

Lead–Chromium Carbonyl Complexes Incorporated with Group 8 Metals: Synthesis, Reactivity, and Theoretical Calculations

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The trichromium–lead complex $[\text{Pb}\{\text{Cr}(\text{CO})_5\}_3]^{2-}$ (**1**) was isolated from the reaction of PbCl_2 and $\text{Cr}(\text{CO})_6$ in a KOH/MeOH solution, and the new mixed chromium–iron–lead complex $[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Fe}(\text{CO})_4\}_2]^{2-}$ (**3**) was synthesized from the reaction of PbCl_2 and $\text{Cr}(\text{CO})_6$ in a KOH/MeOH solution followed by the addition of $\text{Fe}(\text{CO})_5$. X-ray crystallography showed that **3** consisted of a central Pb atom bound in a trigonal-planar environment to two $\text{Fe}(\text{CO})_4$ and one $\text{Cr}(\text{CO})_5$ fragments. When complex **1** reacted with 1.5 equiv of $\text{Mn}(\text{CO})_5\text{Br}$, the $\text{Cr}(\text{CO})_4$ -bridged dimeric lead–chromium carbonyl complex $[\text{Pb}_2\text{Br}_2\text{Cr}_4(\text{CO})_{18}]^{2-}$ (**4**) was produced. However, a similar reaction of **3** or the isostructural triiron–lead complex $[\text{Pb}\{\text{Fe}(\text{CO})_4\}_3]^{2-}$ (**2**) with $\text{Mn}(\text{CO})_5\text{Br}$ in MeCN led to the formation of the Fe_3Pb_2 -based trigonal-bipyramidal complexes $[\text{Fe}_3(\text{CO})_9\{\text{PbCr}(\text{CO})_5\}_2]^{2-}$ (**6**) and $[\text{Fe}_3(\text{CO})_9\{\text{PbFe}(\text{CO})_4\}_2]^{2-}$ (**5**), respectively. On the other hand, the Ru_3Pb_2 -based trigonal-bipyramidal complex $[\text{Ru}_3(\text{CO})_9\{\text{PbCr}(\text{CO})_5\}_2]^{2-}$ (**7**) was obtained directly from the reaction of PbCl_2 , $\text{Cr}(\text{CO})_6$, and $\text{Ru}_3(\text{CO})_{12}$ in a KOH/MeOH solution. X-ray crystallography showed that **5** and **6** each had an Fe_3Pb_2 trigonal-bipyramidal core geometry, with three $\text{Fe}(\text{CO})_3$ groups occupying the equatorial positions and two $\text{PbFe}(\text{CO})_4$ or $\text{PbCr}(\text{CO})_5$ units in the axial positions, while **7** displayed a Ru_3Pb_2 trigonal-bipyramidal geometry with three equatorial $\text{Ru}(\text{CO})_3$ groups and two axial $\text{PbCr}(\text{CO})_5$ units. The complexes **3**–**7** were characterized spectroscopically, and their nature, formation, and electrochemistry were further examined by molecular orbital calculations at the B3LYP level of density functional theory.

Introduction

Cooperative bimetallic synergism is central to some of today's most important applications in catalysts, nanoscience, material science, and magnetism.^{1–5} The cooperative properties of mixed-metal clusters are of great interest. Very few, however, are understood in detail. One important direction in this area is to rationalize the synthesis of symmetric metal or mixed-metal carbonyl clusters bearing a protective shell of CO

or main-group ligands, which can be expected to exhibit reversible or quasi-reversible redox reactions without structural rearrangement of their metal skeletons.^{1,3a,6}

Our research group has an ongoing interest in chromium,⁷ iron,⁸ and ruthenium⁹ carbonyl clusters and has recently explored the lead-containing chromium system.¹⁰ Although group 14-incorporated, group 6 and/or group 8 metal carbonyl complexes have been reported, their lead-incorporated complexes remain rare, mainly because of a lack of applicable synthetic routes.^{2d,f,10–14} Furthermore, the effect and nature of homo- or heteronuclear metal–metal interactions on the formation of such carbonyl complexes remain little explored.^{15,16}

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The study and comparison of lead-containing homo- or heterometallic complexes are, therefore, of great interest and are challenging in view of their cooperative chemical properties and potential applications.

In the present study, facile routes were discovered to a series of lead-containing homo- or heteronuclear chromium and group 8 trigonal-planar complexes, $[\text{Et}_4\text{N}]_2[\text{Pb}\{\text{Cr}(\text{CO})_5\}_3]$ ($[\text{Et}_4\text{N}]_2[\mathbf{1}]$)^{11a} and $[\text{Bu}_4\text{N}]_2[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Fe}(\text{CO})_4\}_2]$

($[\text{Bu}_4\text{N}]_2[\mathbf{3}]$), and trigonal-bipyramidal clusters, $[\text{Et}_4\text{N}]_2[\text{Fe}_3(\text{CO})_9\{\text{PbFe}(\text{CO})_4\}_2]$ ($[\text{Et}_4\text{N}]_2[\mathbf{5}]$), $[\text{Et}_4\text{N}]_2[\text{Fe}_3(\text{CO})_9\{\text{PbCr}(\text{CO})_5\}_2]$ ($[\text{Et}_4\text{N}]_2[\mathbf{6}]$), and $[\text{Et}_4\text{N}]_2[\text{Ru}_3(\text{CO})_9\{\text{PbCr}(\text{CO})_5\}_2]$ ($[\text{Et}_4\text{N}]_2[\mathbf{7}]$), in which the mixed chromium–group 8 trigonal-planar complexes could readily undergo coupling reactions to give rise to their corresponding trigonal-bipyramidal complexes. Clusters **3**, **6**, and **7** represent the first examples of lead-containing mixed chromium–group 8 metal clusters. In addition, the formation, structural transformation, and electrochemistry of the resultant clusters were investigated and discussed on the basis of density functional theory (DFT) calculations.

Results and Discussion

The class of Pb–Cr–CO complexes has previously been rare because of the lack of a feasible methodology and their reactive nature.^{10,11a–g} By varying of the sources of lead(II)

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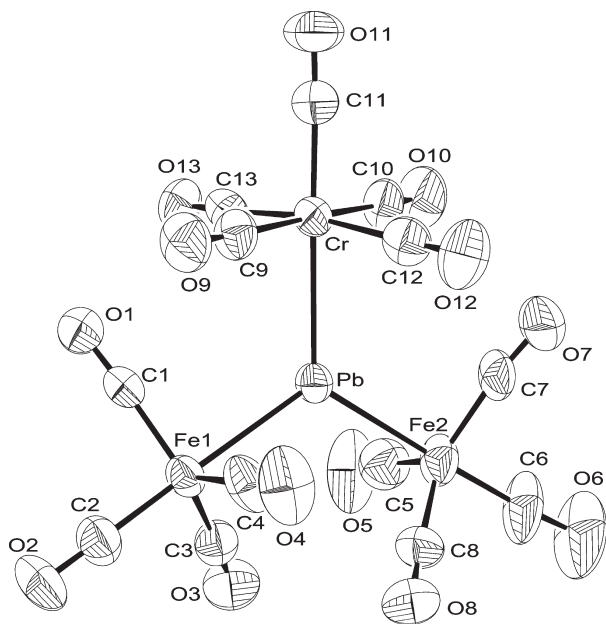
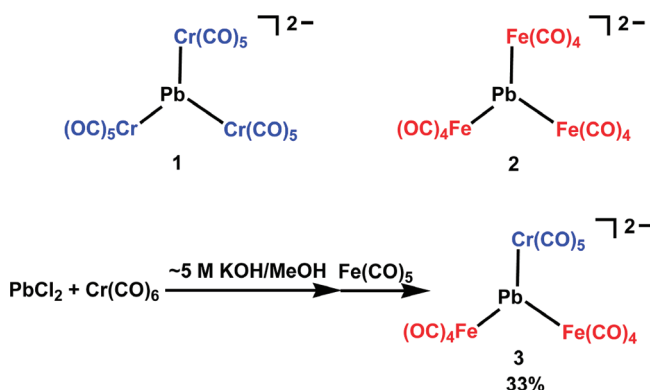


Figure 1. ORTEP diagram of anion **3**, showing 30% probability thermal ellipsoids. Selected bond lengths (Å) and angles (deg): Pb1–Cr1 2.714(2), Pb1–Fe1 2.656(2), Pb1–Fe2 2.636(2); Cr1–Pb1–Fe1 118.67(6), Cr1–Pb1–Fe2 123.67(7), Fe1–Pb1–Fe2 117.03(7).

Scheme 1



and the bases, as well as the concentration of the bases applied, we have discovered a facile route to the known lead–chromium carbonyl complex $[\text{Pb}\{\text{Cr}(\text{CO})_5\}_3]^{2-}$ (**1**)^{11a} from the reaction of PbCl_2 with $\text{Cr}(\text{CO})_6$ in a KOH/MeOH solution. In pursuit of novel heteronuclear clusters, group 8 metals were introduced into the lead–chromium carbonyl system. It was found that when PbCl_2 was treated with $\text{Cr}(\text{CO})_6$ in a KOH/MeOH solution, followed by the addition of $\text{Fe}(\text{CO})_5$, the mixed chromium–iron–lead complex $[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Fe}(\text{CO})_4\}_2]^{2-}$ (**3**) could be obtained in moderate yields (Scheme 1). Complex **3** was quite air-sensitive and was isolated as the salt $[\text{Bu}_4\text{N}]^+$. Complex **3** was characterized by spectroscopic methods, elemental analysis, and single-crystal X-ray diffraction analysis. An ORTEP diagram of the molecular structure of the anion **3** is shown in Figure 1.

The anion **3** is structurally similar to complex **1**^{11a} and the previously known triiron–lead complex **2**.^{12a} The average Pb–Fe bond length in **3** [2.65(1) Å] is comparable to those in **2** [2.624(4) Å] and $[\text{Et}_4\text{N}]_2[\text{Pb}\{\text{Fe}_2(\text{CO})_8\}\{\text{Fe}(\text{CO})_4\}_2]$ [2.7(1) Å]^{12c} (Table 1). The Pb–Cr bond length of **3**, 2.714(2) Å, is close to those in **1** [average 2.729(4) Å] and

the metallocplumbylene complex $[(\text{FcN})_2\text{PbCr}(\text{CO})_5]$ [2.7157(4) Å; $\text{FcN} = [\text{Fe}(\eta^5\text{-C}_5\text{H}_5)(\eta^5\text{-C}_5\text{H}_3(\text{CH}_2\text{NMe}_2)_2\text{-C}, \text{N})]$.^{11e} Complex **3** is an isoelectronic analogue of **1** and **2**, which contain three unsaturated Pb–M (M = Cr, Fe) bonds in terms of the electron-counting rule.^{11a,12a} These three isostructural complexes, **1–3**, thus provided good precursors for the preparation of polynuclear clusters.

Previous studies showed that the trigonal-planar $\text{PbCr}_3\text{--}1$ complex could react with the nucleophilic reagent PMe_3 to form its base adduct $[\text{Pb}\{\text{Cr}(\text{CO})_5\}_3(\text{PMe}_3)]^{2-}$.^{11a} For comparison, and to explore possible metal expansion reactions, the reactions of the trigonal-planar compounds **1–3** with $\text{Mn}(\text{CO})_5\text{Br}$ were studied. When complex **1** was treated with $\text{Mn}(\text{CO})_5\text{Br}$ at ambient temperature, a new ring complex, $[\text{Pb}_2\text{Br}_2\text{Cr}_4(\text{CO})_{18}]^{2-}$ (**4**), was produced with the elimination of $\text{Mn}_2(\text{CO})_{10}$ and $[\text{HCr}_2(\text{CO})_{10}]^-$ (identified by IR spectroscopy). The results suggested that the central Pb atom was nucleophilically attacked by Br^- accompanied with the loss of $[\text{Cr}(\text{CO})_5]^{2-}$, followed by a coupling reaction to give complex **4**. Complex **4** could be further transformed to the known tetrahedral compound $[\text{PbBr}_2\text{Cr}_2(\text{CO})_{10}]^{2-}$ ^{11b} via Pb–Cr bond breakage and Pb–Br bond formation, upon the addition of $\text{Mn}(\text{CO})_5\text{Br}$ in a MeCN solution (Scheme 2).

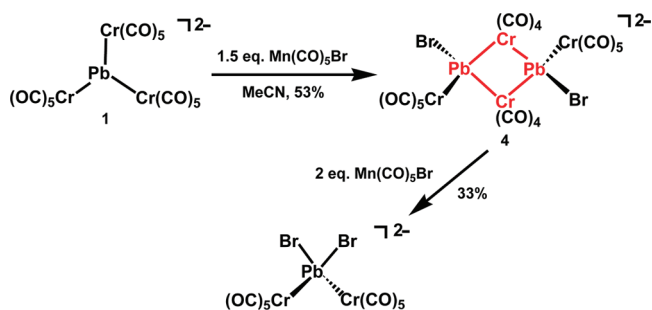
Compound **4** was characterized by IR spectroscopy, elemental analysis, and single-crystal X-ray diffraction. As shown in Figure 2, the anion **4** contains an inversion center at the center of the Pb_2Cr_2 plane, in which the Pb1–Cr2–Pb1a and Cr2–Pb1–Cr2a angles are 109.79(5) and 70.21(5)°, respectively. The Pb atom is four-coordinated but severely distorted from the ideal tetrahedral geometry [70.21(5)–135.81(6)°]. With respect to the bonding, complex **4** may be considered to be composed of two $[(\text{CO})_5\text{Cr}]\text{PbBr}\{\text{Cr}(\text{CO})_4\}^-$ species, which are dimerized by the Pb–Cr bonding to form a four-membered Pb_2Cr_2 ring due to the electron deficiency of the Pb atoms. According to a search of the Cambridge Crystallographic Data Centre, complex **4** is the first example of a four-membered Pb_2M_2 ring in the lead–group 6 metal system. The Pb–Cr(CO)₄ distances in **4** [average 2.72(1) Å] are slightly longer than the Pb–Cr(CO)₅ distances [2.679(2) Å], indicating that the Pb–Cr(CO)₄ bonds in the Pb_2Cr_2 ring are weaker than the pendent Pb–Cr(CO)₅ bonds. As a result, complex **4** was susceptible to the attack of Br^- onto the central Pb_2Cr_2 ring, causing Pb–Cr bond cleavage to give $[\text{PbBr}_2\text{Cr}_2(\text{CO})_{10}]^{2-}$.

For comparison, an isostructural trigonal-planar complex $[\text{Pb}\{\text{Fe}(\text{CO})_4\}_3]^{2-}$ (**2**)^{12a} was synthesized. The treatment of complex **2** with $\text{Mn}(\text{CO})_5\text{Br}$ in MeCN at 80 °C led to the formation of the trigonal-bipyramidal cluster $[\text{Fe}_3(\text{CO})_9\{\text{PbFe}(\text{CO})_4\}_2]^{2-}$ (**5**; Scheme 3). Complex **5** was considered as a result of the coupling reaction of **2** via Pb–Fe and Fe–Fe bond formation and the loss of one $[\text{Fe}(\text{CO})_4]^{2-}$ and three CO units. In a similar fashion, reaction of the chromium–iron planar complex **3** with $\text{Mn}(\text{CO})_5\text{Br}$ in MeCN at 80 °C yielded the mixed chromium–iron trigonal-bipyramidal cluster $[\text{Fe}_3(\text{CO})_9\{\text{PbCr}(\text{CO})_5\}_2]^{2-}$ (**6**; Scheme 3). Likewise, cluster **6** could be regarded as a result of the dimerization of **3** via Pb–Fe and Fe–Fe bond formation and the loss of one $[\text{Fe}(\text{CO})_4]^{2-}$ and three CO units. Both clusters **5** and **6** were isolated as the salt $[\text{Et}_4\text{N}]^+$, and they were fully characterized by IR spectroscopy, elemental analyses, and single-crystal X-ray diffraction. It is of great interest that clusters **5** and **6** were interconvertible under suitable conditions. Cluster **5** readily transformed to cluster **6** in good yields upon treatment with $\text{Cr}(\text{CO})_6$ in a

Table 1. Average Bond Distances (Å) of [PPh₄]₂[1], [Et₄N]₂[2], [Bu₄N]₂[3], [Et₄N]₂[4], [Et₄N]₂[5], [Et₄N]₂[6], [Et₄N]₂[7], and Related Complexes

| complex | Fe—Fe | Ru—Ru | Pb—Cr | Pb—Fe | Pb—Ru | ref |
|---|----------|----------|---|---|-----------|-----|
| [PPh ₄] ₂ [1] | | | 2.729(4) | | | 11a |
| [Et ₄ N] ₂ [2] | | | | 2.624(4) | | 12a |
| [Bu ₄ N] ₂ [3] | | | 2.714(2) | 2.65(1) | | a |
| [Et ₄ N] ₂ [4] | | | 2.72(1) ^b 2.679(2) ^c | | | a |
| [Et ₄ N] ₂ [5] | 2.83(4) | | | 2.60(1) ^d 2.546(2) ^e | | a |
| [Et ₄ N] ₂ [6] | 2.79(6) | | 2.67(1) | 2.65(3) | | a |
| [Et ₄ N] ₂ [7] | | 2.97(1) | 2.680(4) | | 2.77(2) | a |
| [Et ₄ N] ₂ [Pb{Fe ₂ (CO) ₈ }{Fe(CO) ₄ } ₂] | 2.617(5) | | | 2.7(1) | | 12c |
| (FcN) ₂ PbCr(CO) ₅ ^f | | | 2.7157(4) | | | 11e |
| [PPh ₄] ₂ [PbBr ₂ Cr ₂ (CO) ₁₀] | | | 2.70(2) | | | 11b |
| Fe ₃ (CO) ₉ (μ ₃ -CF) ₂ | 2.540(2) | | | | | 12g |
| Fe ₃ (CO) ₉ [μ ₃ -SiFeCp(CO) ₂] ₂ | 2.67(2) | | | | | 12j |
| Fe ₃ (CO) ₉ (μ ₃ -GeEt) ₂ | 2.73(1) | | | | | 12j |
| Fe ₃ (CO) ₉ [μ ₃ -GeFeCp(CO) ₂] ₂ | 2.73(1) | | | | | 12j |
| Fe ₃ (CO) ₉ [μ ₃ -GeRe(CO) ₅] ₂ | 2.719(4) | | | | | 12i |
| Fe ₃ (CO) ₉ [μ ₃ -GeCo(CO) ₄] ₂ | 2.753(6) | | | | | 12h |
| Fe ₃ (CO) ₉ [μ ₃ -SnRe(CO) ₅] ₂ | 2.807(4) | | | | | 12l |
| Fe ₃ (CO) ₉ [μ ₃ -SnFeCp(CO) ₂] ₂ | 2.792(6) | | | | | 12k |
| Fe ₃ (CO) ₉ [μ ₃ -SnMn(CO) ₅] ₂ ^g | 2.81 | | | | | 12m |
| Fe ₃ (CO) ₉ [μ ₃ -PCr(CO) ₅] ₂ | 2.616(6) | | | | | 17c |
| Fe ₃ (CO) ₉ [μ ₃ -PMnCp(CO) ₂] ₂ | 2.636(6) | | | | | 17c |
| Fe ₃ (CO) ₉ (μ ₃ -As) ₂ | 2.623(4) | | | | | 17a |
| Fe ₃ (CO) ₉ [μ ₃ -AsCr(CO) ₅] ₂ | 2.688(2) | | | | | 17c |
| Fe ₃ (CO) ₉ [μ ₃ -AsMnCp(CO) ₂] ₂ | 2.693(2) | | | | | 17c |
| Fe ₃ (CO) ₉ [μ ₃ -SbCr(CO) ₅] ₂ | 2.782(2) | | | | | 17c |
| Fe ₃ (CO) ₉ [μ ₃ -SbMnCp(CO) ₂] ₂ | 2.768(2) | | | | | 17c |
| Fe ₃ (CO) ₉ (μ ₃ -Bi) ₂ | 2.75(1) | | | | | 17b |
| [PtRu ₅ (CO) ₁₅ (μ-PbPh ₂)(μ ₆ -C)] | | | | | 2.6668(5) | 13a |
| Ru(CO) ₂ (PbPh ₃)(ⁱ Pr-DAB)Cl | | | | | 2.7028(8) | 13b |
| [Ru ₃ {μ-PbR ₂ } ₂ (μ-CO)(CO) ₉] ^h | | 2.92(6) | | | 2.78(1) | 13c |
| Ru ₃ (CO) ₉ (μ ₃ -Bi) ₂ | | 2.940(7) | | | | 19a |
| Ru ₃ (CO) ₉ [μ ₃ -GeRu(CO) ₂ (η ⁵ -C ₅ Me ₄ H)] ₂ | | 2.90(3) | | | | 19b |
| Ru ₃ (CO) ₁₂ | | 2.854(5) | | | | 20 |

^a This work. ^b Pb—Cr(CO)₄. ^c Pb—Cr(CO)₅. ^d Equatorial. ^e Terminal. ^f FcN = [Fe(η⁵-C₅H₅)(η⁵-C₅H₃(CH₂NMe₂)-2)-C,N]. ^g Estimated standard deviations are not reported. ^h R = CH(SiMe₃)₂.

Scheme 2

KOH/MeOH solution. Conversely, cluster **6** could be reconverted to **5** upon its reaction with Fe(CO)₅/KOH/MeOH. The DFT calculations also supported the observation that these transformations were reversible (discussed later).

The ORTEP diagram of cluster **5** is illustrated in Figure 3. The anion lies on a site that includes two crystallographic mirror planes, one going through the equatorial Fe₃ plane

and the other including apical PbFe(CO) fragments and bisecting this plane. There is disorder across the latter plane, leading to 50% occupancy for each equatorial Fe atom that brings out two related orientation Fe₃ triangles by rotation of ca. 32° (see Figure S1 in the Supporting Information). Moreover, C8 and C8', which are coordinated to the Fe₄ atom, are disordered with 35% occupancy and 15% for C8a and C8a'. Further, the O8 and O8a atoms are also disordered, with 50% occupancy. The related trigonal-bipyramidal clusters [Fe₃(μ₃-CF)₂(CO)₉],^{12g} [Fe₃(μ₃-GeCo(CO)₄]₂(CO)₉],^{12h} [Fe₃(μ₃-As)₂(CO)₉],^{17a} and [Fe₃(μ₃-Bi)₂(CO)₉]^{17b} crystallized in the higher-symmetry space group and showed similar disorder. Thus, cluster **5** exhibits the expected near-equilateral triangle of Fe atoms with both sides capped by Pb atoms, in which each Pb is further bonded to an Fe(CO)₄ group in a distorted tetrahedral environment and each Fe atom in the trigonal plane is bonded to three CO ligands.

Cluster **6** is isoelectronic and structurally similar to **5**, which consists of an equatorial Fe₃(CO)₉ triangular cluster with both sides capped by the PbCr(CO)₅ fragments (Figure 4). In accordance with the electron-counting rule,¹⁸ three Fe—Fe bonds are observed for a 48-electron Fe₃-based cluster in both **5** and **6**. The average Pb—Fe(equatorial) bond of **5** [2.60(1) Å] is slightly shorter than that in **6** [2.65(3) Å], whereas the average Fe—Fe bond of **5** [2.83(4) Å] is comparable to that in **6** [2.79(6) Å],

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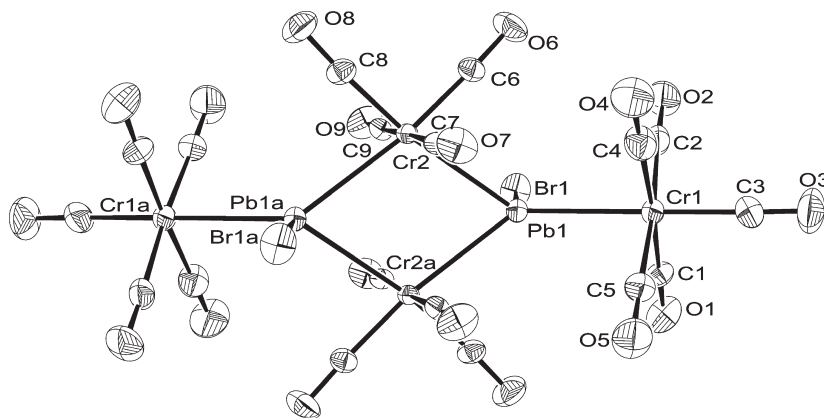
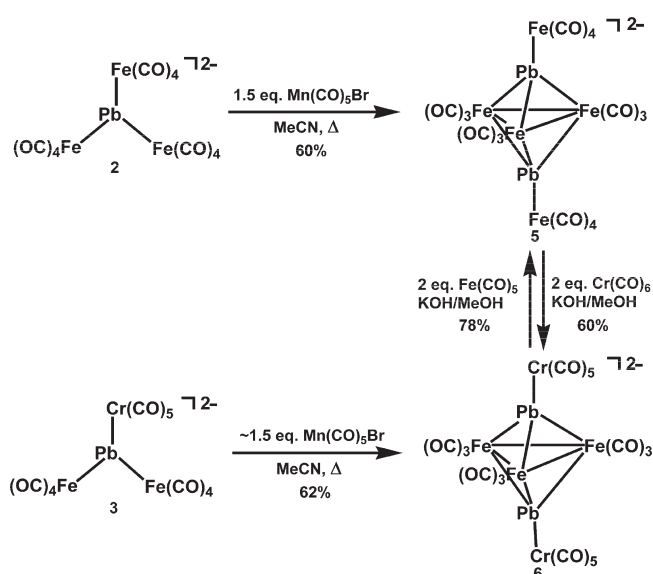


Figure 2. ORTEP diagram of anion **4**, showing 30% probability thermal ellipsoids. Selected bond lengths (Å) and angles (deg): Pb1–Cr1 2.679(2), Pb1–Cr2 2.732(2), Pb1–Br1 2.724(2), Pb1–Cr2a 2.716(2); Cr1–Pb1–Cr2 133.89(6), Cr1–Pb1–Cr2a 135.81(6), Cr2–Pb1–Cr2a 70.21(5), Cr1–Pb1–Br1 102.40(6), Cr2–Pb1–Br1 103.61(5), Cr2a–Pb1–Br1 105.66(5), Pb1–Cr2–Pb1a 109.79(5).

Scheme 3



probably because of the lower steric hindrance of $\text{Fe}(\text{CO})_4$ versus $\text{Cr}(\text{CO})_5$ groups bonded to the Pb atoms (Table 1). The average $\text{Fe}(\text{terminal})\text{--Pb--Fe}(\text{equatorial})$ and $\text{Fe}(\text{equatorial})\text{--Pb--Fe}(\text{equatorial})$ angles of **5** are 140.99 and 66.02°, respectively. In **6**, the average $\text{Cr}(\text{terminal})\text{--Pb--Fe}(\text{equatorial})$ and $\text{Fe}(\text{equatorial})\text{--Pb--Fe}(\text{equatorial})$ angles are 142.43 and 63.51°, respectively. Although analogous E_2Fe_3 trigonal-bipyramidal clusters with $\text{E} = \text{C}, \text{Si}, \text{Ge}, \text{Sn}, \text{P}, \text{As}, \text{Sb}, \text{and Bi}$ are known,^{12g–m,17} clusters **5** and **6** represent the first Pb-capped Fe_3 examples. As shown in Table 1, the average Fe--Fe bonds in **5** [2.83(4) Å] and **6** [2.79(6) Å] are longer than those in the related clusters because of the larger size of the main-group Pb atom.

A more interesting finding was that the reaction of PbCl_2 with $\text{Cr}(\text{CO})_6$ in a KOH/MeOH solution followed by the addition of $\text{Ru}_3(\text{CO})_{12}$ could directly yield the lead-containing mixed chromium–ruthenium trigonal-bipyramidal complex $[\text{Ru}_3(\text{CO})_9\{\text{PbCr}(\text{CO})_5\}_2]^{2-}$ (**7**), which was isolated as

the salt $[\text{Et}_4\text{N}]^+$ (Scheme 4). Unlike the mixed chromium–iron–lead cluster **3**, the analogous chromium–ruthenium–lead trigonal-planar complex $[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Ru}(\text{CO})_4\}_2]^{2-}$ was not observable under similar reaction conditions, probably because of the reactive nature of the Pb–Ru bond (discussed later).

Cluster **7** is fully structurally characterized by spectroscopic methods, elemental analysis, and single-crystal X-ray diffraction. A diagram of the dianion **7** is shown in Figure 5. Cluster **7** is an analogue of **5** and **6**, with the core geometry based on a trigonal-bipyramidal arrangement of three equatorial $\text{Ru}(\text{CO})_3$ groups and two axial $\text{PbCr}(\text{CO})_5$ units. The average $\text{Cr}(\text{terminal})\text{--Pb--Ru}(\text{equatorial})$ and $\text{Ru}(\text{equatorial})\text{--Pb--Ru}(\text{equatorial})$ angles are 141.54 and 64.88°, respectively. As shown in Table 1, the average Pb–Cr length of **7** [2.680(4) Å] is close to that in **6** [2.67(1) Å], whereas the average Pb–Ru length of **7** [2.77(2) Å] is within the range of those in the reported compounds $[\text{PtRu}_5(\text{CO})_{15}(\mu\text{-PbPh}_2)(\mu_6\text{-C})]$ [2.6668(5) Å],^{13a} $[\text{Ru}(\text{CO})_2(\text{PbPh}_3)(\text{Pr-DAB})\text{Cl}]$ [2.7028(8) Å],^{13b} and $[\text{Ru}_3\{\mu\text{-PbR}_2\}_2(\mu\text{-CO})(\text{CO})_9]$ [2.78(1) Å, $\text{R} = \text{CH}(\text{SiMe}_3)_2$].^{13c} Moreover, the average Ru–Ru distance of **7** is 2.97(1) Å, a bit longer than those in the related Ru_3 clusters $[\text{Ru}_3(\text{CO})_9(\mu_3\text{-Bi})_2]$ [2.940(7) Å],^{19a} $[\text{Ru}_3(\text{CO})_9[\mu_3\text{-GeRu}(\text{CO})_2(\eta^5\text{-C}_5\text{Me}_4\text{H})]_2]$ [2.90(3) Å],^{19b} and $\text{Ru}_3(\text{CO})_{12}$ [2.854(5) Å],²⁰ indicative of the small effect of the apical atoms. A search of the Cambridge Crystallographic Data Centre failed to yield any structurally characterized compounds with the Cr--Pb--Fe or Cr--Pb--Ru fragment. Clusters **6** and **7** represent the first examples of lead-containing mixed chromium and group 8 clusters.

DFT Calculations. The computational method of DFT was employed in an attempt to better understand the nature of complexes **1–7**. These calculations were carried out using the B3LYP DFT method^{21,22} with LanL2DZ/6-31G(d) basis sets. The geometries of the X-ray-determined and proposed intermediates in this study were fully optimized, and the natural bond order (NBO)²³ and natural population analyses (NPA)²⁴ for complexes **1–7** are displayed in Table 2.

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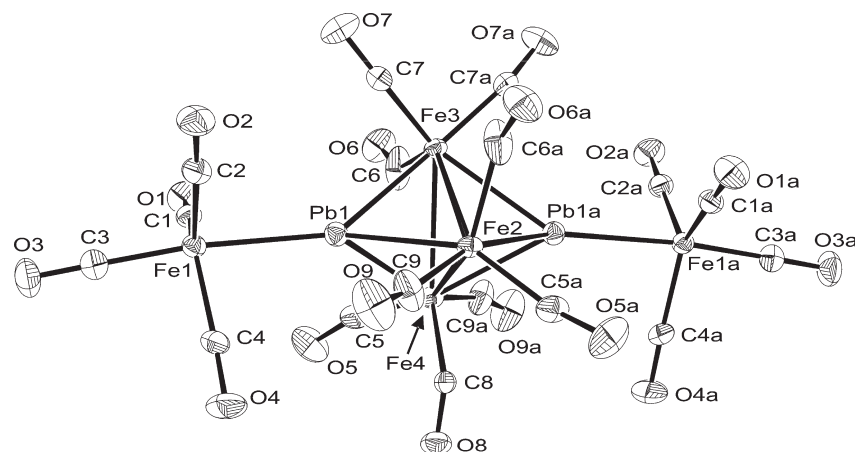


Figure 3. ORTEP diagram of anion **5**, showing 30% probability thermal ellipsoids. Selected bond lengths (Å) and angles (deg): Pb1–Fe1 2.546(1), Pb1–Fe2 2.609(2), Pb1–Fe3 2.588(2), Pb1–Fe4 2.598(2), Fe2–Fe3 2.812(3), Fe3–Fe4 2.805(4), Fe4–Fe2 2.875(4); Fe3–Fe2–Fe4 59.10(9), Fe2–Fe3–Fe4 61.6(1), Fe2–Fe4–Fe3 59.33(8).

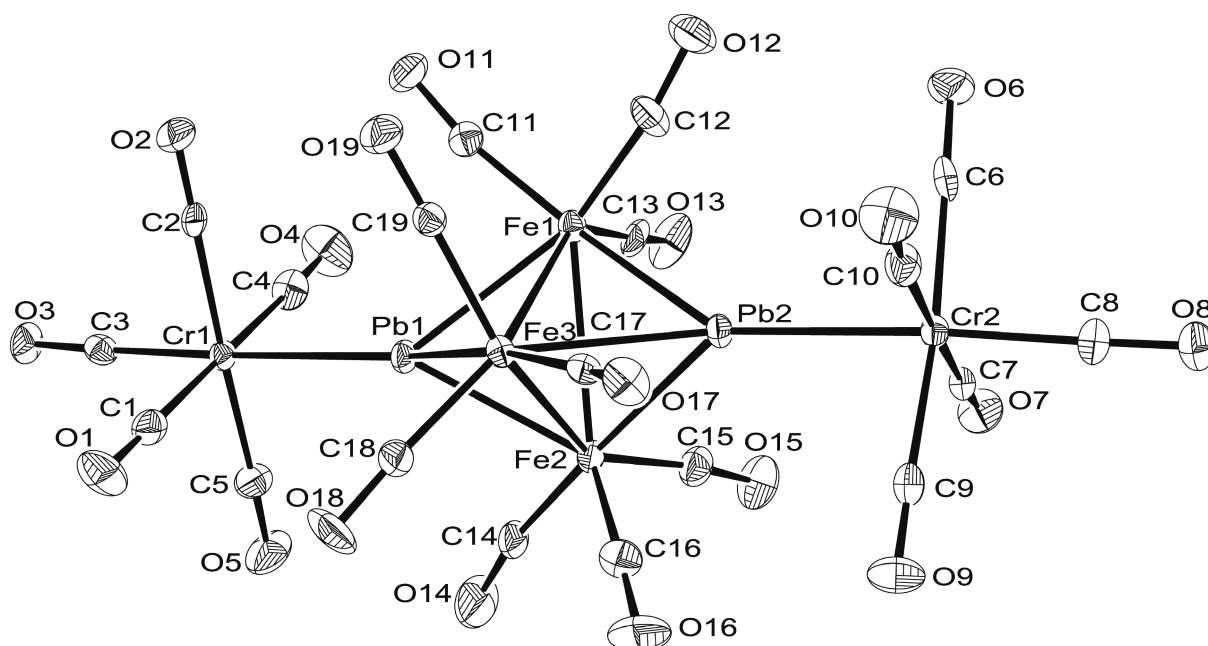
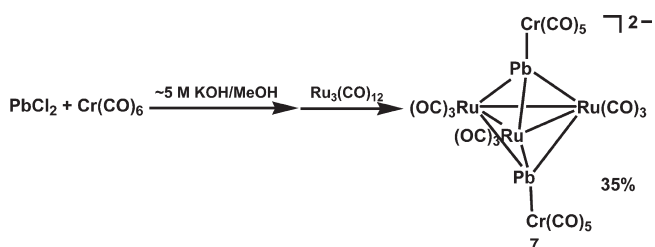


Figure 4. ORTEP diagram of anion **6**, showing 30% probability thermal ellipsoids. Selected bond lengths (Å) and angles (deg): Pb1–Fe1 2.653(1), Pb1–Fe2 2.640(1), Pb1–Fe3 2.661(1), Pb1–Cr1 2.684(1), Pb2–Fe1 2.611(1), Pb2–Fe2 2.626(1), Pb2–Fe3 2.699(1), Pb2–Cr2 2.664(1), Fe1–Fe2 2.859(2), Fe2–Fe3 2.748(2), Fe3–Fe1 2.754(2); Fe1–Fe2–Fe3 58.78(4), Fe2–Fe3–Fe1 62.63(5), Fe3–Fe1–Fe2 58.59(4).

Scheme 4



The calculation showed that the active sites of complexes **1–3** for the formation of complexes **4–6** could be related to the frontier orbitals of complexes **1–3**. As shown in Figure 6, the lowest unoccupied molecular orbital (LUMO) and LUMO+1 of complexes **1–3** had a significant contribution from the p_z or s orbital of the Pb atom, with almost the same energy. It was noted that the LUMO of complex **1**

was similar to the LUMO+1 of **2** and **3**, while the LUMO+1 of complex **1** was similar to the LUMOs of complexes **2** and **3**. Thus, it was reasonable to propose that the incoming Br^- could interact with the Pb atom of complex **1** first, accompanied by a further coupling reaction to give rise to complex **4**. This process also postulated that Br^- attacked the positively charged Pb atom (+0.646) of **1** followed by the loss of $[\text{Cr}(\text{CO})_5]^{2-}$ to give the proposed intermediate “ $[\text{BrPb}\{\text{Cr}(\text{CO})_5\}_2]^-$ ”, where the negatively charged Cr atom (−1.443) further interacted with the Pb atom (+0.917) of another molecule of the intermediate, accompanied by the release of CO units to give the dimeric product (Scheme 5). This assumption was supported by the fact that the As center of the isoelectronic $\text{ClAs}\{\text{Cr}(\text{CO})_5\}_2$ was susceptible to nucleophilic species.²⁵ Further reaction

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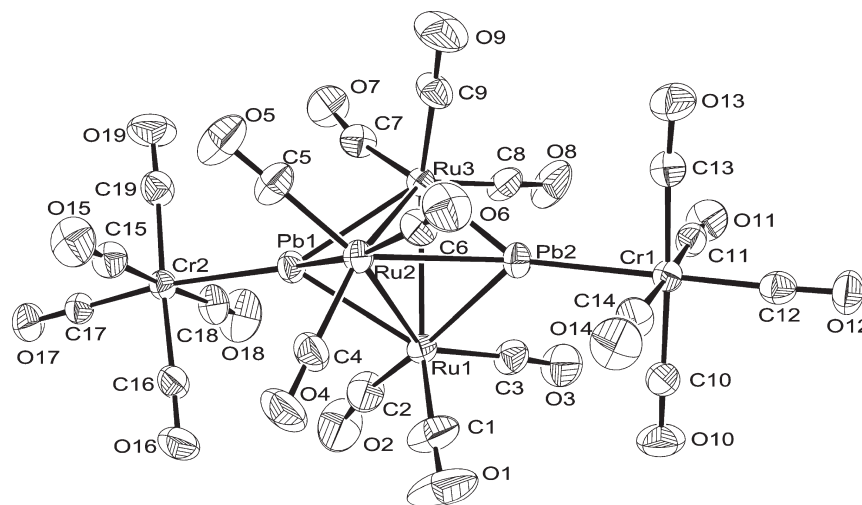


Figure 5. ORTEP diagram of anion **7**, showing 30% probability thermal ellipsoids. Selected bond lengths (Å) and angles (deg): Pb1–Ru1 2.7715(6), Pb1–Ru2 2.7452(6), Pb1–Ru3 2.7809(6), Pb1–Cr2 2.682(1), Pb2–Ru1 2.7458(6), Pb2–Ru2 2.8020(6), Pb2–Ru3 2.7594(6), Pb2–Cr1 2.677(1), Ru1–Ru2 2.9805(8), Ru2–Ru3 2.9526(8), Ru3–Ru1 2.9744(8); Ru1–Ru2–Ru3 60.17(2), Ru2–Ru3–Ru1 60.38(2), Ru3–Ru1–Ru2 59.45(2).

Table 2. Results of NBO and NPA of Complexes **1–7**, the Proposed Intermediates $[\text{BrPb}\{\text{Cr}(\text{CO})_5\}_2]^-$ and $[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Ru}(\text{CO})_4\}_2]^{2-}$, and $\text{Ru}_3(\text{CO})_{12}$

| complex | Wiberg bond index | | | | | natural charge | | | |
|--|--|-------|-------|--------------------|----------------------------------|----------------|--|--|--------|
| | Pb–Cr | Pb–Fe | Pb–Ru | Pb–M _{eq} | M _{eq} –M _{eq} | Pb | Cr | Fe | Ru |
| 1 | 0.317 | | | | | 0.646 | –1.401 | | |
| 2 | | 0.313 | | | | 0.619 | | –0.554 | |
| 3 | 0.302 | 0.321 | | | | 0.619 | –1.389 | –0.555 | |
| $[\text{BrPb}\{\text{Cr}(\text{CO})_5\}_2]^-$ | 0.385 | | | | | 0.917 | –1.443 | | |
| 4 | 0.258 ^a 0.377 ^b | | | | | 0.763 | –1.068 ^a –1.424 ^b | | |
| 5 | | 0.340 | | 0.270 | 0.131 | 0.492 | | –0.469 ^a –0.530 ^b | |
| 6 | 0.326 | | | 0.278 | 0.127 | 0.463 | –1.374 | –0.455 | |
| 7 | 0.328 | | | 0.287 | 0.112 | 0.320 | –1.379 | | –0.406 |
| $[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Ru}(\text{CO})_4\}_2]^{2-}$ | 0.310 | | 0.277 | | | 0.576 | –1.389 | | –0.325 |
| $\text{Ru}_3(\text{CO})_{12}$ | | | | | 0.193 | | | | –0.342 |

^a Ring. ^b Terminal.

of complex **4** with $\text{Mn}(\text{CO})_5\text{Br}$ also involved the attack of Br^- on the positively charged Pb atom (+0.763) of **4**, with Pb–Cr bond cleavage. In addition, the calculated Wiberg bond indices (also called the bond orders) of complex **4** reflected a weak Pb–Cr(CO)₄ bond, which could have facilitated further Br^- attacks on Pb atoms, leading to the formation of the known complex $[\text{PbBr}_2\text{Cr}_2(\text{CO})_{10}]^{2-}$ (Table 2).^{11b}

The formation of the trigonal-bipyramidal clusters **5** and **6** from the trigonal-planar complexes **2** and **3**, respectively, was then examined. The highest occupied molecular orbital (HOMO) and HOMO–1 of complex **2** were nearly degenerate with the significant contributions from the p and d orbitals of the Fe atoms (Figure 6). It was also found that the HOMO of **2** was similar to the HOMO–1 of **3**, while the HOMO–1 of **2** was similar to the HOMO of complex **3**. As mentioned above, the LUMO and LUMO+1 of complexes **2** and **3** were also close in energy with the significant contribution from the s or p_z orbital of the Pb atom (Figure 6). Therefore, it is reasonable to suggest that the negatively charged Fe atom of complex **2** (–0.554) or **3** (–0.555) underwent nucleophilic attack on the positively charged Pb atom of another molecule of complex **2** (+0.619) or **3** (+0.619), with Pb–Fe and Fe–Fe bond formation and the loss of one $[\text{Fe}(\text{CO})_4]^{2-}$ and three CO ligands. As shown in Table 2, it

should be mentioned that the Pb atom in complexes **5** and **6** became less positively charged (+0.492 or +0.463) and the Fe atom became less negatively charged (–0.469, –0.530, or –0.455) compared to **2** and **3**, indicating the stronger covalent character of the Pb–Fe bonds in **5** and **6**. Moreover, calculations gave $\Delta G = -5.502$ kcal/mol for the conversion of **5** to **6**, which supported the experimental results that the conversion was reversible upon the addition of $[\text{Cr}(\text{CO})_5]^{2-}$ or $[\text{Fe}(\text{CO})_4]^{2-}$ but was easier for the transformation of complex **5** to **6**. Further NPA showed that the Pb atom in **5** carried a positive charge of +0.492 and the Fe atom in the $\text{Fe}(\text{CO})_4$ fragment carried a charge of –0.530, while the Pb atom in **6** carried a charge of +0.463 and the Cr atom in the $\text{Cr}(\text{CO})_5$ group carried a negative charge of –1.374, suggestive of the larger coulomb attraction between the Pb atom and $\text{Cr}(\text{CO})_5$ in cluster **6**.

The failure to observe the trigonal-planar intermediate $[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Ru}(\text{CO})_4\}_2]^{2-}$ from the reaction of PbCl_2 , $\text{Cr}(\text{CO})_6$, and $\text{Ru}_3(\text{CO})_{12}$ was probably because of weak Pb–Ru bonds. This phenomenon was supported by the calculation that the Wiberg bond index of the Pb–Ru bond (0.277) for the proposed intermediate “ $[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Ru}(\text{CO})_4\}_2]^{2-}$ ” was significantly smaller than those for the Pb–Cr (0.302–0.317) and Pb–Fe (0.313–0.321) bonds in the corresponding triangular-planar lead complexes

Scheme 5

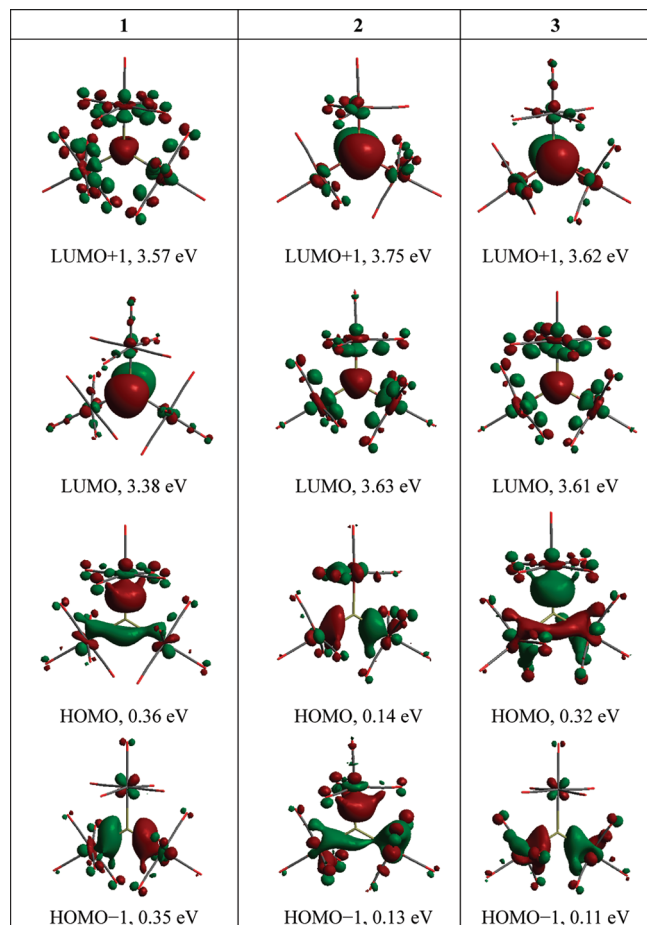
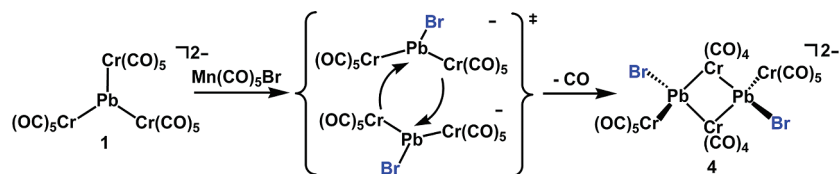


Figure 6. Spatial graphs (isovalue = 0.04–0.05) of the selected frontier molecular orbitals of complexes 1–3.

1–3. In addition, the Wiberg bond index of the Ru–Ru bond in cluster **7** was 0.112, smaller than that in $\text{Ru}_3(\text{CO})_{12}$ (0.193) but comparable to that (0.09) for $[\{\text{PPh}_4\}_2\{\text{Te}_2\text{Ru}_4(\text{CO})_{10}\text{Cu}_4\text{Br}_2\text{Cl}_2\} \cdot \text{THF}]_\infty$ (THF = tetrahydrofuran) in the Te–Ru system.^{9a}

Electrochemistry of 5–7. In view of the highly symmetric Pb_2M_3 ($\text{M} = \text{Fe}, \text{Ru}$) metal cores of the analogous complexes **5–7**, their electrochemical properties were investigated using cyclic voltammetry (CV) and differential pulse voltammetry (DPV) in MeCN under N_2 , which could be further compared with the related clusters $\text{M}_3(\text{CO})_{12}$ ($\text{M} = \text{Fe}, \text{Ru}$).²⁶ The electrochemical data for each complex studied are listed in Tables 3 and S1 in the Supporting Information.

The cyclic voltammograms of Fe_3Pb_2 trigonal-bipyramidal clusters **5** and **6** exhibited three quasi-reversible reductions (around –1.31 to –2.21 V), in which some of

Table 3. DPV of **5–7**, $\text{Fe}_3(\text{CO})_{12}$, and $\text{Ru}_3(\text{CO})_{12}$

| complex | reduction process $E_p^{\text{red}}/\text{V}^a$ ($W_{1/2}/\text{mV}^b$) | | |
|-------------------------------|--|--------------|--------------|
| 5 | –1.238 (107) | –1.550 (171) | –2.102 (110) |
| 6 | –1.430 (97) | –1.694 (150) | –2.074 (93) |
| $\text{Fe}_3(\text{CO})_{12}$ | –0.440 (92) | –0.904 (200) | –2.028 (120) |
| 7 | –1.748 (132) | –2.016 (88) | –2.244 (150) |
| $\text{Ru}_3(\text{CO})_{12}$ | –0.898 (–) ^c | –1.106 (160) | –2.078 (104) |

^a E_p^{red} = reductive peak potential. ^b $W_{1/2}$ = width at half-height. ^c Difficult to determine.

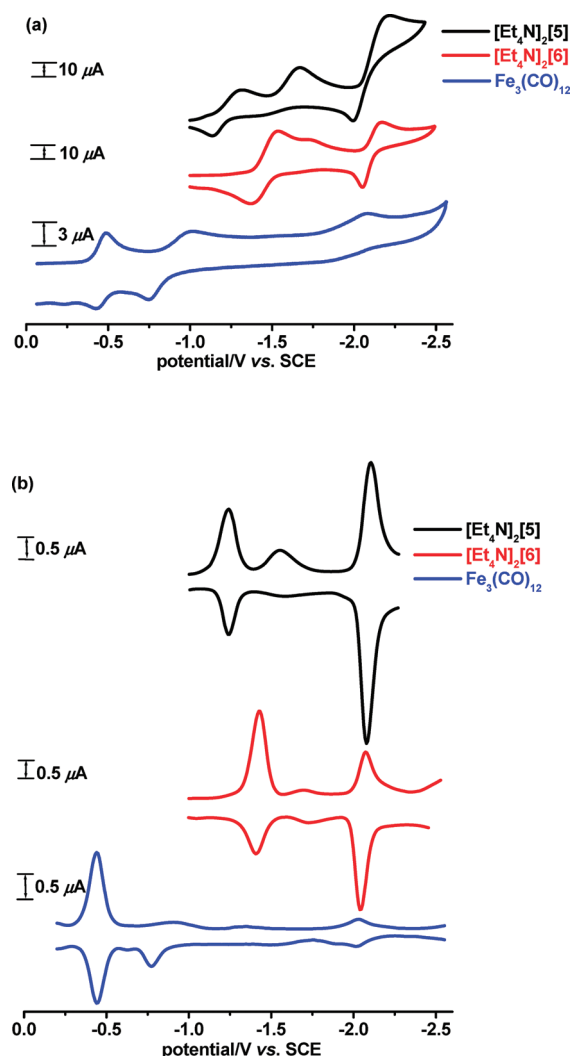


Figure 7. CV (a) and DPV (b) in MeCN for $[\text{Et}_4\text{N}]_2[\text{5}]$, $[\text{Et}_4\text{N}]_2[\text{6}]$, and $\text{Fe}_3(\text{CO})_{12}$. Conditions: electrolyte, 0.1 M Bu_4NClO_4 ; working electrode, glassy carbon; scan rate, 100 mV/s. Potentials are versus SCE.

the peaks were unresolved (Figure 7a). Further DPV results revealed that **5** and **6** each had three quasi-reversible redox reductions at –1.238, –1.550, and –2.102 V and at –1.430, –1.694, and –2.074 V, respectively (Figure 7b). In light of

(26) Bond, A. M.; Dawson, P. A.; Peake, B. M.; Robinson, B. H.; Simpson, J. *Inorg. Chem.* **1977**, *16*, 2199–2206.

Table 4. Reduction Potentials and Electronic Affinities of Complexes 5–7

| complex | reduction potential E^a (V) | EA ^b (kcal/mol) |
|---------|-------------------------------|----------------------------|
| 5 | −1.238 | 78.13 |
| 6 | −1.430 | 81.57 |
| 7 | −1.748 | 86.09 |

^a From DPV. ^b The EA can be estimated by $EA \approx E_{\text{elec}}(N+1) - E_{\text{elec}}(N)$, where $E_{\text{elec}}(N+1)$ and $E_{\text{elec}}(N)$ represent the electronic energies of molecules with $N+1$ and N electrons, respectively.²⁷

the DFT calculations, the LUMO contributions of **5** and **6** predominantly came from the p and d orbitals of the equatorial Fe_3 moieties, indicating that the reduction processes of **5** and **6** occurred in the triangular $\text{Fe}_3(\text{CO})_9$ ring (Figure S2 in the Supporting Information). The results suggested that complexes **5** and **6** [$[\text{Fe}_3(\text{CO})_9\{\text{PbM}(\text{CO})_x\}_2]^{2-}$ ($M = \text{Fe}$, $x = 4, 5$; $M = \text{Cr}$, $x = 5, 6$)] could be reduced to form the square-pyramidal complexes $[\text{Fe}_3(\text{CO})_9\{\text{PbM}(\text{CO})_x\}_2]^{4-}$, with Fe–Fe bond breakage, which paralleled the chemistry of $[\text{Fe}_3(\mu_3\text{-Bi})_2(\text{CO})_9]$.^{17f} As shown in Figure 7b, $\text{Fe}_3(\text{CO})_{12}$ revealed a similar pattern of three redox couples at −0.440, −0.904, and −2.028 V. Compared with $\text{Fe}_3(\text{CO})_{12}$, the reduction potentials of clusters **5** and **6** were shifted to more negative values, basically because of the negative charges of the complexes. Furthermore, the first quasi-reversible reduction of complex **6** (−1.430 V) revealed a significant cathodic shift, in contrast with complex **5** (−1.238 V), because of the larger electron-donating ability of the apical $\text{Cr}(\text{CO})_5$ versus $\text{Fe}(\text{CO})_4$ fragments. This result was also supported by calculations that indicated that complex **6** had a smaller electron affinity, compared with complex **5** [$EA \approx E_{\text{elec}}(N+1) - E_{\text{elec}}(N)$,²⁷ 81.57 vs 78.13 kcal/mol; Table 4].

On the other hand, as shown in Figure 8, the DPV study of the Ru_3Pb_2 trigonal-bipyramidal cluster **7** exhibited three quasi-reversible reductions (−1.748, −2.016, and −2.244 V) that should have occurred around the equatorial $\text{Ru}_3(\text{CO})_9$ ring, in view of the major contribution of the LUMO from the triangular ruthenium moiety (Figure S2 in the Supporting Information). Similar to the case for the Fe_3Pb_2 trigonal-bipyramidal clusters **5** and **6**, the Ru_3Pb_2 complex **7** also showed significant cathodic shifts compared with those of $\text{Ru}_3(\text{CO})_{12}$. Moreover, the first quasi-reversible reduction of **7** (−1.748 V) versus **6** (−1.430 V) revealed that **7** was more difficult to reduce than the isostructural cluster **6**, which is also supported by the smaller electron affinity calculated for **7** compared with **6** (Table 4).

Conclusion

The facile synthesis of the first examples of mixed-metal carbonyl lead-containing trigonal-planar complex **3** and trigonal-bipyramidal clusters $[\text{M}_3(\text{CO})_9\{\text{PbCr}(\text{CO})_5\}_2]^{2-}$ ($M = \text{Fe}, \text{Ru}$) was demonstrated. The nature, formation, and structural transformation of these new complexes were systematically described and compared in terms of the characteristics of transition metals. In addition, three Fe_3Pb_2 - or Ru_3Pb_2 -based trigonal-bipyramidal closo clusters demonstrated a wide range of quasi-reversible redox ability. More interestingly, pronounced cathodic shifts were observed from the Fe_3 to the Ru_3 ring or as the apical fragment changed from an $\text{Fe}(\text{CO})_4$ fragment to a $\text{Cr}(\text{CO})_5$ fragment, which was illustrated well by the results of DFT calculations. This

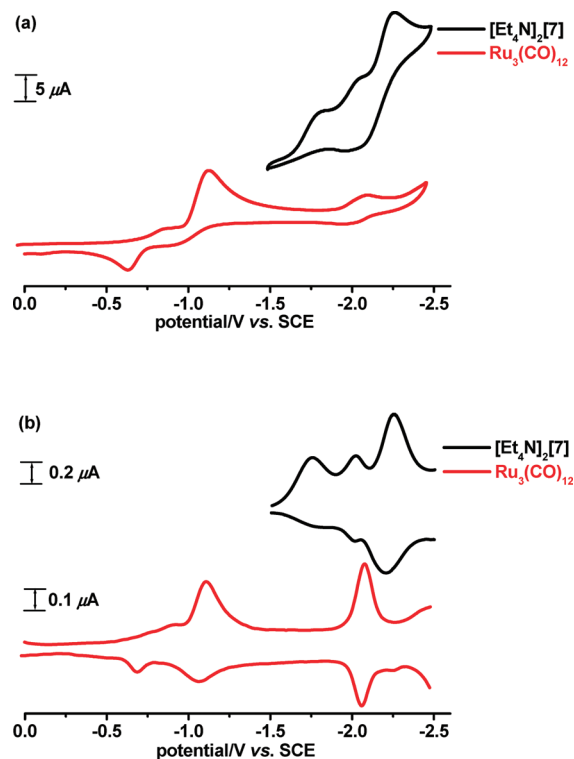


Figure 8. CV (a) and DPV (b) in MeCN for $[\text{Et}_4\text{N}]_2[\mathbf{7}]$ and $\text{Ru}_3(\text{CO})_{12}$. Conditions: electrolyte, 0.1 M Bu_4NClO_4 ; working electrode, glassy carbon; scan rate, 100 mV/s. Potentials are versus SCE.

study provides a new avenue to a systematic comparison of the complicated electronic effects of mixed-metal clusters on the basis of electrochemical and theoretical studies.

Experimental Section

All reactions were performed under an atmosphere of pure nitrogen using standard Schlenk techniques.²⁸ Solvents were purified, dried, and distilled under nitrogen prior to use. $\text{Cr}(\text{CO})_6$ (Strem), $\text{Fe}(\text{CO})_5$ (Strem), $\text{Ru}_3(\text{CO})_{12}$ (Strem), $\text{Mn}(\text{CO})_5\text{Br}$ (Strem), PbCl_2 (Osaka), and KOH (Showa) were used as received. The compound $[\text{Et}_4\text{N}]_2[\text{Pb}\{\text{Fe}(\text{CO})_4\}_3]$ ($[\text{Et}_4\text{N}]_2[\mathbf{2}]$) was prepared according to the published method.^{12a} IR spectra were recorded on a Perkin-Elmer Paragon 500 IR spectrometer as solutions in CaF_2 cells. Elemental analyses for C, H, and N were performed on a Perkin-Elmer 2400 analyzer at the NSC Regional Instrumental Center at National Taiwan University, Taipei, Taiwan. Electrochemical measurements were performed at room temperature under a nitrogen atmosphere and recorded using a BAS-100W electrochemical potentiostat. A glassy carbon working electrode, a platinum wire auxiliary electrode, and a non-aqueous Ag/Ag^+ electrode were used in a three-electrode configuration. Tetra-*n*-butylammonium perchlorate was used as the supporting electrolyte, and the solute concentration was $\sim 10^{-3}$ M. The redox potentials were calibrated with a ferrocenium/ferrocene (Fc^+/Fc) couple in the working solution and referenced to a saturated calomel electrode (SCE).

Synthesis of $[\text{Et}_4\text{N}]_2[\text{Pb}\{\text{Cr}(\text{CO})_5\}_3]$ ($[\text{Et}_4\text{N}]_2[\mathbf{1}]$). To a MeOH solution (15 mL) of $\text{Cr}(\text{CO})_6$ (0.68 g, 3.09 mmol) and KOH (4.21 g, 75.0 mmol) was added PbCl_2 (0.28 g, 1.01 mmol) in an ice/water bath. The resulting solution was stirred at ambient temperature for 4 h to give a reddish-brown solution. The solution was filtered and concentrated, and a MeOH solution of $[\text{Et}_4\text{N}]\text{Br}$ (0.35 g,

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(28) Shriver, D. F.; Drezdon, M. A. *The Manipulation of Air-Sensitive Compounds*; Wiley-VCH Publishers: New York, 1986.

Table 5. Crystallographic Data for [Bu₄N]₂[3], [Et₄N]₂[4], [Et₄N]₂[5], [Et₄N]₂[6], and [Et₄N]₂[7]

| | [Bu ₄ N] ₂ [3] | [Et ₄ N] ₂ [4] | [Et ₄ N] ₂ [5] | [Et ₄ N] ₂ [6] | [Et ₄ N] ₂ [7] |
|--|--|---|---|---|---|
| empirical formula | C ₄₅ H ₇₂ CrFe ₂ -N ₂ O ₁₃ Pb | C ₃₄ H ₄₀ Br ₂ Cr ₄ -N ₂ O ₁₈ Pb ₂ | C ₃₃ H ₄₀ Fe ₅ -N ₂ O ₁₇ Pb ₂ | C ₃₅ H ₄₀ Cr ₂ -Fe ₃ N ₂ O ₁₉ Pb ₂ | C ₃₅ H ₄₀ Cr ₂ -Ru ₃ N ₂ O ₁₉ Pb ₂ |
| fw | 1219.94 | 1546.89 | 1430.30 | 1478.62 | 1614.31 |
| cryst syst | monoclinic | monoclinic | monoclinic | monoclinic | monoclinic |
| space group | <i>P</i> 2 ₁ / <i>n</i> | <i>P</i> 2 ₁ / <i>c</i> | <i>C</i> 2/ <i>c</i> | <i>P</i> 2 ₁ / <i>c</i> | <i>P</i> 2 ₁ / <i>n</i> |
| crystal dimens, mm | 0.40 × 0.25 × 0.20 | 0.40 × 0.05 × 0.03 | 0.41 × 0.24 × 0.07 | 0.25 × 0.22 × 0.18 | 0.20 × 0.12 × 0.10 |
| <i>a</i> , Å | 12.281(3) | 9.5681(2) | 28.3677(6) | 12.9995(2) | 12.1695(1) |
| <i>b</i> , Å | 33.391(1) | 24.6016(6) | 10.6192(3) | 18.8599(3) | 18.2147(3) |
| <i>c</i> , Å | 13.865(4) | 11.3668(3) | 17.6363(4) | 20.5628(4) | 23.5738(3) |
| α, deg | | | | | |
| β, deg | 90.32(2) | 101.805(1) | 120.414(2) | 107.506(1) | 103.0966(6) |
| γ, deg | | | | | |
| <i>V</i> , Å ³ | 5686(3) | 2619.1(1) | 4581.7(2) | 4807.9(1) | 5089.5(1) |
| <i>Z</i> | 4 | 2 | 4 | 4 | 4 |
| <i>D</i> (calcd), g/cm ³ | 1.425 | 2.007 | 2.074 | 2.043 | 2.107 |
| μ, mm ⁻¹ | 3.692 | 8.799 | 8.926 | 8.362 | 7.935 |
| color, habit | black, prism | black, prism | black, prism | black, prism | red, prism |
| diffractometer | Nonius (CAD4) | Nonius (Kappa CCD) | Nonius (Kappa CCD) | Nonius (Kappa CCD) | Nonius (Kappa CCD) |
| radiation (λ), Å | 0.710 73 | 0.710 73 | 0.710 73 | 0.710 73 | 0.710 73 |
| temp, K | 298 | 298 | 200 | 298 | 298 |
| θ range for data collcn, deg | 1.59–24.92 | 3.53–27.71 | 2.09–25.33 | 2.08–25.34 | 4.10–27.47 |
| <i>T</i> _{min} / <i>T</i> _{max} | 0.32/0.53 | 0.30/0.41 | 0.12/0.57 | 0.17/0.20 | 0.29/0.39 |
| no. of indep reflns [<i>I</i> > 2 σ(<i>I</i>)] | 4646 (<i>R</i> _{int} = 0.0633) | 4481 (<i>R</i> _{int} = 0.0928) | 3521 (<i>R</i> _{int} = 0.0798) | 6745 (<i>R</i> _{int} = 0.0747) | 7441 (<i>R</i> _{int} = 0.0690) |
| no. of param | 572 | 280 | 286 | 568 | 568 |
| <i>R</i> ¹ / <i>wR</i> ² [<i>I</i> > 2 σ(<i>I</i>)] | 0.060/0.138 | 0.071/0.172 | 0.057/0.144 | 0.042/0.106 | 0.045/0.082 |
| <i>R</i> ¹ / <i>wR</i> ² (all data) | 0.165/0.179 | 0.095/0.184 | 0.072/0.165 | 0.068/0.140 | 0.089/0.094 |

^a The functions minimized during least-squares cycles were $R1 = \sum ||F_o| - |F_c|| / \sum |F_o|$ and $wR2 = [\sum [w(F_o^2 - F_c^2)^2] / \sum [w(F_o^2)^2]]^{1/2}$.

1.67 mmol) was added dropwise, precipitating the solid. The solid was washed with deionized water in an ice/water bath several times and dried under vacuum. The THF extract was recrystallized from Et₂O/THF to give a reddish-brown sample of [Et₄N]₂[1] (yield 0.29 g, 0.28 mmol, 28% based on PbCl₂). IR (ν_{CO}, THF): 2010 (w), 1977 (s), 1909 (vs), 1864 (m) cm⁻¹.

Synthesis of [Bu₄N]₂[Pb{Cr(CO)₅}₂{Fe(CO)₄}₂] ([Bu₄N]₂[3]). A MeOH solution (20 mL) of PbCl₂ (0.77 g, 2.77 mmol), Cr(CO)₆ (0.61 g, 2.77 mmol), and KOH (5.20 g, 92.7 mmol) was stirred in an ice/water bath. The solution was stirred at ambient temperature for 2 h, and Fe(CO)₅ (0.73 mL, 5.55 mmol) was added. The reaction solution turned brown and was allowed to stir for 12 h. The resulting solution was filtered and concentrated, and an aqueous solution of [Bu₄N]Br (0.70 g, 2.17 mmol) was added dropwise, precipitating the solid. The solid was washed with deionized water several times and dried under vacuum. The THF extract was recrystallized from Et₂O/THF to give a reddish-brown sample of [Bu₄N]₂[3] (yield 1.12 g, 0.92 mmol, 33% based on PbCl₂). IR (ν_{CO}, THF): 2025 (m), 1972 (s), 1957 (vs), 1910 (s), 1878 (s) cm⁻¹. Anal. Calcd for [Bu₄N]₂[3]: C, 44.30; H, 5.95; N, 2.30. Found: C, 44.24; H, 6.19; N, 2.57. Crystals of [Bu₄N]₂[3] suitable for X-ray diffraction were grown from Et₂O/THF. [Et₄N]₂[3] can be isolated by following a similar procedure.

Reaction of [Et₄N]₂[1] with Mn(CO)₅Br. To a MeCN solution (30 mL) of [Et₄N]₂[1] (0.36 g, 0.34 mmol) was added Mn(CO)₅Br (0.14 g, 0.51 mmol) at ambient temperature. The resulting solution was stirred for 30 min to give a yellow-brown solution. The solution was filtered, and the solvent was evaporated under vacuum. The residue was washed with Et₂O, which was shown to contain Mn₂(CO)₁₀ and [HCr₂(CO)₁₀] by IR spectroscopy. The residue was extracted with CH₂Cl₂ to give a sample of [Et₄N]₂[Pb₂Br₂Cr₄(CO)₁₈] ([Et₄N]₂[4]) (yield 0.14 g, 0.09 mmol, 53% based on Pb). IR (ν_{CO}, CH₂Cl₂): 2036 (s), 2012 (m), 1984 (s), 1959 (m), 1943 (m), 1925 (s), 1903 (m) cm⁻¹. Anal. Calcd for [Et₄N]₂[4]: C, 26.40; H, 2.61; N, 1.81. Found: C, 26.41; H, 2.80; N, 1.66. Crystals of [Et₄N]₂[4] suitable for single crystal X-ray diffraction were grown from Et₂O/CH₂Cl₂.

Reaction of [Et₄N]₂[4] with Mn(CO)₅Br. To a MeCN solution (10 mL) of [Et₄N]₂[Pb₂Br₂Cr₄(CO)₁₈] ([Et₄N]₂[4]) (0.14 g, 0.09 mmol) was added Mn(CO)₅Br (0.05 g, 0.18 mmol). The mixed solution was stirred at ambient temperature for 21 h. The solution was filtered, and the solvent was evaporated under vacuum. The residue was extracted with CH₂Cl₂ to give a sample of [Et₄N]₂[PbBr₂Cr₂(CO)₁₀]^{11b} (yield 0.06 g, 0.06 mmol, 33% based on [Et₄N]₂[Pb₂Br₂Cr₄(CO)₁₈]). IR (ν_{CO}, CH₂Cl₂): 1990 (m), 1953 (vs), 1893 (m), 1859 (m) cm⁻¹.

Synthesis of [Et₄N]₂[Fe₃(CO)₉{PbFe(CO)₄}₂] ([Et₄N]₂[5]). **Method 1.** A MeCN solution (30 mL) of [Et₄N]₂[Pb{Fe(CO)₄}₃] ([Et₄N]₂[2]) (2.02 g, 2.08 mmol) and Mn(CO)₅Br (0.86 g, 3.13 mmol) was stirred and heated at 80 °C for 42 h. The resultant yellowish-brown solution was filtered, and the solvent was evaporated under vacuum. The residue was washed several times with deionized water and Et₂O, which was shown to contain Mn₂(CO)₁₀ by IR spectroscopy. The CH₂Cl₂ extract was recrystallized from Et₂O/MeOH/CH₂Cl₂ to give a sample of [Et₄N]₂[5] (yield 0.89 g, 0.62 mmol, 60% based on Pb). IR (ν_{CO}, CH₂Cl₂): 2011 (s), 1955 (vs), 1911 (m) cm⁻¹. Anal. Calcd for [Et₄N]₂[5]: C, 27.71; H, 2.82; N, 1.96. Found: C, 27.66; H, 2.49; N, 2.05. Crystals of [Et₄N]₂[5] suitable for single-crystal X-ray diffraction were grown from Et₂O/CH₂Cl₂.

Method 2. To a MeCN solution (20 mL) of [Et₄N]₂[Fe₃(CO)₉{PbCr(CO)₅}₂] ([Et₄N]₂[6]) (0.61 g, 0.41 mmol) was added Fe(CO)₅ (0.11 mL, 0.84 mmol), KOH (4.14 g, 73.8 mmol), and MeOH (30 mL). The mixed solution was stirred at ambient temperature for 40 h. The solution was filtered, and the solvent was evaporated under vacuum. The residue was washed several times with Et₂O and extracted with CH₂Cl₂ to give a sample of [Et₄N]₂[5] (yield 0.46 g, 0.32 mmol, 78% based on Pb).

Synthesis of [Et₄N]₂[Fe₃(CO)₉{PbCr(CO)₅}₂] ([Et₄N]₂[6]). **Method 1.** To a MeCN solution (40 mL) of [Et₄N]₂[3] (1.00 g, 1.00 mmol) was added Mn(CO)₅Br (0.36 g, 1.31 mmol). The mixed solution was stirred and heated to 80 °C for 2 days. The solution was filtered, and the solvent was evaporated under vacuum. The residue was washed several times with Et₂O, which was shown by IR spectroscopy to contain Mn₂(CO)₁₀. The residue

was then extracted with CH_2Cl_2 to give a sample of $[\text{Et}_4\text{N}]_2[\mathbf{6}]$ (yield 0.46 g, 0.31 mmol, 62% based on Pb). IR (ν_{CO} , CH_2Cl_2): 2034 (s), 1949 (vs), 1918 (m), 1886 (m) cm^{-1} . Anal. Calcd for $[\text{Et}_4\text{N}]_2[\mathbf{6}]$: C, 28.43; H, 2.73; N, 1.89. Found: C, 28.40; H, 2.81; N, 1.86. Crystals of $[\text{Et}_4\text{N}]_2[\mathbf{6}]$ suitable for single-crystal X-ray diffraction were grown from $\text{Et}_2\text{O}/\text{CH}_2\text{Cl}_2$.

Method 2. A MeOH solution (30 mL) of $\text{Cr}(\text{CO})_6$ (0.10 g, 0.45 mmol) and KOH (4.23 g, 0.075 mmol) was stirred in an ice/water bath and warmed to ambient temperature. After 4 h, $[\text{Et}_4\text{N}]_2[\mathbf{5}]$ (0.29 g, 0.20 mmol) was added to the solution, which was stirred for another 14 h to give a yellowish-brown solution. The solution was filtered, and the solvent was evaporated under vacuum. The residue was washed with deionized water and Et_2O several times and dried under vacuum. The CH_2Cl_2 extract was recrystallized from $\text{Et}_2\text{O}/\text{MeOH}/\text{CH}_2\text{Cl}_2$ to give a sample of $[\text{Et}_4\text{N}]_2[\mathbf{6}]$ (yield 0.18 g, 0.12 mmol, 60% based on Pb).

Synthesis of $[\text{Et}_4\text{N}]_2[\text{Ru}_3(\text{CO})_9\{\text{PbCr}(\text{CO})_5\}_2]$ ($[\text{Et}_4\text{N}]_2[\mathbf{7}]$). A MeOH solution (20 mL) of $\text{Cr}(\text{CO})_6$ (0.15 g, 0.68 mmol) and KOH (5.31 g, 94.6 mmol) was stirred in an ice/water bath and warmed to ambient temperature. After 3 h, PbCl_2 (0.20 g, 0.72 mmol) was added to the solution, which was stirred for another 4 h to give a reddish-brown solution. $\text{Ru}_3(\text{CO})_{12}$ (0.22 g, 0.34 mmol) was then added to the solution and stirred for 36 h at ambient temperature to yield a brown solution, which was filtered and concentrated, and an aqueous solution of $[\text{Et}_4\text{N}]\text{Br}$ (0.31 g, 1.48 mmol) was added dropwise, precipitating the solid. The solid was washed with deionized water several times and dried under vacuum. The CH_2Cl_2 extract was recrystallized from $\text{Et}_2\text{O}/\text{MeOH}/\text{CH}_2\text{Cl}_2$ to give a reddish-brown sample of $[\text{Et}_4\text{N}]_2[\mathbf{7}]$ (yield 0.20 g, 0.12 mmol, 35% based on Ru). IR (ν_{CO} , CH_2Cl_2): 2033 (m), 1970 (vs), 1923 (m), 1887 (m) cm^{-1} . Anal. Calcd for $[\text{Et}_4\text{N}]_2[\mathbf{7}]$: C, 26.04; H, 2.50; N, 1.74. Found: C, 26.01; H, 2.71; N, 1.60. Crystals of $[\text{Et}_4\text{N}]_2[\mathbf{7}]$ suitable for X-ray diffraction were grown from $\text{Et}_2\text{O}/\text{CH}_2\text{Cl}_2$.

X-ray Structural Characterization of $[\text{Bu}_4\text{N}]_2[\mathbf{3}]$, $[\text{Et}_4\text{N}]_2[\mathbf{4}]$, $[\text{Et}_4\text{N}]_2[\mathbf{5}]$, $[\text{Et}_4\text{N}]_2[\mathbf{6}]$, and $[\text{Et}_4\text{N}]_2[\mathbf{7}]$. Some selected crystallographic data for $[\text{Bu}_4\text{N}]_2[\mathbf{3}]$, $[\text{Et}_4\text{N}]_2[\mathbf{4}]$, $[\text{Et}_4\text{N}]_2[\mathbf{5}]$, $[\text{Et}_4\text{N}]_2[\mathbf{6}]$, and $[\text{Et}_4\text{N}]_2[\mathbf{7}]$ are given in Table 5. All crystals were mounted on glass fibers with epoxy cement. Data collection for $[\text{Bu}_4\text{N}]_2[\mathbf{3}]$ was carried out using a Nonius (CAD-4) diffractometer with graphite-monochromated Mo K α radiation at 298 K in the 2θ range of $2.0\text{--}50^\circ$ employing $\theta\text{--}2\theta$ scans, and an empirical absorption correction by azimuthal (Ψ) scans was applied.²⁹ Data collection for $[\text{Et}_4\text{N}]_2[\mathbf{4}]$, $[\text{Et}_4\text{N}]_2[\mathbf{5}]$, $[\text{Et}_4\text{N}]_2[\mathbf{6}]$, and $[\text{Et}_4\text{N}]_2[\mathbf{7}]$ was carried out using a Bruker-Nonius Kappa CCD diffractometer with graphite-monochromated Mo K α radiation at 298 K, and an empirical absorption correction by multiscans was applied. The structures of $[\text{Bu}_4\text{N}]_2[\mathbf{3}]$, $[\text{Et}_4\text{N}]_2[\mathbf{4}]$, $[\text{Et}_4\text{N}]_2[\mathbf{5}]$, $[\text{Et}_4\text{N}]_2[\mathbf{6}]$, and $[\text{Et}_4\text{N}]_2[\mathbf{7}]$ were refined by *SHELXL* packages.³⁰ All of the non-H atoms of $[\text{Bu}_4\text{N}]_2[\mathbf{3}]$, $[\text{Et}_4\text{N}]_2[\mathbf{4}]$, $[\text{Et}_4\text{N}]_2[\mathbf{5}]$,

$[\text{Et}_4\text{N}]_2[\mathbf{6}]$, and $[\text{Et}_4\text{N}]_2[\mathbf{7}]$ were refined with anisotropic temperature factors. Additional crystallographic data in the form of CIF files are available as Supporting Information.

Computational Details. All calculations reported in this study were performed via DFT with Becke's three-parameter (B3) exchange functional and the Lee–Yang–Parr (LYP) correlation functional (B3LYP)^{21,22} using the *Gaussian 03* series of packages.³¹ The geometries of complexes **1–7**, $[\text{Cr}(\text{CO})_5]^{2-}$, $[\text{Fe}(\text{CO})_4]^{2-}$, and $\text{Ru}_3(\text{CO})_{12}$ and the proposed intermediates $[\text{BrPb}\{\text{Cr}(\text{CO})_5\}_2]^-$ and $[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Ru}(\text{CO})_4\}_2]^{2-}$ were fully optimized with the Stuttgart/Dresden relativistic effective core potential³² and with the LanL2DZ basis set for Pb, Br, Cr, Fe, and Ru atoms and the 6-31G(d) basis set for C and O atoms. The optimized geometries were further confirmed by harmonic vibrational frequency analysis, obtained via analytical energy second derivatives. Wiberg bond indices³³ and natural charges²⁴ were evaluated using the Weinhold NBO method.²³ Graphical representations of the molecular orbitals were obtained using CS Chem3D 5.0.

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Supporting Information Available: X-ray crystallographic files in CIF format for $[\text{Bu}_4\text{N}]_2[\mathbf{3}]$, $[\text{Et}_4\text{N}]_2[\mathbf{4}]$, $[\text{Et}_4\text{N}]_2[\mathbf{5}]$, $[\text{Et}_4\text{N}]_2[\mathbf{6}]$, and $[\text{Et}_4\text{N}]_2[\mathbf{7}]$, CV, equatorial plane of the structure of **5**, spatial graphs, and computational details for **1–7**, $[\text{Cr}(\text{CO})_5]^{2-}$, $[\text{Fe}(\text{CO})_4]^{2-}$, and $\text{Ru}_3(\text{CO})_{12}$ and the proposed intermediates $[\text{BrPb}\{\text{Cr}(\text{CO})_5\}_2]^-$ and $[\text{Pb}\{\text{Cr}(\text{CO})_5\}\{\text{Ru}(\text{CO})_4\}_2]^{2-}$. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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