Effects of graphene oxide on the performance, microbial community dynamics and antibiotic resistance genes reduction during anaerobic digestion of swine manure

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\textbf{A B S T R A C T}

The role of graphene oxide (GO) on anaerobic digestion (AD) of swine manure concerning the performance, microbial community and antibiotic resistance genes (ARGs) reduction was investigated. Results showed that methane production was reduced by 13.1%, 10.6%, 2.7% and 17.1% at GO concentration of 5 mg/L, 50 mg/L, 100 mg/L and 500 mg/L, respectively, but propionate degradation was enhanced along with GO addition. Both bacterial and archaeal community changed little after GO addition. AD could well reduce ARGs abundance, but it was deteriorated at the GO concentration of 50 mg/L and 100 mg/L and enhanced at 500 mg/L, while no obvious changes at 5 mg/L. Network and SEM analysis indicated that changes of each ARG was closely associated with variation of microbial community composition, environmental variables contributed most to the dynamics of ARGs indirectly, GO influenced the ARGs dynamics negatively and (heavy metal resistance genes (MRGs)) influenced the most directly.

1. Introduction

Antibiotic resistance threatened the public health more and more seriously, and ARGs was generally considered as an emerging pollutant (Dang et al., 2017). The use of antibiotics in the farming made a huge contribution to the improvement of the human living standard, because antibiotics could not only preserve the disease but also promote the growth and the feed efficiency in the livestock, which warranted the increasing demand for animal protein (Boeckel et al., 2015). China is the biggest antibiotics consumer and producer, and it was estimated that the total antibiotics consumption was 162,000 tons and 84,240 tons (ca. 52%) were used for the animals (Zhang et al., 2015). Thus, manure became an important reservoir of antibiotic resistance genes (ARGs), especially in Chinese swine farms (Larson, 2015).

China raised and consumed half the planet’s pigs, which produced an estimated 618 billion kilograms of manure every year (Larson, 2015). Considering the huge amounts of swine manure, anaerobic digestion (AD) was widely adopted due to its effectiveness on the pathogens removal, volume reduction and resource utilization, especially the biogas production, and luckily, it was demonstrated that AD could also realize some ARGs reduction in swine manure (Youngquist et al., 2016). But the digestate still contributed a lot to the spread and proliferation of ARGs in the environment (Sui et al., 2016) and there was much space for the improvement of AD efficiency (Astals et al., 2012), which indicated that some work could be done to simultaneously enhance the AD efficiency and ARGs removal.

Graphene oxide (GO) has attracted significant attention due to its remarkable structural, chemical, mechanical, and electronic properties and has shown great application potential in many fields (Zou et al., 2016). GO was reported to enhance the growth of human gut bacteria while inhibit the proliferation of pathogens (Chen et al., 2014), and the activity of anammox bacteria for the nitrogen removal could be enhanced by GO addition (Wang et al., 2013a,b; Wang et al., 2014). Also, it was demonstrated that the enhancement of microbes proliferation under methanogenic conditions was elucidated through the electron shuttle effects of GO, and graphene could boost the biomethane yield and production rate due to the enhancement of direct interspecies electron transfer in AD (Colunga et al., 2015; Lin et al., 2017).
However, GO could also cause acute toxicity on activated sludge, and the toxic effects were dose dependent (Ahmed and Rodrigues, 2013). While the effects of GO addition on the performance and its microbial mechanisms of swine manure AD have never been clarified.

Moreover, It has been demonstrated that GO has the potential ability to control the antibiotics uptake and ARGs transfer because of the high surface area and the specific sp²/sp³ structure (Zou et al., 2016), and GO nanosheets exhibited excellent removal efficiency on ARGs in aquatic environments due to the physical absorption and the GO-ARG noncovalent combination (Yu et al., 2016). But the ARGs types and abundances in mouse gut increased after the GO exposure (Xie et al., 2016), and Guo and Zhang (2017) indicated that GO concentration above 10 mg/L can damage resistant plasmids, but GO concentrations lower than 1 mg/L led to almost no damage. Further research is needed to evaluate the overall effects of GO on the ARGs control in various environments. Then, considering the increasing ARGs pollution in swine manure, whether the GO addition during swine manure AD could favor the enhancement of ARGs removal deserved to be elucidated.

In this study, batch experiments of swine manure AD at the GO addition of 0 mg/L, 5 mg/L, 50 mg/L, 100 mg/L, and 500 mg/L were established to find out the effects of GO on the performance and ARGs reduction. Microbial community dynamics were studied to explore the microbial mechanisms. Besides, swine manure not only contained high amounts of antibiotics but also heavy metals and human potential pathogens (Pal et al., 2015; Zhang et al., 2016). The co-selection from heavy metals to ARGs has been widely demonstrated (Pal et al., 2015), and the co-occurrence of virulence factors in human potential pathogens with ARGs generally existed (Zhang et al., 2016). Thus, heavy metal resistance genes (MRGs) to reflect the heavy metal selective pressure and virulence factors (VFs) to reflect the human pathogens were investigated to link the factors influencing the ARGs profiles.

2. Materials and methods

2.1. Materials

Swine manure was collected from an intensive pig farming facility in Beijing, China, and the farm run an up-flow solid reactor heated by solar and controlled at 37 °C. The solid solids (TS) and volatile solids (VS) of the swine manure were 28.3% and 80.4%, respectively. Seed sludge was collected from the same facility, and the TS and VS of the seed sludge were 4.9% and 55.8%, respectively. GO nanosheets (XF002-1) were obtained from the Nanjing XFNANO Materials Tech Co., Ltd., China. The synthesis method was Hummers, diameter was 500 nm–5 μm, Thickness was 0.8–1.2 nm, the single layer ratio was ca. 99% and the purity was above 99 wt%. The GO nanosheets were suspended in deionized water to prepare the stock solutions at the concentration of 2 mg/mL by sonication dispersion method for ca. 2 h.

2.2. Batch experiments set-up

Batch experiments were set up by using the Automatic Methane Potential Test System II (Bioprocess Control, Sweden) in which digesters were a series of serum bottles (working volume: 0.4 L) equipped with a sampling tap and plastic caps including agitators and rubber stoppers. The mixing was semi-continuous at the rate of 150 rpm in the cycle of 1 min mixing and 1 min resting. Swine manure and seed sludge used for the batch experiments were mixed thoroughly at a ratio of 3:1 with a sampling tap and plastic caps including agitators and rubber stoppers. The mixing was semi-continuous at the rate of 150 rpm in the Pressure and virulence factors (VFs) to reflect the potential of HGT and multi-resistance (Engelstädter et al., 2016). The functional gene mcrA involved the methane production was quantified to reflect the effects of GO addition on the methanogenesis (Ziganshin et al., 2016).

MRGs including merA, peaA, arsC and cexA were chosen to represent a water bath to control temperature at ca. 37 °C. The methane production were automatically measured excluding CO₂ which was removed by a gas-washing bottle containing 3 M NaOH solution. The batch experiments lasted for 22 days, and sampling was conducted on days 0, 5, 13 and 22 for further analysis.

2.3. Chemical analysis

TS was determined by drying a 10-mL sub-sample for 24 h at 105 °C, and dried solids were incinerated for 1 h at 600 °C to constant weight for VS measurement. Samples were centrifuged at 4000 rpm for 10 min and filtered through 0.45-μm cellulose membrane. The filtrate was analyzed for pH, NH₄⁺-N, Total and soluble chemical oxygen demand (TCOD and SCOD), proteins, polysaccharides and VFAs as previously described (Zhang et al., 2016).

2.4. DNA extraction

0.5 mL of each sludge sample was used for DNA extraction using FastDNA Spin Kit for Soil (MP Biomedicals, USA) according to manufacturer’s instructions, and quality and concentration of the extracted DNA were determined through 1% agarose gel electrophoresis and NanoDrop ND-1000 (NanoDrop, USA), respectively. The three resulting extracts from the same treatment on each sampling day were composited to get a representative DNA sample for further analysis.

2.5. Microbial community analysis

Both bacterial and archaearial community changes were followed through high-throughput sequencing method, and the primers 515F/806R were selected mainly for bacterial community analysis, while archaearial community was analyzed using the nested PCR by the primers Arch340F/Arch1000R and Arch349F/Arch806R as previously described (Zhang et al., 2016). Amplicons were then sent out to Sangon Co., Ltd., in Shanghai for small-fragment library construction and pair-end sequencing using the Illumina MiSeq sequencing system (Illumina, USA).

Pair-end reads were firstly merged using PEAR (–x 0.1), and then sequencing reads were assigned to each sample according to the unique barcode. PRINSEQ was used for the quality control of these merged reads. PCR chimera were filtered out through UCHIME to get the clean sequences. The clean sequences were normalized to the same sequencing depth (ca. 24,000) for further analysis. The clean sequences were submitted to the NCBI Sequence Read Archive (SRA) under the project number of PRJNA388021. The taxonomic classification of the sequences was carried out using the Ribosomal Database Project (RDP) Classifier, and the classifier data was denoised through the package ad4e in R. The OTUs with an abundance below 0.01% were removed. The alpha diversity index including Shannon, ACE, Chao1 and Simpson were calculated through the Mothur software package (https://www.mothur.org/).

2.6. qPCR

Nine frequently detected and widely existed ARGs including sulI, sulII, ermB, ermF, blaTEM, blaCTX-M, tetM, tetG and tetX were selected to represent the changes of ARGs in swine manure (Luby et al., 2016). Also, the recently emerged plasmid-mediated polymixin resistance gene, mcr-1, with high dissemination risk was chosen, while it has never been quantified in swine manure (Liu et al., 2015). The changes of intI1 representing the mobile genetic elements (MGEs) was followed to reflect the potential of HGT and multi-resistance (Engelstädter et al., 2016). The functional gene mcrA involved the methane production was quantified to reflect the effects of GO addition on the methanogenesis (Ziganshin et al., 2016).
the selection pressure from Hg, Ca, As, Co, Zn and Cd, respectively, which belonged to the typical heavy metals existed in swine manure (Wang et al., 2013a,b). VFs in typical potential pathogens of swine manure including *Escherichia coli* (*eaeA*) and *Enterococcus faecium* (*asa1*) were chosen to present the changes of real pathogens with pathogenicity (Paton and Paton, 1998; Vankerckhoven et al., 2004). The qPCR process was described as previously described (Zhang et al., 2017), and the primers, annealing temperature and the corresponding amplification efficiencies in this study were summarized in Tables S1 and S2.

2.7. Data analysis

Free ammonia concentration was calculated using the formula (Hansen et al., 1998): $\text{Free ammonia} = \text{(Total } \text{NH}_4^+ - \text{N}) \times (1 + 10^{-\text{pH}/10^{-0.09018 + \frac{2729.92}{T}}})^{-1}$, and $T$ indicated the temperature (Kelvin). The heat map of the top 10 genus in each sample based on log2 transformed of the abundance was built through the HemI (http://hemi.biocuckoo.org/). Principal component analysis (PCA) was performed using Canoco 5.0 (http://www.canoco5.com/). Network based on spearman analysis and structural equation model (SEM) based on the correlation matrix were conducted through Gephi (https://gephi.org/) and AMOS (SPSS Inc., Chicago, IL, USA), respectively. The abundance indicated the specific gene copies divided by the corresponding 16 s rRNA gene copies.

3. Results and discussion

3.1. Effects of GO addition on the performance

3.1.1. Methane production

The methane production reached 297.66 ± 4.02 mL CH$_4$ g$^{-1}$VS$_{added}$ for the Control, which was comparable to other studies concerning the swine manure AD (Abdelsalam et al., 2017), while GO addition reduced the accumulative methane production. The methane reduction rate was 13.06 ± 6.96%, 10.60 ± 5.98%, 2.68 ± 1.21% and 17.07 ± 1.51% for the GO-5, GO-50, GO-100 and GO-500, respectively, compared to the Control (Fig. 1A). The methane reduction was not dose-dependent, and the maximum reduction was at 5 mg/L and 500 mg/L, while the least methane reduction at the GO addition of 100 mg/L.

There existed two peaks (D2 and D11~12) on the daily methane production shown in Fig. 1B. The first peak period lasted for ca. 3 days, while the second peak, ca. 10 days, and the reduction of methane production due to GO addition mainly happened at the second peak. It was deemed that the first peak was attributed to the fast degradation of easily degradable organics, while the second peak derived from the poorly biodegradable organics.

3.1.2. Chemical parameters

Interestingly, although accumulative methane production was reduced after GO addition, the VS reduction was increased by 5.71–16.77%, and the TS reduction also increased in all the treatments (maximum, 7.79%) except GO-5, which was in accordance with the changes of TCOD (Table 1). These indicated that although methane production was reduced after GO addition, the volume reduction was enhanced. The experimental set-up excluded the CO$_2$ production, and it was speculated that GO addition should have increased the CO$_2$ production which was absorbed by the gas-washing bottles, while this needed further detail investigation.

The GO addition had little influence on the changes of pH and NH$_4^+$-N concentration, and they increased to the maximum at D13 and D5, respectively, while free ammonia reached its maximum at D13. The increase of NH$_4^+$-N was due to the degradation of organics like proteins, while the delay of the maximum of pH could be attributed to the balance of VFAs accumulation and NH$_4^+$-N concentration.

3.1.3. VFAs

During the first peak, the methane production was mainly from the acetate and butyrate fermentation, while it was iso-valerate and propionate fermentation for the methane production at the second peak (Fig. 2). There was no much difference at the first peak between treatments, while the propionate fermentation was increasingly

![Fig. 1. The profiles of accumulative (A) and daily (B) methane production during swine manure AD.](image-url)
enhanced along with the GO addition concentrations at the second peak. The propionate accumulation removal rate was 2.71%, 3.40%, 5.34% and 10.51% for the GO-5, GO-50, GO-100 and GO-500, respectively, while it was 3.63% for the Control. Propionate is an important intermediate of the degradation of organic matter during AD, and due to thermodynamic constraints, the oxidation of propionate requires syntrophic cooperation of propionate-fermenting proton-reducing bacteria and H2-consuming methanogens (Narihiro et al., 2012). Graphene had been demonstrated to enhance the methane production during ethanol (also a key intermediate produce for the methane production) AD due to the enhancement of the direct interspecies electron transfer (DIET) (Lin et al., 2017). It seemed that the DIET of the propionate fermentation was also enhanced during swine manure AD, but what was different was that the oxygenic groups in the GO might also delivered the oxygen to the methanogens, which led to higher CO2 production during swine manure AD. This could well explain the enhancement of VS, TS and TCOD removal but reduction of methane production after GO addition, while this needed further investigation to elucidate the hypothesis.

### Table 1

The characteristics of materials and samples during anaerobic digestion in this study.

| Samples | pH     | TCOD (g/L) | SCOD (g/L) | NH4+–N (g/L) | Free ammonia (g/L) | Proteins (g/L) | Polysaccharides (g/L) | TS (%) | VS (%) |
|---------|--------|------------|------------|--------------|--------------------|----------------|-----------------------|--------|--------|
| Manure  | 7.20 ± 0.01 | --         | --         | --           | --                 | --             | 28.28 ± 0.04          | 80.41 ± 0.15 |
| Seed    | 7.52 ± 0.04 | 35.52 ± 2.47 | 5.44 ± 0.45 | 3.32 ± 0.08  | 0.14               | --             | 4.90 ± 0.01           | 55.78 ± 0.17 |
| D0      | 7.48 ± 0.09 | 75.57 ± 2.77 | 13.73 ± 0.56 | 2.01 ± 0.06  | 0.07               | --             | 5.02 ± 0.68           | 7.66 ± 0.03  | 73.99 ± 0.04 |
| Control | 7.61 ± 0.04 | 61.31 ± 5.40 | 13.60 ± 0.23 | 3.22 ± 0.09  | 0.16               | 5.35 ± 0.58     | --                    | --      |
| D13     | 7.69 ± 0.02 | 48.32 ± 2.01 | 11.99 ± 0.59 | 2.97 ± 0.07  | 0.18               | 5.12 ± 0.50     | --                    | --      |
| D22     | 7.52 ± 0.02 | 44.69 ± 3.61 | 7.19 ± 0.08  | 3.00 ± 0.09  | 0.12               | 4.46 ± 0.41     | 4.88 ± 0.14           | 62.44 ± 1.95 |
| GO-5    | 7.59 ± 0.01 | 66.03 ± 1.53 | 13.08 ± 0.21 | 3.06 ± 0.02  | 0.15               | 5.60 ± 0.56     | --                    | --      |
| D13     | 7.71 ± 0.03 | 49.59 ± 1.10 | 11.75 ± 0.76 | 2.84 ± 0.18  | 0.18               | 4.85 ± 0.59     | --                    | --      |
| D22     | 7.57 ± 0.04 | 46.72 ± 0.99 | 7.01 ± 0.38  | 3.13 ± 0.06  | 0.14               | 4.59 ± 0.46     | 5.22 ± 0.05           | 59.50 ± 0.06 |
| GO-50   | 7.61 ± 0.03 | 59.52 ± 5.18 | 12.69 ± 0.36 | 3.05 ± 0.10  | 0.15               | 5.29 ± 0.68     | --                    | --      |
| D13     | 7.73 ± 0.03 | 47.52 ± 1.31 | 11.92 ± 0.17 | 3.00 ± 0.09  | 0.19               | 5.35 ± 0.51     | --                    | --      |
| D22     | 7.55 ± 0.05 | 44.50 ± 3.69 | 7.83 ± 1.05  | 3.13 ± 0.03  | 0.14               | 5.21 ± 0.38     | 5.01 ± 0.12           | 61.74 ± 1.66 |
| GO-100  | 7.60 ± 0.01 | 62.40 ± 2.79 | 12.73 ± 0.25 | 3.04 ± 0.06  | 0.15               | 5.71 ± 0.67     | --                    | --      |
| D13     | 7.76 ± 0.09 | 45.61 ± 1.20 | 10.38 ± 1.00 | 2.70 ± 0.02  | 0.19               | 5.51 ± 0.41     | --                    | --      |
| D22     | 7.51 ± 0.01 | 43.37 ± 3.48 | 7.21 ± 0.14  | 3.10 ± 0.12  | 0.12               | 4.29 ± 0.37     | 5.05 ± 0.16           | 62.22 ± 2.19 |
| GO-500  | 7.54 ± 0.01 | 58.19 ± 5.66 | 12.67 ± 0.22 | 3.02 ± 0.09  | 0.13               | 6.32 ± 0.66     | --                    | --      |
| D13     | 7.81 ± 0.02 | 47.48 ± 3.92 | 8.75 ± 0.45  | 2.75 ± 0.04  | 0.21               | 5.05 ± 0.40     | --                    | --      |
| D22     | 7.59 ± 0.06 | 43.25 ± 2.18 | 6.83 ± 0.10  | 2.96 ± 0.05  | 0.14               | 5.03 ± 0.32     | 5.28 ± 0.03           | 60.97 ± 1.25 |

### 3.2. Effects of GO addition on microbial community dynamics

#### 3.2.1. Changes of bacterial community

The microbial diversity decreased after AD, and there existed little influence on the changes of microbial diversity after GO addition even at 500 mg/L (Table S3 and Fig. S1). *Firmicutes* and *Bacteroidetes* dominated throughout the AD, and the abundance of *Firmicutes* reached to 91% in the manure, while *Firmicutes* and *Bacteroidetes* accounted for 57% and 32% in the seed sludge, respectively. All the treatments showed the same trends that *Firmicutes* increased, while *Bacteroidetes* decreased along with AD between D5 and D22 (Fig. S2). There was no much difference at the phylum level between treatments. AD was carried out by different groups of microorganisms involved in hydrolysis, acidogenesis, acetogenesis and methanogenesis, and they were not separated steps but unified processes. In order to find out which step GO addition impacted, the microbial community was further analyzed at genus level. The genus of *unclassified Ruminococcaceae* (10–25%), *Clostridium sensu stricto* (10–19%), *Clostridium III* (2–10%) and *unclassified Bacteroidales* (6.25–23.28%) were famous for the...
Pelotomaculum belonged to the phylum of itself but accelerate the propionate-degradation rate (Chen and Dong, 2005). Besides, (2) interestingly, there existed considerable abundance of Proteiniphilum for the syntrophic propionate-degrading that has not been known. In-0.01%. These indicated that there existed some new species accounting the AD and were reduced to below 1% after AD. The genus tipes (6%) which dominated in swine manure could not well adapt to lower abundance (< 0.01%). The genus trophobacter and (Narihiro et al., 2012), its syntrophic bacterium like teria and methanogens, because of its highest energetics di As for the central intermediate, propionate, which can be converted to enhancement of VFAs accumulation removal through the GO addition. addition was generally higher than the Control. This well explained the (Fig. 2), and interestingly, the abundance of Syntrophomonas in the GO addition was generally higher than the Control. This well explained the central intermediate, propionate, which can be converted to VFAs accumulation through the GO addition. As for the central intermediate, propionate, which can be converted to methane and acetate only by the concerted action of syntrophic bacteria and methanogens, because of its highest energetics difficulties (Narihiro et al., 2012), its syntrophic bacterium like Smithella, Syn- ticiphitobacter and Pelotomaculum etc., were not detected or at much lower abundance (< 0.01%). The genus Smithella and Syntrophobacter belonged to the phylum of Proteobacteria whose abundance was below 1%. Pelotomaculum belonged to Firmicutes, but its abundance was below 0.01%. These indicated that there existed some new species accounting for the syntrophic propionate-degrading that has not been known. In-0.01%. These indicated that there existed some new species accounting for the syntrophic propionate-degrading that has not been known. Interestingly, there existed considerable abundance of Proteiniphilum (2~5%) which could not consume propionate of synthesize methane itself but accelerate the propionate-degradation rate (Chen and Dong, 2005). Besides, Streptococcus (32%), Lactobacillus (15%), and Atopos-tipes (6%) which dominated in swine manure could not well adapt to the AD and were reduced to below 1% after AD. The genus Streptococcus hydrolysis and acidogenesis. They were dominant throughout the AD, which indicated that the hydrolysis and acidogenesis happened all the time. This was associated with the co-occurrence of readily, poorly and non-biodegradable organics in swine manure. The genus Alkalifex existed in high abundance at D5 and D13, and it utilized several carbohydres, in particular those that may result from the hydrolysis of cellulose, hemicelluloses and other natural polysaccharides (Zhilina et al., 2004). The main fermentation products were propionate and acetate, which contributed to the propionate accumulation. Acetate, H2 and CO2 etc., could be directly used by methanogens for the biogas production, while VFAs like propionate, butyrate needed further degradation by the syntrophic bacterium. In this study, the dominant syntrophic bacteria was Syntrophomonas which degraded butyrate in co-culture with methanogens or hydrogen-utilizing sulfate-reducing bacteria, and the genus reached the maximum at D13 (2~4%). This was in accordance with the rapid removal of VFAs accumulation (Fig. 2), and interestingly, the abundance of Syntrophomonas in the GO addition was generally higher than the Control. This well explained the enhancement of VFAs accumulation removal through the GO addition. As for the central intermediate, propionate, which can be converted to methane and acetate only by the concerted action of syntrophic bacteria and methanogens, because of its highest energetics difficulties (Narihiro et al., 2012), its syntrophic bacterium like Smithella, Syn- trophobacter and Pelotomaculum etc., were not detected or at much lower abundance (< 0.01%). The genus Smithella and Syntrophobacter belonged to the phylum of Proteobacteria whose abundance was below 1%. Pelotomaculum belonged to Firmicutes, but its abundance was below 0.01%. These indicated that there existed some new species accounting for the syntrophic propionate-degrading that has not been known. Interestingly, there existed considerable abundance of Proteiniphilum (2~5%) which could not consume propionate of synthesize methane itself but accelerate the propionate-degradation rate (Chen and Dong, 2005). Besides, Streptococcus (32%), Lactobacillus (15%), and Atopos-tipes (6%) which dominated in swine manure could not well adapt to the AD and were reduced to below 1% after AD. The genus Streptococcus...
Methanospirillum was the key methanogens for the removal of VFAs accumulation, and the symbiosis of Syntrophobacter and Syntrophomonas with Methanospirillum was widely observed for the propionate or butyrate degradation for methane production (Chen et al., 2005; Zhang et al., 2004). Besides, in the presence of high amounts of ammonia (above 2.8–3 g/L, Table 1), acetate sink for the methane production via the activity of syntrophic acetate oxidizing bacteria coupled with hydrogen utilizing methanogen like Methanobrevibacter and Methanospirillum should not be overlooked (Fotidis et al., 2014; Schnirer and Nordberg, 2008).

3.2.3. Effects of GO addition

There existed no significant difference between treatments as revealed by the PCA analysis considering the bacterial and archaeal community dynamics (Fig. 3), and this indicated that the reduction of methane production and enhancement of VS reduction may be due to that the GO impacted the microbial activity more than the microbial community composition. In order to elucidate the hypothesis, the key functional gene mcrA was quantified, the abundance of mcrA increased along with swine manure AD in all treatments, and it was increased significantly along with GO addition compared to the Control (Fig. S3). The mcrA gene abundance had been demonstrated to be able to well correlate with the activity measurements of methanogenic H₂/CO₂ enriched anaerobic biomass (Venkiteshwaran et al., 2017), thus, this confirmed that the GO addition enhanced the key functional enzymes not changing the microbial community composition.

There was no much difference of mcrA gene abundance at D5 between treatments, but the mcrA gene abundance changed significantly at D13, which could well explain the main methane production due to GO addition happened at the second peak. Interestingly, the methane production correlated significantly negatively with mcrA gene abundance, while GO-100 was a special point. As for the special point of GO-100, this could be explained by the three action mode between GO and microbes including cutting, wrapping and trapping (Palmieri et al., 2017). Cutting and trapping did not kill the methanogens, but enhanced the DIET between syntrophic bacterium and methanogen, while wrapping could cause the membrane and oxidative stress which further killed or inhibited the methanogens, and it was speculated that GO showed the cutting effects at the concentration of 5 mg/L and 50 mg/L, while it was wrapping and trapping effects at 100 mg/L and 500 mg/L, respectively. At the end of the swine manure AD, mcrA gene abundance still increased, and this could be due to that methanogens could stand the poor nutrition environments compared to other microbes (Garcia et al., 2000).

3.3. Effects of GO addition on ARGs, MRGs, VFAs and intI1 reduction

3.3.1. Changes of total ARGs

There existed high abundance of ARGs, MRGs, VFAs and intI1 in both swine manure and seed (Fig. S5), and the total ARGs abundance in the seed (3.01 × 10⁻³) was higher than the swine manure (1.91 × 10⁻³). This was because the seed sludge were collected from the AD system treating the swine manure long-termy, but the MRGs and intI1 in the seed (8.12 × 10⁻⁴ and 1.80 × 10⁻⁵) were much lower than swine manure (1.89 × 10⁻⁴ and 1.04 × 10⁻⁵).

As shown in Fig. 4, AD could effectively realize the ARGs, MRGs, VFAs and intI1 gene abundance reduction. Total ARGs abundance changed little at D5 in all treatments, that is the hydrolysis and acidogenesis, but it decreased significantly as AD progressed to the acetogenesis and methanogenesis, which indicated that ARGs reduction was realized mainly through the acetogenesis and methanogenesis phase.

Total ARGs abundance was reduced by 33.7% for the Control, while GO-50 and GO-100 deteriorated the ARGs reduction, and the total ARGs abundance reduction was 3.7% and 23.9%, respectively. GO-5 influenced the reduction little (32.8%), but GO-500 enhanced the reduction to 40.2%. The deterioration of ARGs abundance reduction could be attributed to the enhancement of HGT by GO, which could be reflected by the higher intI1 abundance at GO-50 and GO-100 (Fig. 4). Castrillón et al. (2015) reported that GO would make cells produce oxidative stress and the oxidative stress might change cell membrane fluidity, and then promote the conjugative transfer of resistance genes. It has been demonstrated that when GO concentration reached 50 mg/L, the HGT frequency was the largest and increased by more than 10 times compared with the normal, but at higher GO concentrations (> 50 mg/L), the frequency gradually decreased (Guo and Zhang, 2017). Considering the complexity of swine manure AD system compared to the model experiment, the cutoff value may be 100 mg/L. The enhancement of ARGs reduction at GO-500 could be explained by the absorbance and inhibitory proliferation of ARGs by GO, especially the extracellular ARGs (Yu et al., 2016; Zou et al., 2016). Thus, there existed a balance between the ARGs proliferation by the enhancement of HGT and ARGs destruction as for the ARGs changes caused by GO. However, GO addition had little influence on the MRGs reduction, but the intI1 reduction was deteriorated especially for GO-50.

3.3.2. Changes of each ARG

The abundance of some selected ARGs including tetM, tetX, blaCTX-M, ermF and sulII in the seed sludge was generally higher than swine manure, and the most abundant ARG was ermB (1.56 × 10⁻⁴) and tetM (1.32 × 10⁻⁴) for swine manure and the seed sludge, respectively. Except pcoA, other MRGs abundance including mera, arsc and caza in the seed sludge was also generally higher than swine manure. The abundance of VFAs was comparable, while intI1 was higher in swine manure.

Although total ARGs abundance was reduced after AD, the fate of each ARG varied significantly (Table 2). Results showed that ermB, tetM, tetG, sulII, blaTEM and mcr-1 were reduced but tetX, sulII, ermF and blaCTX-M were enriched after AD. The dominant ARGs like ermB and tetM in swine manure could be effectively reduced by 63.85% and 36.88%, respectively, but other ARGs should be paid more attention. Besides, AD could effectively reduce the mcr-1 gene abundance by 90.3–95.8%. As for the MRGs, only mera was enriched, which indicated that the selection from Hg should be paid more attention. VFs including aacI and eaeA were both largely reduced after AD.

Interestingly, the dynamics of each ARG varied significantly. For instance, sulII was enriched the most at the hydrolysis and acidogenesis, while tetX was enriched the most at the acetogenesis and methanogenesis, and the reduction of ermB was realized at the acidogenesis with little changed as AD progressed. Although tetG and blaTEM were enriched at acidogenesis, the reduction was realized at the methanogenesis phase, and tetM changed little at acidogenesis while it was reduced after the acetogenesis phase. Also, it seemed like that the tendency of enrichment or reduction was determined at the hydrolysis and acidogenesis phase. There was no such variety for the changes of intI1, MRGs and VFAs, and the reduction was realized at the hydrolysis and acidogenesis phase.

3.4. Network analysis interpreting the discrepant dynamics of each ARG

It was speculated that the discrepant dynamics of each ARG was attributed to their changes of host bacteria, and changes of microbial community contributed most to the changes of ARGs as previously supported (Forsberg et al., 2014; Su et al., 2015; Zhang et al., 2016b). Results of Mantel test and Procrustes analysis showed that there existed significant correlation between dynamics of ARGs and changes of microbial community (R = 0.6956, p = 0.0001), and microbial community changes accounted for 63.8% of variance of ARGs profiles (Fig. S6).

Network analysis showing the potential host bacteria was conducted (Fig. 5), and the potential host bacteria of each ARG were listed (Table S6). The network analysis was divided into three modules: the first modules contained blaCTX-M, tetX and their potential host bacteria, which was gradually enriched as AD progressed; the second module
included \textit{ermB}, \textit{tetM} and \textit{mcr-1} which were reduced a lot after AD, and the reduction was realized at the hydrolysis and acidogenesis phase; the third module contained those that was enriched at hydrolysis and acidogenesis phase and then the enrichment was reduced as AD progressed like \textit{bla}\textsubscript{TEM}, \textit{sulI}, \textit{sulII} and \textit{tetG}. Some ARGs were even reduced to below the initial level like \textit{bla}\textsubscript{TEM}, \textit{sulI} and \textit{tetG}, while \textit{ermF} was an exception whose enrichment changed little as AD progressed.

The potential host bacteria could well explain the discrepant changes of each gene during swine manure AD. For instance, the increasing enrichment of \textit{tetX} and \textit{bla}\textsubscript{CTX-M} could be associated with \textit{Methanospirillum} which also increased along with AD, and \textit{Lactobacillus}, \textit{Streptococcus} and \textit{Atopostipes} were all depicted to be the common host bacteria of \textit{ermB}, \textit{tetM} and \textit{mcr-1}, and these bacteria mainly originated from swine manure and were reduced significantly after AD. These potential host bacteria may be not a real ARB yet, but the link was just functional, but it would make sense considering that the functional link make it have the best chance of becoming ARB (Zhang et al., 2016a).

Besides, the identified host bacteria of \textit{mcr-1} (\textit{Escherichia}) was also distinguished through network analysis, and \textit{Escherichia} was also identified to be the host of \textit{ermB} and \textit{tetM}, while its abundance was reduced to below detection after AD, which could also well explain the changes of \textit{ermB}, \textit{tetM} and \textit{mcr-1}.

Not only the changes of its potential host influenced the ARGs dynamics, but also other factors like environmental variables (EVs), MRGs and \textit{mcrA} (the potential for biomethane production). Interestingly, the significant correlation between \textit{intI1} and any ARG considered was not observed, which indicated that AD could effectively reduce the role of HGT on ARGs. However, there existed significantly positive correlation between \textit{mcrA} and \textit{tetX} and \textit{bla}\textsubscript{CTX-M}, while significantly negative correlation between \textit{mcrA} and other ARGs including \textit{ermB}, \textit{tetM}, \textit{mcr-1}, \textit{tetG} and \textit{sulI}, which indicated that the increasing abundance of archaeal community for biogas production in fluenced ARGs abundance and the influence varied from ARG patterns. The most important MRGs concerning the ARGs profiles was \textit{pcOA}, and it has significantly positive correlation with six ARGs, while the effects of EVs like TCOD, SCOD, proteins and polysaccharides on specific ARG dynamics could also not

| Sample | \textit{ermB}\textsubscript{TEM} | \textit{ermB}\textsubscript{TEM} | \textit{ermB}\textsubscript{TEM} | \textit{ermB}\textsubscript{TEM} | \textit{ermB}\textsubscript{TEM} | \textit{ermB}\textsubscript{TEM} | \textit{ermB}\textsubscript{TEM} | \textit{ermB}\textsubscript{TEM} | \textit{ermB}\textsubscript{TEM} |
|--------|------------------|------------------|------------------|------------------|------------------|------------------|------------------|------------------|------------------|
| Control | 1.00 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 |
| D0 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 |
| GO-5 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 |
| GO-10 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 |
| GO-50 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 |
| GO-100 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 |
| GO-500 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 | 0.99 |

* Values were achieved by the temporary abundance divided by the abundance on D0. If the value > 1, it meant that the gene was enriched. Or, it meant that the gene was reduced, and (1-minus the value) was the removal efficiency. The redder, the more it was reduced. The greener, the more it was enriched.

Fig. 4. The effects of GO addition on the changes of ARGs, MRGs, VFs and \textit{intI1}.
be overlooked.

3.5. SEM ranking the factors influencing the ARGs dynamics

SEM could separate multiple pathways of influence and view them as a system, and is useful to explore the complex networks of relationships found in ecosystems (Hu et al., 2016). The a priori and theoretical assumptions were established before SEM, and the matrix values derived from the pairwise correlation between ARGs and the influencing factors were imported into the model for the construction of SEM. The overall goodness-of-fit of model fits was indicated by a non-significant chi-square test (P > 0.05), high goodness-of-fit index (GFI > 0.90), high adjust GFI (AGFI > 0.90) and low root square mean errors of approximation (RMSEA < 0.05) (Hu et al., 2016).

SEM results shown in Fig. 6 indicated that the standardized effects of GO and mcrA on the ARGs was negative, that is, the increasing of GO and mcrA abundance could avail the ARGs reduction, and GO indirectly affected the ARGs reduction while mcrA influenced the ARGs directly. While other factors like EVs, Biomass, microbial community (MC), MRGs and MGEs affected the ARGs positively, and EVs contributed the most to the ARGs in terms of standardized total effects followed by MRGs and MC, while the most direct effects come from the co-selection of heavy metals reflected by MRGs and the most indirect effects also come from EVs. GO could negatively influence the EVs, VFs and MC and then ARGs were negatively affected, while GO could directly and positively affected the MGEs (R = 0.22), which was in accordance with the hypothesis that GO could boost the HGT of ARGs.
4. Conclusions

Results in this study showed that the biological methane potential of swine manure was reduced due to GO addition. There existed limited influence on changes of bacterial and archaeal community composition by GO. AD could realize the reduction of ARGs abundance and higher concentration of GO could further enhance the reduction, while 50–100 mg/L of GO deteriorated ARGs reduction. Network analysis showed that microbial community variation could well explain the changes of each ARG, while SEM indicated that EVs contributed most to the changes of ARGs indirectly, GO influenced the ARGs changes negatively and MRGs influenced the most directly.

Conflict of interest

The authors declare no conflict of interest.

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Appendix A. Supplementary data

Supplementary data associated with this article can be found, in the online version, at http://dx.doi.org/10.1016/j.biotech.2017.08.217.

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