

Correction to “EPR Spectroscopic Studies of the Fe–S Clusters in the O₂-Tolerant [NiFe]-Hydrogenase Hyd-1 from *Escherichia coli* and Characterization of the Unique [4Fe–3S] Cluster by HYSORE”

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Page 15585. In Table 1, two sets of published data obtained by other researchers were inadvertently placed in the wrong columns.

Table 1, including a new footnote *c*, should read as follows:

Table 1. Midpoint Potentials of the EPR-Active Fe–S Clusters Observed in Native Hyd-1, the P242C, and C19G/C120G Variants Compared to Other Native O₂-Tolerant Hydrogenases^a

enzyme	[4Fe–3S] ^{5+/4+} proximal	[4Fe–3S] ^{4+/3+} proximal	[3Fe–4S] ^{+/0} medial	[3Fe–4S] ^{+/0} _{app.} medial	[4Fe–4S] ^{2+/+} distal
native Hyd-1	230 ± 15	30 ± 30	190 ± 30	130 ± 15	–
P242C	175 ± 15 ^b	90 ± 20	–	–	–
C19G/C120G	–	–	215 ± 10	–	–
<i>Aa</i> Hase I ¹⁴	232 ± 20	98 ± 20	–	78 ± 20	–65 ± 20
<i>Re</i> -MBH ^{23,c}	160	–60 ^c	–	25	–180 ^c
<i>Rm</i> CH34 ^{23,c}	240	50 ^c	–	100	–80 ^c

^aThe midpoint potentials are given in mV vs SHE, were obtained as detailed in Methods section, and reflect the ‘Nernst plots’ given in Figure 2B. The potentials for *Aa* Hase I were obtained at pH 6.4 vs the normal hydrogen electrode,¹⁴ and those for *Re*-MBH and *R. metallidurans* CH34 were obtained at pH 7.0.²³ All potentials for the Hyd-1 enzymes were obtained at pH 6.0. The apparent midpoint potential (‘app’) refers to the potential at which the uncoupled [3Fe–4S]⁺ cluster signal is at half its maximum intensity (Figure S6A). ^bMonitoring peak intensities at different field positions resulted in a spread of reduction potentials of ca. 55 mV (Figure S6B). ^cIn ref 23, the higher midpoint potential (–60 mV and 50 mV) was assigned to cluster I, and the lower potential (–180 mV and –80 mV) was assigned to cluster II; it is assumed here that the lower potential belongs to the distal cluster.