

Aryl Oligogermanes as Ligands for Transition Metal Complexes

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Abstract: The ligand properties of a series of aryl oligogermanes $R_3Ge-GeAr_3$, **3-7** ($Me_3Ge-GePh_3$ (**3**), $Me_3Ge-Ge(pTol)_3$ (**4**), $Ph_3Ge-GePh_3$ (**5**), $(C_6F_5)_3Ge-GePh_3$ (**6**), $Ph_3Ge-GeMe_2GePh_3$ (**7**)), for the synthesis of transition metal carbonyl complexes such as $R_3Ge-GeAr_2(R'C_6H_4-\eta^6)M(CO)_3$ ($M = Cr$, **3a-7a**; $M = Mo$, **3b**; $M = W$, **3c**) were investigated. The complexes were obtained, using several different synthetic approaches, in moderate yields. The physicochemical properties of these newly synthesised complexes were investigated by IR, UV/vis, NMR spectroscopy, electrochemistry and DFT calculations. The molecular structures of **3c**, **4a** and **5a** were studied by single crystal X-ray diffraction analysis. A comparative analysis of the donor- and acceptor properties of aryl oligogermanes, as ligands for transition metal carbonyl complexes, is reported.

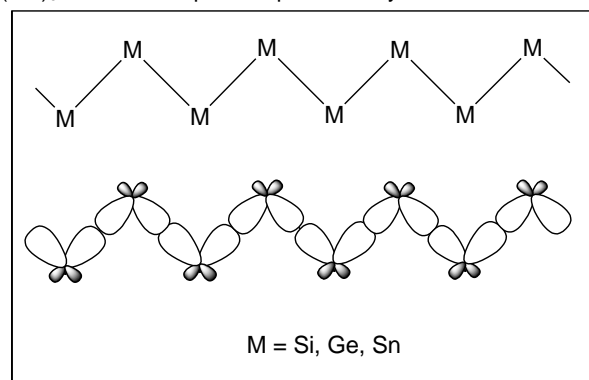
Introduction

Recently, derivatives of Group 14 bearing element – element bonds ($E(IV) = Si, Ge, Sn, Pb$) have attracted significant attention due to their unique physical properties (conductivity, light absorbance, luminescence etc.). These unique properties are due to the presence of a σ -conjugation between the E atoms within the framework of the molecule (Scheme 1).^[1] The number of E atoms in the chain, their nature, the electronic properties of their substituents and the conformation of the molecule significantly affect on the σ -conjugation. The ability to finely tune the electronic properties of these new catenated derivatives

would open the door to the preparation materials displaying new physical properties. Biding transition metals (M) to catenated Group 14 derivatives should impact on the σ -delocalisation and therefore, also on the physical properties of the complex.^[2] In addition, several possible modes of bonding between the metal and the Group 14 ligand could be expected, each of them having a different impact on the σ -conjugation. A few compounds, containing a covalent bond between the E atom and a transition metal, have been reported in the literature (leading references for Si derivatives with $M =$ rare earth metals;^[3] Ti, Zr, Hf;^[4] Ta;^[5] Cr, Mo, W;^[6] Fe;^[7] Pd;^[8] Pt;^[9] Au;^[10] for special interest to Ge derivatives for Hf, Zr;^[4] W, Cr;^[11] Pt;^[9b, 12] Pd;^[13] Au;^[14] for Sn derivatives with Ti;^[4, 15] Mo, W;^[16] Fe^[17] etc.). In the present study, we are reporting the impact of the different types of bonding, including when the metallic fragment is connected to an organic group (η^1 -coordination), on the physical properties of the new compounds. Only a few studies on $\eta^5-C_5R_5$ (Cp) catenated Ge derivatives^[18] and $Cr[\eta^6-C_6H_5GeMe_2]_2$ ^[19] have been reported to date while molecular oligostannanes (Cp^[28] derivatives). Polysilanes^[29] and oligosilane^[20] complexes have been extensively studied (complexed with η^4 -butadiene,^[21] η^5 -half-sandwich,^[6c, 22] ferroceny^[23] or other Cp ligands,^[24] (η^6 -aryl) $M(CO)_3$,^[25] sandwich Cr complexes,^[26] η^7 -derivatives^[27]). Only molecular oligogermanium complexes bearing transition metals remain to be studied to complete the series. To the best of our knowledge, only 6 articles describing complexes of arylmonogermanes with $M(CO)_3$ have been reported up to this day.^[19, 30]

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Supporting information for this article (Crystallographic data, NMR and UV/vis spectra of the compounds obtained, and data of DFT calculations) is given via a link at the end of the document.

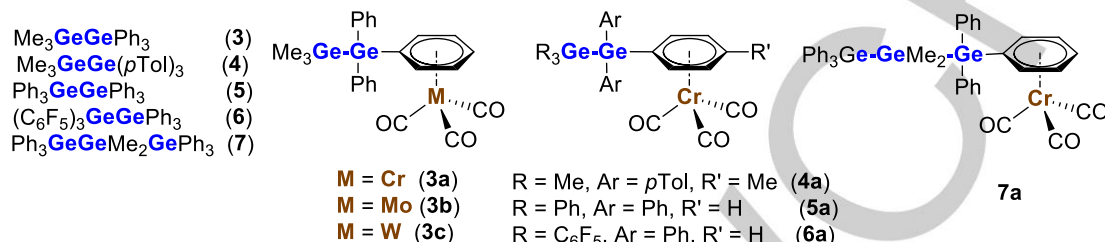


Scheme 1. Schematic representation of σ -conjugation between Group 14 E atoms.

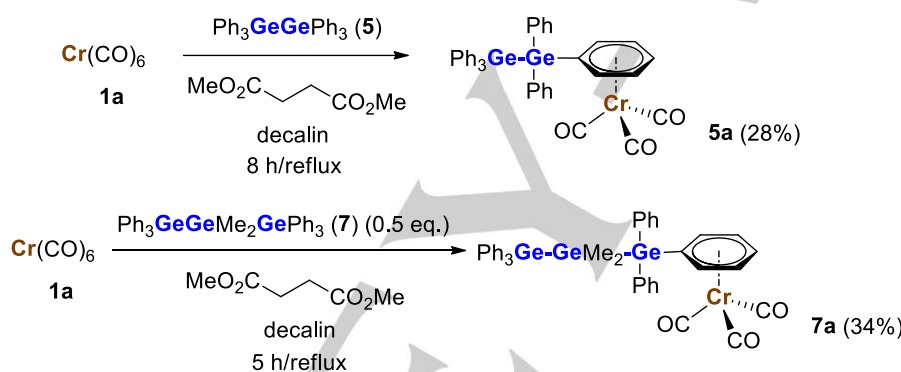
Arene chromium tricarbonyl complexes are valuable compounds that are used in organic synthesis and catalysis,^[31] the preparation of materials with non-linear properties,^[32] etc. In general, a $Cr(CO)_3$ group, coordinated to an arene, acts as a strong electron withdrawing substituent which reduces the electron density of the aryl group. A priori, $M(CO)_3$ would exhibit similar electron-withdrawing properties with a magnitude

comparable to that of a nitro group.^[33] When the arene bears a functional group Y, such as in $(\eta^6\text{-C}_6\text{H}_5\text{Y})\text{Cr}(\text{CO})_3$, the extent of the charge transfer to the metal strongly depends on the electron donating or electron withdrawing properties of the group Y. This has been shown by infrared spectroscopy and is supported by theoretical calculations. In this work, we are reporting the synthesis of the aryloligogermane carbonyl

complexes **3a-c**, **4a-7a** (Scheme 2) as well as an investigation about their structure – physical properties relationships. Oligogermanes $\text{Me}_3\text{Ge-GePh}_3$ (**3**), $\text{Me}_3\text{Ge-Ge}(p\text{Tol})_3$ (**4**), $\text{Ph}_3\text{Ge-GePh}_3$ (**5**), $(\text{C}_6\text{F}_5)_3\text{Ge-GePh}_3$ (**6**), and $[\text{Ph}_3\text{Ge}]_2\text{GeMe}_2$ (**7**) were used as simple and easily available model-compounds, differing in their number of Ge atoms and in the steric and electronic properties of the substituents on their Ge centers.



Scheme 2. Structures of the compounds under investigation.



Scheme 3. Synthesis of **5a** and **7a** using $\text{Cr}(\text{CO})_6$ (**1a**) in decalin.

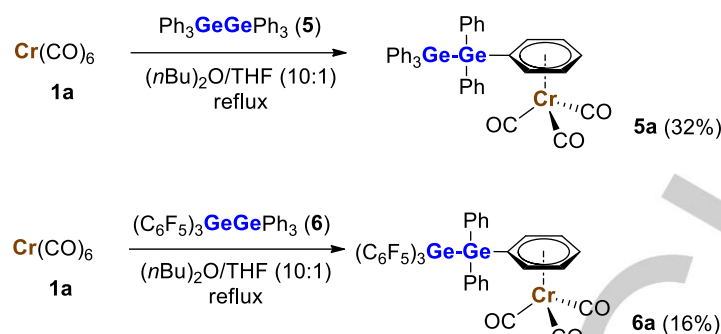
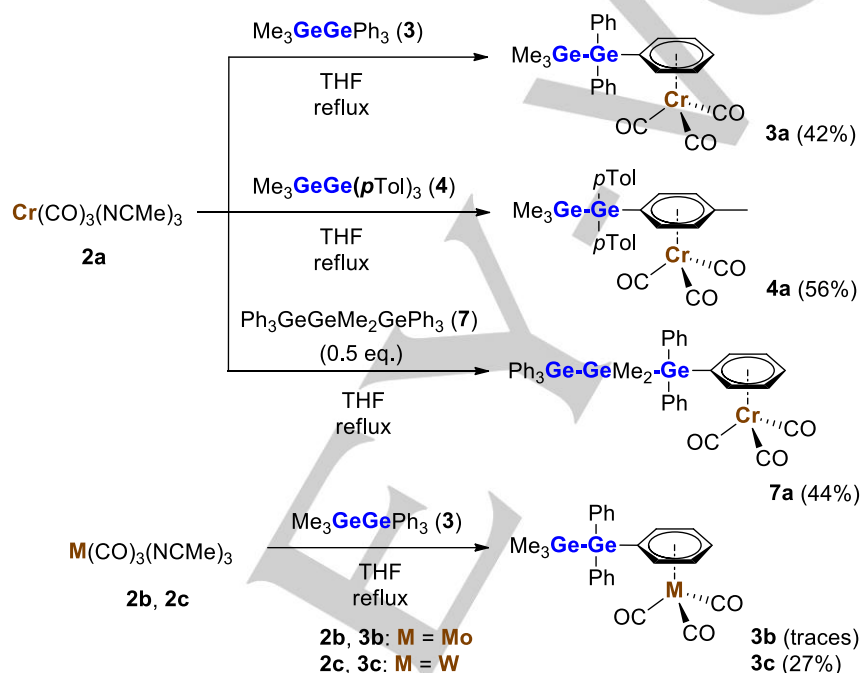
Results and Discussion

Synthesis. The synthesis of the new organometallic complexes required the use of different metallic precursors, $\text{Cr}(\text{CO})_6$ (**1a**) and $\text{M}(\text{CO})_3(\text{NCMe})_3$ ($\text{M} = \text{Cr}$ (**2a**), $\text{M} = \text{Mo}$ (**2b**), $\text{M} = \text{W}$ (**2c**)).

As a first attempt to prepare the new compounds, the thermolysis of $\text{Cr}(\text{CO})_6$, in a solvent with a high boiling point such as decalin,^[34] in the presence of the corresponding arene (aryloligogermane **5** and **7**) and dimethyl succinate as an agent promoting the abstraction of the CO the chromium center,^[35] was investigated (Scheme 3).

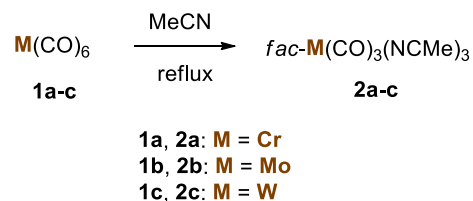
Unfortunately, a significant amount of unreacted starting material was recovered at the end of the reaction, even when the reaction time was extended to 12h and only a low conversion was achieved (29-33%) resulting in low yields in **5a** and **7a**. Using an excess of **1a** did not improve the yield either. The thermolysis approach has shown to have serious drawbacks. In addition to giving low yields in **5a** and **7a**, harsh reaction conditions have to be used to overcome the low reactivity of $\text{Cr}(\text{CO})_6$ and tedious chromatographic separations are often needed to get the pure compounds.

In order to improve the yields and simplify the isolation of the desired compounds, a new set of conditions was investigated. $\text{Cr}(\text{CO})_6$ (**1a**) was reacted with $\text{Ph}_3\text{Ge-GePh}_3$ (**5**) and $(\text{C}_6\text{F}_5)_3\text{Ge-GePh}_3$ (**6**) in a refluxing mixture of $(n\text{Bu})_2\text{O}/\text{THF}$ (10:1)^[34, 36] (Scheme 4). Similar conditions have been reported before for the synthesis of Group 14 derivatives using DME^[37] or dyglime.^[38] When the unsymmetrical donor-acceptor digermane **6** was used, the product **6a** was exclusively formed, where the $\text{Cr}(\text{CO})_3$ coordinates to the electron-rich aromatic ring. The isolation of pure **5a** and **6a** were easily performed by column chromatography (for details, see Experimental part). Unfortunately, once again, the target compounds were isolated in low yields. When benzene was used as the eluent for the chromatographic separation of **6a**, a transmetalation was observed leading to the formation of $(\eta^6\text{-C}_6\text{H}_6)\text{Cr}(\text{CO})_3$; therefore, a mixture of ether and petroleum ether should be used instead of benzene for the isolation of the pure complexes. The yield of the desired complex also depends on the structure of the oligogermane used in the complexation reaction. The use of an oligogermane bearing at least one electron withdrawing group generally results in even lower yields.

Scheme 4. Synthesis of 5a and 6a using Cr(CO)_6 (1a) in $(n\text{Bu})_2\text{O}$.Scheme 6. Synthesis of complexes 3a-c, 4a and 7a using $\text{M(CO)}_3(\text{NCMe})_3$ (2a-c).

When W(CO)_6 was subjected to the same reaction conditions, in the presence of $\text{Me}_3\text{GeGePh}_3$ (3), the desired complex $\text{Me}_3\text{GeGePh}_2(\eta^6\text{-Ph})\text{Cr(CO)}_3$ (3c) was isolated only in a trace amount; the major compound being the initial ligand 3. A possible reason would be the decomposition of the W complex under high-temperature conditions. Therefore, in order to perform the reaction at lower temperatures, 2a-c, $\text{M(CO)}_3(\text{NCMe})_3\text{M(CO)}_6$, in which three carbonyl groups have been substituted by more labile MeCN were used instead of 1a-c (M = Cr, Mo, W).^[30b, 35b, 39]

The compounds 2a-c are easy to prepare (Scheme 5) and can either be isolated in high yields or used without isolation.^[40]



Scheme 5. Synthesis of 2a-c.

As expected, using $\text{M(CO)}_3(\text{NCMe})_3$ as more reactive starting materials, allowed to prepare the desired complexes, with improved yields, by refluxing 2a-c, in THF for 1 h, in the

presence of aryl oligogermanes (Scheme 6). For these reactions, the conversion of the starting material proved to be much higher (54–68 %). Increasing the reaction time (up to 3 h) resulted only in a decrease of the yields, most probably, again, due to the thermal instability of Cr complexes. No conversion was observed when the reaction was carried out either in MeCN (under reflux for 12 h) or in THF at room temperature (for up to 3 days), indicating that acetonitrile is a strong donating ligand than the arenes (see below).

Using an excess of **2a** (up to 2 eq.) resulted in the sole formation of mono-Cr complexes (in the case of **7a**) without any traces of di-Cr complexes. In the case of Mo, the complex **3b** has shown to be unstable and to decompose quickly during the various attempts to isolate it. Mo derivatives are known for their low stability.^[41] When other derivatives, such as $M(\text{CO})_3(\text{Py})_3$ ^[35b] were used as starting materials, under the same conditions, in the presence of $\text{Ph}_3\text{GeGeMe}_3$ (**3**), only traces of the desired compounds were observed.

When comparing the three methods used for the synthesis of **3a-c** and **4a-7a**, namely thermolysis of $M(\text{CO})_6$ in decalin/dimethyl succinate, reflux of $M(\text{CO})_6$ in $(n\text{Bu})_2\text{O}/\text{THF}$ and reflux of $M(\text{CO})_3(\text{NMe})_3$ in THF. Among these three options, the latest one seems to be the most successful, leading to the formation of the desired complexes in higher yields.

All studied the complexes, **3a**, **3c**, **4a-7a**, are yellow powders, which are stable under an inert atmosphere, but get slowly oxidised (and become greenish) upon exposure to air for several months. In stark contrast, solutions of these compounds have shown to be much more air-sensitive. All of the newly synthesized derivatives are soluble in aromatic hydrocarbons (benzene, toluene), acetone, ethers (THF, diethyl ether) but are insoluble in *n*-hexane and decompose very quickly in chloroform (or CDCl_3). Among the possible decomposition pathways, the rupture of the Cr-Ar bond is the most probable, especially under harsh conditions. The rupture of the Ge-Ge bond was also observed when NMR was used to monitor the stability of the solutions of **6a** in C_6D_6 . After 3 days, the formation of **6**, $(\text{C}_6\text{F}_5)_3\text{GeGePh}_3$, and $(\text{C}_6\text{F}_5)_3\text{GeH}$ was observed.

All compounds were characterized by elemental analysis, IR and NMR spectroscopy, mass-spectrometry^[42] and in the case of **3c**, **4a** and **5a** by single crystal X-ray diffraction (XRD) analysis. The deviations in elemental analysis data are explained by the sensitivity of the compounds and by the partial combustion due to the presence of transition metals.

Solid state structure. A search in the CSD (Cambridge Structural Database) (June 2018) for structures of aryloligogermanes coordinated with $M(\text{CO})_3$ returned no result. Therefore, the structural investigation of **3c**, **4a** and **5a** by XRD reported in this work (Figures 1–3; Supporting Information, Table S1) represents a significant contribution to the chemistry of Group 14 derivatives. Moreover, only one structure of monogermanium compound, $\text{N}(\text{CH}_2\text{CH}_2\text{O})_3\text{Ge}(\eta^6\text{-Ph})\text{Cr}(\text{CO})_3$,^[30d] has ever been studied by XRD to date.

For all of the structures studied in this work, the metal-ligand core shown to adopt the well-known “piano stool” geometry.^[43]

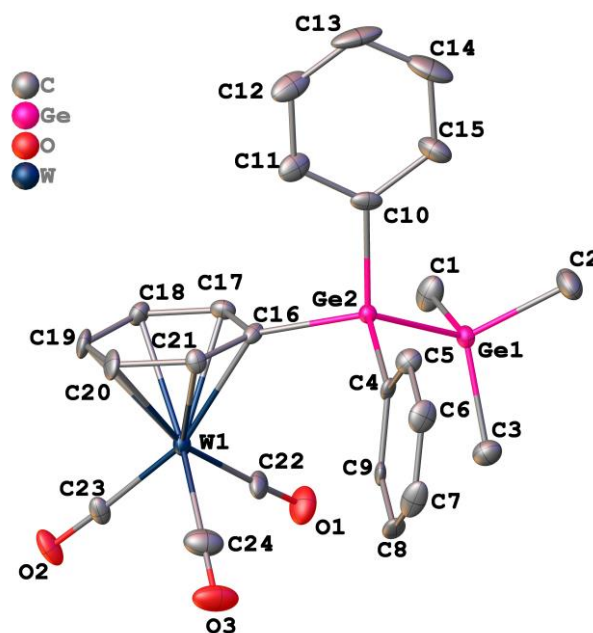
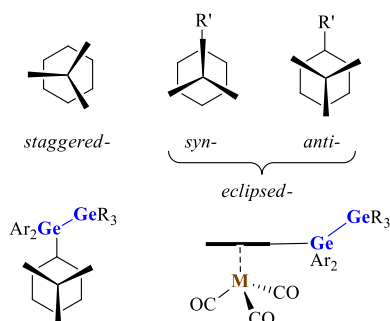


Figure 1. Molecular structure of $\text{Me}_3\text{GeGePh}_2(\eta^6\text{-Ph})\text{W}(\text{CO})_3$ (**3c**). Displacement ellipsoids are shown at 50% probability level. Hydrogen atoms are omitted for clarity. Selected bond lengths (Å) and bond angles (deg): Ge(1)–Ge(2) 2.4266(9), W–C_{av}(arene) 2.341(6), W–C_{av}(centroid) 1.867(6), W–CO_{av} 1.967(7), Ge(1)–C_{av} 1.949(6), Ge(2)–C_{av} 1.957(6), Ge(2)–C(16) 1.972(6), C–O_{av} 1.155(8); C–W–C_{av}(CO) 87.4 (3), C–Ge(1)–Ge(2)_{av} 110.8(2), C–Ge(1)–C_{av} 108.1(3), C–Ge(2)–Ge(1)_{av} 112.64(19), C–Ge(2)–C_{av} 106.2(2).

In **3c**, the $\text{W}(\text{CO})_3$ group in relation to the coordinated C_6H_5 ring adopts an *anti-eclipsed*-conformation (Scheme 6; torsion angles $\text{C}_{\text{Ge}}\text{-C}_{\text{centroid}}\text{-W-C}_\text{O}$ are 55.4(7) and 71.1(7) $^\circ$; 60 $^\circ$ for ideal *anti*- and 0 $^\circ$ for ideal *syn*-). Due to the electronic interactions, a *syn-eclipsed*-conformation is typical for monosubstituted bearing an electron donating group R' ($\eta^6\text{-R}'\text{C}_6\text{H}_5$) $M(\text{CO})_3$. An electron-donating R' increases the electron density in *ortho*- and *para*-positions of the $\eta^6\text{-Ar}$ ring, where the octahedral set of Cr orbitals are overlapping.^[44] Usually, an *anti-eclipsed* conformation is due to steric reasons, but in our case, additional orbital interactions between the *d* orbitals of the Cr and the σ -orbital of Ge-Ar and Ge-Ge bonds may be observed (Scheme 7). The maximal overlapping (torsion angle $\text{M-C}_{\text{ArGe}}\text{-Ge-Ge}$ 180 $^\circ$ is the most profitable) is observed in **5a** (see below); in **3c** a torsion angle of 55.2(7) $^\circ$ is found, indicating that they are electronic interactions between all parts of the molecule. The conformation along the Ge-Ge bond is staggered (torsion angles C–Ge–Ge–C_{av} are 78.27/41.71 $^\circ$).

In **4a**, the conformation of the $\text{Cr}(\text{CO})_3$ group in relation to the coordinated C_6H_5 ring is close to be *anti-eclipsed* (torsions $\text{C}_{\text{ArGe}}\text{-C}_{\text{centroid}}\text{-Cr-C}_\text{O}$ are 52.9(8) and 68.7(8) $^\circ$). The torsion angle of $\text{Cr-C}_{\text{ArGe}}\text{-Ge-Ge}$ is 46.5(8) $^\circ$, indicating weak interactions between Cr and Ge2. DFT calculations for **4a** (see below) showed that in the gas phase this torsion angle is 155 $^\circ$, indicating an effect of the crystal packing on the conformation. The conformation along the Ge-Ge bond is staggered (torsion angles C–Ge–Ge–C_{av} are 70.5(8)/49.5(8) $^\circ$).



Scheme 7. Conformations of substituents in relation to η^6 -benzene.

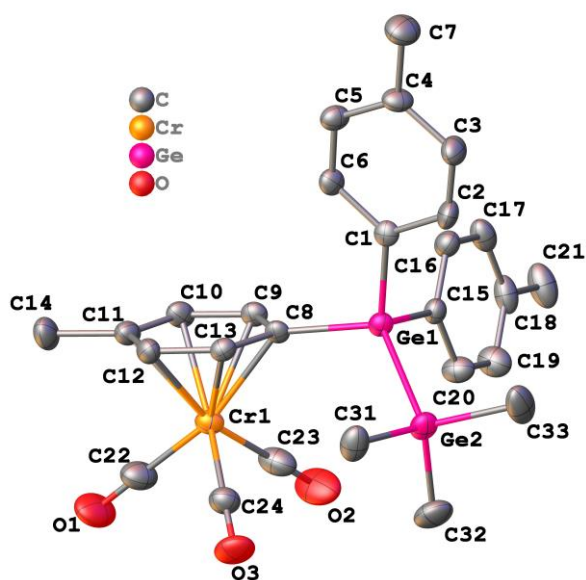


Figure 2. Molecular structure of compound $\text{Me}_3\text{GeGe}(p\text{Tol})_2(\eta^6\text{-}p\text{Tol})\text{Cr}(\text{CO})_3$ (**4a**). Displacement ellipsoids are shown at 50% probability level. Hydrogen atoms are omitted for clarity. Selected bond lengths (Å) and bond angles (deg): Ge(1)-Ge(2) 2.4284(6), Ge(1)-C_{av} 1.952(5), Ge(1)-C(8) 1.970(4), Ge(2)-C_{av} 1.932(7), Cr(1)-CO_{av} 1.828(9), Cr(1)-C_{av}(arene) 2.212(5), Cr-C_{av}(centroid) 1.715(5), C-O_{av} 1.163(8); C-Cr-C_{av}(CO) 88.2(3), C-Ge(1)-Ge(2)_{av} 112.66(13), C-Ge(1)-C_{av} 106.1(2), C-Ge(2)-Ge(1)_{av} 110.49(19), C-Ge(2)-C_{av} 108.5(3).

The structure of the complex **5a** differs from the ones of **3c** and **4a**. The conformation of the $\text{Cr}(\text{CO})_3$ group in relation to the coordinated C_6H_5 ring is distorted and *staggered* (torsion angles $\text{C}_{\text{ArGe}}\text{-C}_{\text{centroid}}\text{-Cr-C-O}$ are 27.6(9) and 95.6(9) $^\circ$), indicating electron interactions between $\text{Cr}(\text{CO})_3$ and Ge2. The torsion angles $\text{Cr-C}_{\text{ArGe}}\text{-Ge-Ge}$ is 159.7(9) $^\circ$, showing a very high interaction between Cr, η^6 -Ph and Ge2. This significant observed difference, in **5a**, may be explained by the presence of Ph groups at both Ge atoms, what is known to affect the electronic structure of the molecule strongly.^{[45], [46]} The conformation along the Ge-Ge

bond is *staggered* (torsion angles $\text{C-Ge-Ge-C}_{\text{av}}$ are 59.6(9)/60.4(9) $^\circ$).

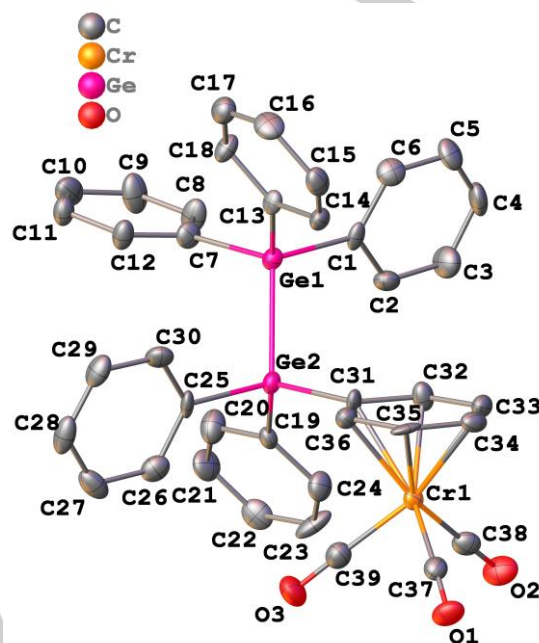


Figure 3. Molecular structure of compound $\text{Ph}_3\text{GeGePh}_2(\eta^6\text{-Ph})\text{Cr}(\text{CO})_3$ (**5a**). Displacement ellipsoids are shown at 50% probability level. Hydrogen atoms are omitted for clarity. Selected bond lengths (Å) and bond angles (deg): Ge(1)-Ge(2) 2.4316(13), Ge(1)-C_{av} 1.955(9), Ge(2)-C_{av} 1.953(10), Ge(2)-C(31) 1.974(9), Cr(1)-C_{av}(arene) 2.209(11), Cr-C_{av}(centroid) 1.698(11), Cr-CO_{av} 1.842(11), C-O_{av} 1.157(13); C-Cr-C_{av}(CO) 88.1(5), C-Ge(1)-Ge(2)_{av} 110.0(3), C-Ge(1)-C_{av} 108.9(4), C-Ge(2)-Ge(1)_{av} 109.0(3), C-Ge(2)-C_{av} 109.9(4).

In all of the structures studied, **3c**, **4a**, and **5a**, the η^6 -coordinated arene ring is almost planar (the maximal deviations from the mean square plane are 0.022(9) Å in **3c** (C(19)-*para* to Ge toward to W), 0.008(9) Å in **4a** (C(8)-*ipso* and C(9)-*ortho* to Ge out of Cr) and 0.012(9) Å in **5a** (C(35)-*meta* to Ge out of Cr).^[43] In every case, the nearest Ge atom is slightly moved out of the plane of the η^6 -Ar in the direction above $\text{M}(\text{CO})_3$ (0.228(9) Å for **3c**, 0.131(9) Å for **4a**, 0.101(9) Å for **5a**; the angles of deviation α are 6.4(7), 3.6(8), 3.0(9) $^\circ$, respectively) due to the electronic effects (Scheme 7), indicating a π -donor effect of the oligogermyl substituent. In **4a**, the methyl group of the *p*-tolyl ring is almost in the plane of aryl group (the deviation is only 0.001(9) Å).

The presence of the $\text{M}(\text{CO})_3$ increases the nonequivalence of the aryl C-C bonds (approaching a hexadienyl character) and leads to an elongation of C1-C2 bond (Scheme 8), whereas the electron density is partially transferred from the Ge-Ge bond (HOMO-3, HOMO-4, HOMO-5; Figs. S26-S28, ESI) to the η^6 -arene ring (LUMO+3, LUMO+4, LUMO+5; Figs. S32-S34, ESI). This was observed for each of the new complexes studied (Scheme 9), indicating that the presence of a $\text{M}(\text{CO})_3$ moiety

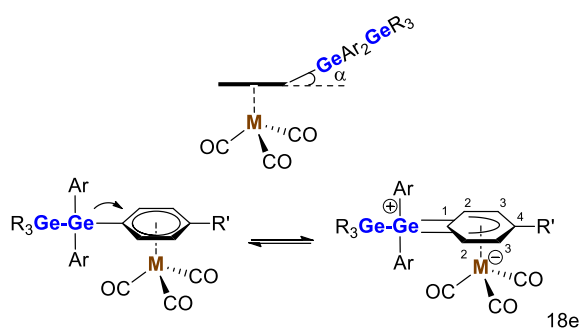
lowers the bond order of the aromatic ring due to its acceptor properties.

Furthermore, a deviation from the ideal sp^2 hybridization for the arene carbon atoms bonded to Ge is observed.

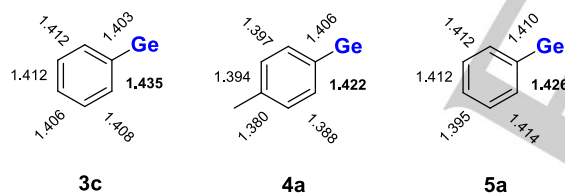
Table 1. Comparison structural parameters of complexes **3c**, **4a**, **5a** with parent oligogermanes **3**, **4** and **5**.

Compound [a]	$d(\text{Ge-Ge})$	$d(\text{Ge-C})$	$d(\text{Ge}_M\text{-C})$	C-Ge-Ge _M	C-Ge-C	C-Ge _M -Ge	C-Ge _M -C
Me ₃ GeGePh ₃ (3) ^[47]	2.418(1)	1.943(4)	1.957(2)	110.2(1)	108.7(1)	110.3(1)	108.6(1)
3c	2.4266(9)	1.949(6)	1.972(6)	110.8(2)	108.1(3)	112.64(19)	106.2(2)
Me ₃ GeGe(<i>p</i> Tol) ₃ (4) ^[48]	2.4292(7)	1.936(7)	1.948(4)	115.6(2)	107.2(4)	110.90(12)	107.97(17)
4a	2.4284(6)	1.932(7)	1.970(4)	110.49(19)	108.5(3)	112.66(13)	106.1(2)
Ph ₃ GeGePh ₃ (5) ^[49]	2.437	1.959	1.959	110.80	108.10	110.80	108.10
5a	2.4316(13)	1.955(9)	1.974(9)	110.0(3)	108.9(4)	109.0(3)	109.9(4)

[a] Bond lengths (Å) and bond angles (deg); Ge_M is a nearest to M(CO)₃ germanium atom; average values are used



Scheme 8. Donor effects of oligogermeryl substituent.



Scheme 9. Values of $d(\text{C-C})$ bond lengths in η^6 -arene for **3c**, **4a** and **5a**.

The Ge-Ge bond is *cisoid*- in relation to the $\text{M}(\text{CO})_3$ group in **3c** (the value of the torsion angle for $\text{Ge-Ge-C}(16)_{\text{Ar}}\text{-W}$ is $55.2(7)^\circ$) and **4a** (torsion $\text{Ge-Ge-C}(8)_{\text{Ar}}\text{-Cr}$ is $46.5(8)^\circ$), whereas in **5a** this same bond is *transoid*- (torsion $\text{Ge-Ge-C}(31)_{\text{Ar}}\text{-Cr}$ is $159.6(9)^\circ$). These data, along with the conformation of $\text{M}(\text{CO})_3$ along the Ar group, indicate that there is a stronger electron interaction between the octahedrally directed metal's orbitals and the η^6 -arene in **5a**. The values of the torsion angles for $\text{Ge-Ge-C}_{\text{Ar}}\text{-C}_{\text{Ar}}$ and $\text{Ge-Ge-C}_{\text{Ar}}\text{-M}$ ($-39.2(7)/55.2(7)^\circ$ for **3c**; $46.0(8)/46.5(8)^\circ$ for **4a**; $-67.1(9)/-159.6(9)^\circ$ for **5a**) indicate the presence of a σ - π -conjugation between the Ge-Ge bond and the η^6 -Ar fragments (90° for an ideal conjugation).

In all of the structures **3c**, **4a** and **5a** the values of $d(\text{C-O})$ are very similar (1.155(8) vs. 1.163(8) vs. 1.157(13) Å), showing acceptor properties across the different oligogermanes and metals. The values of $d(\text{Cr-C}_O)$ in **4a** and **5a** are almost identical

(1.828(9) vs. 1.842(11) Å). Variation of the substituents on the Ge atom or of the nature of M resulted only in small changes of the main structural parameters, i.e. Ge-Ge bond length (2.4266(9) vs. 2.4284(6) vs. 2.4316(13) Å in **3c**, **4a**, **5a**, respectively) and Ge-C bond length (for example, Ge-C_{ArM} 1.972(6) vs. 1.970(6) vs. 1.974(9) Å). The length of $d(\text{Ge-C}_{\text{ArM}})$ is longer than the other Ge-C bonds; this could be due to steric reasons (introduction of a more voluminous $\text{M}(\text{CO})_3$) and electronic effects similar to what has been earlier observed for donor-acceptor oligogermanes.^[46, 50] In general, the geometry at each of Ge atom may be described as a slightly distorted tetrahedral.

A comparison between the structural parameters of complexes **3c**, **4a**, **5a** and the ones of the parent oligogermanes **3**,^[47] **4**^[48] and **5**,^[49] respectively (Table 1), shows minimal differences, indicating a strong mutual influence of the Ge₂, aryl and $\text{M}(\text{CO})_3$ fragments in the molecule, where the acceptor properties of $\text{M}(\text{CO})_3$ fragments are counterbalanced by a Ge₂ donation.

Structure in solution. The NMR spectra of complexes **3a-3c** and **4a-7a** were recorded in C_6D_6 or acetone- d_6 . Due to the weaker π -acceptor properties of the η^6 -coordinated arene ligand in comparison with CO, a quenching of the ring current is observed in such compounds. The electron density is transferred from arene to the $\text{M}(\text{CO})_3$. This phenomenon is reflected in the NMR spectra by upfield shifts of the coordinated aryl group signals in comparison with the parent oligogermanes **3-7**. In ^1H NMR (C_6D_6) the signals of the η^6 -coordinated Ar appeared between δ 4.0-5.3 ppm (an upfield shift of ca. 2 ppm), and the signals of the uncoordinated Ar were shifted toward downfield by ca. 0.2 ppm in comparison with the free oligogermane (for details, see Experimental Part and Supporting Information, Figs. S1-S16). A comparison with the parent ($\eta^6\text{-C}_6\text{H}_6$) $\text{Cr}(\text{CO})_3$ (vs. δ 4.31 ppm)^[51] showed that for oligogermanes **3a-7a** the proton signals are shifted toward lower fields. Furthermore, the signals for each position of the coordinated Ar group (*ortho*-, *para*- and *meta*-) appeared as well resolved multiplets (doublets, triplets *etc.*, $^3J_{\text{H-H}} = 6.6$ Hz, $^4J_{\text{H-H}} = 1.0$ Hz). The weakest upfield shifts were observed for Mo and the largest ones for W (**3b** vs. **3a** vs. **3c**).

The ^{13}C NMR spectra of Cr compounds have shown a single signal for CO (δ 232.1-234.1 ppm), indicating the rapid (on the

NMR time scale) rotation along the M-Ar_{centroid} axis resulting in the equivalence of the 3 carbonyls. The ¹³C NMR spectra are characterized by the presence of 4 signals typical for

coordinated carbon arene at δ 110.6-90.8 ppm (upfield shift of ca. 35 ppm in comparison with non-coordinated Ar); *ipso*,

Table 2. Data of the UV/vis spectra according to experiment and DFT Calculations for the compounds **4**, **4'**, **4''**, **4'''** and **4a**.

compound	λ_{exp} , nm ^[a]	λ_{calc} , nm ^[b]	transition
Me ₃ GeGe(<i>p</i> Tol) ₃ (4) ^[c]	232 (2.3)	229 (0.282)	HOMO→LUMO+2
(η^6 - <i>p</i> Tol)Cr(CO) ₃ (4')	220 (1.74) 253 (0.76)	-	-
		286 (0.2228) 321 (0.0017)	HOMO→LUMO+1, HOMO-1→LUMO
	317 (0.95) ^[d]	327 (0.0030) 343 (0.0010)	HOMO-2→LUMO+1 HOMO-2→LUMO HOMO-1→LUMO+1
(η^6 -Me ₃ Ger)Cr(CO) ₃ (4'')	-	288 (0.2091) 322 (0.0022) 327 (0.0019)	HOMO→LUMO+1, HOMO-1→LUMO HOMO-2→LUMO+1 HOMO→LUMO+1, HOMO-2→LUMO
MeGe(<i>p</i> Tol) ₂ (η^6 - <i>p</i> Tol)Cr(CO) ₃ (4''')	-	295 (0.1044) 318 (0.0037) 323 (0.0032) 333 (0.0028) 337 (0.0027) 350 (0.0025) 382 (0.0025) 391 (0.0071)	HOMO-1→LUMO+1 HOMO-2→LUMO+2 HOMO-1→LUMO+2 HOMO-1→LUMO+1 HOMO-2→LUMO+1 HOMO→LUMO+2 HOMO-1→LUMO HOMO→LUMO
Me ₃ GeGe(<i>p</i> Tol) ₂ (η^6 - <i>p</i> Tol)Cr(CO) ₃ (4a)	233 (4.55) 267 (0.80) 321 (0.87)	- 279 (0.0132) 298 (0.1452) 326 (0.0029) 328 (0.0156) 336 (0.0092) 357 (0.0016) 383 (0.0024) 394 (0.0107)	- HOMO-3→LUMO HOMO-1→LUMO+1 HOMO-2→LUMO+1 HOMO-1→LUMO+2 HOMO-2→LUMO+1 HOMO→LUMO+1 HOMO-1→LUMO HOMO→LUMO

[a] Absorptivity ϵ (10^4 M⁻¹ cm⁻¹) in parentheses; data in CH₂Cl₂. [b] Oscillator strength in parentheses. [c] Data from ^[48]. [d] Data from ^[52].

ortho-meta- and *para*-carbon signals are easily distinguished. In general these signals are shifted toward lower fields in comparison with the parent C₆H₆Cr(CO)₃ (vs. δ 92.36 ppm).^[51]

DFT calculations. Theoretical investigation has been performed for the model compounds Me₃GeGe(*p*Tol)₂(η^6 -*p*Tol)Cr(CO)₃ (**4a**), (η^6 -TolH)Cr(CO)₃ (**4'**), Me₃Ge(η^6 -*p*Tol)Cr(CO)₃ (**4''**), MeGe(*p*Tol)₂(η^6 -*p*Tol)Cr(CO)₃ (**4'''**) and Me₃GeGe(*p*Tol)₃ (**4**) (Supporting Information, Figs. S17-S62, Tables S2-S4); the results obtained are typical for all series of compounds **3a-7a**.

According to the calculations using B3LYP/6-311+G(2d,p) in THF, **4a** has three bands, active in IR for the carbonyl groups, 1960 and 1884/1855 cm⁻¹. This correlates well with the experimental results (1961, 1887 cm⁻¹, see below), confirming the validity of the model used. In the gas phase, using 6-31G(d), a different set of values (2107, 2056/2052 cm⁻¹) was obtained, showing the high dependence of these values on the solvent used. These calculations have also shown that the presence of a Ge2 fragment increases the donor properties of the aryl group (vs. 2117, 2066/2060 cm⁻¹ for (η^6 -TolH)Cr(CO)₃ (**4'**)).

For **4a**, due to the presence of the metal, an electron acceptor, the HOMO is mainly localised on the Cr(CO)₃ fragment (bonding Cr-CO), whereas the HOMO-1, HOMO-2 are mainly localised on the TolCr(CO)₃. The lower energy HOMO-3 orbital is bonding and localised on the Ge-Ge fragment; the other bonding orbitals, HOMO-4, HOMO-5, HOMO-6, HOMO-7 are all located on the non-coordinated Tol rings. Furthermore, the LUMO as well as the LUMO+1, are π -antibonding orbitals are localised on the non-coordinated Tol rings. The LUMO+2 is characterised by a contribution to the antibonding orbital on TolCr(CO)₃. The higher energy orbital LUMO+3 is mainly

localised on the (Tol)Cr(CO)₃ fragment. All of this indicates a full conjugation between the different parts of the molecule and a distribution of the electron density over the whole molecule. This situation differs significantly from what is observed for the parent aryloligogermanes, where the HOMO is mainly localised on the Ge-Ge bond and the LUMO on Ar substituents,^{[45], [46]} as it was confirmed for **4**. It means that the electron density is transferred from Ge2 to ArCr(CO)₃ through an intramolecular conjugation.

Calculations of the UV/vis spectra were performed for **4**, **4'**, **4''**, **4'''** and **4a** (Table 2).

The UV/vis spectra of (η^6 -C₆H₆)Cr(CO)₃^[25c, 52-53] exhibits 3 bands, corresponding respectively to a π - π^* transition (~ 220 nm with ϵ ~1-3x10⁴ M⁻¹ cm⁻¹), a M→ArH charge transfer (~250 nm with ϵ <10⁴ M⁻¹ cm⁻¹) and a M→ π^* CO charge transfer (~ 315 nm with ϵ ~10⁴ M⁻¹ cm⁻¹).^[38a, 53c]

The most hypsochromic bands of the UV/vis absorption spectrum (220 nm for **4'**; 233 nm for **4a**) correspond to an intraligand charge transfers (ILCT) for Ar, where the transition is occurring from a low HOMO (such as HOMO-4 and lower) to a higher LUMO (such as LUMO+5 and higher), which are located on the Ar groups. Due to DFT restrictions, these transitions have not been taken into account during the calculations.

The data obtained for **4a** show an orbital mixing and all bands are significantly bathochromically shifted in comparison with the model compounds. The following bands attributions could be made: 267 nm (metal-to-arene charge transfer, MACT, from Ge2 to Ar, and MACT from TolCr(CO)₃ to the antibonding orbital of Tol; in general from a lower HOMO to LUMO and near) and 321 nm (metal-to-carbonyl charge transfer, MCCT,^[54] from TolCr(CO)₃ to the antibonding orbital of Cr(CO)₃; in general from a lower HOMO to higher LUMO orbitals).

UV/vis spectroscopy. UV/vis spectroscopy is an important method for the characterization of catenated Group 14 compounds and for the estimation of their HOMO-LUMO transitions gap. For the compounds **3a-7a**, one might expect a shift on the bands compared to the parent compounds. The introduction of aromatic groups within an oligogermane framework is known to lead to a bathochromic shift of the UV/vis

absorbance bands due to the σ - π -conjugation. The effect of the insertion of transitional metal fragments remains less evident to predict.

The σ - σ^* transition in the Ge-Ge bonds is usually observed around 230-250 nm for di- and trigermanes with a very high absorptivity ($\epsilon \sim 2\text{-}5 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1}$).

Table 3. UV/vis spectroscopy data for **3a-7a** and related compounds.

compound [a]	λ (nm), $\sigma \rightarrow \sigma^*$ [b]	λ (nm), ILCT [b]	λ (nm), MACT [b]	λ (nm), MCCT [b]	reference
(η^6 -EtO ₂ CC ₆ H ₅)Cr(CO) ₃		223 (2.23)	255 (0.82)	323 (1.00)	[52]
(η^6 -C ₆ H ₆)Cr(CO) ₃		213 (3.00)	256 (0.75)	317 (1.06)	[52]
(η^6 -MeC ₆ H ₅)Cr(CO) ₃		220 (1.74)	253 (0.76)	317 (0.95)	[52]
(η^6 -Et ₃ SiC ₆ H ₅)Cr(CO) ₃		212 (2.17)	256 (0.71)	317 (1.03)	[52]
Me ₃ SiSiMe ₂ (η^6 -C ₆ H ₅)Cr(CO) ₃		231 (1.9)	256	318 (1.1)	[25c]
Me ₃ SiSiMe ₂ (η^6 -C ₆ H ₅)Mo(CO) ₃		232 (1.9)	286 (4.1)	326 (2.1)	[25c]
Me ₃ SiSiMe ₂ (η^6 -C ₆ H ₅)W(CO) ₃		232 (2.1)	278 (5.1)	321 (1.9)	[25c]
PhSiMe ₂ SiMe ₂ (η^6 -C ₆ H ₅)Mo(CO) ₃ [c]		213 (2.87)		327 (1.56)	[53d]
Me ₃ GeGePh ₃ (3)	230				[55]
Me ₃ GeGePh ₂ (η^6 -C ₆ H ₅)Cr(CO) ₃ (3a)		232 (2.47)	263 (0.53)	320 (0.67)	this work
Me ₃ GeGePh ₂ (η^6 -C ₆ H ₅)W(CO) ₃ (3c)		233 (1.96)	291 (0.38)	322 (0.46)	this work
Me ₃ GeGe(<i>p</i> Tol) ₃ (4)	232 (2.3)				[48]
Me ₃ GeGePh ₂ (η^6 -C ₆ H ₅)Cr(CO) ₃ (4a)		233 (4.55)	267 (0.80)	321 (0.87)	this work
Ph ₃ GeGePh ₃ (5)	241				[56]
Ph ₃ GeGePh ₂ (η^6 -C ₆ H ₅)Cr(CO) ₃ (5a)		238 (3.57)	267 (1.64) (sh)	322 (0.64)	this work
(C ₆ F ₅) ₃ GeGePh ₃ (6)	226 (3.70)				[46]
(C ₆ F ₅) ₃ GeGePh ₂ (η^6 -C ₆ H ₅)Cr(CO) ₃ (6a)		234 (2.15)	261 (0.78)	322 (0.41)	this work
Ph ₃ GeGeMe ₂ GePh ₃ (7) [d]	245 (3.02)				[57]
Ph ₃ GeGeMe ₂ GePh ₂ (η^6 -C ₆ H ₅)Cr(CO) ₃ (7a)		244 (3.60)	261 (2.10) (sh)	321 (0.38)	this work

[a] Performed in solution in CH₂Cl₂ unless otherwise stated. [b] ϵ ($10^4 \text{ M}^{-1} \text{ cm}^{-1}$) in parentheses. [c] Solution in THF. [d] Solution in cyclohexane.

Table 4. IR spectroscopy data for arene chromium tricarbonyl complexes.

Compound	ν_{CO} , cm ⁻¹	solvent	reference	Compound	ν_{CO} , cm ⁻¹	solvent	reference
Cr(CO) ₆	1987	hydrocarbons	[58]	Cr(CO) ₃ (η^6 -C ₆ H ₅ Cl)	1991, 1929, 1925	cyclohexane	[34]
<i>fac</i> -Cr(CO) ₃ (PF ₃) ₃	2062, 1998	not given	[59]	Cr(CO) ₃ (η^6 -C ₆ H ₅ NMe ₂)	1969, 1894, 1888	cyclohexane	[34]
<i>fac</i> -Cr(CO) ₃ [(MeO) ₃ P] ₃	1962, 1875, 1966, 1888/1879	CS ₂ , hexadecane	[60], [61]	Cr(CO) ₃ (η^6 -C ₆ H ₅ COOMe)	1990, 1927	cyclohexane	[34]
<i>fac</i> -Cr(CO) ₃ [Me ₃ P] ₃	1939, 1849	not given	[62]	Cr(CO) ₃ (η^6 -C ₆ H ₅ COOMe)	1992, 1929	heptane	[54]
<i>fac</i> -Cr(CO) ₃ [NH ₃] ₃	1935, 1842, 1911, 1887	hexadecane, MeCN	[61], [63]	Cr(CO) ₃ (η^6 -PhSiMe ₃)	1987, 1911	cyclohexane	[30a]
<i>fac</i> -Cr(CO) ₃ [NCMe] ₃	1910, 1782	nujol	[40b]	Cr(CO) ₃ (η^6 -PhGeMe ₃)	1985, 1910	cyclohexane	[30a]
<i>fac</i> -Cr(CO) ₃ [Py] ₃	1910, 1778	not given	[64]	Cr(CO) ₃ (η^6 -PhGeMe ₂ Ph)	1980, 1915/1870	nujol	[30c]
Cr(CO) ₃ (η^6 -C ₆ H ₅ CH ₃)	1983, 1914	cyclohexane	[53b]	Cr(CO) ₃ (η^6 -PhSnMe ₃)	1984, 1908	cyclohexane	[30a]
	1962, 1893	nujol	[65]	Cr(CO) ₃ (η^6 -PhSiMe ₂ SiMe ₃)	1977, 1911	isooctane	[25c]
Cr(CO) ₃ (η^6 -C ₆ H ₅ OCH ₃)	1980, 1908	cyclohexane	[34]	Cr(CO) ₃ (η^6 -C ₆ H ₅ CH ₃)	1972, 1906	hexane	[25d]
	1975/1945, 1903/1857	nujol	[66]	PhSiMe ₂ SiMe ₂ SiMe ₂ C≡CPh			
Cr(CO) ₃ (η^6 -C ₆ H ₆)	1982, 1915	cyclohexane	[34]	Cr(CO) ₃ (η^6 -C ₆ H ₅)	1975, 1909	hexane	[25d]
	1978, 1910	nujol	[67]	PhSiMe ₂ SiMe ₂ SiMe ₂ C≡CPh			
Cr(CO) ₃ (η^6 -C ₆ H ₅ F)	1990, 1929, 1926	cyclohexane	[34]	Cr(CO) ₃ (η^6 -PhSiMe ₂ SiMe ₃)	1974, 1897	nujol	this work
<i>fac</i> -Mo(CO) ₃ (NCMe) ₃	1915, 1783	nujol	[40b], [68]	Cr(CO) ₃ (η^6 -C ₆ H ₅ SiMe ₃)	1970, 1894	nujol	this work
<i>fac</i> -Mo(CO) ₃ (PMe ₃) ₃	1927, 1828	not given	[69]	Cr(CO) ₃ (η^6 -PhGePh ₂ GeMe ₃)	1975, 1900	nujol	this work
Mo(CO) ₃ (η^6 -C ₆ H ₆)	1987, 1916	heptane	[70]	Cr(CO) ₃ (η^6 -PhGePh ₂ GePh ₃)	1975, 1900	nujol	this work
	1985, 1912	nujol	[30e]	Cr(CO) ₃ (η^6 -PhGePh ₂ GePh ₃)	1975, 1900	nujol	this work
Mo(CO) ₃ (η^6 -C ₆ H ₅ Me)	1980, 1910	nujol	[30e]	Cr(CO) ₃ (η^6 -PhGePh ₂ Ge(C ₆ F ₅) ₃)	1976, 1901	nujol	this work
Mo(CO) ₃ (η^6 -C ₆ H ₅ SiMe ₃)	1986/1944, 1916/1874	nujol	[30e]	Cr(CO) ₃ (η^6 -PhGePh ₂ Ge(C ₆ F ₅) ₃)	1972, 1896	nujol	this work
<i>fac</i> -W(CO) ₃ (NCMe) ₃	1885, 1778	nujol	[40b]	PhGePh ₂ GeMe ₂ GePh ₃ (7a)			
<i>fac</i> -W(CO) ₃ [(MeO) ₃ P] ₃	1973, 1894/1880	hexadecane	[61]	Mo(CO) ₃ (η^6 -C ₆ H ₅ SiMe ₃)	1983, 1913	petroleum ether	[53d]
<i>fac</i> -W(CO) ₃ [Me ₃ P] ₃	1940, 1845	hexadecane	[61]	Mo(CO) ₃ (η^6 -PhSiMe ₂ SiMe ₃)	1981, 1912	isooctane	[25c]
				Mo(CO) ₃ (η^6 -C ₆ H ₅ SiMe ₃)	1953, 1871	KBr	[53d]
				Mo(CO) ₃ (η^6 -C ₆ H ₅ GeMe ₃)	1986/1958, 1909/1867	nujol	[30e]
				Mo(CO) ₃ (η^6 -C ₆ H ₅ SnMe ₃)	1979/1951, 1909/1862	nujol	[30e]
				Mo(CO) ₃ (η^6 -C ₆ H ₅ SiMe ₃)	1978, 1900	nujol	this work
				W(CO) ₃ (η^6 -C ₆ H ₆)	1968, 1886	CH ₂ Cl ₂	[71]
				W(CO) ₃ (η^6 -MeC ₆ H ₅)	1985, 1910	heptane	[72]
				W(CO) ₃ (η^6 -PhSiMe ₂ SiMe ₃)	1980, 1909	isooctane	[25c]

W(CO) ₃ (η ⁶ -PhGePh ₂ GeMe ₃) (3c)	1980, 1904	njol	this work
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Due to solubility reasons, all UV/vis spectra in this work were recorded in CH₂Cl₂ which allows getting data at λ > 230 nm. For oligogermane **3a-7a**, three major bands were observed. Other bands were present but were either weak or overlapped with the main absorptions or with the solvent.

The comparison of UV/vis spectra of the compounds obtained within this study with the spectra of the initial oligogermanes shows that the introduction of a Cr(CO)₃ group has a strong influence on the absorption (Table 3; DFT section above). The bands corresponding to the σ → σ* transition in the parent oligogermanes did not appear due to a mixing of the MO. The two bands corresponding to an ILCT and a MACT in the parent chromium complexes are significantly bathochromically shifted in **3a-7a**; at the same time, the bands located at λ 320-322 nm (MCCT) are weakly redshifted. This indicates an increase of the donor properties of the η⁶-Ar ligand (HOMO increase) due to the presence of a Ge₂ group. The weak bands at 260-270 nm (MACT) appeared as a shoulder for Cr and as a distinct peak for the heavier Mo^[25c] and W analogues.^[73] This indicates an orbital mixing of the Ge-Ge σ-bond, π-aryl and π-CO, where Ge-Ge acts as a strong electronic donor. This band may be attributed to a transfer of electronic density from Ge-Ge to Ar.

For the complexes studied, the nature of the metal M (compare **3a** and **3c**) has shown to have little effect on the UV properties; only the MACT is significantly bathochromically shifted when replacing Cr by Mo or W.^[74]

In contrast with the free molecular oligogermanes^{[75], [48]} all complexes showed a total absence of luminescence both in solution and in solid form. The presence of transition metal fragments, M(CO)₃ (M = Cr, Mo, W), within the complexes, results in a quenching of the emissive properties. The electronic conjugation between Ge₂, η⁶-arene and M(CO)₃ occurs through direct interaction between a filled Cr d-orbitals and a σ* of Ge₂ bond, metal-ring σ-π-delocalization (π-conjugation) and a back-donation from Cr d-orbitals to a π*-orbitals of the arene and a donation from Ar to Cr (due to the withdrawing effect of Cr(CO)₃). This resulted in a bathochromic shift of the metal absorption bands (at 220 nm) with a small increase in absorbance.

IR spectroscopy and donor properties. For each of the compounds studied, the IR spectrum contains the expected 2 bands in the carbonyl region: a non-degenerate symmetric A₁ (at higher wave numbers, ν_{CO} ~2000-1970 cm⁻¹) and a doubly-degenerated asymmetric E (ν_{CO} ~1950-1900 cm⁻¹) vibration (local C_{3v} symmetry)^[76] (Table 4). Experimentally, there is no possible splitting of the E bands, which may be observed due to a small perturbation of the general symmetry of the arene substituent^[77] (compared with DFT calculations, see above). The analysis of the IR data (mainly, A₁ band) could be used to assess the donor (a decrease of the IR value) or the acceptor (an increase of the IR value) properties of a ligand. A strong dependence of the IR bands on the solvent has also been observed. Broadened red shifted bands are observed in polar solvents due to Lewis acidic interaction between the solvent and the carbonyl ligands.^[51]

Comparing the IR data for the compounds **3a-c**, **4a-7a** with the parent compound (η⁶-C₆H₆)Cr(CO)₃ shows a decrease of the stretching absorption due to the electron donating properties of the catenated Group 14 fragment; its insertion within the Ar framework results in a small increase in the σ-donating properties and a decrease of the π-acceptor ability. Using data from the literature and the ones obtained through this work (Table 4) helps to place the aryloligogermanes among ligands that increase the π-back acceptor properties: [PY]₃ < [MeCN]₃ < [NH₃]₃ < [PMe₃]₃ < [P(OMe)₃]₃ < **ArGeAr₂GeR₃** < [CO]₃ < [PF₃]₃. The catenated germanium fragment may be considered as a donor group: NMe₂ < **Ar₃GeGeR₂GeR₃** < Ar₃GeGeR₃ ≤ Ar(SiR₂)₂SiR'₃ < OMe < ArSiR₂SiR'₃ < Ar₂GeMe₂ < Me < H < PhSnMe₃ < PhGeMe₃ < PhSiMe₃ < CO₂Me < F < Cl. Also, the Ge atoms have shown to exhibit greater electron donating abilities than Si. Increasing the length of the Ge chain increased the donor properties of the aryloligogermane. This suggests that Ge₂ fragments have σ- as well as π-donating properties.

In comparison, it is known that the introduction of disilane fragment decreases the donor properties of the Ar ligand (Me₃Si(η⁶-Ph)Mo(CO)₃ and PhSiMe₂SiMe₂(η⁶-Ph)Mo(CO)₃ in KBr: 1946, 1865 vs. 1953, 1871 cm⁻¹).^[53d]

The comparison of the IR data between **3a-7a** (a weak increase of the donor properties in **6a** < **5a** < **3a** < **7a** < **4a**) shows a similar trend as the one observed in other spectroscopies. A similar trend could be observed when comparing the NMR chemical shifts of the ¹³CO groups, where increased donor properties correspond to a deshielding.^[78] Increasing the donor properties in η⁶-coordinated Ar groups results to a decrease of the acceptor properties (**4a** in Tol vs. Ph in **3a**); the same effect is observed when increasing the Ge chain length (Ge₃ in **7a** vs. Ge₂ **3a**) and during the metal transition W < Mo < Cr. Varying the nature of the substituent on the Ge atom, neighbouring the η⁶-coordinated Ar group also influence on the donor properties, but this effect remains relatively weak; therefore, the introduction of an electron withdrawing groups resulted only in a small decrease of the donor properties (**3a** vs. **5a**; **5a** vs. **6a**). Changing the nature of substituent on the central Ge atom has a more significant effect (when comparing PhGeMe₃ with Ph₂GeMe₂, Table 3). Therefore, in order to maximise the donor properties of the complex, one has to use long chained oligogermane, containing aryl groups bearing electron donating groups.

Electrochemistry. In this work, compound **4a** was studied as a typical example of the new complexes (Figure 4).

Compound **4a** displays a reversible one-electron oxidation of the organometallic centre Cr(0) → Cr(+1)^[44, 79] that leads to the formation of a stable cation-radical (on the voltammetric experiment timescale);^[80] the presence of the a Ge-Ge donor group offer an additional stabilization to the cationic oxidized product;^[81] no further oxidation has been observed. This differs from the parent oligogermanes, where an irreversible Ge-Ge oxidation is observed usually. The oxidation of the chromium center into a cationic specie is pushing any possible further oxidation towards higher potentials, making a second oxidation

of the compound much more difficult. The oxidation of **4a** has the shape of a typical slow electron transfer and occurs at $E_{1/2} = 0.32$ V vs. Fc/Fc⁺. The oxidation of the parent compound (η^6 -MeC₆H₅)Cr(CO)₃ has been reported to occur at $E_{1/2} = 0.38$ V vs. Fc/Fc⁺,^[53b, 78-79] indicating that the presence of the Ge₂ fragment in conjugation with (η^6 -MeC₆H₅)Cr(CO)₃ framework resulted in a slight HOMO destabilization.^[82] This decrease of the oxidation potential differs only slightly from what was observed for monogermanium fragments ($E_{1/2}$ 0.827 V vs. Ag/AgCl (0.377 V vs. Fc/Fc⁺) for Me₂Ge[(η^6 -Ph)Cr(CO)₃].^[44] Based on the electrochemical data, the effect of a Ge₂ fragment is comparable to the introduction of four methyl groups on the η^6 -benzene ring (for (η^6 -Me₄C₆H₂-1,2,4,5)Cr(CO)₃ $E_{1/2} = 0.33$ V vs. Fc/Fc⁺). Furthermore, these data indicate that the HOMO orbitals of Cr(CO)₃ are located on of the whole molecule. The parent digermene **4** is oxidized, irreversibly, at a much higher potential E_{ox} 1.65 V vs. Ag/AgCl^[48] (1.20 V vs. Fc/Fc⁺).

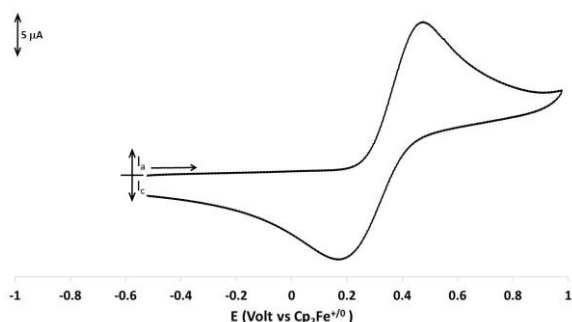


Figure 4. Electrochemical data for Me₃GeGe(pTol)₂(η^6 -pTol)Cr(CO)₃ (**4a**) 1.00 mM in [TBA][PF₆] in CH₂Cl₂. Sweeping rate: 250 mV/s.

Conclusions

Several synthetic routes for the synthesis **3a-3c**, **4a-7a**, R₃Ge-GeAr₂(R'⁶C₆H₄- η^6)M(CO)₃, were developed and studied. Among them, the reaction between aryloligogermene and M(CO)₃(NCMe)₃ was found to be the most efficient one. The analysis of the experimental and literature data for (η^6 -arene)M(CO)₃ complexes, including derivatives of catenated Group 14 compounds, has shown that electronic interactions (π -, d - and σ -interactions) between all parts of the molecule (Ge-Ge, arene, M(CO)₃) are observed and affect the physical properties of the whole molecule even though it remains weak. Oligogermanyl substituents on the aromatic ring have shown to behave similarly to usual electron donating substituents. Therefore complexes such as R₃Ge-GeAr₂(R'⁶C₆H₄- η^6)M(CO)₃ are typical donor-acceptor compounds. Introduction of oligogermanyl groups bathochromically shifted UV/vis absorption bands, due to a significant mixing of the atomic orbitals. The electrochemical oxidation of (η^6 -ArGe₂R₅)M(CO)₃ is reversible since to the oxidation is happening on the metallic centre M rather than on the Ge-Ge bond.

Experimental Section

Experimental Details. All manipulations were performed under a dry, oxygen-free argon atmosphere using standard Schlenk techniques. ¹H NMR (400.130 MHz), ¹³C NMR (100.613 MHz) and ¹⁹F (376.498 MHz) spectra were recorded with a Bruker 400 or Agilent 400MR spectrometers at 298 K. Chemical shifts are given in ppm relative to internal Me₄Si (¹H and ¹³C NMR spectra) or external CFCI₃ (¹⁹F NMR spectra). Mass spectra (EI-MS, 70 eV) were recorded on a quadrupole mass spectrometer FINNIGAN MAT INCOS 50 with direct insertion; all assignments were made with reference to the most abundant isotopes. High-resolution mass spectra (HRMS) were measured on a Bruker micrOTOF II instrument using electrospray ionization (ESI). Elemental analyses were carried out in the Microanalytical Laboratory of the Chemistry Department of the Moscow State University using HeraeusVarioElementar instrument. UV/visible spectra were obtained using two ray spectrophotometer Evolution 300 «Thermo Scientific» with cuvette of 1.00 cm long. The IR spectra were recorded by using a 200 Thermo Nicolet apparatus. Flash chromatography was performed using SiO₂ (0.015-0.040 mm).

Solvents were dried by standard methods and distilled prior to use. Tetrahydrofuran, diethyl ether were stored under solid KOH and then distilled over sodium/benzophenone. Toluene and *n*-hexane were refluxed and distilled over sodium. Dichloromethane and acetonitrile were distilled over CaH₂. C₆D₆ was distilled over sodium under argon.

Starting materials, Me₃GeGePh₃ (**3**),^[56b] Me₃GeGe(pTol)₃ (**4**),^[48] Ph₃GeGePh₃ (**5**),^[56b, 83] (C₆F₅)₃GeGePh₃ (**6**)^[46] were synthesized according to the literature procedures. Metal carbonyls, M(CO)₆ (M = Cr (**1a**), M = Mo (**1b**), M = W (**1c**)), were sublimed under vacuum before using. Other reagents were used as supplied.

Attention! The work with metal carbonyl compounds should be performed under special conditions (under fume hood) due to volatility and formation of CO.

X-Ray crystallography. Crystal data for **3c**, **4a** and **5a** are given in Table S1. These data have been deposited with the Cambridge Crystallographic Data Centre as supplementary publications no. 1865890 (**3c**), 1865473 (**4a**), 1865889 (**5a**). They can be obtained free of charge from the Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif.

Electrochemistry. Electrochemical measurements were carried out using an Autolab 302N potentiostat interfaced through Nova 2.0 software to a personal computer. Electrochemical measurements were performed in a glovebox under oxygen levels of less than 5 ppm using solvent that had been purified by passing through an alumina-based purification system. Diamond-polished glassy carbon electrodes of 3 mm diameter were employed for cyclic voltammetry (CV) scans. CV data were evaluated using standard diagnostic criteria for diffusion control and for chemical and electrochemical reversibility. The experimental reference electrode was a silver wire coated with anodically deposited silver chloride and separated from the working solution by a fine glass frit. The electrochemical potentials in this article are referenced to ferrocene/ferrocenium couple. The ferrocene potential was obtained by its addition to the analyte solution.

Data for initial compounds

Me₃GeGePh₃ (3). ¹H NMR (400.130 MHz, C₆D₆): δ = 7.55-7.50 (m, 6H), 7.16-7.10 (m, 9H) (aromatic protons), 0.36 (s, 9H, GeMe₃) ppm. ¹³C NMR

(100.613 MHz, C₆D₆): δ = 138.49 (*ipso*-C₆H₅), 135.61 and 128.67 (*o*- and *m*-C₆H₅), 128.84 (*p*-C₆H₅), -0.88 (GeMe₃) ppm.

Ph₃GeGePh₃ (5). ¹H NMR (400.130 MHz, C₆D₆): δ = 7.60-7.56 (m, 12H), 7.09-7.01 (m, 18H) (aromatic protons) ppm. ¹³C NMR (100.613 MHz, C₆D₆): δ = 137.92 (*ipso*-C₆H₅), 135.98 and 128.74 (*o*- and *m*-C₆H₅), 129.19 (*p*-C₆H₅) ppm.

(C₆F₅)₃GeGePh₃ (6). ¹H NMR (400.130 MHz, C₆D₆): δ = 7.41-7.33 (m, 6H), 7.06-6.98 (m, 3H), 6.95-6.87 (m, 6H) (aromatic protons) ppm. ¹³C NMR (100.613 MHz, C₆D₆): δ = 135.08 and 128.73 (*o*- and *m*-C₆H₅), 133.21 (*ipso*-C₆H₅), 130.13 (*p*-C₆H₅) ppm. ¹⁹F NMR (376.498 MHz, C₆D₆): δ = -124.52 (-124.60) (2F), 149.03 (-149.13) (1F), -159.68 (-159.78) (2F) ppm.

Synthesis of Ph₃GeGeMe₂GePh₃ (7). At -78 the solution of *n*BuLi in hexane (2.5 M, 2.0 mL, 5.00 mmol) was added dropwise to the solution of Ph₃GeH (1.52 g, 5.00 mmol) in ether (50 mL). After stirring at the same temperature for 2 h the solution of Ph₃GeLi generated *in situ* was used further without isolation. Then Me₂GeCl₂ (0.43 g, 2.50 mol) was added to the solution of Ph₃GeLi obtained as described above, reaction mixture was slowly warmed to room temperature and stirred overnight. Then water (50 mL) was added, aqueous phase was extracted with ether (3x20 mL), combined organic phases were dried over MgSO₄, and then all volatile materials were removed under reduced pressure. After recrystallization from toluene/*n*-heptane mixture compound **7** (1.49 g, 84 %) was isolated as white crystals. ¹H NMR (400.130 MHz, C₆D₆): δ = 7.46-7.39 (m, 12H), 7.14-7.03 (m, 18H) (aromatic protons), 0.70 (s, 6H, GeMe₂) ppm. ¹³C NMR (100.613 MHz, C₆D₆): δ = 138.10 (*ipso*-C₆H₅), 135.85 and 128.61 (*o*- and *m*-C₆H₅), 128.84 (*p*-C₆H₅), -2.00 (GeMe₂) ppm. The ¹³C NMR data in CDCl₃ corresponds to known from the literature.^[64] HRMS (ESI), *m/z* [M-H]⁺, calcd. 709.6045, found 709.6036.

Synthesis of the initial complexes

Synthesis of fac-(MeCN)₃M(CO)₃ (M = Cr (2a), Mo (2b), W (2c)). Synthesis was performed using literature procedure,^[40] by refluxing corresponding M(CO)₆ in MeCN, what allows to obtain (MeCN)₃M(CO)₃ (72 % for **2a**; 67 % for **2b**; 76 % for **2c**) as a yellow powder.

Synthesis of the target arylogogermane carbonyl complexes

Me₃GeGePh₂(Ph- η^6)Cr(CO)₃ (3a). Solid (MeCN)₃Cr(CO)₃ (**2a**) (0.34 g, 1.31 mmol) was added to the solution of Me₃GeGePh₃ (**3**) (0.46 g, 1.09 mmol) in THF (20 mL), the mixture obtained was refluxed for 1 h. Then all volatile materials were removed under reduced pressure and the residue was purified by flash-chromatography (SiO₂, petroleum ether/toluene 4:1) giving target compound **3a** (*R_f* 0.3, 0.25 g, 42 %) as a yellow powder. ¹H NMR (400.130 MHz, C₆D₆): δ = 7.59-7.53 (m, 4H), 7.26-7.17 (m, 6H) (aromatic protons), 5.03 (dd, ³J_{H-H} = 6.6 Hz, ⁴J_{H-H} = 1.0 Hz, 2H, *o*-C₆H₅*Cr(CO)₃), 4.56 (tt, ³J_{H-H} = 6.3 Hz, ⁴J_{H-H} = 1.0 Hz, 1H, *p*-C₆H₅*Cr(CO)₃), 4.23 (t, ³J_{H-H} = 6.3 Hz, 2H, *m*-C₆H₅*Cr(CO)₃), 0.49 (s, 9H, GeMe₃) ppm. ¹³C NMR (100.613 MHz, C₆D₆): δ = 233.52 (Cr(CO)₃), 136.94 (*ipso*-C₆H₅), 135.61 and 128.84 (*o*- and *m*-C₆H₅), 129.43 (*p*-C₆H₅), 100.25 (*ipso*-C₆H₅*Cr(CO)₃), 100.11 and 90.79 (*o*- and *m*-C₆H₅*Cr(CO)₃), 94.80 (*p*-C₆H₅*Cr(CO)₃), -0.49 (GeMe₃) ppm. MS (EI) *m/z* (%): 558 ([M]⁺, 2), 474 ([M - 3CO]⁺, 100), 422 ([M - Cr(CO)₃]⁺, 42), 412 ([M - CO - GeMe₃]⁺, 22), 356 ([M - 3CO - GeMe₃]⁺, 4), 304 ([Ph₃Ge]⁺, 34), 227 ([Ph₂Ge]⁺, 24), 52 ([Cr]⁺, 3). UV/vis (CH₂Cl₂), λ_{max} , nm (ϵ , M⁻¹ cm⁻¹): 232 (sh) (2.47x10⁴), 263 (sh) (0.53x10⁴), 320 (0.67x10⁴). IR (KBr) ν , cm⁻¹: 1971 (s) (A₁, CO), 1910/1894 (s) (E, CO). IR (nujol) ν , cm⁻¹: 1974 (s) (A₁, CO), 1897 (s) (E, CO). HRMS (ESI), *m/z* [M-H]⁺, calcd. 556.7121, found 556.7107. C₂₄H₂₄CrGe₂O₃ (557.7200): Calcd. C 51.68, H 4.34; Found: C 51.26, H 4.12 %.

Me₃GeGePh₂(Ph- η^6)Mo(CO)₃ (3b). Analogously to **3a** using Mo(CO)₃(NCMe)₃ (**2b**) (0.70 g, 2.30 mmol) and Me₃GeGePh₃ (**3**) (0.89 g, 2.11 mmol). The reaction was accompanied by a noticeable decomposition (precipitation of a black precipitate). After evaporation, dissolving in ether and passing through a pad of SiO₂, ¹H NMR analysis of the mixture obtained indicates the presence of initial compound **3** (major) and complex **3b** (trace amounts). For complex **3b**: ¹H NMR (400.130 MHz, C₆D₆): δ = 5.22 (dd, ³J_{H-H} = 6.5 Hz, ⁴J_{H-H} = 0.8 Hz, 2H, *o*-C₆H₅*Mo(CO)₃), 4.75 (tt, ³J_{H-H} = 6.5 Hz, ⁴J_{H-H} = 0.8 Hz, 1H, *p*-C₆H₅*Mo(CO)₃), 4.44 (t, ³J_{H-H} = 6.5 Hz, 2H, *m*-C₆H₅*Mo(CO)₃) ppm. Other signals are overlapped with the signals of Me₃GeGePh₃ (**3**). IR (nujol) ν , cm⁻¹: 1978 (s) (A₁, CO), 1900 (s) (E, CO). HRMS (ESI), *m/z* [M-H]⁺, calcd. 600.6565, found 600.6548.

Me₃GeGePh₂(Ph- η^6)W(CO)₃ (3c). *Method 1.* Analogously to **3a** using W(CO)₃(NCMe)₃ (**2c**) (0.78 g, 1.99 mmol) and Me₃GeGePh₃ (**3**) (0.59 g, 1.40 mmol). After chromatography (SiO₂) initial **3** (0.27 g, *R_f* 0.6, ether/petroleum ether 1:4) and target complex **3c** (0.14 g, 27 %, conversion 54 %, *R_f* 0.3, ether/petroleum ether 1:3, orange powder) were isolated.

Method 2. Solid W(CO)₆ (**1c**) (0.42 g, 1.19 mmol) was added to the solution of Me₃GeGePh₃ (**3**) (0.50 g, 1.19 mmol) in a mixture of (*n*Bu)₂O/THF (10:1) (30 mL). The mixture obtained was frozen in liquid nitrogen, evacuated in high pressure and warmed to room temperature (procedure was repeated three times). Then it was refluxed on oil bath (150 °C) for 5 h, whereas significant decomposition (black precipitate) was observed. Then all volatile materials were removed under reduced pressure, the residue was dissolved in ether and the residue was passed through a pad of SiO₂ indicating the decomposition. ¹H NMR (400.130 MHz, C₆D₆): δ = 7.57-7.53 (m, 6H), 7.24-7.17 (m, 4H) (aromatic protons), 4.98 (dd, ³J_{H-H} = 6.3 Hz, ⁴J_{H-H} = 1.2 Hz, 2H, *o*-C₆H₅*W(CO)₃), 4.50 (tt, ³J_{H-H} = 6.3 Hz, ⁴J_{H-H} = 1.2 Hz, 1H, *p*-C₆H₅*W(CO)₃), 4.17 (tt, ³J_{H-H} = 6.3 Hz, ⁴J_{H-H} = 1.2 Hz, 2H, *m*-C₆H₅*W(CO)₃), 0.45 (s, 9H, GeMe₃) ppm. ¹³C NMR (100.613 MHz, C₆D₆): δ = 209.95 (W(CO)₃), 136.98 (*ipso*-C₆H₅), 135.50 and 128.91 (*o*- and *m*-C₆H₅), 129.54 (*p*-C₆H₅), 97.63 (*ipso*-C₆H₅*W(CO)₃), 97.23 and 89.19 (*o*- and *m*-C₆H₅*W(CO)₃), 91.86 (*p*-C₆H₅*W(CO)₃), -0.57 (GeMe₃) ppm. MS (EI) *m/z* (%): 690 ([M]⁺, 4), 606 ([M - 3CO]⁺, 22), 544 ([M - CO - GeMe₃]⁺, 16), 488 ([M - 3CO - GeMe₃]⁺, 2), 422 ([M - W(CO)₃]⁺, 46), 304 ([Ph₃Ge]⁺, 100), 227 ([Ph₂Ge]⁺, 27), 52 ([Cr]⁺, 2). UV/vis (CH₂Cl₂), λ_{max} , nm (ϵ , M⁻¹ cm⁻¹): 233 (sh) (1.96x10⁴), 291 (0.38x10⁴), 322 (0.46x10⁴). IR (nujol) ν , cm⁻¹: 2924 (s), 2862 (m), 1980 (s) (A₁, CO), 1904 (s) (E, CO), 1460 (m). HRMS (ESI), *m/z* [M-H]⁺, calcd. 688.5565, found 688.5567. C₂₄H₂₄Ge₂O₃W (689.5634): Calcd. C 41.80, H 3.51; Found: C 39.68, H 2.87 %.

Me₃GeGe(pTol)₂(pTol- η^6)Cr(CO)₃ (4a). Analogously to **3a** using Cr(CO)₃(NCMe)₃ (**2a**) (0.27 g, 1.04 mmol) and Me₃GeGe(pTol)₃ (**4**) (0.48 g, 1.04 mmol). After chromatography (SiO₂) initial **4** (0.15 g, *R_f* 0.8, toluene/petroleum ether 1:3) and target complex **4a** (0.24 g, 56 %, *R_f* 0.3, toluene/petroleum ether 1:3, conversion 68 %) were isolated as yellowish crystals. ¹H NMR (400.130 MHz, C₆D₆): δ = 7.58, 7.11 (2d, *J_{H-H}* = 6.7 Hz, each 4H, 2*p*-C₆H₄), 5.23, 4.23 (2d, *J_{H-H}* = 5.1 Hz, each 2H, *p*-C₆H₄*Cr(CO)₃), 2.10 (s, 6H, 2C₆H₄Me), 1.58 (s, 3H, C₆H₄Me*Cr(CO)₃), 0.58 (s, 9H, GeMe₃) ppm. ¹³C NMR (100.613 MHz, C₆D₆): δ = 234.06 (Cr(CO)₃), 139.17, 133.52 (2 *ipso*-C₆H₄Me-*p*), 135.64, 129.72 (2 *o*-C₆H₄Me-*p*), 110.60, 98.21 (2 *ipso*-C₆H₄Me-*p**Cr(CO)₃), 101.05, 91.98 (2 *o*-C₆H₄Me-*p**Cr(CO)₃), 21.31 (C₆H₄Me-*p*), 20.26 (C₆H₄Me-*p**Cr(CO)₃), -0.46 (GeMe₃) ppm. MS (EI) *m/z* (%): 600 ([M]⁺, 2), 516 ([M - 3CO]⁺, 100), 464 ([M - Cr(CO)₃]⁺, 454 ([M - CO - GeMe₃]⁺, 18), 398 ([M - 3CO - GeMe₃]⁺, 1), 38), 346 ([pTol]₃Ge]⁺, 14), 255 ([pTol]₂Ge]⁺, 32), 52 ([Cr]⁺, 4). IR (THF) ν , cm⁻¹: 1961 (s) (A₁, CO), 1887 (s) (E, CO). IR (nujol) ν , cm⁻¹: 1970 (s) (A₁, CO), 1894 (s) (E, CO). UV/vis (CH₂Cl₂), λ_{max} , nm (ϵ , M⁻¹ cm⁻¹): 233 (sh) (4.55x10⁴), 267 (sh) (0.80x10⁴), 321 (0.87x10⁴). HRMS

(ESI), m/z [M-H]⁺, calcd. 598.7924, found 598.7911. C₂₇H₃₀CrGe₂O₃ (599.7998): Calcd. C 54.07, H 5.04; Found: C 52.78, H 4.22 %.

Ph₃GeGePh₂(Ph-η⁶)Cr(CO)₃ (5a). *Method 1.* Dimethyl succinate (0.46 g, 3.10 mmol) and Cr(CO)₆ (1a) (0.76 g, 3.44 mmol) were added to the suspension of Ph₃GeGePh₃ (5) (1.90 g, 3.10 mmol) in decalin (20 mL). Reaction mixture was frozen in liquid nitrogen, evacuated in vacuum and warmed to room temperature (three times); then it was heated at 190 °C for 2 h. Then all volatile materials were removed under high vacuum, residue was purified by chromatography (SiO₂, petroleum ether, then petroleum ether/benzene 3:2), giving target compound 5a (*R_f* 0.4, 0.18 g, 28 %, conversion 29 %) as a yellow powder and unreacted Ph₃GeGePh₃ (5a) (*R_f* 0.7 in benzene, 1.37 g).

Method 2. Analogously to 3c (Method 2) using Cr(CO)₆ (1a) (0.31 g, 1.40 mmol) and Ph₃GeGePh₃ (5) (0.80 g, 1.32 mmol) in (*n*Bu)₂O/THF (10:1) mixture (20 mL), heating for 26 h. The residue was purified by chromatography (SiO₂, *R_f* 0.3, benzene/petroleum ether 1:3) giving 5a (0.31 g, 32 %) as a yellow powder. ¹H NMR (400.130 MHz, C₆D₆): δ = 7.75-7.70 (m, 4H), 7.61-7.54 (m, 6H), 7.14-7.06 (m, 15H) (aromatic protons), 5.05 (d, *J_{H-H}* = 6.3 Hz, 2H, *o*-C₆H₅*Cr(CO)₃), 4.48 (t, *J_{H-H}* = 6.3 Hz, 1H, *p*-C₆H₅*Cr(CO)₃), 4.14 (t, *J_{H-H}* = 6.3 Hz, 2H, *m*-C₆H₅*Cr(CO)₃) ppm. ¹³C NMR (100.613 MHz, C₆D₆): δ = 233.10 (Cr(CO)₃), 136.84 (Ge(*p*-C₆H₅)₂(Ph*Cr(CO)₃)), 136.14 (Ge(*p*-C₆H₅)₃), 135.94 and 128.90 (Ge(*o*- and *m*-C₆H₅)₃), 135.79 (Ge(*ipso*-C₆H₅)₃), 129.89 and 129.52 (Ge(*o*- and *m*-C₆H₅)₂(Ph*Cr(CO)₃)), 128.53 (Ge(*ipso*-C₆H₅)₂(Ph*Cr(CO)₃)), 100.60 (Ge(*ipso*-C₆H₅)Cr(CO)₃), 99.82 and 90.74 (Ge(*o*- and *m*-C₆H₅)*Cr(CO)₃), 94.27 (Ge(*p*-C₆H₅)*Cr(CO)₃) ppm. MS (EI) m/z (%): 744 ([M]⁺, 2), 660 ([M - 3CO]⁺, 43), 608 ([M - Cr(CO)₃]⁺, 38), 412 ([M - CO - GePh₃]⁺, 2), 357 ([M - 3CO - GePh₃]⁺, 2), 304 ([Ph₃Ge]⁺, 100), 227 ([Ph₂Ge]⁺, 21), 151 ([PhGe + H]⁺, 18), 52 ([Cr]⁺, 2). UV/vis (CH₂Cl₂), λ_{max}, nm (ε, M⁻¹ cm⁻¹): 238 (sh) (3.57x10⁴), 267 (sh) (1.04x10⁴), 322 (0.64x10⁴). IR (nujol) ν, cm⁻¹: 2932 (s), 1975 (s) (A₁, CO), 1900 (s) (E, CO), 1454 (s), 1370 (m). HRMS (ESI), m/z [M-H]⁺, calcd. 742.9208, found 742.9211. C₃₉H₃₀CrGe₂O₃ (743.9282): Calcd. C 62.97, H 4.06; Found: C 60.72, H 3.66 %.

(C₆F₅)₃GeGePh₂(Ph-η⁶)Cr(CO)₃ (6a). Analogously to 3c (Method 2) using Cr(CO)₆ (1a) (0.15 g, 0.69 mmol) and (C₆F₅)₃GeGePh₃ (6) (0.58 g, 0.66 mmol) in (*n*Bu)₂O/THF (10:1) mixture (20 mL), heating for 30 h. The residue was purified by chromatography (SiO₂, *R_f* 0.3, ether/petroleum ether 1:3), giving 6a (0.11 g, 16 %) as an orange powder. ¹H NMR (400.130 MHz, C₆D₆): δ = 7.58-7.54 (m, 4H), 7.04-7.00 (m, 6H) (aromatic protons), 4.91 (d, ³*J_{H-H}* = 6.3 Hz, 2H, *o*-C₆H₅*Cr(CO)₃), 4.35 (t, ³*J_{H-H}* = 6.3 Hz, 1H, *p*-C₆H₅*Cr(CO)₃), 4.06 (t, ³*J_{H-H}* = 6.3 Hz, 2H, *m*-C₆H₅*Cr(CO)₃) ppm. ¹³C NMR (100.613 MHz, C₆D₆): δ = 232.17 (Cr(CO)₃), 149.85-149.66, 147.42-147.24, 144.88-144.78, 142.39-142.13, 139.16-138.85, 136.63-136.27 (6m, Ge(C₆F₅)₃), 135.19 and 129.04 (*o*- and *m*-C₆H₅), 131.25 (*ipso*-C₆H₅), 130.86 (*p*-C₆H₅), 97.79 and 90.94 (*o*- and *m*-C₆H₅*Cr(CO)₃), 95.16 (*ipso*-C₆H₅*Cr(CO)₃), 93.51 (*p*-C₆H₅*Cr(CO)₃) ppm. ¹⁹F NMR (376.498 MHz, C₆D₆): δ = -124.29 – (-124.34) (2F), 148.61 – (-148.72) (1F), -159.46 – (-159.56) (2F) ppm. MS (EI) m/z (%): 1014 ([M]⁺, 1), 930 ([M - 3CO]⁺, 32), 878 ([M - Cr(CO)₃]⁺, 22), 575 ([C₆F₅)₃Ge]⁺, 8), 304 ([Ph₃Ge]⁺, 100), 167 ([C₆F₅]⁺, 16), 52 ([Cr]⁺, 2). UV/vis (CH₂Cl₂), λ_{max}, nm (ε, M⁻¹ cm⁻¹): 234 (sh) (2.15x10⁴), 261 (0.78x10⁴), 322 (0.41x10⁴). IR (nujol) ν, cm⁻¹: 2852 (s), 1976 (s) (A₁, CO), 1901 (s) (E, CO), 1650 (m), 1462 (s), 1372 (m), 1082 (s). HRMS (ESI), m/z [M-H]⁺, calcd. 1012.7777, found 1012.7782. C₃₉H₁₅CrF₁₅Ge₂O₃ (1013.7851): Calcd. C 46.20, H 1.49; Found: C 44.64, H 1.07 %.

Ph₃GeGeMe₂GePh₂(Ph-η⁶)Cr(CO)₃ (7a). *Method 1.* Dimethyl succinate (0.30 g, 2.02 mmol, 2.0 eq.) and Cr(CO)₆ (1a) (0.49 g, 2.22 mmol, 2.2 eq.) were added to the suspension of [Ph₃Ge]₂GeMe₂ (7) (0.72 g, 1.01 mmol, 1.0 eq.) in decalin (20 mL). Reaction mixture was frozen in liquid

nitrogen, evacuated in vacuum and warmed to room temperature (three times); then it was heated at 190 °C for 2 h. Then all volatile materials were removed under high vacuum, residue was purified by chromatography (SiO₂), giving unreacted [Ph₃Ge]₂GeMe₂ (7) (0.48 g, *R_f* 0.6, ether/petroleum ether 1:3) and target compound 7a (0.10 g, 34%, *R_f* 0.3, ether/petroleum ether 1:2, conversion 33 %) as a yellow powder.

Method 2. Trigermane [Ph₃Ge]₂GeMe₂ (7) (0.24 g, 0.33 mmol) was added to the solution of Cr(CO)₃(NCMe)₃ (2a) (0.19 g, 0.73, 2.2 eq.) in THF (20 mL), the solution was refluxed for 2 h, then all volatile materials were removed under reduced pressure. The residue was purified by chromatography (SiO₂, ether/petroleum ether 1:2) giving initial compound 7 (*R_f* 0.7, 0.09 g) and target complex 7a as a yellow powder (*R_f* 0.2, 0.08 g, 44 %, conversion 63 %). ¹H NMR (400.130 MHz, C₆D₆): δ = 7.44-7.34 (m, 10H), 7.15-7.07 (m, 15H) (aromatic protons), 4.80 (d, *J_{H-H}* = 5.9 Hz, 2H, *o*-C₆H₅*Cr(CO)₃), 4.53 (br t, *J_{H-H}* = 5.9 Hz, 1H, *p*-C₆H₅*Cr(CO)₃), 4.17 (br t, *J* = 5.9 Hz, 2H, *m*-C₆H₅*Cr(CO)₃), 0.81 (s, 6H, GeMe₂) ppm. ¹³C NMR (100.613 MHz, C₆D₆): δ = 233.36 (Cr(CO)₃), 137.62 (Ge(*ipso*-C₆H₅)₃), 136.52 (Ge(*ipso*-C₆H₅)₂Ph*Cr(CO)₃), 135.91 and 128.81 (Ge(*o*- and *m*-C₆H₅)₂Ph*Cr(CO)₃), 135.78 and 128.71 (Ge(*o*- and *m*-C₆H₅)₃), 129.53 (Ge(*p*-C₆H₅)₂Ph*Cr(CO)₃), 129.01 (Ge(*p*-C₆H₅)₃), 100.54 (Ge(*ipso*-C₆H₅)Cr(CO)₃), 99.95 and 90.80 (Ge(*o*- and *m*-C₆H₅)*Cr(CO)₃), 94.52 (Ge(*p*-C₆H₅)*Cr(CO)₃), -1.68 (GeMe₂) ppm. UV/vis (CH₂Cl₂), λ_{max}, nm (ε, M⁻¹ cm⁻¹): 244 (sh) (3.60x10⁴), 261 (sh) (2.10x10⁴), 321 (0.38x10⁴). IR (nujol) ν, cm⁻¹: 2942 (s), 1972 (s) (A₁, CO), 1896 (s) (E, CO), 1452 (m). MS (EI) m/z (%): 847 ([M]⁺, 1), 763 ([M - 3CO]⁺, 22), 711 ([M - Cr(CO)₃]⁺, 100), 407 ([Ph₃GeGeMe₂]⁺, 12), 304 ([Ph₃Ge]⁺, 22), 52 ([Cr]⁺, 2). HRMS (ESI), m/z [M-H]⁺, calcd. 845.6298, found 845.6281. C₄₁H₃₆CrGe₃O₃ (846.6372): Calcd. C 58.16, H 4.29; Found: C 60.12, H 4.52 %.

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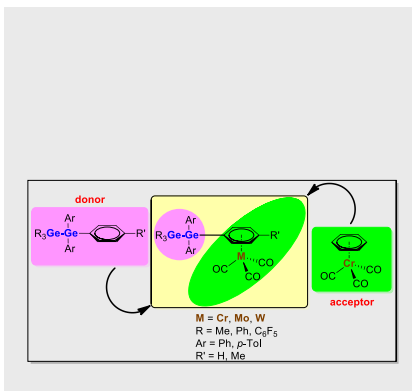
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Entry for the Table of Contents

FULL PAPER

A series of transition metal carbonyl complexes bearing aryloligogermanes ligand was obtained and thoroughly investigated through NMR, IR, UV/vis spectroscopy, electrochemistry, DFT, and XRD. Oligogermanyl substituents have been found to have an electron donor behaviour, whereas the $M(CO)_3$ group is behaving as an electron withdrawing group. The dependences of the physical properties on the nature of M, R, Ar and on the number of Ge atoms were studied.



Oligogermanium Complexes

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