


 Cite this: *RSC Adv.*, 2020, **10**, 9582

Unprecedented formation of methylsilylcarbonates from iridium-catalyzed reduction of CO₂ with hydrosilanes†

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The iridium complex [Ir(μ-CF₃SO₃)(κ²-NSi^{Me2})₂]₂ (**3**) (NSi^{Me2} = {4-methylpyridine-2-yloxy}dimethylsilyl) has been prepared by reaction of [Ir(μ-Cl)(κ²-NSi^{Me2})₂]₂ (**1**) with two equivalents of AgCF₃SO₃. The solid structure of **3** evidenced its dinuclear nature, being a rare example of an iridium species with triflate groups acting as bridges. The **3**-catalyzed reduction of CO₂ with HSiMe(OSiMe₃)₂ affords a mixture of the corresponding silylformate and methoxysilane together with the silylcarbonate CH₃OCO₂SiMe(OSiMe₃)₂ (**4a**). This is the first time that the formation of silylcarbonates has been observed from the catalytic reduction of CO₂ with silanes. Analogous behaviour has been observed when HSiMe₂Ph and HSiMePh₂ were used as reductants.

 Received 8th January 2020
 Accepted 23rd February 2020

DOI: 10.1039/d0ra00204f

rsc.li/rsc-advances

Introduction

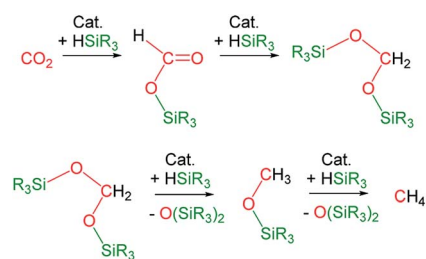
The catalytic reaction of CO₂ with silicon-hydrides has proven to be an effective and thermodynamically favoured methodology for its reduction to formate, formaldehyde, methanol or methane (Scheme 1).^{1,2}

However, to the best of our knowledge the formation of methylsilylcarbonates, MeOCO₂SiR₃ (Fig. 1),³ as products of the catalytic reduction of CO₂ with silicon-hydrides has not been reported so far.

On the other hand, hydrosiloxanes are obtained in large scale as side products of the silicone industry and therefore, the development of catalytic processes effective for the selective reduction of CO₂ based on hydrosiloxanes, instead of hydrosilanes, as reductants is of great interest.⁴ In this context, in recent years we have focused our research on the application of iridium species as catalysts for the reduction of CO₂ with 1,1,1,3,5,5,5-heptamethyltrisiloxane, HSiMe(OSiMe₃)₂.^{5,6}

Recently, we have described the synthesis and catalytic behavior of the iridium(III) complexes [Ir(μ-Cl)(κ²-NSi^{Me2})₂]₂ (**1**) and [Ir(CF₃CO₂)(κ²-NSi^{Me2})₂]₂ (**2**) (NSi^{Me2} = {4-methylpyridine-2-yloxy}dimethylsilyl) (Scheme 2).⁶ Species **2** catalyzed the selective reduction of CO₂, depending on the reaction conditions, to silylformate or methoxysilane with HSiMe(OSiMe₃)₂.⁶ As

a continuation of our studies on the chemistry of iridium complexes with pyridine-2-yloxyI-silyl ligands, we have found that the triflate derivative [Ir(μ-CF₃SO₃)(κ²-NSi^{Me2})₂]₂ (**3**) (Scheme 2) catalyzes the reaction of CO₂ (3 bar) with HSiMe(OSiMe₃)₂ to give a mixture of the corresponding silylformate and methoxysilane along with a new compound that has been characterized by multinuclear NMR spectroscopy as CH₃OCO₂SiMe(OSiMe₃)₂ (**4a**). To the best of our knowledge, this is the first time that the formation of silylcarbonates from the catalytic reduction of CO₂ with silanes has been observed. The results from these studies are described below.



Scheme 1 Products from the catalytic reduction of CO₂ with silicon-hydrides reported so far.

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† Electronic supplementary information (ESI) available: NMR spectra and crystallographic information. CCDC 1972218. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d0ra00204f

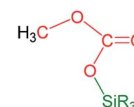
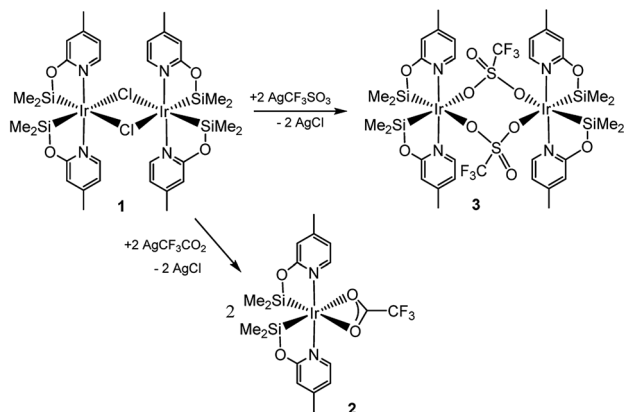


Fig. 1 Methylsilylcarbonates.





Scheme 2 Preparation of the catalytic precursors 2 and 3.

Results and discussion

Synthesis and characterization of the catalyst precursor

The reaction of complex **1** with two equivalents of AgCF_3SO_3 leads to the formation of $[\text{Ir}(\mu\text{-CF}_3\text{SO}_3)(\kappa^2\text{-NSi}^{\text{Me}_2})_2]_2$ (**3**), which has been isolated as a white solid in 89% yield. The solid structure of complex **3** evidenced its dinuclear nature (Fig. 2). As far as we know, **3** is the first example of an iridium species with triflate groups acting as bridges. The molecule exhibits a crystallographic inversion center, with half the molecule as the independent part. Each metal atom shows an octahedral geometry like that found for **1**.⁶ Main difference concerns the separation between the iridium atoms (6.0870(2) Å) in **3**, significantly longer than that observed for **1** with chlorine bridges (4.0718(12) Å).

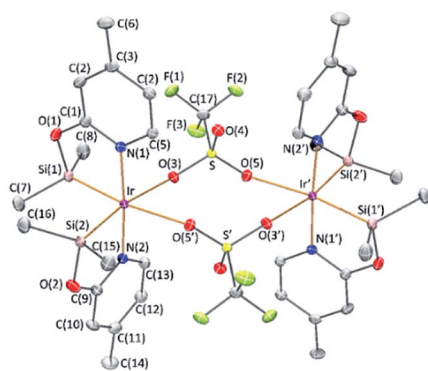


Fig. 2 Molecular structure of compound **3**. For clarity hydrogen atoms have been omitted. Primed atoms are related to unprimed ones through $1 - x, 1 - y, -z$ symmetry operation. Selected bond distances (Å) and angles (°): Ir–Si(1), 2.2570(5); Ir–Si(2), 2.2615(5); Ir–O(3), 2.3653(12); Ir–O(5'), 2.4331(13); Ir–N(1), 2.0590(15); Ir–N(2), 2.0583(15); Si(1)–Ir–Si(2), 91.68(5); Si(1)–Ir–O(3), 95.18(3); Si(1)–Ir–O(5'), 172.22(3); Si(1)–Ir–N(1), 81.57(4); Si(1)–Ir–N(2), 94.94(4); Si(2)–Ir–O(3), 170.27(3); Si(2)–Ir–O(5'), 93.26(3); Si(2)–Ir–N(1), 95.76(4); Si(2)–Ir–N(2), 82.22(4); O(3)–Ir–O(5'), 80.67(4); O(3)–Ir–N(1), 92.06(5); O(3)–Ir–N(2), 90.32(5); O(5')–Ir–N(1), 91.95(5); O(5')–Ir–N(2), N(1)–Ir–N(2), 175.94(6).

^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra of **3** in C_6D_6 show only one pattern of resonances for the four NSi^{Me_2} ligands, which agrees with its high symmetry in solution. $^{29}\text{Si}\{^1\text{H}\}$ NMR spectra of **3** show a resonance at δ 38.2 ppm, corresponding to the four silicon atoms, which compares well to the value of δ 39.7 ppm reported for **2**.⁶ In addition, a singlet resonance at δ –76.97 ppm in the $^{19}\text{F}\{^1\text{H}\}$ NMR spectra confirm the presence of the triflate ligand.

3-Catalyzed reduction of $^{13}\text{CO}_2$ with silicon hydrides

Preliminary studies on the **3**-catalyzed reaction of CO_2 (3 bar) with $\text{HSiMe}(\text{OSiMe}_3)_2$ at 298 K showed that 24 hours are required to achieve the conversion of 94% of the starting hydrosiloxane into a mixture of the corresponding silylformate (53.0%) and methoxysilane (21.4%) along with a new compound (25.6%) ($\text{TOF} = 3.9 \text{ h}^{-1}$). This behavior differs from that previously reported for **2**, which under the same reaction conditions catalyzed the full conversion of $\text{HSiMe}(\text{OSiMe}_3)_2$ into silylformate (93.0%) and methoxysilane (7.0%) in 3.5 h ($\text{TOF} = 28.6 \text{ h}^{-1}$).⁶ Therefore, complex **2** is not only more active but also more selective than **3**.

To determine the nature of the new species observed when **3** is used as catalyst for the hydrosilylation of CO_2 , we decided to study this reaction using isotopically labeled $^{13}\text{CO}_2$. Thus, ^1H and ^{13}C NMR studies of the **3**-catalyzed (1.0 mol%) reaction of $^{13}\text{CO}_2$ (2.7 bar, 323 K, C_6D_6) with $\text{HSiMe}(\text{OSiMe}_3)_2$ were performed. These experiments evidenced, since early reaction stages, the presence of a mixture of $^{13}\text{CH}_3\text{O}-^{13}\text{CO}_2\text{-SiMe}(\text{OSiMe}_3)_2$ (**4a**), together with the corresponding silylformate, $\text{H}^{13}\text{CO}_2\text{SiMe}(\text{OSiMe}_3)_2$ (**5a**), and the methoxysilane, $^{13}\text{CH}_3\text{OSiMe}(\text{OSiMe}_3)_2$ (**6a**) (Table 1).

The most characteristic resonance in the ^1H NMR spectra of **4a** is a doublet of doublets centered at δ 3.33 ppm ($^1J_{\text{C-H}} = 146.9 \text{ Hz}$; $^3J_{\text{C-H}} = 4.1 \text{ Hz}$) corresponding to the $^{13}\text{CH}_3\text{O}$ protons

Table 1 Results from **3**-catalyzed reduction of $^{13}\text{CO}_2$ (2.7 bar) with $\text{HSiMe}(\text{OSiMe}_3)_2$ ^a

Entry	<i>T</i> (K)	4a ^b	5a ^b	6a ^b	Conversion ^c	Time (h)
1	323	26.7	65.2	8.1	70	3
2	323	18.8	70.0	11.2	98	12
3	323	19.0	70.0	11.0	98	24
4	358	11.7	68.7	19.6	100	24 ^d
5	358	8.5	68.0	23.5	100	48 ^d
6 ^e	323	0	0	0	0	24

^a General conditions: **3** (1.0 mol%) in 0.5 mL of dry C_6D_6 . ^b Calculated by ^1H NMR and expressed in mol%. ^c Calculated by ^1H NMR relative to the starting hydrosiloxane and expressed in mol%. ^d After 24 h at 323 K. ^e Control experiment without iridium catalyst (Fig. S24 see ESI).



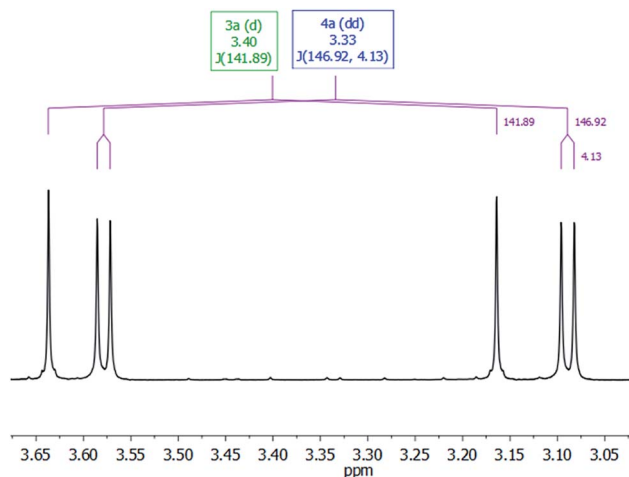
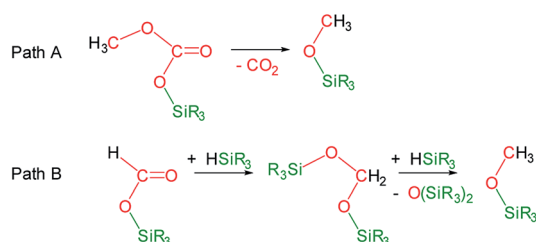


Fig. 3 CH_3O -region of the ^1H NMR spectra in C_6D_6 of the **3**-catalyzed reaction of $^{13}\text{CO}_2$ (2.7 bar) with $\text{HSiMe}(\text{OSiMe}_3)_2$ at 323 K.

(Fig. 3). This resonance shows a direct C–H bond correlation in the ^1H – ^{13}C HSQC spectra with a doublet resonance centered at δ 54.1 ppm ($^3J_{\text{C-C}} = 1.7$ Hz), assigned to the $^{13}\text{CH}_3\text{O}$ carbon in the ^{13}C APT NMR spectra, and also a C–H bond correlation in the ^1H – ^{13}C HMBC spectra with a doublet resonance that appeared centered at δ 153.0 ppm ($^3J_{\text{C-C}} = 1.8$ Hz), assigned to the $^{13}\text{CO}_3$ carbon in the ^{13}C APT NMR spectra. These data compare well with those reported for alkyl-silyl-carbonates³ and support the structure proposed for **4a** in Table 1 (see ESI†).

Under the above described reaction conditions the slow consumption of $\text{HSiMe}(\text{OSiMe}_3)_2$ was observed (Table 1, entries 1 and 2). Thus, after 12 h the reaction is complete and ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra evidenced the consumption of all the starting $^{13}\text{CO}_2$ and of the 98% of $\text{HSiMe}(\text{OSiMe}_3)_2$ to give a mixture of **4a** (18.8 mol%), **5a** (70.0 mol%) and **6a** (11.2 mol%) (Table 1, entry 2). Heating this sample for another 12 h at 323 K did not evidence changes in the composition of the mixture (Table 1, entry 3). Interestingly, increasing the temperature at 358 K the slow transformation of **4a** into **6a** occurs (Table 1 entries 4 and 5), in addition, the formation of $^{13}\text{CO}_2$ was observed. This outcome agrees with the known thermal behavior of alkyl-silyl-carbonates, $\text{ROCO}_2\text{SiR}_3$,^{3d,e} which thermally decompose to give the corresponding methoxysilane and CO_2 (Scheme 3, Path A).



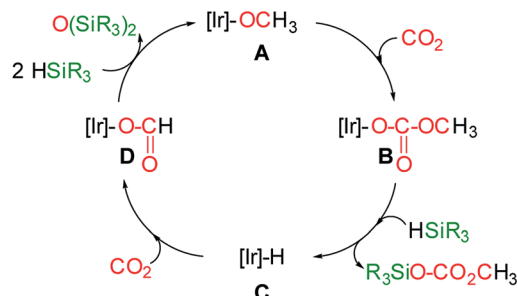
Scheme 3 Reaction paths for the formation of methoxysilanes from the **3**-catalyzed hydrosilylation of CO_2 .

The formation of methoxy-silanes from the catalytic reduction of CO_2 with hydrosilanes has been so far explained by reaction of bis(silyl)acetals with the corresponding hydro-silane (Scheme 3, Path B).^{6,7} Indeed, since it has been observed that at 323 K only traces of **4a** are transformed into **6a** (Table 1 entry 3) it is reasonable to assume that under these conditions **6a** is formed by reactions of the bis(silyl)acetal $^{13}\text{CH}_2\{\text{OSiMe}(\text{OSiMe}_3)_2\}_2$ (**7a**) with $\text{HSiMe}(\text{OSiMe}_3)_2$. In agreement with this we have observed the presence of traces of **7a** in the NMR spectra of this reaction (ESI†). However, after 24 h of reaction 98% of the starting hydrosiloxane was consumed and therefore, under these conditions, the formation of **6a** at 358 K can only be explained by thermal decomposition of **4a** (Scheme 3, Path B).

Analogous behavior has been observed when HSiMe_2Ph and HSiMePh_2 were used as reducing agents. Thus, the formation of the mixtures of methyl-silyl-carbonate $^{13}\text{CH}_3\text{O}-^{13}\text{CO}_2\text{SiR}_3$ ($\text{SiR}_3 = \text{SiMe}_2\text{Ph}$, **4b**; SiMePh_2 , **4c**) and the corresponding silylformate and methoxysilane from the **3**-catalyzed reduction of $^{13}\text{CO}_2$ with HSiR_3 ($\text{SiR}_3 = \text{SiMe}_2\text{Ph}$, SiMePh_2) was also observed (ESI†).

It should be mentioned, that the formation of the methyl-silyl-carbonates was not observed along the **2**-catalyzed reduction of CO_2 with $\text{HSiMe}(\text{OSiMe}_3)_2$.⁶ In this context, even though species **2** and **3** seem very similar, their reactivity against silanes is very different. Thus, while complex **3** is stable in presence of one equivalent of $\text{HSiMe}(\text{OSiMe}_3)_2$ at 323 K, the trifluoroacetate ligand in **2** is reduced to give the corresponding silylether $\text{CF}_3\text{CH}_2\text{OSiR}_3$.^{5d,6} Therefore, it is reasonable to think that the stability of the triflate ligand could play a role stabilizing reaction intermediates, which could open new paths of reaction, no accessible in **2**-catalyzed CO_2 hydrosilylation reactions.

At this point the question arises, how methylsilylcarbonates are formed from the **3**-catalyzed hydrosilylation of CO_2 . A plausible mechanism proposal is shown in Scheme 4. The formation of methylcarbonates has been explained by insertion of CO_2 into the Ir–O bond of iridium– OCH_3 species.⁸ Therefore, we propose that the reaction of an iridium-methoxy intermediate (**A** in Scheme 4) with CO_2 could lead to the methylcarbonate species **B**, which by reaction with HSiR_3 would afford the corresponding methylsilylcarbonate and the Ir–H species **C**. The next step, is the reaction of **C** with CO_2 to give the iridium-formate intermediate **D**, which according to our previous results has been found to be a thermodynamically favoured



Scheme 4 Plausible mechanism proposal for the formation of methyl-silyl-carbonates from the **3**-catalyzed hydrosilylation of CO_2 .



process.^{5d,6} Finally, the reaction of the iridium–formate **D** with excess of silane could regenerate **A** closing the catalytic cycle.

Experimental

General considerations

All manipulations were carried out under an argon atmosphere by Schlenk-type techniques or in a Glovebox MBraun Unilab. Organic solvents were dried by standard procedures and distilled under argon prior to use or obtained oxygen- and water-free from a solvent purification system (Innovative Technologies). ¹H, ¹³C, ²⁹Si and ¹⁹F NMR spectra were obtained on a Bruker AV-300, AV-400 or AV-500 spectrometer. Chemical shifts (δ), reported in ppm, are referenced to the residual solvent peaks and coupling constants (J) are reported in Hz.

Synthesis of 3. Toluene (10 mL) was added to a light-protected Schlenk containing [Ir(μ -Cl)(κ^2 -NSi^{Me2})₂]₂ (**1**) (300 mg, 0.268 mmol) and silver triflate (151 mg, 0.590 mmol). The mixture was stirred at room temperature for 5 hours and then filtered through Celite. Solvent was removed under reduced pressure and the solid was washed with pentane (3 \times 8 mL) to afford a white solid. Yield: 320 mg (89%). ¹H NMR (300 MHz, C₆D₆, 298 K): δ 8.73 (d, J_{H-H} = 6.2 Hz, 2H, py), 6.32 (s, 2H, py), 6.02 (d, J_{H-H} = 6.2 Hz, 2H, py), 1.51 (s, 6H, Me-py), 0.72 (s, 6H, Si-Me), 0.41 (s, 6H, Si-Me). ¹³C NMR plus APT and ¹H-¹³C HSQC (75 MHz, C₆D₆, 298 K): δ 168.7 (s, py), 152.6 (s, py), 148.7 (s, py), 118.4 (s, py), 111.9 (s, py), 20.6 (s, CH₃-py), 3.8 (s, CH₃-Si), 2.4 (s, CH₃-Si). ²⁹Si{¹H} NMR plus ¹H-²⁹Si HMBC (60 MHz, C₆D₆, 298 K): δ 38.2 (Ir-Si). ¹⁹F NMR (282 MHz, C₆D₆, 298 K): δ -76.97 (s, CF₃SO₃). High resolution mass spectrometry (ESI⁺): calc. m/z = 525.1006; found m/z = 525.1004 (M⁺-CF₃SO₃).

3-Catalyzed (1.0 mol%) reaction of ¹³CO₂ with HSiR₃. A Young cap NMR tube was charged with **3** (2.83 mg, 0.0021 mmol), 0.42 mmol of the corresponding silane (114 μ L, HSiMe(OSiMe₃)₂; 64.4 μ L, HSiMe₂Ph; 83.7 μ L, HSiMePh₂) and 0.5 mL of C₆D₆. Argon gas was evacuated by three freeze-pump-thaw cycles. Then the tube was pressurized with ¹³CO₂ (2.7 bar), heated at 323 K and monitored by NMR spectroscopy.

Selected data for 4a. ¹H NMR plus HSQC ¹H-¹³C (300 MHz, C₆D₆, 298 K): δ 3.33 (dd, $^1J_{H-C}$ = 146.9 Hz, $^3J_{H-C}$ = 4.1 Hz, 3H, CH₃OCO₂). ¹³C{¹H} plus HSQC and HMBC ¹H-¹³C (75 MHz, C₆D₆, 298 K): δ 153.0 (d, $^2J_{C-C}$ = 1.7 Hz, CO₃), 54.1 (d, $^2J_{C-C}$ = 1.7 Hz, CH₃O). Selected data for **4b**: ¹H NMR plus HSQC ¹H-¹³C (300 MHz, C₆D₆, 298 K): δ 3.30 (dd, $^1J_{H-C}$ = 147.1 Hz, $^3J_{H-C}$ = 4.1 Hz, 3H, CH₃OCO₂). ¹³C{¹H} plus HSQC and HMBC ¹H-¹³C (75 MHz, C₆D₆, 298 K): δ 154.1 (d, $^2J_{C-C}$ = 1.7 Hz, CO₃), 54.2 (d, $^2J_{C-C}$ = 1.7 Hz, CH₃O). Selected data for **4c**: ¹H NMR plus HSQC ¹H-¹³C (300 MHz, C₆D₆, 298 K): δ 3.23 (dd, $^1J_{H-C}$ = 147.1 Hz, $^3J_{H-C}$ = 4.1 Hz, 3H, CH₃OCO₂). ¹³C{¹H} plus HSQC and HMBC ¹H-¹³C (75 MHz, C₆D₆, 298 K): δ 153.9 (d, $^2J_{C-C}$ = 1.8 Hz, CO₃), 54.3 (d, $^2J_{C-C}$ = 1.8 Hz, CH₃O).

Crystal structure determination of complex 3. Single crystal X-ray diffraction data were collected at 100(2) K with graphite-monochromated Mo K α radiation (λ = 0.71072 Å) using narrow frame rotation ($\Delta\omega$ = 0.3°) on a Bruker Smart APEX diffractometer. Measured intensities were integrated and corrected for absorption effects with SAINT⁹ and SADABS¹⁰

programs, included in APEX2 package. The structure was solved with direct methods with SHELXS-2013 (ref. 11) and refined by full-matrix least-squares refinement on F^2 with SHELXL-2018 (ref. 12) program, included in WingX package.¹³ The disordered solvent region has been analyzed with SQUEEZE program.¹⁴

CCDC 1972218 contains the supplementary crystallographic data for this paper.

Crystal data compound **3**. C₃₄H₄₈F₆Ir₂N₄O₁₀S₂Si₄·0.5(H₂O); M = 1364.65; colourless prism 0.140 \times 0.240 \times 0.330 mm³; triclinic $P\bar{1}$, a = 9.0468(5), b = 12.0453(6), c = 12.0619(7) Å, α = 66.4720(10), β = 88.5460(10), γ = 88.7690(10)°, V = 1204.65(11) Å³; Z = 1; D_c = 1.881 g cm⁻³; μ = 5.783 mm⁻¹; T_{\min}/T_{\max} : 0.1676/0.5996; 23 425/5849 reflections measured/unique (R_{int} = 0.0179), number of data/restraint/parameters 5849/0/286, $R_1(F^2)$ = 0.0131 (5771 reflections, $I > 2\sigma(I)$) and $wR(F^2)$ = 0.0324 (all data), final GoF = 1.034, largest difference peak: 0.988 e Å⁻³.

Conclusions

In conclusion, the use of triflate, instead of trifluoroacetate, as ancillary ligand in the chemistry of Ir-(NSi^{Me2})₂ species allows the preparation of **3**, which is a rare example of a dinuclear iridium complex with triflate ligands acting as bridges.

¹H NMR studies of the 3-catalyzed reduction of CO₂ with hydrosilanes evidenced the unprecedented formation of methylsilylcarbonates as reaction products, together with the corresponding silylformate and methoxysilane.

The results of this investigation show that the formation of methoxysilanes during the catalytic reduction of CO₂ with silanes, which traditionally has been explained by the catalytic reaction of bis(silyl)acetals with silanes, could also be consequence of thermal decomposition of the corresponding methylsilylcarbonate.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We gratefully acknowledge MICINN/FEDER projects PGC2018-099383-B-100 and the Regional Government of Aragon/FSE 2014–2020 “Building Europe from Aragón” (group E42_17R) for funding.

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