Methods to characterize type, relevance, and interactions of organic matter and microorganisms in fluids along the flow path of a geothermal facility

Alessio Leins[1,2], Danaé Bregnard[3], Andrea Vieth-Hillebrand[1], Stefanie Poetz[1], Florian Eichinger[4], Guillaume Cailleau[3], Pilar Junier[3], and Simona Regenspurg[1]

[1]GFZ German Research Centre for Geosciences, Telegrafenberg, 14473, Potsdam, Germany
[2]Friedrich Schiller University, Institute of Geosciences, Applied Geology, Burgweg 11, 07749 Jena, Germany
[3]University of Neuchâtel, Laboratory of Microbiology, Institute of Biology, Rue Emile-Argand 11, 2000 Neuchâtel, Switzerland
[4]Hydroisotop GmbH, Woelkestraße 9, 85301 Schweitenkirchen, Germany

Correspondence: Alessio Leins (leins@gfz-potsdam.de)

Abstract. ~~Dissolved organic matter and microorganisms were analyzed~~ The analysis of dissolved organic matter (DOM) and microorganisms was conducted along the flow path of a geothermal facility in Austria. Various analytical methods were used to characterize and differentiate between natural and synthetic organic matter, characterize the ~~microbial community composition, and determine~~ composition of the microbial community, and assess the implications of microorganisms in ~~an operating~~ the operation of a geothermal site. ~~Dissolved organic carbon (DOC) concentrations were in the range of 8.4–10.3 and typically decreased from the production to the injection side. Carbonate scalings are avoided in the facility by the injection of a chemical scaling inhibitor within the production well at 500 depth. It was calculated that the inhibitor contributes approximately 1 DOC to the produced fluids.~~ Ion chromatography (IC), liquid chromatography — organic carbon detection (LC-OCD), and Fourier-transform ion cyclotron resonance mass spectrometry (FT-ICR-MS) in negative electrospray ionization (ESI(-)) and positive atmospheric pressure photoionization (APPI(+)) mode were applied to the fluid samples to characterize the ~~dissolved organic matter (DOM) composition~~ composition of DOM and distinguish between the inhibitor and the natural DOM. The concentration of dissolved organic carbon (DOC) ranged from 8.5 to 10.4 $mg\,C\,L^{-1}$. To prevent carbonate scaling, a chemical scaling inhibitor is injected into the production well at a depth of 500 m. The inhibitor adds approximately 1 $mg\,C\,L^{-1}$ DOC to the produced fluids. Depending on the applied ionization mode, the FT-ICR-MS results show that between 31 % and 65 % of the macromolecular formulas (150–1000 Da) detected in the fluid samples ~~seem~~ appear to originate from the inhibitor. However, the DOM ~~is mainly composed~~ primarily consists of low molecular weight acids (LMWA), especially acetate with up to 7.4 $mg\,C\,L^{-1}$. ~~The microbial community composition~~ Targeted amplification of the 16S rRNA gene was performed to assess the bacterial diversity. Based on this, metabolic pathways related to the presence of acetate in the samples were predicted using PICRUSt2. The composition of the microbial community varied along the ~~flowpath with dominant phyla being~~ flow path, with Firmicutes, Proteobacteria, and Thermotogae ~~. Based on the microorganisms found in the sample, the metabolic pathways have been assessed. Acetate might be produced by microorganisms~~ being the dominant phyla. Microorganisms may produce acetate through various fermentation processes~~(e.g.~~, such as from lysine, pyruvate~~and hexitol),~~ and hexitol. Assessing the presence and

interaction of organic compounds and microorganisms in geothermal fluids ~~provides~~ can provide a broader understanding of processes within the geothermal facility. This understanding could be beneficial for the efficient ~~use~~ operation of a geothermal

25 power plant.

1 Introduction

Deep hydrothermal energy production is becoming increasingly important as an alternative energy source. Geothermal power plants extract heat from subsurface fluids to produce electricity and heat. The depth at which these fluids are extracted can vary from a few hundred meters to a few kilometers. Geothermal power plants face various operational challenges, such as mineral precipitation (scaling) or corrosion of the well casing and the components of the geothermal plant (Regenspurg et al., 2016; Demir et al., 2014). Scaling and corrosion are caused by hydrochemical reactions related to changes in fluid pressure and temperature during transport or the presence of metabolic by-products of microorganisms present in fluids (Inagaki et al., 2003; Little and Lee, 2015). Therefore, analyzing the fluids and their composition is crucial for understanding, predicting, and mitigating possible chemical reactions that could compromise the functioning of a geothermal plant. However, the role of organic components has rarely been considered in the analyses performed to characterize geothermal power stations. In recent studies, corrosion and declining injectivity in power plants have been attributed to biofilm formation, microbially induced corrosion (MIC), or microbially induced mineral scaling (Alawi et al., 2011; Vetter, 2012; Lerm et al., 2013; Little and Lee, 2015; Westphal et al., 2019; Brehme et al., 2020; Leins et al., 2022; Madirisha et al., 2022). The growth of these microorganisms may be sustained by organic compounds in the fluids. These compounds can originate from natural sources in the geothermal aquifer or may have been introduced artificially, such as from detergents used during drilling or injection of organic scaling inhibitors.

Organic acid anions have been reported in various deep subsurface systems, including oil-field waters (Carothers and Kharaka, 1978; Hatton and Hanor, 1984; Kharaka et al., 1985, 1997), waters from fractured crystalline rock (Sherwood Lollar et al., 2021; Kieft et al., 2018), hydrothermal vents (Lang et al., 2010, 2018; McDermott et al., 2015), and fluids from geothermal sites in the Molasse Basin (Alawi et al., 2011; Vetter, 2012; Leins et al., 2022). Their concentrations in fluids can vary by several orders of magnitude, ranging from a few $\mathrm{mg\,L^{-1}}$ to $10,000\,\mathrm{mg\,L^{-1}}$ (Kharaka et al., 1997). The highest concentrations are usually found in oil-field waters and are dominated by acetate, followed by propionate, butyrate, and valerate (Carothers and Kharaka, 1978; Fisher and Boles, 1990; Kharaka et al., 1987).

For information on the presence of specific organic acid anions in the fluids and the scaling inhibitor, IC was applied. However, these organic acid anions may only represent a small fraction of the detectable DOC present in the fluids. In this study, the characterization of DOC was performed via LC-OCD to quantify the DOC content and characterize its distribution into fractions with size exclusion chromatography (SEC). These fractions range from low molecular weight compounds, such as low molecular weight acids (LMWA) and neutral compounds (LMWN), to high molecular weight compounds, such as humic sub-

stances and biopolymers). The analyses were performed on fluids and a scaling inhibitor sample from the geothermal power plant located in Bad Blumau, Austria. Additionally, Fourier transform ion cyclotron resonance mass spectrometry (FT-ICR-MS) was used to conduct a detailed characterization of medium to high molecular weight organic compounds within a mass range of 150 to 1000 Da. Both APPI(+) and ESI(-) ionization modes were used to detect polar and less polar compounds, since the chemical properties of the inhibitor were unknown beforehand. FT-ICR-MS provides accurate masses of molecules, enabling the determination of elemental formulas. This helps to reveal the influence of chemical scaling inhibitors and biomarkers for microorganisms in geothermal fluids on a molecular level. To the best of our knowledge, this is the first time that FT-ICR-MS has been used to characterize DOM in fluids from a geothermal power plant. ESI(-) FT-ICR-MS has already been applied to a variety of water systems such as groundwater (McDonough et al., 2020), deep fracture water (Kieft et al., 2018), pore water (D'Andrilli et al., 2010; Rossel et al., 2016; Schmidt et al., 2009), hydrothermal vents (Noowong et al., 2021; Rossel et al., 2017; Gomez-Saez et al., 2016), and marine as well as terrestrial waters (D'Andrilli et al., 2010; Koch et al., 2008; Minor et al., 2012; Sleighter and Hatcher, 2008).

In addition to the analyses of the DOM, a parallel analysis of bacterial diversity in the fluids was conducted. Targeted amplification of the 16S rRNA gene allowed to assess the bacterial diversity present at the three same sampling points. In order to explain the presence of certain organic compounds, we predicted bacterial metabolic pathways based on the known metabolisms of the most dominant microorganisms found in our samples. The study evaluated the bacterial metabolic pathways associated with the consumption or production of acetate to determine if changes in the microbial community are associated with changes in the presence of particular organic acids. Changes in the bacterial community and associated metabolic pathways, as well as microbial growth, may impact the efficacy of power plants. Therefore, it is important to assess both the organic compounds and the microbial composition of geothermal fluids.

This study aims to (1) characterize the DOM and microbial community of a deep geothermal fluid using various methods; (2) distinguish between natural and synthetic DOM and determine its origin; (3) determine the metabolic pathways related to the consumption or production of acetate, and (4) assess whether the composition of DOM correlates with a change in microbial diversity.

2 Material and methods

2.1 Site description

The geothermal site Bad Blumau is a geothermal power plant and thermal spa located in south-east Austria (Upper Styrian Basin as part of the Pannonian Basin). The targeted geothermal system is also utilized by several other spas and heat

usages in the area~~, and~~ . This area was discovered during a hydrocarbon exploration campaign in the second half of the 20th century (Alt-Epping et al., 2013). The Styrian Basin is of Miocene age, composed of Tertiary siliciclastic basin-fills ~~underlain by~~ that overlie Paleozoic carbonates and phyllites of the Grazer Paleozoicum, which in turn overlie the crystalline basement (Goldbrunner, 2000; Alt-Epping et al., 2013). The geothermal reservoir in this area is hosted by carbonate rocks, ~~which consist of~~ specifically Devonian limestones and dolomites ~~originating~~ that originated from Paleozoic reef development (Hubmann et al., 2006). ~~Tectonic deformation caused intense fracturing of the carbonate rocks , which therefore~~ The carbonate rocks exhibit good aquifer properties due to intense fracturing caused by tectonic deformation (Goldbrunner, 2000).

In addition to its balneological purpose, the site is also used for heat and electricity production, as well as the commercial production of liquefied CO_2 (Alt-Epping et al., 2013). The system ~~is operated~~ operates as a geothermal doublet with an internal distance of 2300 m between the production and injection well~~and targets~~ , targeting the Paleozoic (Devonian) carbonate ~~formation in~~ at a depth of 2800 m (Goldbrunner, 2000). The reservoir temperature was reported to be 124 °C (Goldbrunner, 2005), while the ~~produced~~ fluids at the production well head reach 107 °C and are reinjected with approximately 65–50 °C after ~~the~~ heat extraction (Westphal et al., 2019). The geothermal fluid ~~ascends~~ rises via natural gas lift of CO_2 ~~with~~ at an average flow rate of 20 L s^{-1}. The CO_2 is reported to ascend from the mantle regions along fault zones and is ~~regarded to be~~ considered a product of the Neogene volcanism of the Styrian Basin (Goldbrunner, 2000). The presence of CO_2 in the system ~~poses~~ presents a major challenge ~~for the plant operation . At~~ to the operation of the plant. At a depth of approximately 300–350 m below the surface, ~~degassing of the CO2 takes place leading to~~ CO_2 degassing occurs, resulting in carbonate precipitation (Alt-Epping et al., 2013).

To prevent carbonate scaling, the inhibitor named hydrin 45.3 is injected into the fluid within the production well at a depth of approximately 500 m ~~,~~ and at a pressure before ~~degassing of CO2~~ CO_2 degassing. The current inhibitor consists of organic polyelectrolytes. Generally, the exact chemical composition of the inhibitor falls under the protection of commercial and industrial secrecy.

The geothermal fluids at Bad Blumau were described to be of $NaHCO_3$ type with a salinity of approximately 20 g L^{-1}, and a slightly alkaline pH of around 8.0 (Westphal et al., 2019). The CO_2 concentration makes up 99 % of the gases and is estimated to be approximately 5 L L^{-1}.

2.2 Sample collection

Fluid samples were collected during two sampling campaigns in March and June 2021. ~~In both campaignsone sample was taken from each sampling point from the surface installation during regular operation of the geothermal plant,~~ During both campaigns, the geothermal plant was in regular operation. Samples were taken from the surface installations of the plants. The sampling points were at the production well GB2 (P) before ~~CO2~~ CO2 extraction, at the heat central (HC), and at the injection well GB1 (I) (Fig. 1). The respective ~~fluid where~~ fluids were at approximately 108 °C (P), 50 °C (HC), and ~~42.2~~ 42 °C (I). Unfiltered fluids were collected in 500 ml Duran glass bottles with screw caps containing teflon-coated septa inside. The bottles were pre-rinsed with the fluid and ~~afterwards filled completely~~ then completely filled to avoid contact with air. The samples
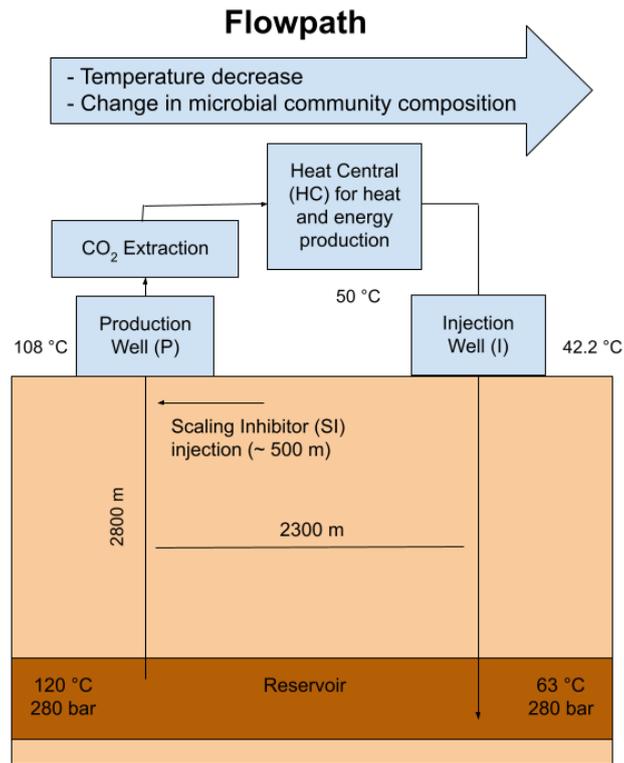
Figure 1. Schematic of the geothermal power plant in Bad Blumau. Modified from Westphal et al. (2019).

were stored at 4 °C until shipment to the laboratory, where they were ~~again stored~~ stored again at 4 °C until further analyses. In addition to the fluid samples, a sample of the ~~used scaling inhibitor~~ scaling inhibitor used was obtained from the site operators.

For microbial analysis, 40 L of fluids were sampled at ~~the same~~ each of the three sampling points (P, HC, I) ~~during the sampling campaign~~ in June 2021. The 40 L of fluids were directly filtered through 0.22 μm nitrocellulose membrane filters (Merck Millipore, Germany) under sterile conditions. ~~Six independent filters were prepared simultaneously as independent replicates~~ Filtering of each sample was done using an EZ-Stream pump (Merck Millipore, Germany) and six glass filtration stations mounted on a manifold (Merck Millipore, Germany). This resulted in six filters for each sample that were prepared simultaneously and served as independent replicates. Filters were transported ~~at~~ to the laboratory at a temperature of 4 °C ~~to the laboratory, where they were~~ and were later stored at -20 °C until further processing.

2.3 Analytical Methods

2.3.1 Ion chromatography

The quantification of organic anions (formate, acetate, propionate, butyrate, valerate, oxalate) and inorganic anions (F⁻, Cl⁻, Br⁻, SO_4^{2-}) from both sampling campaigns was ~~conducted via~~ performed by IC (ICS 3000, Thermo Fisher Scientific) using an AS-AP autosampler, AS11 HC column and a conductivity detector. KOH solutions with varying concentrations over time were used as eluent for the samples. The initial KOH concentration was 1.4 mM and ~~stepwise increased towards~~ gradually increased to 60 mM ~~within~~ over a period of 32 min. After 32 min the concentration was reduced to the initial value of 1.4 mM and equilibrated for 12 min. The flow-rate was 0.38 ml min⁻¹. The column temperature was ~~at~~ 35 °C and 10 μl of the sample was injected for each run. The quality of the measurements was verified daily using standards that contain the analytes in different concentrations. The concentrations were 0.02; 1.0; 10 and 100 mg L⁻¹. For samples with high chloride concentrations (>1 g L⁻¹), the chloride was reduced prior to ~~the analysis of the~~ analyzing organic anions using OnGuard II AG/H cartridges (Thermo Fischer Scientific).

2.3.2 Liquid chromatography – organic carbon detection (LC-OCD)

The ~~characterization and quantification of the~~ DOC and its fractions ~~from both sampling campaigns were determined by~~ were characterized and quantified using SEC with subsequent UV (λ = 254 nm) and IR detection by ~~a~~ an LC-OCD system (Huber and Frimmel, 1996) ~~. Phosphate~~ in both sampling campaigns. A mobile phase of phosphate buffer (pH 6.85; 2.7 g L⁻¹ KH_2PO_4, 1.6 g L⁻¹ Na_2HPO_4) was used ~~as mobile phase and a flow of~~, and the flow was adjusted to 1.1 mL min⁻¹ ~~was adjusted~~ (Huber et al., 2011). The samples passed a 0.45 μm membrane syringe filter before entering the chromatographic column (Toyopearls HW 50 S, 30 μm 250 mm x 20 mm). Here, the DOC will be separated into different fractions according to their molecular masses: Macro.1 (biopolymers), Macro.2 (humic substances), Macro.3 (building blocks), low molecular weight acids (LMWA), and low molecular weight neutrals (LMWN) (Huber et al., 2011). See Table 1 for properties and description of the fractions. ~~The~~ DOC was quantified by IR detection of the released CO_2 after UV oxidation (λ = 185 nm) in a Gräntzel thin-film reactor. Humic and fulvic acids standards of the Suwannee River, provided by the International Humic Substances Society (IHSS), were used for molecular mass calibration. Solutions of known amounts of potassium hydrogen phthalate were used for external calibration of the CO_2-quantification.

2.3.3 Solid phase extraction (SPE)

Salts are known to cause ~~ionization suppression~~ suppression of ionization (King et al., 2000) and have to be eliminated prior to FT-ICR-MS analysis. ~~On the other hand, also~~ Additionally, the concentrations of DOM in natural geothermal water samples ~~is~~ are too low for ~~being analyzed directly~~ direct analysis by FT-ICR-MS. Therefore, geothermal fluids had to be pretreated by solid phase extraction (SPE) on SPE cartridges (PPL Bond Elut 1 g, 6 ml cartridge; Agilent Technologies, Germany) (Dittmar et al., 2008) to obtain salt free samples and accumulate 1 mg of DOC for the FT-ICR-MS analysis. The ~~cartridges were pre-rinsed~~

Table 1. Description of LC-OCD fractions (Zhu et al., 2015). Modified from Huber et al. (2011), Penru et al. (2013).

Fraction	Molecular Mass Range	Properties	Description
Hydrophobic organic carbon (HOC)		Hydrophobic	lipids (fats) released from bacteria and algae, hydrocarbons
Biopolymers (Makro.1)	>10,000 Da	Not UV-absorbable, hydrophilic	Polysaccharides and proteins
Humic substances (Makro.2)	∼1000 Da	Highly UV-absorbable, hydrophobic	Calibration based on Suwannee River standard from IHSS
Building blocks (Makro.3)	350–500 Da	UV-absorbable	Breakdown products of humic substances
Low molecular weight organic acids (LMWA)	<350 Da	Negatively charged	aliphatic acids
Low molecular weight neutrals (LMWN)	<350 Da	Weakly or uncharged hydrophilic, amphiphilic	Alcohols, aldehydes, ketones, amino acids

~~with methanol and acidified deionized water (pH2, hydrochloric acid) for cleaning. The samples~~ sample amount for SPE was adjusted to approximately contain 1 mg of DOC based on the DOC concentration determined by LC-OCD. The fluid samples from March 2021 and the inhibitor sample were filtered with 0.45 μm membrane syringe filters, diluted with deionized water (1:1) and acidified up to pH 2 with hydrochloric acid (suprapur) and passed through the cartridges. The ~~sample amount was~~

170 ~~adjusted to approximately contain 1 of DOC. After the~~ cartridges were pre-rinsed with methanol and acidified deionized water (pH2, hydrochloric acid) to activate the resins. After absorption, the cartridges were rinsed with 3 x 6 ml of acidified deionized water (pH 2) to remove any remaining salts. The cartridges were dried by vacuum pump for 5 minutes. Finally, the DOM was eluted with 6 ml of methanol into pre-combusted glass vials, dried ~~under~~ in N_2 atmosphere and weighed. The dried samples were then stored in the dark at -24 °C until FT-ICR-MS analysis.

175 **2.3.4 Fourier transform ion cyclotron resonance mass spectrometry (FT-ICR-MS)**

FT-ICR-MS, with its ultra-high resolution in combination with atmospheric pressure ionization modes, can provide the elemental composition of thousands of individual medium- to high-molecular weight organic compounds. All ~~the DOM samples as well as the~~ DOM samples and inhibitor were dissolved in methanol (MeOH) to give a stock solution with a final concentration of 1 $mg\,ml^{-1}$. The samples were analyzed on a Bruker Solarix FT-ICR-MS with a 12 T refrigerated actively shielded

180 superconducting magnet. For ESI(-) analysis, the stock solutions were spiked with 4 μL of 25 % aqueous ammonia solution. Measurement solutions of 100 $μg\,ml^{-1}$ ~~in MeOH were prepared~~ were prepared in MeOH. Ionization was realized with an Apollo II ESI source from Bruker Daltonik GmbH (Bremen, Germany) in negative ion mode. Samples were infused at a flow rate of 150 $μl\,h^{-1}$ using a syringe pump (Hamilton). The capillary voltage was set ~~to~~ at 3000 V and an additional collision-induced dissociation (CID) voltage of 70 V ~~in the source was applied to avoid cluster and adduct formation~~ was applied at the source to

185 avoid the formation of clusters and adducts. Nitrogen was used as a drying gas at a flow rate of 4.0 L min^{-1} and a temperature of 220 °C and a nebulizing gas at 1.4 bar. The spectra were recorded in broadband mode using 4 megaword data sets. The ion accumulation time was set at 0.05 s and 200 scans were collected and added to each mass spectrum. Ions were detected in a *m/z* range between 150 and 1000.

For the APPI(+) analyses, solutions of 20 $\mu\text{g ml}^{-1}$ in MeOH were prepared from the stock solutions. Measurement
190 solutions of 100 $\mu\text{g ml}^{-1}$ were prepared in MeOH:n-hexane (9:1). The ion source used was an APPI-II from Bruker Daltonik GmbH (Bremen, Germany). Samples were introduced into the MS at an infusion flow rate of 20 $\mu\text{l min}^{-1}$ with a syringe pump (Hamilton). The capillary voltage was set to -1000 V and CID to 30 V. Nitrogen was used as a drying gas at a flow rate of 3.0 L min^{-1} and a temperature of 210 °C, as well as a nebulizing gas at 2.3 bar and a temperature of 350 °C. The spectra were recorded in broadband mode using 4 megaword data sets. The ion accumulation time was set to 0.05 s and 300 scans
195 were collected and added to each mass spectrum. Ions were detected in a *m/z* range between 147 and 1500.

In ESI(-), the DOM samples were internally recalibrated using O_x compounds, while for the inhibitor both O_x and S_1O_x compounds were used. In APPI(+), both the fluid samples and the inhibitor sample were internally recalibrated using O_x compounds. The root mean square deviations of the eight internal calibrations ranged between 0.013 and 0.018. Method blanks covering sample preparation steps (SPE) and the FT-ICR-MS measurement were prepared for both modes and blank
200 signals were removed from the DOM signal list of the fluid and the inhibitor samples.

Data evaluation was performed using Data Analysis 4.0 SP5 (Bruker Daltonik GmbH, Germany), Excel 2019 (Microsoft Corporation, Redmont, WA), and the statistical data analysis tool R 4.0.1 (R Core Team, 2020) using the tidyverse package (Wickham et al., 2019). Only *m/z* values with a signal to noise ratio ≥ 9 were exported for formula assignment. The molecular formulas were calculated considering the isotopes ^{12}C and ^{13}C
205 with upper elemental thresholds of O $\leq$ 32, S $\leq$ 2, and N $\leq$ 2, C and unlimited H. The mass tolerance was set to $\pm$ 0.5 ppm, and formulas containing ^{13}C were excluded from the final dataset.

The abundance of inhibitor-derived compounds in the fluids was determined by adding up the total monoisotopic ion abundance (TMIA) of the formulas derived from the inhibitor for each fluid sample.

The double bond equivalent (DBE) is a measure to express the number of double bonds and rings. With the respective
210 molecular formula, it can be calculated from the number of carbon (C), hydrogen (H), and nitrogen (N) atoms as follows in Eq. (1):

$$DBE = C - \frac{H}{2} + \frac{N}{2} + 1 \tag{1}$$

Since the DBE counts all double bonds with at least one carbon as a bonding partner, it is not well suited to describe aromaticity of DOM compounds that contain a high number of double bonds within carboxy groups. Therefore, the DOM
215 adapted modified aromaticity index (AI_{mod}) described by Koch and Dittmar (2006) has been used, as expressed in Eq. (2), to evaluate the proportion of aromatic compounds in the dataset. It is a measure of the double bond density in a molecule

considering ~~,~~ taking into account the contribution of heteroatoms. The AI_{mod} is based on the assumption that 50 % of the oxygen is bound with double bonds in carboxyl groups.

$$AI_{mod} = \frac{1 + C - 0.5O - S - 0.5(N + P + H)}{C - 0.5O - N - S - P} \tag{2}$$

Three ranges were established to describe the aromaticity of a given DOM compound. AI_{mod} values ≤ 0.5 are described as aliphatic (Zhu et al., 2019), AI_{mod} between 0.5 and 0.67 represent aromatic compounds, and $AI_{mod} \geq 0.67$ describes condensed aromatic compounds (Koch and Dittmar, 2006).

Intensity-weighted averages for DBE, AI_{mod}, O/C ratio, H/C ratio, carbon, hydrogen, and oxygen number in each sample were calculated after Bae et al. (2011) in Eq. (3)

$$var_{average} = \frac{\Sigma_i Ii * (var)i}{\Sigma_i Ii} \tag{3}$$

where *Ii* and *(var)i* are the relative abundances and respective variable value of peak *i*.

2.3.5 Microbial analysis

Five of the six filters were processed for DNA extraction. The last filter was kept as a backup. DNA extraction was ~~done~~ performed with the FastDNA®SPIN kit for soil (MP Biomedicals, USA) using three bead-beating rounds and pooling the three independent DNA extracts at the final step (Wunderlin et al., 2013). In parallel, a DNA blank extract was prepared by performing the same procedure without any cellular material. DNA was quantified with the Qubit®dsDNA HS Assay Kit and Qubit®2.0 Fluorometer (Invitrogen, Carlsbad, CA, USA). The DNA extracts were sent to Fasteris SA (Geneva, Switzerland) for amplicon sequencing ~~in~~ on an Illumina MiSeq sequencing platform (Illumina, San Diego, CA, USA). The V3-V4 region of the 16S rRNA gene for bacteria was amplified using the Bakt_341F (CCTACGGGNGGCWGCAG) and Bakt_805R (GAC-TACHVGGGTATCTAATCC) primers (Herlemann et al., 2011). Sequences, provided as pre-trimmed and pre-demultiplexed, were processed with Qiime2 (Bolyen et al., 2019) using the dada2 pipeline (Callahan et al., 2016). Sequences were truncated and joined to full ~~denoised~~ de-noised sequences of 464 bp. These sequences were grouped as amplicon sequences variants (ASVs). Taxonomy was assigned using the Silva database release 132 (Quast et al., 2012) and the vSEARCH-based consensus taxonomy classifier (Rognes et al., 2016). Further analysis was performed in R Studio V3.6.3 using ~~the~~ R version 4.2.2 with the phyloseq package (McMurdie and Paulson, 2016), the vegan package Oksanen et al. (2022), and the ggplot2 package Wickham (2016). ~~In order to model metabolic capabilities~~ To represent the metabolic capacities of the community~~present, ASVs matching those present~~, ASVs corresponding to those found in the DNA blank extract, mitochondria, and chloroplast signals were ~~removed~~ excluded from the database ~~before~~ prior to exporting the representative sequences and the ASVs count table. A taxonomy was then assigned to the PYCRUSt2 (Douglas et al., 2020) pipeline outputs using the SILVA database (Quast et al., 2012) to highlight organisms' involvement in metabolic functionalities, such as those related to acetate production or degradation. These steps were performed in R using the biomformat (McMurdie and Paulson, 2016), tidyverse (Wickham et al., 2019), and ggplot2 (Wickham, 2016) packages.

3 Results

3.1 Organic and inorganic anions

IC analyzes show relatively constant concentrations of Cl^- and F^- throughout the power plant with around 4 g L^{-1} and 10 mg L^{-1}, respectively (Table 2). SO_4^{2-} values ranged from 500–570 mg L^{-1}. Bromide concentrations ranged from 2.6–15.2 mg L^{-1}. Cl^- and SO_4^{2-} were slightly lower, with around 3.4 g L$^-$ and 490 mg L$^-$, respectively, compared to Westphal et al. (2019). Acetate was the predominant organic acid anion with concentrations ranging from 5.8 to 7.4 mg C L^{-1}. Propionate was found slightly above the detection limit with 0.6 mg C L^{-1} in the injection side sample from June. Formate, butyrate, valerate, and oxalate were not detected (< 0.6 mg C L^{-1}). The inhibitor analyses showed 220 mg C L^{-1} of acetate, suggesting that with the reported addition of 10 mg L^{-1} of inhibitor to the fluids, 2.2 µg C L^{-1} acetate could have been added from the inhibitor.

3.2 DOC and bulk fractions

The DOC in the fluid samples ranged from 8.5–10.4 mg C L^{-1} (Table 2). In March, the DOC decreased from the production to the injection side, whereas in June the concentrations were quite similar in the production and injection sides and slightly higher at the heat central. According to Westphal et al. (2019), the concentration of DOC was found to be 14.5 mg C L^{-1} in the production sample and 4 mg C L^{-1} in the injection sample. A decrease in DOC concentration was observed along the pathway. However, a stronger one compared to this study. The inhibitor DOC comprises 102 g C L^{-1}. With the reported dosage, the inhibitor contributes approximately 1 mg C L^{-1} to the total DOC of the fluids. The DOC fractions measured by size exclusion chromatography show a predominant LMWA fraction in all samples (Table 2), which is likely due to the high acetate concentrations in the fluid. The Makro fraction is the second most abundant (16–19 %), followed by the LMWN fraction (8–12 %). In this study, we were unable to distinguish the Makro.1, Makro.2 and Makro.3 fractions since the chromatograms display only one peak spanning across the retention times for all three Makro fractions. Therefore, this peak represents the whole Makro fraction (10,000–350 Da). HOC was detected only in the March samples (13–19 %). The relative abundance of the LMWA fraction along the flowpath in both campaigns correlates with the respective organic acid anion trends of the samples. In the inhibitor sample, the Makro fraction represents 99 % of the DOC. By utilizing the Makro fraction provided by the SEC in the fluid samples and the inhibitor dosage, we determined that approximately 60–74 % of the Makro fraction in the fluid samples originates from the inhibitor.

Table 2. DOC, concentrations of organic and inorganic anions, and relative abundance of the DOC fractions of the production (P), heat central (HC), injection (I) fluid, and inhibitor (SI) samples for the sampling campaign in March 2021 and June 2021 measured by LC-OCD and IC. Fluid data was compiled from Leins et al. (2023).

	March 2021			June 2021			
	P	HC	I.	P	HC	I	SI
Cl^- (g L^{-1})	3.9	4.1	4.0	4.0	3.9	3.7	0.14
SO_4^{2-} (mg L^{-1})	544	570	554	545	520	500	14.7
F^- (mg L^{-1})	10.1	10.3	10.2	10.5	10.6	10.4	2
Br^{-1} (mg L^{-1})	2.7	2.6	4.5	12.4	12.2	15.2	<1
Acetate							
(mg L^{-1})	17.2	16.7	14.2	17.9	18.2	17.9	542
(mg C L^{-1})	7	6.8	5.8	7.3	7.4	7.3	220
Propionate							
(mg L^{-1})	<1	<1	<1	<1	<1	1.3	<1
(mg C L^{-1})						0.6	
Σ Organic acid anions							
(mg L^{-1})	17.2	16.7	14.2	17.9	19.5	19.1	542
(mg C L^{-1})	7	6.8	5.7	7.3	7.4	7.9	220
DOC (mg C L^{-1})	10.4	10	8.6	8.8	9.2	8.5	102,180
DOC Fractions							
HOC %	13	13	19	0	0	0	0
Makro %	16	17	19	17	17	19	99
LMWA %	59	61	54	72	72	72	0
LMWN %	12	9	8	11	11	9	1

3.3 Molecular composition of the DOM

280 While LC-OCD analyzes provide general information about the molecular size distribution of the DOM, FT-ICR-MS enables highly-resolved insight into the molecular composition of DOM compounds within a mass range from approximately 150 to 1000 Da. Acidic compounds are detected in ESI(-), and low polarity neutral compounds are characterized using APPI(+) mode.
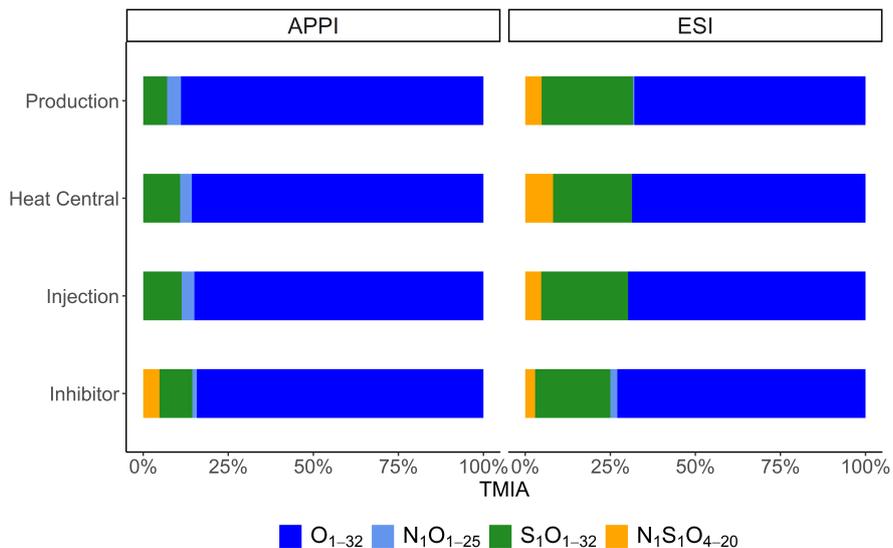
Figure 2. Relative abundances of assigned compound classes (O_x, N_1O_x, S_1O_x, and $N_1S_1O_x$) in APPI and ESI ~~Mode~~mode.

3.3.1 ESI(-)-FT-ICR-MS

285 Negative mode ESI FT-ICR-MS of the Bad Blumau samples provided several hundred assigned formulas in the mass to charge (*m/z*) range of 157–987 (Table 3). The main compound classes in the DOM of the fluid samples were, in decreasing abundance, oxygen containing compounds (O_x) with around 67 % of the ~~total monoisotopic ion abundance (TMIA)~~TMIA, sulfur and oxygen containing compounds (S_1O_x) with ~~22.7–26.6~~23–27 %TMIA, and nitrogen, sulfur and oxygen containing compounds ($N_1S_1O_x$) with 4.5–4.7 %TMIA. N_1O_x compounds were ~~only present~~ found only in the production side sample (0.4 %TMIA).

290 ~~DOM from~~ The DOM of the inhibitor sample shows a similar distribution with O_x accounting for ~~72.9~~73 %TMIA, S_1O_x with ~~22.1~~22 %TMIA, $N_1S_1O_x$ with 2.9 %TMIA, and N_1O_x with 2 %TMIA (Fig. 2). ~~The fluid samples are affected by inhibitor signals ranging from~~ In direct comparison, 31–65.~~3~~ % of the TMIA in the fluid samples derive from formulas that are also present in the inhibitor samples. The abundance of inhibitor derived compounds decreases from the production side (65 %TMIA) to the heat central (31 %TMIA), followed by an increase to the injection side (42 %TMIA) (Table 3).

295 The sample with the highest ~~numbers of assigned formulas~~ number of formulas assigned is the heat central (Table 3). The mean number of atoms ~~across~~ in all samples ranges from 22.7–24.2 (C), 29.3–31.9 (H), and 15.2–16.8 (O). The M_n and M_w range from ~~554.9~~549.9–595 and 623.3–670.7, respectively, and DBE values are around 8.5 to 8.9 (Table 4). The AI_{mod} ~~shows~~ indicates that the majority of the signals ~~are~~ consist of aliphatic compounds ($AI_{mod} \leq 0.5$), with a few aromatic ($0.5 < AI_{mod} < 0.67$) and condensed aromatic ($AI_{mod} \geq 0.67$) compounds (Table 4).

300 The ~~Van Krevelen diagrams were used to visualize compositional differences in the samples by presenting the molecular ratio of H/C and O/C atoms. To simplify the comparison, we followed the differentiation into four groups with different H/C and O/C ranges after Zhu et al. (2019) (I: H/C > 1 and O/C < 0.5 including lipids, proteins and part of the lignins, II: H/C~~

Table 3. Total and unique numbers of assigned monoisotopic signals within their sample group including the distribution of the main elemental classes, total mass range, percentage of TMIA derived from inhibitor signals, elemental numbers (carbon, hydrogen, oxygen), and molecular weight.

Well	No. of formulas						Mass range	SI amount	Mean			M_n	M_w
	Total	O_x	N_1O_x	S_1O_x	$N_1S_1O_x$	Unique	(Da)	(TMIA %)	C	H	O	Total	Total
ESI negative													
P	669	521	5	80	63	106	171–995	~~65.3~~ 65 (280)	23	29.3	16.8	582.1	629.6
HC	809	619	0	82	108	198	179–981	~~31.1~~ 31 (241)	24.2	31.9	16.4	595	639.9
I	573	469	0	61	43	16	165–987	~~42.4~~ 42 (222)	22.9	29.8	15.9	569.3	623.3
SI	472	409	7	40	16	148	157–987	-	22.7	29.9	15.2	554.9	670.7
APPI positive													
P	700	594	55	51	0	46	209–1019	64 (496)	20.8	24.5	10	436.6	467.6
HC	649	547	39	63	0	52	193–1019	~~58.1~~ 58 (460)	21.2	25.8	9.74	440.6	473.6
I	649	555	43	50	0	40	209–912	~~57.5~~ 58 (463)	21	25	9.52	433.8	461.3
SI	1741	1276	82	189	194	1262	181–1175	-	22.7	28.8	12.8	511.9	543.2

M_n: number-averaged molecular weight; M_w: weight-averaged molecular weight

~~> 1 and O/C > 0.5 including amino sugars, carbohydrates and part of the tannins, III: H/C < 1 and O/C < 0.5 including condensed hydrocarbons and part of the lignins and IV: H/C < 1 and O/C > 0.5 including partly condensed hydrocarbons and tannins). The majority of the signals in the fluid and inhibitor samples of the ESI data are present in area II (Fig. 3)~~ O_x class (x = 1–32) shows a roughly Gaussian distribution of relative abundances, dominating in the range of O_{12-24} in the fluid samples (P, HC, I) (Fig. 3), especially, the even numbered compounds. S_1O_x compounds in the inhibitor are mainly composed of compounds with odd numbers of oxygen in the range of S_1O_{13-27}. They exhibit a higher relative abundance compared to the other signals. These distinct peaks coincide with the dominant peaks in the inhibitor sample. The few N_1O_x compounds, all exhibit odd oxygen numbers as the inhibitor. These compounds were introduced by the inhibitor and are only present in the production side sample. $N_1S_1O_x$ compounds, appear to be less affected by the inhibitor. Mainly $N_1S_1O_4$ and likely the even numbered oxygen compounds in the range of $N_1S_1O_{8-20}$ are introduced by the inhibitor.

3.3.2 APPI(+)-FT-ICR-MS

APPI FT-ICR-MS of the Bad Blumau fluid samples provided assigned formulas in the *m/z* range of 193–1019 (Table 3). The main compound classes in the DOM of the fluid samples were, in order of predominance, O_x with 69–76.~~1~~ %TMIA, S_1O_x with 6–9.2 %TMIA, and N_1O_x with 2.8–3.5 %TMIA. For the inhibitor sample, 1741 signals ~~were assigned~~ with a similar compound class distribution of O_x with ~~81.8~~ 82 %TMIA and S_1O_x with 9.3 %TMIA were assigned. $N_1S_1O_x$ compounds were assigned with 4.7 %TMIA. N_1O_x compounds have the lowest abundance with 1.3 %TMIA (Fig. 2). The sample with the highest ~~numbers~~

number of assigned formulas is the inhibitor (Table 3). Up to 64 % of the TMIA in fluid samples originate from formulas that are also present in the inhibitor sample. The abundance decreases from the production side (64 %TMIA) to the injection side (58 %TMIA). Comparing the fluid samples only, the heat central sample has a slightly higher number of unique formulas. The mean number of atoms in all samples ranges from 20.8–22.7 (C), 24.5–28.8 (H), and 9.5–12.8 (O). The mean oxygen numbers decrease from the production to the injection side (10 to 9.52) and are lower than the mean oxygen number of the inhibitor (12.8). The M_n and M_w range from 433.8–543.2 and 461.3–543.2, respectively. The DBE values are found around 8.8 to 9 and the AI_{mod} with 0.23 shows that the compounds are aliphatic (Table 4). $N_1S_1O_x$ compounds are only present in the inhibitor and were not detected in the fluid samples.

The O_x class shows a Gaussian distribution centered around O_{9-12} in the fluid samples and O_{11-13} in the inhibitor sample (Fig. 3). A comparison of the assigned O_x signals in the fluids with those of the inhibitor shows that the inhibitor introduces a large amount of up to 63.2 % of the signals in the fluids.
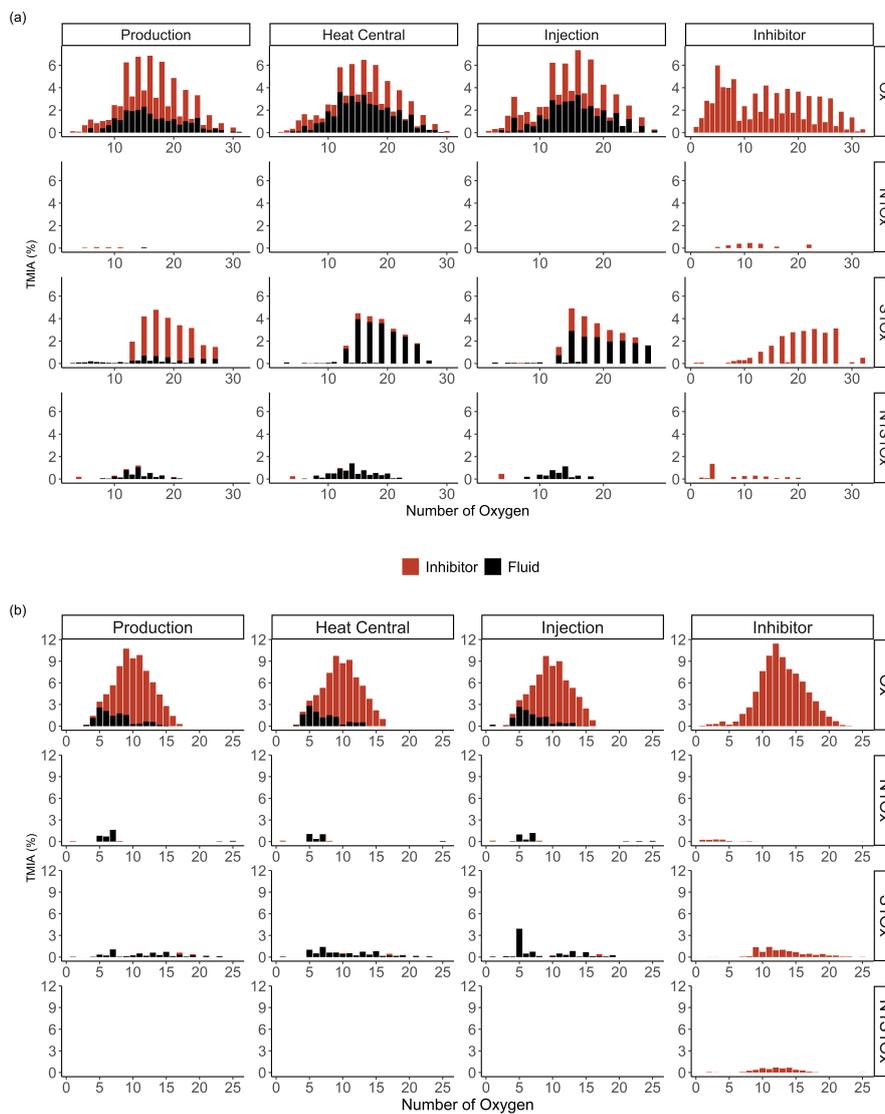
Figure 3. ~~Van Krevelen diagrams~~ TMIA vs. number of ~~the fluid~~ oxygen for every compound class and ~~inhibitor samples~~ sample in (a) ~~ESI~~APPI(~~-~~+) and (b) ~~APPI~~ESI(+~~-~~) mode~~color-coded by~~. Data colored red within the ~~assigned compound classes~~fluid samples represent the fluid signals that correspond to the signals found in the inhibitor sample.

3.4 Microorganisms in the fluids

The composition of the bacterial communities in the fluids was investigated by amplicon sequencing of the 16S rRNA gene. The relative abundance of the 10 most abundant bacterial phyla present in the different fluids of Bad Blumau is shown in Fig. 4. A very significant shift in community composition was detected consistently in the replicate samples of the three sampling points. In all three, the Firmicutes phylum was dominant. Amplicon sequence variants (ASVs) assigned to Proteobacteria were present in a small relative abundance in the production well but became highly abundant (co-dominant with ASVs from Firmicutes) at the heat central. However, at the injection site, Proteobacteria were not detected and replaced by the *Thermotogae* phylum, which dominated the community together with Firmicutes. The *Thermotogae* phylum was also detected in all replicates from the heat central, but at a much lower relative abundance. In addition to the dominant phyla, there was also a low abundance of *Actinobacteria*, *Cyanobacteria* and *Planctomycetes*, followed by few *Nitrospirae* (in one sample), *Thermotogae* (in one sample), and *Verrucomicrobia* in the production side. Low abundance of *Thermotogae*, *Actinobacteria*, and *Nitrospirae* were shown in the heat central and low abundance of *Thermotogae*, as well as an extremely low relative abundance of *Actinobacteria* and *Bacteroidetes* were observed in the injection side.

The same analysis was performed at a lower taxonomic rank. At the genus level, drastic changes in the composition of the bacterial community at different sampling points were observed more easily. The relative abundance of the 10 most abundant genera per sampling point is represented in Fig. 5. The genus *Bacillus* (Firmicutes) dominated the fluids from the production well. *Caulobacter* (Proteobacteria) and SCADC1-2-3 (Firmicutes) were dominant in the heat central, while the sulfate-reducing bacteria (SRB) *Desulfotomaculum* (Firmicutes) was dominant in the injection side. The top five genera represent around 60 % of the detected community in the production well and the heat central, while the top five genera represent around 75 % of the community at the injection site. This suggests that the community at the injection site was less diverse than the communities present at the other two sites.

The metabolic potential of the bacterial communities in Bad Blumau was assessed based on the phylogenetic composition using PICRUSt2 (Douglas et al., 2020). It predicts the metabolic capacities of a given bacterial genus based on the conserved V3-V4 regions of the bacterial 16S rRNA gene. After predicting
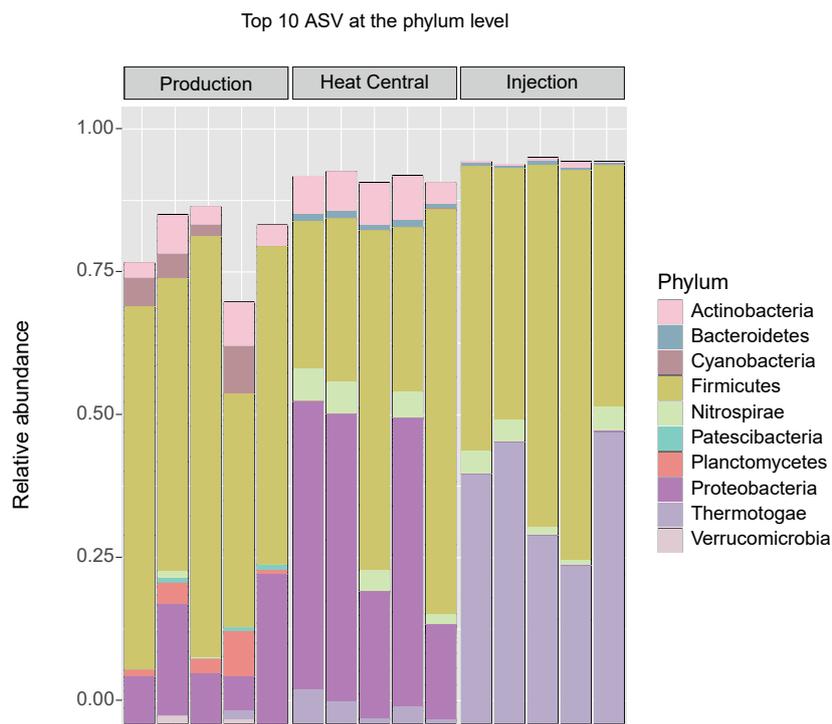
Figure 4. Top 10 bacterial phyla in relative abundance (in all samples). The ASVs not assigned to any phylum (unclassified) are not shown.

365   the pathways that can be present, the software also ~~predicts~~ estimates the contribution of bacteria from a given genus to the predicted pathway. The potential metabolisms ~~resulting in acetate production~~ that resulted in the production of acetate were the fermentation of lysine to acetate and butyrate, the fermentation of pyruvate to acetate and lactate, and the fermentation of hexitol, which can also lead to the production of lactate, formate, ethanol, and acetate (Figure 6). Acetate can then be used in methanogenesis (Figure 6). ~~At the production site, Bacillus~~ *Bacillus* is the dominant genus ~~contributing to the different~~

370   ~~metabolic pathways identified here, while at the injection site,~~ that contributes to the identified metabolic pathways at the production side, while *Desulfotomaculum* was the dominant genus at the injection side. The heat central, depending on the targeted metabolism, ~~either~~ represented a transition between the production and injection points ~~, (Figure 6)~~ or can be dominated by a different genus, ~~like the~~ such as the ~~*Caulobacter*~~*Cauulobacter* genus in the case of ~~the~~ lysine fermentation (Figure 6).
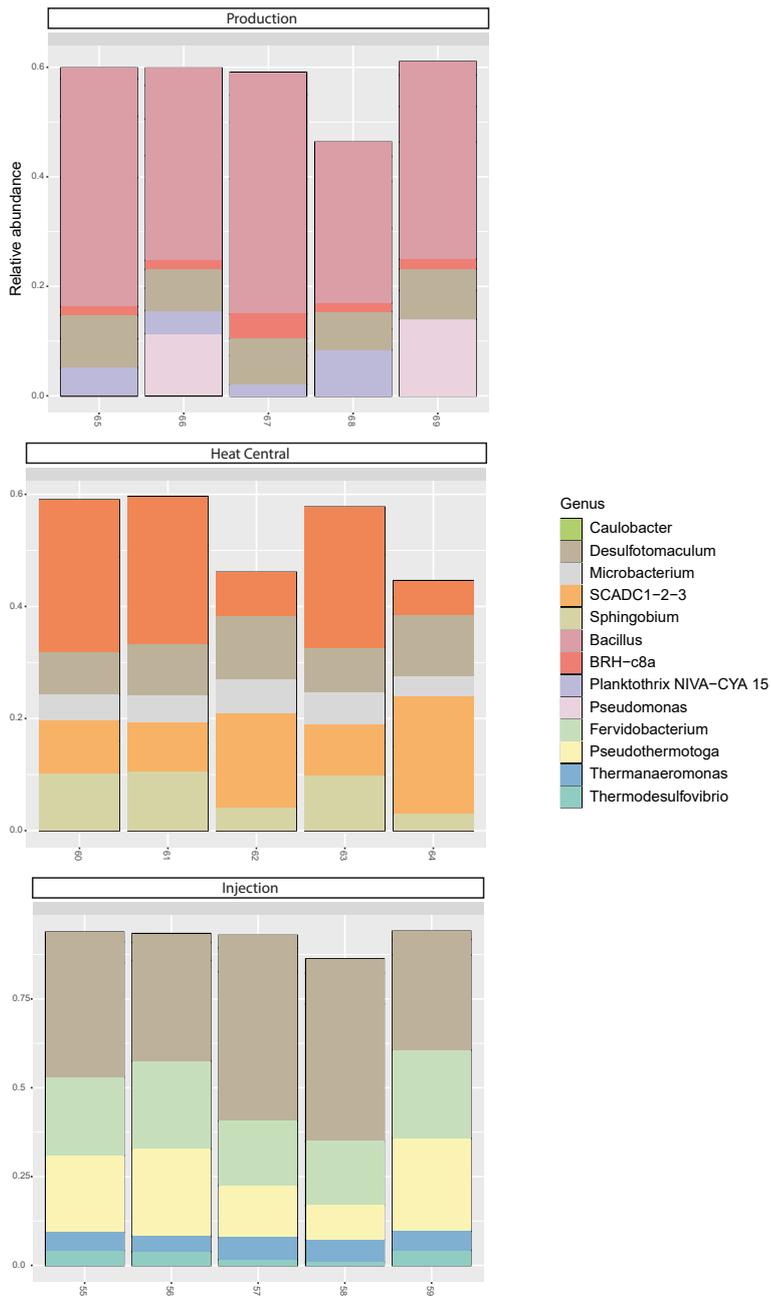
Figure 5. Top 5 bacterial genera in relative abundance (calculated for each sampling point separately). The ASVs not assigned to any phylum (unclassified) are not shown.
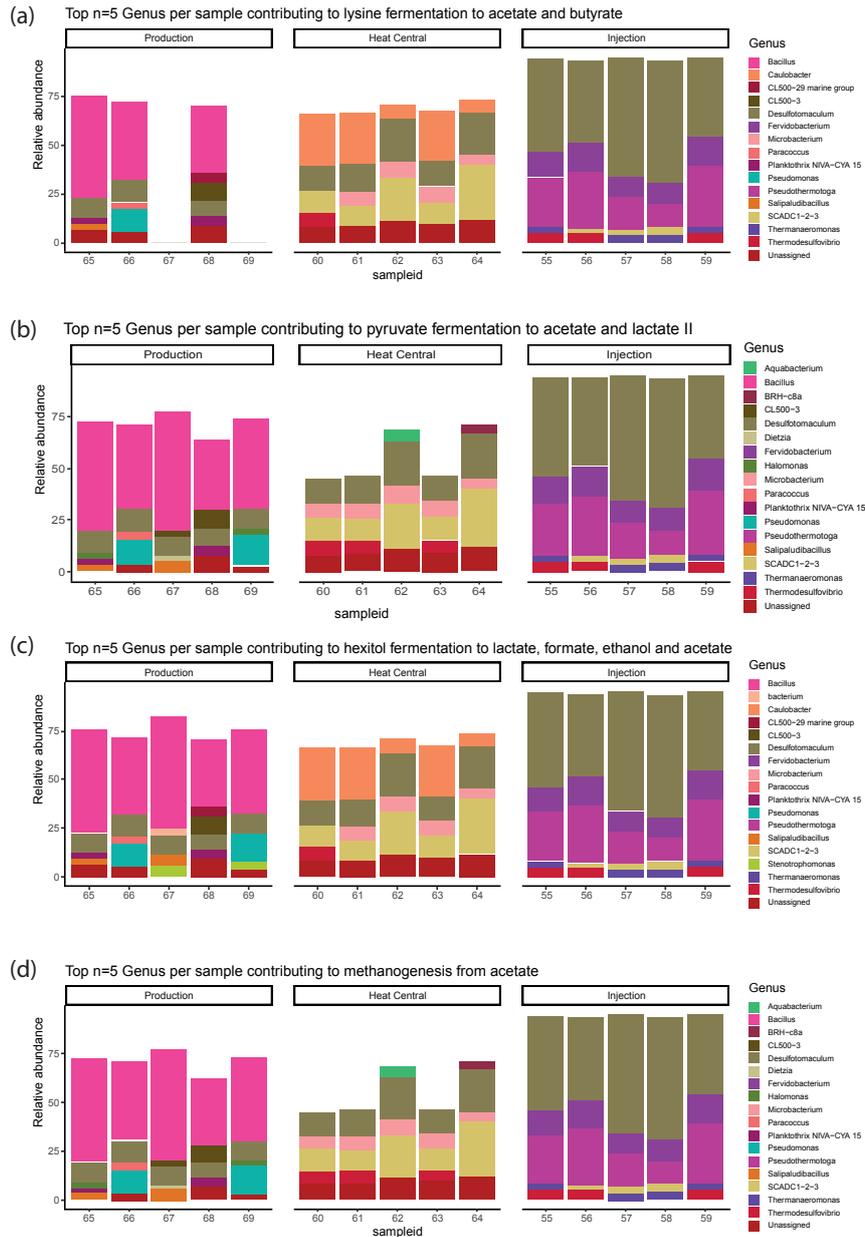
Figure 6. Top 5 Genus per sample ~~contributing to~~ and their relative abundance in (a) lysine to acetate and butyrate fermentation, (b) pyruvate fermentation to acetate and lactate II, (c) hexitol fermentation to lactate, formate, ethanol and acetate, (d) methanogenesis from acetate. Sampleid (55 – 69) represents the numbers assigned to each filter.

4 Discussion

LC-OCD analyses provided a good general overview of the DOC fractions in the geothermal fluids, showing that most of the DOM is present in the LMWA fraction. IC results confirmed that this acid fraction is mainly composed of acetate. With FT-ICR-MS we were able to analyze the molecular formulas of the macromolecular inventory and characterize the organic compounds in the fluids that derive from the inhibitor. This arises the following questions: (1) is it possible to characterize and subtract synthetic organic matter to obtain an indication of the natural organic matter in geothermal fluids? (2) Why does DOM and microbial diversity change along the fluid pathway and is there a correlation between the two? (3) Where does the acetate derive from and what is the metabolic pathway for it? These three questions are discussed below.

4.1 DOM composition and impact of the inhibitor in the fluid of the deep geothermal site Bad Blumau

The overall decrease in abundance of inhibitor derived signals from the production to the injection side in ESI(-) mode could be an indication for degradation, alteration, or dilution of the inhibitor compounds along the fluid pathway. The strong overlap of odd numbered S_1O_{13-27} compounds in the fluid and inhibitor samples suggests that these compounds derive mainly from the inhibitor. The assigned signals from the fluid and inhibitor samples show a strong influence in the production side sample with up to 65 %. Both the heat central and injection side samples do not show such a strong overlap with the inhibitor sample. However, it is likely that these S_1O_x compounds have undergone chemical alteration along the flow path, and are therefore no longer detectable in the geothermal fluid samples from the heat central and injection side. A similar result can be seen for the O_x compound class after subtracting the inhibitor signals. Another strong influence of the inhibitor can be observed with the intensity-weighted average of the molecular O/C ratio in the production side (Table 4). The O/C ratio is higher with inhibitor components and could be explained by the

S_1O_{13-27}, which were mainly identical to inhibitor signals in the production side but not in the heat central and the injection side~~(Table 4). Approximately half of the signals in the fluid samples in area II of the Van Krevelen diagrams match with the signals of the inhibitor (Fig. 8). With the exception of the production side sample, the inhibitor signals in the fluid samples in~~

410 ~~area I are mainly present at lower .~~

The van Krevelen diagrams were used to visualize compositional differences in the samples by presenting the molecular ratio of H/C and O/C atoms. To simplify the comparison, we followed the differentiation into four groups with different H/C (and O/C ranges after Zhu et al. (2019) (I: H/C > 1 and O/C < ~~1.6) values. Two groups of S_1O_x signals are mainly present in the production side. One with high H~~0.5 including lipids, proteins and part of the lignins, II: H/C ~~values around 1.6 and one~~

415 ~~with~~ > 1 and O/C > 0.5 including amino sugars, carbohydrates and part of the tannins, III: H/C ~~around 0.8 and both groups at approximately~~ < 1 and O/C ~~0.2. These signals correspond to S_1~~< 0.5 including condensed hydrocarbons and part of the lignins and IV: H/C < 1 and O~~ₓ (x~~/C > 0.5 including partially condensed hydrocarbons and tannins). Most of the signals in the fluid and inhibitor samples of the ESI data are present in area II (Fig. 7). Approximately half of the signals from the fluid samples that are plotted in area II of the van Krevelen diagram are consistent with the inhibitor signals. In area I, the fluid samples from

420 heat central and injection side show inhibitor-derived formulas mainly with lower H/C (< ~~1~~1). ~~This further suggests that these compounds may not derive from the inhibitor but are part of the natural DOM or are coming from other sources.~~1.6) values (Fig. 8).

Similar to the ESI(-) dataset, the APPI(+) data ~~shows~~ show that the proportion of the TMIA in the fluid samples ~~deriving~~ derived from the inhibitor decreases from the production to the injection side. ~~Here, the inhibitor-derived TMIA shows a~~

425 ~~minimal and gradual decrease over the three sampling points from 64 to 57.5 (Table 3). The mean oxygen numbers decrease from the production to the injection side and differ clearly from the mean oxygen number of the inhibitor . The mean number of carbon and hydrogen, as well as the mean $M_{\bar{n}}$, and M_w, increase from the production side to the heat central, followed by a slight decrease to the injection side. An opposite trend is shown for the mean DBE values (Table 4). In terms of mean DBE, carbon, and hydrogen number, the production side sample is most similar to the inhibitor sample. The~~ The inhibitor affects

430 especially the O_x class ~~shows a Gaussian distribution centered around O_{9-12} in the fluid samples, and O_{11-13} in the inhibitor sample~~ (Fig. 7). ~~A comparison of the assigned O_x signals in the fluids with those in the inhibitor show that a large amount with up to 63.2 of the signals in the fluids are introduced by the inhibitor. Removing the~~ 3). Removing inhibitor signals results in a shift ~~of~~ in the O_x distribution from O_{9-12} to O_{3-9} centered around O_5 in the fluid samples. O_x compounds with more than 15 oxygen atoms are ~~solely found~~ found exclusively in the inhibitor (Fig. ~~7~~3). The introduction of molecules with high numbers

435 of oxygen is also shown by the decrease of average O/C ratios ~~decreasing if the~~ if inhibitor signals are removed from the data (Table 4).

The APPI data shows distinct patterns for the fluid and ~~the~~ inhibitor samples in the ~~Van~~ van Krevelen diagrams (Fig. 8). ~~The majority~~ Most of the signals in the fluid samples ~~are accumulated~~ accumulate in area I, while the inhibitor sample shows a strong accumulation in area II and a weaker accumulation in area I at O/C < 0.2. Signals that were only found in the fluid samples

440 are mainly present in the O/C and H/C range of 0.2–0.4 and 1–1.7, respectively. They represent the O_x as well as distinct N_1O_x and S_1O_x compound classes compared to the inhibitor compounds. Typically, signals in this range are attributed to compounds
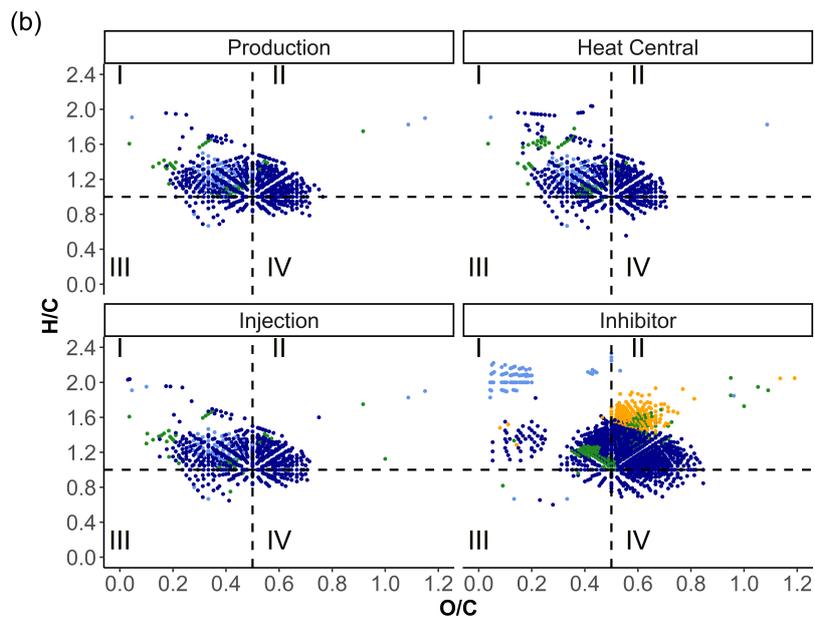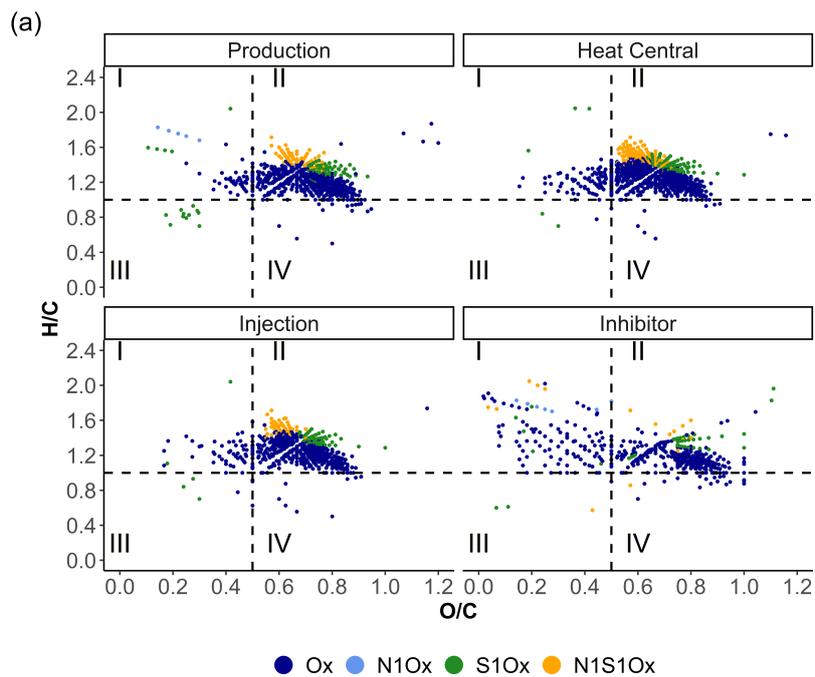
Figure 7. ~~TMIA vs. number~~ Van Krevelen diagrams of ~~oxygen for every compound class~~ the fluid and ~~sample~~ inhibitor samples in (a) ~~APPI~~ESI(~~+~~-) and (b) ~~ESI~~APPI(~~-~~+) mode ~~. Data colored in red within~~ color-coded by the ~~fluid samples represent the fluid signals that match with the signals found in the inhibitor sample~~assigned compound classes.
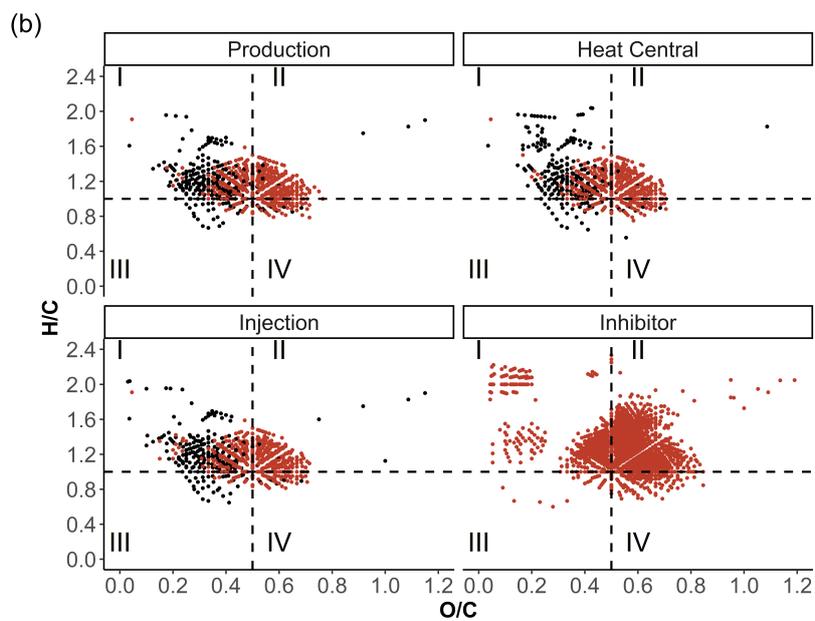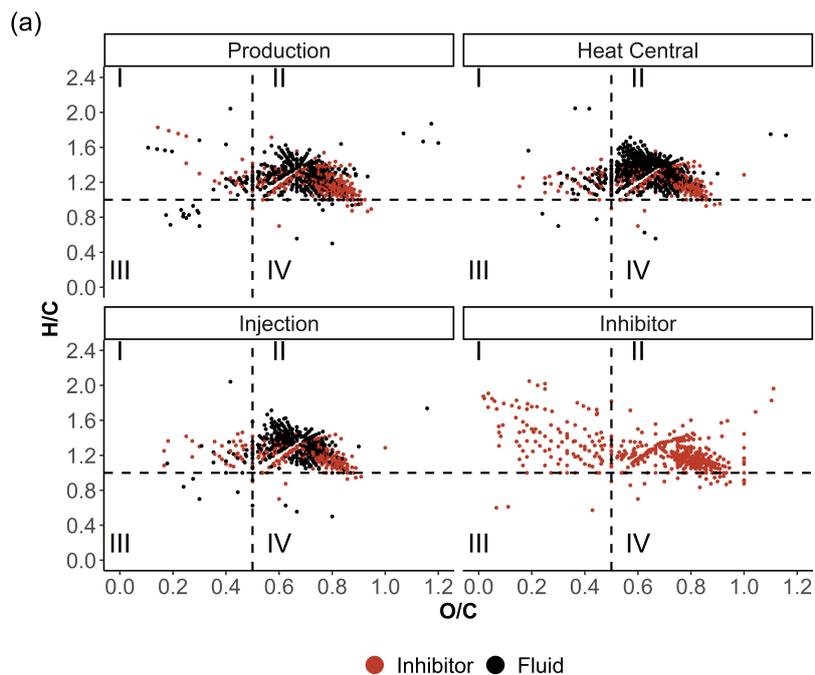
Figure 8. Van Krevelen diagrams of the fluid and inhibitor samples in (a) ESI(-) and (b) APPI(+) mode color-coded by matching signals in inhibitor and fluid samples.

~~deriving~~ derived from proteins (Sleighter and Hatcher, 2007). ~~Since microorganisms were reported for the~~ These signals may

Table 4. Comparison of the average and standard deviations of intensity-weighted averages of DBE, AI_{mod}, H/C, and O/C ratios of all detected APPI(+) and ESI(-) signals and signals where the inhibitor was removed from the dataset (w/o Inh.). The proportion of aromatic compounds is given as %TMIA.

	ESI						APPI					
	Production		Heat Central		Injection		Production		Heat Central		Injection	
	All	w/o Inh.	All	w/o Inh.	All	w/o Inh.	All	w/o Inh.	All	w/o Inh.	All	w/o Inh.
H/C ratio	1.26	1.29	1.31	1.36	1.29	1.33	1.17	1.23	1.20	1.30	1.18	1.22
O/C ratio	0.73	0.68	0.67	0.65	0.69	0.67	0.49	0.35	0.47	0.33	0.46	0.33
DBE	8.93	8.9	8.75	8.62	8.47	8.58	9.05	9.7	8.85	9.07	9.05	9.78
AI_{mod}	0.04	0.04	0.04	0.01	0.05	0.02	0.22	0.24	0.23	0.21	0.24	0.23
Aromatics %	0.9	0.8	0.7	0.5	1.6	1.3	3	1.6	3	1.6	2.6	1.8

represent microbial activity, since microorganisms have been reported for Bad Blumau fluids (Westphal et al., 2019), these signals may represent microbial activity. Generally, signals within the lipid group of the van Krevelen diagrams (H/C around 2 and O/C below 0.2) represent compounds deriving from cell membranes of microorganisms (Sleighter and Hatcher, 2007) and are therefore therefore are also good indicators for of microbial activity. Especially, the heat central sample shows an accumulation of signals in this area. It was assumed that the According to Westphal et al. (2019), microorganisms in the Bad Blumau fluids were likely to feed on the macromolecular organic matter introduced by the scaling inhibitor(Westphal et al., 2019). FT-ICR-MS data shows certain compound classes that are present in the inhibitor sample but absent in the fluid samples (Fig. 4). This is shown observed in the APPI(+) N_1O_x, S_1O_x, and $N_1S_1O_x$ as well as ESI(-) N_1O_x, and $N_1S_1O_x$ compound classes(Fig. 5). . The absence of these inhibitor specific compounds in the fluid samples could be explained by: (1) alteration in the form of chemical reactions with the fluid forming complexes, (2) degradation after the injection into the fluids, (3) no detection in the fluid samples due to strong dilution of the inhibitor, and (4) degradation by microorganisms that target these specific compounds.

The major absence of signals in area III and IV of the van Krevelen diagrams in both ionization modes suggests a low amount of highly condensed and aromatic compounds (Kim et al., 2003; Sleighter and Hatcher, 2007). For example the DOM of a dismal swamp presented by ESI FT-ICR-MS (Sleighter and Hatcher, 2007) and pore water of a Mangrove (?) reported a A wider range of signals with H/C ratios below 1 compared to the samples in this studywere observed in deep groundwater DOM (McDonough et al., 2022). A strongly different distribution of DOM signals was reported for coal water extracts (kerogen type III) where the majority of the signals was found at H/C ratios below 1 in area III and IV (Zhu et al., 2019).

Both ionization modes show only a few aromatic compounds that derive derived from the inhibitor. Those However, those deriving from the inhibitor however, have a notable effect on the TMIA in the APPI (+) datasetsince aromatic abundance is decreasing , since the aromatic abundance decreases in the inhibitor filtered data (Table 4). Aromatic and condensed aromatic compounds may be of natural origin from the reservoir or from other sources. For example, aromatic S_1O_x compounds detected

by ESI(-) mode in the fluid samples could be polystyrene-, naphthalene-, and lignosulfates which are used as superplasticizers in cement (Flatt and Schober, 2012; Hewlett et al., 2019) and could derive from the casing of the borehole. These data suggest the absence or only a very low impact of petroleum hydrocarbons on the fluids, since only a low proportion of DOM signals are present in area III and IV. A complete lack of signals was observed in area III and IV in the deep fracture water DOM of the Witwatersrand Basin, which was explained by the low contribution of hydrocarbons from the organic rich reefs of the Witwatersrand Supergroup (Kieft et al., 2018).

A major difference of the intensity-weighted average O/C values is shown when the Bad Blumau fluid samples are compared to other natural DOM detected by ESI(-) mode. The Bad Blumau samples exhibit high average O/C and H/C values from 0.65 to 0.68 and 1.26 to 1.33, respectively. The average values for marine, pore and river waters were reported in the O/C and H/C range of 0.32 to 0.52 and 1.13 to 1.29, respectively (Koch et al., 2008; Sleighter and Hatcher, 2008). Studies investigating the DOM of hydrothermal fluids reported average O/C values of 0.25 to 0.35 and generally high H/C ratios above 1.35 (Noowong et al., 2021; Gomez-Saez et al., 2016). Higher O/C values are generally attributed to tannine like compounds in area II and IV at O/C greater 0.5 and H/C around 1 (Sleighter and Hatcher, 2007) originating from terrestrial plant matter and some algae (De Leeuw and Largeau, 1993). However, compound types deriving from algal detritus and/or microbial biomass in marine sediments are reflected by lower O/C and higher H/C ratios (Sleighter and Hatcher, 2008), which is shown in our APPI(+) data but not the ESI(-) results. A contribution from higher plants is unlikely, since the Bad Blumau reservoir rock was formed during Palaeozoic reef development. However, data on deuterium and oxygen isotopes indicate an influence of meteoric water for the deep thermal Bad Blumau fluids (Goldbrunner, 2000). The higher O/C ratios in the ESI(-) data could be the result of terrestrial DOM being transported into the subsurface. However, within a geothermal setting, the DOM is likely to undergo thermal degradation. Elevated temperature experiments with marine DOM showed a preferential loss of high-molecular weight and oxygen rich molecules within two weeks of run time (Hawkes et al., 2016). The experiments were conducted in the temperature range of 100–380 °C and suggested that abiotic hydrothermal alteration may start at temperatures above 68 °C. It is unlikely that at the Bad Blumau reservoir temperatures of 124 °C the intensity-weighted average O/C ratios calculated for the ESI(-) formulas would show such high values for DOM originating from reservoir fluids. Another explanation for the still high O/C ratios could be the strong influence of artificial DOM, even after filtering all signals deriving from the inhibitor. Contrary to the ESI(-) data, the inhibitor filtered APPI(+) data show average O/C and H/C values from 0.33 to 0.35 and 1.23 to 1.3, respectively, which are in good agreement with hydrothermal vent DOM detected by ESI(-) mode (Noowong et al., 2021; Gomez-Saez et al., 2016). The average DBE values of both APPI(+) and ESI(-) measurements are generally in the same range as diffuse hydrothermal fluid DOM with temperatures up to 170 °C (Noowong et al., 2021; Gomez-Saez et al., 2016). Hotter fluids (>300 °C) were reported with much higher average DBE values. The low grade of aromaticity in the Bad Blumau DOM with average AI_{mod} values below 0.25 coincide with the hydrothermal vent and thermally altered marine DOM (Noowong et al., 2021; Gomez-Saez et al., 2016; Hawkes et al., 2016). It was stated that

500 ~~the elevated temperatures at~~ elevated temperatures under hydrothermal conditions likely resulted in a decrease ~~of~~ in molecular formulas detected by FT-ICR-MS due to a loss of thermally unstable DOM (Noowong et al., 2021; Hawkes et al., 2016; Rossel et al., 2017; Longnecker et al., 2018). This assumption could explain the relatively low number of formulas in the Bad Blumau fluid DOM ($< 10^3$), ~~since~~ as a similar range of detected formulas was reported for the hydrothermal DOM (Noowong et al., 2021).

505 Answering the first question posed for the discussion, we were able to characterize the major parts of synthetic DOM in the fluid and present a DOM fingerprint that better reflects the natural composition of the Bad Blumau geothermal fluids.

4.2 Variation of the composition of DOM and microbial community along the flow path

The DOM composition does not show much variation along the ~~flowpath. Most~~ flow path. The most notable are the signals in area I and III of the production side sample at H/C 1.7 and 0.8 in ESI(-) mode and the cluster in area I of the heat central at H/C
510 1.7 in APPI(+) mode (Fig. 8). The ESI(-) signals ~~, as previously discussed,~~ are either N_1O_x ~~compounds that derive from the inhibitor~~ or S_1O_x that ~~are likely introduced from~~ derive from the inhibitor or from other artificial sources. The distinct APPI(+) signals in the heat central ~~are~~ derived solely from the inhibitor. It ~~has to be taken into consideration that the samples from the three sampling points do not represent the same body of water. Thus, slight variations and differences such as inhibitor signals being present in the heat central and not in the other samples do not necessarily represent processes linked to the flowpath~~ is
515 important to note that FT-ICR-MS is not a quantitative approach, and therefore no assumptions regarding concentrations along the flow path can be made with these data.

~~In terms~~ However, a decrease of the DOC concentration ~~however, a decrease~~ along the flowpath was observed not only in this study but also in Westphal et al. (2019) ~~. This DOC decrease~~ and could be indicative ~~for~~ of microbial degradation. Especially ~~,~~ since a variety of microorganisms were detected in this study. ~~At the phylum level, the~~ The detection of Firmicutes,
520 Proteobacteria ~~,~~ and *Thermotogae* under the extreme environmental conditions in the three sampling points is not unlikely. ~~Indeed,~~ Firmicutes are known to form endospores (Cano and Borucki, 1995; Nicholson et al., 2002), which are highly resistant structures known to withstand conditions such as those in the power plant. Members of the Firmicutes (Filippidou et al., 2016) and Proteobacteria (Dib et al., 2008) have been detected in several extreme environments, showing their potential to withstand ~~the~~ environmental conditions in deep geothermal reservoirs. Members of the phylum *Thermotogae* can be either mesophilic,
525 thermophilic and hyperthermophilic, and most of the cultivated representatives have been obtained from extreme environments (Bhandari and Gupta, 2014).

~~At the production site, in~~ In addition to *Bacillus*, some *Pseudomonas*, *Desulfotomaculum*, *Planktothrix*, and ASVs related to the clade BRH-c8a were also detected in the production side sample. Species of *Bacillus* spp. are ~~able to form~~ capable of forming highly resistant endospores (Nicholson et al., 2000). ~~Thus~~ Therefore, the dominance of *Bacillus* at the production
530 site, ~~where the temperature is higher than at the other sampled sites~~ with the highest fluid temperatures, is not surprising. A previous bacterial diversity study performed in Bad Blumau (Westphal et al., 2019) already reported the detection of members of the genus *Desulfotomaculum*. ~~Moreover~~ Furthermore, this genus has been detected in other geothermal systems, such as different terrestrial hot springs (Amin et al., 2013; Poratti et al., 2016) or in geothermal ~~ground water (Daumas et al., 1988)~~

groundwater (Daumas et al., 1988). Therefore, it is also not surprising to find *Peptococcacea* closely related to *Desulfotomaculum* (BRH-c8a) (Sousa et al., 2018) in a geothermal system. Representatives of the genus *Pseudomonas* have been reported in the oxic zone of a geothermal system (Burté et al., 2019). The presence of Cyanobacteria belonging to the genus *Planktothrix* is more surprising. Cyanobacteria are known for their phototrophic metabolism and would need at least occasional light exposure for active growth (Puente-Sánchez et al., 2018). However, Cyanobacteria have recently been found in different environments that are not exposed to light, such as the deep subsurface or close to hydrothermal vents (Hubalek et al., 2016; Puente-Sánchez et al., 2018; Chen et al., 2022). Moreover, Cyanobacteria are commonly found in geothermal environments (Ward et al., 2012). This indicates that the presence of Cyanobacteria in different ecosystems, such as the deep geothermal system, is possible.

The dominant genera in the heat central corresponded to *Caulobacter*, *Desulfotomaculum* (also detected at the production and injection sites), *Sphingobium*, SCADC1-2-3, and *Microbacterium*. Members of the genus *Caulobacter* and *Sphingobium* were detected in the oxic and anoxic zone, respectively, of the same geothermal system in which *Pseudomonas* was reported (Burté et al., 2019). The SCADC1-2-3 is a group of uncultured organisms that belongs to the *Desulfisporaceae* family (Gavrilov et al., 2022). The SCADC1-2-3 group is part of a family of thermophilic sulfate-reducing bacteria and has been detected in subsurface waters (Gavrilov et al., 2022). The *Microbacterium* species described so far are known to be tolerant to extreme conditions, for instance, to the presence of arsenic (Achour-Rokbani et al., 2010), and have been isolated notably from a heated aquifer bore well (Adelskov and Patel, 2017), surface hot springs (Mehetre et al., 2019), and from the Atacama Desert (Mandakovic et al., 2020).

In addition to *Desulfotomaculum*, which was highly dominant in the injection well, *Fervidobacterium*, *Pseudothermotoga*, and with a low relative abundance, *Thermanaeromonas* and *Thermodesulfovibrio* were detected. The described species belonging to the genus *Fervidobacterium* are all anaerobic and extremely thermophilic, fermenting glucose to acetate and reducing sulfur to H$_2$S (Huber and Stetter, 2015). *Pseudothermotoga* are known thermophilic and anaerobic bacteria, isolated from hot springs and oil reservoirs (Farrell et al., 2021). *Thermanaeromonas* species are thermophilic anaerobes that are capable of forming spores (Mori and Hanada, 2015). They were notably isolated from a geothermal aquifer (Mori et al., 2002) and a subterranean clay environment (Gam et al., 2016). *Thermodesulfovibrio* are also anaerobic thermophilic organisms that are able to oxidize organic substrates to acetate (Maki, 2015) and have been isolated from hydrothermal vent waters (Henry et al., 1994) or hot springs (Sonne-Hansen and Ahring, 1999).

The results of our study are largely consistent with those of a previous study investigating bacterial diversity in the fluids collected in 2011 and 2013 at Bad Blumau (Westphal et al., 2019). In this previous study, samples were also taken at the production well and at the injection well. The third sampling point was located after CO$_2$ extraction, before the heat central. In our case, the third sampling point was located at the heat central itself. However, some differences were detected. In Westphal et al. (2019), the alpha-proteobacteria and beta-proteobacteria classes were found in the production fluids. In this study, alpha-proteobacteria were detected in the production fluids and after the heat central, but no beta-proteobacteria were detected as part of the most abundant classes (data not shown). *Clostridia* were detected in all

fluids by Westphal et al. (2019), which was also the case in this study. The classes Actinobacteria, Clostridia, and Thermotogae were the dominant classes detected in the injection well by Westphal et al. (2019). The Clostridia class was detected in the different samples in both studies, and the Clostridia and Thermotogae classes became dominant in the injection well, displacing Actinobacteria, which were only present in a low relative abundance. Our observations also confirm the presence of the Bacilli class as a dominant class at the production site. Previously, Ignavibacteria and the Nitrospira class were detected in the injection fluids, but in our case, Ignavibacteria were detected in a very low abundance in the fluids from the heat central and the injection well, and Nitrospira was not detected. However, the class Thermodesulfovibriona was detected in low abundance at the heat central and the injection fluids and was previously part of the Nitrospira class (Rabus et al., 2015; Umezawa et al., 2021). Therefore, the difference in the Nitrospira class in our analysis compared to Westphal et al. (2019) may only reflect a change in the bacterial taxonomy. The genus *Desulfotomaculum* was consistently detected in the production well, after CO_2 removal, and in the injection well in both studies. In the production well, Westphal et al. (2019) and the present study detected the class Bacilli and the genus *Pseudomonas*. Several genera, such as *Comamonas* and *Ralstonia*, were detected in the production fluids by Westphal et al. (2019), but were not among the top 5 genera observed in our study. Westphal et al. (2019) also detected *Thermodesulfovibrio*, *Desulfovirgula*, *Desulfovibrio*, *Fervidobacterium* and *Thermanaeromonas* at the injection well. In our case, *Thermodesulfovibrio*, the *Fervidobacterium* and *Thermanaeromonas* genera were also detected. One difference between the two studies lies in the detection of Cyanobacteria (*Planktothrix*) and *Planctomycetes* at the production well, but none of these phyla are dominant.

Overall, both studies highlight the changes in bacterial communities along the power plant system, from the production to the injection well. Such differences in communities highlight the impact of fluid conditions on bacterial communities, with some bacteria replacing others as conditions change within the system (Alawi et al., 2011; Lerm et al., 2013; Westphal et al., 2019). Furthermore, this is consistent with tests to change bacterial communities by adding nitrate to the fluids, which led to changes in the community. More importantly, the consistency between two independent studies demonstrates that the communities are relatively stable over several years. Differences in results may be due to differences in detection methods or small changes in the community. However, the dominant genera were those detected previously, and the changes in the communities from production to injection well followed the same patterns as observed by Westphal et al. (2019). This confirms that the presence of these microorganisms in the Bad Blumau system was not only temporary, as a byproduct of maintenance work, for instance. Furthermore, despite a relatively high flow rate, changes in the microbial community show that these conditions are compatible with the reactivation and development of a microbial community.

Answering the second question posed for the discussion, (a) the decrease in DOC content in our March 2021 samples is likely due to microbial degradation, and (b) the composition of the microbial community along the flow path is mainly driven by the given fluid temperature at the sampling points.

4.3 The role and origin of acetate: link to active microbial community

Acetate forms the dominant part of the DOC in the fluid samples. Several possible sources for its presence in our samples were considered: abiotic (e.g. water-oil contact or inhibitor derived) and biotic as a byproduct from active microbial communities. The contact of water and oil is a known source of organic acid anions in the fluid and was suggested to increase the LMWA content due to the release of hydrophilic acids (Reinsel et al., 1994). Oil-field waters are generally described to contain predominantly acetate, but also detectable amounts of formate, propionate, butyrate, and valerate (Carothers and Kharaka, 1978; Hatton and Hanor, 1984; Kharaka et al., 1985, 1997). The organic acid anion content is less likely to be associated with the water-oil contact in our samples due to the lack of butyrate, valerate, and only sporadic detection of propionate. However, the consumption by microorganisms could be another reason for the absence of these organic acids.

Thermodynamic conditions were found to be favorable for the abiotic generation of formate and acetate in deep fracture waters (Sherwood Lollar et al., 2021). However, the temperatures (25 °C) were significantly different from those in Bad Blumau. Abiotic acetate production has also been shown to occur at a temperature of 60 °C at 2 bar and in the presence of a greigite (Fe$_3$S$_4$) catalyst (Preiner et al., 2020). However, abiotic production of acetate due to water-rock reactions appears to be correlated with the presence of formate (Kieft et al., 2018; Sherwood Lollar et al., 2021; McDermott et al., 2015; Lang et al., 2010), which is absent in the Bad Blumau samples. It is unlikely that abiotic reactions have a significant effect on the production of the acetate in Bad Blumau, since no formate was detected.

Acetate is therefore most likely biogenic and is formed as a primary breakdown product of complex organic matter due to the microbial degradation by fermentative anaerobes. Metabolic pathways were predicted in our study using PICRUSt2 (Douglas et al., 2020) based on the composition of the microbial community. These predictions do not necessarily prove the existence of a given metabolism, but can be a useful aid to guide future enrichment or the selective detection of specific microbial groups/metabolic processes. As acetate was present in Bad Blumau in large amounts (14.2–18.2 mg L^{-1}), all pathways related to acetate were evaluated (Fig. 6). Many microorganisms have the potential to use or produce acetate. This is also the case within the Bad Blumau fluid systems, where the production and use of acetate by microorganisms would be possible at the production well, the heat central and the injection well.

Relatives of fermentative bacteria (*Thermoanaerobacter brockii*) were detected in the produced fluids by Westphal et al. (2019), which are described to produce lactic acid, acetic acid, H$_2$ and CO$_2$ as fermentation products. Macromolecular components of the scaling inhibitor are likely to act as an energy source and a carbon source for this fermentation process, as already suggested by Westphal et al. (2019). Some studies suggest that propionate and even butyrate may also be produced by the fermentation of organic matter (Sørensen et al., 1981; Lovley and Klug, 1986; Lovley

and Phillips, 1989). The sporadic detection of propionate in the samples could be indicative ~~for~~ of this process. The degradation products can be used as ~~substrate~~ substrates by other microbial communities. Cross-feeding interactions were described including sulfate reducing bacteria using lactate for sulfate reduction and~~in turn~~, in turn, providing hydrogen for hydrogenotrophic bacteria in the Bad Blumau fluids, leading to the formation of acetate as a degradation product~~(Westphal et al., 2019)~~. A gradual decrease ~~of sulfate was only observed in the~~ in sulfate was observed only in samples from June 2021~~samples, which might~~, which could be related to ~~the~~ sulfate reduction in the power plant (Westphal et al., 2019). However, it is more likely that these concentrations represent ~~its natural variability in the fluids~~~~since~~ the natural variability of the fluids, as the high fluid flow provides a continuous supply of sulfate.

~~Water-oil~~ Answering the third question posed for the discussion, water-oil contact and an abiotic origin can likely be excluded due to the absence of other LMWA's, while the inhibitor only induces a minimal amount of acetate (2.2 µg C L^{-1}). ~~To conclude~~In conclusion, a microbial origin of the acetate ~~due to~~ as a result of fermentative processes is likely to account for the ~~major part~~ majority of the acetate present in our fluid samples.

5 Summary and Conclusions

With regard to the ~~aims~~ objectives of this study, it was shown that 1) various methods ~~to characterize the DOM~~ were successfully used to ~~give~~ characterize DOM to provide a detailed description of the DOM present in the fluids. It was shown with LC-OCD that the DOC consists predominantly of ~~LMWAs and~~ LMWA's and was confirmed by IC that it is~~in fact~~, in fact, acetate. Using FT-ICR-MS, ~~the macromolecular DOM~~ macromolecular DOM was revealed to consist mainly of O_x and S_1O_x compounds ~~, deriving mostly~~ that are mainly derived from the synthetic scaling inhibitor~~and meeting the aim~~. Also objective 2) has been reached by FT-ICR-MS, to distinguish between natural and synthetic OM. ~~Overall, the scaling inhibitor adds approximately 1 of artificial DOC to the fluids.~~ DOM compounds that were found in the inhibitor but not in the fluids such as N_1O_x and $N_1S_1O_x$ might be absent due to (a) alteration ~~in form of~~ by chemical reactions with the fluid, forming complexes, (b) degradation after ~~the~~ injection into the fluids, (c) no detection in the fluid samples due to strong dilution of the inhibitor, or (d) degradation by microorganisms that target these specific compounds. With respect to objective 3)~~The acetate is likely biogenic~~, it can be concluded that acetate present in the fluids is likely of biogenic origin as a product of fermentative bacteria, turning lysine, pyruvate, or hexitol into acetate. The dominant genera involved in these processes were *Bacillus* and *Desulfotomaculum*. Last but not least, with view to objective 4) ~~It was observed that both~~~~the DOC concentration~~ it can be stated that both, the concentration of DOC and the microbial communities change along the flow path. The change in DOC content is probably caused by microbial degradation. However, in ~~this case, the temperature is likely the main driver of microbial community composition, since high temperatures are more limiting for microbial activity compared to the DOC concentration found here. However, in~~ terms of DOM composition~~no significant~~, no specific change was observed along the ~~flowpath~~ flow path and does not correlate with ~~the~~ changes in microbial diversity.

Additional insight ~~in~~ with regard to chemical fluid composition, processes ~~along~~ that occur along the flow path and the ~~flowpath and~~ origin of the ~~organic matter~~ dissolved organic matter was gained in the Bad Blumau geothermal power plant ~~was~~

670 ~~gained by assessing both ,~~ by evaluating both organic compounds and ~~microbial~~ the microbial community composition in the fluids. ~~These findings showed that both, the organic composition of a geothermal fluid and the microbial diversity~~ Both are rather complex. So far~~no indications were~~, no indications have been found that they contribute to changes in well productivity or injectivity~~of the wells~~. However, on a larger time scale the impact of the ~~organics~~ organic compounds can still be relevant, e.g. when decomposition processes are not possible in the inhibitor enriched fluids. Overall, this information ~~adds significantly~~

675 significantly adds to the understanding of ~~processes in the fluids of~~ fluid processes at a geothermal site and ~~might prove helpful to mitigate operational~~problems that could arise could be helpful in minimizing operational, such as biofilm formation and/or microbially induced corrosion.

Author contributions. AVH, PJ and SR designed the project. FE and DB took the samples. AL analyzed the organic compound data. DB analyzed the microbial data. GC performed the modeling of the metabolic capabilities of the present microorganisms. AL prepared and wrote
680 the major part the original draft with contributions from DB. AVH, FE, SP, PJ, and SR reviewed and edited the manuscript. All authors read and approved the final manuscript.

Competing interests. The authors declare that they have no conflict of interest.

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