Fluid Chemistry Evolution in Deep-Sea Hydrothermal Environments: Unraveling Mineral-Fluid-Microorganism Interactions through Continuous Culture Experiment

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Keywords: Gaz-lift Bioreactor; Lithium isotopes; Strontium isotopes; Microbial diversity; Geochemical modeling; Lucky strike hydrothermal field

Highlights

- Integrating Li/Sr isotopes tracers, microbial diversity and geochemical modeling.
- Microbial diversity of the bioreactor fluid reflects the sulfate-based chimney one.
- Microorganisms and minerals shape elemental and isotopic evolution in the fluid.
- $^{87}Sr/^{86}Sr$ ratio trace mineral-fluid-microorganism interactions, unlike $\delta^{7}Li$.

Abbreviations:

LSHF, Lucky Strike Hydrothermal Field

Buoyant HF, Buoyant Hydrothermal Fluid

Abstract

This study investigates minerals and microorganisms effects on fluid chemistry through a continuous enrichment culture in a gas-lift bioreactor during the MoMARsat'19 cruise. A sulfate-based chimney and buoyant hydrothermal fluid, both collected *in situ* at the Aisics vent of the Lucky Strike hydrothermal field, were incubated for 18 days under physico-chemical conditions mimicking those of *in situ* diffuse vents. We present the evolution of elemental and Sr, and Li isotopic compositions of the bioreactor fluid, alongside Bacteria and Archaea diversity, and analyze the mineral saturation state of the fluid through geochemical modeling. Our results reveal that the microbial diversity in the bioreactor reflects that of the sulfate-based chimney. During the initial 168 h, minerals precipitation/dissolution primarily controlled the elemental and Sr isotopic composition of the fluid. From 168 h to 264 h, sulfate-reducing Archaea (Archaeoglobi) disappeared in favor of sulfur-reducing Archaea (Thermoprotei and Thermococci). This coincides with a drastic increase in trace element concentrations and less radiogenic $^{87}Sr/^{86}Sr$ ratios, showcasing microbial influence on the fluid. From 264 h onwards, with stable sulfur-reducing archaeal diversity, mineral saturation state primarily controls the elemental composition of the fluid. However, we attribute the observed increase in the $^{87}Sr/^{86}Sr$ ratio and δ^7Li to changes in bacterial diversity, notably increasing Deinococci abundance. This study reveals that in diffuse vent environments related to the sulfur cycle: (i) microorganism and mineral influence fluid chemistry over time, (ii) microbial diversity affects trace metal concentrations and isotopic signatures, and (iii) the $^{87}Sr/^{86}Sr$ ratio trace mineral-fluid-microorganism interactions, unlike δ^7Li.

1 Introduction

Hydrothermal vents are distributed along the 67,000 km long mid-ocean ridge system and have a global impact on ocean chemistry, particularly on trace elements and their isotopes (Elderfield & Schultz, 1996; German et al., 2016). These environments, characterized by chemical and physical gradients, offer habitats that support microorganisms growth, making them among the most biologically active regions in the deep ocean (Holden et al., 2012; Zeng et al., 2021). This biological activity involves interactions not only between organisms and chemical species but also between organisms and minerals (Breier et al., 2010; Edwards et al., 2005; Holden et al., 2012; Rogers et al., 2003). Despite significant progress in understanding hydrothermal biogeochemical processes, further research is crucial, particularly in unraveling the complexities of mineral-fluid-microorganism interactions and their influence on biogeochemical cycles (Holden et al., 2012).

To better comprehend water-rock interaction processes, lithium (Li) and strontium (Sr) concentrations and isotopes are widely used to trace geochemical processes notably in geothermal and hydrothermal systems (Araoka et al., 2016; Barker et al., 2008; Chavagnac, Leleu, et al., 2018; Millot et al., 2010; Wang et al., 2023). However, their complex oceanic budget remains unresolved (Davis et al., 2003; Teagle et al., 2003; Tomascak et al., 2016; Vance et al., 2009). Moreover, despite their bioaccumulation in marine organisms (Chowdhury & Blust, 2011; Thibon et al., 2021), few studies investigate the Sr and Li elements in relation to aquatic ecosystems (Burger & Lichtscheidl, 2019; Thibon et al., 2021; Thibon et al., 2023). It is essential to study the impact of Li and Sr on marine biota, especially considering the growing economic interest on these elements, which leads to studies on their extraction from seawater (Hong et al., 2018; Ryu et al., 2020; Vikström et al., 2013), particularly Li from black smokers hydrothermal vents, which contain 10–20 times more Li than seawater (Chavagnac, Leleu, et al., 2018; *European Commission, Study on the EU's List of Critical Raw Materials – Final Report*, 2020).

This study combines for the first time the elemental and Li and Sr isotopic tracers (δ^7Li, and ^{87}Sr/^{86}Sr ratio) of the fluid with mineral saturation state obtained by thermodynamical modeling (PHREEQC), and microbial diversity analysis. This approach provides new insights into mineral-fluid-microorganism interactions. Studying microbial diversity at hydrothermal systems usually involves two approaches, the first one involves deploying *in situ* deep-sea devices (Alain et al., 2004; McCliment et al., 2006; Reysenbach et al., 2000; Rommevaux et al., 2019), while the second employs continuous enrichment culture in a laboratory. The second approach, used in this study, is the only one enabling the follow-up of microorganism interactions with dissolved chemical compounds and minerals overtime while controlling environmental conditions. Previous gas-lift bioreactor experiments were conducted to study deep-sea microbial communities evolution with constant fluid medium renewal (Callac et al., 2015; Godfroy et al., 2000; Godfroy et al., 2006; Postec, Pignet, et al., 2005; Postec et al., 2007; Raven et al., 1992). Only Callac et al (2015) collected both the inoculum and the culture medium *in situ* at a hydrothermal vent. In this study, we explore mineral-fluid-microorganism interactions through a continuous enrichment culture experiment conducted in a gas-lift bioreactor during the MoMARsat'19 cruise. Here, the inoculum is a portion of a hydrothermal chimney collected at the Aisics chimney at the foot step of the Tour Eiffel vent in the Lucky Strike Hydrothermal Field, and the culture medium is the Aisics' buoyant hydrothermal fluid, collected *in situ* between 100 and 150 °C. The sample is the top youngest part of the Aisics chimneys and is mainly composed of anhydrite. Anhydrite is commonly found in black smokers and impact marine biogeochemical cycle of calcium and sulfate. This study investigates the chemical evolution of the fluid once in contact with a sulfate-based chimney over an 18-day period, by analysing major and trace elements concentration, as well as lithium (Li) and strontium (Sr) isotopes, alongside mineralogy and microbial diversity. Furthermore, we discuss the impact of microorganisms on the Sr and Li concentrations and isotopic signatures of the fluid medium, unveiling new perspectives on the Li and Sr oceanic biogeochemical cycles.

2 Materials and Methods

2.1 Study area

The Lucky Strike Hydrothermal Field (LSHF) is located on the Mid-Atlantic Ridge at 37°17'N and 32°20'W, approximately 400 km to the Southwest of the Azores archipelago (Langmuir et al., 1997; Von Damm et al., 1998). This 1 km^2 hydrothermal vent field lies on a basaltic substratum and comprises 20 to 30 active vents distributed around a fossil lava lake (apart from Capelinhos vent) surrounded by three ancient volcanic cones (Charlou et al., 2000; Escartin et al., 2015; Fouquet et al., 1995; Langmuir et al., 1997; Ondréas et al., 2009; Von Damm et al., 1998). Fig. 1 presents the LSHF bathymetric map, at depths ranging between ~1550 and 1750 m below sea level (mbsl), with 12 active hydrothermal sites. Of specific relevance for this study is the Aisics chimney, located southeast of the fossil lava lake, at the base of the Tour Eiffel hydrothermal edifice.

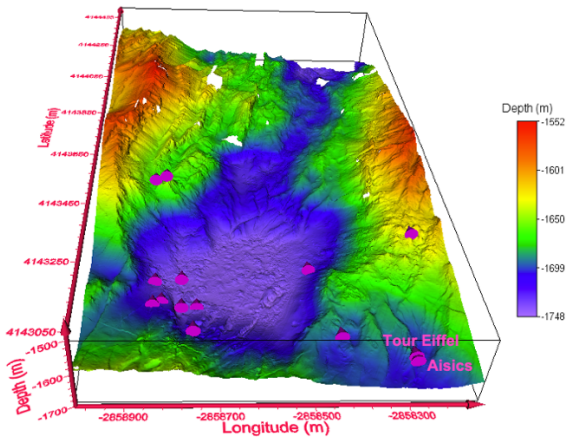

Fig. 1 3D bathymetric map of the Lucky Strike Hydrothermal Field (LSHF, Ondréas et al., 2009). Active vent locations are indicated by pink 3D cones.

2.2 Sample collection and onboard processing

During the MoMARsat'19 EMSO-Azores maintenance cruise aboard the *R.V. Pourquoi Pas?* (June - July 2019, Sarradin & Legrand, 2019), hydrothermal materials were collected at the LSHF using the hydraulic arm of the Human Operated Vehicle (HOV Nautile 6000). Successively, samples of hydrothermal chimney (Fig. 2a), buoyant hydrothermal fluid (buoyant HF, Fig. 2b), and high temperature hydrothermal fluid (end-members, Fig. 2C) were collected at the Aisics vent site. Upon recovery of the HOV Nautile on the research vessel, all samples were processed immediately in a shipboard chemical laboratory (Class 100 000, ISO 8).

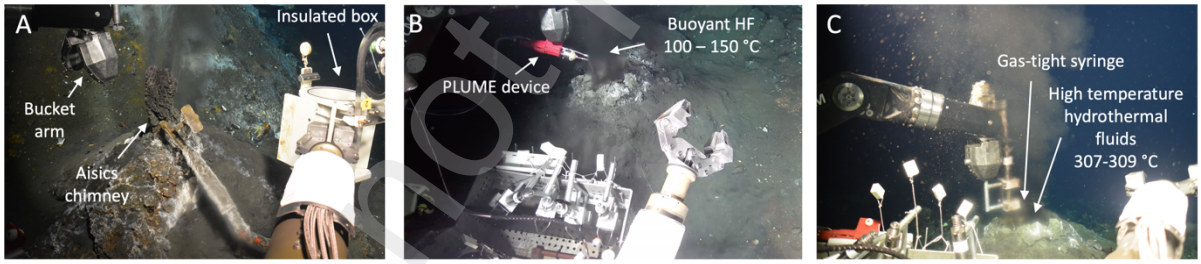

Fig. 2 Chronological overview of scientific operations conducted at Aisics vent site. Snapshots of the HOV Nautile (Sarradin & Legrand, 2019) (A) Collection of Aisics' chimney sample using the bucket arm and an insulated box, (B) Sampling of the buoyant hydrothermal fluids (buoyant HF) with the PLUME device, and (C) Sampling of high-temperature hydrothermal fluids via gas-tight titanium syringe.

The anhydrite-bearing top of the Aisics chimney (sample number MOM19_Aisics1, PL 1939-1, June 12th 2019) was collected using the bucket arm of the HOV Nautile and then dropped into a decontaminated insulated box (Fig. 2a). Before use, the insulated box was cleaned, disinfected with ethanol, and then filled with sterile distilled water (30 min, 121°C). To prevent atmospheric contamination before the dive and seawater contamination during the descent and ascent in the water column, the insulated box was opened and closed at depth upon chimney collection. Upon recovery of the HOV Nautile on the research vessel, the hydrothermal chimney was transferred into a sterile mortar under a laminar flow hood, and immediately

ground in a controlled anaerobic chamber under an N_2/H_2 (90:10) atmosphere. The ground chimney was then stored in a glass flask until its introduction into the gas-lift bioreactor tank.

The buoyant HF was collected unfiltered into 5L PVC/DEHP blood bags (Promepla, , sterilized by ethylene oxide) via the PLUME fluid pumping system implemented on the HOV Nautile (Fig. 2b). Prior to each dive, the sampling tubes and cannulas of the PLUME device were rinsed with Milli-Q water, then filled with a small volume of Milli-Q water to prevent depression during descent. The temperature sensor, attached to the snorkel of the PLUME device, guided the HOV pilot in collecting the buoyant HF within the mixing gradient at temperature between 100 and 150°C. All buoyant HF used in this study were collected at the Aisics vent at similar *in situ* temperatures during dives numbered 1939 on June 12th (2 blood bags; samples number MOM19_PL1939-1_PLUME3 and MOM19_PL1939-1_PLUME2), 1941 on June 14th (2 blood bags, > 4L sample number MOM19_PL1941-3_PLUME2 and MOM19_PL1941-3_PLUME3), and 1955 on June 30th (1 blood bag, > 4L sample number MOM19 Aisics PL1955-17_PLUME3). Upon recovery, each filled sterile blood bag was closed and stored at 4 °C in a dark room prior to connection to the gas-lift bioreactor.
Prior to high temperature hydrothermal fluid sampling (Fig. 2c), the fluid temperature was measured *in situ* at 307 – 309 °C by inserting the HOV high temperature probe into the chimney. A total of eight high temperature hydrothermal fluid were sampled during two dives, on dive 1939 (June 12th, samples M19FLU01 to M19FLU04) and dive 1955 (June 30th, samples M19FLU49 to M19FLU52, Supplementary Material Table S1). These fluids were collected using 200 mL gas-tight titanium syringes. Prior to each dive, gas-tight titanium syringes were washed with diluted hydrochloric acid, then rinsed with ultrapure milli-Q water. The syringe snorkel was inserted into the chimney and operated individually by the hydraulic arm of the HOV Nautile (Fig. 2c). Upon recovery, the high temperature hydrothermal fluids were extracted from gas-tight titanium syringes, filtered through 0.22 µm Millipore filters, split into distinct aliquots for onboard and onshore analysis, and stored at 4 °C in a dark room. Their chemical composition, analyzed using onshore instrumental and analytical facilities, allows the characterization of pure hydrothermal fluid end-member after extrapolation to Mg-zero prior to its dilution with the surrounding North Atlantic Deep Water (NADW).

For chemical analysis aboard the research vessel, pH, conductivity, salinity, Total Dissolved Solids (TDS), and redox potential (Eh) were measured immediately after fluid extraction using the Consort C562 multi-parameter analyzer. Total Sulfur (TS) and hydrogen sulfide (H_2S) contents were measured with an amperometric micro-sensor (AquaMS, France), connected to both temperature and pH electrodes to establish equilibrium. Dissolved Fe (dFe) concentrations were measured with the HI96721 Iron High Range Photometer (range from 0 to 5 mg/L, Hanna instruments). The instrument was calibrated and validated using Hanna CAL CHECK™ Standards. Prior to each day's measurements, the instrument was zeroed with a blank solution. The measurement accuracy is ± 0.04 mg/L ± 2% of reading.

2.3 Gas-lift bioreactor

A continuous enrichment culture experiment was conducted aboard the *R.V. Pourquoi Pas?* during the MoMARsat'19 cruise (Sarradin & Legrand, 2019), using a gas-lift bioreactor as shown on Fig. 3 (Callac et al., 2015; Godfroy et al., 2006; Postec, Urios, et al., 2005; Postec et al., 2007).

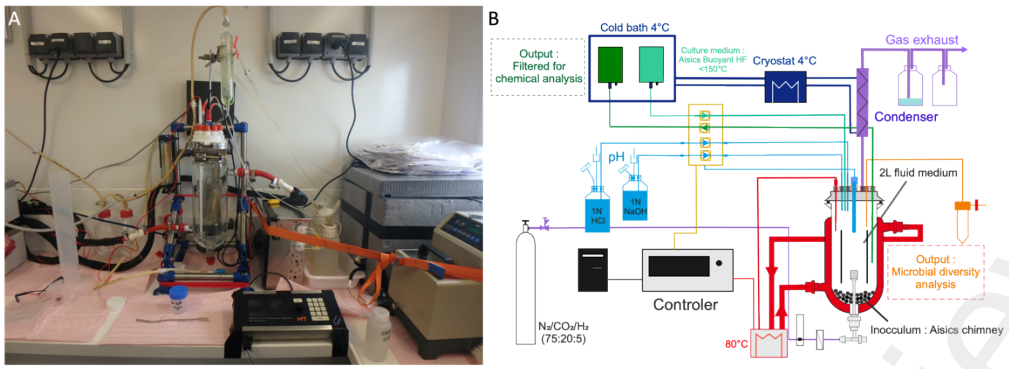

Fig. 3 Set-up of the continuous enrichment culture experiment conducted aboard the *R.V. Pourquoi Pas?*. (A) Aboard photograph of the experiment in the laboratory. (B) Schematic illustration of the experimental setup, adapted from Godfroy et al (2006).

The aim of this experiment was to gain understanding on the chemical evolution of the fluid medium correlatively with hydrothermal microbial diversity and/or mineral precipitation/dissolution in a setting closely mimicking the *in situ* conditions of the diffuse hydrothermal environment.

The culture medium used in this study is the buoyant HF collected at the Aisics vent, with temperatures ranging between 100 and 150°C (see section 2.2). On June 13th, 2 L of the buoyant HF (sample MOM19_PL1939-1_PLUME3) were introduced into the gas-lift bioreactor tank. The conditions inside the bioreactor maintain a temperature of ~80 °C, a pressure of 1 atm, and anaerobic conditions thanks to a continuous gas flow of N_2:CO_2:H_2 (75:20:5 proportions, 10 cm^3/minute flow rate). This continuous gas flow provides H_2 as the electron donor and CO_2 as the carbon source. During the experiment, the pH was controlled around 6.5 at 80°C (dead zone +/- 0.2) by adding either a 1 N HCl or 1N NaOH solution (Godfroy et al., 2006). After an hour and a half to reach incubation temperature and gas equilibrium, a sample of the culture medium (MOM19_FERM_T-1, Table 1) was filtered through a 0.22 µm Sterivex filter and split into different aliquots for subsequent aboard and onshore chemical analyses. Then, a 150 ml portion of the Aisics hydrothermal chimney (MOM19_Aisics1, see section 2.2) was used as the inoculum and introduced into the bioreactor tank; this corresponds to the start of the experiment at T0. By then, we will use the term "fluid medium" to refer to the aqueous phase collected all along the experiment (samples labeled MOM19 _FERM_T1 to T+18 for chemical analysis, Table 1, and MOM19.FerT0 to T18 for microbial analysis, Supplementary Material Table S2). The water/rock ratio is equal to 13 in the gas-lift bioreactor. The chimney was allowed to react with the buoyant HF for an 1 h 15 min before sampling for microbial diversity (MOM19.FerT0). Fifteen minutes after this sample, the continuous injection and withdrawal of fresh fluid began at a rate of 0.025 L/hour, maintaining a 2 L culture tank capacity. Throughout the 18-day/432-hour experiment, a total of four buoyant HF blood bags (collected on dives PL1939 and PL1941, see section 2.2) were used to maintain the capacity of the culture tank and were connected to the bioreactor at 0 h, 5 h, 120 h, and 288 h, respectively.

The mineralogical composition of the chimney sample was determined by X-Ray Diffraction analysis (XRD) both upon collection and at the end of the experiment. Upon collection, the inoculum was composed of 93 % of anhydrite ($CaSO_4$), 3 % of pyrite (FeS_2), 2 % halite (NaCl) and 1 % of chalcopyrite ($CuFeS_2$). By the end of the experiment, the inoculum was composed of 91 % of anhydrite, 3 % of pyrite, 3 % halite, and 2 % of chalcopyrite (François, 2021).

During the course of the experiment it was not possible to sample the inoculum in the bioreactor.

For chemical analysis (notably elemental and isotope analysis), aliquots of (i) culture medium were extracted before inoculation (MOM19_FERM_T-1, Table 1), and (ii) fluid medium 24 h after the beginning of the experiment and then at 48-hour intervals over the following 18 days, resulting in a total of 10 additional samples (samples labeled MOM19_FERM_T1 to T18, Table 1). Each of these 10 samples corresponds to a 24h bioreactor withdrawal, and was collected in 600ml sterile Terumo blood bags, stored at 4°C (dark green output bags in Fig. 3B) and then filtered through 0.22 µm Sterivex filter prior to onboard and onshore chemical analyses.

For microbial diversity analysis (bacterial and archaeal diversity), both the chimney sample (sample MOM19_Aisics1) and the buoyant HF fluid (sample M19PL1955_PLUME3 filtered on Sterivex) were stored at - 80°C for subsequent onshore analysis. During the experiment, 50 ml aliquots of the fluid medium were sampled daily (orange falcon on Fig. 3B) and stored at -80°C for microbial diversity analysis, resulting in a total of 19 samples (samples labeled MOM19.FerT0 to T18, Supplementary Material Table S2). Additionally, 1ml aliquots stored in a 9ml Sea water/2% formaldehyde solution and then were filtered through 0.22µm Nuclepore filters, stained with Sybr Goldfor cell counting with an Axio Imager Z2 Apotome microscope (Carl Zeiss MicroImaging GmbH, Göttingen, Allemagne).

2.4 Microbial diversity analysis

The microbial diversity analysis detailed below was conducted on the buoyant HF (M19PL1955 Plume 3 labeled 19.Ais.100.150 for molecular analysis), the chimney sample (MOM19_Aisics1 labeled M19.Ais1 for molecular analysis), and on eighteen fluid medium samples (samples labeled MOM19.FerT0 to T18, Supplementary Material Table S2). A nested-PCR approach was used to amplify the variable regions V3-V4 of the archaeal 16S rRNA genes. The full-length archaeal 16S rDNA was amplified using the primers A24F-1492R (CGGTTGATCCTGCCGGA ; GGCTACCTTGTTACGACT, Lepage et al., 2004; Teske et al., 2002). The PCR products were gel purified and used as a template to amplify the V3-V4 region by using the primers A344F-archaea806R (AYGGGGYGCASCAGGSG ; GGACTACVSGGGTATCTAAT, Stahl, 1991; Takai & Horikoshi, 2000). The archaeal 16S rRNA genes libraries were sequenced with Illumina MiSeq at MR DNA (Shallowater, TX, USA). The V3-V4 bacterial 16S rRNA genes libraries were prepared and sequenced with Illumina MiSeq at MR DNA (Shallowater, TX, USA) using the primers whoi341-whoi785R (CCTACGGGNGGCWGCAG ; GACTACHVGGGTATCTAATCC, Herlemann et al., 2011). The metabarcoding data were processed using the pipeline SAMBA (https://github.com/ifremer-bioinformatics/samba) which is based on QIIME 2 (Bolyen et al., 2019). Primers and barcode were removed using cutadapt (Martin, 2011), with the following parameters (errorRate = "0.1" ; overlap = "5"). Trimming of short reads and low quality sequences, ASVs inference and removal of chimeric sequences were performed using DADA2 (Callahan et al., 2016) with the following parameters (FtrimLeft = "20" and RtrimLeft = "80" for *Archaea* ; FtrimLeft = "30" and RtrimLeft = "90" for *Bacteria* ; FmaxEE = "6" ; RmaxEE = "6" ; minQ = "3" ; chimeras = "consensus"). An additional step of ASV clustering has been performed using the dbOTU3 algorithm to avoid an overestimation of the diversity (Olesen et al., 2017). Taxonomic assignment of processed sequences was performed using the SILVA v138 reference database (Quast et al., 2012).

2.5 Elemental and isotopic analysis

All the chemical analyses were conducted at the Observatoire Midi-Pyrénées (Toulouse, France).

The analytical methods used for major dissolved cations (dCa, dK, dMg, dNa, dSi), anions (dCl, dBr, dSO_4), and trace element (dBa, dFe, dMn, dLi, dSr) analyses are detailed in Besson et al (2014), Leleu (2017), Chavagnac et al (2018), and Artigue et al (2022) and will be briefly described below.

Bioreactor samples (samples labeled MOM19_FERM_T-1 to T18, Table 1) were diluted with Milli-Q water 30-fold for dCa, dK, dNa, and 10-fold for dMg and dSi concentration measurements. Other trace element concentrations were measured in pure solutions. All these element concentrations (except for dFe) were measured using an inductively coupled plasma atomic emission spectrometer (ICP-AES) Horiba Ultima2 instrument, with an analytical precision better than 2%. The ICP-AES was calibrated using mono elemental solution and an IAPSO seawater standard solution (OSIL Ltd. UK) diluted 10 to 200-fold with Milli-Q water. The analytical drift was quantified by the standard bracketing method every 8 samples. Detection limits were determined through daily repeated blank solutions (n = 10) at 0.2 µmol/L for dCa, 2 µmol/L for dK, 0.3 µmol/L for dMg, 20 µmol/L for dNa, 0.14 µmol/L for dSi, 0.01 µmol/L for dBa, 0.02 µmol/L for dMn, 0.3 µmol/L for dLi, and 0.01 µmol/L for dSr.

Anion concentrations were measured in 10-fold diluted samples, and determined by anionic chromatography (Dionex ICS-2000) equipped with a specific column for a highly charged matrix (DIONEX IC AS19). The instrument was calibrated with IAPSO seawater standard diluted 10 to 50 folds with Milli-Q water. The instrument's error is 0.0001 ppm.

For isotopic measurements, all fluids were processed in a clean laboratory to isolate Li and Sr from their matrix using conventional liquid chromatography. For each element, 1 mL of individual fluid samples was evaporated to dryness in a Savillex beaker on a hot plate at 70 °C. The IAPSO and NASS 6 international standards were processed in the same manner and used alongside our samples.

Dissolved Sr was separated from the matrix using Sr-Spec resin (Eichrom, USA) following Pin et al (2014) protocol. Sr isotopic ratio ($^{87}Sr/^{86}Sr$) was measured using a Thermo Fisher Triton+ Thermal Ionization Mass Spectrometer. The $^{87}Sr/^{86}Sr$ ratio was defined as the average values of 150 measurements of ion intensities in the static multi-collection mode. The $^{87}Sr/^{86}Sr$ ratios were corrected from mass fractionation using the $^{86}Sr/^{88}Sr$ normalization ratio of 0.1194. Repeated measurements of the NBS 987 Sr standard gave a mean ratio of 0.710259 ± 0.000013 (2 SD; n = 24; 2SE = 0.000003). The $^{87}Sr/^{86}Sr$ ratios of our samples were corrected from the deviation of the measured NBS 987 to the recommended NBS 987 value of 0.710248.

The $^{87}Sr/^{86}Sr$ ratios of international standards were measured to verify the accuracy of the measurements: 1. IAPSO seawater with a measured value of 0.709174 ± 0.000003 (2SD; n = 4), consistent with published values of 0.709179 ± 0.000007 (2SD; n = 7; El Meknassi et al., 2020), and 2. NASS-6 seawater with a measured value of 0.709174 ± 0.000005 (2SD; n = 3), consistent with published values of 0.709179 ± 0.000014 (2SD; n = 8; Neymark et al., 2014).

Dissolved Li was separated from the NaCl-rich matrix using two steps ion exchange columns made of AGW-X12 200-400 mesh cation resin bed and eluted with 1N HCl (Protocol adapted to NaCl-solution from James & Palmer, 2000). The Li isotopic composition of each fluid sample was measured on a Thermo Fisher Triton+ Thermal Ionisation Mass Spectrometer at

the Observatoire Midi-Pyrénées. Additional information regarding sample loading and mass spectrometer setup can be found in Artigue et al (2022). The $^{7}Li/^{6}Li$ ratios are expressed in the $\delta^{7}Li$ ‰ notation relative to the IRMM-16 Li standard (Li_2CO_3) at similar Li concentration. Repeated measurements of the IRMM-16 standard gave a mean $^{7}Li/^{6}Li$ ratio of 12.082 ± 0.012 (2SD; n = 17), an internal precision of 0.25 ‰ (2SE) and an external precision of 1.03 ‰ (2SD; n = 17). The accuracy of our technique was verified against the measured ratios of the international IAPSO seawater standard with a measured value of +29.5 ± 0.2 ‰ (2SE), consistent with published values of +30.8 ± 0.1 ‰ (2SE, with external precision ≤ 1‰, Rosner et al., 2007).

2.6 Geochemical modeling

Geochemical modeling was performed with the PHREEQC software package developed by USGS (Graphical User Interface Version 3, www.usgs.gov/software/phreeqc-version-3, Parkhurst & Appelo, 2013). PHREEQC can be used as a speciation program, particularly to calculate the distribution of aqueous species, and the possibility of mineral dissolution/precipitation. To perform these speciation calculations, PHREEQC requires the major elements total concentrations of the solution (user input) and the specific equilibrium constants from the PHREEQC databases. The saturation state of the fluid regarding to minerals is given by its saturation index (SI), which is calculated as the logarithm of the chemical activities of the dissolved ions (ion activity product, IAP) over their solubility constant (K). The possibility of a mineral to dissolve or precipitate is characterized by either undersaturation (SI < 0) or oversaturation (SI > 0).

In this study, the PHREEQC program was used along with the "llnl.dat" database (Johnson et al., 1992). This database provides logarithms of equilibrium constants (log K) along with thermodynamical data available up to 300 °C. The speciation modeling was run twice on the buoyant HF: first at its *in situ* temperature (126 °C), and then after reaching the incubation temperature (80 °C) and gas equilibrium of the gas-lift bioreactor. Subsequently, it was run on each aliquot of fluid medium extracted after inoculation from the gas-lift bioreactor (10 samples, MOM19_FERM_T1 to T18). The chemical composition of the fluids was input as total concentrations of all previously analyzed chemical elements. To model the continuous gaseous flush of N_2:CO_2:H_2, thermodynamic equilibrium was established between the fluid and a gas phase at a total pressure of 1 atm and at the N_2:CO_2:H_2 proportions (75:20:5) of the gas-lift bioreactor setup.

In the modeling, the oxidation potential was calculated regarding the H_2S/SO_4^{2-} redox couple. Li and Sr aqueous species distribution was calculated as well as the saturation indices (SI) for both the buoyant HF and the fluid medium throughout the entire duration of the bioreactor experiment.

3 Results

3.1 Geochemistry of pure hydrothermal fluids

The geochemical features of the high temperature hydrothermal fluids collected during MoMARsat'19 cruise are reported in Supplementary Material Table S1. High temperature hydrothermal fluids exhibit pH values at 25 °C ranging between 3.58 and 4.40, and chemical enrichment in dissolved Ca, K, Si, Fe, Mn, Si, and Li (dCa, dK, dSi, dFe, dMn, dLi) compared to seawater. Since pure hydrothermal fluid should be totally dMg-depleted, the end-member composition is obtained by linear extrapolation to zero-Mg of the least-square regression method (Von Damm et al., 1998). The result is similar to previous end-member chemical features obtained at this site (Chavagnac, Leleu, et al., 2018; Leleu, 2017). However, H_2S concentrations of 4.1 to 11.9 mmol/L, are much higher than previous values of 2 - 4 mmol/L (Charlou et al., 2000; Chavagnac, Saleban Ali, et al., 2018; Pester et al., 2012; Von Damm et al., 1998).

3.2 Elemental and isotopic composition of bioreactor fluid medium

The elemental and isotopic composition of the buoyant HF, and all fluid medium extracted from the gas-lift bioreactor are reported in Table 1 and are shown in Fig. 4.

	Buoyant HF (80 °C)	Fluid medium (80 °C, sample labeled MOM19_FERM_T1 to T18)									
	MOM19_FERM_T-1	T1	T3	T5	T7	T9	T11	T13	T15	T17	T18
	Before Inoculation	24 h	72 h	120 h	168 h	216 h	264 h	312 h	360 h	408 h	432 h
Aboard analysis											
pH	6.21	6.66	6.42	6.73	6.60	6.93	6.97	7.15	7.02	7.10	6.52
Total S (mg/L)	190	25	13	22.1	15	17	145	132	54	92	98
H_2S (mmol/L)	1.0	0.6	0.1	0.5	0.2	0.4	0.2	1.3	0.8	1.1	1.8
Eh (mV)	3	-92	-18	118	-47	124	129	138	215	46	88
Conductivity (mS/cm)	43.5	46.6	47.3	46.7	46.4	45.1	45.9	44.6	47.1	47.2	47.2
TDS (g/L)	25.4	24.6	27.5	26.5	26.9	26.3	26.6	26	27.4	27.4	27.5
Onshore analysis											
dMg (mmol/L)	28.43	30.79	31.38	30.98	31.47	30.43	30.11	32.33	30.88	32.91	33.9
dCa (mmol/L)	20.37	24.39	23.12	23.01	23.28	24.23	23.94	22.78	22.03	21.5	20.88
dK (mmol/L)	13.97	14.12	14.96	14.1	14.57	16.59	15.65	15.27	14.75	15.94	15.55
dNa (mmol/L)	417.4	429.5	424.7	409.6	419.8	428.1	418	407.2	408.3	421.2	409.6
dFe (µmol/L)	75.1	24.8	20.5	16	21.8	10.8	12.2	16.2	7.6	5.4	4.7
dMn (µmol/L)	65.7	50.2	46.5	45.7	48.3	37.5	90.4	87.7	66.1	46.4	40.8
dSi (mmol/L)	1.68	1.67	1.76	1.82	1.93	1.96	1.94	1.92	2.1	2.32	2.22
dCl (mmol/L)	498.3	511.3	512.1	508.2	505.5	502.3	506.1	509.2	515.8	519.1	520.8
dSO_4^{2-} (mmol/L)	18.25	24.03	23.22	22.93	22.38	21.95	22.05	22.93	24.65	25.56	26.35
dBa (µmol/L)	1.5	1.4	2.9	2.6	2.4	3.1	4.5	4.3	3	2.4	2.5
dBr (µmol/L)	812	827	823	820	818	812	823	827	832	839	836
dSr (µmol/L)	79	77	75	72	68	85	142	145	133	118	106
$^{87}Sr/^{86}Sr$	0.707513	0.707488	0.707492	0.707483	0.707404	0.707363	0.707337	0.707439	0.707646	0.707760	0.707794
± 2SE	± 0.000004	± 0.000004	± 0.000004	± 0.000004	± 0.000004	± 0.000005	± 0.000005	± 0.000004	± 0.000005	± 0.000005	± 0.000004
dLi (µmol/L)	208	162	158	149	166	226	369	387	251	214	179
δ^7Li (‰)	7.2	4.6	7	6.3	4	7.3	4.6	6.1	7.3	10.3	6.3
Ratios											
dNa/dCl	0.84	0.84	0.83	0.81	0.83	0.85	0.83	0.8	0.79	0.81	0.79
dNa/dLi	2.01	2.66	2.68	2.74	2.52	1.89	1.13	1.05	1.62	1.97	2.28
dMg/dLi	0.14	0.19	0.20	0.21	0.19	0.13	0.08	0.08	0.12	0.15	0.19

Table 1 Geochemical compositions in the gas-lift bioreactor of the buoyant hydrothermal fluid (buoyant HF) before inoculation (MOM19_FERM_T-1), of the fluid medium 24h after inoculation (MOM19_FERM_T1), and of fluid medium sampled at a 48-hour intervals until the end of the experiment (MOM19_FERM_T3 to T18).

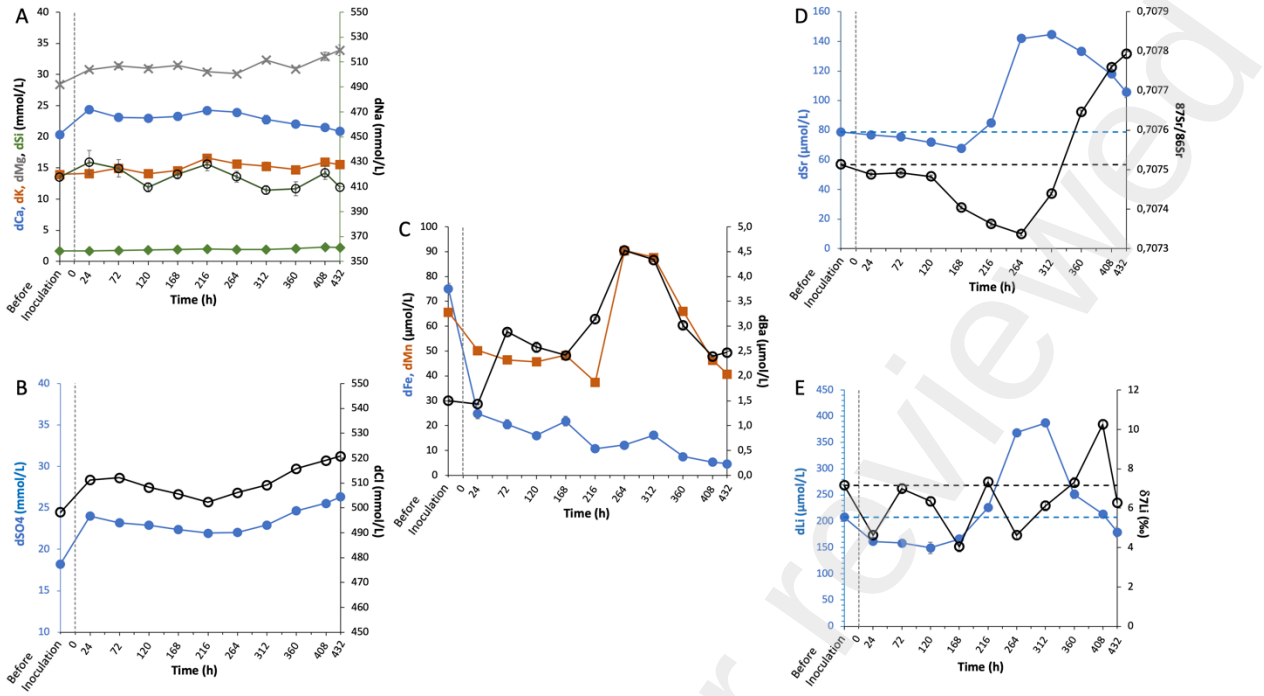

Fig. 4 Temporal evolution of the geochemical composition in the gas-lift bioreactor of the buoyant hydrothermal fluid (Buoyant HF) before inoculation, and of the fluid medium 24h after inoculation and at 48-hour intervals until the end of the experiment. The start of the experiment i.e. inoculation time is indicated at 0 h. (A) Major cation concentrations, (B) Major anion concentrations, (C) dFe, dMn, dBa concentrations, (D) dSr concentrations and $^{87}Sr/^{86}Sr$ ratios, and (E) dLi concentrations and $\delta^{7}Li$ ‰ values. For (D) and (E), horizontal lines denote the initial values, aiding in visualizing fluctuations from the collection time to the end of the experiment. All data are plotted with their respective uncertainties.

The pH of the buoyant HF in the bioreactor before inoculation is 6.21. Throughout the experiment, pH values ranged between 6.42 and 7.15 (Table 1). Dissolved Na and dCl vary from 407 to 430 mmol/L and 502 to 521 mmol/L, respectively, and the dNa/dCl ratios show little variation at 0.79 – 0.85 compared to a standard seawater ratio of 0.86 (Millero et al., 2008). Between the sampling of the buoyant HF in the gas-lift bioreactor before inoculation (MOM19_FERM_T-1, Table 1) and the first sampling of the fluid medium for chemical analysis 24 hours after inoculation (MOM19_FERM_T1, Table 1), element concentrations exhibit various trends. Concentrations of dNa, dK, dCl, dBr, dBa, and dSr remain fairly constant, while dSO_4, dCa, and dMg increase by up to 32%, 20%, and 8%, respectively. Conversely, concentrations of dFe, dMn, and dLi decrease by 67%, 24%, and 22%, respectively (Fig. 4). Then, from 24 h to 168h, element concentrations are overall constant, with slight reductions of dCa, dSO_4, dMn, and dSr concentrations by up to 5%, 7%, 4%, and 12%, respectively (Fig. 4a, b, and d). Between 168 h and 264 h, major element concentrations remain stable; however, all trace element concentrations, except dFe, doubled (dMn, dBa, dSr, and dLi). Subsequently, from 264 h to the end of the experiment, trace elements, along with dCa, decreased progressively: dMn by 55%, dLi by 51%, dBa by 45%, dSr by 26%, and dCa by 13%, while dSO_4 and dMg concentrations increased by up to 20% and 13%, respectively. Regarding the variability in dFe, it continuously decreases over time, by up to a factor of 5 by the end of the experiment (Fig. 4C).

The $^{87}Sr/^{86}Sr$ ratio of the gas-lift bioreactor culture medium before inoculation was measured at 0.707513 ± 0.000004 (MOM19_FERM_T-1, Table 1). From this sampling time to 24 h after the experiment start, the $^{87}Sr/^{86}Sr$ ratio of the fluid medium decreases down to 0.707488, remaining relatively stable until 120 h, before decreasing to its minimum value of 0.707337 at 264 h. From 264 h onwards, the $^{87}Sr/^{86}Sr$ ratios continuously increase reaching a maximum value of 0.707794 by the end of the experiment (Fig. 4d). Regarding the Li isotopic composition, the δ^7Li values show a large variation between +4.0 and +10.3 ‰ with a median value of +5.5 ‰ without any specific temporal evolution (Fig. 4e).

3.4 PHREEQC geochemical modeling: saturation state and aqueous speciation

PHREEQC geochemical modeling is controlled by the physico-chemical conditions of the experiment. It allows a thermodynamic diagnosis of the reactivity in the bioreactor fluid medium throughout the experiment, but doesn't account for kinetics. The mineral saturation indexes and element species abundances calculated by PHREEQC are presented in Table 2. Considering the observed minerals in the inoculum, thermodynamic calculations evidence the continuous undersaturation state of the medium fluid regarding anhydrite ($CaSO_4$) throughout the experiment, suggesting anhydrite dissolution. However, the saturation indices of sulfides such as pyrite (FeS_2) and chalcopyrite ($CuFeS_2$), as well as oxides such as hematite (Fe_2O_3), calcite ($CaCO_3$) and other carbonates, and even barite ($BaSO_4$) are all positive. This suggests either their stability in the fluid medium (for chalcopyrite) or their potential precipitation. Note that the continuous flushing of N_2:CO_2:H_2 gaseous phase ensures anaerobic conditions which preserve sulfur mineralization.

Dissolved Sr speciation in solution is essentially composed of Sr^{2+}, $SrCl^+$ and $SrSO_4$ species at ~85 %, ~10 % and ~4 %, respectively. The $SrCO_3$ species is present at the percent level when pH of the fluid medium is close to 7 or above. Dissolved Li speciation consists of 98 % of Li^+ species with a contribution of ~1 % for LiCl and $LiSO_4^-$ species.

	Buoyant HF (126 °C) MOM19-FERM T-1 *In situ*	Buoyant HF (80 °C) MOM19-FERM T-1 Before Inoculation	Fluid medium (80 °C, samle labeled MOM19_FERM_T1 to T18) T1 24 h	T3 72 h	T5 120 h	T7 168 h	T9 216 h	T11 264 h	T13 312 h	T15 360 h	T17 408 h	T18 432 h
Mineral saturation index (SI)												
Anhydrite ($CaSO_4$)	0.10	-0.30	-0.13	-0.17	-0.18	-0.18	-0.17	-0.19	-0.19	-0.18	-0.19	-0.16
Gypsum ($CaSO_4$:$2H_2O$)	-0.58	-0.63	-0.46	-0.50	-0.51	-0.51	-0.50	-0.52	-0.52	-0.51	-0.53	-0.49
Barite ($BaSO_4$)	-0.38	-0.02	0.06	0.35	0.31	0.26	0.37	0.54	0.53	0.42	0.32	0.34
Pyrite (FeS_2)	8.15	9.57	7.58	7.00	7.31	7.10	6.94	8.55	8.65	7.60	7.81	7.88
Pyrrhotite (FeS)	3.18	2.79	1.50	1.17	1.49	1.18	1.13	2.33	2.35	1.68	1.81	1.30
Hematite (Fe_2O_3)	3.68	1.04	0.54	0.55	0.79	0.37	0.25	0.82	0.83	0.49	0.33	-1.22
Quartz (SiO_2)	0.09	0.48	0.47	0.49	0.50	0.54	0.54	0.52	0.52	0.56	0.59	0.60
Strontianite ($SrCO_3$)	-0.12	0.04	0.20	0.27	0.67	0.07	0.56	1.38	1.17	1.33	1.45	0.20
Calcite ($CaCO_3$)	1.05	0.88	1.12	1.17	1.58	1.02	1.43	1.99	1.76	1.93	2.08	0.91
Rhodochrosite ($MnCO_3$)	-0.20	0.03	0.07	0.12	0.53	-0.02	0.27	1.23	1.00	1.06	1.08	-0.16
Species abundances (%)												
Lithium												
Li^+	98.0	98.2	98.0	98.0	98.0	98.1	98.1	98.1	98.0	97.9	97.9	97.9
LiCl	1.7	1.1	1.1	1.1	1.1	1.1	1.1	1.1	1.1	1.1	1.1	1.1
$LiSO_4^-$	0.4	0.7	0.9	0.9	0.9	0.8	0.8	0.9	0.9	1.0	1.0	1.0
LiOH	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
Strontium												
Sr^{2+}	82.3	87.4	86.5	86.5	86.3	86.8	86.8	85.9	86.1	85.3	84.9	85.9
$SrCl^+$	16.1	9.0	9.0	9.1	9.1	9.0	8.9	8.9	9.0	9.1	9.0	9.2
$SrSO_4$	1.5	3.5	4.4	4.2	4.3	4.1	4.0	4.1	4.2	4.7	4.7	4.7
$SrCO_3$	0.1	0.1	0.1	0.1	0.4	0.1	0.3	1.0	0.6	1.0	1.4	0.1
Sulfide S(-II)												
HS^-	70.8	65.5	69.9	71.7	80.3	67.8	76.9	86.9	83.8	86.6	88.8	66.5
H_2S	29.1	34.5	30.1	28.3	19.6	32.2	23.1	13.1	16.2	13.4	11.2	33.5
Sulfate S (+VI)												
$MgSO_4$	58.5	37.3	36.6	37.4	37	38	36.6	35.3	37.7	35.6	36.1	39.0
SO_4^{2-}	26.1	39.0	38.9	38.6	39.3	38.2	38.8	40.2	39	40.7	40.3	38.3
$NaSO_4^-$	9.3	17.0	17.1	16.9	16.7	16.6	17.1	17.2	16.4	17.0	17.2	16.1
$CaSO_4$	5.2	5.7	6.5	6.1	6.1	6.2	6.4	6.1	5.9	5.7	5.2	5.5
KSO_4^-	0.9	0.9	0.9	0.9	0.9	0.9	1.0	1.0	1.0	1.0	1.0	1.0

Table 2 Mineral saturation indexes and element species abundances of (i) the buoyant hydrothermal fluid (Buoyant HF) at *in situ* temperature (126 °C), (ii) the buoyant HF after reaching incubation temperature (80 °C) and gas equilibrium (MOM19_FERM_T-1), and (iii) the fluid medium 24 hours after inoculation (MOM19_FERM_T1), and then at 48 h intervals until the end of the experiment (MOM19_FERM_T3 to T18). All values were calculated using PHREEQC aqueous geochemical modeling.

3.5 Microbial diversity evolution in the bioreactor

The microbial diversity (bacterial and archaeal diversity) found throughout the experiment is presented in Fig. 5 and in Supplementary Material Table S2. These data show the relative abundance of the most representative taxa based on their sequence proportions. It is important to note that these data indicate the detected archaeal and bacterial taxa for each incubation time, thus conveying semi-quantitative variations.

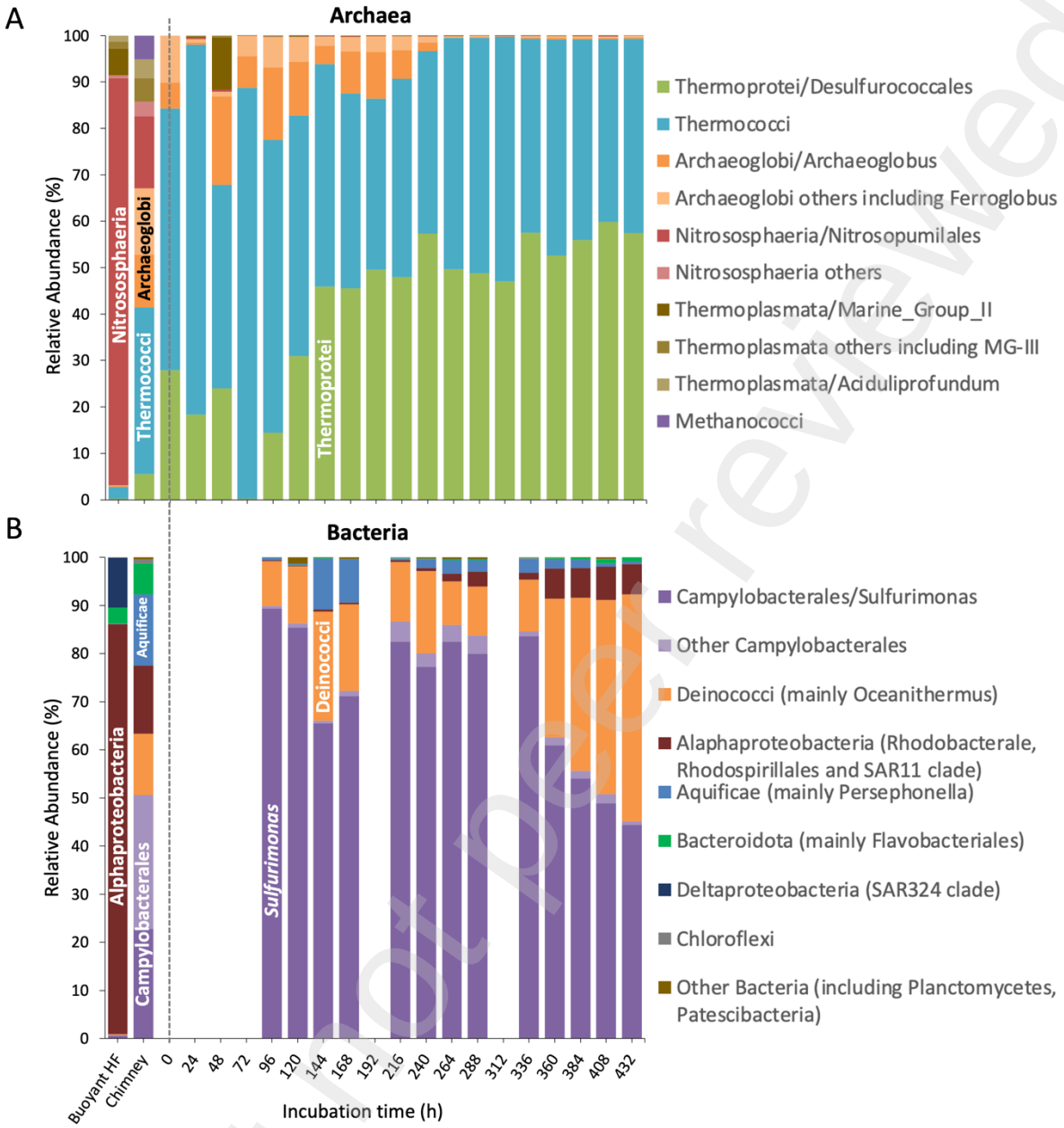

Fig. 5 Barplots presenting the relative abundance of (A) Archaea and (B) Bacteria enriched in the continuous culture over the incubation time (x-axis, hours). The microbial diversity in both the Aisics chimney sample (used as the inoculum) and the buoyant hydrothermal fluid (buoyant HF, used as the culture medium) is also presented. All taxa names are in the legend, with main taxa labeled on corresponding bars for clarity.

The cell counts of microbial communities show an overall increase throughout the experiment, reaching 2 x 10^5 cell/mL at 120 h, then varying around 6 x 10^6 cell/mL at 240 h before reaching a maximum of 8.5 x 10^6 cell/mL at 360 h, then cells concentration slightly decrease to reach 4 x 10^6 cells/mL at the end of the experiment (François, 2021). At all sampling times, archaeal sequences were detected, whereas no bacterial sequences were obtained until 96 hours, at 192 hours, and at 312 hours. This could be due to the sequencing reaction failing because of too low bacterial abundance, or the number of sequences being too low once contaminants were

removed for bacterial diversity analysis. Due to the length of 16S RNA gene sequences obtained using illumina sequencing, identification of enriched microorganisms was possible up to the order level for Archaea and up to the genus level for some Bacteria.

The archaeal diversity within the buoyant HF is mainly dominated by the class Nitrososphaeria (88%), with the *Nitrosopumulales* being the most represented order (Fig. 5a, Supplementary Material Table S2). Nitrososphaeria was also detected in the chimney sample but at significantly lower abundance (19%) compared to the buoyant HF. Within the chimney, Thermococci is the most abundant class (36%), followed by the Archaeoglobi class (26%) which includes *Archaeoglobus* and other genera such as *Ferroglobus*. Additionally, the class Thermoprotei, with the *Desulfurococcales* being its most represented order and the class Methanococci were detected in low abundance in the chimney (6% and 5%, respectively, Fig. 5a). The class Thermoplasmata, including the genus *Aciduliprofundum* and Marine Group-III (MG-III), was detected at a few percent's in both buoyant HF and chimney samples (3% and 9%, respectively), while the Thermoplasmata Marine Group-II (MG-II) was only detected in the buoyant HF (6%). Regarding bacterial diversity, the buoyant HF is largely dominated by the class Alphaproteobacteria (85%) including the orders *Rhodobacterales* and *Rhodospirillales*, and the SAR11 clade (Fig. 5b). Additionally, the class Deltaproteobacteria (SAR324 clade) and the Phylum Bacteroidota (mainly *Flavobacteriales* order) are also present in the buoyant HF, albeit at low (10%) and very low abundances (3%), respectively. By contrast, the chimney sample display higher bacterial diversity with the *Campylobacterales* order (including *Sulfurimonas* genus and other *Campylobacterales*) being dominant (51%), and classes such as Aquificae (mainly *Persephonella* genus), Deinococci (mainly *Oceanithermus* genus), Alphaproteobacteria, and Bacteroidota being detected at 15%, 13%, 14%, 6%, respectively. Throughout the 432 hours experiment, both archaeal and bacterial diversities of the fluid medium showed closer similarity to the chimney sample communities rather than to the buoyant HF ones. In the fluid medium, Thermococci and Themoprotei are the most abundant archaea classes, except at 72 h where no Thermoprotei were detected. Archaeoglobi represent between 3% and 22% of the Archaeal diversity during the first 240 hours (except at 24h) but afterward decrease to less than 1% until the end of the experiment. Regarding bacterial diversity, the *Campylobacterales* genus *Sulfurimonas* dominates until 360 h (between 61% and 89%), after which it shares prevalence with the Deinococci class (mainly *Oceanithermus*). Both Aquificae (mainly *Persephonella*) and Alphaproteobacteria classes were detected in almost all samples at very low abundance. However, Aquificae peaked at approximately 10% at 144 and 168 hours and Alphaproteobacteria reached ~6% from 360 hours to the end of the experiment (Fig. 5b, Supplementary Material Table S2).

4. Discussion

In the gas-lift bioreactor, the chemical and isotopic composition of the fluid medium results from interactions with minerals, microorganisms (archaea and bacteria) and gases (bioreactor setup conditions). In section 4.1, we will first characterize the collected materials, i.e., the Aisics chimney (used as the inoculum) and the buoyant HF (used as the culture medium), along with their microbial diversity. Once these materials are introduced into the gas-lift bioreactor, the experiment starts. In section 4.2, we will discuss how the elemental and isotopic chemical composition of the fluid medium respond to the evolution of microbial diversity and the mineral precipitation/dissolution processes.

4.1 Characterization of the collected materials

The buoyant HF was collected *in situ* between 100 and 150°C (François, 2021). Both the buoyant HF and the end-member high temperature hydrothermal fluid exhibit enrichment in dCa, dK, dSi, dFe, dMn, dLi compared to seawater (Table 1 and S1, Leleu, 2017; Millero et al., 2008). The measured chemical composition of the buoyant HF can be modeled by an adiabatic and conservative mixing between Aisics hydrothermal end-member and NADW. Major cation and anion concentrations correspond to a contribution of 35 to 44 % of hydrothermal end member and a temperature range of 110 to 141 °C, consistent with the in-situ temperature of buoyant HF collection (100 – 150 °C, Supplementary Material Fig. S1). The buoyant HF is characterized by a $^{87}Sr/^{86}Sr$ ratio of 0.707513 ± 0.000004 and a δ^7Li value of +7.2‰. While the Sr isotopic signature is fully coherent with those obtained through conservative mixing of 35 to 44% of the hydrothermal end-member with NADW, and the Li isotopic value corresponds to a 54% end-member contribution and a temperature of 170 °C (Supplementary Material Fig. S1). Contrarily to cations and anions, the end-member Sr and Li isotopic values were taken from AISICS end-member measurements between 2013 and 2015, as the 2019 values are not available (Artigue et al., 2022; Chavagnac, Leleu, et al., 2018; Leleu, 2017). Thus, this discrepancy might be attributed to a slight change in the Aisics end-member δ^7Li value in 2019, or to potential fractionation processes during mixing between end-member and seawater. Indeed, while the instrument analysis of the $^{87}Sr/^{86}Sr$ ratio eliminates mass-dependent Sr isotope fractionation occurring before or during the analysis (Andrews et al., 2016), the δ^7Li values are affected by Li isotopes fractionation notably controlled by minerals precipitation/dissolution (Hindshaw et al., 2019; Vigier et al., 2008; Wang et al., 2023).

The inoculum is mainly composed of anhydrite (at 93%) which has a retrograde solubility at temperatures below 150 °C (Bischoff & Seyfried, 1978). Sulfur bearing minerals, pyrite and chalcopyrite, are present in smaller proportions (at 1 and 3%, respectively). These minerals can provide element and energy for sulfur or sulfate reducer or oxidizer microorganisms. Indeed, archaeal and bacterial development depends not only on the physico-chemical state of the buoyant HF or inoculum but also on the bioavailable energy sources provided through chemical components releasing and/or accepting electrons.

The microbial diversity (bacteria and archaea) is summarized in Supplementary Material Table S3, along with their origins in the gas-lift bioreactor (buoyant HF or chimney), their occurrence or absence in the fluid medium, optimal growth temperature, and main known metabolism.

The most abundant archaeal class found in the buoyant HF are Nitrosophaeria (mainly *Nitrosopumulales*), which are also present in the inoculum but at a lower abundance (Fig. 5a, Supplementary Material Table S2). Nitrosophaeria were already detected at hydrothermal environments (Takai et al., 2004; Teske et al., 2021) and commonly retrieved in deep sea water. At LSHF they were identified in microbial mats (associated or not with mussel assemblages) either located at the base of the Tour Eiffel vent at LSHF (< 10°C, Crépeau et al., 2011; Rommevaux et al., 2019) or at diffuse venting at the same site (40-55°C, Astorch-Cardona et al., 2023). However, Teske et al. (2021) suggest that Nitrosophaeria thrive is inhibited by acidic, anaerobic and high temperature conditions typical of such extreme environments. Indeed, this archaeal class finds ideal conditions of development in sea water environment characterized by aerobic, neutrophilic, and mesophilic conditions (Baker et al., 2012; Supplementary Material Table S3, Könneke et al., 2005; Qin et al., 2016; Qin et al., 2017). The class Thermoplasmata was also detected in the Buoyant HF, including the mesophilic marine groups MG-II and MG-III, as well as the thermoacidophilic genus *Aciduliprofundum*

(Supplementary Material Table S2, Reysenbach et al., 2006; Santoro et al., 2019). While *Aciduliprofundum* is commonly found in deep-sea vents, MG-II and MG-III are considered to be low in abundance in the deep sea and are rarely present in hydrothermal chimneys (François, 2021; Haro-Moreno et al., 2017; Rinke et al., 2019; Zhang et al., 2015).

In the chimney sample, a few percent of *Aciduliprofundum* and MG-III, were also detected in the chimney but no MG-II. The most abundant archaeal classes in the chimney sample (inoculum) are Thermococci and Archaeoglobi, with Thermoprotei present to a lower extent (Fig. 5a and 6a). These taxa have been previously identified in deep-sea hydrothermal environments, notably at the Tour Eiffel site of the LSHF (Huber et al., 1997; Huber et al., 2006; Reysenbach et al., 2000; Rommevaux et al., 2019). These taxa have been described as anaerobes (or facultative anaerobes), (hyper)thermophilic, slightly acidophilic to alkaline (Supplementary Material Table S3 and reference therein). The Archaeoglobi class includes *Archaeoglobus* and other genera such as *Ferroglobus* (Fig. 5a, Supplementary Material Table S2). *Archaeoglobus* genus is able to reduce sulfate, sulfite, or thiosulfate compounds to H_2S using organic substrate and/or H_2 as electron donors depending on the strain (dissimilatory sulfate reduction, Barton et al., 2014; Burggraf et al., 1990; Hartzell & Reed, 2006; Liu et al., 2012; Mori et al., 2008; Offre et al., 2013). *Ferroglobus* genus is known as an Fe(II) oxidizer or an Fe(III) reducer (Hafenbradl et al., 1996; Tor & Lovley, 2001). Both the Thermoprotei class, with *Desulfurococcales* as its most represented order, and the Thermococci class, including the *Thermococcales* order, have species involved in the sulfur cycle. Some *Desulfurococcales* can growth autotrophically by oxidizing hydrogen using sulfur, nitrate, or nitrite compounds as electron acceptor, and CO_2 as a carbon source. Organotrophic growth can also occur through aerobic respiration, anaerobic sulfur respiration with organic compounds as electron donors, or fermentation of organic compounds with elemental sulfur as the electron acceptor (Huber & Stetter, 2006; Liu et al., 2012). Note that some members of *Desulfurococcales* cannot use elemental sulfur or sulfur compounds (Huber & Stetter, 2006). *Thermococcales* species are organotrophic thermophiles that can also growth through fermentation of organic compounds with or without elemental sulfur (Bertoldo & Antranikian, 2006; Liu et al., 2012). While elemental sulfur stimulates the growth of *Thermococcales*, it is not always essential, and some thrive without elemental sulfur. In the presence of elemental sulfur, it is reduced to H_2S; in its absence, H_2 is produced by proton reduction (Schut et al., 2014). Some Methanococci were detected in the chimneys samples at a low abundance (5%). Methanococci are methanogenic archaea that produce methane from H_2 and CO_2. While they are frequently detected at hydrothermal vents, they have not been previously found in Lucky Strike chimneys until now (Flores et al., 2011; Jeanthon et al., 1999; Jones et al., 1983; Jones et al., 1989; Whitman & Jeanthon, 2006).

The most abundant bacterial class identified in the buoyant HF is Alphaproteobacteria, including the orders *Rhodobacterales* and *Rhodospirillales*, and the SAR11 clade (Fig. 5B, Supplementary Material Table S2, Garrity et al., 2005). This class is also present to a lesser extent in the chimney sample, probably originating from the surrounding seawater. This class is described as ubiquitously distributed in the marine environment (Morris et al., 2002; Rappé et al., 2002) and notably found in black smoker chimneys (Voordeckers et al., 2008), microbial mats from the LSHF (Astorch-Cardona et al., 2023; Crépeau et al., 2011) and sediments (Cerqueira et al., 2017). SAR324, an uncultivated clade of Deltaproteobacteria, is only present in the buoyant HF (Fig. 5b, Fig. 6a). Known for its metabolic flexibility (Sheik et al., 2014; Swan et al., 2011; Wright et al., 1997), SAR324 clade can thrive in the full water column (Boeuf et al., 2021) and marine environments in the vicinity of hydrothermal sites (Dick et al., 2013; Dick & Tebo, 2010; François, 2021). Some Bacteroidota (mainly *Flavobacteriales* order) were detected at a few percents in both the buoyant HF and the chimney. Species of the *Flavobacteriales* order are primarily known to inhabit surface cold water environments

(Gómez-Pereira et al., 2010); however, one species was isolated from a biofilm on the surface of a black smoker chimney on the Arctic Mid-Ocean Ridge (Wissuwa et al., 2017). In the chimney sample, the bacterial abundance is dominated by *Campylobacterales* with the *Sulfurimonas* genus, and other *Campylobacterales*. Then, the rest of the bacterial diversity is shared between Alphaproteobacteria, Aquificae, Deinococci, and with a lower abundance Bacteroidota (mainly *Flavobacteriales*, Fig. 5b, Fig. 6a). All of these taxa have been previously found in deep-sea hydrothermal environments (Li et al., 2020; Miroshnichenko et al., 2003; Molari et al., 2023; Reysenbach et al., 2000; Sievert et al., 2000; Waite et al., 2017; Zeng et al., 2021; Zhang et al., 2016). *Sulfurimonas* and other *Campylobacterales* and Aquificae (mainly *Persephonella*) were specifically identified at the LSHF and at the Aisics chimney of the Tour Eiffel site (François, 2021; François et al., 2021; Mino et al., 2017; Rommevaux et al., 2019). The *Sulfurimonas* genus is described as mesophilic chemolithoautotroph bacteria that grow at an optimum pH range from 4.5 to 8.6, relying on the presence of hydrogen, elemental sulfur or thiosulfate as the sole energy source, carbon dioxide as the sole carbon source, and ammonium or nitrate as the sole nitrogen source (François et al., 2021; Hu et al., 2021; Supplementary Material Table S3, Takai et al., 2006; Zeng et al., 2021). Bacteria from the Aquificea class (mainly *Persephonella*) are thermophilic and grow in microaerophilic conditions, oxidizing hydrogen, elemental sulfur, or thiosulfate using oxygen as an electron acceptor. In anaerobic conditions, they can also perform anaerobic nitrate reduction using hydrogen as electron donor and nitrate as electron acceptor (François et al., 2021; Supplementary Material Table S3, Reysenbach et al., 2001; Zeng et al., 2021). A novel strain of the *Sulfurimonas* genus, designated as MO1340^T and exhibiting similar metabolic properties to other *Persephonella* species, was isolated from the Aisics chimney (François, 2021; François et al., 2021). Deinococci (mainly *Vulcanithermus* genus) were detected in the chimney samples (Fig. 5b). *Vulcanithermus* genus grow between pH 5.5 and 8.4 and are capable of anaerobic growth by nitrate reduction, as well as lithoheterotrophic growth with molecular hydrogen (Supplementary Material Table S3, Miroshnichenko et al., 2003).

The main archaeal and bacterial taxa identified in the collected buoyant HF and the chimney samples are illustrated in Fig. 6A, along with the mineral and chemical composition of the collected materials.

4.2 Elemental and isotopic evolution of the fluid medium: Impact of Mineral-fluid-microorganism interactions

Based on our results, we identify four time periods during which obvious and significant chemical variations are observed in trace element concentrations (except dFe) and Sr isotopic composition of the fluid medium. These four time intervals are: 1. First fluid-inoculum interactions (0 to 24 h), 2. Stabilization (24 to 168 h), 3. Drastic change (168 to 264 h), and 4. Return to stabilization (264 to 432 h, marking the end of the experiment). Note that the changes in blood bags throughout the experiment at 0 h, 5 h, 120 h, and 288 h (see section 2.3) are unrelated to the observed chemical, mineral, or microbial diversity changes. Fig. 6B is a schematic conceptual model illustrating the mineral-fluid-microorganism interactions occurring within the gas-lift bioreactor during these four-time intervals.

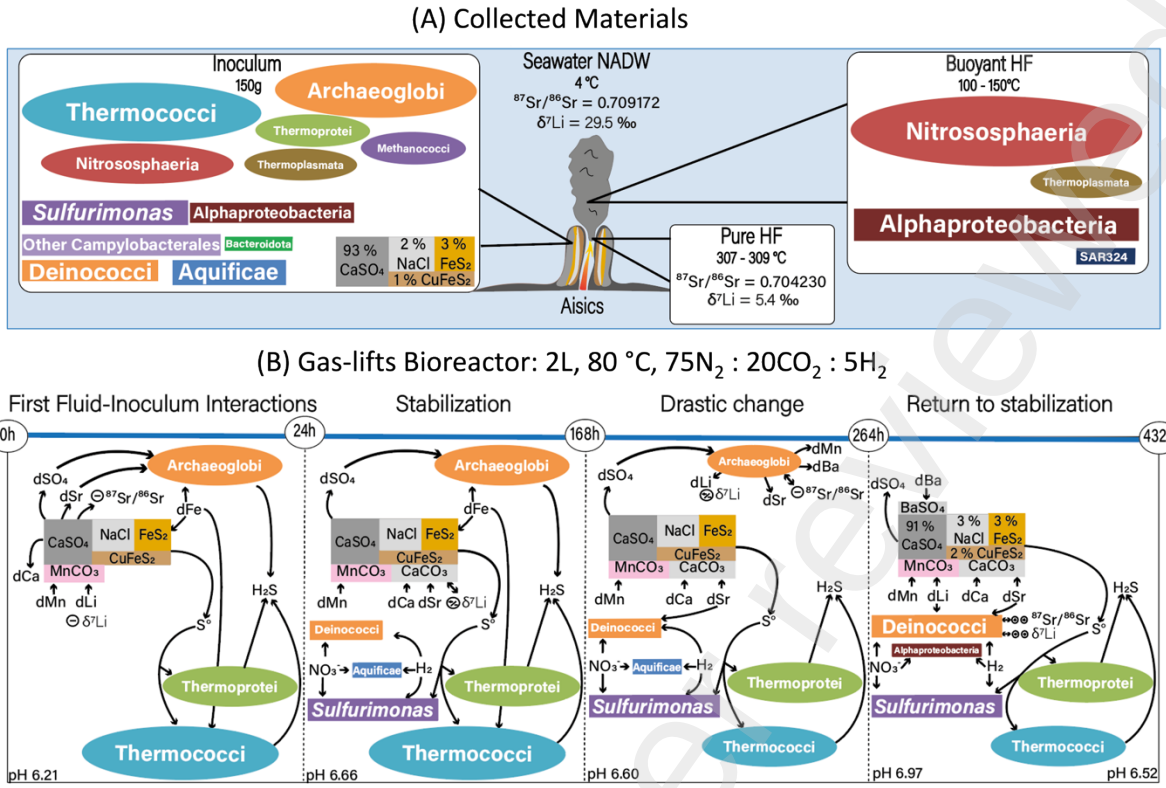

Fig. 6 Conceptual model illustrating (A) the mineral, microbial, and chemical composition of the collected materials: the high-temperature pure hydrothermal fluid (Pure HF), the buoyant hydrothermal fluid (Buoyant HF, culture medium), and the Aisics chimney (Inoculum). (B) The model showcases the impacts of mineral-fluid-microorganism interactions on the evolution of the fluid medium within the gas-lift bioreactor across four distinct time intervals: first fluid-inoculum interactions, stabilization, drastic change, and return to stabilization.

4.2.1 First fluid-inoculum interactions: first 24 h

During this time interval, (i) the pH slightly increases from 6.21 to 6.66, (ii) the chemical concentrations in the fluid medium remain either constant (dNa, dK, dSi, dCl, dBr, and dBa), increase (dCa, dSO_4, and dMg), or decrease (dFe, dMn, and dLi), (iii) the $^{87}Sr/^{86}Sr$ ratio and δ^7Li value decrease, respectively, from 0.707513 and +7.2 ‰ in the buoyant HF down to 0.707488 and +4.6 ‰ in the fluid medium at 24 h (Table 1, Fig. 4), and (iv) Nistrosophaeria class has disappeared in the gas-lift bioreactor, with sulfur-reducing archaea classes Thermoprotei and Thermococci prevailing over the sulfate-reducing archaea of the *Archaeoglobus* genus (Fig. 5a) and no bacteria were detected suggesting there are present at very low concentration (Fig. 5b).

Between the collection time of the buoyant HF and the beginning of the gaz-lift bioreactor experiment (0h), the temperature decreases from 126 °C to 80 °C, and the anhydrite saturation index shifts from being oversaturated (SI > 0) to undersaturated (SI < 0, Table 2). Consequently, anhydrite can dissolve which can explain increasing dCa and dSO_4 concentrations by up to 20 and 32%, respectively. Additionally, the increased dSO_4 concentration also change the dSr and dLi speciation to higher $SrSO_4$ and $LiSO_4^-$ concentrations (from 3.5 to 4.4% and 0.7 to 0.9%, respectively, Table 2). The dCa concentration increase of 4.02 mmol/L corresponds to the dissolution of 1.09 g of anhydrite, i.e., 0.8% of the inoculum. As the chimney sample was collected at immediate contact with high temperature hydrothermal fluid, we assume that the $^{87}Sr/^{86}Sr$ ratio of anhydrite is similar

to that of the end-member hydrothermal fluid at the same site (0.704230 ± 0.00004, Chavagnac, Leleu, et al., 2018; Leleu, 2017). Thus, its dissolution should induce a less radiogenic $^{87}Sr/^{86}Sr$ ratio in the fluid medium, as observed here.

Contrarily to anhydrite, pyrite and chalcopyrite as well as hematite remain oversaturated in the fluid medium (SI > 0; Table 2), allowing their potential precipitation and the drastic decrease of dFe concentrations (- 67% of its initial concentration; Table 1, Fig. 4c).

Other trace metals such as dMn, and dLi also decrease in concentrations, by 24 and 22%, respectively (Fig. 4c, e). We attribute these variations to mineral precipitation, as the fluid medium is oversaturated (SI > 0) regarding to carbonate minerals, notably rhodochrosite ($MnCO_3$). The δ^7Li value decreases as well from +7.2 to +4.6 ‰ in the fluid medium. Li isotope fractionation is known to be controlled by minerals precipitation/dissolution (Hindshaw et al., 2019; Vigier et al., 2008; Wang et al., 2023). Thus, even if the dLi concentration and δ^7Li can be disrupt by adsorption on oxide surfaces (especially Mn-oxides, Chan & Hein, 2007), we cannot decipher here which minerals and processes can explain a lighter δ^7Li signature. Li isotopes fractionation also depends on factors such as the temperature (Millot et al., 2010; Taylor et al., 2019), the water/rock ratio (Verney-Carron et al., 2015), the pH (Li & Liu, 2020), and CO_2 concentrations in the solutions (Ji et al., 2022). However, all these factors are controlled and maintained in the bioreactor throughout the experiment.

The physico-chemical conditions in the gas-lift bioreactor (anaerobic environment maintained at 80 °C and pH ~6.5) are drastically unfavorable for the development of Nistrosophaeria, which are the most abundant archaea in the buoyant HF. They disappear from the fluid medium after 24 hours of incubation (Fig. 5a, Fig. 6b). In contrast, these conditions are favorable for the preservation and growth of Archaeglobi, Thermococci, and Thermoprotei. It is noticeable that most enriched archaea that developed in the gas-lift bioreactor originated from the chimney sample (inoculum) rather than the buoyant HF, despite continuous feeding with fresh buoyant HF (Fig. 5a, Fig. 6b). With chemical features evidencing anhydrite dissolution, we anticipate that sulfate-reducing archaea such as *Archaeoglobus* would develop in such anaerobic conditions. However, *Archaeoglobus* and other Archaeoglobi are absent at 24 h which could be explained by their lowest growth rate compared to both Thermococci and Thermoprotei (Huber & Stetter, 2015a; Huber & Stetter, 2015b; Zillig & Reysenbach, 2015). The occurrence of Thermoprotei and Thermococci is coherent with the physico-chemical setup of the experiment. Indeed, both classes contain (hyper)thermophilic, anaerobic, and circumneutral species capable of organotrophic growth with or without elemental sulfur. Many *Thermococcales* grow better in the presence of elemental sulfur (see section 4.1, Huber & Stetter, 2006; Le Guellec et al., 2021; Liu et al., 2012). In the gas-lift bioreactor, elemental sulfur can be provided by sulfur-bearing minerals, i.e. pyrite and chalcopyrite, present in the chimney sample (Fig. 6).
The presence of genera such as *Ferroglobus* from the Archaeoglobi class and *Thermococcus* from the Thermococci class, known for their roles in iron metabolism, could also contribute to the observed significant decrease in dissolved iron (dFe). Indeed, *Thermococcus* reduces Fe(III), while *Ferroglobus* is capable of both oxidizing Fe(II) and reducing Fe(III) (Hafenbradl et al., 1996; Kashefi et al., 2002; Lim et al., 2020; Slobodkina et al., 2009; Tor & Lovley, 2001; Zeng et al., 2021).

4.2.2 Stabilization: 24 h to 144 h

During this time interval, (i) the pH remains mainly stable around 6.6, (ii) the chemical concentrations in the fluid medium remain essentially constant, with slight reductions of dCa, dSO_4, dMn, and dSr concentrations (Fig. 4a, b, and d), (iii) the $^{87}Sr/^{86}Sr$ ratios remain stable at 0.707488 ± 0.000005, contrary to the δ^7Li value of the fluid medium, which fluctuates between +4.0 and +7.0 ‰ (Table 1, Fig. 4d and 4e), (iv) Thermococci are the most abundant archaea and Thermoprotei and Archaeoglobi are present, (v) only bacteria from the genus *Sulfurimonas* and Deinococci class are significantly abundant, with *Sulfurimonas* being the most prevalent (Fig. 6b).

The observed covariations of dCa, dSO_4 and dSr concentrations would suggest anhydrite ($CaSO_4$) precipitation. However, the fluid medium is undersaturated regarding this mineral (SI < 0), precluding its precipitation (Table 2). Concomitant dCa and dSr concentrations decrease could be related to calcite precipitation (SI > 0, Table 2). In another gas-lift bioreactor study, Callac et al (2015) also observed a concomitant dCa and dSr concentrations evolution attributed to calcite - fluid medium interaction. The presence of calcite within the interstice of natural anhydrite chimney walls was already identified by Pagé et al (2008). Calcite precipitation induces Li isotope fractionation with a factor ranging from -8 to +2 ‰ depending on pH values and calcite growth rate (Füger et al., 2019; Füger et al., 2022; Marriott, Henderson, Belshaw, et al., 2004; Marriott, Henderson, Crompton, et al., 2004; Seyedali et al., 2021). This large range of fractionation factor could explain the δ^7Li variability between +4.0 and +7.0 ‰. Moreover, the presence of *Archaeoglobus* genus of the Archaeoglobi class is coherent with calcite precipitation as they contribute to the total carbon mineralization process in marine sediments as sulfate reducing prokaryotes (Barton et al., 2014). Moreover, *Archaeoglobus* use dSO_4 in their metabolism which is coherent with the dSO_4 decrease observed here. Another carbonate mineral that can precipitate here is Rhodochrosite (SI > 0), whereby its precipitation could explain the observed decrease in dMn concentrations (Fig. 4C).

Data on bacterial diversity in the fluid medium are available starting 96 hours after the start of the experiment (Fig. 5b). Among the bacteria identified in the buoyant HF and in the chimney sample (see section 4.1), only, *Sulfurimonas* genus from the *Campylobacterales* order and Deinococci class, are significantly abundant with *Sulfurimonas* largely dominating the diversity (Fig. 5b, Supplementary Material Table S2). As for the archaeal diversity, the most enriched bacteria thriving in the gas-lift bioreactor originates from the chimney sample rather than the buoyant HF. Mesophilic bacteria from *Sulfurimonas* genus was not expected to grow at such temperature (~ 80 °C) in the bioreactor (Supplementary Material Table S3). Nonetheless, *Sulfurimonas* genus was identified in various mesophilic and thermophilic hydrothermal habitats, including plumes, sediments, chimneys, and diffuse-flow fluids (Akerman et al., 2013; Hu et al., 2021; Inagaki, 2003; Li et al., 2020; Mino et al., 2017; Nakagawa et al., 2005). The presence of species at temperatures significatively above their known optimal growth temperature, such as *Sulfurimonas*, has previously been described in bioreactor enrichment culture experiments (Callac et al., 2015; Postec, Urios, et al., 2005). Aquificae, mainly *Persephonella* species, originating from the chimney sample, are initially detected at extremely low abundance (< 1%) in the fluid medium. However, their abundance increases to approximately 10% by 144 hours, comparable to that of the chimney sample (Supplementary Material Table S2, Fig. 5b). This is consistent with the known growth conditions of *Persephonella* species, which are suitable for our culture system (François et al., 2021). Here, the gas-lift bioreactor physico-chemical conditions (gas-flux composition and pyrite and chalcopyrite in the inoculum) provides all the essential growth prerequisites for Deinococci, and Aquificae.

4.2.3 Drastic change: 168h to 264h

During this time interval, drastic changes are observed regarding trace element concentrations and their isotopes. Indeed, (i) the pH slightly increasing from 6.60 to 6.97; (ii) major element concentrations remain overall constant; (iii) all trace element concentrations (apart from dFe), drastically increase by a factor of two; (iv) the $^{87}Sr/^{86}Sr$ ratio drastically decreases reaching a minimum value of 0.707337 at 264 h, while δ^7Li values continue to fluctuate between +4.0 and 7.3‰, (v) Archaeoglobi class progressively disappears; and (vi) the bacterial diversity (mainly *Sulfurimonas* and Deinococci) remains overall unchanged (Fig. 6b).

The mineral dissolution/precipitation does not explain the drastic trace elements increase. Indeed, the observed drastic increase of dMn, dLi, and dBa concentrations cannot be due to rhodochrosite and barite dissolution as the fluid medium is oversaturated regarding to both of them. The observed increase of dSr concentrations cannot be due to anhydrite dissolution even if anhydrite could dissolve (SI < 0), and supply dSr in the fluid medium. The increase of dSr concentrations by up to 74 µmol/L would imply the dissolution of 9 g of anhydrite (assuming a Sr concentration ~1500 ppm in anhydrite (Humphris & Bach, 2005), leading to a dCa and dSO4 increase by up to 32 mmol/L, which is not observed here. Therefore, an effect of microbial diversity has to be addressed.

The bacterial diversity (mainly *Sulfurimonas* and Deinococci) and their relative abundance (66-82 % and 9-23%, respectively), remain overall constant. However, Archaeoglobi abundance seems to decrease over time, while sulfur-reducing archaea classes, i.e. Thermoprotei and Thermococci, are much more abundant, reaching both a relative abundance of 50% at 264 h. We observed that the progressive Archaeglobi class low abundance is concomitant with an increase in dSr (as well as dMn, dLi, and dBa, Fig. 4) and less radiogenic $^{87}Sr/^{86}Sr$ ratios from 0.707483 down to 0.707337 in the fluid medium. We suggest that Archaeoglobi has the potential to store many trace metals via its metabolic pathway. Indeed, to sustain their metabolic demands, bacterial and archaeal cells have to ensure the supply of the right metals to the right proteins (Waldron & Robinson, 2009). This process, known as metal homeostasis, is based on the involvement of specific genes capable of sensing, transporting and storing any metals inside and between cells (Chandrangsu et al., 2017; Waldron & Robinson, 2009). Thus, it is possible that Archaeoglobi have stored dSr in their cells with other trace metals (dMn, dLi, and dBa), and progressively release them into the fluid medium when they vanish. Such process could lead to the observed less radiogenic $^{87}Sr/^{86}Sr$ ratios from 0.707483 down to 0.707337. The δ^7Li values fluctuates between +4.0 and +7.3 ‰ without any specific trend according to time. Poet et al (2023) showed that membrane transporters and channels transport ^6Li faster than ^7Li. Such process associated to Archaeoglobi should result in lower δ^7Li one when they progressively disappear, which is not clearly evidenced here. Further studies on the link between Archaeoglobi class, trace metal storage and associated isotopic fractionation are therefore needed.

4.2.4 Return to stabilization: 264 h to 432 h

We observed that (i) the pH remains stable at around 7 (ii) the chemical concentrations in the fluid medium either decrease (dCa, dMn, dBa, dSr , dLi), or increase (dSO_4, and dMg), (iii) the $^{87}Sr/^{86}Sr$ ratio and δ^7Li values increases reaching a maximum of 0.707794 at 432 h and +10.3 ‰ at 408 h, respectively, (iv) the abundance of archaea classes remain stable and fully dominated by sulfur-reducing ones (Thermoprotei and Thermococci, each at 50%), (iv) the abundance of the Deinococci class increases, and significant abundance of Alphaproteobacteria is observed, while the relative abundance of *Sulfurimonas* decreases (Fig. 6b).

Contrarily to the previous time intervals, the observed decrease of dCa (-13%), dSr (-26%), dMn (-55%), dLi (-51%), and dBa (-45%) concentrations is coherent with calcite, rhodochrosite and barite possible precipitation, as the fluid medium is still oversaturated regarding these minerals (SI > 0). The increase of dSO_4 (+20%) could be attributed to anhydrite dissolution (SI < 0), and no more consumption by the *Archaeoglobus* genus. However, assuming that $^{87}Sr/^{86}Sr$ ratio of anhydrite is similar to that of the end-member (see section 4.2.1), its dissolution should deliver less radiogenic $^{87}Sr/^{86}Sr$ ratio to the fluid medium which is not consistent with the drastic increase of $^{87}Sr/^{86}Sr$ ratio from 0.707337 to 0.707794 (Fig. 4D).

The microbial influence needs to be address. The relative abundance of Deinococci increases, reaching a level of abundance similar to that of *Sulfurimonas*, and is correlated with a decrease in dSr concentrations and an increase in $^{87}Sr/^{86}Sr$ ratios. We suggest that Deinococci trap dSr through either biosorption or bioaccumulation, and can potentially fractionate the Sr isotope distribution via its metabolic pathway, leading to progressive radiogenic ^{87}Sr in the fluid medium (Fig. 6b). We observe a similar behavior between Sr and Li elemental and isotopic composition (Fig. 4d and 4e). The δ^7Li values show a significant and progressive increase over time, i.e. from +4.6 to +10.3 ‰ (apart at 432 h). This is concomitant with dLi decrease and Deinococci class taking over the *Sulfurimonas* genus (Fig. 5b). dLi plays a role in many physiological and biochemical functions of many living organisms (Jakobsson et al., 2017). Also, dLi could be exchanged through inward flux of H+ in a regulation of intracellular toxic metal process (Swartz et al., 2005). Moreover, Poet et al (2023) showed that Li incorporation through membrane ion channels and Na^+-Li^+/H+ exchangers fractionate Li isotopes transporting ^6Li faster than ^7Li. Such fractionation process should lead to a heavier δ^7Li signature in the fluid which is observed here. Therefore, we suggest that Deinococci trap dLi leading to a decrease of dLi concentrations and a heavier δ^7Li signature in the fluid medium.

5 Conclusion

The chemical and isotopic composition of the fluid medium in the gas-lift bioreactor reflects mineral-fluid-microorganism interactions (Fig. 6). Since the first interaction between buoyant hydrothermal fluid and the sulfate-based chimney (inoculum, 93% anhydrite), the microorganism diversity present in both collected material is depleted in favor of that of the inoculum. Throughout the experiment, Archaeoglobi, Thermoprotei, Thermococci are the main archaea present in the bioreactor, while *Sulfurimonas* and Deinococci are the main bacteria. Over the course of the 18-days experiment, the control exerted by microorganisms and minerals over the chemical composition of the fluid medium evolves. This evolution delineates four distinct temporal phases: first fluid-inoculum interactions (0 to 24 h), stabilization (24 to 168 h), drastic change (168 to 264 h), and return to stabilization (264 to 432 h). During the first fluid-inoculum interactions, sulfur-reducing archaea classes (Thermoprotei and Thermococci) prevail over sulfate-reducing ones (Archaeoglobi). Mineral precipitation (sulfide minerals, rhodochrosite) and dissolution (anhydrite) seem to control the elemental and isotopic chemical composition of the fluid medium. During the stabilization (24 to 168 h), bioreactor conditions, microbial diversity, and most chemical concentrations as well as the Sr isotopic signature of the fluid remained stable. The slight decrease of dCa, dSO_4, and dSr concentrations can be explained by mineral (calcite) precipitation and are coherent with the sulfate-reducing metabolism of Archaeoglobi. During the drastic change, while Archaeoglobi were less abundant and the microbial diversity is dominated by sulfur-reducing microorganisms, major element concentration remains overall constant, trace element concentrations drastically increase (except dFe), and Sr isotopic ratio drastically decreases. We suggest that Archaeoglobi

would have released previously stored trace metals (dSr, dMn, dLi, and dBa) into the fluid medium leading to less radiogenic $^{87}Sr/^{86}Sr$ ratios. During the return to stabilization interval, the elementary chemical composition of the fluid is controlled by the carbonate precipitation and anhydrite dissolution. Increases in the $^{87}Sr/^{86}Sr$ ratio and $\delta^{7}Li$ isotopic signature are concomitant with the increased abundance of Deinococci, reaching a similar level of abundance as *Sulfurimonas*. We suggest that Deinococci likely absorb and accumulate dSr and dLi, leading to the gradual increase in radiogenic $^{87}Sr/^{86}Sr$ ratios and a heavier $\delta^{7}Li$ signature in the fluid medium.

Overall, the evolution of major element concentrations in the fluid medium is controlled by mineral-fluid interactions, the trace element concentrations are controlled both by minerals and microorganisms, and the Sr and Li isotopic variations seem to be mainly controlled by microorganisms (Fig. 6). This underscores the necessity for cautious interpretation when utilizing Sr isotopes as tracers for paleo hydrothermal records, as they might be affected by biological isotopic fractionation. Note that Sr isotopic variation effectively highlight mineral-fluid-microorganisms interactions, whereas the consistent fluctuations in $\delta^{7}Li$ signatures throughout the experiment complicate its use as a reliable tracer. Further studies focusing on measuring isotopic fractionation factors in these three compartments are needed to enhance our understanding of mineral-fluid-microorganism interactions.

Author Contribution Statement:

AG, DF and FL did the bioreactor experiment and sampling onboard, and the microbial diversity study onshore. VC, and CD performed the chemical analysis onboard. LA performed the elemental and isotopic analyses onshore, the PHREEQC geochemical modeling in collaboration with CD, integrated all data, and performed the visualization/data presentation work with inputs from the other authors. LA, VC, and CD conducted the interpretation work and drafted the manuscript with LA leading the writing and with significant contributions from AG and DF.

Funding

The Institut Carnot ISIFoR through the ADERA provided funding for LA, and VC. The CNRS/INSU TELLUS 2021 call provided funding for the AMINO project. The EU project Emso Eric (http://www.emso-eu.org/) provided funding for the maintenance of the EMSO-Azores observatory.

Data Availability Statement

The original data presented in the study are included in the main article and in the Supplementary Material.

Acknowledgments

We thank the *R.V. Pourquoi Pas?* and Genavir crew for their tremendous work during the MoMARsat'19. We are grateful to the chemistry facility of the Géoscience Environment Toulouse laboratory (GET), the clean room facility of the Laboratoire d'Etudes en Geophysique et Océanographie Spatiales (LEGOS), and the mass spectrometry facility of the PANGEE platform at the Observatoire Midi-Pyrénées (OMP). We thank Céline Rommevaux for her assistance on board and her feedback on the first draft of this paper.

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