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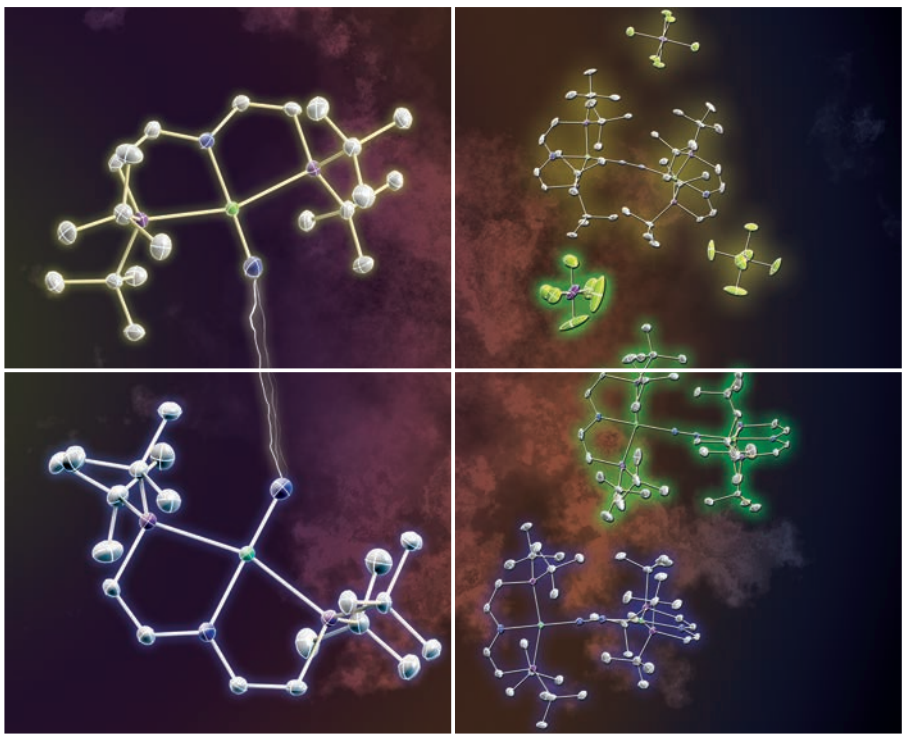

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Coupling of terminal iridium nitrido complexes†

Josh Abbenseth,[a] Markus Finger,[a] Christian Würtele,[a] Müge Kasanmascheff[b] and Sven Schneider*[a]

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The oxidative coupling of nitride ligands (N^{3-}) to dinitrogen and its microscopic reverse, N_2-splitting to nitrides, are important elementary steps in chemical transformations, such as selective ammonia oxidation or nitrogen fixation. Here an experimental and computational evaluation is provided for the homo- and heterocoupling of our previously reported iridium(IV) and iridium(V) nitrides $[IrN(PNP)]^n$ (n = 0, +1; PNP = $N(CHCHPtBu_2)_2$). All three formal coupling products $[(PNP)IrN_2Ir(PNP)]^n$ (n = 0–+2) were structurally characterized. While the three coupling reactions are all thermodynamically feasible, homocoupling of $[IrN(PNP)]^+$ is kinetically hindered. The contributing parameters to relative coupling rates are discussed providing qualitative guidelines for the stability of electron rich transition metal nitrides.

Introduction

The splitting of dinitrogen at ambient temperatures is a highly attractive reaction in the context of nitrogen fixation, yet still represents a challenging goal.[1] Ever since the seminal work of Cummins and co-workers about 20 years ago,[2] several other systems were reported to undergo the splitting of dinitrogen into well-defined nitrido complexes.[3] The first reported and intensively examined example proceeds through diazenido dimer $[(ArtBuN)_3Mo(NN)Mo(NtBuAr)_3]$. Importantly, this linear μ-N_2 complex holds 10 π-electrons in the {MNNM}-core (Fig. 1a), finally resulting in the closed-shell nitride $[NMo(NtBuAr)_3]$.[4,5] In comparison, the dimers $[(ArtBuN)_3Mo(NN)Mo(NtBuAr)_3]^{n+}$ (n = 1–2) exhibit stronger degrees of N_2 activation due to depletion of a molecular orbital that is N–N antibonding in character. However, N_2-splitting is not observed due to destabilization of the nitrides upon oxidation. Similarly,

[a]*Institut für Anorganische Chemie, Georg-August-Universität, Tammannstraße 4, 37077 Göttingen, Germany. E-mail: sven.schneider@chemie.uni-goettingen.de*
[b]*Max Planck Institute for Biophysical Chemistry, 37077 Göttingen, Germany*

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Sven Schneider

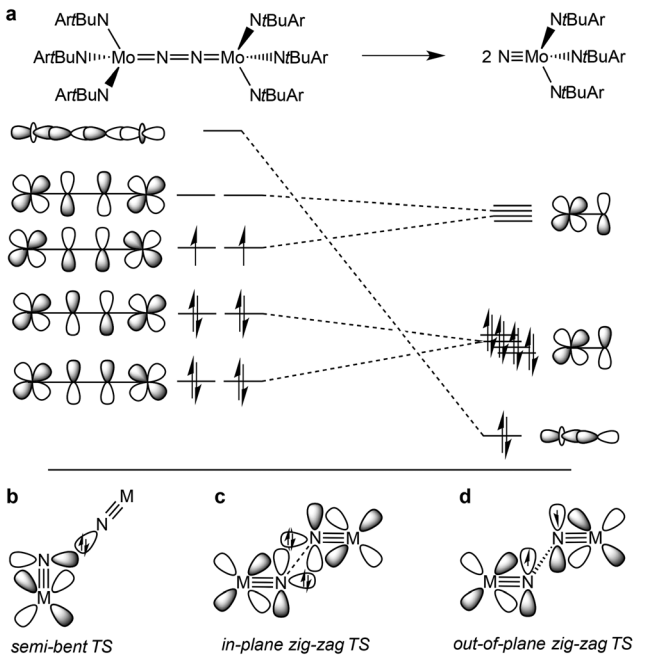

Fig. 1 Qualitative molecular orbitals relevant for N_2 splitting with Cummins' complex (a) and schematic transition states that were proposed for N_2-splitting/coupling (b–d).

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Nishibayashi and co-workers examined the redox series $[(depf)_2Cp^*Mo(NN)MoCp^*(depf)_2]^{n+}$ (n = 0–2; depf = 1,1′-bis(diethylphosphino)ferrocene), observing N_2 splitting for the 10 π-electron system (n = 0) and the reverse, nitride coupling, after oxidation of the resulting nitride.[3d]

A similar picture arises for the reverse reaction, *i.e.* the coupling of terminal nitrides. Several authors reported the formation of dinitrogen bridged complexes upon reductive coupling of octahedral M^V nitrides (M = Fe, Ru, Os).[6] These transient nitrides are typically formed *in situ*, *e.g.* by photolysis of M^{III}-azides, reduction of stable M^{VI} nitrides or oxidation of low-valent ammine complexes to give divalent ($\{M^{II}N_2\}$ or $\{M^{II}N_2M^{II}\}$) or mixed valent $\{M^{II}N_2M^{III}\}$ dinitrogen complexes. Except for a few cases, it is not fully clear whether the reaction proceeds *via* $M^V{\equiv}N$ homocoupling or coupling of $M^V{\equiv}N$ with parent $M^{VI}{\equiv}N$. Importantly, within the simplified MO-picture (Fig. 1a) the $M^{II/II}$ and $M^{II/III}$ dinitrogen bridged complexes represent 12 and 11 π-electron {MNNM}-cores, respectively. However, some closed-shell nitrides also decay by nitride coupling underlining that the electronic configuration is not the only predictor for the thermochemistry of N_2-splitting/coupling.[7–9]

In comparison, the relationship of electronic structure and kinetics received less attention. Ware and Taube examined the thermal decomposition of $[Os^{VI}NCl_3(py)_2]$ by nitride coupling to $[Os^{III}Cl_3(py)_3]$.[7] They pointed out that donation from a nitride lone-pair into an empty $M{\equiv}N$ π*-orbital should be favored over a collinear approach of the two monomers and therefore proposed a semi-bent transition state (Fig. 1b). In line with such a polar transition state (TS), Seymore and Brown showed that heterocoupling of electrophilic Os^{VI}- and nucleophilic Mo^{VI}-nitrides is considerably faster than the respective homocoupling reactions.[10] However, Burger and coworkers emphasized that nitride homocoupling should favor a symmetrical "non least-motion pathway" or in-plane zig-zag TS, which features mutual σ/π donor–acceptor interactions of the two monomer fragments (Fig. 1c).[11]

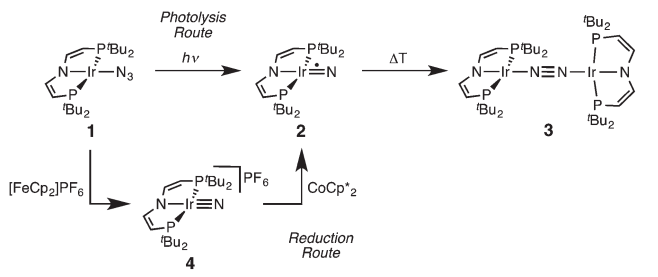
Recently, we reported that photolysis of the iridium(II) azide $[Ir(N_3)(PNP)]$ (**1**, PNP = $N(CHCHP^tBu_2)_2$) results in the formation of iridium(IV) nitride [IrN(PNP)] (**2**, Scheme 1).[12] Spectroscopic and computational analysis suggested that **2** and the analogous rhodium compound are π-radicals with strong delocalization of the unpaired electron over the $M{\equiv}N$ core, *i.e.* considerable "nitridyl" character.[12,13] In solution, these transient nitrides undergo clean coupling, *e.g.* to 12-π-electron N_2-complex $[N_2\{Ir(PNP)\}_2]$ (**3**) in case of **2** (Scheme 1). The computed TS for a PMe_2-truncated model featured a mutual π/π MO-interaction as the most favorable pathway with an out of plane zig-zag arrangement of the two monomer radicals (Fig. 1d).[12] Alternatively, oxidation of **1** results in formation of isolable iridium(V) nitride $[IrN(PNP)]^+$ (**4**), which in turn also gives **2** upon reduction (Scheme 1). In this context, we were interested whether the enhanced thermal stability of **4** can be attributed to thermodynamic or kinetic arguments. Here, we present a systematic experimental and computational study that addresses the stability of the iridium nitrides with respect to oxidative coupling.

Results and discussion

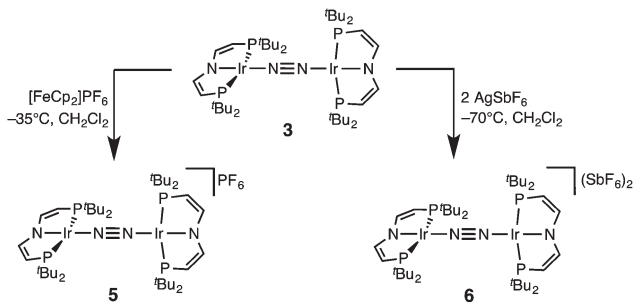
Synthesis of the $[N_2\{Ir(PNP)\}_2]^{n+}$ (n = 0–2) redox series

In a slightly improved synthetic protocol, the neutral dimer **3** can be obtained analytically pure in 66% yield on the iridium(V) nitride reduction route using Na/Hg with strict exclusion of light. Complex **3** was characterized by cyclic voltammetry (CV, see ESI†). The CV reveals three oxidative redox processes at $E_{1/2}$ = −0.27, 0.18, 0.35 V (*vs.* $FeCp_2/FeCp_2^+$), respectively. While the $Ir^{I/I}/Ir^{I/II}$ couple is fully reversible at r.t., the higher oxidations seem pseudo-reversible, as evidenced by the observation of small additional peaks upon cyclic back to low potentials. Hence, the electrochemical results suggest, that one- and two-electron oxidation of **3** could give isolable products.

Chemical oxidation of **3** with one equivalent of $[FeCp_2]PF_6$ in dichloromethane at −35 °C affords the isolation of $Ir^{I/II}$ N_2-complex $[N_2\{Ir(PNP)\}_2]PF_6$ (**5**) as a temperature, light, and air sensitive dark green solid (Scheme 2). The X-band EPR spectrum of **5** in frozen solution (4 K) reveals a rhombic signal corresponding to an S = 1/2 system (ESI†). The large anisotropy of the g-tensor (g_x = 2.66, g_y = 2.12, g_z = 1.62) without resolved hyperfine coupling was similarly observed for several Ir^{II}(PNP) complexes with square-planar geometry.[5,14] From the CV data a comproportionation constant K_C is estimated to be around 10^7. A large K_C like this is frequently stressed as a simple predictor for electronic delocalization in the mixed valent complex indicating stabilization due to resonance exchange.[15] Note that K_C data should be treated with caution and is easily



Scheme 1 Syntheses of iridium(IV) and iridium(V) PNP nitrides **2** and **4** and coupling to iridium(I)/iridium(I) N_2-complex **4**.[12]



Scheme 2 Chemical oxidation of iridium(I)/iridium(I) N_2-complex **3**.

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over-interpreted due to the many other parameters possibly contributing to electrochemical potentials.[16] In fact, the bond parameters in the solid state indicate valence localization in the crystal (see below) and the DFT model of **5** also exhibits localization of the spin density on one of the two Ir(PNP) moieties (ESI†). In contrast, the ^{1}H NMR spectrum of **5** exclusively shows one set of paramagnetically shifted and broadened pincer ligand peaks (ESI†), suggesting rapid electron transfer between the N_2-bridged moieties on the NMR timescale (1–10^{-4} s). Unfortunately, detailed analysis of the NIR spectrum was hampered by the thermal instability of **5** and dication **6** (see below) and did not allow for a reliable assignment of intervalence charge transfer *vs.* interconfigurational electronic transitions. However, IR spectroscopy reveals the presence of a relatively intense band assignable to the N_2 stretching vibration both in solution (CH_2Cl_2: 1959 cm^{-1}) and in the solid state (nujol: 1960 cm^{-1}). The assignment was confirmed by DFT computations (ESI†) scaled to monomeric complex [Ir(N_2)(PNP)] (2077 cm^{-1}):[12] The experimental shift $\Delta\nu_{N_2}$ = 117 cm^{-1} is well reproduced by DFT ($\Delta\nu_{N_2}$ = 115 cm^{-1}). This comparison also excludes the formation of monomeric [Ir(N_2)-(PNP)]$^+$, which should be blue shifted with respect to [Ir(N_2)-(PNP)]. Importantly, the observation of an N_2-stretch confirms the presence of a permanent dipole, hence valence localization on the IR-timescale (~10^{-14} s) also in solution providing an upper limit for the charge transfer rate. Similar properties between localized and delocalized mixed-valence (Class II–III) were found for N_2-bridged $Os^{II/III}$ complexes.[17] Unequivocal assignment of complex **5** requires further investigations.

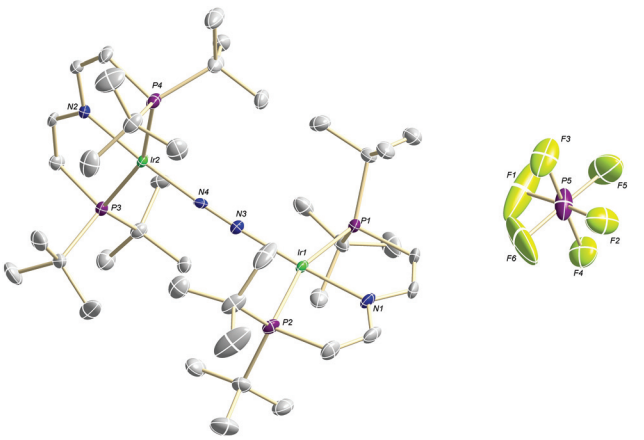

Fig. 2 Molecular structure of **5** in the solid state (ellipsoids at 50% probability level, hydrogen atoms and solvent molecules omitted for clarity).

Confirmation of the molecular structure of **5** is provided by single-crystal X-ray diffraction (Fig. 2 and Table 1). Both Ir ions are in distorted square-planar coordination geometries and bridged by a linear, end-on bound N_2, as in parent **3**. The structure features two crystallographically distinctly different Ir(PNP) fragments. For example, the Ir–N_{PNP} and Ir–N_{N_2} distances differ by 0.06 Å and 0.11 Å, respectively. Ir2 exhibits a longer bond to the PNP π-donor ligand and a shorter bond to the π-acceptor N_2 than Ir1. In that respect, Ir2 behaves similar to Ir in **3**. Hence, the structural parameters suggest trapped oxidation states in the solid state with Ir1 and Ir2 being associated with the +2 and +1 valence states, respectively. Importantly, the N–N distance (1.136(6) Å) is identical with that in **3** (1.135(4) Å) and in [N_2{Ir(PCP)}$_2$] (1.134(2) Å; PCP = C_6H_3-2,6-(CH_2PtBu_2)$_2$)[18] and close to free N_2 (1.097 Å) revealing a weak degree of N_2-activation upon oxidation.

The reaction of **3** with two equivalents of $AgSbF_6$ at −70 °C in dichloromethane under the exclusion of light results in the immediate precipitation of metallic silver and formation of a deep red solution (Scheme 2). Monitoring by ^{1}H NMR spectroscopy reveals the formation of a new paramagnetic compound with one set of signals assignable to a PNP pincer ligand, yet not as clean as in case of **5** (ESI†). No signals were found by X-band EPR spectroscopy as expected for a non-Kramers system. Decomposition in solution above −50 °C into several unidentified paramagnetic compounds is indicated by new ^{1}H NMR signals upon warming. Importantly, the nitride **4** was not found as decomposition product.

The high lability of the product from double oxidation with respect to ambient temperatures and light hampered a more detailed spectroscopic characterization, so far. However, crystallization at low temperatures afforded red crystals of the dinitrogen bridged $Ir^{II/II}$ complex [(N_2){Ir(PNP)}$_2$][SbF_6)$_2$ (**6**) that were suitable for X-ray diffraction. In contrast to mixed-valent complex **5**, the molecular structure of **6** (Fig. 3 and Table 1)

Table 1 Comparison of selected experimental and computed bond lengths and angles of **3**, **5**, and **6**

	3[a]		**5**[a]		**6**[a]	
	Exp	DFT	Exp	DFT	Exp	DFT
Bond lengths (Å)						
Ir–N_2	1.937(3)/1.933(3)	1.922	1.984(5)/1.878(5)	1.988/1.855	1.954(3)	1.928
Ir–N_{PNP}	2.041(3)/2.035(3)	2.055	1.973(4)/2.031(4)	1.969/2.059	1.983(3)	1.995
N–N	1.135(4)	1.136	1.136(6)	1.141	1.138(6)	1.140
Bond angles (°)						
Ir–N–N	174.6(2)/172.3(3)	172.8	175.3(4)/174.6(4)	172.0/173.2	172.2(3)	173.2
N–Ir–N	173.6(2)/171.0(1)	172.0	172.8(2)/173.3(2)	169.9/171.6	168.0(1)	169.0

[a] Computed spin ground–states: ^{1}**3**, ^{2}**5**, ^{3}**6**.

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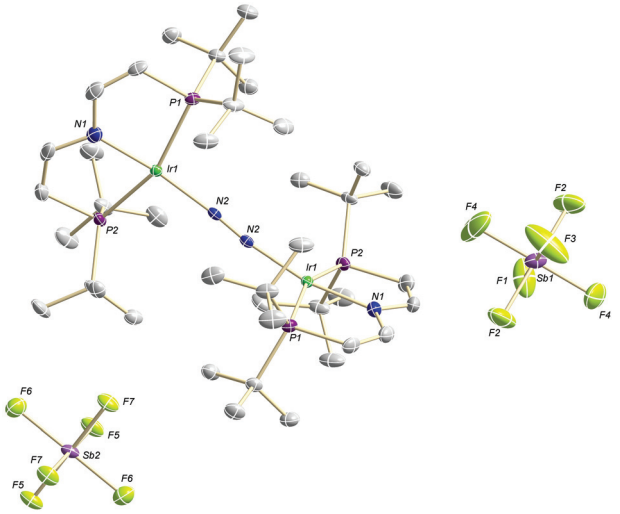
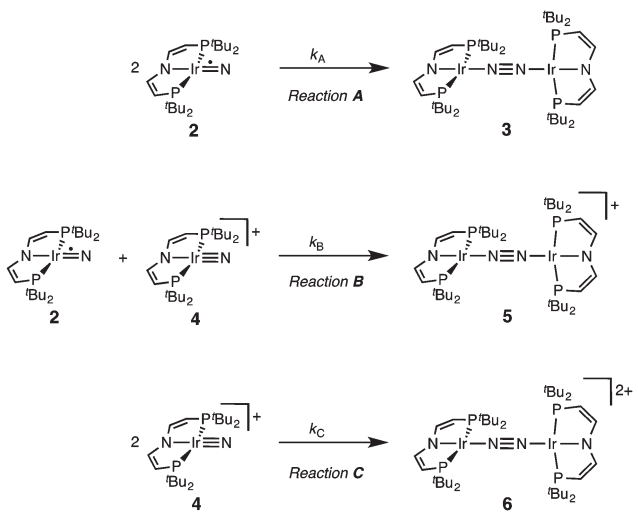


Fig. 3 Molecular structure of **6** in the solid state (ellipsoids at 50% probability level, hydrogen atoms and solvent molecules omitted for clarity).

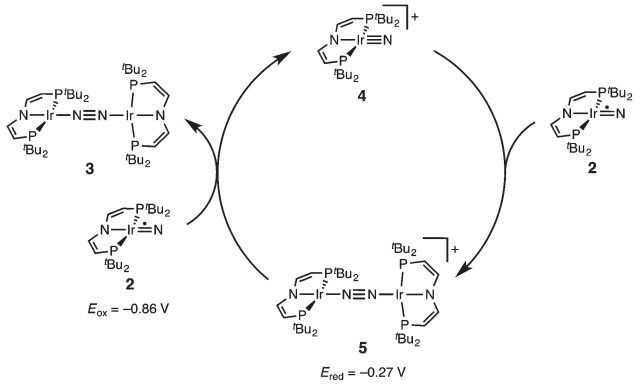
features two crystallographically equivalent Ir(PNP) fragments. The Ir–N_{PNP} (1.983(3) Å) and Ir–N_{N_2} (1.954(3) Å) bond lengths are very close to those of Ir1 in **5**, supporting the valence state assignments for the structure of **5** and full oxidation of **6** to $Ir^{II/II}$. The N_2 bond length (1.138(6) Å) is identical within error compared to the parent $Ir^{I/I}$ (1.135(4) Å) and $Ir^{I/II}$ (1.136(6) Å) complexes, indicating that N_2 activation is negligible within the $[N_2\{Ir(PNP)\}_2]^{n+}$ (n = 0–2) redox series.

Nitride coupling to $[N_2\{Ir(PNP)\}_2]^{n+}$ (n = 0–2)

The successful preparation of **5** and **6** and the absence of N_2 activation upon oxidation of **3** suggest that the iridium(v) nitride **4** is only kinetically stabilized with respect to coupling. We previously reported the kinetics of iridium(iv) nitride coupling (Scheme 3, **A**): A second order rate-law in **2** was found with a rate constant at room temperature k = 1.1(2) M^{-1} min^{-1}.



Scheme 3 Nitride coupling reactions examined.



Scheme 4 Proposed mechanism of redox catalysis for the coupling of nitride **2**.

Hence, this reaction is accompanied with a moderate kinetic barrier around $\Delta G^{\ddagger}_{298\ K}$ = 22 kcal mol^{-1}. In contrast, very slow decay is observed for iridium(v) nitride **4** over several days at elevated temperatures in dichloromethane, into several unidentified products. Furthermore, in contrast to **2**, decomposition of **4** follows a rate law that is first-order in **4** (k = 0.474(9) d^{-1}; ESI†). Hence, the nitride coupling rate (Scheme 3, **C**) could not be directly derived, but the decomposition rate allows for an estimate of a lower limit for the kinetic barrier of nitride coupling ($\Delta G^{\ddagger}_{313\ K}$ > 25 kcal mol^{-1}).

Coupling of the iridium(v) and iridium(iv) nitrides to mixed-valent **5** was also examined (Scheme 3, **B**). Spectro-electrochemical reduction of **4** (E_{red} = −0.86 V) at room temperature results in clean conversion to $Ir^{I/I}$ complex **3** (E_{ox} = −0.27 V). The observation of an isosbestic point at 560 nm indicates the absence of long-lived intermediates on the experimental time-scale (ESI†). Moreover, the equimolar addition of **2** to **4** at −70 °C, where both pure nitrides are kinetically stable, results in immediate coupling to **5** (ESI†) confirming much more rapid coupling to **5** than to **3** or **6**. Even the addition of a substoichiometric amount of **4** (10 mol%) to **2** at −70 °C results in the immediate formation with dimer **3** as the main product (ESI†). This observation can be rationalized with a redox-catalytic cycle (Scheme 4): Nitrides **2** and **4** undergo rapid coupling. The resulting dimer **5** is then reduced by starting material **2** to restore catalyst **4**, which is in line with the potentials of redox couples **2**/**4** and **3**/**5**, respectively. This mechanism also explains a previously reported observation for the two routes to iridium(iv) nitride **2** (Scheme 1): Unlike the photolysis route, the reduction route also yields considerable amounts of coupling product **3**, even at low temperatures where coupling of pure **2** is not observed.

Computational results

Our experimental results demonstrate a clear succession in nitride coupling rates (k_B > k_A > k_C). This result was rationalized by means of density functional theory (DFT). The ener-

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getics of the three coupling reactions (Scheme 3, **A–C**) were examined by DFT computations on the D3BJ-PBE0(Cosmo (THF))/def2-TZVP//D3BJ-PBE0/def2-SVP level of theory. The molecular structures of dimers **3**, **5** and **6** are well reproduced with singlet (**3**), doublet (**5**) and triplet (**6**) ground states, respectively (Tables 1 and 2 and ESI†). Ferromagnetic coupling (J = 39 cm^{-1}) of the iridium(II) ions in **6** results in the triplet state being slightly more stable than the open shell singlet state by 0.4 kcal mol^{-1}. The computed electronic ground states resemble the simplified picture that arises from symmetry considerations (Fig. 1) for 12 (**3**), 11 (**5**) and 10 (**6**) π-electron configurations, respectively. Accordingly, the geometry of **5** is distorted along the Ir–N–N–Ir axis avoiding uneven occupation of degenerate π-MOs. Similarly, coplanar arrangement of the two Ir(PNP) fragments would also lift π-MO degeneracy due to mixing of the Ir–N–N–Ir π-manifold with PNP nitrogen lone-pairs. While such a coplanar conformation is not accessible with bulky P*t*Bu$_2$ groups (see below), the PMe$_2$-truncated model of **5** favors such a conformation, emphasizing the importance of considering the full model.

Computational evaluation of reactions **A–C** (Scheme 3) fully reproduces our experimental results (Table 2 and ESI†). Nitride coupling is thermodynamically strongly favored for all three reactions. However, the driving force within the redox series **3** (ΔG_{298} = −101.5 kcal mol^{-1}), **5** (ΔG_{298} = −97.1 kcal mol^{-1}), and **6** (ΔG_{298} = −73.9 kcal mol^{-1}) decreases, as expected from the destabilizing population of an Ir–N π*-antibonding orbital in **2**. For reaction **A**, a kinetic barrier was computed ($\Delta G^{\ddagger}_{298}$ = 21.0 kcal mol^{-1}) that is in excellent agreement with experiment (≈ 22 kcal mol^{-1}).[12] It mainly arises from the entropic contribution with only a small electronic fraction. The transition state, $^{BS(1,1)}$**TS$_A$** (Fig. 4A), was located on the open shell singlet surface at a rather long N–N distance (2.06 Å). The spin is eventually quenched at significantly smaller N–N distances, leading to the closed shell ground state of dimer **3**. The TS structure features the two Ir(PNP) monomer fragments being essentially perpendicular to each other. The two Ir–N groups are offset from collinearity along both relative orientations of the Ir(PNP) planes resulting in an out-of-plane zig-zag transition state. The orthogonalized sets of natural frontier orbitals represent hybrid orbitals from

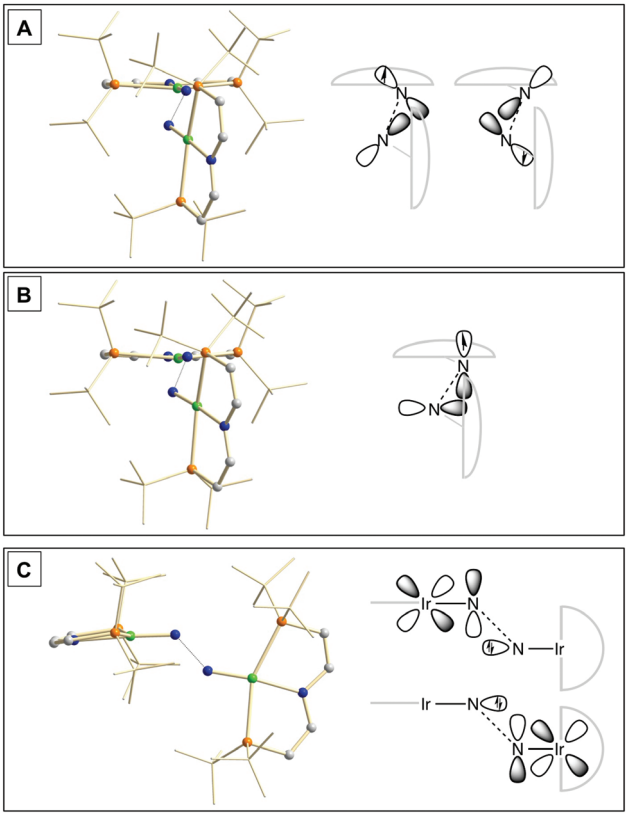

Fig. 4 Transition state structures of $^{BS(1,1)}$**TS$_A$** (**A**), ^{2}**TS$_B$** (**B**) and ^{1}**TS$_C$** (**C**) and schematic representations of the NBO interactions.

SOMO/LUMO mixing of the nitride monomers. π/π-Orbital overlap between the two monomers is further increased by slightly tilting the two monomer fragments towards each other (Ir–N–N–Ir dihedral angle: 152°). The previously computed PMe$_2$-truncated model of reaction **A** also gave a good agreement for the driving force (ΔG_{298} = −108.2 kcal mol^{-1}) but underestimation of the reaction barrier ($\Delta G^{\ddagger}_{298}$ = 13.7 kcal mol^{-1}).[12] In fact, the smaller model featured a distinctly different conformation for the TS with almost coplanar relative orientation of the two monomers. However, the minute electronic contributions to biradical coupling is almost identical for both models, while the entropic contribution seems underestimated in case of the small model presumably due to steric constraints for vibrational and rotational degrees of freedom of the *t*Bu groups in the TS.

Coupling of ^{2}**2** and ^{1}**4** to ^{2}**5**, was computed to be essentially electronically barrierless with a free energy of activation $\Delta G^{\ddagger}_{298}$ = 10.1 kcal mol^{-1}, confirming the experimentally observed rapid coupling reaction **B**. The transition state ^{2}**TS$_B$** (Fig. 4B) exhibits strong structural similarities with $^{BS(1,1)}$**TS$_A$** concerning the N–N distance (D_{N-N} = 2.00 Å) and the out-of-plane zig-zag relative orientation of the two nitride fragments (Ir–N–N–Ir dihedral angle: 142°). Accordingly, ^{2}**TS$_B$** also features a π/π-interaction between the two monomers, here arising from orbital overlap of the SOMO (**2**) and LUMO (**4**) monomer orbitals, respectively. The spin density is equally distributed over

Table 2 Calculated reaction electronic energies (ΔE including zero point energy) and free energies (ΔG) for the coupling reactions **A–C** in kcal mol^{-1} and N–N distances in the transition states (D_{N-N}) in Å

	A		**B**		**C**	
	^{2}**2** + ^{2}**2** → ^{1}**3**		^{2}**2** + ^{1}**4** → ^{2}**5**		^{1}**4** + ^{1}**4** → ^{3}**6**	
Reaction	ΔE	ΔG_{298} [b]	ΔE	ΔG_{298} [b]	ΔE	ΔG_{298} [b]
TS	6.9[a]	21.0	1.2[a]	10.1	13.8[a]	29.3
Product	−121.8[b]	−101.5	−115.6[b]	−97.1	−91.3[b]	−73.9
D_{N-N}	2.057		2.003		1.841	

[a] With respect to encounter complexes (ESI). [b] With respect to starting complexes.

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the mixed-valent dimer indicating strong electronic coupling and charge delocalization in the TS.

In contrast, reaction **C** is associated with a considerably higher computed kinetic barrier ($\Delta G^{\ddagger}_{298}$ = 29 kcal mol^{-1}), which is in agreement with the enhanced thermal stability of **4**. The transition state ^{1}**TS$_C$** (Fig. 4C) is located on the singlet surface. At a later stage there is a change to an open shell singlet and eventually the triplet ground state of ^{3}**6** (ESI†). ^{1}**TS$_C$** exhibits significant differences compared with $^{BS(1,1)}$**TS$_A$** and ^{2}**TS$_B$**: The two Ir–N bond vectors are almost parallel (Ir–N–N–Ir dihedral angle: 178°) and the offset from collinearity only along one of the Ir(PNP) planes results in an in-plane zig-zag transition state. The closed-shell σ/π-orbital interaction between the two monomers resembles the TS previously considered by Burger.[11] In the present case though different Ir–N π*-orbitals are involved on the two monomers, respectively, due to the perpendicular orientation of the two monomer fragments. Importantly, the N–N distance in ^{1}**TS$_C$** (1.84 Å) is considerably shorter compared with those in $^{BS(1,1)}$**TS$_A$** and ^{2}**TS$_B$** (both >2.00 Å) indicating that a stronger orbital interaction is necessary to overcome the kinetic barrier.

Hence, the computational evaluation confirms the order of relative coupling rates that was found experimentally ($k_B > k_A > k_C$). Albeit partitioning of TS energies into the contributing terms was not done within this study, the TS structures allow for a qualitative rationalization that is in line with the observed order: Coulombic repulsion of the two positively charged nitrides is expected to contribute to the higher barrier of reaction **C** *vs.* **A**. In turn, the TS of reaction **B** exhibits strong charge delocalization over both monomer moieties, tantamount a stabilizing Coulombic effect. Besides these charge considerations, barrierless coupling of **2** and **4** involves direct overlap of the respective SOMO and LUMO orbitals while diradical coupling reaction **A** necessitates some electronic rearrangement within $^{BS(1,1)}$**TS$_A$** with respect to ^{2}**2**. However, in all three TSs relatively large N–N distances were found and NBO analysis of ^{1}**TS$_C$** reflects only a donor–acceptor interaction between electronically weakly perturbed monomers. Hence, these results suggest for the Coulombic interaction to be the main contributor to the computed order of reaction barriers.

Conclusions

We previously reported the coupling of iridium(IV) nitride **2** to the Ir$^{I/I}$ N_2-bridged dimer **3**.[12] In contrast, iridium(V) nitride **4** does not undergo coupling at room temperature. In the present paper, the other products of formal nitride coupling besides **3**, *i.e.* the N_2-bridged Ir$^{I/II}$ and Ir$^{II/II}$ complexes **5** and **6**, were synthesized and structurally characterized. None of the dimers reveal structural or spectroscopic features indicative of an appreciable degree of N_2-activation. These results suggest that nitride **4** is also inherently thermodynamically unstable with respect to N_2-coupling. This notion is supported by DFT computations, though indicating a considerably smaller driving force for homocoupling of **4** *vs.* **2**. Examination of the kinetics of homo- and heterocoupling reactions **A**–**C** revealed the order $k_B > k_A > k_C$, also supported by DFT. Analysis of the TS structures indicate that the Coulombic interactions between the nitrides favor this order, leading to stabilization of ^{2}**TS$_B$** by charge delocalization and repulsion within ^{1}**TS$_C$**. Hence, this work indicates some guidelines for thermodynamic and kinetic stabilization of electron rich transition metal nitrides.

Experimental section

Materials and methods

All experiments were carried out using Schlenk (argon atmosphere) and glove-box (argon atmosphere) techniques. All solvents were dried by passing through columns packed with activated alumina. Deuterated solvents were obtained from Euriso-Top GmbH, dried over Na/K (Toluene-d_8) or CaH_2 (CD_2Cl_2), distilled by trap-to-trap transfer *in vacuo*, and degassed by three freeze–pump–thaw cycles, respectively. $[FeCp_2]PF_6$ (SigmaAldrich) and $AgSbF_6$ (SigmaAldrich) were used as purchased. Silica gel 60 silanized was purchased from Merck KGaA and heated *in vacuo* for 5 days prior to use. **1**, **4**, and Na/Hg were prepared according to published procedures.[12,19]

Irradiation was carried out with a Lot Oriel Xe-Short arc lamp. Cyclic voltammograms were recorded with a Metrohm Autolab PGSTAT101 using Ag/Ag$^+$ reference-, glassy-carbon working- and Pt-wire counter-electrodes. Pt-net working-electrodes and a spectro-electrochemistry cuvette (0.1 cm optical pathway) were used for UV/VIS-spectroelectrochemical measurements in combination with an Avantes AvaSpec-2048×14 UV/VIS-spectrometer. Otherwise UV/VIS-spectra were recorded with an Agilent Cary 300 UV/VIS-spectrometer at room temperature with a 0.1 mM concentration in quartz cuvettes (1 cm optical pathway). NMR-spectra were recorded on a Bruker Avance 300/400 spectrometer and calibrated to the residual proton resonance of the solvent (CD_2Cl_2: δ_H = 5.32 ppm; δ_C = 54.00 ppm; Toluene-d_8: δ_H = 7.09, δ_C = 137.86). ^{31}P-NMR chemical shifts are reported relative to external phosphoric acid (δ = 0.0 ppm). Signal multiplicities are abbreviated as: s (singlet), h (heptet), br (broad). Experimental X-band EPR spectra were recorded on a Bruker ELEXSYS-II E500 CW-EPR. The spectra were simulated by iteration of the anisotropic *g*-values, (super)hyperfine coupling constants, and line widths using the EPR-simulation program W95EPR developed by Prof. Dr Frank Neese. ESI-MS-experiments employed a micrOTOF-Q II instrument (Bruker Daltonik). IR spectra were recorded with a Thermo Scientific Nicolet iZ10 FT/IR spectrometer at r.t. IR spectra in solution were recorded with a thin layer IR cell.

Improved synthesis of [IrCl(PNP)]. To a suspension of $[Ir(COE)_2Cl]_2$ (53.8 mg, 60.0 μmol, 1.00 eq.) in THF (50 mL) ($^{t}Bu_2PCH_2CH_2)_2NH$ (43.4 mg, 120 μmol, 2.00 eq.) in THF (5 mL) is added. The reaction is stirred for 15 min at room temperature and benzoquinone (32.4 mg, 300 μmol, 5.00 eq.) is added. After stirring for 16 h at room temperature the sus-

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pension is extracted with THF (3 × 5 mL), silanized silica is added and the solvent is removed *in vacuo* under intense stirring. The solid is extracted with pentanes (5 × 5 mL), the solvent removed *in vacuo* and the remaining residue dissolved in benzene and lyophilized. (Yield: 56 mg, 95.9 μmol, 80%). NMR data were identical with the previously reported synthesis. Anal. Calc. for $C_{20}H_{40}IrNP_2$ (584.16): C, 41.12; H, 6.90; N, 2.40. Found: C, 41.41; H, 7.05; N, 2.15.

Improved synthesis of [(N$_2$){Ir(PNP)}$_2$] (3). Under the exclusion of light, [IrN(PNP)][PF$_6$] (50.0 mg, 70.7 μmol, 1.00 eq.) is dissolved in THF (3 mL) and NaHg (1 M, 1.24 g, 91.9 μmol, 1.30 eq.) is added and the solution is stirred for 1 min at room temperature. The solvent is removed *in vacuo* and the residue is extracted with benzene (6 × 0.8 mL). The product is isolated by column chromatography (2 cm silanized silica gel, 1.5 cm diameter; benzene). (Yield: 27.0 mg, 23.4 μmol, 66%). NMR data were identical with the previously reported synthesis. Anal. Calc. for $C_{40}H_{80}Ir_2N_4P_2$ (1125,44): C, 42.69; H, 7.17; N, 4.98. Found: C, 43.44; H, 7.32; N, 4.30.

Synthesis of [(N$_2$){Ir(PNP)}$_2$]PF$_6$ (5). Under the exclusion of light, **3** (5.0 mg, 4.4 μmol, 1.0 eq.) and [Fe(C$_5$H$_5$)$_2$]PF$_6$ (1.5 mg, 4.4 μmol, 1.0 eq.) are dissolved in DCM (2 mL) at −35 °C and stirred for 5 min. The solvent is removed *in vacuo*. The residue is exhaustively washed with toluene (−35 °C, 8 × 2 mL) to remove ferrocene, extracted with THF (−35 °C, 4 × 2 mL) and the solvent is removed in *in vacuo*. ^{1}H-NMR: (300 MHz, CD_2Cl_2, RT): δ = 39.1 (s br, 4H, C***H***), 5.21 (s br, 72H, PCC***H***$_3$) −129.0 (s br, 4H, C***H***). ^{1}H-NMR: (300 MHz, THF-d$_8$, RT): δ = 42.7 (s br, 4H, C***H***), 5.45 (s br, 72H, PCC***H***$_3$) −140.1 (s br, 4H, C***H***). ^{31}P{^{1}H}-NMR: (121.49 MHz, CD_2Cl_2, RT): δ = −145.0 (h, $^2J_{PF}$ = 711 Hz, ***P***F$_6$). ESI-MS (*m*/*z*): [M]$^+$ calcd for $C_{40}H_{80}N_4P_4Ir_2$ 1124.457; found: 1124.406.

Synthesis of [(N$_2$){Ir(PNP)}$_2$](SbF$_6$)$_2$ (6). Under the exclusion of light, **3** (4.0 mg, 3.6 μmol, 1.0 eq.) and $AgSbF_6$ (2.4 mg, 7.1 μmol, 2.0 eq.) are dissolved in DCM (−70 °C, 0.5 mL) and the solution is shaken for 1 min. ^{1}H-NMR: (400 MHz, CD_2Cl_2, −70 °C): δ = 29.09 (s br, 4H, C***H***), 21.78 (s br, 72H, PCC***H***$_3$) −11.25 (s br, 4H, C***H***).

Coupling of 2 in the presence of catalytic amounts of 4. [Ir(N$_3$)(PNP)] (5.0 mg, 8.5 mmol, 1.0 eq.) is dissolved in THF-d$_8$ (0.4 mL) and irradiated at −80 °C for 20 minutes. The solution is frozen with liq. N_2 and **4** (0.6 mg, 0.85 mmol 0.1 eq.) in DCM-d$_2$ (0.1 mL) is added to the frozen solid. The solution is thawed and NMR spectra were recorded at −70 °C (For an amount of 1 mol% of 3 only 0.01 mL of the DCM-d$_2$ solution is added).

Thermal stability of 4. **4** (5.3 mg, 7.5 mmol, 1.0 eq.) is dissolved in DCM-d$_2$ (0.54 mL) and naphthalene (1.9 mg, 15 mmol, 2.0 eq.) is added. The solution is heated to 40 °C for 5 days.

Crystallographic details

Suitable single crystals for X-ray structure determination were selected from the mother liquor under an inert gas atmosphere and transferred in protective perfluoro polyether oil on a microscope slide. The selected and mounted crystals were transferred to the cold gas stream on the diffractometer. Intensity data for **5** and **6** were obtained at 100 K on a Bruker D8 three-circle diffractometer, equipped with a PHOTON 100 CMOS detector and an INCOATEC microfocus source with Quazar mirror optics (λ = 0.71073 Å). The data obtained were integrated with SAINT and a multi-scan absorption correction with SADABS was applied. Both structures were solved and refined using the Bruker SHELX 2014 software package.[20] All non-hydrogen atoms were refined with anisotropic displacement parameters. All C–H hydrogen atoms were refined isotropically on calculated positions by using a riding model with their U_{iso} values constrained to 1.5 U_{eq} of their pivot atoms for terminal sp^3 carbon atoms and 1.2 times for all other carbon atoms. Crystal structure determination of $C_{40}H_{80}F_6Ir_2N_4P_5$ (**5**): CCDC 1437698, M = 1270.33, orthorhombic, a = 13.1762(5), b = 13.3606(5), c = 28.3330(11) Å, U = 4987.8(3) Å^3, T = 100 K, space group $P2_12_12_1$ (no. 19), Z = 4, 78 343 reflections measured, 12 410 unique (R^{int} = 0.0589), which were used in all calculations. The final w$R(F^2)$ was 0.0454 (all data). Crystal structure determination of $C_{52}H_{104}F_{12}Ir_2N_4O_3P_4Sb_2$ (**6**): CCDC 1437699, M = 1813.17, monoclinic, a = 15.7367(8), b = 15.1562(7), c = 15.2970(8) Å, b = 113.649(2)°, U = 3342.1(3) Å^3, T = 100 K, space group $P2/c$ (no. 13), Z = 2, 55 887 reflections measured, 8353 unique (R^{int} = 0.0446), which were used in all calculations. The final w$R(F^2)$ was 0.0600 (all data).

Computational details

DFT calculations were performed with the Turbomole 7.0 package[21] applying the PBE0[22] functional and Grimme's dispersion correction with Becke–Johnson damping (D3BJ).[23] Ahlrich's revised basis sets were used (def2-SVP for structure optimization and def2-TZVP for single point energies) with a full basis for all elements but Ir for which a Stuttgart–Dresden 60 electron core potential has been used, replacing the inner shell 1s–4f orbitals.[24] No symmetry restrains were imposed and the optimized structures were defined as minima (no negative eigenvalue) or transition states (one negative eigenvalue) by vibrational analyses at the same level of theory. In case of ^{1}**TS$_C$** a small second imaginary mode at −13 cm^{-1} was observed. Solvent effects were accounted for by applying the COSMO model (ε = 7.6 for THF) in the single point calculations.[25] Final energies were obtained by adding zero point vibrational energies or enthalpies from the optimizations to the SCF energies, in short notified as D3BJ-PBE0(COSMO(THF))/def2-TZVP//D3BJ-PBE0/def2-SVP. The electronic structures of $^{BS(1,1)}$**TS$_A$** and **6** were evaluated by the broken symmetry protocol and the open shell singlet (OSS) structures (BS1,1) were located. The energies of the (multi-determinant) OSS were estimated from the energy ε_0 of the optimized single-determinant broken symmetry solution and the energy ε_1 from a separate unrestricted triplet (m_s = 1) calculation at the same geometry with the same functional and basis set, using the approximate spin correction formula proposed by Yamagushi:[26]

$$\varepsilon_S \approx \frac{S_1{}^2\varepsilon_0 - S_0{}^2\varepsilon_1}{S_1{}^2 - S_0{}^2}$$

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NBO analyses at the D3BJ-PBE0/def2-SV(P) level were performed with Gaussian 09 RevD.01[27] coupled to NBO6.0[28] using the structures optimized as described above.

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