Global mRNA decay and 23S rRNA fragmentation in *Gluconobacter oxydans* 621H

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**Abstract**

**Background:** *Gluconobacter oxydans* is a strictly aerobic Gram-negative acetic acid bacterium used industrially for oxidative biotransformations due to its exceptional type of catabolism. It incompletely oxidizes a wide variety of carbohydrates regio- and stereoselectively in the periplasm using membrane-bound dehydrogenases with accumulation of the products in the medium. As a consequence, only a small fraction of the carbon and energy source enters the cell, resulting in a low biomass yield. Additionally, central carbon metabolism is characterized by the absence of a functional glycolysis and absence of a functional tricarboxylic acid (TCA) cycle. Due to these features, *G. oxydans* is a highly interesting model organism. Here we analyzed global mRNA decay in *G. oxydans* to describe its characteristic features and to identify short-lived mRNAs representing potential bottlenecks in the metabolism for further growth improvement by metabolic engineering.

**Results:** Using DNA microarrays we estimated the mRNA half-lives in *G. oxydans*. Overall, the mRNA half-lives ranged mainly from 3 min to 25 min with a global mean of 5.7 min. The transcripts encoding GroES and GroEL required for proper protein folding ranked at the top among transcripts exhibiting both long half-lives and high abundance. The F-type H+-ATP synthase transcripts involved in energy metabolism ranked among the transcripts with the shortest mRNA half-lives. RNAseq analysis revealed low expression levels for genes of the incomplete TCA cycle and also the mRNA half-lives of several of those were short and below the global mean. The mRNA decay analysis also revealed an apparent instability of full-length 23S rRNA. Further analysis of the ribosome-associated rRNA revealed a 23S rRNA fragmentation pattern exhibiting new cleavage regions in 23S rRNAs which were previously not known.

**Conclusions:** The very short mRNA half-lives of the H+-ATP synthase, which is likely responsible for the ATP-proton motive force interconversion in *G. oxydans* under many or most conditions, is notably in contrast to mRNA decay data from other bacteria. Together with the short mRNA half-lives and low expression of some other central metabolic genes it could limit intended improvements of *G. oxydans*’ biomass yield by metabolic engineering. Also, further studies are needed to unravel the multistep process of the 23S rRNA fragmentation in *G. oxydans*.

**Keywords:** *Gluconobacter oxydans*, mRNA decay, ATP synthase, Tricarboxylic acid cycle, Ribosome, 23S rRNA fragmentation, Intervening sequence

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Background

*Glucobacter oxydans* is a Gram-negative, strictly aerobic acetic acid bacterium industrially used for oxidative biotransformations of carbohydrates. Important products are e.g. L-sorbose, a precursor for vitamin C production, dihydroxyacetone, a chemical used for tanning lotions, or 6-amino-L-sorbose, a precursor of the antidiabetic drug miglitol [1–6]. The beneficial ability of G. *oxydans* is the incomplete oxidation of a variety of substrates (e.g. sugars and sugar alcohols) in the periplasm by membrane-bound dehydrogenases and release of resulting products into the cultivation medium [7–9]. Correspondingly, only a small amount of substrate is taken up by the cell and channeled into the cytoplasmic metabolism for growth [10]. Genome sequencing revealed the absence of genes coding for enzymes of the central metabolism, such as 6-phosphofructokinase, succinyl-CoA synthetase, and succinate dehydrogenase [11]. Accordingly, the Embden-Meyerhof-Parnas pathway (glycolysis) and the tricarboxylic acid (TCA) cycle are incomplete. Both the dominant incomplete periplasmic oxidation and the incomplete cytoplasmic sugar metabolism contribute to the limited assimilation of carbohydrates into new cell material and therefore to a low biomass yield. Industrial use of G. *oxydans* for oxidative biotransformations is therefore costly. To overcome these hindrances, metabolic engineering was performed to complete the TCA cycle by introducing heterologous genes for succinate dehydrogenase and succinyl-CoA synthetase into the genome with simultaneous deletion of the genes for the membrane-bound and soluble glucose dehydrogenase, thus abolishing periplasmic and cytoplasmic glucose oxidation [12, 13]. Furthermore, the NADH oxidation capacity was increased by introducing an additional NADH dehydrogenase gene [14]. These steps led to an increase of the biomass yield on glucose by up to 60%, thereby reducing the costs for biomass formation. Although this is already very advantageous for industrial applications, other still unrecognized bottlenecks in G. *oxydans* naturally evolved partially incomplete metabolism might exist.

Therefore, in this study we conducted genome-wide mRNA decay analysis to get further insights into the physiology of G. *oxydans*. In living cells, the abundance of mRNAs is a result of the balance between gene expression and degradation of mRNAs. Global mRNA decay analysis in prokaryotes was already described, for example, for the model microorganisms *Escherichia coli* [15], *Bacillus subtilis* [16], and *Mycobacterium tuberculosis* [17]. These studies used rifampicin for inhibition of transcription at different time points during growth to measure the changes of relative mRNA levels using DNA microarrays. These changes reflect the degradation of transcripts and allow the calculation of mRNA half-lives [15, 18]. In bacteria, mRNA half-lives typically range from around 1 min or shorter up to 30 min [15, 16, 19]. Generally, a correlation between the half-lives of transcripts and the cellular function of the encoded proteins was observed in these studies. Transcripts associated to housekeeping functions such as cell envelope and ion transport exhibited relatively long mRNA half-lives, whereas genes involved in stress responses and in regulatory functions exhibited a faster transcript turnover to adapt to environmental changes in a short time [20, 21]. Likewise, mRNA half-lives are not fixed and can be affected by, for example, alterations in the growth rate when environmental conditions change [22, 23]. Since mRNA half-lives are affected by different growth conditions, regulation of their stability and degradation is quite diverse. It is dependent on secondary structures of the 5′ and 3′ untranslated regions, posttranscriptional modifications such as polyadenylation, abundance of ribonucleases, presence of cleavage sites recognized by ribonucleases, interaction with small regulatory RNAs, as well as location of the mRNAs in the cell [20, 24].

Here, we measured temporal RNA level changes in G. *oxydans* 621H in response to rifampicin and estimated the mRNA half-lives. The mRNA decay analysis additionally revealed an apparent instability of the full-length 23S rRNA. Enrichment of ribosomes and further analysis of the associated rRNAs uncovered that the 23S rRNA is fragmented in G. *oxydans*.

**Methods**

**Bacterial strain and cultivation conditions**

In this study, the wild type *Glucobacter oxydans* 621H strain (DSM 2343) from the German Collection of Microorganisms and Cell Cultures (DSMZ) was used. The cells were cultivated in mannitol medium containing 220 mM (4% w/v) mannitol, 5 g L⁻¹ yeast extract, 1 g L⁻¹ KH₂PO₄, 1 g L⁻¹ (NH₄)₂SO₄, 2.5 g L⁻¹ MgSO₄ × 7 H₂O, and 50 μg mL⁻¹ cefoxitin as antibiotic. Cells were grown in 500 mL shaking flasks with three baffles containing 50 mL of the mannitol medium (30 °C, 140 rpm). Cell growth in liquid culture was followed by measuring the optical density at 600 nm (OD₆₀₀) using a spectrophotometer (UV-1800, Shimadzu). If rifampicin was applied, it was added to a cell culture with an OD₆₀₀ of 0.6 to 0.8 from a stock solution (50 mg mL⁻¹ in methanol) to obtain the final concentration as indicated, while the appropriate volume of methanol without rifampicin was added to the control culture. For isolation of total RNA from cells and purification of ribosomes, cells were harvested at the indicated OD₆₀₀ as described below and stored at −20 °C until use [10].

**Isolation of RNA**

For determination of mRNA half-lives by DNA microarray analysis, total RNA was isolated from harvested
cells as described [10]. The RNA fraction of ribosomes was isolated by phenol-chloroform-isoamyl alcohol (25:24:1) and chloroform-isoamyl alcohol (24:1) extractions followed by ethanol precipitation [25]. RNA concentrations were determined photometrically using a Nanodrop ND-1000 and fluorometrically using a Qubit® RNA BR Assay Kit (Life Technologies). Both RNAs are present in mix A and mix B in a 1:1 ratio A/B (Agilent Technologies). Accordingly, the normalization factor was calculated that the log base 2 of the normalized ratio of median values of the 1:1 control RNAs is 0 on average. Subsequently, the ratio of median value for each gene was normalized with the calculated factor. All microarray data including the normalized ratio of medians were stored for further analysis, quality filtering and mRNA half-life calculation in the in-house DNA microarray database [27]. A normalized ratio value was included in the calculation of the average of the triplicates for each time point if the following quality filter was fulfilled by the spot data (GenePix Pro 6.0): i) Flags ≥0 and ii) signal/noise ≥3 for Cy5 (F635Median / B635Median) or Cy3 (F532Median / B532Median). The resulting data matrix with the average values of the four time points was used to calculate the mRNA half-lives and R² in Excel (Microsoft) by linear fit as described for E. coli data [18]. For further analysis, genes were functionally grouped according to the assigned product functions [11]. Significant differences between functional groups were identified via a one-way ANOVA test using Excel (Microsoft). Subsequently, a post-hoc t-test was performed to identify mean mRNA half-lives of functional groups, which differ significantly from the overall average half-life. p values were adjusted using Bonferroni correction, which allows adjustment of p values after multiple comparisons [28].

**DNA microarray analysis**

The DNA microarray analysis aimed at the determination of the time-dependent mRNA level changes in G. oxydans after addition of rifampicin. The analysis was performed three times with independent cultures for each time point. G. oxydans cells were harvested for isolation of total RNA directly before (t0) and 2 min (t2), 5 min (t5), 10 min (t10) as well as 15 min (t15) after addition of rifampicin. Each RNA sample isolated after addition of rifampicin (tx) was compared to the t0 RNA sample. For pairwise comparisons, the A mix and the B mix of the Agilent Spike-In Kit (Agilent Technologies) was used to spike t0 and tx RNA samples accordingly. The synthesis of labeled cDNA from the RNA samples was carried out as described [10]. Custom-made 4 × 44 K DNA microarrays for genome-wide gene expression analysis were obtained from Agilent Technologies and were designed using Agilent’s eArray platform (https://earray.chem.agilent.com/earray). The array design comprised oligonucleotides for the annotated protein-coding genes and the structural RNA genes of G. oxydans 621H (CP000009, and CP000004 to CP000008) [11], as well as Agilent’s control spots. After hybridization according to the manufacturer’s instructions, the arrays were washed using Agilent’s wash buffer kit. Subsequently, the fluorescence of DNA microarrays was determined at 532 nm (Cy3-dUTP) and 635 nm (Cy5-dUTP) at 5 μm resolution with a GenePix 4000B laser scanner and GenePix Pro 6.0 software (Molecular Devices). Raw data files of fluorescence images were saved in TIFF format followed by quantitative image analysis (GenePix Pro 6.0) using the corresponding Agilent’s gene array list (GAL) file. The results were saved as GPR file containing the non-normalized ratio of median values (GenePix Pro 6.0).

**Data normalization and calculation of mRNA half-lives**

For calculation of mRNA half-lives, first the ratio of median values (GenePix Pro 6.0) reflecting the relative mRNA level changes were normalized using a factor. This factor was calculated for each hybridization based on the log base 2 of the non-normalized ratio of median values of the Spike-In (+)E1A_r60_1 and (+)E1A_r60_a20 RNAs each having 32 array spots randomly scattered (Agilent Technologies). Both RNAs are present in mix A and mix B in a 1:1 ratio A/B (Agilent Technologies). Accordingly, the normalization factor was calculated that the log base 2 of the normalized ratio of median values of the 1:1 control RNAs is 0 on average. Subsequently, the ratio of median value for each gene was normalized with the calculated factor. All microarray data including the normalized ratio of medians were stored for further analysis, quality filtering and mRNA half-life calculation in the in-house DNA microarray database [27]. A normalized ratio value was included in the calculation of the average of the triplicates for each time point if the following quality filter was fulfilled by the spot data (GenePix Pro 6.0): i) Flags ≥0 and ii) signal/noise ≥3 for Cy5 (F635Median / B635Median) or Cy3 (F532Median / B532Median). The resulting data matrix with the average values of the four time points was used to calculate the mRNA half-lives and R² in Excel (Microsoft) by linear fit as described for E. coli data [18]. For further analysis, genes were functionally grouped according to the assigned product functions [11]. Significant differences between functional groups were identified via a one-way ANOVA test using Excel (Microsoft). Subsequently, a post-hoc t-test was performed to identify mean mRNA half-lives of functional groups, which differ significantly from the overall average half-life. p values were adjusted using Bonferroni correction, which allows adjustment of p values after multiple comparisons [28].

**Determination of FPKM expression values**

For determination of FPKM values reflecting mRNA abundance, we used FASTQ files that were generated previously for comprehensive RNAseq analysis of G. oxydans (NCBI accession number: ERR2232412) [29]. Sequencing reads were trimmed and strand-specifically mapped to the genome reference (CP0000009) and the five plasmids pGOX1 to pGOX5 (CP000004 – CP000008) using the RNAseq analysis tool of the CLC Genomics Workbench (Qiagen Aarhus A/S) to determine absolute FPKM values [30]. Only uniquely mapped reads with ≤1% mismatches were considered for this analysis. Linear regression analysis of transcript abundances and mRNA half-lives was performed with GraphPad Prism 7.00 using default settings.

**Purification of ribosomes**

For preparation of lysates, frozen cell pellets were thawed and resuspended to 0.2 g mL⁻¹ in lysis buffer (70 mM KCl, 10 mM MgCl₂, 10 mM Tris-HCl, pH 7.4). Resuspended cells were disrupted in a French press (SLM Amino) at 15,000 psi (3 passages) using 10 mL of cell suspension. Remaining intact cells and debris were removed by centrifugation (20,000 g; 20 min; 4 °C) and the supernatant was filtered through a 0.22 μm filter. The protein concentration was determined using Pierce

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BCA protein assay kit (Thermo Fisher Scientific). If required, the volume was adjusted by addition of lysis buffer to obtain a concentration of 1.5 to 4.5 mg\text{protein} \text{mL}^{-1}. Ribosomes were purified based on the use of a strong anion exchange quaternary amine (QA) monolithic column [31]. Therefore, 0.5 mL of the prepared cell lysate was injected into an ÄKTA pure FPLC system (GE Healthcare Life Sciences) equipped with two CIM\textsuperscript{+} QA-0.34 mL monolithic disks (BIA separations) encased within a polyetheretherketone (PEEK) housing and pre-equilibrated in lysis buffer. The chromatography was performed at a flow rate of 2 mL min\textsuperscript{−1} using lysis buffer (buffer A) and lysis buffer containing 1 M NaCl (buffer B). The loaded QA monolithic disks were washed with 5 column volumes of buffer A and sequentially eluted with 7 column volumes of 40%, 56% and 100% of buffer B. During the step-wise elution the online chromatogram was visually inspected for upcoming peaks to manually collect from the start to the end of a peak into one elution fraction. The collected fractions were further analyzed using mass spectrometry to identify proteins.

Protein identification

For the identification of proteins in relevant elution fractions, proteins were precipitated with trichloroacetic acid (10%) and resuspended in 50 \text{µL} of digestion buffer followed by tryptic digestion using the Trypsin Singles Proteomics Grade Kit according to the manufacturer's instructions for solution digestion in a volume of 100 \text{µL} (Sigma-Aldrich). After tryptic digestion, the peptides were separated chromatographically on a nanoLC Eksigent ekspert \textsuperscript{™} 425 system (Sciex) coupled with a quartz emitter Tip (New Objective) to a TripleTof\textsuperscript{™} 6600 mass spectrometer (Sciex). Digested samples were loaded on a pre-column (ChromXP C18–3 \text{µm}, 350 \text{µm} × 0.5 mm, Sciex) for desalting and enrichment using a flow of 3 \text{µL/min} (10 min) of buffer A (0.1% formic acid in HPLC grade water). The separation of peptides followed on an analytical column (ChromXP 3C18-CL-120, 0.075 × 150 mm, Sciex) with a gradient method (125 min) using buffer A and buffer B (0.1% formic acid in acetonitrile) at 40 °C and a flow of 300 nL/min. The gradient conditions were 5% of buffer B for 1 min, 5–9% for 9 min, 9–20% for 50 min, 20–40% for 40 min, 40–80% for 5 min and 80% for 4 min. The mass spectrometer was operated with a “top 50” method: Initially, a 250 ms survey scan (TOF-MS mass range 400–1500 amu, high resolution mode) was collected from which the top 50 precursor ions were automatically selected for fragmentation, whereby each MS/MS event (mass range 100–1700 amu, high sensitivity mode) consisted of a 75 ms fragment ion scan. The source and gas settings were 2200 V spray, 40 psi curtain gas, 6 psi ion source gas, and 75 °C interface heater. The mass spectrometry data obtained were processed with ProteinPilot\textsuperscript{™} using the Paragon algorithm (V4.5 beta, Sciex) for identification of peptides (99% confidence) and proteins (1% false discovery rate).

rRNA sequencing

RNA purified from ribosomes was sequenced to analyze the rRNAs. Sequencing libraries were generated using the TruSeq stranded mRNA sample preparation kit (Illumina) according to the manufacturer’s instructions, yet without the fragmentation step. cDNA libraries were quantified and sequenced as described [29]. Sequencing reads were processed and mapped to both the four 23S rRNA genes (GOX0221, GOX1159, GOX1319, and GOX1467) and 16S rRNA genes (GOX0224, GOX1156, GOX1316, and GOX1464) using CLC Genomics Workbench (Qiagen Aarhus A/S). The default mapping parameters were changed to consider only reads, which mapped over their complete length with an identity of at least 99%. The coverage per base was extracted and manually inspected in Excel (Microsoft) to identify regions with a coverage < 5% of the average gene coverage and visualized with Origin (OriginLab).

Results

Global mRNA half-lives in \textit{G. oxydans} ranged from 2 to 25 min

As outlined in the introduction, \textit{G. oxydans} has an unusual type of metabolism characterized by an incomplete periplasmic oxidation of carbon sources, no glycolysis, lack of a full TCA cycle, and low biomass yield. It thus represents an interesting model organism for the acetic acid bacteria. In a recent study, we employed RNAseq approaches to determine global gene expression, operon structures, transcription start sites, promoter motifs, and ribosome binding sites [29]. Here, we extended these genome-wide studies by determining the mRNA half-lives using DNA microarrays to improve the understanding of gene expression in \textit{G. oxydans}. Typically, mRNA half-lives are determined in rifampicin experiments. Rifampicin inhibits the activity of the RNA polymerase [32], thereby enabling the measurement of the time-dependent decrease of mRNA levels by turnover in the absence of mRNA de novo synthesis. We first tested the influence of different concentrations of rifampicin on the growth of \textit{G. oxydans} in liquid culture. Based on data reported for \textit{Escherichia coli}, \textit{Bacillus subtilis}, \textit{Mycobacterium tuberculosis} and two \textit{Sulfolobus spp}, we focused on a range up to 250 \text{µg/mL} [15–17, 19]. With 250 \text{µg/mL} and 200 \text{µg/mL} of rifampicin a notable drop of the cell density within 30 min was observed, suggesting cell damage and lysis to some extent (Fig. 1a). With 150 \text{µg/mL} of rifampicin a growth inhibition within 30 min was observed without a drop of the cell density (Fig. 1a). Although apparently similar results in growth stop were obtained with 100 \text{µg/mL} and 50 \text{µg/mL} of
rifampicin (data not shown) as with 150 μg/mL, we chose 150 μg/mL of rifampicin to analyze the global mRNA decay in *G. oxydans*. For comparison of transcriptomes, total RNA was isolated from cells directly before addition of rifampicin and 2, 5, 10, and 15 min afterwards. Formaldehyde agarose gels were used to size-separate and visualize the isolated RNA to assess its quality (Fig. 1b). The 23S rRNA band was generally much weaker compared to the 16S rRNA and almost disappeared after 15 min, suggesting a decay or further processing of the expected mature full-length 23S rRNA transcripts with 2709, 2710 and 2711 nt in *G. oxydans* 621H. These results suggest that for *G. oxydans* the rRNAs should better not be used for microarray data normalization, as it was done in other mRNA decay studies where the rRNAs of the host were considered as stable enough to be used for data normalization [18]. We therefore used the Spike-In RNA mixtures A and B (Agilent Technologies), which contain several RNA transcripts for which known ratios between mix A and mix B can be expected. For the pairwise DNA microarray comparisons of transcriptomes corresponding to the time point before addition of rifampicin (t0) and 2, 5, 10, and 15 min after addition of 150 μg/mL rifampicin were pairwise mixed and hybridized on Agilent 4-plex arrays. mRNA half-lives were calculated based on three biological replicates including dye-swap labeling.

**Fig. 1** Overview of global mRNA decay analysis. a Growth of *G. oxydans* 621H in the presence of different concentrations of rifampicin as well as the controls methanol and water. The addition of rifampicin or a control is indicated by the arrow. b Formaldehyde agarose gel (1.8%) analysis to inspect the quality of total RNA isolated from *G. oxydans* cells before and after addition of 150 μg/mL of rifampicin. Aliquots of 1 μg of total RNA were loaded. Bands corresponding to the 23S rRNAs (2709 to 2711 nt) and the 16S rRNAs (1478 nt) are indicated as well as the sizes of the RNA fragments present in the Ribofl Ruler Low Range RNA Ladder (M). w/o Rif, without rifampicin. c Schema of the experimental design. RNA isolated at the given time points were mixed with Spike-In RNA mix A or B and used for cDNA synthesis and Cy3 or Cy5 labeling. cDNA synthesized from RNA isolated from cells before addition of rifampicin (t0) and 2, 5, 10, and 15 min after addition of 150 μg/mL rifampicin were pairwise mixed and hybridized on Agilent 4-plex arrays. mRNA half-lives were calculated based on three biological replicates including dye-swap labeling. d Histogram showing the distribution of calculated mRNA half-lives of 2500 genes from *G. oxydans* where R² was > 0.7 when using all four or the first three sampling times.
the respective hybridization experiments were normalized. The normalized ratio data were then used to calculate the average mRNA ratio for each gene at each time point (t0/t2, t0/t5, t0/t10, t0/t15) from three independent biological replicate experiments. Subsequently, the mRNA half-lives can be estimated via linear regression of the log ratios over time. Based on the four time points of the analysis and filtering for $R^2 > 0.7$ of the linear fit [18], we obtained half-life values for 1193 transcripts. That number corresponds to 44% of all protein-coding genes of *G. oxydans* 621H [11] and is in the range reported for other mRNA decay studies, e.g. for *B. subtilis* (35%), *E. coli* (53%), *M. tuberculosis* (53%), or *S. acidocaldarius* (70%) [15–17, 19]. Nevertheless, there is no half-life estimation for the remaining genes which still represent a significant proportion (56% for *G. oxydans*). The uniform calculation by always including the latest time point may not be adequate for all genes. For example, genes with shorter half-lives may exhibit the decrease in the mRNA level at earlier sampling times, while at later sampling times there is no further decrease for several reasons including inherent limits of the DNA microarray technology used. Therefore, many genes exhibiting this type of mRNA decay kinetics will likely not fulfill a well-intentioned $R^2$ criterion for filtering results of the linear regression analysis when later time points are included. Depending on the range of the absolute sampling times it is legitimate to check the outcome of half-life estimations by linear regression with a focus on earlier time points. Accordingly, for the 1446 remaining genes (56%), where $R^2 > 0.7$ was not fulfilled when using all sampling times, we omitted the last time point (t0/t15) from half-life calculation. Based on the first three sampling times (t0/t2, t0/t5, t0/t10) further half-life estimations with an $R^2 > 0.7$ were obtained for 1307 transcripts. Together, this resulted in mRNA half-life estimations for 2500 (95%) of the protein-coding genes of *G. oxydans* (Additional file 1: Table S1). Overall, the calculated mRNA half-lives of the 2500 genes mainly ranged from approximately 2 min to 25 min. Solely two operons consisting of 8 genes encoding hypothetical and phage-related proteins exhibited apparent mRNA half-lives from 25 to 75 min. The global mean was 5.7 min and the median 5.1 min, which is also reflected by the high number of transcripts exhibiting half-lives between 3 and 7 min (Fig. 1d).

**mRNA half-life characteristics in *G. oxydans***

In previous studies, correlations between the stability of mRNAs and the functional group of the gene products were observed [15, 16, 19]. For *G. oxydans*, one-sided ANOVA test revealed significant differences between functional groups ($p < 0.0001$). Taken into account the within-group variance subsequent *post hoc* analysis identified 2 categories with significantly shorter mean half-lives than the overall average half-life. These categories were ATP-proton motive force interconversion and RNA metabolism (Table 1, Fig. 2).

According to observations in *E. coli* and the archaeon *Sulfolobus solfataricus*, mRNAs exhibiting higher transcript abundance were on average less stable [15, 19]. In *G. oxydans* we also observed such an inverse relationship (Fig. 3a and Additional file 1: Table S2). On average genes with higher expression values showed slightly shorter apparent mRNA half-lives and *vice versa*. However, certain transcripts exhibited both high abundance and long half-life, such as the transcripts for chaperonin GroEL (GOX1902) and co-chaperonin GroES (GOX1901), nucleoside diphosphate kinase (GOX1927), 50S ribosomal protein L17 (GOX3055), coenzyme PQ synthesis protein PqqA (GOX0987), as well as a number of putative/hypothetical proteins. On the other hand, certain transcripts exhibited both low abundance and short half-life, such as the transcripts for the cell division proteins FtsE (GOX0273) and FtsX (GOX0274), putative fatty acyl CoA synthetase (GOX1394), ubiquinol-cytochrome c reductase cytochrome b subunit (GOX0566), NAD(P)H-dependent glycerol-3-phosphate dehydrogenase (GOX1880), and cytochrome c subunit and small subunit of membrane-bound aldehyde dehydrogenase (GOX0585, GOX0586). Plotting the mRNA half-lives versus the ORF lengths showed no correlation for *G. oxydans* (Fig. 3b). Similarly, for *E. coli* such a correlation was also not found [15, 33].
Table 1 Functional categories or subcategories comprising 1582 assigned protein-coding ORFs with mean mRNA half-lives based on data from 1526 (96%) transcripts (Additional file 1: Table S1)

| Functional category / pathway | number of genes assigned with a calculated mRNA half-life | proportion (%) | half-life (min) |
|------------------------------|-----------------------------------------------------------|----------------|----------------|
| ATP-proton motive force interconversion | 18 | 18 | 100 | 4.0** |
| RNA metabolism | 23 | 23 | 100 | 4.4* |
| Cell division | 26 | 26 | 100 | 4.6 |
| DNA degradation | 9 | 9 | 100 | 4.7 |
| Transcription | 13 | 10 | 77 | 4.7 |
| Mono/dioxygenase | 10 | 10 | 100 | 4.7 |
| Fatty acid and phospholipid metabolism | 36 | 35 | 100 | 4.7 |
| Antibiotics resistance | 12 | 12 | 100 | 4.7 |
| tRNA metabolism | 13 | 13 | 100 | 4.7 |
| Nucleotide metabolism | 59 | 58 | 98 | 4.8 |
| Signal transduction | 30 | 29 | 97 | 4.8 |
| Cell envelope | 134 | 131 | 98 | 4.9 |
| Amino acid metabolism | 120 | 117 | 98 | 5.0 |
| Central intermediary metabolism | 33 | 31 | 94 | 5.0 |
| Degradation of proteins and peptides | 51 | 48 | 94 | 5.0 |
| Ribosome assembly | 67 | 52 | 78 | 5.0 |
| DNA replication | 26 | 26 | 100 | 5.1 |
| Pyruvate metabolism | 12 | 11 | 92 | 5.1 |
| Regulatory functions | 99 | 94 | 95 | 5.1 |
| DNA repair | 34 | 34 | 100 | 5.2 |
| Transport | 232 | 227 | 98 | 5.2 |
| Aminoacyl-tRNA biosynthesis | 33 | 33 | 100 | 5.3 |
| Pentose phosphate pathway | 13 | 13 | 100 | 5.3 |
| Biosynthesis of cofactors | 85 | 84 | 99 | 5.4 |
| Translation factors | 20 | 20 | 100 | 5.4 |
| DNA recombination | 18 | 17 | 94 | 5.5 |
| Electron transport | 53 | 53 | 100 | 5.5 |
| Unknown function | 41 | 40 | 98 | 5.5 |
| Detoxification | 30 | 27 | 90 | 5.6 |
| Tricarboxylic acid cycle | 9 | 9 | 100 | 5.6 |
| Uncharacterized oxidoreductase | 66 | 63 | 96 | 5.7 |
| Adaptations to atypical conditions | 19 | 19 | 100 | 6.1 |
| Protein folding and stabilization | 28 | 28 | 100 | 6.3 |
| Sugar and alcohol degradation | 25 | 24 | 96 | 6.3 |
| Biosynthesis and degradation of polymers | 11 | 10 | 91 | 6.4 |
| Glycolysis / Gluconeogenesis | 17 | 16 | 94 | 6.5 |
| Cell motility | 41 | 39 | 95 | 6.6 |
| Entner-Doudoroff pathway | 2 | 2 | 100 | 6.7 |
| Ion homeostasis | 6 | 6 | 100 | 6.7 |
| DNA restriction and modification | 8 | 8 | 100 | 7.0 |

a) Functional categories and number of genes are based on assigned gene product functions [11]

b) Functional categories with significantly shorter or longer mean half-lives than the overall mean half-life (5.7 min) are highlighted by ** (p < 0.001) or * (p < 0.05)
Fig. 2 Distribution of mRNA half-lives (○) with mean value indicator (|) based on the functional categories or subcategories and number of genes given in Table 1.

Fig. 3 mRNA half-lives versus FPKM expression values and ORF lengths. a Linear regression analysis showed a statistically significant slightly negative correlation (R = −0.24) between the abundance of transcripts and their half-lives. Plots are related to the half-life data obtained for 2500 transcripts (○) with R² > 0.7 based on the 4 or 3 sampling times as described. Transcripts of the molecular chaperones GroES (GOX1901) and GroEL (GOX1901) exhibited high expression values as well as long half-lives (○). The operons of the F1F0-type ATP synthase encoded by atpBEFF (GOX1110–13) and atpHAGDC (GOX1310–14) (○) belong to the operons/genes with the shortest mRNA half-lives in G. oxydans. In comparison, the transcripts of the second F1F0-type ATP synthase encoded by GOX2167–75 (○) exhibited almost 2-fold longer half-lives and approximately 80-fold lower expression values. Among the genes of the incomplete TCA cycle (○) the transcript of aconitase of (acn, GOX1335) was the most stable, while the transcripts of malatequinone oxidoreductase (mqo, GOX2070), dihydrolipoamide succinyl transferase (E2) of 2-oxoglutarate dehydrogenase (adhb, GOX1073), and dihydrolipoamide dehydrogenase (lpd, GOX2292) exhibited the shortest mRNA half-lives. FPKM values were obtained with cells grown on mannitol.

b The mRNA half-lives and ORF lengths of 2500 transcripts (○) did not correlate (R = −0.06).
operon (GOX2167–75) which is an ortholog of the Na⁺-translocating F₁Fₒ-type ATP synthases present always in addition to the H⁺-ATP synthase in the archaea Methanosarcina barkeri and M. acetivorans, in a number of marine and halotolerant bacteria and in pathogenic Burkholderia species [35–37]. These genes exhibited on average almost 2-fold longer mRNA half-lives (5.1 min) which is at the global median, yet the apparent expression levels were extremely low (80-fold lower) compared to the H⁺-ATP synthase, which presented transcript levels ranked above the global expression median (Fig. 3a). Thus, the H⁺-ATP synthase with

### Table 2

Top 20 operons exhibiting the shortest mRNA half-lives in G. oxydans and their assigned gene expression values FPKM. The arrow lines indicate operon genes and genomic orientation.

| locus tag / operon | annotation | operon half-life Δ (min) | half-life (min) | FPKM |
|-------------------|------------|--------------------------|----------------|------|
| GOX1310a | F₁Fₒ ATP synthase subunit delta | 3.1 | 3.7 | 6320 |
| GOX1311a | F₁Fₒ ATP synthase subunit alpha | 3.0 | 1780 |
| GOX1312a | F₁Fₒ ATP synthase subunit gamma | 3.1 | 1232 |
| GOX1313a | F₁Fₒ ATP synthase subunit beta | 2.9 | 1651 |
| GOX1314a | F₁Fₒ ATP synthase epsilon chain | 3.1 | 6950 |
| GOX1737 | rod shape-determining protein MreB | 2.8 | 2.4 | 908 |
| GOX1738 | rod shape-determining protein MreC | 2.8 | 383 |
| GOX1739 | hypothetical protein | 3.0 | 93 |
| GOX1110a | F₁Fₒ ATP synthase subunit b' | 2.8 | 2.9 | 937 |
| GOX1111a | F₁Fₒ ATP synthase subunit b | 2.9 | 1600 |
| GOX1112a | F₁Fₒ ATP synthase C chain | 2.9 | 1743 |
| GOX1113a | F₁Fₒ ATP synthase subunit a | 2.4 | 1911 |
| GOX0149 | cell division protein MraZ | 2.8 | 3.0 | 2124 |
| GOX0150 | SAM-dependent methytransferase | 2.8 | 528 |
| GOX0151 | hypothetical protein | 2.6 | 255 |
| GOX1427 | FAD-dependent thymidylate synthase | 2.7 | 2.7 | 1121 |
| GOX0476 | putative oxidoreductase | 2.7 | 2.9 | 658 |
| GOX0477 | hypothetical protein | 2.5 | 847 |
| GOX0478 | putative oxidoreductase | 2.8 | 1202 |
| GOX1596 | hypothetical protein | 2.7 | 2.7 | 119 |
| GOX2462 | transcriptional regulator | 2.7 | 2.7 | 4866 |
| GOX0607 | D-alanyl-D-alanine carboxypeptidase | 2.7 | 2.7 | 1540 |
| GOX1430 | serine protease | 2.7 | 2.8 | 170 |
| GOX1431 | hypothetical protein | 2.6 | 336 |
| GOX0024 | undecaprenyl pyrophosphate phosphatase | 2.6 | 2.8 | 198 |
| GOX0025 | amino acid permease | 2.5 | 142 |
| GOX2207 | methylenetetrahydrofolate reductase | 2.6 | 2.6 | 547 |
| GOX0649 | sugar-proton symporter | 2.6 | 2.6 | 364 |
| GOX2378 | short chain alcohol dehydrogenase | 2.6 | 2.6 | 301 |
| GOX1151 | hypothetical protein | 2.5 | 2.5 | 2854 |
| GOX1696 | hypothetical protein | 2.5 | 2.5 | 35 |
| GOX2067 | manganese transport protein MntH | 2.5 | 2.5 | 526 |
| GOX0942 | hypothetical protein | 2.5 | 2.5 | 178 |
| GOX1652 | heme exporter protein C | 2.4 | 2.4 | 323 |
| GOX0666 | outer membrane receptor for ferric iron uptake | 2.4 | 2.4 | 222 |
| GOX1426 | hypothetical protein | 2.4 | 2.4 | 577 |

* For the F₁Fₒ ATP synthase encoded by two operons the data of the operon GOX1310–14, which was close to the top 20, were also included.
the very short-lived mRNAs (atpBEFF and atpHAGDC) is likely the one which is involved in the energy metabolism under the conditions tested.

A characteristic of G. oxydans are the membrane-bound dehydrogenases (mDHs), often with a broad substrate spectrum, incompletely oxidizing a wide variety of carbohydrates in the periplasm with accumulation of the products in the medium [8, 11]. Many of their transcripts exhibited half-lives below the global mean of 5.7 min (Table 3). For the membrane-bound gluconate-2 dehydrogenase mGlDH (GOX1230–2) the longest mDH transcript half-lives were found (8 to 9 min), while expression levels were moderate. Polyal dehydrogenase SldAB (GOX0854–5) required for growth on mannitol and sorbitol, and membrane-bound alcohol dehydrogenase mADH (GOX1067–8) exhibited by far the highest transcript levels. Their transcript half-lives close to 4 min follow the trend of the inverse relationship between FPKM and half-life. The transcripts of the PQQ-dependent mDHs 1 (GOX1857), 3 (GOX1441), and 4 (GOX0516), sorbitol dehydrogenase (GOX2094–97), and aldehyde dehydrogenase mAcDH (GOX0585–7) exhibited very low expression values and half-lives from 4 min to 9 min.

Since G. oxydans has an unusual central metabolism, we were also interested in the stability of transcripts encoding enzymes of the central carbon metabolism including the pentose phosphate pathway (PPP), the Entner-Doudoroff pathway (EDP), and the pyruvate metabolism. Therefore, we mapped the estimated mRNA half-lives to the central carbon metabolism [38]. Overall, mRNA half-lives ranged from 3.6 min for aceEa (GOX2289) encoding pyruvate dehydrogenase E1 component subunit α to 12.4 min for one of two annotated triosephosphate isomerases (tpi, GOX2217) (Fig. 4). According to the FPKM values this tpi transcript showed almost the lowest abundance for genes involved in central carbon metabolism (Fig. 4, Additional file 1: Table S2). This apparently very low expression might be compensated by the longer mRNA half-life. Besides this tpi transcript, which is among the top 3% with the longest mRNA half-lives overall, the transcript for dihydroxyacetone kinase (GOX2222), which is linked to central carbon metabolism via the product dihydroxyacetone phosphate (DHAP), is with 19.1 min among the top 1% of the most-stable transcripts in G. oxydans. The transcripts of the PPP genes exhibited half-lives below the global mean ranging from 3.8 min to 5.1 min. Besides

Table 3 mRNA half-lives and expression values (FPKM) of genes encoding annotated membrane-bound dehydrogenases. The arrow lines indicate operon genes and genomic orientation.

| locus tag / operon | annotation | half-life (min) | FPKM |
|--------------------|------------|----------------|------|
| ↓ GOX0265          | membrane-bound glucose dehydrogenase (PQQ) | 4.4 | 146 |
| ↑ GOX0516          | PQQ-dependent dehydrogenase 4 | 8.3 | 22 |
| ↓ GOX0585          | cytochrome c subunit of aldehyde dehydrogenase | 3.7 | 32 |
| ↓ GOX0586          | membrane-bound aldehyde dehydrogenase, small subunit | 3.7 | 38 |
| ↓ GOX0587          | membrane-bound aldehyde dehydrogenase, large subunit | 9.0 | 31 |
| ↑ GOX0854          | polyl dehydrogenase subunit SldA | 4.2 | 2147 |
| ↑ GOX0855          | polyl dehydrogenase subunit SldB | 4.6 | 5992 |
| ↑ GOX1067          | alcohol dehydrogenase cytochrome c subunit precursor | 4.0 | 1807 |
| ↑ GOX1068          | alcohol dehydrogenase large subunit | 3.4 | 1706 |
| ↑ GOX1230          | gluconate 2-dehydrogenase, cytochrome c subunit | 9.0 | 208 |
| ↑ GOX1231          | gluconate 2-dehydrogenase alpha chain | 8.5 | 791 |
| ↑ GOX1232          | gluconate 2-dehydrogenase gamma chain | 7.8 | 684 |
| ↓ GOX1253          | D-lactate dehydrogenase | 5.0 | 159 |
| ↑ GOX1441          | PQQ-dependent dehydrogenase 3 | 4.7 | 31 |
| ↑ GOX1857          | PQQ-containing dehydrogenase 1 | 7.1 | 122 |
| ↑ GOX1968          | hypothetical protein | 3.6 | 496 |
| ↑ GOX1969          | alcohol dehydrogenase large subunit | 3.5 | 540 |
| ↓ GOX1970          | GTP-binding protein EngA | 4.3 | 408 |
| ↑ GOX2094          | sorbitol dehydrogenase cytochrome c subunit | 5.8 | 10 |
| ↑ GOX2095          | sorbitol dehydrogenase large subunit | 5.3 | 13 |
| ↑ GOX2096          | sorbitol dehydrogenase large subunit | 4.1 | 51 |
| ↑ GOX2097          | sorbitol dehydrogenase small subunit | 3.5 | 74 |
Fig. 4 (See legend on next page.)
the rpi transcript (GOX1708, 4.4 min) encoding ribose-5-phosphate isomerase, the transcript of GOX2218 also annotated to encode a ribose-5-phosphate isomerase, yet without strong sequence similarity to the GOX1708 protein, exhibited a 3-fold longer half-life (13.3 min). The two EDP gene transcripts exhibited a half-life of 5 min (edd) and 8.3 min (eda). Genes of the incomplete TCA cycle exhibited the lowest apparent expression in the central metabolism based on the FPKM values, with the aconitate transcript (GOX1335) as the most stable (9.3 min) above the global mean, and with the transcripts of transcript (GOX1316 and GOX1464 encode for sequence-identical α, pyruvate dehydrogenase E1 component alpha subunit (GOX2289); aceE, pyruvate dehydrogenase E1 component beta subunit (GOX2290); aceF, dihydrolipoamide acetyltransferase component of pyruvate dehydrogenase (GOX2291); acn, acacitate hydratase (GOX1335); ald, aldehyde dehydrogenase (GOX2018); eda, KDPG aldolase (GOX0430); edd, phosphogluconate dehydrogenase (GOX0431); eno, enolase (GOX2279); fba, fructose-bisphosphate aldolase (GOX0780); ftp, fructose 1,6-bisphosphatase (GOX1516); frk, fructokinase (GOX0284); furc, fumarate hydratase (GOX1643); gap, glyceraldehyde-3-phosphate dehydrogenase (GOX0508); gdhM, membrane-bound glucose dehydrogenase (GOX0265); gdhS, soluble glucose dehydrogenase (GOX2015); gik, glucokinase (GOX2419); gldt, citrate synthase (GOX1182); gnd, 6-phosphogluconate dehydrogenase (GOX1705); gmk, gluconokinase (GOX1709); gno, gluconate 5-dehydrogenase (GOX2187); gpg, phosphoglyceromutase (GOX3030); icd, isocitrate dehydrogenase (GOX1336); lpd, dihydrolipoamide dehydrogenase (GOX2292); mqa, malate:quinone oxidoreductase (GOX2070); ndhA, 2-oxoglutarate dehydrogenase E1 component (GOX0882); ndhB, dihydrolipoamide succinyltransferase E2 component of 2-oxoglutarate dehydrogenase complex (GOX1073); pdc, pyruvate decarboxylase (GOX1081); pgk, phosphoglycerate kinase (GOX0507); pgl, 6-phosphogluconolactonase (GOX1707); ppc, phosphoenolpyruvate carboxylase (GOX1022); pyk, pyruvate kinase (GOX2250); rpe, ribulosephosphate 3-epimerase (GOX1352); rpi, ribose 5-phosphate isomerase (GOX1708); sda, polyol dehydrogenase subunit Sda (GOX0854); sdb, polyol dehydrogenase subunit Sdb (GOX0855); tal, tpgi, bifunctional transaldolase (GOX1704); tx, transketolase (GOX1703); tpi, triosephosphate isomerase (GOX2217); zwf, glucose-6-phosphate-1-dehydrogenase (GOX0145). Metabolites: 1,3-BPG, 1,3-bisphosphoglycerate; 2-KFU, 2-ketofructose; 2-OG, 2-oxoglutarate; 2PGA, 2-phosphoglycerate; 3PGA, 3-phosphoglycerate; 5-KGLCNT, 5-ketogluconate; 6PG, 6-phosphogluconate; 6PGK, 6-phosphogluconolactonate; ACD, aconitate hydratase; ACE, acetate; AcCoA, acetyl coenzyme A; CIT, citrate; CoA, coenzyme A; DHAP, dihydroxyacetone phosphate; E4P, erythrose-4-phosphate; F6P, fructose 6-phosphate; FBP, fructose 1,6-bisphosphate; FRU, fructose; FUM, fumarate; G-1,5-LC, glucono-1,5-lactone; G6P, glucose 6-phosphate; GAP, glyceraldehyde 3-phosphate; GLC, glucose; GLCNG, glucuronate; ICT, isocitrate; KDPG, 2-keto-3-deoxy-6-phosphogluconate; MAL, malate; MAN, mannitol; OAA, oxaloacetate; PEP, phosphoenolpyruvate; PYR, pyruvate; RSP, ribose 5-phosphates; RUSP, ribulose 5-phosphate; S7P, sedoheptulose 7-phosphate; SucCoA, succinyl coenzyme A; XSP, xylulose 5-phosphate.

Mapping analysis revealed fragmentation regions in 235 rRNAs from G. oxydans

During the global mRNA decay analysis the visualization results of total RNA samples indicated that the full-length 23S rRNA from G. oxydans was underrepresented and appeared very unstable compared to the 16S rRNA (Fig. 1a). In the genome of G. oxydans 621H four mRNA operons are present [11]. In these operons, GOX0224, GOX1156, GOX1316 and GOX1464 encode for sequence-identical 16S rRNA transcripts with 1478 nt. GOX0221 and GOX1467 encode sequence-identical 23S rRNA transcripts with 2710 nt. The C at position 2119 is absent in the 23S rRNA transcript of GOX1159 with 2709 nt. At position 1746 a C is inserted in the 23S rRNA transcript of GOX1139 with 2711 nt. The underrepresentation of processed full-length 23S rRNA in G. oxydans RNA samples always raised questions on the quality of the prepared RNA, for example in DNA microarray studies. To check whether the 23S rRNA in G. oxydans maybe indeed rapidly degraded or rather is fragmented as found in several bacteria including α-proteobacteria other than Gluconobacter [39, 40], we analyzed the rRNA fraction obtained from G. oxydans ribosomes. Therefore, we chromatographically enriched ribosomes from G. oxydans cells grown in mannitol medium without rifampicin. We analyzed two time points, one in the exponential growth phase (OD600 = 1 after 4.25 h of cultivation) and one in the early stationary phase (OD600 = 2.5 after 9.5 h of cultivation). Cell pellets were used to obtain crude protein extracts. The sample volumes were adjusted to obtain the protein concentration suitable for the chromatography runs on the monolithic disks. According to the elution
profile we always obtained four major peaks termed P1 to P4 (Additional file 1: Figure S1A). The sum of the protein content in the elution fractions of the four peaks typically comprised 85% to 95% of the total protein applied to the column (Additional file 1: Table S4). According to nanoLC-based mass spectrometry, 933 G. oxydans proteins were identified overall in the four protein fractions (Additional file 1: Table S5). 449 to 865 proteins were identified in peaks P1 and P2, with the apparent overall content of ribosomal proteins being rather low. The highest abundance of ribosomal proteins was found in peak P3 as judged by SDS-PAGE analysis and MALDI-ToF mass spectrometry (data not shown) as well as by the overall high numbers of detected peptides per ribosomal protein in the nanoLC-MS/MS analysis (Additional file 1: Table S6). Moreover, by far the largest amount of RNA could be isolated from the protein fraction of peak P3 (Additional file 1: Table S4), which is expected when ribosomes are enriched according to the method used. All 30S ribosomal proteins were detected in peak P3 (Additional file 1: Table S6). From the 50S ribosomal proteins, the protein L32 (GOX0117), L36 (GOX0732), and L34P (GOX1825) could not be detected at all. Overall, the chromatogram and elution profile of ribosomes in peak P3 is typical for this method as already described elsewhere for other bacteria [31].

In the RNA isolated from peak P3 a mature full-length 23S rRNA could not be detected in the formaldehyde agarose gel analysis at ~ 2710 nt, yet the RNA samples exhibited a specific pattern of smaller fragments (Additional file 1: Figure S1B). This suggested that the 23S rRNA is fragmented in G. oxydans ribosomes. The pattern showed three fragments at approximately 1500 nt, 900 nt and 800 nt, and two shorter fragments at approximately 300 nt and 400 nt. By size the longest fragment of ~ 1500 nt corresponds to the 16S rRNA. To narrow down the regions of fragmentation in the 23S RNA transcripts, the ribosome-associated RNA was sequenced via next-generation sequencing and the paired-end reads were mapped to the sequences of the 16S and 23S RNA gene loci from G. oxydans using very stringent mapping parameters. To identify possible fragmentation positions based on the mapping coverage we searched for regions with less than 5% coverage of the average coverage for the entire gene locus (Fig. 5). In the RNA samples from the exponential phase, three regions with such a low or almost no coverage were found for the 23S rRNA transcripts suggesting fragmentation sites in GOX0221, GOX1319, and GOX1467. The same three regions were also found in the shortest 23S rRNA (GOX1159; 2709 nt), but in this case a fourth potential fragmentation region was present according to the very low mapping coverage (Fig. 5b). The three very low coverage regions found for GOX1159 were at positions 1140–1159 (20 nt), 1456–1465 (10 nt), and 1717–1770 (55 nt) (Fig. 5, Additional file 1: Tables S7-S10). For the other three 23S rRNA genes, the positions of two of the three regions differed by maximal two nucleotides: GOX0221 (1139–1160; 1456–1465; 1716–1770), GOX1319 (1140–1159; 1456–1465; 1715–1771), GOX1467 (1140–1159; 1456–1465; 1717–1770). The fourth region only found for GOX1159 was at position 2022–2136 (Additional file 1: Table S8). In this fourth region GOX1159 exhibits two nucleotide differences compared to the other three 23S rRNA genes at nt 2034 and nt 2119. The absence of mapping coverage in this region raises the question whether

![Fig. 5](#) Graphical representation of the 23S rRNA (a, b) and 16S rRNA (c) mapping coverage. The mapping is based on the Illumina reads obtained with RNA samples isolated from enriched ribosomes (peak P3) from cells in the exponential growth phase. The y axes are zoomed to a maximum of 150 to better illustrate the very low coverage regions (< 5%) indicated by the arrows. a Mapping coverage of the 23S rRNA locus GOX1319 (2711 nt). The same coverage pattern was obtained for the 23S rRNA loci GOX0221 and GOX1467 (2710 nt). b Mapping coverage of the 23S rRNA locus GOX1159 (2709 nt) which indicates an additional fragmentation site. c Mapping coverage obtained for the 16S rRNA locus GOX1156. The same coverage pattern was observed for the other 16S rRNA loci GOX0224, GOX1316, and GOX1464.
GOX1159 was actually transcribed into RNA. It should be mentioned that GOX1159 exhibits another nucleotide difference at nt 936 (T). According to the mapped reads 871-fold (Additional file 1: Table S8) and 8301-fold (Additional file 1: Table S16) coverage/reads were obtained at nt 936 in GOX1159. This shows the presence of the T and suggests that GOX1159 was very likely transcribed into RNA. For the 16S rRNA genes we did not find such low coverage regions indicating the presence of the mature full-length transcripts in the ribosomes (Fig. 5c, Additional file 1: Tables S11-S14). In the RNA samples from the stationary phase, the Illumina read mappings showed very similar results (Additional file 1: Tables S15-S22). Only for the second and fourth fragmentation position an extension of the low coverage regions by 55 nt (1401–1465) and by 31 nt (1991–2136) was observed.

Comparison of G. oxydans 23S rRNA fragmentation with other 23S rRNAs
Fragmentation of 23S rRNA and the presence of intervening sequences (IVSs) in 23S rRNAs were found in several bacteria [39]. We compared the 23S rRNAs GOX1319 and GOX1159 of G. oxydans with selected 23S rRNA sequences from such bacteria and also included E. coli where the 23S rRNA is not fragmented and IVSs are absent (Additional file 1: Figure S2). For Salmonella typhimurium [41], Rhodobacter sphaeroides, Bradyrhizobium japonicum, Rhodopseudomonas palustris [40, 42], Rhizobium leguminosarum, and Agrobacterium radiobacter [43], it was shown that fragmentation of 23S rRNA occurs by RNA cleavage to remove IVSs. For example, according to the literature IVSs can be found at the positions 131–168, 543–550, and 1176 relative to the E. coli sequence (Additional file 1: Figure S2). Close to the first very low coverage region in the mapping for G. oxydans an IVS is present in S. typhimurium and R. sphaeroides. These regions only partially overlap with the relevant G. oxydans region and the sequence similarity is low (Additional file 1: Figure S2). For all bacteria shown here and independent of fragmentation, the sequences are quite diverse in this region. For A. radiobacter and R. leguminosarum fragmentation without the presence of IVS was observed close to position 1500. Close to this region we also observed fragmentation of 23S rRNA in G. oxydans (Additional file 1: Figure S2). Sequence similarities among the three bacteria in this region are approximately 70%. The third fragmentation position close to 1750 appears to be present only in G. oxydans 23S rRNA (Additional file 1: Figure S2). Sequence similarities between rRNAs with and without fragmentation in this region are quite high, yet there are a few differences in G. oxydans compared to the other bacteria. The fourth region with a very low coverage was found only for GOX1159 and not for the other three 23S rRNA copies of G. oxydans. In this region GOX1159 differs only at the nucleotide positions 2034 (C for T) and 2119 (the C is absent in GOX1159) compared to the three other 23S rRNAs in G. oxydans. This suggests that these positions in the transcript of GOX1159 are specifically relevant for binding and/or cleavage by the processing RNase, likely RNase III as described for others [40, 44]. This fourth fragmentation position is also not known from the other bacteria (Additional file 1: Figure S2). Altogether, no known IVS region described in the literature were found in the 23S rRNAs from G. oxydans.

**Discussion**
In this study, we estimated the mRNA half-lives in G. oxydans 621H on a global scale and detected and analyzed fragmentation of the 23S rRNA transcripts. The mRNA decay plays an important role in the metabolism of nucleic acids in both prokaryotic and eukaryotic cells and also affects fluctuations in protein synthesis and growth [45, 46]. The mRNA abundance in cells is determined by its synthesis and degradation rates. For E. coli and the archaeeon S. solfataricus an inverse relationship between transcript abundance and half-lives was reported [15, 19, 21, 47, 48]. We also observed such an inverse relationship for G. oxydans. Many highly abundant transcripts are among the least stable and transcripts with longer half-lives are less abundant. This also supports the view that overall mRNA stability does not play a major role to obtain high mRNA abundance and that a rapid mRNA turnover facilitates fast adaptation to environmental changes. For example, transcripts of highly expressed genes essential for growth can be rapidly degraded in the case of a cell cycle arrest [19]. Nevertheless, as in other studies, not all transcripts follow the inverse relationship between abundance and half-life. For example, the molecular chaperones GroES and GroEL transcripts exhibited high abundance as well as long half-lives in G. oxydans. GroES and GroEL are required for proper folding of many proteins [49]. As their activity is required almost independent of environmental conditions, in such cases it is probably advantageous for the cell to support high transcript levels also by long half-lives. Similarly, ATP-proton motive force interconversion is a cellular function that actually needs to be maintained under many conditions. However, in G. oxydans this functional category exhibited by far the shortest mRNA half-lives among all categories. With a mean half-life of 3 min the mRNAs of the H+-ATP synthase were among the transcripts with the shortest half-lives. The Na⁺-ATP synthase with a mean RNA half-life of 5.1 min exhibited only very low transcript abundance compared to the H+-ATP synthase (80-fold lower). Comparative DNA microarray analysis of the effects of
oxygen limitation in *G. oxydans* revealed an approximately 2-fold decreased expression of the H⁺-ATP synthase and 2- to 3-fold increased expression of the Na⁺-ATP synthase [34]. This expression pattern and the abundance of their transcripts suggest that the assumed Na⁺-ATP synthase might play an important role under oxygen limitation and that the H⁺-ATP synthase with the very short mRNA half-lives is the major one under many other conditions. Actually, one would expect that genes of the energy metabolism including ATP synthases are among the medium or most stable transcripts due to their classification as housekeeping genes [50]. Indeed, for *E. coli* the mRNA half-lives of genes associated with energy metabolism (mean 6.3 min) rank in the top 3 categories with the longest half-lives among all groups reported with mean values ranging from 3.8 min to 6.4 min [15]. The *E. coli* ATP synthase transcripts exhibited a mean half-life of 6.2 min in M9 medium, which is twice as long compared to *G. oxydans* (3 min). This 2-fold difference in mRNA half-lives is even more remarkable when also considering the inverse and almost 2-fold difference in the typical doubling time of *E. coli* (~60 to 70 min) and *G. oxydans* (~100 to 110 min). Thus, the H⁺-dependent ATP synthase could represent a bottleneck in *G. oxydans* under certain conditions, for example in strains engineered for improved growth [13, 14, 51].

The average mRNA half-lives of genes involved in central carbon metabolism did not differ significantly from the global mean (5.7 min). This is attributed to partially high variation of half-lives within functional groups. There is also partially high variation of half-lives associated with the central carbon metabolism in *E. coli* [15], while in *S. solfatarius* transcripts of central metabolic pathways generally showed shorter half-lives [19]. In *S. cerevisiae* transcripts encoding the enzymes that participate in central metabolism are characteristically among those with the longest half-lives [48]. Thus, a general trend is not obvious and the mRNA half-lives in the central metabolism appear to be a species-specific characteristic.

Another species-specific characteristic is fragmentation of 23S rRNA and the absence or presence of segments termed intervening sequence (IVS). While this phenomenon is already known and studied in several bacteria including α-proteobacteria [39, 40], it was not yet reported for *Glactobacter*. In the γ-proteobacterium *S. typhimurium* fragmentation occurs by RNase III-dependent excision of IVS elements [41, 52]. In α-proteobacteria, for example *Rhizobiales*, *Bradyrhizobiales*, and *Rhodobacteriales*, processing of an IVS located close to the 5′-end of the 23S rRNA likely also involves RNase III in an early step. The multistep process generates a fragment of approximately 130 nt from the 5′-end, which is separated by the IVS from the main segment of the 23S rRNA [40, 42, 43]. According to our mapping coverage analysis and sequence comparisons with other 23S rRNAs, no IVS element close to the 5′-end were found, whose excision could possibly lead to a 130 nt fragment. Three potential fragmentation regions were found in all four 23S rRNAs and a fourth fragmentation position only in the 23S rRNA transcript of GOX1159. The potential fragmentation region at nt 1456–1465 is within the same fragmentation region of 23S rRNA from *A. radiobacter* and *R. leguminosarum*. For the other three fragmentation regions no fragmentation was reported in the 23S rRNAs of other α-proteobacteria for which fragmentations at other sites were reported. One of the three is found at nt 1139–1160 within a region without conservation, where also one IVS is present in the 23S rRNAs of *S. typhimurium* and *R. sphaeroides*. The sequences of this region exhibit no or only partial similarity to the region in *G. oxydans*. The third fragmentation region at nt 1716–1770 is in a highly conserved region according to the alignment, yet a fragmentation in other bacteria was not reported. This may be attributed to single nucleotide differences in this conserved region that can enable or disable cleavage by RNase activity, as can be seen at the fourth fragmentation position found only in GOX1159. In GOX1159 this region contains two specific single nucleotide variations compared to the other three 23S rRNA sequences from *G. oxydans*, which may be responsible for recognition and cleavage by RNase activity. Since these variations are located within the fragmentation region and not at their ends or in the flanking regions, the results suggest further rRNA processing of the fragments after the cleavage. Indeed, such secondary maturation steps which further process 23S rRNA 5′- and 3′-ends after RNase III cleavage were already suggested for α-proteobacteria [40].

Generally, fragmentation of 23S rRNA appeared to be phenotypically silent and processing of IVS elements was not required for the production of functional ribosomes [52]. Due to missing phenotypes there is still no convincing explanation for the physiological role of 23S rRNA fragmentation. For *Yersinia enterocolitica* and *S. typhimurium* it was suggested that removing the IVS elements may protect the resulting 23S rRNA fragments from unknown bacteriocins in the gut [53]. For *S. typhimurium* it was also suggested that fragmentation may provide an advantage in stationary phase to quickly adapt to the rapidly changing growth environment experienced in animal hosts [54]. A more general role for rRNA fragmentation was speculated recently in a study of the mammal naked mole-rat in the context of translational fidelity [55]. Naked mole-rat fibroblasts exhibited significantly increased translational fidelity while having comparable translation rates with mouse fibroblasts. In the
naked mole-rat the 28S rRNA is also fragmented while in mouse not. It was speculated that rRNA fragmentation may change the folding or dynamics of the large ribosomal subunit, altering the rate of GTP hydrolysis and/or interaction of the large subunit with tRNA during accommodation, thus finally affecting the fidelity of protein synthesis. It will be interesting to see if the biological function of IVS elements is indeed related to translational fidelity of ribosomes by fragmentation of 23S or 28S rRNAs.

Conclusion
In conclusion, the mRNA decay data from *G. oxydans* showed many similarities to the results obtained in other bacteria and also exhibited some specific differences. Overall, the slightly inverse relationship of transcript abundance and mRNA stability supports the view that mRNA stability does not play a general role to obtain high mRNA abundance. Rather, a quick mRNA turnover for fast adaptation of the cell to environmental changes is made possible by the shorter half-lives of highly expressed genes. The very short mRNA half-lives of the H⁺-ATP synthase, which is likely responsible for the ATP-proton motive force interconversion in *G. oxydans* under many or most conditions according to expression data, is notably in contrast to mRNA decay data from other bacteria. Together with the short mRNA half-lives and the relatively low expression of some genes of the incomplete TCA cycle, which exhibited the lowest expression values in the central carbon metabolism, these could be bottlenecks in *G. oxydans* at some point. This should be considered in future metabolic engineering approaches to further improve growth and biomass yield of *G. oxydans*. The consequences of 23S rRNA fragmentation on growth and fitness of *G. oxydans* are unknown. Our Illumina sequencing and mapping analysis of the RNA fragments isolated from enriched ribosomes revealed three potential fragmentation regions which were previously not known from 23S rRNA fragmentation studies in other bacteria. Further studies are needed to unravel the multistep process of fragmentation in the 23S rRNAs of *G. oxydans*. Although fragmentation of 23S rRNAs appears to be phenotypically silent, it could be relevant under some conditions or for some aspect, for example the fidelity during protein synthesis.

Additional file

**Additional file 1**: Figure S1. Chromatogram of ribosome enrichment and rRNA obtained. Figure S2; 23S rRNA sequence alignment. Table S1. mRNA decay data. Table S2. FPKM expression values and mRNA half-lives. Table S3. mRNA half-lives of operons and monocistronic transcripts. Table S4. Amounts of protein and RNA in peak P1, P2, P3, P4. Table S5. Proteins identified in chromatographic elution fractions. Table S6. Ribosomal proteins with trypsin peptides in P1, P2, P3, P4. Tables S7-S14. rRNA mapping coverage in early stationary phase. (ZIP 220 kb)

**Abbreviations**
PPP: Pentose phosphate pathway; TCA: Tricarboxylic acid; EDP: Entner-Doudoroff pathway; mDH: Membrane-bound dehydrogenase; IVS: Intervening sequence.

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**Availability of data and materials**
The microarray data are accessible in NCBI’s Gene Expression Omnibus through accession number GSE103428. The RNAseq data are publicly available in the European Nucleotide Archive under accession number PRJEB18739.

**Authors’ contributions**
AX performed RNA sequencing and analyzed the RNAseq data. AS and TP performed mRNA decay data processing and GEO upload. UD carried out cell cultivations and DNA microarray experiments. SM developed the protocol for ribosome enrichment. UD and SM purified ribosomes. AMK performed LC-MS/MS analysis. TP designed and supervised the study. AK, MB, and TP analyzed the data and wrote the manuscript. All authors read and approved the final manuscript.

**Ethics approval and consent to participate**
Ethics approval and consent to participate are not applicable because no humans or their data, and no animals, were used in this study. The source of the wild type *Gluconobacter oxydans* 621H strain (DSM 2343) used in this study was the German Collection of Microorganisms and Cell Cultures (DSMZ, Braunschweig, Germany).

**Consent for publication**
Not applicable.

**Competing interests**
The authors declare that they have no competing interests.

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