Differences in microbial community composition between injection and production water samples of water flooding petroleum reservoirs

P. K. Gao\textsuperscript{1,2}, G. Q. Li\textsuperscript{1,2}, H. M. Tian\textsuperscript{1,2}, Y. S. Wang\textsuperscript{1,2}, H. W. Sun\textsuperscript{3}, and T. Ma\textsuperscript{1,2}

\textsuperscript{1}Key Laboratory of Molecular Microbiology and Technology, Ministry of Education, Tianjin 300071, China
\textsuperscript{2}College of Life Sciences, Nankai University, Tianjin 300071, China
\textsuperscript{3}College of Environmental Science and Engineering, Nankai University, Tianjin 300071, China

Correspondence to: T. Ma (tingma@nankai.edu.cn)

Received: 23 October 2014 – Published in Biogeosciences Discuss.: 5 December 2014
Revised: 5 May 2015 – Accepted: 10 May 2015 – Published: 5 June 2015

Abstract. Microbial communities in injected water are expected to have significant influence on those of reservoir strata in long-term water flooding petroleum reservoirs. To investigate the similarities and differences in microbial communities in injected water and reservoir strata, high-throughput sequencing of microbial partial 16S rRNA of the water samples collected from the wellhead and downhole of injection wells, and from production wells in a homogeneous sandstone reservoir and a heterogeneous conglomerate reservoir were performed. The results indicate that a small number of microbial populations are shared between the water samples from the injection and production wells in the sandstone reservoir, whereas a large number of microbial populations are shared in the conglomerate reservoir. The bacterial and archaeal communities in the reservoir strata have high concentrations, which are similar to those in the injected water. However, microbial population abundance exhibited large differences between the water samples from the injection and production wells. The number of shared populations reflects the influence of microbial communities in injected water on those in reservoir strata to some extent, and show strong association with the unique variation of reservoir environments.

1 Introduction

Water flooding is an efficient and inexpensive oil recovery process that is employed worldwide. Water flooding is believed to be a continued reinoculation of a reservoir with surface microorganisms. After long-term water flooding, microbial populations possessing exceptional survival abilities in injected water are expected to flow into oil-bearing strata, where exogenous and indigenous microbial populations form a new complex ecosystem (Zhang et al., 2012a). When injecting nutrients and air into reservoir strata, these microbial populations can be stimulated, and produce metabolites, such as polysaccharide, surfactants, acid, alcohol, and biogas. Because these metabolites can improve reservoir properties by blocking preferred water flow paths, by lowering interfacial tension between brine and the oil phase, and also by decreasing oil viscosity (Youssef et al., 2009), microbial enhanced oil recovery (MEOR) has been applied to the petroleum industry and is currently being studied extensively (Abdel-Waly, 1999; Zhang et al., 2012b; Bao et al., 2009; Gao et al., 2013; Li et al., 2014).

Microbial populations inhabiting petroleum reservoirs play critical roles in the microbial enhancing of the oil recovery process. As a result, an increasing number of studies, especially those based on culture-independent methods, have been performed, and has broadened our knowledge of microbial diversity in oil reservoirs (Al-Bahry et al., 2013; Kumaraswamy et al., 2011; Lenchi et al., 2013; Okoro et al., 2014; Wang et al., 2012). However, to the best of our knowledge, detailed studies about the effects of the microbial
communities in injected water on those in reservoir strata remains poorly understood. Based on the 16S rRNA gene clone library method, several studies to date have suggested that despite being flooded by the same injected water, there is a significant difference in the communities between each production well (Tang et al., 2012; Ren et al., 2011). Zhang et al. (2012a) compared microbial communities in samples of injection and production wells from reservoirs with different in situ temperatures, and pointed out that the effects of microorganisms in the injected waters on microbial community compositions in produced waters are strongly associated with reservoir temperature. However, because of the low throughput of the clone library method, many infrequent microbial taxa may not be detected, making it difficult to compare microbial communities in detail.

If microbial populations in injected water can flow into reservoir strata and reach production wells, is the microbial community in the injected water expected to have a similar community composition to those in the production wells? If there is a large difference in community composition, what is the difference and how many microbial populations are shared? To explore these issues, we investigated the microbial populations and their abundance in injection and production wells in a homogeneous sandstone petroleum reservoir with a permeability of $522 \times 10^{-3} \, \mu m^2$ and inter-well spacing of 300–425 m using 16S rRNA pyrosequencing and real-time fluorescent quantitative PCR (qPCR). At the same time, we analyzed microbial communities in water samples collected from the wellhead and downhole of injection wells, and from production wells in a heterogeneous conglomerate water-flooding petroleum reservoir with a permeability of $362 \times 10^{-3} \, \mu m^2$ and inter-well spacing of 100–150 m. High-throughput sequencing provides the opportunity to compare microbial populations with unprecedented levels of coverage and detail. The variation in permeability, inter-well spacing, and heterogeneity of the reservoirs is of benefit to exploring the influence of reservoir physical properties on microbial distribution in injected water and reservoir strata. The simi-

### Table 1. Reservoir characteristics of Lu and Liu field block.

<table>
<thead>
<tr>
<th>Reservoir characteristics</th>
<th>Lu field block</th>
<th>Liu field block</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oil Reservoir</td>
<td>Sandstone</td>
<td>Conglomerate</td>
</tr>
<tr>
<td>Formation lithology</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Average depth (m)</td>
<td>1200</td>
<td>1088</td>
</tr>
<tr>
<td>Pressure (MPa)</td>
<td>10.2</td>
<td>7.2</td>
</tr>
<tr>
<td>Stratal temperature (°C)</td>
<td>37</td>
<td>22.6</td>
</tr>
<tr>
<td>Average water content, %</td>
<td>80.8</td>
<td>86.8</td>
</tr>
<tr>
<td>Interwell distances, m</td>
<td>300–425</td>
<td>100–150</td>
</tr>
<tr>
<td>Average permeability, $\mu m^2$</td>
<td>$522 \times 10^{-3}$</td>
<td>$362 \times 10^{-3}$</td>
</tr>
<tr>
<td>Effective porosity, %</td>
<td>29.9</td>
<td>18.96</td>
</tr>
<tr>
<td>Water flooding (yr)</td>
<td>13</td>
<td>30</td>
</tr>
</tbody>
</table>

### Table 2. Chemical properties of the water samples obtained from the Lu and Liu field blocks (Unit: mg L$^{-1}$).

<table>
<thead>
<tr>
<th>Samples</th>
<th>3084</th>
<th>1039</th>
<th>2180</th>
<th>3073</th>
<th>3095</th>
<th>T86–0</th>
<th>T86–8</th>
<th>T93–0</th>
<th>T93–7</th>
<th>T90</th>
<th>T95</th>
<th>T96</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nitrogen</td>
<td>15.1</td>
<td>11.5</td>
<td>10.6</td>
<td>12.7</td>
<td>11.6</td>
<td>7.6</td>
<td>6.8</td>
<td>10.2</td>
<td>11.5</td>
<td>5.7</td>
<td>6.3</td>
<td>8.5</td>
</tr>
<tr>
<td>Phosphorus</td>
<td>20.2</td>
<td>19.1</td>
<td>17.5</td>
<td>18.8</td>
<td>19.5</td>
<td>16.8</td>
<td>15.1</td>
<td>22.6</td>
<td>21.2</td>
<td>18.5</td>
<td>16.2</td>
<td>12.1</td>
</tr>
<tr>
<td>$Na^+, K^+$</td>
<td>4525</td>
<td>4803</td>
<td>4565</td>
<td>4309</td>
<td>4487</td>
<td>3364</td>
<td>3630</td>
<td>3802</td>
<td>4349</td>
<td>4014</td>
<td>3097</td>
<td>3139</td>
</tr>
<tr>
<td>$Mg^{2+}$</td>
<td>21.7</td>
<td>32.1</td>
<td>31.6</td>
<td>26.0</td>
<td>28.8</td>
<td>33.1</td>
<td>63.1</td>
<td>28.4</td>
<td>63.1</td>
<td>68.1</td>
<td>17.5</td>
<td>50.2</td>
</tr>
<tr>
<td>$Ca^{2+}$</td>
<td>191.3</td>
<td>281.9</td>
<td>284.7</td>
<td>181.6</td>
<td>216.4</td>
<td>70.2</td>
<td>77.9</td>
<td>72.8</td>
<td>78.0</td>
<td>96.2</td>
<td>86.4</td>
<td>108.7</td>
</tr>
<tr>
<td>$Cl^-$</td>
<td>5640</td>
<td>6125</td>
<td>5820</td>
<td>5160</td>
<td>5850</td>
<td>3010</td>
<td>3630</td>
<td>2922</td>
<td>3453</td>
<td>3099</td>
<td>3816</td>
<td>3406</td>
</tr>
<tr>
<td>$SO_4^{2-}$</td>
<td>116.2</td>
<td>14.04</td>
<td>4.86</td>
<td>23.13</td>
<td>8.86</td>
<td>483.4</td>
<td>89.4</td>
<td>523</td>
<td>362.4</td>
<td>89.9</td>
<td>65.6</td>
<td>165.1</td>
</tr>
<tr>
<td>$HCO_3^-$</td>
<td>356</td>
<td>434</td>
<td>464</td>
<td>846</td>
<td>511</td>
<td>3140</td>
<td>3823</td>
<td>4052</td>
<td>5687</td>
<td>5837</td>
<td>1915</td>
<td>2841</td>
</tr>
</tbody>
</table>
larity among microbial communities was investigated using hierarchical clustering and Principal coordinate analysis. Microbial populations were also clustered according to injection and production wells to highlight the populations that showed the highest variability.

2 Materials and methods

2.1 Sampling locations

The Lu and Liu field block reservoirs are located in the Xinjiang Oil Field, in the Junggar Basin of the Xinjiang Uygur Autonomous Region, Northwest China. The Lu field block is a homogeneous sandstone reservoir that has been water flooded since 2001. The depth of the sampling horizon is approximately 1200 m with a temperature of 37 °C. The porosity of the reservoir is 29.9 %, with an average permeability of 522 × 10^{-3} \mu m^2. The density of the crude oil is 0.846 g cm^{-3}, with an oil viscosity of 18 mPa × s (Table 1). In the investigated well group (an injection well and four production wells), injection well Lu3084, located in the centre of the production wells, has a direct influence on the neighbouring producers, with inter-well distances of 300–425 m. The Liu field block is a heterogeneous conglomerate reservoir that has been water flooded for approximately 30 years. The depth of the block horizon is approximately 1088 m, with a temperature of 22.6 °C. The porosity of the reservoir is 18.96 %, with an average permeability of 362 × 10^{-3} \mu m^2. The oil density is 0.912 g cm^{-3}, with an oil viscosity of 80 mPa × s. The selected well group includes two injection and three production wells, with an inter-well spacing of 100–150 m. Production well T90 is located at the centre of injection wells T86 and T93, while production wells T95 and T96 are located at the edge of the field block and are mainly flooded by injection well T93 (Fig. 1).

The concentrations of potential nutrient factors, including crude oil properties, total nitrogen (TN), total phosphorus (TP), and ion concentration of formation brines, are listed in Table 2. The differences in geochemical parameters between crude oil samples from the two blocks are not obvious, indicating similar oil characteristics and maturity. The crude oil in both blocks had a higher content of saturates and aromatics, which favour the growth of hydrocarbon-degrading bacteria (HDB), and some anaerobes, such as sulfate-reducing bacteria. The cations and anions of the water samples in the two blocks were similar, with lower nitrogen and phosphorus content, which are essential for the survival and growth of microorganisms. The lack of nitrogen and phosphorus implies a low metabolism level of microorganisms in the reservoir.

2.2 Water samples collection and DNA extraction

Based on tracer techniques, the time interval for injected water to flow from an injection well into neighbouring production wells was approximately 30–45 days in the sandstone reservoir, and 7–10 days in the conglomerate reservoir (data provided by the Xinjiang Oil Field Company). Injected water from the sandstone reservoir was collected on three occasions every 15 days between October 2012 and November 2012, and the produced water samples (from the reservoir strata) were collected along with the second injected water sample on three occasions at a 30-day interval. All of the injected and produced water samples were collected randomly from sampling valves located on the wellhead. In the conglomerate reservoir, the injected water samples were collected in November 2011 from the wellhead and the zone close to downhole (obtained by backflow, that is, the injected water flowed upward through the injection well) of the injection wells. The water samples in reservoir strata were collected from the wellhead of production wells. T86, T93, and Lu3084 are the selected injection wells, and T90, T95, T96, Lu1039, Lu2180, Lu3073, and Lu3095 are the production wells.

For DNA extraction, the residual oil was first removed by heating the sample to 60 °C for 30 min and by phase separation in sterilized separatory funnels. Microbial cells
were then collected from 5 L of each water sample by centri
duggage at 4 °C for 15 min at 10 000 x g in a high-speed cen
trifuge (Beckman, CA 92821, USA). The cell deposits
collected from the same sampling location were mixed and
resuspended with TE buffer (Tris 80, EDTA 40 mM, pH
8.0), and then lysed using a mini bead-beater (BioSpec,
Bartlesville, OK 74005, USA) at 200 rpm for 1 min at room
temperature with 0.1 mm glass beads. After bead beating,
lysozyme was added (final concentration of 1 mg mL
−1), and the samples were incubated at 37 °C for 1 h. Following the
lysozyme treatment, 120 μL sodium-dodecyl sulphate (20 %
SDS, W/V) was added and the samples were incubated at
65 °C for 60 min. Total genomic DNA was then extracted
from the suspension solution using an AxyPrepTM Genomic
DNA miniprep kit (Axygen Biosciences, Tewksbury, MA
01876, USA) according to the manufacturer’s instructions
and stored at −80 °C for subsequent study.

2.3 Pyrosequencing of partial 16S rRNA genes and
sequence analysis

Broadly conserved primers 27F (5′-AGA GTT TGA TCC
TGG CTC AG-3′) and 533R (5′-TTC CCG CCG CTG CTG
GCA C-3′) were used to amplify the bacterial 16S rRNA
gene, while primers 344F (5′-ACG GGT GGC AGC AGG
CGC GA-3′) and 915R (5′-GTG TCT CCC CGC CAA TTC
CT-3′) were used to amplify the archaeal 16S rRNA gene.
PCR reactions were performed following the protocol de
scribed in the Supplement. Replicate PCR products of the
same sample were mixed in a PCR tube. Then, the amplicons
from each reaction mixture were pooled in equimolar ratios
based on concentration and subjected to emulsion PCR to
generate amplicon libraries. Amplicon pyrosequencing was
performed on a Roche Genome Sequencer GS FLX+ plat
form at Majorbio Bio-Pharm Technology, Shanghai, China.

Sequences generated from pyrosequencing were analyzed
using default settings in the open-source software package
mothur (Schloss et al., 2009). The detailed process is de
scribed in the SI. Alpha diversity analyses, including rarefac
tion and computation of the Shannon, Simpson, Chao1 met
ric, and phylogenetic diversity (PD) were used to assess biod
ersity. The similarity among microbial communities was
determined using UniFrac analysis in which weighted and
unweighted principal coordinate analysis (PCoA) were per
formed based on OTUs (Operational taxonomic units) abun
dance or phylogenetic relationships. Specific differences in
community composition of samples were visualized using heatmaps, ggplot, and Venn diagrams using the R software
package.

2.4 Miseq-sequencing of partial 16S rRNA genes and
sequence analysis

The bacterial and archaean 16S rRNA gene V4 region (300–
350 bp) were amplified using primer set 515f (GTG CCA
GCM GCC GCG GTAA) and 806r (GGA CTA CHV GGG
TWT CTA AT) with the protocol described by Caporaso et al.
(2011, 2012). A composite sample for sequencing was cre
ated by combining equimolar ratios of amplicons from the
individual samples, followed by gel purification and ethanol
precipitation to remove any remaining contaminants and
PCR artifacts. Amplicon sequencing was conducted on an
Illumina MiSeq platform at Novogene Co., Beijing, China.

Pairs of reads from the original DNA fragments were
merged using FLASH (fast length adjustment of short reads)
(Magoc and Salzberg, 2011). Sequences were then analyzed
using the Quantitative Insights Into Microbial Ecology (QI
ME) software package (Caporaso et al., 2010), and an OTU
clustering pipeline, UPARSE (Edgar, 2013). The detailed
process is described in the SI. The similarity among microb
ial communities was determined using UniFrac analysis in
which weighted PCoA was performed based on OTUs com
position and phylogenetic relationships. Specific differences
in community composition of samples were visualized using
heatmaps, ggplot, and Venn diagrams using the R package.

2.5 Quantification of community abundance

Evaluation of community abundance by real-time fluorescent
qPCR was performed using the 16S rRNA gene as a molec
ular marker. Reactions were performed using the FastStart
Universal SYBR Green Master PCR mix in a Bio-Rad iQ5
Sequence detection system. The primer set 8F (5′-AGA GGT
TGA T(CT)(AC) TGG CTC-3′)/338R (5′-GCT GCC TCC
CTG AGG AGT-3′) were used to quantify the bacterial com
munity, while 806F (5′-ATT AGA TAC CCS BGT AGT CC
3′)/958R (5′-YCC GGC GTT GAM TCC AAT T-3′) were
used to quantify the archaean community (Gittel et al., 2009).
Tenfold serial dilutions of a known copy number of plasmid
DNA containing the target gene were subjected to real-time
PCR in triplicate to generate an external standard curve. The
PCR efficiency and correlation coefficients for the standard
curves were determined by the melting curve. Gene copy num
bers in unknown samples were determined based on standard
curves.

2.6 Sequence accession numbers

The raw reads were deposited in the National Center for
Biotechnology Information (BioProject ID: PRJNA246768;
3 Results

3.1 Microbial community composition in the sandstone reservoir

Up to 4016–5060 bacterial and 2688–2857 archaeal sequences were obtained by pyrosequencing in the sandstone reservoir. These sequences were assigned into 249–538 bacterial and 45–130 archaeal OTUs at a 3% cutoff (Table S1 in Supplement). The individual rarefaction, Shannon, and Phylogenetic diversity curves tended to approach the saturation plateau (Fig. S1 in Supplement). The results of qPCR indicated that the copy number of bacterial 16S rRNA in water samples of the injection well was 8.25 × 10⁴ copies mL⁻¹, while 1.5 × 10⁶ to 2.75 × 10⁶ copies mL⁻¹ in the water samples of the production wells. Compared with the bacteria, the number of archaea was about 1% of the bacterial number, with 3.75 × 10⁴ 16S rRNA copies mL⁻¹ in injection well and 8.5 × 10³ to 5.75 × 10⁵ copies mL⁻¹ in the production wells.

Phylogenetic analysis indicated that the injected water (Lu3084) was dominated by Proteobacteria (50.43%), Cyanobacteria (15.51%), and Chloroflexi (9.12%). Among the Proteobacteria, Betaproteobacteria (20.42%) and Alphaproteobacteria (19.63%) were numerically dominant, while a small quantity of Deltaproteobacteria (5.49%), Gammaproteobacteria (4.44%), and Epsilonproteobacteria (0.32%) were detected (Fig. 2aI). The produced water from Lu3073 was dominated by Proteobacteria (65.35%) Spirochaetes (13.38%), and Bacteroidetes (12.38%). Gammaproteobacteria (23.96%), Deltaproteobacteria (22.16%), Alphaproteobacteria (13.47%), and Spirochaetes (13.38%) dominated at class level (Fig. 2aI). In the produced water from Lu3095, Lu1039, and Lu2180, Proteobacteria composed 78.58–95.75% of the bacterial communities. Alphaproteobacteria (15.43), Deltaproteobacteria (23.48, 50.57, 12.94%), and Epsilonproteobacteria (2.79, 4.38, 25.54%) were dominant (Fig. 2aII).

To archaeal communities, more than 95% of the archaeal sequences were assigned to Methanobacteria, Methanococci, and Methanomicrobia (Fig. 2aII). In the injected water, 87% sequences were classed into Methanomicrobia, and the dominant genera were Methanoseta (42.39%), Methanomethylovorans (25.57%), and Methanolobus (10.96%). Among them, Methanoseta uses only acetate to produce CH₄. Methanomicrobia accounted for 84.03% in the produced water of Lu1039, and Methanolobus (83.46%) and Methanococcus (11.23%) were the dominant genera. The archaeal communities were much more conserved in the produced water at Lu2180, Lu3073, and Lu3095, with Methanococcus accounting for 95.34, 90.79, and 86.79%, respectively. The Methanolobus and Methanococcus are methylotrophic and hydrogenotrophic methanogens.

3.2 Microbial community composition in the conglomerate reservoir

Between 52,719 and 129,106 16S rRNA gene sequences were analyzed and assigned to 2623 – 3414 genus-level OTUs (Table S2). In combination with the relative abundance, the number of bacterial and archaeal sequences was calculated, with the number of sequences per sample ranging in size from 51,273 to 128,980 and 85 to 1445, respectively (Fig. S2). Based on the results of qPCR, the copy number of bacterial 16S rRNA in the water samples ranged from 1.5 × 10⁷ to 6.5 × 10⁷ copies mL⁻¹, while archaeal 16S rRNA ranged from 4.5 × 10⁵ to 8.5 × 10⁵ copies mL⁻¹.

In contrast to the sandstone reservoir, Proteobacteria, Bacteroidetes, Firmicutes, Spirochaetes, and Synergistes were simultaneously detected in the water samples from the injection and production wells, composing 85.7–94.1% of all bacterial communities. Similar to the sandstone reservoir, more Proteobacteria were detected in water samples from production wells. At the class level, Gammaproteobacteria, Epsilonproteobacteria, Alphaproteobacteria, Betaproteobacteria, Deltaproteobacteria, Bacteroidia, Bacilli, and Clostridia composed 74.5–83.7% of the bacterial communities in the water samples from the injection and production wells (Fig. 2bI). The archaea were mainly assigned to Methanomicrobia, Methanococci, Methanobacteriales, Thaumarchaeota, Parvarchaeae, and Thermoplasmata (Fig. 2bII). Among them, Methanobacteria, Methanococci, and Methanomicrobia were simultaneously detected, and composed 64.3–94.6% of the archaeal communities in the water samples from the injection and production wells. Compared with the injected water collected from the wellhead of the injection wells (T86-0 and T93-0), more Methanomicrobia were detected in the downhole of injection wells (T86-8 and T93-7) and production well T90. At genus level, Methanocorpusculum, Methanococcus, and Methanocalculus were dominant, accounting for 60.3–88.5% of the archaeal communities in the injection wells and production well T90. The three taxa can use H₂ and formate as carbon sources to produce CH₄.

3.3 Shared microbial populations between injected water and reservoir strata

The shared microbial OTUs and genera between communities in the water samples from the injection and production wells were investigated using Venn diagrams, histograms, and a heatmap. Based on the Venn diagrams, 16.3–32.81% of bacterial OTUs and 13.73–51.61% of archaeal OTUs were shared between the water samples from the injection well and those of the production wells in the sandstone reservoir (Fig. 3a). These shared bacterial OTUs accounted for 4.6–24.71% of the total bacterial community in water samples from injection well, and 43.23–76.18% in the production wells (Fig. 4aI). Furthermore, only 13 bacterial and 3
archaeal OTUs were shared by both the water samples from the injection and production wells (Fig. 3a). Accordingly, the shared genera only accounted for 2.26% of the bacterial community in the injected water, but dominated each production well (12.02–36.5%; Fig. 4aII). Similar to the bacteria, three archaeal genera belonging to Methanobacterium, Methanococcus, and Methanobrevibacter were simultaneously detected, comprising 13.58% of the total archaea in the water samples from the injection well, and 90.4–96.89% in the water samples from the production wells (Fig. 4a III).

In the conglomerate reservoir, most of the OTUs and genera were simultaneously detected in the water samples from the injection and production wells (Figs. 3b and 4b). Similar to the sandstone reservoir, these shared populations accounted for a minor proportion of the communities in the water samples collected from the wellhead of injection wells, but dominated the water samples obtained from the downhole of injection wells and each production well (Fig. 3b).

### 3.4 Microbial population distribution in injected water and reservoir strata

Microbial populations were clustered according to injection and production wells to highlight the populations that showed the most variability (Fig. 5). In the sandstone reservoir, more *Sphingomonas* and *Azospirillum* were detected in the water samples from the injection well, while *Arcobacter*, *Marinobacterium*, *Pseudomonas*, *Hyphomonas*, *Novispirillum*, *Proteiniphilum*, *Spirochaeta*, and *Rhizobium* were highly abundant in the water samples from the production wells. In the conglomerate reservoir, higher amounts of *Paracoccus*, *Bacillus*, *Ochrobactrum*, *Parabacteroides*, *Sphaerochaeta*, *Thauera*, *Halomonas*, and *Alcanivorax* were detected in the water samples from the injection well, while *Arcobacter*, *Marinobacterium*, *Pseudomonas*, *Bacteroides*, *Oleibacter*, *Marinobacter*, and *Shewanella* were dominant in the downhole of the injection and production wells. Among them, *Marinobacterium*, *Paracoccus*, *Ochrobactrum*, *Sphingomonas*, *Alcanivorax*, and *Azospirillum* are aerobic bacteria, while *Pseudomonas*, *Rhizobium*, *Arcobacter*, *Halomonas*, *Spirochaeta*, *Bacillus*, *Thauera*, *Halomonas*, and *Bacteroides* are microaerophilic bacteria, facultative anaerobes, or anaerobes.

To further investigate the microbial distribution in injected water and reservoir strata, hierarchical clustering and Unifrac PCoA were performed based on microbial OTUs abundance and phylogenetic relationships. In the sandstone reservoir, hierarchical clustering showed that the community in the water samples from the injection well was distinct from those of the production wells (Fig. S3). Weighted PCoA distinguished the bacterial community of the injected water from those of the production wells, while communities of the production wells were placed at a comparatively decentralized position...
Figure 3. Venn diagrams of the bacterial and archaeal OTUs in the injection and production wells. (a) Sandstone and (b) conglomerate reservoirs. (I) Bacterial OTUs; (II) archaeal OTUs. Venn diagrams indicate the shared microbial OTUs between communities in the injected and produced water samples.

(Fig. 6a bacteria). Similar to the bacterial communities, hierarchical clustering and PCoA distinguished the archaeal community of the injected water from those of the production wells, whereas production wells were placed at a close proximity (Fig. 6a archaea). In the conglomerate reservoir, communities of water samples collected from the wellhead of injection wells clustered into a group in the PCoA plot, indicating that communities remained unchanged before injected water flowed into the injection wells (Fig. 6b). Communities in the water samples collected from the downhole of injection wells and neighbouring production well T90 clustered into one group, while production well T95 and T96 clustered into another (Fig. 6b). This shows that the microbial community reassembled during the process of the injected water flowing into the reservoir strata and each production well.

4 Discussion

MEOR technique is generally classified into exogenous MEOR and indigenous MEOR (Youssef et al., 2009). The former includes injection of exogenous microorganisms and injection of ex situ produced products into reservoirs to enhance oil recovery (Zobell, 1947). This is an effective way to quickly improve oil recovery. However, because of the sieve effect of strata on microbial cells, the injected microorganisms are generally difficult to migrate into reservoir strata (Youssef et al., 2009; Brown, 2010). Diverse microbial populations inhabit petroleum reservoirs. Among them, HDB, nitrate-reducing bacteria (NRB), sulfate-reducing bacteria (SRB), and methanogens are the most important functional populations in reservoir ecosystems (Youssef et al., 2009).

When injecting nutrients into reservoir, these microbial populations can be stimulated and produce biosurfactants, organic acids, and gas, which improve oil recovery (Belyaev SS et al., 1998). Compared with exogenous MEOR, indigenous microorganisms are more adapt to the environmental conditions present in reservoirs. Additionally, because nutrients can be easier to migrate into reservoir strata, indigenous MEOR has higher oil displacement efficiency. Despite the validity in field trial, indigenous MEOR also has some limitations, in particular, the uneven oil displacement efficiency in different production wells in the same reservoir block. The community composition and diversification have been found to have a significant influence on oil displacement efficiency (Li et al., 2014). Therefore, it is needed to investigate the relationship between microbial communities in injected water and reservoir strata, because microbial communities in injected water are expected to flow into oil-bearing strata, and produce a significant influence on those of reservoir strata in long-term water flooding petroleum reservoirs (Youssef et al., 2009; Dahle et al., 2008).

Molecular methods have been widely used to assess the microbial diversity of petroleum reservoirs. Compared to the traditional 16S rRNA gene clone library and sequencing, high-throughput sequencing has generated hundreds of thousands of short sequences, and significantly improved our ability to compare microbial populations with unprecedented levels of coverage and detail (Caporaso et al., 2012). In the conglomerate reservoir, Miseq-sequencing produced approximately 52,719 to 129,106 16S rRNA gene sequences. The sequencing reads were approximately 10–20-folds of those (obtained by pyrosequencing) of the sandstone reservoir, and 50–400-folds of the 16S rRNA gene clone library (assuming 300 clones per library). We simultaneously sequenced the bacterial and archaeal V4 region of 16S rRNA gene, obtaining a total of 51,273–128,980 bacterial sequences per sample, but only 85–1,445 archaeal sequences. This is consistent with the count result for archaea, which are about one percent of bacterial number. The result implies that deeper sequencing is needed for detecting rare archaeal populations using Miseq-sequencing based on bacterial and archaeal universal primer set 515f and 806r. In contrast, the bacterial and archaeal communities were sequenced independently using pyrosequencing in the sandstone and we obtained 4,016–5,060 bacterial and 2,688–2,857 archaeal sequences. The rarefaction curves, Shannon diversity estimates, and observed species suggest that this sequencing depth was enough for the investigation of the bacterial and archaeal communities.

If the microbial populations in the injected water could flow into the reservoir strata and reach the production wells along with the injected water, is the microbial community...
in the injected water expected to have a similar community composition with those in the production wells? In the homogeneous sandstone reservoir, we found that the number of shared bacterial and archaeal populations between the water samples from the injection and production wells was different. As shown in Fig. 3a, 16.3–32.81 % of bacterial OTUs and 13.73–51.61 % of archaeal OTUs were shared between the water sample of injection well and those of production wells. It is reasonable to speculate that microbial populations in the injected water produce different levels of impact on those in production wells. Based on the previous research, the main reason may be the sieve effect that can be enhanced by the long inter-well spacing (Ren et al., 2011). Because of this effect on microbial cells when injected fluid passes through a subsurface formation, it is more difficult for microbial cells to migrate in reservoir strata. In contrast, almost all OTUs and genera detected in the injected water were also observed in the downhole of the injection and neighbouring production wells in the heterogeneous reservoir. Compared with the sandstone reservoir, this reservoir has a similar permeability, but shorter inter-well spacing. It appears that most microbial populations in the injected water flowed into the oil-bearing...
strata and reached the production wells during water flooding process. Additionally, despite lacking sufficient nutrients, bacterial and archaeal communities in the reservoir strata have high concentrations, which are similar to those in the injected water. This phenomenon implies some correlations of microbial communities in injected water and reservoir strata. However, we appreciate that it is less rigorous to delineate the transport of microbial populations in the reservoir strata simply by detecting the shared microbial populations in the injection and production wells using 16S rRNA sequencing, because this method is not able to demonstrate whether the species detected in the produced water are the same ones as in the injected water. Therefore, labelled strains, such as ones containing green fluorescent protein, may be a suitable way to investigate microbial migration in petroleum reservoirs.

The number of shared microbial populations reflects the influence of microbial communities in injected water on those in reservoir strata to some extent, and show strong association with the unique variation of reservoir environments. In contrast to the sandstone reservoir, a large number of microbial populations were shared between the water samples of injection wells and those of production wells in the conglomerate reservoir. However, the community structure, particularly the abundance of the shared OTUs and genera, exhibited a large difference between the injected water and reservoir strata. The environmental variables, such as salinity, pH, and nutrients were supposed to be the primary drivers for the community diversification (Kuang et al., 2013). However, few differences in cations and anions among the injected and produced water samples were observed. Petroleum reservoir represents an anaerobic environment with multiphase fluids of oil, gas, and water. Therefore, except for the sieve effect of reservoir strata on microbial migration, dissolved oxygen, which is known to be strongly related to reservoir microbial growth and metabolism (Gao et al., 2013), may be the main factor influencing the community structures. In both of the reservoirs, aerobic bacteria, including *Sphingomonas*, *Azospirillum*, *Paracoccus*, *Ochrobactrum*, *Alcanivorax*, and *Hydrogenophilaceae* were more frequently detected in water samples from the injection wells, while microaerophilic bacteria, facultative anaerobes, and anaerobes including *Pseudomonas*, *Rhizobium*, *Arcobacter*, *Halomonas*, *Spirochaeta*, and *Bacteroides*, were found to have higher relative abundance in reservoir strata (Fig. 5).

Apart from the dissolved oxygen, crude oil, particularly the saturated and aromatic hydrocarbon, may also strongly influence the microbial distribution in injected water and reservoir strata. Petroleum reservoirs represent oligotrophic environments. Although diverse microbial populations inhabit the reservoirs, only hydrocarbon-degrading bacteria and some anaerobes, such as sulfate-reducing bacteria, could
grow with crude oil as the carbon source. This is consistent with the observed results that more hydrocarbon-degrading bacteria, including *Marinobacterium*, *Pseudomonas*, *Rhizobium*, *Halomonas*, and *Oleibacter*, were detected in the downhole of injection and production wells.

This study compared the differences in microbial community composition between injected water and reservoir strata using microbial genomes obtained from the aqueous phase. In fact, each component of the reservoir multiphasic fluid, including crude oil, gases, and insoluble particles, may act as an important habitat for microbial growth in addition to the water phase within the petroleum reservoir (Kryachko et al., 2012; Kobayashi et al., 2012). Recent research has also compared microbial communities in aqueous and oil phases of water-flooded petroleum reservoirs, and found that the oil phase also harboured a large number of microorganisms, with large differences in the bacterial community between the aqueous and oil phases of the reservoir fluid (Wang et al., 2014). Therefore, simultaneous analysis of DNA extracted from both aqueous and oil phases may provide a better understanding of the microbial communities in injection and production water samples.

In summary, this study investigated the relationship shared by microbial communities in injected water and reservoir strata in two long-term water flooding reservoirs. However, the results cannot provide any reliable information on the indigenous microbial community. The indigenous microbial populations may be those in newly drilled wells without water flooding in the same oil-bearing block. However, the two reservoirs have been water-flooded for decades. Due to the introduction of exogenous microorganisms in injected water and other sources of contaminations by enhanced oil recovery processes, determining whether a microorganism is indigenous to a petroleum reservoir becomes increasingly difficult. This study implies that the number of shared populations reflects the influence of microbial communities in injected water on those in reservoir strata to some extent, and shows strong association with the unique variation of reservoir environments. However, it cannot make a conclusion on the transport of microbial populations in the reservoir strata by detecting the shared microbial populations in injected water and produced water samples using 16S rRNA sequencing. To further investigate the relationship shared by microbial communities in injection and production wells, injecting labelled strains containing marked gene (e.g., green fluorescent protein coded gene) into reservoirs may bring novel insight and greater predictive power. Therefore, further research on microbial diversification and transferability as injected water flows into reservoir is needed. Solving these problems is significant to guide the application of MEOR approaches based on injecting nutrients or microbial populations into reservoirs.
5 Conclusions

Using high-throughput sequencing, this study revealed the similarities and differences in microbial communities in water samples from injection wells and those from reservoir strata in a homogeneous sandstone reservoir and a heterogeneous conglomerate reservoir. The bacterial and archaeal communities in the reservoir strata have high concentrations, which are similar to those in water samples from injection wells. However, microbial community compositions exhibited large differences between the water samples from injection and production wells. The number of shared populations reflects the influence of microbial communities in injected water on those in reservoir strata to some extent, and show strong association with the unique variation of reservoir environments. Additionally, aerobic bacterial populations were more frequently detected in water samples from injection wells, while microaerophilic bacteria, facultative anaerobes, and anaerobes dominated the reservoir strata.

The Supplement related to this article is available online at doi:10.5194/bg-12-3403-2015-supplement.

Acknowledgements. This study was supported by the National High Technology Research and Development Program of China (grant No. 2013AA064402), the National Natural Science Foundation of China (grant No. 41373074), the Chinese Postdoctoral Science Foundation (grant No. 2014MS1175), and the Key Technology Research and Development Program of Tianjin (grant No. 14ZCZDSY00016).

Edited by: Z. Jia

References


