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Flexibility of Syntrophic Enzyme Systems in *Desulfovibrio* Species Ensures Their Adaptation Capability to Environmental Changes

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The mineralization of organic matter in anoxic environments relies on the cooperative activities of hydrogen producers and consumers obligately linked by interspecies metabolite exchange in syntrophic consortia that may include sulfate reducing species such as *Desulfovibrio*. To evaluate the metabolic flexibility of syntrophic *Desulfovibrio* to adapt to naturally fluctuating methanogenic environments, we studied *Desulfovibrio alaskensis* strain G20 grown in chemostats under respiratory and syntrophic conditions with alternative methanogenic partners, *Methanococcus maripaludis* and *Methanospirillum hungatei*, at different growth rates. Comparative whole-genome transcriptional analyses, complemented by G20 mutant strain growth experiments and physiological data, revealed a significant influence of both energy source availability (as controlled by dilution rate) and methanogen on the electron transfer systems, ratios of interspecies electron carriers, energy generating systems, and interspecies physical associations. A total of 68 genes were commonly differentially expressed under syntrophic versus respiratory lifestyle. Under low-energy (low-growth-rate) conditions, strain G20 further had the capacity to adapt to the metabolism of its methanogenic partners, as shown by its differing gene expression of enzymes involved in the direct metabolic interactions (e.g., periplasmic hydrogenases) and the ratio shift in electron carriers used for interspecies metabolite exchange (hydrogen/formate). A putative monomeric [Fe-Fe] hydrogenase and Hmc (high-molecular-weight-cytochrome *c*₃) complex-linked reverse menaquinone (MQ) redox loop become increasingly important for the reoxidation of the lactate/pyruvate oxidation-derived redox pair, DsrC_{red} and Fd_{red}, relative to the Qmo-MQ-Qrc (quinone-interacting membrane-bound oxidoreductase; quinone-reducing complex) loop. Together, these data underscore the high enzymatic and metabolic adaptive flexibility that likely sustains *Desulfovibrio* in naturally fluctuating methanogenic environments.

In anoxic environments depleted in inorganic electron acceptors (e.g., freshwater and marine sediments, flooded soils, landfills and sewage digestors) the complete mineralization of complex organic matter to CO₂ and methane relies on the cooperative activities of phylogenetically and metabolically distinct microbial groups assembled in syntrophic consortia (1–3). In these assemblages, sulfate-reducing bacteria (SRB) such as *Desulfovibrio* species function as secondary fermenters obligately linked via interspecies electron transfer to the metabolic activity of methanogenic archaea, since the oxidation of common substrates (organic acids and alcohols) only yields sufficient energy for cell maintenance and growth when the methanogens maintain low concentrations of the SRB metabolites (1–3). Hydrogen and formate are considered to be the primary shuttle compounds for interspecies electron transfer in methanogenic consortia (1–3). Exclusive use of H₂ as an electron carrier has been proven for several syntrophs, while a combined formate/H₂ electron shuttling has been confirmed in propionate- and butyrate-degrading cocultures (2, 4–10). Since syntrophic communities exist in conditions close to thermodynamic equilibrium and metabolite exchange is considered to occur primarily via diffusion (1–4, 11), intermicrobial distances influence the efficiency of interspecies transfer (especially for H₂), which affects the growth and biodegradation rates of these consortia (2, 6, 11–13). To ensure close proximity, cell aggregates of high complexity are often formed (e.g., sludge granules in anaerobic reactors) (11–13) with flagellar proteins having a putative role in triggering syntrophic interaction (14). In a previous comparative study of two *Desulfovibrio* species (*Desulfovibrio alaskensis* strain G20 and *Desulfovibrio vulgaris* strain Hildenborough) growing in lactate-based continuous coculture with *Methanococ-*

cus maripaludis, we demonstrated that their various syntrophic growth characteristics are genetically and metabolically based on the involvement of different electron-transferring enzyme systems and interspecies electron carriers used for syntrophic growth with the same methanogenic partner (15). Faster growth of the strain G20 pairing was associated with two factors: (i) a kinetic advantage conferred by electron transfer through a redox loop consisting of a quinone-interacting membrane-bound oxidoreductase, Qmo (16), and a quinone-reducing complex, Qrc (17), that permits higher lactate oxidation rates at elevated H₂ levels, thereby enhancing growth of *M. maripaludis*, and (ii) use of formate as the main electron exchange mediator (>70% electron flux) via periplasmic formate dehydrogenases (as opposed to the exclusively H₂-based exchange used by strain Hildenborough). Thus, *Desulfovibrio* species have evolved different adaptive strategies for growth in environments that fluctuate in electron acceptor availability, as shown by the absence of a conserved pool of genes that are essential to sustain syntrophy (15).

Differences in adaptive strategy may also in part be associated with the syntrophic partner. At low substrate levels, the specific

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growth rate of mutualistic cultures were shown to depend upon the hydrogen affinity of the methanogen, as given by the substrate saturation constant (K_s) for H_2 and its maximum specific growth rate in axenic culture on hydrogen (18). At low H_2 concentrations, methanogenesis becomes fully coupled to growth yield (19, 20). The importance of the methanogenic partner was indicated by our initial batch culture analyses, showing that each *Desulfovibrio* species varied significantly in its growth characteristics (growth rate, yield, and population structure) when syntrophically paired with different hydrogenotrophic methanogens (B. Meyer, unpublished results). Thus, these physiological differences among methanogens, which may in part be determined by variation in electron transfer complexes coupled to the use of either H_2 or formate at low concentrations (21–24), are expected to influence the metabolism of their syntrophic partner and, by inference, the fitness of the pairing.

Several model organisms (*Syntrophobacter*, *Syntrophomonas* [3–5, 9, 10], and *Desulfovibrio* [25] species) have been investigated to develop an understanding of the biochemistry and genetics of syntrophic growth. However, with only two notable exceptions (26, 27), these studies have been based almost exclusively on batch culture growth. However, the complex dynamics of growth in batch culture complicates the resolution of processes related to growth rate from those of syntrophic association. Thus, in the present study we used chemostats to examine both the impact of energy source availability (controlled by dilution rate) and the methanogenic partner on the syntrophic metabolism and energetics of *Desulfovibrio* species: *D. alaskensis* strain G20 was grown in sulfate-limited monoculture, as well as in coculture, at different growth rates (0.047 and 0.027 h^{-1}) and with two different methanogenic partners, *Methanococcus maripaludis* and *Methanospirillum hungatei*, in chemostats. Comparative whole-genome transcriptional analyses, complemented by G20 mutant strain growth experiments and physiological data, revealed distinctive differences in the electron transfer systems, ratios of interspecies electron carriers, and energy conservation mechanisms associated with growth rate and methanogenic partner. Together, these data underscore the high enzymatic and metabolic adaptive flexibility that likely sustains *Desulfovibrio* in naturally fluctuating methanogenic environments.

MATERIALS AND METHODS

Microorganisms. The wild-type strains *D. alaskensis* G20, *M. maripaludis* S2 (28, 29), and *M. hungatei* JF-1 (30, 31) used to establish the initial cocultures were obtained from the American Type Culture Collection (ATCC). A total of 26 *D. alaskensis* G20 transposon mutant strains were investigated in comparative phenotypic growth experiments (Table 1; for a detailed list of the mutant strains used in the present study, see Table S1 in the supplemental material).

Mono- and coculture medium composition for chemostat and batch cultures. The basal salt solution of the media was described earlier (26) and contained 30 mM sodium D,L-lactate or sodium pyruvate, resazurin as a redox indicator, and 10 mM Na_2SO_4 (chemostat monoculture media) or 30 mM Na_2SO_4 (batch monoculture media). The media were prepared in bioreactor glass vessels (2 liters), medium reservoir bottles (10 liters), and Widdel flasks (1 and 2 liters) anoxically under an N_2 - CO_2 (80:20, vol/vol) gas atmosphere according to the method of Widdel and Bak (32). The media were buffered using 1.1 mM KH_2PO_4 and 30 mM $NaHCO_3$. Nonchelated trace element mixture 1 and selenite-tungstate solution, as well as vitamin mixture, riboflavin, vitamin B_1 and B_{12} solutions and 1 M Na_2S solution (32), were added aseptically in 1-ml aliquots per liter of medium. All stock solutions were prepared under anoxic conditions as

described previously (32). For batch culture cultivation, the media were dispensed anoxically in sterile N_2 - CO_2 (80:20 [vol/vol]) pregassed 250-ml serum bottles or 25-ml Balch tubes in 130- and 13-ml aliquots, respectively. After preparation, the bioreactor and medium reservoir bottles were connected to a N_2 - CO_2 (90:10 [vol/vol]) gas mixture tank.

Chemostat operation conditions. A 0.5-ml inoculum from a glycerol stock of a previously grown coculture (*D. alaskensis* G20 paired either with *M. maripaludis* or *M. hungatei*, tenth transfer generation) or monoculture was used to inoculate 130 ml of medium in 250-ml serum bottles. Cultures were incubated at 37°C in a shaking incubator (300 rpm) until reaching an optical density at 600 nm (OD_{600}) of 0.3 or 0.26 ± 0.02 (for *M. hungatei* coculture pairing). A 100-ml inoculum was added to a 2.45 liter bioreactor vessel (FairMenTec 2.3 l model, Göttingen, Germany, and Electrolab FerMac 300 Series, Tewkesbury, United Kingdom) containing 2 liters of culture media. Three biological replicates of each coculture pairing and sulfate-limited *D. alaskensis* G20 monocultures were grown in chemostats. After inoculation, chemostats were initially run in batch mode at 37°C with stirring at 250 rpm. The headspace of the bioreactor was flushed through a sterile 0.2 micron syringe filter with N_2 - CO_2 (90:10, [vol/vol]) at a constant rate of 1 ml/min using a mass and volumetric precision gas flow controller (MC-20 Series; Alicat Scientific, Tucson, AZ). The pH was maintained at 7.0 ± 0.1 by using bicarbonate buffer and small additions of 0.1 M NaOH or HCl as needed. Headspace gas levels of N_2 , O_2 , CH_4 , H_2 , and CO were measured at time intervals of 1 to 10 h using an SRI 8610SC gas chromatograph (GC) with thermal conductivity and reducing gas detectors (analytical column: molecular sieve MS-5A, 6 ft; carrier gas: 4.7 laser grade helium at 20 ml/min). Organic acid concentrations (lactate, acetate, and formate) were determined using a Dionex DX500 ionic chromatograph model LC20-1 (analytical column IonPac AS11-HC, eluent 0.75 to 40 mM NaOH, CD20 conductivity detector), whereas ethanol and glycerol were measured enzymatically as described elsewhere (33). The chemostat mode was initiated after the optical density OD_{600} reached 0.29 or 0.25 ± 0.01 (for *M. hungatei* containing coculture) and maintained at a dilution rate of 0.027 h^{-1} . Coculture samples were taken regularly for direct cell counting (Petroff-Hauser counting chamber). Cells were harvested after three retention periods with an OD_{600} variance of <10% by using an ice-chilled sterile stainless steel tube connected to the chemostat medium exhaust line that allowed a cell suspension transfer with minimal O_2 -contact into pregassed, ice-chilled Falcon tubes (50 ml). After centrifugation for 15 min at $3,220 \times g$ at 4°C, the supernatant was carefully poured off, and the cell pellets were immediately frozen at -80°C .

For the qualitative and quantitative investigation of the interspecies electron mediators in coculture, the methanogen was specifically inhibited by manually adding 5 mM bromoethanesulfonate (BES) to the chemostat and medium reservoir bottle after three volume turnovers at constant OD_{600} values. The OD_{600} , organic acids, and gas levels were monitored at short time intervals for 24 h after BES addition, as described above. Based on the data of the first 3 h after BES addition, the electron flow ratio through each electron carrier during syntrophic lactate conversion has been calculated according to the following equation (simplified): % electron flow through electron carrier ($HCOO^-/H_2/CO$) from lactate oxidation = $[\Delta n_{\text{electron}}$ of electron carrier formed / (n_{electron} of lactate inflow - n_{electron} of lactate outflow - $\Delta n_{\text{electron}}$ of methane formed)] $\times 100$, where n_{electron} is the number of electrons in moles and $\Delta n_{\text{electron}}$ is the difference of electrons in moles. Only electrons available from conversion of lactate to acetate (four electrons per lactate molecule) were taken into consideration. The parameters for the chemostat runs were 30 mM lactate in the culture media, a dilution rate of 0.027 h^{-1} , and a 1-ml/min gas flow rate.

Whole-genome transcriptional analysis. Microarray gene expression measurements were performed on three biological replicates of coculture and monoculture chemostats. Total RNA was extracted from frozen cell paste (stored at -80°C) using the RNeasy kit according to the manufacturer's protocol but omitting the lysozyme cell wall digestion step. Total RNA concentration and quality were determined using an RNA nanochip

TABLE 1 Gene expression of hydrogenases, formate dehydrogenases, membrane-bound electron transfer complexes, and putative lactate oxidation pathway enzymes in *Desulfovibrio alaskensis* G20 during syntrophic growth with *Methanococcus maripaludis* and *Methanospirillum hungatei*

Redox proteins in <i>D. alaskensis</i> G20 ^a	Gene locus tag(s) ^b	Mutant growth in coculture ^c	Microarray data					
			Log ₂ <i>R</i> (syntrophy vs sulfate respiration) ^d			Absolute log ₂ ^e		
			<i>M. maripaludis</i>		<i>M. hungatei</i>	<i>M. maripaludis</i>		<i>M. hungatei</i>
			0.047 h ⁻¹	0.027 h ⁻¹	0.027 h ⁻¹	0.047 h ⁻¹	0.027 h ⁻¹	0.027 h ⁻¹
Hydrogenases								
Periplasmic								
HynBA [NiFe]	Dde_2137-Dde_2138	+/-	++	++	-	+	+	+/-
HynBAC ₃	Dde_3754-Dde_3756		-	+/-	+/-	-	-	-
HysBA [NiFeSe]	Dde_2134-Dde_2135	†	-	-	-	-	-	+
HydAB [FeFe]	Dde_0081-Dde_0082	+/-	+/-	+	+	+	+	-
	Dde_2280-Dde_2281	+/-	+/-	+	+	+/-	+/-	-
Cytoplasmic								
[FeFe] monomeric Fd-dependent	Dde_0725		-	+	+	+	+	+
Carbon monoxide dehydrogenase								
Cytoplasmic								
CooSC	Dde_3028-Dde_3029	+/-	-	+/-	+	+	+	++
Formate dehydrogenases								
Periplasmic								
FdhAB	Dde_0716-Dde_0718	-	++	++	++	+	+	++
	Dde_0812-Dde_0813	+/-	+/-	+/-	-	-	-	-
	Dde_3513-Dde_3514	+/-	-	+/-	+/-	-	-	-
FdhABC ₃	Dde_0680-Dde_0683	+/-	+/-	+/-	+/-	+/-	+/-	+/-
Cytoplasmic								
FhcABCD	Dde_0473-Dde_0476	+/-	-	+/-	+/-	-	-	-
Periplasmic cytochromes								
Cytochrome <i>c</i> ₃ (TpIc ₃) CycA	Dde_3182		-	++	++	++	++	++
Split-Soret cytochrome <i>c</i>	Dde_3211		++	++	++	+	+	++
METC								
QrcABCD	Dde_2932-Dde_2935	-	+	+/-	+	+	+	+
HmcABCDEF	Dde_0648-Dde_0653	-/†	+/-	+/-	+	+	+	+
DhcARnfCDGEAB	Dde_0580-Dde_0586		+/-	-	-	+/-	+/-	+/-
QmoABC-D	Dde_1111-Dde_1114	-	+	-	-	+	+	+
DsrMKJOP	Dde_2271-Dde_2275		+/-	-	-	+	+/-	+/-
TmcABCD	Dde_3707-Dde_3710		-	-	-	-	-	-
Carbon metabolism enzymes								
Pyruvate-oxidoreductase								
Por	Dde_3237		-	+	+	++	++	++
PorAB	Dde_1792-Dde_1793		+/-	-	-	+	+	+/-
D-Lactate permease								
LctP-1	Dde_1074	†	++	++	++	+	+	+
LctP-2	Dde_3238	†	++	+	+	+	+	+
L-Lactate dehydrogenase								
L-Ldh	Dde_0750		-	-	+/-	+/-	+/-	+/-
D-Lactate dehydrogenase (monomeric)								
D-Ldh	Dde_3604		+/-	-	-	+/-	-	-
GlcD	Dde_0182		-	+/-	+/-	-	-	+/-
D-LdhA	Dde_1681		+/-	-	-	+/-	+/-	+/-
D/L-Lactate dehydrogenase (with HdrD-like subunits)								
LldEFG	Dde_1842-Dde_1844		+	-	-	+	+	+/-
LdhA + Ldh-HdrD-1 (LdhB-1)	Dde_3239-Dde_3240	†	++	+	++	++	++	++
Ldh3	Dde_1085-Dde_1087		-	+/-	+/-	-	-	-
Phosphotransacetylase Pta	Dde_3241		++	+/-	+	++	++	++
Acetate kinase AckA	Dde_3242		+	+	+	+	++	++
Pyruvate formate lyase Pfl-I	Dde_1273	+/-	-	-	-	-	-	-
NADH oxidase Nox	Dde_0374	-	-	-	-	+/-	+/-	+/-

(Continued on following page)

TABLE 1 (Continued)

Redox proteins in <i>D. alaskensis</i> G20 ^a	Gene locus tag(s) ^b	Mutant growth in coculture ^c	Microarray data					
			Log ₂ <i>R</i> (syntrophy vs sulfate respiration) ^d			Absolute log ₂ ^e		
			<i>M. maripaludis</i>		<i>M. hungatei</i>	<i>M. maripaludis</i>		<i>M. hungatei</i>
			0.047 h ⁻¹	0.027 h ⁻¹	0.027 h ⁻¹	0.047 h ⁻¹	0.027 h ⁻¹	0.027 h ⁻¹
Hdr-like proteins								
HdrABC-FloxABCD	Dde_1207-Dde_1213		–	–	–	+/-	+/-	+/-
	Dde_3523-Dde_3530		–	+/-	+/-	–	–	–
Ldh-HdrD-2 (LdhB-2)	Dde_3245	†	++	+	+	+		++
FAD-HdrD	Dde_0312	–	+/	–	–	+/-	+/-	+/-
Mop-HdrD	Dde_3540		–	–	–	–	–	–
Cytoplasmic electron carrier								
Ferredoxin II	Dde_0286		++	++	+/	++	++	+
DsrC (thiol/disulfide redox pair)	Dde_0762		–	–	–	++	++	++

^a Gene annotation according to <http://www.microbesonline.org> (87), Pereira et al. (39), and Keller and Wall (38).

^b *D. alaskensis* G20 transposon insertion mutants of respective genes investigated in comparative growth experiments are indicated in boldface.

^c Mutant growth in coculture with *M. hungatei* on lactate is summarized as follows: +/-, growth identical to reference mutant strain Dde_1072; –, inhibited growth relative to the reference strain; and †, no growth observed.

^d Log₂*R* of gene expression of *D. alaskensis* G20 grown in coculture with *M. maripaludis* or *M. hungatei* (30 mM lactate) versus *D. alaskensis* G20 grown in monoculture (30 mM lactate, 10 mM sulfate) at dilution rates of 0.047 h⁻¹ and 0.027 h⁻¹ (log₂*R* values for enzymes or protein complexes are summarized as follows: ++, log₂ *R* > 1; +, log₂ *R* > 0.5; +/-, log₂ *R* > 0; and –, log₂ *R* < -0.5).

^e Absolute log₂ values (normalized with a median of 0) of *D. alaskensis* G20 grown in coculture with *M. maripaludis* or *M. hungatei* at dilution rates of 0.047 h⁻¹ and 0.027 h⁻¹ (++, absolute log₂ > 4; +, absolute log₂ > 2; +/-, absolute log₂ > 0; –, absolute log₂ < -0.5).

run on a Bioanalyzer (Agilent, Santa Clara, CA). We generated first strand cDNA using the SuperScript Indirect cDNA Labeling System (Life Technologies, Carlsbad, CA) with random hexamers. The cDNA was labeled with Alexa 555 (Life Technologies). Labeled cDNA (~1 µg per slide) was hybridized to a custom Nimblegen microarray containing four independent microarrays comprised of 72,000 probes each (60-mers). The vast majority of genes are represented by nine unique oligonucleotide probes, with each probe arrayed in duplicate. After overnight incubation in a hybridization station (Maui), slides were washed using the standard Nimblegen protocol. Microarrays were scanned using a GenPix Pro 4200A according to the Nimblegen recommendations. Arrays were scanned at a minimum of three PMT (photomultiplier tube) gains. The scanned array images were analyzed for probe intensities using the Nimblegen Scan v2.4, with the local alignment turned off and a border of -1. Feature files containing the probe intensities and probe identifiers were exported for further analysis. PMT settings were used with ~1% of probes being above half of the maximum value. Each scan was normalized by forcing the distribution of values to conform to the distribution of log₂ intensities of monoculture control chemostats (average of two monoculture chemostats). After excluding probes that were in the lowest 2% of the control, initial log₂ levels were computed by averaging across replicate spots of each probe, and the log₂ ratio median was set to zero. To remove intensity-dependent effects, we used local regression (Loess in R) of each probe's log₂ ratio versus the log₂ level and subtracted out these estimated effects. To assess whether genes had statistically significant changes in expression, we used OpWise (34), which uses the correlation of expression of genes in predicted operons to estimate the reliability of bacterial expression data.

Phenotypic growth analyses of strain G20 transposon mutants. The complete description and validation of a *D. alaskensis* G20 transposon mutant collection is in preparation and will be described elsewhere (A. Deutschbauer and A. Arkin, unpublished data). For each of the 26 mutant strains used here, the insertion of the transposon was determined by a two-step arbitrary PCR protocol (35) and further confirmed by PCR amplification with a transposon- and genome-specific primer pair.

Endpoint dilution monocultures from glycerol stocks of *M. hungatei*, *D. alaskensis* G20 wild-type and mutant strains were used to initiate all pairings (methanogen was grown in media containing only 5 mM acetate

with an H₂-CO₂ [80:20, vol/vol] headspace at an overpressure of 20 lb/in²). Cocultures were established by mixing 1 ml of the highest dilution of the *D. alaskensis* G20 wild type or an insertion mutant strain with 1 ml of the highest dilution of *M. hungatei* in 13 ml of coculture medium in Balch tubes. Initial batch mono- and cocultures were transferred at least two times (1% [vol/vol]) before inoculation of triplicate tubes for monitoring growth (OD₆₀₀) using uninoculated medium as a reference. All cultures were incubated at 37°C with horizontal shaking at 300 rpm.

Microarray data accession number. The microarray data are available under GEO reference [GSE46547](https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE46547).

RESULTS

Growth characteristics of *D. alaskensis* G20 chemostat cocultures with different methanogenic partners and growth rates. In a previous study (15), continuous cultures of *D. alaskensis* G20 were grown on 30 mM lactate in monoculture (10 mM sulfate added) and coculture with *M. maripaludis* at a high dilution rate of 0.047 h⁻¹ (the maximum growth rate of this coculture pairing in chemostat is 0.051 h⁻¹). The impact of growth rate and methanogenic partner on syntrophic assemblies was investigated by running additional chemostats with *D. alaskensis* G20 grown in monoculture or coculture with different methanogens (*M. maripaludis* or *M. hungatei*) at a low dilution rate of 0.027 h⁻¹ (close to the maximum growth rate of the *M. hungatei* coculture pairing at 0.031 h⁻¹ [see Table 2]). Notably, the latter pairing showed an extended lag phase by ~50 h compared to the *M. maripaludis* containing coculture and reached only a maximum OD₆₀₀ of 0.25 ± 0.01 before the start of the chemostat growth (Fig. 1 and 2). The coculture cell ratios stabilized at ratios of ~1.0 (*Desulfovibrio: Methanococcus*) and ~1.2 (*Desulfovibrio: Methanospirillum*), respectively (Fig. 1). Scanning electron microscopy (SEM) imaging of cell suspensions at steady state demonstrated the *Desulfovibrio*-methanogen interactions to be localized predominantly at the *Desulfovibrio* cell poles, whereas a general increase in cell aggregation (EPS formation) was observed compared to growth at a rate of

TABLE 2 Growth characteristics of *D. alaskensis* G20 cocultures on 30 mM lactate in chemostat with different methanogens and dilution rates^a

Parameter	Coculture pairing/dilution rate		
	G20 + <i>M. maripaludis</i>		G20 + <i>M. hungatei</i>
	0.047 h ⁻¹	0.027 h ⁻¹	0.027 h ⁻¹
Batch mode growth			
Maximum growth rate	0.051 h ⁻¹	0.051 h ⁻¹	0.031 h ⁻¹
Mean maximum OD ₆₀₀ ± SD	0.30 ± 0.01	0.30 ± 0.01	0.25 ± 0.01
Chemostat			
Start after inoculation (h)	60–65	60–65	105–120
Mean OD ₆₀₀ at steady state ± SD	0.29 ± 0.01	0.25 ± 0.01	0.21 ± 0.01
Cell ratio (<i>Desulfovibrio</i> /methanogen)	1.4	1.0	1.2
Organic acid concn (mM)			
Lactate	< 0.05	< 0.05	< 0.05
Acetate	26–28	26–28 mM	26–28
Formate	ND ^b	ND	ND
Gas content			
Methane (%)	50–60	30–40	30–40
Hydrogen (ppm)	350–550	280–400	70–110
Carbon monoxide (ppm)	10–15	7–9	4–6

^a Average values of three replicate chemostats for each pairing/growth rate are shown.

^b ND, no data (below the detection limit of IC).

0.047 h⁻¹ (see Fig. S2 in the supplemental material). The lactate level remained close to, or below, the detection limit (0.05 mM) during chemostat growth, indicating complete utilization of the added lactate (Fig. 2). Acetate was the dominant fermentation product (26 to 28 mM), while no other organic acids, ethanol, or glycerol were detected. After an initial hydrogen burst (fermentation burst [1–3], prolonged in the *M. hungatei* coculture pairing with occasionally measurable formate levels), the H₂ concentration dropped to constant levels of between 280 and 400 ppm for the *M. maripaludis* coculture pairings and between 70 and 110 ppm for the *M. hungatei* coculture pairings. In contrast, the carbon monoxide concentration increased slowly during batch growth to final concentrations of 8 ppm (*M. maripaludis*) or 5 ppm (*M. hungatei*) throughout the run (Fig. 3). Measurable methane formation started within 10 to 15 h (*M. maripaludis*) or 45 to 50 h (*M. hungatei*) after inoculation comprising ~35% of the headspace gas phase during chemostat operation (Fig. 3; gas level fluctuations are most likely attributable to manual sampling and GC injection). Notably, in the coculture chemostats in which the methanogen metabolic activity was inhibited by BES (following growth stabilization after at least three retention periods), formate could be detected within the first hour after treatment (0.1 and 0.05 mM for the *M. maripaludis* and *M. hungatei* cocultures, respectively), accumulating to more than 0.3 and 0.2 mM, respectively, within the next 2 h (Fig. 2). The inhibition of the methanogen by BES was confirmed by an immediate drop in the OD, a decrease in acetate, an increase in lactate, and an exponential decline in methane levels (Fig. 2 and 3). The later was consistent with the calculated gas exchange rate based on a 1-ml/min gas flow through the bioreactor headspace. The accumulation of formate so shortly after BES treatment is a strong indicator that formate is an important mediator in the interspecies electron exchange in cocultures of *D. alaskensis* G20 irrespective of the growth rate and

methanogenic partner. An approximate calculation of the syntrophic electron flow at 0.027 h⁻¹ growth rate (based on the formate, H₂, and CO production immediately following methanogen inhibition [data were obtained from at least two replicate experiments]) points to a transfer of reducing equivalents of >63.5% as formate (less than 36% and 0.5% as H₂ and CO, respectively). Notably, the *M. hungatei* coculture shuttled more electrons via H₂ compared to *M. maripaludis* (Table 3). Both methanogens are capable of growing on formate-acetate; however, their growth rates on formate are generally slower compared to growth on H₂-acetate (Fig. 4). No methane production or significant growth was observed for either methanogen on carbon monoxide-acetate. Irrespective of the substrate, *M. hungatei* showed an extended lag phase and lower growth rate and biomass yield, but a slightly higher total methane production compared to *M. maripaludis*.

Differential gene expression during syntrophic growth: impact of growth rate and methanogenic partner. Syntrophically grown *D. alaskensis* G20 had 172 and 119 differentially regulated genes in coculture with *M. maripaludis* and *M. hungatei*, respectively, compared to respiratory growth in monoculture at identical low growth rate (genes with an absolute log₂ *R* value of >0.5 and an absolute *Z* score of >2.0 were included; see Tables S3 and S4 in the supplemental material). The numbers of differentially regulated genes are lower relative to the number of genes of the *M. maripaludis* pairing grown at a higher growth rate (400 up- or

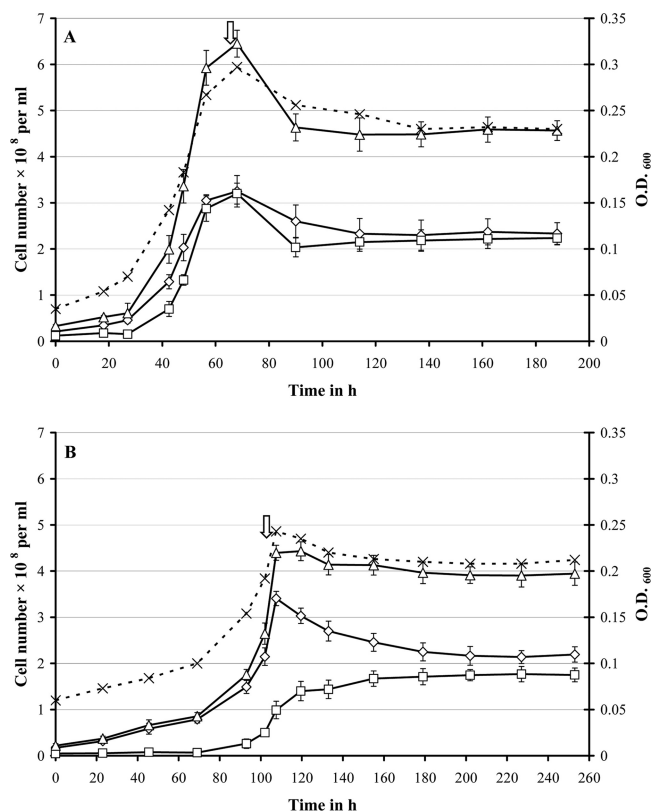


FIG 1 Growth characteristics of *D. alaskensis* G20 grown in coculture with either *M. maripaludis* (A) or *M. hungatei* (B) in chemostats at a dilution rate of 0.027 h⁻¹. The optical density (dashed line, ×), *D. alaskensis* G20 (◇), methanogen (□), and total cell number (△) are shown for a single coculture replicate. The continuous coculture was started at 65 or 110 h, respectively (indicated by arrow).

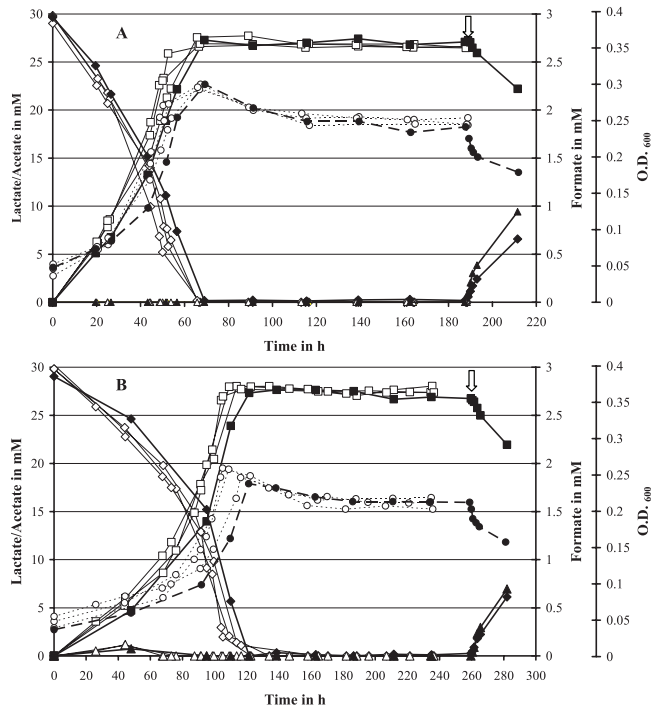


FIG 2 Optical density (○, ●), lactate (◇, ◆), acetate (□, ■), and formate (△, ▲) profiles for three coculture replicates of *D. alaskensis* G20 grown in coculture with either *M. maripaludis* (A) or *M. hungatei* (B) in chemostats (open symbols). The coculture chemostat of each pairing that was treated with 5 mM BES after three volume turnovers at 0.027 h⁻¹ dilution rate is marked by closed symbols (time point of BES addition is marked by arrow).

downregulated genes [15]). The COG functional category of energy metabolism was highest among both up- and downregulated genes for both coculture pairings. Significant increases in transcript abundance were observed for general function prediction, inorganic ion transport (*M. maripaludis*) and coenzyme metabolism categories (*M. hungatei*), whereas several genes of signal transduction, intracellular trafficking, and inorganic ion transport were downregulated (see Fig. S5 in the supplemental material). Notably, energy source availability (as controlled by dilution rate) had a much greater impact on syntrophic gene regulation in *D. alaskensis* G20 (see Fig. S6A and B in the supplemental material) than the methanogenic partners (see Fig. S6C in the supplemental material). The significant growth rate effect was reflected by expression changes in 268 genes (including 33 operons) in the *M. maripaludis* cocultures (absolute Z score > 2.0; absolute $\Delta\log_2 R > 0.5$; Fig. 5A and see Table S7 in the supplemental material for details). Their affiliation with almost every COG category demonstrates the global transcriptional response to energy limitation. Although several of the significantly up- or downregulated genes (absolute $\log_2 R$ value > 0.5) were exclusively differentially expressed in only one of the coculture pairings (66 and 27, respectively), only 22 genes showed a significant change in expression (absolute $\Delta\log_2 R > 0.5$) between both pairings when grown at the same rate (Fig. 5B; see also Table S8 in the supplemental material for details). Interestingly, a large fraction of these genes (41%) was associated with energy metabolism (COG category C) and included periplasmic formate dehydrogenase (*fhd-1*, Dde_0716-Dde_0718), periplasmic [NiFeSe] and [NiFe] hydrogenases

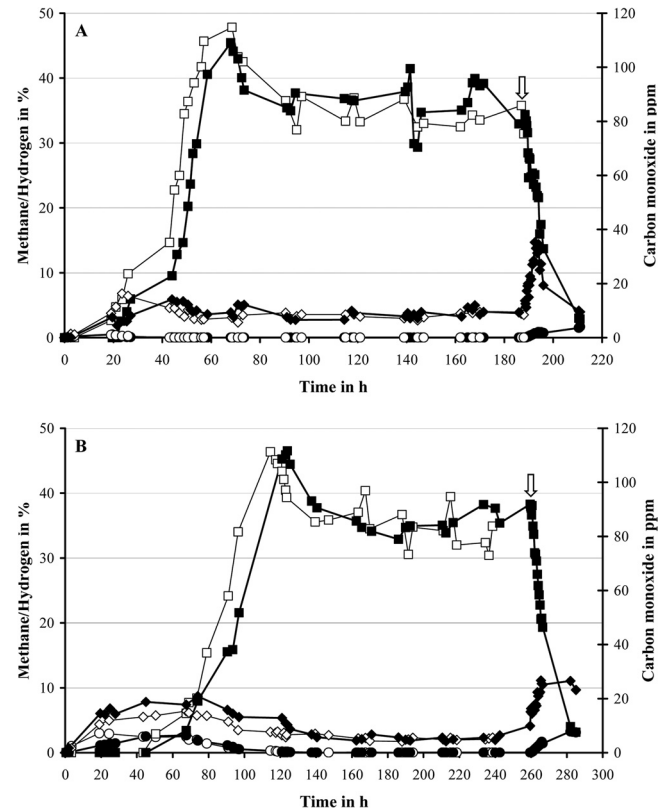


FIG 3 Methane (□, ■), hydrogen (◇, ◆), and carbon monoxide (○, ●) profiles for three coculture replicates of *D. alaskensis* G20 grown in coculture with either *M. maripaludis* (A) or *M. hungatei* (B) in chemostats (open symbols). The coculture chemostat treated with 5 mM BES after three volume turnovers at 0.027 h⁻¹ dilution rate is shown in closed symbols (the time point of BES addition is marked by an arrow).

(*hysAB*, Dde_2134-Dde_2135; *hynAB*, Dde_2137-Dde_2138), ferredoxin (Dde_0286), and CO dehydrogenase (Dde_3028-Dde_3029), as well as putative cofactor biosynthesis-associated genes (Dde_0719, Dde_0805-Dde_0806). Thus, the results clearly demonstrated that this *Desulfovibrio* also has the capacity to adapt to different methanogenic partners.

The gene expression data sets were merged to identify putative syntrophy-specific genes in *D. alaskensis* G20. Notably, only 68 genes were commonly differentially expressed under the different syntrophic growth conditions syntrophy relative to sulfate respiration on lactate (Fig. 6A; for a detailed list, see Table S9A in the supplemental material). Of these genes, 54 were commonly downregulated, while only 14 genes are upregulated in strain G20 during syntrophy. The most highly downregulated (lowest $\log_2 R$ values under all syntrophic conditions studied) comprised 11 operons (33 genes) of the Fur regulon (e.g., biopolymer transport complexes Dde_0447-Dde_0457, Dde_2152-Dde_2157; iron-repressed flavodoxin *fld*, Dde_3667; ferrous iron uptake systems *feo*, Dde_2669-Dde_2673; and regulatory proteins *hdd*, Dde_1511; *araC*, Dde_1611; *rsbU*, Dde_3047 [36, 37]). In addition, the *tmc* operon (Dde_3707-Dde_3710), a gene cluster of regulators and signal transduction components (Dde_3711-Dde_3714, TmcR regulated) and two uncharacterized operons (Dde_3197-Dde_3202, RpoN regulated [36]) were significantly repressed in syntrophy compared to sulfate respiration (COG groups: C, P, U,

TABLE 3 *D. alaskensis* G20 electron flow ratio through the electron carriers formate, hydrogen, and carbon monoxide during syntrophic lactate conversion^a

Electron carrier	Mean electron flow ratio (%) in coculture \pm SD					
	G20 + <i>M. maripaludis</i>				G20 + <i>M. hungatei</i>	
	0.047 h ⁻¹		0.027 h ⁻¹		0.027 h ⁻¹	
	1 h	3 h	1 h	3 h	1 h	3 h
Formate	83.5 \pm 2.4	78.0 \pm 2.1	84.5 \pm 3.0	81.0 \pm 2.5	80.3 \pm 2.8	63.7 \pm 1.7
Hydrogen	16.5 \pm 1.7	21.5 \pm 2.3	15.5 \pm 2.5	18.5 \pm 2.5	19.5 \pm 3.2	36.3 \pm 1.7
Carbon monoxide	0.02 \pm 0.1	0.02 \pm 0.01	0.5 \pm 0.1	0.03 \pm 0.01	0.3 \pm 0.1	0.07 \pm 0.01

^a Calculations are based on biochemical data from the first 3 h after BES inhibitor treatment of coculture chemostats. Average values of two to three replicate experiments for each growth rate/methanogen pairing are shown.

M, and T). In contrast, most of the commonly upregulated genes during syntrophy were associated with energy metabolism and comprised genes encoding for lactate uptake (LctP-1 and -2) and conversion (lactate-oxidation pathway operon [38, 39], Dde_3237-Dde_3245), split-soret cytochrome *c* (Ssc, Dde_3211), Fdh-1 (all regulated by RpoN and ModE [36]) and the carbon starvation protein CstA (Dde_0227).

Notably, there appears to be a pool of 19 “*M. maripaludis* syntrophy-relevant” genes that are only differentially expressed when strain G20 is grown in coculture with this methanogen (including HynAB, HysAB, ferredoxin, cofactor biogenesis-associated genes Dde_0805-Dde_0806, Dde_2018, and the 2-hydroxyglutaryl-coenzyme A (CoA) dehydratase operon Dde_0089-Dde_0090; see Fig. 6B and see Table S9B in the supplemental material). Interestingly, most of these genes also showed the highest change in gene expression between both coculture pairings (Fig. 5B). The transcriptional data set analysis further indicate 18 genes to be relevant in *D. alaskensis* G20 for syntrophic growth at low growth rate (e.g., signal transducing GGDEF-domain protein, Dde_3048, and uncharacterized histidine kinase, Dde_3715; Fig. 6C and see Table S9C in the supplemental material for details).

Comparative phenotypic growth experiments of mutant strains. Phenotypic growth analyses of 26 *D. alaskensis* G20 transposon mutants with gene insertions in formate dehydrogenases, hydrogenases, membrane-bound electron transfer complexes

(METCs), lactate metabolism enzymes were carried out in monoculture and coculture with *M. hungatei* to evaluate the function of the respective genes for syntrophy at a low growth rate (this methanogen is a slower H₂/formate scavenger relative to *M. maripaludis*) (Fig. 7). Surprisingly, all mutants showed an extended lag phase in lactate-based coculture growth compared to the wild type even after four transfers (1% inoculum), whereas no differences were observed between the mutant and wild-type strains when grown in coculture on pyruvate and in monoculture on both substrates. The mutant phenotypic growth behavior in coculture on lactate was therefore evaluated in reference to mutant strain Dde_1072, which carries a syntrophy-irrelevant transposon insertion. Among the mutants of formate dehydrogenases or hydrogenases, the HysA mutant (Dde_2135) showed no syntrophic growth on lactate, while both mutants for Fdh-1 (Dde_0716-Dde_0718) were slightly impaired (lower growth rates and maximum cell densities than the reference strains). Notably, the *hysAB* and *fdh-1* genes were highly expressed during coculture growth at steady-state conditions (see Table 1). None of the investigated formate dehydrogenase and hydrogenase mutants showed growth inhibition during sulfate respiration on lactate or pyruvate.

Among the METC subunit mutants investigated, the QrcD (Dde_2935) and HmcF (Dde_0648) showed no growth in coculture on lactate, whereas mutants for HmcE (Dde_0649), HmcB (Dde_0652), and QmoD (Dde_1114; the structural component of

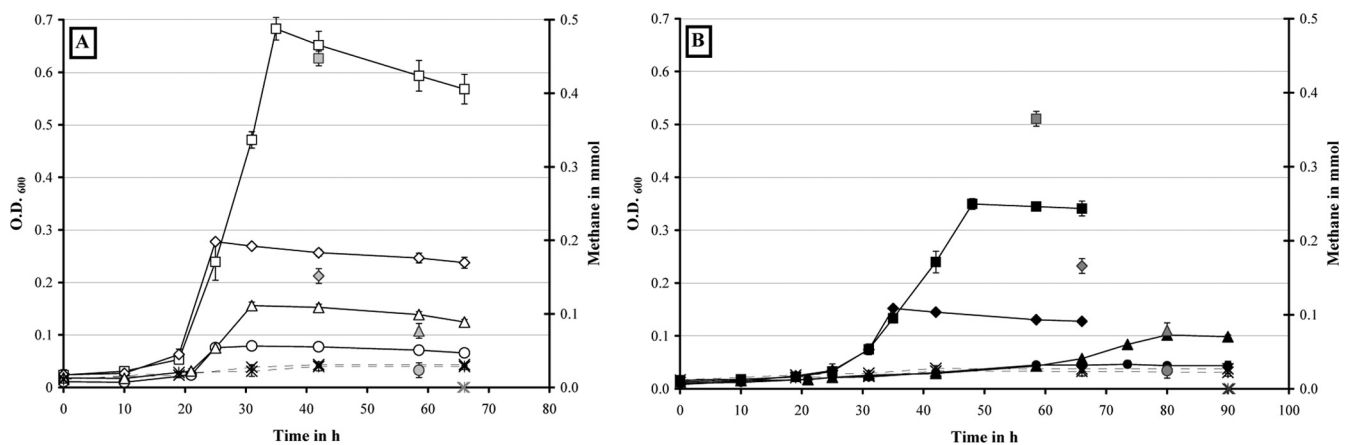


FIG 4 Growth curves of *M. maripaludis* (A) and *M. hungatei* (B) in batch culture on the following substrates: hydrogen at 30 lb/in² (□, ■) or 1 lb/in² (◇, ◆) overpressure, formate at 60 mM (△, ▲) or 20 mM (○, ●), and carbon monoxide at 0.38% (dashed lines, *) or 0.2% (dashed lines, ×) headspace concentration (5 mM acetate was added to all cultures). The total amount of methane produced is shown in gray symbols for each substrate. Error bars indicate the standard deviations for triplicate cultures.

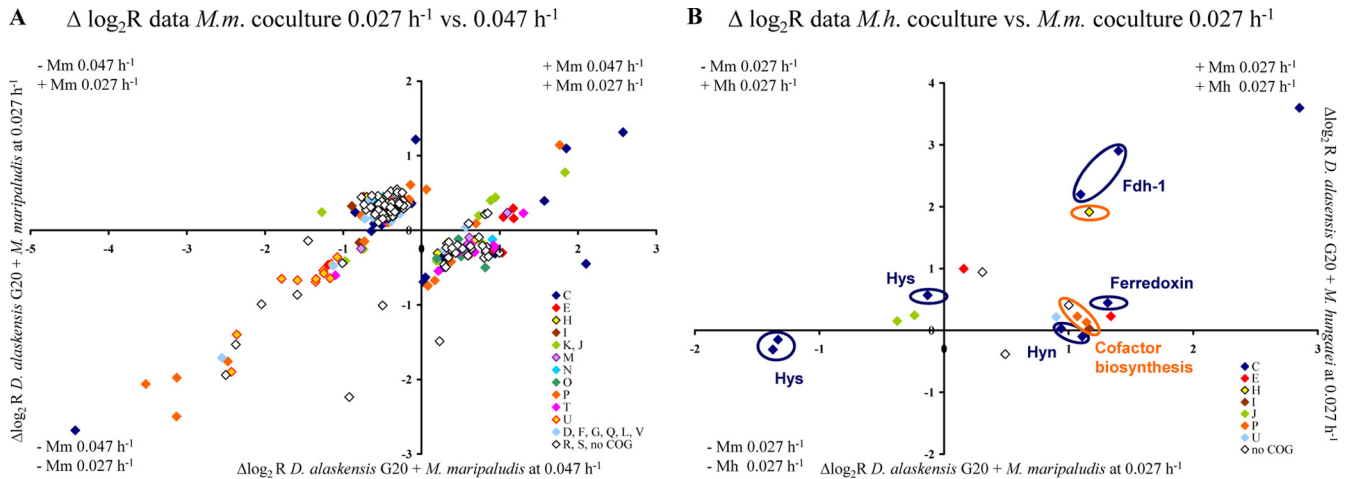


FIG 5 (A) Plot of the $\log_2 R$ values of the 268 genes of *D. alaskensis* G20 differentially expressed during syntrophic growth with *M. maripaludis* between different growth rates. (B) Plot of the $\log_2 R$ values of the 22 genes of *D. alaskensis* G20 differentially expressed during syntrophic growth between different methanogens at the same growth rate. Genes are differently colored according to their COG functional affiliation (single-letter code: amino acid transport, E; carbohydrate transport and metabolism, G; cell division and chromosome partitioning, D; cell motility and secretion, N; cell envelope biogenesis, M; chromatin structure and dynamics, B; coenzyme metabolism, H; defense mechanism, V; energy production and conservation, C; function unknown, S; general function prediction only, R; inorganic ion transport and metabolism, P; intracellular trafficking and secretion, U; lipid metabolism, I; nucleotide transport and metabolism, F; posttranslational modification, protein turnover, chaperones, O; DNA replication, recombination and repair, L; secondary metabolites biosynthesis, transport and catabolism, Q; signal transduction mechanisms, T; transcription, K; and translation, ribosomal structures and biogenesis, J).

the Qmo complex [40]) were slightly inhibited with lower growth rates and maximum cell densities relative to the reference strains (80 to 73%). The QmoD mutant demonstrated inhibited respiratory growth on both lactate and pyruvate (extended lag phase), while the QrcD mutant was only impaired in monoculture on pyruvate (27% lower maximum cell density than the wild type). Among the lactate metabolism enzymes, mutants of the pyruvate: formate lyase (Dde_1273) demonstrated slightly higher biomass production than the wild type in monoculture and pyruvate-grown coculture. Mutants for the NADH oxidase, Nox, and an uncharacterized FAD-HdrD homologue were inhibited in coculture on lactate with extended lag phases and lower growth rates relative to the reference strains. The mutants of both putative lactate permeases (LctP-1, Dde_1074, and LctP-2, Dde_3238) and the HdrD-like putative beta subunits of the lactate dehydrogenase (LdhB-1, Dde_3240, and LdhB-2, Dde_3245) showed no growth in coculture on lactate, confirming their essential function during syntrophy. However, as demonstrated by their slightly inhibited (LdhB-1/2) and impaired (LctP-2) growth or reduced maximum cell density (LctP-1 with <20% than the wild type) in monoculture on lactate, they also are involved in lactate uptake/oxidation processes during sulfate respiration. Notably, the mutant for the lactate-oxidation-operon integral permease LctP-2 also showed inhibited growth in mono- and coculture on pyruvate.

DISCUSSION

The transcriptional data of this study demonstrated a significant shift in the gene expression profile upon change of syntrophic growth conditions. The limitation of energy availability had a major impact on the physiology of syntrophic *D. alaskensis* G20, as shown by the differential expression of 268 genes (including 41 operons) of a wide range of COG functional categories between different coculture growth rates with the same methanogen. Besides energy metabolism, many of the encoded proteins are involved in cell surface biogenesis (19 genes, e.g., LpxDC, KdsA,

Gmd, and pilus assembly proteins), transport (18 genes, e.g., LivFK, CysWA, PstAC, and Amt), and signal transduction and transcriptional regulation (13 and 9, respectively, e.g., *hmc* operon-associated Rrf2 [41]) that might result in structural and physiological modifications of the cell. The methanogenic partner has only a minor influence on the gene expression of strain G20, as shown by significant transcriptional level variations in only 22 genes between the coculture pairings. However, only 68 genes were commonly differentially expressed in strain G20 under syntrophic versus respiratory lifestyle. The majority of the 54 down-regulated genes in syntrophy are part of the Fur regulon [36, 37] and reflect physiological adaptations of the *Desulfovibrio* to changes in the environmental conditions associated with the absence of sulfide production in syntrophy, e.g., different metal availability. Notably, only 14 genes are upregulated under all syntrophic conditions investigated which include the lactate oxidation pathway operon, genes for an additional lactate permease and periplasmic cytochrome *c* (Ssc), the Fdh-1 formate dehydrogenase (RpoN and ModE regulon [36]), and the proposed dipeptide transporter, CstA (induced in, for example, *Escherichia coli* during starvation periods to enhance its metabolic potential [42, 43]). Their significant upregulation is most simply explained by the requirement for increased lactate uptake and oxidation rates to sustain growth through fermentation (lower energy availability relative to sulfate respiration). Thus, there are no truly syntrophy-specific but only syntrophy-relevant genes present in *D. alaskensis* G20.

In a previous study examining the metabolic and genetic basis for syntrophy in two different *Desulfovibrio* species (grown with *M. maripaludis* at their respective maximum growth rates), we demonstrated that the faster coculture growth of *D. alaskensis* G20 relative to *D. vulgaris* Hildenborough was attributed to two factors: (i) an alternative electron transfer system conferring a kinetic advantage by permitting higher lactate oxidation rates under ele-

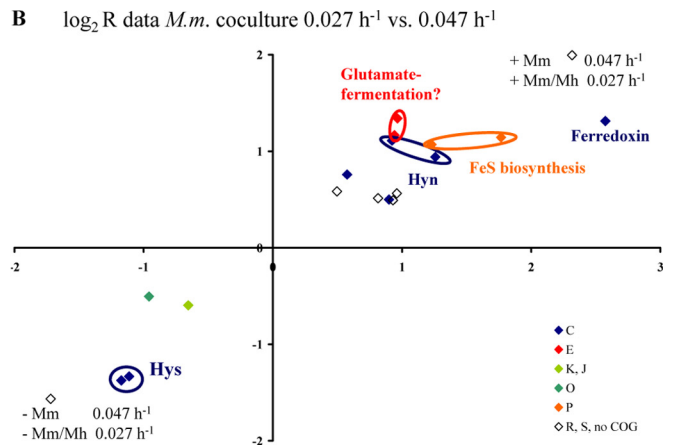
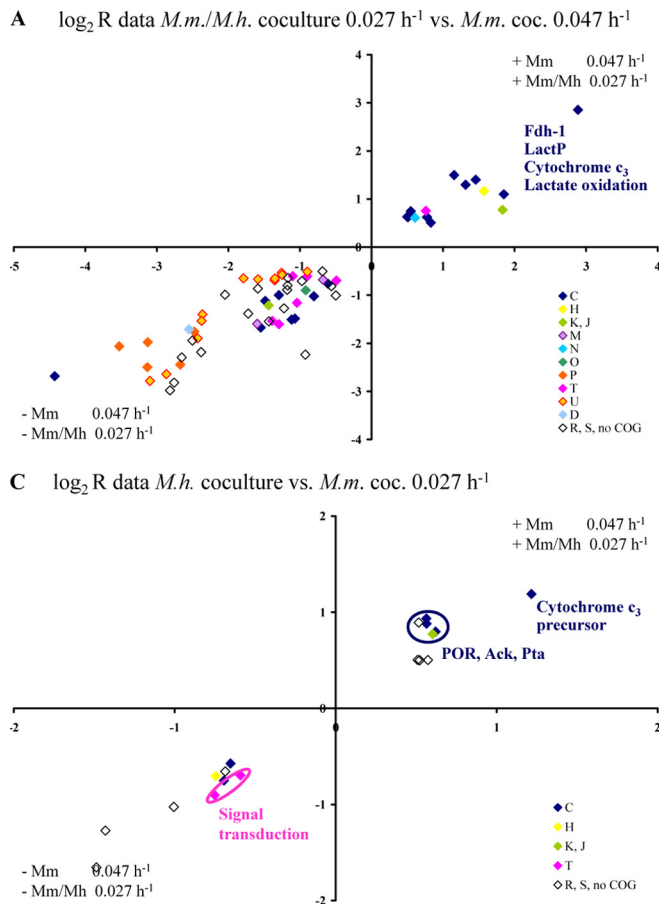


FIG 6 Plot of the $\log_2 R$ values of *D. alaskensis* G20 genes commonly differentially expressed in strain G20 during syntrophic growth irrespective of growth rate and methanogenic partner (“syntrophy-relevant,” 68 genes) (A), only in coculture with *M. maripaludis* irrespective of growth rate (putatively “*M. maripaludis* syntrophy-relevant,” 19 genes) (B), and only at a low growth rate irrespective of the methanogenic partner (putatively “low syntrophic growth rate-relevant,” 18 genes) (C). Genes are differently colored according to their COG functional category affiliation (see the legend of Fig. 5 for the definitions of the single-letter code).

ated H₂ levels and (ii) the use of formate as a main interspecies electron exchange mediator (15), which could sustain a 100-fold-higher conversion rate than interspecies hydrogen transfer based on diffusion kinetics (6, 7). Divergent growth characteristics of syntrophic cocultures of individual *Desulfovibrio* species in batch culture studies were indicative of a high level of intraspecies metabolic flexibility that was evaluated in the present study by varying the environmental parameters, e.g., methanogenic partner and energy source availability (controlled by dilution rate), for *D. alaskensis* G20 cocultures.

The most general effect of energy limitation imposed by the lower growth rate was a significant reduction in the proportion of *Desulfovibrio* (Table 2). A drop in total cell numbers in the *M. maripaludis* and *M. hungatei* cocultures of 77 or 67%, respectively, at the lower growth rate was primarily associated with reduced *Desulfovibrio* cell numbers (70 and 65% of the values observed at the higher growth rate). The coculture structure shifted from a *Desulfovibrio* dominated to an evenly populated community, indicating the nonsustainability of the initial coculture populations at lower energy availability. The lower methanogen cell number and higher cell ratio in the *M. hungatei* pairing relative to the *M. maripaludis* coculture might be attributable to the cell size/morphology differences between both species (28–31). At a lower growth rate, the community was engaged in physical interspecies association. The interspecies cell-cell contact appears to be initiated by a tight association of the methanogen to the poles of individual *Desulfovibrio* cells, most likely as a result of the involvement

of spatially varying surface structures that promote the interactions, e.g., polar chemoreceptor clusters (44–47) or flagella (demonstrated to function in cell aggregation and biofilm formation by inducing expression of genes required for syntrophic growth [14]).

In mutualistic cocultures, the methanogen determines the specific growth rate for a pairing since both partners have to equalize their individual growth rates to achieve a balanced exchange of electron carriers to allow exponential growth of the culture (18, 48–50). The higher maximum growth rate of the *M. maripaludis*-*D. alaskensis* G20 coculture (0.051 h⁻¹) relative to the *M. hungatei* pairing (0.031 h⁻¹) correlates with higher growth rates of *M. maripaludis* in monoculture (e.g., on formate, the rates are 0.16 h⁻¹ versus 0.04 h⁻¹ for *M. hungatei*). Its higher growth rate in coculture might reflect the structural (e.g., sheath structure of *M. hungatei* [47, 51, 52] and type IV pili in *M. maripaludis* [53, 54]) or physiological (facultative autotrophic *M. maripaludis* [28, 29] and mixotrophic *M. hungatei* [31, 55]) differences. *M. maripaludis* utilizes H₂ and formate faster than *M. hungatei* under nonlimiting conditions (see Fig. 4) (30, 56), which might be the result of the various enzyme systems used for methanogenesis and energy conservation (21–24). In *M. maripaludis*, an electron-bifurcating complex (consisting of heterodisulfide reductase [HdrABC], F₄₂₀-reducing formate dehydrogenase [Fdh-1], and F₄₂₀-nonreducing hydrogenase [Vhu]) couples the reduction of heterodisulfide and CO₂ and turns the process of methanogenesis into a cyclic pathway (57–59); the respective genes are also upregulated in syn-

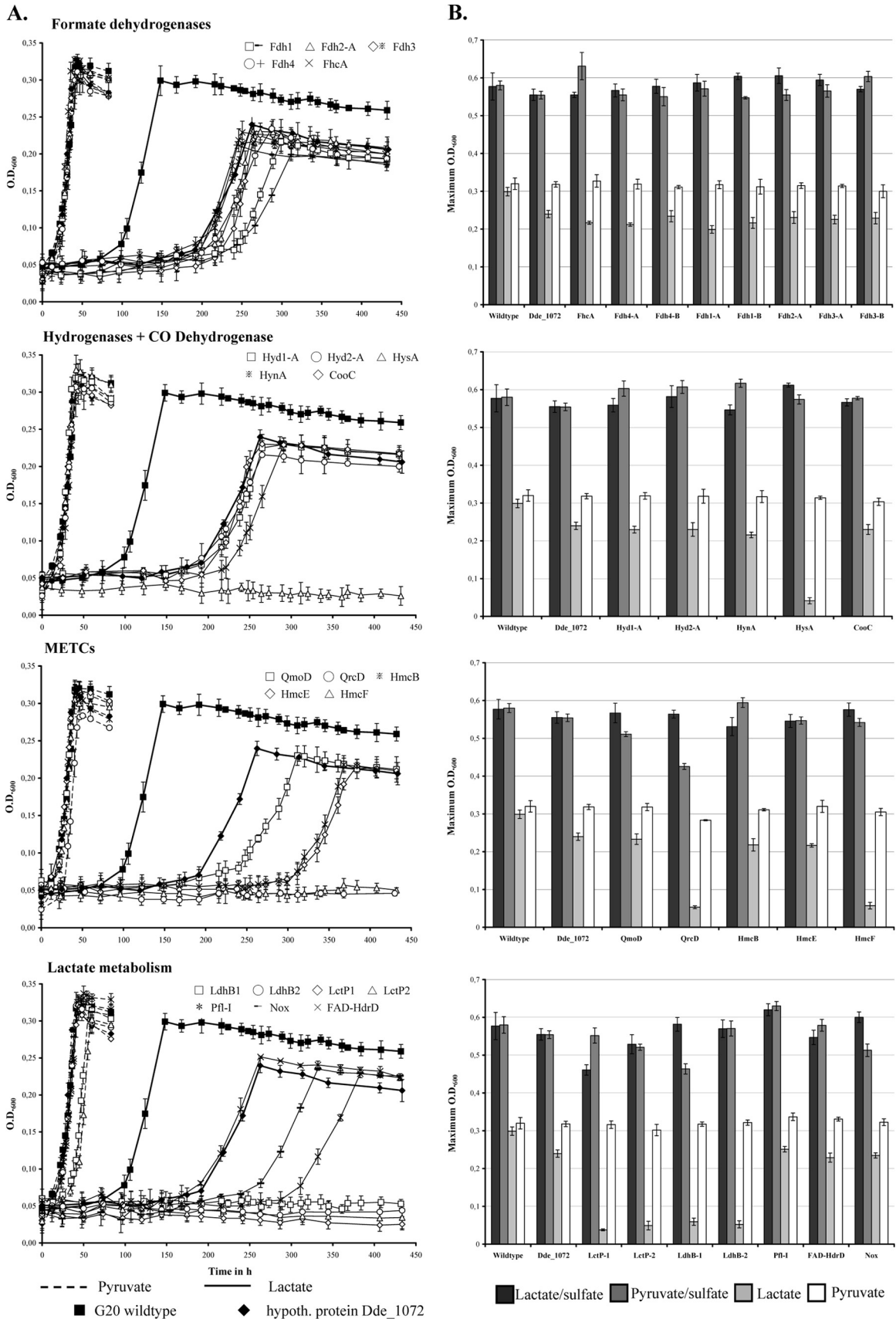


FIG 7 (A) Growth curves for *D. alaskensis* wild-type and mutant strain cocultures with *M. hungatei* on pyruvate and lactate. The wild type and the Dde_1072 insertion mutant strain are indicated by filled symbols, whereas other mutant strains are indicated by open symbols (mutant strain data have been clustered into groups according to their proposed gene function, as shown in Table 1). (B) Maximum ODs of wild-type and mutant strains grown in monoculture (with sulfate) or coculture (without sulfate) on either 30 mM lactate or 30 mM pyruvate. The error bars indicate the standard deviations for triplicate cultures.

trophic *M. maripaludis* (60). In contrast, the heterodisulfide/CO₂ reductions in *M. hungatei* are proposed to be coupled to transmembrane proton or sodium ion transport (22). Genes for Fdh-1 and energy-converting hydrogenases (Ech and Mbh) are highly expressed in syntrophic *M. hungatei* (9), which could result in lower syntrophic growth rates since ferredoxin-linked methanogenesis requires an energized membrane. In *D. alaskensis* G20 cocultures, formate is used as the main interspecies electron carrier irrespective of growth conditions (at least 64% of syntrophic electron flow was via formate). Notably, *M. hungatei* utilizes H₂ preferentially (the monoculture growth rates were 0.04 h⁻¹ on formate and 0.11 h⁻¹ on H₂) and maintained a 4-fold-lower H₂ level in coculture than *M. maripaludis* (the average concentrations of dissolved H₂ in chemostats were 70 and 265 nM, respectively) due to its high affinity for H₂ relative to formate (K_s values of ~6 μM and ~0.22 mM, respectively [61, 62]). This preferential use of H₂ by *M. hungatei* possibly also contributed to the reduction in the syntrophic growth rate of this coculture pairing. Remarkably, *D. alaskensis* G20 adapted to its partner's preference for H₂ by shifting the ratio of electron carriers used for interspecies metabolite exchange (81% as formate with *M. maripaludis* versus only 63.7% with *M. hungatei*). The low K_s values of *M. hungatei* for H₂ and formate relative to other hydrogenotrophic methanogens (H₂ K_s , 14 to 66 μM [61, 63, 64]; formate K_s , 0.58 mM [62]) suggests that *M. hungatei* might have a competitive advantage when substrates supplied by *Desulfovibrio* are below the saturation constant(s) of the other species (i.e., at low dilution rates). Thus, *M. maripaludis* might be adapted to an opportunistic lifestyle, growing rapidly in the presence of occasional high nutrient concentrations, whereas *M. hungatei* is adapted to life under continuous nutrient limitation (a copiotrophic versus an oligotrophic lifestyle [18, 65]).

Transcription and mutant phenotype analyses revealed a significant influence of both energy source availability (as controlled by dilution rate) and methanogenic partner on the electron transfer enzyme system of syntrophically grown *D. alaskensis* G20. The data now serve to refine and extend our previous metabolic model for the syntrophic growth of strain G20 (15) and highlight the flexibility of *Desulfovibrio* species to adapt enzymatically and energetically to natural fluctuations in the environment (Fig. 8). At high growth rates sustained by association with *M. maripaludis*, strain G20 links lactate oxidation with periplasmic H⁺/CO₂ reduction by coupling the oxidation of high redox potential DsrC_{red} (thiol/disulfide redox pair, lactate oxidation derived) energetically with the oxidation of the low redox potential Fd-II_{red} (pyruvate oxidation derived). This involves a flavin-based electron confurcation process and a MQ redox loop-mediated reverse electron flow by the membrane-bound Qmo and Qrc complexes. The Qmo-MQ-Qrc loop serves as the main electron transfer system permitting high lactate oxidation rates under elevated H₂ levels (460 ppm, on average, in the gas phase), thereby enhancing methanogenic growth. Although energetically costly (coupling of DsrC_{red} oxidation to Fd_{red} and reverse MQ cycling), the high lactate throughput presumably generates sufficient energy by substrate level phosphorylation to sustain growth.

The transcription profiles also revealed a shift in the gene expression of METCs in response to lower energy availability (lower dilution rate or pairing with *M. hungatei*). The Qmo complex was significantly downregulated at a low growth rate, whereas the Hmc complex was slightly upregulated, pointing to a possible shift in electron channeling through these two electron transfer sys-

tems. The Hmc closely resembles the DsrMKJOP complex (66, 67) in terms of subunit structure, composition, and location (39, 68), consisting of one periplasmic cytochrome *c* subunit (HmcA) and [4Fe-4S] cluster protein (HmcB), two integral membrane proteins of the NrfD (HmcC) and NarI (HmcE) families, and a cytoplasmic HdrD homologue (HmcF) (69). Being reduced by TpIc₃ (70), the Hmc was initially postulated to be involved in H₂ uptake of *Desulfovibrio* species (41, 71), but expression levels are low during respiratory growth with H₂ (72, 73) and lactate (38). Our recent work indicates a functional role in electron transfer during syntrophic growth of *D. vulgaris* Hildenborough (15, 26). Thus, based on the data presented here, we postulate that under energy source limitation the Hmc complex is increasingly involved in the reoxidation of the LdhAB-1 derived DsrC_{red} and transfer of electrons via TpIc₃/Ssc to the periplasmic W-formate dehydrogenase Fdh-1 and hydrogenases (Hyn or Hys) in strain G20. This linkage would be achieved using an MQ-based reverse redox loop involving an HmcEF and HmcABCD module operating as menaquinone reductase and menaquinol oxidase, respectively. The HmcE is homologous to the cytochrome *b* integral membrane subunit HdrE (21, 74), while HmcC belongs to the NrfD family of proteins that interact with the quinone pool without the involvement of heme b, as for QrcD (17). Indeed, a bidirectional electron transport between cytoplasm and periplasm was demonstrated for the complex (71, 75). The shift in electron channeling from the Qmo-Qrc to the HmcEF-HmcABCD redox loop for the reoxidation of the reduced DsrC electron carrier might be energetically beneficial since it preserves the low redox potential Fd_{red} for coupling periplasmic H⁺/CO₂ reduction with cytoplasmic lactate oxidation. A proton-consuming cytoplasmic H₂ generation from Fd_{red} mediated by the putative monomeric [Fe-Fe] hydrogenase (Dde_0725; slightly upregulated under low-energy conditions) could contribute to proton motive force and allow additional chemiosmotic energy conservation besides the substrate level phosphorylation step. Thus, the Hmc complex and monomeric [Fe-Fe] hydrogenase likely represent a linked system that becomes increasingly important (besides the Qmo-MQ-Qrc loop) under low-energy conditions when maintenance energy becomes an increasing component of total energy demand (76). An additional pathway for thiol/disulfide conversion reactions supporting syntrophic growth was also suggested by coculture growth results of mutants for the uncharacterized HdrD homologue, Dde_0312 (additional FAD group containing), and the NADH oxidase, Nox. In association, both proteins could operate as a soluble flavin-based electron-bi-/confurcating enzyme that reversibly converts any remaining ferredoxin, thiol/disulfide redox pairs, and NADH in the cytoplasm. NADH could be reoxidized by a Qmo complex-associated Nox (77). The upregulation of the Ssc encoding gene during syntrophy irrespective of growth rate and methanogenic partner is indicative of an as-yet-uncharacterized functional role in linking electron transport between METCs and periplasmic enzymes in addition to TpIc₃. Although its reduction by periplasmic dehydrogenases has been experimentally confirmed (78), a direct interaction with the Qrc and Hmc complex has not yet been demonstrated.

Despite the overall similarity of their transcriptional profiles, 22 genes showed significant transcriptional level variation between both coculture pairings at a low growth rate. *D. alaskensis* G20 adapted to the metabolism of the respective methanogenic partner, as shown by the differing gene expression and mutant

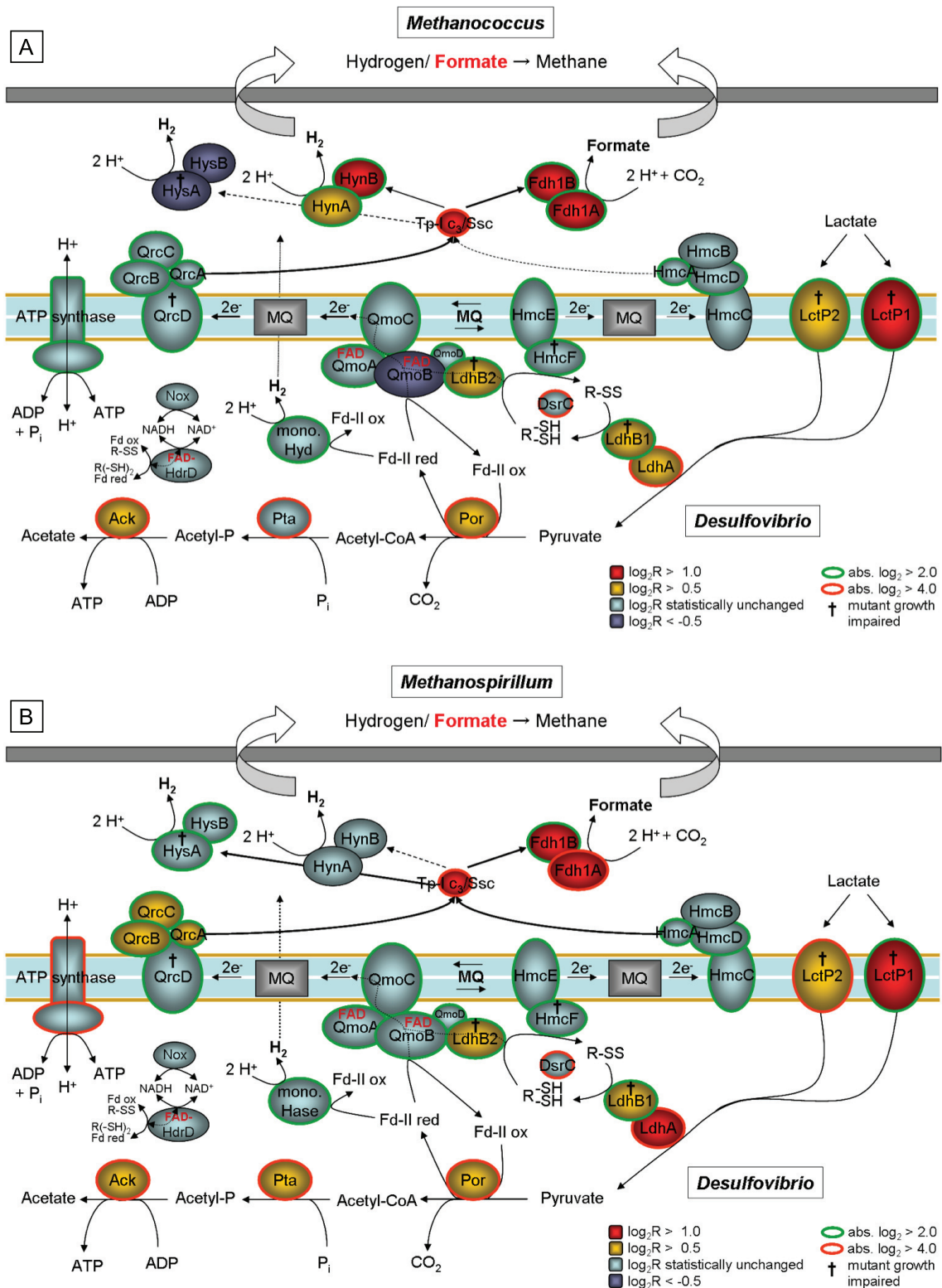


FIG 8 Proposed metabolic models for syntrophic growth of *D. alaskensis* G20 at the low growth rate of 0.027 h^{-1} with either *M. maripaludis* (A) or *M. hungatei* (B). Colors indicate transcriptional changes in individual genes during coculture growth relative to a sulfate-limited monoculture (the main electron transfer pathway in respective *Desulfovibrio* species is highlighted with red arrows); *D. alaskensis* G20 mutants with no growth in coculture on lactate are marked with the symbol “†” (the coculture mutant phenotypes with *M. maripaludis* were determined in an earlier study [15]). Abbreviations used: LctP, lactate permease; Ldh, lactate dehydrogenase; R-(SH)₂ and R-SS, reduced and oxidized form of unknown thiol/disulfide redox pair (probably DsrC) interacting as electron carrier with Ldh; Por, pyruvate:ferredoxin oxidoreductase; Fd_{red} and Fd_{ox}, reduced and oxidized form of ferredoxin; Qmo, quinone-interacting membrane-bound oxidoreductase; MQ, menaquinone; Qrc, quinone reduction complex; Pta, phosphate acetyltransferase; Ack, acetate kinase; Aor, aldehyde:ferredoxin oxidoreductase; Adh, alcohol dehydrogenase; Hdr-Flox, putative electron-bifurcating complex coupling NADH oxidation with reduction of thiol/disulfide redox pair (probably DsrC) and ferredoxin and vice versa (electron confurcation); Hmc, high-molecular-weight cytochrome c complex; Hyn, [NiFe] hydrogenase; Fdh, formate dehydrogenase; Hyd, [Fe] hydrogenase; TplC₃, type I cytochrome c₃; Ssc, split-soret cytochrome c; CoA, coenzyme A.

coculture phenotypes of the periplasmic hydrogenases. Strain G20 possesses four periplasmic hydrogenases (HysAB, HynAB, and HydAB-1/-2) with unique kinetic properties (e.g., different substrate affinities) that have been demonstrated to be differently regulated in *D. vulgaris* Hildenborough in dependence on medium composition and growth conditions during sulfate respiration (72, 79, 80). Both [NiFe] hydrogenase (Hyn) and [NiFeSe] hydrogenase (Hys) were suggested to function bidirectionally (H_2 oxidizing/producing) (81). Our results show that irrespective of growth rate, Hyn seems to be the main enzyme responsible for periplasmic H_2 production in *D. alaskensis* G20 when grown syntrophically with *M. maripaludis*. The recently described Hyn-Qrc-TpIc₃ supercomplex formation (82) likely ensures an optimized electron transfer required for periplasmic H_2 production at elevated levels (the average concentration of dissolved H_2 is 265 nM). Since *M. hungatei* keeps the dissolved H_2 in coculture at a lower level (70 nM, on average, reflecting its low K_s value [61, 83]), strain G20 responds by using a different periplasmic hydrogenase (Hys) for H_2 generation. Notably, under all syntrophic conditions, the W-formate dehydrogenase Fdh-1 appears to be the main enzyme for the periplasmic formate generation in strain G20, most likely because of tungsten availability in the media; the same effect on transcription has been reported for its respiratory growth on formate (84, 85).

Among the 22 genes with a significant change in expression between the coculture pairings, 11 belong to the pool of putatively “*M. maripaludis* syntrophy-relevant” genes that are exclusively upregulated during syntrophic growth with this methanogen (irrespective of energy availability). These include genes for the 2-hydroxyglutaryl-CoA dehydratases (Dde_0089 and Dde_0090), which catalyze a key step in the 2-hydroxyglutarate pathway in which glutamate is fermented to ammonia, CO_2 , acetate, butyrate, and H_2 (60). Operating in reverse, this process might be an additional pathway in *D. alaskensis* G20 for consuming the end products of lactate fermentation, with glutamate being excreted and transported to *M. maripaludis* as another exchanged metabolite. Recently, an interspecies alanine transfer has been reported for *D. vulgaris* Hildenborough-*M. maripaludis* cocultures (86).

In summary, the comparative transcriptional analyses, complemented by individual mutant growth experiments and physiological data, established distinctive differences in the population structures, electron transfer systems, ratios of interspecies electron carriers (H_2 /formate), enzyme systems, and energy conservation mechanisms that sustain the syntrophic growth of *D. alaskensis* G20 when grown under different environmental conditions (energy source availability, different methanogenic partner). The results emphasize the high enzymatic and metabolic flexibility of syntrophic *D. alaskensis* G20, which ensures its ability to adapt to the frequent fluctuations of growth conditions in nature. Adaptive strategies may also involve architectural changes, e.g., cell attachment and aggregation (observed in coculture at low growth rate), which might allow for efficient metabolite transfer to the methanogen and therefore more energy-efficient energy coupling.

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