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Hybrid material by anchoring a ruthenium(II) imine complex to SiO₂: preparation, characterization and DFT studies†‡

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Ruthenium–silica hybrid material (RuCl₂(PR₃)₂-2-PyCH-AMPTSi/SiO₂) was prepared and characterized by various spectroscopic techniques. A deconvolution procedure was applied to the spectroscopic data to deconstruct the overlapped bands. A density functional theoretical approach was applied to get insights into the electronic structure of the ruthenium coordination site and the functional RI-PBE-D3/Def2TZVP basis set was used for the optimization. Relativistic effects were considered using the zero-order regular approximation (ZORA). The anchoring process, evinced for each step of the synthesis of the hybrid material, was tracked by FT-IR analyses. The transitions observed in the FT-IR spectra were verified by DFT analyses, which agree with the experimental data. In the DRS-UV-Vis spectra, three main bands were detected by the deconvolution procedure that correspond to the charge transfer transitions, with the main contributions from ruthenium–chlorine and imine–pyridine fragments. TD-DFT results reveal that ruthenium–chlorine antibonding orbitals act as main charge donors, while pyridine–imine is the main charge acceptor.

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Introduction

Hybrid materials, formed from heterogeneous solids mixed with homogeneous fractions, are generating considerable interest in the pharmacology and catalysis fields. The main advantage of hybrid materials is that they can combine the properties of a solid and a homogeneous fraction in a novel material. Furthermore, new properties may be generated; for instance, the interaction of two components affects the electronic properties of the hybrid material if electron transfer is enabled by the interaction.¹

Synthesis of these hybrid materials can be achieved *via* covalent or non-covalent immobilization of either organic or

inorganic compounds on the solid support by encapsulation techniques, absorption, and covalent anchoring.^{2–5} In particular, in the covalent immobilization of a coordination/organometallic complex, the reactivity of the material may be influenced by the functional groups that anchor the compound to the solid as reported by Silva *et al.*⁵ Moreover, the anchored compound provides solubility to the hybrid material, suspending the solid in the solvent, whereas, the solid support contributes to the heterogeneous properties.

Technological uses of hybrid materials are broad. For instance, materials synthesized by non-covalent anchoring of hydrophobic molecules contained in hydrophilic cages are applied in catalysis, drug delivery as reported by Gibb–Bohne *et al.*³ Likewise, covalent anchoring is applied to synthetase catalytic materials. Later, Freire–Pires group⁴ reported a modified Jacobson type catalytic material, with application in alkene asymmetric epoxidation. The modified catalyst was anchored on hexagonal meso-porous amine functionalized solid.

Hybrid materials syntheses by covalent anchoring are reported by two methodologies.^{2–5} The first one involves the anchoring of previously synthesized metal complex containing ligands with functional groups that react with the functionalized support active sites. The second methodology consists of constructing *in situ* the metallic complex, by functionalizing the solid support by a ligand, followed by the formation of the metallic complex.

A key aspect in hybrid materials is the difficulty in the correct determination of appropriate anchoring of the guest with the solid support through a covalent bond by spectroscopic

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† This paper is dedicated to the memory of Prof. Dr Armando Cabrera (July 1944–August 2014) and Mr Elias Mendoza Maca (May 1937–January 2021).

‡ Electronic supplementary information (ESI) available: General experimental details, FT-IR spectra, Raman spectra, XPS analyses, atomic absorption (AA), DRS-UV-Vis spectra and spectra deconvolution, thermogravimetric analysis (TGA), superficial analysis (BET and BJH) and NTOs contour plots. See DOI: 10.1039/d0ra09282g

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techniques guest–solid signal overlap. An issue with those experimental techniques is the guest–solid signal overlap. To overcome this limitation, deconvolutions⁶ are applied to the FT-IR and UV-Vis spectrums.

To get an insight into the nature and properties of the new hybrid material, knowledge of electronic and structural properties of the guest complexes is also desirable. Density Functional Theory (DFT), including exchange and correlation energies, is a suitable approach to study the electronic structure of medium size systems.^{7–12} The dispersion interactions become crucial in molecules with stacking and $\text{CH}\cdots\pi$ interactions, which in general were not accounted by DFT. A widely used approach to alleviate this DFT flaw was proposed by Grimme (DFT-D),⁹ which consists of adding a dispersion term to the KS-DFT total energy. DFT-D approach has an advantage because the computation time to calculate the dispersion correction term is negligible and thus geometry converges smoothly. The DFT-D can be applied for the structural optimization and to calculate the theoretical IR-spectra. Likewise, including dispersion corrections, UV-Vis will be computed using TD-DFT approach. Non-negligible ruthenium relativistic effects were considered by calculations with the ZORA Hamiltonian. The resolution of identity (RI) approximation was used in all the calculations.

In the present work, we are presenting the preparation and characterization of covalently anchored ruthenium complexes with triphenylphosphine to silica (SiO_2) solid surface ($\text{RuCl}_2(\text{-PR}_3)_2(\text{L})/\text{SiO}_2$). The anchoring process was tracked by applying deconvolutions to the FT-IR and UV-Vis spectra. DFT calculation was applied to compute the structures of ruthenium complexes, IR and UV-Vis spectra allowed to gain insight into the experimental observations. With this, we focussed on the purpose of showing experimental and computational techniques for the identification of characteristic bands in a Ruthenium complex, which is covalently anchored *in situ* on the surface that generates overlaps between its signals and those of the compound to be characterized.

Experimental

Solvents and reagents

(3-Aminopropyl)triethoxysilane, 2-pyridinecarboxaldehyde, triphenylphosphine, triphenylphosphite, ruthenium(III) chloride trihydrate, and the solvents used in the modification of silicas (Degussa and MCM-41), and the anchoring of the ruthenium complex were obtained from Sigma-Aldrich.

Functionalization of SiO_2 with 3-aminopropyltriethoxysilane (AMPTSi)

Initially, the Degussa and MCM-41 silicas were functionalized with 3-aminopropyltriethoxysilane (AMPTSi) (separately); 3 g of silica were suspended in toluene (50 mL) and AMPTSi (8.83 mmol, 2.10 mL) was added, and the mixture was heated to reflux for 24 hours. The final material (AMPTSi/ SiO_2) was filtered under vacuum and washed with toluene (3×3 mL). Finally, the solid was dried at 85 °C overnight and the solids AMPTSi/Degussa **1a** (4.85 g) and AMPTSi/MCM-41 **1b** (4.92 g) were obtained.

Activation of AMPTSi/ SiO_2 with 2-pyridinecarboxaldehyde (2-PyCHO)

AMPTSi/ SiO_2 (1.0 g) was mixed with 2-PyCHO (0.3 mL, 3.12 mmol) in ethanol (10.0 mL), the mixture was refluxed with vigorous agitation for 3 h. The active solid was filtered under vacuum and washed with ethanol/diethyl ether (1 : 1) (3×5 mL). The solid 2-PyCH-AMPTSi/ SiO_2 was dried at 50 °C for 1 h. Through this methodology the solids (2-PyCH)AMPTSi/Degussa **2a** (1.17 g) and (2-PyCH)-AMPTSi/MCM-41 **2b** (1.18 g) were obtained.

Anchorage of $\text{RuCl}_2(\text{PR}_3)_2$ to 2-PyCH-AMPTSi/ SiO_2 solid

The complex $[\text{RuCl}_2(\text{PR}_3)_3]$ can be obtained *in situ* by the addition of $\text{RuCl}_3 \cdot 3\text{H}_2\text{O}$ and ligand PR_3 (PR_3 = triphenylphosphine or triphenylphosphite) in the reaction mixture. To solution of ligand PR_3 (triphenylphosphine 1.5 g, 5.7 mmol, or triphenylphosphite 1.8 g, 5.7 mmol) in methanol (15.0 mL) in a Schlenk flask under nitrogen atmosphere, the activated solid (300 mg, **2a** or **2b**) and $\text{RuCl}_3 \cdot 3\text{H}_2\text{O}$ (80.0 mg, 0.3 mmol) were added, and the mixture was refluxed for 4 h. A dark red-wine solid was obtained (Scheme 1), which was filtered under vacuum, even when the solvent was hot, subsequently, the solid was washed with hot methanol (3×5 mL) and diethyl ether (3×5 mL). Finally, the solid $\text{RuCl}_2(\text{PR}_3)_2(2\text{-PyCH})\text{-AMPTSi}/\text{SiO}_2$ was dried at 40 °C for 30 minutes. A red solid of the type $\text{RuCl}_2(\text{PR}_3)_2(2\text{-PyCH})/\text{SiO}_2$ red was obtained.^{13–17} The solids $\text{RuCl}_2(\text{PPh}_3)_2(2\text{-PyCH})/\text{Degussa}$ **3a** (302.7 mg), $\text{RuCl}_2\{\text{P}(\text{OPh})_3\}_2(2\text{-PyCH})/\text{Degussa}$ **3b** (305.5 mg), $\text{RuCl}_2\{\text{P}(\text{OPh})_3\}_2(2\text{-PyCH})/\text{MCM-41}$ **3c** (302.4 mg), and of $\text{RuCl}_2(\text{PPh}_3)_2(2\text{-PyCH})/\text{MCM-41}$ **3d** (301.4 mg) were obtained respectively. In polar solvents, the solids present a uniform dispersion, contrary to apolar solvents such as heptane and toluene generate viscous agglomerates.

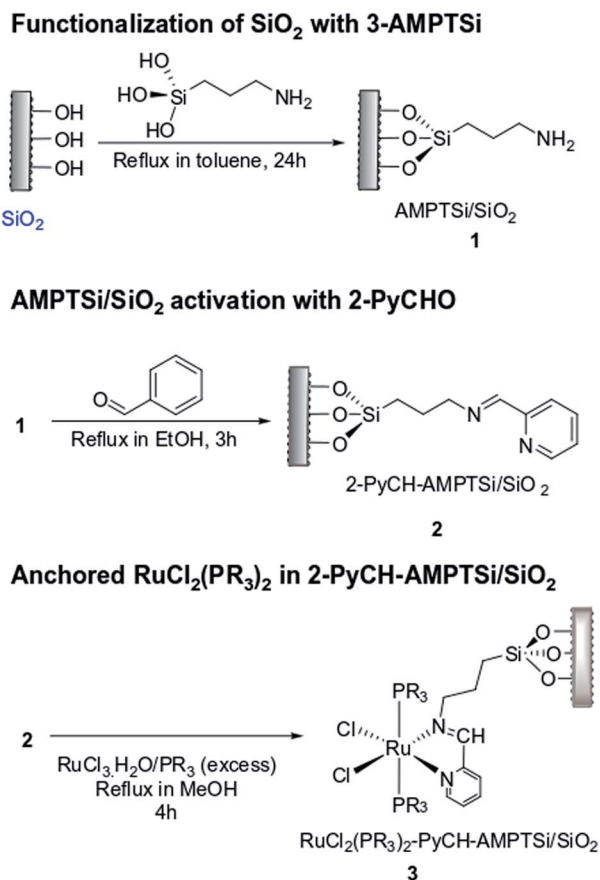
Preparation of the $\text{RuCl}_2(\text{PPh}_3)_2(2\text{-PyCH})\text{-AMPTSi}$ **3e**

In a three-necked round-bottom flask 2-PyCHO (151.22 mg, 1.4 mmol) and 3-AMPTSi (346 μL , 1.45 mmol) in ethanol (10.0 mL) were added, the product (2-PyCH)-AMPTSi was concentrated and washed with dry toluene. The product was dried and characterized by FT-IR.

Subsequently, in a Schlenk flask under nitrogen atmosphere was added $\text{RuCl}_3 \cdot 3\text{H}_2\text{O}$ (12.0 mg, 0.04 mmol) in dry methanol (15 mL). The mixture was heated to reflux for 10 min. Then, it added PPh_3 (51.0 mg, 0.19 mmol) and heated to reflux for 2 h. After, on the hot mixture was added (2-PyCH)-AMPTSi (15.0 mg, 0.04 mmol) and left in reflux for another 3 h. Finally, the product was concentrated under vacuum, was washed with diethyl ether (3×5 mL), toluene (2×3 mL) and ethanol/diethyl ether (3×3 mL) and was dried under a nitrogen gas flux. The compound **3e** (28.7 mg obtained) was characterized for FT-IR.

Instrumentation

A FT-IR Nicolet IR-200 spectrophotometer was used to record the spectra with 32 scans and a resolution of $16 \text{ cm}^{-1} \text{ s}^{-1}$. The RAMAN spectra were obtained using a Raman Perkin Elmer NIR



Scheme 1 Synthetic route for the formation of hybrid materials RuCl₂(PR₃)₂-(2-PyCH) AMPTS/SiO₂

FT-Raman SpectrUM GX spectrophotometer, a small amount of sample (a few milligrams) was deposited on a glass slide. The analyses were carried out according to the determined operating conditions. XPS analyses were performed using VG S-Probe XPS monochromatic spectrometer, with a monochromatic aluminum AlK α (1486.6 eV) anode X-ray source, using 45° take off angle ($q = 45^\circ$) and the source voltage of 10 kV with 200 W power. DRS-UV-Vis spectra were recorded on a UV-Vis-NIR spectrophotometer (Varian-Cary 500). Thermogravimetric analyses (TGA) were performed using the SDT Q600 instrument. Brunauer–Emmett–Teller (BET) and Barrett–Joyner–Halenda (BJH) analyses were obtained on the TRISTAR 3000 instrument. The atomic absorption analysis was performed using an Atomic Absorption Spectrophotometer Solaar Brand 5. The deconvolution process was recorded using Fityk 0.9.8. Software (A curve fitting and data analysis program).

Theoretical methods

Ruthenium complexes were studied through all-electron DFT calculations within the generalized gradient approximation (GGA) of Perdew, Burke and Ernzerhof (PBE).⁷ As the studied compounds are large, the resolution of identity (RI)

approximation was used in all the calculations.⁸ Non-negligible ruthenium relativistic effects were treated efficiently with the ZORA Hamiltonian along with the ZORA-Def2-TZVP basis set and SARC/J auxiliary basis set for the RI approach.^{8,12}

The structural and electronic relaxation procedures were performed without imposing symmetry constraints. The located optimized states were confirmed to be true local minima on the Potential Energy Surface (PES) by estimating the normal vibrations within the harmonic approximation with following the computational protocol. Tight convergence was required for the total energy minimization to 10^{-9} a.u.; while the geometries were optimized with 10^{-7} a.u. and 3×10^{-4} a.u. the thresholds for the root-mean square error of forces and distances, respectively. The structures reported in this work are true minima on the PES, because of the positive frequencies.

Dispersion interactions were included in the calculations for the phenyl group's contribution. A widely used approach to include dispersion on DFT was proposed by Grimme(DFT-D),⁹ it consists of adding a dispersion term to the KS-DFT energy. DFT-D has the advantage that the computation time to calculate the dispersion correction term is negligible and geometry converges smoothly. DFT dispersion correction was applied to optimize the structures and calculate the theoretical IR and UV-Vis spectra.

The ORCA 4.0.1.2 version electronic structure package were used for the geometry optimization,⁷ IR spectra and TD-DFT calculations.

Results and discussion

Amine functionalization of the Degussa and MCM-41 silica

In the IR spectra of 3-aminopropyl triethoxysilane functionalized silica (Degussa and MCM-41), OH- and Si–O vibrations are observed in $3500\text{--}3200\text{ cm}^{-1}$ and $1200\text{--}1000\text{ cm}^{-1}$ regions, respectively (Fig. 1a and b). In the functionalized silica, AMPTS/Degussa (**1a**), new vibrations appeared between $3000\text{--}2800\text{ cm}^{-1}$ and $1750\text{--}1300\text{ cm}^{-1}$ regions (Fig. 1a, spectrum 2) can be attributed to N–H and C–H stretching vibrations, which are verified through a deconvolution process with a symmetrical Gaussian shape centered on the respective FT-IR bands (Fig. 1b), as reported earlier in the literature.^{20–28} Fig. 1c displays H–N–H and H–C–H bending vibrations, visualized as vibrational components in the deconvolution plot at 1568.4 cm^{-1} and 1487.3 cm^{-1} , respectively confirming the activation of Degussa silica. Similar results are reported for the functionalized and activated MCM-41 silica **1b** material (see ESI†).

The computed vibrational signal of propylamine is in very good agreement with the experimental one, well reproducing both the peak frequency position and the bandwidth. The IR spectrum of propylamine was computed to model the observed bands of the FT-IR data.

Covalent attachment of 2-pyridine carboxaldehyde (2-PyCHO) onto the functionalized silica AMPTS/SiO₂

Functionalized silicas **1a** and **1b** were activated *via* covalent attachment with 2-pyridine carboxaldehyde. The infrared

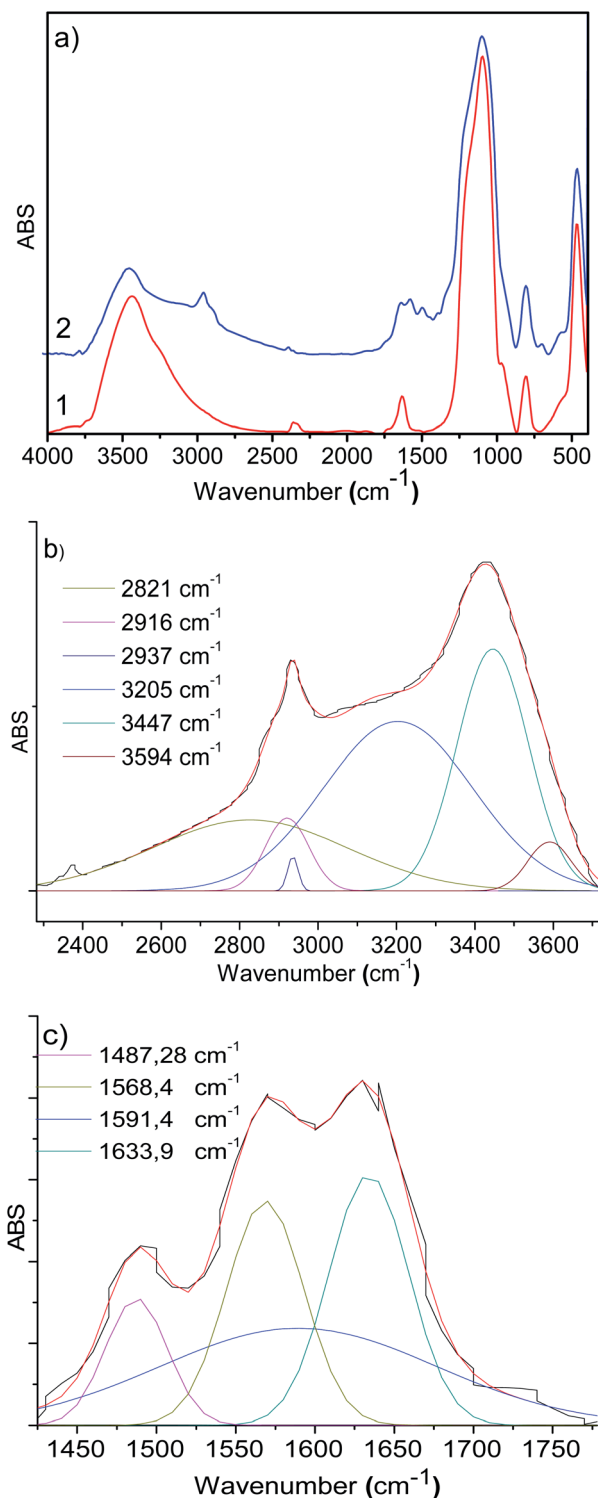


Fig. 1 (a) FT-IR spectra of the (1) Degussa silica, and (2) 3-Amino-propyltriethoxysilane functionalized Degussa silica (**1a**). (b and c) Deconvolution FT-IR spectra plots of the material **1a** in the regions 3700–2300 cm^{-1} (b) and 1750–1300 cm^{-1} (c).

spectra of the 2-PyCH-AMPTSi/Degussa (**2a**) and 2-PyCH-AMPTSi/MCM-41 (**2b**) materials show changes 3300–2700 cm^{-1} (Fig. 2a), which correspond to the 2-PyCH- motif. An expected imine -C=N- signal (**1a** and **1b**) in the FT-IR

region (1390 and 1650 cm^{-1}) is not visible because of the overlapping of the signals of the complex matrix formed by the silica. Nevertheless, vibrational bands at 1652.3 and 1653.9 cm^{-1} are present for **2a** and **2b**, respectively; this was possible by the deconvolution process of the specific band (Fig. 2b and c). These results verify the covalent attachment

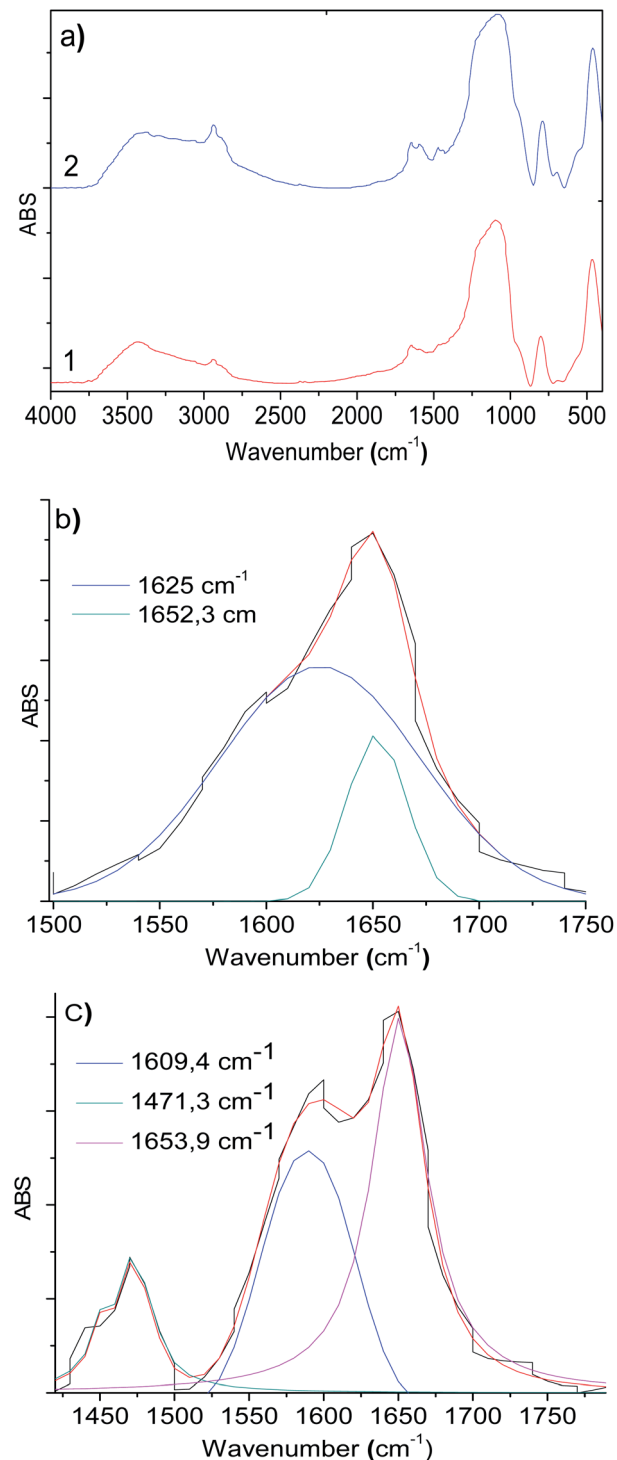


Fig. 2 (a) FT-IR spectra of the (1) material **2a** and (2) material **2b**, (b and c) deconvolution FT-IR spectra plots of the material **2a** (b) and **2b** (c) in the 1750–1500 cm^{-1} and 1750–1300 cm^{-1} regions, respectively.

of the Schiff base ligand to the functionalized silica. Vibrational components, below 1650 cm^{-1} , corresponding to the $-\text{NH}_2$ group may be assigned to the non-reacting 2-PyCHO fraction.

The computed vibrational signal of the pyridyl imine ligand is in very good agreement with the experimental one, well reproducing both the peak frequency position and the bandwidth. The theoretical frequency obtained for the imine bond vibration agrees with the experimental result (1664.1 cm^{-1}). Four C–H stretching vibrations were found for the pyridine group at 3064.6 cm^{-1} , 3095.3 cm^{-1} , 3110.0 cm^{-1} , and 3124.0 cm^{-1} . The lowest energy C–H stretching vibrations were asymmetric, while the last one is symmetric C–H stretching.

In situ coordinate covalent anchoring of Ru complex ($\text{RuCl}_2(\text{PR}_3)_2\text{-2-PyCH-AMPTS}/\text{SiO}_2$) onto functionalized silica

The complex $[\text{RuCl}_2(\text{PR}_3)_2(2\text{-PyCH-AMPTS})]$ was formed *in situ* on activated silica 2-PyCH-AMPTS/ SiO_2 (2). Table 1 shows the percentage of ruthenium anchored to the solids, which is low in concentration, and this correlates to the low increase in weight within the covalent anchoring process seen in the experimental section (302.7 mg of **3a**, 305.5 mg **3b**, 302.4 mg of **3c**, and 301.4 mg of **3d** from 300 mg of the respective solid precursors **2a** or **2b** derivated from mesoporous solids), however, these solid products were evaluated to verify if the physical properties were modified when they were transformed into a hybrid solid.

Then, synthesized materials (**3a–3d**) are of type IV with H1 hysteresis, the surface area, pore volume, pore size, and isotherm (see ESI†) are typical of mesoporous materials with cylindrical geometry opened at the ends of cylinders^{17–19} (Table 1). Furthermore, these materials show initial decomposition between $155.6\text{--}223.4\text{ }^\circ\text{C}$ temperature range, and the TGA analysis show weight losses corresponding to Cl_2 , HCl , N_2 and NO_2 , which indicates the thermal degradation of the covalently anchored complex and the organic compound 2-PyCH-AMPTS anchored on the silica. Due to these decomposition temperatures, despite its low content of ruthenium, this solid could be promising for use in different catalytic processes.

Table 1 Physical characterization of $\text{RuCl}_2(\text{PR}_3)_2\text{-}(2\text{-PyCH})\text{AMPTS}/\text{Silica}$ ($\text{R} = \text{OPh}, \text{Ph}$) hybrid materials^{a,b}

Material	AA Ru%		BJH ^c ($\text{cm}^3\text{ g}^{-1}$)	BJH ^c (\AA)	TGA ^c ($^\circ\text{C}$)
	P/P	BET ^c ($\text{cm}^3\text{ g}^{-1}$)			
3a	0.35	63	0.225	240	223
3b	0.18	45	0.155	288	177
3c	0.16	34	0.078	233	155
3d	0.18	47	0.139	140	198

^a The BET and BJH analysis are presented in the ESI (see ESI). ^b The TGA plots are shown in the ESI (see ESI). ^c The BET ($\text{cm}^3\text{ g}^{-1}$), BJH ($\text{cm}^3\text{ g}^{-1}$), BJH (\AA), and TGA ($^\circ\text{C}$) corresponds to the surface area, pore-volume, pore size, and the initial decomposition temperature respectively.

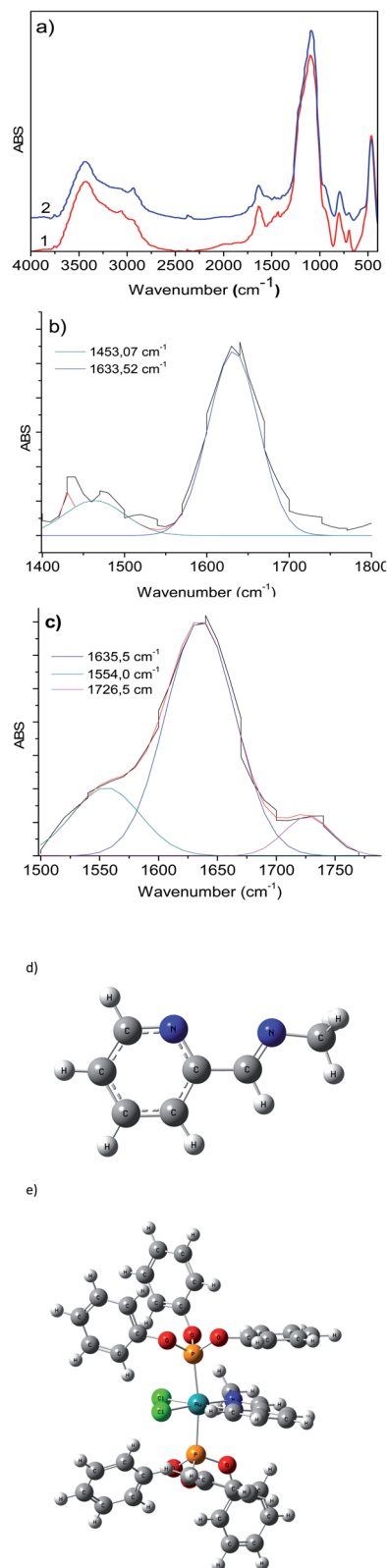


Fig. 3 (a) FT-IR spectra of the (1) material **3a** and (2) material **3d**, and deconvolution FT-IR spectra plots of the materials (b) **3a** and (c) **3d** in the $1700\text{--}1400\text{ cm}^{-1}$ region (d) modeling of the structure of the Schiff base ligand and (e) modeling of $\text{RuCl}_2(\text{PR}_3)_2\text{-2-PyCH-AMPTS}/\text{SiO}_2$ complex.

On the other hand, spectroscopic studies were carried out to verify and study the complex anchored to silica, and due to the difficulty in scrutinizing various spectroscopic signals of the silica and the ruthenium complex (especially in DRS-UV-Vis and FT-IR), because of their overlapping, UV-Vis and IR spectra were studied by employing DFT calculations.

These spectral studies can define key features of the metal-ligand interactions of these hybrid synthesized materials. Other reported studies have also addressed the study of UV-Vis and IR for similar compounds.^{14–39}

The infrared spectra display slight changes in the 1750–1300 cm^{-1} region (Fig. 3a) for $\text{RuCl}_2(\text{PPh}_3)_2(2\text{-PyCH-AMPTSi})/\text{Degussa}$ and $\text{RuCl}_2(\text{PPh}_3)_2(2\text{-PyCH-AMPTSi})/\text{MCM-41}$, **3a** and **3d** materials that correspond to solids, respectively. These changes are most evident through the deconvolution process (Fig. 3b–c), as mentioned earlier by Georgieva *et al.*²⁹ a shift in the FT-IR spectra of the coordinate imine. The coordinate imine bond $-\text{C}=\text{N}-$ vibrations appeared at 1633.5 cm^{-1} and 1635.5 cm^{-1} for **3a** and **3d** materials (Fig. 3b and c), respectively. Whereas, the equivalent deconvolution plot for 2-PyCH-AMPTSi determined an imine vibrational component in 1650.2 cm^{-1} (see in ESI†), which makes evident the shift in the FT-IR from 1650 cm^{-1} to 1635 cm^{-1} , in the free and coordinated ligand. These results are similar for **3b** and **3c** materials (see ESI†).

These results are comparable to those for the free compound $\text{RuCl}_2(\text{PPh}_3)_2(2\text{-PyCH-AMPTSi})$ **3e**. Fig. 4 displays the FT-IR spectrum (the deconvolution plot between 1850–1550 cm^{-1} is presented in the ESI†), where the vibrational band at 1636.8 cm^{-1} corresponds to the coordinated imine vibration, as reported by M. Yáñez *et al.*³⁰

To calculate appropriately the IR shifts originated by imine–ruthenium bond formation, the IR spectra of imine displayed in Fig. 3d was compared to the ruthenium coordination compound (Fig. 3e). Two imine vibrational modes were found in the 1600–1500 cm^{-1} region (1551.8 cm^{-1} and 1504.1 cm^{-1} for $\text{PR}_3 = \text{PPh}_3$, 1577.1 cm^{-1} and 1530.5 cm^{-1} for $\text{PR}_3 = \text{P}(\text{OPh})_3$). The vibrations at 1560–1600 cm^{-1} are due to PPh_3 ,

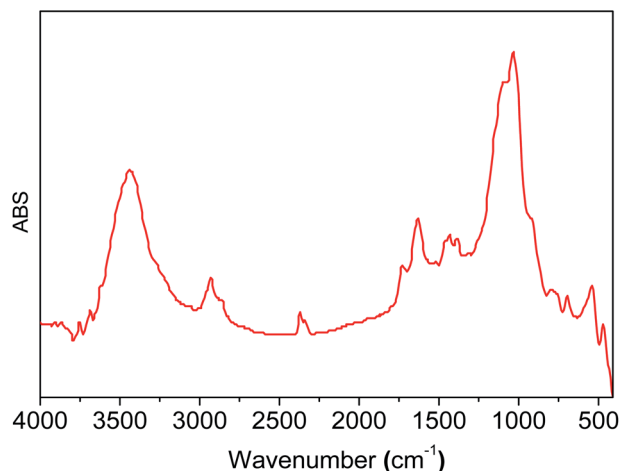


Fig. 4 FT-IR spectrum of the material **3e**.

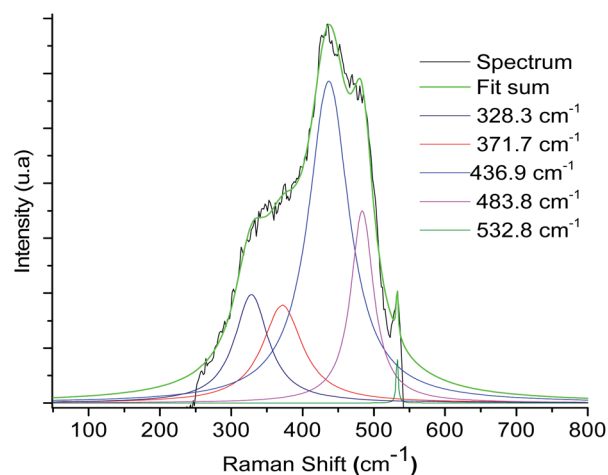


Fig. 5 Raman spectrum deconvolution plot of the material **3a** of the broadband between the 500–300 cm^{-1} regions.

Table 2 Component for bond Ru–N through Raman spectroscopy

Sample	Component ν Ru–N (cm^{-1})
3a	371.7
3b	371.9
3c	375.4
3d	373.1

$\text{P}(\text{OPh})_3$ or pyridine in-plane ring distortions. The calculated imine vibration frequency in the modeled imine is 1670.1 cm^{-1} (Fig. 3d). The IR red shifts in vibration frequencies 118.3–166.0 cm^{-1} and 93–139.6 cm^{-1} are observed for PPh_3 and $\text{P}(\text{OPh})_3$ respectively while the left shoulder of **3b** and **3c** appears because of imine and the highest intensity vibration ($\approx 1635 \text{ cm}^{-1}$) originates from in-plane aromatic ring distortions.

In addition, Raman spectroscopy shows bands at 1520–1540 cm^{-1} (see ESI†) and 371.7 cm^{-1} that correspond to coordinate imine bond and Ru–N vibrations respectively, a similar observation was presented by Chavez-Gil *et al.*³¹ In the solids **3a–3d**, the Ru–N band overlapped with the other bands of the solid matrix, so the deconvolution process over

Table 3 Atomic percent by XPS analysis on the surface of the materials **3b–3d**^a

	Atomic (%)						
	C	Cl	N	O	P	Ru	Si
3a	55.37	1.80	3.33	23.48	2.26	0.93	15.10
3b	36.84	0.72	4.22	34.98	2.87	0.31	20.05
3c	46.19	0.87	4.85	27.02	1.72	0.39	18.97
3d	46.15	1.04	4.65	27.52	0.89	0.39	19.35

^a The XPS plots are in the ESI (see ESI).

Table 4 Cl/Ru, N/Ru, P/Ru and C/Ru ratios from the results obtained in XPS^a

	Cl/Ru		N/Ru		P/Ru		C/Ru	
	Exp	Cal	Exp	Cal	Exp	Cal	Exp	Cal
3a	1.9	2	3.6	2	2.4	2	59.6	45
3b	2.3	2	13.6	2	9.2	2	118.2	45
3c	2.2	2	12.6	2	4.4	2	120.1	45
3d	2.7	2	12.0	2	2.3	2	119.0	45

^a The XPS plots are in the ESI (see ESI).

the broadband was appropriate for this observation (Fig. 5 and Table 2).

A complementary analysis of surface composition of the solids by the X-ray Photoelectron Spectroscopy (XPS) detected 0.93% of ruthenium on the surface of the solid **3a** and around 0.35% for the solids **3b–3d** (Table 3). The C/Ru ratio in **3a–3d** solids is above 59.6 (Table 4), the excess in the C/Ru ratio agrees with the excess in the N/Ru ratio. Carbon and nitrogen excess represent the non-coordinated anchoring ligand and the inactive AMPTSi/SiO₂ (free amine groups on the functionalized silica surface). The Cl/Ru and P/Ru ratios agree with the expected values, except for **3b** and **3c** solids where P(OPh)₃ is used as a ligand. Unlike PPh₃ ligand, P(OPh)₃ ligand is expected to show higher retention on silica; because of its excess during the preparation of the solids and the oxygen present in its structure.

Regarding the Binding Energies (BE) of the metallic ruthenium (Ru⁰), the states Ru 3d_{5/2} and Ru 3d_{3/2} are 280 and 284 eV, respectively.³² For Ru(II) complex with pyridine ligands, the BE (Ru 3d_{5/2}) is between 279.5–282 eV; whereas for the Ru(III) complex, the binding energy is 284 eV.^{33,34} The BE for the Ru 3d_{5/2} state, reported in Table 5, is 279 eV; we also observe the state Ru 3d_{3/2} is between 284.3–284.7 eV (Table 5), characteristic for Ru(II) compounds. Likewise, the states Ru 3p_{3/2} and Ru 3p_{1/2} are visualized in Table 5, which are adjusted to ruthenium complex with an oxidation state +2.³³

According to XPS analyses, a Ru(II) complex has been anchored, so studying the solids through diffuse reflectance

Table 6 DRS-UV-Vis absorption spectrum data^a

	Electronic spectra data			
	λ ₁	λ ₂	λ ₃	λ ₄
3a	<236	275	339	438 (480)
3b	<236	275	330	511 (453)
3c	<250	275	333	525 (439)
3d	236	278	361	494

^a Values in parenthesis are the shoulders in the given band.

UV-Vis spectroscopy (DRS-UV-Vis), a band related to metal–ligand charge transfer was identified for each solid. Table 6 presents the DRS-UV-Vis data (λ_{nm}). The band λ₁ may arise from the π → π* transition of PPh₃ and P(OPh)₃ aromatic rings^{8,35,36} as reported earlier in the literature for similar complexes *e.g.* [RuCl₂(PPh₃)₃] [(NH₃)₄Ru(Py)₂]²⁺ and [RuCl₂(PPh₃)₂(2,6-dmpz)₂].^{8,37–39} The band λ₂ could be assigned to the transition π → π* of the pyridine ring (2-PyCH-AMPTSi) ligand; these bands are allowed by Laporte and spin, elucidating their high intensities. The moderate intensity UV band λ₃ is attributed to an inter ligand transition n → π of the non-activated AMPTSi/SiO₂ excess with 2-PyCHO.

Regarding the visible region, the band λ₄ with shoulder or partially overlapped was detected (Fig. 6 and Table 6). These band intensities are higher than the λ₁ and λ₂ bands. Thus, we assumed visible region bands, allowed by Laporte, correspond to metal–ligand charge transfer (MLCT). Bolaños *et al.* reported the transitions B₁(d_{xz}) → B₁(pπ*) and A₂(d_{xy}) → A₂(pπ*),^{14,35–39} for a similar [RuCl₂(PPh₃)₂(2,6-dmpz)₂] coordination compound,^{8,35} which contains two 2,6-dimethyl pyrazine ligands in a *cis* position arranged in octahedral coordination. Indeed, the solids **3a–3d** are coordinated, in a similar fashion, by a Schiff base derived from pyridine carboxaldehyde (Fig. 7).

Group theory analysis allowed us to give an insight into the possible metal–ligand charge transfer of transitions. Ruthenium coordination compound was assigned with C_{2v} symmetry and hence, considering the first neighbor atoms,

Table 5 Curving fitting data of the XPS spectra in the C 1s, O 1s, N 1s, Si 2p, P 2p, Cl 2p, Ru d and Ru 3p

	C 1s ^a	O 1s	N 1s	Si 2p	P 2p	Cl 2p	Ru 3d _{5/2} ^a	Ru 3d _{3/2}	Ru 3p _{3/2}	Ru 3p _{1/2}
3a	280.8 281.9	528.2	424.4	99.3	127.4	194.2	279.7	284.5	458.3	480.5
3b	281.9 283.1	529.9	397.3	101.1	130.5	195.3	279.41	284.3	458.8	482
3c	282.9 284.7	532.2	399.4	103.0	130.5	197.9	279.4	284.7	461.5	483.5
3d	281.8 282.1	529.4	396.4	100.2	128.2	194.8	279.6	284.6	458.8	481

^a The C 1s and Ru 3d_{5/2} binding energies were differentiated because the intensity in C1's peaks is greater than for the **3d** of the ruthenium because of the low concentration of metal in comparison to the organic material present in the analysis.

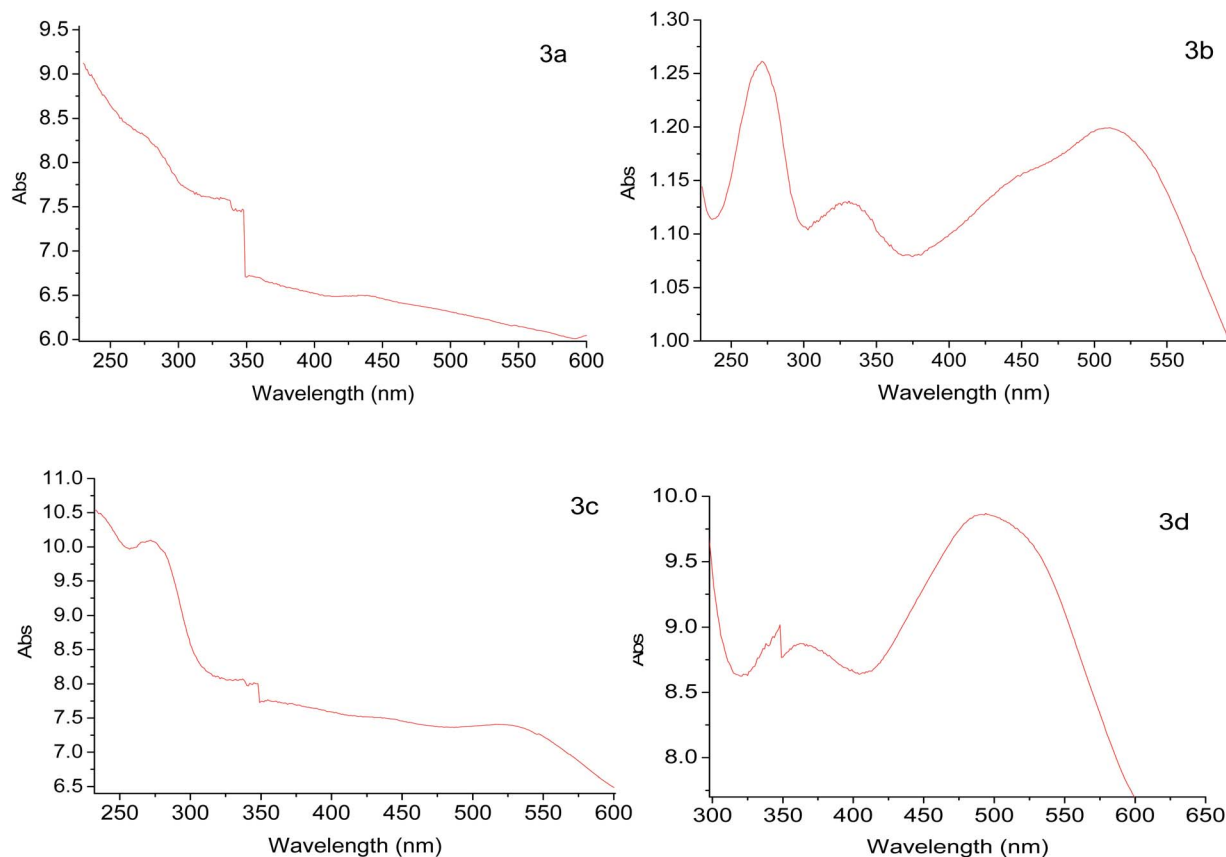


Fig. 6 Electronic spectra of **3a**, **3b**, **3c** and **3d** solids.

involving the π symmetry orbitals of 2-PyCH-AMPTS*i* ligand were transformed into C_{2v} point group. Applying these simplifications, A_2 and B_1 representations were assigned to the pyridine ring and A_1 to the iminic nitrogen, the orbitals $d_{xy}(A_2)$ and $d_{xz}(B_1)$ may be involved in the transitions, because the Terminal Atom Symmetry Orbital $p\pi^*$ (**TASO** $p\pi^*$) are with $B_1(\pi)$ and $A_2(\pi)$ symmetries. Probably the orbital d_{yz} is not involved in the metal–ligand transition, because d_{yz} orbital is B_2 .

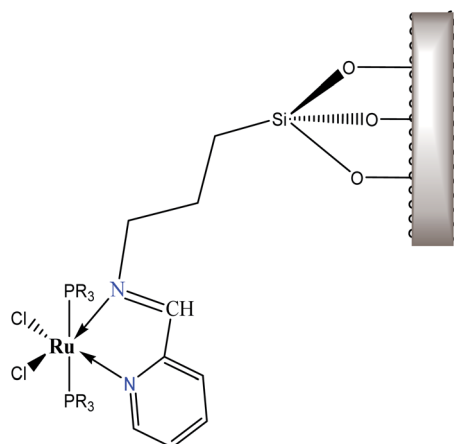


Fig. 7 Covalently anchored Ru(II) coordination compound.

The possible transitions for Ru– N_{py} bond are $B_1(d_{xz}) \rightarrow B_1(p\pi^*)$ and $A_2(d_{xy}) \rightarrow A_2(p\pi^*)$. The former is higher in energy and intensity because **TASO** B_1 is located over the coordinated nitrogen and the pyridine ring carbon atoms. Whereas **TASO** A_2 is near to the pyridine carbons only and is relatively far from the metal center; thus, less overlapping with the metal orbitals. A transition $A_2(d_{xy}) \rightarrow A_2(p\pi^*)$ of Ru– $N_{(Shiff\ base)}$ bond is also expected. This bond should be weaker and a larger bond length is expected in comparison to the Ru–pyridine bond. Therefore, the following three transitions, $B_1(d_{xz}) \rightarrow B_1(p\pi^*) > A_2(d_{xy}) \rightarrow A_2(p\pi^*) > A_2(d_{xy}) \rightarrow A_2(p\pi^*)$. In descending order of energy, are expected in the electronic spectrum of the studied materials (Fig. 8).

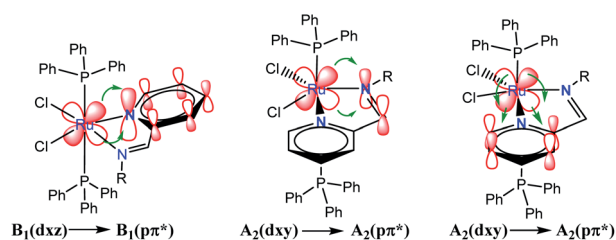


Fig. 8 Electronic transitions of metal to ligand charge transfer (MLCT) in $RuCl_2(PR_3)_2(2-PyCH)-AMPTS*i*/SiO_2$ (Ph and O–Ph ligands were omitted for clarity).

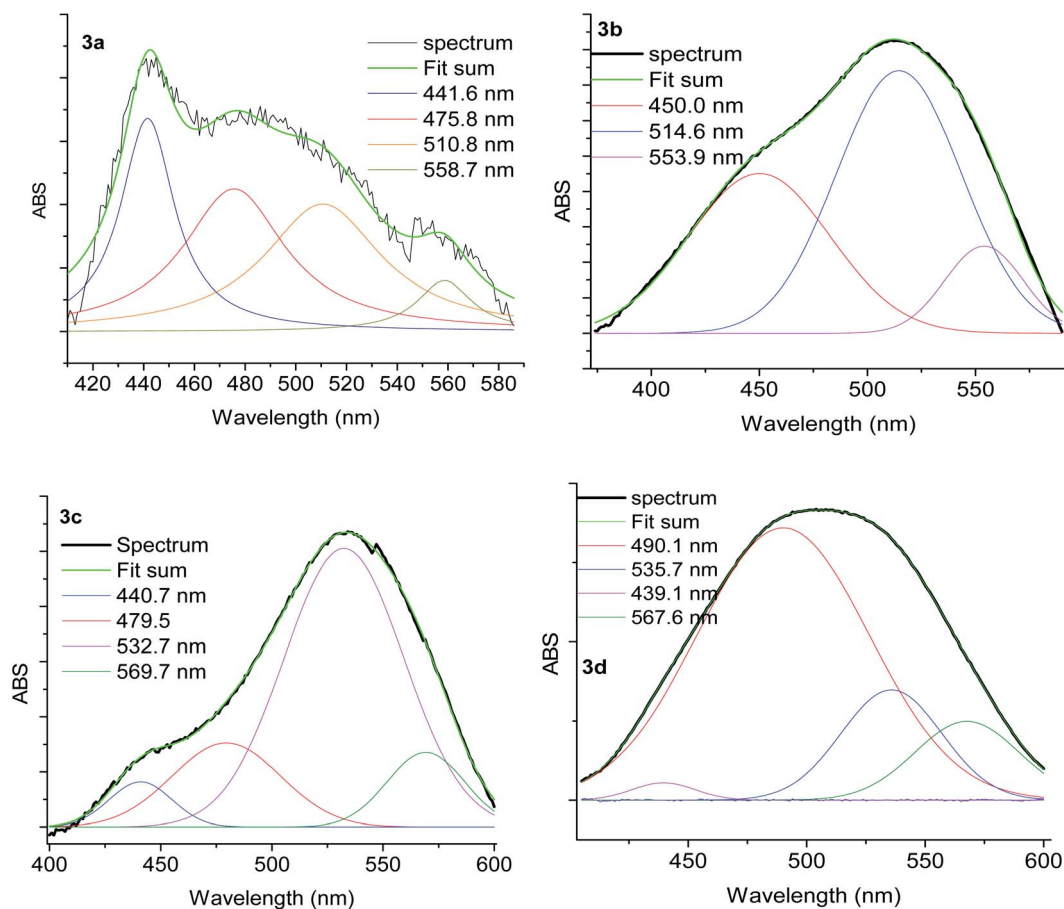


Fig. 9 Deconvolution plot of the solids **3a–3d** of the spectra band in the visible region.

Nevertheless, the observed widespread band in the electronic spectra of **3a–3d** materials suggests that these transitions are overlapped. The components related to the expected transitions were found by applying a deconvolution procedure using the symmetric Gaussian function (Fig. 9). Such components are displayed in Table 7 as λ_a , λ_b and λ_c , which correspond to the $B_1(d_{xz}) \rightarrow B_1(p\pi^*)$, $A_2(d_{xy}) \rightarrow A_2(p\pi^*)$ and $A_2(d_{xy}) \rightarrow A_2(p\pi^*)$ metal–ligand transitions respectively. The transition $A_2(d_{xy}) \rightarrow A_2(p\pi^*)$ is less intense, which may be due to the overlapping of metal ion orbitals on XY axis and the ligand orbital $A_2(p\pi^*)$ (Fig. 8)³ as suggested earlier by Day and Sanders.^{5,35–39} Meanwhile the interaction $B_1(d_{xz}) \rightarrow B_1(p\pi^*)$ is of higher intensity and shorter wavelength because

Table 7 Individual components on the spectra band in the visible region

	λ (nm)		
	$\lambda_a, B_1(d_{xz}) \rightarrow B_1(p\pi^*)$	$\lambda_b, A_2(d_{xy}) \rightarrow A_2(p\pi^*)$	$\lambda_c, A_2(d_{xy}) \rightarrow A_2(p\pi^*)$
3a	442	476–511	558
3b	450	515	554
3c	441–479	533	570
3d	490	536	568

of main back donation, as expected.²⁸ The pattern of the intensities ($\lambda_a > \lambda_b > \lambda_d$) was observed in materials with PPh_3 ligands (**3a** and **3d**), whereas a different pattern in the intensities was detected ($\lambda_b > \lambda_a > \lambda_d$) in the materials with $P(OPh)_3$ ligands. This behaviour could be attributed to lesser donor ability of the $P(OPh)_3$ ligand in comparison to PPh_3 ligand, which affects the back donation $B_1(d_{xz}) \rightarrow B_1(p\pi^*)$. An appropriate orbital combination was allowed to build the qualitative molecular orbital scheme, as illustrated in Fig. 10.³⁵

For further insight into the qualitative analysis of the UV-Vis spectra, the visible range theoretical spectra were also computed by evaluating singlet excited states using TD-DFT at the RI-PBE/Def2TZVP ZORA level of theory in the gas phase. Forty excitations in each complex were evaluated to compare the TD-DFT results with the experimental visible spectra and Natural transition orbitals (NTO) were analyzed. The theoretical visible spectra of $[Ru(II)Cl_2(PR_3)_2(2-PyCH_3)]$ ($PR_3 = PPh_3$ and $P(OPh)_3$) are displayed in Fig. 11. (see ESI[†]).

$RuCl_2(P(Ph)_3)_2(2-Py-CH)=N-CH_3$. The transition at 579.2 nm corresponds to the deconvoluted experimental peaks at 558.7 nm (**3a**) and 567.6 nm (**3d**) ($PR_3 = PPh_3$). This excitation may be represented in terms of two NTO transitions: $HntoS5 \rightarrow LntoS5$ and $HntoS5-LntoS5 \rightarrow LntoS5+1$.

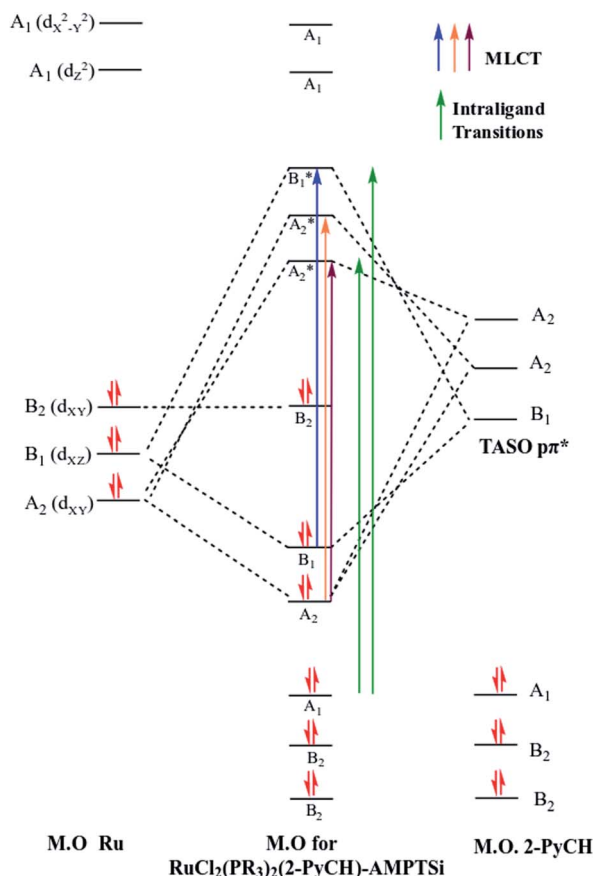


Fig. 10 Simplified molecular orbitals diagram for RuCl₂(PR₃)₂(2-PyCH)-AMPTS/SiO₂ complex.

The former is a transition from ruthenium–phosphine bonding orbital and chlorine atoms to metal–phosphine and pyridine molecules. The second transition is from metal–chlorine antibonding orbital to phosphine molecules (MLLCT) and thus, this transition can be assigned as an MLMLCT excitation. To a lesser extent, S6 state contributes to the lower energy peak, which is attributed to an MLLCT (ruthenium–chlorine to pyridine). The band located in between the highest and lowest energies bands, at 509.8 nm may be assigned to the experimental peak detected at 476–511 (3a) and 536 (3d). Such transition is an MLMLCT from ruthenium–chlorine antibonding orbital to ruthenium–phosphine antibonding orbital. The main contribution to the higher energy band is from state 15, which is composed mainly of three NTOs; these transitions are MLMLCT, where the strongest contribution is from ruthenium–chlorine antibonding orbital to ruthenium–pyridine antibonding orbital.

RuCl₂(P(OPh)₃)₂(2-Py-CH)=N-CH₃. The lowest energy transition of the triphenyl-phosphite coordination compound is MLMLCT between ruthenium–chlorine antibonding orbital and the ruthenium–pyridine orbital. The next two transitions at 554.4 nm and 532.6 nm are from phosphite to ruthenium–pyridine charge transfers (LMLCT). Higher energy bands may be mainly attributed to 488.7 nm which is composed of three

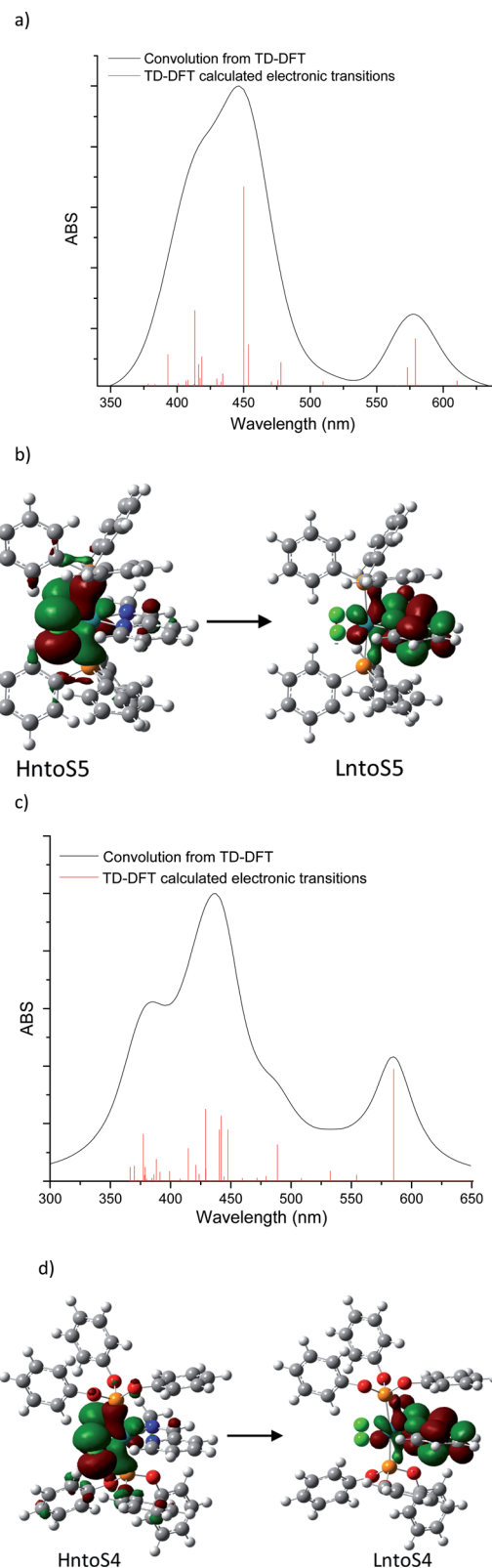


Fig. 11 TD-DFT absorption spectra and natural transition orbitals calculated with RI-PBE-D3/ZORA-Def2TZVP; (a) RuCl₂(PPh₃)₂(2-Py-CH)=N-CH₃ TD-DFT UV-Vis spectra, (b) HntoS5 → LntoS5, λ = 579.2 nm, (c) RuCl₂(P(OPh)₃)₂(2-Py-CH)=N-CH₃ TD-DFT UV-Vis spectra, (d) HntoS4 → LntoS4, λ = 585.1 nm.

NTOs: a metal–chlorine to pyridine (MLLCT), metal–phosphine–chlorine to metal–pyridine (MLMLCT) and metal–chlorine to metal–phosphine (MLMLCT) charge transfers.

Conclusions

The hybrid material $\text{RuCl}_2(\text{PR}_3)_2\text{-2-PyCH-AMPTSi/SiO}_2$ ($\text{PR}_3 = \text{PPh}_3$ or P(OPh)_3) were synthesized *in situ* and characterized by FT-IR, Raman, XPS, and DRS-UV-Vis spectroscopies. Density functional theory, at the RI-PBE/Def2TZVP ZORA level, was used to model the electronic properties at the ruthenium coordination site.

An oxidation state 2+ for the ruthenium was verified by solid characterization. Natural transition orbital analyses provided by TD-DFT calculations suggest that ruthenium 4d orbitals are involved in almost all the transitions. Electrons are mostly transferred to the ruthenium–pyridine–imine from the ruthenium–chlorine antibonding orbitals (MLMLCT). To a lesser extent, P(OPh)_3 and PPh_3 ligands may also contribute to the electronic transitions in the visible region. Notwithstanding the theoretical deviation from the experimentally UV-Vis spectrum determination, the visible region agrees with the intensities hierarchy observed in the experiment. That is, the intensity of the lowest energy is higher with the P(OPh)_3 ligand in comparison to PPh_3 ligand; while the intensity of the highest energy band is lower for the PPh_3 ligand than for P(OPh)_3 ligand.

The changes in the IR spectra observed in the synthesis pathway, allowed to track the anchorage process of the hybrid material. A theoretical approach by DFT confirmed the observed shifts in the FT-IR spectra.

Abbreviations

FT-IR	Fourier-transform infrared spectroscopy
DRS-UV-Vis	UV-Vis diffuse reflectance spectroscopy
XPS	X-ray photoelectron spectroscopy
ZORA	Zero-order regular approximation
AMPTSi	Aminopropyltriethoxysilane
Ru–Npy	bond Ru–N pyridinic
TASO	Terminal atom symmetry orbital

Conflicts of interest

There are no conflicts to declare.

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