Catalyzed M–C coupling reactions in the synthesis of $\sigma$-(pyridylethynyl) dicarbonylcyclopentadienyliiron complexes†

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† Electronic supplementary information (ESI) available. CCDC 1973678, 1973917, 1973260 and 1982510. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d0ra02333g

The reactions between terminal ethynylpyridines, (trimethylsilyl)ethynylpyridines and cyclopentadienyliiron dicarbonyl iodide were studied under Pd/Cu-catalyzed conditions to develop a synthetic approach to the $\sigma$-alkynyl iron complexes $\text{Cp(CO)}_2\text{Fe} \equiv \text{C} \equiv \text{C} \equiv \text{R}$ ($\text{R} = \text{ortho-}, \text{meta-}, \text{para-}$-pyridyl). Depending on the catalyst and reagents used, the yields of the desired $\sigma$-pyridylethynyl complexes varied from 40 to 95%. In some cases the reactions with ortho-ethynylpyridine gave as byproduct the unexpected binuclear FePd $\mu$-pyridylvinylidyne complex $[\text{Cp(CO)}\text{Fe}(\eta^1\text{C}_6\text{H}_4\text{N})\equiv\text{C}(\equiv\text{C}(\equiv\text{Fe}(\eta^1\text{C}_6\text{H}_4\text{N}))\equiv\text{Fe}(\eta^1\text{C}_6\text{H}_4\text{N})\equiv\text{C})]$. The conditions, catalysts, and reagents that provide the highest yields of the desired $\sigma$-pyridylethynyl iron compounds were determined. The methods developed allowed the synthesis of the corresponding $\sigma$-4-benzothiadiazolylethynyl complex $\text{Cp(CO)}_2\text{Fe} \equiv \text{C} \equiv \text{C} \equiv \{4\text{-C}_{6}\text{H}_{4}\text{N}_{2}\} \equiv \text{S}$ as well. Eventually, synthetic approaches to $\sigma$-alkynyl iron complexes of the type $\text{Cp(CO)}_2\text{Fe} \equiv \text{C} \equiv \text{C} \equiv \text{R}$ ($\text{R} = \text{ortho-}, \text{meta-}, \text{para-}$-pyridyl, 4-benzothiadiazol-2,1,3-yl) based on the Pd/Cu-catalyzed cross-coupling reactions were elaborated.

Introduction

Metal $\sigma$-alkynyl complexes displaying such peculiar characteristics as linear geometry, high stability, and $\pi$-unsaturated character have been demonstrated to constitute promising building blocks for the design of materials, which can possess such properties as optical nonlinearity,4–5 light-emission,6–12 and electrical conductivity.13–18 Moreover, they are an important class of coordination compounds because of their relevance in synthetic chemistry19–24 and proton reduction catalysis.25–27

A variety of methods for the synthesis of transition metal acetylides have been developed.28 The most common synthetic route to them is transmetalation, where a generated $[\text{M} \equiv \text{C} \equiv \text{C} \equiv \text{R}]$ species ($\text{M} = \text{Cu}(i), \text{Ag}(i), \text{Au}(i)$, alkali-metal (Li, Na), or an alkaline-earth-metal (MgX, etc.)) acts as an alkynyl transfer reagent to transition-metal halide complexes $\text{L}_n\text{MX}$ ($\text{X} = \text{I}, \text{Br}$, Cl).28–31 However, in some cases, such reactions may give low yields of desired products and a range of by-products, for example, complexes in which the copper or the silver fragments are $\pi$-coordinated to the transferred alkynyl ligand.35–38

Other general strategy to the preparation of transition metal acetylides takes advantage of the facility of some transition metal complexes to catalyze M–C coupling reactions. The most general route to group 10 metal $\sigma$-alkynyl derivatives is based on copper(0)-catalyzed dehydrohalogenation reactions between an appropriate metal halide complex and a terminal alkyn in an amine solvent.39–42 This method is applicable to the synthesis of alkynyl, polynyls, and polyynyls of tungsten, molybdenum, iron, ruthenium, rhodium, and iridium.54–47 In some cases, the Cu-catalyzed reactions of transition metal halides with stannyl acetyl enes may be performed in the absence of amines.48–51

Palladium catalysts, currently being an indispensable tool of organic synthesis, can also be used in the $\text{M} \equiv \text{C} \equiv \text{C} \equiv \text{R}$ bond formation. The team of Claudio Lo Sterzo demonstrated that, similarly to organic electrophiles, transition metal iodides undergo coupling with trialkyltin acetylides, in the presence of palladium to form alkynyl complexes of ruthenium, iron, tungsten, and molybdenum.52–55 The role of palladium catalysts in promoting these transformations was also investigated.52,56–59

Although, the Lo Sterzo approach was shown to be a valuable route to alkynyl complexes, the need for preparation of tin reagents and removal of tin impurities limits the appeal of this method. To overcome these disadvantages, one can use an organometallic analogue of the Sonogashira protocol,60,61 where...
transition metal halides react with terminal alkynes to form metal alkynyls via the Pd/Cu- or Pd-catalyzed dehydrohalogeneration route, which provides milder reaction conditions and facilitates purification of products. Despite the seeming availability of this approach for the synthesis of metal acetylides, it was exploited only once by Oshima in the synthesis of σ-alkynyl iron complexes \( \text{Cp(CO)}_2\text{Fe-C≡C-Ar} \).

As a part of our studies on proton reduction catalysis, we searched for a facile synthetic route to a series of σ-pyridylethynyl iron complexes \( \text{Cp(CO)}_2\text{Fe-C≡C-(n-C}_5\text{H}_4\text{N)} \) \( n = \text{ortho (1), meta (2), para (3)} \) containing two Lewis base centers \( C_\text{6}\text{H}_4\text{N} \) of ethynyl and \( N \) atom of pyridine. The efficient preparation of the iron arylethynyls complexes \( \text{Cp(CO)}_2\text{Fe-C≡C-}\text{Ar} \) by Pd/Cu-catalyzed cross-coupling of \( \text{Cp(CO)}_2\text{FeI} \) with terminal arylacetylenes inspired us to apply the Oshima protocol for the preparation of 1–3. Unfortunately, our first attempt to obtain the \( \text{ortho-pyridylethynyl iron complex \( \text{Cp(CO)}_2\text{Fe-C≡C-(2-C}_5\text{H}_4\text{N) \) (1 \( \text{using the Oshima conditions did not give the target substance. Upon an increase in the reaction temperature to 60 °C an unexpected binuclear \( \text{FePd \ μ-pyridylvinylidene complex \[ \text{[Cp(CO)}_2\text{Fe(μ-\text{η}^1\text{C}_5\text{H}_4\text{N})\text{-η}^1\text{C}_5\text{H}_4\text{N}\text{-κ}^2\text{N}\text{-C}_5\text{H}_4\text{N})\text{-C}_6\text{H}_4\text{H}\text{[μ-\text{CO]}\text{Pd]} \right\] (4 \)) \) \) \) \( \text{was obtained in yield of 2% \text{, but still without traces of 1 (Scheme 1). The 12% yield of complex 4 \) \) \( \text{was achieved by using one equivalent of \( \text{PdCl}_2 \), without copper iodide and pure diisopropylamine as a solvent (Scheme 1). Attempts to synthesize the iron derivatives of meta- and para-pyridylethynyls under described conditions were also unsuccessful; the iron acetylide compounds were obtained in very low yields (less than 5%) although no formation of the side binuclear \( \text{FePd products was observed in these cases. For that reasons, we became interested in developing a simple and reliable synthetic approach to σ-acetylide iron complexes 1–3 based on the Pd/Cu-catalyzed cross-coupling reactions. We proposed that the inability to obtain the pyridylethynyl iron complexes 1–3 from \( \text{Cp(CO)}_2\text{FeI} \) and ethynylpyridines under the Oshima conditions was primarily caused by a suppression of the formation and transfer of pyridylethynyl moieties to the palladium atom (Pd/Cu-catalyzed reactions) or of the formation of this moiety at the Fe-Pd center from the \( \text{π} \)-coordinated alkyne (Cu-free reaction). Here we report on the cross-coupling of terminal ethynylpyridines and (trimethylsilyl) ethynylpyridines with cyclopentadienyliron dicarbonyl iodide under different Pd/Cu-catalyzed conditions to obtain the target iron pyridylethynyl complexes.

Results and discussion

Development of Pd/Cu-catalyzed approaches to the synthesis of \( \text{Cp(CO)}_2\text{Fe-C≡C-(n-C}_5\text{H}_4\text{N)} \) \( n = \text{ortho (1), meta (2), para (3)} \) and \( \text{Cp(CO)}_2\text{Fe-C≡C-(4-C}_5\text{H}_4\text{N}_2\text{S) \) \( (5 \) \) \) \)

Following on from the hypotheses reasons of the inability to obtain the ethynylpyridyl iron complexes \( \text{Cp(CO)}_2\text{Fe-C≡C-(n-C}_5\text{H}_4\text{N)} \) \( n = \text{ortho (1), meta (2), para (3)} \), we realized that facilitation of the transmetallation step (the formation and the transfer of acetylide species) in Pd/Cu-catalyzed reaction between \( \text{Cp(CO)}_2\text{FeI} \) and ethynylpyridines should allow the preparation of the desired products 1–3. Therefore, two approaches were proposed to solve this problem: (i) the use of (trimethylsilyl)ethynylpyridines that in the presence of fluoride ion generate anionic pentacoordinate silicate species \( \text{Me}_3\text{SiF-C≡C-Pyr}^- \), in which the alkynyl group is more nucleophilic and should be smoothly transferred to a palladium catalyst (Hiyama coupling); (ii) the application of stronger base than secondary and tertiary amines such as 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) to facilitate the acetylide species formation in Pd/Cu-catalyzed reactions of \( \text{Cp(CO)}_2\text{FeI} \) with terminal ethynylpyridines. Our preliminary experiments suggested that both approaches can be used.

The palladium(II) complexes bis(triphenylphosphine)palladium(II) dichloride \( \text{PdCl}_2(\text{PPh}_3)_2 \) and bis(acetonitrile)palladium dichloride \( \text{PdCl}_2(\text{NCMe})_2 \), as well as tris(dibenzylideneacetone) dipalladium(0) \( \text{Pd}_2(\text{dba})_2 \) were tested as catalysts in this study. However, in our first experiments an application of \( \text{PdCl}_2(\text{PPh}_3)_2 \) as catalyst was found to results in an unexpected substitution of CO ligands by PPh$_3$ in \( \text{[Cp(CO)}_2\text{Fe]} \) fragments to give the side-products \( \text{Cp(CO)}(\text{PPh}_3)_2\text{FeI} \) and \( \text{Cp(CO)}(\text{PPh}_3)_2\text{Fe-Pd-C≡C-(2-C}_5\text{H}_4\text{N)} \), thereby making the separation of the reaction mixture more difficult. Therefore, in our following experiments we applied only such palladium catalysts that do not contain ligands capable to substitute carbonyl groups in the initial and the target compounds, namely \( \text{Pd}_2(\text{dba})_2 \) and \( \text{PdCl}_2(\text{NCMe})_2 \). The influence of the presence of CuI cocatalyst on the reaction outcome was also examined.

The application of the “(trimethylsilyl)ethynylpyridine” approach resulted in only moderate yields (up to 60%) of the target complexes 1–3; the radical dimerization product \( \text{[Cp(CO)}_2\text{Fe]}^2 \) (in 23–37% yield) and the corresponding 1,4-di(\( n \)-pyridyl)buta-1,3-dienes (trace amounts) were also obtained. The nature of the palladium catalyst \( \text{PdCl}_2(\text{NCMe})_2 \) or \( \text{Pd}_2(\text{dba})_2 \) didn’t affect the yields of 1–3 here, as well as when the presence of CuI as cocatalyst (Table 1). However, a source of...
The presence of CuI in these conditions is important for the catalytic formation of the complexes [Cp(CO)₂Fe]₂ (3%), [Pd(CO)]₂[PPh₃]FeI (3%), and also, [μ-pyridylvinylidene] complexes (entries 3, 8, 11 in Table 1). It should be noted that for the reactions between Cp(CO)₂Fe and ethynylpyridines we decided to apply the same conditions of catalytic formation of complex 1. The proposed catalytic cycles were obtained in the ethynylation of Cp(CO)₂FeI with [(trimethylsilyl)ethynyl]pyridines (entries 3, 8, 11 in Table 1).

Therefore, all reactions given in Table 3 were conducted at 36 °C with 10 mol% of Pd₂(db₃). To emphasize the applicability of the Pd/Cu-catalyzed pyridylethynylation of Cp(CO)₂FeI to the synthesis of similar compounds, we tried to prepare the corresponding σ-4-benzothiadiazolylenyl complex Cp(CO)₂Fe–C≡C–(4-C₆H₄N₂S) (5). Following the developed strategies, we investigated the coupling reactions of Cp(CO)₂FeI with 4-(trimethylsilyl)ethynyl-2,1,3-benzothiadiazole and 4-ethyl-2,1,3-benzothiadiazole. In all experiments performed using both approaches complex 5 was obtained. However, its highest yield 94% was achieved only in the coupling of Cp(CO)₂FeI and H–C≡C–(4-C₆H₄N₂S) at 60 °C using 1 mol% of PdCl₂(NCMe)₂ and 5 mol% of CuI (Scheme 2, entry 3).

**The proposed catalytic cycles**

Previously, the Lo Sterzo group clearly demonstrated a close analogy between the Pd-catalyzed carbon–carbon and metal–carbon bond formation in the coupling of cyclopentadienyliron dicarbonyl iodide and [(trimethylsilyl)ethynyl]pyridines to form amidyl complexes.
Table 2  Pd(II)/CuI- and Pd(II)-catalyzed coupling reactions of cyclopentadienyliron dicarbonyl iodide and ethynylpyridines

| #  | Pyr | [Pd], mol% | Cul, mol% | Conditions | Conversion, % | Target complex | Byproducts |
|----|-----|------------|-----------|------------|---------------|----------------|------------|
| 1  | | PdCl2(PPh3)2, 10 mol% | 10 | 40 °C; 30 min | 86 | 1 – 66% | Cp(CO)(PPh3)Fe (3%); Cp(CO)(PPh3)Fe-C≡C–(2-C6H4N) (8%) |
| 2  | | PdCl2(NCMe)2, 10 mol% | 10 | 24 °C; 20 min | 98 | 1 – 91% | [Cp(CO)2Fe]2 (5%); |
| 3  | | PdCl2(NCMe)2, 2 mol% | 20 | 24 °C; 20 min | 97 | 1 – 94% | [Cp(CO)2Fe]2 (2%); |
| 4  | | PdCl2(NCMe)2, 1 mol% | 5 | 60 °C; 30 min | 98 | 1 – 92% | [Cp(CO)2Fe]2 (4%); |
| 5  | | PdCl2(NCMe)2, 5 mol% | — | 60 °C; 30 min | 29 | 1 – 15% | [Cp(CO)2Fe]2 (9%); 4 – 3% [Fe], 55% [Pd]b |
| 6  | | PdCl2(NCMe)2, 10 mol% | 10 | 24 °C; 20 min | 97 | 3 – 91% | — |
| 7  | | PdCl2(NCMe)2, 2 mol% | 20 | 24 °C; 20 min | 96 | 3 – 92% | — |
| 8  | | PdCl2(NCMe)2, 1 mol% | 5 | 60 °C; 30 min | 99 | 3 – 96% | — |
| 9  | | PdCl2(NCMe)2, 5 mol% | — | 60 °C; 90 min | 75 | 3 – 67% | [Cp(CO)2Fe]2 (7%); |
| 10 | | PdCl2(NCMe)2, 10 mol% | 20 | 24 °C; 20 min | 99 | 2 – 83% | [Cp(CO)2Fe]2 (13%); |
| 11 | | PdCl2(NCMe)2, 2 mol% | 20 | 24 °C; 20 min | 98 | 2 – 93% | — |
| 12 | | PdCl2(NCMe)2, 1 mol% | 5 | 60 °C; 30 min | 99 | 2 – 95% | — |
| 13 | | PdCl2(NCMe)2, 5 mol% | — | 60 °C; 90 min | 74 | 2 – 66% | [Cp(CO)2Fe]2 (6%); |

a All reactions were carried out with use of 1 equiv. Cp(CO)2FeI, 1.5 equiv. H–C≡C–(n-C6H4N), in the presence of 1.5 equiv. DBU in THF. b Isolated yield of 4 based on Cp(CO)2FeI and PdCl2(NCMe)2 consumption.

Table 3  Pd(0)/CuI- and Pd(0)-catalyzed coupling reactions of cyclopentadienyliron dicarbonyl iodide and ethynylpyridines

| #  | Pyr | [Pd], mol% | Cul, mol% | Conditions | Conversion, % | Target complex | Byproducts |
|----|-----|------------|-----------|------------|---------------|----------------|------------|
| 1  | | PdCl2(PPh3)2, 10 mol% | 10 | 36 °C; 90 min | 96 | 1 – 84% | [Cp(CO)2Fe]2 (8%); [C≡C–(2-C6H4N)]2 (3%); |
| 2  | | PdCl2(NCMe)2, 10 mol% | 10 | 36 °C; 90 min | 95 | 1 – 64% | [Cp(CO)2Fe]2 (17%); 4 – 7% [Fe], 72% [Pd]b |
| 3  | | PdCl2(NCMe)2, 2 mol% | 10 | 36 °C; 90 min | 95 | 3 – 86% | [Cp(CO)2Fe]2 (6%); [C≡C–(4-C6H4N)]2 (2%); |
| 4  | | PdCl2(NCMe)2, 1 mol% | 10 | 36 °C; 90 min | 96 | 3 – 67% | [Cp(CO)2Fe]2 (20%); [C≡C–(4-C6H4N)]2 (4%); |
| 5  | | PdCl2(NCMe)2, 5 mol% | 10 | 36 °C; 90 min | 96 | 2 – 87% | [Cp(CO)2Fe]2 (7%); [C≡C–(3-C6H4N)]2 (3%); |

a All reactions were carried out with use of 1 equiv. Cp(CO)2FeI, 1.5 equiv. H–C≡C–(n-C6H4N), in the presence of 1.5 equiv. DBU in THF. b Isolated yield of 4 based on Cp(CO)2FeI and PdCl2(NCMe)2 consumption.

carbon bond formation mechanisms through reactions between \([\eta^3\text{-1-diphenylphosphino-2,4-diphenyl)cyclopentadienyl]}(CO)\)_2-
MI (M = Mo, W) and PdCl2(NCMe)2 with a following treatment of the
resulting binuclear MPd oxidation addition complexes with
4-nitro-1-[2-(tributhylstannyl)ethyl]benzene.\(^{27,28}\) The mecha-
nism of the Pd/Cu- and Pd-catalyzed pyridylethylation of
cyclopentadienyliron dicarbonyl iodide should be apparently
analogous to that of the carbon–carbon coupling reactions. Some
additional clues to the mechanisms of the formation of the
complexes 1–3, 5 may be provided by a consideration of the
reactions between Cp(CO)2FeI and 2-ethynylpyridine. The formation of
the binuclear FePd μ-pyridylvinilidenic complex 4 in
the reactions under Cu-free conditions (Table 2 entries 5, 9, 13
and Table 3 entries 2, 4) clearly shows that the reaction pathways
leading to 1 and 4 may have common intermediates. Moreover,
one can assume that in the reaction performed with the sto-
ichiometric ratios of Pd2(dbaj3, Cp(CO)2FeI and H–C≡C–(2-
C6H4N), the ratio of 1 to 4 should change in favor of the latter.
Indeed, the reaction of Cp(CO)2FeI with 2-ethynylpyridine in
the presence of 1 equiv. of Pd2(dbaj3 in triethylamine as a solvent
gave complex 4 in yield of 78% (Scheme 3). However, the trace amount (3% yield) of the 2-pyridylethynyl complex 1 was obtained, thereby confirming that the catalytic 2-pyridylethynylation of Cp(CO)2FeI occurs even under these conditions, although in a much lesser degree.

Obviously, one of these common intermediates in the reaction pathways to 1 and 4 is binuclear FePd species (Scheme 4, A) formed by an oxidative addition of Cp(CO)2FeI to catalytically active Pd0 species. In the absence of an efficient transmetallating agent, such as the copper acetylide or anionic pentacoordinate silicate adduct [Me3Si(F)-C≡C-(2-C5H4N)]NBu4, a π-coordination of the alkyne to A can occur resulting in complex A’. Without a strong base A’ rearrange into the μ-pyridylvinylnidene complex 4. To reveal the mechanism of this process, an additional study is needed, since there are several possible reaction pathways for its proceeding. It is worth noting that the examples of the acetylene-to-vinylidene tautomerism involving two adjacent metal centers are known and described in several works,66-70 but they are much less studied in contrast to the well-understood acetylene-to-vinylidene rearrangement mediated by a single metal center.71,72 It is also worth to note that the ortho-position of the nitrogen atom in 2-ethynylpyridine plays an important role in the process of formation of the binuclear FePd μ-pyridylvinylnidene complex 4, as the Pd-catalyzed reactions of Cp(CO)2FeI with 3- and 4-ethynylpyridines simply resulted in a decrease of yields of 2 and 3, and didn’t give any binuclear products (Table 2 entries 9 and 13).

Taking into account the data obtained and the known mechanistic aspects of the cross-coupling reactions, a noncontradictory mechanistic scheme of the pyridylethynylation of cyclopentadienylinidron dicarboxyliodide was proposed (Scheme 4), which explains the reaction pattern and the possibility of alternative reaction pathway. For the Pd/Cu-catalyzed reactions (pathway A), the first step of the catalytic cycle would be an oxidative addition of the Cp(CO)2FeI to the catalytically active Pd0 species to give a binuclear FePd intermediate A. A subsequent transmetallation, where transfer of the pyridylacetyliden moiety from the Cu-acetylide formed in the copper-cycle to the FePd complex A led to a complex B, which undergoes trans to cis isomerization to give a complex B’, a following reductive elimination of the latter results in the Fe–C cross-coupling products Cp(CO)2Fe–C≡C–(4-C6H4N2S) (n = ortho (1), meta (2), para (3)). The initial oxidative addition and the final reductive elimination steps of a mechanism for the Cu-free reaction (pathway B) are the same as for the Pd/Cu-catalyzed one. However, an alternative ligand L substitution in the intermediate A by the alkyne can take place resulting in π-alkyne complex A’. Its subsequent deprotonation then occurs to give the complex B’. The generation of the complex 4 in this context can be considered as a side pathway C of the Pd-catalyzed reaction between Cp(CO)2FeI and 2-ethynylpyridine.
that is facilitated by the absence of transmetallating agents or appropriate base, and ortho-position of the nitrogen atom in the pyridine ring of the alkyne. The catalytic cycle for the complex formation should be the same as for 1–3.

The results obtained in the case of the “(trimethylsilyl)ethynylpyridine” demonstrated that pyridylethynylation of Cp(CO)₂FeI should proceed according to the reaction pathway commonly accepted for the Hiyama coupling, i.e. transfer of the pyridylacetylide moiety to the FePd complex A should go through anionic pentacoordinate silicate species (Scheme 5).

### Characterization of the complexes

The IR and the \(^1\text{H}\) and \(^{13}\text{C}\{^1\text{H}\} NMR data for the complexes 1–3 and 5 were obtained. The NMR signals were assigned on the basis of \(^1\text{H}\)–\(^{13}\text{C}\) correlations measured through HSQC and HMBC experiments, respectively (Table 4). The structure of the complexes can be deduced from the combined NMR and IR data.

The IR spectra of 1–3 in CH₂Cl₂ solution show two very strong absorptions at about 2043 and 1996 cm⁻¹ that

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Scheme 4 Proposed mechanism of Pd/Cu- and Pd-catalyzed coupling reactions of cyclopentadienyliron dicarbonyl iodide and ethynlypyridines.

Scheme 5 Proposed mechanism of Pd-catalyzed coupling reactions of cyclopentadienyliron dicarbonyl iodide and [((trimethylsilyl)ethynyl] pyridines.
The proximity of crystals 1 and 2 cell parameters (Table S3†) and small differences in the sequence of atoms in their molecules suggest that there are small differences in a packing of molecules. Indeed, the structures 1 and 2 are mirror-like each other, and the mirror plane is perpendicular to the cell axis b. The iron atom in 1–3 and 5 is coordinated by a cyclopentadienyl

The molecular structures of the complexes Cp(CO)2Fe–C≡C–(n-C5H5N) [n = ortho (1), meta (2), para (3)] and Cp(CO)2Fe–C≡C–(4-C5H4N2S) (5) were solved on the basis of X-ray diffraction data. Suitable crystals of 1–3 and 5 were grown from dichloromethane/hexane mixtures. The views of the structures are shown in Fig. 1, selected bond lengths and angles are given in Table 5. The crystal data and refinement parameters are included in the ESL†

[Table 4] IR and NMR (δ, ppm [J, Hz]) spectroscopic data for the Cp(CO)2Fe–C≡C–(n-C5H5N) [n = ortho (1), meta (2), para (3)] and Cp(CO)2Fe–C≡C–(4-C5H4N2S) (5)

| NMR (δ, ppm [J, Hz], CD2Cl2) |   |   |   |   |   |
|-----------------------------|--|--|--|--|--|
| 1H                         | 13C |   |   |   |   |
| C5H5 | Cα | Cβ | Cα | CO | IR, cm⁻¹ (CD2Cl2) |
| 1   | 5.14 | 116.7 | 95.4 | 85.4 | 212.2 | 2112 (v(νC=O), 2043, 1997 (νC=O) |
| 2   | 5.13 | 112.3 | 95.7 | 85.3 | 212.4 | 2111 (ν(νC=O), 2044, 1996 (νC=O) |
| 3   | 5.13 | 114.3 | 102.0 | 85.6 | 212.3 | 2110 (ν(νC=O), 2044, 1998 (νC=O) |
| 4   | 5.20 | 112.3 | 102.9 | 85.6 | 212.3 | 2100 (ν(νC=O), 2042, 1997 (νC=O) |

The 13C nuclei of α- and β-alkynyl carbons of 1–3 and 5 resonate in the regions δ 112–117 and 95–103 ppm, respectively. It is noteworthy that these signals slightly vary with the nature of the substituents in σ-alkynyl ligands. The downfield shift of Cα nuclei by approximately 2 ppm is observed on moving from the meta- to para- and to ortho-pyridylethynyl iron complexes. However, this trend doesn’t hold for the Cβ nuclei, here only the signal of 3 is downfield shifted by 7 ppm compared with those of 1 and 2. These correlations are apparently due to changes in π-electron distribution induced by the electronic effect of the pyridyl fragments. However, these effects can’t be reduced to a single parameter (like only inductive or mesomeric effects), as was showed by Claude Lapinte team for cationic complexes of the type [Cp*(dppe)Fe–C≡C–(n-C5H5N)[PF6]] (n = 2, 3, 4). At the same time, the 13C chemical shifts of the cyclopentadienyl and carbonyl groups coordinated to the iron atom are almost independent of their nature (Table 4). The presence of the pyridyl substituents in 1–3 is indicated by signals between 7.02 and 8.50 ppm in 1H NMR spectra, and between 119 and 152 ppm in 13C NMR spectra of the complexes. The carbon and hydrogen atoms of the 2,1,3-benzothiadiazole group in 5 resonate in the regions δ 118–156 and δ 7.48–7.76 ppm in 13C and 1H NMR spectra, respectively. So, overall the IR and NMR spectra parameters of 1–3 and 5 are similar to those found for analogous cyclopentadienyliron dicarbonyl complexes with different σ-alkynyl ligands.

The molecular structures of the complexes Cp(CO)2Fe–C≡C–(2-C5H5N) (1), Cp(CO)2Fe–C≡C–(3-C5H5N) (2), Cp(CO)2Fe–C≡C–(4-C5H5N) (3), and Cp(CO)2Fe–C≡C–(4-C5H4N2S) (5) were showed by Claude Lapinte team for cationic complexes of the type [Cp*(dppe)Fe–C≡C–(n-C5H5N)[PF6]] (n = 2, 3, 4). At the same time, the 13C chemical shifts of the cyclopentadienyl and carbonyl groups coordinated to the iron atom are almost independent of their nature (Table 4). The presence of the pyridyl substituents in 1–3 is indicated by signals between 7.02 and 8.50 ppm in 1H NMR spectra, and between 119 and 152 ppm in 13C NMR spectra of the complexes. The carbon and hydrogen atoms of the 2,1,3-benzothiadiazole group in 5 resonate in the regions δ 118–156 and δ 7.48–7.76 ppm in 13C and 1H NMR spectra, respectively. So, overall the IR and NMR spectra parameters of 1–3 and 5 are similar to those found for analogous cyclopentadienyliron dicarbonyl complexes with different σ-alkynyl ligands.

The proximity of crystals 1 and 2 cell parameters (Table S3†) and small differences in the sequence of atoms in their molecules suggest that there are small differences in a packing of molecules. Indeed, the structures 1 and 2 are mirror-like each other, and the mirror plane is perpendicular to the cell axis b. The iron atom in 1–3 and 5 is coordinated by a cyclopentadienyl
ring in \( \eta^1 \)-fashion, two terminal CO ligands, and by \( \sigma \)-alkynyl ligand to adopt a typical pseudo-octahedral geometry. The bond lengths and angles in 1–3 and 5 are close to each other and to those found in the known complexes of the type Cp(CO)\(_2\)FeC\( \equiv \)C-\( C\equiv \)C-R.\(^{7,4-77}\) The differences in geometry of the complexes’ molecules are associated with the position of C\( \equiv \)C-R moiety relative to the iron fragment. The pyridylethyln ligands in 1 and 2, and the 4-ethynyl-2,1,3-benzothiadiazolyl ligand in 5 lean slightly toward one of the carbonyl group of the iron atom in such a way that the planes between ring of the substituent and [Fe(CO)\(_2\)] fragment exhibit angles about 81°. At the same time, in the complex 3 the \( \eta^2 \)-pyridylethyln ligand is almost perpendicular to the plane formed by two carbonyl ligands and the iron atom. Moreover, the angles between the planes of the substituent’s ring in the \( \sigma \)-alkynyl and the cyclopentadienyl ligands are about 70° in 3 and 5, whereas the analogous angles in 1 and 2 are ca. 85°. It is also of interest that the pyridyl substituents in 1 and 2 differ in their orientations: when the nitrogen atom in 1 is oriented to Cp ring, that in the complex 2 the nitrogen is placed opposite, \textit{i.e.} oriented to carbonyl groups.

## Conclusions

In this paper, two approaches were developed for the pyridylethynylation of cyclopentadienyliron dicarbonyl iodide based on Pd/Cu- and Pd-catalyzed Fe-C coupling of \((\text{trimethylsilyl})\text{ethynylpyridines and ethynylpyridines with C} \text{p(CO)} \text{2Fe} \). Although these two approaches differ only in the nature of species participating in the transmetallation step (transfer of the pyridylacetylide moiety to catalytically active species), the second “ethynylpyridine” approach was found to be much more effective compared with the first “\((\text{trimethylsilyl})\text{ethynylpyridine} \)” one. For example, the \( \sigma \)-pyridylethynyl iron complexes Cp(CO)\(_2\)Fe-C\( \equiv \)C-(n-C\(_5\)H\(_4\)) \( n = \text{ortho} (1), \text{meta} (2), \text{para} (3) \) were obtained in highest yields (about 95%) in the reaction of ethynylpyridines with Cp(CO)\(_2\)FeI in THF in the presence of 1,8-diazabicyclo[5.4.0]undec-7-ene as a base upon the catalysis with 1 mol% of PdCl\(_2\)(NCMe)\(_2\) and 5 mol% of CuI and heating at 60 °C for 30 minutes. In case of the Cu-free coupling between Cp(CO)\(_2\)FeI and 2-ethynylpyridine, there was an additional side-reaction, which resulted in traces of the binuclear complex [Cp(CO)FeC\( \equiv \)C] with 1 mol% of PdCl\(_2\)(NCMe)\(_2\) and 5 mol% of CuI and heating at 60 °C for 30 minutes. In case of the Cu-free coupling between Cp(CO)\(_2\)FeI and 2-ethynylpyridine, there was an additional side-reaction, which resulted in traces of the binuclear complex [Cp(CO)FeC\( \equiv \)C] with 1 mol% of PdCl\(_2\)(NCMe)\(_2\) and 5 mol% of CuI and heating at 60 °C for 30 minutes. In case of the Cu-free coupling between Cp(CO)\(_2\)FeI and 2-ethynylpyridine, there was an additional side-reaction, which resulted in traces of the binuclear complex [Cp(CO)FeC\( \equiv \)C] with 1 mol% of PdCl\(_2\)(NCMe)\(_2\) and 5 mol% of CuI and heating at 60 °C for 30 minutes. In case of the Cu-free coupling between Cp(CO)\(_2\)FeI and 2-ethynylpyridine, there was an additional side-reaction, which resulted in traces of the binuclear complex [Cp(CO)FeC\( \equiv \)C] with 1 mol% of PdCl\(_2\)(NCMe)\(_2\) and 5 mol% of CuI and heating at 60 °C for 30 minutes. In case of the Cu-free coupling between Cp(CO)\(_2\)FeI and 2-ethynylpyridine, there was an additional side-reaction, which resulted in traces of the binuclear complex [Cp(CO)FeC\( \equiv \)C]. The oxidative addition species Cp(CO)\(_2\)Fe-PdL\(_2\)H \((A \text{ in Scheme 4}) \) are supposed to be a key intermediate in the catalytic cycle of the \( \sigma \)-alkynyl complexes formation, however in the absence of effective transmetallating agents or appropriate base \( A \) is assumed to become a precursor to 4.

The developed methodology can be extended to the synthesis of other \( \sigma \)-alkynyl complexes of iron. So, the coupling of cyclopentadienyliron dicarbonyl iodide and 4-ethynyl-2,1,3-benzothiadiazole under the same condition as above was found to proceed efficiently resulting in the complex Cp(CO)\(_2\)Fe-C\( \equiv \)C-(4-C\(_6\)H\(_5\)N\(_2\)) \((5) \) of the spectroscopic and structural features of 1–3, 5 were described.

## Experimental

### General considerations

All operations and manipulations were carried out under an argon atmosphere. Solvents (dichloromethane, petroleum ether, ethyl acetate, hexane, triethylamine) were purified by distillation from the appropriate drying agents and stored under argon. THF was dried by refluxing over sodium/benzophenone ketyl and freshly distilled prior to use. The course of reactions was monitored by TLC on Silica gel (Alu foils, Sigma-Aldrich) and IR spectroscopy. Neutral silica gel (silica 60, 0.2–0.5 mm, Macherey-Nagel) was used for column chromatography. Pd(Cl)\(_2\)(PPh\(_3\))\(_2\)\(^{82}\), Pd(Cl)\(_2\)(NCMe)\(_2\)\(^{79}\), Pd\(_2\)C\(_3\)(dba)\(_2\), CHCl\(_3\)\(^{80}\) and Cp(CO)\(_2\)Fe\(^{84}\) were prepared according to literature procedures. Ortho, meta, \textit{para} \( -[(\text{trimethylsilyl}) \) ethynyl]pyridines\(^{82}\) and 4-[(\text{trimethylsilyl})ethyl-2,1,3-benzothiadiazole\(^{83}\) were prepared from ethynyltrimethylsilane. Ortho- and \textit{para}-ethynylpyridines were synthesised from 2-methyl-4(\text{pyridyl})but-3-yn-2-ols by elimination of acetone, according to

| Distances, Å | 1 | 2 | 3 | 5 |
|--------------|---|---|---|---|
| Fe–C3        | 1.9159(17) | 1.919(2) | 1.906(3) | 1.917(3) |
| C3–C9        | 1.204(2) | 1.200(3) | 1.203(5) | 1.208(4) |
| C9–C10       | 1.436(2) | 1.437(3) | 1.423(4) | 1.435(4) |
| Fe–Cp        | 1.7223(8) | 1.7200(11) | 1.7186(19) | 1.7222(16) |

| Bond angles, † | Fe–C3-C9 | C3–C9-C10 | Fe–C3–Fe–C1 | C3–Fe–C2 |
|----------------|----------|-----------|-------------|----------|
| 175.94(16)     | 176.54(19) | 89.26(8)  | 81.38(6)   |
| 174.3(2)       | 176.8(3)  | 86.8(1)   | 80.97(7)   |
| 179.4(4)       | 177.7(5)  | 89.68(17) | 88.27(8)   |
| 178.6(3)       | 174.6(4)  | 89.07(16) | 88.0(13)   |
| 178.4(3)       | 178.9(4)  | 89.07(16) | 89.56(14)  |

| Angles between planes, ‡ | \( \eta^1 \)-Cp:ring | (O1–C1–Fe–C2–O2):ring |
|--------------------------|---------------------|----------------------|
| \( \eta^1 \)-Cp:ring     | 84.51(7)            | 81.38(6)             |
| (O1–C1–Fe–C2–O2):ring   | 84.58(8)            | 80.97(7)             |

\( \eta^1 \)-Cp:ring denotes the pyridylacetylide moiety to catalytically active species, \( \eta^1 \)-ethynylpyridine denotes the pyridylacetylide moiety to catalytically active species, and \( \eta^1 \)-alkynyl denotes the pyridylacetylide moiety to catalytically active species. The selected bond distances, bond lengths, and angles in 1–3 and 5 are given.

\( \eta^1 \)-Cp:ring denotes the pyridylacetylide moiety to catalytically active species, \( \eta^1 \)-ethynylpyridine denotes the pyridylacetylide moiety to catalytically active species, and \( \eta^1 \)-alkynyl denotes the pyridylacetylide moiety to catalytically active species.
published procedures. Meta-ethenylpyridine and 4-ethenyl-2,1,3-benzothiadiazole were obtained by desilylation of meta-[(trimethylsilyl)ethyl]pyridine and 4-(trimethylsilyl)ethyl-2,1,3-benzothiadiazole. Tetrabutylammonium fluoride solution (1 M solution in THF, Aldrich), 1,8-diazabicyclo[5.4.0]undec-7-ene (Aldrich), and catalyst Cul (“Vekton-M” Ltd.) were purchased and used directly. The synthetic procedures for the preparation of 1–3 and 5 providing the highest yields are described in the experimental part (Table 2, entries 4, 8, and 12; Scheme 2, entry 3), the detailed isolation procedures and characterization of the products can be found in the ESL.

Physical-chemical characteristics were obtained in the Krasnoyarsk Regional Centre of Research Equipment, Siberian Branch of the Russian Academy of Sciences. The IR spectra were recorded on the Shimadzu IR Tracer-100 spectrometer (Japan). 

**Synthesis of Cp(CO)2Fe-C≡C-(2-C5H4N) (1)**

Cyclopentadienyliron dicarbonyl iodide (315 mg, 1.036 mmol) and 2-ethenylpyridine (0.155 mL, 158 mg, 1.534 mmol) were dissolved in benzene (10 mL), then DBU (0.23 mL, 234 mg, 1.539 mmol), PdCl2(NCMe)2 (3 mg, 0.012 mmol), and Cul (10 mg, 0.052 mmol) were added. The reaction mixture was stirred at 60 °C for 30 minutes, and then was evaporated to dryness; the residue was dissolved in dichloromethane and passed through a pad (0.5 cm) of silica gel by using ethyl acetate as an eluent. The filtrate was concentrated in vacuo and chromatographed on silica gel (9 × 2 cm). A dark-yellow fraction containing the σ-alkenylpyridine iron complex was eluted with petroleum ether-ethyl acetate (9 : 1), (4 : 1), (3 : 2) mixtures and subsequently with ethyl acetate. The complex Cp(CO)2Fe-C≡C-(2-C5H4N) (1) was isolated as brown-yellow solid after evaporation of the solvent (266 mg, 0.953 mmol, 92%). $^1$H NMR (CD2Cl2, 25 °C) δ ppm [J, Hz]: 5.14 (s, 5H, C5H4N); 7.02 (dd, 1H, $^3$JHH = 7.6 Hz, $^3$JHH = 5.3 Hz, $^4$JHH = 0.8 Hz, H$a$ of (2-Pyr) [H$^a$]); 7.32 (d, 1H, $^3$JHH = 8.0 Hz, H$a$ of (2-Pyr) [H$^a$]); 7.53 (dd, 1H, $^3$JHH = 7.7 Hz, $^4$JHH = 1.6 Hz, H$\eta$ of (2-Pyr) [H$\eta$]); 8.40 (d, 1H, $^3$JHH = 5.3 Hz, N-C-H$\eta$ of (≡C)-(2-Pyr) [H$^a$]); 13C{1H} NMR (CD2Cl2, 25 °C) δ ppm [J, Hz]: 85.4 (s, C1H); 95.4 (s, C2H; 116.7 (s, C3H; 119.9 (s, C$\eta$ of (≡C)-(2-Pyr) [C$^\eta$]); 125.9 (s, N-C$\eta$ of (≡C)-(2-Pyr) [C$^\eta$]); 135.3 (s, C$\eta$ of (≡C)-(2-Pyr) [C$^\eta$]); 145.8 (s, C$\eta$ of (≡C)-(2-Pyr) [C$^\eta$]); 149.0 (s, C$\eta$ of (≡C)-(2-Pyr) [C$^\eta$]); 212.2 (s, 2Fe-CO-). IR (CHCl3, ν/cm$^{-1}$): 2112s (vC≡C), 2043vs, 1997vs (vC≡C), 1582m, 1556w, 1460m (vC≡C and vC=N). Anal. found: C, 60.39%; H, 3.24%; N, 5.00%. Calc. for C14H9FeNO2 (279): C, 60.25%; H, 2.35%; N, 5.02%.

**Synthesis of Cp(CO)2Fe-C≡C-(3-C5H4N) (2)**

Following the procedure described for preparation of 1, complex Cp(CO)2Fe-C≡C-(3-C5H4N) (2) was obtained with 95% yield (218 mg, 0.781 mmol) from cyclopentadienyliron dicarbonyl iodide (250 mg, 0.822 mmol) and 3-ethenylpyridine (127 mg, 1.223 mmol) with the use of THF (8 mL), DBU (0.18 mL, 183 mg, 1.204 mm), PdCl2(NCMe)2 (2 mg, 0.008 mmol), and Cul (8 mg, 0.042 mmol). $^1$H NMR (CD2Cl2, 25 °C) δ ppm [J, Hz]: 5.13 (s, 5H, C5H4N); 7.14 (dd, 1H, $^3$JHH = 7.3 Hz, $^3$JHH = 5.0 Hz, H$\eta$ of (3-Pyr) [H$^a$]); 7.54 (d, 1H, $^3$JHH = 8.0 Hz, H$a$ of (3-Pyr) [H$^a$]); 8.30 (d, 1H, $^3$JHH = 3.2 Hz, H$\eta$ of (3-Pyr) [H$^a$]); 8.49 (s, 1H, N-C-H$\eta$ of (3-Pyr) [H$^a$]). 13C{1H} NMR (CD2Cl2, 25 °C) δ ppm [J, Hz]: 85.3 (s, C1H); 95.7 (s, C2H; 112.3 (s, C3H; 122.4 (s, C$\eta$ of (3-Pyr) [C$^\eta$]); 121.6 (s, C$\eta$ of (3-Pyr) [C$^\eta$]); 145.3 (s, C$\eta$ of (3-Pyr) [C$^\eta$]); 152.1 (s, N-C$\eta$ of (3-Pyr) [C$^\eta$]); 212.4 (s, 2Fe-CO-). IR (CHCl3, ν/cm$^{-1}$): 2111s (vC≡C), 2049vs, 1996vs (vC≡C), 1570w, 1550w, 1479w (vC≡C and vC=N). Anal. found: C, 60.21%; H, 3.26%; N, 5.00%. Calc. for C14H9FeNO2 (279): C, 60.25%; H, 3.25%; N, 5.02%.
(0.014 mmol, 3% yield) of Cp(CO)₂Fe–C≡C–(2-C₅H₄N) (1). A recrystallization of Cp(CO)₂Fe[μ–C≡C–CH(2-C₅H₄N)]Pd from CHCl₃–hexane (1 : 2) mixture gave 176 mg of red-brown microcrystals. ¹H NMR (CD₂Cl₂, 25 °C) δ, ppm [J, Hz]: 5.25 (s, C₅H₅); 9.41 (d, J₁HH = 8.2, H₀₀₀ of (2-Pyr)); 7.07 (t, J₁HH = 5.9, H₀₀₀ of (2-Pyr)); 7.54 (d, J₁HH = 7.3, H₀₀₀ of (2-Pyr)); 9.41 (d, J₁HH = 3.4, N–C–H₀₀₀ of (2-Pyr)).¹³C (¹H) NMR (CD₂Cl₂, 25 °C) δ, ppm [J, Hz]: 88.3 (s, C₅H₅); 115.2 (s, C₀₀₀ of (2-Pyr)); 112.5 (s, C₀₀₀ of (2-Pyr)); 121.0 (s, C₀₀₀ of (2-Pyr)); 139.1 (s, C₀₀₀ of (2-Pyr)); 140.2 (s, C₀₀₀ of (2-Pyr)); 1599 (m, 1583 (m, 1546 (m, 1464 (m, 129.4 (s, Fe(2-Pyr)); 212.3 (s, 2Fe(2-Pyr)); 212.3 (s, 2Fe(2-Pyr)).

The structures were solved by the use of THF (9 mL), DBU (0.23 mL, 234 mg, 1.539 mmol) with the use of column chromatography on silica gel (9 cm). A 2031s, 1977vs (CH); 1532 (s, 2Fe(2-Pyr)); 2065 (s, Fe–CO); 230.77 (s, Fe–CObridging); 312.8 (s, C₀₀₀ of (2-Pyr)); 102.9 (s, C₀₀₀ of (2-Pyr)); 1599, 1553, 1546, 1464 (s, C₀₀₀ of (2-Pyr)). Anal. found: C, 32.78%; H, 1.95%; N, 2.73%. Calc. for C₁₄H₁₀FeINO₂Pd (513): C, 32.75; H, 1.96%; N, 2.73%.

**Synthesis of Cp(CO)₂Fe–C≡C–(4-C₅H₄N₂S) (5)**

Following the procedure described for preparation of 1–3, complex Cp(CO)₂Fe–C≡C–(4-C₅H₄N₂S) (5) was obtained from cyclopentadienyliron dicarbonyl iodide (309 mg, 1.016 mmol) and 4-ethyl-2,1,3-benzothiadiazole (245 mg, 1.531 mmol) with the use of THF (9 mL), DBU (0.23 mL, 234 mg, 1.539 mmol), PdCl₂(NCMe)₂ (3 mg, 0.012 mmol), and Cul (10 mg, 0.052 mmol). The product was isolated from the reaction mixture by column chromatography on silica gel (9 × 2 cm). A dark-yellow fraction containing the 6-4-benzothiadiazole-5-yl complex was eluted with petroleum ether-ethyl acetate (4 : 1 and 3 : 2) mixture. The complex Cp(CO)₂Fe–C≡C–(4-C₅H₄N₂S) (5) was obtained as brown-yellow solid after evaporation of the solvent (319 mg, 0.949 mmol, 94%). ¹H NMR (CD₂Cl₂, 25 °C) δ, ppm [J, Hz]: 5.21 (s, 5H, C₅H₅); 7.48 (dd, J₁HH = 7; J₁HH = 1.5, 1H, H₀₀₀ of (4-C₅H₄N₂S)); 7.50 (dd, J₁HH = 8.5; J₁HH = 7.0, H₀₀₀ of (4-C₅H₄N₂S)); 7.77 (dd, J₁HH = 8.4, J₁HH = 1.5, 1H, H₀₀₀ of (4-C₅H₄N₂S)).¹³C (¹H) NMR (CD₂Cl₂, 25 °C) δ, ppm [J, Hz]: 85.6 (s, C₅H₅); 102.9 (s, C₀₀₀); 112.3 (s, C₀₀₀); 118.1 (s, C₀₀₀ of (4-C₅H₄N₂S)); 121.1 (s, C₀₀₀ of (4-C₅H₄N₂S)); 129.4 (s, C₀₀₀ of (4-C₅H₄N₂S)); 131.0 (s, C₀₀₀ of (4-C₅H₄N₂S)); 1548 (s, N–C₀₀₀ of (4-C₅H₄N₂S)); 1557 (s, N–C₀₀₀ of (4-C₅H₄N₂S)); 212.3 (s, 2Fe–CO). IR (CH₂Cl₂, cm⁻¹): 2095 (s, Pd(0)); 2055s, 1848s (s, CO). Anal. found: C, 75.89%; H, 1.95%; N, 8.33%. Calc. for C₁₄H₁₀FeINO₂Pd (513): C, 75.89; H, 1.96%; N, 8.33%.

**X-ray diffraction studies**

Crystal data and X-ray experimental details for the complexes 1–3 and 5 are given in Table S4 (the ESI†). Brown-yellow crystals of the complexes suitable for X-ray crystallography analysis were obtained by evaporation of a solution of the complexes in a dichloromethane : hexane mixture = 1 : 2 under argon atmosphere at +5 °C. The experimental data were collected on a Smart Photon II diffractometer (Bruker AXS) with CCD area detector and graphite monochromator using Mo Kα radiation at room temperature. Absorption corrections have been applied using multisiscan procedure.⁶⁸ The structures were solved by direct methods and refined by full-matrix least squares on F², using SHELX programs.⁶⁶–⁷⁸ Hydrogen atoms have been placed in calculated positions and taken into account in the final stages of refinement in the “riding model” approximation.

**Author contributions**

Victor V. Verpekin: conceptualization, data curation, funding acquisition, investigation, methodology, supervision, visualization, writing – original draft, writing – review & editing; Oleg V. Semeikin: conductive investigation on Pd(0)-catalyzed coupling reactions of Cp(CO)₂Fe and [(trimethylsilyl)ethynyl] pyridines; Alexander D. Vasiliev: X-ray diffraction investigation; Alexander A. Kondrasenko: NMR investigation; Yuri A. Belousov: resources; Nikolai A. Ustynyuk: conceptualization, writing – review & editing. All authors have read and agreed to the published version of the manuscript.

**Conflicts of interest**

There are no conflicts to declare.

**Acknowledgements**

This research was funded by a grant from the Russian Science Foundation (project No. 18-73-00150).

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