (δ , CDCl₃) ¹H 7.39 (m, 16H of 8 Ph), 7.29 (m, 8H of 8 Ph), 7.21 (m, 16H of 8 Ph), 2.71 (m, 8H, PCH₂), 2.09 (m, 8H, PCH₂CH₂), 1.56–1.23 (m, 28H, remaining CH₂); ¹³C{¹H}^{53,54} 145.8 (d, ¹J_{CF} = 228 Hz, *o* to Pt), 136.4 (dm, ¹J_{CF} = 247 Hz, *m/p* to Pt), 132.9 (virtual t, ²J_{CP} = 5.6 Hz, *o* to P), 131.9 (virtual t, ¹J_{CP} = 27.7 Hz, *i* to P), 130.3

^{(52) (}a) DSC and TGA data were treated as recommended by Cammenga, H. K.; Epple, M. Angew. Chem., Int. Ed. Engl. 1995, 34, 1171; Angew. Chem. 1995, 107, 1284. The T_e values best represent the temperature of the phase transition or exotherm. (b) Except in cases of desolvation, DSC measurements were not continued beyond the onset of mass loss (TGA).

^{(53) (}a) In most ¹³C NMR spectra, the *i*-C₆F₅ signal was not observed. (b) For virtual triplets (Hersh, W. H. *J. Chem. Educ.* **1997**, *74*, 1485), the *J* values represent the *apparent* couplings between adjacent peaks. (c) The PtC≡ CC≡C signals were assigned according to trends established earlier.¹⁴

⁽⁵⁴⁾ Complexes with PtPCH₂CH₂CH₂CH₂ linkages exhibit a characteristic pattern of ¹³C NMR signals. A ¹H, ¹H COSY spectrum of **4** was used to make definitive assignments of the corresponding ¹H signals, and a ¹H, ¹³C COSY spectrum in turn gave definitive assignments of the ¹³C signals. Analogous experiments have been conducted with closely related compounds.³⁴
(55) This coupling represents a satellite (d; ¹⁹⁵Pt = 33.8%) and is not reflected

⁽⁵⁵⁾ This coupling represents a satellite (d; ¹⁵⁵Pt = 33.8%) and is not reflected in the peak multiplicity given.

⁽⁵⁶⁾ UV-visible spectra were recorded in CH_2Cl_2 (1.25 × 10⁻⁶ M). Absorptions are in nm (ϵ , M⁻¹ cm⁻¹).

⁽⁵⁷⁾ FAB (3-nitrobenzyl alcohol matrix); m/z for the most intense peak of the isotope envelope.

⁽⁵⁸⁾ Additional data are available from the Cambridge Crystallographic Data Centre via the following CCDC numbers: 171482 (3 (toluene)_{1.5}), 176209 (3 (benzene)_{1.5}), 171483 (4), 171486 (5 (toluene)_{5.5}), 606935 (5 (toluene)₂), 171485 (7 (toluene)₄), 171484 (7 (CHCl₃)₂), 606936 (9 (CH₂Cl₂)_{2.5}), 171481 (11 (CH₂Cl₂)_{0.67}).

(s, *p* to P), 127.9 (virtual t, ${}^{3}J_{CP} = 5.1$ Hz, *m* to P), 99.4 (s, PtC≡), 93.7 (s, PtC≡C), 63.4 (s, PtC≡CC), 58.5 (s, PtC≡CC=C), 31.3 (virtual t, ${}^{3}J_{CP} = 7.9$ Hz, PCH₂CH₂CH₂), 30.8 (s, CH₂), 30.4 (s, CH₂), 29.4 (s, CH₂), 28.7 (virtual t, ${}^{1}J_{CP} = 18.0$ Hz, PCH₂), 26.3 (s, PCH₂CH₂); ${}^{31}P{}^{1}H{}$ 14.6 (s, ${}^{1}J_{PPt} = 2576$ Hz).⁵⁵ IR (cm⁻¹, powder film): $\nu_{C=C}$ 2146 (m), 2003 (w). UV−vis:⁵⁶ 275 (76 000), 291 (118 400), 316 (148 800), 352 (9600), 379 (8800), 410 (4800). MS:⁵⁷ 1869 (8⁺, 100%), 1791 ([8−C₆F₅]⁺, 2%), 1701 ([8−2C₆F₅]⁺, 6%), no other significant peaks > 825.

trans,trans-(C₆F₅)(Ph₂P(CH₂)₁₂PPh₂)Pt(C=C)₄Pt(Ph₂P(CH₂)₁₂PPh₂)-

(C₆F₅) (9). Complex 2 (0.103 g, 0.050 mmol), CH₂Cl₂ (100 mL), and a solution of Ph₂P(CH₂)₁₂PPh₂ (0.065 g, 0.12 mmol)^{27,28} in CH₂Cl₂ (120 mL) were combined in a procedure analogous to that for **3**. After 14 h, the solvent was removed by oil pump vacuum. The residue was chromatographed (25 cm silica gel column, 30:70 v/v CH₂Cl₂/hexanes). The solvent was removed from the product-containing fraction by rotary evaporation and oil pump vacuum to give **9** as a yellow solid (0.034 g, 0.018 mmol, 36%), mp >210 °C dec.

NMR (δ , CDCl₃) ¹H 7.37 (m, 16H of 8 Ph), 7.28 (m, 8H of 8 Ph), 7.19 (m, 16H of 8 Ph), 2.68 (m, 8H, PCH₂), 2.04 (m, 8H, PCH₂CH₂), 1.56–1.46 (m, 8H, PCH₂CH₂CH₂), 1.45–1.23 (m, 24H, remaining CH₂); ¹³C{¹H}^{53,54} 145.7 (dd, ¹J_{CF} = 225 Hz, ²J_{CF} = 22 Hz, *o* to Pt), 136.3 (dm, ¹J_{CF} = 240 Hz, *m/p* to Pt), 132.8 (virtual t, ²J_{CP} = 6.0 Hz, *o* to P), 131.9 (virtual t, ¹J_{CP} = 28.0 Hz, *i* to P), 130.2 (s, *p* to P), 127.9 (virtual t, ³J_{CP} = 5.2 Hz, *m* to P), 98.8 (s, PtC=), 94.8 (s, ²J_{CPt} = 272 Hz, ⁵⁵ PtC=C), 63.4 (s, PtC=CC), 58.4 (s, PtC=CC=C), 31.2 (virtual t, ³J_{CP} = 7.2 Hz, PCH₂CH₂CH₂), 30.1 (s, 2CH₂), 28.5 (s, CH₂), 28.4 (virtual t, ¹J_{CP} = 18.4 Hz, PCH₂), 25.5 (s, PCH₂CH₂); ³¹P{¹H} 13.7 (s, ¹J_{PPt} = 2573 Hz).⁵⁵ IR (cm⁻¹, powder film): $\nu_{C=C}$ 2142 (m), 1999 (w). MS:⁵⁷ 1897 (**9**⁺, 100%), 1729 ([**9**-C₆F₅]⁺, 6%).

trans,trans-(C₆F₅)(Ph₂P(CH₂)₁₈PPh₂)Pt(C=C)₆Pt(Ph₂P(CH₂)₁₈PPh₂)-(C₆F₅) (11). A Schlenk flask was charged with *trans,trans*-(C₆F₅)(*p*tol₃P)₂Pt(C=C)₆Pt(P*p*-tol₃)₂(C₆F₅) (10;^{13a} 0.105 g, 0.050 mmol) and CH₂Cl₂ (2.0 L). A solution of Ph₂P(CH₂)₁₈PPh₂ (0.060 g, 0.106 mmol)²⁷ in CH₂Cl₂ (10 mL) was added over 5 h. The solution was concentrated by rotary evaporation (100 mL), and ethanol (20 mL) was added. The solution was further concentrated until a yellow powder precipitated (oligomers). The powder was removed by filtration (G3 frit) and washed with ethanol. The filtrate was stored at -18 °C. Yellow needles formed overnight and were collected by filtration, washed with ethanol (5 mL), and dried by oil pump vacuum to give **11**•(CH₂Cl₂)_{0.67} (0.029 g, 0.014 mmol, 27%), mp >230 °C dec. Calcd for C₁₀₈H₁₁₂F₁₀P₄Pt₂•(CH₂-Cl₂)_{0.67}: C, 60.13; H, 5.26. Found: C, 59.40; H, 5.25. DSC: desolvation endotherm with *T*_i, 45.3 °C; *T*_e, 71.4 °C; *T*_p, 99.5 °C; *T*_c, 112.9 °C; *T*_f, 131.2 °C; exotherm with *T*_i, 172.9 °C; *T*_e, 190.9 °C; *T*_p, 203.1 °C; *T*_c, 211.8 °C; *T*_f, 215.2 °C; endotherm with *T*_i, 215.3 °C; *T*_e, 227.9 °C; *T*_p, 241.1 °C; *T*_c, 246.1 °C; *T*_f, 246.1 °C.⁵² TGA: mass loss 1.9% between 60 °C and 102 °C (calculated for (CH₂Cl₂)_{0.67}, 1.7%); onset of further mass loss (*T*_e), 236.3 °C.

NMR (δ , CDCl₃) ¹H 7.38 (m, 16H of 8 Ph), 7.29 (m, 8H of 8 Ph), 7.21 (m, 16H of 8 Ph), 5.28 (s, CH₂Cl₂), 2.69 (m, 8H, PCH₂), 2.08 (m, 8H, PCH₂CH₂), 1.52 (m, 8H, PCH₂CH₂CH₂), 1.36 (m, 8H, PCH₂CH₂-CH₂CH₂), 1.21 (m, 40H, remaining CH₂); ¹³C{¹H}^{53,54} 143.8 (dd, ¹J_{CF} = 225 Hz, ²J_{CF} = 30 Hz, *o* to Pt), 136.5 (dm, ¹J_{CF} = 242 Hz, *m/p* to Pt), 132.8 (virtual t, ²J_{CP} = 6.0 Hz, *o* to P), 131.6 (virtual t, ¹J_{CP} = 28.2 Hz, *i* to P), 130.4 (s, *p* to P), 127.9 (virtual t, ³J_{CP} = 5.0 Hz, *m* to P), 103.8 (s, PtC=), 93.6 (s, PtC=C), 65.0, 62.7, 61.0, 57.3 (4 s, PtC=CH₂CH₂CH₂), 30.2 (s, CH₂), 30.1 (s, CH₂), 30.0 (s, CH₂), 30.0 (s, CH₂), 28.4 (virtual t, ¹J_{CP} = 18.2 Hz, PCH₂), 25.7 (s, PCH₂CH₂); ³¹P{¹H} 14.5 (s, ¹J_{PPt} = 2554 Hz).⁵⁵ IR (cm⁻¹, powder film): $\nu_{C=C}$ 2131 (m), 2092 (s), 1996 (m). UV-vis:⁵⁶ 312 (79 200), 331 (233 600), 354 (360 800). MS:⁵⁷ 2113 ([**11**-H]⁺, 100%), 1945 ([**11**-C₆F₅]⁺, 8%), no other significant peaks >825.

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Supporting Information Available: Details of general methods including cyclic voltammetry, crystallographic data collection and refinement,⁵⁸ and associated tables. This material is available free of charge via the Internet at http://pubs.acs.org.

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