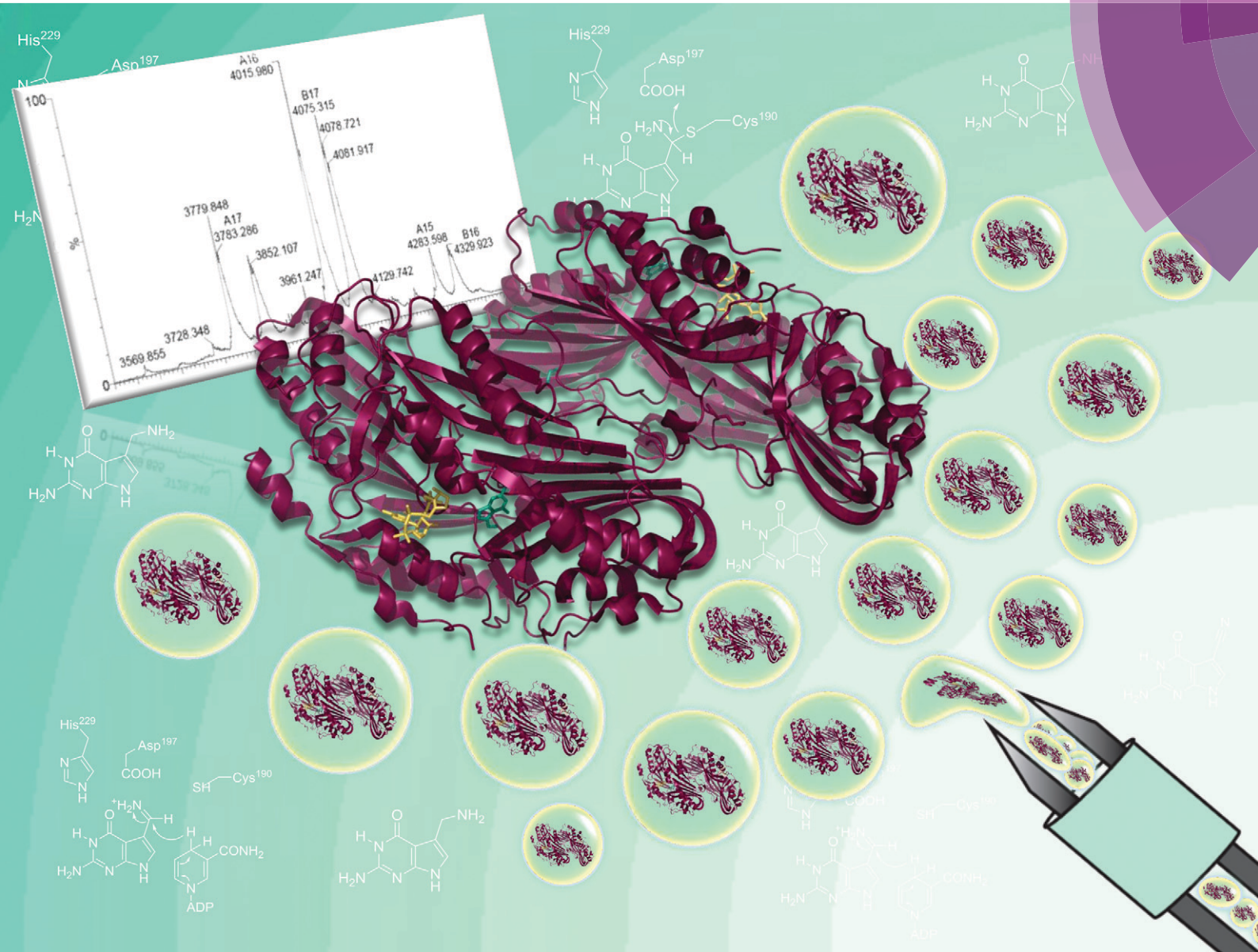


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Substrate and cofactor binding to nitrile reductase: a mass spectrometry based study

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Substrate and cofactor binding to nitrile reductase: a mass spectrometry based study†

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Nitrile reductases catalyse a two-step reduction of nitriles to amines. This requires the binding of two NADPH molecules during one catalytic cycle. For the nitrile reductase from *E. coli* (*EcoNR*) mass spectrometry studies of the catalytic mechanism were performed. *EcoNR* is dimeric and has no Rossmann fold. It was demonstrated that during catalysis each active site binds one substrate molecule. NADPH binds independent of the substrate. The PreQ₀ binding pocket of the active site is not involved in the binding of NADPH; this is in conflict with an earlier hypothesis.

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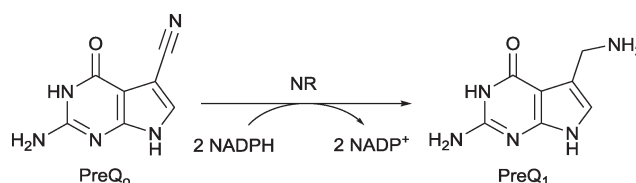
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Introduction

Nitrile reductases (NR) have been discovered as participants in the biosynthetic pathway of queuosine, displaying an unusual activity. They are the first enzymes able to catalyse the reduction of a nitrile group to a primary amine.¹ NR has emerged as a potential tool to overcome and replace traditional chemical approaches and harsh conditions needed to accomplish nitrile hydrogenation.² The nitrile reductase is a NADPH-dependent enzyme which catalyses the reduction of the nitrile group of 7-cyano-7-deazaguanine (PreQ₀) to an amino group, 7-aminomethyl-7-deazaguanine (PreQ₁) (Scheme 1). This chemoselective enzyme has been characterised in several organisms confirming this particular activity.³ The crystal structures of NR from *Bacillus subtilis* (*BsNR*)⁴ and *Vibrio cholerae* (*VcNR*)⁵ have been resolved, providing more information about the mechanism of the catalytic reaction. While the active site in all known NR's is highly conserved, the overall structure of the enzymes can be different. To date two types of structure have been elucidated. In the case of *BsNR* the enzyme forms a homodecamer (dimer of pentamers) and for *VcNR* a homotetramer (dimer of dimers) was observed (Fig. 1). In the former the active site is established at the interfaces of the five subunits, in the latter the active sites are within one protein and two monomers face each other in the dimer. Nonetheless, the active sites are virtually identical in both types of structures (Fig. 1). The

covalently bound substrate is clearly visible in the active sites. The reaction mechanism for all NRs is proposed to proceed in two reduction steps.^{3–6} In order to reduce one molecule of PreQ₀, two consecutive molecules of NADPH will act as hydride donor within the active site. The first overall step involves the formation of the covalent adduct of the PreQ₀ within the active site. By introducing mutations in the active site of the NR from *Escherichia coli* (*EcoNR*) in the positions previously elucidated for *VcNR* and *BsNR*, the essential amino acids for the catalytic cycle were confirmed.^{3c} They are shown to be Cys190, Asp197 and His229 in the schematic representation of the catalytic cycle (Scheme 2).

Remarkably, no determined structure of an NR contains a Rossmann fold, characteristic for NAD(P)H dependent enzymes. Moreover, no structures with a complete NADPH bound to the enzyme are available (see also pdb 3UXJ and 3UXV). This raises the question how the two NADPH cofactor molecules per catalytic cycle bind to the NR. Molecular dynamic simulation performed on *VcNR* suggest that the cofactor will be positioned in one of the active sites of the dimeric enzyme.⁵ The adenine moiety of the NADPH is believed to bind within the PreQ₀ pocket of one monomer of the dimer. The docking studies suggest that this occurs due to great similarity between the adenine moiety and PreQ₀. This is supposed to occur in combination with PreQ₀ docking in the



Scheme 1 Catalytic reduction of PreQ₀ with NADPH catalysed by NR.

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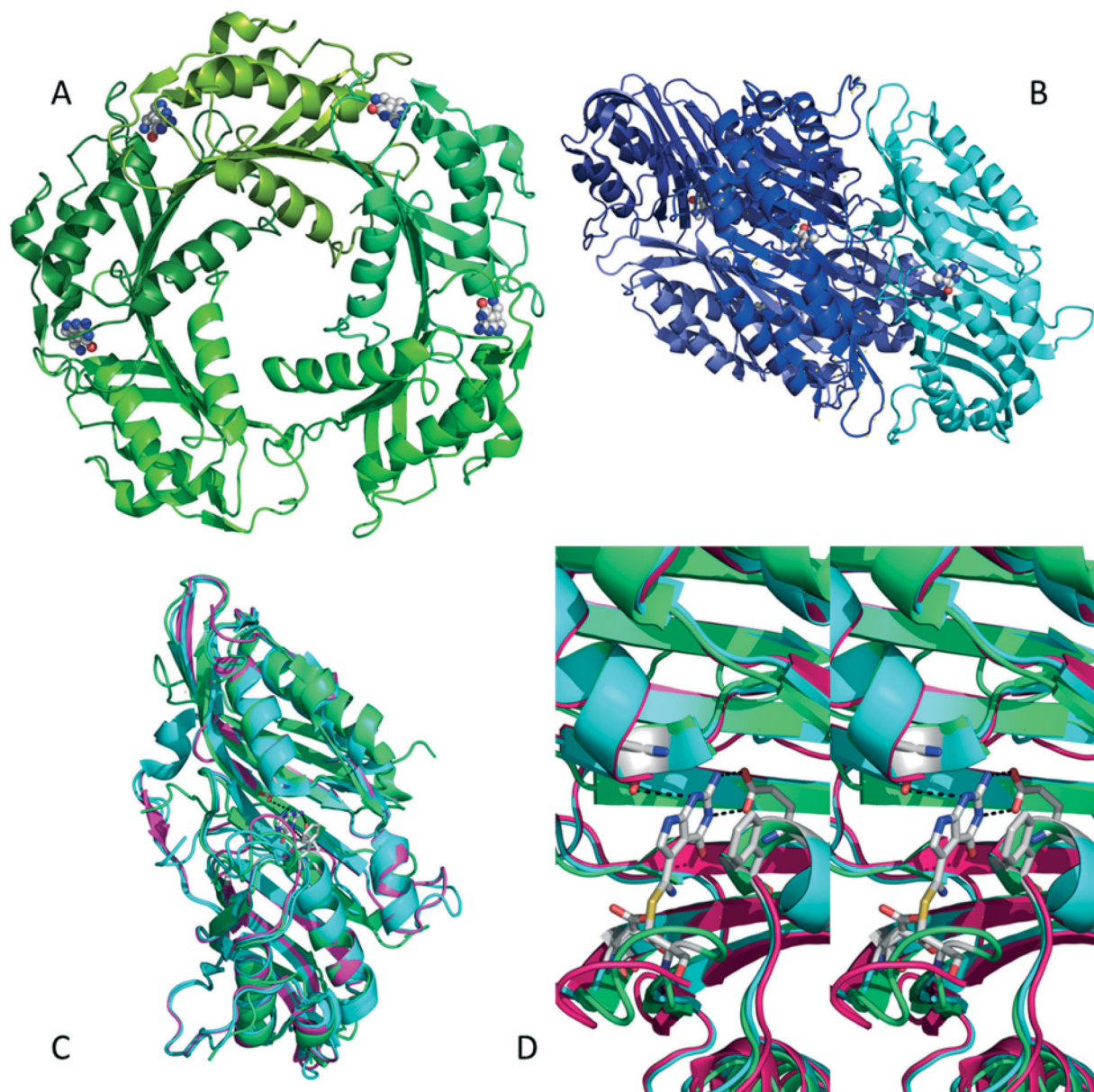


Fig. 1 A) *BsNR* is pentameric, with the active sites located at the interfaces of the subunits, the substrate is visible in four active sites (pdb 4F8B). B) *VcNR* is dimeric, with two dimers assembling to a tetramer, the substrate is visible in the active sites which are within the subunits (pdb 3BP1). C) Overlay of *VcNR* (blue, pdb 3BP1), *BsNR* (green, pdb 4F8B) and *EcoNR* (magenta), indicating the very high degree of similarity, even though the subunits might be different. D) Stereoview of the three different active sites, with PreQ₀ covalently bound; colours as in C. All pictures were generated using the PyMOL molecular graphics system.

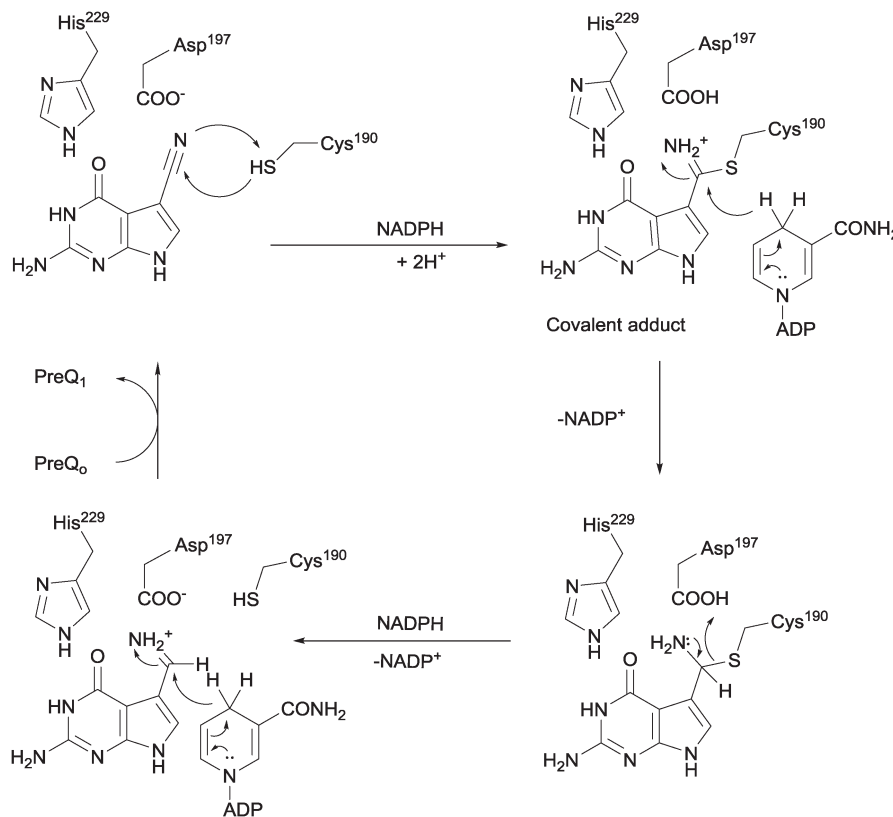
second active site of the dimer.⁵ This implies that per dimer of *VcNR* only one substrate molecule can bind in an active site as the other active site acts as anchor for the NADPH.

To probe this, direct detection of the binding of PreQ₀ and NADPH needs to be monitored. With mass-spectrometry the proposed intermediates can ideally be identified directly.^{7,8} This would allow insight into the mechanism of the reduction reaction and the binding mode of the cofactor. In our previous research a production system for *EcoNR* was successfully established^{3d} *EcoNR* was purified *via* His-tag affinity column. We found for PreQ₀ a K_m value of 2.84 μM and $k_{\text{cat}} = 5.24 \text{ min}^{-1}$ and for the cofactor NADPH $K_m = 6.42 \mu\text{M}$ and k_{cat}

$= 5.2 \text{ min}^{-1}$ for the enzyme monomer, in line with previous results (ESI[†] Fig. S1 and S2).^{1,3}

Based on the sequence analysis, *EcoNR* shows a 63.6% homology with *VcNR* (Fig. 1). As for *VcNR*, dimer and tetramer are the main conformations of *EcoNR* (Fig. S3[†]). This allows for electrospray ionisation mass-spectrometry (ESI-MS) and detection of the covalent adduct (Scheme 2), particularly as the active site is inside one monomer. The pentameric and decameric enzymes such as that from *BsNR* are less suitable for this purpose. Indeed, attempts with ESI-MS to characterise the covalent PreQ₀ adduct with *BsNR* had earlier been unsuccessful.^{3a}





Scheme 2 Cys190 is the nucleophile that attacks the nitrile, Asp197 and His229 have been suggested to take part in the catalytic mechanism of NR, His229 in particular in the positioning of the substrate.

Results and discussion

Feasibility studies

As expected the dimeric structure of *EcoNR* is essentially not affected by ESI and the MS shows the enzyme to occur predominantly as a dimer (see Fig. 2A). In ESI-MS experiments multiple charged ions of the proteins are obtained. Therefore the dimeric *EcoNR* that has a large mass (>60 000 Da) can still be measured straightforwardly. The signals represent different mass to charge ratios (m/z). All peaks in the spectra represent an intact protein or dimer in our case. From the m/z value the molecular weight of the protein can be determined. As shown in Fig. 2A, the dominate peak $m/z = 4015$ ($z = 16^+$) is correlated to the mass of dimeric *EcoNR*, in which the His-tag is lost from both of the monomers during ionisation. Also the m/z peak ($m/z = 4079$, $z = 17^+$) of the dimeric *EcoNR* with one tag still attached to the N-terminus of one of the monomers is clearly visible. The reason for the loss of the His-tag is unknown. The other peaks relate to the same dimers of *EcoNR* yet with a higher or lower number of charges (for the complete spectrum up to $m/z = 9000$ see Fig. S4[†]). Unlike the pentameric *BsNR*, *EcoNR* is thus a suitable system to study the binding of the substrates. This being PreQ₀ bound to the Cys190 as an imidothioate and NADPH of which the mode of binding still needs to be unravelled. In the currently proposed mechanism PreQ₀ and NADPH would each bind to one active site of the dimer.

For *BsNR*, adduct formation with PreQ₀ and its reduction were followed by UV-vis spectroscopy.^{3a,4b} While inconclusive for the NADPH binding, this allows ESI-MS independent proof of adduct formation. In line with those results the characteristic absorbance of the new band appearing at 376 nm was observed for *EcoNR*. Using 2–3 fold excess of PreQ₀ per active site of *EcoNR* (concentration 40–60 μ M), gentle shaking for 3 h, followed by dialysis to remove the excess of the substrate the adduct formation was confirmed by UV-vis spectrometry. Based on these results the PreQ₀ *EcoNR* should be detectable by ESI-MS (Fig. 3).

ESI-MS characterisation of the intermediates of the catalytic cycle

Initial experiments to detect the PreQ₀ *EcoNR* adduct by ESI-MS were inconclusive. Conditions of the UV-vis experiments were not suitable for ESI-MS measurements, possibly due to hydrolysis of the delicate imidothioate bond of the covalent adduct. To circumvent this, the nitrile reductase was treated with one equivalent PreQ₀ per active site and analysed immediately. No mass peak of free *EcoNR* was found. Instead a new peak $m/z = 4037$ is observed, the multiply charged ion ($z = 16^+$) of the dimeric *EcoNR* with one molecule of PreQ₀ covalently bound to each of the monomers. Complete saturation of the enzyme with the substrate (Fig. 2B) is thus achieved by treatment with one equivalent of substrate per



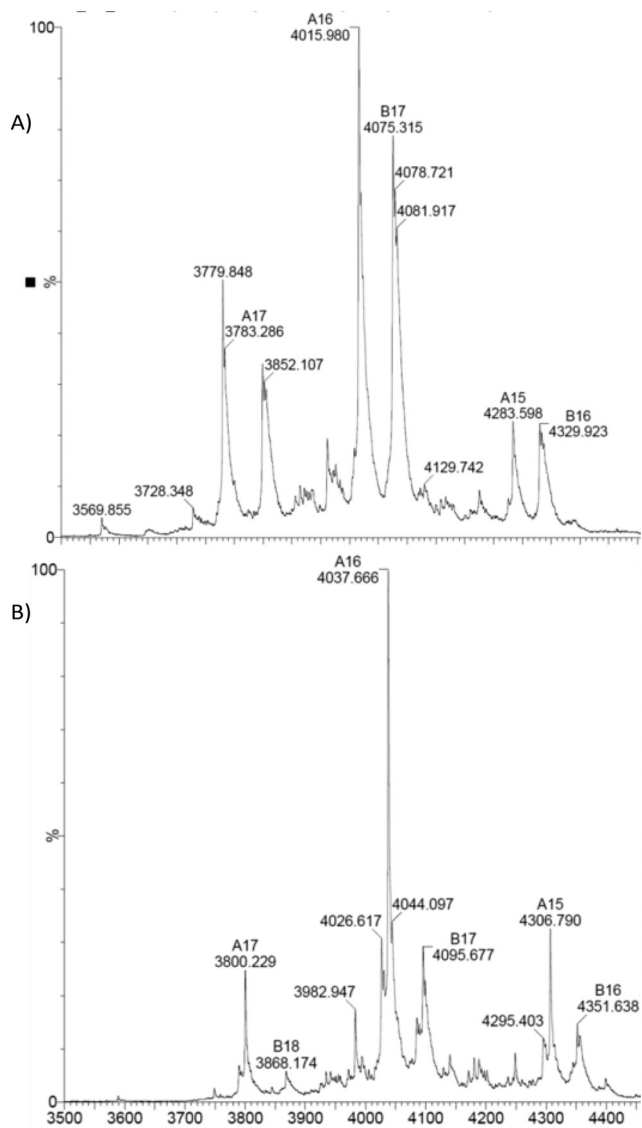


Fig. 2 A) Native *EcoNR* and B) *EcoNR* treated with one equivalent of substrate $PreQ_0$ per monomer ($m/z = 4037$).

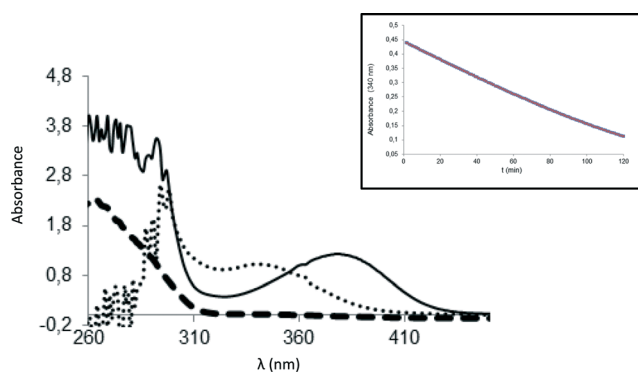


Fig. 3 UV-vis characterisation of the covalent $PreQ_0$ adduct. Solid line: covalent adduct; (–) covalent adduct after addition of an excess of NADPH; (·) *EcoNR*. Inset: Consumption of the substrate over time in presence of NADPH.

active site. This confirms the earlier crystallographic results for *VcNR* and *BsNR* (Fig. 1).

Based on the similar K_m values for NADPH and $PreQ_0$ one equivalent of NADPH per active site was added to *EcoNR*. Molecular modelling and crystal structures had suggested that only one molecule of cofactor is bound to the dimeric enzyme, as the minimum catalytic unit.⁵ Analysis of the mass-spectra revealed the appearance of a new signal ($m/z = 4062$, $z = 16^+$), with a deconvoluted mass 752 amu higher than the native *EcoNR* (Fig. 4). The new peak corresponds to the dimeric enzyme with one molecule of NADPH bound to it. This would be in line with the proposed binding of one molecule of NADPH in one active site per dimer but contradict our results from the $PreQ_0$ binding studies.⁵ Alternatively, the NADPH could bind to the dimeric *EcoNR*, however, in a different binding pocket than that of $PreQ_0$.

To probe the mode of binding of NADPH and $PreQ_0$ to *EcoNR* both were added. First, *EcoNR* was treated with one equivalent of $PreQ_0$ per monomer followed by the addition of one equivalent of NADPH per active site/monomer. In a second experiment, NADPH and $PreQ_0$ were added in the reverse order to *EcoNR*. Both experiments led to the same mass spectra (Fig. 5 and Fig. S5 and S6, ESI[†]). When interpreting ESI mass-spectra it needs to be taken into account that spectra are recorded over time. From the moment the enzyme is treated with $PreQ_0$ and cofactor, the catalytic reaction will proceed. This includes the time needed for sample preparation and the measurements, which represent several scans over the entire measured mass range. Therefore, signals for

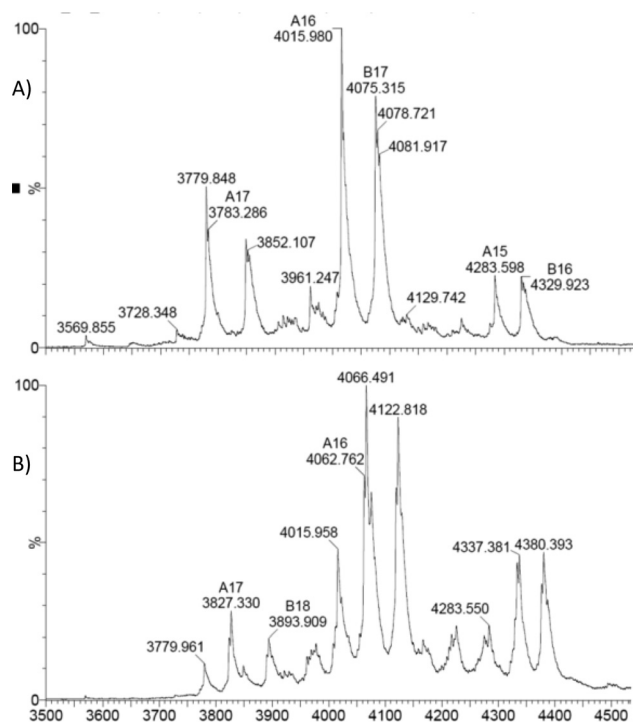


Fig. 4 A) native *EcoNR* and B) *EcoNR* treated with one equivalent of NADPH per monomer ($m/z = 4062$).



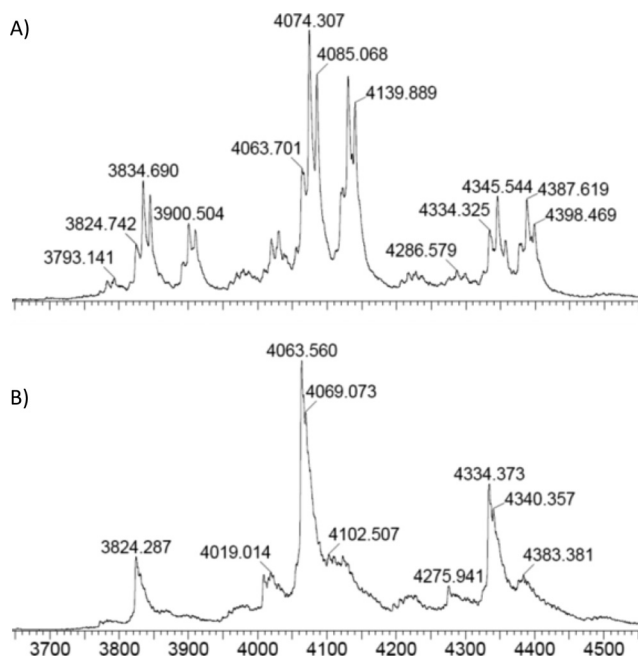


Fig. 5 A) *EcoNR* treated with one equivalent of PreQ₀ and NADPH each per monomer; three main peaks are present: $m/z = 4085$ (dimeric enzyme + 2 PreQ₀ + NADPH); $m/z = 4074$ (dimeric enzyme + PreQ₀ + NADPH). B) The mass peak $m/z = 4063$ is correlated to the mass of the dimeric *EcoNR* + NADPH when an excess of cofactor was used.

various species can be expected to be present in the spectra. In more detail (Fig. 5A), the difference between the molecular weight of the new peaks appearing in the spectra compared to the mass of the free *EcoNR*, led to the identification of the following multiple charge ions: a mass peak correlating to two molecules of PreQ₀ and one molecule of NADPH bound to the dimeric enzyme ($m/z = 4085$, $z = 16^+$). This multiple-charged ion $m/z = 4085$ proves our earlier hypothesis that the cofactor and the two substrate molecules bind, independent of each other to the enzyme. NADPH cannot bind to the PreQ₀ binding pocket, as suggested earlier.⁵ Therefore, they must bind to two different sites and not both to the PreQ₀ binding pocket. The second peak correlates to one substrate and one cofactor molecule bound to the dimer ($m/z = 4074$, $z = 16^+$). Additionally, a small peak correlating to only a NADPH molecule bound to the enzyme ($m/z = 4063$, $z = 16^+$) is visible. It may be assumed that each of the m/z peaks identified represents a step of the catalytic cycle. However, given the time delay in the measurements it is not clear whether both active sites act in concert or not. Initially a substrate molecule adds per active site. A possible intermediate with also two NADPH molecules per dimer is not visible. Although unlikely, since only one NADPH molecule bound per dimer was detected when only NADPH was added (see above, Fig. 4), it cannot be ruled out as it might have already reacted before measurements. The signal corresponding to two PreQ₀ molecules and one molecule of NADPH in the dimer might represent a mixture of two substrate molecules bound and

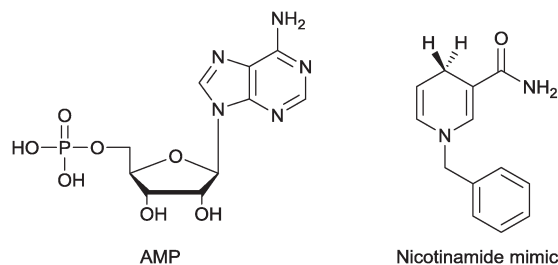


Fig. 6 Cofactor mimics.

one equivalent of NADPH or already represent an intermediate in which one preQ₀ or both are partially reduced. This is followed by the complete reduction of one of the substrates resulting in the peak at $m/z = 4074$. Subsequently complete reduction of the other preQ₀ leaves only one NADPH molecule bound to the dimeric enzyme ($m/z = 4063$, $z = 16^+$).

In the following experiments, an excess of NADPH was added to an aliquot of the reaction mixture used to acquire the mass-spectrometry data reported in Fig. 5A. Complete consumption of PreQ₀ (Fig. 5B) verified that the enzyme remains active under the reaction conditions. Remarkably, even with the excess of NADPH remaining at the end of this experiment only one NADPH molecule is bound to the dimeric enzyme.

The absence of the Rossmann fold in the NRs left the mode of NADPH binding unanswered, in particular since the suggested binding to one active site can now be ruled out. Therefore, even the importance of the adenosine moiety of NADPH is in question.

In order to elucidate whether the cofactor binds *via* the nicotinamide and/or the adenosine portion, *EcoNR* was treated with the AMP and/or a nicotinamide analogue (Fig. 6), followed by ESI-MS. No adduct was observed when the enzyme was treated with AMP using equimolar or higher concentrations per monomer of *EcoNR*. Subsequently, NADPH and substrate were added to the mixture of enzyme and AMP (Fig. S7[†]). After the addition, the same mass-spectra were obtained as in the absence of AMP (Fig. 4A). This demonstrated that the presence of AMP has no influence on the NADPH binding to the enzyme. Thus the binding of the cofactor seems to be independent of the adenosine moiety. Binding to the *EcoNR* *via* the nicotinamide moiety of NADPH was probed *via* nicotinamide cofactor analogues. NR was treated with a nicotinamide mimic (Fig. 6).⁹ Again no change in the mass of *EcoNR* was observed, nor did any reaction occur. Also, no difference was seen when a mixture of AMP and nicotinamide analogue was employed, ruling out a co-operation of both of these moieties in binding.

Conclusion

Here, we provide new evidence for the mechanism of action of nitrile reductases. Mass-spectrometry proved to be a successful technique to gain insight in the binding mode of the cofactor and the natural substrate. Additionally, it allowed



surveying the steps of the catalytic cycle. Indeed, the first step of the catalytic cycle is the covalent binding of the substrate, one to each monomer of the dimeric enzyme. This dimeric enzyme was the only catalytic unit we detected. ESI-MS analysis suggested that only one molecule of NADPH binds to the dimeric *EcoNR*. This is independent of the other two molecules of substrate bound to the two active sites. This would contradict the previously proposed docking of one molecule of substrate and one molecule of NADPH per dimeric enzyme. Furthermore, the binding of NADPH is independent of the adenosine moiety.

Experimental section

Materials and methods

UV-vis measurements were performed on a UV/vis spectrometer UV-2401 PC from Shimadzu with a Julano type F12 thermostat. Liquid chromatography purification was done on a UFLC equipment from Shimadzu. Ni-NTA spin nickel column were provided from Qiagen. Bicinchoninic acid assay (BCA) kit was purchased from ThermoFisher Scientific. All equipment was used according to the provided instructions. The chemicals were purchased from Sigma-Aldrich, if not mentioned otherwise. PreQ₀ was synthesised as described earlier.^{3b,c}

Gene synthesis. The revised mature protein-coding sequence of *EcoNR* (GeneID: 947270), was synthesised by GeneArt (ThermoFisher Scientific) after the optimisation of the codon usage for *E. coli*.^{3d}

***E. coli* expression.** The coding sequence was cloned in the expression vector pET-30a and the resulting plasmid (pET30a-*EcoNR*) was used for expression. *EcoNR* was produced in *E. coli* BL21(DE3). Cells were grown for 3 h at 37 °C in LB, induced with 1 mM isopropyl-β-D-thiogalactopyranoside (IPTG), and grown further for 4 h at 37 °C, purification was performed as described earlier.^{3d}

Kinetic measurements. The activity of the enzyme was measured using a UV-vis spectrophotometer, by monitoring the reduction in time of NADPH at 340 nm. All the measurements were recorded at 37 °C for 120 in duplicate or triplicate. The data were analysed with the software package Origin. For the kinetics measurements stock solutions were prepared. All the solutions were diluted in buffer (100 mM Tris, 100 mM KCl, 1.0 mM DTT, pH = 7.5), if not mentioned otherwise. The final volume is 1.0 mL. 500 μM PreQ₀ stock (1.75 mg per 20 mL, DMSO:buffer, 7:3), which was further diluted in Tris buffer mentioned above, to obtain the desired final concentration. Fresh stock solution of NADPH [100 μM in buffer] and PreQ₀ (final concentration of 500 μM containing 30% DMSO in buffer) were prepared prior the experiments. In order to determine the *K_m* for PreQ₀, the concentration of the cofactor was kept constant at 100 μM, while changing the substrate concentration from 1–75 μM. In order to determine the *K_m* for NADPH, the concentration of PreQ₀ was kept constant at 50 μM, while changing the cofactor concentration from 1–400 μM.

Protein mass spectrometry

Native protein mass spectra were recorded on a Waters quadrupole time-of-flight mass spectrometer (Q-TOF premier, Waters, Manchester, UK) equipped with an 8 kDa quadrupole mass filter. Mass calibration in the range of *m/z* 1000 to 10 000 was done by electrospraying a solution of cesium iodide (2 mg ml⁻¹ in 50% 2-propanol and 50% water). Prior to analysis, proteins were buffer-exchanged to 50 mM ammonium acetate in MilliQ water (pH = 6.8) using ultra centrifugation filters (Amicon® Ultra) with a molecular weight cut-off of 10 kDa. Protein ions were formed in the positive ion mode from a ~2–20 μM solution mode *via* nanoelectrospray using borosilicate spray capillaries (ES383 from ThermoFisher Scientific, Breda, The Netherlands). The applied voltage was optimised for each capillary (range 0.9 to 1.5 kV). The pressure in the first stage of the source region of the MS system was set to 6.0 mbar (standard 1.8 mbar) by reducing the pumping efficiency of the scroll pump (Edwards XDS 35i) for optimised transmission of ions with high *m/z* values. The sample cone voltage was set to 150 V, the extraction cone was set to 5 V and the quadrupole was set as a broad-band mass filter, transmitting ions from *m/z* 2000 to 10 000. The scan rate was set to 3 s with a 0.1 s interscan time. Protein mass spectra were analyzed using the MassLynx 4.1 software package (Waters).

Sample preparation for mass spectrometry experiments.

Final concentration of the pure *EcoNR* was 10 μM. Fresh stock solutions of 500 μM (DMSO:ammonium acetate 50 mM pH = 7.2, 7:3) of PreQ₀ and the other compounds tested were made. They were further diluted in ammonium acetate buffer (50 mM) until the final concentration of 10–40 μM was reached. A stock solution of NADPH [1.0 mM] was freshly made prior to each set of experiments and further diluted until the desired final concentration was reached (10–40 μM). The final concentration of DMSO was kept the same in all the samples. The enzyme was treated with equimolar amount of PreQ₀ or any of the other compounds, gently mixing for 15 min. Prior to analysis, the samples were buffer-exchanged to 50 mM ammonium acetate in MilliQ water (pH = 6.8) using ultra centrifugation filters (Amicon® Ultra) with a molecular weight cut-off of 10 kDa.

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