Rhodocenium Functionalization Enabled by Half-Sandwich Capping, Zincke Reaction, Diazoniation and Sandmeyer Chemistry

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In continuation of our exploration of metalloccenium chemistry we report here on innovative ways toward monofunctionalized rhodocenium salts applying half-sandwich capping reactions of cyclopentadienyl rhodium(III) halide synthons with cyclopentadienyl ylides containing pyridine, phosphine or dinitrogen leaving groups, followed by Zincke and Sandmeyer reactions. Thereby amino, diazonio, bromo, azido and iodo rhodocenium salts containing valuable functional groups are accessible for the first time. Target compounds were characterized by spectroscopic (¹H/¹³C/¹⁹F-NMR, IR, HR-MS), structural (single crystal XRD) and electrochemical (CV) methods and their properties were compared to those of isoelectronic cobaltoce- nium compounds. These new functionalized rhodocenium complexes significantly expand the so far extremely limited chemical space of rhodocenium salts with promising options for the future development in the area of rhodocenium chemistry.

Introduction

Metallocenes in general constitute a huge class of important organometallic compounds with examples throughout the periodic table of elements. By far, the most representatives are ferrocenes with their many useful applications in coordination chemistry and catalysis, materials science, redox chemistry, medicinal chemistry, etc. In contrast, the isoelectronic 18 valence electron cationic metallocenes of group 9-cobaltoce- nium, rhodocenium and iridocenium-are very much less studied, due to less convenient synthetic accessibility and functionalization caused by their deactivating positive charge preventing “standard” organic derivatization protocols. Chemically, cobaltoce- nium, rhodocenium and iridocenium salts are interesting for a number of reasons: (i) They are highly stable and non-air-sensitive, (ii) they have a rich electrochemistry displaying usually reversible (Co) or irreversible (Rh, Ir) redox events, (iii) they are highly polar with high solubility in polar solvents, including water, (iv) they are strongly electron-withdrawing causing unusual reactivities with nucleophiles or of their appended functional groups, and-last but not least-(v) they represent a synthetic challenge even after 70 years of metallocene chemistry. Conceptually, we believe that function- alized Co/Rh/Ir metallocenium compounds will be valuable new building blocks in organometallic chemistry in general and in coordination chemistry in polar solvents, redox catalysis and redox sensing, or medicinal chemistry.

To this end, we have developed in the last couple of years new synthetic strategies in the area of cobaltoce- nium chemistry[2] and now we put further efforts into rhodocenium functionalization. As a first contribution, we recently published our results on rhodocenium monocarboxylic acid hexafluorido- phosphate and its derivatives[2] and here we report on nitrogen and phosphorus derivatives, including the very important, synthetically very valuable amino and diazonio rhodocenium salts.

Results and Discussion

Synthesis

In comparison to cobalt organometallic chemistry, a distinctive feature of rhodium chemistry is the higher stability of half-sandwich CpRhX₂ compounds (X = halogen)[3] that are obvious precursors for rhodocenium complexes by metathesis reactions with prefunctionalized Cp synthons (Scheme 1). To facilitate these transformations, the CpRhX₂ synthons were first con-
Scheme 1. Synthesis of functionalized mono and dicationic rhodocenium salts 2–10.

verted with AgPF$_6$ to substitution-labile [CpRh]$^{2+}$-solvato complexes, driven by the precipitation of insoluble AgX salts. Reaction of cyclopentadienyldiodorhodium dimer (1) with two equivalents of silver hexafluoridophosphate followed by two equivalents of pyridinium cyclopentadienide afforded air-stable pyridiniorhodocenium bis(hexafluoridophosphate (2a) in 82% yield. In a similar manner, pyridiniorhodocenium dinitrate (2b) was generated using silver nitrate instead of silver hexafluoridophosphate. From this compound, aminorhodocenium hexafluoridophosphate (3) was prepared in 66% yield by a Zincke-type reaction with aqueous ammonia. This is an operationally simple but mechanistically remarkable synthetic transformation: In the classical Zincke reaction, N-substituted pyridinium salts are generated from 1-(2,4-dinitrophenyl)pyridinium chloride with secondary amines (usually aniline derivatives) by a nucleophilic addition (AN), ring opening and ring closing (ANRORC) mechanism under formation of 2,4-dinitroaniline as “waste” product. Essentially, an amino group is formally exchanged from the more electron-rich aniline derivative to the electron-poor dinitrophenyl group. Here we are not interested in the usual Zincke-pyridinium product but in the amino side product. In an organometallic Zincke-type reaction, we use the strongly electron-withdrawing rhodocenium moiety as an electron-acceptor of similar strength to the Zincke salt, thereby realizing an atom-economic amino transfer from aqueous ammonia to rhodocenium with formation of unsubstituted pyridine and/or the Zincke aldehyde as the waste product(s). This synthetic protocol using pyridiniorhodocenium (2b) as an amino precursor was designed by us because unsubstituted CpNH$_2$ as a potential reagent for half-sandwich capping reactions does not exist, in contrast to unsubstituted CpNR$_2$ compounds. Potentially, this Zincke-amination protocol might be also applicable in the future to other organometallic salts, likely candidates would be dicaticonic pyridinio-substituted benzenetricarbonylmanganese or benzenecyclopentadienyliron complexes.

Aminorhodocenium hexafluoridophosphate (3) was selectively oxidized with the HgO/I$_2$ reagent in 89% yield to orange, water-soluble, dicationic dirhodoceniumdiazene (4), an azorhodocenium compound analogous to azocobaltocenium that became recently available by our previous work. Amino-rhodocenium hexafluoridophosphate (3) is clearly a highly valuable compound due to the very rich chemistry of the amino functional group in organic chemistry. Diazoniation under standard conditions in water afforded beige-colored diazonio-rhodocenium bis(hexafluoridophosphate) (5) in 60% yield, similar as the recently published diazoniation of aminocobaltocenium hexafluoridophosphate. Diazonium salts in general are privileged synths in organic chemistry allowing various nucleophilic and free-radical substitution reactions enabled by the superior leaving group N$_3$. Dicationic 5 is a highly reactive, special diazonium salt with similar unique properties.
as its cobaltocenium congener.\textsuperscript{14a} Reactions in solution are more or less restricted to nitromethane as solvent, other more common solvents lead to decomposition. Therefore, small-scale Sandmeyer reactions are preferably performed without solvent by ball-milling, e.g. reaction of solid 5 with cuprous bromide afforded bromorhodocenium hexafluoridophosphate (6) in 86\% yield, or in nitromethane solution, e.g. reaction with sodium azide gave azidorrhodocenium hexafluoridophosphate (7) in 62\% yield and with potassium iodide iodorhodocenium iodide (8b) in 84\% yield. Clearly many other synthetic applications of 5 may be envisaged, for example aza-coupling with arenes would give access to redox-responsive rhodocenium azo dyes.

An alternative and shorter route to iodo-substituted rhodocenium salts was developed by the reaction of starting material cyclopentadienylidiodorhodium dimer (1) with one equivalent of silver hexafluoridophosphate per rhodium atom and with diazocyclopentadiene or its more stable and easier to handle triphenylphosphine-adduct\textsuperscript{9} under insertion of the transient cyclopentadienylidene into the rhodium-iodine bond, in analogy to such reactions with ruthenium\textsuperscript{10} or manganese and rhodium\textsuperscript{11} halide complexes, affording thereby iodorhodocenium hexafluoridophosphate (8a) in 55\% isolated yield. Furthermore and conceptionally similar to the preparation of pyridiniorhodocenium 2, half-sandwich capping reaction of 1 with triphenylphosphonium cyclopentadienide\textsuperscript{8a,12} gave slightly air-sensitive, dicatonic phosphonio-substituted rhodocenium salt 9 in 90\% yield, potentially useful for Pd-catalyzed cross-coupling reactions.\textsuperscript{13} Finally, Staudinger reaction of azidorrhodocenium hexafluoridophosphate (7) with triphenylphosphine gave access to monocatonic iminophosphorane (10) in 91\% yield as an orange, air-stable complex. In contrast to the classical Staudinger reaction, no hydrolysis to aminorhodocenium (3) was observed, and, synthetically much more important, no aza-Wittig reaction\textsuperscript{14} with aldehydes proved possible. However, it is known that electron-poor azides, e.g. perfluoroalkyl azides, afford air and water-stable iminophosphoranes that have current applications in bioorthogonal derivatization.\textsuperscript{15} Obviously, the catonic rhodocenium moiety is similarly electron-withdrawing, as was also observed in the cobaltoceniumiminophosphorane congener.\textsuperscript{16}

Overall, these synthetic methods allowed access to new valuable functionalized rhodocenium salts with prospects for further development of this class of compounds. However, a general problem with this chemistry is the difficulty to obtain pure compounds: chromatography is usually not possible and separation of various rhodocenium species, e.g. undesired unsubstituted rhodocenium from target substituted rhodocenium salts, by selective precipitation and/or crystallization is not easy to achieve and results in loss of material. Therefore, carefully optimized reaction conditions are mandatory. Moreover, rhodium chemicals are quite expensive, therefore mostly small-scale reactions are feasible.

Physical, Spectroscopic, and Structural Properties

Mono or dicatonic rhodocenium derivatives 2–10 are highly polar, air-stable salts (except slightly air-sensitive 9) of yellow to orange color, soluble in polar solvents like acetonitrile, nitromethane, methanol, acetone, dichloromethane and water.

Positive mode electrospray high-resolution mass spectra gave signals of their most abundant monoisotopic peaks of the cationic moieties in excellent agreement with calculated values, except for azido derivative 7 where matrix-induced degradation via the nitrene intermediate to the amino derivative 3 was evident.

ATR-IR spectra are dominated by the very strong absorptions of the PF\textsubscript{6} anions (\(\approx 820\) and 550 cm\(^{-1}\)) and show diagnostic bands for dipolar functional groups (compare Experimental Section), e.g. the diazonio substituent of 5 gave a weak absorption at 2289 cm\(^{-1}\) and the azido substituent of 7 showed a medium strong band at 2025 cm\(^{-1}\).

\(^1\)H-NMR spectra of these rhodocenium compounds display the common principal pattern of monosubstituted metalloenes (one singlet, 5 H, unsubstituted Cp and two pseudo-triplets, 2×2H, substituted Cp) in the range of 5.4–7.4 ppm with additional \(\text{J}(^1\text{H},^{13}\text{Rh})\) couplings of approximately 1 Hz to the \(^{10}\)Rh nucleus (S\(=\frac{1}{2}\), 100\% natural abundance), thereby each unsubstituted Cp is observed as a doublet and each hydrogen signal of the substituted Cp is observed as a poorly resolved multiplet (see Supporting Information). As a representative example, Figure 1 shows the \(^1\)H and \(^{13}\)C spectra of amino-rhodocenium hexafluoridophosphate (3). In addition, hydrogen signals of the attached substituents are observed where applicable, e.g. amino resonances for 3 and pyridine or phenyl resonances for compounds 2, 9 and 10. As expected, these latter signals are shifted to lower field in comparison to those of “normal” (het)aryl substituents, due to the positive charge of these pyridinio or phosphonium substituents.

In the \(^{13}\)C-NMR spectra of 2–10, carbon resonances of the rhodocenium sandwich moieties are easily identified as doublets with \(\text{J}(^{13}\text{C},^{10}\text{Rh})\) coupling constants of 6–9 Hz; see for example Figure 1 (bottom) for 3. The chemical shift range of the rhodocenium signals is in the range from 70–100 ppm, a spectral area rarely covered by organic compounds. Furthermore, characteristic \(^{31}\)P-NMR signals with corresponding couplings to \(^{13}\)C (see Supporting Information) are observed for compounds 9 and 10.

\(^{103}\)Rh-NMR is a special case: On the one hand, \(^{103}\)Rh is a nucleus with “good” NMR properties (S\(=\frac{1}{2}\), 100\% natural abundance), but on the other hand its extremely low gyromagnetic ratio (\(\lambda=-0.8420\times10^7\) radT\(^{-1}\)s\(^{-1}\)), poor receptivity (0.2\% of that of \(^{13}\)C) and huge chemical shift range (\(\approx 13000 \) ppm) makes it in practice very difficult to obtain NMR data. Moreover, most standard multinuclear NMR spectrometers are not equipped for detecting \(^{103}\)Rh signals. However, in diamagnetic Rh(I) or Rh(III) complexes with resolved scalar couplings to other nuclei of spin \(\frac{1}{2}\), application of magnetization transfer pulse sequences allow considerable increase in sensitivity with concomitant shorter data acquisition periods.\textsuperscript{17} Therefore we set out to try this technique on our family of new
monosubstituted rhodocenium compounds having \( J(\text{H}^1,^{103}\text{Rh}) \) couplings of approximately 1 Hz, aiming at a possible correlation of their \( ^{103}\text{Rh} \) chemical shifts with inductive or resonance substituent parameters.\(^{18}\) In our first paper on rhodocenium chemistry,\(^{2}\) we experienced some difficulties obtaining reliable \( ^{103}\text{Rh}\)-NMR data, but here in a second attempt we succeeded using carefully optimized spectrometer settings on selected compounds (3, 4, 7, 8a, 9, see Experimental Section). For comparison, we include also data of some rhodocenium carboxylic acid derivatives obtained earlier\(^{2}\) (Table 1). Figure 2 shows the HMQC spectrum of 9 as a representative example with nicely resolved \( J(\text{H}^1,^{103}\text{Rh}) \) couplings to all hydrogens of both cyclopentadienyl ligands.

Overall, the \( ^{103}\text{Rh} \) chemical shifts of the parent unsubstituted rhodocenium and of these nine monosubstituted rhodocenium complexes span a range of approximately 300 ppm with aminorhodocenium hexafluoridophosphate (3) showing the most deshielded and unsubstituted rhodocenium hexafluoridophosphate the most shielded value, respectively. This is quite counterintuitive to simple donor/acceptor considerations, indicating that 3 is an unusual amine, as we will also see in its solid-state structure discussed below. Neglecting the \( ^{103}\text{Rh}\)-NMR value of 3 and attempting to correlate the other data with Hammett substituent parameters\(^{18}\) gave no linear regression at all, reflecting that no simple resonance and inductive phenomena are operating in this family of compounds.

Single crystal X-ray structure analyses are available for complexes 2, 3 (chloride instead of hexafluoridophosphate as counterion), 7, 8b, 9 and 10. Molecular structures of the cations are depicted in Figure 3 and details of the refinement and other structural data are available in the Supporting Information. Overall, these simple rhodocenium salts show the expected regular structures with normal, undistorted sandwich moieties containing parallel cyclopentadienyl ligands. Rhodium-carbon bond lengths are in the usual range of 2.15-2.19 Å\(^{18}\) and the appended functional groups display common structural metrics. There are only two details that need further comment: First, the carbon-nitrogen bond length in aminorhodocenium 3' is shortened [C10-N1 = 1.34 Å] in comparison to a standard C–N single bond, indicative of partial iminium/fulvene bonding, a structural motif generally observed in aminometalocenium salts.\(^{14}\) This is in line with the unusual \( ^{103}\text{Rh}\)-NMR shift of 3 (vide supra) and with a reduced Brønsted-basicity of the amino group of 3. Second, whereas the cationic part of iodorhodocenium iodide 8b is quite normal, the counterion iodide shows an interesting halogen-bonding\(^{19}\) to a diiodine molecule with 50% occupancy in the unit cell (see Supporting Information).

All in all, these structures give further definite proof of the identity of these new rhodocenium compounds.

### Electrochemistry

Due to the limited amount of material available for some compounds, only selected materials were studied by cyclic voltammetry. All rhodocenium salts investigated (2a, 3, 4, 9, 10) show irreversible rhodocenium centered redox-processes \( E_{\text{Rc}} \) between –1.83 and –2.20 V vs. the ferrocene/ferrocenium couple (Figure 4, Table 2). Compared to our previous reported rhodocenium complexes,\(^{2}\) these potentials are slightly anodically shifted to lower potentials, which can be easily explained by the more electron donating character of the cyclopentadienyl substitution patterns. In addition to the expected rhodocene/rhodocenium redox event, the complexes 2a, 4, 9 and 10 show also further redox processes depending on their substitution.

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**Table 1.** \(^{103}\text{Rh}\)-NMR data of monosubstituted rhodocenium salts.

| Compound | Formula | \( [\text{ppm}] ^{103}\text{Rh})^{c} \) |
|----------|---------|---------------------------------|
| \([\text{Rh} = \text{H}]\text{PF}_6\) | –10127 |
| \([\text{Rh} = \text{CH}_3]\text{PF}_6\) | –10059 |
| \([\text{Rh} = \text{CO}_2\text{CH}_3]\text{PF}_6\) | –9997 |
| \([\text{Rh} = \text{CO}_2\text{H}]\text{PF}_6\) | –9991 |
| \([\text{Rh} = \text{CO}_2\text{NH}]\text{PF}_6\) | –10000 |
| \([\text{Rh} = \text{NH}]\text{PF}_6\) | –9820 |
| \([\text{Rh} = \text{N} = \text{N} = \text{Rh}]\text{PF}_6\) | –9921 |
| \([\text{Rh} = \text{N}]\text{PF}_6\) | –9949 |
| \([\text{Rh} = \text{I}]\text{PF}_6\) | –9976 |
| \([\text{Rh} = \text{P(C}_2\text{H}_5)]\text{PF}_6\) | –9986 |

[a] Unnumbered rhodocenium carboxylic acid derivatives were synthesized according to reference 2. [b] \( \text{Rc} = \text{rhodoceniumy}, [\text{C}_9\text{H}_7\text{Rh} (\text{C}_9\text{H}_6)]^{+} \). [c] Chemical shifts (ppm) are referenced against external Rh(acac).
pattern: Complex 2a shows a reduction at −1.03 V (E1) which is coupled to two redox events E2 and E3 and is most likely centered on the pyridinium substituent. Complex 4 shows a broad redox feature at −1.04 V and −0.72 V due to the azo bridge between the two rhodocenium centers, less resolved in comparison to its cobaltocenium congener. Complex 9 shows a second reduction at −2.89 V which is most likely a P(V)/P(IV) redox couple. The irreversible redox processes E1 and E2 in the CV of 9 may be explained similarly, reducing the P(V) center to P(IV).

Conclusion

The synthesis of a range of new functionalized rhodocenium salts was achieved by capping reactions of half-sandwich CpRhX₂ starting materials with zwitterionic pyridinium/phosphonium-Cp synthons followed by an unusual application of the Zincke reaction, diazoniation, Sandmeyer reactions, and Staudinger reaction. The new mono-functionalized rhodocenium complexes are air-stable materials that were fully characterized by high-resolution mass spectrometry, IR spectroscopy, ¹H, ¹³C and most notably also by ¹⁰³Rh NMR spectroscopy. Single crystal structure analyses of all target compounds show the expected regular sandwich structure of simple monofunctionalized metallocenes. Cyclic voltammetry of selected representatives show irreversible rhodocenium/rhodocene reduc-

Table 2. Overview of redox potential of selected rhodocenium salts.[a]

| Complex | E₁   | E₂   | E₃   | E₄   |
|---------|------|------|------|------|
| 2a      | −1.83 V | −1.03 V | −0.19 V | 0.34 V |
| 3       | −1.96 V | −     | −     | −    |
| 4       | −2.04 V | −1.04 V | −0.72 V | −    |
| 9       | −2.01 V | −2.89 V | −     | −    |
| 10      | −2.20 V | −1.54 V | −2.72 V | −0.54 V |

[a] Potentials are referenced versus an external ferrocene/ferrocenium couple. E₁ indicates rhodocenium centered redox processes, E₂ to E₄ indicate follow up redox events.
tions and additional redox events dependent on the substitution pattern. Overall, with this research we give access to chemically useful new rhodocenium salts, thereby fostering the future development of this so far rarely studied class of compounds.

**Experimental Section**

**General procedures:** Synthetic operations were performed by standard organometallic methods in Schlenk glassware under an inert atmosphere of Ar. Solvents were purified and dried in a solvent purification system with molecular sieves as water scavengers under an atmosphere of Ar. Chemicals were obtained commercially and used as received. Starting material cyclopentadienylidiodorhodium dimer (1) was synthesized according to a published procedure.[3] 

\[ \text{3} \]

\[ \text{H,}^{13}\text{C,}^{31}\text{P NMR spectra were recorded on a Bruker DPX 300 NMR spectrometer and signals were referenced internally against} \ 1\text{H/}^{13}\text{C residual solvents peaks or externally (}^{31}\text{P)} \text{ against 85% aqueous phosphoric acid.} \]

\[ \text{E}_{\text{Rh}} \text{ indicates rhodocenium centered redox processes,} \ E_1 \text{ to} \ E_3 \text{ indicate follow up redox events.} \]

![Figure 3](image-url) Molecular structures of rhodocenium salts with thermal ellipsoids at a 50% probability, counterions PF$_6$/Cl/I and solvent molecules omitted for clarity. Selected bond lengths [Å]: 2a: C10-N1 = 1.437(8); 3': C10-N1 = 1.331(13); 7: C10-N1 = 1.398(3), N1-N2 = 1.253(3), N2-N3 = 1.124(3); 8b: C10-I1 = 2.095(7); 9: C10-P1 = 1.799(5); 10: C10-N1 = 1.351(3), N1-P1 = 1.5983(18).

![Figure 4](image-url) Cyclic voltammograms of complexes 2a, 3, 4, 9 and 10 in acetonitrile. Measurements conditions: 0.2 M NBu$_4$PF$_6$ electrolyte and 0.001 M analyte; referenced vs. ferrocene/ferrocenium couple as an external standard; scan rate 100 mV s$^{-1}$. $E_{\text{m}}$ indicates rhodocenium centered redox processes, $E_1$ to $E_3$ indicate follow up redox events.
comparison for $^{103}$Rh-NMR data, rhodocenium hexafluoridophosphate ([Rh(CN)$_3$]$_3$), methylrhodocenium hexafluoridophosphate ([Rh(C$_3$H$_7$)F$_6$]$_3$), rhodoceniumcarboxylic acid hexafluoridophosphate ([RCO$_2$H]PF$_6$), methoxyacetylenoborylidenium hexafluoridophosphate ([RCO$_2$CH$_2$CF$_3$]PF$_6$) and rhodoceniumcarboxylic acid amide hexafluoridophosphate ([RCO$_2$NH]PF$_6$) were obtained as recently published.$^{29}$ Mass spectrometric data were measured on a Thermo Finnigan Q Exactive Orbitrap spectrometer, IR spectra were performed on a Bruker ALPHA IR spectrometer, single crystal X-ray diffraction data were collected on a Bruker D8 Quest diffractometer with graphite-monochromated Mo–K$_\alpha$ radiation ($\lambda$=0.71073 Å) and structures were solved by direct methods. Cyclovoltammetry measurements were performed inside an Ar-filled glove-box using a BioLogic SP 150 potentiostat with a three-electrode setup: glassy carbon (working electrode), platinum (counter electrode), silver (pseudo reference).

Pyridiniorhodocenium bis(hexafluoridophosphate) (2a): A Schlenk tube was charged under an atmosphere of Ar with 234.8 mg (0.5566 mmol, 1 equiv) [CpRhI]$_3$ (1), 281.4 mg (1.113 mmol, 2 equiv) AgPF$_6$, and 25 mL of dry acetone. Under exclusion of light the mixture was stirred for half an hour, resulting in an orange solution and a slightly yellow precipitate of AgI. To this mixture, 79.7 mg (0.5566 mmol, 1 equiv) pyridiniumcyclopentadienide$^{40}$ was added under protection from air and stirring was continued at room temperature overnight. The resulting purple solution was then evaporated on a rotary evaporator, diethyl ether was added to precipitate the crude product, the mixture was filtered and the residue was dissolved in acetone, filtered again, the solution was evaporated on a rotary evaporator, and the oily residue was dried in vacuo, resulting in 2.57 mg (0.4586 mmol, 82.4% yield) of yellow air-stable 2a.

Diazoniorhodocenium bis(hexafluoridophosphate) (5): In a round bottom flask, 112.9 mg (0.2872 mmol, 1 equiv) aminorhodocenium hexafluoridophosphate (3), 235.0 mg (1.085 mmol, 2.5 equiv) HgO, 275.4 mg (1.085 mmol, 2.5 equiv) I$_2$, and 50 mL of reagent grade dichloromethane. After stirring at ambient conditions for 24 hours, 60 mL of diethyl ether was added and the resulting dispersion was filtered through a paper filter. The solid material in the paper filter was eluted with acetonitrile, then the solution was evaporated on a rotary evaporator, and the resulting orange product is dried in vacuo, affording 150.5 mg (0.1925 mmol, 88.7% yield) of 5. HRMS (ESI pos): C$_{33}$H$_{33}$NRh, calcd. most abundant isotope peak 247.9941 (M - PF$_6$)$_2$ +, found m/z=247.9936. Single crystal structure XRD analysis: Single crystals of 3’ (chloride and the hexafluoridophosphate) were obtained from an aqueous solution in diluted hydrochloric acid at 4°C (Figure 3, Supporting Information).

Diazeniorhodocenium bis(hexafluoridophosphate) (6): A round bottom flask was charged with 170.6 mg (0.4340 mmol, 1 equiv) aminorhodocenium hexafluoridophosphate (3), 235.0 mg (1.085 mmol, 2.5 equiv) HgO, and 25 mL of dry acetone. Under exclusion of light the mixture was stirred for half an hour, resulting in an orange solution and a slightly yellow precipitate of AgI. To this mixture, 79.7 mg (0.4340 mmol, 1 equiv) AgPF$_6$, and 50 mL of reagent grade dichloromethane. After stirring at ambient conditions for 24 hours, 60 mL of diethyl ether was added and the resulting dispersion was filtered through a paper filter. The solid material in the paper filter was eluted with acetonitrile, then the solution was evaporated on a rotary evaporator, and the resulting orange product is dried in vacuo, affording 150.5 mg (0.1925 mmol, 88.7% yield) of 6. HRMS (ESI pos): C$_{33}$H$_{33}$NRh, calcd. most abundant isotope peak 247.9941 (M - PF$_6$)$_2$ +, found m/z=247.9936. Single crystal structure XRD analysis: Single crystals of 3’ (chloride and the hexafluoridophosphate) were obtained from an aqueous solution in diluted hydrochloric acid at 4°C (Figure 3, Supporting Information).

Dirhodoceniumylidiane (‘Azorhodocenium’) bis(hexafluoridophosphate) (4): A round bottom flask was charged with 170.6 mg (0.4340 mmol, 1 equiv) aminorhodocenium hexafluoridophosphate (3), 235.0 mg (1.085 mmol, 2.5 equiv) HgO, and 25 mL of dry acetone. Under exclusion of light the mixture was stirred for half an hour, resulting in an orange solution and a slightly yellow precipitate of AgI. To this mixture, 79.7 mg (0.4340 mmol, 1 equiv) AgPF$_6$, and 50 mL of reagent grade dichloromethane. After stirring at ambient conditions for 24 hours, 60 mL of diethyl ether was added and the resulting dispersion was filtered through a paper filter. The solid material in the paper filter was eluted with acetonitrile, then the solution was evaporated on a rotary evaporator, and the resulting orange product is dried in vacuo, affording 150.5 mg (0.1925 mmol, 88.7% yield) of 4. HRMS (ESI pos): C$_{33}$H$_{33}$NRh, calcd. most abundant isotope peak 247.9941 (M - PF$_6$)$_2$ +, found m/z=247.9936. Single crystal structure XRD analysis: Single crystals of 3’ (chloride and the hexafluoridophosphate) were obtained from an aqueous solution in diluted hydrochloric acid at 4°C (Figure 3, Supporting Information).
product contains approx. 14% unsubstituted rhodocenium salts (hexafluoridophosphate and/or bromide), a common impurity in ball mill reactions of metalocenium diazonium salts.\(^{16}\) \(\text{H-NMR (300 MHz, CD}_2\text{CN, 25}^\circ\text{C): }\delta = 5.82\) (unresolved m, 2 H, \(\text{CP}_{\text{subt}}\)), 5.91 (d, \(\text{J}^{(131}\text{C}-\text{P}) = 1.0\) Hz, Cp), 6.22 (unresolved m, 2 H, \(\text{CP}_{\text{subt}}\)) ppm. Note: A \(\text{13C-NMR spectrum could not be obtained due to the very limited amount of material available. HRMS (ESI pos): } \text{C}_3\text{H}_8\text{BrRh, calcld. most abundant isotope peak 310.8937 (M+P}^{+}\text{Rh)}^+\))

Azidorhodocenium hexafluoridophosphate (7): A small Schlenk tube charged with 17.3 (0.0315 mmol, 1 equiv) diazonio-rhodocenium hexafluoridophosphate (5) and 3 mL of dry nitromethane. The resulting solution was cooled to \(4^\circ\text{C and 5.7 mg (0.0944 mmol, 3 equiv) NaN}_3\) was added. After stirring at \(4^\circ\text{C for 20 minutes, the mixture was allowed to warm to room temperature. Workup under ambient conditions: Solvent was removed on a rotary evaporator, the residue was dissolved in dichloromethane, and the solution was filtered through a paper filter, solvent was removed on a rotary evaporator and the resulting product was dried in vacuo, affording 8 mg (0.0194 mmol, 61.6% yield of dark yellow 7. H-NMR (300 MHz, acetone-\(d_6\), 25°C): \(\delta = 5.98\) (unresolved m, 2 H, \(\text{CP}_{\text{subt}}\)), 6.17 (d, \(\text{J}^{(131}\text{C}-\text{P}) = 1.0\) Hz, Cp), 6.28 (unresolved m, 2 H, \(\text{CP}_{\text{subt}}\)) ppm. Note: A \(\text{13C-NMR spectrum could not be obtained due to the very limited amount of material available.}

Triphenylphosphorohodocenium hexafluoridophosphate (8 a): A Schlenk vessel was charged under an atmosphere of Ar with 51 mg (0.1209 mmol, 1 equiv) \(\text{CPBr}_3\) (1, 6.11 mg (0.0240 mmol, 2 equiv) AgPF\(_6\)) and 20 mL of dry acetonitrile. After stirring at room temperature for 30 minutes, 39.5 mg (0.1209 mmol, 1 equiv) of triphenylphosphinocyclopentadienide\(^{12}\) was added and stirring was continued overnight. Workup under ambient conditions: The mixture was filtered, the solution was evaporated to dryness and the beige crystalline solid was dried in vacuo, affording 85.6 mg (0.291 mmol, 93% yield) \(8\text{ a}\). The product was eluted from the filter with dichloromethane, the residue was stirred further overnight. Workup under ambient conditions: Solvent was evaporated nearly to dryness, then the same volume of diethyl ether was added and the precipitate was filtered off. The product was eluted from the filter with dichloromethane, the solvent was evaporated on a rotary evaporator and the residue was dried in vacuo. Yield: 5 mg (9.9 mmol, 84%) \(8\text{b}\). H-NMR (300 MHz, acetone-\(d_6\), 25°C): \(\delta = 6.07\) (unresolved m, 2 H, \(\text{CP}_{\text{subt}}\)), 6.13 (d, \(\text{J}^{(131}\text{C}-\text{P}) = 1.0\) Hz, Cp), 6.42 (unresolved m, 2 H, \(\text{CP}_{\text{subt}}\)) ppm. IR (ATR): \(\nu\) = 3084, 2957, 2926, 2857, 1724, 1601, 1460, 1405, 1377, 1268, 1120, 1072, 1040, 1014, 998, 839, 741, 704, 570, 412 cm\(^{-1}\). Single crystal structure XRD analysis: Suitable crystals were obtained by slow evaporation of a dichloromethane solution of \(8\text{ b}\) at room temperature. (Figure 3, Supporting Information).
The authors declare no conflict of interest.

Conflict of Interest

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Keywords: Metallocenes · Molecular electrochemistry · Rhodium · Sandwich complexes · Structure elucidation

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