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Nickel-responsive regulation of two novel *Helicobacter pylori* NikR-targeted genes†

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Nickel is an essential transition metal for the survival of *Helicobacter pylori* in the acidic human stomach. The nickel-responsive transcriptional regulator *HpNikR* is important for maintaining healthy cytosolic nickel concentrations through the regulation of multiple genes, but its complete regulon and role in nickel homeostasis are not well understood. To investigate potential gene targets of *HpNikR*, ChIP sequencing was performed using *H. pylori* grown at neutral pH in nickel-supplemented media and this experiment identified HPG27_866 (*frpB2*) and HPG27_1499 (*ceuE*). These two genes are annotated to encode a putative iron transporter and a nickel-binding, periplasmic component of an ABC transporter, respectively. *In vitro* DNA-binding assays revealed that *HpNikR* binds both gene promoter sequences in a nickel-responsive manner with affinities on the order of $\sim 10^{-7}$ M. The recognition sites of *HpNikR* were identified and loosely correlate with the *HpNikR* pseudo-consensus sequence (TATTATT-N₁₁-AATAATA). Quantitative PCR experiments revealed that HPG27_866 and HPG27_1499 are transcriptionally repressed following growth of *H. pylori* G27 in nickel-supplemented media, and that this response is dependent on *HpNikR*. In contrast, iron supplementation results in activation of HPG27_1499, but no impact on the expression of HPG27_866 was observed. Metal analysis of the $\Delta 866$ strain revealed that HPG27_866 has an impact on nickel accumulation. These studies demonstrate that HPG27_866 and HPG27_1499 are both direct targets of *HpNikR* and that HPG27_866 influences nickel uptake in *H. pylori*.

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Introduction

The transition metal nickel plays an important role in the survival of many organisms because it serves as a catalytic co-factor for a variety of enzymes.^{1–3} A case in point is *H. pylori*, the persistent human pathogen that is estimated to infect 50% of the global population.⁴ This microorganism is the causal agent of chronic gastritis, peptic ulcer disease, and gastric mucosa associated lymphoid tissues (MALT) lymphoma, and is the most important risk factor for gastric cancer.^{5,6} *H. pylori* are able to survive and thrive under the acidic conditions of the stomach lining, where they encounter pH fluctuations that range from 6 to as low as 1.^{7–9} A primary component of *H. pylori* acid defence is the nickel metalloenzyme urease, which generates ammonia and bicarbonate from urea to buffer the cytoplasm and periplasm.^{5,10–12} Nickel is also a component of [NiFe]-hydrogenase, an enzyme needed for efficient colonization because it allows *H. pylori* to use the hydrogen gas produced

by other host flora as an energy source.¹³ Although nickel is a nutritional requirement in this organism and a substantial amount of the metal metabolism of *H. pylori* is devoted to the production of these two obligate nickel enzymes,¹⁴ the metal can be toxic to *H. pylori* if it accumulates inside the cell at high concentrations.¹⁵ Intracellular nickel distribution is tightly controlled in *H. pylori* through the efforts of metalloproteins that function in uptake, efflux, storage, and metalloenzyme assembly.^{1,14} Regulation of these metal homeostasis systems is primarily at the transcriptional level, controlled by a nickel-responsive transcription factor called *HpNikR*.^{16,17}

HpNikR is a pleiotropic regulator, acting as either a repressor or an activator of a variety of genes.^{16,18,19} Ni(II)-*HpNikR* regulates transcription upon binding to the promoters of genes that encode factors involved in nickel import (*nixA*, *feca*, *frpB4*, *exbB*), nickel storage (*hpn*), enzymatic activity (*ureAB*), auto-regulation (*nikR*), as well as genes involved in iron homeostasis (*fur*, *pfr*) and possibly other cellular processes.^{18,20–23} *HpNikR* is also involved in the global response of *H. pylori* to acidic conditions,^{24,25} and there is evidence that the activity of *HpNikR* is directly modulated by changes in pH.²¹ The mechanism by which nickel or protons activate DNA binding by *HpNikR* remains unresolved.^{20,21,24,26,27} Furthermore, a strong consensus DNA-binding sequence for *HpNikR* has not been identified. Several iterations of alignments of the DNA recognition

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0.01% IGEPAL). Following incubation, 1 μL of 0.4 U of an RNase-free DNase I enzyme (Invitrogen) was added to each 20 μL sample and quenched after 4 min at room temperature by the addition of 80 μL DNase stop buffer (20 mM Tris, pH 8, 40 mM EDTA, 1% (w/v) SDS, 0.1 $\mu\text{g } \mu\text{L}^{-1}$ herring sperm DNA). The proteins were extracted from the samples by using a 24:25:1 phenol:chloroform:isoamyl alcohol mixture and the DNA was precipitated with ethanol at -20°C overnight. The samples were then centrifuged, and the DNA was resuspended in 4 μL formamide loading dye and denatured at 90°C for 3 min before loading onto a pre-run 8% denaturing polyacrylamide gel. The samples were then resolved at 1700 V for 2 hours in TBE buffer (89 mM Tris-HCl, pH 7.5, 87 mM borate and 20 mM EDTA). The gel was vacuum-dried and exposed overnight to a phosphor screen, scanned with a Pharos FX Plus Molecular Imager (Bio-Rad) and analysed using Quantity One software. The degree of protection was determined by quantifying the intensity of 2 bands within the recognition sequence relative to a reference band. Half maximal binding was calculated by fitting the data to the Hill equation with a variable Hill coefficient as described for the electrophoretic mobility shift assays.

H. pylori growth conditions and nickel exposure

Wild-type *H. pylori* G27 and its isogenic ΔnikR strain were plated on Columbia agar plates containing 5% horse blood (Thermo Scientific) under a microaerobic environment (5% O_2 , 10% CO_2 and 85% N_2) at 37°C for two days, then replated on new Columbia blood agar plates and grown for a further 22 hours. The colonies were then cultured in 30 mL of Brucella broth with 10% fetal bovine serum (Gibco BRL) and 25 $\mu\text{g mL}^{-1}$ Helicobacter Selective Supplement (Oxoid) while shaking at 100 rpm at 37°C under the microaerobic environment for 10–15 hours up to an OD_{600} of 0.5–0.7. For overnight exposure to nickel, the medium was supplemented with 10 μM NiCl_2 . Samples for qPCR from the overnight exposure were collected by pelleting an aliquot of the cells such that the final number of bacteria per sample was $\sim 4 \times 10^7$. For transient nickel exposure, *H. pylori* grown overnight without added nickel were transferred to 5 mL of fresh Brucella broth at pH 7.0 to a final OD_{600} of 0.2, supplemented with 50 mM urea and with or without 100 μM NiCl_2 . The cells were incubated for 15 minutes at 37°C in a microaerobic environment and then pelleted at 4000 rpm at 4°C for 10 minutes. The supernatant was removed using a vacuum pump and the pellets were either immediately stored at -80°C or lysed for RNA isolation. After the exposure but prior to centrifugation, 10 μL aliquots of the cultures were collected and serially diluted ten-fold in Brucella broth three times. An aliquot of 5 μL from each dilution was spotted on a Columbia blood agar plate and incubated for 3 days to measure cell viability.

Chromatin immunoprecipitation

The *Helicobacter pylori* strain G27 was grown as a liquid culture at pH 7.0 supplemented with 20 μM NiCl_2 as described above. The ChIP experiment was carried out using the One-Day Chromatin Immunoprecipitation Kit (Millipore), with a bacterial sample size of 2×10^8 cells and five thirty-second pulses in a

bath sonicator to lyse the cells and shear the DNA to an average fragment size of 200–500 bp. The remainder of the protocol was as prescribed by the kit. Rabbit polyclonal anti-*HpNikR* antibody (raised against purified *HpNikR*, Division of Comparative Medicine, University of Toronto) was purified from serum by Western blotting using purified *HpNikR* protein as previously described.⁴⁰ The recovered genomic DNA was sequenced by Illumina sequencing at ACTG Corporation (Toronto, ON). The reads for ChIP and control DNA isolated with IgG were 35 284 339 and 39 211 483, respectively. The reads were aligned to the *H. pylori* G27 genomic DNA sequence using the Novoalign short sequence alignment software (novocraft.com). The location information of uniquely mapped reads was converted to BED (Browser Extensible Data) format and used as input for the peak finding software SSSRs (Site Identification from Short Sequence Reads).⁴¹ SSSRs was run for binding site identification with the parameters $D = 0.001, 0.01, 0.1$; $e = 1, 5, 10$; $p = 0.1, 0.01, 0.001$ and $w = 2, 4, 8, 16$. A php script was used to merge the identified sequences. These hits were matched with promoters for various genes in *H. pylori* G27 and used to generate a list of potential regulation targets of *HpNikR*.

H. pylori knockout construction

To generate a knockout of the HPG27_866 gene in *H. pylori* G27, a DNA construct containing the *aphA-3* kanamycin resistance cassette from *Campylobacter coli* flanked by two homologous regions corresponding to the 500 bases upstream and downstream of the HPG27_866 gene was synthesized (BioBasic Canada, Inc.). The construct was transformed into wild-type *H. pylori* G27 via natural transformation, as previously described.⁴² Briefly, a lawn of *H. pylori* was collected from a Columbia blood agar plate grown for two days, resuspended in Brucella broth, spotted onto another blood agar plate and incubated under microaerophilic conditions at 37°C for 4 hours. At that point, 2 μg of the knockout construct DNA was applied to the spot, given time to dry and then further incubated for 12 hours. The spot was then resuspended in Brucella broth and plated onto blood agar plates containing 50 $\mu\text{g mL}^{-1}$ kanamycin. Colonies of transformants were collected after 3 days. Successful incorporation of the resistance cassette was checked by amplifying the genomic DNA of the transformants around the HPG27_866 gene by PCR using the primers sKO866_F and sKO866_R (Table S1, ESI[†]). The upstream and downstream junctions of the recombination region were also amplified using the primers sKO866up_F, sKO866up_R, sKO866dwn_F and sKO866dwn_R (Table S1, ESI[†]) and the products were sequenced to ensure that the genes flanking the *aphA-3* cassette were not disrupted.

RNA isolation and reverse transcription

H. pylori G27 RNA was isolated by resuspending and lysing the bacterial pellet for 5 min in 1 mL of the TRIzol reagent (Invitrogen), according to product protocols. The aqueous layer extracted from the TRIzol was washed twice with chloroform and the RNA precipitated with isopropanol for at least 2 hours at -20°C . The RNA was then pelleted and washed with 70%



An In-112 standard (5 ppb in 35% HNO₃) was run before each set of samples to correct for signal fluctuations in the instrument. The Ni-60 signals were measured from the neat samples and Mg-24 signals from the 1/10 dilutions, because the Mg-24 signal saturated the detector in the neat samples. The raw counts for each isotope were collected as an average of six readings in five sweeps. The Mg-24 signal was used as an internal control to correct the Ni-60 signals for the loss of cells during washing and pipetting.

Quantitative PCR

Identifying and analyzing new *HpNikR* targets

Several novel promoters were also identified as putative targets of *HpNikR* (Table S2, ESI[†]). Two genes identified in this initial experiment are HPG27_1499 and HPG27_866, referred to as 1499 and 866 in this report, which respectively encode CeuE, the periplasmic protein component of a putative transporter of the ABC superfamily^{33,34} and FrpB2, an iron-responsive outer membrane protein.^{31–33} Both of these genes were identified as possible targets of *HpNikR* in a previous microarray study of the SS1 strain of *H. pylori*,⁴⁵ but direct regulation of these genes by *HpNikR* has not been demonstrated. Given this precedence, the possibility of nickel-responsive regulation of these genes by *HpNikR* was explored.

Nickel-activated *HpNikR* directly binds to the target promoters

To validate the results of the ChIP-seq experiment, we examined whether *HpNikR* can bind directly to the two promoter sequences by using electrophoretic mobility shift assays (EMSAs). In the presence of 800 μM NiSO_4 , *HpNikR* binds to the 1499 sequence at half-maximal saturation with a protein concentration of 140 nM (Fig. 1, Table 1). Under the same conditions, *HpNikR* also binds to the 866 promoter with an affinity of approximately 60 nM (Fig. 1, Table 1). In both cases, no visible shift was observed when the experiment was run with 75 μM NiSO_4 (data not shown). DNA binding by holo-*HpNikR* (loaded with 1 : 1 nickel) was also observed when 3 mM MgSO_4 was included in the gel and running buffer (Fig. 2). The calculated



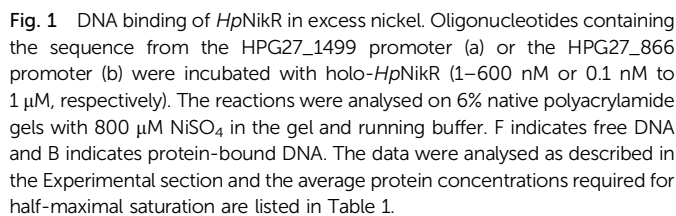
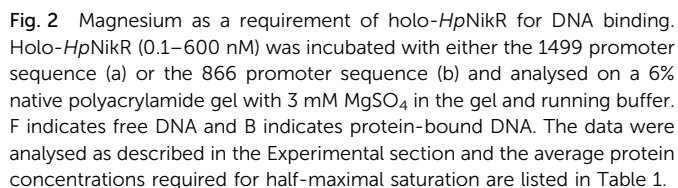


Table 1 Apparent DNA-binding affinities ($\times 10^{-9}$ M)^a

	EMSA		DNase footprint	
	Holo- <i>Hp</i> NikR		Holo- <i>Hp</i> NikR	Excess Ni(n)
	800 μ M NiSO ₄	3 mM MgSO ₄	1 : 1 Ni(n)	35 μ M NiCl ₂
1499	140 $\pm$ 30	130 $\pm$ 80	230 $\pm$ 80	330 $\pm$ 40
Hill coefficient	1.3 $\pm$ 0.3	1.9 $\pm$ 0.8	2.1 $\pm$ 0.3	2.02 $\pm$ 0.04
866	60 $\pm$ 50	22 $\pm$ 5	210 $\pm$ 70	N/D
Hill coefficient	1.4 $\pm$ 0.2	1.9 $\pm$ 0.4	1.7 $\pm$ 0.4	N/D

N/D – Not determined. ^a Values are the average protein concentrations required for half-maximal saturation and standard deviation from at least three experiments.



Ni(II)-*Hp*NikR affinities for the 1499 and 866 sequences that were measured in excess Mg(II) were similar to those measured in excess Ni(II) and revealed that approximately 100 nM protein was required for half-maximal binding on both promoters (Table 1). These results are consistent with previous observations that an excess of divalent cations are required to detect clear DNA binding by nickel-loaded *Hp*NikR.^{26,37,48}

In contrast, apo-*Hp*NikR did not interact with either promoter in the presence of 1 mM EDTA, which was added to the gel

and buffer to ensure that *Hp*NikR was metal-free (data not shown). Mobility shift assays performed with apo-*Hp*NikR in the presence of 3 mM MgSO₄ in the gel and running buffer revealed weak but detectable binding to the 866 and 1499 promoters (Fig. S1, ESI[†]). Apo-*Hp*NikR binding in the presence of excess Mg(II) was only detected with >50 μM protein, a concentration several orders of magnitude higher than that required for DNA binding by holo-*Hp*NikR.

To locate the binding sites of *HpNikR* on the 1499 and 866 promoters, DNase I footprinting assays with holo-*HpNikR* were performed. These assays revealed a 37 bp protected region on the 1499 sequence (Fig. 3) and titration of Ni(II)-*HpNikR* into the reactions resulted in half-maximal DNA binding with 230 nM protein, comparable to the affinity measured by EMSA. Similarly, a 37 bp footprint was detected on the 866 promoter with a calculated affinity of 210 nM (Fig. 4). Both recognition sequences are AT-rich and loosely follow the pseudo-palindromic consensus sequence (TATTATT-N₁₁-AATAATA) proposed by Delany *et al.* (Fig. 5),²⁸ as well as the modified version suggested by Stoof *et al.* (TRWYA-N₁₅-TRWYA).²⁹ Furthermore, the recognition sequence on the 866 promoter matches the region predicted by bioinformatic analysis of the *frpB2* promoter in *H. pylori* 26695, albeit with some minor changes in the flanking and bridging sequences.²⁹ Finally, the same footprint region and binding constant were observed when *HpNikR* was incubated with the 1499 promoter in the presence of excess nickel (Fig. S2, ESI[†]), which suggests that excess nickel is not required for tight binding of the promoter in this footprinting experiment. Altogether, these results demonstrate that the promoters of 866 and 1499 are directly bound by *HpNikR* in a nickel-dependent manner.

In vivo regulation of 1499 and 866 by *HpNikR*

To determine the influence of nickel on the transcription of 1499 and 866 and the role of *HpNikR* in this response, quantitative PCR (qPCR) was performed. Wild-type *H. pylori* G27 and an isogenic *ΔnikR* strain were grown in the absence or presence of nickel. Two different conditions were applied, either an overnight incubation in media supplemented with 10 μM nickel, or a shorter 15 min exposure to higher nickel concentrations (100 μM). As a positive control for our *in vivo* analysis, *ureA* gene transcription was also examined. The *ureA* gene encodes the large subunit of the *H. pylori* urease enzyme, which is an abundant nickel metalloenzyme in *H. pylori* that is crucial for acid resistance and is a critical virulence factor for *H. pylori* infection.^{11,49–51} Upregulation of *ureA* upon nickel exposure has been observed previously, attributed to the activity of *HpNikR*, and recently it was discovered that *ureA* is among the first genes activated by *HpNikR* when *H. pylori* are exposed to increased nickel concentrations.^{22,23,27,52}

In this study, the qPCR data reveal a statistically significant upregulation of *ureA* upon exposure of wild-type *H. pylori* to 10 μ M nickel (Fig. 6), in agreement with previous studies.^{22,27,52} No significant difference in *ureA* transcription was detected in the Δ *nikR* strain upon nickel supplementation, consistent with

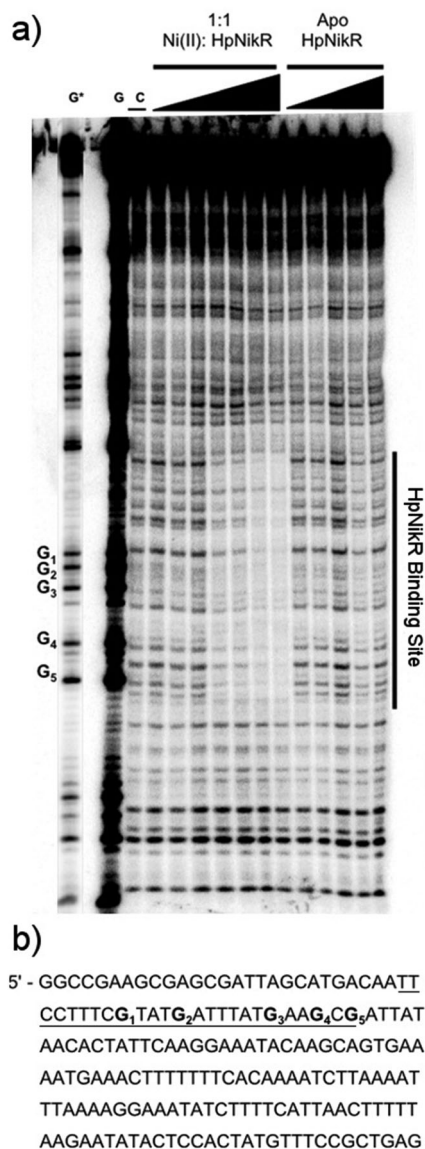


Fig. 3 DNA binding by *HpNikR* on the 1499 promoter sequence. (a) Apo-*HpNikR* (50 nM–5 μ M) and holo-*HpNikR* (50 nM–5 μ M) were incubated with 32 P-radiolabeled 179 bp DNA probe containing the HPG27_1499 promoter sequence before cleavage with DNase I. The reactions were resolved on an 8% denaturing polyacrylamide gel and the area of protection is indicated. Key: G and G*, Maxam–Gilbert G reaction with longer and shorter reaction times, respectively; C, control DNase I reaction in the absence *HpNikR*. The apparent DNA-binding affinity, determined to be 230 ± 80 nM with a Hill coefficient of 2.1 ± 0.3 , is an average from four independent experiments. (b) DNA sequence of the 1499 promoter. The binding region is underlined and the guanines within the footprint are bolded and noted on the footprint gel. Note that the 1499 probe used is the reverse complement of the genomic promoter sequence, thus the start codon is upstream of the indicated binding site.

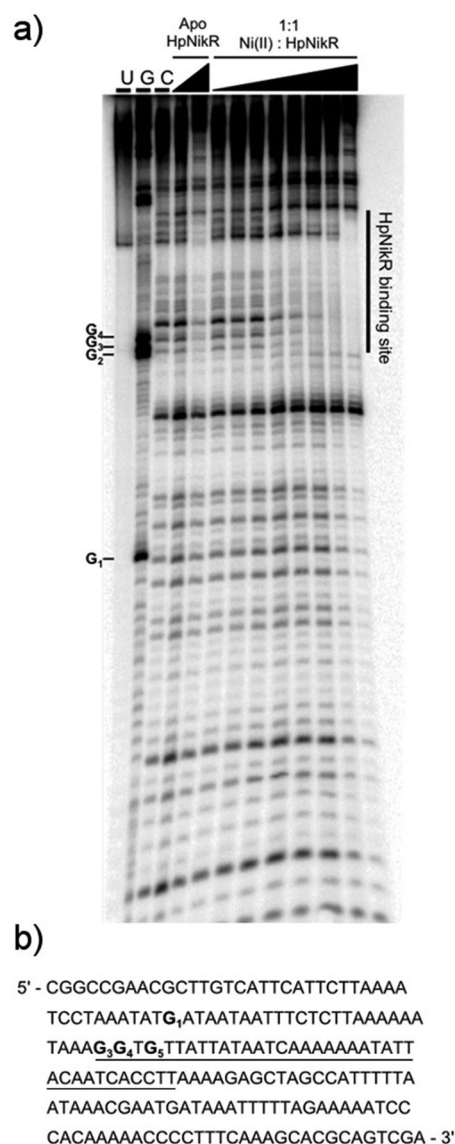


Fig. 4 DNA binding by *HpNikR* on the 866 promoter. (a) Apo-*HpNikR* (100 nM, 10 μ M) and holo-*HpNikR* (100 nM–10 μ M) were incubated with 32 P-radiolabeled 185 bp DNA probe containing the HPG27_866 promoter sequence prior to the addition of DNase I. The reactions were resolved on an 8% denaturing polyacrylamide gel and the area of protection is indicated. Key: U, undigested 190 bp probe; G, Maxam–Gilbert G reaction; C, control DNase I reaction of 185 bp probe with no *HpNikR*. The apparent DNA-binding affinity was determined to be 210 ± 70 nM with a Hill coefficient of 1.7 ± 0.4 , calculated as an average from three independent experiments. (b) DNA sequence of the 866 promoter region. The binding region is underlined and the guanines within the footprint are bolded and noted on the footprint gel. Note that the 1499 probe used is the reverse complement of the genomic promoter sequence, thus the start codon is upstream of the indicated binding site.

the role of *HpNikR* as the primary nickel-responsive regulator in the cell. In contrast to the impact on *ureA*, the transcription levels of 866 and 1499 were significantly repressed when the wild-type bacteria were grown in nickel-supplemented media (Fig. 6), whereas no change in expression was observed for either gene when nickel was added to the Δ *nikR* knockout

strain. These results suggest that 866 and 1499 are directly regulated by nickel-activated *HpNikR*, consistent with the *in vitro* DNA-binding data.

Similar trends were observed when the bacteria were grown with higher levels of nickel but for a shorter time (Fig. S3, ESI†). Nickel supplementation results in activation of *ureA* gene

			1	2	3	4	5	6	7			8	9	10	11	12	13	14												
Consensus 1	-	-	-	T	A	T	W	A	T	N	N	N	N	N	N	N	N	N	A	A	T	W	A	T	-	-	-			
866	-	-	GG	T	G	T	T	A	T	A	T	C	A	A	A	A	A	A	A	A	T	A	T	T	A	C	A	A	T	
1499	T	T	C	G	T	A	T	G	A	T	T	A	T	G	A	A	G	C	G	A	T	T	A	A	C	A	C	T	A	T
ureA	G	A	T	A	T	A	A	C	A	C	T	A	A	T	C	A	T	T	T	T	A	A	T	A	A	T	A	A	T	-
nixA	A	A	T	A	T	T	A	C	A	A	T	T	A	C	C	A	A	A	A	A	A	G	T	A	T	T	A	T	T	T
frpB4	A	A	G	T	A	T	T	A	T	A	A	T	A	G	A	A	T	A	A	T	A	G	T	A	A	T	A	A	T	-
fecA3	C	G	C	A	T	T	A	T	T	A	A	G	T	T	T	T	T	T	T	T	T	G	T	T	T	T	A	T	T	A
Consensus 2	-	-	-	T	R	W	Y	A	N	N	N	N	N	N	N	N	N	N	N	N	T	R	W	Y	A	-	-	-	-	-

Fig. 5 Alignment of the *HpNikR* binding sites from select promoters. The two parts of the consensus sequence are bolded, in red, and numbered according to Evans *et al.* for the aligned sequences.⁶² Consensus 1 and Consensus 2 represent the DNA recognition sites proposed by Delany *et al.*²⁸ and Stoof *et al.*,²⁹ respectively. N represents any nucleotide and W represents either an A or T nucleotide. Clustal Omega was used to align the promoters and generate the image.

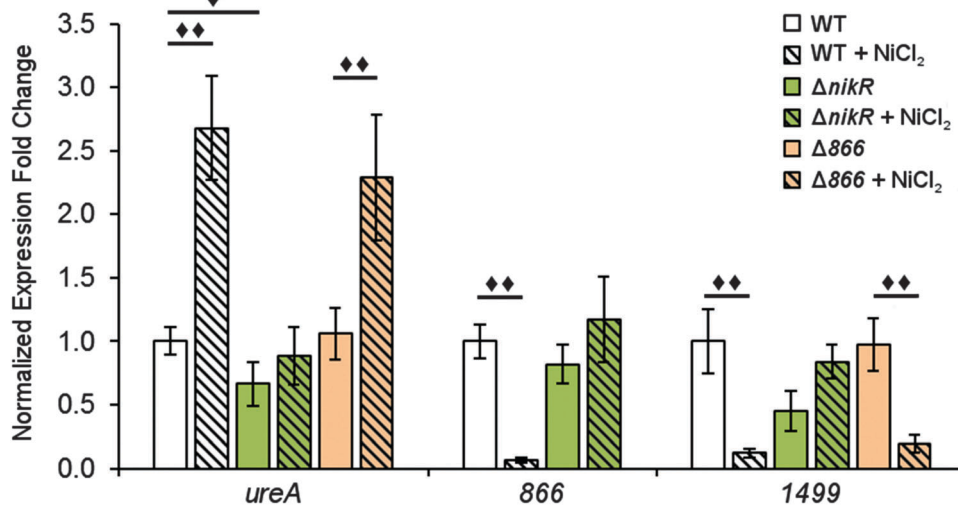


Fig. 6 Expression of *H. pylori* G27 genes after overnight nickel exposure. Quantitative PCR was used to analyse *H. pylori* G27 (wild-type, Δ *nikR* or Δ 866) grown overnight to midlog phase in the presence or absence of a 10 μ M NiCl₂ media supplement. Expression of the *ureA*, 866 and 1499 genes were normalized to 16S and *gyr* expression levels and relative to wild-type expression of the gene in the absence of supplemental nickel. Statistically significant differences are represented by diamonds ($\diamond = P < 0.05$, $\blacklozenge = P < 0.01$). All data represent the average values from at least three biological replicates with three technical replicates each and error bars represent one standard error of the mean.

transcription and repression of the 866 and 1499 genes in the wild-type strain. A small but significant reduction in *ureA* expression was observed in Δ *nikR* strain under these growth conditions, and while this latter observation could suggest a nickel-responsive component in *ureA* regulation that is not directly controlled by *HpNikR*, as previously suggested,²² the exact physiological basis behind this effect is not known. In addition, under the short exposure conditions, 1499 transcription was upregulated in the Δ *nikR* strain *versus* the wild-type strain when grown in unsupplemented media. One possible explanation is that apo-*HpNikR* is regulating 1499 *in vivo*, but this model is not consistent with the extremely weak DNA binding observed *in vitro* with apo-*HpNikR* on the 1499 promoter. Instead, it is more likely that trace levels of holo-*HpNikR* cause some repression, which is alleviated in the knock-out strain, or that the 50 mM urea added to the cultures for the short exposures affected relative transcription levels, as urea was previously observed to affect global transcription levels in *H. pylori*.⁵³

There is evidence for overlap between the *HpNikR* and *HpFur* regulons.^{18,28} Furthermore, both 866 and 1499 are regulated in an iron-responsive fashion in some strains,^{32,33,35}

but not in others,^{54–56} so iron-mediated regulation was also investigated in this study (Fig. 7). No significant change was observed in *ureA* transcription when either the wild-type or Δ *nikR* G27 bacteria were exposed to iron-supplemented media, in agreement with previous reports.^{52,55} Similarly, the expression of 866 was unaffected by iron. In contrast, the expression of 1499 was observed to increase significantly in wild-type *H. pylori* when the bacteria were exposed to FeCl₃. This effect was not observed with the Δ *nikR* mutant, suggesting that the iron-dependent component of 1499 regulation may be mediated through *HpNikR*.

Metal uptake through HPG27_866

Given the nickel-responsive regulation of 866, we examined the effects of an 866 knockout on gene transcription and nickel uptake by *H. pylori* G27. The Δ 866 strain did not exhibit any significant changes in the growth rate *versus* the wild type, with or without nickel supplementation of the media (data not shown), suggesting that 866 is not critical for *H. pylori* growth under standard laboratory conditions. Furthermore, the baseline and nickel-dependent regulation of *ureA* and 1499



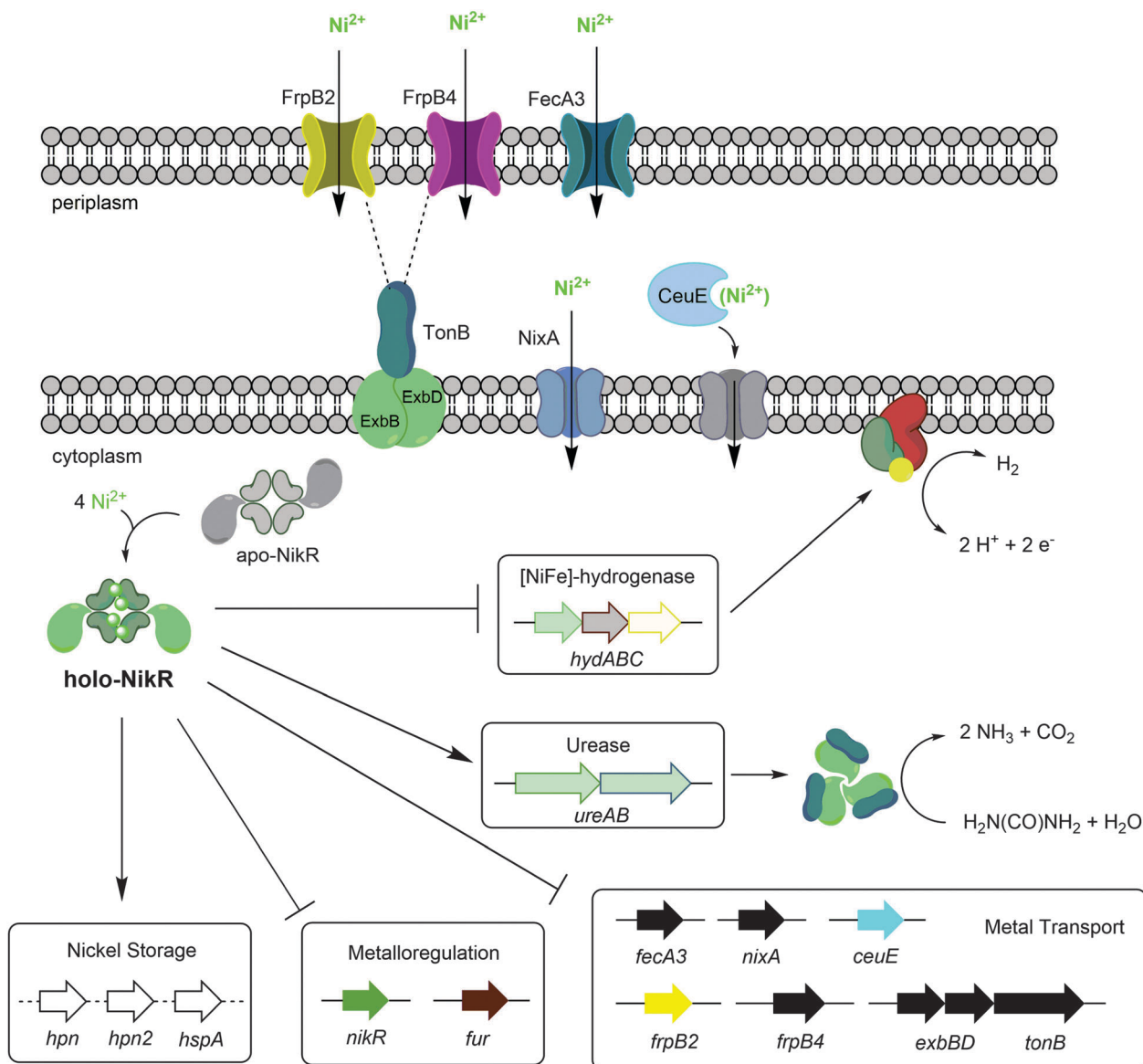


Fig. 9 Regulation by Ni(II)-HpnNikR *in vivo*. The transmembrane components of the CeuE nickel transporter have not been explicitly defined, although there is some evidence that FecD/FecE fill these roles. The TonB–ExbB–ExbD complex is believed to provide energy to the FrpB2 outer membrane protein,³³ and may also power the FrpB4 homolog as well. Note that although urease is depicted as a cytoplasmic protein, it is present in the cytoplasm, outer membrane and is also excreted outside the cell.⁷² The exact nature of the nickel transported is not clear for all systems, and only genes regulated by HpnNikR in response to nickel are shown.

that direct regulation of these genes by HpnNikR had not been demonstrated and that the ChIP-seq strategy tends to produce false positives through non-specific binding,⁶¹ we examined whether HpnNikR regulates these two genes in response to nickel. *In vitro* DNA-binding assays were used to demonstrate nickel-responsive HpnNikR binding to both promoter sequences, the recognition sequences were mapped out, and *in vivo* transcriptional analysis indicated that HpnNikR represses transcription of both genes in response to an increase in the levels of environmental nickel. Altogether, these results suggest that both 866 and 1499 are components of the HpnNikR regulon.

Analysis of the Ni(II)-HpnNikR–DNA complexes by using both EMSAs and DNase footprinting assays revealed half-maximal

saturation with ~100 nM protein on the promoter sequences of both HPG27_866 and HPG27_1499, a value comparable to the affinities of HpnNikR for other DNA recognition sites.^{20,21,26,37,48} Some HpnNikR–DNA complexes are significantly weaker, and it was proposed that the promoters regulated by HpnNikR can be divided into two classes, with the tighter complexes being observed for the genes that encode proteins that directly utilize nickel (K_d values of low-to-mid nM) and weaker complexes on the promoters of genes of other types of proteins (K_d values of low μ M and weaker).²⁰ The affinity of HpnNikR for 866 and 1499 promoter sequences places these genes in the former class, along with the genes encoding other nickel membrane transporters as well as urease. However, it is unclear how this



two-tiered mode of recognition observed by the *in vitro* studies translates into the physiological activity of *HpNikR*.²²

DNase footprinting assays revealed that the sequences of the *HpNikR* binding sites on the 866 and 1499 promoters are 37 bp, AT-rich sequences (Fig. 5). Although neither is an exact match for the initially proposed consensus sequence TATTATT-N₁₁-AATTATA,²⁸ some resemblance is clear, and the 866 site is in full agreement with the more inclusive and recently derived version TRWYA-N₁₅-TRWYA.²⁹ This newer version of the *HpNikR* consensus was generated by analysing known *NikR* binding sites from several *Helicobacter* species. It was then applied to predict *NikR* targets in *Helicobacter* genomes, and one of the operators identified was that of the 866 analog in *H. pylori* 26695, *frpB2* (HP0916) (Fig. S4, ESI†).²⁹ It has also been suggested that several specific DNA bases in the *HpNikR* recognition sequences are more crucial for protein binding than others, with thymine 10 at the 3' half-site serving a role in distinguishing the tight binders from the weak ones.⁶² In support of this proposal, both the 1499 and 866 recognition sequences have thymine at position 10, consistent with the relatively tighter affinity of these *HpNikR*-DNA complexes.

In the *H. pylori* strain 26695, the gene orthologous to 866 is split by a stop codon caused by a frame-shift deletion, forming *frpB2* and the pseudogene *frpB3* (HP0916 and HP0915, respectively), which are repressed by the iron metalloregulator *HpFur* in an iron-dependent manner.³² However, iron-responsive regulation of this gene is not always observed. Iron-dependent changes in 866 expression were not observed in the G27 wild-type or Δ *nikR* strains under the conditions used in this study, a result consistent with previous experiments with *H. pylori* G27⁵⁴ as well as with array-based studies of other strains.^{35,55} Furthermore, neither of the promoters of HPG27_866 and HP0916/HP0915 contain the putative *HpFur* recognition sequence and evidence that *HpFur* binds directly to these promoters is lacking.^{54,63} In contrast, the promoters in both strains contain a *HpNikR* binding site (as observed in this work and previously²⁹), albeit with slight variations in the bridging and flanking sequences of each (Fig. S4, ESI†), and the *in vitro* and *in vivo* experiments reported here demonstrate that regulation of this gene is directly controlled by *HpNikR* in response to nickel. Although iron-responsive regulation of 866 is not consistently observed, it is possible that *frpB2* expression is modulated indirectly by iron under some conditions through *HpNikR* activity. It has been previously suggested that the *HpNikR* and *HpFur* regulons overlap;^{18,28} for example both proteins have been demonstrated to directly regulate common genes such as the *hydABC* cassette^{22,63} and *HpNikR* directly represses the transcription of *HpFur*.^{20,22}

Deletion of HPG27_866 negatively affects nickel accumulation in *H. pylori* grown under standard laboratory conditions, suggesting that this outer membrane transporter may be involved in nickel uptake. No significant impact was observed on the expression of the *ureA* or 1499 genes in the Δ 866 mutant strain, indicating that *HpNikR* activity is not affected. The lack of impact on regulation despite lower nickel levels could arise from partitioned pools of nickel within *H. pylori*, as proposed

for *E. coli*.⁶⁴ It has also been suggested that *FrpB2* contributes to iron uptake as a haemoglobin receptor. Overexpression of the *FrpB2* protein encoded by *H. pylori* J99 allows *E. coli* to survive in an iron-deficient medium that was supplemented with Hb, and analysis of the crude *E. coli* lysates suggested that *FrpB2* could bind human haemoglobin.³¹ A connection between nickel and heme uptake was also made with the observation that *E. coli* *NikA*, the periplasmic component of the *NikABC* transporter, can bind heme,⁶⁵ although how this activity contributes to heme utilization is not clear.⁶⁶ Future work on the biochemistry and cell biology of *FrpB2* will be necessary to pinpoint the specific role that it plays in metal uptake in *H. pylori*.

HPG27_1499 was also originally annotated as an iron transport protein, possibly due to the homology with the *Campylobacter coli* periplasmic iron-binding protein.^{67,68} Despite the weak binding affinity of *HpFur* to the 1499 binding site,⁵⁶ iron-responsive regulation of the corresponding gene in *H. pylori* SS1 was observed only in the stationary growth phase and not in other strains.^{32,35,55,56} Furthermore, mutation of the genes encoding the putative transmembrane components of this ABC transporter, *fecD* and *fecE*, did not dramatically impact the iron uptake in *H. pylori* 11637,⁶⁷ suggesting that this transporter is involved in the uptake of a different metal. In this study, weak but significant upregulation of HPG27_1499 was observed upon a brief exposure to high concentrations of iron in the wild-type bacteria ($P = 0.049$) but not in the Δ *nikR* mutant strain. As discussed above for HPG27_866, it is possible that iron-responsive regulation of HPG27_1499 is a result of cross-regulation between *HpFur* and *HpNikR*. As previously suggested, the sigma factor *RpoD* (σ^{80}) could also be affecting the transcription of 1499 through an unknown indirect iron-dependent pathway.⁶⁹ While not yet validated, the *rpoD* gene was identified as a putative target of *HpNikR* by our ChIP-seq results and contains the *HpNikR* consensus binding sequence in its promoter, but further experiments would be required to test this hypothesis.

The assignment of HPG27_1499 as a component of the *HpNikR* regulon is consistent with the recent re-classification of this protein as a factor involved in nickel acquisition. Interruption of the corresponding *ceuE* or *fecD* genes in *H. mustelae* resulted in decreased total nickel and cobalt.³³ Furthermore, recent structural analysis of *H. pylori* *CeuE* revealed that it matches the fold of a periplasmic-binding-protein component of an ABC metal transporter and that it binds Ni(L-His)₂, similar to *E. coli* *NikA*,⁷⁰ but not heme, vitamin B12 or enterobactin.³⁴ These observations suggest a model for a biological connection between *HpNikR* and *CeuE* that is analogous to *E. coli* *NikR* and the *NikABCDE* inner membrane nickel transporter, where *HpNikR* represses transcription of *CeuE* to halt nickel import in response to high concentrations of extracellular nickel.^{19,71} The genes encoding *FecD* and *FecE*, HPG27_842 and HPG27_841, respectively, are in a separate operon from 1499 and seemingly lack an *HpNikR* recognition sequence in the promoter region. However, the *fecE* gene was also identified as an *HpNikR* target in our ChIP-Seq data



(Table S2, ESI[†]), suggesting that the transmembrane and periplasmic components of the CeuE/FecD/FecE transporter could be regulated by H_pNikR in concert.

Conclusions

Ni(II)-H_pNikR binds to the promoters of HPG27_1499 (*ceuE*) and HPG27_866 (*frpB2*) *in vitro* with strong, sub-micromolar binding affinities, and DNase I footprinting revealed binding sites within both promoters that resemble the established pseudo consensus sequences. Furthermore, the transcription of both genes is significantly downregulated when *H. pylori* is exposed to media supplemented with NiCl₂, a response that is dependent on the presence of H_pNikR, linking the *in vivo* transcription of 866 and 1499 to the *H. pylori* nickel response. These results allow these genes to be assigned to the H_pNikR regulon, and the impact of the 866 deletion on nickel accumulation is consistent with a role of the gene product in nickel transport. Future work will establish the mechanism of how these proteins contribute to metal homeostasis in *H. pylori*. Finally, these experiments provide a snapshot of the H_pNikR regulon under a single set of growth conditions, but it is clear that multiple variables impact the activity of this protein, including time and pH.^{21,22,24,25} Analysis of bacteria exposed to different conditions will provide details of how H_pNikR supports *H. pylori* as it adapts to the challenges of its niche environment.

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