An Ancient Pathway Combining Carbon Dioxide Fixation with the Generation and Utilization of a Sodium Ion Gradient for ATP Synthesis

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Abstract

Synthesis of acetate from carbon dioxide and molecular hydrogen is considered to be the first carbon assimilation pathway on earth. It combines carbon dioxide fixation into acetyl-CoA with the production of ATP via an energized cell membrane. How the pathway is coupled with the net synthesis of ATP has been an enigma. The anaerobic, aceticogenic bacterium *Acetobacterium woodii* uses an ancient version of this pathway without cytochromes and quinones. It generates a sodium ion potential across the cell membrane by the sodium-motive ferredoxin:NAD oxidoreductase (Rnf). The genome sequence of *A. woodii* solves the enigma: it uncovers Rnf as the only ion-motive enzyme coupled to the pathway and unravels a metabolism designed to produce reduced ferredoxin and overcome energetic barriers by virtue of electron-bifurcating, soluble enzymes.

Introduction

The atmosphere in the early days of our planet was highly reducing and did not contain oxygen but gases such as molecular hydrogen and carbon dioxide. How life evolved under these conditions is a matter of debate but some researchers favour a pathway that combines two essential features: carbon fixation into organic molecules and, at the same time, generation of ATP [1]. The Wood-Ljungdahl pathway of carbon dioxide fixation is such a pathway and, therefore, could be considered to have evolved on earth very early [2,3]. It is present in strictly anaerobic sulfate reducing bacteria and archaea, in methanogenic archaea and acetogenic bacteria [4]. Only in the latter it fulfills the dual function mentioned which is growth on molecular hydrogen and carbon dioxide according to:

\[
4 \text{H}_2 + 2 \text{CO}_2 \rightarrow \text{CH}_3\text{COO}^- + \text{H}^+ + 2 \text{H}_2\text{O}
\]

\[\Delta G^o = -95 \text{kJ/mol}\]  

The pathway involves reduction of carbon dioxide to formate, binding of formate to the cofactor tetrahydrofolate (THF) and subsequent reduction of the formyl group to methyl tetrahydrofolate. Another mol of carbon dioxide is reduced to enzyme-trapped carbon monoxide that is then condensed with a methyl group and coenzyme A to yield acetyl-CoA (Fig. 1). In the anabolic route, acetyl-CoA is the precursor molecule for biosynthetic reactions and in the catabolic route the precursor of acetate [5,6,7]. Acetyl-CoA allows for the synthesis of one mol ATP via substrate level phosphorylation. However, this ATP is required for formate activation to formyltetrahydrofolate leaving no ATP. Therefore, acetogenesis from 4 H2 + 2 CO2 must yield additional ATP by a mechanism other than substrate level phosphorylation. How this is achieved is unknown. The amount of energy gained from equation 1 would theoretically allow the generation of only 1.5 mol of ATP per mol acetate produced [8]. This calculation is based on a partial pressure of molecular hydrogen of 1 bar. However, at the low hydrogen partial pressures observed in the environment in which the acetogenic bacteria thrive the free energy change associated with reaction 1 is much more positive and sufficient to generate only a fraction of an ATP [9]. Therefore, acetogens clearly live at the thermodynamic limit of life.

Acetogens can be divided in two groups with respect to their energy metabolism [10]. One group contains cytochromes and quinones that are considered to be involved in a membrane-bound electron transport process that generates a proton potential across the membrane; they use a proton potential-driven ATP synthase. This can be considered as the evolutionary more advanced version of the pathway since it required the evolution of cytochromes [11].
The second group, to which the Gram positive bacterium Acetobacterium woodii belongs, represents a more ancient version of the pathway. It does not involve cytochromes and quinones but at least one ion pump, a sodium ion-translocating ferredoxin:NAD$^+$ oxidoreductase (Rnf) [12]. The Rnf complex is a membrane-bound electron transfer complex containing iron-sulfur centers and flavins. Exergonic electron transfer from reduced ferredoxin (Fd$_2^-$) to NAD$^+$ is found to be coupled with vectorial, electrogenic Na$^+$ transport in inverted membrane vesicles of A. woodii [13].

Immunological studies revealed that the Rnf complex is present in cells grown on H$_2$+CO$_2$, underlining its importance in the bioenergetics of the Wood-Ljungdahl pathway [14]. The complex is also present in cells grown on formate, methanol, betaine or fructose and its synthesis was also independent of the electron acceptor used and, therefore, is constitutive [14]. The sodium ion potential across the membrane drives the synthesis of ATP via a membrane-integral Na$^+$F$_1$F$_0$ ATP synthase [15,16].

Another argument that this version of the pathway has evolved early is that it uses Na$^+$ as coupling ion. Membranes are more tight for Na$^+$ than for H$^+$ [17] which is essential for organisms living in ecosystems that have a high concentration of organic acids that function as “proton ferries” allowing protons to backenter the cell and thus destroying part of the already very little power available to fuel the ATP synthase. Therefore, sodium bioenergetics using a DpNa is considered an early step in the evolution of cellular bioenergetics [18].

A. woodii employs the Wood-Ljungdahl pathway without cytochromes and is one of the very few organisms known to completely rely on a sodium ion potential for energetic reactions [10]. It grows well in the absence of a proton potential across its membrane, and apparently uses a primary Na$^+$ potential to fuel ATP synthesis, transport processes and flagellar rotation [16]. Thus it is a prime candidate to unravel the bioenergetics of what may have been one of the first pathways in coupling exergonic and endergonic reactions in life. Therefore, we sequenced the entire genome of A. woodii and performed a few key experiments to fill the gaps in understanding how the pathway is coupled to the synthesis of ATP.

Results and Discussion

Genetic organization and predicted properties of enzymes of the Wood-Ljungdahl pathway

General genome features of A. woodii are listed in Table 1. The genome of A. woodii consists of a single circular 4,044,785 bp chromosome with a GC content of 39.3%. 3473 protein-encoding ORFs, 5 rRNA clusters and 60 tRNA genes account for 85.1% of the genomic DNA. 71.1% of the predicted ORFs are with assigned functions (NCBI database, accession number CP002987). The genes encoding proteins of the Wood-Ljungdahl pathway are mainly found in three clusters on the chromosome (Fig. 1).

In cluster I, two genes for formate dehydrogenase (fdhF) are present. The formate dehydrogenase catalyzes the first
The Rnf complex is the only membrane-bound electron transfer system of the Wood-Ljungdahl pathway

Inspection of the genome sequence ruled out that any of the orfs encoding for enzymes involved in carbon flow in the Wood-Ljungdahl pathway are membrane-bound and potential Na⁺ pumps. This was a surprise since it was speculated for decades that the methyl-THF:corrinoid-iron sulfur protein methyltransferase is a membrane integral, Na⁺-translocating enzyme [10]. This assumption was based on the finding of membrane-bound corrinoids in A. woodii [20] and the finding of a corrinoid-containing, Na⁺-translocating methyltransferase in methanogens [21]. A. woodii encodes 30 different methyltransferase systems involved in channeling methyl groups from various substrates into the central pathway, but none is predicted to be membrane-bound. The corrinoid-containing proteins found in the membrane fraction of A. woodii [20] may thus stem from “sticky” cytoplasmic methyltransferases.

Thus, the genome sequence revealed that Rnf is probably the only ion-pumping enzyme active during litho-autotrophic growth of A. woodii. The use of Rnf as the only coupling site in the pathway leads to important consequences. The $E_0^\circ$ of the $\text{Fd}_{\text{red}}/\text{Fd}⁺$ couple is probably $-500 \text{ mV}$ and that of the $\text{NAD}⁺/\text{NADH}$ couple is $-320 \text{ mV}$ (the $E_0^\circ$ of the $\text{Fd}_{\text{red}}/\text{Fd}⁺$ couple is assumed to be near $-500 \text{ mV}$ because in A. woodii reduced ferredoxin is used as electron donor for the reduction of CO₂ to CO, the $E_0^\circ$ of the $\text{CO}_2/\text{CO}$ couple being $-520 \text{ mV}$ [20]). The higher the $\text{Fd}⁺/\text{NAD}⁺$ concentration ratio, the higher the energy that can be used for building up a sodium ion gradient. Therefore, the entire catabolic metabolism has to be optimized to produce a maximal $\text{Fd}⁺/\text{NAD}⁺$ ratio. A maximal number of oxidation steps should produce reduced ferredoxin whereas a maximal number of reduction steps should use NADH as electron donor. This leads to a metabolism focused on ferredoxin.

Overcoming a first energy barrier in the electron pathway

In the first step of the electron transfer pathway, ferredoxin is reduced with $\text{H}_2$ as reductant. This reaction is highly endergonic with a $\Delta G^\circ = +18.5 \text{ kJ/mol}$ $[\text{Fd}⁺/\text{Fd}_{\text{red}} = -500 \text{ mV}; E_0^\circ (\text{H}^+/\text{H}_2) = -414 \text{ mV}]$ and thus requires the input of energy. One possibility to overcome this step energy barrier is to drive the endergonic electron transfer from molecular hydrogen to ferredoxin by the electrochemical ion potential across the membrane through ion influx into the cell. Such a kind of reverse electron flow is catalyzed by Ech-type or related hydrogenases [22] but genes encoding such enzymes are absent from the chromosome. Moreover, the genome of A. woodii does not harbour genes encoding [NiFe]-hydrogenases but a putative operon consisting of 5 genes (hydAI-hydE, Awo_c26970-Awo_c27010, Fig. 2) that potentially encode a cytoplasmic multimeric [FeFe]-hydrogenase. hydAI encodes a putative catalytic hydrogenase subunit with a complete H-cluster, one [2Fe-2S] and three [4Fe-4S] clusters, based on sequence similarities. hydB codes for a putative iron sulfur protein with one [2Fe-2S], three [4Fe-4S] and a flavin binding site. hydC is the first gene of the cluster and codes for a protein with one predicted [2Fe-2S] binding site. The protein encoded by hydD has apparently no cofactor binding sites. hydE has no similarity to hydrogenase subunits but is similar to ATP–binding domains of histidine kinases. Proteomic analysis of cells grown with H₂ shows that the H₂-reducing proteins do not change in concentration in the presence of $\text{H}_2$ but none is predicted to be membrane-bound. The corrinoid-containing proteins found in the membrane fraction of A. woodii may thus stem from “sticky” cytoplasmic methyltransferases.

Table 1. General features of the A. woodii genome.

| Feature                        | Value |
|--------------------------------|-------|
| Genome size (Mbp)              | 4.044785 |
| protein encoding orfs          | 3472  |
| percent coding (%)             | 85.1  |
| G+C content (mol%)             | 39.34 |
| rRNA cluster                   | 5     |
| tRNA                            | 59    |
| orfs with assigned functions    | 2469  |
| orfs without assigned functions | 1004  |

doi:10.1371/journal.pone.0033439.t001

...step in the pathway, the conversion of CO₂ to formate. FdhF1 and FdhF2 are 79% identical and FdhF2 seems to be a selenocystein-containing protein. Apparently, it is the only selenocystein-containing protein encoded on the genome of A. woodii. The genes encoding selenocystein biosynthesis proteins are next to the cluster upstream of fdfF1. Furthermore, there are genes present that potentially encode a formate dehydrogenase accessory protein (fdhD), three copies of a putative FeS-containing electron transfer protein (hydB) and a subunit (hydA2) harbouring the active site characteristic of [FeFe]-hydrogenase. HycB1 and HycB2 are 82% identical and the encoding genes are downstream of fdfF1 and fdfF2, respectively. Therefore, we suggest that HycB1 and HycB2 are electron transfer proteins specific for FdhF1 and FdhF2, respectively. HycB3 is only 34 and 33% identical to HycB1 and HycB2, respectively, and its localization in the gene cluster indicates that it is the electron output module for the hydrogenase HydA2.

Cluster II encodes the enzymes catalyzing the reduction of the formyl-group to the methyl-group, the formyl-tetrahydrofolate (THF) synthetase, methenyl-THF cyclohydrolase, methylene-THF dehydrogenase and methylene-THF reductase. The genetic organization is different from Moorella thermoacetica where these genes are spread all over the genome [19]. A second formyl-THF synthetase is found in A. woodii (Awo_c08040), whereas in M. thermoacetica and Clostridium ljungdahlii the genes for formyl-thymine nucleosidase, Geobacter sulfurreducens or Methanococcus acetivorans. In contrast, homologues of Awo_c09290 are only found in this region in Ruminococcus obeum, Blautia hydrogenotropha and Bryantella formategens, and three members of the genus Clostridium. This possible indication for the presence of these proteins is discussed below.

Acetyl-CoA is formed from methyl-THF and a second CO₂ by the CO dehydrogenase/acetoyl-CoA synthase complex. Genes coding for the subunits (acsAB) are found in cluster III (Awo_c10670-Awo_c10760) together with the genes coding for the corrinoid-iron sulfinic acid (acsCD) and the methyltransferase (acsE). Additionally there are two genes in the cluster encoding for a CODH nickel-insertion accessory protein (ccoC) and a corrinoid activation/regeneration protein (acsF) (Fig. 1). The same genes are also clustered in M. thermoacetica, Acellulibacter arbutanicum and C. ljungdahlii. The genes for the phosphotransacetylase (Awo_c190620) and the acetate kinase (Awo_c21260) are located in other regions of the genome.

The Rnf complex is the only membrane-bound electron transfer system of the Wood-Ljungdahl pathway...
The first step in CO₂ reduction is an endergonic reaction assuming that electrons are passed from molecular hydrogen to CO₂ via NADH+H⁺ as a reductant (the standard redox potential of the CO₂/formate couple is −430 mV). Electrons may be delivered via NADPH generated from NADH via the CO₂/formate couple is thermodynamically highly unfavourable and Schut and Adams [23] proposed an flavin-based electron bifurcating mechanism to overcome the energetic barrier. Flavin-based electron bifurcation was first shown in anaerobic clostridia as a new coupling mechanism in bioenergetics: the energy liberated during energetic “downhill” transfer of one electron from a donor to an acceptor is used to transfer another electron “uphill” to an acceptor with a more negative redox potential. That was first experimentally proven for the exergonic NADH+H⁺-driven reduction of crotonyl-CoA that is coupled with endergonic ferredoxin reduction [25,26]. For the hydrogen-evolving hydrogenase of T. maritima, it is suggested that the energetic “downhill” transfer of two times one electron from ferredoxin (E₀' approx. −500 mV) to two protons (2 H⁺/H₂; E₀' = −414 mV) drives the energetic “uphill” transport of two times one electron from NADH+H⁺ (E₀' = −320 mV) to two protons. In sum, the reaction is:

\[
\text{Fd}^{2-} + \text{NADH} + \text{H}^+ \rightarrow 2 \text{H}_2 + \text{Fd}_{\text{ox}} + \text{NAD}^+ \quad (2)
\]

Due to the similarity of the enzymes from A. woodii and T. maritima we suggest that the [FeFe]-hydrogenase of A. woodii catalyzes a reversal of equation 2, namely the reduction of one ferredoxin and one NAD⁺ with two H₂ (Fig. 2). The finding that cell free extracts of A. woodii grown on fructose catalyze hydrogen-dependent NAD⁺ reduction only after the addition of FMN and ferredoxin purified from Clostridium pasteurianum is consistent with this hypothesis.

Overcoming a first energy barrier in the carbon pathway

The first step in CO₂ reduction is an endergonic reaction assuming that electrons are passed from molecular hydrogen to CO₂ via NADH+H⁺ as a reductant (the standard redox potential of the CO₂/formate couple is −430 mV). Electrons may be delivered via NADPH generated from NADH+H⁺ in an energy-dependent reaction driven by either a membrane-integral transhydrogenase [27] or an electron bifurcating soluble transhydrogenase [28]. However, homologues of these enzymes are apparently not encoded in A. woodii. But how are the electrons transferred from molecular hydrogen to carbon dioxide? As mentioned above, the formate dehydrogenase gene cluster harbors one gene (Awo_c08260; hydA1) encoding a protein with high similarity (43%) to HydA1 (Awo_c26970) from the [FeFe]-hydrogenase of A. woodii. HydA2 is predicted to contain a complete H-Cluster. The genetic organization is similar to a gene cluster of Treponema primitia that was reported to belong to the same class as the hydrogenase-coupled formate dehydrogenase (FdhF) from Escherichia coli, which oxidizes formate and produces H₂ during sugar fermentation. Based on the similarity of FdhF₁₁ to FdhF of E. coli and the presence of the hydrogenase genes it was suggested for the enzyme of T. primitia that it directly uses molecular hydrogen as reductant [29]. A related gene cluster was also found in Eubacterium acidaminophilum [30]. This gene cluster consists of genes encoding hydrogenase and formate dehydrogenase subunits but the encoded hydrogenase is more complex and similar to the multimeric hydrogenase from A. woodii. Graetzkoedleffer et al. [30] proposed the most simple type of a formate hydrogen lyase system. Based on the analogy we suggest that Awo_c08190-Awo_c08260 encode a hydrogenase-coupled formate dehydrogenase that oxidizes hydrogen and donates the electrons via the electron transfer subunits HycB3 and HycB1/2 to CO₂ which is reduced to formate (Fig. 3).

Re-gaining reduced ferredoxin in the carbon pathway

The methylene-THF reductase catalyzed reaction is the only considerably exergonic reaction of the pathway and, therefore, was already considered in 1977 [8] as one of the energy conserving reactions in acetogens. The methylene-THF reductase was purified from the two acetogens Clostridium formicoaceticaicum [31] and Blautia producta [32]. The latter one consists of a single subunit with a molecular mass of 32 kDa, contains FAD as cofactor and is NADH-dependent. In contrast, the enzyme from C. formicoaceticaicum

Figure 2. Genetic organization and subunit composition of the [FeFe]-hydrogenase of A. woodii. Awo_c27000 (HydE) is not a component of the active enzyme but may be involved in assembly or signalling. doi:10.1371/journal.pone.0033439.g002
consists of two subunits (26 and 35 kDa), has been proposed to be ferredoxin-dependent and contains FAD, FeS cluster and zinc. In *A. woodii*, there is one gene (Awo_c09310) that encodes a soluble, 33 kDa protein with similarity to methylene-THF reductases (MetF) with a conserved flavin binding site. The preceding gene (Awo_c09300) encodes a protein with 205 amino acids and the region from amino acid 104 to 200 is similar to the C-terminal domain of methylene-THF reductases. This domain is encoded in many organisms next to *metF*, in several organisms it is even fused to *metF*. Awo_c09300 encodes for a 23 kDa protein that contains 8 conserved cysteine residues, which could coordinate two iron-sulfur clusters. Therefore it seems likely that this protein is the small subunit of the methylene-THF reductase of *A. woodii* and the entry point for electrons. Thus, the methylene-THF reductase is suggested to have a small (MetV) and large subunit (MetF) encoded by Awo_c09300 and Awo_c09310 with flavin and FeS cluster as cofactors.

The reduction of methylene-THF ($E^\circ_{\text{methylene-THF/methyl-THF}} = -200 \text{ mV}$) with NADH is exergonic [33] and was suggested to drive the reduction of ferredoxin with NADH [34] by electron bifurcation [26,35]. This is supported by the predicted flavin and FeS centers (Fig. 4). Reduction of ferredoxin in this step is beneficial for the overall bioenergetics of the pathway (see below).

Interestingly, the genes encoding the small and large subunit of the methylene-THF reductase are cotranscribed with the preceding gene, Awo_c09290, indicating that it is also involved in the reduction of methylene-THF. Awo_c09290 encodes a protein with similarity (61%) to the RnfC-subunit of the Rnf complex, and the residues for binding flavin and the FeS-cluster are conserved. Thus, it is named RnfC2. In contrast to RnfC1 an extension of 220 amino acids is found at the N-terminus, which is surprisingly similar (59%) to the small subunit of the methylene-THF reductase. It is, therefore, conceivable that RnfC2 could substitute for the small subunit in the methylene-THF reductase and bind directly to the Rnf complex at the RnfC binding site. The electrons derived from the Rnf complex could then be transferred to the methylene-THF reductase without the intermediate step via NADH (Fig. 4, right panel). Reduced ferredoxin from the methylene-THF reductase could be reoxidized immediately at RnfB. This would fit to previous findings that this step may be membrane-associated and involved in ion translocation [36,37].

**Energy conservation during autotrophic growth: a quantitative model**

Based on the experimental data and the predictions from the genome sequence, a model for the bioenergetics of acetogenesis from H$_2$ + CO$_2$ is proposed (Fig. 5). Oxidation of hydrogen is catalyzed by the electron bifurcating [FeFe]-hydrogenase, six mol of hydrogen are used to reduce three mol of NAD$^+$ and three mol of ferredoxin. The three mol of ferredoxin are oxidized by the Rnf complex to reduce another three mol of NAD. Assuming a Na$^+$/e$^-$ stoichiometry of 1, electron transfer is accompanied by the

**Figure 3.** Subunit composition of the formate dehydrogenase of *A. woodii*. *fdhF1* and *fdhF2* code for sulfur and selenium containing isoenzymes, respectively. HycB1 and HycB2 may be specific for FdhF1 and FdhF2, respectively. The electron transfer subunits HycB1, HycB2 and HycB3 each have four conserved tertacysteine motifs.

**Figure 4.** Subunit composition and model of the methylene-THF reductase of *A. woodii*. Panel A depicts an indirect coupling via NADH+H$^+$ and the small subunit of the methylene-THF reductase (MetV) as electron input module, panel B a direct coupling of the methylene-THF reductase to the Rnf complex via RnfC2.
to formate with H$_2$ in *A. woodii* is catalyzed by a cytoplasmic enzyme complex composed of a [FeFe]-hydrogenase, two electron transfer iron-sulfur proteins and a molybdenum/tungsten-dependent formate dehydrogenase.

The free energy change associated with 2 CO$_2$ reduction with 4 H$_2$ to acetate under physiological conditions (CO$_2$ partial pressure of 0.2×10$^5$ Pa, a H$_2$ partial pressure of 250 Pa and an acetate concentration of 10 mM at pH 7) is about ~40 kJ/mol which is sufficient to drive the phosphorylation of only less than 1 mol ADP considering that under the irreversible conditions in cells between 50 and 80 kJ/mol are required for the synthesis of 1 mol ATP [8].

The hydrogen threshold

To determine the minimal hydrogen concentration that can be used by *A. woodii* cell suspensions were incubated under different hydrogen partial pressures and the hydrogen consumption was monitored by gas chromatography. *A. woodii* was able to oxidize hydrogen down to 2500 ppm (250 Pa) which leads to a redox potential of about ~540 mV (Fig 6). Therefore, we can conclude that the minimal hydrogen concentration in the ecosystem must be above 250 Pa to allow litho-autotrophic growth on H$_2$+CO$_2$. Acetogenesis from H$_2$+CO$_2$, has a free energy change of ~95 kJ per mol which leads to a theoretical hydrogen threshold (ΔG = 0 kJ/mol) of about 10 Pa. If we assume an ATP gain of n ATP/acetate this leads to the equation:

$$4 \text{H}_2 + 2 \text{CO}_2 + n \text{ADP} + n \text{P}_i \rightarrow \text{CH}_3\text{COO}^- + H^+ + n \text{ATP} + 2 \text{H}_2\text{O}$$

With a ΔG of ~45 kJ/mol (ΔG for ADP phosphorylation ~50 kJ/mol) and n = 0.5 the theoretical threshold for H$_2$ increases to approximately 100 Pa. For n = 1 the threshold is reached at approximately 1100 Pa. The measured and theoretical hydrogen concentrations are in very good accord leading to an ATP gain in acetogenesis of 0.5 to 1 mol ATP/mol acetate. It has to be considered that the measured threshold is always lower than the theoretical one due to a partial uncoupling of metabolism and energy conservation. The threshold for H$_2$ of 250 Pa is very high compared to the thresholds of methanogens without cytochromes which is one order of magnitude lower [38]. Therefore *A. woodii* can not compete with those methanogens in its natural habitat for H$_2$+CO$_2$ as carbon and energy source.

Conclusions

The acetogenic bacterium *A. woodii* is a prime candidate for an organism that catalyzes what may be one of the first life sustaining processes on earth based on CO$_2$ fixation. Acetate formation from hydrogen and carbon dioxide by the Wood-Ljungdahl pathway is catalyzed by cytoplasmic, soluble enzymes. Energy conservation is via a chemiosmotic mechanism of the simplest type with just two enzyme complexes: the ATP synthase and the Rnf complex. The Rnf complex couples oxidation of reduced ferredoxin by electron bifurcation that are used to reduce two mol of CO$_2$ to CO. The model gives an even redox balance and postulates an ATP gain of 0.75 ATP per mol of acetate formed.

The reduction of CO$_2$ to formate with H$_2$ is endergonic by +3 kJ/mol under physiological standard conditions (H$_2$ and CO$_2$ as gases at 10$^5$ Pa partial pressure, formate at 1 M concentration and pH = 7.0). As will be shown below, the threshold concentration of *A. woodii* for H$_2$ at a CO$_2$ partial pressure of 0.2×10$^5$ Pa is 250 Pa. Under these conditions the equilibrium concentration of formate is about 0.1 mM which is in the range of the $K_m$ for formate of formyl-THF synthase, which catalyzes the second step in CO$_2$ reduction to acetate, namely the formation of formyl-THF from formate, THF and ATP. The reduction of CO$_2$ to formate with H$_2$ in *A. woodii* is therefore most likely not energy dependent. Consistent with this hypothesis, the reduction of CO$_2$ and translocation of six mol of Na$^+$. Recentrity of the six mol of Na$^+$ allows for the production of 1.5 mol ATP by the ATP synthase, assuming a stoichiometry of 4H$^+$/ATP for the ATP synthase. Two mol NADH+H$^+$ are channelled into the methylene-THF dehydrogenase reaction, four mol into the methylene-THF reductase reaction. The latter may yield two mol of reduced ferredoxin by hydrogenase reaction, four mol into the methylene-THF reductase reaction. The latter may yield two mol of reduced ferredoxin by hydrogenase reaction. Four mol of Na$^+$ may be used bioenergetically, to reduce four mol of CO$_2$. The model gives an even redox balance and postulates an ATP gain of 0.75 ATP per mol of acetate formed.

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Conclusions

The acetogenic bacterium *A. woodii* is a prime candidate for an organism that catalyzes what may be one of the first life sustaining processes on earth based on CO$_2$ fixation. Acetate formation from hydrogen and carbon dioxide by the Wood-Ljungdahl pathway is catalyzed by cytoplasmic, soluble enzymes. Energy conservation is via a chemiosmotic mechanism of the simplest type with just two enzyme complexes: the ATP synthase and the Rnf complex. The Rnf complex couples oxidation of reduced ferredoxin by electron bifurcation that are used to reduce two mol of CO$_2$ to CO. The model gives an even redox balance and postulates an ATP gain of 0.75 ATP per mol of acetate formed.

The reduction of CO$_2$ to formate with H$_2$ is endergonic by +3 kJ/mol under physiological standard conditions (H$_2$ and CO$_2$ as gases at 10$^5$ Pa partial pressure, formate at 1 M concentration and pH = 7.0). As will be shown below, the threshold concentration of *A. woodii* for H$_2$ at a CO$_2$ partial pressure of 0.2×10$^5$ Pa is 250 Pa. Under these conditions the equilibrium concentration of formate is about 0.1 mM which is in the range of the $K_m$ for formate of formyl-THF synthase, which catalyzes the second step in CO$_2$ reduction to acetate, namely the formation of formyl-THF from formate, THF and ATP. The reduction of CO$_2$ to formate with H$_2$ in *A. woodii* is therefore most likely not energy dependent. Consistent with this hypothesis, the reduction of CO$_2$ to formate with H$_2$ in *A. woodii* is catalyzed by a cytoplasmic enzyme complex composed of a [FeFe]-hydrogenase, two electron transfer iron-sulfur proteins and a molybdenum/tungsten-dependent formate dehydrogenase.

The free energy change associated with 2 CO$_2$ reduction with 4 H$_2$ to acetate under physiological conditions (CO$_2$ partial pressure of 0.2×10$^5$ Pa, a H$_2$ partial pressure of 250 Pa and an acetate concentration of 10 mM at pH 7) is about ~40 kJ/mol which is sufficient to drive the phosphorylation of only less than 1 mol ADP considering that under the irreversible conditions in cells between 50 and 80 kJ/mol are required for the synthesis of 1 mol ATP [8].
limitations. The metabolism is optimized to (i) get as much as possible reduced ferredoxin to fuel the Rnf complex and (ii) not to use the sodium motive force to overcome energy barriers but the soluble electron bifurcating [FeFe]-hydrogenase and methylene-THF reductase and a hydrogen-coupled formate dehydrogenase. Altogether, this allows for the synthesis of about 0.5–1 mol ATP per mol of acetate produced.

Materials and Methods

Growth of A. woodii

A. woodii was cultivated at 30°C. The medium was prepared as described [40]. Fructose was used as carbon source at a final concentration of 20 mM. Growth was followed by measuring the optical density at 600 nm (OD600).

Sequencing strategy

Whole-genome sequencing was performed using the 454 FLX pyrosequencing system (Roche 454, Branford, USA). Total DNA of A. woodii was extracted by MasterPure™ complete DNA purification kit (Epicentre, Madison, USA). The isolated DNA was used to create 454-shotgun libraries following the GS FLX general library protocol (Roche 454, Branford, USA). One medium lane of a Titanium picotiter plate was used for sequencing of the library, resulting in 232121 shot gun reads. The reads were de novo assembled using the Roche Newbler assembly software 2.3 (Roche 454).

Gene prediction and annotation

Annotation was done by using the ERGO tool from Integrated Genomics with a two-step approach. Initially, all proteins were screened against Swiss-Prot and NCBI databases and available protein sequences from other completed genomes by using FASTA3. All predictions were verified and modified manually by comparing the protein sequences with Pfam, GenBank, ProDom, COG, and Prosite public databases. All coding sequences were searched for similarities to protein families and domains using CD-search [41].

Proteome analysis

A. woodii was grown either on fructose or on hydrogen and carbon dioxide. Cells were harvested during early exponential, late exponential and stationary growth. 2D gel electrophoresis (500 µg soluble protein, pH 4–7) was done according to [42]. Staining and analysis of the gels and identification of proteins by mass spectrometry were done as described earlier [43] except that a Proteome Analyzer 4800 was used and the signal-to-noise ratio of the TOF-TOF measurement was raised to 10. An A. woodii data base was used for searching the peak lists.

Transcriptional organization

Cells were grown on fructose and harvested in the exponential growth phase. Isolation of RNA and analysis of transcriptional organization was performed as described previously [44].

Determination of the H2 threshold concentration

H2-threshold concentrations were determined with 1 g cells (wet mass) harvested in the exponential growth phase and resuspended in 1 ml medium. The 2 ml cell suspensions were transferred to 150 ml H2-free Müller–Krempel flasks (Bulach, Germany) sealed by Perbunan rubber stoppers (Deutsch & Naumann, Berlin, Germany) and repeatedly evacuated and filled with 80% N2/20% CO2 to remove remaining traces of H2. Then H2 was added up to 2000 Pa. The flasks were then rapidly shaken at 30°C. Gas samples were taken with a gas-tight syringe (Hamilton, Bonaduz, Switzerland) at first every 10 min and then later every hour. After the H2 concentration remained constant, H2 was added again to ascertain that the same H2 threshold was reached a second time. The samples were analyzed by gas chromatography using a stainless steel separation column (0.5 nm molecular sieve: 80/100 mesh; length =2 m×4 mm) kept at 120°C. H2 was detected with a thermal conductivity detector (GC-8A; Shimadzu) (carrier...
Supporting Information

Figure S1 The soluble proteome of A. woodii grown either on fructose (greenimage) or on H₂+CO₂ (redimage). The dual channel image was created with the Delta 2D software (Decodon GmbH, Greifswald, Germany). Proteins were prepared during early exponential growth, separated in a pH gradient 4–7 and stained with colloidal Coomassie Brilliant Blue.

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