Evidence for microbial iron reduction in the methanogenic sediments of the oligotrophic SE Mediterranean continental shelf

Hanni Vigderovich1, Lewen Liang2, Barak Herut3, Fengping Wang2, Eyal Wurgaft1,4, Maxim Rubin-Blum3 and Orit Sivan1

1 The Department of Geological and Environmental Sciences, Ben-Gurion University of the Negev, Beer-Sheva, 8410501, Israel.
2 School of Life Sciences and Biotechnology, Shanghai JiaoTong University, Shanghai, 200240, P.R.China.
3 Israel Oceanographic and Limnological Research, Haifa, 31080, Israel.
4 Currently: The Department of Marine Chemistry and Biochemistry, Woods-Hole Oceanographic Institution, Woods-Hole, USA

Correspondence to: Orit Sivan (oritsi@bgu.ac.il)

Abstract. Dissimilatory iron reduction is probably one of the earliest metabolisms, which still participates in important biogeochemical cycles such as carbon and sulfur. Traditionally, this process is thought to be limited to the shallow part of the sediment column, as one of the energetically favorable anaerobic microbial respiration cascade, usually coupled to the oxidation of organic matter. However, in the last decade iron reduction has been observed in the methanogenic depth in many aquatic sediments, suggesting a link between the iron and the methane cycles. Yet, the mechanistic nature of this link has yet to be established, and has not been studied in oligotrophic shallow marine sediments. In this study we present first geochemical and molecular evidences for microbial iron reduction in the methanogenic depth of the oligotrophic Southern Eastern (SE) Mediterranean continental shelf. Geochemical pore-water profiles indicate iron reduction in two zones, the traditional zone in the upper part of the sediment cores and a deeper second zone located in the enhanced methane concentration layer. Results from a slurry incubation experiment indicate that the iron reduction is microbial. The Geochemical data, Spearman correlation between microbial abundance and iron concentration, as well as the qPCR analysis of the mcrA gene point to several potential microorganisms that could be involved in this iron reduction via three potential pathways: H2/organic matter oxidation, an active sulfur cycle or iron driven anaerobic oxidation of methane.

1 Introduction

Iron (Fe) is the fourth most abundant element in the Earth's crust. It appears as elemental Fe, Fe(II) and Fe(III), and has an important geobiological role in natural systems (Roden, 2006). Dissimilatory microbial iron reduction is likely one of the first evolutionary metabolisms, and plays a key role in the reductive dissolution of Fe(III) minerals in the natural environment (Weber et al., 2006), in the mineralization of organic matter in freshwater sediments (Roden and Wetzel, 2002), and as a redox wheel that drives the biogeochemical cycles of carbon, nitrogen, sulfur and phosphorous (Li et al., 2012). Dissimilatory iron reduction is part of the anaerobic respiration cascade, in which different organic substrates are used for energy by microorganisms and oxidized to dissolved inorganic carbon (DIC). This is accomplished by reduction of electron acceptors, other than oxygen, according to their availability and...
energy yield. Denitrification is the first respiratory process in anoxic sediments, followed by manganese and iron reduction and then sulfate reduction. Methane (CH$_4$) production (methanogenesis) by archaeal methanogens is traditionally considered to be the terminal process of microbial organic matter mineralization in anoxic environments, after the other electron acceptors have been exhausted (Froelich et al., 1979). When outward diffusing methane meets an electron acceptor it can be consumed by microbial oxidation (methanotrophy). In anoxic marine sediments anaerobic oxidation of methane (AOM) is coupled mainly to sulfate reduction (Hoehler et al., 1994). This process was found to consume up to 90% of the methane that diffuses upwards to the sulfate methane transition zone (SMTZ) (Valentine, 2002).

The classical process of dissimilatory iron reduction is coupled to the oxidation of organic matter (organoclastic iron oxidation, Eq. 1) (Lovley, 1991; Lovley et al., 1996). However, iron reduction can be coupled to other processes as well, such as hydrogen (H$_2$) oxidation (hydrogenotrophic iron reduction) (Eq. 1). Besides H$_2$ oxidation, Fe(III) can be reduced microbially (and also abiotically) by pyrite oxidation (Eq. 2) (Bottrell et al., 2000), leading to S intermediates, and followed by their disproportionation to sulfate and sulfide via a "cryptic" sulfur cycle (e.g. Holmkvist et al., 2011).

$$Fe^{3+} + \text{organic matter}/H_2/humic acids \rightarrow Fe^{2+} + HCO_3^-/CO_2$$  \hspace{1cm} (1)

$$FeS_2 + 14Fe^{3+} + 8H_2O \rightarrow 15Fe^{2+} + 2SO_4^{2-} + 16H^+$$  \hspace{1cm} (2)

Another, recently discovered, pathway of iron reduction is by AOM (Eq. 3).

$$CH_4 + 8Fe(OH)_3 + 15H^+ \rightarrow HCO_3^- + 8Fe^{2+} + 21H_2O$$  \hspace{1cm} (3)

This process in marine sediments was evident through incubations of marine seeps (Beal et al., 2009; Sivan et al., 2014). It was suggested through the modeling of geochemical profiles in deep sea sediments (Sivan et al., 2007; Riedinger et al., 2014) and in brackish coastal sediments (Slomp et al., 2013; Segarra et al., 2013; Egger et al., 2015; Egger et al., 2016; Rooze et al., 2016). In freshwater, it was suggested to occur in lakes (Crowe et al. 2011; Norði et al., 2013), and was shown in enriched, denitrifying cultures from sewage, where it was performed by methanogens (Ettwig et al., 2016). Iron-coupled AOM in natural lake sediments was indicated using isotope pore-water depth profiles (Sivan et al., 2011), rate modeling based on these profiles (Adler et al., 2011), microbial profiles (Bar-Or et al., 2015), and directly by a set of sediment slurry incubation experiments using several methods (Bar-Or et al. 2017). The few microbial studies about iron-coupled AOM (mainly in cultures) showed the involvement of methanogenic/methanotrophic archaea (Scheller et al., 2016; Ettwig et al., 2016; Cai et al., 2018; Yan et al., 2018; Rotaru and Thamdrup, 2016) or cooperation between methanotrophic archaea and iron reducing bacteria (Bar-Or et al., 2017).

Whereas Fe(II) is highly soluble, Fe(III) appears as low solubility minerals, and is the most abundant species of iron under natural conditions close to neutral pH. This makes iron usage a challenge to microorganisms, which need to respire low-solubility iron oxide minerals, thus rendering many of iron-oxide minerals effectively unavailable for reduction and leading to the dominance of sulfate reducing bacteria beyond a certain depth.
Therefore, the observation of iron reduction below its traditional depth, in the methanogenic zone, where iron-oxides are assumed to be inactive, is surprising. Moreover, this reduction is occasionally accompanied by a depletion in methane concentrations, suggesting a possible link between the iron and the methane cycles. The coupling can be through a competition between methanogens and iron reducing bacteria, methanogens switching from methanogenesis to iron reduction metabolism, and/or iron coupled AOM, as mentioned above. Previous observations in other environments demonstrated the inhibition of methanogenesis under iron-reducing conditions due to competition between methanogens and iron-reducing bacteria for the common acetate and hydrogen substrates (Conrad, 1999; Lovley and Phillips, 1986; Roden and Wetzel, 1996; Roden, 2003). Different methanogens can also utilize iron directly, by reducing Fe(III). This was shown in pure cultures with the amorphous Fe(III) oxyhydroxide (Bond and Lovley., 2002), in pure cultures close to natural sedimentary conditions (Sivan et al., 2016), in natural lake sediments with different iron oxides (i.e. amorphous iron, goethite, hematite and magnetite) (Bar-or et al., 2017), and in iron-rich clays (Liu et al., 2011; Zhang et al., 2012; Zhang et al., 2013).

Despite the above studies, the nature of the link between the iron and the methane cycles in marine methanogenic zone, which reactivates iron oxides and making them available for reduction, has not been determined yet. Furthermore, microbial iron reduction in methanogenic zones has not been shown in oligotrophic shallow marine environments. In this study we report observations of microbial iron reduction in the methanogenic depth in marine sediments of the oligotrophic SE Mediterranean continental shelf. We show both geochemical pore-water profiles and microbial investigation at three different stations combined with a basic incubation experiment with slurry from the methanogenic zone. These profiles and the incubation, including the related microorganisms, are discussed in terms of the possible links between iron and methane cycles.

2 Methods

2.1 Study site

The Levantine Basin of the SE Mediterranean Sea is one of the most oligotrophic nutrient-poor marine environment in the world (Kress and Herut, 2001), including the Israeli continental shelf (Herut et al., 2000). The continental shelf narrows from south to north and is built mainly of Pliocene-Quaternary Nile-derived sediments, whose rate of sedimentation decreases with increasing distance from the Nile Delta and from the shoreline (Nir, 1984; Sandler and Herut, 2000). While the highest levels of total organic carbon (TOC) (1 – 2%) in sediments were found in the Western Mediterranean Basin and offshore the Nile River delta, the central and eastern deep water regions of the Levantine Basin have relatively low TOC levels (0 – 0.5%) (Astrahan et al., 2017). Along the Egyptian coast, maximal contents of TOC in surface sediments on the shelf is up to 1.5% (Aly Salem et al., 2013), while in the Israeli shelf sediments (< 100 m depth) the TOC levels vary between < 0.1 – 1% (Almogi-Labin et al., 2009). The discovery of a 'gas front' from seismic profiles within the sediments of the continental shelf of Israel (Schattner et al., 2012), led to the findings of biogenic methane formation at some locations in shallow sediments (Sela-Adler et al., 2015). The bottom seawater across the shelf is well oxygenated therefore sulfate concentration in the water-sediment interface is ~30 mmol L⁻¹ (Sela-Adler et al., 2015).
2.2 Sampling

Seven sediment cores (~5 – 6 m long) were collected using a Benthos 2175 piston corer, from the undisturbed seafloor sediments of the SE Mediterranean continental shelf of Israel at water depths of 81 – 88 m from three stations; SG-1 (32°57.82’ N 34°55.30’ E), PC-3 (32°55.30’ N 34°54.14’ E) and PC-5 (32°55.47’ N 34°55.01’ E). The cores were sampled by the R.V. Shikmona between 2013 to 2017, and by the R.V. Bat-Galim on January 2017. The sediment cores were sliced on board every 25 – 35 cm within minutes upon retrieval from the seafloor. These stations were investigated previously with other focuses, such as the sulfate reduction in the SMTZ (Antler et al., 2015; Wurgaft et al., unpublished), and methanogenesis characteristics (Sela-Adler et al., 2015).

From each interval, a 2.5 mL sediment sample was collected and inserted into an anaerobic 10 mL glass bottle filled with 5 mL NaOH 1.5 N for headspace measurements of methane concentration (after Nusslein et al, 2003). In addition, another 2.5 mL sediment sample was taken from each segment of the cores and transferred into a 20 mL glass bottle filled with NaCl saturated solution for H\(_2\) concentrations measurements. Sediment samples from each segment of the cores were centrifuged on board if possible or in the lab within a day by Sorval centrifuge at 9500 RPM under 4 °C and Ar atmosphere in order to extract pore-water for chemical analysis. The supernatant was filtered (0.22 µm) and analyzed for Fe(II), sulfate, sulfide and the isotope composition of the DIC (\(\delta^{13}C_{DIC}\)). After the pore-water extraction, the sediment was analyzed for the content of the different iron oxides. In addition, sediment sub-sample from each segment of the January 2017 core from SG-1 station was kept in -20 °C for molecular analysis. Due to high water content in the upper part of the sediments, the top 20 cm of the piston cores pore-water measurements might be affected by sediment movement and mixing; to avoid this, two ~30 cm cores were collected during the September 2015 and January 2017 cruises via a 0.0625 m\(^2\) box corer (Ocean Instruments BX 700 Al). The cores were stored at 4 °C and cut in the lab less than 24 hours after their collection. Sediment and pore-water samples were measured for CH\(_4\), Fe(II), sulfate and \(\delta^{13}C_{DIC}\) measurements.

2.3 Slurry incubation experiment

The experimental set-up consisted of 11 bottles with sediment from the methanogenic zone, 260 cm below sea floor level from SG-1 station, where iron reduction was apparent in the pore-water profiles. The sediment from the designated depth was homogenized in an anaerobic bag under N\(_2\) atmosphere. It was then transferred under anaerobic conditions to a 250 mL glass bottle with the addition of synthetic sea water without sulfate to reach 1:1 sediment – water slurry ratio for 3 months incubation period. Then the slurry was sub-divided anaerobically to the 60 mL experiment bottles, and synthetic sea water was added for final sediment – water ratio of 1:3. The bottles were sealed with a crimp cap and were flushed with N\(_2\) for 5 minutes, shaken vigorously and flushed again, (repeated 3 times). The experiment bottles were amended with iron oxides (hematite (Fe\(_2\)O\(_3\)) or magnetite (Fe\(_3\)O\(_4\)) with final concentration of 10 mmol L\(^{-1}\) with/without hydrogen (H\(_2\)) (to the final concentration of ~4% of the Head space volume). The experiment bottles were sampled several times for dissolved Fe(II) concentrations during the 14 day experiment period.
2.4 Analytical methods

2.4.1 Pore-water analyses
Methane concentrations were analyzed by Focus Gas – Chromatograph (GC; Thermo) equipped with FID detector with detection limit of 50 μmol L⁻¹. H₂ concentrations were analyzed in a Reducing Compound Photometer Gas-Chromatograph (RCP-GC; Peak Laboratories). Dissolved Fe(II) concentrations were measured using the ferrozine method (Stookey, 1970) by a spectrophotometer at 562 nm wavelength with detection limit of 1 μmol L⁻¹. Sulfide was measured using the Cline (1969) method by a spectrophotometer at 665 nm wavelength with detection limit of 1 μmol L⁻¹. Sulfate concentrations were measured in an inductive coupled plasma atomic emission spectrometer (ICP-AES), Perkin Elmer Optima 3300, with an analytical error of ±1% (average deviations from repeated measurements of a seawater standard). The δ¹³C values were measured on a DeltaV Advantage Thermo® isotope-ratio mass-spectrometer (IRMS) at a precision of ±0.1 ‰. Results are reported versus VPDB standard. Several pore-water profiles were performed during the study, and all of them are presented (and not their average). For each profile, the error bar is that of the average deviation of the mean of the duplicates, in cases where they were taken, otherwise it is that of the analytical error (if larger than the symbol).

2.4.2 Sediment analysis
Reactive Fe(III) in the sediments was measured according to Poulton and Canfield (2005) definition and sequential extraction procedure. The different reactive iron oxides were separated to (1) carbonate-associated Fe; (2) easily reducible oxides; (3) reducible oxides and (4) magnetite. At the end of each extraction stage, the extractant was transferred to a 15 mL falcon tube with 0.1 mL ascorbic acid and 0.1 mL ferrozine solution to reduce all the Fe(III) to Fe(II) and fix it, then it was measured spectrophotometrically. The results are presented as "reactive Fe(III)", which was the sum of the easily reducible oxides, reducible oxides and magnetite.

2.4.3 Quantitative PCR and 16S rRNA gene V4 amplicon pyrosequencing
DNA was extracted from the sediment core of station SG-1 from January 2017 using PowerSoil DNA Kit (MoBio Laboratories, Inc., Carlsbad, CA, USA) following manufacturer’s instructions. Copy numbers of selected genes were estimated with quantitative PCR (qPCR) as described previously (Niu et al., 2017) using specific primers: Uni519f/Arc908R and bac341f/519r for archaeal and bacterial 16S rRNA genes, respectively, and mlas/mcrA-rev for the mcrA gene, which encodes the α-subunit of methyl-coenzyme M reductase. The amplification efficiency was 94.5%, 106.3% and 92.4% for the archaeal 16S rRNA, bacterial 16S rRNA and the mcrA gene, respectively (the respective R² of the standard curve was 0.998, 0.998 and 0.995). The V4 regions of bacterial and archaean 16S rRNA genes were amplified using barcoded 515FB/806RB primers (Walters et al., 2015) and Arch519/Arch806 primers (Song et al., 2013), respectively. PCR mixture contained 6 – 10 ng total DNA, 5 μL 10x Ex Taq buffer, 4 μL 2.5 mmol L⁻¹ dNTP mix, 1 μL of each primer, 0.25 μL Ex Taq polymerase (Ex-Taq; TaKaRa, Dalian, China) and 5 μL bovine serum albumin (25 mg mL⁻¹) in a total volume of 50 μL. DNA was sequenced as 2x150 bp reads using Illumina MiSeq platform (Illumina, USA). Sequence quality assessments, chimera detection and down-stream
phylogenetic analyses were conducted in QIIME (Caporaso et al., 2010). Taxonomical assignments for each OTU were performed in QIIME using the BLAST method and the SILVA128 reference database. 24056 to 132042 high quality sequences were obtained per sample, with the proportion of high-quality sequence versus total sequence between 81.97 – 99.89%. Spearman correlation was performed using the online calculator (http://www.sthda.com/english/rsthdacorrelation.php) to test the relevance of microbial abundance and communities with Fe(II) concentration along the depth of the sediment core from 185 cm to the bottom 575 cm, which is the methanogenic zone of the sediment core according to the geochemical profile (see the results below).

3 Results

3.1 Geochemical profiles

Geochemical pore-water profiles of several sediment cores from the three stations (SG-1, PC-3 and PC-5) were performed in order to test the possibility of the iron reduction process in the methanogenic zone of the SE Mediterranean continental shelf and its potential sources. Pore-water analyses of sulfate concentrations show complete depletion at approximately 150 cm depth in all cores at station SG-1 (Fig. 1). Sulfide concentrations were below the detection limit in all cores and therefore are not presented. Methane profiles show an increase in concentration immediately after the consumption of sulfate. The maximal methane concentration was the saturation level (Sela-Adler et al., 2015) of about 10 nmol L\(^{-1}\) at ~140 cm depth in June 2015 (station SG-1), probably due to intensive methane production at the exact location of the core collected at that time. The other methane depth-profiles show high concentrations of methane of approximately 2 nmol L\(^{-1}\) all the way to the bottom of the cores (~600 cm). Detected dissolved Fe(II) concentrations were found in the traditional iron reduction zone in the upper part of the cores (between 30 – 90 cm depth). However, a second peak was found in the deeper part of the sediment, at the methanogenic zone (below 180 cm depth). Maximum dissolved Fe(II) concentrations reached 84 \(\mu\)mol L\(^{-1}\) in the traditional iron reduction zone of the sediment cores, and 65 \(\mu\)mol L\(^{-1}\) in the methanogenic zone (Fig. 1). It should be noted that iron species are highly sensitive to environmental changes such as shifts in local pH, the different types of electron shuttles, and organic compounds that are present in the surroundings. These changes affect the net dissolved Fe(II) observed; consequently the dissolved Fe(II) results show variability between the cores that were extracted and analyzed from the same station. The \(\delta^{13}C_{\text{DIC}}\) values were the lowest (-35 ‰) at the SMTZ depth, as expected from the intensive sulfate-coupled AOM process there, which uses the isotopically light carbon of the methane as a carbon source with only small fractionation. The \(\delta^{13}C_{\text{DIC}}\) values were the highest in the methanogenic zone, as expected (the carbon source for the methane comes from the CO\(_2\), leaving the residual DIC heavier by about 60 ‰ (Whiticar, 1999)). Fitting the intensive methane profile from June 2015, the \(\delta^{13}C_{\text{DIC}}\) showed the most dramatic decrease and increase that date as well. The SMTZ was also the shallowest in this core because of the intensive methane oxidation, thus the traditional iron reduction is missing in the sampled pore-water. H\(_2\) concentrations decreased to a minimum peak of 17 nmol L\(^{-1}\) at 155 cm depth, and then increased to a maximum of 147 nmol L\(^{-1}\) at 485 cm depth.
Pore-water analyses from station PC-3 on all three sampling dates show similar patterns to SG-1 with less methane activity (Fig 1). Sulfate was completely depleted within the upper 300 cm depth. Sulfide concentrations were below the detection limit at this station as well. The methane profiles show an increase in methane concentration immediately after the consumption of sulfate. The maximum methane concentration reached 0.8 mmol L$^{-1}$ at 450 cm depth in the Aug-13 core. Fe(II) profiles show two peaks, one in the upper part of the cores with maximum of 32 µmol L$^{-1}$ at 177 cm depth, and another one with maximum of 64 µmol L$^{-1}$ at 390 cm depth at the methanogenic depth. The δ$^{13}$C$_{DIC}$ decreased from approximately -10 ‰ at the water-sediment interface to -20 ‰ at the SMTZ. Below that zone there was an increase in δ$^{13}$C$_{DIC}$ values to about -5 ‰ due to methanogenesis. H$_2$ concentrations remained around 2 µmol L$^{-1}$ along the core. The few deviating points do not fit a clear pattern and therefore not discussed.

The H$_2$ concentrations at the PC-3 station are higher by one order of magnitude than the concentrations at the SG-1 station. This is probably due to the more intensive methanogenesis process at SG-1 station, as shown by the higher methane concentrations than those at PC-3 station.

Pore-water analyses from the core collected at station PC-5 resembles the profiles of PC-3 station. Sulfate was depleted at approximately 300 cm, and methane concentrations increased below that depth to 0.3 mmol L$^{-1}$. The Fe(II) profile shows two peaks in this core as well, one in the upper part of 20 µM at 150 cm depth and the second of 30 µmol L$^{-1}$ in the methanogenic zone. The δ$^{13}$C$_{DIC}$ value decreased from -5 % at the water-sediment interface to -25 % at the SMTZ, and below that depth δ$^{13}$C values increased to -17 %.

The reactive Fe(III) oxide profile from SG-1 (Fig. S1) shows a general decrease from 3 dry wt % at 13 cm depth to 2.3 dry wt % at 507 cm depth, with two minimum peaks of 2.4 dry wt % at 103 cm and of 1.9 dry wt % at 312 cm. PC-3 profile shows no significant trends in the reactive Fe(III) concentrations.

The values vary between 2.1 dry wt %, which is the maximum point at 167 cm, and 1.8 dry wt %, which is the minimum point at 137 cm depth.

3.2 Abundance and diversity of bacteria and archaea

The qPCR of bacterial and archaeal 16S rRNA genes from the SG-1 core revealed that the abundance of bacterial genes was between 1.46 – 9.45×10$^6$ copies per g wet sediment, while that of archaea was between 8.15×10$^5$ – 2.25×10$^7$ copies per g wet sediment (Fig. 2). These abundances are typical to oligotrophic marine sediments (e.g. South China Sea that contain ~0.5 – 1 % TOC (Yu et al., 2018a). The abundance of bacteria and archaea decreased gradually in the top 95 cm, increased sharply at 125 cm within the SMTZ, remained relatively stable with high abundance at 185 – 245 cm (the top layer of the methanogenic zone), and then decreased. Notably, the abundance of both bacteria and archaea peaked within the methanogenic zone at 245 cm in correspondence with a Fe(II) concentration peak. However, it is not feasible to compare the abundance of archaea and bacteria by this method due to bias caused by the PCR primers used (Buongiorno et al., 2017). The abundance of the mcrA gene increased sharply from the surface layer to the SMTZ, peaked at 155 cm and remained stable at 155 – 245 cm, indicative of active anaerobic methane metabolism in the SMTZ and active methanogenic zone (Fig. 2). Spearman correlation test shows that the abundance of the bacteria and archaea 16S rRNA genes and mcrA genes
correlated with Fe(II) concentration in the methanogenic zone, where mcrA gene correlated the most significantly ($r = 0.5429$, p value = 0.04789).

Illumina-sequencing of the 16S rRNA gene revealed diverse bacterial and archaeal communities throughout the SG-1 core. Although no clear plateau was observed on species rarefaction curve for the current sequencing depth (Fig. S2), Shannon diversity indices reached stable values, indicating that those sequences well covered the diversity of bacterial and archaeal populations in the samples (Fig. S3). Shannon index, based on 16S rRNA gene sequences, showed higher diversity in the top layers of the sediment along with similar values through the core using the bacterial primers, while for sequences using archaeal primers, the values varied in different layers (Table S1). The bacterial sequences were affiliated with the following phyla: Planctomycetes (25.7%), Chloroflexi (23.2 %), Proteobacteria (12.9%), Deinococcus-Thermus (9.9 %), Acidobacteria (3.5%), Aminicenantes (3.3 %), Spirochaetes (2.3%), Deferribacteres (1.7%), Elusimicrobia (1.6%), Nitrospira (1.4%), Firmicutes (1.4 %), Actinobacteria (1.4 %), TM6 (Dependentiae) (1.2%), Marinimicrobia (SAR406 clade) (1.0%), and other taxa with less than 1% of the bacterial communities (Fig. 3a). Bathyarchaeota were the predominant archaea in all the sediment layers, based on the high relative abundance of their 16S rRNA gene sequences (91.0%). The remaining archaeal phyla comprised Euryarchaeota (3.2%), Thaumarchaeota (2.4%), Lokiarchaeota (1.0%), and other phyla with less than 1% of the archaeal communities (Fig. 3b). Spearman correlation analysis revealed that uncultured SBR1093 (r = 0.6176, p value = 0.01859) from bacterial Candidate Phylum SBR1093, subgroup 26 of Acidobacteria (r = 0.5841, p value = 0.02828), the uncultured bacterium from TK10 Class of Chloroflexi phylum (r = 0.5297, p value = 0.0544) and uncultured Bathyarchaeota sp. (archaea) (r = 0.5516, p value = 0.04388) correlated significantly with Fe(II) iron concentration.

3.3 Incubation experiment

Sediment from the observed deep iron reduction zone of SG-1 station from January 2017 core was taken for a basic short-term (few weeks) slurry incubation experiment in order to characterize the iron reduction process in the methanogenic zone. The results of the experiment are shown in figure 4. Dissolved Fe(II) concentrations show significant increase from 11 µmol L$^{-1}$ to approximately 90 µmol L$^{-1}$ during the first three days in all the experimental bottles, except for the killed bottles, implying that the reduction is microbially mediated. Another observation was that the microorganisms were able to reduce both hematite and magnetite to the same extent. In addition, no difference in the Fe(II) concentrations between bottles with and without the addition of H$_2$ was observed.

4 Discussion

This study was performed in the SE Mediterranean in the area of the recently discovered ‘gas front’ (Schattner et al., 2012), where biogenic methane was found at some locations in shallow sediments with low TOC content (Sela-Adler et al., 2015). SG station is located at the center of this area, while PC-3 and PC-5 at the edges, and indeed methane involved processes seem more intensive at this station (SG-1) (Fig. 1). At this station methane reaches higher concentrations, and the intensive methanogenesis also leads to intensive methane oxidation by sulfate at the SMTZ, causing it to occur at shallower depth with
lower δ¹³C DIC values, as observed in previous studies (e.g., Sivan et al., 2007). The shallower SMTZ values also interfered with the ability to observe the traditional iron reduction zone in our SG-1 sampling resolution.

Despite the pore-water profiles variability between the stations, they show a resemblance in their trends. All geochemical pore-water and reactive Fe(III) profiles suggest that the sediments in this area of the SE Mediterranean shelf can be classified into three general depth-zones (Fig. 1): zone 1 is the upper part of the sediment, where the traditional classical iron reduction occurs, probably coupled to organic matter oxidation, with sulfate reduction below it; zone 2 is the SMTZ, where methane starts to increase with depth, sulfate is completely depleted, sulfide is absent and Fe(II) is either present in low concentrations or absent as well (probably due to the precipitation of iron-sulfide minerals); zone 3 is the methanogenic zone, where methane concentrations increased to the highest values in all stations. At this zone, local maxima of Fe(II) concentrations in the pore-water were found in all cores, indicating iron oxides reactivation and reduction. The results of the slurry experiment show only a slight increase in Fe(II) concentrations in the killed bottles compared to the non-killed bottles, indicating that most of the iron reduction in zone 3 is microbial (Fig. 4).

The observed intensive iron reduction in the methanogenic sediments is the first in the Southeastern Mediterranean shelf. The phenomenon of iron reduction in the methanogenic depth has been observed before in other marine provinces (Egger et al., 2016; Jørgensen et al., 2004; März et al., 2008; Riedinger et al., 2014; Slomp et al., 2013; Treude et al., 2014), however, the type of link to the methane cycle is complex. Usually, iron reduction is coupled to oxidation of organic matter (Lovley and Phillips, 1988) and is performed by iron reducing bacteria, which is probably the case in zone 1. It is however questionable if this also stands for zone 3 and if not, what process is responsible for the reactivation of iron oxides at this depth and its relation to methane.

The oligotrophic nature of the studied area would suggest that intensive bacterial iron reduction coupled to the oxidation of organic matter in zone 3 is less likely. The present low nutrient and low chlorophyll concentrations in the water results in low amount of TOC in the sediments, reaching up to ~1% (Sela-Adler et al., 2015). However, we observe in situ biogenic methane formation in the shallow shelf sediments based on the geochemical (δ¹³C DIC) and microbial profiles (population and functional mcrA gene). This indicates that regardless the area’s present oligotrophic nature, the TOC substrate is enough to sustain all the microbial activity up to methanogenesis. These environmental conditions are hypothetically attributed to the Last Glacial Maximum or Mid-Pleistocene sources (Schattner et al., 2012). At the methanogenic zone and below, it might be that the microbial communities present at these depths are used as a food source.

Another potential process that can be coupled to iron reduction in the methanogenic zone is H₂ oxidation. H₂ is an important intermediate in anaerobic aquatic sediments. In this type of environment, it is produced mainly by fermentation of organic matter (Chen et al., 2006), and can be involved in different microbial processes; where each process would need a certain amount of H₂ in order to occur (Lovley and Goodwin, 1988). The H₂ levels at stations SG-1 and PC-3 (Fig. 1) are relatively high (Lilley et al., 1982; Novelli et
al., 1987), suggesting that there is enough H$_2$ to sustain the iron reduction process. The increase in H$_2$ concentration at the methanogenic zone in SG-1 station could be explained by the occurrence of fermentation processes, which enables H$_2$ to accumulate (Chen et al., 2006).

H$_2$ involvement was tested by injecting 1 mL of this gas to the experimental bottles in the methanogenic iron reduction process (Fig. 4). We observed that the increase of Fe(II) concentration was similar in the bottles with H$_2$ addition compared to the bottles without H$_2$. This could mean that either there is enough H$_2$ in the sediments as it is, as implied by the H$_2$ pore-water profiles, or that at the methanogenic depth H$_2$ is not involved in the iron reduction process.

A different way to reactivate the iron reduction process in zone 3 is to have an active sulfur cycle at this depth. In this scenario, Fe(III) is reduced by pyrite oxidation (Eq. 3) (Bottrell et al., 2000), which triggers the sulfur "cryptic" cycle (Holmkvist et al., 2011). In this cycle elemental sulfur and eventually, by disproportionation, sulfide and sulfate are produced, the sulfide reacts with iron-oxide and precipitates as FeS. The sulfate can inhibit methanogenesis (Mountfort et al., 1980; Mountfort and Asher, 1981), which can result in the enhancement of the iron reduction process due to competition for substrate with the methanogenesis process.

The recently discovered iron coupled AOM process (Eq. 3) is another potential process that involves iron oxides reduction in the methanogenic zone. Fe(III) as an electron acceptor for AOM provides a greater free energy yield than sulfate (Zehnder and Brock, 1980), and its global importance was emphasized (Sivan et al., 2011; Lovley, 1991a; Kappler and Straub, 2005; Roden, 2003). In our profiles AOM could be a valid option, as can be inferred from figure 5, where inverse association was observed between the dissolved Fe(II) concentrations in zone 3 and the methane concentrations. In most cores presented in the figure it is apparent that at high concentrations of Fe(II), methane concentrations are low (relatively to the specific core), and at high methane concentrations, Fe(II) concentrations are low (relatively to the specific core). This could be a result of iron-coupled AOM that uses methane to reduce Fe(III)-oxides, releasing dissolved Fe(II) to the pore-water. This can also suggest a type of competitive relationship between methanogenesis and microbial iron reduction; perhaps over substrate, or microbial population switching from methanogenesis to iron reduction metabolism (e.g. Sivan et al., 2016). It should be noted that our experiment was not designed to test AOM due to its short time scale of a few weeks, hence another long experiment with the addition of the $^{13}$C-labeled methane will enable us to shed more light on this association.

Our data profiles and incubation indicate that iron reduction is performed and stimulated by microbial activity. This is despite the fact that the known potential bacterial iron reducers, such as Alicyclobacillus, Sulfoacidilus, Desulfomaculum genera (Firmicutes), Acidiphilium (Alphaproteobacteria); Desulfobacter, Desulfoaromas, Geobacter, Geothermobacter, Anaeromyxobacter (Deltaproteobacteria) and Shewanella (Gammaproteobacteria) (Weber et al., 2006) comprise less than 0.1% of bacteria detected in the methanogenic zone (from 185 cm and below). This is because it appears that both the microbial abundance and the Fe(II) concentration peaked at the methanogenic zone. Cultivation efforts indicated that archaeal methanogens may also play a role in iron reduction within sediments (Sivan et al.,
Moreover, the relative abundance of methane-metabolizing archaea was shown to correlate with Fe(II) concentrations in Helgoland muds from the North Sea, where microbial abundance and the Fe(II) concentrations peaked at the methanogenic zone, similarly to the Mediterranean sediments (Oni et al., 2015). It is possible that methane-metabolizing archaea were involved in the iron reduction in the Mediterranean sediments, as the highest mcrA gene copies per gram wet sediment were detected in the SMTZ and in the top of the methanogenic zone where the Fe(II) concentrations are high. Methanotrophs, such as ANMEs, were found to be involved in iron coupled AOM in marine and freshwater cultures (Scheller et al., 2016; McGlynn et al., 2015; Ettwig et al., 2016; Cai et al., 2018). ANMEs were found with relatively low frequencies (ANME1, below 1% in most samples, circa 5% in the 185 cm layer), and their role in iron reduction within the Mediterranean sediments remains to be tested. It should be noted that even though, the microbial population was tested only on the sediment core that was extracted on January 2017 at SG-1 station, we believe that it represents the general microbial population abundance in the SE Mediterranean continental shelf.

In our study, Spearman correlation analysis revealed that bacterial phyla SBR1093 (candidate Phylum), Acidobacteria and Chloroflexi, as well as archaean Phylum Bathyarchaeota showed significant correlation with a Fe(II) concentration in the methanogenic zone. The Candidate Phylum SBR1093 was firstly identified in phosphate-removing activated sludge from a sequencing batch reactor (Bond et al., 1995), and continuously detected in a short-chain fatty acid rich environment such as wastewater treatment, and marine sediments (Wang et al., 2014). It was thought to be capable of growing autotrophically, but the metabolic capabilities related to iron reduction remain unclear. Strains of Acidobacteria and Chloroflexi phylum were found to be capable of iron reduction (Kawaichi et al., 2013; Kulichevskaya et al., 2014). In addition, members of Acidobacteria were found in iron-coupled AOM enrichment (Beal et al., 2009). The metabolic properties of Subgroup 26 from Acidobacteria and TK10 Class of Chloroflexi are still not known. Bathyarchaeota are globally distributed and account for a considerable fraction of the archaean communities in the marine sediments, particularly, in the Mediterranean Pleistocene sapropels (Coolen et al., 2002; Zhou et al., 2018). While Bathyarchaeota have diverse metabolic capabilities (Lloyd et al., 2013; Meng et al., 2014; Evans et al., 2015; He et al., 2016; Yu et al., 2018b; Feng et al., 2019), their role in iron reduction warrants further studies, as suggested from their high abundance here. Therefore, iron reduction and methane cycling within the deep methanogenic zone may be facilitated by an interplay among bacterial and archaeal groups, whose physiology and functions needs further investigation.

The geochemical and microbial data from the profiles and the slurry incubation experiment suggest that deep iron reduction is occurring in the methanogenesis depth, and that both Bacteria and archaea can be involved in the process. The geochemical profiles show Fe(II) peaks in the deep part of the sediments, indicating iron reduction. The iron reduction was shown also in the incubation experiment, where microbial involvement was evident. The Spearman correlation pointed out several potential microbial players (bacterial and archaean) that correlate to the dissolved Fe(II) profiles (e.g. Bathyarchaeota, Acidobacteria and Chloroflexi). The geochemical conditions lead to three possible microbial iron reduction pathways: a) H2 or organic carbon oxidation, b) an active sulfur cycle, or c) iron driven AOM.
To verify the main iron reduction process at the methanogenic depth of the Mediterranean shelf sediments further incubations and microbial work are needed.

**Author contribution**

H.V and O.S designed research; B.H was the PI of the cruises and M.RB was part of the scientific crew on the ship; H.V, E.W and L.L performed research and analyzed the data; H.V, O.S, B.H, F.W, M.RB and L.L synthesized the data and wrote the paper.

The authors declare that they have no conflict of interest.

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Figures captions:

Figure 1: Geochemical pore-water profiles of sediment cores collected from the three stations SG-1 (top), PC-3 (middle) and PC-5 (bottom) in the Eastern Mediterranean. The profiles are divided roughly to three zones according to the dominant processes: upper microbial iron and sulfate reduction, sulfate-methane transition zone (SMTZ), and the methanogenic zone at the deep part. The dashed line in the CH$_4$ graph at SG-1 station represents the CH$_4$ saturation value in the pore-water. The error bars for CH$_4$ are presented where duplicate sediment samples were collected. The error bars for Fe(II), $\delta^{13}$C$_{DIC}$ and H$_2$ are presented where measurement repetition of each sample was taken (at least twice). The analytical errors were smaller than the symbols.

Figure 2: Sedimentary depth profiles of bacterial and archaeal 16S rRNA and mcrA functional genes of Station SG-1 from January 2017. Triplicates were made for each sample with error bars smaller than the symbols.

Figure 3: Phyla level classification of bacterial (a) and archaeal (b) diversity in the sediments of Station SG-1 from January 2017.

Figure 4: Dissolved Fe(II) results of the sediment slurry incubation experiment from SG-1 core. The sediment was collected on January 2017 from sediment depth of 260 cm. The error bars were smaller than the symbol.

Figure 5: The relationship between dissolved Fe(II) concentrations and methane concentrations in zone 3 sediments at (a) Station SG-1 and (b) Station PC-3. An inverse association is observed between the two species, suggesting a relationship of competition or Fe(III)-coupled anaerobic methane oxidation.
Figures:

Figure 1

![Graphs showing SO₄²⁻, CH₄, δ¹⁵N, Fe(II), and H₂ concentrations over depth for different sampling dates in OG-1, PC-3, and PC-5 zones.](image-url)
Figure 2

Gene copy number per g wet sediment

Depth [cm, baf]

10^3  10^5  10^7

- mcrA
- Bacteria
- Archaea
Figure 3
Figure 4

![Graph showing dissolved Fe(II) concentration over days for different samples.](image)

- Hematite
- Hematite + H₂
- Magnetite
- Magnetite + H₂
- Killed + hematite + H₂
- Killed + magnetite + H₂
Figure 5