Widening the Window of Spin-Crossover Temperatures in Bis(formazanate)iron(II) Complexes via Steric and Noncovalent Interactions

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ABSTRACT: Bis(formazanate)iron(II) complexes undergo a thermally induced $S = 0$ to $S = 2$ spin transition in solution. Here we present a study of how steric effects and π-stacking interactions between the triarylformazanate ligands affect the spin-crossover behavior, in addition to electronic substituent effects. Moreover, the effect of increasing the denticity of the formazanate ligands is explored by including additional OMe donors in the ligand (7). In total, six new compounds (2–7) have been synthesized and characterized, both in solution and in the solid state, via spectroscopic, magnetic, and structural analyses. The series spans a broad range of spin-crossover temperatures ($T_{1/2}$) for the LS ← HS equilibrium in solution, with the exception of compound 6 which remains high-spin ($S = 2$) down to 210 K. In the solid state, 6 was shown to exist in two distinct forms: a tetrahedral high-spin complex (6a, $S = 2$) and a rare square-planar structure with an intermediate-spin state (6b, $S = 1$). SQUID measurements, $^{57}$Fe Mössbauer spectroscopy, and differential scanning calorimetry indicate that in the solid state the square-planar form 6b undergoes an incomplete spin-change-coupled isomerization to tetrahedral 6a. The complex that contains additional OMe donors (7) results in a six-coordinate (NNO)$_2$Fe coordination geometry, which shifts the spin-crossover to significantly higher temperatures ($T_{1/2} = 444$ K). The available experimental and computational data for 7 suggest that the Fe···OMe interaction is retained upon spin-crossover. Despite the difference in coordination environment, the weak OMe donors do not significantly alter the electronic structure or ligand-field splitting, and the occurrence of spin-crossover (similar to the compounds lacking the OMe groups) originates from a large degree of metal–ligand π-covalency.

INTRODUCTION

The geometry of transition metal complexes is dependent on the electronic structure, and it is often the case that the geometry preferred on steric grounds is overridden in favor of a different one by electronic effects. In four-coordinate complexes two extreme geometries can be observed: the sterically favored tetrahedral and the electronically stabilized square-planar structure. While complexes with a d$^5$ configuration have been thoroughly investigated, the balance between steric and electronic effects on the geometry of compounds with a lower d-electron count is not well established. In the case of first-row transition metals such as Fe(II), the electronic stabilization is typically small, and therefore these compounds tend to adopt a tetrahedral configuration. Therefore, to observe square-planar Fe(II) complexes, specific requirements are usually needed that result in intermediate-spin ($S = 1$) compounds: (i) macrocyclic ligands that enforce a planar geometry around the metal center or (ii) strong field ligands, e.g., phosphines, that provide a greater ligand field stabilization energy compared to nitrogen and oxygen donors (in the case of mono- and bidentate ligands), often in combination with ortho-substituted aryl coligands. Exceptions to this where Fe(II) square-planar structures were observed have been sporadically reported. Furthermore, while isomerization between tetrahedral and square-planar geometries is a well-established phenomenon for cobalt(II), nickel(II), and copper(II), it is rare for iron(II).

In the simplistic terms of crystal field theory, the spin state of a complex in a certain geometry is determined by the orbital splitting (Δ) and the pairing energy (PE). When the values of these two parameters are comparable, various electronic states can be observed. The balance between the stabilizing electronic effects of the ligand field and the destabilizing electronic effects of the geometry must be optimized for the square-planar geometry to be favored. The complexes reported here illustrate that the geometry of bis(formazanate)iron(II) complexes can be tuned by the addition of ortho-methoxy groups (OMe) in the triarylformazanate ligands.
The SCO properties, such as the spin-crossover temperature ($T_{1/2}$), is well recognized. However, predicting the effect of changes in steric/electronic properties of the ligand and spin-crossover energetics remains very challenging due to the small energy differences involved.

Following our report of a four-coordinate Fe(II) spin-crossover complex with formazanate ligands, we recently established that spin-crossover is a general feature of this class of compounds. The stability of the low-spin ($S = 0$) state for these compounds is ascribed to an unusual splitting pattern of the d-orbitals in this geometry. Specifically, the formazanate ligands, which are good $\pi$-acceptor ligands, are engaged in $\pi$-backdonation with the metal, and this allows the formation of a highly covalent metal–ligand bond, stabilizing one of the d-orbitals (the antibonding $d_{\sigma}$ orbital that belongs to the $t_2$ set in a conventional tetrahedral complex), which gives rise to an "inverted" ligand field with an approximate "two-over-three" splitting pattern (Figure 1D). We demonstrated that it is possible to tune the SCO properties of bis(formazanate)iron(II) complexes by substituent effects that are purely electronic in nature. In the present work, we extend these studies to include steric effects as well as $\pi$-stacking interactions between the triarylformazanate ligands. Included in this analysis are nonsymmetric ligands that have two different N–Ar substituents. In addition, we describe the effect of additional OMe donor groups in the ligand.

### RESULTS AND DISCUSSION

The bis(formazanate) iron complexes 2–7 were synthesized following a procedure previously reported by us starting from the iron precursor Fe[N(SiMe$_3$)$_2$]$_2$ as depicted in Scheme 1. Complex 1 has already been extensively studied in our previous work, and it is therefore included in the discussion as reference compound. Besides compounds 1 and 2, all the others feature nonsymmetric ligands that have two different N–Ar substituents. The effect of an electron-withdrawing perfluorinated ring (Ar = C$_6$F$_5$) is studied either in the $C$–Ar position (2 and 4), as the N–Ar$_1$ group (5), or in both positions (6). At the same time the influence of the electron-donating, sterically demanding mesityl group (Ar$_2$ = Mes) is investigated in the N–Ar position alone (3) or in combination with the perfluorinated ring (4, 5, and 6). Furthermore, the ortho-anisyl group (Ar$_2$ = o-An) is introduced in the N–Ar position in compound 7, increasing the coordination ability of the formazanate to a tridentate monocationic ligand.

### Solid-State Characterization

While attempts to obtain single crystals suitable for X-ray diffraction were not successful for 2 and 5, the other compounds could be obtained in crystalline form. Single-crystal X-ray diffraction studies for complexes 3, 4, 6, and 7 allowed determination of their molecular structure, and pertinent metrical parameters are collected in Table 1. Overall, the structure of compound 3 is very similar to 1: it has relatively short Fe–N distances averaging to 1.831 Å and a flattened tetrahedral geometry...
Table 1. Selected Bond Lengths (Å) and Angles (deg) in Compounds 1, 3, 4, 6, and 7 at 100 K (Unless Stated otherwise)

|       | 1   | 3   | 4   | 6a  | 6b  | 7   |
|-------|-----|-----|-----|-----|-----|-----|
| Fe(1)−N(1) | 1.8278(15) | 1.8192(14) | 1.9946(11) | 2.030(2) | 1.9259(9) | 1.877(2) |
| Fe(1)−N(4) | 1.8207(15) | 1.8351(13) | 1.9610(12) | 1.9851(19) | 1.9461(9) | 1.883(2) |
| Fe(1)−N(5) | 1.8330(16) | 1.8242(13) | 1.9864(12) | 2.035(2) | 1.874(2) | 1.895(2) |
| Fe(1)−N(6) | 1.8174(16) | 1.8449(13) | 1.9916(12) | 1.9966(19) | 1.885(2) | 2.1128(18) |
| Fe(1)−O(1) | 1.9946(11) | 1.9461(9) | 1.9461(9) | 1.895(2) | 2.1039(19) | 1.9946(11) |
| \(\angle(NFeN)/\angle(NFeN)\) | 60.97(10) | 64.06(9) | 83.21(7) | 89.31(12) | 0.00(0) | 0.00(0) |
| Fe out-of-plane | 0.001 | 0.018 | 0.116 | 0.582 | 0.700 | 0.220 |
| distances of 6b | 1.9851(19) | 2.030(2) | 1.9851(19) | 2.030(2) | 1.9851(19) | 2.030(2) |
| distances of 6a | 1.9851(19) | 2.030(2) | 1.9851(19) | 2.030(2) | 1.9851(19) | 2.030(2) |

Data taken from ref 14a. Structure measured at 200 K. Dihedral angle between the coordination planes defined by the N−Fe−N atoms. Displacement of the Fe atom out of the plane defined by the four N atoms of each ligand backbone.

Figure 2. Crystal structure of compound 3 showing the π-stacking interactions between the mesityl rings. The Fe center, ligand backbone, and the mesityl rings are shown as 50% probability ellipsoids and the remaining atoms as wireframe; hydrogen atoms are removed for clarity.

Figure 3. Molecular structure of compounds 6a and 6b showing 50% probability ellipsoids; hydrogen atoms omitted for clarity. The inset for each shows the Fe(NNCNN)\(_2\) core of the structure with the N−Fe−N planes and the dihedral angle.

Figure 4. Crystal structure of compound 6b illustrating the π-stacking interactions between the aromatic rings, showing 50% probability ellipsoids. Parts of the molecule are shown as wireframe, and hydrogen atoms are removed for clarity.
planar angle of 10.14° and centroid-to-plane distance of 3.459 Å are observed. While the FeN₄ fragment is planar (enforced by the crystallographic symmetry), the FeNCCN bond angles are six-membered chelate rings are puckered with the Fe center displaced out of the ligand plane. The Fe–N bonds in square-planar 6b (1.9259(9)–1.9461(9) Å) are shorter than those found in high-spin FeN₄ complexes, such as tetrahedral 6a and in distorted-planar iron bis(amidinate) complexes reported by Hessen et al. (2.0528–2.0697 Å). The similarity of the geometrical parameters in 6b to those in intermediate-spin Fe(II) porphyrins (e.g., 1.972(4) Å in Fe(TPP)) suggests that 6b also has an S = 1 ground state. To the best of our knowledge, this is the first example of an intermediate-spin Fe(II) complex with bidentate nitrogen donor ligands which adopts a square-planar geometry in the solid state. Although solution studies (vide infra) indicate that 6 is high-spin in toluene, the accessibility of a square-planar polymorph for 6 suggests that controlling the strength of π-stacking interactions is a viable approach to change the geometric preference and thus spin state in this class of compounds.

57Fe Mössbauer spectroscopy was employed to elucidate the electronic structure of 6. A quadrupole doublet with isomer shift δ = 0.75 mm/s and quadrupole splitting ΔEₚ = 1.21 mm/s was observed for a batch of crystals for the tetrahedral complex 6a (Figure 5B). In contrast, crystals of square-planar 6b have a lower isomer shift (δ = 0.54 mm/s) and a higher quadrupole splitting (ΔEₚ = 2.73 mm/s) (Figure 5A). The Mössbauer spectra of both batches differ significantly from low-spin (S = 0) bis(formazanate)iron compounds, which have isomer shifts (δ) around 0 mm/s and an ΔEₚ of ca. 2 mm/s. The isomer shift of 6a is indicative of a high-spin state (S = 2) and indeed is comparable to that in the high-spin bis(formazanate)iron complex Fe(PhNCPPhNPPh) (δ = 0.60 mm/s). On the other hand, the isomer shift for 6b is in agreement with an intermediate spin state (S = 1), similar to the one reported for Fe(TPP) (δ = 0.50 mm/s). A crude powder of a pristine sample of 6 (i.e., not purified by crystallization) shows a Mössbauer spectrum identical with that of 6b and remains unchanged between 7 and 300 K (Figure 5C and Figure S4a–c). The magnetic susceptibility measurement of the powder sample of 6 recorded on a SQUID magnetometer gave χM(T) ≈ 1.1 cm³ mol⁻¹ K, supporting the assignment of an intermediate-spin state (Figure 6). The magnetic susceptibility in the solid state stays constant up to 390 K, and then it suddenly increases, approaching a value of 2.5 cm³ mol⁻¹ K at 400 K, which is lower than the expected value for a high-spin state S = 2 but could be an indication of an incomplete spin transition. To further probe this, the sample used for the SQUID measurement was subsequently analyzed by Mössbauer spectroscopy (Figure 5D). After 6b was heated to 400 K, the major species (82%) has a quadrupole doublet with δ = 0.74 mm/s and ΔEₚ = 1.17 mm/s, which are in good agreement with the values obtained for 6a. Thus, this indicates that square-planar, intermediate-spin 6b switches at least partially to tetrahedral, high-spin 6a in the solid state. Differential scanning calorimetry analysis of a fresh powder sample of 6 shows an endothermic transition at 412 K with an onset temperature around 397 K (followed by subsequent decomposition) and corroborates a spin transition in the solid state at high temperature.

Lastly, compound 7 containing formazanate ligands with an additional OMe donor moiety was characterized. Single-crystal X-ray diffraction allowed determination of the molecular structure as shown in Figure 7. It shows a distorted octahedral geometry where both the formazanate moieties act as...
Compounds of the metal complexes are collected in Table 3, and a plot of the high-spin fraction as a function of temperature is shown in Figure 8. A complex that has a high-spin ground state based on the lowest temperature that could be reached inside the NMR cell indicates an equilibrium between the high- and low-spin states. The changes in these values in the solid state at 80 K result from electron-donating substituents at that position, since the N–Ar groups predominantly influence metal–ligand π-bonding. However, it is clear from the crystallographic data of compound 3 (vide supra) that the N–Mes rings are engaged in noncovalent interactions (stacking), and we conclude that these attractive forces act to stabilize the more compact low-spin state.

Variable-Temperature NMR and UV/Vis Spectroscopy in Solution. We subsequently studied the spin-crossover behavior in solution by monitoring the spectral changes as a function of temperature. The NMR chemical shifts for all compounds are found to be temperature dependent but at low temperature do not follow the Curie behavior that is expected for a paramagnet: instead, the NMR resonances of all compounds except 6 converge into the dihedral range of the spectrum, suggestive of population of the $S = 0$ state. With 1 as reference, the change induced by the different ligand substituents is discussed below. The enthalpy and entropy differences ($\Delta H / \Delta S$) that describe the LS $\leftrightarrow$ HS equilibrium as well as the spin-crossover temperature ($T_{1/2}$) for the series of compounds are collected in Table 3, and a plot of the high-spin fraction as a function of temperature is shown in Figure 8.

For compound 2, which has a symmetrical ligand with a highly electron-donating C–Ar group, the variable-temperature $^1$H and $^{19}$F NMR spectra in toluene-$d_8$ are indicative of an equilibrium between the high- and low-spin states. Although the former is predominant even at 207 K (the lowest temperature that could be reached inside the NMR probe), and fitting the temperature dependence of the chemical shifts thus is somewhat less accurate, it is clear from the data that the thermodynamic values that describe the spin equilibrium are much decreased in 2 ($\Delta H = 8.5 \pm 0.4$ kJ mol$^{-1}$, $\Delta S = 45 \pm 4$ J mol$^{-1}$ K$^{-1}$) compared to 1. This can be attributed to the decrease in $\sigma$-donor strength of the ligands, which results in a smaller ligand-field splitting and destabilization of the low-spin state.

The introduction of an electron-rich, sterically demanding mesityl ring as an N–Ar group in compound 3 resulted in larger differences between both spin states, with $\Delta H = 26.3 \pm 0.1$ kJ mol$^{-1}$ and $\Delta S = 78 \pm 1$ J mol$^{-1}$ K$^{-1}$ from fitting the NMR data. The increase in these values stands in contrast to the expected effect of electron-donating substituents at that position, since the N–Ar groups predominantly influence metal–ligand π-bonding. However, it is clear from the crystallographic data of 3 (vide supra) that the N–Mes rings are engaged in noncovalent interactions (stacking), and we conclude that these attractive forces act to stabilize the more compact low-spin state.

The two effects discussed above were subsequently combined in compound 4. While the crystallographic data indicate that 4 is high spin in the solid state, the solution data clearly indicate that the $S = 0$ state is populated at low temperature. The combination of two opposing effects on the relative stability of the low-spin state results in thermodynamic parameters for the spin-state equilibrium in 4 ($\Delta H = 12.6 \pm 1.0$ kJ mol$^{-1}$; $\Delta S = 67 \pm 5$ J mol$^{-1}$ K$^{-1}$) that are intermediate between those of compounds 2 and 3.

Subsequently, we evaluated the influence of a highly electron-withdrawing N–C$_6$F$_5$ substituent that is present in compounds 5 and 6. Changing the N–Ph group in 3 to N–C$_6$F$_5$ in 5 results in a noticeable decrease in $\Delta H$ and $\Delta S$ to values of $19.0 \pm 0.4$ kJ mol$^{-1}$ and $70 \pm 1$ J mol$^{-1}$ K$^{-1}$, respectively. In the absence of structural data for 5, we refrain from a detailed interpretation of these values. It is noted, however, that this result runs counter to the expectation that an electron-withdrawing N–Ar group leads to increased $\Delta H / \Delta S$ due to stronger metal–ligand π-bonding.

For compound 6, which has an additional C$_6$F$_5$ substituent at the C–Ar position, the solution characterization data are

| $\delta$ ($\text{mm s}^{-1}$) | $\Delta E_{Q}$ ($\text{mm s}^{-1}$) |
|----------------------------|----------------------------|
| 0.03                       | 2.05                       |
| 0.05                       | 1.99                       |
| 0.75                       | 1.21                       |
| 0.54                       | 2.73                       |
| 0.55                       | 2.72                       |

*Measured in the solid state at 80 K. *$^b$Powder sample of the crude product before crystallization.

Table 2. $^{57}$Fe Mössbauer Parameters (δ = Isomer Shift in mm s$^{-1}$; |$\Delta E_{Q}$| = Quadrupole Splitting in mm s$^{-1}$) for Compounds 1, 3, and 6$^a$

|        | 1 | 2a | 3 | 6a | 6b |
|--------|---|----|---|----|----|
| δ      | 0.03 | 0.05 | 0.75 | 0.54 | 0.55 |
| |$\Delta E_{Q}$| 2.05 | 1.99 | 1.21 | 2.73 | 2.72 |

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For compound 6, which has an additional C$_6$F$_5$ substituent at the C–Ar position, the solution characterization data are

|        | 1 | 2 | 3 | 4 | 5 | 6 | 7 |
|--------|---|---|---|---|---|---|---|
| $\Delta H$ (kJ mol$^{-1}$) | 22.2 ± 0.3 | 8.5 ± 0.4 | 26.3 ± 0.1 | 12.6 ± 1.0 | 19.0 ± 0.4 | − | 37.5 ± 1.6 |
| $\Delta S$ (J mol$^{-1}$ K$^{-1}$) | 64 ± 1 | 45 ± 4 | 78 ± 1 | 67 ± 5 | 70 ± 1 | − | 85 ± 5 |
| $T_{1/2}$ (K) | 345 ± 7 | 192 ± 18 | 340 ± 2 | 188 ± 21 | 271 ± 8 | − | 444 ± 34 |

*Data reproduced from ref 14a. *$^b$Estimated from fitting a limited temperature range. *$^c$The uncertainty in $T_{1/2}$ is obtained by using error propagation from $\Delta H$ and $\Delta S$. 

Figure 8. Temperature dependence of the high-spin fraction ($\gamma_{HS}$) of compounds 1–5 in toluene-$d_8$, including error bars for $T_{1/2}$ ($\gamma_{HS} = 0.5$). The liquid range for toluene is indicated with the color gradient at the temperature axis.
indicative of a high-spin ground state also at low temperature, as shown by a magnetic moment of 4.9−5.1 μB across the temperature range studied (217−348 K). The spectral changes in the VT 1H NMR studies provide evidence for a spin equilibrium at the lowest temperatures (a departure from Curie behavior), which may indicate population of either the intermediate-spin (S = 1) state that is observed by crystallography or a low-spin (S = 0) complex similar to that present for the other compounds. However, the population of a different spin state is too small at these temperatures to allow an unambiguous interpretation of the changes that occur.

Finally, the spin-crossover properties of compound 7 were evaluated in solution. At room temperature, the 1H NMR spectrum of 7 shows resonances in the diamagnetic range, and the number of signals is indicative of C₃v symmetry. While most peaks are sharp, those corresponding to the 1H of the OMe groups appear broadened, suggesting that also 7 may show a temperature-dependent equilibrium between a LS (S = 0) diamagnetic state and a HS (S = 2) paramagnetic state. Indeed, when the temperature was increased, the resonances of 7 broaden substantially and shift away from their diamagnetic values (Figure 9).

Figure 9. 1H NMR spectra of 7 recorded between 247 and 397 K (toluene-d₈, 500 MHz).

The variable-temperature NMR data can be modeled with the equilibrium parameters ΔH = 37.5 ± 1.6 kJ mol⁻¹ and ΔS = 85 ± 5 J mol⁻¹ K⁻¹. The increase in ΔH compared to the other compounds discussed above indicates that there is a substantial additional enthalpic penalty upon changing the spin state from singlet to quintet. A key question surrounding the spin-crossover in 7 is whether or not the Fe···OMe interaction is retained in solution; that is, does it involve a change in the coordination sphere around the Fe center, or does the ligand maintain the same coordination mode in both spin states? Several lines of experimental and computational evidence point toward retention of the tridentate NNO coordination mode of the ligand in both spin states, resulting in an octahedral geometry for 7 throughout. First, although 7 is predominantly low-spin at room temperature, its OMe resonance is somewhat broadened. This is likely because it is in close proximity to the paramagnetic center and is thus noticeably affected, also when the population of high-spin 7 is still very low. In addition, the spin-state equilibrium in 7 is characterized by a value of ΔS (85 ± 5 J mol⁻¹ K⁻¹) that is only marginally larger than that of the others; loss of the Fe···OMe interaction in the high-spin state is expected to lead to a much larger entropy change. Finally, we performed density functional theory calculations on 7 in both spin states, with and without the Fe···OMe interaction (7_calc and 7'_calc, respectively; see the Supporting Information for details). The results of geometry optimizations with a def2-TZVP basis set using either pure (BP86) or hybrid functionals (TPSSH, B3LYP) all indicate that structures with the Fe···OMe interaction are favored over those in which the OMe group points away from the metal center. The optimized geometries for 7_calc in the low-spin state have short Fe−O bonds of 2.14−2.21 Å, which are elongated to 2.43−2.51 Å in the S = 2 minima; the shortest bonds are found for the TPSSH geometries and the longest for BP86. Although, as expected, there are large differences between these functionals for the computed energy differences between the different spin states, it is important to note that the calculations indicate that coordination of the OMe groups is stabilizing in both spin states, regardless of the functional used (ΔG_calc > 23.7 kJ mol⁻¹). Analysis of the frontier molecular orbitals of (low-spin) 7_calc shows that the additional interaction with the weak OMe donor groups does not lead to a substantial change in ligand-field strength in comparison to a structure in which the OMe groups are rotated away from the metal center (7'_calc; see Figures S54 and S55 for a comparison of the canonical DFT orbitals). In fact, the HOMO−LUMO energy gap at the BP86/def2-TZVP level is somewhat smaller in the structure with the Fe···OMe interaction (LS−7_calc: 9227 cm⁻¹) compared to without (LS−7′_calc: 10466 cm⁻¹). Analysis of the intrinsic bonding orbitals (Figure 10). A similar analysis using the B3LYP and TPSSH functionals at the corresponding minima for 7_calc shows that the intrinsic bonding orbitals are qualitatively similar for TPSSH. On the other hand, the B3LYP results show less covalent Fe−N bonds, which is reflected by a lower N-contribution to the relevant bonding orbitals.

Figure 10. Representation of intrinsic bonding orbitals at the BP86/def2-TZVP minima, both with (7_calc: (A)) and without Fe−O interaction (7′_calc: (B)).
IBOs and a smaller Wiberg bond index for the Fe–N bonds (Table S6).

To corroborate the NMR data, we subsequently performed variable-temperature UV–vis spectroscopic measurements on all compounds. Dilute solutions in toluene (ca. 10⁻⁵ M) were analyzed at temperatures down to 183 K. The fact that we could access lower temperatures in the UV–vis spectrometer was particularly helpful in the analysis of compounds 2 and 4, for which spin-crossover has a relatively low \( T_{1/2} \). The thermodynamic parameters obtained from the fitting of the UV–vis data are congruent with those found from the NMR analysis (see Table S4).

Although the UV–vis spectra of the compounds in this series often are equilibrium mixtures that contain both spin states, the data at the extremes of the temperature range represent predominantly low- or high-spin (at low or high temperature, respectively), and these were taken to extract the absorption maxima of the other spin state by scaled subtraction (see the Supporting Information for details). The only exception is compound 6, the UV–vis spectrum of which does not change appreciably with temperature, and 6 is predominantly found in the high-spin state (Figure 11A). The reported in this work allowed crystallographic characterization of structures with different coordination geometries and spin states: pseudo-tetrahedral low-spin (3), tetrahedral high-spin (4, 6a), square-planar intermediate-spin (6b), and octahedral low-spin (7). Moreover, 6b is shown to thermally switch in the solid state to 6a, undergoing an incomplete spin-change-coupled square-planar–tetrahedral isomerization, which is rare for iron(II) compounds. The combination of steric, \( \pi \)-stacking interactions, and electronic effects provides a plethora of tools that can be used to substantially affect spin-crossover behavior in this class of compounds. Overall, we were able to tune the system to obtain solution spin-crossover properties that range from very low \( T_{1/2} \) (∼190 K in 2 and 4) to well above room temperature (444 K in 7). Computational data suggest that the spin-crossover in the six-coordinate bis(formazanate)iron(II) complex (7) is of similar nature to that previously described for the four-coordinate derivatives 14 and originates from a large degree of covalency in the Fe–N bonds due to metal → ligand \( \pi \)-back-donation. Given the relevance of understanding and tuning spin-state-dependent reactivity, we anticipate that this study provides useful insight into ways to fine-tune the spin-state energetics in Fe(II) complexes.

**EXPERIMENTAL SECTION**

**General Considerations.** All manipulations were performed under nitrogen or argon by using standard glovebox, Schlenk, and vacuum-line techniques. THF (Aldrich, anhydrous, 99.8%) was dried by percolation over columns of Al₂O₃ (Fluka); toluene, hexane, and pentane (Aldrich, anhydrous, 99.8%) were passed over columns of Al₂O₃ (Fluka), BASF R3-11-supported Cu oxygen scavenger, and molecular sieves (Aldrich, 4 Å). THF-\( d₆ \) (Euroisotop-top) and Tol-\( d₄ \) (Aldrich) were vacuum transferred from Na/K alloy and stored under nitrogen.

The compounds 2H, 3H, 4H, 5H, 6H, and Fe[N-(SiMe₃)₂]₃ were synthesized according to the literature procedures. Ligand 7H was prepared according to a slightly adapted version of a literature method 26,27 (see the Supporting Information for a detailed description). Sodium carbonate (Merck), tetrabutylammonium bromide (Sigma-Aldrich, 99%), \( \alpha \)-anisidine (Sigma-Aldrich, >99%), hydrochloric acid (Boom B.V., 37–38%), and sodium nitrite (Sigma-Aldrich, 99%) were used as received.

NMR spectra were recorded on a Varian Mercury 400, Inova 500, or Bruker 600 MHz spectrometer. The \(^1\)H and \(^{13}\)C NMR spectra were referenced internally by using the residual solvent resonances and reported in ppm relative to TMS (0 ppm). The assignments of NMR resonances were aided by COSY, HMQC, HSQC, and HMBC experiments using standard pulse sequences.

Elemental analyses were performed by the analytical laboratory of the Institute of Inorganic Chemistry at the University of Göttingen using an Elementar Vario EL III instrument.

**Synthesis of Fe[PhN(NC(C₆F₅))₂NPh]₂ (2).** A dark-orange solution of L₂H (516.5 mg, 1.32 mmol) in THF (20 mL) was added to a green solution of Fe[N(SiMe₃)₂]₃ (247.9 mg, 0.66 mmol) in THF (10 mL). The reaction mixture was stirred overnight at room temperature, leading to a red-brick-colored solution. The volatiles were removed under a vacuum, and the product was extracted in THF. Slow diffusion of hexane into the THF solution at ∼30 °C resulted in 303.8 mg of dark-brown powder (0.36 mmol, 55%).

**Synthesis of Fe[PhN(NC(p-Tol)NMe₃)]₂ (3).** Fe[N(SiMe₃)₂]₃ (0.71 g, 1.88 mmol) was dissolved in THF (15 mL), and a solution of

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**Figure 11.** UV/vis spectra in toluene for (A) compound 6 recorded between 183 and 293 K and (B) compound 7 recorded between 293 and 383 K.

LS spectra for 2–5 show two intense bands in the visible range with absorption maxima between 375–445 and 515–575 nm which are assigned to ligand-based \( \pi \)-\( \pi \) transitions. 14 In the HS state, the two bands are bathochromically shifted (around 390–510 and 580–630 nm, respectively), and the lowest energy band shows a significantly lower intensity (Figure S24).

The UV–vis spectrum of compound 7 is distinct from the others as it shows three absorption maxima in the LS state (\( \lambda = 459, 608, \) and 828 nm; see Figure 11B). The lowest-energy transition in 7 is much broader and occurs at significantly lower energy than in the other compounds. Thus, the presence of the additional OMe donor groups in 7 results in an additional low-lying excited state that is a distinguishing feature of this compound.

**CONCLUSIONS**

In this work, we have extended the series of bis(formazanate)-iron complexes to systems featuring nonsymmetric ligands with two different N–Ar substituents. We have demonstrated that the spin-crossover behavior of this class of compounds may be modulated via modification of the ligand using different strategies: electronic effects, steric effects, \( \pi \)-stacking interactions, and ligand denticity. The formazanate ligands...
Table 4. Crystallographic Data for Compounds 3, 4, 6a, 6b, and 7

| 3        | 4         | 6a           | 6b         | 7          |
|----------|-----------|--------------|------------|------------|
| M         | C₂₄H₁₃N₇Fe | C₂₄H₁₃N₇Fe₂  | C₂₄H₁₃N₇Fe₂ | C₂₄H₁₃N₇Fe₂ |
| crys syst | triclinic | monoclinic   | monoclinic | monoclinic |
| color, habit | red, black | brown, black | red, black | green, needle |
| size (mm) | 0.23 × 0.18 × 0.11 | 0.42 × 0.26 × 0.09 | 0.30 × 0.17 × 0.15 | 0.34 × 0.20 × 0.09 |
| ρ, abs g cm⁻³ | 1.308 | 1.425 | 1.609 | 1.843 |
| radiation [Å] | Mo Kα 0.71073 | Mo Kα 0.71073 | Mo Kα 0.71073 | Cu Kα 1.54178 |
| μ(Mo Kα), mm⁻¹ | 0.432 | 0.407 | 0.458 | 0.525 |
| μ(Cu Kα), mm⁻¹ | 2.026Fᵣ₀ | 2.026Fᵣ₀ | 2.026Fᵣ₀ | 2.026Fᵣ₀ |
| F(000) | 808 | 1036 | 2192 | 548 |
| temp (K) | 100(2) | 100(2) | 200(2) | 100(2) |
| θ range (deg) | 2.76–29.176 | 3.04–27.92 | 2.88–26.38 | 2.88–27.94 |
| data collected (h, k, l) | −10:10–13:14 | −16:16–16:16 | −15:15–25:25 | −9:9–16:16–16:16–14:14–28:28 |
| no. of rns collected | 62040 | 70899 | 54918 | 47685 |
| no. of indpndt collected | 8538 | 11211 | 9089 | 4755 |
| observed rns | 7238 | 9251 | 6346 | 5557 |
| R(F) (%) | 3.46 | 3.36 | 4.32 | 2.55 |
| wR(F2) (%) | 8.03 | 8.05 | 9.93 | 2.55 |
| GooF | 1.045 | 1.040 | 1.030 | 7.07 |
| weighting a, b | 0.0267, 1.5260 | 0.0311, 1.6277 | 0.0324, 3.4691 | 0.0372, 0.6125 |
| params refined | 504 | 687 | 664 | 334 |
| min, max resid dens | −0.427, 0.295 | −0.280, 0.375 | −0.260, 0.264 | −0.451, 0.379 |

L3H (1.31 g, 3.66 mmol) in THF (15 mL) was added. The reaction mixture was stirred for 2 days at room temperature, leading to a dark-red solution. The solution was filtered, and the volatiles were removed under a vacuum. Recrystallization by slow diffusion of hexane into a THF solution gave 0.62 g of brown powder (0.83 mmol, 44% yield). ¹H NMR (600 MHz, THF-d₅, 25 °C): δ 12.48 (2H, Ph-m-CH), 10.94 (2H, p-Tol-m-CH), 10.56 (1H, Mes m-CH), 9.76 (3H, p-Tol-CH₃), 8.66 (6H, Mes o-CH₃), 8.45 (1H, Mes p-CH₃), 3.96 (2H, p-Tol o-CH₂), 0.42 (1H, Ph-p-CH), −0.70 (3H, Mes p-CH₃), −1.44 ppm (2H, Ph-p-CH). ¹³C NMR (500 MHz, THF-d₅, −55 °C): δ 8.06 (2H, p-Tol p-CH), 7.37 (2H, p-Tol m-CH), 7.25 (2H, p-Me CH), 7.11 (1H, Mes m-CH²), 7.05 (1H, Ph-p-CH), 6.65 (1H, Mes m-CH²), 6.47 (2H, Ph-p-CH), 2.58 (6H, p-Tol o-CH₃ and Mes o-CH₃), 1.69 (3H, Mes o-CH₃), 0.41 ppm (3H, Mes o-CH₃). ¹³C NMR (125 MHz, THF-d₅, −55 °C): δ 144.5 (Ph pico-C), 142.3 (Mes pico-C), 141.4 (NCN-C), 137.8 (pico-C), 136.5 (pico-C), 134.2 (pico-C), 133.9 (Mes m-C), 133.7 (Ph pico-C), 133.2 (p-tol o-C), 132.9 (p-tol m-C), 131.9 (Ph m-C), 247 (Mes o-CH₃), 242 (p-tol p-CH₂), 228 (Mes p-CH₃), 220 ppm (Mes o-CH₃). Anal. Calcd for C₂₄H₁₃N₇Fe: C, 72.06; H, 0.65; N, 14.61. Found: C, 72.54; H, 5.82; N, 14.12.

**Synthesis of Fe[PhNNC(p-Tol)NMMes]₅** (5). A red-brick-colored solution of LSH (1.41 g, 3.16 mmol) in toluene (40 mL) was added to a green solution of Fe[N(SiMe₃)₂]₃ (0.60 g, 1.58 mmol) in toluene (10 mL). The reaction mixture was stirred overnight at room temperature, leading to a brown solution. The solution was filtered, and the volatiles were removed under a vacuum; the obtained dark solid was quickly washed with cold hexane, giving 1.28 g (1.35 mol, 85%) of crude product. Any attempt to recrystallize the product was unsuccessful. ¹H NMR (400 MHz, CD₂Cl₂, 25 °C): δ 24.23 (3H), 17.32 (2H), 15.52 (3H), 14.15 (2H), 12.84 (6H, Mes o-CH₃), −0.62 ppm (2H). ¹⁹F NMR (375 MHz, CD₂Cl₂, 25 °C): δ −91.63 (1F, p-CF₃), −163.25 ppm (2F, m-CF₃). The signal of C₃F₃ o-CF was not visible due to line broadening.

**Synthesis of Fe[C₆F₅NNC(p-Tol)NMMes]₅** (6). A dark-orange solution of 6H (1.352 g, 2.587 mmol) in THF (40 mL) was added to a green solution of Fe[N(SiMe₃)₂]₃ (0.489 g, 1.299 mmol) in THF (20 mL). The reaction mixture was stirred for 5 h, leading to a brown solution. The volatiles were removed under a vacuum, and a brown solid was collected in 70% yield (1.009 g, 0.919 mmol). The solid was recrystallized from refluxing hexane which afforded crystals of 6a suitable for X-ray diffraction. Alternatively, diffusion of hexane into a THF solution afforded single crystals of 6b. ¹H NMR (500 MHz, toluene-d₅, 25 °C): δ 20.94 (3H, Mes p-CH₃), 14.77 (2H, Mes m-CH), 11.22 ppm (6H, Mes o-CH₃). ¹⁹F NMR (375 MHz, toluene-d₅, 25 °C): δ −93.78 (1F, p-CF₃), −106.37 (2F, o-CF₃), −130.67 (1F, p-CF₃), −154.58 (2F, m-CF₃), −160.23 ppm (2F, m-CF₃). The signal of o-
CF of N–CF$_2$ was not visible due to line broadening. Note: NMR spectra are identical for 6a and 6b. Anal. Calcd for C$_{64}$H$_{62}$N$_8$F$_4$Fe: C, 48.11; H, 2.02; N, 10.20. Found: C, 48.25; H, 1.85; N, 10.04.

**Synthesis of Fe[PhNHC(p-Tol)NN(o-An)]$_2$·0.5(THF)** (7). A fuchsia THF (10 mL) solution of L$_7$H (96.4 mg, 0.28 mmol) was added to a green solution of Fe[N(SiMe$_3$)$_2$]$_2$ (52.7 mg, 0.14 mmol) in 5 mL of THF. The reaction mixture was stirred for 3 days at room temperature, leading to a brown solution that was filtered through a 0.2 μm syringe filter, and slow diffusion of hexane into the THF solution afforded 7 as dark needles in 70% yield (76.6 mg, 0.098 mmol). $^1$H NMR (400 MHz, C$_6$D$_6$, 25 °C): $\delta = 8.43$ (d, $J = 7.6$ Hz, 1H, o-An $^6$CH), 8.36 (d, $J = 7.9$ Hz, 2H, p-tolyl $^o$CH), 7.37 (d, $J = 7.7$ Hz, 2H, p-tolyl m-CH), 7.08 (t, $J = 7.2$ Hz, 1H, o-An $^4$CH), 6.84 (d, $J = 6.7$ Hz, 2H, Ph m-CH), 6.75 (t, $J = 7.5$ Hz, 1H, o-An $^6$CH), 6.64 (t, $J = 7.3$ Hz, 1H, Ph p-CH), 6.41 (m, 1H, Ph o-CH), 6.30 (d, $J = 7.9$ Hz, 1H, o-An $^4$CH), 3.58 (m, 1H, THF)$^a$, 2.78 (s, 3H, o-An OCH$_3$), 2.41 (s, 3H, p-tolyl CH$_3$), 1.42 ppm (m, 1H, THF)$^a$. $^{13}$C NMR (151 MHz, C$_6$D$_6$, 25 °C): $\delta = 169.7$ (Ph iso-C), 152.8 (NCN), 151.6 (o-An iso-COCH$_3$), 150.4 (o-An iso-C), 137.0 (p-tolyl iso-CCH$_3$), 136.4 (p-tolyl iso-C), 128.8 (p-tolyl m-CH), 127.4 (p-tolyl o-CH), 127.3 (Ph m-CH), 126.6 (Ph p-CH) 124.6 (Ph o-CH), 124.4 (o-An $^6$CH), 123.2 (o-An $^4$CH), 118.6 (o-An $^6$CH), 112.0 (o-An $^4$CH), 67.8 (THF)$^a$, 56.5 (o-An OCH$_3$), 25.8 (THF)$^a$, 21.0 ppm (p-tolyl CH$_3$). Anal. Calcd for C$_{64}$H$_{62}$N$_8$O$_2$.5Fe: C 67.87, H 5.44, N 14.39. Found: C 68.02, H 5.43, N 14.02.

**X-ray Crystallography.** Single crystals of compounds 3, 4, 6a, 6b, and 7 (directly obtained from the mother liquor) were mounted on top of a cryoloop and transferred into the cold nitrogen stream (100 K; 200 K for 6a) of a Bruker-AXS D8 Venture diffractometer. Data collection and reduction was done by using the Bruker software suite APEX3.$^{37}$ The final unit cell was obtained from the xyz centroids of 9772 (3), 9845 (4), 9892 (6a), 9813 (6b), and 9876 (7) reflections after integration. A multiscan absorption correction was applied for compounds 3, 4, 6a, and 6b based on the intensities of symmetry-related reflections measured at different angular settings (SADABS). For compound 7, a numerical absorption correction was applied after indexing of the crystal faces in APEX3. The structures were solved by direct methods using SHELXS$^{34}$ and refinement of the structure was performed by using SHELXL.$^{35}$ From the refinement of 4 it was clear that the hexane solvent molecule was disordered. A two-site disorder model was used to describe this. The site-occupancy factor for the major disorder component refined to 0.72. Several of the atoms in the disordered solvent molecule gave nonpositive definite displacement parameters when refined freely, and ultimately DFIX and ISOR instructions were applied. The structure of 6a was measured at 200 K because the data at 100 K indicated an (incomplete) phase transition (not further investigated). For all structures, the hydrogen atoms were generated by geometrical considerations, constrained to idealized geometries and allowed to ride on their carrier atoms with an isotropic displacement parameter related to the equivalent displacement parameter of their carrier atoms. Crystal data and details on data collection and refinement are presented in Table 4.

**ASSOCIATED CONTENT**

**Supporting Information**

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/acs.inorgchem.0c03593.

Full experimental and characterization data, computational details (PDF)

**Accession Codes**

CCDC 2036148–2036152 contain the supplementary crystallographic data for this paper. These data can be obtained free of charge via www.ccdc.cam.ac.uk/data_request cif, or by emailing data_request@ccdc.cam.ac.uk, or by contacting The Cambridge Crystallographic Data Centre, 12 Union Road, Cambridge CB2 1EZ, UK; fax: +44 1223 336033.

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**Notes**

The authors declare no competing financial interest.

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**ADDITIONAL NOTE**

“Crystals of 7 contain one THF per iron complex, but drying results in loss of part of the THF solvate molecules.

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