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rsc.li/daltonSynthesis, characterisation, and magnetic
properties of a permethylindenyl manganocene†Alexander Evans,^a Thomas A. Q. Arnold,^a Paul Ransom,^a Samantha C. Binding,^a
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The heptamethylindenyl ligand ($C_9Me_7^-$; Ind*) has been used to synthesise the manganese complex, (η^5 -Ind*)₂Mn, by lithiation of the proligand and subsequent salt metathesis with $MnCl_2$. The solid state molecular structure indicates a low-spin ($S = \frac{1}{2}$) ground state for the Mn^{2+} metal centre at 100 K. Solution EPR spectroscopy, solid state magnetic susceptibility measurements, and computational studies suggest the complex undergoes a thermally accessible spin-crossover to a high-spin ($S = 5/2$) state.

Introduction

Following the discovery of the prototypical metallocene complex, ferrocene,^{1–3} the cyclopentadienyl anion (Cp) has become ubiquitous in the isostructural transition metal metallocenes (η^5 -Cp)₂M (e.g.: M = V–Ni),^{4–6} and further existing in complexes exhibiting a range of hapticities.⁷ Cp₂M metallocene sandwich complexes, were later expanded to include s-block (Cp₂Mg)^{8,9} and p-block (Cp₂Sn and Cp₂Pb) metals.^{10–12} More recently, homo- and heterobimetallic sandwich complexes have been reported in the literature from across the periodic table utilising Cp or sterically and electronically tuned derivatives.^{13–16} Additionally, increased steric bulk provided by Cp permethylation, Cp*, yields increased kinetic stability compared to their Cp analogues.¹⁷

The indenyl anion ($C_9H_7^-$; Ind), formally a benzannulated derivative of Cp, displays significantly altered chemistry from Cp, predominantly due to its flexibility in binding modes, and has become an important ligand framework in organometallic chemistry. Transition metal bis(η^5 -Ind) complexes are well documented, with Ind₂Fe and Ind₂Co first reported in 1954;¹⁸ however, it is notable that Ind₂Mn has eluded characterisation as a non-THF adduct. To date η^6 -, η^3 -, η^2 - and η^1 -indenyl transition metal complexes have also been reported^{19,20} in addition to mixed complexation modes with Nesmeyanov *et al.*

reporting the synthesis of a tungsten complex displaying both η^3 - and η^5 -bound Ind ligands to the same metal centre.²¹ An array of substituted manganocenes were reported by Hanusa and co-workers in 2010 and later by Maekawa *et al.*^{22,23} A spectrum of binding modes was reported, but five compounds (Ind^R₂Mn, R = 2-SiMe₃, 1,3-SiMe₃, 1,3-ⁱPr, 1,3-^tBu and 1,3-Cy) displayed η^5 binding. Magnetic and (DFT) studies clearly illustrate that the above species all contain high-spin manganese, mirroring the original discovery concerning Cp₂Mn.

In many cases, indenyl metallocene, and indeed indenyl metallocenophane, complexes, exhibit significantly divergent reactivities compared to their Cp counterparts.²⁴ This is reasoned to be caused by the ability of the indenyl group to undergo η^5, η^3 -haptotropic ring slip while maintaining the aromaticity of the benzene ring. This phenomenon is commonly referred to as the “indenyl effect”.²⁵ The η^5 – η^3 ring slip is formally a two-electron oxidation of the metal, which becomes electrophilic and coordinatively unsaturated, allowing concomitant associative substitution processes not common for 18 electron complexes,²⁶ leading to a several-thousand-fold increase in the rate of ligand substitution.²⁷ This differing reactivity is classically exemplified in the S_N2 (associative) substitution of PPh₃ into the rhodium complex, (η^5 -Ind)Rh(CO)₂, which proceeds 10⁸ times faster than its Cp analogue.²⁸

Similarly to Cp, permethylation confers kinetic stability to permethylindenyl (Ind*) complexes through increased steric bulk. This reduces the propensity for reactive collision, and the positive inductive effect of the methyl groups through $\sigma_{CH} \rightarrow \pi_{CC}^*$ hyperconjugation generally increases the thermodynamic stability.²⁹ The synthesis of the bulky Ind*H ligand precursor from (2*E*)-2-methylbut-2-enoic acid is well established,²⁹ enabling its use in the preparation of Ind* metallocenes (Ind*₂M) for a variety of d- and f-block elements through reactions with alkali metal Ind* salts and metal sources (such

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as $\text{Co}(\text{acac})_2$ or $\text{FeCl}_2 \cdot x\text{THF}$.^{30–36} The electronic properties of the Ind* ligand in Ind^*_2Fe have been studied by D. O'Hare *et al.* – a combined electrochemical and photoelectron spectroscopic approach showed quantitatively the electron donating nature of the Ind* ligand by a reduction in the first ionisation energies and redox potentials compared to Cp^*_2Fe .³⁰

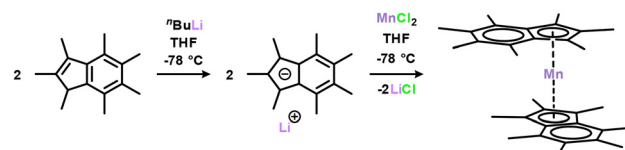
Manganocene, and its methyl-substituted derivatives, have been extensively studied due to their unique property of having two energetically accessible spin states (Fig. 1).^{37–41} These can be readily interconverted, induced by external stimuli such as light, pressure, or temperature, in a phenomenon known as spin-crossover.⁴² It has been found that while Cp_2Mn is a high-spin ($^6\text{A}_{1g}$) species and its permethylated counterpart Cp^*_2Mn occupies an $S = \frac{1}{2}$ ($^2\text{E}_{2g}$) ground state, whereas a singularly methylated Cp complex $\text{Cp}'_2\text{Mn}$ varies the occupation of the two spin states quite significantly with temperature.^{43–48} Work by Walter *et al.* has examined the high temperature spin-crossover behaviours of a variety of manganocenes using both EXAFS and solid-state magnetic measurements,⁴⁹ whilst recent studies have investigated the low-spin d^5 metallocene as a photoredox agent.⁵⁰ Herein, we report the synthesis and characterisation of the manganese sandwich complex, Ind^*_2Mn – the first indenyl-containing manganocene to display thermally induced variation in the Mn^{2+} spin state.

Results and discussion

Synthesis and characterisation of Ind^*_2Mn

Two procedures for the preparation of proligand ($\text{C}_9\text{Me}_7\text{H}$; Ind*H) were used in this work: (1) methylation of hexamethylindene ($\text{C}_9\text{Me}_6\text{H}_2$; Ind*H) by the action of MeI on the lithiated precursor (Ind^*Li), and (2) an improved synthesis following the previously reported route, minimising reaction steps and optimising reagents (ESI†).¹³ In both cases Ind*H was subsequently deprotonated with $n\text{BuLi}$ at -78°C in THF and isolated as a solid powder. Synthesis of Ind^*_2Mn was achieved by adding two equivalents of Ind*Li to a stirred THF solution of MnCl_2 at -78°C , allowing to warm to room temperature overnight (Scheme 1). Work-up afforded Ind^*_2Mn as a purple solid.

The solution phase ^1H NMR spectrum was paramagnetically broadened as expected for a d^5 Mn^{2+} centre (Fig. S4†). On further investigation by variable temperature ^1H NMR spectroscopy (Fig. 2), the manganocene's wing tip methyl (Me_2)



Scheme 1 Synthesis of Ind^*_2Mn .

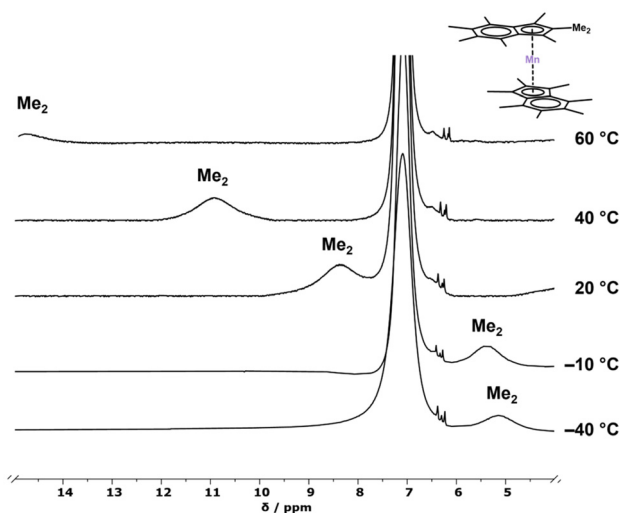


Fig. 2 Variable-temperature ^1H NMR spectra of Ind^*_2Mn in toluene- d_8 solution.

resonance demonstrated a dramatic response to temperature ($\Delta\delta = 9.8$ ppm between 333 and 233 K) whilst other Me resonances varied slightly ($\Delta\delta = 0.18$ ppm) over the same temperature range. The temperature dependence of the chemical shifts did not obey the Curie law ($\delta \propto T^{-1}$) and we were unable to extract meaningful quantitative data (*vide infra*; Fig. S12†). This has been similarly observed by Ashley *et al.* for the paramagnetic bimetallic chromium permethylpentalene complex and Cloke and co-workers for an asymmetric manganese pentalene complex; where variation in ligand field and sterics confound solution phase thermodynamic interpretation.^{51–53}

Single crystals suitable for X-ray diffraction were obtained from a concentrated hexanes solution after cooling to -24°C overnight. The molecular structure is depicted in Fig. 3. The molecule crystallises in the triclinic space group $P\bar{1}$; alternate views are shown in Fig. S7† and relevant bond lengths and angles are given in Table S2.† Scrutinising the molecular structure; the Ind* ligands are nearly coplanar, as shown by the low value of the fold angles about the $\text{C}_6\text{--C}_5$ ring junctions ($1.91(6)^\circ$ and $2.68(6)^\circ$) and a tilt angle = $4.80(5)^\circ$. Crystal structures previously reported for neutral Ind^*_2M complexes of the first row transition-metals are those of Ind^*_2Cr ,³¹ and Ind^*_2Fe .⁵⁴ The expected trend in decreasing average M–C distance is observed across the period ($\text{Cr} = 2.182 \text{ \AA}$, $\text{Mn} = 2.1272(13) \text{ \AA}$, $\text{Fe} = 2.074 \text{ \AA}$). The average Mn–C distance is significantly different from the values observed for all other structurally characterised alkyl manganocenes, Ind^*_2Mn where $\text{R} = 2\text{-SiMe}_3$, 1,3-

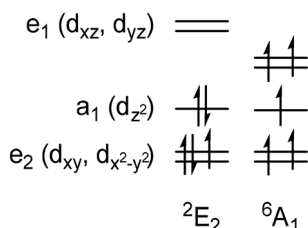


Fig. 1 $\text{Mn } d^5$ ground states in distorted O_h fields.^{37–41}

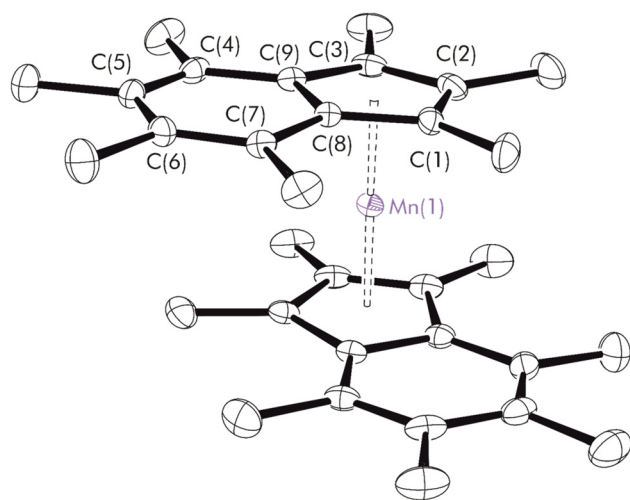


Fig. 3 Molecular structure of Ind*₂Mn. For clarity, hydrogens atoms were omitted (displacement ellipsoids are drawn at 30% probability). Selected distances (Å) and angles (°): Mn–Cp_{cent} 1.7400(6), Δ_{M–C} 0.073, Δ_{M–C} 0.077, RA 85.2(2), C₆–C₅ 1.443(2), C₂₁–C₂₂ 1.441(2). Ring-slip parameter (Δ_{M–C}) defined in eqn (S1).[†] Rotation angle (RA) based on adapted torsion angle, visually defined in Fig. S8.^{†55}

SiMe₃, 1,3-ⁱPr, 1,3-^tBu and 1,3-cyclohexyl (Table 1), where the reported average Mn–C distance is 2.41 Å.^{22,23} Furthermore, the average Mn–Cp_{cent} distance is 1.7400(6) Å for Ind*₂Mn compared to 2.05 Å for the other Ind*₂Mn complexes. By both measures, the Ind*₂Mn distance is 0.3 Å shorter, clearly indicating a major change in the electronic structure.

In fact, the measured Mn–Cp_{cent} distance much more closely matches the value for Cp*₂Mn (1.734 Å), a low-spin manganocene.⁵⁶ A DFT analysis conducted by Maekawa *et al.* on the effect of the spin state of Mn in their bis(indenyl) analogue suggested dramatically different values for the M–Cp_{cent} distance in the low- and high-spin cases, encouraging us to investigate the effect of Ind* computationally.²³ Interestingly, the rotation angle (RA) for Ind*₂Mn is 85.2(2)°, whereas that for the chromium and iron species are 180° and 151° respectively; this value is similar to that found for the cationic cobalt and chromium species, and does not fit previous conclusions that the RA for permethylindenyl complexes fall broadly into two regions: 180° for the neutral species and 90° for the cations.³¹ However, it is in good agreement with the observed and calculated values for

Ind*₂Mn, where R = 1,3-^tBu and 1,3-SiMe₃ are reported as 86.0° and 83.7° respectively.^{22,23} It was noted that the ring-slip parameter values (Δ_{M–C} = 0.073 and 0.077 Å) were within the range of literature values (0.07–0.156 Å).

Electron paramagnetic resonance (EPR) studies

The temperature dependence of the X-band CW-EPR signal for Ind*₂Mn was studied between 4.5–160 K in a toluene glass.

Representative spectra are shown in Fig. 4. At 12 K, the signal is characteristic of low-spin Mn²⁺. As the temperature approaches 100 K, the unresolved outer manifold transitions of high-spin Mn²⁺, $M_S = \pm 5/2 \leftrightarrow M_S = \pm 3/2$ and $M_S = \pm 3/2 \leftrightarrow M_S = \pm 1/2$, are apparent on either side of the sextet of ⁵⁵Mn hyperfine lines at $g = 2$ from the low-spin Mn²⁺ and $M_S = \pm 1/2 \leftrightarrow M_S = \pm 1/2$ of high-spin Mn²⁺. Double integration of the total EPR signal over the temperatures studied reveals an increasing amplitude with temperature as the high-spin state is populated. The low-spin signal was simulated, shown in Fig. S13,[†] providing $g_{||} = 2.56$ and $g_{\perp} = 1.975$, while the ⁵⁵Mn hyperfine simulations were $A_{||}(\text{⁵⁵Mn}) = 206$ MHz and $A_{\perp}(\text{⁵⁵Mn}) = 80$ MHz. The theory of Maki and Berry has a separation of $g_{||}$ and g_{\perp} defined as follows:⁵⁷

$$\begin{aligned} g_{||} &= 2 + 4\kappa'(1 - \zeta^2)/(1 + \zeta^2) \\ g_{\perp} &= 4\zeta/(1 + \zeta^2) \\ \zeta &= (\delta/\xi)/\left(1 + \sqrt{1 + (\delta/\xi)^2}\right) \end{aligned} \quad (1)$$

Here, the observed g -values of $g_{||}$ and g_{\perp} are described with respect to the orbital reduction factor, κ' , effective spin orbit

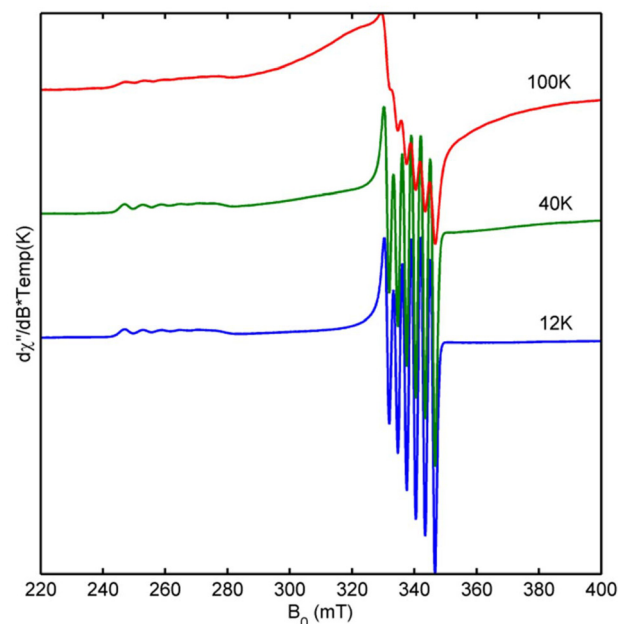


Fig. 4 X-band CW-EPR of the Ind*₂Mn complex at 12, 40 and 100 K. Conditions were a microwave power of 50 W, 100 kHz modulation amplitude of 0.8 mT, microwave frequency of 9.372(4) GHz and a time constant of 20.48 ms.

Table 1 Comparison of relevant parameters for other crystallographically characterised indenyl manganocenes

Ind* ₂ Mn	M–C avg. (Å)	Cp _{cent} –M (Å)	Δ _{M–C} ^a (Å)	RA ^b (°)
R = 1,2,3,4,5,6,7-Me	2.127	1.740	0.075	85.2
R = 2-SiMe ₃ ²²	2.409	2.079	0.14	180
R = 1,3-SiMe ₃ ²²	2.42	2.085	0.12	83.7
R = 1,3- ⁱ Pr ²²	2.4	2.04	0.07	176
R = 1,3- ^t Bu ²³	2.42	2.10	0.156	86.0
R = 1,3-cyclohexyl ²³	2.383	1.96	0.148	91.35

^a Ring-slip parameter (Δ_{M–C}) defined in eqn (S1).[†] ^b Rotation angle (RA) based on adapted torsion angle, visually shown in Fig. S8.^{†55}



coupling parameter, $\xi = \kappa' \xi_0$, and an axial distortion parameter, δ . It was found that this molecule has an axial distortion of $|\delta| = 1700 \text{ cm}^{-1}$, with $\kappa' = 0.89$, using the literature spin orbit coupling of $\xi_0 = 305 \text{ cm}^{-1}$.

According to eqn (1), the Ind* ligand provides a new extreme of an axial distortion, shown in Fig. S8.† Comparison of the g -values of Ind*₂Mn with those of Cp₂Mn, Cp*₂Mn, and Cp' ₂Mn reveals a clear trend.⁴⁸

The g -value of the lowest symmetry ligand, Cp', is most similar to that of Ind*, followed by Cp* and Cp. On a lesser note, samples in frozen solution provide g -values that are closer than samples in magnetically-dilute solids.

Magnetic susceptibility measurements

The solid state magnetic susceptibility of Ind*₂Mn has been measured between 6–300 K (Fig. 5). No difference was observed between the magnetic susceptibility data of a zero-field cooled (ZFC) compared to a field cooled (FC) sample over this temperature range. Attempts to fit the data to the Curie–Weiss law either gave very poor agreement or unrealistic values. For example, a least squares fit to the magnetic susceptibility data between 50 and 200 K gives a Weiss constant, $\theta = -76 \text{ K}$, and an effective magnetic moment, $\mu_{\text{eff}} = 4.18 \mu_{\text{B}}$. Such a large value for the Weiss constant, implied antiferromagnetic behaviour, is not reasonable for a paramagnetic metallocene (Fig. S10†).⁵⁸

In the solid state, Ind*₂Mn undergoes a thermally induced spin-crossover. At 6 K, $\mu_{\text{eff}} = 1.67 \mu_{\text{B}}$ which is close to the value predicted on the basis of the low temperature EPR ($\mu_{\text{eff}} = 1.89 \mu_{\text{B}}$, $S = \frac{1}{2}$, $\langle g \rangle = 2.19$). Between 6 K and 300 K, the effective magnetic moment of the sample steadily increases, reaching a $\mu_{\text{eff}} = 3.85 \mu_{\text{B}}$ at 300 K, which is consistent with a thermally induced partial population of a high-spin, $S = 5/2$ state. Assuming an ideal high-spin excited state with a $\mu_{\text{eff}} = 5.92$, the magnetic moment at 300 K is consistent with a 36% population of this high temperature form.⁴⁷ A derived Arrhenius plot ($\ln K$ vs. T^{-1}) over the studied temperature range enabled reasonable extraction of thermodynamic parameters *via* a

linear fit regression (Fig. S11;† $R^2 = 0.992$, $\Delta H = 0.485 \pm 0.007 \text{ kJ mol}^{-1}$, and $\Delta S = -7.98 \pm 0.17 \text{ J mol}^{-1}$).⁴⁹ Negligible enthalpic contributions in the solid state suggest an entropy driven process towards the thermally induced spin-crossover. Variable-temperature single-crystal X-ray diffraction was also employed to probe the system; all crystal structures exhibit largely the same unit cell parameters, with only minor variations attributed to thermal expansion. Notably, there were no significant changes to the RA, suggesting that any geometry induced spin-state transition does not manifest in the crystal-line phase; however, the complex may behave differently in a toluene glass/solution due to reduced conformational rigidity. Consequently, we believe the magnetometry studies isolate the thermodynamic data associated with purely the spin-crossover phenomena, whilst the NMR spectroscopic data includes a second regime related to the ligand geometry, supported by DFT (*vide infra*), confounding the extraction of thermodynamic parameters.

The solution phase effective magnetic moment was measured at 298 K using the Evans method⁵⁹ and found to be $2.53 \mu_{\text{B}}$ in a range of hydrocarbon solvents. Negligible variation of the moment over the accessible temperature range, 193–333 K, was observed. The existence of a low-spin configuration contrasts all previously reported manganese bis(indenyl) complexes which have high-spin, $S = 5/2$ ground states from 4–300 K: (Ind^R₂)₂Mn (R = ^tPr, ^tBu, or Cy), $\mu_{\text{eff}} = 5.25\text{--}5.9 \mu_{\text{B}}$.^{4,5}

The magnetic susceptibility behaviour of Ind*₂Mn reflects the trend observed with bis(indenyl)chromium(II) complexes studied by Hanusa, Yee and co-workers, where the staggered and eclipsed conformations favour high-spin ground states, and the *gauche* conformations are generally associated with low-spin complexes.⁶⁰ In addition, they also report a general correlation between increasing methylation and a propensity for low-spin ground states; this trend is mirrored in magnetic studies on the manganocenes Cp₂Mn, Cp*₂Mn, and Cp' ₂Mn.^{43–48} Extrapolating from these observations, Ind*₂Mn would be expected to favour a low-spin configuration.

Computational studies

Magnetometry measurements at 300 K indicate that 36% of the complex exists in the high-spin state.⁴⁷ At 130 K, EPR spectroscopy reveals the loss of hyperfine splitting, suggesting the presence of a high-spin complex. This observation is consistent with the expectation that a significant high-spin population would overwhelm the signal from the low-spin state and may likewise explain the lack of variation of the effective magnetic moment in solution using the Evans method. Given the discrepancies between observed spin-crossover behaviours in the NMR and EPR spectroscopic, and SQUID magnetometric measurements, the complex was computationally investigated in order to probe the ligand influence on the Mn spin-state.

Accurate computational prediction of relative spin state energy levels for metal complexes are notoriously difficult to obtain.^{61,62} Screening a variety of DFT functionals from the crystallographically-defined coordinates of Ind*₂Mn yielded a wide range of electronic energy difference between the low-

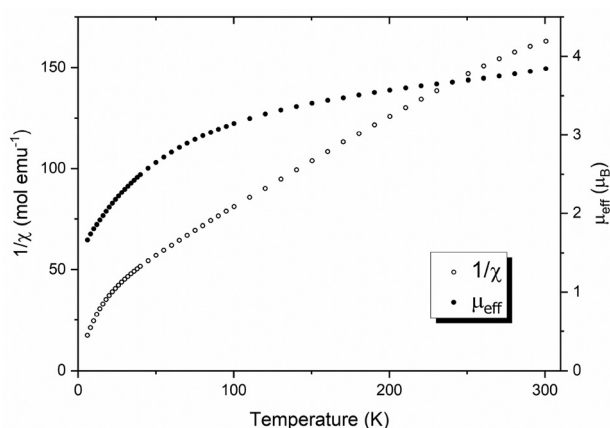


Fig. 5 Temperature dependence of reciprocal molar magnetic susceptibility ($1/\chi_{\text{mol}}$) and effective magnetic moment (μ_{eff}) for Ind*₂Mn.



and high-spin states (approximately $420 \text{ kJ mol}^{-1} \Delta E_{\text{HL}}$; Table S4†). Given the disparity in ΔE_{HL} , similarly observed in the literature for manganese spin-crossover complexes,⁴² more accurate methods were utilised to ascertain an appropriate functional for the reactive surface. Consequently, complete active space self-consistent field with second-order *N*-electron valence state perturbation theory (CASSCF/NEVPT2) along with domain-based local pair natural orbital coupled-cluster theory (DLPNO-CCSD(T)) were selected due to their increased reliability in determining ΔE_{HL} : CASSCF-NEVPT2 $\Delta E_{\text{HL}} = 117 \text{ kJ mol}^{-1}$ and CCSD(T) $\Delta E_{\text{HL}} = 134 \text{ kJ mol}^{-1}$.⁴² Reasonable agreement between the methods ($\Delta E = 17 \text{ kJ mol}^{-1}$) prompted selection of the DFT functional MN15 ($\Delta E = 138 \text{ kJ mol}^{-1}$), due to its alignment with CCSD(T).⁴² Consequently, optimised geometries were performed, with a scan across the RA for the optimised low- and high-spin structures where single point calculations were performed at DLPNO-CCSD(T) (Fig. 6). The potential energy surfaces of the high-spin and low-spin states were found to intersect, supporting the experimental observation that Ind^*_2Mn undergoes thermally induced spin-crossover.

In order to probe the differences in the nature of the Cp^* and Ind^* ligand architectures, supplementary energy decomposition analysis (EDA), *via* sobEDA, of L_2Mn ($\text{R} = \text{Ind}^*$ and Cp^*) was carried out. The total energy interaction was found to be broadly similar (Table S7†). While both ligands stabilise the manganese centre through electrostatic and orbital (E_{orb}) interactions, the Ind^* ligand contributes both larger Pauli repulsive (E_{rep}) and orbital attractive forces, markedly counteracting one another. Increased E_{rep} is reasonably explained by the imposed steric requirements of the benzanulated ring. Further decomposition of E_{orb} was undertaken using EDA-NOCV. The pairwise deformation densities reveal that, in the Ind^* complex, the benzene ring contributes additional electron density to the Mn centre through its extended π -system (Fig. S16†).

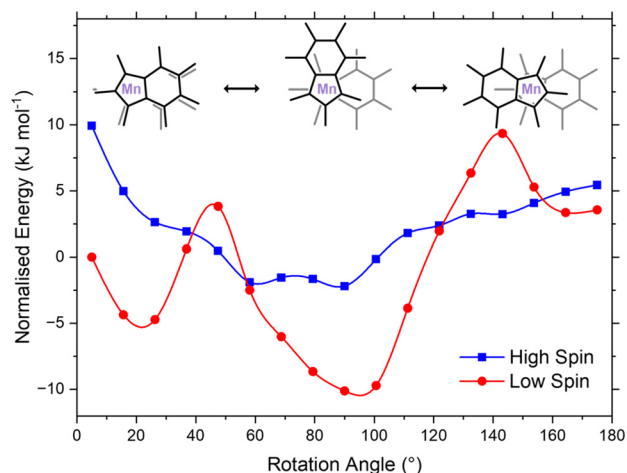


Fig. 6 Single point calculations performed at CCSD(T) for both high- and low-spin Ind^*_2Mn , energies normalised to the low-spin complex at 5° .

Conclusions

The solid state structure and the low temperature magnetic responses of Ind^*_2Mn suggest that it is the first indenyl manganese complex with a low-spin ground state. At room temperature the solid state magnetic moment suggest that a high-spin ($S = 5/2$) configuration is significantly populated. Given the ubiquity of Cp^* , certain circumstances likely demand increased steric constraint; consequently use of Ind^* as an alternate ligand framework may provide attractive complex properties and we hope this puts into greater focus the work being done already within the group.^{63–65}

Experimental

Synthesis of Ind^*H

7.40 g 1,2,4,5,6,7-hexamethyl-1*H*-indene (Ind^*H , 36.9 mmol, 1 eq.) was dissolved in 150 mL THF and was cooled, in a 300 mL schlenk under N_2 , to -78°C before fast dropwise addition of 23.1 mL *n*-butyllithium (1.6 M in hexanes; 36.93 mmol, 1 eq.). The reaction mixture was allowed to warm to room temperature over 16 h before cooling to -78°C and 2.53 mL MeI (40.63 mmol, 1.1 eq.) added dropwise. The reaction mixture was allowed to warm to room temperature over 2 h. THF was removed *in vacuo* and the residue dissolved in DCM and transferred to a separating funnel containing brine solution. Product was extracted in $3 \times 50 \text{ mL}$ DCM, dried over MgSO_4 , DCM removed by rotary evaporation before placing the oil in a -4°C freezer. Crystalline solid product was isolated by Buchner filtration and recrystallised from EtOH at -4°C yielding 7.35 g (92%) yellow, crystalline 1,2,3,4,5,6,7-heptamethylindene (Ind^*H). ^1H NMR (Benzene- d_6 , 298 K, 400 MHz) δ (ppm): 3.03 (q, 1H, $^3J_{\text{HH}} = 7.5$, CH), 2.44 (s, 3H, Ar-Me), 2.21 (s, 3H, Ar-Me), 2.18–2.13 (m, 9H, Ar-Me and Cp-Me), 1.82 (s, 3H, Cp-Me), 2.43 (d, 3H, $^3J_{\text{HH}} = 7.5$, Cp-Me).

Synthesis of Ind^*_2Mn

One equivalent of MnCl_2 (0.27 g, 2.18 mmol) was stirred in an ampoule in 30 mL THF for 16 hours, yielding a white suspension. This mixture was cooled to -78°C and a slurry of two equivalents of Ind^*Li (1.00 g, 4.36 mmol) in 30 mL THF added. The yellow-brown mixture was allowed to warm to room temperature and stirred for 15 hours, affording a brown-orange solution. The solvent was removed under vacuum to give a brown-orange solid. Extraction with hexane gave a purple solution, which was filtered through Celite and dried *in vacuo* affording a purple solid in 60% yield (0.543 g, 1.30 mmol). ^1H NMR (benzene- d_6 , 298 K, 499.9 MHz) δ (ppm): 9.06, 3.05, 2.44, 2.16, 1.82, 1.18 (all broad). $^{13}\text{C}\{^1\text{H}\}$ NMR (benzene- d_6 , 298 K, 499.9 MHz) δ (ppm): 145.3, 142.3, 141.2, 133.5, 132.7, 130.6, 128.6, 126.5, 46.6, 16.8, 16.4, 16.3, 16.3, 16.1, 15.5, 12.3. MS (EI): calc.: 481.2667 found: 481.2301. UV/Vis: $\lambda_{\text{max}}(n\text{-hexanes}) = 533.1 \text{ nm}$ ($\epsilon = 5.30 \times 10^2 \text{ mol}^{-1} \text{ dm}^3 \text{ cm}^{-1}$). FT-IR (KBr, cm^{-1}): 2965 (s), 1616 (m), 1447 (m), 1385



(s), 1261 (s), 1093 (m), 1022 (m), 801 (s). CHN analysis (calculated, found; %): C = 79.79, 79.91; H = 8.80, 8.83.

X-ray crystallography details

Colourless single crystals of Ind*H were isolated from concentrated pentane solution at room temperature. $C_{16}H_{22}$, $M_r = 214.33$, monoclinic, $P2_1/c$, $a = 13.0353(4)$ Å, $b = 10.9285(4)$ Å, $c = 9.0453(3)$ Å, $\alpha = 90^\circ$, $\beta = 91.765(3)^\circ$, $\gamma = 90^\circ$, $V = 1287.95(7)$ Å³, $Z = 4$, $T = 150$ K, block, colourless, 5161 independent reflections, $R_1 = 0.0470$, $wR_2 = 0.1368$ [$I > 2\sigma(I)$]. CCDC deposition number: 2421030† Dark purple single crystals of Ind*₂Mn were isolated from a concentrated solution (hexanes) after cooling to -24°C overnight. $C_{32}H_{42}Mn$, $M_r = 481.59$, triclinic, $P\bar{1}$, $a = 9.05120(10)$ Å, $b = 10.43980(10)$ Å, $c = 15.81240(10)$ Å, $\alpha = 73.2730(10)^\circ$, $\beta = 84.7780(10)^\circ$, $\gamma = 63.9150(10)^\circ$, $V = 1284.09(2)$ Å³, $Z = 2$, $T = 100^\circ\text{K}$, block, purple, 5238 independent reflections, $R(\text{int}) = 0.0279$, $R_1 = 0.0290$, $wR_2 = 0.0801$ [$I > 2\sigma(I)$]. CCDC deposition number: 2421029.†

Author contributions

A. E. carried out the investigation and formal analysis unless otherwise stated. W. K. M. performed the investigation and formal EPR analysis. A. E. C performed the investigation and formal computational analysis. S. C. B. assisted the SQUID investigation. T. A. Q. A. developed the improved ligand synthesis. A. E. managed the project, and drafted the manuscript. All authors revised the manuscript. D. O'H supervised the research and acquired funding.

Conflicts of interest

There are no conflicts to declare.

Data availability

The data supporting this article have been included as part of the ESI.† Crystallographic data for Ind*H and Ind*₂Mn has been deposited at the CCDC repository under 2421030 and 2421029.†

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References

- 1 T. J. Kealy and P. L. Pauson, *Nature*, 1951, **168**, 1039–1040.
- 2 S. A. Miller, J. A. Tebbboth and J. F. Tremaine, *J. Chem. Soc.*, 1952, 632–635.
- 3 H. Werner, *Angew. Chem., Int. Ed.*, 2012, **51**, 6052–6058.
- 4 P. J. Chirik and J. E. Bercaw, *Metallocenes: Synthesis Reactivity Applications*, Wiley-VCH, Weinheim, 1998.
- 5 P. L. Pauson, *Q. Rev., Chem. Soc.*, 1955, **9**, 391–414.
- 6 G. Wilkinson, P. L. Pauson and F. A. Cotton, *J. Am. Chem. Soc.*, 1954, **76**, 1970–1974.
- 7 J. M. O'Connor and C. P. Casey, *Chem. Rev.*, 1987, **87**, 307–318.
- 8 E. O. Fischer and W. Hafner, *Z. Naturforsch., B*, 1954, **9**, 503–505.
- 9 G. Wilkinson and F. A. Cotton, *Chem. Ind.*, 1954, 307.
- 10 L. Wirtz and A. Schäfer, *Chem. – Eur. J.*, 2021, **27**, 1219–1230.
- 11 E. O. Fischer and U. Piesbergen, *Z. Naturforsch., B*, 1956, **11**, 758–759.
- 12 E. O. Fischer and H. Grubert, *Z. Anorg. Allg. Chem.*, 1956, **286**, 237–242.
- 13 I. Resa, E. Carmona, E. Gutierrez-Puebla and A. Monge, *Science*, 2004, **305**, 1136–1138.
- 14 J. T. Boronski, A. E. Crumpton, L. L. Wales and S. Aldridge, *Science*, 2023, **380**, 1147–1149.
- 15 I.-A. Bischoff, S. Danés, P. Thoni, B. Morgenstern, D. M. Andrada, C. Müller, J. Lambert, E. C. J. Gießelmann, M. Zimmer and A. Schäfer, *Nat. Chem.*, 2024, **16**, 1093–1100.
- 16 T. Piou, F. Romanov-Michailidis, M. Romanova-Michaelides, K. E. Jackson, N. Semakul, T. D. Taggart, B. S. Newell, C. D. Rithner, R. S. Paton and T. Rovis, *J. Am. Chem. Soc.*, 2017, **139**, 1296–1310.
- 17 D. C. Calabro, J. L. Hubbard, C. H. Blevins, A. C. Campbell and D. L. Lichtenberger, *J. Am. Chem. Soc.*, 1981, **103**, 6839–6846.
- 18 P. L. Pauson and G. Wilkinson, *J. Am. Chem. Soc.*, 1954, **76**, 2024–2026.
- 19 I. A. Lobanova and V. I. Zdanovich, *Russ. Chem. Rev.*, 1988, **57**, 967–980.
- 20 M. Stradiotto and M. J. McGlinchey, *Coord. Chem. Rev.*, 2001, **219–221**, 311–378.
- 21 A. N. Nesmeyanov, N. A. Ustynyuk, L. G. Makarova, V. G. Andrianov, Y. T. Struchkov, S. Andrae, Y. A. Ustynyuk and S. G. Malyugina, *J. Organomet. Chem.*, 1978, **159**, 189–199.
- 22 J. A. Crisp, R. M. Meier, J. S. Overby, T. P. Hanusa, A. L. Rheingold and W. W. Brennessel, *Organometallics*, 2010, **29**, 2322–2331.
- 23 M. Maekawa, C. G. Daniliuc, M. Freytag, P. G. Jones and M. D. Walter, *Dalton Trans.*, 2012, **41**, 10317–10327.
- 24 V. Cadierno, J. Díez, M. Pilar Gamasa, J. Gimeno and E. Lastra, *Coord. Chem. Rev.*, 1999, **193–195**, 147–205.
- 25 E. O. Fischer, D. Seus and R. Jira, *Z. Naturforsch., B*, 1953, **8**, 692–693.



- 26 A. J. Hart-Davis and R. J. Mawby, *J. Chem. Soc. A*, 1969, 2403–2407.
- 27 A. J. Hart-Davis, C. White and R. J. Mawby, *Inorg. Chim. Acta*, 1970, **4**, 441–446.
- 28 M. E. Rerek, L.-N. Ji and F. Basolo, *J. Chem. Soc., Chem. Commun.*, 1983, 1208–1209.
- 29 T. K. Miyamoto, M. Tsutsui and L.-B. Chen, *Chem. Lett.*, 1981, **10**, 729–730.
- 30 D. Ohare, J. C. Green, T. Marder, S. Collins, G. Stringer, A. K. Kakkar, N. Kaltsoyannis, A. Kuhn, R. Lewis, C. Mehnert, P. Scott, M. Kurmoo and S. Pugh, *Organometallics*, 1992, **11**, 48–55.
- 31 D. O'Hare, V. J. Murphy and N. Kaltsoyannis, *J. Chem. Soc., Dalton Trans.*, 1993, 383–392.
- 32 T. M. Frankcom, J. C. Green, A. Nagy, A. K. Kakkar and T. B. Marder, *Organometallics*, 1993, **12**, 3688–3697.
- 33 D. Ohare, V. Murphy, G. M. Diamond, P. Arnold and P. Mountford, *Organometallics*, 1994, **13**, 4689–4694.
- 34 T. M. Trnka, J. B. Bonanno, B. M. Bridgewater and G. Parkin, *Organometallics*, 2001, **20**, 3255–3264.
- 35 M. Tsutsui, L. B. Chen, D. E. Bergbreiter and T. K. Miyamoto, *J. Am. Chem. Soc.*, 1982, **104**, 855–856.
- 36 D. O'Hare, V. Murphy, G. M. Diamond, P. Arnold and P. Mountford, *Organometallics*, 1994, **13**, 4689–4694.
- 37 N. Hebenanz, F. H. Koehler, G. Mueller and J. Riede, *J. Am. Chem. Soc.*, 1986, **108**, 3281–3289.
- 38 D. Cozak and F. Gauvin, *Organometallics*, 1987, **6**, 1912–1917.
- 39 J. H. Ammeter, R. Bucher and N. Oswald, *J. Am. Chem. Soc.*, 1974, **96**, 7833–7835.
- 40 S. Evans, M. L. H. Green, B. Jewitt, G. H. King and A. F. Orchard, *J. Chem. Soc., Faraday Trans. 2*, 1974, **70**, 356–376.
- 41 J. A. Bandy, F. G. N. Cloke, G. Copper, J. P. Day, R. B. Girling, R. G. Graham, J. C. Green, R. Grinter and R. N. Perutz, *J. Am. Chem. Soc.*, 1988, **110**, 5039–5050.
- 42 M. Drosou, C. A. Mitsopoulou and D. A. Pantazis, *Polyhedron*, 2021, **208**, 115399.
- 43 F. Engelmann, *Z. Naturforsch., B*, 1953, **8**, 775–776.
- 44 E. O. Fischer and H. Leipfinger, *Z. Naturforsch., B*, 1955, **10**, 353–355.
- 45 L. T. Reynolds and G. Wilkinson, *J. Inorg. Nucl. Chem.*, 1959, **9**, 86–92.
- 46 G. Wilkinson, F. A. Cotton and J. M. Birmingham, *J. Inorg. Nucl. Chem.*, 1956, **2**, 95–113.
- 47 M. E. Switzer, R. Wang, M. F. Rettig and A. H. Maki, *J. Am. Chem. Soc.*, 1974, **96**, 7669–7674.
- 48 J. L. Robbins, N. M. Edelstein, S. R. Cooper and J. C. Smart, *J. Am. Chem. Soc.*, 1979, **101**, 3853–3857.
- 49 M. D. Walter, C. D. Sofield, C. H. Booth and R. A. Andersen, *Organometallics*, 2009, **28**, 2005–2019.
- 50 A. M. May, M. Deegbey, K. Edme, K. J. Lee, R. N. Perutz, E. Jakubikova and J. L. Dempsey, *Inorg. Chem.*, 2024, **63**, 1858–1866.
- 51 A. E. Ashley, A. R. Cowley and D. O'Hare, *Eur. J. Org. Chem.*, 2007, 2239–2242.
- 52 A. E. Ashley, R. T. Cooper, G. G. Wildgoose, J. C. Green and D. O'Hare, *J. Am. Chem. Soc.*, 2008, **130**, 15662–15677.
- 53 G. Balazs, F. G. N. Cloke, L. Gagliardi, J. C. Green, A. Harrison, P. B. Hitchcock, A. R. M. Shahi and O. T. Summerscales, *Organometallics*, 2008, **27**, 2013–2020.
- 54 S. A. Westcott, A. K. Kakkar, G. Stringer, N. J. Taylor and T. B. Marder, *J. Organomet. Chem.*, 1990, **394**, 777–794.
- 55 M. J. Calhorda, V. Félix and L. s. F. Veiros, *Coord. Chem. Rev.*, 2002, **230**, 49–64.
- 56 D. P. Freyberg, J. L. Robbins, K. N. Raymond and J. C. Smart, *J. Am. Chem. Soc.*, 1979, **101**, 892–897.
- 57 A. H. Maki and T. E. Berry, *J. Am. Chem. Soc.*, 1965, **87**, 4437–4441.
- 58 S. Mugiraneza and A. M. Hallas, *Commun. Phys.*, 2022, **5**, 95.
- 59 D. F. Evans, *J. Chem. Soc.*, 1959, 2003–2005.
- 60 M. B. Meredith, J. A. Crisp, E. D. Brady, T. P. Hanusa, G. T. Yee, M. Pink, W. W. Brennessel and V. G. Young Jr., *Organometallics*, 2008, **27**, 5464–5473.
- 61 J. Cirera, M. Via-Nadal and E. Ruiz, *Inorg. Chem.*, 2018, **57**, 14097–14105.
- 62 S. Gómez-Coca and E. Ruiz, *Dalton Trans.*, 2024, **53**, 11895–11902.
- 63 C. G. Collins Rice, L. J. Morris, J.-C. Buffet, Z. R. Turner and D. O'Hare, *Chem. Commun.*, 2023, **59**, 12128–12131.
- 64 P. Ransom, A. E. Ashley, N. D. Brown, A. L. Thompson and D. O'Hare, *Organometallics*, 2011, **30**, 800–814.
- 65 C. G. Collins Rice, A. Evans, Z. R. Turner, J. Wattoom and D. O'Hare, *Ind. Chem. Mater.*, 2025, **3**, 178–190.

