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Journal

Journal of the American Chemical Society, 138(24)

ISSN

0002-7863

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Publication Date

2016-06-22

DOI

10.1021/jacs.6b03329

Peer reviewed



Published in final edited form as:

J Am Chem Soc. 2016 June 22; 138(24): 7468–7471. doi:10.1021/jacs.6b03329.

Electron Paramagnetic Resonance Characterization of Three Iron–Sulfur Clusters Present in the Nitrogenase Cofactor Maturase NifB from *Methanocaldococcus infernus*

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Abstract

NifB utilizes two equivalents of S-adenosyl methionine (SAM) to insert a carbide atom and fuse two substrate [Fe–S] clusters forming the NifB cofactor (NifB-co), which is then passed to NifEN for further modification to form the iron–molybdenum cofactor (FeMo-co) of nitrogenase. Here, we demonstrate that NifB from the methanogen *Methanocaldococcus infernus* is a radical SAM enzyme able to reductively cleave SAM to 5'-deoxyadenosine radical and is competent in FeMo-co maturation. Using electron paramagnetic resonance spectroscopy we have characterized three [4Fe–4S] clusters, one SAM binding cluster, and two auxiliary clusters probably acting as substrates for NifB-co formation. Nitrogen coordination to one or more of the auxiliary clusters in NifB was observed, and its mechanistic implications for NifB-co dissociation from the maturase are discussed.

NifB belongs to the superfamily of radical S-adenosyl methionine (SAM) enzymes.¹ These enzymes are present in all forms of life and provide the most common way in biology to perform directed radical based chemistry by utilizing a [4Fe–4S] cluster to coordinate SAM and generate a 5'-deoxyadenosyl radical (5'dA[•]). The 5'dA[•] then typically abstracts an H atom from the substrate to initiate catalysis.² In NifB, simple [Fe–S] clusters are converted into a nitrogenase-specific [Fe–S] cluster of high nuclearity, termed NifB-co.³ NifB-co is the source of most (or all) Fe and S present in FeMo-co along with its central C atom (Scheme 1).^{4–7} The insertion of carbide to fuse [Fe–S] clusters is a unique reaction among radical SAM enzymes.¹

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Supporting Information: The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/jacs.6b03329.

Materials and Methods (PDF)

Supporting Schemes, Tables and Figures (PDF)

Notes: The authors declare no competing financial interest.

NifB proteins, first purified from *Azotobacter vinelandii* and *Klebsiella oxytoca*, exhibited spectroscopic and catalytic properties typical of radical SAM proteins and were shown to be competent in FeMo-co maturation.^{3,8} Further investigations were limited due to instability of the purified NifB when not fused with the downstream maturase NifEN.^{9–11} NifB from methanogens was shown to be more stable target for spectroscopic studies.^{12,13} Purification of NifB from the methanogens *Methanocaldococcus infernus*, *Methanosarcina acetivorans*, and *Methanobacterium thermoautotrophicum* confirmed their ability to use SAM as substrate to form dAdoH and SAH, the products of the radical SAM and demethylation reactions, respectively.^{12,14} These homologues were competent in FeMo-co maturation and were proposed to each have three [4Fe–4S] clusters based on the Fe content of fully reconstituted enzymes.

Here, we expand our understanding of NifB by characterizing the three [Fe–S] clusters of *M. infernus* NifB (*M*NifB) through the use of site-directed variants and electron paramagnetic resonance (EPR) spectroscopy. Further, we detect N ligand(s) to at least one of these clusters and discuss the implications this may have on the release of NifB-co to NifEN.

The metin_0554 gene from *M. infernus* was heterologously expressed as a His₁₀-tagged protein in *E. coli* as described.¹³ Anaerobically purified *M*NifB ran as a single 35–kDa band in SDS-PAGE (Figure S1A), in good agreement with its predicted molecular weight of 37 543 Da and TOF/TOF mass spectrometry analysis (37 483 Da; Figure S2). The UV–vis spectrum of as-isolated *M*NifB displayed features of [Fe–S] proteins (Figure S1B). Among these features, the shoulder at 320 nm and the peak at 400–420 nm are the most defining, as is the broad tail extending beyond 700 nm. This absorption feature is assigned to S → Fe charge transfer transitions characteristic of [4Fe–4S]²⁺ clusters. Consistently, as isolated *M*NifB contained, on average, 3.3 Fe and 4 acid-labile S atoms per monomer (Table S1). *In vitro* reconstitution of [Fe–S] clusters was carried out either on as-isolated *M*NifB or on samples previously stripped of Fe and S (apo-*M*NifB). Incorporation of Fe and S was higher on apo-*M*NifB samples (Table S1), and these preparations exhibited UV–vis spectra with the most prominent 320 and 420 nm features (Figure S1B).

As-isolated *M*NifB was active in the NifB-dependent *in vitro* FeMo-co synthesis and apo-NifDK activation assay. Apo-NifDK activation was strictly dependent on addition of *M*NifB or NifB-co (isolated from *K. oxytoca* cells and used here as control of NifB activity; Figure S1C). Addition of either apo-*M*NifB or aerobically purified *M*NifB did not activate apo-NifDK (data not shown). The capacity of *M*NifB to transfer a methyl group from SAM, generate 5′ dAdo•, and initiate FeMo-co synthesis were determined by measuring concomitant production of *S*-deoxyadenosyl-L-homocysteine (SAH), 5′-deoxyadenosine (dAdoH), and reconstituted NifDK in a single assay. SAH and AdoH production correlated with *M*NifB-dependent activation of nitrogenase over time (Figure S1D). Interestingly, demethylation of SAM proceeded at higher rates than dAdoH formation (3:2 SAH to dAdoH ratio), and this difference was maintained over time. Incubation with ¹³CD₃-SAM resulted in production of deuterated 5′-deoxyadenosine (dAdoD), demonstrating abstraction of a H atom from the methyl group of SAM (Figure S3).

The Fe and S content of reconstituted wild-type (WT) *MNifB* (10 ± 0.2 Fe and 11.2 ± 0.4 S atoms per monomer, Table S1) was consistent with the presence of, at minimum, two [Fe–S] clusters in addition to the SAM-binding [4Fe–4S] cluster. The SAM-binding cluster is anchored to the enzyme by three cysteine thiolate side chains, identified by the canonical CX₃CX₂C motif. Substitution of these Cys residues (C36A/C40A/C43A) in variant SM (Table 1) resulted in reduction of Fe and S content, changes in the UV–vis spectrum, and inability to activate apo-NifDK (Figure 1A–D and Table S1). NifDK activation ability was also lost in the AM variant carrying amino acid substitutions in the C-terminal CXXC U-type motif, which is similar to those shown to bind [Fe–S] clusters for storage and transfer^{15,16} and in the DM variant with amino acid substitutions in both domains (Table 1 and Figure 1D). [Fe–S] cluster reconstituted DM preparations exhibited UV–vis features, as well as Fe and S content consistent with a [4Fe–4S] cluster (Figure 1C and Table S1), suggesting the presence of one additional [Fe–S] cluster-binding motif in *MNifB*. Anaerobic size-exclusion chromatography of reconstituted WT and the SM, AM, and DM *MNifB* variants indicated these proteins were monomers (Table S2), suggesting that each *MNifB* monomer is able to coordinate all observed clusters.

To investigate the composition of the [Fe–S] clusters present in WT *MNifB*, continuous wave \times -band (9.4 GHz) EPR analysis was performed on reconstituted enzyme preparations. WT and the SM and AM variants reduced by sodium dithionite (DTH), in the presence or absence of SAM, exhibited spectra with components representative of multiple $S = 1/2$ [Fe–S] clusters (Figure 2A–D and Figure S4). The EPR spectrum of the DM variant was consistent with a single [4Fe–4S] cluster (Figure 2E), in agreement with its Fe and S content (Table S1). The experimental spectra of WT and all the variants were simulated using the DM variant as starting point (see Table S3 for simulation parameters, Table S4 for spin quantitations, and Figure S5 for individual component spectra).

Temperature-dependent EPR, which has proven useful to establish the nuclearity of [Fe–S] clusters,^{17,18} was utilized to identify individual species present. Data shown in Figure S6 suggest that the three clusters present in *MNifB* are [4Fe–4S] clusters. Additionally, the g_1 feature of each cluster becomes clear when examining the temperature dependence of the different spectra, providing spectroscopic handles for these clusters. Deconvolution through spectral subtractions was performed to support individual cluster assignments. Unique features for each of the species present, and the positions of g_1 and g_3 for each of the clusters were verified (Figure S7). The cluster present in the DM variant, termed AC2, showed features typical of a [4Fe–4S] cluster, such as a fairly axial g-tensor ($g = [2.039 \ 1.923 \ 1.886]$) and signal disappearance above 40 K. Three clusters were identified in the SM variant. The [4Fe–4S] cluster described above for the DM variant, an additional [4Fe–4S] cluster, termed AC1, with g-values of $[2.062 \ 1.917 \ 1.875]$, and a third cluster that was only observed above 50 K and had a much more rhombic g-tensor of $[2.058 \ 1.985 \ 1.909]$ consistent with a [2Fe–2S]. This [2Fe–2S] cluster was only apparent in the SM variant and was tentatively assigned to a degraded cluster, termed ACx. The EPR spectrum of the AM variant could be simulated with a combination of AC2 and a new [Fe–S] cluster with g-values of $[2.038 \ 1.931 \ 1.916]$, which has been assigned to the SAM-binding cluster (RS). The g-values and temperature dependence of the EPR spectrum for RS are in agreement with reported SAM binding clusters.¹

Examination of WT *Mn*NifB by three pulse electron spin echo envelope modulation (3P-ESEEM) revealed nitrogen modulations to the echo from one or more of the [Fe–S] clusters (Figure 3, black trace). Incubation with isotopically enriched ^{15}N -SAM, carried out to investigate changes in spectral features attributed to SAM coordination, resulted in no change in modulations (Figure 3, blue trace) as would be expected if this modulation derived from coupling to SAM nitrogen. This lack of $^{14}\text{N}/^{15}\text{N}$ SAM isotope sensitivity of the ESEEM spectra suggests proteinaceous nitrogen coordination to one or more of the [Fe–S] clusters. Due to its simplified EPR spectrum and cluster content, the DM variant was selected to investigate nitrogen coordination from the protein to the AC2 cluster. The nitrogen modulations observed in the DM variant (Figure 3, red trace) were very similar to those of the WT, suggesting that AC2 is coordinated by at least one nitrogen ligand. While the nitrogen modulations arising from AC2 are present in the spectra of WT *Mn*NifB with peaks at 2 and 0.7 MHz (Figure 3), there was also evidence for additional nitrogen modulations from a second cluster (peaks at 2.3 and 1 MHz), which was tentatively assigned to AC1.

NifB from *M. infernus* is functional in nitrogenase cofactor maturation and represents a simplified homologue (lacking the NifX domain) of NifB proteins from model diazotrophs *A. vinelandii* and *K. oxytoca*. These results are in agreement with those recently reported for *M. acetivorans* and *M. thermoautotrophicum*.¹⁴ EPR and biochemical analyses indicate the presence of three [4Fe–4S] clusters in *Mn*NifB. One of these clusters is identified as the [4Fe–4S] SAM binding cluster (RS) by the canonical $\text{CX}_3\text{CX}_2\text{C}$ motif and dAdoH formation assays. While the auxiliary clusters AC1 and AC2 may be present as conduits for electron transfer, our preferred model is that they are in-fact precursors to NifB-co. Based on the temperature behavior of EPR spectra, reporting information on the relaxation dynamics of the cluster, we propose that both auxiliary clusters are present as [4Fe–4S] clusters. While the temperature dependence of an [Fe–S] cluster can help discriminate between a [4Fe–4S] and [2Fe–2S] cluster (a [4Fe–4S] cluster having faster relaxation rates than a [2Fe–2S] cluster), this analysis can become clouded when multiple clusters are present and near each other causing increased relaxation rates from dipolar coupling between them. Therefore, the presence of one [2Fe–2S] cluster cannot be completely ruled out. The assignment of NifB-co as a fusion of two [4Fe–4S] clusters or one [4Fe–4S] and one [2Fe–2S] units is consistent with the previous proposal of NifB-co comprising at least the [6Fe–9S–C] core of FeMo-co.⁶

Based on the number of strictly conserved cysteines in *Mn*NifB, there are not enough to fully coordinate the SAM-binding cluster plus two additional [4Fe–4S] clusters (Table S5). This would indicate that the two auxiliary clusters are either not saturated in Fe coordination or that some noncysteinyll residues may coordinate the auxiliary clusters as suggested by 3P-ESEEM. *Mn*NifB has two conserved His residues as well as several conserved Arg residues. Coordination by His or Arg residues is well-known for [2Fe–2S] clusters and has precedence for an auxiliary [4Fe–4S] cluster of Cp1 Fe–Fe hydrogenase, as well as in the hydrogenase cofactor maturase HydF.^{18–20} Additionally, in BioB, IscU and MitoNEET nitrogen coordination is utilized to bind transient [Fe–S] clusters.²¹ Nitrogen coordination to the cluster would provide a convenient path for cluster release through a simple protonation of the nitrogen ligand and release of NifB-co to NifEN or NifX.

Like NifB from *M. acetivorans* and *M. thermoautotrophicum*,¹⁴ *Mi*NifB produces more SAH than AdoH in turnover assays (a 3:2 ratio). We note that SAM cleavage to AdoH has never been observed to be in excess of SAH in NifB suggesting a single H abstraction followed by oxidation/deprotonation events to remove the remaining H from the transferred methyl to form carbide. Interestingly, we observed a shift in the g_3 feature at 352 mT of the WT EPR spectrum in the presence of SAM (Figure 2B), but no indication of binding of SAM to the RS cluster in the 3P-ESEEM, suggesting the existence of additional steps in the reaction mechanism to form NifB-co. In the mechanism that has been proposed for NifB,^{10,22} SAM transfers a methyl group to one of the clusters that will be fused to form NifB-co (Scheme S1). We suggest the observed perturbation to WT NifB EPR spectrum in the presence of SAM is due to methyl transfer to one of the auxiliary [Fe-S] clusters, poised for NifB-co formation, thus ensuring that, when 5' dAdo* is formed, the site of hydrogen abstraction is present preventing unwanted side reactions from the highly reactive primary carbon radical.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

We thank Juan Fontecilla for providing pRSFiscMetKDuet-1, and Etienne Mulliez and Phan  lie Perche-Letuv  e for providing ¹⁵N-SAM and ¹³CD₃-SAM. This work was supported by the European Research Council Starting Grant 205442 (to L.M.R.), MINECO Grant BIO2014-59131-R (to L.M.R.), FPI Fellowship BES-2010-038322 (to A.S.), and National Institutes of Health Grant GM-104543 (to R.D.B.).

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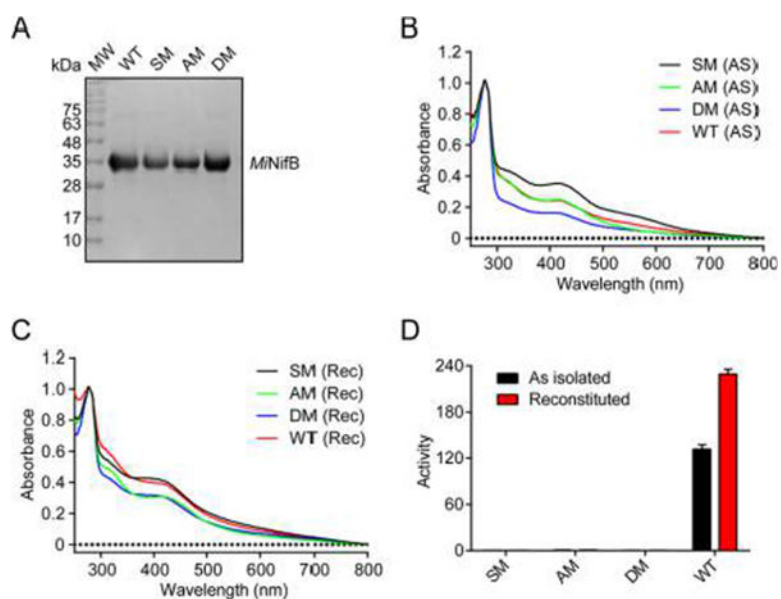


Figure 1. Biochemical properties of *MnNifB* variants lacking [Fe-S] cluster-binding motifs. (A) SDS-PAGE of pure *MnNifB* preparations. (B,C) UV-visible spectra of WT *MnNifB* (red), and the SM (black), AM (green), and DM (blue) variants in as-isolated (AS) (B) and reconstituted (Rec) (C) forms. Spectra were normalized to their absorbance at 279 nm. (D) *MnNifB*-dependent apo-NifDK reconstitution assays with *MnNifB* WT and variants. Activities are nmol of ethylene·min⁻¹·mg⁻¹ NifDK (mean ± SD; *n* = 2).

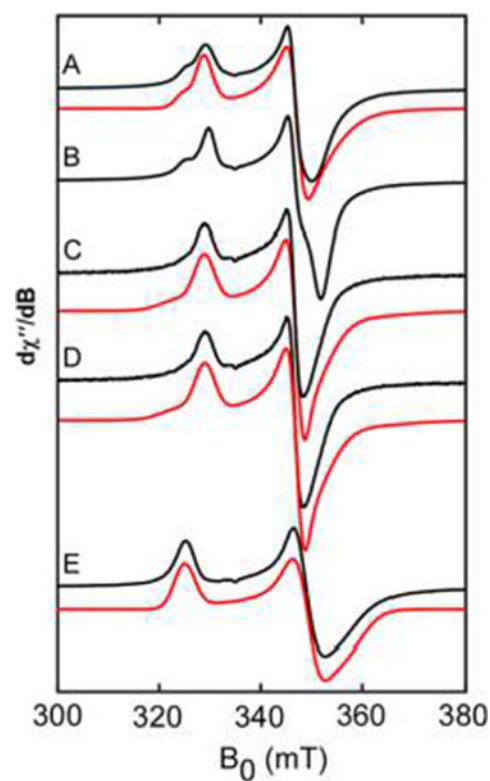


Figure 2. X-band CW EPR spectroscopy of DTH-reduced $MnNifB$ WT and site-directed variants. (A) WT $MnNifB$ (274 μM), (B) WT $MnNifB$ in the presence of 3 mM SAM, (C) SM variant (159 μM), (D) AM variant (159 μM), and (E) DM variant (252 μM). Data are shown in black with composite spectral simulations in red.

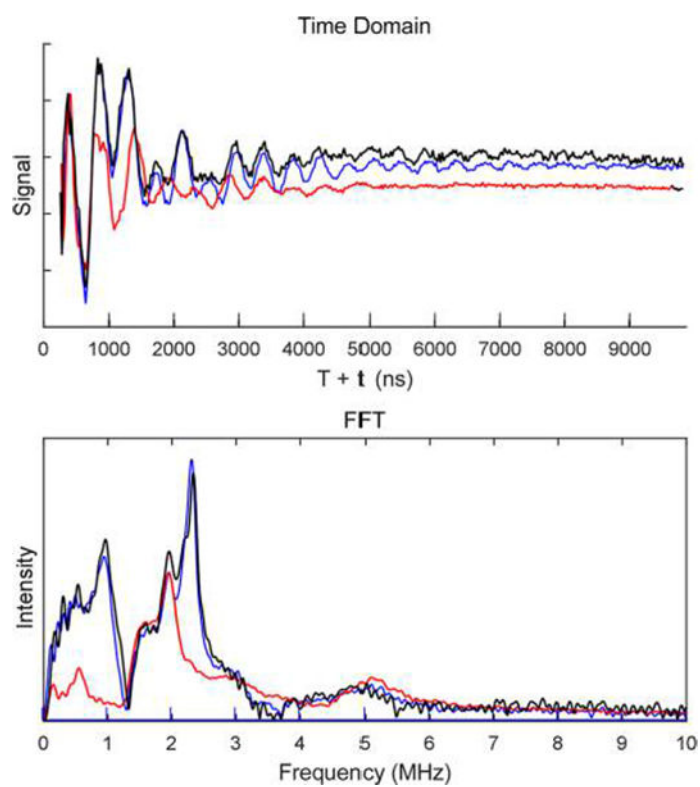
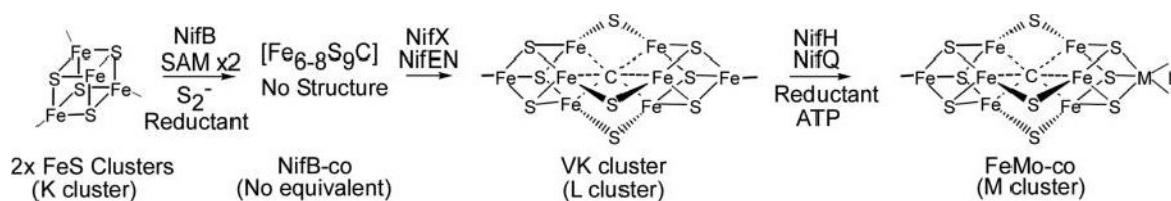


Figure 3. X-band (9.4 GHz) 3P-ESEEM time domain (top) and Fourier transform of the time domain (bottom) of *MnNifB*. Black trace, WT *MnNifB* reduced with 10 mM DTH in the presence of 3 mM ^{14}N -SAM. Blue trace, WT *MnNifB* reduced with 10 mM DTH in the presence of 3 mM ^{15}N -SAM. Red trace, DM variant reduced with 10 mM DTH in the absence of SAM.



Scheme 1. Maturation of Nitrogenase FeMo Cofactor^a

^aRequirements for metallocluster transformation are indicated for each step of FeMo-co biosynthesis. M is Mo, and L represents homocitrate. The two nomenclatures used to refer to these metalloclusters are shown.

Table 1
***MiNifB* Variant Nomenclature**

name	variant	substitutions
WT	wild type	none
SM	SAM motif variant	C36A/C40A/C43A
AM	auxiliary motif variant	C260A/C263A
DM	double variant	C36A/C40A/C43A C260A/C263A

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