



Relevance of NADH Dehydrogenase and Alternative Two-Enzyme Systems for Growth of *Corynebacterium glutamicum* With Glucose, Lactate, and Acetate

Tomoya Maeda^{†*}, Abigail Koch-Koerfges[†] and Michael Bott^{*}

IBG-1: Biotechnology, Institute of Bio- and Geosciences, Forschungszentrum Jülich, Jülich, Germany

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*Correspondence:

Michael Bott
m.bott@fz-juelich.de

[†]These authors have contributed
equally to this work

*Present address:

Tomoya Maeda,
RIKEN Center for Biosystems
Dynamics Research, Osaka, Japan

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The oxidation of NADH with the concomitant reduction of a quinone is a crucial step in the metabolism of respiring cells. In this study, we analyzed the relevance of three different NADH oxidation systems in the actinobacterial model organism *Corynebacterium glutamicum* by characterizing defined mutants lacking the non-proton-pumping NADH dehydrogenase Ndh (Δndh) and/or one of the alternative NADH-oxidizing enzymes, L-lactate dehydrogenase LdhA ($\Delta ldhA$) and malate dehydrogenase Mdh (Δmdh). Together with the menaquinone-dependent L-lactate dehydrogenase LldD and malate:quinone oxidoreductase Mqo, the LdhA-LldD and Mdh-Mqo couples can functionally replace Ndh activity. In glucose minimal medium the Δndh mutant, but not the $\Delta ldhA$ and Δmdh strains, showed reduced growth and a lowered $NAD^+/NADH$ ratio, in line with Ndh being the major enzyme for NADH oxidation. Growth of the double mutants $\Delta ndh\Delta mdh$ and $\Delta ndh\Delta ldhA$, but not of strain $\Delta mdh\Delta ldhA$, in glucose medium was stronger impaired than that of the Δndh mutant, supporting an active role of the alternative Mdh-Mqo and LdhA-LldD systems in NADH oxidation and menaquinone reduction. In L-lactate minimal medium the Δndh mutant grew better than the wild type, probably due to a higher activity of the menaquinone-dependent L-lactate dehydrogenase LldD. The $\Delta ndh\Delta mdh$ mutant failed to grow in L-lactate medium and acetate medium. Growth with L-lactate could be restored by additional deletion of *sugR*, suggesting that *ldhA* repression by the transcriptional regulator SugR prevented growth on L-lactate medium. Attempts to construct a $\Delta ndh\Delta mdh\Delta ldhA$ triple mutant were not successful, suggesting that Ndh, Mdh and LdhA cannot be replaced by other NADH-oxidizing enzymes in *C. glutamicum*.

Keywords: NADH dehydrogenase, malate dehydrogenase, malate:quinone oxidoreductase, lactate dehydrogenase, $NAD^+/NADH$ ratio, respiratory chain, SugR

INTRODUCTION

Corynebacterium glutamicum is a Gram-positive, non-pathogenic soil bacterium that was isolated in a screen for microorganisms which excrete the flavor enhancer L-glutamate (Kinoshita et al., 1957). It is used in industry for the biotechnological production of amino acids and proteins (Eggeling and Bott, 2015; Freudl, 2017). Furthermore, strains for efficient production of many

other metabolites have been constructed, such as organic acids (Wieschalka et al., 2013), diamines (Wendisch et al., 2018), and various other compounds (Becker and Wittmann, 2012). The majority of the corresponding production processes are performed under aerobic conditions and require a functional respiratory chain (Bott and Niebisch, 2003; Matsushita, 2013). Several dehydrogenases oxidize their substrates with concomitant reduction of menaquinone, the only respiratory quinone in *corynebacteria*. Finally, electron transfer from menaquinol to oxygen is catalyzed either by a cytochrome *bc₁-aa₃* supercomplex (Niebisch and Bott, 2003; Graf et al., 2016; Kao et al., 2016) or by cytochrome *bd* oxidase (Kusumoto et al., 2000; Kabus et al., 2007).

As the majority of reducing equivalents from substrate oxidation are initially transferred to NAD⁺, a sufficient rate of NADH reoxidation is crucial to sustain carbon flux in central metabolism. *C. glutamicum* possesses a non-proton-pumping, single-subunit NADH dehydrogenase Ndh (Nantapong et al., 2005), but neither a proton-pumping complex I-type NADH dehydrogenase (Parey et al., 2020) nor a sodium-ion-pumping Nqr-type NADH dehydrogenase (Steuber et al., 2014). Analysis of a Δndh mutant of *C. glutamicum* revealed slightly decreased growth on glucose and acetate minimal agar plates and almost no growth defect in glucose minimal medium (Molenaar et al., 2000; Nantapong et al., 2004). Two alternative systems were proposed that can couple the oxidation of NADH to menaquinone reduction and thus compensate the lack of Ndh (Molenaar et al., 2000; Nantapong et al., 2004). These are the NAD⁺-dependent malate dehydrogenase Mdh (Genda et al., 2003) acting in concert with the membrane-associated malate:quinone oxidoreductase Mqo, and the NAD⁺-dependent L-lactate dehydrogenase LdhA acting in concert with the membrane-associated L-lactate dehydrogenase LldD (Figure 1).

The net reaction of the Mdh-Mqo couple equals that of an Ndh and it could serve as an alternative “NADH dehydrogenase,” as Mdh reduces oxaloacetate with NADH to L-malate ($\Delta G^{0'} = -28.9$ kJ/mol), and the membrane-associated Mqo subsequently re-oxidizes L-malate back to oxaloacetate and reduces MK ($\Delta G^{0'} = -18.5$ kJ/mol). It was shown that Mqo is crucial for oxidation of malate to oxaloacetate during aerobic growth of *C. glutamicum* (Molenaar et al., 2000). Mdh was able to complement the absence of Mqo activity only when the Δmqo mutant was supplied with nicotinamide in order to increase the concentrations of NAD⁺ and NADH (Molenaar et al., 2000). Under these conditions, the NAD⁺/NADH ratio decreased from 4.4 in the absence of nicotinamide to 2.8 in the presence of 1 mg L⁻¹ nicotinamide, which presumably enabled the thermodynamically unfavorable oxidation of malate by Mdh (Molenaar et al., 2000). Similar to the Mdh-Mqo couple, the net reaction of the LdhA-LldD couple also corresponds to that of Ndh, as LdhA reduces pyruvate with NADH to L-lactate ($\Delta G^{0'} = -25.1$ kJ/mol) and LldD re-oxidizes L-lactate to pyruvate and reduces MK ($\Delta G^{0'} = -22.4$ kJ/mol) (Nantapong et al., 2004).

The Mdh-Mqo and LdhA-LldD couples might function as overflow systems that come into play when Ndh activity is insufficient. For example, an F₁F₀-ATP synthase mutant with 25% residual ATPase activity showed enhanced glucose

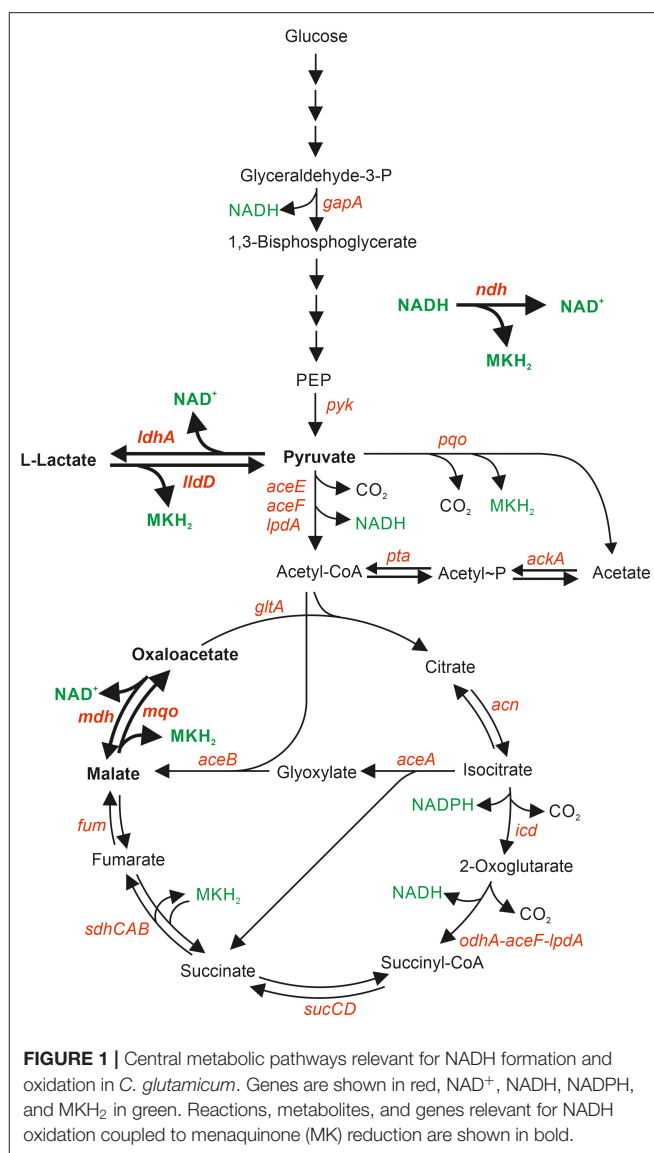


FIGURE 1 | Central metabolic pathways relevant for NADH formation and oxidation in *C. glutamicum*. Genes are shown in red, NAD⁺, NADH, NADPH, and MKH₂ in green. Reactions, metabolites, and genes relevant for NADH oxidation coupled to menaquinone (MK) reduction are shown in bold.

consumption and respiration rates (Sekine et al., 2001), requiring increased NADH oxidation rates. A proteome comparison of this ATPase mutant with the wild type revealed that the amounts of Mqo and Mdh were increased (Li et al., 2007). The authors suggested that the Mdh-Mqo couple enabled the increased NADH oxidation rates in the ATP synthase mutant (Sawada et al., 2012). A ΔF_1F_0 mutant of *C. glutamicum* completely devoid of ΔF_1F_0 -ATP synthase activity also showed increased glucose consumption and respiration rates and both the mRNA level and the protein level of Mdh were increased (Koch-Koerfges et al., 2012).

In this study, we aimed at getting a better understanding of the role of the different NADH oxidation systems for growth and metabolism of *C. glutamicum*. For this purpose, we constructed and characterized six deletion mutants that lack one or two of the NADH oxidation systems. The phenotype of the different mutant

strains strongly depended on the carbon source. In our study, we tested glucose, acetate, and L-lactate. The results indicate that NADH oxidation is predominantly catalyzed by Ndh, but that the Mdh-Mqo and LdhA-LldD couples are also involved and increase metabolic flexibility.

MATERIALS AND METHODS

Bacterial Strains and Culture Conditions

C. glutamicum strains and plasmids used in this work are listed in **Table 1**. For the analysis of growth, organic acid production, carbon source consumption, oxygen consumption, enzyme activities, and NAD⁺/NADH ratios, a 5 ml preculture in brain-heart infusion (BHI) medium was inoculated with colonies from a fresh agar plate (BHI agar) and incubated for 8 h at 30°C and 170 rpm. Cells from the preculture were transferred into 20 ml CGXII minimal medium (Keilhauer et al., 1993) containing either 4% (wt/vol) glucose (1,071 mM C), or 2–4% (wt/vol) sodium L-lactate (536–1,071 mM C), or 2–4% (wt/vol) sodium acetate (488–975 mM C) as carbon and energy source. The cultures were incubated for 16 h at 30°C and 130 rpm. After washing the cells with 0.9% (wt/vol) NaCl, the main culture with 50 ml CGXII minimal medium containing 4% (wt/vol) of the desired carbon source was inoculated to give an optical density at 600 nm (OD₆₀₀) of 1. The CGXII medium was always supplemented with 30 mg/l 3,4-dihydroxybenzoic acid as iron chelator (Frunzke et al., 2008). The main cultivations were performed in baffled 500 ml Erlenmeyer flasks containing a septum for sterile sampling of the cultures at 30°C and 130 rpm. Cells were harvested during the exponential growth phase at an OD₆₀₀ of 4–6 for further analysis. *E. coli* DH5 α , which was used as host for cloning, was cultivated in LB medium or on LB agar plates at 37°C. When appropriate, kanamycin was used at a concentration of 25 μ g/ml (*C. glutamicum*) or 50 μ g/ml (*E. coli*).

Construction of pK19mobsacB Plasmids and Deletion Mutants

In-frame deletion mutants of *C. glutamicum* ATCC 13032 were constructed as described previously (Niebisch and Bott, 2001). For this purpose, the *ndh* or *mdh* upstream region, including the first seven to ten codons of each gene, and the downstream region, including the last seven codons of each gene, were amplified with the Expand High Fidelity kit (Roche Diagnostics, Mannheim, Germany) using the oligonucleotide pairs Δ XXX-1-for/ Δ XXX-2-rev and Δ XXX-3-for/ Δ XXX-4-rev, respectively. The “XXX” stands for the name of the gene to be deleted and is specified in **Table 2**, which lists the oligonucleotides used in this study. The resulting PCR products of about 500 bp were subsequently fused by overlap-extension PCR to products of $\sim$ 1,000 bp. After digestion with *Xma*I and *Xba*I, these fragments were cloned into pK19mobsacB (Schäfer et al., 1994), cut with the same restriction enzymes, to yield pK19mobsacB- Δ ndh and pK19mobsacB- Δ mdh, respectively. DNA sequence analysis revealed that the cloned PCR products contained no unwanted mutations. Subsequently, the plasmids were transferred by electroporation (van der Rest et al., 1999) into *C. glutamicum* strain ATCC 13032 and the transformation

mixture was plated on a BHIS agar plate containing 25 μ g kanamycin/ml. After selection for the first and second recombination event, kanamycin-sensitive and sucrose-resistant clones were analyzed by colony-PCR with the primer pair Δ XXX-fw/ Δ XXX-rev in order to distinguish between wild type and Δ ndh or Δ mdh clones, respectively. For the construction of the double deletion strains lacking two NADH oxidation systems, namely Δ ndh Δ mdh, Δ ndh Δ ldhA, and Δ mdh Δ ldhA, pK19mobsacB- Δ mdh or pK19mobsacB- Δ ldhA were transferred by electroporation into competent Δ ndh or Δ mdh cells and selected and analyzed as described above. For the construction of the triple deletion strain Δ ndh Δ mdh Δ sugR, pK19mobsacB- Δ sugR was transferred by electroporation into competent Δ ndh Δ mdh cells and selected and analyzed as described above.

Construction of pAN6-Based Expression Plasmids

The genes *ldhA*, *mdh*, and *ndh* were amplified from chromosomal DNA of *C. glutamicum* ATCC 13032 with the gene-specific oligonucleotides XXX_F and XXX_R (**Table 2**). The former one introduced an *Nde*I restriction site including the start codon, the latter one an *Nhe*I restriction site. The resulting PCR products, with a size of 960 bp for *ldhA*, 1,001 bp for *mdh*, and 1,418 bp for *ndh*, were cut with *Nde*I and *Nhe*I and cloned separately into the expression plasmid pAN6 (Frunzke et al., 2008) cut with the same enzymes. The correctness of the cloned DNA fragments was confirmed by DNA sequencing. Subsequently, the plasmids were transferred by electroporation into *C. glutamicum* wild type and mutants and the transformation mixture was plated on a BHIS agar plate containing 25 μ g/mL⁻¹ kanamycin. Enzymes used for DNA restriction, ligation or dephosphorylation were obtained either from Roche Diagnostics (Mannheim, Germany) or New England Biolabs (Frankfurt am Main, Germany). Plasmid DNA from *E. coli* or *C. glutamicum* was isolated with the QIAprep Spin Miniprep kit according to the manufacturer's instructions (Qiagen, Hilden, Germany).

Determination of Growth Parameters and Consumption Rates of Glucose, Organic Acids, and Oxygen

Growth was followed by measuring the optical density at 600 nm (OD₆₀₀) with an Ultrospec 500-pro spectrophotometer (Amersham Biotech). The biomass concentration was calculated from OD₆₀₀ values using an experimentally determined correlation factor of 0.25 g_{CDW} L⁻¹ for OD₆₀₀ = 1 (Kabus et al., 2007). Quantitative determination of glucose and organic acids in culture supernatants, and calculation of carbon source uptake rates was carried out as described (Koch-Koerfges et al., 2012).

Oxygen uptake rates (OUR) of non-growing cells were measured with a Clarke-type oxygen electrode using a thermostatically controlled, magnetically stirred 2-ml chamber at 30°C (Oxygraph, Hansatech Instruments, Germany) as described previously (Koch-Koerfges et al., 2013). The cells were harvested during exponential growth in shake flasks and oxygen consumption was followed using a cell suspension with an OD₆₀₀ of 0.5–2.5 in the chamber. The protein content

TABLE 1 | Strains and plasmids used in this study.

Strain or plasmid	Relevant characteristic(s)	Source or references
STRAINS		
<i>C. glutamicum</i> ATCC 13032	Biotin-auxotrophic wild type	Kinoshita et al., 1957
$\Delta ldhA$	13032 derivative with an in-frame deletion of <i>ldhA</i> (cg3219)	This study
Δmdh	13032 derivative with an in-frame deletion of <i>mdh</i> (cg2613)	This study
Δndh	13032 derivative with an in-frame deletion of <i>ndh</i> (cg1656)	This study
$\Delta mdh \Delta ldhA$	13032 derivative with in-frame deletions of <i>mdh</i> and <i>ldhA</i>	This study
$\Delta ndh \Delta mdh$	13032 derivative with in-frame deletions of <i>ndh</i> and <i>mdh</i>	This study
$\Delta ndh \Delta ldhA$	13032 derivative with in-frame deletions of <i>ndh</i> and <i>ldhA</i>	This study
$\Delta ndh \Delta mdh \Delta sugR$	13032 derivative with in-frame deletions of <i>ndh</i> , <i>mdh</i> , and <i>sugR</i> (cg2115)	This study
<i>E. coli</i> DH5 α	F ⁻ Φ 80dlac Δ (<i>lacZ</i>)M15 Δ (<i>lacZYA-argF</i>) U169 <i>endA1recA1 hsdR17</i> (<i>r</i> _K ⁻ , <i>m</i> _K ⁺) <i>deoR thi-1 phoA supE44</i> λ ⁻ <i>gyrA96 relA1</i>	Invitrogen (Karlsruhe, Germany)
PLASMIDS		
pK19mobsacB	Km ^r , <i>oriV_{E.coli}</i> <i>oriT</i> <i>sacB</i> integration vector for allelic exchange in <i>C. glutamicum</i>	Schäfer et al., 1994
pK19mobsacB- $\Delta ldhA$	Kan ^R , pK19mobsacB derivative containing an overlap- extension PCR product covering the up- and downstream regions of <i>ldhA</i>	Litsanov et al., 2012
pK19mobsacB- Δmdh	Kan ^R , pK19mobsacB derivative containing an overlap- extension PCR product covering the up- and downstream regions of <i>mdh</i>	This study
pK19mobsacB- Δndh	Kan ^R , pK19mobsacB derivative containing an overlap- extension PCR product covering the up- and downstream regions of <i>ndh</i>	This study
pK19mobsacB- $\Delta sugR$	Kan ^R , pK19mobsacB derivative containing an overlap- extension PCR product covering the up- and downstream regions of <i>sugR</i>	Engels and Wendisch, 2007
pAN6	Km ^R ; <i>C. glutamicum</i> / <i>E. coli</i> shuttle vector for regulated gene expression (<i>P_{lac}</i> , <i>lacI^q</i> , pBL1 <i>oriV_{C.glutamicum}</i> , pUC18 <i>oriV_{E.coli}</i>)	Frunzke et al., 2008
pAN6- <i>ldhA</i>	Km ^R ; pAN6 derivative containing the gene <i>ldhA</i>	This study
pAN6- <i>mdh</i>	Km ^R ; pAN6 derivative containing the gene <i>mdh</i>	This study
pAN6- <i>ndh</i>	Km ^R ; pAN6 derivative containing the gene <i>ndh</i>	This study

was calculated by assuming that 50% of the cell dry weight corresponds to protein.

Preparation of Cell Fractions for Enzymatic Assays

C. glutamicum cells were harvested by centrifugation (10,000 g, 4°C, 5 min) during the exponential growth phase and washed twice with 20 mM potassium phosphate buffer (PPB), pH 7.5. One ml of the cell suspension was mixed with 1 g zirconia/silica beads (0.1 mm diameter; Biospec, Bartlesville, USA) in a 2-ml Eppendorf tube and the cells were mechanically disrupted by three 30 s shakings in a Silamat S5 (Ivoclar Vivadent, Ellwangen, Germany). Cell debris and unbroken cells were separated by centrifugation for 5 min at 16,000 g and 4°C. The supernatant was then ultracentrifuged at 171,000 g for 60 min at 4°C. The resulting supernatant containing the soluble proteins and the resuspended sedimented membrane fraction (in 20 mM PPB pH 7.5) were used for the assays.

Enzyme Activity of Membrane-Bound Enzymes

LldD and Mqo activities were measured spectrophotometrically at 25°C by following the change in the absorbance of 2,6-dichlorophenolindophenol (DCPIP) at 600 nm (Molenaar et al., 2000; Nantapong et al., 2004). The reaction mixture contained

100 μ g membrane protein, 0.2 mM DCPIP, 0.4 mM phenazine methosulfate (PMS), 1 mM NaN₃, 1 mM substrate (L-lactate or L-malate for LldD and Mqo, respectively), and 50 mM PPB, pH 7.0, in a total volume of 1 ml. The amount of enzyme reducing 1 μ mol/min DCPIP was defined as 1 unit, using an extinction coefficient of 22 mM⁻¹ cm⁻¹. Ndh activity was determined as described previously (Molenaar et al., 2000). The reaction mixture contained 100 μ g membrane fraction, 0.4 mM DCPIP, 1 mM NaN₃, 0.4 mM NADH, and 50 mM PPB, pH 6.5, in a total volume of 1 ml. Before adding the substrate, the reaction mixtures were incubated for 5 min at 25°C. Using a reaction mixture with buffer instead of membrane protein as a control, the decrease in absorption at 340 nm was measured and subtracted from each measurement to calculate the net value. The activity unit was defined as 1 μ mol NADH oxidized per min, using an extinction coefficient of 6.3 mM⁻¹ cm⁻¹.

Enzyme Activity of Cytoplasmic Enzymes

LdhA and Mdh activities were determined spectrophotometrically at 25°C by following the decrease in the absorbance of NADH at 340 nm. The reaction mixture contained 100 μ g soluble protein fraction, 0.2 mM NADH, 1 mM pyruvate or oxaloacetate (for Ldh or Mdh, respectively), and 50 mM PPB (pH 7.0 or pH 7.5, for the Ldh or Mdh activity assay, respectively) in a total volume of 1 ml. The activity unit was defined as 1 μ mol/min NADH oxidized, using an extinction

TABLE 2 | Oligonucleotides used in this study.

Oligonucleotide	Sequence	Application
Δmdh -1-for	5'-TAATCTAGACGCTTGGACATGCCAGATGCCTT-3'	Amplification of <i>mdh</i> upstream region, includes XmaI restriction site
Δmdh -2-rev	5'-CCCCGTAACATAAATCTAAACAGACGTTCTGCGGGGAATTCAT-3'	Amplification of <i>mdh</i> upstream region
Δmdh -3-for	5'-TGTTTAAGTTTAGTTACGGGGGCAGTGCAGCTTGTCTAA-3'	Amplification of <i>mdh</i> downstream region
Δmdh -4-rev	5'-TAACCGGGGCTTGATAAATCCACGCTGGG-3'	Amplification of <i>mdh</i> downstream region, includes XbaI restriction site
Δmdh -fw	5'-CCTTCTTATCGCCAAAGTGA-3'	Control of <i>mdh</i> deletion via colony PCR
Δmdh -rev	5'-GCGGGTCGATTCCACG-3'	Control of <i>mdh</i> deletion via colony PCR
<i>mdh</i> -F	5'-TGGCATATGAATCCCCGAGAACGTC-3'	Amplification of <i>mdh</i> for expression plasmid pAN6- <i>mdh</i> , includes NdeI restriction site
<i>mdh</i> -R	5'-GTGCTAGCTTAGAGCAAGTCGCGCACTGCC-3'	Amplification of <i>mdh</i> for expression plasmid pAN6- <i>mdh</i> , includes NheI restriction site
Δndh -1-for	5'-TAATCTAGAACCCAGGCCACTCTTCC-3'	Amplification of <i>ndh</i> upstream region, includes XmaI restriction site
Δndh -2-rev	5'-CCCCATCCACTAAACTTAAACAGCCTTCGGGGCGGGTTGGTT-3'	Amplification of <i>ndh</i> upstream region
Δndh -3-for	5'-TGTTTAAGTTTAGTTACGGGGCAGCGTTTCAGCGGAAAGTAA-3'	Amplification of <i>ndh</i> downstream region
Δndh -4-rev	5'-TAACCGGGGCAGAAAGAGTCCCGCATTGA-3'	Amplification of <i>ndh</i> downstream region, includes XbaI restriction site
Δndh -fw	5'-CCAGCAAACGCTAGGTTGGG-3'	Control of <i>ndh</i> deletion via colony PCR
Δndh -rev	5'-TGCACCCCTCAATGGCCTTGGT-3'	Control of <i>ndh</i> deletion via colony PCR
<i>ndh</i> -F	5'-GCGCATATGTCAGTTAACCCAAACC-3'	Amplification of <i>ndh</i> for expression plasmid pAN6- <i>ndh</i> , includes NdeI restriction site
<i>ndh</i> -R	5'-GCGCTAGCTTACTTTCCGCTGAAACG-3'	Amplification of <i>ndh</i> for expression plasmid pAN6- <i>ndh</i> , includes NheI restriction site
$\Delta ldhA$ -fw	5'-GCACCAAGTTCGATGTGGGTGG-3'	Control of <i>ldhA</i> deletion via colony PCR
$\Delta ldhA$ -rev	5'-CGTTGTCGATCATCTGCTTCCAG-3'	Control of <i>ldhA</i> deletion via colony PCR
<i>ldhA</i> -F	5'-AAACATATGAAAGAAACCGTCGTAACAAGATTG-3'	Amplification of <i>ldhA</i> for expression plasmid pAN6- <i>ldhA</i> , includes NdeI restriction site
<i>ldhA</i> -R	5'-AAAGCTAGCTTAGAAGAACTGCTTCTGAATTCGCGCAG-3'	Amplification of <i>ldhA</i> for expression plasmid pAN6- <i>ldhA</i> , includes NheI restriction site

Restriction sites are underlined, other functional regions are italic.

coefficient of $6.3 \text{ mM}^{-1} \text{ cm}^{-1}$. All enzyme activity results for cells grown on glucose, acetate or L-lactate minimal medium were performed in triplicates.

Determination of Intracellular NAD^+/NADH Ratios

The concentrations of the pyridine nucleotides NADH and NAD^+ were determined with the EnzyChrom™ NAD^+/NADH assay kit (BioAssay Systems, Hayward, USA) following the manufacturer's instructions and as described previously (Siedler et al., 2011). Cells in the exponential growth phase were cooled with ice water, harvested by centrifugation (10,000 g, 4°C, 5 min), and washed twice with cold phosphate-buffered saline (PBS), pH 7.5. The sedimented cells were resuspended either in acid extraction buffer or in base extraction buffer (both part of the kit) to extract oxidized pyridine nucleotides or reduced pyridine nucleotides, respectively. The concentrations of NAD^+ and NADH in each extract were then quantified by the lactate dehydrogenase cycling reaction, in which the formed NADH reduces a formazan (MTT) reagent.

RESULTS

Growth of the *C. glutamicum* Strains Δndh , Δmdh , and $\Delta ldhA$ on Glucose and Acetate

In order to determine the relevance of NADH dehydrogenase Ndh, malate dehydrogenase Mdh and L-lactate dehydrogenase LdhA for aerobic growth of *C. glutamicum*, we characterized the deletion mutants Δndh , Δmdh , and $\Delta ldhA$ during cultivation in shake flasks with glucose, L-lactate, or acetate, as carbon source. These substrates were chosen as their oxidation is linked to different numbers of NADH-forming reactions. Glucose oxidation to 6 CO_2 involves three NADH-forming reactions [glyceraldehyde 3-phosphate dehydrogenase, pyruvate dehydrogenase complex, and unusual 2-oxoglutarate dehydrogenase complex (Niebisch et al., 2006; Hoffelder et al., 2010; Bruch et al., 2020; Kinugawa et al., 2020)] and is coupled to the formation of 6 NADH, 2 NADPH, and 4 MQH_2 if the oxidative pentose phosphate pathway is neglected. L-Lactate oxidation to 3 CO_2 involves two NADH-forming reactions (pyruvate dehydrogenase complex, and 2-oxoglutarate dehydrogenase complex) and leads to the formation of 2 NADH, 1 NADPH, and 3 MQH_2 . Acetate oxidation to 2

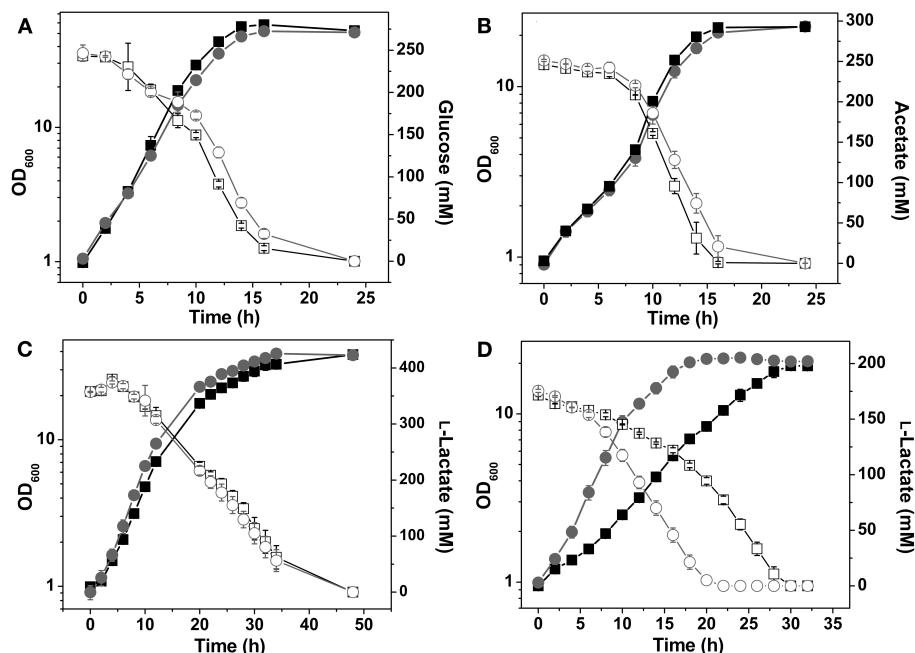


FIGURE 2 | Growth and carbon source consumption of *C. glutamicum* wild type (■) and the Δndh mutant (●). Cells were grown in CGXII minimal medium containing 4% (wt/vol) glucose (A), 2% (wt/vol) acetate (B), 4% (wt/vol) L-lactate (C), or 2% (wt/vol) L-lactate (D). In (C) cells were precultivated in L-lactate minimal medium, in (D) cells were precultivated in glucose minimal medium. The concentrations of the individual carbon sources in the culture supernatant are shown by open squares (□) for the wild type and by open circles (○) for the Δndh mutant. Average values from at least three independent cultivations and standard deviations are shown.

CO₂ involves only one NADH-forming reaction (2-oxoglutarate dehydrogenase complex) and is linked to the formation of 1 NADH, 1 NADPH, and 2 MQH₂ (Figure 1).

In glucose minimal medium the Δndh mutant showed an 18% decreased growth rate and a 12% reduced biomass formation compared to the wild type (Figure 2A, Table 3). NADH dehydrogenase activity was not detectable in extracts of the Δndh mutant, whereas it was high (2.4 U mg⁻¹) in the wild type (Table 4). The Δndh mutant showed a 28% decreased glucose uptake rate (GUR, Table 3) and a 27% decreased oxygen uptake rate compared to the wild type (OUR, 185 ± 13 vs. 254 ± 6 nmol O₂ mg_{protein}⁻¹ min⁻¹), suggesting that glycolytic flux and consequently also respiration were reduced due to an insufficient rate of NADH oxidation in the absence of Ndh. In agreement with this assumption, the NAD⁺/NADH ratio was reduced by more than 50% in the Δndh mutant compared to the wild type (2.0 vs. 4.7). Furthermore, the Δndh mutant secreted 22% more lactate than the wild type (Table 3). Pyruvate reduction to lactate by LdhA represents an alternative path for NADH oxidation. The enzyme activities of LdhA, Mdh, and Mqo were unchanged in the Δndh mutant, whereas LldD activity was increased 3.3-fold (Table 4).

In acetate minimal medium, the growth rate of the Δndh mutant was slightly decreased by 7% (Figure 2B), the acetate uptake rate (AUR) was reduced by 13%, and the biomass yield was unchanged compared to the wild type (Table 3). The Mdh and Mqo activities were reduced by about 20% (Table 4). The

NAD⁺/NADH ratio during growth on acetate was reduced by 18% in the Δndh mutant compared to the wild type (2.3 vs. 2.8 in the wild type, Table 4). It is noticeable that the NAD⁺/NADH ratio of the wild type on acetate was 40% lower than on glucose.

In contrast to the Δndh strain, growth of the Δmdh and $\Delta ldhA$ mutants on glucose or acetate was comparable to that of the wild type and no changes of the NAD⁺/NADH ratio and of any enzyme activity tested except for the one deleted were observed (data not shown).

Growth of the Δndh Mutant on L-lactate

When cultivated in CGXII medium with L-lactate as sole carbon source, the Δndh mutant grew slightly faster than the wild type ($\mu = 0.22$ vs. 0.19 h⁻¹), whereas biomass formation was unchanged (Figure 2C). While the wild type required pre-cultivation in L-lactate medium to reach optimal growth rates on L-lactate, the Δndh mutant showed optimal growth irrespective of the carbon source used in the preculture (Figures 2C,D). The growth rate of non-adapted wild-type cells on L-lactate ($\mu = 0.13$ h⁻¹) was 34% lower compared to that of L-lactate-adapted wild-type cells ($\mu = 0.19$ h⁻¹). The improved growth of the Δndh mutant on L-lactate could be reversed by expression of *ndh* using the expression plasmid pAN6-*ndh* (data not shown). This indicates that the absence of Ndh has a positive influence on L-lactate utilization. The NAD⁺/NADH ratio on lactate was 2.8 for the wild type, comparable to acetate-grown cells, and 2.1 for the Δndh mutant. Similar to glucose-grown cells, the LldD activity was 1.6-fold increased in a Δndh mutant grown in L-lactate

TABLE 3 | Growth rate, biomass formation (max. OD600), carbon source uptake rates, and maximal organic acid formation of the *C. glutamicum* wild type, and its Δndh , $\Delta mdh\Delta ldhA$, $\Delta ndh\Delta mdh$ and $\Delta ndh\Delta ldhA$ mutants.

Carbon source	Strain	μ (h ⁻¹)	Max. OD ₆₀₀	Carbon source uptake rates (mmol min ⁻¹ mg _{CDW} ⁻¹)	Maximal organic acid formation (mM)			
					L-Lactate	Acetate	Succinate	Pyruvate
Glucose	WT	0.39 ± 0.01	62.2 ± 0.9	88 ± 9 (528 ± 54) ^a	96 ± 5	24 ± 2	4.3 ± 0.5	0.4 ± 0.4
	Δndh	0.32 ± 0.01	50.9 ± 1.5	64 ± 2 (384 ± 12)	117 ± 4	28 ± 11	1.9 ± 0.3	0.5 ± 0.3
	$\Delta mdh\Delta ldhA$	0.36 ± 0.01	49.3 ± 0.9	76 ± 4 (456 ± 24)	N.D. ^b	12 ± 1.8	6.4 ± 2.8	4.1 ± 0.3
	$\Delta ndh\Delta ldhA$	0.25 ± 0.01	37.4 ± 2.6	53 ± 2 (318 ± 12)	N.D.	85 ± 14	14 ± 3	11 ± 1
	$\Delta ndh\Delta mdh$	0.24 ± 0.01	52.9 ± 0.4	59 ± 3 (354 ± 18)	121 ± 3	5.7 ± 0.1	1.4 ± 0.3	2.5 ± 1.1
L-Lactate	WT	0.19 ± 0.01	36.6 ± 0.2	98 ± 5 (294 ± 15)	–	N.D.	0.5 ± 0.1	3.9 ± 0.7
	Δndh	0.22 ± 0.01	38.2 ± 0.2	115 ± 5 (345 ± 15)	–	N.D.	N.D.	0.8 ± 0.2
	$\Delta mdh\Delta ldhA$	0.19 ± 0.02	36.7 ± 1.0	105 ± 8 (315 ± 24)	–	N.D.	1.0 ± 0.2	2.7 ± 0.3
	$\Delta ndh\Delta ldhA$	0.18 ± 0.01	36.0 ± 0.4	99 ± 3 (297 ± 9)	–	N.D.	N.D.	15 ± 1
	$\Delta ndh\Delta mdh$	N.G. ^c ; M.A. ^d	–	95 ± 4 (285 ± 12)	–	N.D.	N.D.	21 ± 2
Acetate	WT	0.29 ± 0.01	23.1 ± 0.8	231 ± 4 (462 ± 8)	0.8 ± 0.1	–	N.D.	N.D.
	Δndh	0.27 ± 0.01	23.5 ± 0.4	200 ± 10 (400 ± 20)	0.8 ± 0.1	–	N.D.	N.D.
	$\Delta mdh\Delta ldhA$	0.28 ± 0.01	21.3 ± 0.5	230 ± 11 (460 ± 22)	N.D.	–	N.D.	N.D.
	$\Delta ndh\Delta ldhA$	0.26 ± 0.01	22.9 ± 0.3	192 ± 3 (384 ± 6)	N.D.	–	N.D.	N.D.
	$\Delta ndh\Delta mdh$	N.G.	–	–	–	–	–	–

Cells were cultivated in shake flasks as described in Materials and Methods. Mean values and standard deviations from three independent cultivations are shown.

^aNumbers in parenthesis give uptake rate as mmol C min⁻¹ mg_{CDW}⁻¹.

^bN.D., not detected.

^cN.G., no growth.

^dM.A., metabolically active.

medium and the oxygen consumption rate of the Δndh mutant with L-lactate as substrate was increased by 17% compared to the wild type (335 ± 17 vs. 285 ± 15 nmol O₂ mg_{Protein}⁻¹ min⁻¹). The increased LldD activity of the Δndh mutant is presumably responsible for the increased growth rate and the increased oxygen consumption rate. The wild type secreted up to 3.9 mM pyruvate during the early exponential phase, which was metabolized during mid and late exponential phase, whereas the Δndh mutant secreted below 1 mM pyruvate (Table 3).

Characteristics of a $\Delta mdh\Delta ldhA$ Double Mutant

To explore how the absence of more than one NADH oxidation system influences growth and other parameters, we constructed the *C. glutamicum* strains $\Delta mdh\Delta ldhA$, $\Delta ndh\Delta ldhA$, and $\Delta ndh\Delta mdh$. Attempts to construct the triple mutant $\Delta ndh\Delta mdh\Delta ldhA$ failed, suggesting that under the conditions tested no further efficient NADH oxidation system exists in *C. glutamicum* that can compensate the lack of Ndh, LdhA, and Mdh.

In glucose minimal medium, the $\Delta mdh\Delta ldhA$ mutant, which can oxidize NADH via Ndh, showed an 8% reduced growth rate, a slowed transition to the stationary phase, 21% reduced biomass formation, and a 14% reduced GUR (Figure 3A, Table 3). The slightly reduced growth rate might be due to the inability of the $\Delta mdh\Delta ldhA$ mutant to form L-lactate, which is transiently formed by the wild type as NADH oxidation product and subsequently consumed again (Koch-Koerfges et al., 2012). The

NAD⁺/NADH ratio (Table 4) of the $\Delta mdh\Delta ldhA$ mutant (2.5 ± 0.2) was lower than in the wild type (4.7 ± 0.3), but higher than in the Δndh mutant (2.0 ± 2.0), suggesting that the enzyme couples LdhA-LldD and/or Mdh-Mqo contribute to NADH oxidation. In minimal medium with L-lactate or acetate as carbon sources, growth of the $\Delta mdh\Delta ldhA$ mutant was comparable to that of the wild type (Figure 3), suggesting that both LdhA and Mdh are irrelevant for NADH oxidation on these substrates.

Characteristics of a $\Delta ndh\Delta ldhA$ Double Mutant

The $\Delta ndh\Delta ldhA$ mutant, which presumably can oxidize NADH only via the Mdh-Mqo couple, showed a stronger growth defect in glucose minimal medium than the Δndh mutant (Figure 3A). The growth rate was 0.25 ± 0.1 (Δndh mutant 0.32 ± 0.1) and the final OD₆₀₀ was 37.4 ± 2.6 (Δndh mutant 50.9 ± 1.5). In agreement with the reduced growth rate, the GUR was reduced by 17% compared to the Δndh mutant (53 ± 2 vs. 64 ± 2 nmol min⁻¹ mg_{CDW}⁻¹). These results suggest that the LdhA-LldD couple plays a role in NADH oxidation on glucose besides Ndh. As expected, due to the absence of LdhA, the $\Delta ndh\Delta ldhA$ mutant did not form L-lactate. Instead, the formation of acetate, succinate, and pyruvate strongly increased, with acetate being the major by-product (Table 3). This suggests that due to limited NADH reoxidation the pyruvate dehydrogenase activity and the TCA cycle activity is limited, causing an increased overflow metabolism. The NAD⁺/NADH ratio of the $\Delta ndh\Delta ldhA$ mutant

TABLE 4 | NAD⁺/NADH ratios and enzyme activities in *C. glutamicum* wild type and its Δndh , $\Delta mdh\Delta ldhA$, $\Delta ndh\Delta ldhA$, and $\Delta ndh\Delta mdh$ mutants.

Carbon source	Strain	NAD ⁺ /NADH ratio	Enzyme activity (Units/mg _{Protein})				
			LdhA	LldD	Mdh	Mqo	Ndh
Glucose	WT	4.7 ± 0.3	0.35 ± 0.01	0.03 ± 0.01	2.02 ± 0.24	0.10 ± 0.03	2.37 ± 0.29
	Δndh	2.0 ± 0.1	0.37 ± 0.01	0.10 ± 0.02	1.93 ± 0.11	0.10 ± 0.02	N.D. ^a
	Δmdh	4.7 ± 0.1	0.35 ± 0.01	0.02 ± <0.01	0.01 ± <0.01	0.10 ± 0.02	2.10 ± 0.10
	$\Delta ldhA$	5.4 ± 0.3	0.01 ± <0.01	0.01 ± <0.01	2.06 ± 0.12	0.13 ± 0.01	2.32 ± 0.21
	$\Delta mdh\Delta ldhA$	2.5 ± 0.2	0.01 ± 0.01	0.01 ± 0.01	0.01 ± 0.01	0.08 ± 0.04	2.46 ± 0.13
	$\Delta ndh\Delta ldhA$	1.8 ± 0.1	0.01 ± 0.01	0.01 ± 0.01	2.09 ± 0.20	0.10 ± 0.01	N.D.
	$\Delta ndh\Delta mdh$	1.3 ± 0.1	0.40 ± 0.01	0.31 ± 0.04	0.02 ± 0.01	0.14 ± 0.01	N.D.
L-Lactate	WT	2.8 ± 0.1	0.03 ± 0.01	0.13 ± 0.01	3.00 ± 0.26	0.12 ± 0.05	0.93 ± 0.17
	Δndh	2.1 ± 0.1	0.03 ± 0.01	0.21 ± 0.01	2.80 ± 0.27	0.11 ± 0.02	N.D.
	$\Delta mdh\Delta ldhA$	2.4 ± 0.1	0.01 ± 0.01	0.17 ± 0.01	0.01 ± 0.01	0.10 ± 0.02	0.96 ± 0.31
	$\Delta ndh\Delta ldhA$	2.2 ± 0.1	0.01 ± 0.01	0.15 ± 0.01	3.04 ± 0.22	0.12 ± 0.01	N.D.
	$\Delta ndh\Delta mdh$	N.G. ^b	N.G.	N.G.	N.G.	N.G.	N.G.
Acetate	WT	2.8 ± 0.1	0.03 ± 0.01	0.03 ± 0.01	2.69 ± 0.13	0.29 ± 0.02	1.41 ± 0.18
	Δndh	2.3 ± 0.3	0.03 ± 0.01	0.03 ± 0.01	2.10 ± 0.08	0.23 ± 0.03	N.D.
	$\Delta mdh\Delta ldhA$	2.4 ± 0.2	0.01 ± 0.01	0.01 ± 0.01	0.01 ± 0.01	0.24 ± 0.01	1.21 ± 0.27
	$\Delta ndh\Delta ldhA$	2.1 ± 0.3	0.01 ± 0.01	0.01 ± 0.01	2.88 ± 0.16	0.25 ± 0.03	N.D.
	$\Delta ndh\Delta mdh$	N.G.	N.G.	N.G.	N.G.	N.G.	N.G.

Cells were cultivated in shake flasks as described in Materials and Methods and harvested in the early exponential growth phase before the onset of oxygen limitation. Mean values and standard deviations from three independent cultivations are shown.

^aN.D., not detected.

^bN.G., no growth.

was 1.8 ± 0.1 and thus lower than the one of the Δndh mutant (2.0 ± 0.1).

The growth of the $\Delta ndh\Delta ldhA$ mutant in acetate minimal medium was comparable to that of the wild type (Figure 3C) and the Δndh mutant (Figure 2B) and therefore LdhA is not relevant for this substrate. Interestingly, growth of the $\Delta ndh\Delta ldhA$ mutant in L-lactate minimal medium was comparable to that of the wild type (Figure 3E). Thus, the additional deletion of *ldhA* abolished the positive growth effect of the *ndh* deletion on L-lactate utilization (see above). The positive influence of the *ndh* deletion on L-lactate correlated with an increased LldD activity, which was not observed in the $\Delta ndh\Delta ldhA$ mutant (Table 4). This indicates that the formation of L-lactate from pyruvate by LdhA is necessary to trigger increased *lldD* expression, most likely via relieve of *lldD* repression by the transcriptional regulator LldR (Gao et al., 2008; Georgi et al., 2008).

Characteristics of a $\Delta ndh\Delta mdh$ Double Mutant

In glucose minimal medium, the $\Delta ndh\Delta mdh$ mutant, which presumably can oxidize NADH only via the LdhA-LldD couple, showed a growth rate of 0.24 h^{-1} , a GUR of $59 \pm 3 \text{ nmol min}^{-1} \text{ mg}_{\text{CDW}}^{-1}$, and final OD₆₀₀ of 52.9 ± 0.4 , corresponding to reductions of 38, 40, and 33% compared to the wild type (Table 3). The $\Delta ndh\Delta mdh$ mutant secreted 26% more L-lactate than the wild type (Table 3), which supports the importance of LdhA for NADH oxidation. With a value of 1.3 ± 0.1 , the NAD⁺/NADH ratio of the $\Delta ndh\Delta mdh$ mutant was the lowest one observed in our study (Table 4). This suggests that NADH oxidation by the LdhA-LldD couple is less efficient than by Ndh or Mdh-Mqo, although the activities of LdhA and LldD were

highest in the $\Delta ndh\Delta mdh$ mutant with values of 0.40 and 0.31 U mg⁻¹, respectively (Table 4).

In contrast to the $\Delta mdh\Delta ldhA$ and $\Delta ndh\Delta ldhA$ double mutants, the $\Delta ndh\Delta mdh$ mutant was unable to grow with acetate or L-lactate (Figure 3). The growth defect on acetate and L-lactate could be complemented by plasmid-based expression of either *ndh* (pAN6-*ndh*) or *mdh* (pAN6-*mdh*) (data not shown). Whereas, acetate was not consumed at all by the $\Delta ndh\Delta mdh$ mutant (Figure 3D), a decrease of the L-lactate concentration was observed within the first 20 h of incubation (Figure 3F) and pyruvate was secreted (Table 3). This suggests that the cells possessed LldD activity, allowing oxidation of L-lactate to pyruvate with concomitant reduction of menaquinone to menaquinol, which was then reoxidized by the terminal oxidases. The observation that pyruvate was excreted rather than further oxidized is probably caused by an insufficient pyruvate dehydrogenase activity due to the limited availability of the cofactor NAD⁺. The phenotype of the $\Delta ndh\Delta mdh$ mutant suggests that NADH oxidation during growth on acetate and L-lactate requires either Ndh or the Mdh-Mqo couple, which cannot be replaced by the LdhA-LldD couple.

Restoration of Growth on L-lactate of the $\Delta ndh\Delta mdh$ Mutant

It was shown that expression of *ldhA* in *C. glutamicum* is tightly repressed by the transcriptional regulator SugR in the absence of sugars (Engels et al., 2008; Toyoda et al., 2009). As LdhA is probably the only catabolic NADH oxidation enzyme left in the $\Delta ndh\Delta mdh$ mutant, repression of *ldhA* by SugR might be responsible for the inhibited growth on L-lactate. To test this hypothesis we constructed a $\Delta ndh\Delta mdh\Delta sugR$ mutant, which

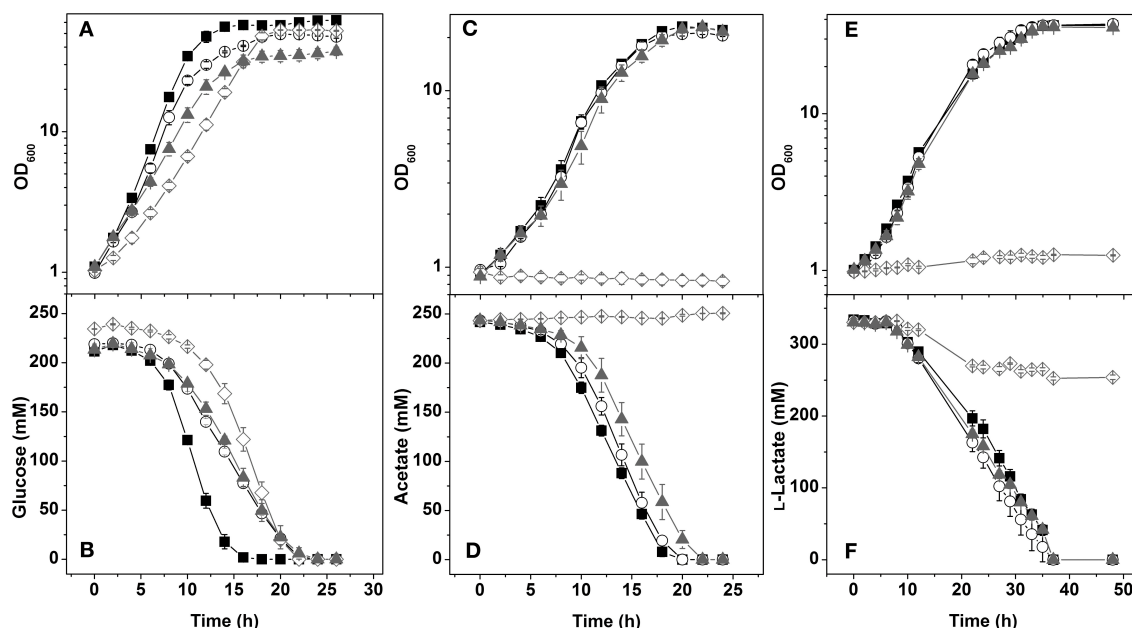


FIGURE 3 | Growth and carbon source consumption of the *C. glutamicum* wild type (■) and the mutants $\Delta mdh\Delta ldhA$ (○), $\Delta ndh\Delta ldhA$ (▲), and $\Delta ndh\Delta mdh$ (◇). Cells were cultivated in CGXII minimal medium containing either 4% (wt/vol) glucose (A,B), 2% (wt/vol) acetate (C,D), or 4% (wt/vol) L-lactate (E,F). Growth (A,C,E) and carbon source concentration in the supernatants (B,D,F) are shown. Average values from at least three independent cultivations and the standard deviations are shown.

indeed was able to grow on L-lactate as sole carbon and energy source, although at a much lower growth rate (Figure 4A). The triple mutant secreted high amounts of pyruvate and later also malate (Figure 4B). Growth of the $\Delta ndh\Delta mdh$ mutant in L-lactate minimal medium was also possible by plasmid-based expression of *ldhA* using plasmid pAN6-*ldhA* (data not shown). Thus, insufficient LdhA activity due to *ldhA* repression by SugR is an important reason for the growth defect of the $\Delta ndh\Delta mdh$ mutant in L-lactate medium. Neither the $\Delta ndh\Delta mdh\Delta sugR$ mutant nor the $\Delta ndh\Delta mdh$ mutant carrying pAN6-*ldhA* were able to grow on acetate (data not shown), indicating that there must be additional reasons for the inability to utilize acetate.

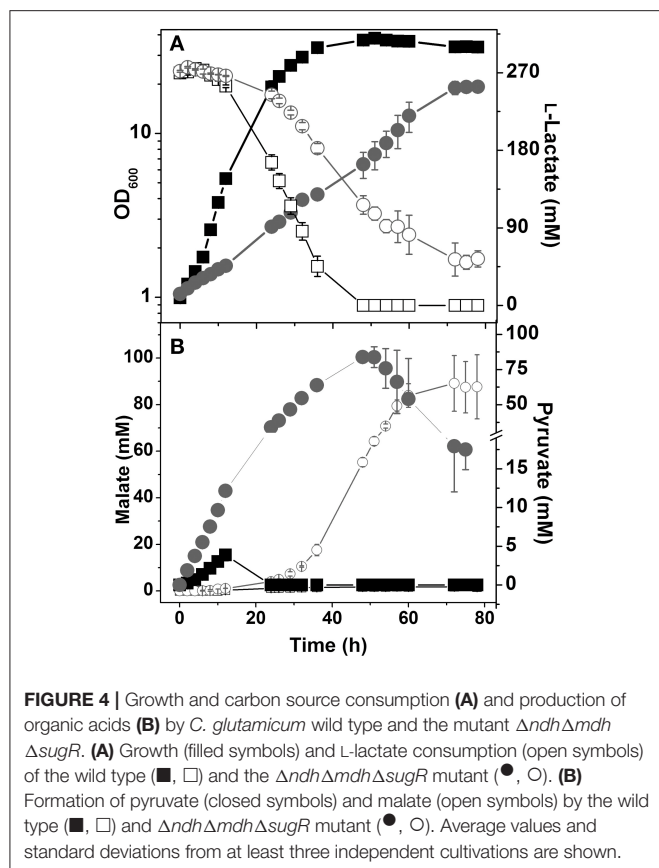
DISCUSSION

In the present work, we investigated a set of six mutants, namely Δndh , $\Delta ldhA$, Δmdh , $\Delta mdh\Delta ldhA$, $\Delta ndh\Delta ldhA$, and $\Delta ndh\Delta mdh$, that lack one or two of the three known catabolic NADH oxidation enzymes in *C. glutamicum* to explore their role for aerobic growth and metabolism on glucose, acetate, and L-lactate. As our attempts to construct the triple mutant $\Delta ndh\Delta mdh\Delta ldhA$ mutant failed, it appears likely that there is no further enzyme enabling catabolic NADH oxidation in *C. glutamicum* besides Ndh, Mdh, and LdhA.

In two previous studies, it was reported that an *ndh* integration mutant grows at a slightly lower rate than the wild type ATCC 13032 on glucose and acetate agar plates (Molenaar et al., 2000)

and that the growth rate of an *ndh* disruption mutant of the lysozyme-sensitive strain KY9714 in glucose minimal medium was not much altered compared to the parent strain (Nantapong et al., 2004). Our studies with the in-frame deletion mutant Δndh of strain ATCC 13032 revealed a slightly reduced growth rate and biomass formation in glucose minimal medium along with reduced glucose and oxygen consumption rates. A strong effect was observed for the $NAD^+/NADH$ ratio, which decreased from 4.7 in the wild type to 2.0 in the Δndh mutant. As growth and the $NAD^+/NADH$ ratio of the strains $\Delta ldhA$ and Δmdh in glucose minimal medium were comparable to the wild type (Table 4), our results show that Ndh is the most important enzyme for NADH oxidation during aerobic growth of *C. glutamicum* on glucose.

The observation that the *C. glutamicum* Δndh mutant did not show a stronger growth defect on glucose was explained by the existence of alternative NADH oxidation systems formed by the couples Mdh-Mqo and LdhA-LldD, which catalyze the same net reaction as Ndh, namely the oxidation of NADH and reduction of menaquinone when acting in concert (Molenaar et al., 2000; Nantapong et al., 2004). In line with this explanation, the $\Delta ndh\Delta mdh$ and $\Delta ndh\Delta ldhA$ double mutants showed stronger growth defects and a stronger reduction of the $NAD^+/NADH$ ratio than the Δndh mutant in glucose minimal medium. Therefore, neither the Mdh-Mqo couple alone nor the LdhA-LldD couple alone is able to compensate the lack of the other two NADH oxidation systems adequately. The $\Delta mdh\Delta ldhA$ mutant, which still can oxidize NADH via Ndh, grew better than the $\Delta ndh\Delta mdh$ and $\Delta ndh\Delta ldhA$ mutants, but not as good as the



wild type (Table 3), suggesting that the Mdh-Mqo and LdhA-LldD couples contribute to NADH oxidation also when Ndh is present.

Growth on L-lactate was improved by the *ndh* deletion (Figures 2C,D). This positive effect is presumably due to a 2-fold increased activity of LldD (Table 4), which was observed also previously in the study of the *ndh* mutant of strain KY9714 (Nantapong et al., 2005). It has been shown that expression of the *cg3226-ldd* operon is regulated by the FadR-type transcriptional regulator LldR, which represses the two genes in the absence of L-lactate (Gao et al., 2008; Georgi et al., 2008). The gene *cg3226* encodes a putative L-lactate permease, which is not essential for growth on L-lactate, however (Stansen et al., 2005). The reason for the increased LldD activity in the Δndh mutant is most likely the increased formation of L-lactate (Table 3), which relieves repression of the *cg3226-ldd* operon by LldR. In line with this explanation, the additional deletion of *ldhA* in the Δndh strain reversed the improved growth and the increased LldD activity (Figures 3C,D, Tables 2, 3). The reason why even L-lactate-adapted wild-type cells grew not as good as the Δndh mutant

(Figure 2C) is not clear yet. The absence of Ndh might allow for an increased availability of menaquinone as electron acceptor of LldD.

Compared to glucose, growth on acetate of the strains Δndh , $\Delta ndh\Delta ldhA$, and $\Delta mdh\Delta ldhA$ was only minimally affected (Figures 2, 3, Table 3). This difference can be explained by the fact that glucose oxidation is coupled to the formation of up to 6 NADH/glucose, while acetate oxidation is linked to the formation of only 1 NADH/acetate (Figure 1). Thus, glucose oxidation has a much higher NADH reoxidation demand than lactate or acetate oxidation. As essentially no L-lactate is formed during growth on acetate (Table 3) and *ldhA* is repressed by SugR, the LdhA-LldD cycle is probably not active and NADH oxidation is accomplished via the Mdh-Mqo cycle in the mutants Δndh and $\Delta ndh\Delta ldhA$ and via Ndh in strain $\Delta mdh\Delta ldhA$. In agreement with this interpretation, the $\Delta ndh\Delta mdh$ strain was unable to grow on acetate (Figure 3).

In summary, our results show that *C. glutamicum* uses a set of three systems for NADH oxidation coupled to menaquinone reduction. Ndh is the dominant enzyme, but the Mdh-Mqo couple and LdhA-LldD couples complement or substitute it and thereby enable NADH oxidation in situations with high catabolic fluxes or a defective Ndh enzyme. Such backup systems exist not only for NADH oxidation, but also for menaquinol oxidation, where cytochrome *bd* oxidase can substitute or complement the cytochrome *bc₁-aa₃* supercomplex. This redundancy likely contributes to the metabolic flexibility and the survival of the organism in nature.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/supplementary material, further inquiries can be directed to the corresponding author/s.

AUTHOR CONTRIBUTIONS

MB and AK-K designed the study. TM and AK-K performed the experiments and prepared the figures and tables. MB wrote the manuscript. All authors contributed to the interpretation of the data.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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