Anaerobic Codigestion of Municipal Wastewater Treatment Plant Sludge with Food Waste: A Case Study

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1. Introduction

Sludge production from municipal wastewater treatment plants (MWTPs) is expected to continue to increase with the increasing number of treatment plants being constructed or upgraded due to the growing population connected to the sewage networks of Australia. The disposal of sludge generated at the MWTPs is a problem of increasing importance, representing up to 50% of the current operating costs of a wastewater treatment plant [1]. In Australia, MWTPs produce approximately 360,000 dry tonnes of stabilised sewage sludge to dispose of which costs about $100 M per year [2, 3]. Hence, water authorities operating these plants in Australia have been actively investigating alternative sustainable and economic sludge management pathways [4]. Although different disposal routes are possible, anaerobic digestion (AD) appears to be the most promising sludge management alternative due to its ability to generate bioenergy by the reduction of the sludge volumes to be disposed of [5–8].

Sewage sludge (SS) contains a high percentage of organic matter (60–70% of the dry matter) and nutrients, and typically comprises primary sludge (PS) and waste activated sludge (WAS) [9, 10]. However, since WAS has low biodegradability; the AD of WAS has low efficiency from both processing and economic standpoints [11]. One of the different strategies to enhance the performance of AD is the codigestion of sludge with other organic wastes as it increases biodegradable organic matter and provides a feedstock with an optimum C/N ratio [1, 4–12]. Among the factors that limit the codigestion are the selection and type of new organic wastes and the transportation cost of cosubstrates to the MWTPs [9–13].

Food wastes (FW) from different sources, for example, residential and commercial, are being produced at an increasing rate due to the growing population and rising living standards [8]. FW is available all year round and accounts for a significant proportion of municipal solid waste. In Victoria Australia, FW contributes 35.6% of the total municipal solid waste.
wastes when source separated, usually referred to as organic fraction of municipal solid waste [14]. However, due to its high biodegradability and volatile acids, AD as a single substrate may encounter various potential inhibitors including accumulation of volatile fatty acids’ (VFAs) accumulations [15]. Therefore, these FW could be beneficial in anaerobic codigestion for high energy recovery as well as solid waste reduction.

The application of anaerobic codigestion for the treatment of SS has been receiving growing attention for improving the biogas yield, solid destruction, and the production of digestate of a suitable quality to use as a fertilizer [16]. Full-scale applications of anaerobic codigestion of SS with FW can become an environmentally sound renewable energy source by creating opportunities to recover the energy potential from these very low or zero cost FW and obtain the benefit of high organic matter to increase the methane yield. This will result in significantly less biosolids’ disposal and a reduction in municipal solid wastes as well as the operating costs of the plants.

Many authors have reported increased biogas yields from the codigestion of SS with different types of food and/or food processing wastes. For example, codigestion of sludge mix with fat, oil and grease (FOG) from a meat processing plant (46% VS added to the feed) increased the methane yield by 60% [17]. Similarly, methane yield was 2.6 times higher when SS was codigested with oil and grease (48% of total VS load) from restaurants [18]. Under mesophilic conditions, the highest methane production rate was observed when FW was mixed in the range of 30–40% VS with SS [16, 19].

An MWTP in Melbourne, Australia, produces about 3600 kg solids/day of which 627 kg is WAS and the remaid is PS. Since, this plant is in the progress of upgrading the existing old anaerobic digestion reactor, the management was interested in assessing the feasibility of the codigestion of sludge with two streams of wastes, namely, grease trap waste collected from food businesses in the area, referred to in this study as FOG, and waste from a food products manufacturing factory at a small ratio. The MWTP interest is to maximise methane yield, enhance solids removal, and maintain or improve biosolids quality.

The aim of this study, therefore, was to assess the effect of the sludge: waste ratio on the biogas yield and the quality of digestate and supernatant nutrient produced under semicontinuous conditions and, to monitor process performance and stability during codigestion experiments.

2. Materials and Methods

2.1. Characteristics of Substrates and Inoculum. The sludge feedstocks used in this study were thickened PS and WAS collected from Melton Recycled Water Treatment Plant, Victoria, Australia. The PS and WAS were mixed at a ratio consistent with their flow rates such that the final mixed SS’s total solids (TS) is 4%. The raw PS and WAS were collected several times while running the experiment and each time they were characterised and mixed as described. The characteristics of different SS samples are reported in Table 1. The SS was stored in a sealed plastic container at 4°C.

The FW used were (i) thickened grease trap, referred to in this paper as GT, obtained from a commercial business that collects FOG from restaurants and food businesses in the western area of Melbourne and (ii) wastes from processed food products manufacturing factory denoted as PF. These FW are mostly comprised cooking oil, butter, cheese, meat, bread, meat fat and bones, mayonnaise, salad dressing, and so forth. The food wastes were collected regularly, homogenised using a high speed homogeniser, and then characterised by the parameters shown in Table 1. The TS of the substrates (SS, PF, and GT) was adjusted such that the AD reactors received a feedstock of consistent TS and chemical oxygen demand (COD) concentration throughout the duration of the experiments. The inoculum used in this experiment was collected from the mesophilic anaerobic digester at Melton wastewater treatment plant. The characteristics of the feedstocks (SS, PF, and GT) and the inoculum are shown in Table 1.

2.2. Batch Experiments. Batch tests were performed to determine the biochemical methane potential (BMP) of the individual substrates (SS, PF, and GT) and mixtures of the SS and FW (a mixture of PF and GT at 50:50 w/w) at different ratios. The experimental design is shown in Table 2. All the BMP tests were performed in 500 mL glass bottles at 37°C according to the guideline of Angelidaki et al. [20]. Each reactor contained 4000 mg VS with VS in substrate:VS in inoculum ratio of 0.25. In addition, two reactors received only inoculum as a control. The headspace of the bottles was flushed with nitrogen gas for 2 minutes and the bottles were closed with a rubber Suba-Seal. All batch tests were performed in duplicate. The bottles were kept at 37 ± 1°C in an incubated shaker at a constant rotational speed of 100 rpm. The volume of biogas produced was measured using a water displacement unit and the biogas composition was monitored using gas chromatography. The volume of biogas (or methane) from the control was subtracted from the volume of biogas (or methane) produced in each reactor to obtain the net production of biogas or BMP from the substrates fed into the reactor.

2.3. Semicontinuous Experiments. SS was mixed with the wastes from the processed food products manufacturing and/or FOG, at the designated ratio of sludge to waste (SS:PF, SS:GT). The experiments were performed in 500 mL glass reactors, designed to allow feeding and nitrogen flushing simultaneously, at 37 ± 1°C in an incubated shaker at a constant rotational speed of 100 rpm. The reactors received the substrates at a concentration of 4% TS and operated at an organic loading rate of 2.0 kg TS/m²-d. The experiment comprised duplicate reactors for each condition. The reactors were operated at a sludge retention time of 20 days (equivalent to hydraulic retention time, HRT, in this case) and were fed and wasted once a day. The biogas was collected before feeding the reactors every day. The biogas measurement, feeding, and wasting were done within a 15 min window out of the incubator. The reactors were monitored weekly for biogas quality, and the wastage was analysed every ten days for pH, TS, and VS, total COD (tCOD), and soluble COD (sCOD). The feedstock to the reactors was prepared from different substrates at the ratios shown in Table 2.
Table 1: Characteristics of substrates and inoculum.

| Parameters | Unit | SS 1st sample | SS 2nd sample | PF 1st sample | PF 2nd sample | GT 1st sample | GT 2nd sample | Inoculum 1st sample | Inoculum 2nd sample |
|------------|------|---------------|---------------|---------------|---------------|---------------|---------------|---------------------|---------------------|
| TS         | %    | 3.7 ± 0.1     | 18.77 ± 0.8   | 7 ± 0.2       | 26.1 ± 0.2    | 1.85 ± 0.2    |                |                     |                     |
| VS         | %    | 3.13 ± 0.11   | 18.06 ± 0.7   | 6.8 ± 0.16    | 25.55 ± 0.2   | 1.32 ± 0.12   |                |                     |                     |
| tCOD       | g/L  | 53.73 ± 8.2   | 239.1 ± 0.91  | 405.3 ± 50    | 475.5 ± 10    | 12.9 ± 2.8    |                |                     |                     |
| sCOD       | g/L  | 3.95 ± 0.6    | 3.42 ± 0.04   | 2.98 ± 0.9    | 3.8 ± 0.7     | 1.4 ± 0.7     |                |                     |                     |
| Total N    | g/L  | 2.6 ± 0.1     | 3.55 ± 0.15   | 3.5 ± 0.2     | 3.54 ± 0.2    | 1.86 ± 0.003  |                |                     |                     |
| Ammonium   | g/L  | 0.11 ± 0.01   | 0.11 ± 0.003  | 0.14 ± 0.01   | 0.26 ± 0.007  | 0.48 ± 0.007  |                |                     |                     |
| Total PO$_4^{3-}$ | g/L | 1.5 ± 0.05 | 1.1 ± 0.04 | 2.56 ± 0.06 | 2.58 ± 0.1 | 0.9 ± 0.3 | | | |
| Total VA   | g acetic acid/L | 0.6 ± 0.01 | 1.98 ± 0.15 | 1.9 ± 0.2 | 2.03 ± 0.2 | 0.17 ± 0.013 | | | |
| Alkalinity | g CaCO$_3$/L | 2.7 ± 0.001 | 1.42 ± 0.001 | 1.3 ± 0.001 | 2.1 ± 0.01 | 4.1 ± 0.002 | | | |
| pH         |      | 6.36 ± 0.09   | 5.54 ± 0.01   | 5.0 ± 0.6     | 6 ± 0.3       | 7.55 ± 0.13   |                |                     |                     |

Table 2: Composition of the feedstocks used in the BMP and semicontinuous tests.

| Experiment type | Substrates in feedstock | Substrates Composition (w/w) | Nomenclature |
|-----------------|-------------------------|------------------------------|--------------|
| Batch Single    | SS                      | 100                          | 100% SS      |
|                 | PF                      | 100                          | 100% PF      |
|                 | GT                      | 100                          | 100% GT      |
| Batch Two       | SS : PF                 | 99 : 01                      | 1% PF        |
|                 | SS : PF                 | 98 : 02                      | 2% PF        |
|                 | SS : PF                 | 90 : 10                      | 10% PF       |
|                 | SS : PF                 | 75 : 25                      | 25% PF       |
|                 | SS : GT                 | 99 : 01                      | 1% GT        |
|                 | SS : GT                 | 98 : 02                      | 2% GT        |
|                 | SS : GT                 | 90 : 10                      | 10% GT       |
|                 | SS : GT                 | 75 : 25                      | 25% GT       |
|                 | SS : GT                 | 50 : 50                      | 50% GT       |
| Batch Three     | SS : PF : GT            | 95 : 25 : 2.5                | 5% FW$^e$    |
|                 | SS : PF : GT            | 80 : 10 : 10                 | 20% FW$^e$   |
|                 | SS : PF : GT            | 50 : 25 : 25                 | 50% FW$^e$   |
|                 | SS : PF : GT            | 33.3 : 33.3 : 33.3           | 66.67% FW$^e$ |
| Semicontinuous Single | SS               | 100                          | 100% SS      |
| Semicontinuous Two | SS : PF                | 99 : 1                       | 1% PF        |
|                 | SS : PF                | 98 : 2                       | 2% PF        |
|                 | SS : GT                | 99 : 1                       | 1% GT        |
|                 | SS : GT                | 98 : 2                       | 2% GT        |
| Semicontinuous Three | SS : PF : GT         | 95 : 25 : 2.5                | 5% FW$^e$    |

FW = mixture of PF and GT at ratio 50 : 50 (w/w).

2.4. Analytical Methods. TS and VS were measured by gravimetric analysis according to the Standard Methods 2540B and 2540E, respectively [21]. tCOD and sCOD were measured according to HACH method 8000. The total phosphorus (TP), total nitrogen (TN), ammonium, and volatile acids (VAs) were measured by colorimetric techniques using a HACH spectrophotometer (Model DR/4000 U) according to the methods 10127, 10072, 10031, and 8196, respectively. The samples were centrifuged (Eppendorf 5702, Germany) at 4.4 rpm for 15 mins and then filtered through 0.45 μm filter paper (mixed cellulose esters membrane filter, Advantec, Japan) to measure the soluble constituents. The measurement of pH was carried out using a calibrated pH meter (ThermoOrion, Model 550A) and alkalinity was measured by the APHA method 2320B.

The volume of biogas was normalised to standard conditions compromising dry gas, standard temperature, and pressure (0°C and 1 bar) according to the method described by Strömberg et al. [22] and the results are presented as norm-litre (L$_N$). The headspace was corrected for methane (CH$_4$) and carbon dioxide (CO$_2$) to 100% according to VDI 4630 (2006) [23]. The composition of the biogas was analysed according to APHA method 2720C using gas chromatography (Varian 450-GC, Varian Australia Pty Ltd., Netherlands).
equipped with a packed column (GS-CarbonPLOT I13–3132, 1.5 microns, 30 m × 0.320 mm, stainless steel, Agilent Technologies Inc., Australia) and a thermal conductivity detector. The carrier gas used was helium at a flow rate of 28 mL/min. The temperatures of the column, detector, and injector were 70°C, 200°C, and 100°C, respectively. The biogas was collected and manually injected using a 50 mL FORTUNA® Optima glass syringe (Poulten & Graf, Germany). Calibration was done using three points and five levels of CH₄, CO₂, and nitrogen (BOC, Australia). Screening of the metals in the digestate samples was tested for sodium (Na) to cerium (Ce) by a commercial laboratory (ALS Environmental Division: Water Research Group).

2.5. Statistical Analysis. Predictions of the optimum mixture ratio for two and three substrates from batch tests were obtained using MATLAB R2013b. Furthermore, a predictive model for optimum FW incorporation was prepared with surface and contour plots. To determine the significance of difference in cumulative methane yields over the digestion period, each set of codigestion feedstock was statistically analysed with 100% SS using one-way analysis of variance (ANOVA) at α = 0.05 in MATLAB R2013b.

3. Results and Discussion

3.1. Batch Experiments. Batch experiments were carried out to investigate the optimum ratio of FW for incorporation in SS. The effects of two substrates and three substrates were also investigated at different mixture ratios. The cumulative methane yields and the daily biogas yields during the anaerobic codigestion are shown in Figures 1(a)–1(d) and 1(e)–1(h), respectively. The BMP tests continued for 46 days until little or no biogas production was observed. The results presented are the net biogas and methane yield from the feedstock after subtracting the control yield.

According to Figure 1(a), the BMP of 100% SS was 192 ± 12.3 mL NCH₄/g VS added, whereas the processed food wastes, 100% PF and 100% GT, had a BMP of 466.2 ± 0.73 and 408.7 ± 6.6 mL NCH₄/g VS added, respectively, which is 1.42 and 1.12 times higher than 100% SS alone. For 100% SS, the biogas production started after 2 days and reached the first peak at day 8 with a rate of 21.5 mL N biogas/g VS added/d (Figure 2(e)). The second peak occurred at day 17 with a peak value of 46.1 mL N biogas/g VS added/d and after 21 days slowly decreased. Both food wastes started biogas production after day one and reached the first peak at day 17 with daily biogas yields of 54.3 and 45.4 mL N biogas/g VS added/d, respectively, for PF and GT. The second peak values were 56.3 and 25.3 mL N biogas/g VS added/d for PF and GT, respectively, at days 28 and 36. The technical digestion time, that is, T80–90 (the time for 80–90% of the maximum biogas production), was calculated to be between 20 and 27, 31 and 35, and 37 and 40 days for SS, PF, and GT, respectively. The technical digestion time can be used as a HRT for continuous anaerobic digestion for these substrates [24].

The codigestion of SS with PF enhanced the BMP from 199.6 ± 20.6 to 616.8 ± 30.2 mL N CH₄/g VS added for PF fractions of 1% to 50%, that is, 4% to 287% increase in methane yield compared with 100% SS alone (Figure 1(b)). However, with 1% PF to 10% PF incorporation, a lag phase of 2 days was observed, and the 25–50% PF mixture with SS immediately started biogas production. For 1% PF to 25% PF, a single peak in daily biogas yield was observed at day 17 with peak values of 36.4 ± 0, 43.9 ± 7.1, 67.2 ± 4, and 86.7 ± 2.7 mL N biogas/g VS added/d, respectively, for 1% PF, 2% PF, 10% PF, and 25% PF (Figure 1(f)). A rising trend was observed in peak value with increasing PF ratio. The production of biogas was decreased after 20 days and almost ceased after 36 days. However, for 50% PF, an inhibition in biogas production was obtained with two peaks. At day 15, the first peak of 42.4 ± 0 mL N biogas/g VS added/d with easily degradable organic materials was noted and, at day 28, a small second peak of 29.6 ± 8.7 mL N biogas/g VS added/d with slow degradation were observed. T80–90 was calculated between 20 and 26, 21 and 27, 21 and 27, and 27 and 33 days, respectively, for 1–50% PF incorporation.

SS mixed with GT enhanced the BMP of SS from 200 ± 2.6 to 561.3 ± 16.9 mL N CH₄/g VS added/d, that is, there was a 5% to 260% increase in methane production, by adding up to 50% GT during codigestion (Figure 1(c)). It was observed that increasing the GT fraction in the feedstock from 1% to 50% caused an increase in BMP up to 17 days and it started decreasing until completely ceased at around 36 days (Figure 1(g)). The peak values were 47.3 ± 2.4, 47.7 ± 3.1, 42.3 ± 1.8, 75.4 ± 13.8, and 89.8 ± 22.8 mL N biogas/g VS added/d, respectively, for 1% GT, 2% GT, 10% GT, 25% GT, and 50% GT. No inhibition was observed with T80–90 between 20 and 26, 20.5 and 26, 21 and 28, 21 and 28, and 25 and 32 days, respectively, for 1–50% GT incorporation.

For three substrates, biogas production improved with up to 50% FW addition (632.8 ± 10.1 mL N CH₄/g VS added) and decreased for the mixture ratio of 66.7% FW (603.3 ± 6.7 mL N CH₄/g VS added) (Figure 1(d)). An early peak at day 8 was observed for 5% FW with a peak value of 40.7 ± 7.9 mL N biogas/g VS added/d. It was, however, observed at day 17 for 20–66.7% FW with peak values of 63.6 ± 0.5, 74.8 ± 6.8 and 73.3 ± 0 mL N biogas/g VS added/d, respectively (Figure 1(h)). T80–90 was calculated between 24 and 35, 25 and 32, 24 and 30, and 26 and 37 days, respectively, for 5%–66.7% FW incorporation.

Therefore, the addition of FW with SS decreased the technical digestion time with a single peak. The VAs usually associated with the GT appear to be below inhibition up to 50%. However, the inhibition effect at 50% PF indicates that there is NH₄ that reached a threshold (2.1 ± 0.1 g/L). Ammonia which is an important indicator of AD, is produced by the hydrolysis of proteins and urea [25, 26] and accumulates in the AD process [27]. FW was incorporated with SS.

The BMP assay can be utilised to calculate the synergic effect of codigestion as additional methane yield over the weighted average of the individual feedstock’s methane yield [28]. The weighted experimental methane was calculated from single substrate using the following formula

\[
\text{Weighted EMY}_{FW} = \text{EMY}_{100\%SS} + \text{EMY}_{100\%PF} + \text{EMY}_{100\%GT}
\]

\[
\text{Weighted EMY}_{FW} = \text{EMY}_{100\%SS} + \text{EMY}_{100\%PF} + \text{EMY}_{100\%GT} (1)
\]

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Figure 1: Accumulative methane production (a–d) and daily biogas yield (e–f) from batch experiments of single, two, and three substrates.
Weighted EMY\textsubscript{PF} = EMY\textsubscript{100\% SS} * P\textsubscript{100\% SS} + EMY\textsubscript{100\% PF} * P\textsubscript{100\% PF}, \hspace{1 cm} (2)

Weighted EMY\textsubscript{GT} = EMY\textsubscript{100\% SS} * P\textsubscript{100\% SS} + EMY\textsubscript{100\% GT} * P\textsubscript{100\% GT}, \hspace{1 cm} (3)

where weighted EMY\textsubscript{FW}, EMY\textsubscript{PF}, and EMY\textsubscript{GT} represent the weighted average of the experimental methane yield of the substrates FW, PF, and GT, respectively. P\textsubscript{100\% SS}, P\textsubscript{100\% PF}, and P\textsubscript{100\% GT} refer to the percentage composition and EMY\textsubscript{100\% SS}, EMY\textsubscript{100\% PF}, and EMY\textsubscript{100\% GT} are the experimental methane yield for substrates SS, PF, and GT, respectively, in the cosubstrates mixture. According to Li et al. (2013) if the difference (EMY – weighted EMY) was higher than the standard deviation of EMY, synergic effect could be observed [29]. The EMYs of the codigestion substrates during the digestion period were analysed statistically with respect to the EMYs of 100% SS. As Table 3 shows, 1-2% PF and GT did not have very significant synergistic effects; however, increasing the amount of food wastes resulted in a very significant (p < 0.05) increase in methane yield compared to the digestion of SS alone.

A synergic effect was found in almost all of the cases when food wastes were added to SS representing higher biodegradability. This is possibly due to the adjustment in C/N ratios during codigestion [29] compared to the single substrate. The C/N ratio is a good indicator of the efficiency of AD that can be limited by inadequate amount and diversity of waste from a single resource. For example, high carbon content of a sample can cause rapid acidification and methanogenesis will be inhibited by the low pH. The optimum C/N ratio is waste specific over a range from 9 to 30 [31]. The C/N ratio of SS used in this study was 8.16 which is lower than the C/N ratio of PF and GT (17.64 and 15.5, resp.). Incorporating 50% FW in the feedstock with SS increased the C/N ratio of the reactors up to 12-13. Antagonism (probably due to inhibition) was observed for 50% PF. In case of three substrates, 5% FW showed the highest increase in methane yield. Luostarinen et al. (2009) also reported inhibition with the addition of grease trap sludge to SS of more than 50% [17]. However, these inhibitory effects were only deduced from the pattern of methane production and the synergistic effects and will require further investigations.

To investigate the optimum mixture ratio of FW and SS with respect to methane yield, a trend was predicted using MATLAB (Figures 2(a)–2(c)). The $R^2$ correlation values were 0.999, 0.993, and 0.885 for %PF, %GT, and %FW incorporation with SS, respectively, indicating a good fit between experimental and predicted values. The results showed that methane yield obtained maximum values of 614.6, 562, and 651.1 mL\textsubscript{N} CH\textsubscript{4}/g VS\textsubscript{added} when 47% PF, 61.4% GT, and 48% FW were incorporated with SS improving the C/N ratio of 12.5. Figure 3 shows the 3D model of optimum FW incorporation with SS, where %PF and %GT with SS on the x- and y-axis with methane yield on z-axis. The dark red area represents the maximum methane yield region. FW incorporation up to 48% with the mixture of GT and PF according to the dark red region will produce the maximum volume of biogas. Considering SS as the main substrate, batch
Table 3: Synergistic effect evaluation of codigestion of SS with PF, GT, and FW (mixture of PF : GT).

| Substrates ratio<sup>a</sup> | EMY  | SD  | Weighted EMY | Difference | Increase in EMY (%) | p value | Synergistic effect |
|-------------------------------|------|-----|--------------|------------|--------------------|---------|--------------------|
| 1% PF                         | 199.6| 20.6| 195.7        | 3.9        | 2.0                | 0.9310  | Not clear          |
| 2% PF                         | 226.6| 16.3| 198.4        | 28.2       | 14.2               | 0.6106  | Not significant    |
| 10% PF                        | 383.1| 22.9| 220.3        | 162.8      | 73.9               | 0.0462  | Synergistic        |
| 25% PF                        | 537.5| 12.3| 261.3        | 276.2      | 105.7              | 0.0084  | Synergistic        |
| 50% PF                        | 616.8| 30.2| 329.6        | 287.2      | 87.2               | 0.0066  | Synergistic        |
| 1% GT                         | 200.8| 2.6 | 195.1        | 5.16       | 2.7                | 0.9067  | Not clear          |
| 2% GT                         | 230.6| 10.3| 197.3        | 33.32      | 16.9               | 0.5423  | Not significant    |
| 10% GT                        | 317.3| 14.8| 214.5        | 102.8      | 47.9               | 0.0467  | Synergistic        |
| 25% GT                        | 413.2| 16.9| 246.7        | 166.3      | 67.4               | 0.0259  | Synergistic        |
| 50% GT                        | 561.3| 16.9| 300.8        | 260.5      | 86.6               | 0.0081  | Synergistic        |
| 5% FW                         | 433.7| 72.7| 205.2        | 228.5      | 114.4              | 0.0176  | Synergistic        |
| 20% FW                        | 508.9| 70.1| 241.8        | 267.1      | 110.4              | 0.0110  | Synergistic        |
| 50% FW                        | 632.8| 16.1| 315.2        | 317.6      | 100.8              | 0.0038  | Synergistic        |
| 66.67% FW                     | 603.3| 6.7 | 352.4        | 250.9      | 71.2               | 0.0066  | Synergistic        |

EMY: experimental methane yield (mL/g VS added); SD: standard deviation; and weighted EMY: weighted average of experimental methane yield for cosubstrates.

<sup>a</sup>Percentage of food wastes (PF, GT, and FW) mixed with SS.

Figure 3: 3D prediction of optimum FW incorporation: (a) surface plot and (b) contour plot.

Experiments indicated that mixtures of more than 50% of SS with other substrates can be performed with no risk of inhibition. However, inhibition under continuous operation of a plant also depends on factors such as organic loading rate (OLR), HRT, and reactor configurations. Therefore, a small pilot scale continuously fed anaerobic digester should be operated before incorporating the mixture ratio.

3.2. Semicontinuous Experiments. According to the requirement of the plant only 5% or less food waste incorporation was tested for process performances under semicontinuous conditions for six HRT cycles of 20 days each. Figure 4 shows the specific biogas and methane production from the four cycles (20–100 days) reported as mL<sub>N</sub>/g VS added fed to the reactor. The average daily methane yield from SS (100% SS) and different mixture ratios of SS with PF and GT (1% to 2%) varied between 212 and 415 mL<sub>N</sub>/g VSadded. For small amounts of FW incorporation, biogas production was proportional to the percentage of FW and the biogas yield for 5% FW was the highest throughout the experiment duration which is consistent with the BMP assays [11].

For 100% SS, the average SBP was 284±9.7 mL<sub>N</sub>/g VS with methane content in the range of 64% and 66%. The average TS, VS, and tCOD removal for 100% SS was 41%, 50%, and 58%, respectively, which was in agreement with COD and VS removal of 55% and 36%, respectively, reported by Silvestre et al. for continuous AD of sludge mix of 70% PS and 30% WAS at an OLR of 1.5 to 1.7 kg VS/m<sup>3</sup>⋅d and HRT of 20 days [9]. A low SBP of 236±6.6 mL<sub>N</sub>/g VS was observed during the third HRT cycle (40–60 days) compared to HRT cycle two (20–40 days) when a new batch of feed was prepared with newly collected sample. Low TS, VS, and tCOD removal was also found during the period. This lag phase might be because of the biomass adaptation with the new feed [9]. The pH varied between 6.9 and 7.1 during the whole experiment.

The average SBP of 1% PF and 2% PF was 359±9 and 367±11 mL<sub>N</sub>/g VS added which is 25% and 32% higher than the SBP from 100% SS alone. Similarly, 23% and 47% increase in SBP were observed for 1% GT and 2% GT with an average SBP of 355±9 and 367±3 mL<sub>N</sub>/g VS added. As FOG has high biodegradability and BMP value (when added below 20% of the influent COD) [13], codigestion with a small
proportion of GT produced more biogas than the other food wastes in the same amount. The codigestion of three wastes SS:PF:GT at 95:2.5:2.5 (5% FW) produced an average SBP of 424 ± 10 mL\textsubscript{N2}/g VS (methane yield 327 mL\textsubscript{N2}/g VS) which is 50% higher than 100% SS (single substrate). These results are in agreement with the results reported by Luostarinen et al. (2009) [17] and Davidsson et al. (2008) [32]. They worked with SS and grease trap sludge (95:5 w/w) and reported...
Table 4: Biogas production and process performance in terms of TS, VS, and COD removal.

| Feedstocks | Parameters     | Period I (0–20d) | Period II (20–40d) | Period III (40–60d) | Period IV (60–80d) | Period V (80–100d) | Period VI (100–120d) |
|------------|----------------|------------------|--------------------|---------------------|--------------------|--------------------|----------------------|
|            | Avg biogas     | 256±16           | 337±14             | 320±1.5             | 284±8              | 339±7              | 355±9                |
| 1% PF      | CH4%           | 69±3.2           | 65±7.81            | 69±2.8              | 77±2.8             | 71±6.7             | 71±6.7               |
|            | TS removal%    | 43±0.01          | 43±0.03            | 40±0.04             | 40±0.4             | 46±1               | 41±4.5               |
|            | VS removal%    | 49±0             | 51±0.02            | 49±0.03             | 45±0.1             | 57±3.3             | 50±2.1               |
|            | COD removal%   | 61±0.06          | 58±0.01            | 53±0.02             | 59±0.03            | 59±0.03            | 55±2.03              |
|            | pH             | 7.4±0.4          | 7.0±0.08           | 7.1±0.08            | 7.01±0.06          | 7.02±0.05          | 7.07±0.03            |
|            |                |                  |                    |                     |                    |                    |                      |
|            | Avg biogas     | 252±14           | 335±9              | 334±2               | 322±1              | 364±2              | 367±3                |
| 2% PF      | CH4%           | 69±2.4           | 66±3.4             | 69±2.4              | 74±1               | 69±4.5             | 69±5.4               |
|            | TS removal%    | 43±0.02          | 45±0.01            | 41±0.05             | 45±2.5             | 46±0.06            | 45±2.2               |
|            | VS removal%    | 51±0.01          | 52±0.01            | 50±0.04             | 52±0.05            | 55±0.03            | 53±1.6               |
|            | COD removal%   | 58±0.1           | 54±0.01            | 55±0.06             | 59±0.03            | 57±0.03            | 57±2.05              |
|            | pH             | 7.3±0.4          | 7±0.04             | 7.1±0.06            | 7.01±0.06          | 7.03±0.02          | 7.05±0.04            |
|            |                |                  |                    |                     |                    |                    |                      |
|            | Avg biogas     | 281±1            | 376±2              | 386±3               | 393±2              | 415±19             | 424±10               |
| 5% FW      | CH4%           | 69±5.5           | 68±7.5             | 69±7.8              | 77±4.9             | 72±5.1             |                      |
|            | TS removal%    | 49±0.01          | 52±0.02            | 52±0.08             | 44±0.06            | 50±1.0             | 52±0.07              |
|            | VS removal%    | 56±0.01          | 60±0.02            | 60±0.02             | 55±0.08            | 54±1.01            | 59±4.05              |
|            | COD removal%   | 58±0.1           | 54±0.01            | 55±0.02             | 60±0.04            | 54±0.08            | 58±1.01              |
|            | pH             | 7.3±0.5          | 7.09±0.06          | 7.08±0.08           | 7.05±0.07          | 7.1±0.3            | 7.05±0.04            |
|            |                |                  |                    |                     |                    |                    |                      |
|            | Avg biogas     | 253±36           | 285±15             | 308±7               | 348±5              | 345±12             | 361±1                |
| 1% GT      | CH4%           | 69±2.51          | 67±2.3             | 69±2.12             | 75±2.8             | 69±7.1             |                      |
|            | TS removal%    | 44±0.01          | 45±0.01            | 45±0.06             | 43±0.01            | 44±0.06            | 45±3.05              |
|            | VS removal%    | 52±0.02          | 52±0.02            | 46±0.06             | 50±1.02            | 48±0.04            | 48±5.09              |
|            | COD removal%   | 63±0.04          | 59±0.01            | 63±0.04             | 65±1.06            | 56±2.3             | 59±1.06              |
|            | pH             | 7.3±0.5          | 7±0.06             | 7.02±0.04           | 7±0.05             | 7±0.12             | 7.08±0.13            |
|            |                |                  |                    |                     |                    |                    |                      |
|            | Avg biogas     | 284±9            | 336±5              | 312±3               | 329±7              | 405±4              | 395±8                |
| 2% GT      | CH4%           | 68±2             | 66±5.1             | 69±4.2              | 76±4.9             | 72±4.0             |                      |
|            | TS removal%    | 46±0.02          | 45±0.01            | 46±0.04             | 45±0.01            | 44±0.02            | 46±2.04              |
|            | VS removal%    | 54±0.01          | 54±0.02            | 53±0.03             | 53±0.01            | 57±0.08            | 53±0.03              |
|            | COD removal%   | 65±0.06          | 60±0.01            | 60±0.05             | 60±0.04            | 56±1.01            | 59±5.04              |
|            | pH             | 7.25±0.42        | 7±0.09             | 7.03±0.05           | 7±0.06             | 7.01±0.2           | 7.06±0.07            |
|            |                |                  |                    |                     |                    |                    |                      |
|            | Avg biogas     | 212±1.7          | 271±5.8            | 236±6.6             | 264±3.16           | 269±3.5            | 284±9.7              |
| 100% SS    | CH4%           | 64±4.5           | 62±1.5             | 64±7.8              | 66±2.8             | 66±9.6             |                      |
|            | TS removal%    | 43±0.02          | 41±0.03            | 40±0.06             | 40±0.01            | 39±0.03            | 40±6.08              |
|            | VS removal%    | 50±0.01          | 54±0.02            | 46±0.06             | 50±1.3             | 51±2.6             | 53±1.7               |
|            | COD removal%   | 60±0.03          | 58±0.03            | 55±0.06             | 56±0.6             | 55±0.8             | 55±1.01              |
|            | pH             | 7±0.12           | 6.99±0.09          | 7.04±0.09           | 6.93±0.02          | 7.03±0.05          | 7.05±0.04            |

methane yield of 374 and 295–308 mL/g VS corresponding to the organic loading of 1.67–2.23 and 2.5 kg VS/m³⋅d for HRT of 16 and 13 days, respectively. The addition of food wastes also increased the methane content and the average methane content was 69–72% in this experiment.

The TS removal for 1-2% food wastes (GT, PF) was between 42% and 49% and the corresponding VS removal was found to be between 50% and 56% (Table 4). This is similar to the VS removal reported in previous studies [17,32]. At the start of the second HRT cycle (20–40 days), the pH was between 6.8 and 6.9 for all the reactors, possibly because of high VA production at the beginning. The pH started increasing after that, indicating the consumption of produced VA due to acidification and inoculum acclimatisation [33]. However, when a new feed was prepared in the fourth HRT cycle (60–80 days), a lag phase was observed with low organic content removal, and low pH as well as low biogas production. However after the lag phase the reactors produced stable biogas production in the last two HRT cycles of the codigestion.
Methane production was increased significantly from 2% GT after the lag phase, possibly because the methanogens were acclimated to inoculum [4]. However, GT which is mainly lipid-rich material [34] has been found to have wide variation in characteristics (from Table 1, where characteristics' results from two different sample collections are shown).

The daily biogas production was observed to fluctuate, although the feedstocks were prepared by homogenising to constant TS loading throughout the experiments. As the FW had high variations in their characteristics, feeding a very small proportion in the reactors every day (from a batch of prepared feedstock) resulted in variations. Therefore, the average biogas production over each HRT cycle is shown in Table 4 and a rising trend was observed because of the acclimatisation of the inoculum to the feedstock.

The biogas and methane production potential of the food wastes was very high because of its high fat and protein content. Therefore, the incorporation of FW at very small ratios (1–5%) with SS in codigestion significantly improved biogas production from the SS alone. Although the biogas production improved greatly, VS and COD removal was not improved significantly (Table 4). This was likely due to the huge amount of more slowly degradable and/or inert material in the SS (60% degradable) [17]. The biodegradability of FW on the other hand was probably close to 100% due to the dilution with SS which caused the high biogas production.

Although SBP and methane yield depend on the origin of the substrates, composition, and operational conditions (SRT, temperature), the results reported by Silvestre et al. (2011) [9] and Davidsson et al. (2008) [32] showed a methane yield lower than this study when a small percentage of wastes from the dissolved air flotation unit of a wastewater treatment plant and kitchen grease wastes were added to SS. In addition to the biogas yield, different parameters were monitored at the end of each cycle to assess the quality of the supernatant and digestate (Table 5). It was observed that the pH value remained relatively stable at around 7 throughout the operation of the reactors. The alkalinity in all the reactors was around 2.5 to 2.75 g/L which also indicates no accumulation of VAs and the highest VA was observed from 2% GT (0.315 g/L) which is well below the threshold of inhibition (4 g/L) [35]. The VA accumulation might cause the instability of the process and an inhibition of acetotrophic methanogenesis [13]. However, the VAs in the reactors indicate stable process conditions. Luostarinen et al. (2009) observed total VA accumulation of not more than 0.43 g/L with a high ratio of grease trap to sludge in the feedstock (71% of the feed VS) while working with a mixture of PS, WAS, and grease trap sludge [17]. The ammonia-N content in all the reactors was between 0.6 and 0.71 g/L which is below the inhibition range (1.5–2.0 g/L) [4].

The last aspect to consider in anaerobic codigestion is the possibility of producing high quality compost (or fertilizer). In this case, the dewatered digestate characteristics for heavy metal contents need to be considered when assessing the effect of codigestion [7].

In Australia, the concentration of contaminants present in the biosolids and the microbial quality are two important parameters for biosolids' classifications. Contaminant grade (C1 and C2) and treatment grade (T1, T2, and T3) are the classifications of biosolids based on the factors described where C1/T1 are high quality products and can be used without restriction. According to the EPA guideline, biosolids from wastewater treatment plants are categorised as C2/T3 [30]. Integration among the AD and composting is possible where composting can play the role of curing step to overcome the phytotoxicty limit for VA and ammonia [36]. In the AD reactors, no inhibition of VA and ammonia N was observed. The digestate characteristics were adequate for the production of good quality compost by integrating a simple aerobic poststabilisation and dewatering for biological stability.

In Australia, the regulation of heavy metals in fertilizers of organic origin is governed by the Fertilizer Working Group, Department of Agriculture, AU Government (http://www.agriculture.gov.au/). The concerned heavy metals are zinc (Zn), copper (Cu), nickel (Ni), cadmium (Cd),

### Table 5: Bench scale AD reactors' performance at the end of the experiment.

| Parameter | Unit | 1% PF | 2% PF | 5% FW | 1% GT | 2% GT | 100% SS |
|-----------|------|-------|-------|-------|-------|-------|---------|
| TS g/L    |      | 21.15±2.43 | 20.62±2.57 | 21.13±5.1 | 20.02±3.46 | 20.45±0.26 | 20.34±0.97 |
| VS g/L    |      | 15.67±2.1 | 17.72±1.61 | 17.16±1.97 | 15.33±0.28 | 14.61±0.26 | 14.90±2.92 |
| tCOD g/L  |      | 28.025±0.25 | 26.65±0.07 | 28.9±0.21 | 26.075±7.88 | 28.55±0.21 | 29.025±5.69 |
| sCOD g/L  |      | 2.05±0.10 | 1.765±0.06 | 3.24±0.10 | 1.755±0.02 | 2.285±0.04 | 1.925±0.11 |
| TS removal % |     | 45±2 | 47±3 | 52±5 | 48±3 | 48±1 | 46±1 |
| VS removal % |     | 52±2.1 | 53±1.6 | 55±2 | 53±0.3 | 57±0.3 | 53±3 |
| COD removal % |    | 57±2.5 | 59±0.7 | 59±2.1 | 60±7.8 | 59±2.1 | 55±5.7 |
| TP g/L    |      | 0.36±0.03 | 0.38±0.04 | 0.44±0.04 | 0.4±0.02 | 0.42±0.01 | 0.41±0.001 |
| TN g/l    |      | 0.93±0.035 | 0.895±0.04 | 0.98±0.035 | 0.877±0.018 | 0.965±0.014 | 0.945±0.035 |
| TKN* g/l  |      | 1.9±0.3 | 2±0.42 | 2.4±0.3 | 2.2±0.2 | 2.3±0.15 | 2.2±0.3 |
| NH4-N g/l |      | 0.62±0.014 | 0.575±0.035 | 0.685±0.05 | 0.7±0.014 | 0.71±0.014 | 0.67±0.014 |
| VA g/l    |      | 0.147±0.004 | 0.148±0.01 | 0.266±0.07 | 0.253±0.05 | 0.315±0.014 | 0.208±0.005 |
| pH        |      | 7±0 | 7.08±0 | 7.13±0.02 | 7.05±0.02 | 7.09±0.05 | 7.04±0.02 |
| Alkalinity g/L | 24.98±0.29 | 2.7±0.06 | 2.756±0.08 | 2.711±0.05 | 2.678±0.03 | 2.671±0.05 |

*Analyses were carried out at a commercial laboratory (ALS, Australia).
Table 6: Digestate heavy metals concentration* in mL/g at the end of the experiment (after six HRT cycle of 20 days).

| Parameter | 1% PF | 2% PF | 5% FW | 1% GT | 2% GT | 100% SS | Limit** (mg/kg) |
|-----------|-------|-------|-------|-------|-------|---------|-----------------|
| Ca        | 430 ± 4 | 455 ± 5 | 480 ± 6 | 450 ± 5 | 440 ± 4 | 450 ± 7 | 450 ± 7        |
| Mg        | 82 ± 1 | 85 ± 0.5 | 89 ± 0.7 | 89 ± 0.4 | 85 ± 0.3 | 87 ± 0.5 | 87 ± 0.5       |
| Ca hardness | 1100 ± 2 | 1100 ± 0.5 | 1200 ± 1 | 1100 ± 0 | 1100 ± 0.5 | 1100 ± 0 | 1100 ± 0      |
| Mg hardness | 340 ± 0 | 350 ± 0 | 370 ± 1 | 360 ± 2 | 350 ± 5 | 360 ± 5 | 360 ± 5       |
| Al        | 170 ± 0 | 170 ± 0 | 180 ± 0 | 170 ± 0 | 170 ± 0 | 200 ± 0 | 200 ± 0       |
| As        | <1     | <1     | <1     | <1     | <1     | <1     | 20             |
| Cd        | <0.1   | <0.1   | <0.1   | <0.1   | <0.1   | <0.1   | 1              |
| Cr        | 0.5 ± 0.1 | 0.5 ± 0.1 | 0.5 ± 0.1 | 0.5 ± 0.1 | 0.5 ± 0.1 | 0.5 ± 0.1 | 400            |
| Cu        | 11 ± 1 | 11 ± 1 | 12 ± 0 | 12 ± 1 | 12 ± 0 | 14 ± 0 | 100            |
| Fe        | 160 ± 10 | 160 ± 5 | 170 ± 10 | 170 ± 5 | 170 ± 5 | 170 ± 5 | 170 ± 5       |
| Pb        | <0.5   | <0.5   | <0.5   | <0.5   | 0.5 ± 0.1 | 0.6 ± 0.1 | 300            |
| Hg        | <1     | <1     | <1     | <1     | <1     | <1     | 1              |
| Ni        | 0.4 ± 0.1 | 0.4 ± 0.1 | 0.5 ± 0.1 | 0.4 ± 0.1 | 0.4 ± 0.1 | 0.4 ± 0.1 | 60             |
| Zn        | 19 ± 0.5 | 20 ± 0.5 | 21 ± 1  | 20 ± 0.5 | 21 ± 1  | 22 ± 1.5 | 200            |
| Si        | 170 ± 5 | 180 ± 5 | 170 ± 5 | 170 ± 5 | 180 ± 5 | 210 ± 10 | 210 ± 10       |
| Si-SiO2   | 350 ± 10 | 380 ± 5 | 370 ± 5 | 380 ± 5 | 380 ± 5 | 440 ± 10 | 440 ± 10      |
| S         | 160 ± 5 | 170 ± 0 | 180 ± 5 | 170 ± 5 | 170 ± 5 | 200 ± 5 | 200 ± 5        |
| S-SO4     | 480 ± 10 | 520 ± 5 | 540 ± 5 | 520 ± 5 | 520 ± 5 | 600 ± 10 | 600 ± 10       |

* Heavy metal screening of the digestate samples was carried out by a commercial laboratory (ALS Environmental Division: Water Research Group).

** Contaminant upper limits for biosolids as grade C1 [30].

lead (Pb), chromium (Cr), and mercury (Hg) and their allowable limits are shown in Table 6. These heavy metals may be present in concentrations above the legal limits which can potentially harm environment and affect crop quality, crop yield, and soil fertility. Heavy metal concentration may increase during AD due to the microbial mineralization and loss of volatile solids [37]. Most national regulations prohibit the use of organic fertilizers, for example, digestate, if the concentrations of one or more heavy metals are higher than the threshold concentrations. There is also evidence suggesting that AD increases the complexation of heavy metals with organic ligands and hence lowers the mobility of heavy metals in the digestate [38, 39]. However, the metal contents found in these experiments were less than the allowable limit used in Australia for high quality amendments. Table 6 shows the concentration of heavy metal in the digestate collected reactors after six HRT cycles (at the end of the experiment).

4. Conclusions

FW is a suitable cosubstrate for the anaerobic codigestion of SS. The addition of 5% FW to the SS increased the SBP by up to 50% during semicontinuous experiments. Although the TS, VS, and tCOD removal slightly increased with codigestion, the methane content of the biogas improved significantly. The reactors showed stable pH and performance with no inhibitory effect. Based on the results from batch assays and the use of surface modelling, FW can be added at ratios up to 47%-48% (v/v) without inhibition to the AD process. Overall these results reveal the high potential of codigestion FW with SS to enhance biogas yield and quality.

Competing Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

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References

[1] L. Appels, J. Baeyens, J. Degrève, and R. Dewil, “Principles and potential of the anaerobic digestion of waste-activated sludge,” Progress in Energy and Combustion Science, vol. 34, no. 6, pp. 755–781, 2008.

[2] A. J. Gale, “The Australasian biosolids partnership and public perceptions,” Water Practice and Technology, vol. 2, no. 4, Article ID wpt2007081, 2007.

[3] D. L. Pritchard, N. Penney, M. J. McLaughlin, H. Rigby, and K. Schwarz, “Land application of sewage sludge (biosolids) in Australia: risks to the environment and food crops,” Water Science and Technology, vol. 62, no. 1, pp. 48–57, 2010.

[4] S. Woon and M. Othman, “Anaerobic digestion of meat wastes,” in Proceedings of the 2012 International Conference on Clean and Green Energy (IPCBEE ’12), vol. 27, pp. 36–40, IACSIT Press, 2012.

[5] X. Wang, G. Yang, F. Li, Y. Feng, G. Ren, and X. Han, “Evaluation of two statistical methods for optimizing the feeding composition in anaerobic co-digestion: mixture design and central composite design,” Bioresource Technology, vol. 131, pp. 172–178, 2013.
[6] I. Shizas and D. M. Bagley, "Experimental determination of energy content of unknown organics in municipal wastewater streams," *Journal of Energy Engineering*, vol. 130, no. 2, pp. 45–53, 2004.

[7] C. Cavinato, D. Bolzonella, P. Pavan, F. Fatone, and F. Cecchi, "Mesophilic and thermophilic anaerobic co-digestion of waste activated sludge and source sorted biowaste in pilot- and full-scale reactors," *Renewable Energy*, vol. 55, pp. 260–265, 2013.

[8] X. Dai, N. Duan, B. Dong, and L. Dai, "High-solids anaerobic co-digestion of sewage sludge and food waste in comparison with mono digestions: stability and performance," *Waste Management*, vol. 33, no. 2, pp. 308–316, 2013.

[9] G. Silvestre, A. Rodriguez-Abalde, B. Fernández, X. Flotats, and A. Bonmati, "Biomass adaptation over anaerobic co-digestion of sewage sludge and trapped grease waste," *Bioresource Technology*, vol. 102, no. 13, pp. 6830–6836, 2011.

[10] H. Bouallagui, L. Marouani, and M. Hamdi, "Performances comparison between laboratory and full-scale anaerobic digesters treating a mixture of primary and waste activated sludge," *Resources, Conservation and Recycling*, vol. 55, no. 1, pp. 29–33, 2010.

[11] N. D. Park, R. W. Thring, and S. S. Helle, "Comparison of methane production by co-digesting fruit and vegetable waste with first stage and second stage anaerobic digester sludge from a two stage digester," *Water Science and Technology*, vol. 65, no. 7, pp. 1252–1257, 2012.

[12] Y. Kalogo, H. Monteith, and P. Eng, *State of Science Report: Energy and Resource Recovery from Sludge*, Water Environment Research Foundation (WERF), 2008.

[13] R. Girault, G. Bridoux, F. Nauleau et al., "Anaerobic co-digestation of waste activated sludge and greasy sludge from flotation process: batch versus CSTR experiments to investigate optimal design," *Bioresource Technology*, vol. 105, pp. 1–8, 2012.

[14] *Food Waste in the Garbage Bin 2013*, Sustainability Victoria, Melbourne, Australia, 2014.

[15] D. Brown and Y. Li, "Solid state anaerobic co-digestion of yard waste and food waste for biogas production," *Bioresource Technology*, vol. 127, pp. 275–280, 2013.

[16] K. Koch, M. Plabst, A. Schmidt, B. Helmreich, and J. E. Drewes, "Co-digestion of food waste in a municipal wastewater treatment plant: comparison of batch tests and full-scale experiences," *Waste Management*, vol. 47, pp. 28–33, 2016.

[17] S. Luostarinen, S. Luste, and M. Sillanpää, "Increased biogas production at wastewater treatment plants through co-digestion of sewage sludge with grease trap sludge from a meat processing plant," *Bioresource Technology*, vol. 100, no. 1, pp. 79–85, 2009.

[18] J. C. Kabouris, U. Tezel, S. G. Pavlostathis et al., "Methane recovery from the anaerobic codigestion of municipal sludge and FOG," *Bioresource Technology*, vol. 100, no. 15, pp. 3701–3705, 2009.

[19] H.-W. Kim, S.-K. Han, and H.-S. Shin, "The optimisation of food waste addition as a co-substrate in anaerobic digestion of sewage sludge," *Waste Management and Research*, vol. 21, no. 6, pp. 515–526, 2003.

[20] I. Angelidaki, M. Alves, D. Bolzonella et al., "Defining the biomethane potential (BMP) of solid organic wastes and energy crops: a proposed protocol for batch assays," *Water Science and Technology*, vol. 59, no. 5, pp. 927–934, 2009.

[21] E. W. Rice, L. Bridgewater, and A. P. H. Association, *Standard Methods for the Examination of Water and Wastewater*, American Public Health Association, Washington, DC, USA, 2012.

[22] S. Strömberg, M. Nistor, and J. Liu, "Towards eliminating systematic errors caused by the experimental conditions in Biocatalyzed Methane Potential (BMP) tests," *Waste Management*, vol. 34, no. 11, pp. 1939–1948, 2014.

[23] *VDI Standard Procedures 4630: Fermentation of Organic Materials. Characterisation of the Substrate, Sampling, Collection of Material Data. Fermentation Tests*, Verein Deutscher Ingenieure, Beuth Verlag, Berlin, Germany, 2006.

[24] G. K. Kafle and S. H. Kim, "Anaerobic treatment of apple waste with swine manure for biogas production: batch and continuous operation," *Applied Energy*, vol. 103, pp. 61–72, 2013.

[25] S. Uładag-Derimir, G. N. Derimir, C. Frear, and S. Chen, "Anaerobic digestion of dairy manure with enhanced ammonia removal," *Journal of Environmental Management*, vol. 86, no. 1, pp. 193–200, 2008.

[26] Q. Niu, W. Qiao, H. Qiang, T. Hojo, and Y.-Y. Li, "Mesophilic methane fermentation of chicken manure at a wide range of ammonia concentration: stability, inhibition and recovery," *Bioresource Technology*, vol. 137, pp. 358–367, 2013.

[27] Z.-G. Liu, X.-F. Zhou, Y.-L. Zhang, and H.-G. Zhu, "Enhanced anaerobic treatment of CSTR-digested effluent from chicken manure: the effect of ammonia inhibition," *Waste Management*, vol. 32, no. 1, pp. 137–143, 2012.

[28] R. A. Labatut, L. T. Angenent, and N. R. Scott, "Biochemical methane potential and biodegradability of complex organic substrates," *Bioresource Technology*, vol. 102, no. 3, pp. 2255–2264, 2011.

[29] Y. Li, R. Zhang, C. Chen, G. Liu, Y. He, and X. Liu, "Biogas production from co-digestion of corn stover and chicken manure under anaerobic wet, hemi-solid, and solid state conditions," *Bioresource Technology*, vol. 149, pp. 406–412, 2013.

[30] *EPAVictoria, Guideline for Environmental Management-Biosolids Land Application*, EPAVictoria, Melbourne, Australia, 2004.

[31] Z. Siddiqui, N. J. Horan, and K. Anaman, "Optimisation of C:N ratio for co-digested processed industrial food waste and sewage sludge using the BMP test," *International Journal of Chemical Reactor Engineering*, vol. 9, article S4, 2011.

[32] Å. Davidsson, C. Lövstedt, J. Jansen, C. Gruberger, and H. Aspégren, "Co-digestion of grease trap sludge and sewage sludge," *Waste Management*, vol. 28, no. 6, pp. 986–992, 2008.

[33] M. Kawai, N. Nagao, N. Tajima, C. Niwa, T. Matsuyama, and T. Toda, "The effect of the labile organic fraction in food waste and the substrate/inoculum ratio on anaerobic digestion for a reliable methane yield," *Bioresource Technology*, vol. 157, pp. 174–180, 2014.

[34] J. H. Long, T. N. Aziz, E. L. de los Reyes III, and J. J. Ducoste, "Anaerobic co-digestion of fat, oil, and grease (FOG): a review of gas production and process limitations," *Process Safety and Environmental Protection*, vol. 90, no. 3, pp. 231–245, 2012.

[35] I. Siegert and C. Banks, "The effect of volatile fatty acid additions on the anaerobic digestion of cellulose and glucose in batch reactors," *Process Biochemistry*, vol. 40, no. 11, pp. 3412–3418, 2005.

[36] T. Di Stefano, M. Drennan, and J. VerNooy, "Laboratory-scale investigation of the curing process for anaerobic digestate," in *Proceedings of the 5th International Symposium on Anaerobic Digestion of Solid Wastes and Energy Crops*, Hammamet, pp. 25–28, Hammamet, Tunisia, 2008.

[37] C. Ciavatta, M. Govi, A. Simoni, and P. Sequi, "Evaluation of heavy metals during stabilization of organic matter in compost..."
produced with municipal solid wastes,” *Bioresource Technology*, vol. 43, no. 2, pp. 147–153, 1993.

[38] R. S. Lavado, M. B. Rodriguez, and M. A. Taboada, “Treatment with biosolids affects soil availability and plant uptake of potentially toxic elements,” *Agriculture, Ecosystems & Environment*, vol. 109, no. 3-4, pp. 360–364, 2005.

[39] C.-E. Marcato, E. Pinelli, M. Cecchi, P. Winterton, and M. Guiresse, "Bioavailability of Cu and Zn in raw and anaerobically digested pig slurry,” *Ecotoxicology and Environmental Safety*, vol. 72, no. 5, pp. 1538–1544, 2009.