



Cite this: *J. Mater. Chem. C*, 2016, **4**, 2319

Chemical vapor deposition of ruthenium-based layers by a single-source approach†

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A series of ruthenium complexes of the general type $\text{Ru}(\text{CO})_2(\text{P}(n\text{-Bu})_3)_2(\text{O}_2\text{CR})_2$ (**4a**, R = Me; **4b**, R = Et; **4c**, R = *i*-Pr; **4d**, R = *t*-Bu; **4e**, R = CH_2OCH_3 ; **4f**, R = CF_3 ; **4g**, R = CF_2CF_3) was synthesized by a single-step reaction of $\text{Ru}_3(\text{CO})_{12}$ with $\text{P}(n\text{-Bu})_3$ and the respective carboxylic acid. The molecular structures of **4b**, **4c** and **4e–g** in the solid state are discussed. All ruthenium complexes are stable against air and moisture and possess low melting points. The physical properties including the vapor pressure can be adjusted by modification of the carboxylate ligands. The chemical vapor deposition of ruthenium precursors **4a–f** was carried out in a vertical cold-wall CVD reactor at substrate temperatures between 350 and 400 °C in a nitrogen atmosphere. These experiments show that all precursors are well suited for the deposition of phosphorus-doped ruthenium layers without addition of any reactive gas or an additional phosphorus source. In the films, phosphorus contents between 11 and 16 mol% were determined by XPS analysis. The obtained layers possess thicknesses between 25 and 65 nm and are highly conformal and dense as proven by SEM and AFM studies.

Received 23rd November 2015,
Accepted 22nd February 2016

DOI: 10.1039/c5tc03930d

www.rsc.org/MaterialsC

Introduction

The ongoing miniaturization of devices in semiconductor industry has introduced new manufacturing and materials challenges.¹ A promising candidate which can replace current materials in many microelectronic applications in future technology nodes is ruthenium, as it possesses a high thermal and chemical stability, a low electrical resistance and a negligible solid solubility with copper.^{1,2} Thin films of ruthenium are of particular interest as diffusion barriers for seedless copper interconnects in the manufacturing of integrated circuits by the damascene process.^{3–7} Thereby, one key destination is the development of a single material liner for future copper interconnects. However, pure ruthenium layers cannot act as sole

diffusion barriers for copper.^{8–10} Due to its high surface energy, ruthenium follows a 3D (Volmer–Weber) growth mechanism and hence leads to the formation of polycrystalline, columnar films.¹¹ These polycrystalline structures allow copper diffusion at unacceptably low temperatures through the grain boundaries.^{12,13} For this reason the need for the development of nanocrystalline or amorphous ruthenium-based films arises, *e.g.* obtained by incorporation of phosphorus.^{14,15} Such layers have shown to provide better copper diffusion barrier properties than pure polycrystalline ruthenium coatings.¹⁶

To date, only few approaches for the deposition of Ru(P) layers by chemical vapor deposition (CVD) or atomic layer deposition (ALD) have been described.^{11,16–19} Most commonly, $\text{Ru}_3(\text{CO})_{12}$ and a phosphine PR_3 (R = H, Me, Ph) are deposited simultaneously forming a Ru(P) layer.^{15–18} In this dual-source approach, the films contain 10–50 mol% C with decreasing C contents in the order $\text{PPh}_3 > \text{PMe}_3 > \text{PH}_3$.^{13,15,17,18} Furthermore, precise control and reproducibility of the film stoichiometry are difficult to achieve, because of the different vapor pressures and reactivities of both components.¹⁸ Much better reproducibilities are obtained in the single-source approach that uses precursors containing both elements in the same molecule. Moreover, a better homogeneity is provided as the desired elements are premixed at the molecular level.²⁰ The only single-source precursor that has been examined in the deposition of Ru(P) films so far is air-sensitive $\text{RuH}_2(\text{PMe}_3)_4$.^{11,16} Therein, the undesired incorporation of C still results in a significant increase of the film resistivity

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† Electronic supplementary information (ESI) available: Additional SEM, EDX, XPS and XRPD data. Crystallographic data of **4b**, **c** and **4e**, **f**. CCDC 1430280 (**4b**), 1430281 (**4c**), 1430282 (**4e**), 1430283 (**4f**) and 1430284 (**4g**). For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c5tc03930d



and reduces its applicability as a directly plateable diffusion barrier for Cu.¹³ The quality of the films deposited from RuH₂(PMe₃)₄ can be improved by the addition of H₂ as a reactive gas.¹⁵

We herein present the synthesis of ruthenium precursors of the general type Ru(CO)₂(P(*n*-Bu)₃)₂(O₂CR)₂ (R = Me, Et, *i*-Pr, *t*-Bu, CH₂OCH₃, CF₃, CF₂CF₃) and their use as single-source CVD precursors for the preparation of thin and conformal phosphorus-doped ruthenium layers.²¹ The influence of the carboxylate ligands on the thermal behavior and vapor pressure is discussed. The layers obtained were characterized by SEM, EDX, AFM and XPS measurements.

Experimental section

Instruments and materials

All synthesis procedures were performed under an atmosphere of argon with the solvents degassed prior to use. All reagents were obtained from commercial suppliers and used without further purification. For column chromatography, silica with a particle size of 40–60 μm (230–400 mesh (ASTM), Fa. Macherey-Nagel) was used.

NMR spectra were recorded using a Bruker Avance III 500 spectrometer operating at 500.3 MHz for ¹H, 125.7 MHz for ¹³C{¹H} and 202.5 MHz for ³¹P{¹H} in the Fourier transform mode at 298 K. Chemical shifts are reported in δ (ppm) downfield from tetramethylsilane with the solvent as a reference signal (¹H NMR, CHCl₃ δ 7.26; ¹³C{¹H} NMR, CDCl₃ δ 77.16; ³¹P{¹H} NMR, standard external relative to 85% H₃PO₄ δ 0.0). FT-IR spectra were recorded using a Thermo Nicolet IR 200 instrument. The melting points were determined using a Gallenkamp MFB 595 010 M melting point apparatus. Elemental analyses were performed using a Thermo FlashAE 1112 instrument. High-resolution mass spectra were recorded using a Bruker Daltonics microTOF-QII spectrometer using electrospray ionization (ESI⁺).

TG experiments were performed using a Mettler Toledo TGA/DSC1 1100 system with a UMx1 balance. CVD experiments were carried out using a home-built vertical cold-wall CVD reactor with a heater dimension of 20 × 20 mm (BACH Resistor Ceramics GmbH). Heating could be adjusted up to 773 K and was controlled by a Gefran 600 module connected with a Pt100 thermosensor. The carrier gas (N₂) was controlled by MKS type 247 mass flow controllers connected to the reactor by heated copper lines. The CVD system was attached to a rotary vane pump RZ 6 (Vacuubrand). The pressure of the reactor system was controlled by a Vacuubrand vacuum controller (CVC 3000) in combination with an external Pirani vacuum sensor (VSP 3000).

The surface morphology was investigated by field-emission scanning electron microscopy using a ZEISS Supra60 SEM. Cross-sectional SEM investigations were carried out to determine the film thickness. All AFM measurements were accomplished using NanoWizard I and II devices (JPK Instruments AG, Berlin, Germany) under ambient conditions using JPK SPM Control Software provided by the manufacturer. RMS values were determined using JPK Image Processing. The measurements were carried out in tapping mode using silicon AFM tips with a cantilever length

of 125 μm and a spring constant of approximately 42 N m^{−1} (Pointprobe[®] NCH, NanoWorld AG, Neuchâtel, Switzerland). Energy-dispersive X-ray analysis using a Bruker Quantax 400 system attached to a SEM was applied to determine the chemical composition of the films. The composition of the Ru samples was investigated using a PREVAC XPS system. Monochromatic aluminum Kα radiation (1486.6 eV) was provided by a VG Scienta MX 650 X-ray source and a monochromator. The energy distribution of the photoelectrons was measured using a VG Scienta EW3000 XPS/UPS/ARPES analyzer. This analyzer was operated at 200 eV pass energy with a step size of 100 meV and a measurement time of 3.0 s for each data point. Casa XPS 2.3.16 Pre-rel 1.4 software was used for the deconvolution of the XPS peaks. For the calculation of the atomic concentration, Scofield relative sensitivity factors (RSFs) were used. These RSFs were corrected for a monochromator-analyzer angle of 52.55°. For the escape depth correction in Casa XPS, a value of −0.7 was applied. XRPD patterns were collected on a STOE-STADI P diffractometer using Cu Kα (1.5405 Å) radiation and a Ge(111) monochromator.

Diffraction data were collected with an Oxford Gemini S diffractometer using graphite-monochromated Mo Kα radiation (**4b,c,f,g**) (λ = 0.71073 Å) or Cu Kα radiation (**4e**) (λ = 1.54184 Å) at 110 K using oil-coated shock-cooled crystals. The structures were solved by direct methods and refined by full-matrix least-squares procedures on F².^{22,23} All non-hydrogen atoms were refined anisotropically, and a riding model was employed in the refinement of the hydrogen atom positions. Graphics of the molecular structures were created by using SHELXTL²³ and ORTEP.²⁴

Precursor synthesis and characterization

General synthesis procedure for ruthenium complexes 4a–g.

A solution of Ru₃(CO)₁₂ (200 mg, 313 μmol) and P(*n*-Bu)₃ (738 mg, 3.65 mmol) in 4-methylpentan-2-one (18 mL) was refluxed for 8 h. Subsequently, the respective carboxylic acid (2.50 mmol) was added and the solution refluxed for further 6 h. After solvent removal in vacuum the residue was subjected to silica-gel column chromatography (dimension 3 × 20 cm) or purified by recrystallization from *n*-pentane at −78 °C.

Synthesis of Ru(CO)₂(P(*n*-Bu)₃)₂(O₂CCH₃)₂ (4a). The title compound was synthesized according to the general procedure by using acetic acid (150 mg, 2.50 mmol). The raw product was recrystallized from *n*-pentane at −78 °C to give the respective product as a colorless, crystalline solid. Yield: 494 mg (727 μmol, 78%).

M.p. 108 °C; anal. calcd for C₃₀H₆₀O₆P₂Ru: C, 53.00; H, 8.90; found: C, 53.07, H, 8.96; IR data (KBr, ν/cm^{−1}): 2961 m, 2930 m, 2874 w, 2863 w, 2036 vs, 1970 vs, 1621 s, 1467 w, 1460 w, 1420 w, 1368 m, 1321 s, 1214 w, 1183 w, 1092 w, 1055 w, 1009 w, 967 w, 907 w, 895 w, 867 w, 806 w, 774 w, 746 w, 721 w, 683 w, 669 w, 625 w, 602 m; ¹H-NMR (500 MHz, CDCl₃, δ): 0.92 (t, ³J_{HH} = 7.2 Hz, 18H, CH₂CH₃), 1.33–1.50 (m, 24H, PCH₂CH₂, CH₂CH₃), 1.77–1.86 (m, 12H, PCH₂), 1.98 (s, 6H, CH₃); ¹³C-NMR (126 MHz, CDCl₃, δ): 13.8 (s, CH₂CH₃), 23.5 (t, J_{CP} = 12.9 Hz, PCH₂), 23.6 (s, CH₃), 24.5 (t, J_{CP} = 6.5 Hz, CH₂CH₃), 25.1 (s, PCH₂CH₂), 177.0 (s, O₂C), 198.0 (t, J_{CP} = 11.5 Hz, CO); ³¹P-NMR (203 MHz, CDCl₃, δ): 16.9



(s, P(*n*-Bu)₃); HRMS (ESI-TOF, *m/z*): calcd for C₂₈H₅₇O₄P₂Ru: 621.2778; found 621.2828 [M–O₂CCH₃]⁺.

Synthesis of Ru(CO)₂(P(*n*-Bu)₃)₂(O₂CCH₂CH₃)₂ (4b). The title compound was synthesized according to the general procedure by using propionic acid (185 mg, 2.50 mmol). The raw product was subjected to silica-gel column chromatography using a mixture of *n*-hexane/ethyl acetate (ratio 1:2, v:v) as an eluent to give the respective product as a colorless, crystalline solid. Yield: 366 mg (517 μmol, 55%).

M.p. 93 °C; anal. calcd for C₃₂H₆₄O₆P₂Ru: C, 54.30; H, 9.11; found: C, 54.43, H, 9.19; IR data (KBr, ν/cm^{−1}): 2962 m, 2931 m, 2872 m, 2037 vs, 1969 vs, 1607 s, 1465 m, 1424 m, 1373 m, 1340 m, 1303 w, 1270 m, 1212 w, 1094 m, 1056 w, 905 w, 723 w, 604 w; ¹H-NMR (500 MHz, CDCl₃, δ): 0.89 (t, ³J_{HH} = 7.2 Hz, 18H, CH₂CH₃), 1.06 (t, ³J_{HH} = 7.6 Hz, 6H, O₂CCH₂CH₃), 1.30–1.48 (m, 24H, PCH₂CH₂, CH₂CH₃), 1.73–1.81 (m, 12H, PCH₂), 2.22 (q, ³J_{HH} = 7.6 Hz, 2H, O₂CCH₂CH₃); ¹³C-NMR (126 MHz, CDCl₃, δ): 10.9 (s, O₂CCH₂CH₃), 13.7 (s, CH₂CH₃), 23.5 (t, J_{CP} = 12.8 Hz, PCH₂), 24.5 (t, J_{CP} = 6.5 Hz, CH₂CH₃), 25.1 (s, PCH₂CH₂), 30.2 (s, O₂CCH₂CH₃), 179.8 (s, O₂C), 198.0 (t, J_{CP} = 11.7 Hz, CO); ³¹P-NMR (203 MHz, CDCl₃, δ): 16.5 (s, P(*n*-Bu)₃); HRMS (ESI-TOF, *m/z*): calcd for C₂₉H₅₉O₄P₂Ru: 635.2935; found 635.2956 [M–O₂Cet]⁺.

Crystal data for 4b. C₃₂H₆₄O₆P₂Ru, *M*_r = 707.84, monoclinic, *P*2/c, λ = 0.71073 Å, *a* = 8.7593(4) Å, *b* = 13.9460(5) Å, *c* = 15.6865(5) Å, β = 95.930(3)°, *V* = 1905.97(12) Å³, *Z* = 2, ρ_{calcd} = 1.233 g cm^{−3}, μ = 0.531 mm^{−1}, *T* = 110 K, θ range 3.63–26.00°, 8850 reflections collected, 3736 independent reflections (*R*_{int} = 0.0247), *R*₁ = 0.0318, *wR*₂ = 0.0696 (*I* > 2σ(*I*)).

Synthesis of Ru(CO)₂(P(*n*-Bu)₃)₂(O₂CCH(CH₃)₂)₂ (4c). The title compound was synthesized according to the general procedure by using isobutyric acid (220 mg, 2.50 mmol). The raw product was subjected to silica-gel column chromatography using a mixture of *n*-hexane/diethyl ether (ratio 1:1, v:v) as an eluent to give the respective product as a colorless, crystalline solid. Yield: 626 mg (851 μmol, 91%).

M.p. 74 °C; anal. calcd (%) for C₃₄H₆₈O₆P₂Ru: C, 55.49; H, 9.31; found: C, 55.51, H, 9.55; IR data (KBr, ν/cm^{−1}): 2958 m, 2931 m, 2871 m, 2036 vs, 1969 vs, 1600 s, 1465 m, 1322 m, 1386 m, 1360 w, 1339 s, 1302 w, 1271 m, 1209 w, 1092 m, 1055 w, 906 w, 892 w, 802 w, 762 w, 725 w, 604 m; ¹H-NMR (500 MHz, CDCl₃, δ): 0.90 (t, ³J_{HH} = 7.2 Hz, 18H, CH₂CH₃), 1.11 (d, ³J_{HH} = 7.0 Hz, 12H, CH(CH₃)₂), 1.31–1.50 (m, 24H, PCH₂CH₂, CH₂CH₃), 1.72–1.84 (m, 12H, PCH₂), 2.43 (sep, ³J_{HH} = 7.0 Hz, 2H, CH(CH₃)₂); ¹³C-NMR (126 MHz, CDCl₃, δ): 13.8 (s, CH₂CH₃), 20.5 (s, CH(CH₃)₂), 23.5 (t, J_{CP} = 12.8 Hz, PCH₂), 24.5 (t, J_{CP} = 6.5 Hz, CH₂CH₃), 25.3 (s, PCH₂CH₂), 36.1 (s, CH(CH₃)₂), 182.2 (s, O₂C), 198.1 (t, J_{CP} = 11.6 Hz, CO); ³¹P-NMR (203 MHz, CDCl₃, δ): 15.5 (s, P(*n*-Bu)₃); HRMS (ESI-TOF, *m/z*): calcd for C₃₄H₆₈O₆P₂Ru + Na: 759.3432; found 759.3380 [M + Na]⁺.

Crystal data for 4c. C₃₄H₆₈O₆P₂Ru, *M*_r = 735.89, monoclinic, *I*2/a, λ = 0.71073 Å, *a* = 22.4889(8) Å, *b* = 9.0891(4) Å, *c* = 38.7408(12) Å, β = 102.185(3)°, *V* = 7740.4(5) Å³, *Z* = 8, ρ_{calcd} = 1.263 g cm^{−3}, μ = 0.526 mm^{−1}, *T* = 110 K, θ range 3.03–25.00°, 20 311 reflections collected, 6793 independent reflections (*R*_{int} = 0.0271), *R*₁ = 0.0272, *wR*₂ = 0.0625 (*I* > 2σ(*I*)).

Synthesis of Ru(CO)₂(P(*n*-Bu)₃)₂(O₂CC(CH₃)₃)₂ (4d). The title compound was synthesized according to the general procedure by using pivalic acid (256 mg, 2.50 mmol). The raw product was subjected to silica-gel column chromatography using a mixture of *n*-hexane/diethyl ether (ratio 2:1, v:v) as an eluent to give the respective product as a colorless, crystalline solid. Yield: 493 mg (645 μmol, 69%).

M.p. 48 °C; anal. calcd (%) for C₃₆H₇₂O₆P₂Ru: C, 56.60; H, 9.50; found: C, 56.83; H, 9.61; IR data (KBr, ν/cm^{−1}): 2960 m, 2930 m, 2872 m, 2864 sh, 2039 vs, 1975 vs, 1606 s, 1565 w, 1478 m, 1466 m, 1461 m, 1420 w, 1390 m, 1333 s, 1213 m, 1093 w, 1048 w, 1007 w, 968 w, 911 w, 888 w, 799 w, 785 w, 727 w, 602 m; ¹H-NMR (500 MHz, CDCl₃, δ): 0.91 (t, ³J_{HH} = 7.1 Hz, 18H, CH₂CH₃), 1.14 (s, 18H, C(CH₃)₃), 1.32–1.49 (m, 24H, PCH₂CH₂, CH₂CH₃), 1.76–1.87 (m, 12H, PCH₂); ¹³C-NMR (126 MHz, CDCl₃, δ): 13.8 (s, CH₂CH₃), 23.3 (t, J_{CP} = 12.7 Hz, PCH₂), 24.5 (t, J_{CP} = 6.4 Hz, CH₂CH₃), 25.4 (s, PCH₂CH₂), 28.9 (s, C(CH₃)₃), 39.5 (s, C(CH₃)₃), 183.5 (s, O₂C), 198.2 (t, J_{CP} = 11.7 Hz, CO); ³¹P-NMR (203 MHz, CDCl₃, δ): 14.7 (s, P(*n*-Bu)₃); HRMS (ESI-TOF, *m/z*): calcd for C₃₁H₆₃O₄P₂Ru: 663.3248; found 663.3376 [M–O₂C^tBu]⁺.

Synthesis of Ru(CO)₂(P(*n*-Bu)₃)₂(O₂CCH₂OCH₃)₂ (4e). The title compound was synthesized according to the general procedure by using 2-methoxyacetic acid (225 mg, 2.50 mmol). The raw product was subjected to silica-gel column chromatography using a mixture of *n*-hexane/ethyl acetate (ratio 1:2, v:v) as an eluent to give the respective product as a colorless, crystalline solid. Yield: 578 mg (781 μmol, 83%).

M.p. 96 °C; anal. calcd (%) for C₃₂H₆₄O₈P₂Ru: C, 51.95; H, 8.72; found: C, 52.13; H, 9.14; IR data (KBr, ν/cm^{−1}): 2959 m, 2930 m, 2871 m, 2815 w, 2037 vs, 1969 vs, 1645 s, 1615 m, 1467 w, 1422 w, 1397 w, 1376 m, 1334 w, 1293 m, 1277 w, 1213 w, 1196 w, 1128 s, 1093 w, 1055 w, 948 w, 904 w, 797 w, 776 w, 723 w, 605 w, 594 w; ¹H-NMR (500 MHz, CDCl₃, δ): 0.92 (t, ³J_{HH} = 7.1 Hz, 18H, CH₂CH₃), 1.33–1.49 (m, 24H, PCH₂CH₂, CH₂CH₃), 1.79–1.87 (m, 12H, PCH₂), 3.41 (s, 6H, OCH₃), 3.89 (s, 4H, CH₂O); ¹³C-NMR (126 MHz, CDCl₃, δ): 13.8 (s, CH₂CH₃), 23.6 (t, J_{CP} = 13.0 Hz, PCH₂), 24.5 (t, J_{CP} = 6.6 Hz, CH₂CH₃), 25.2 (s, PCH₂CH₂), 59.0 (s, OCH₃), 72.1 (s, CH₂O), 175.5 (s, O₂C), 197.5 (t, J_{CP} = 11.6 Hz, CO); ³¹P-NMR (203 MHz, CDCl₃, δ): 16.7 (s); HRMS (ESI-TOF, *m/z*): calcd for C₃₂H₆₄O₈P₂Ru + K: 779.2760; found 779.2831 [M + K]⁺.

Crystal data for 4e. C₃₂H₆₄O₈P₂Ru, *M*_r = 739.84, monoclinic, *C*2/c, λ = 1.54184 Å, *a* = 8.3401(8) Å, *b* = 31.639(3) Å, *c* = 15.0063(16) Å, β = 91.517(11)°, *V* = 3958.4(7) Å³, *Z* = 4, ρ_{calcd} = 1.241 g cm^{−3}, μ = 4.306 mm^{−1}, *T* = 110 K, θ range 4.06–67.22°, 10 925 reflections collected, 3485 independent reflections (*R*_{int} = 0.0431), *R*₁ = 0.0488, *wR*₂ = 0.1303 (*I* > 2σ(*I*)).

Synthesis of Ru(CO)₂(P(*n*-Bu)₃)₂(O₂CCF₃)₂ (4f). The title compound was synthesized according to the general procedure by using trifluoroacetic acid (285 mg, 2.50 mmol). The raw product was subjected to silica-gel column chromatography using a mixture of *n*-hexane/diethyl ether (ratio 9:1, v:v) as an eluent to give the respective product as a colorless, crystalline solid. Yield: 441 mg (560 μmol, 60%).

M.p. 100 °C; anal. calcd (%) for C₃₀H₅₄F₆O₆P₂Ru: C, 45.74; H, 6.91; found: C, 45.76; H, 6.94; IR data (KBr, ν/cm^{−1}): 2966 m,



2934 m, 2878 m, 2052 vs, 1988 vs, 1688 s, 1467 m, 1411 m, 1384 w, 1194 s, 1145 s, 1090 w, 1055 w, 967 w, 906 w, 848 m, 806 w, 792 m, 775 w, 728 m, 600 m; $^1\text{H-NMR}$ (500 MHz, CDCl_3 , δ): 0.92 (t, $^3J_{\text{HH}} = 7.1$ Hz, 18H, CH_2CH_3), 1.33–1.49 (m, 24H, PCH_2CH_2 , CH_2CH_3), 1.77–1.87 (m, 12H, PCH_2); $^{13}\text{C-NMR}$ (126 MHz, CDCl_3 , δ): 13.6 (s, CH_2CH_3), 23.5 (t, $J_{\text{CP}} = 13.2$ Hz, PCH_2), 24.3 (t, $J_{\text{CP}} = 6.8$ Hz, CH_2CH_3), 25.0 (s, PCH_2CH_2), 115.6 (q, $^1J_{\text{CF}} = 290.1$ Hz, CF_3), 162.7 (q, $^2J_{\text{CF}} = 36.9$ Hz, O_2C), 196.4 (t, $J_{\text{CP}} = 11.2$ Hz, CO); $^{31}\text{P-NMR}$ (203 MHz, CDCl_3 , δ): 15.9 (s, $\text{P}(n\text{-Bu})_3$); HRMS (ESI-TOF, m/z): calcd for $\text{C}_{30}\text{H}_{54}\text{F}_6\text{O}_6\text{P}_2\text{Ru} + \text{Na}$: 811.2244; found 811.2187 $[\text{M} + \text{Na}]^+$.

Crystal data for 4f. $\text{C}_{30}\text{H}_{54}\text{F}_6\text{O}_6\text{P}_2\text{Ru}$, $M_r = 787.74$, monoclinic, $P2_1/c$, $\lambda = 0.71073$ Å, $a = 18.0972(4)$ Å, $b = 14.0559(3)$ Å, $c = 15.2263(3)$ Å, $\beta = 94.708(2)^\circ$, $V = 3860.08(14)$ Å³, $Z = 4$, $\rho_{\text{calcd}} = 1.355$ g cm⁻³, $\mu = 0.554$ mm⁻¹, $T = 110$ K, θ range 2.90–25.00°, 19 426 reflections collected, 6781 independent reflections ($R_{\text{int}} = 0.0271$), $R_1 = 0.0366$, $wR_2 = 0.0779$ ($I > 2\sigma(I)$).

Synthesis of $\text{Ru}(\text{CO})_2(\text{P}(n\text{-Bu})_3)_2(\text{O}_2\text{CCF}_3)_2$ (4g). The title compound was synthesized according to the general procedure by using pentafluoropropionic acid (411 mg, 2.50 mmol). The raw product was subjected to silica-gel column chromatography using a mixture of *n*-hexane/diethyl ether (ratio 9:1, v:v) as an eluent to give the respective product as a colorless, crystalline solid. Yield: 713 mg (802 μmol, 86%).

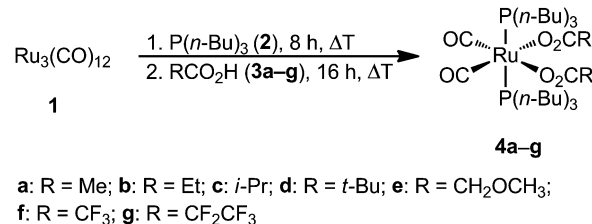
M.p. 93 °C; anal. calcd (%) for $\text{C}_{32}\text{H}_{54}\text{F}_{10}\text{O}_6\text{P}_2\text{Ru}$: C, 43.29; H, 6.13; found: C, 43.26, H, 6.23; IR data (KBr, ν/cm^{-1}): 2962 m, 2937 m, 2876 m, 2052 vs, 1987 vs, 1681 s, 1469 m, 1425 m, 1383 m, 1337 m, 1219 s, 1170 s, 1145 s, 1093 m, 1053 w, 1030 m, 906 w, 821 m, 801 w, 776 w, 736 m, 721 m, 599 m; $^1\text{H-NMR}$ (500 MHz, CDCl_3 , δ): 0.91 (t, $^3J_{\text{HH}} = 7.0$ Hz, 18H, CH_2CH_3), 1.33–1.47 (m, 24H, PCH_2CH_2 , CH_2CH_3), 1.78–1.87 (m, 12H, PCH_2); $^{13}\text{C-NMR}$ (126 MHz, CDCl_3 , δ): 13.6 (s, CH_2CH_3), 23.4 (t, $J_{\text{CP}} = 13.2$ Hz, PCH_2), 24.3 (t, $J_{\text{CP}} = 6.7$ Hz, CH_2CH_3), 25.0 (s, PCH_2CH_2), 106.3 (tq, $^1J_{\text{CF}} = 264.0$ Hz, $^2J_{\text{CF}} = 37.9$ Hz, CF_2), 118.7 (qt, $^1J_{\text{CF}} = 285.5$ Hz, $^2J_{\text{CF}} = 35.1$ Hz, CF_3), 163.1 (t, $^2J_{\text{CF}} = 26.0$ Hz, O_2C), 196.5 (t, $J_{\text{CP}} = 11.2$ Hz, CO); $^{31}\text{P-NMR}$ (203 MHz, CDCl_3 , δ): 15.1 (s, $\text{P}(n\text{-Bu})_3$); HRMS (ESI-TOF, m/z): calcd for $\text{C}_{32}\text{H}_{54}\text{F}_{10}\text{O}_6\text{P}_2\text{Ru} + \text{Na}$: 911.2181; found 911.2090 $[\text{M} + \text{Na}]^+$.

Crystal data for 4g. $\text{C}_{32}\text{H}_{54}\text{F}_{10}\text{O}_6\text{P}_2\text{Ru}$, $M_r = 887.76$, monoclinic, $I2/a$, $\lambda = 0.71073$ Å, $a = 23.1780(5)$ Å, $b = 9.2295(3)$ Å, $c = 38.6832(8)$ Å, $\beta = 102.737(2)^\circ$, $V = 8071.5(4)$ Å³, $Z = 8$, $\rho_{\text{calcd}} = 1.461$ g cm⁻³, $\mu = 0.552$ mm⁻¹, $T = 110$ K, θ range 2.97–25.00°, 35 399 reflections collected, 7072 independent reflections ($R_{\text{int}} = 0.0365$), $R_1 = 0.0307$, $wR_2 = 0.0780$ ($I > 2\sigma(I)$).

Results and discussion

Synthesis and characterization

Ruthenium complexes $\text{Ru}(\text{CO})_2(\text{P}(n\text{-Bu})_3)_2(\text{O}_2\text{CR})_2$ (**4a**, R = Me; **4b**, R = Et; **4c**, R = *i*-Pr; **4d**, R = *t*-Bu; **4e**, R = CH_2OCH_3 ; **4f**, R = CF_3 ; **4g**, R = CF_2CF_3) were prepared by reacting $\text{Ru}_3(\text{CO})_{12}$ (**1**) with $\text{P}(n\text{-Bu})_3$ (**2**) and the respective carboxylic acid (**3**) analogously to a synthetic methodology described by Bianchi²⁵ (Scheme 1). After an appropriate work-up, organometallic compounds **4a–g** could be isolated in good to excellent yields as air and moisture stable colorless solids, which dissolve in most of the common



Scheme 1 Synthesis of ruthenium complexes **4a–g**.

organic solvents including methanol, dichloromethane, toluene, diethyl ether and *n*-hexane.

The identities of all compounds were confirmed by elemental analysis, IR and NMR (^1H , $^{13}\text{C}\{^1\text{H}\}$, $^{31}\text{P}\{^1\text{H}\}$) spectroscopy and high resolution ESI mass spectrometry (Experimental section). The molecular structures of **4b**, **4c** and **4e–g** in the solid state were determined by single crystal X-ray structure analysis. In addition, the thermal behavior was studied by vapor pressure and thermogravimetric measurements.

The $^{31}\text{P}\{^1\text{H}\}$ NMR spectra of ruthenium complexes **4a–g** exhibit one characteristic singlet for the $\text{P}(n\text{-Bu})_3$ groups ranging from 14.7 to 16.7 ppm (Experimental section). As the respective resonance signals are shifted to higher fields as compared to **2** (−30.9 ppm), $^{31}\text{P}\{^1\text{H}\}$ NMR spectroscopy is suitable to monitor the progress of the reactions. The ^1H and $^{13}\text{C}\{^1\text{H}\}$ NMR spectra of **4a–g** are in accordance with the proposed structures. Unique to the $^{13}\text{C}\{^1\text{H}\}$ NMR spectra is the splitting of the α - and β - CH_2 groups of the $\text{P}(n\text{-Bu})_3$ ligand into triplets (Experimental section), which is a common phenomenon for complexes containing *trans*-phosphine ligands.^{26,27} This finding is confirmed by the calculations of Metzinger²⁸ and Harris.²⁹

In the IR spectra of ruthenium complexes **4a–g**, two strong stretching vibrations for the terminal carbonyl groups are observed between 1969 and 2057 cm⁻¹ (Experimental section). The number of CO vibrations reveals that the carbonyl groups have to adopt a *cis*-arrangement in the octahedral coordination sphere of the Ru(II) ion, as for a *trans* isomer just one strong carbonyl stretching is expected.³⁰ From the difference of the characteristic asymmetric ($\bar{\nu}_{\text{asym}}$) and symmetric ($\bar{\nu}_{\text{sym}}$) carboxylate stretching vibrations ($\Delta\bar{\nu}$, $\Delta\bar{\nu} = \bar{\nu}_{\text{asym}} - \bar{\nu}_{\text{sym}}$) one can estimate the structural bonding motif of the carboxylate ligands.³¹ For ruthenium complexes **4a–g** large $\Delta\bar{\nu}$ values of over 250 cm⁻¹ indicate a monodentate coordination of the carboxylate groups to the Ru(II) ion, which was confirmed by single X-ray structure determination (see below).

The structures of **4b,c** and **4e–g** in the solid state were determined by single crystal X-ray diffraction analysis. Suitable crystals were obtained from a concentrated *n*-hexane solution at 5 °C. The ORTEP diagrams of complexes **4b,c,e** are presented in Fig. 1. The molecular structures of complexes **4f,g** along with key structural data can be found in the ESI.† The crystal and structure refinement data are presented in the Experimental section.

All compounds crystallize in monoclinic space groups (**4b**: $P2_1/c$; **4c,h**: $I2/a$; **4e**: $C2/c$; **4f**: $P2_1/c$) with one molecule in the asymmetric unit, except for **4b** and **4e** with one half of the compound and a



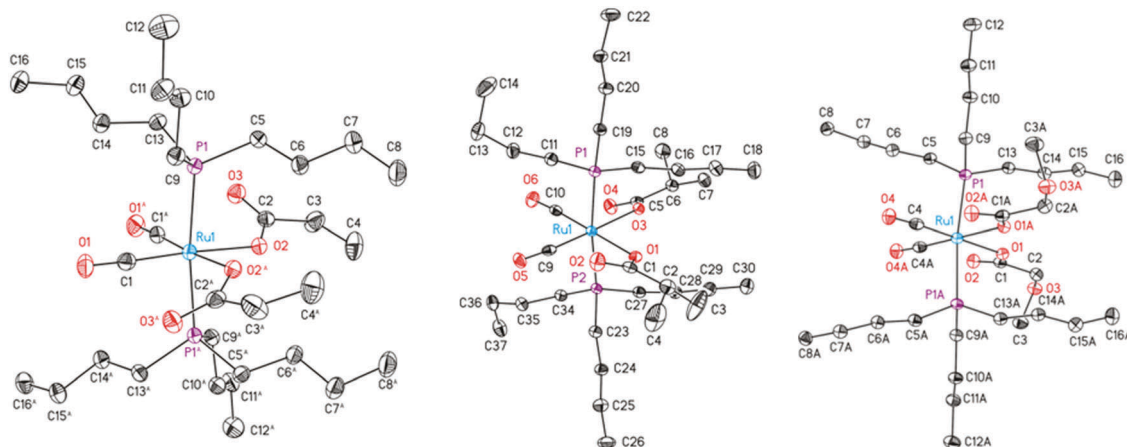


Fig. 1 ORTEP diagrams (30% probability level) of the molecular structures of **4b** (left), **4c** (middle) and **4e** (right, 50% probability level) with the atom numbering scheme. All hydrogen and disordered atoms have been omitted for clarity (symmetry code for **4b**: $A = -x + 1, y, -z + 3/2$; for **4e**: $A = -x + 2, y, -z + 1/2$).

C_2 -symmetry axis through the Ru atom. The complexes consist of a slightly distorted octahedral coordinated ruthenium atom with the two *trans*-positioned tri-*n*-butylphosphines (P1 and P2) in the apical positions, two *cis*-oriented carbonyls and two *cis*-monodentate O-bonded carboxylates in the equatorial plane (Fig. 1).

The tri-*n*-butyl coordinated complexes **4b** and **4c** and **4e–g** exhibit a similar structural behavior, compared to the recently reported triphenyl derivatives,^{26,27} e.g. a slight bending of the phosphines towards the carboxylate moieties (P–Ru–P angles between 171.11(2) in **4g** and 176.23(3)° in **4f**). The CO₂-planes are rotated out of the central coordination plane by 24.1(2) up to 32.9(3)° except for the more coplanar **4e** (10.6(3)°) and all carbonyl oxygens are directed towards the carbonyl substituents. The angles between the carboxylates and carbonyls differ between each derivative (O–Ru–O: 79.27(11) in **4e** to 83.75(6)° in **4g**; C–Ru–C: 87.13(19) in **4e** to 89.94(13)° in **4f**). However, a trend based on electronic or steric effects is not present. The Ru–P bond distances of electronically poor complexes **4f** and **4g** are increased to at least 2.3993(6) Å compared to electron-rich **4b, c, e** in which they do not exceed 2.3910(5) Å. The *n*-butyl moieties of both phosphine substituents reveal a staggered rotation between 171.47(15) (**4f**) and 179.53(15)° (**4b**).

Thermal behavior and vapor pressure measurements

To gain first information on the thermal behavior of the designed precursor complexes, compounds **4a–g** were studied by thermogravimetric (TG) analysis. The physical properties of **4a–g** are summarized in Table 1. The melting points of the complexes could be tuned by changing the carboxylate ligands. In this respect, the melting points decrease with the increasing chain length or branching of the carboxylate ligand from 108 °C in **4a** to 48 °C in **4d**. TG measurements were carried out in a nitrogen carrier gas flow of 60 mL min^{−1}. From the respective TG traces it can be seen that all compounds show a weight loss between 220 and 350 °C which results from an overlapped process of precursor evaporation and decomposition (Fig. 2).

Table 1 Physical properties of Ru precursors **4a–g**

Precursor	M.p. [°C]	Onset temp. [°C]	$\log p$ [bar] = $A - B/T^a$			$\Delta H_{\text{vap}}(\text{exp.})$ [kJ mol ^{−1}]
			A	B	R ²	
4a	108	247	7.93	4918	0.9986	122.4
4b	93	244	8.64	5216	0.9962	107.8
4c	74	254	9.04	5330	0.9782	119.1
4d	48	268	7.42	4602	0.9868	123.7
4e	96	258	9.52	5349	0.9768	95.9
4f	100	277	9.19	5331	0.9777	100.3
4g	93	275	8.54	5135	0.9828	111.1

^a A and B = Antoine parameters; T = absolute temperature; R² = coefficient of determination.

Thereby, the appearance of the TG traces is strongly dependent on the applied heating rate (ESI†). A heating rate of 10 K min^{−1} was employed to ensure that the precursor does not evaporate completely and hence information on the decomposition temperature of the precursor can be received. A lower heating rate leads to the evaporation of the preferred compounds, which is beneficial for the vapor pressure measurements (see below).^{32,33} The TG traces also indicate that marginal changes in the periphery of the complex, for example, the modification of the carboxylate ligands, do not have a significant effect on the evaporation and decomposition process. This observation is also confirmed by the similarity of the respective onset temperatures ranging from 244 °C for **4b** to 277 °C for **4f** (Table 1).

Vapor pressure measurements of ruthenium complexes **4a–g** were carried out over a temperature range from 100 to 240 °C under atmospheric pressure using a TGA system with a horizontal balance. The details of the applied method have been published previously.² All vapor pressure measurements were performed thrice to provide a statistic validation of the experimental data. From the respective vapor pressure traces it can be seen that the different carboxylate ligands influence the vapor pressure of the resulting precursor complex (Fig. 3). The highest volatility was observed for compound **4e**, featuring a CH₂OCH₃ substituent. As expected, the CF₃ groups in **4f** also increase the volatility due



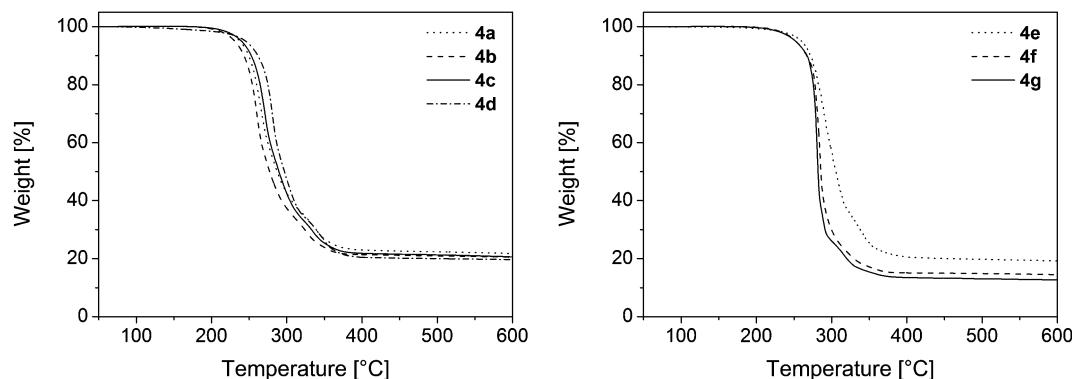


Fig. 2 TG traces of **4a–d** (left) and **4e–g** (right); gas flow, N_2 60 mL min^{-1} , heating rate, 10 K min^{-1} .

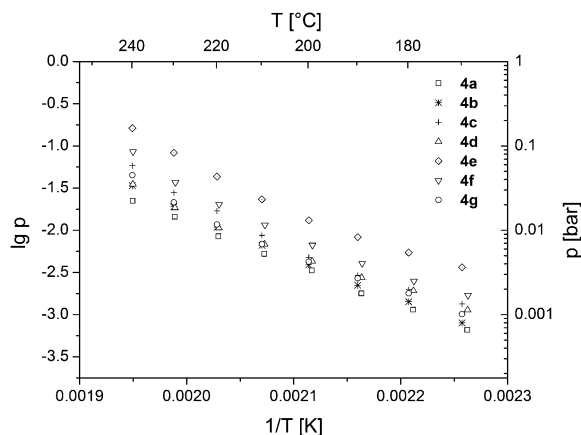


Fig. 3 Vapor pressure traces of ruthenium complexes **4a–g**.

to the low polarizability of the C–F bonds, which leads to reduced intermolecular interactions between the complexes.³⁴ However, the prolonged CF_2CF_3 substitution in **4g** resulted in a decrease of the volatility. The vapor pressures of the remaining compounds **4a–d** hardly differ from each other. In comparison to the previously reported vapor pressures of substituted ruthenocenes and half-open ruthenocenes,² the reported ruthenium complexes **4a–g** possess quite similar volatilities.

Chemical vapor deposition experiments

Ruthenium complexes **4a–f** have been successfully applied in CVD experiments for the deposition of phosphorus-doped ruthenium layers. The experiments were carried out in a home-built vertical cold-wall reactor equipped with a continuous evaporation system. Deposition was performed using nitrogen as a carrier gas (30–50 mL min^{-1} ; 0.6–0.8 mbar working pressure). The ruthenium films were deposited on Si wafers, which were covered by a continuous 100 nm thick thermal SiO_2 layer. Additionally, precursor **4e** was deposited on a trench patterned SiO_2 wafer in order to study the step coverage properties.

For appropriate evaporation of the precursor complexes, temperatures of 120–135 °C in the vaporizer unit were needed. Furthermore, the glass lines were heated to approximately 100 °C. The substrate temperatures were selected according to the results

of the TG analyses. Decomposition of the precursor complexes starts at around 270 °C, but a minimum substrate temperature of 350 °C is required in all cases for complete precursor deposition and formation of ruthenium layers. Therefore, deposition was carried out using substrate temperatures between 350 and 400 °C. The MOCVD deposition parameters of the obtained layers A–H are summarized in Table 2. The deposition was monitored visually and terminated after the formation of metallic layers. Depending on the experimental conditions, 25–65 nm thick uniform layers were obtained. The 40–55 nm thin films are reflective metallic layers, whereas thinner films possess a slightly bluish appearance and thicker films show a yellowish tinge.

Layer characterization

The scanning electron microscopy (SEM) images of the films deposited from **4a–f** evidence the formation of dense and conformal layers for all precursors (Fig. 4). In general, the morphology of the produced layers A–F is very similar, so that no optical differences in surface roughness or homogeneity could be observed. The film thicknesses between 25 and 65 nm were determined by cross-sectional SEM images (ESI†). Thereby, the highest growth rates were obtained for **4e** and **4f**, possessing the highest volatilities according to the vapor pressure measurements (Fig. 3). The deposition rate can be enhanced by higher deposition temperatures and carrier gas flow rates (layers E and F, Table 2).

Table 2 Deposition parameters of Ru layers A–H deposited from **4a–f**

Layer	Compd	$\vartheta_{(Prec.)}$ [°C]	$\vartheta_{(Dep.)}$ [°C]	N_2 flow [mL min^{-1}]	Pressure [mbar]	Dep. time [min]	Layer thickness ^a [nm]
A	4a	130	400	50	0.8	60	40
B	4b	130	400	50	0.8	60	40
C	4c	135	400	50	0.8	60	25
D	4d	125	400	50	0.8	60	25
E	4e	135	400	50	0.8	60	55
F	4f	130	400	50	0.8	60	65
G	4e	130	350	30	0.6	60	25
H ^b	4e	120	400	50	0.8	30	10

^a Determined by cross-sectional SEM images. ^b Deposited on a trench patterned SiO_2 wafer.



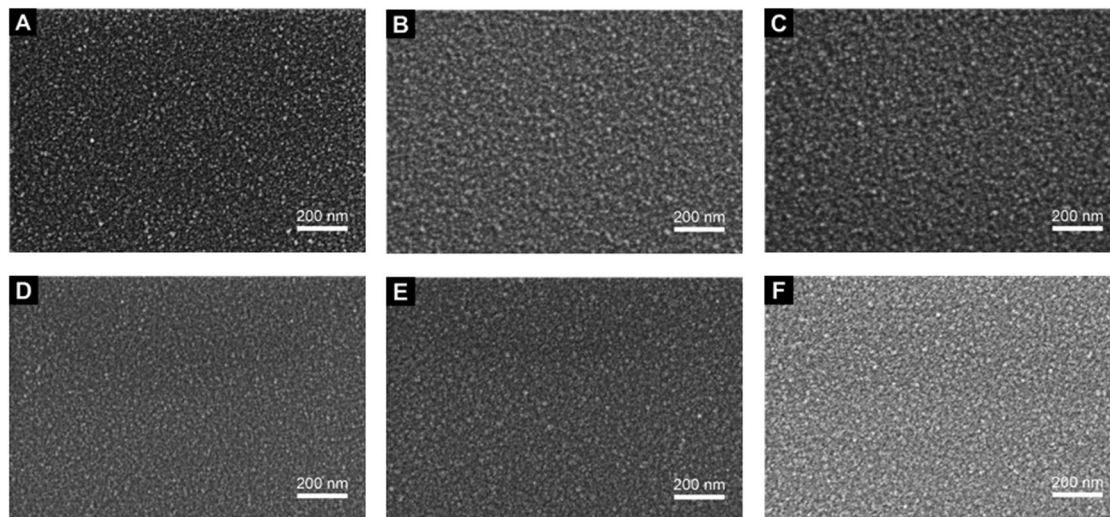


Fig. 4 SEM images (magnification: 80 000 \times) of ruthenium films A–F deposited on SiO₂ using the parameters given in Table 2. A: **4a**, B: **4b**, C: **4c**, D: **4d**, E: **4e**, F: **4f**.

The step coverage property of ruthenium precursor **4e** was investigated by the deposition on a trench patterned Si/SiO₂ wafer with an aspect ratio of 2.5 (ESI[†]).³⁵ The SEM images reveal that a complete and conformal coverage of the patterned substrate was achieved for the deposition of ruthenium complex **4e** (layer H, Table 2).

In order to study the surface roughness, the deposited ruthenium films A–F were examined by atomic force microscopy (AFM). A representative AFM image of a ruthenium film obtained from complex **4e** is depicted in Fig. 5. In comparison, the AFM images taken for all samples show very similar structures with only minor variations in height. In all cases, the resulting layer topography is very homogeneous and characterized by well-interconnected globular grains. However, the consistent bead-like shapes indicate that the actual structures are too small to be accurately imaged using a standard AFM tip. Nevertheless, all RMS (root mean square) roughness values in this study were

found to be in the range of 1.0 nm (Table 3), corresponding to relatively smoother surfaces than previously reported.^{36–40}

The film composition was analyzed by energy-dispersive X-ray spectroscopy (EDX) and X-ray photoelectron spectroscopy (XPS). In EDX spectroscopic measurements, the presence of the characteristic pattern of ruthenium was found in all samples (ESI[†]). In addition, the presence of phosphorus, silicon, oxygen and carbon was detected. Silicon and parts of oxygen originate from the applied Si/SiO₂ substrate. Hence, the intensity of these signals decreases by reducing the energy of the electron beam (penetration depth) and increasing the film thickness.

In order to determine the film composition without penetration of the Si/SiO₂ wafer surface sensitive XPS measurements were carried out. Thereby no silicon could be detected. Instead the presence of ruthenium, phosphorus, carbon and oxygen was confirmed. It is also necessary to note that layer F (Table 2) is fluorine-free as no fluorine could be detected by EDX or XPS analyses. In liner materials, the absence of fluorine is of particular importance as it may penetrate into the oxide layer and hence lead to device damaging.⁴¹

XPS measurements were performed on the surface as well as in the layer after argon ion sputtering (4.0 keV; 330 s) with the intention to remove surface contaminations (Fig. 6). The elemental contributions are summarized in Table 4. Due to the spin-orbit

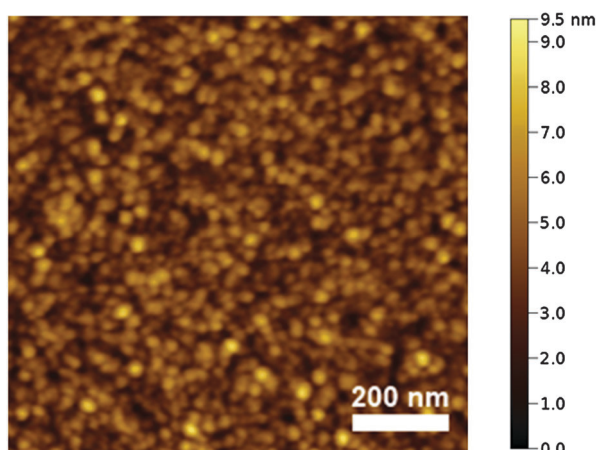


Fig. 5 AFM height image of layer E (Table 2) deposited from **4e**. The RMS roughness value is 0.9 nm.

Table 3 Film properties of Ru layers A–F from Table 2

Layer	Compd	Layer thickness [nm]	rms roughness ^a [nm]
A	4a	45	1.0
B	4b	40	0.9
C	4c	25	0.8
D	4d	30	0.8
E	4e	55	0.9
F	4f	65	0.7

^a Measured by AFM.



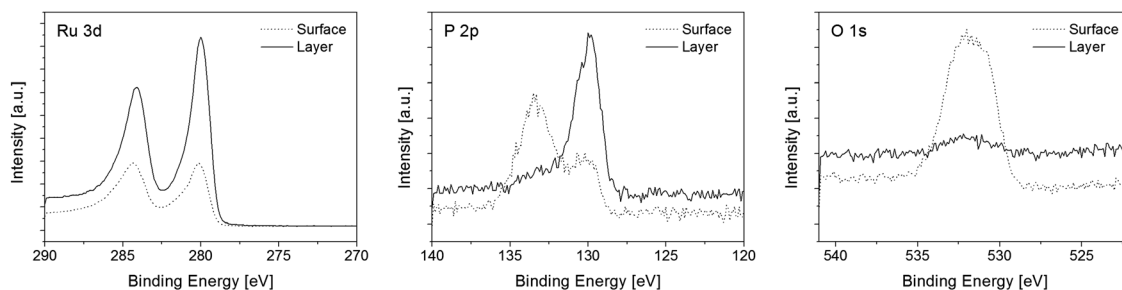


Fig. 6 Detailed XPS spectra of the surface and layer composition of film E; Ru 3d (left), P 2p (middle), O 1s (right).

interaction two peaks for Ru 3d were found (Ru 3d_{3/2} at 284.2 eV and Ru 3d_{5/2} at 280.0 eV), which are in agreement with XPS binding energy databases and correspond to metallic ruthenium.⁴² These ruthenium peaks overlap with the C 1s peak at 284.5 eV,⁴³ preventing the direct measurability of the carbon content.⁴⁴ To determine the carbon content a peak deconvolution was carried out. The applied deconvolution parameters regarding the peak shape and peak position are summarized in the ESI.[†] For all peaks a maximum full width at half maximum (FWHM) of 2.0 eV was allowed. Furthermore, the area ratio of both metallic and non-metallic Ru 3d_{3/2} and Ru 3d_{5/2} peaks was fixed to 2:3 and the peak separation between 3d_{3/2} and 3d_{5/2} to 4.2 eV.⁴⁵ For the identification of peak splitting with respect to P 2p and O 1s a Gaussian–Lorentzian peak shape was applied with a Lorentzian contribution of 25%.

The comparison of the surface and film composition reveals that the ruthenium and phosphorus contents increase in all layers, whereas the carbon and oxygen contents decrease. In general, the layers themselves consist of approximately 48–58 mol% ruthenium, 11–16 mol% phosphorus, 7–31 mol% carbon and 4–29 mol% oxygen (Table 4). The nature of the ruthenium exhibits that beside metallic species ruthenium oxide compounds are also present which can be attributed to RuO₂ (RuO₂ 3d_{5/2} at 281.0 eV).⁴³ The RuO₂ in the surface layer most probably arises mainly from oxygen adsorption and surface oxidation of ruthenium during the transport of the sample from the CVD reactor to the vacuum system for XPS analysis. This estimation is supported by the observation that the oxygen content clearly decreases during sputtering and the films B–F then only contain 4–7 mol% oxygen. Since the deposition was carried out in an inert gas atmosphere, the remaining oxygen most likely originates from the dissociation of the carboxylate or CO ligands.^{46,47} For the phosphorus content

different species in the layers compared to the surface were found (Fig. 6). On the film surface signals, which can be assigned to P(III) of undecomposed phosphine ligands (P 2p at 130.2 eV)⁴² and P(V) of phosphine oxide (P 2p at 133.3 eV)^{48,49} were observed. However, within the layers only one signal was monitored, which refers to elemental phosphorus (P 2p at 129.9 eV).⁴² Due to the small energy difference of P(0) and P(III) species of 0.3 eV, it cannot be excluded that P(III) is present within the layers.⁴² The carbon impurities arise from the catalytic properties of the ruthenium surface, which lead to C–H and C–C activation of the adsorbed ligands resulting in the incorporation of elemental carbon.^{50,51}

To evaluate the crystallinity of the deposited films X-ray powder diffraction (XRPD) measurements were performed (Fig. S6, ESI[†]). The XRPD pattern shows some diffraction arising from the Si/SiO₂ substrate. As no peaks originating from the Ru(P) film could be determined the layers are most probably amorphous.

Conclusion

Ruthenium complexes Ru(CO)₂(P(*n*-Bu)₃)₂(O₂CR)₂ (**4a**, R = Me; **4b**, R = Et; **4c**, R = *i*-Pr; **4d**, R = *t*-Bu; **4e**, R = CH₂OCH₃; **4f**, R = CF₃; **4g**, R = CF₂CF₃) were successfully applied as single-source precursors for the deposition of phosphorus-doped ruthenium layers by the CVD process. The ruthenium compounds were synthesized by a “one-pot” synthetic methodology reacting Ru₃(CO)₁₂ with P(*n*-Bu)₃ and the respective carboxylic acid. All precursors are stable against air or moisture and allow a liquid precursor delivery in a continuous CVD process due to their low melting points. Additional advantages of **4a–g** are that their thermal properties including melting points, vapor pressures and decomposition behaviors can easily be modified by the introduction of different carboxylate ligands. Deposition was carried out in a cold-wall CVD reactor at deposition temperatures between 350 and 400 °C in an inert gas atmosphere without the need of any additional phosphorus source or reactive gas. The highest growth rates were observed for **4e** and **4f**, which also exhibited the highest vapor pressures. All received films were dense and conformal as proven by SEM images and were also uniform in the deposition on patterned wafers with an aspect ratio of 2.5. Furthermore, they possess a very low surface roughness with an RMS value of approximately 1.0 nm as determined by AFM. The elemental composition of the layers was analyzed by EDX and XPS measurements.

In conclusion, ruthenium complexes **4a–f** show promising properties for the fabrication of homogeneous and conformal

Table 4 Elemental contribution of layers A–F from Table 2

Layer	Surface composition [mol%]				Layer composition ^a [mol%]			
	Ru 3d	P 2p	C 1s	O 1s	Ru 3d	P 2p	C 1s	O 1s
A	12.4	9.8	36.3	41.5	48.0	16.1	6.9	29.0
B	15.8	10.0	37.7	36.5	53.9	12.5	29.1	4.5
C	17.1	33.3	38.9	10.7	51.9	12.9	29.6	5.6
D	15.1	9.7	40.5	34.7	49.7	13.5	31.2	5.6
E	17.6	11.0	37.1	34.3	58.1	11.8	25.5	4.6
F	17.1	9.2	46.2	27.5	53.6	10.8	31.5	4.1

^a After argon ion sputtering (330 s, 4.0 keV).



phosphorus-doped ruthenium layers, as all complexes are easy to synthesize, stable in air and do not need a reactive gas or an additional phosphorus source during deposition. The relatively high phosphorus contents of up to 16 mol% make the resulting layers attractive as single material liners for copper interconnects in the damascene process. In the future, the application of phosphine ligands with shorter alkyl chains will lead to significantly decreased carbon contents. Investigations concerning this matter are currently in progress.

Acknowledgements

This work was supported by the Fonds der Chemischen Industrie. J. J. and M. K. thank the Fonds der Chemischen Industrie for Chemiefonds fellowships. We thank Heraeus for the generous gift of Ru₃(CO)₁₂. In addition, we acknowledge Cornelia Kowol and Dirk Rittrich for performing the SEM/EDX measurements, Lutz Mertens and Prof. Michael Mehring for the XRPD analyses and Prof. Robert Magerle for permitting us to perform the AFM studies on his devices.

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