Syngas as Electron Donor for Sulfate and Thiosulfate Reducing Haloalkaliphilic Microorganisms in a Gas-Lift Bioreactor

Caroline M. Plugge\(^1\), João A. B. Sousa\(^1\), Stephan Christel\(^3\), Mark Dopson\(^3\), Martijn F. M. Bijmans\(^2\), Alfons J. M. Stams\(^1\) and Martijn Diender\(^1\)

Abstract: Biodesulfurization processes remove toxic and corrosive hydrogen sulfide from gas streams (e.g., natural gas, biogas, or syngas). To improve the efficiency of these processes under haloalkaline conditions, a sulfate and thiosulfate reduction step can be included. The use of H\(_2\)/CO mixtures (as in syngas) instead of pure H\(_2\) was tested to investigate the potential cost reduction of the electron donor required. Syngas is produced in the gas-reforming process and consists mainly of H\(_2\), carbon monoxide (CO), and carbon dioxide (CO\(_2\)). Purification of syngas to obtain pure H\(_2\) implies higher costs because of additional post-treatment. Therefore, the use of syngas has merit in the biodesulfurization process. Initially, CO inhibited hydrogen-dependent sulfate reduction. However, after 30 days the biomass was adapted and both H\(_2\) and CO were used as electron donors. First, formate was produced, followed by sulfate and thiosulfate reduction, and later in the reactor run acetate and methane were detected. Sulfide production rates with sulfate and thiosulfate after adaptation were comparable with previously described rates with only hydrogen. The addition of CO marginally affected the microbial community in which Tindallia sp. was dominant. Over time, acetate production increased and acetogenesis became the dominant process in the bioreactor. Around 50% of H\(_2\)/CO was converted to acetate. Acetate supported biomass growth and higher biomass concentrations were reached compared to bioreactors without CO feed. Finally, CO addition resulted in the formation of small, compact microbial aggregates. This suggests that CO or syngas can be used to stimulate aggregation in haloalkaline biodesulfurization systems.

Keywords: acetogenesis; biomass retention; carbon monoxide; carrier material; formate; haloalkaline biodesulfurization; hydrogen; sulfate reduction; thiosulfate reduction

1. Introduction

Hydrogen sulfide is a toxic and corrosive compound present in gas streams such as natural gas or biogas. In biodesulfurization systems, sulfide is removed from the gas and elemental sulfur is formed.
Biodesulfurization systems that produce elemental sulfur operated under haloalkaline conditions have been studied and applied for more than 25 years [1,2]. The efficiency of converting sulfide to elemental sulfur in laboratory experiments is close to 97% [2] and models suggest that 98% efficiency is possible [3,4]. The remaining part of the sulfide is biologically oxidized to sulfate or chemically to thiosulfate. These compounds are soluble and accumulate in the system causing a pH decrease that has a negative impact because high pH (>8.5) is required for efficient hydrogen sulfide gas absorption. To solve this problem, part of the liquid content of the system is removed, generating a bleed stream with high pH and high salinity and containing sulfate and thiosulfate [1].

This bleed stream can be treated by biological reduction of sulfate and thiosulfate to sulfide in a separate bioreactor. This allows the recirculation of the bleed stream into the sulfide-oxidizing bioreactor of the biodesulfurization process [5,6]. Previous studies showed that this is possible under haloalkaline conditions in different bioreactor types using a variety of electron donors [7–9]. Hydrogen gas (H₂) is considered a suitable electron donor for biodesulfurization systems applied for natural gas as it can be produced on-site by gas reforming of methane. However, syngas is produced in the gas reforming process that consists mainly of H₂, carbon monoxide (CO), and carbon dioxide (CO₂). Purification of syngas to obtain pure H₂ implies higher costs because of additional post-treatment. The application of syngas as an electron donor without removal of CO is thus preferred and was previously applied successfully at neutral pH and low salinity conditions [10]. On the other hand, CO is known to be toxic to microorganisms [11,12]. Because of its strong reductive properties (E° = –520 mV), CO reacts with metals in the active centres of enzymes [13]. Via this mechanism CO is known to inhibit both [Fe-Fe] and [Ni-Fe] hydrogenases [14] of which the former appears more sensitive to inhibition by CO [15,16]. Presence of CO can therefor inhibit H₂-driven sulfate reduction, and other potential hydrogen-dependent metabolisms such as formate, acetate, and methane production.

Despite its toxicity, several types of microbial metabolism are able to use CO as substrate. Acetogens are relatively well-known CO-utilizing microorganisms, and their use for production of fuels and chemicals from syngas is currently considered at an industrial scale [17]. CO-driven metabolism is also found in sulfate reducers and methanogens [11,12,18]. Sulfate reducers are mainly found to convert CO via the water-gas shift reaction to produce H₂ (Table 1, Equation (1)), that is subsequently used as electron donor for sulfate reduction [19–21]. Previously, it was shown that sulfate-reducing biomass from bioreactors at neutral and high temperature conditions could be adapted to CO as substrate [18,22]. Additionally, it was observed that CO had an effect on the formation of granules, making them smoother [22]. This was probably caused by the development of different layers on the granule, with a potential external layer containing acetogenic bacteria that convert CO while the inner layer contained sulfate-reducing bacteria. These sulfate-reducing bacteria likely used the produced acetate as an additional carbon source [22]. Granules with a smoother shape settle better than irregular shaped aggregates, and in this way CO could induce the formation of aggregates with enhanced settling properties.

To date, the effect of CO on haloalkaphilic microorganisms has not been studied in pure or mixed cultures. In this study, we investigated syngas as an electron donor for a sulfate and thiosulfate-reducing bioreactor operated under haloalkaline conditions. The impact of CO on sulfate and thiosulfate reduction as well as on formate, acetate, and methane production was investigated. In addition, biomass attachment to sand as well as microbial aggregation was studied in order to understand if CO can be used to influence the adhesion of microorganisms similar to what was observed at neutral pH and low salinity conditions [22]. The feasibility of industrial application of syngas as electron donor for sulfate and thiosulfate reducing bioreactors was evaluated and the main advantages and disadvantages of the use of syngas versus pure hydrogen gas are discussed.

\[
\text{CO} + 2\text{H}_2\text{O} \rightarrow \text{HCO}_3^- + \text{H}_2 + \text{H}^+ \tag{1}
\]

\[
4\text{CO} + 4\text{H}_2\text{O} \rightarrow \text{C}_2\text{H}_3\text{O}_2^- + 2\text{HCO}_3^- + 3\text{H}^+ \tag{2}
\]
4H₂ + 2HCO₃⁻ + H⁺ → C₂H₃O₂⁻ + 4H₂O  
4CO + SO₄²⁻ + 4H₂O → 4HCO₃⁻ + HS⁻ + 3H⁺  
4H₂ + SO₄²⁻ + H⁺ → HS⁻ + 4H₂O  
CO + H₂O → HCO₂⁻ + H⁺  
H₂ + HCO₃⁻ → HCO₂⁻ + H₂O  
4CO + 5H₂O → CH₄ + 3HCO₃⁻ + 3H⁺  
4H₂ + HCO₃⁻ + H⁺ → CH₄ + 3H₂O

Table 1. Gibbs free energy under the bioreactor actual conditions (ΔG') for different CO and H₂ consuming reactions at starting and continuous operation conditions.

| Reaction | ΔG' Starting (kJ mol⁻¹) | ΔG' Continuous Operation (kJ mol⁻¹) | Equation No. |
|----------|-------------------------|------------------------------------|--------------|
| CO + 2H₂O → HCO₃⁻ + H₂ + H⁺ | -22.7 (±10.5) | -23.1 (±10.5) | (1) |
| 4CO + 4H₂O → C₂H₃O₂⁻ + 2HCO₃⁻ + 3H⁺ | -199.6 (±26.6) | -182.9 (±26.6) | (2) |
| 4H₂ + 2HCO₃⁻ + H⁺ → C₂H₃O₂⁻ + 4H₂O | -108.9 (±26.9) | -90.5 (±26.9) | (3) |
| 4CO + SO₄²⁻ + 4H₂O → 4HCO₃⁻ + HS⁻ + 3H⁺ | -240.6 (±34.3) | -218.8 (±34.3) | (4) |
| 4H₂ + SO₄²⁻ + H⁺ → HS⁻ + 4H₂O | -150.0 (±25.0) | -126.4 (±25.0) | (5) |
| CO + H₂O → HCO₂⁻ + H⁺ | -38.0 (±6.5) | -32.3 (±6.5) | (6) |
| H₂ + HCO₃⁻ → HCO₂⁻ + H₂O | -15.2 (±8.9) | -9.2 (±8.9) | (7) |
| 4CO + 5H₂O → CH₄ + 3HCO₃⁻ + 3H⁺ | -215.4 (±30.6) | -198 (±30.6) | (8) |
| 4H₂ + HCO₃⁻ + H⁺ → CH₄ + 3H₂O | -124.4 (±25.1) | -105.6 (±25.1) | (9) |

Conditions used for calculations: pH 9, 1.5 ionic strength, 825 mM HCO₃⁻. Starting conditions: 50 mM sulfate, 1 mM sulfide, 1 mM acetate, 1 mM formate, 150 mbar CO, 850 mbar H₂, 10 mbar methane. Continuous operation conditions (average of days 190 to 216): 5.7 mM sulfate, 36.9 mM sulfide, 37.8 mM acetate, 4.5 mbar formate, 88 mbar CO, 333 mbar H₂, 575 mbar methane. Error for each ΔG' estimation is presented in brackets because formation energy of the compounds under high pH and ionic strength are estimated based on models.

2. Materials and Methods

2.1. Bioreactor Set-Up

A 4.4-L glass gas lift reactor with an internal three phase separator was used [7] (Figure S1). The temperature was maintained at 35 °C using a water jacket connected to a thermostat bath (DC10-P5/U, Haake, Dreiich, Germany). The influent feeding was performed by a membrane pump (Stepdos 08 RC, KNF-Verder, Utrecht, The Netherlands). The H₂, CO, and CO₂ gas supply was controlled using digital mass flow controllers (F-201CV-020-AGD-22-V and F-201CV-020-AGD-22-Z, Bronkhorst, Ruurlo, the Netherlands). The gas was recycled using a vacuum pump (Laboport®, KNF, Trenton, NJ, USA) and the gas flow was measured using a calibrated flow meter (URM, Kobold, Amsterdam, The Netherlands). A pH and a redox potential sensor (CPS11D and CPS12D, Endress + Hauser, Naarden, the Netherlands) connected to a controller (Liquiline CM44x, Endress + Hauser, Naarden, the Netherlands) were used to monitor the conditions inside the reactor. The pH was controlled at pH 9 by supplying CO₂ via the mass flow controller. As biomass carrier material, 0.5 L of acid washed sea sand with a particle size of 0.1–0.3 mm (VWR, Amsterdam, The Netherlands) was added as described in [23].

2.2. Inoculum

The inoculum was a 1/1 ratio mixture from a sulfate and thiosulfate reducing gas-lift bioreactors fed with H₂ and CO₂ [7,8,23]. For the inoculation, 100 mL of concentrated biomass from each bioreactor was used, obtained by centrifuging the content of the previously operated bioreactors at the end of the experiments.
2.3. Medium

A mineral medium was used that was buffered with sodium carbonate and sodium bicarbonate at pH 9 ± 0.05, that contained a total of 1.5 M Na⁺. The medium composition was as follows: Na₂CO₃ (33.6 g L⁻¹), NaHCO₃ (69.3 g L⁻¹), KHCO₃ (1 g L⁻¹), K₂HPO₄ (1 g L⁻¹), NH₄Cl (0.27 g L⁻¹), MgCl₂.6H₂O (0.1 g L⁻¹), CaCl₂.2H₂O (0.01 g L⁻¹), and 10 mL L⁻¹ of vitamin solution [24]. Two trace element solutions were added [7]. As electron acceptors, 7.1 g (25 mM) of sodium sulfate and 3.95 g (12.5 mM) of sodium thiosulfate were added.

2.4. Experimental Design

After filling it with medium, the bioreactor was flushed with H₂ gas overnight to lower the redox potential. The gas recirculation was set at 5 L min⁻¹ (±0.5). Then the H₂ gas supply was set at 20 mL.min⁻¹ and the pH control (set at 9) was turned on. The inoculum was added (time 0) that initiated the start-up phase. Hereafter, the only parameter changed was the gas composition by adding CO (Table 2).

| Period (Days) | Mode          |
|--------------|---------------|
| Start-up     | 0–6           |
| 0% CO        | 7–45          |
| 5% CO        | 46–123        |
| 15% CO       | 124–216       |
| 60% CO spike * | 217–235   |

* See experimental design for a detailed description.

On day 217 a CO spike experiment was performed, where the CO supply was increased to 60% of the gas phase and maintained for 48 h before being switched to 15% on day 219 until the end of the experiment.

2.5. Batch Experiments

Two batch experiments were performed to assess the metabolic capacity of the biomass at different stages of the bioreactor operation. All experiments were performed with the same medium composition used for the bioreactor.

To study the activity of the biomass after CO addition, 125 mL of biomass containing bioreactor liquid from day 54 of operation was transferred to 250 mL serum bottles. The serum bottles headspace was replaced with N₂ gas. CO was added to make three different gas compositions: 30, 55, and 80% CO (at 1 bar of total pressure). The bottles were incubated at 37 °C and shaking at 150 rpm for 14 days. Each CO condition was tested in triplicate. For each sampling point, 5 mL of liquid content and 1 mL of gas content were collected from the bottles.

To study the effect of formate and acetate on sulfate/thiosulfate reduction in adapted biomass in the presence of CO, 120 mL serum bottles were filled with 70 mL fresh medium and N₂ headspace. Three different conditions were tested: no electron donor, with 25 mM sodium formate, and with 12.5 mM sodium acetate. Biomass containing bioreactor liquid (50 mL) was collected at day 90, centrifuged for 10 min at 10,000× g, re-suspended with 10 mL buffer (pH 9, 1.5 M Na⁺), and added as inoculum. CO was added at approximately 25% CO in the gas phase (1.4 bar total pressure). Each set of batch cultures was tested in triplicate. For each sampling point, 2 mL of liquid content and 1 mL of gas content were collected from the bottles.
2.6. Analytical Procedures

Liquid samples for volatile fatty acids, sulfate, thiosulfate, and sulfide analysis were prepared and analyzed as described previously [7]. The H₂, CO, CO₂, N₂, and CH₄ in the gas phase were quantified by gas chromatography using a CP-4900 microGC (Varian, Palo Alto, CA) as previously described [7].

To measure the biomass concentration, the sand-attached biomass was separated from the suspended biomass by settling for 30 s and both fractions were transferred to a new separate tube. One milliliter of sand was collected and the samples were washed three times with a carbonate/bicarbonate buffer with lower salinity (LS buffer; pH 9, 0.5 M Na⁺ instead of 1.5 M Na⁺). In subsequent washing steps, sand was separated from the buffer by 30 s settling. Two milliliters of the suspended biomass was centrifuged (10 min, 10,000× g) and washed three times with LS buffer. Finally, the total nitrogen content was determined using a cuvette test (LCK238, Hach Lange, Düsseldorf, Germany).

The particle size of the bioreactor content (including sand) was measured using laser measurement in a particle size and shape analyzer (Eyetech, Doner technologies, Or Akiva, Israel) with the Dipa 2000 software (Doner technologies, Or Akiva, Israel). Each sample was analyzed in triplicate and each measurement was performed continuously for 120 s with stirring. Microscopy pictures were taken using a light microscope (DMI6000B, Leica, Biberach, Germany). The sand particles were not measured because of settling in the mixing chamber, being only small aggregates analyzed for particle size.

2.7. Scanning Electron Microscopy

Biomass samples were fixed in 2.5% (w/v) glutaraldehyde overnight at 4 °C. The fixed samples were separated and washed following the same procedure for attached fraction and suspended fraction described above for biomass measurement. Then the samples were dehydrated in a series of ethanol solutions, (10%, 25%, 50%, 75%, 90% and twice with 100%) with 20 min in each step and then dried in a desiccator. The samples were coated with gold and analyzed in a JEOL JSM-6480LV scanning electron microscope (JEOL Benelux, Nieuw-Vennep, The Netherlands).

2.8. DNA Isolation and Bacteria Community Profiling

Samples (10 mL) for DNA analysis were collected on days 0, 46, 96, 123, 216, and 218 of bioreactor operation and stored at −80 °C. The samples were separated into the attached and suspended fractions and washed following the same protocol as for biomass measurements. Total genomic DNA from the suspended fraction of all samples and attached fraction of samples from day 123 was extracted using the PowerBiofilm™ DNA Isolation Kit (MoBio, Carlsbad, CA, USA) following the manufacturer’s instructions. DNA was stored at −20 °C.

A fragment of the 16S rRNA gene of bacteria, including the V3-V5 regions, was amplified with primers 341F and 805R [25]. Sequences were submitted to the ENA database (http://www.ebi.ac.uk/ena) under the accession number PRJEB11708. The PCR protocol and sequencing using the Illumina Miseq platform were performed as previously described at the Science for Life Laboratory, Sweden (www.scilifelab.se) [26,27]. The sequencing data were processed with the UPARSE pipeline and annotated against the SINA/SILVA database (SILVA 119) [28,29]. Finally, the data were analyzed using Explicet 2.10.5 [30].

2.9. Calculations

Calculations and assumptions:
Thermodynamic calculations under actual bioreactor conditions were performed using the online tool eQuilibrator 2.0 (equilibrator.weizmann.ac.il) [31]. The relative electron donor use calculations were based on the molar quantities indicated by the reactions described in Table 1. The assumptions used in the calculations were as follows: (i) Evaporation does not cause a major loss of liquid from the bioreactor because it is a closed system; (ii) the bioreactor liquid volume was assumed to be constant.
during the operation meaning that the liquid flow going into the bioreactor was equal to the liquid flowing out of the bioreactor \( Q_{\text{in}} = Q_{\text{out}} \); (iii) accumulation of sulfur compounds by incorporation in biomass and formation of sulfide precipitates was assumed to play a negligible role. This was due to the high sulfate and thiosulfate concentration in the influent compared to the low concentration of metals added to the medium; (iv) hydrogen used for biomass synthesis was assumed to play a negligible role in the bioreactor; (v) the N fraction value of 0.2 was used to calculate biomass concentration based on total N, following the biomass molecular formula: \( C_{1.8}H_{1.8}O_{0.5}N_{0.2} \); and (vi) the attached biomass fraction was composed of the sand particles with attached microorganisms and microbial aggregates without support material.

\[
C_x = \frac{C_{\text{total}N}}{0.2} M_x \\
C_{tx} = C_{x,sp} + C_{x,ar} \\
r_{vs} = \frac{Q C_{s,in} - Q C_{s,out}}{V_r} \\
r_{sp} = \frac{Q C_{p,out} - Q C_{p,in}}{V_r}
\]

3. Results and Discussion

3.1. Bioreactor Performance

The use of syngas as an electron donor for haloalkaliphilic sulfate and thiosulfate reducing microorganisms was possible up to 15% CO. However, a period of adaptation of the biomass to CO was required. All \( H_2 \)-driven microbial processes, including sulfate/thiosulfate reduction and formate production, were inhibited by the presence of 5% CO (Figure 1A,B). This is in agreement with similar studies performed at neutral conditions where the sulfate reduction activity decreased from 140 mmol L\(^{-1}\) d\(^{-1}\) to 98 mmol L\(^{-1}\) d\(^{-1}\) with 5% CO [22]. During this inhibition period (day 47 to 75), thiosulfate disproportionation to sulfate and sulfide was not inhibited. This can be observed by the increase of sulfate and sulfide concentrations in an approximately 1:1 ratio (Figure 1). During this period, in which no acetate was formed, the pH decreased and the CO\(_2\) fraction in the gas phase increased that was indicative of the water-gas shift reaction (Equation (1), Table 1, and Figure 1C). After 38 days of operation with 5% CO (day 84), the sulfate/thiosulfate reduction activity recovered. This activity was not further affected by an increase to 15% CO on day 124 (Figure 1A). Thiosulfate was completely reduced while 87 ± 3% of the sulfate was reduced during the stable phase with 15% CO (day 200 to 216). The sulfidogenic rate was similar to that achieved in a study with only \( H_2 \) \((42.3 \pm 2.2 \text{ mmol L}^{-1} \text{ d}^{-1})\), when applying the same sulfate/thiosulfate loading rate [23]. The sulfidogenic rates \((r_{vs})\) achieved almost matched the loading rates of sulfate and thiosulfate (Figure 2A). This indicated that sulfidogenic rates could be higher if loading of sulfate and thiosulfate was increased as described for other sulfate and thiosulfate reducing bioreactors (Table 3). Moreover, increasing the length of the different stages may contribute to a further increase in the sulfidogenic rates.
Besides sulfate and thiosulfate reduction, acetate, formate, and methane were produced (Figure 1a,b and Figure 2b). Acetate production started after the inhibition period on day 76, probably using hydrogen and/or CO as electron donors. It was not possible to distinguish if either H₂ or CO were specifically used as H₂ and CO were simultaneously consumed (Figure 2c). However, looking at the overall electron donor consumption (H₂ + CO), acetate production was the main electron sink in the system. Approximately 49% (±5%) of the supplied electron donors in the stable phase with 15% CO
Besides sulfate and thiosulfate reduction, acetate, formate, and methane were produced (Figure 1a,b and Figure 2b). Acetate production started after the inhibition period on day 76, probably using hydrogen and/or CO as electron donors. It was not possible to distinguish if either H₂ or CO were specifically used as H₂ and CO were simultaneously consumed (Figure 2c). However, looking at the overall electron donor consumption (H₂ + CO), acetate production was the main electron sink in the system. Approximately 49% (±5%) of the supplied electron donors in the stable phase with 15% CO was converted to acetate (Figure 3). This was considerably higher than reported in bioreactors fed with H₂ or formate as electron donors [9,23]. Furthermore, high acetate concentrations might have contributed to increased biomass growth compared to previous results (Table 3, Figure 4). This effect of acetate was observed when no other carbon sources were added to the media [23].
Microorganisms 2020, 8, x FOR PEER REVIEW 11 of 19

Table 3. Conditions and performance of sulfate and thiosulfate reducing bioreactors operated under haloalkaline conditions.

|                                    | This Study              | Sousa et al. 2020       | Zhou and Xing 2015    | Sousa et al. 2017       |
|------------------------------------|-------------------------|-------------------------|-----------------------|-------------------------|
| Reactor type                       | Gas lift with 3 phase   | Gas lift with 3 phase   | Anaerobic filter      | Gas lift with 3 phase   |
| separator                          | separator               | separator               |                       | separator               |
| e⁻ acceptor                        | Sulfate/thiosulfate     | Sulfate/thiosulfate     | Sulfate               | Thiosulfate             |
| e⁻ donor                           | H₂/CO                   | H₂                      | Formate               | H₂                      |
| pH                                 | 9                       | 9                       | 9.5                   | 9                       |
| Na⁺ conc. (M)                      | 1.5                     | 1.5                     | 1                     | 1.5                     |
| Temperature (°C)                   | 35                      | 35                      | 37                    | 35                      |
| HRT (d)                            | 1                       | 1                       | 1                     | 1.7                     |
| Max loading rate (mmol L⁻¹ d⁻¹)    | 50                      | 100                     | 88.5                  | 50                      |
| Sulfidogenic rate (mmol L⁻¹ d⁻¹)   | 46.8 (±0.8)             | 85.45 (±3)              | 85.05 (±0.2)          | 28.7 (±0.8)             |
| Side products                      | Formate/Acetate/Methane | Formate/Acetate/Methane | Acetate               | Formate                 |
| Biomass conc. (mg L⁻¹)             | 197 (±39)               | 127 (±41)               | N.D.                  | 14 (±2.2)               |

N.D.—No data available.

Figure 3. Fraction of electron donor (H₂ + CO) used by the different metabolisms: sulfate and thiosulfate reduction and formate, acetate and methane production. For calculation, all values of electron donors and acceptors were converted to mmol/d. Vertical dashed lines represent the beginning of CO experiments: 5% CO (1st), 15% CO (2nd), and 60% CO spike (3rd).

Even though high and stable sulfate and thiosulfate reduction rates were achieved with 15% CO in the gas supply (Figures 1 and 2), temporary CO increases could still have a negative and disturbing effect on the bioreactor operation. To study this, 60% CO was fed to the bioreactor from day 217 until day 219 before returning the CO to 15%. The addition of 60% CO led to a steep decrease in sulfate reduction, formate and acetate production, and biomass concentration (Figures 1 and 4). However, thiosulfate disproportionation was not affected and methane production even increased after the 60% CO spike (Figures 1 and 2). This indicated that either methanogenesis could compete for H₂ after inhibition of the other hydrogenotrophic microorganisms or that the methanogens present were using CO as the electron donor. Generally, methanogens are rapidly inhibited upon CO exposure, and are considered more sensitive to CO compared to acetogens [32,33]. However, methanogenesis with CO has never been reported under haloalkaline conditions and it is thermodynamically favorable at the conditions present in the bioreactor (Table 1).
Figure 3. Fraction of electron donor (H₂ + CO) used by the different metabolisms: sulfate and thiosulfate reduction and formate, acetate and methane production. For calculation, all values of electron donors and acceptors were converted to mmol/d. Vertical dashed lines represent the beginning of CO experiments: 5% CO (1st), 15% CO (2nd), and 60% CO spike (3rd).

Figure 4. Biomass in the bioreactor. (A) Total biomass concentration (Cₜₓ) in the bioreactor during its operation. (B) Particle diameter distribution of the biomass on days 46 (without CO), 99 (5% CO), 125 (just after increase to 15%), and 216 (stable phase with 15% CO).

3.2. Inhibition by CO

As observed in previous studies, hydrogen-driven metabolism can be rapidly inactivated upon CO exposure, and is usually thought to be related to hydrogenase inhibition [34,35]. The results here show that upon exposure of the biomass to CO, H₂-driven sulfate reduction activity dropped rapidly, as well as formate and acetate production (Figure 1). This is potentially related to the inhibition of hydrogenases. The observation that thiosulfate disproportionation remained active supports this hypothesis, as thiosulfate disproportionation does not require the action of hydrogenases. However, some hydrogenases have been reported to be highly resistant to CO like some [Ni-Fe]-hydrogenases and a single [Fe-Fe] hydrogenase [14,36–38]. Microorganisms capable of producing such CO-resistant hydrogenases would be able to metabolize H₂ in CO-rich environments. Some hydrogenogenic organisms operating the water gas-shift reaction employ such hydrogenases [37], explaining potential hydrogenogenic activity during the CO-inhibited phase.

Despite the seeming inhibition of many hydrogenase-dependent conversions, biomass levels remained relatively constant during this phase, indicating microbial growth (Figure 4). Besides thiosulfate disproportionation and the water-gas shift reaction, acetate production from CO might also have occurred during this phase. Even though acetate was under the detection limit, its production in lower amounts might have supported the biomass growth. Slow oxidation of CO was observed in batch bottles incubated with biomass from the period just after CO was added (Figure 5). This suggests that the microbial community present during this operation period had the ability to convert CO.
With further analysis, ~99% of these sequences were closely related to the genus Tindallia with glucose as substrate, that was first converted into ethanol, lactate, acetate and formate in the lower region of the bioreactor. Several related bacteria were previously detected as dominant bacteria in H₂-driven microbial processes, at day 76 acetogenic activity was observed that correlated with the removal of CO from the headspace (Figures 1 and 2). Several acetogens are able to utilize CO as substrate via the Wood-Ljungdahl pathway [34]. In volcanic environments that contain high levels of CO, carboxydotrophic organisms play a role in the removal of CO, thereby creating a viable environment for non-CO-tolerant microbes [39]. The CO-driven acetogenic activity observed in the bioreactor could have created favorable conditions for hydrogenotrophic sulfate reducers and formate producers. Additionally, adaptation to CO by microorganisms already present in the biomass, such as via production of CO-resistant enzymes, also might have contributed to the restoration of activity. Some hydrogenases have been reported to be highly resistant to CO like some [Ni-Fe]-hydrogenases and O₂ tolerant hydrogenases [14,36,37]. Microorganisms capable of producing such CO-resistant hydrogenases would be able to metabolize H₂ in CO-rich environments.

The effect of different CO gas fractions on the activity of adapted biomass was assessed in batch experiments. Formate and acetate production from CO as sole electron donor occurred, but higher formate formation and lower acetate formation were observed with increasing CO fractions (Figure 5). Formate is produced by some bacteria and archaea that grow acetogenically on CO, such as Clostridium ljungdahlii, Methanosarcina acetivorans, and Archaeoglobus fulgidus [40–42]. Currently, Fuchsiella alkaliacetigena and F. ferrireducens are the only isolates representing haloalkaliphilic hydrogenotrophic acetogen, but both are unable to grow in the presence of CO [43,44]. Bacteria belonging to Clostridiaceae family dominated the microbial community in the bioreactor (Figure 6). With further analysis, ~99% of these sequences were closely related to the Tindallia genus. Tindallia related bacteria were previously detected as dominant bacteria in H₂ fed sulfate and thiosulfate-reducing bioreactors and their role in formate production was hypothesized [7,8,23]. Tindallia related bacteria were also detected in high relative abundance in a 12 L bench-scale haloalkaliphilic bioreactor that was continuously operated for the treatment of high concentrations of sulfate [45]. The bioreactor was fed with glucose as substrate, that was first converted into ethanol, lactate, acetate and formate in the lower region of the bioreactor by fermentative microbes, and then sulfate reducers metabolized the organic...
acids coupled to sulfate reduction, both in the lower as well as in the higher regions of the bioreactor. Formate was detected in all regions of the bioreactor as well as in the effluent. Despite the low energy yield of H₂-driven formate production (Equation (7), Table 1), one of the Tindallia isolates was capable of H₂-driven formate production coupled to growth [46]. As Tindallia related bacteria were dominant during the whole bioreactor operation period, it was likely that they had adapted to CO.

**Figure 6.** Microbial 16S rRNA relative abundance on family level in the inoculum and biomass samples from the end of the runs without CO (day 46), with 5% CO (day 123), with 15% CO (day 216), and after the spike with 60% CO (day 218). For the samples of day 123, the suspended and settling fractions were separated and analyzed separately. A and B represent duplicates for the corresponding day. OTUs with less than 0.5% relative abundance were grouped in “others.”

CO-oxidation coupled to sulfate reduction at pH neutral, thermophilic conditions by mixed or pure cultures, such as Desulfotomaculum carboxydivorans and Archaeoglobus fulgidus, is well documented [20–22,40,47,48]. In the present study, sulfate reduction only started after the removal of CO and no CO-driven sulfate reduction was detected in batch experiments (Figure 7). This suggested that hydrogen and/or formate were the electron donors for sulfate reduction. Interestingly, incubations with CO and 50 mM formate did show sulfate reduction. This indicated that formate could act as an electron donor for sulfate reduction in the presence of CO, possibly circumventing hydrogenases. The sulfate reducers detected in both the suspended and attached biomass were related to the Desulfohalobiaceae family (Figure 6). Zooming in to genus level of Desulfohalobium related sequences detected in the libraries, all were closely related to Desulfonatronovibrio. The studied isolates belonging to this genus can use both H₂ and formate to reduce sulfate [49]. However, to date there is no information available on the effect of CO on these bacteria and how formate can be used as an electron donor in the presence of CO.
Even though inhibition depending on the feedstock used and production method, from 0 up to higher than 50% [53,54].

### 3.5. Application in Gas Biodesulfurization

Focusing on the microbial composition of the aggregates, the effect of pH on the hydrophobicity of cells surfaces or absence of soluble divalent cations for EPS stabilization [50–52].

The observed aggregates were more compact than those in a bioreactor without addition of CO. This might be connected to a metabolic relationship between CO oxidizers and sulfate reducers as was observed for pH neutral conditions [22]. Focusing on the microbial composition of the aggregates in settled biomass on day 123; ~58% consisted of Desulfohalobiaceae and more specifically Desulfitronovibrio-like bacteria (Figure 6). All studied Desulfitronovibrio sp. isolates use acetate as carbon source for growth [49]. Thus, aggregation of sulfate reducers together with CO-oxidizing microbes, such as acetogens, could have enhanced their growth due to acetate production. Additionally, the lower in situ CO concentrations generated by CO oxidizing acetogens might also have been more favorable to the sulfate reducers.

### 3.4. Biomass Aggregation

The addition of CO to the bioreactor influenced the formation of biomass aggregates that were not attached to sand. After addition of CO, small compact biomass aggregates were observed while no biofilm formation on sand particles was detected (Figure S2). A similar effect was previously reported in bioreactors operated at neutral conditions [22]. The formation of biomass aggregates under haloalkaline conditions was previously noted in a bioreactor fed with H2, but their appearance was not as compact as observed with CO (Figure S2) [23]. The particles in the bioreactor were mainly dominated by biomass aggregates that decreased in diameter upon CO addition (Figure 4). The maximum size achieved after CO addition did not increase with time as can be seen by the similar particle size distribution at days 125 and 216 (Figure 4). This phenomenon might be related to the haloalkaline conditions and more specifically, the effect of pH on the hydrophobicity of cells surfaces or absence of soluble divalent cations for EPS stabilization [50–52].

The observed aggregates were more compact than those in a bioreactor without addition of CO. This might be connected to a metabolic relationship between CO oxidizers and sulfate reducers as was observed for pH neutral conditions [22]. Focusing on the microbial composition of the aggregates in settled biomass on day 123; ~58% consisted of Desulfohalobiaceae and more specifically Desulfitronovibrio-like bacteria (Figure 6). All studied Desulfitronovibrio sp. isolates use acetate as carbon source for growth [49]. Thus, aggregation of sulfate reducers together with CO-oxidizing microbes, such as acetogens, could have enhanced their growth due to acetate production. Additionally, the lower in situ CO concentrations generated by CO oxidizing acetogens might also have been more favorable to the sulfate reducers.

### 3.5. Application in Gas Biodesulfurization

Syngas can be used as electron donor for sulfate and thiosulfate-reducing bioreactors operated under haloalkaline conditions. However, the CO concentration in the syngas varies considerably depending on the feedstock used and production method, from 0 up to higher than 50% [53,54]. Even though inhibition effects were observed during the spike of 60% CO performed in this study,
such spikes can be controlled by a proper production of syngas or pre-treatment of syngas [54]. Thus, despite using syngas without removing CO would decrease the cost of the electron donor, the capacity of biomass to withstand CO fractions only up to 15% might make the application of bleed stream treatment using syngas and recycling an interesting option for biodesulfurization systems (Figure 8).

Figure 8. Biological process for gas desulfurization using an anaerobic sulfate/thiosulfate reducing bioreactor to enable bleed stream recycling. The sulfide (H$_2$S) present in the gas is dissolved in an alkaline solution as HS$^-$ using a scrubber (1). This HS$^-$ rich solution goes to an aerobic bioreactor (2) where it is biologically oxidized under controlled microaerophilic conditions to mostly elemental sulfur (SO$_0$) and a minor fraction to sulfate (SO$_4^{2-}$) followed by chemical oxidation via polysulfides, to thiosulfate (S$_2$O$_3^{2-}$). The SO$_0$ is separated in a settler (3) and most of the liquid is recycled to the scrubber (1). Part of the liquid from the settler (3) goes to an anaerobic bioreactor (4) where SO$_4^{2-}$ and S$_2$O$_3^{2-}$ are reduced to HS$^-$ using syngas as an electron donor. The HS$^-$ produced is recycled back to the aerobic bioreactor (2). With time, this prevents accumulation of SO$_4^{2-}$ and S$_2$O$_3^{2-}$ in the whole system and theoretically reduces the amount of caustic required to increase the pH to almost zero. Additionally, it will prevent the disposal of bleed stream into the environment and maximize the S$^0$ production.

Acetate production might be a drawback for the application of such bioreactor in biodesulfurization systems. For the sulfate and thiosulfate reducing bioreactor, excessive acetate production consumes additional electron donor, and lowers the pH. This implies the requirement of extra syngas and alkalinity to compensate for acetate production, increasing the costs of operation. After treatment, the sulfide- and acetate-rich stream from the sulfate/thiosulfate-reducing bioreactor (Figure 8, n$^0$ 4) could be recycled back into the sulfide-oxidizing bioreactor (Figure 8, n$^0$ 2) [55]. In the sulfide-oxidizing bioreactor, acetate leads to organic contamination of the system resulting in growth of unwanted, heterotrophic microorganisms. Acetate oxidation requires extra consumption of O$_2$, increasing aeration costs, and production of CO$_2$, which decreases the pH. The pH decrease affects the sulfide absorption process (Figure 8, n$^0$ 1) that increases the caustic required to increase pH.

4. Conclusions

Syngas, containing up to 15% CO, can be used as electron donor for a sulfate/thiosulfate-reducing bioreactor operated at haloalkaline conditions. Adaptation of the biomass to CO is required as it
inhibits hydrogen-dependent microbial processes, such as sulfate reduction, formate production, and acetate production in non-adapted biomass. After adaptation, the biomass sulfate/thiosulfate reduction activity was comparable to previous studies using other electron donors, such as H₂ or formate. Acetate production was the dominant conversion in the bioreactor when CO was supplied. Acetate production seemed to enhance the growth of sulfate-reducing bacteria that require acetate as the carbon source. The high acetate concentration in the treated bleed stream has consequences for the sulfide oxidation step and sulfide absorption step of the biodesulfurization process after it is recycled.

Supplementary Materials: The following are available online at http://www.mdpi.com/2076-2607/8/9/1451/s1: Figure S1. Scheme of the gas-lift bioreactor set-up, Figure S2. Microscopic observations of the biomass from the bioreactor.

Author Contributions: Conceptualization, C.M.P., J.A.B.S., A.J.M.S., and M.D. (Martijn Diender); methodology, J.A.B.S., M.D. (Mark Dopson), and S.C.; formal analysis, C.M.P., J.A.B.S., S.C., M.D. (Martijn Diender); investigation, J.A.B.S., S.C., and M.D. (Martijn Diender); data curation, J.A.B.S.; writing—original draft preparation, C.M.P. and J.A.B.S.; writing—review and editing, C.M.P., J.A.B.S., S.C., M.D. (Mark Dopson), M.F.M.B., A.J.M.S., and M.D. (Martijn Diender); supervision, C.M.P., M.D. (Mark Dopson), M.F.M.B., and A.J.M.S.; project administration, C.M.P.; funding acquisition, C.M.P., M.F.M.B. and A.J.M.S. All authors have read and agreed to the published version of the manuscript.

Funding: This work was performed in the TTIW-cooperation framework of Wetsus, European Centre of Excellence for Sustainable Water Technology (www.wetsus.nl). Wetsus is funded by the Dutch Ministry of Economic Affairs, the European Union Regional Development Fund, the Province of Fryslân, the City of Leeuwarden and the EZ/Kompas program of the “Samenwerkingsverband Noord-Nederland”. Research of AJMS is financed by ERC grant project 323009 and by Gravitation grant project 024.002.002 from the Netherlands Ministry of Education, Culture and Science.

Acknowledgments: The authors like to thank the participants of the research theme “Sulfur,” namely Paqell, for fruitful discussions.

Conflicts of Interest: The authors declare no conflict of interest.

Nomenclature

- \( C_x \): Biomass concentration (g L\(^{-1}\))
- \( Ctx \): Total biomass concentration (g L\(^{-1}\))
- \( M_x \): Biomass molecular weight (g mol\(^{-1}\))
- \( V_r \): Bioreactor volume (L)
- \( Q \): Flow rate (L d\(^{-1}\))
- \( C_s \): Sulfate or thiosulfate concentration based on S molar content (mmol L\(^{-1}\))
- \( C_{TotalN} \): Total nitrogen concentration (mol L\(^{-1}\))
- \( r_{es} \): Sulfidogenic volumetric activity (mmol L\(^{-1}\) d\(^{-1}\))
- \( r_{vp} \): Formate, acetate or methane production volumetric activity (mmol L\(^{-1}\) d\(^{-1}\))

Subscript

- \( a \): Attached biomass
- \( x \): Biomass
- \( out \): Effluent
- \( in \): Influent
- \( p \): Products
- \( r \): Reactor
- \( s \): Sulfur
- \( sp \): Suspended biomass
- \( tx \): Total biomass
- \( TotalN \): Total nitrogen
References

1. Janssen, A.J.H.; Lens, P.N.L.; Stams, A.J.M.; Plugge, C.M.; Sorokin, D.Y.; Muyzer, G.; Dijkman, H.; Van Zessen, E.; Luimes, P.; Buismans, C.J.N. Application of bacteria involved in the biological sulfur cycle for paper mill effluent purification. *Sci. Total Environ.* 2009, 407, 1333–1343. [CrossRef] [PubMed]

2. De Rink, R.; Klok, J.B.M.; Van Heeringen, G.J.; Sorokin, D.Y.; Ter Heijne, A.; Zeijlmaker, R.; Mos, Y.M.; de Wilde, V.; Keesman, K.J.; Buismans, C.J.N. Increasing the selectivity for sulfur formation in biological gas desulfurization. *Environ. Sci. Technol.* 2019, 53, 4519–4527. [CrossRef] [PubMed]

3. Klok, J.B.M.; de Graaff, M.; van den Bosch, P.L.F.; Boelee, N.C.; Keesman, K.J.; Janssen, A.J.H. A physiologically based kinetic model for bacterial sulfide oxidation. *Water Res.* 2013, 47, 483–492. [CrossRef] [PubMed]

4. Roman, P.; Bijmans, M.F.M.; Janssen, A.J.H. Influence of methanethiol on biological sulphide oxidation in gas treatment system. *Environ. Technol.* 2016, 37, 1693–1703. [CrossRef] [PubMed]

5. Qian, Z.; Tianwei, H.; Mackey, H.R.; van Loosdrecht, M.C.M.; Guanghao, C. Recent advances in dissimilatory sulfate reduction. *Water Res.* 2019, 150, 162–181. [CrossRef]

6. Mu, T.; Yang, M.; Xing, J. Deep and high-efficiency removal of sulfate through a coupling system with sulfate-reducing and sulfur-oxidizing capacity under haloalkaliphilic condition. *Bioprocess Biosyst. Eng.* 2020, 43, 1009–1015. [CrossRef]

7. Sousa, J.A.B.; Plugge, C.M.; Stams, A.J.M.; Bijmans, M.F.M. Sulfate reduction in a hydrogen fed bioreactor operated at haloalkaline conditions. *Water Res.* 2015, 68, 67–76. [CrossRef]

8. Sousa, J.A.B.; Bijmans, M.F.M.; Stams, A.J.M.; Plugge, C.M. Thiosulfate conversion to sulfide by a haloalkaliphilic microbial community in a bioreactor fed with H2 gas. *Environ. Sci. Technol.* 2017, 51, 914–923. [CrossRef]

9. Zhou, J.; Xing, J. Effect of electron donors on the performance of haloalkaliphilic sulfate-reducing bioreactors for flue gas treatment and microbial degradation patterns related to sulfate reduction of different electron donors. *Biochem. Eng. J.* 2015, 96, 14–22. [CrossRef]

10. Van Houten, B.H.G.W.; van Doesburg, W.; Dijkman, H.; Copini, C.; Smidt, H.; Stams, A.J.M. Long-term performance and microbial community analysis of a full-scale synthesis gas fed reactor treating sulfate- and zinc-rich wastewater. *Appl. Microbiol. Biotechnol.* 2009, 84, 555–563. [CrossRef]

11. Diender, M.; Stams, A.J.M.; Sousa, D.Z. Pathways and bioenergetics of anaerobic carbon monoxide fermentation. *Front. Microbiol.* 2015, 6, 1275. [CrossRef] [PubMed]

12. Parshina, S.N.; Sipma, J.; Henstra, A.M.; Stams, A.J.M. Carbon monoxide as an electron donor for the biological reduction of sulphate. *Int. J. Microbiol.* 2010, 2010. [CrossRef] [PubMed]

13. Jeoung, J.H.; Fesseler, J.; Goetzl, S.; Dobbek, H. Carbon monoxide. Toxic gas and fuel for anaerobes and aerobes: Carbon monoxide dehydrogenases. In *Metal Ions in Life Sciences*; Kroneck, P.M.H., Sosa Torres, M.E., Eds.; Springer: Berlin, Heidelberg, Germany, 2014; pp. 37–69. [CrossRef]

14. Lubitz, W.; Hideaki, O.; Rudiger, O.; Reijerse, E. Hydrogenases. *Chem. Rev.* 2014, 114, 4081–4148. [CrossRef] [PubMed]

15. Adams, M.W.W. The structure and mechanism of iron-hydrogenases. *Biochim. Biophys. Acta Bioenerg.* 1990, 1020, 115–145. [CrossRef]

16. De Lacey, A.L.; Fernández, V.M.; Roussset, M.; Cammack, R. Activation and inactivation of hydrogenase function and the catalytic cycle: Spectroelectrochemical studies. *Chem. Rev.* 2007, 107, 4304–4330. [CrossRef]

17. Takors, R.; Kopf, M.; Mampel, J.; Bluemke, W.; Blombach, B.; Eikmanns, B.; Bengelsdorf, F.R.; Weuster-Botz, D.; Dürr, P. Using gas mixtures of CO, CO2 and H2 as microbial substrates: The do’s and don’ts of successful technology transfer from laboratory to production scale. *Microb. Biotechnol.* 2018, 11, 606–625. [CrossRef]

18. Sipma, J.; Lens, P.N.L.; Stams, A.J.M.; Lettinga, G. Carbon monoxide conversion by anaerobic bioreactor sludges. *FEMS Microbiol. Ecol.* 2003, 44, 271–277. [CrossRef]

19. Robb, F.T.; Techtman, S.M. Life on the fringe: Microbial adaptation to growth on carbon monoxide. *F1000Research* 2018, 7. [CrossRef]

20. Parshina, S.N.; Kijlstra, S.; Henstra, A.M.; Sipma, J.; Plugge, C.M.; Stams, A.J.M. Carbon monoxide conversion by thermophilic sulfate-reducing bacteria in pure culture and in co-culture with *Carboxydothermus hydrogenoformans*. *Appl. Microbiol. Biotechnol.* 2005, 68, 390–396. [CrossRef]
21. Parshina, S.N.; Sipma, J.; Nakashimada, Y.; Henstra, A.M.; Smidt, H.; Lysenko, A.M.; Lens, P.N.L.; Lettinga, G.; Stams, A.J.M. *Desulfotomaculum carboxydovorans* sp. nov., a novel sulfate-reducing bacterium capable of growth at 100% CO. *Int. J. Syst. Evol. Microbiol.* 2005, 55, 2159–2165. [CrossRef]

22. Van Houten, R.T.; van der Spoel, H.; van Aelst, A.C.; Hulshoff Pol, L.W.; Lettinga, G. Biological sulfate reduction using synthesis gas as energy and carbon source. *Biotechnol. Bioeng.* 1996, 50, 136–144. [CrossRef]

23. Esquivel-Elizondo, S.; Delgado, A.G.; Rittmann, B.E.; Krajmalnik-Brown, R. The *Acetobacterium woodii* metabolism by pure and mixed microbial cultures. *Appl. Environ. Microbiol.* 2017, 83, e00301-17. [CrossRef] [PubMed]

24. Rother, M.; Metcalf, W.W. Anaerobic growth of *Clostridium ljungdahlii* represents a microbial production platform based on syngas. *Appl. Microbiol. Biotechnol.* 2009, 80, 141017. [CrossRef] [PubMed]

25. Herlemann, D.P.; Labrenz, M.; Jürgens, K.; Bertilsson, S.; Waniek, J.J.; Andersson, A.F. Transitions in bacterial communities along the 2000 km salinity gradient of the Baltic Sea. *ISME J.* 2011, 5, 1571–1579. [CrossRef] [PubMed]

26. Hugerth, L.W.; Wefer, H.A.; Lundin, S.; Jakobsson, H.E.; Lindberg, M.; Rodin, S.; Engstrand, L.; Andersson, A.F. DegePrime, a program for degenerate primer design for broad-taxonomic-range PCR in microbial ecology studies. *Appl. Environ. Microbiol.* 2014, 80, 5116–5123. [CrossRef]

27. Lindh, M.V.; Figueroa, D.; Sjöstedt, J.; Baltar, F.; Lundin, D.; Andersson, A.; Legrand, C.; Pinhassi, J. Transplant experiments uncover Baltic Sea basin-specific responses in bacterioplankton community composition and metabolic activities. *Front. Microbiol.* 2015, 6, 223. [CrossRef]

28. Edgar, R.C. UPARSE: Highly accurate OTU sequences from microbial amplicon reads. *Nat. Methods* 2013, 10, 996–999. [CrossRef]

29. Quast, C.; Pruesse, E.; Yilmaz, P.; Gerken, J.; Schweer, T.; Yarza, P.; Peplies, J.; Glöckner, F.O. The SILVA ribosomal RNA gene database project: Improved data processing and web-based tools. *Nucleic Acids Res.* 2013, 41, 590–596. [CrossRef]

30. Robertson, C.E.; Harris, J.K.; Wagner, B.D.; Granger, D.; Browne, K.; Tatem, B.; Feazel, L.M.; Pace, N.R.; Frank, D.N. Explicet: Graphical user interface software for metadata-driven management, analysis and visualization of microbiome data. *Bioinformatics* 2013, 29, 3100–3101. [CrossRef]

31. Flamholz, A.; Noor, E.; Bar-Even, A.; Milo, R. eQuilibrator—the biochemical thermodynamics calculator. *Nucleic Acids Res.* 2015, 81, 5949–5956. [CrossRef] [PubMed]

32. Ferry, J.G. CO in methanogenesis. *Ann. Microbiol.* 2010, 60, 1–12. [CrossRef]

33. Esquivel-Elizondo, S.; Delgado, A.G.; Rittmann, B.E.; Krajmalnik-Brown, R. The effects of CO$_2$ and H$_2$ on CO metabolism by pure and mixed microbial cultures. *Biotechnol. Biofuels* 2017, 10, 220. [CrossRef] [PubMed]

34. Bertsch, J.; Müller, V. CO metabolism in the acetogen *Acetobacterium woodii*. *Appl. Environ. Microbiol.* 2015, 81, 5949–5956. [CrossRef] [PubMed]

35. Daniels, L.; Fuchs, G.; Thauer, R.K.; Zeikus, J.G. Carbon monoxide oxidation by methanogenic bacteria. *J. Bacteriol.* 1977, 132, 118–126. [CrossRef] [PubMed]

36. Adams, M.W.W. The metabolism of hydrogen by extremely thermophilic, sulfur-dependent bacteria. *FEMS Microbiol. Rev.* 1990, 6, 219–237. [CrossRef]

37. Fox, J.D.; Kerby, R.L.; Roberts, G.P.; Ludden, P.W. Characterization of the CO-induced, CO-tolerant hydrogenase from *Rhodospirillum rubrum* and the gene encoding the large subunit of the enzyme. *J. Bacteriol.* 1996, 178, 1515–1524. [CrossRef]

38. Ceccaldi, P.; Schuchmann, K.; Muller, V.; Elliott, S.J. The hydrogen dependent CO$_2$ reductase: The first completely CO tolerant Fe$_2$H$_4$-hydrogenase. *Energy Environ. Sci.* 2017, 10. [CrossRef]

39. Tchtmann, S.M.; Colman, A.S.; Robb, F.T. “That which does not kill us only makes us stronger”: The role of carbon monoxide in thermophilic microbial consortia. *Environ. Microbiol.* 2009, 11, 1027–1037. [CrossRef]

40. Henstra, A.M.; Dijkema, C.; Stams, A.J.M. *Archaeoglobus fulgidus* couples CO oxidation to sulfate reduction and acetogenesis with transient formate accumulation. *Environ. Microbiol.* 2007, 9, 1836–1841. [CrossRef]

41. Köpke, M.; Held, C.; Hujer, S.; Liesengang, H.; Wieser, A.; Wollherr, A.; Ehrenreich, A.; Liebl, W.; Gottschalk, G.; Dürr, P. *Clostridium ljungdahlii* represents a microbial production platform based on syngas. *Proc. Natl. Acad. Sci. USA* 2010, 107, 15305. [CrossRef]

42. Rother, M.; Metcalf, W.W. Anaerobic growth of *Methanosarcina acetivorans* C2A on carbon monoxide: An unusual way of life for a methanogenic archaeon. *Proc. Natl. Acad. Sci. USA* 2004, 101, 16929–16934. [CrossRef] [PubMed]
43. Zhilina, T.N.; Zavarzina, D.G.; Panteleeva, A.N.; Osipov, G.A.; Kostrikina, N.A.; Tourova, T.P.; Zavarzin, G.A. Fuchsiella alkaliacetigena gen. nov., sp. nov., an alkaliphilic, lithoautotrophic homoacetogen from a soda lake. Int. J. Syst. Evol. Microbiol. 2012, 62, 1666–1673. [CrossRef] [PubMed]

44. Zhilina, T.N.; Kuznetsov, B.B.; Zavarzina, D.G.; Detkova, E.N.; Patutina, E.O. Fuchsiella ferrireducens sp. nov., a novel haloalkaliphilic, lithoautotrophic homoacetogen capable of iron reduction, and emendation of the description of the genus Fuchsiella. Int. J. Syst. Evol. Microbiol. 2015, 65, 2432–2440. [CrossRef]

45. Mu, T.; Xing, J.; Yang, M. Sulfate reduction by a haloalkaliphilic bench-scale sulfate-reducing bioreactor and its bacterial communities at different depths. Biochem. Eng. J. 2019, 147, 100–109. [CrossRef]

46. Sorokin, D.Y.; Detkova, E.N.; Muyzer, G. Sulfur-dependent respiration under extremely haloalkaline conditions in soda lake “acetogens” and the description of Natroniella sulfidigena sp. nov. FEMS Microbiol. Lett. 2011, 319, 88–95. [CrossRef] [PubMed]

47. Sipma, J.; Lettinga, G.; Stams, A.J.M.; Lens, P.N.L. Hydrogenogenic CO conversion in a moderately thermophilic (55 °C) sulfate-fed gas lift reactor: Competition for CO-derived H₂. Biotechnol. Prog. 2006, 22, 1327–1334. [CrossRef] [PubMed]

48. Sinharoy, A.; Baskaran, D.; Pakshirajan, K. A novel carbon monoxide fed moving bed biofilm reactor for sulfate rich wastewater treatment. J. Environ. Manag. 2019, 249, 109402. [CrossRef]

49. Sorokin, D.Y.; Tourova, T.P.; Abbas, B.; Suhacheva, M.V.; Muyzer, G. Desulfonatronovibrio halophilus sp. nov., a novel moderately halophilic sulfate-reducing bacterium from hypersaline chloride-sulfate lakes in Central Asia. Extremophiles 2012, 16, 411–417. [CrossRef]

50. Otto, K.; Elwing, H.; Hermansson, M. Effect of ionic strength on initial interactions of Escherichia coli with surfaces, studied on-line by a novel quartz crystal microbalance technique. J. Bacteriol. 1999, 181, 5210–5218. [CrossRef]

51. Sousa, J.A.B.; Sorokin, D.Y.; Bijmans, M.F.M.; Plugge, C.M.; Stams, A.J.M. Ecology and application of haloalkaliphilic anaerobic microbial communities. Appl. Microbiol. Biotechnol. 2015, 99, 9331–9337. [CrossRef] [PubMed]

52. Guan, G.; Kaewpanha, M.; Hao, X.; Abudula, A. Catalytic steam reforming of biomass tar: Prospects and challenges. Renew. Sustain. Energy Rev. 2016, 58, 450–461. [CrossRef]

53. Voldsund, M.; Jordal, K.; Anantharaman, R. Hydrogen production with CO₂ capture. Int. J. Hydrog. Energy 2016, 41, 4969–4992. [CrossRef]

54. Van den Bosch, P.L.F. Biological Sulfide Oxidation by Natron-Alkaliphilic Bacteria Application in Gas Desulfurization. Ph.D. Thesis, Wageningen University, Wageningen, The Netherlands, 2008.

© 2020 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).