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Stereochemical Inversion of Rim-Differentiated Pillar[5]arene Molecular Swings

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ABSTRACT: To investigate the dynamic stereochemical inversion behavior of pillar[5]arenes (P[5]s) in more detail, we synthesized a series of novel rim-differentiated P[5]s with various substituents and examined their rapid rotations by variable-temperature NMR (203–298 K). These studies revealed for the first time the barrier of "methyl-through-the-annulus" rotation (ΔG^{\ddagger} = 47.4 kJ·mol⁻¹ in acetone) and indicated that for rim-differentiated P[5]s with two types of alkyl substituents, the smaller rim typically determines the rate of rotation. However, substituents with terminal C=C or C=C bonds give rise to lower inversion barriers, presumably as a result of attractive $\pi - \pi$ interactions in the transition state. Finally, data on a rim-differentiated pentamethyl-penta-propargyl P[5] exhibited the complexity of the overall inversion dynamics.

INTRODUCTION

Pillar[5]arene¹ (P[5]s), consisting of five repeating parasubstituted hydroquinone units connected by methylene bridges, are a class of macrocycles with intriguing dynamic stereochemical properties.² In the solid state, the P[5] scaffold forms its characteristic name-giving hollow pillar-like structure, with all five hydroquinone units aligned, as a result of both steric hindrance and guest inclusion.1 While a near-constant pillar shape is observed with alkoxy groups on both sides, this conformation is not observed anymore if the alkyl substituents are (partially) removed to expose phenolic -OH groups for intramolecular hydrogen bonding,³ or when one of the rims has only C-H functionalization, as shown in recently synthesized tiara[5]arenes.⁴ Such structural variability reflects the rich conformational isomerism of P[5]s,⁵ as the hydroquinone planes may rotate and adopt various angles relative to the overall pentagonal scaffold in solution.

Different from calix[n]arenes,⁶ on which additional substituents have to be grafted to break internal mirror planes, P[5]s are inherently chiral⁷ regardless of their functionalization patterns. The chirality of the P[5] scaffold with respect to a plane of symmetry σ arises from its nonplanar conformations, which contain neither plane nor center of symmetry. Enantiomeric P[5] conformers are interconvertible to each other through a sequence of "substituent-through-the-annulus" inside-out and outside-in rotations (Figure 1). This motion is



Figure 1. Schematic representation of the stereochemical inversion between enantiomeric conformers of pillar[5]arene and the idealized energy curve of the inversion process. The hydroquinone units flip by way of " R^1/R^2 -through-the-annulus" motions: R^1 moves either through the cavity while R^2 goes along the outside (top route in red) or the other way around (bottom route in blue).

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complicated in nature: before the first ring has fully rotated, the second already starts to rotate to minimize any incurred steric hindrance between adjacent rings. Along this interconversion path, multiple transition states and intermediates may be observed, depending on the nature of the substituents.⁸ When the inversion energy barriers are not high enough to hinder such movements, in solution, P[5]s are stereolabile, undergoing rapid enantiomerization⁹ or diastereoisomerization¹⁰ when chiral substituents are attached onto the P[5] scaffold, under thermodynamic equilibrium.

In 2010, Ogoshi and co-workers systematically investigated¹¹ the stereochemical inversion processes of a series of per-functionalized P[5]s with different *n*-alkyl chain lengths by dynamic NMR (DNMR). The rotation barriers derived from the corresponding coalescence temperatures (T_c) range from 46.3 (ethyl) to 63.2 (*n*-dodecyl) kJ·mol⁻¹, showing a monotonic increase of the inversion barrier with the increase of the *n*-alkyl chain length. Nonetheless, the inversion barrier of per-methyl-P[5],^{1a} one of the most widely used P[5]s on account of its high-yielding facile synthesis, could not be accurately determined, since its overall structure does not contain any diastereotopic ¹H probes for DNMR study. The energy barrier for rotation of A1-/A2-dihydroxy-octamethyl $P[5][A1/A2^{3b,12}$ refers to both hydroxy groups being bound to the same aryl ring, and located at different rims] was reported¹³ by Stoddart et al. in 2013 to be 49.7 kJ·mol⁻¹ in CDCl₃. The interactions provided by intramolecular hydrogen bonds were found to significantly impact the stereochemical inversion barriers of these difunctionalized P[5] derivatives. When the substituents on P[5]s are bulky enough to enable successful chiral resolution, inversion energy barriers canapart from by DNMR-also be determined¹⁴ by monitoring the racemization process using circular dichroism spectroscopy or HPLC with chiral stationary phases.

So far, only per-functionalized¹⁵ and A1/A2-disubstituted $P\lceil 5\rceil s^{3b,12}$ have been studied by DNMR for their inversion processes, since most other P[5] functionalization patterns with low symmetry result in complicated NMR spectra whose assignments remain nontrivial. Rim-differentiated P[5]s, whose two rims are decorated with different functionalities, represent another type of highly symmetric P[5] species. Such rim differentiation is desirable for a number of applications, and we thought to make use of this platform for dynamic stereochemical studies on account of its two nondegenerate rotational modes, e.g., R¹-through-the-annulus and R²-throughthe-annulus (Figure 1). However, progress on DNMR studies of such rim-differentiated P[5]s was hampered by their inefficient syntheses and difficult purification steps. In recent years, our research group has developed¹⁷ efficient methods for the synthesis and derivatization of C5-symmetric rim-differentiated P[5]s that largely overcome these limitations.

Taking advantage of this unique molecular scaffold, in this paper, we report rim-differentiated P[5]-based "molecular pentagonal swings" with restricted rocking motions. These specifically synthesized compounds allow for the determination of unattainable inversion barriers by variable-temperature ¹H NMR studies (203–298 K), differentiate the roles of size and presence of π -electrons, and indicate a far greater complexity of the rotation behavior than anticipated.

RESULTS AND DISCUSSION

Synthesis. First, a series of rim-differentiated P[5]s 1a-1e were synthesized (Scheme 1) using the methodology we

Scheme 1. (a) Syntheses of Rim-Differentiated Pillar[5]arenes and (b) Per-Functionalized Pillar[5]arenes under Current Study

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recently developed.¹⁷ Per-functionalized P[5]s 2a-2c were prepared by following the procedures reported in the literature.^{11,18} We briefly describe this procedure for one of the compounds under study, a rim-differentiated P[5] with 4cyanobenzyl moieties on one rim and methyl groups on the other, (4-cyanobenzyl)₅(Me)₅-P[5] 1d (see also the Experimental Section). Compound 1d was obtained through two synthetic routes. The first one involved the FeCl₃-catalyzed cyclization of the corresponding asymmetrically substituted 2,5-dialkoxybenzyl alcohol according to our previously synthesized "preoriented" procedure^{17a} and generated the expected product in 10% yield after careful column chromatography. The other pathway entailed the synthesis of generic building block $(benzyl)_5(Me)_5$ -P[5] $1e^{16c,17b}$ —as this can easily be obtained on a gram scale, with 25% isolated vield-followed by a highly efficient two-step debenzylation/ realkylation^{17b} with 4-cyanobenzyl bromide, affording (4- $(Me)_{5}-P[5]$ 1d in a near-quantitative yield. This superior yield along with a much easier purification step (precipitation from hot EtOAc) makes this indirect route highly preferable.

Dynamic NMR Studies. In our molecular design for $(4-cyanobenzyl)_5(Me)_5-P[5]$ 1d, the 4-cyanobenzyl groups serve as stoppers that are too bulky to enter the P[5] cavity, and its benzylic geminal protons can be employed as the ¹H DNMR probe. On the other hand, the "methyl-through-the-annulus" movement of the other rim seems hardly affected (see Figure 2a). Instead of continuous full rotations, the stereochemical inversion of the P[5] scaffold can still happen through back-and-forth rocking resembling that of swings on the macroscopic scale.

To further verify that the 4-cyanobenzyl group is an effective stopper of the rotational motion, single-crystal samples of $(4-cyanobenzyl)_{5}(Me)_{5}-P[5]$ 1d were obtained and then elucidated (Figure 2b) by X-ray crystallography to provide detailed information of its molecular geometry. In the solid state, the P[5] macrocyclic scaffold adopts the typical pillar-like shape containing a *n*-hexane guest in the cavity. The distance between the center of a hydroquinone unit and the nitrogen



Figure 2. (a) Stereochemical inversions of **1d** can be completed with predominant "methyl-through-the-annulus" rotations, while similar movements of the other 4-cyanobenzyl side were suppressed. (b) Side views of crystal structure of **1d**, where the distance between N atom and center of aromatic unit on the P[5] scaffold is 10.2 Å and the cavity size is 5.5 Å. (c) Partial DNMR spectra of H_a/H_b on **1d** from 298 down to 203 K in acetone- d_6 . (d) Partial DNMR spectra of **1e** from 298 down to 213 K in toluene- d_8 .

atom on the tip of the 4-cyanobenzyl group is 10.2 Å, which is larger than the roughly 5.5 Å opening of the P[5] macrocycle (see Figure 2b). Furthermore, a difunctionalized P[5] with two 4-cyanobenzyl moieties attached on both sides of one particular hydroquinone unit, namely, A1/A2-bis(4-cyanobenzyl)-P[5] **1d**', was also synthesized (see the Experimental Section) for DNMR studies. A clear AB system of ¹H NMR signals of the $-OCH_2-$ methylene protons was still observed (see the Supporting Information) at an elevated temperature of 367 K, indicating that even at that temperature, the geminal $-OCH_2-$ protons exchange only slowly on the NMR time scale. These experiments both suggest that the size of the oxygen-bound 4-cyanobenzyl substituent is sufficient to block the dynamic stereochemical inversion when attached to P[5]s.

The on-average C_5 symmetry of 1d renders the corresponding protons on the five repeating hydroquinone units homotopic, resulting in simple and clear NMR spectra that are easy to interpret. Variable-temperature 400 MHz ¹H NMR spectra were collected in acetone- d_6 from 203 to 298 K (Figure 2c; see the Supporting Information for full spectra). As expected, the geminal protons H_a and H_b of the $-OCH_2$ methylene units are diastereotopic and anisochronous, pubs.acs.org/joc

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appearing as two doublets in the NMR spectra recorded at lower temperatures. The inversion processes become faster and increasingly relevant on the NMR time scale as the temperature increases, and the fast exchange of the chemical environment surrounding protons H_a and H_b causes coalescence of their ¹H NMR signals at 4.69 ppm at 240 K. Above the coalescence temperature (T_c) , the $-OCH_2$ - proton signals eventually converge into a sharp singlet. Since the stereochemical inversion of the "molecular pentagonal swings" is only accessible through "methyl-through-the-annulus" motions, the value of the barrier (ΔG^{\ddagger}) calculated (Table 1) from the DNMR data, 47.3 kJ·mol⁻¹, should therefore also serve as a reasonable estimation of the inversion barrier of permethyl-P[5] in acetone- d_6 . Owing to the poor solubility of 1d, attempts to determine its inversion barrier in toluene- d_8 were not successful. Instead, rim-differentiated (benzyl) (Me) -P[5] **1e** with five benzyl groups¹⁹ was subjected (Figure 2d) to DNMR measurements in toluene- d_{8} , and the corresponding T_{c} and ΔG^{\ddagger} were found (Table 1) to be 250.5 K and 49.6 kJ· mol^{-1} , respectively.

To further understand the dynamic stereochemical inversion behavior of P[5]s, additional DNMR studies were carried out in toluene- d_8 on a series of rim-differentiated P[5]s with methyl on one rim and propargyl/allyl/propyl groups on the other (compounds 1a-1c), as well as the corresponding perfunctionalized P[5]s 2a-2c with propargyl/allyl/propyl substituents.

Their coalescence temperatures and inversion barriers are summarized in Table 1, and full NMR spectra are available in the Supporting Information. The DNMR measurement of perfunctionalized $(propyl)_{10}$ -P[5] 2c shows (Figure 3a) a coalescence temperature at 258 K and a corresponding inversion barrier of 51.8 kJ·mol⁻¹, which is within the ± 1 kJ· mol⁻¹ error range of the DNMR method compared to the value of 50.9 kJ·mol⁻¹ reported¹¹ by Ogoshi on the same compound. Nonetheless, the -OCH₂- geminal proton NMR signals of per-functionalized $(propargyl)_{10}$ -P[5] 2a and (allyl)₁₀-P[5] **2b** did not show (Figure 3b,c) any splitting even down to 203 K, suggesting that rotations of their hydroquinone units still occur faster than the NMR time scale, and their ΔG^{\ddagger} values are even smaller than the lower detection limit around 40 kJ·mol⁻¹. Given the minor difference between propyl and propargyl/allyl groups in terms of size, this indicates that-in addition to steric hindrance and hydrogen bonds—the presence of π electrons might favorably interact with the aryl π electrons during the rotation process, exerting considerable influence on the rate of the rim inversion process. These experimental results further substantiate our previous theoretical investigations on P[5] rotational energy profiles,⁸ which indicated that "the stabilization of the intermediate

Table 1. Dynamic NMR Studies and Derived Kinetic Parameters for the Through-Annulus Rotation of Various Rim-Differentiated (Compounds 1a–1e) and Per-Functionalized (2a–2c) Pillar[5]arenes

P[5]	$1a^a$		1b ^{<i>a</i>}	$1c^{a}$	1d ^b	1e ^a	2a ^{<i>a</i>}	$2b^a$	$2c^{a}$
$T_{\rm c}$ (K)	238 ^c	233 ^d		238	240	250.5			258
$\Delta \nu$ (Hz)	65.6	26.0		80.0	90.4	88.0			65.2
$k_{\rm coal}~({\rm s}^{-1})$	159.6	87.1		182.8	211.7	207.2			150.5
$t_{1/2}$ (ms)	4.3	8.0		3.8	3.3	3.3			4.5
ΔG^{\ddagger} (kJ/mol)	47.6	48.4	<40	47.2	47.4	49.6	<40	<40	51.8

^aSolvent: toluene- d_8 . ^bSolvent: acetone- d_6 . ^cTwo datasets of 1a were obtained from the NMR signal of H_a/H_b . ^dTwo datasets of 1a were obtained from the NMR signal of H_c/H_d .



(a) (b) (c) 1b 1a 1c H_a/H_b H_a/H_d H_/ T (K) T (K) T (K) 298 298 298 282 283 288 273 273 273 263 263 263 253 253 243 253 243 238 238 243 235.5 236 233 233 233 228 223 223 223 🙏 213 213 213 203 203 203 4.0 4.5 4.0 3.5 3.0 4.5 3.5 4.0 Chemical Shift / ppm

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Figure 3. Partial DNMR spectra of (a) 2a, (b) 2b, and (c) 2c from 298 down to 203 K in toluene- d_8 .

structures by noncovalent van der Waals interactions, as well as by hydrogen bonds, constitutes a major factor affecting barrier heights."

From the DNMR studies above, we have established the "Rthrough-the-annulus" ΔG^{\ddagger} values for P[5]-rims functionalized with different short hydrocarbon chains in the following order: propyl (51.8 kJ·mol⁻¹ in toluene- d_8) > methyl (49.6 kJ·mol⁻¹ in toluene- d_8 or 47.4 kJ·mol⁻¹ in acetone- d_6) > allyl/propargyl $(\langle 40 \text{ kJ} \cdot \text{mol}^{-1} \text{ in toluene-} d_8)$. For rim-differentiated P[5]s with two nondegenerate rotational modes, it is anticipated that the substituent with the lower barrier height should dominate the inversion processes. This is indeed the case for 1b and 1c (see Table 1 and Figure 4). The energy barrier of $(n-Pr)_{s}(Me)_{s}$ -P[5] 1c is 47.2 kJ·mol⁻¹, which is 4.6 kJ·mol⁻¹ lower than the "propyl-through-the-annulus" value (observed for $(n-Pr)_{10}$ -P[5] 2c). Similar to the case of per-functionalized $(allyl)_{10}$ -P[5] 2b, the rim-differentiated (allyl)₅(Me)₅-P[5] 1b also displays rapid rotations in the temperature range studied, as a result of the low inversion barrier of the allyl-functionalized rim.

The rim-differentiated (propargyl)₅(Me)₅-P[5] **1a**, nonetheless, displays anomalous behavior. Instead of being dominated by the propargyl groups, which would be expected to provide a low-energy pathway for fast rotation, the inversion slows down significantly at lower temperatures, and coalescence of the $-OCH_2-$ protons was observed (Figure 4a) at 238 K, which corresponds to an energy barrier of 47.6 kJ·mol⁻¹ (Table 1). Moreover, it is noteworthy that the Ar-CH₂-Ar protons of **1a** were also found (Figure 4a) to be anisochronous, splitting into two sets of doublets at lower temperatures. A T_c of 233 K and a ΔG^{\ddagger} value of 48.4 kJ·mol⁻¹ were found for the Ar-CH₂-Ar geminal protons, in which the combination of lower T_c and higher ΔG^{\ddagger} resulted from the significant difference in Δv

Figure 4. Partial DNMR spectra of (a) 1a, (b) 1b, and (c) 1c from 298 down to 203 K in toluene- d_8 .

values for these two sets of protons (at the coalescence temperature, $\Delta v = 26.0$ Hz for the Ar–CH₂–Ar protons, and $\Delta v = 65.6$ Hz for the $-OCH_2 - protons$; see eq 1, Experimental Section). Although the methylene bridge protons of rim-differentiated P[5]s are, in principle, always diastereotopic to each other, so far la is the only compound that shows this phenomenon, which also allows observation of this complex dynamic behavior in our DNMR studies. Both ΔG^{\ddagger} values obtained for 1a are higher than the "propargyl-throughthe-annulus" barrier (<40 kJ·mol⁻¹), suggesting that the enantiomerization process of 1a is more complex than that of the other compounds under current study. This (highly reproducible, but) counterintuitive result is in line with our earlier theoretical findings that the mechanical movements between different P[5] hydroquinone units are not mutually independent,⁸ and it confirms that not only the transition-state energy but also that of starting conformations and unexpectedly stable intermediates need to be taken into account to fully describe the complex dynamics of these materials (see the Supporting Information for the DFToptimized structure of 1a). Given the relatively small size of this effect, a thorough understanding thereof might require indepth theoretical investigations at a high level of theory.

CONCLUSIONS

Using tailor-made rim-differentiated and per-functionalized pillar[5] arenes, we demonstrated that whereas the interconversion of enantiomeric pillar[5] arene conformers typically proceeds by moving the sterically smallest substituent through the cavity, the energy barrier for this process is in fact dominated by the contribution of the "fastest" substituents, i.e., the smallest one or the one that provides most stabilization in the transition state, e.g., by attractive $\pi - \pi$ interactions between the substituent and aryl rings. However, the interconversion is a complex process, in which substituents may interact

competitively or cooperatively in many ways, yielding inextricably intertwined motions during the inversion process and concomitantly occasional "outliers" in the overall activation barriers. These findings thus shed more light on the complex nature of the dynamic process of pillar[5]arene stereochemical inversion and may provide future design guidance for pillararene-based chiral sensors²⁰ and molecular switches and machines.²¹

EXPERIMENTAL SECTION

General Procedure. Starting materials, reagents, and solvents were purchased from commercial vendors and used as received, unless otherwise noted. All reactions that required heating employed an oil bath. Analytical thin-layer chromatography (TLC) was performed on aluminum sheets, precoated with silica gel GF254. Flash column chromatography was performed over silica gel (200–300 mesh or 300–400 mesh). The chemical shifts are listed in parts per million (ppm) on the δ scale, and coupling constants were recorded in hertz (Hz). Chemical shifts are calibrated relative to chloroform (CDCl₃: δ 7.26 ppm for ¹H and 77.16 ppm for ¹³C).

A1/A2-bis(4-cyanobenzyl)-P[5] 1d': In a sealed tube were introduced A1/A2-dihydroxy-octamethyl $P[5]^{12b}$ (200 mg, 0.277 mmol, 1.0 equiv), 4-cyanobenzyl bromide (271 mg, 1.39 mmol, 5.0 equiv), KI (13.8 mg, 0.083 mmol, 0.3 equiv), and K2CO3 (229 mg, 1.66 mmol, 6.0 equiv). Dry MeCN (5 mL) was added, and the resulting mixture was stirred at 115 °C for 12 h. After cooling to 25 °C, H₂O was added and the crude mixture was extracted with EtOAc $(3 \times 30 \text{ mL})$, dried over MgSO₄, filtered, and concentrated to dryness. Column chromatography (EtOAc/n-hexane, 15/85) afforded the target compound as a white solid (152 mg, 0.159 mmol, 57%). ¹H NMR (600 MHz, CDCl₃) δ 7.65 (d, J = 8.2 Hz, 4H), 7.49 (d, J = 8.2Hz, 4H), 6.82 (s, 2H), 6.82 (s, 2H), 6.78 (s, 2H), 6.75 (s, 2H), 6.64 (s, 2H), 5.02 (d, J = 13.3 Hz, 2H), 4.94 (d, J = 13.3 Hz, 2H), 3.92 (s, 1H), 3.90 (s, 1H), 3.81 (s, 2H), 3.77 (s, 4H), 3.74 (s, 1H), 3.72 (s, 1H), 3.71 (s, 6H), 3.69 (s, 6H), 3.50 (s, 6H), 3.38 (s, 6H). ${}^{13}C{}^{1}H{}$ NMR (151 MHz, CDCl₃) δ 150.82, 150.81, 150.7, 150.6, 149.9, 143.5, 132.3, 128.7, 128.6, 128.3, 128.2, 127.43, 127.38, 118.8, 115.2, 114.1, 114.0, 113.9, 111.4, 69.4, 55.97, 55.96, 55.93, 55.92, 55.6, 55.4, 29.9, 29.6, 29.4. HRMS (ESI) m/z [M + Na]⁺ calcd for C₅₉H₅₆O₁₀N₂Na 975.3827, found 975.3788.

3d: To a solution of 2-hydroxy-5-methoxybenzaldehyde (1.00 g, 6.57 mmol, 1.0 equiv) in MeCN (40 mL) was added K₂CO₃ (1.81 g, 13.1 mmol, 2.0 equiv) followed by 4-cyanobenzyl bromide (1.29 g, 6.57 mmol, 1.0 equiv). The resulting mixture was refluxed for 20 h, cooled down to 25 °C, filtered, and concentrated to dryness. The crude aldehyde was dissolved in MeOH (200 mL), to which NaBH₄ (120 mg, 3.3 mmol, 0.5 equiv) was added. The resulting mixture was stirred at room temperature for 2 h and concentrated to dryness. Column chromatography (EtOAc/n-hexane, 20/80 to 40/60) afforded 3d as a white solid (1.51 g, 5.61 mmol, 85%). ¹H NMR (400 MHz, CDCl₃): δ 7.61 (d, J = 8.0 Hz, 2H), 7.49 (d, J = 8.0 Hz, 2H), 6.95 (d, J = 2.9 Hz, 1H), 6.81-6.67 (m, 2H), 5.07 (s, 2H), 4.70 (s, 2H), 3.72 (s, 3H), 2.71 (s, 1H). $^{13}C{^1H}$ NMR (101 MHz, CDCl₃): δ 154.2, 149.6, 142.6, 132.4, 130.8, 127.4, 118.7, 114.6, 112.8, 112.7, 111.5, 69.5, 61.0, 55.6. HRMS (ESI) m/z [M + Na]⁺ calcd for C₁₆H₁₅O₃NNa 292.0944, found 292.0944.

(4-Cyanobenzyl)₅(Me)₅-P[5] **1d**: *Route A*. To a solution of **3d** (700 mg, 2.60 mmol, 1.0 equiv) in Cl(CH₂)₂Cl₂ (300 mL) was added anhydrous FeCl₃ (169 mg, 1.04 mmol, 0.4 equiv). The resulting mixture was stirred at 25 °C for 20 h, quenched with MeOH (5 mL), and concentrated to dryness. Column chromatography (CH₂Cl₂/MeOH, 98/2 to 95/5) followed by slow evaporation from a CHCl₃ solution (40 mL) afforded **1d** as a white solid (60 mg, 0.05 mmol, 10%). *Route B*. To a solution of (benzyl)₅(Me)₅-P[5] **1e**^{16c,17b} (100 mg, 0.088 mmol, 1.0 equiv) in EtOAc (5 mL) was added Pd/C (10% wt, wetted with 55% H₂O, 100 mg). The resulting mixture was stirred under H₂ atmosphere at 25 °C for 4 h, filtered over a pad of celite, and concentrated to dryness to afford the corresponding pentanol as a white solid (60 mg, 0.088 mmol, and concentrated to dryness to afford the corresponding pentanol as a white solid (60 mg, 0.088 mmol, and concentrated to dryness to afford the corresponding pentanol as a white solid (60 mg, 0.088 mmol, 9.088 mmol, 9.0088 mmol, 9.0088

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dissolved in H₂O (1 mL), NaOH (26.5 mg, 0.662 mmol, 7.5 equiv) was added, and the resulting mixture was stirred at 60 °C for 30 min. A solution of 4-cyanobenzyl bromide (173 mg, 0.882 mmol, 10.0 equiv) in MeCN (1 mL) was added, and the mixture was further stirred at 60 °C for 12 h. After being cooled down to rt, the precipitate was filtered out, and dissolved in EtOAc (5 mL) at 60 °C. Hexane (50 mL) was added dropwise. The resulting precipitate was collected and washed with hexane (10 mL). This procedure was repeated once to afford **1d** as a white solid (81 mg, 97%). ¹H NMR (400 MHz, CDCl₃) δ 7.59 (d, *J* = 8.0 Hz, 10H), 7.29 (d, *J* = 8.0 Hz, 10H), 6.76 (s, 5H), 6.61 (s, 5H), 4.50 (s, 10H), 3.81 (s, 10H), 3.62 (s, 15H). ¹³C{¹H} NMR (101 MHz, CDCl₃) δ 151.7, 149.4, 143.1, 132.4, 129.1, 128.5, 127.6, 118.6, 115.8, 114.5, 111.9. HRMS (ESI) *m/z* [M + Na]⁺ calcd for C₈₀H₆₅O₁₀N₅Na 1278.4624, found 1278.4611. (*n*-Pr)₅(Me)₅-P[S] **1c**: To a solution of (allyl)₅(Me)₅-P[S] **1b**^{17a}

(*n*-Pr)₅(Me)₅-P[5] **1c**: To a solution of (allyl)₅(Me)₅-P[5] **1b**^{1/a} (285 mg, 0.323 mmol, 1.0 equiv) in EtOAc (30 mL) was added Pd/C (10% wt, wetted with 55% H₂O, 285 mg). The resulting mixture was stirred under a H₂ atmosphere at 25 °C for 24 h, then filtered over a pad of celite, and concentrated to dryness. Column chromatography (EtOAc/*n*-hexane, 5/95) afforded **1c** as a white solid (256 mg, 0.287 mmol, 89%). ¹H NMR (600 MHz, CDCl₃) δ 6.80 (s, 5H), 6.76 (s, 5H), 3.78 (s, 10H), 3.77 (t, *J* = 6.8 Hz, 10H), 3.67 (s, 15H), 1.75–1.69 (m, 10H), 0.99 (t, *J* = 7.4 Hz, 15H). ¹³C{¹H} NMR (151 MHz, CDCl₃) δ 149.6, 149.1, 127.3, 127.2, 114.0, 113.1, 68.9, 54.8, 28.6, 22.0, 9.7. HRMS (ESI) *m*/*z* [M + NH₄]⁺ calcd for C₅₅H₇₄NO₁₀ 908.5307, found 908.5278.

Variable-Temperature NMR Measurements. All ¹H NMR spectra were recorded using a Bruker Avance III 400 MHz instrument with 32 scans. The ΔG^{\ddagger} values were calculated using the coalescence temperature T_c and the difference in chemical shifts ($\Delta \nu$) measured at 203 and 273 K, respectively. To determine T_c , ¹H NMR spectra were recorded at various temperatures from 203 to 343 K in toluene- d_8 or acetone- d_6 . In a typical VT NMR experiment, a 1 mM solution of P[5] was prepared by dissolving an appropriate amount in 500 μ L of toluene- d_8 and thorough mixing. To accurately determine $\Delta \nu$, the shimming was performed manually wherever poor shimming (i.e., poor resolution, broad or unsymmetrical peaks) was evidenced. To prevent artifacts like spinning sidebands, spinning was turned off wherever needed. The phasing of spectra and baseline were corrected automatically. The calculation of ΔG^{\ddagger} was performed using the following equation²²

$$\Delta G^{\ddagger} = 8.314 T_{\rm c} \left[22.96 + \log \left(\frac{T_{\rm c}}{\Delta \nu} \right) \right] \tag{1}$$

X-ray Crystallography. Single crystals suitable for X-ray diffraction were grown by the vapor-vapor diffusion method, EtOAc/n-hexane for 1c and $CHCl_3/n$ -hexane for 1d, then selected and mounted in inert oil under a cold gas stream, and their X-ray diffraction intensity data were collected on a Rigaku XtaLAB FRX diffractometer equipped with a Hypix6000HE detector, using Cu K α radiation ($\lambda = 1.54184$ Å). By the use of Olex2,²³ the structure was solved either (i) with the ShelXS structure solution program using Direct Methods or (ii) with the ShelXT structure solution program using direct methods or intrinsic phasing,²⁴ refined with the ShelXL refinement package using least-squares minimization.²⁵ The hydrogen atoms were set in calculated positions and refined as riding atoms with a common fixed isotropic thermal parameter. Selected details of the data collection and structural refinement of compounds 1c and 1d can be found in the Supporting Information, and full details are available in the corresponding CIF files (CCDC 1989352 and 1896020).

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.joc.0c01464.

X-ray data for compound 1c (CIF)

X-ray data for compound 1d (CIF)

¹H, ¹³C, and DNMR spectra, and X-ray data (PDF)

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Notes

The authors declare no competing financial interest.

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