

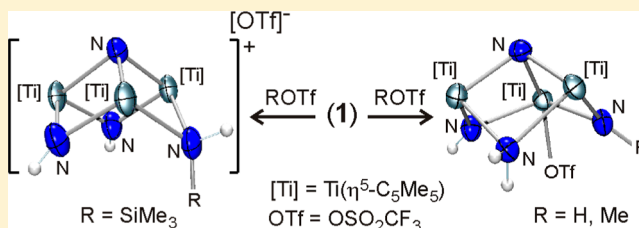
Reactivity with Electrophiles of Imido Groups Supported on Trinuclear Titanium Systems

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Supporting Information

ABSTRACT: Several trinuclear titanium complexes bearing amido μ -NHR, imido μ -NR, and nitrido μ_3 -N ligands have been prepared by reaction of $[\{\text{Ti}(\eta^5\text{-C}_5\text{Me}_5)(\mu\text{-NH})\}_3(\mu_3\text{-N})]$ (**1**) with 1 equiv of electrophilic reagents ROTf (R = H, Me, SiMe₃; OTf = OSO₂CF₃). Treatment of **1** with triflic acid or methyl triflate in toluene at room temperature affords the precipitation of compounds $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})_2(\mu\text{-NH}_2)(\text{OTf})]$ (**2**) or $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})(\mu\text{-NH}_2)(\mu\text{-NMe})(\text{OTf})]$ (**3**). Complexes **2** and **3** exhibit a fluxional behavior in solution consisting of proton exchange between μ -NH₂ and μ -NH groups, assisted by the triflate ligand, as could be inferred from a dynamic NMR spectroscopy study. Monitoring by NMR spectroscopy the reaction course of **1** with MeOTf allows the characterization of the methylamido intermediate $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})_2(\mu\text{-NHMe})(\text{OTf})]$ (**4**), which readily rearranges to give **3** by a proton migration from the NHMe amido group to the NH imido ligands. The treatment of **1** with 1 equiv of Me₃SiOTf produces the stable ionic complex $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})_2(\mu\text{-NHSiMe}_3)]^+[\text{OTf}]^-$ (**5**) with a disposition of the nitrogen ligands similar to that of **4**. Complex **5** reacts with 1 equiv of $[\text{K}\{\text{N}(\text{SiMe}_3)_2\}]$ at room temperature to give $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-N})(\mu\text{-NH})(\mu\text{-NHSiMe}_3)]$ (**6**), which at 85 °C rearranges to the trimethylsilylimido derivative $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})_2(\mu\text{-NSiMe}_3)]$ (**7**). Treatment of **7** with $[\text{K}\{\text{N}(\text{SiMe}_3)_2\}]$ affords the potassium derivative $[\text{K}\{(\mu_3\text{-N})(\mu_3\text{-NH})(\mu_3\text{-NSiMe}_3)\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})\}]$ (**8**), which upon addition of 18-crown-6 leads to the ion pair $[\text{K}(18\text{-crown-6})][\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-N})(\mu\text{-NH})(\mu\text{-NSiMe}_3)]$ (**9**). The X-ray crystal structures of **2**, **5**, **6**, and **8** have been determined.



INTRODUCTION

An extensive chemistry has been developed with transition metal complexes bearing imido or nitrido ligands as a terminal functionality, $\text{M}=\text{NR}$ or $\text{M}\equiv\text{N}$.¹ The reactivity of terminal imido or nitrido ligands has been an area of interest because of their potential to show either electrophilic or nucleophilic character. The behavior of these nitrogen ligands depends on the nature of the metal, its oxidation state, and the ancillary ligands. These factors determine the π interaction between the imido or nitrido moiety and the metal center, and considerations based on frontier molecular orbitals have been used to rationalize the increasing electrophilicity of the nitrogen ligand upon going from early to later transition metals.^{1,2} This experimental trend in the reactivity of nitrido complexes has been recently interpreted in terms of the variation of the partial charge on the nitrogen atom.³ The reaction of imido or nitrido ligands with nucleophiles leads to a two-electron reduction of the metal center and a reduced bond order between the metal and the nitrogen atom.^{4,5} In contrast, when imido or nitrido ligands react with electrophiles, the net result is a reduced bond order between the metal and the nitrogen atom, to produce amido and imido derivatives, respectively, with no change in the oxidation state of the metal.^{6,7}

While those extensive studies have been mainly carried out on mononuclear midtransition metal (groups 6–8) complexes

bearing terminal imido or nitrido ligands, the reported reactivity of polynuclear early transition metal (groups 4 and 5) derivatives containing bridging (μ -NR or μ -N) ligands remains comparatively scarce.^{8,9} Representative examples include nucleophilic attack of a phosphine to a bridging nitrido ligand in a dinuclear titanium derivative,¹⁰ or electrophilic attacks on μ -N groups of dinuclear titanium, niobium, or tantalum complexes generated by dinitrogen activation.¹¹

During the past decade, we have been involved in the study of the reactivity of the trinuclear imido-nitrido titanium(IV) derivative $[\{\text{Ti}(\eta^5\text{-C}_5\text{Me}_5)(\mu\text{-NH})\}_3(\mu_3\text{-N})]$ (**1**). This molecule contains two potentially reactive functionalities: the three μ -NH imido groups and the μ_3 -N nitrido ligand. While complex **1** is capable of acting as a Lewis base through the imido groups toward many metal derivatives to give cube-type adducts $[\text{L}_n\text{M}\{(\mu_3\text{-NH})_3\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})\}]$,¹³ the Lewis base behavior of the apical μ_3 -N nitrido ligand has been only documented with copper and silver MX Lewis acids.¹⁴ This nitrido ligand is quite chemically unreactive, although we have reported the “apparent” nucleophilic attack of an acetylide group $[\text{C}\equiv\text{CR}]^-$ at this site to yield alkynylimido $\mu_3\text{-NCCR}$ ligands with a two-electron reduction of the Ti_3 core.¹⁵ Density

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functional theory (DFT) calculations showed that this reaction involves the formation of an alkynyl titanium intermediate followed by migration of the alkynyl group to the apical nitrido ligand. We were interested in studying the reactivity of **1** toward electrophiles, and here we report on the reaction with **1** equiv of ROTf to generate polynuclear complexes by selective functionalization of the imido groups. The preliminary results on the treatment of complex **1** with MeOTf have been recently communicated.¹⁶

EXPERIMENTAL SECTION

General Considerations. All manipulations were carried out under argon atmosphere using Schlenk line or glovebox techniques. Toluene and hexane were distilled from Na/K alloy just before use. NMR solvents were dried with Na/K alloy (C_6D_6 , C_7D_8) or calcium hydride ($CDCl_3$, C_5D_5N) and vacuum-distilled. Dichloromethane- d_2 was dried over activated molecular sieves and stored under argon. Oven-dried glassware was repeatedly evacuated with a pumping system (ca. 1×10^{-3} Torr) and subsequently filled with inert gas. $ROSO_2CF_3$ ($R = H, Me, Me_3Si$), $[K\{N(SiMe_3)_2\}]$, and 1,4,7,10,13,16-hexaoxacyclooctadecane (18-crown-6) were purchased from Aldrich and used as received. $[Ti(\eta^5-C_5Me_5)(\mu-NH)_3(\mu_3-N)]$ (**1**) was prepared according to a published procedure.^{12b} The syntheses and characterization of complexes **3** and **4** have been reported previously.¹⁶

Samples for infrared spectroscopy were prepared as KBr pellets, and the spectra were obtained using an FT-IR Perkin-Elmer SPECTRUM 2000 spectrophotometer. 1H , $^{13}C\{^1H\}$, and ^{19}F NMR spectra were recorded on Varian Unity-300, Mercury-300, and/or Unity-500 Plus spectrometers. Chemical shifts (δ , ppm) in the 1H and $^{13}C\{^1H\}$ NMR spectra are given relative to residual protons or to carbon of the solvent. Chemical shifts (δ , ppm) in the ^{19}F NMR spectra are given relative to $CFCl_3$ as external reference. Microanalyses (C, H, N, S) were performed in a Leco CHNS-932 microanalyzer.

Synthesis of $[Ti_3(\eta^5-C_5Me_5)_3(\mu_3-N)(\mu-NH)_2(\mu-NH_2)(OSO_2CF_3)]$ (2**).** A 100 mL Schlenk flask was charged with **1** (0.30 g, 0.49 mmol), $HOSO_2CF_3$ (0.089 g, 0.59 mmol), and toluene (10 mL). The reaction mixture was stirred at room temperature for 24 h to give a yellow solid and an orange solution. The solid was isolated by filtration onto a glass frit and vacuum-dried to afford **2** as a yellow powder (0.31 g, 84%). IR (KBr, cm^{-1}): $\tilde{\nu}$ 3374 (w), 3362 (w), 3353 (m), 3250 (m), 2914 (s), 2860 (m), 2727 (w), 1578 (w), 1490 (w), 1436 (m), 1377 (s), 1315 (vs), 1279 (w), 1234 (vs), 1207 (vs), 1199 (vs), 1167 (vs), 1068 (w), 1017 (vs), 779 (vs), 775 (vs), 752 (vs), 708 (m), 678 (s), 634 (s), 548 (m), 508 (m), 485 (w), 461 (w), 420 (w). 1H NMR (300 MHz, $CDCl_3$, 20 °C): δ 12.50 (s br, 2H; NH), 3.28 (s br, 1H; NHH), 2.01 (s br, 45H; C_5Me_5), one resonance signal for the NH_2 ligand was not detected. 1H NMR (500 MHz, CD_2Cl_2 , 20 °C): δ 13.65 (s br, 2H; NH), 4.23 (m br, 1H; NHH), 3.32 (m br, 1H; NHH), 2.09 (s br, 45H; C_5Me_5). 1H NMR (500 MHz, CD_2Cl_2 , -50 °C): δ 13.69 (s br, 2H; NH), 4.37 (d, $^2J(H,H) = 9.5$ Hz, 1H; NHH), 3.20 (d, $^2J(H,H) = 9.5$ Hz, 1H; NHH), 2.04 (s, 30H; C_5Me_5), 1.92 (s, 15H; C_5Me_5). $^{13}C\{^1H\}$ NMR (75 MHz, $CDCl_3$, 20 °C): δ 121.4 (s br, C_5Me_5), 11.9 (C_5Me_5), the CF_3 carbon atom resonance was not detected. ^{19}F NMR (282 MHz, $CDCl_3$, 20 °C): δ -78.3. Anal. Calcd for $C_{31}H_{49}F_3N_4O_3SiTi_3$ ($M_w = 758.41$): C, 49.09; H, 6.51; N, 7.39; S, 4.23. Found: C, 49.88; H, 6.24; N, 7.33; S, 4.26.

Synthesis of $[Ti_3(\eta^5-C_5Me_5)_3(\mu_3-N)(\mu-NH)_2(\mu-NHSiMe_3)]-[O_3SCF_3]$ (5**).** In a fashion similar to the preparation of **2**, the treatment of **1** (0.60 g, 0.98 mmol) with $Me_3SiOSO_2CF_3$ (0.22 g, 0.99 mmol) in toluene (10 mL) for 24 h afforded **5** as an orange powder (0.72 g, 88%). IR (KBr, cm^{-1}): $\tilde{\nu}$ 3271 (m, broad), 2952 (m), 2914 (s), 1489 (w), 1428 (m), 1380 (s), 1252 (vs), 1222 (s), 1200 (m), 1149 (vs), 1031 (vs), 951 (w), 840 (vs), 761 (vs), 736 (vs), 711 (vs), 666 (vs), 637 (vs), 571 (m), 517 (m), 461 (m), 422 (w). 1H NMR (300 MHz, $CDCl_3$, 20 °C): δ 14.10 (s br, 2H; NH), 4.92 (s, 1H; $NHSiMe_3$), 2.12 (s, 30H; C_5Me_5), 2.01 (s, 15H; C_5Me_5), -0.17 (s, 9H; $NHSiMe_3$). $^{13}C\{^1H\}$ NMR (75 MHz, $CDCl_3$, 20 °C): δ 123.9 (C_5Me_5), 123.3 (C_5Me_5), 12.7 (C_5Me_5), 11.9 (C_5Me_5), 4.8 ($SiMe_3$), the CF_3 carbon atom resonance was not detected. ^{19}F NMR (282

MHz, $CDCl_3$, 20 °C): δ -77.9. Anal. Calcd for $C_{34}H_{57}F_3N_4O_3SiTi_3$ ($M_w = 830.59$): C, 49.17; H, 6.92; N, 6.75; S, 3.86. Found: C, 48.83; H, 6.44; N, 6.72; S, 3.48.

Synthesis of $[Ti_3(\eta^5-C_5Me_5)_3(\mu_3-N)(\mu-N)(\mu-NH)(\mu-NHSiMe_3)]$ (6**).** A 100 mL Schlenk flask was charged with **5** (0.30 g, 0.36 mmol), $[K\{N(SiMe_3)_2\}]$ (0.080 g, 0.40 mmol), and toluene (20 mL). The reaction mixture was stirred at room temperature for 30 min to give a red solution and a white fine solid. After filtration, the volatile components of the solution were removed under reduced pressure, and the resultant dark red solid was vacuum-dried for 3 h to give **6** (0.17 g, 69%). IR (KBr, cm^{-1}): $\tilde{\nu}$ 3353 (w), 3273 (w), 2909 (s), 2857 (s), 2721 (w), 1495 (w), 1435 (m), 1374 (m), 1261 (w), 1246 (m), 1167 (w), 1066 (w), 1030 (m), 956 (w), 838 (s), 783 (s), 764 (s), 731 (s), 703 (s), 609 (s), 516 (m), 482 (w), 466 (w), 422 (m). 1H NMR (300 MHz, C_6D_6 , 20 °C): δ 12.84 (s br, 1H; NH), 3.72 (s br, 1H; $NHSiMe_3$), 2.14 (s, 15H; C_5Me_5), 2.03 (s, 15H; C_5Me_5), 1.94 (s, 15H; C_5Me_5), 0.25 (s, 9H; $NHSiMe_3$). $^{13}C\{^1H\}$ NMR (75 MHz, C_6D_6 , 20 °C): δ 118.3 (C_5Me_5), 117.4 (C_5Me_5), 117.1 (C_5Me_5), 12.5 (C_5Me_5), 12.4 (C_5Me_5), 12.0 (C_5Me_5), 5.7 ($SiMe_3$). Anal. Calcd for $C_{33}H_{56}N_4SiTi_3$ ($M_w = 680.51$): C, 58.24; H, 8.29; N, 8.23. Found: C, 57.97; H, 8.08; N, 7.59.

Synthesis of $[Ti_3(\eta^5-C_5Me_5)_3(\mu_3-N)(\mu-NH)_2(\mu-NSiMe_3)]$ (7**).** A 100 mL ampule (Teflon stopcock) was charged with **5** (0.30 g, 0.36 mmol), $[K\{N(SiMe_3)_2\}]$ (0.080 g, 0.40 mmol), and toluene (20 mL). The reaction mixture was stirred at 85 °C for 16 h. After filtration, the volatile components of the solution were removed under reduced pressure to give **7** as an orange solid (0.17 g, 69%). 1H NMR (300 MHz, C_6D_6 , 20 °C): δ 14.22 (s br, 2H; NH), 2.09 (s, 30H; C_5Me_5), 1.88 (s, 15H; C_5Me_5), 0.15 (s, 9H; $NSiMe_3$). $^{13}C\{^1H\}$ NMR (75 MHz, C_6D_6 , 20 °C): δ 118.2 (C_5Me_5), 117.8 (C_5Me_5), 12.5 (C_5Me_5), 11.8 (C_5Me_5), 7.0 ($SiMe_3$). Compound **7** has been previously prepared in 59% yield by a different procedure.¹⁷

Synthesis of $[K\{(\mu_3-N)(\mu_3-NH)(\mu_3-NSiMe_3)Ti_3(\eta^5-C_5Me_5)_3(\mu_3-N)\}]$ (8**).** A 100 mL ampule (Teflon stopcock) was charged with **5** (0.30 g, 0.36 mmol), $[K\{N(SiMe_3)_2\}]$ (0.16 g, 0.80 mmol), and toluene (30 mL). The reaction mixture was stirred at 85 °C for 24 h. After filtration, the volatile components of the solution were removed under reduced pressure to give **8** as a red solid (0.23 g, 89%). IR (KBr, cm^{-1}): $\tilde{\nu}$ 3325 (w), 2905 (s), 2856 (s), 2720 (w), 1496 (w), 1437 (m), 1374 (m), 1257 (m), 1244 (s), 1095 (w), 1065 (w), 1023 (m), 956 (vs), 822 (s), 731 (vs), 702 (s), 662 (s), 629 (s), 593 (w), 548 (w), 510 (w), 473 (m), 415 (w). 1H NMR (300 MHz, C_6D_6 , 20 °C): δ 13.34 (s br, 1H; NH), 2.30 (s, 15H; C_5Me_5), 2.13 (s, 15H; C_5Me_5), 2.11 (s, 15H; C_5Me_5), 0.11 (s, 9H; $NSiMe_3$). 1H NMR (300 MHz, C_5D_5N , 20 °C): δ 13.92 (s br, 1H; NH), 2.28 (s, 15H; C_5Me_5), 2.15 (s, 15H; C_5Me_5), 2.05 (s, 15H; C_5Me_5), 0.30 (s, 9H; $NSiMe_3$). $^{13}C\{^1H\}$ NMR (75 MHz, C_6D_6 , 20 °C): δ 116.6 (C_5Me_5), 115.9 (C_5Me_5), 113.9 (C_5Me_5), 13.0 (C_5Me_5), 12.8 (C_5Me_5), 12.5 (C_5Me_5), 9.0 ($SiMe_3$). $^{13}C\{^1H\}$ NMR (75 MHz, C_5D_5N , 20 °C): δ 115.4 (C_5Me_5), 114.4 (C_5Me_5), 113.3 (C_5Me_5), 12.7 (C_5Me_5), 12.5 (C_5Me_5), 12.0 (C_5Me_5), 8.4 ($SiMe_3$). Anal. Calcd for $C_{33}H_{55}KN_4SiTi_3$ ($M_w = 718.60$): C, 55.16; H, 7.71; N, 7.80. Found: C, 54.89; H, 7.90; N, 7.76.

Synthesis of $[K(18-crown-6)][Ti_3(\eta^5-C_5Me_5)_3(\mu_3-N)(\mu-N)(\mu-NH)(\mu-NSiMe_3)]$ (9**).** A toluene solution (10 mL) of 18-crown-6 (0.074 g, 0.28 mmol) was added to a solution of **8** (0.20 g, 0.28 mmol) in toluene (15 mL). The reaction mixture was stirred at room temperature for 5 min to give an abundant yellow solid. The solid was isolated by filtration onto a glass frit and vacuum-dried to afford **9** as a yellow powder (0.22 g, 80%). IR (KBr, cm^{-1}): $\tilde{\nu}$ 3348 (w), 2901 (vs), 2747 (m), 2715 (m), 1496 (w), 1472 (m), 1452 (m), 1372 (m), 1352 (s), 1286 (w), 1249 (s), 1230 (s), 1105 (vs), 960 (s), 939 (s), 826 (s), 744 (vs), 686 (s), 623 (s), 514 (m), 487 (m), 441 (w). 1H NMR (300 MHz, C_5D_5N , 20 °C): δ 14.09 (s br, 1H; NH), 3.41 (s, 24H; OCH_2), 2.44 (s, 15H; C_5Me_5), 2.31 (s, 15H; C_5Me_5), 2.26 (s, 15H; C_5Me_5), 0.48 (s, 9H; $NSiMe_3$). $^{13}C\{^1H\}$ NMR (75 MHz, C_5D_5N , 20 °C): δ 113.9 (C_5Me_5), 113.0 (C_5Me_5), 112.6 (C_5Me_5), 70.0 (OCH_2), 12.8 (C_5Me_5), 12.6 (C_5Me_5), 12.3 (C_5Me_5), 8.3 ($SiMe_3$). Anal. Calcd for $C_{45}H_{79}KN_4O_6SiTi_3$ ($M_w = 982.91$): C, 54.99; H, 8.10; N, 5.70. Found: C, 55.17; H, 7.91; N, 5.30.

Table 1. Experimental Data for the X-ray Diffraction Studies on Compounds 2, 5, 6, and 8

	2	5	6	8
formula	C ₃₁ H ₄₉ F ₃ N ₄ O ₃ STi ₃	C ₃₄ H ₅₇ F ₃ N ₄ O ₃ SSiTi ₃	C ₃₃ H ₅₆ N ₄ SiTi ₃	C ₃₃ H ₅₅ KN ₄ SiTi ₃
<i>M_r</i>	758.50	830.69	680.61	718.70
<i>T</i> [K]	200(2)	200(2)	200(2)	200(2)
<i>λ</i> [Å]	0.71073	0.71073	0.71073	0.71073
crystal system	monoclinic	orthorhombic	triclinic	monoclinic
space group	<i>P</i> 2 ₁ / <i>c</i>	<i>Pbca</i>	<i>P</i> $\bar{1}$	<i>P</i> 2 ₁ / <i>c</i>
<i>a</i> [Å]; <i>α</i> [deg]	11.864(2)	19.579(14)	11.293(6); 95.11(3)	10.918(9)
<i>b</i> [Å]; <i>β</i> [deg]	19.143(4); 117.84(1)	20.000(6)	12.691(1); 100.16(3)	16.303(8); 115.38(4)
<i>c</i> [Å]; <i>γ</i> [deg]	18.043(3)	21.224(1)	12.788(7); 98.30(3)	22.856(11)
<i>V</i> [Å ³]	3264(1)	8311(6)	1773(1)	3676(4)
<i>Z</i>	4	8	2	4
<i>ρ</i> _{calcd} [g cm ^{−3}]	1.390	1.328	1.275	1.299
<i>μ</i> _{MoKα} [mm ^{−1}]	0.754	0.691	0.720	0.809
<i>F</i> (000)	1584	3488	724	1520
crystal size [mm ³]	0.22 × 0.20 × 0.17	0.16 × 0.15 × 0.14	0.27 × 0.23 × 0.20	0.25 × 0.13 × 0.12
<i>θ</i> range (deg)	3.19–27.51	3.01–26.17	3.26–27.50	3.06–27.51
index ranges	−15 to 15 −24 to 24 −15 to 23	−24 to 24 −24 to 24 −26 to 26	−14 to 14 −15 to 16 −16 to 16	−14 to 14 −21 to 21 −29 to 14
reflns collected	86 270	225 003	71 530	73 443
unique data	8304 [<i>R</i> (int) = 0.051]	8226 [<i>R</i> (int) = 0.197]	8151 [<i>R</i> (int) = 0.073]	8449 [<i>R</i> (int) = 0.097]
obsd data [<i>I</i> > 2σ(<i>I</i>)]	5593	5083	5849	4692
GOF on <i>F</i> ²	1.035	1.038	1.055	1.015
final <i>R</i> ^a indices [<i>I</i> > 2σ(<i>I</i>)]	<i>R</i> 1 = 0.049 w <i>R</i> 2 = 0.110	<i>R</i> 1 = 0.073 w <i>R</i> 2 = 0.172	<i>R</i> 1 = 0.052 w <i>R</i> 2 = 0.136	<i>R</i> 1 = 0.068 w <i>R</i> 2 = 0.133
<i>R</i> ^a indices (all data)	<i>R</i> 1 = 0.090 w <i>R</i> 2 = 0.124	<i>R</i> 1 = 0.131 w <i>R</i> 2 = 0.219	<i>R</i> 1 = 0.083 w <i>R</i> 2 = 0.150	<i>R</i> 1 = 0.144 w <i>R</i> 2 = 0.155
largest diff. peak/hole [e Å ^{−3}]	0.349 and −0.623	0.787 and −0.597	0.504 and −0.565	0.991 and −0.496

$$^a R1 = \sum |F_o| - |F_c| / \sum |F_o|; wR2 = \{[\sum w(F_o^2 - F_c^2)^2] / [\sum w(F_o^2)^2]\}^{1/2}.$$

X-ray Structure Determination of 2, 5, 6, and 8. Crystals of complexes 2 and 6 were grown from toluene solutions at −35 °C. Crystals of complexes 5 and 8 were obtained from toluene solutions at room temperature. The crystals were removed from the Schlenk flasks and covered with a layer of a viscous perfluoropolyether (FomblinY). A suitable crystal was selected with the aid of a microscope, mounted on a cryoloop, and immediately placed in the low temperature nitrogen stream of the diffractometer. The intensity data sets were collected at 200 K on a Bruker-Nonius KappaCCD diffractometer equipped with an Oxford Cryostream 700 unit. Crystallographic data for all of the complexes are presented in Table 1.

The structures were solved, using the WINGX package,¹⁸ by direct methods (SHELXS-97) and refined by least-squares against *F*² (SHELXL-97).¹⁹ In the crystallographic study of compound 2, all non-hydrogen atoms were anisotropically refined. All of the hydrogen atoms were positioned geometrically and refined by using a riding model, except those of the imido (H(12) and H(23)) and amido (H(13a) and H(13b)) groups, which were located in the difference Fourier map and refined isotropically. Moreover, SADI restraints were employed for the lengths N(12)–H(12) and N(23)–H(23).

Anion CF₃SO₃[−] in complex 5 presented disorder, which was treated conventionally by using the PART tool of the SHELXL-97 program and allowing free refinement of the occupancy factors with the FVAR command. The final values of occupancy were 55% and 45%. All non-hydrogen atoms were anisotropically refined, except fluorine and oxygen atoms for the triflate group (F(1), F(2), F(3), F(1'), F(2'), F(3'), O(1), O(2), O(3), O(1'), O(2'), and O(3')), which were refined isotropically. The hydrogen atoms were positioned geometrically and refined using a riding model. Furthermore, DFIX restraints were used for the triflate anion.

Crystals of the complex 6 showed disorder for the carbon atoms C(11)–C(20) of the pentamethylcyclopentadienyl ligand linked to Ti(1). This disorder was also treated using the PART tool, and the

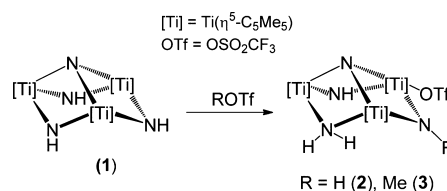
final values of occupancy were 62.0% and 38.0%. All non-hydrogen atoms were anisotropically refined. The amido group hydrogen atom, H(12), was located in the difference Fourier map and refined isotropically, whereas the rest of the hydrogen atoms were positioned geometrically and refined using a riding model. The imido hydrogen atom was distributed over the nitrogen atoms N(13) and N(23) with 50% of occupancy for each position. The nondisordered pentamethylcyclopentadienyl groups, linked to Ti(2) and Ti(3), were restrained with DELU instructions.

Finally, the solid-state structure of compound 8 revealed a one-dimensional polymeric network. All of the non-hydrogen atoms were anisotropically refined, whereas all of the hydrogen atoms were positioned geometrically and refined by using a riding model, except that of the imido group (H(23)), which was located in the difference Fourier map and refined isotropically.

RESULTS AND DISCUSSION

The treatment of [$\{\text{Ti}(\eta^5\text{-C}_5\text{Me}_5)(\mu\text{-NH})\}_3(\mu_3\text{-N})$] (1) with 1 equiv of triflic acid or methyl triflate in toluene at room temperature afforded the precipitation of complexes [$\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})_2(\mu\text{-NH}_2)(\text{OSO}_2\text{CF}_3)$] (2) or [$\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})(\mu\text{-NH}_2)(\mu\text{-NMe})(\text{OSO}_2\text{CF}_3)$] (3)

Scheme 1. Reaction of 1 with 1 equiv of ROTf (R = H, Me)



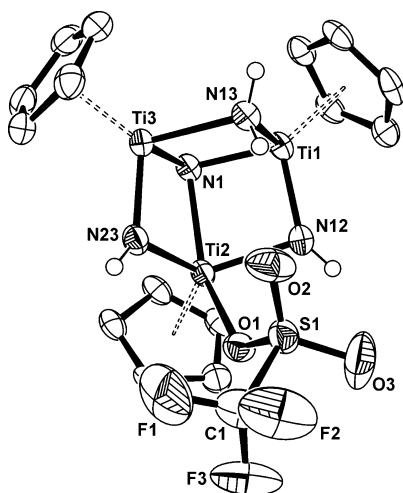


Figure 1. Perspective view of complex **2** (thermal ellipsoids at the 50% probability level). The methyl groups of the pentamethylcyclopentadienyl ligands are omitted for clarity.

(Scheme 1). Compounds **2** and **3** were isolated in good yields (84% and 78%, respectively) as yellow or orange solids, which are very soluble in halogenated solvents, moderately soluble in toluene or benzene, and insoluble in hexane.

Complexes **2** and **3** were characterized by analytical and spectroscopic methods, as well as by X-ray crystal structure determinations. The solid-state structure of **2** is shown in Figure 1, which is almost identical to that reported for complex **3** with the substitution of the methylimido group for a μ -NH ligand.¹⁶ Selected distances and angles for both structures are compared in Table 2, showing very similar values for the two compounds. The crystal structures show six-membered Ti_3N_3 rings in chair conformation with the three titanium atoms also bridged by a further nitrogen atom. Two of the titanium atoms, Ti(1) and Ti(3), have classical three-legged piano-stool arrangements, where the legs are occupied by one μ - NH_2 amido, one μ_3 -N nitrido, and a μ -NR imido ligand. The Ti(2)

Scheme 2. Schematic Representation of the Bonding Situation in Complexes 2 and 3

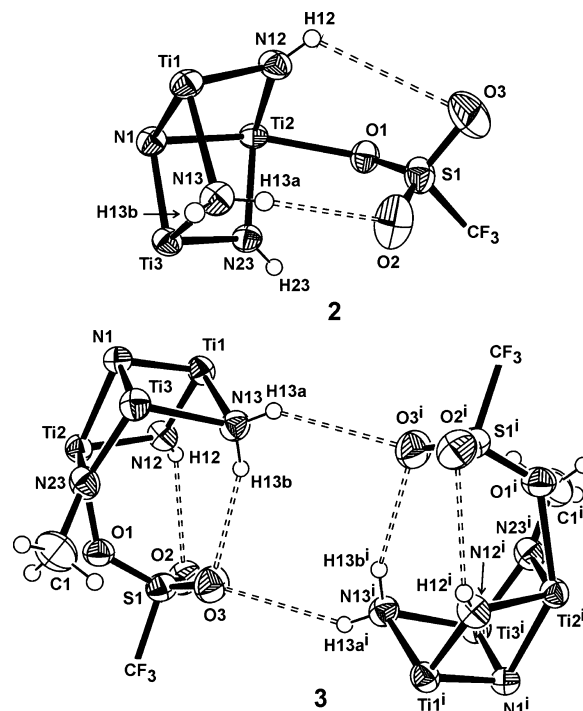
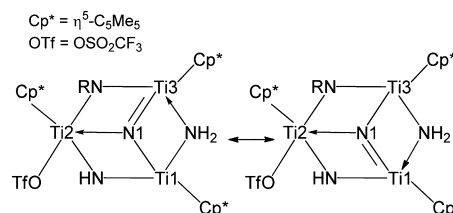


Figure 2. Simplified view of the intra- and intermolecular hydrogen-bonding interactions present in complexes **2** and **3**. Symmetry code: (i) $-x, 2-y, -z$.

Table 2. Selected Lengths (Å) and Angles (deg) for Complexes 2 and 3

lengths	2	3	angles	2	3
Ti(1)–N(1)	1.866(2)	1.868(3)	N(1)–Ti(1)–N(12)	90.3(1)	88.7(1)
Ti(1)–N(12)	1.858(3)	1.831(3)	N(1)–Ti(1)–N(13)	85.3(1)	85.1(1)
Ti(1)–N(13)	2.107(3)	2.102(3)	N(12)–Ti(1)–N(13)	101.0(1)	102.2(1)
Ti(2)–N(1)	2.137(2)	2.084(3)	N(1)–Ti(2)–N(12)	79.6(1)	78.9(1)
Ti(2)–N(12)	1.986(2)	1.983(3)	N(1)–Ti(2)–N(23)	78.4(1)	80.5(1)
Ti(2)–N(23)	1.975(2)	2.000(3)	N(12)–Ti(2)–N(23)	118.9(1)	118.8(1)
Ti(2)–O(1)	2.172(2)	2.157(3)	N(1)–Ti(2)–O(1)	145.0(1)	145.0(1)
Ti(3)–N(1)	1.877(2)	1.868(3)	N(12)–Ti(2)–O(1)	84.2(1)	82.1(1)
Ti(3)–N(13)	2.103(2)	2.099(3)	N(23)–Ti(2)–O(1)	82.9(1)	83.3(1)
Ti(3)–N(23)	1.844(3)	1.856(3)	N(1)–Ti(3)–N(13)	85.1(1)	85.2(1)
Ti(1)···Ti(2)	2.874(1)	2.866(1)	N(1)–Ti(3)–N(23)	88.7(1)	90.3(1)
Ti(1)···Ti(3)	2.910(1)	2.905(1)	N(13)–Ti(3)–N(23)	101.5(1)	101.2(1)
Ti(2)···Ti(3)	2.895(1)	2.853(1)	Ti(1)–N(1)–Ti(2)	91.5(1)	92.8(1)
N(23)–C(1)		1.467(5)	Ti(1)–N(1)–Ti(3)	102.1(1)	102.1(1)
			Ti(2)–N(1)–Ti(3)	92.1(1)	92.3(1)
			Ti(1)–N(12)–Ti(2)	96.8(1)	97.4(1)
			Ti(1)–N(13)–Ti(3)	87.5(1)	87.5(1)
			Ti(2)–N(23)–Ti(3)	98.5(1)	95.4(1)
			Ti(2)–N(23)–C(1)		124.4(3)
			Ti(3)–N(23)–C(1)		140.2(3)

Table 3. Relevant Hydrogen Bonds^a for Compounds 2 and 3

	D–H...A	D...A/Å	H...A/Å	D–H...A/deg
2	N(12)–H(12)...O(3)	3.574(5)	3.15(4)	118(3)
	N(13)–H(13a)...O(2)	3.198(4)	2.45(4)	164(4)
3	N(12)–H(12)...O(2)	3.130(4)		
	N(13)–H(13a)...O(3)	3.417(6)	2.54(5)	159(4)
	N(13)–H(13b)...O(3) ^b	3.576(6)	2.81(4)	146(3)

^aA = acceptor; D = donor. ^bSymmetry code: (i) $-x, 2 - y, -z$.

atoms exhibit four-legged piano-stool arrangements, in which the legs are occupied by one triflate, one nitrido, and two imido ligands.

The NH₂ amido group bridges the titanium(1) and titanium(3) atoms in a symmetric fashion with Ti–N(13) bond lengths, 2.099(3)–2.107(3) Å, which are about 0.25 Å longer than the distances between those titanium atoms and the imido ligands, Ti(1)–N(12) and Ti(3)–N(23), of 1.831(3)–1.858(3) Å. The nitrido group N(1) also bridges symmetrically the Ti(1) and Ti(3) atoms with values of titanium–nitrogen bond lengths (1.866(2)–1.877(2) Å) in the upper limit of those found in titanium dinuclear complexes containing Ti=N=Ti fragments (1.763(5)–1.878(7) Å),^{11d,12b,20} and substantially shorter than the Ti(2)–N(1) separations (2.137(2) Å in 2 and 2.084(3) Å for 3). A plausible interpretation of the bonding in complexes 2 and 3 is illustrated in Scheme 2. The bonding system can be described by two resonance forms in valence bond theory terms involving one titanium–nitrogen(1) double bond within the flat ring containing the Ti(1), Ti(3), N(1), and N(13) atoms. The first of the contributing structure represents a double bond between the Ti(3) and N(1) atoms with concomitant dative N(13)→Ti(3) bond, whereas the second structure comprises Ti(1)=N(1) and N(13)→Ti(1) units.

The triflate group is linked to titanium(2) with Ti(2)–O(1) bond lengths of 2.172(2) Å in 2 and 2.157(3) Å for 3, which are longer than the Ti–O distances found in other titanium derivatives with terminal triflate ligands (1.938(1)–2.142(3) Å).²¹ The remaining oxygen atoms of the trifluoromethanesulfonato groups, O(2) and O(3), are involved in N–H...O intramolecular hydrogen bonds with the *endo* hydrogen of the μ -NH₂ amido and the μ -NH imido ligands (Figure 2, Table 3). Those interactions can be classified as weak according to the criteria on the donor–acceptor distances.²² In addition, complex 3 displays intermolecular hydrogen-bonding interactions between the O(3) atom and the *exo* hydrogen of the μ -NH₂ amido ligand to result in molecules linked in pairs across a crystallographic symmetry center (Figure 2).

The IR spectra (KBr) of compounds 2 and 3 show several bands in the ν_{NH} region, between 3374 and 3250 cm^{−1}, for the NH and NH₂ ligands. In addition, the spectra of 2 and 3 reveal

one absorption at 1578 and 1590 cm^{−1}, respectively, assignable to the NH₂ bending mode.²³ Also, the IR spectra show several strong absorptions in the range 1315–1011 cm^{−1} for the coordinated triflate ligands.²⁴ Surprisingly, the ¹H and ¹³C{¹H} NMR spectra of solutions of crystals of complexes 2 and 3 in benzene-*d*₆ or chloroform-*d*₁ at room temperature are different from those expected from their solid-state structures. Thus, the ¹H NMR spectra of 2 show only one broad singlet for the η^5 -C₅Me₅ ligands, instead of the two 2:1 signals expected from the nearly C_s symmetry in the solid state. Furthermore, while the ¹H NMR spectra of 3 reveal the resonance signals predicted for the C₁ symmetry determined in the crystal structure, the resonances for two η^5 -C₅Me₅ ligands and those assigned to the NH and NH₂ groups are broad.¹⁶ These NMR data suggest a dynamic exchange process in solution, and ¹H NMR spectra of dichloromethane-*d*₂ solutions were taken at low temperatures in a 500 MHz spectrometer. Indeed, the ¹H NMR spectrum of 2 at −50 °C is consistent with a C_s symmetric structure in solution, while the spectrum of 3 at −30 °C reveals sharp resonances for the η^5 -C₅Me₅ and μ -NH₂ ligands.

The dynamic behavior may be the result of the proton transfer from the NH₂ group to the hydrogen-bonded triflate ligand and generation of HOTf, which then delivers that proton to the NH imido group with concomitant coordination of the triflate ligand at the opposite titanium atom. Such rapid exchange would create time-averaged C_{3v} symmetry for 2 and C_s symmetry for 3 by ¹H NMR spectroscopy at high temperatures. Complete dissociation of HOTf can be ruled out because addition of small amounts of 1 to solutions of compounds 2 or 3 does not affect the exchange process, and the activation parameters remain unaltered. A plausible mechanism for the dynamic event is shown in Scheme 3, which is similar to that studied in detail by Limbach and co-workers for the proton exchange process in diaryltriazenes catalyzed by the presence of bases in the medium.²⁵ In our complexes, the function of the basic catalyst could be played by the triflate ligand, which is prone to dissociate from the metal centers.²⁶

The kinetic parameters (Table 4) of the observed process were calculated on the basis of dynamic ¹H NMR spectroscopy data with line shape analysis of the η^5 -C₅Me₅ resonances using the gNMR program (see Tables S1–S4 and Figures S1–S8 in the Supporting Information).²⁷ The results disagree with those expected for an intramolecular nondissociative mechanism because the log A (8.5–9.8) and $\Delta S^\ddagger$ (between −7.7 and −21.3 eu) values differ from the usual parameters for that type of process (log A = 12–14 and $\Delta S^\ddagger \approx 0$ eu).²⁸ However, in accord with the highly negative values of $\Delta S^\ddagger$, the reduction of the log A in several units could be related to the organization of the solvent molecules due to the participation of zwitterionic species in the exchange process,^{25b,29} as shown in Scheme 3.

Scheme 3. Proposed Fluxional Process for Complexes 2 and 3

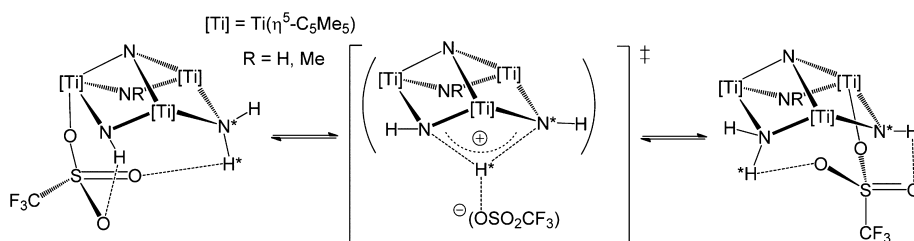


Table 4. Activation Parameters for the Exchange of η^5 -C₅Me₅ Resonances in Complexes 2 and 3

compound	log A	E_a [kcal mol ⁻¹]	$\Delta H^\ddagger$ [kcal mol ⁻¹]	$\Delta S^\ddagger$ [cal mol ⁻¹ K ⁻¹]	$\Delta G^\ddagger_{298K}$ [kcal mol ⁻¹]
2 ^a	8.5 ± 1.2	8.4 ± 0.3	7.9 ± 0.3	-21.3 ± 1.0	14.2
2 ^b	9.8 ± 0.8	9.7 ± 0.2	9.2 ± 0.2	-15.6 ± 0.7	13.9
2 ^c	8.5 ± 2.1	8.4 ± 0.5	11.9 ± 0.5	-7.7 ± 1.8	14.2
3 ^a	8.9 ± 1.7	9.5 ± 0.4	8.9 ± 0.4	-19.7 ± 1.5	14.8

^aCD₂Cl₂, ~1.6 × 10⁻² M. ^bCD₂Cl₂, ~5.3 × 10⁻³ M. ^cC₇D₈, ~1.1 × 10⁻² M.

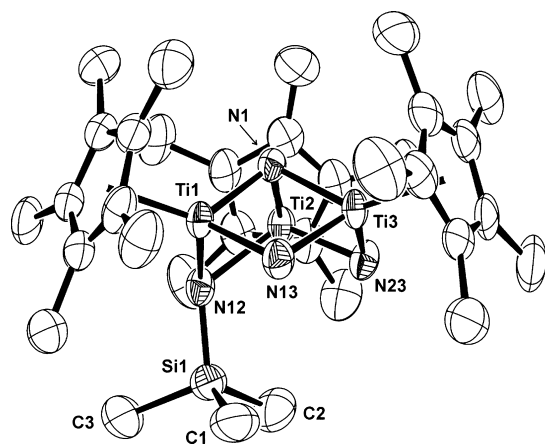
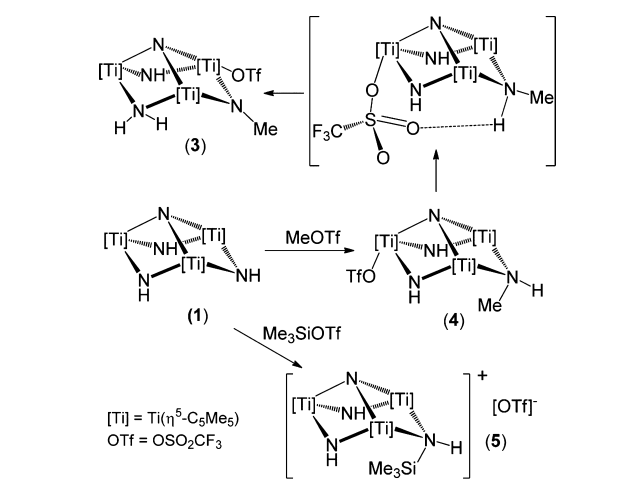
Scheme 4. Reaction of 1 with 1 equiv of ROTf (R = Me, SiMe₃)

Figure 3. Perspective view of the cationic fragment of complex 5 (thermal ellipsoids at the 50% probability level). Hydrogen atoms are omitted for clarity.

Indeed, when the kinetic parameters of complex 2 are obtained in toluene-*d*₈, the entropy of activation falls to -7.7 eu in agreement with that expected for a less polar solvent. Importantly, the $\Delta G^\ddagger$ values calculated in both solvents are comparable, suggesting that the exchange process is the same and the entropy differences are compensated by the enthalpy associated.²⁹

An analogue proton migration within the trinuclear titanium system could explain the formation of complex 3 in the treatment of 1 with 1 equiv of MeOTf (Scheme 4). Monitoring by NMR spectroscopy the reaction course in benzene-*d*₆ showed that the alkylation occurs at the NH imido ligands of 1 to produce the methylamido derivative [Ti₃(η^5 -C₅Me₅)₃(μ_3 -N)(μ -NH)₂(μ -NHMe)(OTf)] (4).¹⁶ Compound 4 readily rearranges at room temperature to form complex 3 (conversion

is ca. 25% after 2 h, and 100% after 24 h). Thus, complex 4 could not be obtained in a pure form and was only characterized by ¹H, ¹³C{¹H}, and ¹⁹F NMR spectroscopy. The NMR data are consistent with a C_s symmetric structure with the methylamido and triflate ligands in the mirror plane of the molecule. NOESY-1D experiments on a solution of 4 in benzene-*d*₆ provided evidence for the *endo* position of the methyl group for the μ -NHMe ligand. The rotation of the amido ligand in complex 4 could place the methyl fragment in the *exo* position, while the new *endo* hydrogen interacts with the triflate ligand. From this undetected intermediate, the migration of the proton from the NHMe amido group to the NH imido ligands, assisted by the triflate moiety, would lead to compound 3.

To isolate a stable compound with a structure similar to 4, we studied the reaction of [Ti(η^5 -C₅Me₅)(μ -NH)]₃(μ_3 -N) (1) with an electrophilic reagent ROTf bearing a bulkier R fragment. Thus, the treatment of 1 with 1 equiv of trimethylsilyl triflate in toluene at room temperature afforded the precipitation of the ionic complex [Ti₃(η^5 -C₅Me₅)₃(μ_3 -N)(μ -NH)₂(μ -NHSiMe₃)] [O₃SCF₃] (5) (Scheme 4). This compound was obtained in 88% yield as an orange solid, which is poorly soluble in toluene or benzene but exhibits a good solubility in halogenated solvents. Solutions of 5 in chloroform-*d*₁ slowly decompose within several days at room temperature to give free trimethylsilylamine and several unidentified products as determined by ¹H NMR spectroscopy. The IR spectrum (KBr) of 5 shows only one broad band centered at 3271 cm⁻¹ for the ν_{NH} vibrations and several strong absorptions in the range 1252–1031 cm⁻¹ for the triflate group.²⁴ In particular, the observed $\nu_{as}(\text{SO}_3)$ band at 1252 cm⁻¹ is typical for the free trifluoromethanesulfonate anion.^{24b} The ¹H NMR spectrum in chloroform-*d*₁ at room temperature reveals two resonances for η^5 -C₅Me₅ ligands in a 2:1 ratio, two broad signals for the μ -NH and μ -NHSiMe₃ protons, and a high-field singlet for the trimethylsilyl group. These data agree with a C_s symmetric structure in solution, which is also consistent with the ¹³C{¹H} NMR spectrum. NOESY-1D experiments confirmed the *endo* disposition of the trimethylsilyl group in a fashion similar to the methyl fragment in 4, but the greater steric bulk of the SiMe₃ group precludes the coordination of the triflate to yield 5 as an ion pair, at least in the solid state.

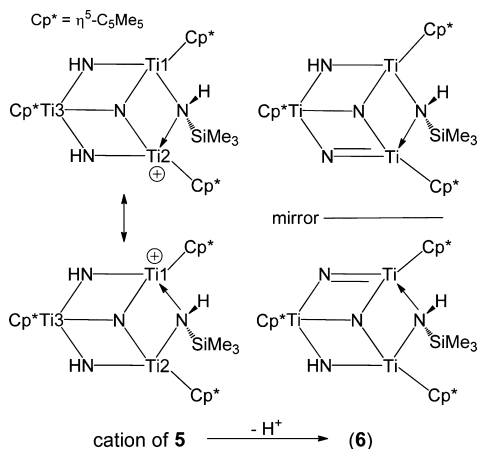
The X-ray crystal structure of 5 shows each trifluoromethanesulfonate anion linking through hydrogen-bonding interactions two trinuclear titanium cations to give polymeric zigzag chains (see Figure S9 in the Supporting Information). Thus, each triflate anion establishes a hydrogen bond between the O(1) atom and the NH ligand of one cation (N(13)⋯O(1) = 3.322(1) Å), while O(3) interacts with the NH group of a contiguous cation (N(23)⋯O(3) = 3.311(1) Å). The cationic fragment of compound 5 is presented in Figure 3, while selected distances and angles are given in Table 5.

The cation of 5 contains a [Ti₃N₃] six-membered ring that adopts a chair conformation similar to that of 1,^{12a} and those

Table 5. Selected Lengths (Å) and Angles (deg) for Complexes 5 and 6

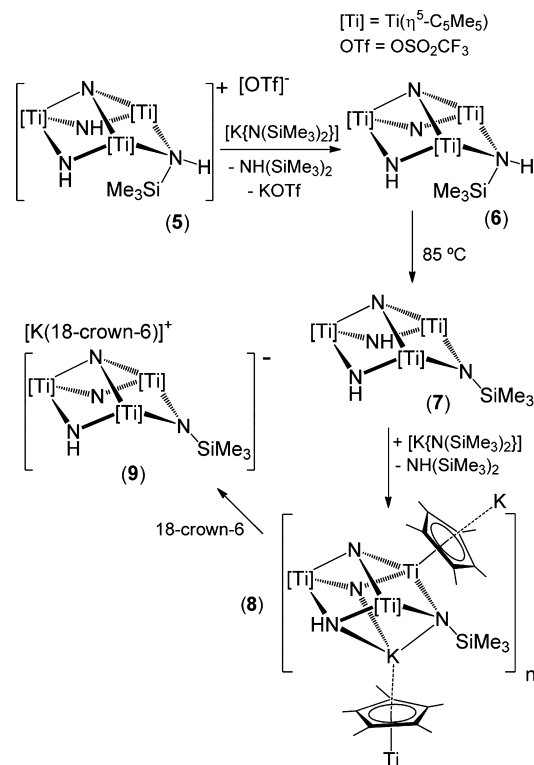
lengths	5	6	angles	5	6
Ti(1)–N(1)	1.921(4)	1.878(2)	N(1)–Ti(1)–N(12)	85.9(2)	86.8(1)
Ti(1)–N(12)	2.121(5)	2.105(2)	N(1)–Ti(1)–N(13)	87.3(2)	89.1(1)
Ti(1)–N(13)	1.866(4)	1.817(2)	N(12)–Ti(1)–N(13)	103.3(2)	103.6(1)
Ti(2)–N(1)	1.917(5)	1.914(2)	N(1)–Ti(2)–N(12)	86.3(2)	85.9(1)
Ti(2)–N(12)	2.111(5)	2.103(2)	N(1)–Ti(2)–N(23)	88.0(2)	88.4(1)
Ti(2)–N(23)	1.868(4)	1.793(3)	N(12)–Ti(2)–N(23)	105.2(2)	103.8(1)
Ti(3)–N(1)	1.938(4)	1.957(2)	N(1)–Ti(3)–N(13)	84.0(2)	83.8(1)
Ti(3)–N(13)	1.971(5)	1.923(2)	N(1)–Ti(3)–N(23)	84.4(2)	84.4(1)
Ti(3)–N(23)	1.978(5)	1.892(2)	N(13)–Ti(3)–N(23)	110.4(2)	109.9(1)
N(12)–Si(1)	1.808(5)	1.761(2)	Ti(1)–N(1)–Ti(2)	99.8(2)	99.6(1)
Ti(1)⋯Ti(2)	2.935(2)	2.896(1)	Ti(1)–N(1)–Ti(3)	93.8(2)	91.7(1)
Ti(1)⋯Ti(3)	2.818(1)	2.753(2)	Ti(2)–N(1)–Ti(3)	93.3(2)	90.4(1)
Ti(2)⋯Ti(3)	2.804(1)	2.746(1)	Ti(1)–N(12)–Ti(2)	87.8(2)	87.0(1)
			Ti(1)–N(12)–Si(1)	124.0(2)	124.5(1)
			Ti(2)–N(12)–Si(1)	120.6(2)	121.0(1)
			Ti(1)–N(13)–Ti(3)	94.5(2)	94.7(1)
			Ti(2)–N(23)–Ti(3)	93.5(2)	96.3(1)

Scheme 5. Schematic Representation of the Bonding Situation in the Cation of 5 and Complex 6



described above for complexes 2 and 3. In contrast to compounds 2 and 3, the cationic fragment of 5 shows the nitrido atom N(1) bridging the titanium atoms with the three Ti–N(1) distances within a narrow range, 1.917(5)–1.938(4) Å, which compares well with that determined in complex 1 (Ti–N(1) 1.912(1) Å). The NHSiMe₃ amido group in 5 bridges the Ti(1) and Ti(2) atoms in a symmetric fashion with Ti–N(12) bond lengths, 2.111(5) and 2.121(5) Å, very similar to the Ti–N separations found for the NH₂ amido ligand in complexes 2 and 3. Each NH imido ligand of 5 bridges asymmetrically two titanium atoms with Ti(1)–N(13) and Ti(2)–N(23) bond lengths, 1.866(4) and 1.868(4) Å, clearly shorter than those found with the Ti(3) atom, Ti(3)–N(13) 1.971(5) and Ti(3)–N(23) 1.978(5) Å. Therefore, the solid-state structure of the cation of 5 is very close to the C_s symmetry determined in solution by NMR spectroscopy. A plausible interpretation of the bonding in the trinuclear cation is illustrated on the left side of Scheme 5. The bonding system can be described by two resonance forms in valence bond theory terms containing the positive charge on titanium(1) or titanium(2) atoms.

The treatment of 5 with 1 equiv of potassium bis(trimethylsilyl)amide in toluene at room temperature leads to

Scheme 6. Reactions with [K{N(SiMe₃)₂}]

the molecular complex [Ti₃(η⁵-C₅Me₅)₃(μ₃-N)(μ-N)(μ-NH)(μ-NHSiMe₃)] (6), NH(SiMe₃)₂, and KOTf (Scheme 6). Compound 6 was isolated as a red solid in 69% yield if the reaction and subsequent workup were performed within 1 h. Reactions carried out at room temperature for longer periods of time afforded mixtures of compound 6 and the trimethylsilylimido derivative [Ti₃(η⁵-C₅Me₅)₃(μ₃-N)(μ-NH)₂(μ-NSiMe₃)] (7). Indeed, complex 7 was isolated as an orange solid in 69% yield if the reaction mixture of 5 and [K{N(SiMe₃)₂}] in toluene was heated at 85 °C for 24 h. Compound 7 has been previously obtained by treatment of [Li(μ₄-N)(μ₃-NH)₂Ti₃(η⁵-C₅Me₅)₃(μ₃-N)]₂ with [SiClMe₃] in toluene at room temperature.¹⁷

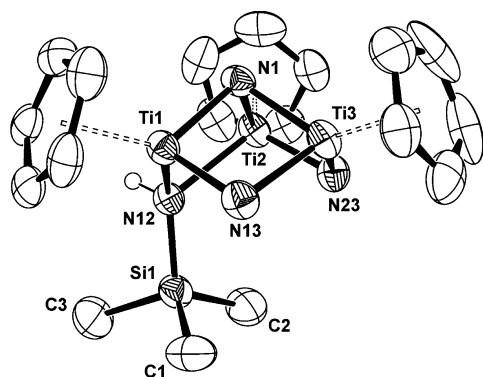


Figure 4. Perspective view of complex **6** (thermal ellipsoids at the 50% probability level). Methyl groups of the pentamethylcyclopentadienyl ligands and hydrogen atoms of the imido groups are omitted for clarity.

Compound **6** was characterized by spectroscopic and analytical methods, as well as by an X-ray crystal structure determination. The IR spectrum (KBr) of **6** shows two ν_{NH} vibrations at 3353 and 3273 cm^{-1} for the NH and NHSiMe_3 ligands. The ^1H NMR spectrum of **6** in benzene- d_6 at room temperature reveals three resonance signals for the $\eta^5\text{-C}_5\text{Me}_5$ ligands in a 1:1:1 ratio, one broad signal for one $\mu\text{-NH}$ imido group, and those assigned to the NHSiMe_3 trimethylsilylamido ligand. The NMR data of **6** are consistent with a C_1 symmetric structure in solution, in contrast to the C_s symmetry observed in the NMR spectra of **7**.¹⁷ The solid-state structure of **6** is presented in Figure 4, while selected distances and angles are given in Table 5. The $[\text{Ti}_3\text{N}_4]$ core of **6** is similar to that described above for the cation of complex **5**. Thus, the nitrido ligand N(1) bridges the three titanium centers with Ti–N(1) bond lengths in the range 1.878(2)–1.957(2) Å, and the amido NHSiMe_3 ligand is bonded to the Ti(1) and Ti(2) atoms with distances Ti(1)–N(12) and Ti(2)–N(12) of 2.105(2) and 2.103(2) Å, respectively. While these bond lengths are very close to those observed in **5**, complex **6** shows a shortening of the Ti–N distances associated with the N(13) and N(23) atoms as expected for the deprotonation of one NH imido group to yield a $\mu\text{-N}$ nitrido ligand. The Ti(1)–N(13) and Ti(2)–N(23) bond lengths, 1.817(2) and 1.793(2) Å, respectively, are about 0.1 Å shorter than the titanium–nitrogen distances associated to the titanium(3) atom, Ti(3)–N(13) 1.923 Å and Ti(3)–N(23) 1.892(2) Å. A plausible interpretation of these distances, and the nearly C_s symmetric

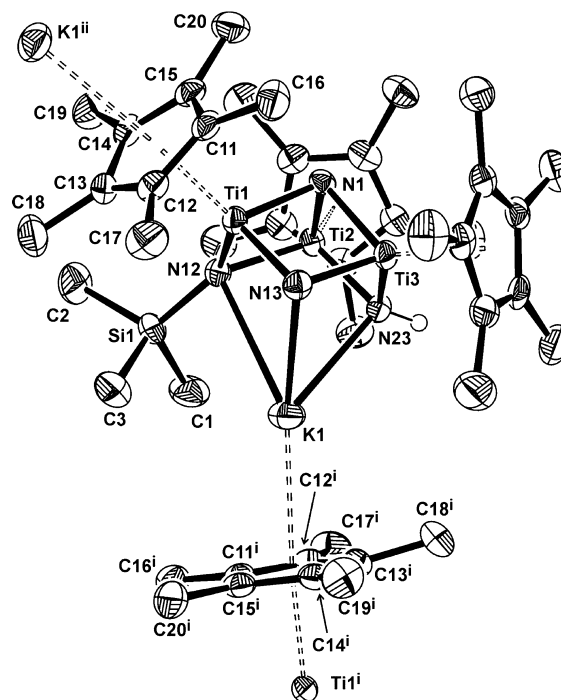


Figure 6. Perspective view of a portion of the polymeric chain of compound **8** (thermal ellipsoids at the 50% probability level). Hydrogen atoms of the pentamethylcyclopentadienyl groups are omitted for clarity. Symmetry code: (i) $-x, 1/2 + y, -1/2 - z$; (ii) $-x, -1/2 + y, -1/2 - z$.

structure of **6** determined by X-ray crystallography, is the existence of an equimolecular mixture of the two enantiomers originated from proton abstraction of complex **5** as shown in the right side of Scheme 5.

The conversion of **6** to complex **7** in solution at room temperature was monitored by ^1H NMR spectroscopy. Once **6** is isolated in a pure form, the rearrangement is slow in benzene- d_6 solution (ca. 10% after 3 days). While addition of tetrahydrofuran (4 equiv) to a benzene- d_6 solution of **6** does not affect the reaction rate, the presence of residual free $\text{NH}(\text{SiMe}_3)_2$ in the solution produces a significant enhancement of the transformation. Furthermore, upon dissolving **6** in chloroform- d_1 , there is an immediate color change from red to orange, and the ^1H NMR spectrum shows resonances assigned to **7**. Complete consumption of **6** in chloroform- d_1 occurs within 24 h to give complex **7**, with partial incorporation of

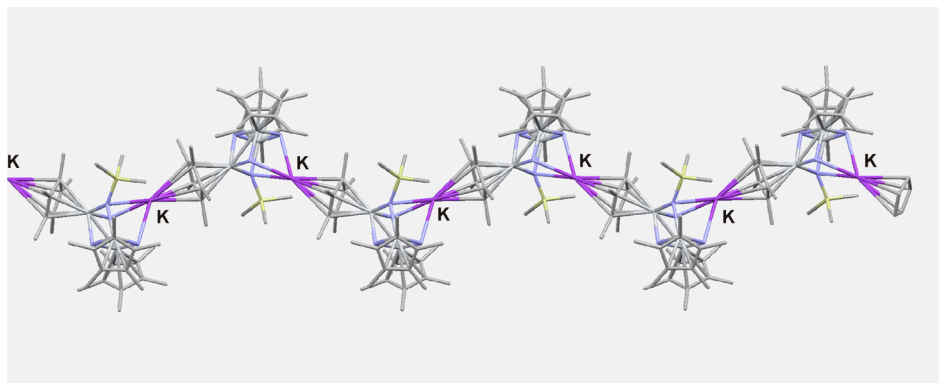


Figure 5. Perspective view of the polymeric chain of **8**. Hydrogen atoms are omitted for clarity.

Table 6. Selected Lengths (Å) and Angles (deg) for Compound 8^a

K(1)–N(12)	3.108(4)	K(1)–N(13)	2.772(4)
K(1)–N(23)	2.776(4)	K(1)–C(11) ⁱ	3.258(4)
K(1)–C(12) ⁱ	3.168(4)	K(1)–C(13) ⁱ	3.006(4)
K(1)–C(14) ⁱ	3.000(4)	K(1)–C(15) ⁱ	3.171(4)
K(1)–Cm(1) ⁱ	2.882	Ti(1)–N(12)	1.980(3)
Ti(1)–N(13)	1.860(3)	Ti(1)–N(1)	1.916(3)
Ti(1)–Cm(1) ^b	2.147	Ti(2)–N(12)	1.961(3)
Ti(2)–N(23)	1.913(4)	Ti(2)–N(1)	1.905(3)
Ti(2)–Cm(2) ^b	2.105	Ti(3)–N(13)	1.883(3)
Ti(3)–N(23)	1.929(4)	Ti(3)–N(1)	1.957(3)
Ti(3)–Cm(3) ^b	2.126	Si(1)–N(12)	1.713(3)
Ti(1)···Ti(2)	2.805(2)	Ti(1)···Ti(3)	2.762(2)
Ti(2)···Ti(3)	2.804(3)		
N(12)–K(1)–N(13)	62.0(1)	N(12)–K(1)–N(23)	61.9(1)
N(13)–K(1)–N(23)	65.2(1)	N(12)–K(1)–Cm(1) ⁱ	143.2
N(13)–K(1)–Cm(1) ⁱ	152.9	N(23)–K(1)–Cm(1) ⁱ	128.9
N(12)–Ti(1)–N(13)	104.7(2)	N(12)–Ti(1)–N(1)	86.5(1)
N(13)–Ti(1)–N(1)	87.7(1)	N(12)–Ti(2)–N(23)	103.4(1)
N(12)–Ti(2)–N(1)	87.4(1)	N(23)–Ti(2)–N(1)	87.3(2)
N(13)–Ti(3)–N(23)	103.3(2)	N(13)–Ti(3)–N(1)	85.9(1)
N(23)–Ti(3)–N(1)	85.4(2)	Ti(1)–N(12)–Ti(2)	90.7(1)
Ti(1)–N(12)–K(1)	89.1(1)	Ti(2)–N(12)–K(1)	90.8(1)
Ti(1)–N(12)–Si(1)	133.6(2)	Ti(2)–N(12)–Si(1)	135.3(2)
K(1)–N(12)–Si(1)	85.1(1)	Ti(1)–N(13)–Ti(3)	95.1(2)
K(1)–N(13)–Ti(1)	102.5(1)	K(1)–N(13)–Ti(3)	92.9(1)
Ti(2)–N(23)–Ti(3)	93.7(2)	K(1)–N(23)–Ti(2)	102.6(1)
K(1)–N(23)–Ti(3)	91.8(1)	Ti(1)–N(1)–Ti(2)	94.5(1)
Ti(1)–N(1)–Ti(3)	91.0(1)	Ti(2)–N(1)–Ti(3)	93.1(2)

^aSymmetry code: (i) $-x, 1/2 + y, -1/2 - z$. ^bCm = centroid of the η^5 -C₅Me₅ groups.

deuterium at the μ -NH ligands, along with minor amounts (~7%) of compound 1. In view of these results, it appears that the rearrangement is catalyzed by a Brønsted acid HA, such as NH(SiMe₃)₂ or CHCl₃, which transfers the proton to the μ -N ligand of 6, and subsequently its conjugate base A[−] accepts the proton from the μ -NHSiMe₃ amido ligand to generate 7.

The reaction of 5 with 2 equiv of potassium bis-(trimethylsilyl)amide in toluene at 85 °C or the treatment of 7 with 1 equiv of [K{N(SiMe₃)₂}] at room temperature afforded the potassium derivative [K{(μ_3 -N)(μ_3 -NH)(μ_3 -NSiMe₃)Ti₃(η^5 -C₅Me₅)₃(μ_3 -N)}] (8) (Scheme 6). Compound 8 was isolated in 89% yield as a red solid, which is very soluble in benzene or toluene, suggesting a molecular structure in solution. ¹H and ¹³C{¹H} NMR spectra in benzene-*d*₆ or pyridine-*d*₅ show resonances for three different η^5 -C₅Me₅ ligands and are consistent with a C₁ symmetric structure in solution. The NMR data could also agree with an edge-linked double-cube structure similar to those documented in alkali metal derivatives [M{(μ_4 -N)(μ_3 -NH)₂Ti₃(η^5 -C₅Me₅)₃(μ_3 -N)}₂] (M = Li, Na, K, Rb, Cs).^{17,30} However, in contrast to those species that are only soluble in pyridine, the high solubility of 8 in aromatic hydrocarbon solvents suggests a different structure. The solid-state structure of 8 determined by X-ray diffraction consists of polymeric zigzag chains made up of cube-type [K{(μ_3 -N)(μ_3 -NH)(μ_3 -NSiMe₃)Ti₃(η^5 -C₅Me₅)₃(μ_3 -N)}] units with the potassium atom linked to the contiguous molecule through a η^5 -C₅Me₅ ligand (Figure 5). Compound 8 crystallized in the centrosymmetric P2₁/c space group, and the crystal lattice shows a regular alternation of the two enantiomers in neighboring chains.

A portion of the polymeric chain of compound 8 is shown in Figure 6, while selected distances and angles are given in Table 6. The crystal structure contains distorted [KTi₃N₄] cube-type cores with the metalloligands [(μ_3 -N)(μ_3 -NH)(μ_3 -NSiMe₃)-Ti₃(η^5 -C₅Me₅)₃(μ_3 -N)] coordinating in a tripodal fashion to the potassium centers. In addition, the alkali metal atoms are bonded to the five carbon atoms of the η^5 -C₅Me₅ ligand linked to the titanium(1) atoms of a contiguous molecule. If the centroid Cm(1) of these pentamethylcyclopentadienyl ligands is considered, the coordination sphere about the potassium atoms may be described as distorted tetrahedral with N–K(1)–N and N–K(1)–Cm(1) ranging from 61.9(1)–65.2(1)° and 128.9–152.9°, respectively. While the potassium–nitrogen bond lengths for the nitrido and NH imido groups are almost identical (K(1)–N(13) 2.772(4) and K(1)–N(23) 2.776(4) Å, respectively), those associated with the bulkier trimethylsilylimido ligands are significantly longer (K(1)–N(12) 3.108(4) Å). The steric repulsion of the SiMe₃ group might also be responsible of the differences in the potassium–carbon(C₅Me₅) bond lengths (range 3.000(4)–3.258(4) Å), the shortest distances being those with the C(13) and C(14) carbon atoms located on the opposite side of the trimethylsilylimido ligand.

Addition of 1 equiv of 18-crown-6 to a toluene solution of 8 led to the immediate precipitation of the well-separated ion pair [K(18-crown-6)][Ti₃(η^5 -C₅Me₅)₃(μ_3 -N)(μ -N)(μ -NSiMe₃)] (9). This compound was isolated in 80% yield as a yellow powder, which is only soluble in pyridine-*d*₅, while it reacts immediately with chloroform-*d*₁ to give complex 7 and free 18-crown-6 according to NMR spectroscopy. ¹H and ¹³C{¹H} NMR spectra of 9 in pyridine-*d*₅ show resonances for

three different η^5 -C₅Me₅ ligands in accord with a C₁ symmetry for the trinuclear titanium anion. In addition, the ¹H NMR spectrum of **9** shows a sharp singlet for the methylene groups of one crown ether ligand per anionic fragment. The three resonances for the *ipso* carbon of the η^5 -C₅Me₅ groups (δ = 114.2, 113.3, and 113.0) in the ¹³C{¹H} NMR spectrum of **9** are slightly shifted upfield with respect to those found in the spectra of **8** in pyridine-*d*₅ (δ = 115.4, 114.4, and 113.3) or benzene-*d*₆ (δ = 116.6, 115.9, and 113.9). Most likely, the structure of compound **8** in those solutions contains a cube-type [KTi₃N₄] core with solvent molecules coordinated to the potassium atom as a result of the rupture of the polymeric association determined in the solid state.

CONCLUSION

The imido-nitrido complex $[\{\text{Ti}(\eta^5\text{-C}_5\text{Me}_5)(\mu\text{-NH})\}_3(\mu_3\text{-N})]$ (**1**) reacts with 1 equiv of electrophilic reagents ROTf (R = H, Me) through selective protonation or methylation at one imido nitrogen to give $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})_2(\mu\text{-NHR})\text{-(OTf)}]$, with coordination of the triflate ligand to the opposite titanium atom of the resultant amido NHR. In those complexes, the triflate is hydrogen-bonded to the NHR group and assists the proton exchange between amido NHR and imido NH ligands. The larger trimethylsilyl fragment of Me₃SiOTf attacks the same nitrogen of **1** but produces a complex $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})_2(\mu\text{-NHSiMe}_3)]\text{[OTf]}$ with the triflate anion not coordinated to the metals. Deprotonation of this ionic compound with $[\text{K}\{\text{N}(\text{SiMe}_3)_2\}]$ occurs at the imido NH ligands rather than involving the NHSiMe₃ group, and the resultant molecular complex $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-N})(\mu\text{-NH})(\mu\text{-NHSiMe}_3)]$ readily rearranges to the trimethylsilylimido derivative $[\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-NH})_2(\mu\text{-NSiMe}_3)]$. The latter can be further deprotonated with $[\text{K}\{\text{N}(\text{SiMe}_3)_2\}]$ to give a potassium complex $[\text{K}\{(\mu_3\text{-N})(\mu_3\text{-NH})(\mu_3\text{-NSiMe}_3)\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})\}]$ and upon addition of 18-crown-6 leads to the well-separated ion-pair $[\text{K}(18\text{-crown-6})][\text{Ti}_3(\eta^5\text{-C}_5\text{Me}_5)_3(\mu_3\text{-N})(\mu\text{-N})(\mu\text{-NH})(\mu\text{-NSiMe}_3)]$. The treatment of **1** with 1 equiv of electrophiles and subsequent reactions on the resultant products described here have shown the possibility of preparing a series of cationic, neutral, or anionic trinuclear titanium complexes bearing bridging amido, imido, and nitrido functionalities. Several of these species undergo interesting rearrangement reactions involving proton exchange between the nitrogen ligands. Current work explores the reactivity of **1** with a variety of electrophiles in higher molar ratios.

ASSOCIATED CONTENT

Supporting Information

Kinetic data and analyses for the dynamic NMR spectroscopy study for compounds **2** and **3**, and the crystal structure of complex **5** showing the hydrogen bonds between ions. X-ray crystallographic files in CIF format for complexes **2**, **5**, **6**, and **8**. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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Notes

The authors declare no competing financial interest.

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REFERENCES

- (1) (a) Nugent, W. A.; Mayer, J. M. *Metal-Ligand Multiple Bonds*; John Wiley & Sons: New York, 1988. (b) Eikey, R. A.; Abu-Omar, M. M. *Coord. Chem. Rev.* **2003**, *243*, 83–124. (c) Berry, J. F. *Comments Inorg. Chem.* **2009**, *30*, 28–66.
- (2) (a) Nugent, W. A.; McKinney, R. J.; Kasowski, R. V.; Van-Catledge, F. A. *Inorg. Chim. Acta* **1982**, *65*, L91–L93. (b) Romo, S.; Antonova, N. S.; Carbó, J. J.; Poblet, J.-M. *Dalton Trans.* **2008**, 5166–5172.
- (3) Hedegård, E. D.; Bendix, J.; Sauer, S. P. A. *J. Mol. Struct. (THEOCHEM)* **2009**, *913*, 1–7.
- (4) For selected examples of the reaction of nitrido complexes with nucleophiles, see: (a) Crevier, T. J.; Bennett, B. K.; Soper, J. D.; Bowman, J. A.; Dehestani, A.; Hrovat, D. A.; Lovell, S.; Kaminsky, W.; Mayer, J. M. *J. Am. Chem. Soc.* **2001**, *123*, 1059–1071. (b) Seymore, S. B.; Brown, S. N. *Inorg. Chem.* **2002**, *41*, 462–469. (c) Bennet, B. K.; Saganic, E.; Lovell, S.; Kaminsky, W.; Samuel, A.; Mayer, J. M. *Inorg. Chem.* **2003**, *42*, 4127–4134. (d) Yi, X.-Y.; Lam, T. C. H.; Sau, Y.-K.; Zhang, Q.-F.; Williams, I. D.; Leung, W.-H. *Inorg. Chem.* **2007**, *46*, 7193–7198. (e) Yi, X.-Y.; Ng, H.-Y.; Williams, I. D.; Leung, W.-H. *Inorg. Chem.* **2011**, *50*, 1161–1163.
- (5) For selected examples of the reaction of imido complexes with nucleophiles, see: (a) Shapley, P. A.; Shusta, J. M.; Hunt, J. L. *Organometallics* **1996**, *15*, 1622–1629. (b) Kogut, E.; Wiencko, H. L.; Zhang, L.; Cordeau, D. E.; Warren, T. H. *J. Am. Chem. Soc.* **2005**, *127*, 11248–11249.
- (6) For selected examples of the reaction of nitrido complexes with electrophiles, see: (a) Shapley, P. A.; Own, Z.-Y.; Huffman, J. C. *Organometallics* **1986**, *5*, 1269–1271. (b) Marshman, R. W.; Shapley, P. A. *J. Am. Chem. Soc.* **1990**, *112*, 8369–8378. (c) Marshman, R. W.; Shusta, J. M.; Wilson, S. R.; Shapley, P. A. *Organometallics* **1991**, *10*, 1671–1676. (d) Sellmann, D.; Wemple, M. W.; Donaubauer, W.; Heinemann, F. W. *Inorg. Chem.* **1997**, *36*, 1397–1402. (e) Leung, W.-H.; Chan, E. Y. Y.; Lai, T. C. Y.; Wong, W.-T. *J. Chem. Soc., Dalton Trans.* **2000**, 51–56. (f) Abram, U.; Voigt, A.; Kirmse, R. *Polyhedron* **2000**, *19*, 1741–1748. (g) Sceats, E. L.; Figueroa, J. S.; Cummins, C. C.; Loening, N. M.; Van der Wel, P.; Griffin, R. G. *Polyhedron* **2004**, *23*, 2751–2768. (h) Lutz, C. M.; Wilson, S. R.; Shapley, P. A. *Organometallics* **2005**, *24*, 3350–3353. (i) Sarkar, S.; Abboud, K. A.; Veige, A. S. *J. Am. Chem. Soc.* **2008**, *130*, 16128–16129. (j) Walstrom, A.; Fan, H.; Pink, M.; Caulton, K. G. *Inorg. Chim. Acta* **2010**, *363*, 633–636. (k) Curley, J. J.; Cozzolino, A. F.; Cummins, C. C. *Dalton Trans.* **2011**, *40*, 2429–2432.
- (7) For selected examples of the reaction of imido complexes with electrophiles, see: (a) Chiu, K. W.; Wong, W.-K.; Wilkinson, G.; Galas, A. M. R.; Hursthouse, M. B. *Polyhedron* **1982**, *1*, 31–36. (b) Powell, K. R.; Pérez, P. J.; Luan, L.; Feng, S. G.; White, P. S.; Brookhart, M.; Templeton, J. L. *Organometallics* **1994**, *13*, 1851–1864. (c) Gountchev, T. I.; Tilley, T. D. *J. Am. Chem. Soc.* **1997**, *119*, 12831–12841. (d) Tonzetich, Z. J.; Schrock, R. R.; Müller, P. *Organometallics* **2006**, *25*, 4301–4306.
- (8) For accounts on the types of imido complexes, see: (a) Nugent, W. A.; Haymore, B. L. *Coord. Chem. Rev.* **1980**, *31*, 123–175. (b) Wigley, D. E. *Prog. Inorg. Chem.* **1994**, *42*, 239–482.
- (9) For reviews on polynuclear nitrido complexes, see: (a) Dehnicke, K.; Strähle, J. *Angew. Chem., Int. Ed. Engl.* **1981**, *20*, 413–426. (b) Dehnicke, K.; Strähle, J. *Angew. Chem., Int. Ed. Engl.* **1992**, *31*, 955–978. (c) Dehnicke, K.; Weller, F.; Strähle, J. *Chem. Soc. Rev.* **2001**, *30*, 125–135.

- (10) Morello, L.; Yu, P.; Carmichael, C. D.; Patrick, B. O.; Fryzuk, M. D. *J. Am. Chem. Soc.* **2005**, *127*, 12796–12797.
- (11) (a) MacKay, B. A.; Munha, R. F.; Fryzuk, M. D. *J. Am. Chem. Soc.* **2006**, *128*, 9472–9483. (b) Hirotsu, M.; Fontaine, P. P.; Epshteyn, A.; Zavalij, P. Y.; Sita, L. R. *J. Am. Chem. Soc.* **2007**, *129*, 9284–9285. (c) Akagi, F.; Matsuo, T.; Kawaguchi, H. *Angew. Chem., Int. Ed.* **2007**, *46*, 8778–8781. (d) Nikiforov, G. B.; Vidyaratne, I.; Gambarotta, S.; Korobkov, I. *Angew. Chem., Int. Ed.* **2009**, *48*, 7415–7419.
- (12) (a) Roesky, H. W.; Bai, Y.; Noltemeyer, M. *Angew. Chem., Int. Ed. Engl.* **1989**, *28*, 754–755. (b) Abarca, A.; Gómez-Sal, P.; Martín, A.; Mena, M.; Poblet, J.-M.; Yélamos, C. *Inorg. Chem.* **2000**, *39*, 642–651.
- (13) (a) Abarca, A.; Martín, A.; Mena, M.; Yélamos, C. *Angew. Chem., Int. Ed.* **2000**, *39*, 3460–3463. (b) Freitag, K.; Gracia, J.; Martín, A.; Mena, M.; Poblet, J.-M.; Sarasa, J. P.; Yélamos, C. *Chem.-Eur. J.* **2001**, *7*, 3644–3651. (c) García-Castro, M.; Gracia, J.; Martín, A.; Mena, M.; Poblet, J.-M.; Sarasa, J. P.; Yélamos, C. *Chem.-Eur. J.* **2005**, *11*, 1030–1041. (d) García-Castro, M.; Martín, A.; Mena, M.; Yélamos, C. *Organometallics* **2007**, *26*, 408–416. (e) Martínez-Espada, N.; Mena, M.; Mosquera, M. E. G.; Pérez-Redondo, A.; Yélamos, C. *Organometallics* **2010**, *29*, 6732–6738. (f) Caballo, J.; García-Castro, M.; Martín, A.; Mena, M.; Pérez-Redondo, A.; Yélamos, C. *Inorg. Chem.* **2011**, *50*, 6798–6808.
- (14) Carbó, J. J.; Martínez-Espada, N.; Mena, M.; Mosquera, M. E. G.; Poblet, J.-M.; Yélamos, C. *Chem.-Eur. J.* **2009**, *15*, 11619–11631.
- (15) Carbó, J. J.; Martín, A.; Mena, M.; Pérez-Redondo, A.; Poblet, J.-M.; Yélamos, C. *Angew. Chem., Int. Ed.* **2007**, *46*, 3095–3098.
- (16) Caballo, J.; González-Moreiras, M.; Mena, M.; Pérez-Redondo, A.; Yélamos, C. *Dalton Trans.* **2012**, *41*, 6069–6071.
- (17) García-Castro, M.; Martín, A.; Mena, M.; Pérez-Redondo, A.; Yélamos, C. *Chem.-Eur. J.* **2001**, *7*, 647–651.
- (18) Farrugia, L. J. *J. Appl. Crystallogr.* **1999**, *32*, 837–838.
- (19) Sheldrick, G. M. *Acta Crystallogr., Sect. A* **2008**, *64*, 112–122.
- (20) (a) Duan, Z.; Verkade, J. G. *Inorg. Chem.* **1996**, *35*, 5325–5327. (b) Carmalt, C. J.; Mileham, J. D.; White, A. J. P.; Williams, D. J. *New J. Chem.* **2000**, *24*, 929–930.
- (21) (a) Schubart, M.; O'Dwyer, L.; Gade, L. H.; Li, W.-S.; McPartlin, M. *Inorg. Chem.* **1994**, *33*, 3893–3898. (b) Ngo, S. C.; Toscano, P. J.; Welch, J. T. *Helv. Chim. Acta* **2002**, *85*, 3366–3382. (c) Kessler, M.; Hansen, S.; Hollmann, D.; Klahn, M.; Beweries, T.; Spannenberg, A.; Brückner, A.; Rosenthal, U. *Eur. J. Inorg. Chem.* **2011**, 627–631. (d) Davidson, M. G.; Johnson, A. L. *Eur. J. Inorg. Chem.* **2011**, 5151–5159.
- (22) (a) Jeffrey, G. A. *An Introduction to Hydrogen Bonding*; Oxford University Press: New York, 1997. (b) Steiner, T. *Angew. Chem., Int. Ed.* **2002**, *41*, 48–76.
- (23) Nakamoto, K. *Infrared and Raman Spectra of Inorganic and Coordination Compounds*, 5th ed.; Wiley: New York, 1997.
- (24) (a) Lawrance, G. A. *Chem. Rev.* **1986**, *86*, 17–33. (b) Johnston, D. H.; Shriver, D. F. *Inorg. Chem.* **1993**, *32*, 1045–1047. (c) Huang, W.; Frech, R.; Wheeler, R. A. *J. Phys. Chem.* **1994**, *98*, 100–110.
- (25) (a) Männle, F.; Limbach, H.-H. *Angew. Chem., Int. Ed. Engl.* **1996**, *35*, 441–442. (b) Limbach, H.-H.; Männle, F.; Detering, C.; Denisov, G. S. *Chem. Phys.* **2005**, *319*, 69–92.
- (26) (a) Lee, K. E.; Gladysz, J. A. *Polyhedron* **1988**, *7*, 2209–2211. (b) Straus, D. A.; Zhang, C.; Quimbata, G. E.; Grumbine, S. D.; Heyn, R. H.; Tilley, T. D.; Rheingold, A. L.; Geib, S. J. *J. Am. Chem. Soc.* **1990**, *112*, 2673–2681. (c) Straus, D. A.; Grumbine, S. D.; Tilley, T. D. *J. Am. Chem. Soc.* **1990**, *112*, 7801–7802. (d) Burns, R. M.; Hubbard, J. L. *J. Am. Chem. Soc.* **1994**, *116*, 9514–9520. (e) Hayashida, T.; Kondo, H.; Terasawa, J.; Kirchner, K.; Sunada, Y.; Nagashima, H. *J. Organomet. Chem.* **2007**, *692*, 382–394.
- (27) (a) Sandström, J. *Dynamic NMR Spectroscopy*; Academic Press: London, 1992. (b) *gNMR, version 4*; Cherwell Scientific Publishing: Oxford, UK, 1997.
- (28) (a) Mann, B. E. Non-Rigidity in Organometallic Compounds. In *Comprehensive Organometallic Chemistry*; Geoffrey, W., Gordon, F., Stone, A., Abel, E. W., Eds.; Pergamon: Oxford, UK, 1982; Vol. 20, pp 89–171. (b) Faller, J. W. Stereochemical Nonrigidity of Organometallic Complexes. In *Encyclopedia of Inorganic Chemistry*, 2nd ed.; King, R. B., Ed.; John Wiley & Sons: Chichester, UK, 2005; Vol. 8, pp 5270–5303.
- (29) (a) Caldin, E. F.; Mateo, S. J. *Chem. Soc., Faraday Trans. 1* **1975**, *71*, 1876–1904. (b) Scherer, G.; Limbach, H.-H. *J. Am. Chem. Soc.* **1994**, *116*, 1230–1239.
- (30) Martín, A.; Mena, M.; Pérez-Redondo, A.; Yélamos, C. *Inorg. Chem.* **2004**, *43*, 2491–2498.