



Article Optimisation of Biogas Production in the Co-Digestion of Pre-Hydrodynamically Cavitated Aerobic Granular Sludge with Waste Fats

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Abstract: The characteristics of excess aerobic granular sludge, related to its structure and chemical composition, limit the efficiency of anaerobic digestion. For this reason, pre-treatment methods and compositions with other organic substrates are used. In earlier work, no attempt was made to intensify the methane fermentation of the excess aerobic granular sludge by adding fatty waste materials. The aim of the research was to determine the effects of co-digestion of pre-hydrodynamically cavitated aerobic granular sludge with waste fats on the efficiency of methane fermentation under mesophilic and thermophilic conditions. The addition of waste fats improved the C/N ratio and increased its value to 19. Under mesophilic conditions, the highest effects were observed when the proportion of volatile solids from waste fats was 25%. The amount of biogas produced increased by 17.85% and CH₄ by 19.85% compared to the control. The greatest effects were observed in thermophilic anaerobic digestion at 55 °C, where a 15% waste fat content in volatile solids was ensured. This resulted in the production of 1278.2 \pm 40.2 mL/gVS biogas and 889.4 \pm 29.7 mL/gVS CH₄. The CH₄ content of the biogas was 69.6 \pm 1.3%. The increase in biogas and CH₄ yield compared to pure aerobic granular sludge anaerobic digestion was 34.4% and 40.1%, respectively. An increase in the proportion of waste fats in the substrate had no significant effect on the efficiency of methane fermentation. Strong positive correlations ($R^2 > 0.9$) were observed between biogas and CH₄ production and the C/N ratio and VS concentration.

Keywords: anaerobic digestion; aerobic granular sludge; waste fats; co-digestion; biogas; methane; process optimisation

1. Introduction

Anaerobic digestion (AD) is a recognised, technologically efficient, economically viable and, therefore, widely used method for the management of sewage sludge (SS). Its use leads to the stabilisation of the SS by reducing the content of organic compounds and susceptibility to putrefaction [1]. AD improves sanitary properties and limits the spread of odours and microbiological aerosols, which determines the possibility of using SS in nature or agriculture [2,3]. So far, AD has been used for the anaerobic degradation of mixed sewage sludge (MSS), which consists of about 60–80% organic material from primary sewage sludge (PSS) and 20–40% biomass from excess sewage sludge (ESS) [4,5]. PSS consists mainly of easily settleable suspensions that enter the sedimentation tanks with the wastewater and are released there. PSS is a substrate that is very susceptible to biodegradation under anaerobic conditions and is characterised by a high potential for biogas and methane production [6].

ESS, on the other hand, is the growing bacterial biomass that is regularly removed from reactors to maintain the appropriate concentration of microorganisms in wastewater



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Copyright: © 2024 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https:// creativecommons.org/licenses/by/ 4.0/). treatment plants (WWTP) [7]. In contrast to PSS, it poses major technological and operational problems in AD chambers [8]. Compared to PSS, the biogas yield of ESS is much lower, ranging from 500 to 700 m³/MgVS [9]. Therefore, it is common practice to mix ESS and PSS to achieve a biogas production of more than 1000 m³/MgVS [10]. ESS consists mainly of carbohydrates, proteins, humic substances and nucleic acids, which are enclosed in bacterial cells. The complex flocculent structure, in combination with extracellular polymeric substances (EPS), degradation-resistant cell walls and other high molecular weight organic compounds in ESS, makes it difficult to hydrolyse [11]. This requires a longer hydraulic retention time (HRT) and a larger digester volume. In addition, the C/N ratio of ESS is usually far below the optimal range for AD, which is between C/N 15–30 [12].

A specific type of ESS is aerobic granular sludge (AGS), whose popularity in wastewater treatment technology has increased dynamically in recent years [13]. In many cases, the use of AGS leads to lower investment and operating costs for wastewater treatment [14]. The main barrier to the dynamic development of AGS technology is the lack of data on the processing of excess sludge in AD [15]. Due to the characteristics and features of AGS that differ from typical ESS, the processes currently in use need to be assessed for their suitability and effectiveness. This may mean that the fundamentals and technological parameters of the AD process need to be verified and adapted to a substrate with a different chemical composition, structure and properties. Another problem related to the possibility of using AD for the biodegradation of AGS arises when this technology is used to treat industrial wastewater. In this case, the characteristics of the pollutants very often preclude the possibility of separating PSS from the wastewater stream to support AD [16]. The only option available to the operator of the technological system is the excessive use of AGS, which, due to its properties, chemical composition, and compact granular cell structure, poses significant technological problems in AD [17].

In such a case, it is necessary to take technological measures and introduce procedures that support the optimisation of AD. These primarily include AGS pretreatment techniques and anaerobic co-digestion (AC-D) through the use of organic substrates that complement the substrate composition [18]. Pretreatment processes lead to the destruction of complex cellular structures and the release of carbonaceous substances into the liquid phase, increasing access to anaerobic bacteria [19]. These include various processes based on mechanical disruption, hydrothermal depolymerisation under elevated temperature and pressure conditions, and chemical, physical and enzymatic hydrolysis processes [20]. Electrochemical technologies such as electrooxidation and electrocoagulation are also used to improve AD efficiency and the quality of the effluent after fermentation [21]. However, the AC-D of AGS with other externally supplied organic substrates is an increasingly common and widespread trend observed in municipal facilities and industrial wastewater treatment plants [22]. This is related to the concept of a utilisation biorefinery, the concept of which fits directly into the currently promoted assumptions of the bioeconomy and circular economy [23]. This type of treatment is fully justified from a technological and economic point of view and, in many cases, contributes to achieving energy self-sufficiency of WWTP and even a positive energy balance [24].

The experiments presented are directly in line with current trends in the development of biorefinery systems and make a significant new contribution to the current state of knowledge on AGS fermentation processes. The work describes the next step in multi-stage AD optimisation research, following the previous selection of the hydrodynamic cavitation process parameters [25]. This is the first report in which the performance of AC-D subjected to AGS pretreatment was analysed with a high lipid concentration of waste fats (WF) from the poultry sector. The main objective of the study was to determine the effects of codigestion of pre-hydrodynamically cavitated AGS with WF on the performance of methane fermentation. The influence of different mass ratios of the tested organic substrates and the temperature of the AD process on the generation efficiency and the composition of the gas products, as well as the kinetics of the anaerobic conversions, was investigated. The experimental data led to an assessment of the correlation between the properties of the biomass and the observed effects of the methane fermentation process.

2. Materials and Methods

2.1. Organisation of the Experiment

The experimental work was carried out under laboratory conditions in batch fermentation reactors. The study was divided into two stages (S). The criterion for the separation of the stages was the temperature of AD. In stage 1 (S1), the process was carried out under mesophilic conditions (38 ± 1 °C), and in stage 2 (S2), under thermophilic conditions (55 ± 1 °C). Four variants (V) were distinguished in each stage. Depending on V, a different proportion of volatile solids from the WF was used in the substrate mixture. In variant 1 (V1), no WF was added to the AGS pre-treated in the HC process (control sample). In variant 2 (V2), the proportion of volatile solids from WF was 8%, in variant 3 (V3) 15% and in variant 4 (V4) 25%. The organigram of the experiment is shown in Figure 1.



Figure 1. Organisation chart of the experimental work.

2.2. Materials

2.2.1. Aerobic Granular Sludge (AGS)

The AGS used in the experiment originates from a sequential biological reactor (SBR) that treats synthetic wastewater. Prior to the AD process, the AGS separated in the secondary settling tank and concentrated to $4.9 \pm 0.2\%$ TS was subjected to hydrodynamic cavitation (HC) for 15 min. Detailed information on the origin of the AGS and the design data of the HC used were presented in a previous research paper by the authors [25]. In these experiments, optimisation work was carried out to determine the most effective pre-treatment time in terms of methane production and energy balance. The highest technological and economic effects were achieved in the variant in which the HC time was 15 min [25]. Therefore, such a disintegrated AGS was used in the AD with WF. The characteristics of the AGS used are shown in Table 1.

| Parameter | Unit | AGS | WF | AS |
|-----------------------------|---------------|--|--|--|
| Total solids (TS) | mg/gFM %FM | $\begin{array}{c} 49.1\pm1.8\\ 4.91\pm0.18\end{array}$ | $101 \pm 5.4 \\ 10.10 \pm 0.54$ | $\begin{array}{c} 47.8 \pm 1.3 \\ 4.78 \pm 0.13 \end{array}$ |
| Mineral solids (MS) | mg/gFM %TS | $13.3 \pm 1.2 \\ 27.1 \pm 0.9$ | $\begin{array}{c} 10.6 \pm 0.5 \\ 9.44 \pm 0.91 \end{array}$ | $\begin{array}{c} 14,\!7\pm0.3\\ 30.7\pm0.2 \end{array}$ |
| Volatile solids (VS) | mg/gFM %TS | 35.8 ± 1.2 72.9 ± 1.1 | $90.3 \pm 1.6 \\ 90.6 \pm 1.3$ | $33.5 \pm 1.4 \\ 69.3 \pm 0.8$ |
| Total nitrogen (TN) | mg/gTS | 40.2 ± 5.8 | 95.11 ± 9.6 | 45.3 ± 3.1 |
| Total carbon (TC) | mg/gTS | 297.2 ± 4.5 | 9149 ± 52.6 | 384.4 ± 19.2 |
| Total organic carbon (TOC) | mg/gTS | 216.7 ± 4.1 | 6164 ± 44.4 | 319.7 ± 21.6 |
| Total inorganic carbon (IC) | mg/gTS | 80.5 ± 1.1 | 2985 ± 16.4 | $64,7\pm20.5$ |
| pH | - | 7.31 ± 0.07 | 7.01 ± 0.12 | 7.43 ± 0.06 |
| Lipid | mg/gTS | 4.2 ± 1.2 | 775.5 ± 22.7 | 1.9 ± 0.4 |
| Protein | mg/gTS | 251.3 ± 12.5 | 594.4 ± 110.2 | 283.1 ± 19.4 |
| C/N (TOC/TN) | - | 5.39 ± 1.1 | 64.8 ± 4.4 | 7.06 ± 0.7 |

Table 1. Characteristics of the initial substrates used in AD (AGS, WF) and the anaerobic sludge (AS) inoculum.

2.2.2. Waste Fat (WF)

The used WF came from an industrial plant specialising in the slaughtering and processing of poultry meat (chickens, turkeys). The substrate came from a plant for the removal of WF from wastewater, which is based on pressure flotation. In the pressure flotation system, the average size of the compressed air microbubbles used was 50 μ m. The floating grease was removed at regular intervals with a scraper into a storage tank. The WF was removed from the storage tank at regular intervals using a flotation pump. The purity of the WF was ensured by a pre-cleaning process based on the use of a sand trap to remove mineral impurities and a drum-type microsieve with a sieve diameter of 1.0 mm. The microsieve is equipped with a cleaning system based on spray nozzles. The impurities separated during drum filtration (sand, stones, suspensions, protein emulsions) are channelled into the sedimentation tank. The characteristics of the WF used are listed in Table 1.

2.2.3. Anaerobic Sludge Inoculum (AS)

The lab-scale AD batch bioreactors were inoculated with anaerobic sludge (AS) from a closed anaerobic chamber (CAC) with an active volume of 5000 m³, which is used for large-scale stabilisation of SS. The applied organic load rate (OLR) of the CAC was 2.4 kg VS/m³·day, the hydraulic retention time (HRT) was 20 days, and the process temperature was 38 °C. Since the thermophilic fermentation process was also used in the research work, it was first necessary to adapt the AS to these conditions. For this purpose, a CAC with an active volume of 5.0 L was operated at a temperature of 55 ± 1 °C for a period of 80 days (4 complete hydraulic replacements of the reactor). The substrate used, and the basic technological parameters corresponded to those of mesophilic fermentation (38 °C), OLR—2.4 kg VS/m³·day, HRT—20 days. The characteristics of the AS used in the study are presented in Table 1.

2.3. Anaerobic Digestion Respirometer Kit

AD was performed in respirometric bioreactors with a total volume of 500 mL (Automatic Methane Potential Test System II—AMPTS II, BPC Instruments AB, Lund, Sweden). Process monitoring was based on continuous measurements of volumetric biogas production. The bioreactors were equipped with agitators with a rotation speed of 80 rpm. The stirring system was started every 9 min and operated continuously for 1 min. At the beginning of the test cycle, the respirometric bioreactors were filled with 200 mL of AS inoculum, and then the tested substrate compositions were added. An initial OLR of 5 g VS/L was used. Anaerobic conditions in the bioreactors were ensured by flushing the inoculum and AS mixture with pure nitrogen at a capacity of 150 L/h for 1 min. Biogas/methane production was monitored once a day using the AMPTS application, which converted the volume to normal conditions (pressure 101.3 kPa, temperature 273 K). To determine the CH₄ yield, the biogas produced was fed into the ex-situ CO₂ absorption unit. The measurement was performed until the available organic compounds were completely biodegraded. Three consecutive measurements of the total volume of biogas produced, between which a difference of less than 1% was observed, determined the end of the experiment. The results obtained were corrected for the endogenous biogas produced by AS, to which no substrates were fed. A diagram of the test rig and the methods used in the respirometric measurements is shown in Figure 2.



Reactor volume: 500 mL Active volume: 200 mL Temperature control: S1 - 38±1°C, S2 - 51±1°C, Mixing of reactors: 80 rpm Ex-situ CO₂ absorption unit Volume of bottles: 100 mL Absorption liquid: 3M NaOH Absorption efficiency: >98% Flow cell array and DAQ unit Measurement resolution: 9 ml Detection capacity: up to 900 L Measuring range: 1 to 6000 mL/h Measuring precision: $CV \le 1\%$

Figure 2. Scheme of the test stand and the parameters used for the respirometric measurements of AD.

2.4. Analytical Methods

Total solids (TS), volatile solids (VS) and mineral solids (MS) were determined in accordance with standard EN 15934:2012. The pH of the homogenised solutions was measured using a pH meter (1000 L, VWR International, Radnor, PA, USA). The samples dried at 105 °C were also determined for total carbon (TC), total organic carbon (TOC) and total nitrogen (TN) using a Flash 2000 analyser (Thermo Scientific, Delft, The Netherlands). The protein concentration was determined using the protein conversion factor of TNx6.25. The fat was determined gravimetrically after evaporation of the solvent and drying of the residue at 105 °C. The biomass was acidified with HCl to pH 1, then evaporated and

transferred to thimbles and placed in a Soxhlet apparatus. Lipid extraction was performed with petroleum ether. The quality of the resulting biogas was analysed using an Agilent 7890A GC gas chromatograph (Agilent, Santa Clara, CA, USA) with a TCD detector. The biogas was collected using a gas-tight syringe (volume 20 mL). The GC was equipped with a Hayesep Q-column (80/100 mesh), a molecular sieve column (60/80 mesh) and a Porapak Q-column (80/100), which was operated at 70 °C. The injection and detector temperatures were 150 °C and 250 °C, respectively. Helium and argon were the carrier gases (flow rate 15 mL/min).

2.5. Calculation and Statistical Methods

The biogas and CH_4 production rate (r) and the rate constants (k) were estimated on the basis of an iterative method with linear regression [26].

$$Y(t) = -Y_{max} (e^{-kt} - 1)$$
(1)

where Y—biogas/methane yield (mL_N/gVS_{added}·day), Y_{max}—maximum biogas/methane yield (mL_N/gVS_{added}), k—kinetic constant (1/day), t—time (day). The biogas/methane r was determined by multiplying k and the maximum methane yield per gramme VS added to the bioreactors, resulting from first-order kinetics. The research works were conducted in four replicates. The significance of the differences between the variables tested was performed at a significance level of $\alpha = 0.05$ (Statistica 13.3, Statsoft, Inc., Tulsa, OK, USA). Normality of the distribution (Shapiro-Wilk test). Significance of the differences between the means (one-way analysis of variance—ANOVA). Homogeneity of variance between the groups (Levene test). Significance of the differences between the analysed variables (Tukey HSD test). The independent variable is the experimental V—the substrate/AD temperature used—and the dependent variables are the monitored and calculated characteristic AD parameters.

3. Results and Discussion

The work includes measures aimed at intensifying the AGS methane fermentation process by supplementing the substrate composition with WF from a poultry processing plant. Since the aim of the research was to put the results into practice, the maximum and limit value of the proportion of WF in the total amount of VS introduced into the AD was set at 25%. This value results from the available scientific and practical knowledge on the biodegradation of fats by anaerobic bacteria. It has been proven and confirmed that the addition of fatty substances to ESS usually improves the final efficiency of AD in terms of the amount of biogas and methane produced, but a high percentage of WF may result in the need to modify the technological parameters of the process [27]. It has been shown that the biodegradation of WF is slow, which means that if a significant amount of this substrate is added, the HRT must be extended, or the process must be carried out at lower OLR values [28]. Lower parameters of process kinetics may also be related to the accumulation of products of the acidic phase of fermentation, mainly volatile fatty acids (VFA), which lower the pH and limit the activity of methanogenic microorganisms [29].

Feeding anaerobic chambers intended for ESS fermentation with excessive amounts of WF also causes operational difficulties related to reduced mixing efficiency, clogging of pipes, reduced efficiency of pumping systems, foaming and scum formation, which limits the effective removal of biogas [30]. Due to the risk of such limitations, the substrate composition based on ESS and WF should be chosen carefully and wisely, which is especially important for existing and currently operating plants. Negative operational phenomena and emerging operational difficulties related to the use of fatty substances have been described by Davidsson et al. (2008) [31]. They showed that monodigestion of SS with grease traps does not ensure stable AD. After 10 days of slow start-up at an OLR of 1.7 kg VS/m³·day, the pH decreased, resulting in reduced CH₄ production. Despite repeated additions of NaHCO₃, it was not possible to stabilise the process [31]. Silvestre et al. (2014) [29], in turn, showed that increasing the WF content to 37% in AC-D with SS under thermophilic conditions led to unstable operation of the reactor, an accumulation of long-chain fatty acids (LCFA) and poor dewaterability of the digestate.

It has been shown that the simplest way to reduce the potential hazards to anaerobic systems arising from the AC-D of ESS with WF is to increase the AD temperature [32]. Changing the conditions from mesophilic to thermophilic helps to increase the biodegradability of fats by increasing the kinetics of hydrolysis and acidogenesis processes [33]. This is confirmed by the studies of Veeken et al. (1999) [34], who found that the rate of substrate hydrolysis accelerates with increasing temperature when comparing mesophilic and thermophilic conditions. In addition, Hao and Wang (2015) [35] found that the activity of extracellular enzymes was almost twice as high under thermophilic conditions (55 °C) than under mesophilic conditions (35 $^{\circ}$ C). Other researchers [36] also reported the activity of extracellular enzymes under different temperature regimes and confirmed that thermophilic temperatures can increase the activity of extracellular enzymes, which increases the rate of hydrolysis compared to mesophilic conditions. It was also found that acetate can be formed not only from pyruvate in the acetyl-CoA pathway but also from H_2 and CO_2 produced by homoacetogens (belonging to proteobacteria) during the acidification step [37]. For example, Hao and Wang (2015) [35] have shown that the abundance of proteobacteria is more than 10% higher under thermophilic conditions (35 °C) than under mesophilic conditions (55 °C) and that higher abundance of proteobacteria can contribute to homoacetogenesis. Based on the observations of operational efficiency and abundance of microbial communities, the likely dominant metabolic pathways under mesophilic and thermophilic conditions indicate that the ability of microbial communities to convert CO_2 to acetate via the acetyl-CoA pathway increases with increasing abundance of homoacetogens under thermophilic conditions [38]. This contributes to an increase in the production rate of simpler compounds and VFAs, which are the initial substrate for methanogenic bacteria [39]. In the studies by Hao and Wang (2015) [35], thermophilic fermentation led to a 10-fold higher accumulation of VFAs compared to mesophilic fermentation. α -Glucosidase and protease showed significantly higher activity in the thermophilic reactor. Illumina sequencing showed that the abundance of Clostridiaceae, Microthrixaceae and Thermotogaceae increased with increasing fermentation temperature, which could facilitate hydrolysis or acidification. Real-time PCR analyses showed that under thermophilic conditions, the relative abundance of homoacetogens increased in batch tests and reached higher values in stable fermentation, while it increased only slightly under mesophilic conditions in batch tests. Higher fermentation temperatures increased the activity of the main hydrolases, increased the proportion of bacteria involved in hydrolysis and acidification and promoted the relative abundance of homoacetogens, all of which led to higher VFA production [35]. Similar observations were made by Al-Sulaimi et al. (2022) [40] in studies on the thermophilic fermentation of SS. It was found that the dominant accumulation of VFAs is acetic acid. It has been shown that acetogens and methanogens can transfer electrons between groups of microorganisms, which promotes the rapid conversion of acetic acid to CH₄. It was found that the biogas produced from primary sludge (PS) and mixed sludge (MS) contained 66.75 \pm 0.5% and 52.29 \pm 0.5% CH₄, respectively. The high CH₄ content in digesters fed with PS under thermophilic conditions resulted from the high accumulation of VFA, which amounted to 824.68 ± 0.5 mg/L [40]. Under thermophilic AD conditions, a significantly higher activity of methanogenic microorganisms was also observed [41]. This is supported by the studies of Banach et al. (2018) [42], which showed that the genotypic structure of methanogenic communities analysed by PCR–DGGE changed under thermophilic conditions. The temperature had the greatest effect on the archaea methanogens in the digester immediately after the temperature was increased. Under thermophilic conditions, a significantly higher biogas yield and a higher average methane content in the produced biogas were observed [42].

The facts presented above formed the basis for selecting the WF dosage and conducting comparative studies on the AC-D of the tested AGS with WF under the commonly used mesophilic conditions and in the thermophilic fermentation process. Increasing the temperature of the AD process in systems where the biogas is combusted in a cogeneration unit is a relatively simple procedure. It is possible to recover and utilise low-temperature heat from combined heat and power (CHP) plants [43]. An additional simplification is the fact that WF from the meat sector must undergo heat treatment in accordance with regulations [44]. The requirements for hygienisation prior to the AD process stipulate that the material is pre-crushed to a maximum substrate particle diameter of 60 mm and heated at 70 °C for 60 min. This complies with the provisions of the European Commission's specification No. 142/2011 of 25 February 2011, which is contained in Regulation (EC) No. 1069/2009 of the European Parliament and of the Council [45].

3.1. Characterisation of the Substrates

In variant 1 (V1), which served as a reference for the subsequent research phases, AD was exposed to the AGS monosubstrate, which was pretreated with HC for 15 min [25]. The basis for the selection of the pretreatment time was a multivariate investigation aimed at obtaining the highest technological effect in terms of CH₄ production, as well as the operational and economic effect resulting from the energy balance performed [25]. The hydration of the AGS used was $4.91 \pm 0.18\%$ with a VS content of $72.9 \pm 1.1\%$ TS. The TN content was 40.2 ± 5.8 mg/gTS, the TOC concentration was 216.7 ± 4.1 mg/gTS, and the ratio of organic carbon to total nitrogen C/N was 5.39 ± 1.1 . The characteristics of the substrate compositions tested in the following variants of the study are shown in Table 2.

Table 2. Characteristics of the substrate mixtures used in the following variants of the experimental work.

| Parameter | Unit | Variant 1 | Variant 2 | Variant 3 | Variant 4 |
|-------------------------------|---------------|--|--|--|--|
| Total solids (TS) | mg/gFM %FM | $\begin{array}{c} 49.1 \pm 1.8 \\ 4.91 \pm 0.18 \end{array}$ | $50.8 \pm 2.5 \\ 5.08 \pm 0.25$ | $52.6 \pm 2.3 \\ 5.26 \pm 0.23$ | $54.3 \pm 1.9 \\ 5.43 \pm 0.19$ |
| Mineral solids (MS) | mg/gFM %TS | $\begin{array}{c} 13.3\pm1.2\\ 27.1\pm0.8\end{array}$ | $\begin{array}{c} 13.2 \pm 1.6 \\ 25.98 \pm 0.9 \end{array}$ | $\begin{array}{c} 13.1\pm1.4\\ 24.90\pm0.8\end{array}$ | $\begin{array}{c} 12.4 \pm 1.5 \\ 22.83 \pm 0.7 \end{array}$ |
| Volatile solids (VS) | mg/gFM %TS | 35.8 ± 1.2 72.9 ± 1.1 | 37.6 ± 1.6 74.02 ± 1.9 | $\begin{array}{c} 39.4\pm1.4\\ 75.10\pm2.0\end{array}$ | $\begin{array}{c} 41.9 \pm 1.5 \\ 77.17 \pm 1.8 \end{array}$ |
| Total nitrogen (TN) | mg/gTS | 40.2 ± 5.8 | 42.0 ± 3.8 | 43.9 ± 4.4 | 46.3 ± 6.5 |
| Total carbon (TC) | mg/gTS | 297.2 ± 4.5 | 592.3 ± 28.4 | 887.3 ± 21.5 | 1280 ± 42.2 |
| Total organic carbon (TOC) | mg/gTS | 216.7 ± 4.1 | 415 ± 15.2 | 613.2 ± 20.3 | 877 ± 22.3 |
| Total inorganic carbon (IC) | mg/gTS | 80.5 ± 1.1 | 177.3 ± 9.5 | 274.1 ± 8.8 | 403 ± 9.7 |
| pH | - | 7.31 ± 0.07 | 7.27 ± 0.06 | 7.19 ± 0.11 | 7.11 ± 0.07 |
| Lipids | mg/gTS | 4.2 ± 1.2 | 29.9 ± 2.1 | 55.6 ± 4.8 | 89.9 ± 4.2 |
| Proteins | mg/gTS | 251.3 ± 12.5 | 262.7 ± 13.3 | 274.1 ± 14.6 | 289.4 ± 18.4 |
| C/N (TOC/TN) | - | 5.39 ± 1.1 | 9.88 ± 1.9 | 13.96 ± 2.2 | 18.94 ± 2.6 |

These values are characteristic of AGS, as previous studies in the literature have shown [46–48]. According to common knowledge, the value of the C/N ratio is well below the optimum value for the efficient operation of the methane fermentation process [49]. According to literature data and operating reports, the value of this parameter should be in the range of 15 to 30 [50]. Many previous works have reported too low values for the C/N ratio in SS. Zheng et al. (2021) [51] reported 6.60 ± 0.22 , and Azarmanesh et al. (2021) [12] showed a value of 7.6. The purpose of introducing WF into the substrate composition was to improve the C/N ratio and ensure a higher supply of organic compounds, the available amount of which determines the amount of biogas and methane produced. This is a typical technological procedure described by Arelli et al. (2021) [52] in their work on the AC-D of

ESS with food waste and by Ahmadi-Pirlou and Mesri Gundoshmian (2021) [53] in their research on the AC-D of ESS with municipal solid waste.

The addition of WF to AGS at a level ensuring 8% VS from fats in V2 made it possible to increase the C/N values to $9.88 \pm 1.9 \ (p < 0.5)$ (Table 2). At this stage, the TOC and TN contents in the biomass were $415 \pm 15.2 \ \text{mg/gTS}$ and $42.0 \pm 3.8 \ \text{mg/gTS}$, respectively. The VS content in the biomass was $74.02 \pm 1.9\%$ TS (Table 2). Increasing the proportion of WF in the substrate led to a further increase in the organic compound content, which had a direct effect on positive changes in the C/N ratio, which was 13.96 ± 2.2 at V3 and reached a level in the optimum range of 18.94 ± 2.6 at V4 (p < 0.5) (Table 2). The composition of the raw WF contained a VS concentration of $90.6 \pm 1.3\%$ TS, a TOC content of $6164 \pm 44.4 \ \text{mg/gTS}$, a TN of 95.11 ± 9.6 and a C/N ratio of 64.8 ± 4.4 (Table 1). The use of the AGS and WF proportions tested in the study had no significant (p > 0.5) effect on the changes in TN concentration in the biomass, which was within a narrow range of $40.2 \pm 5.8 \ \text{mg/gTS}$ to $46.3 \pm 6.5 \ \text{mg/gTS}$ (Table 2). The increase in VS and TOC content in the biomass and the improvement in the C/N ratio due to the introduction of lipids in SS are also confirmed by the work of Silvestre et al. (2011) [54] and Davidsson et al. (2008) [31].

3.2. Anaerobic Digestion

During AD of the AGS monosubstrate at 38 °C (S1V1), biogas production was 760.5 \pm 29.3 mL/gVS with a CH₄ content of 63.4 \pm 1.1%. The biogas production rate (r) was 174.8 mL/day, and the production rate constant (k) was 0.23 1/day (Figure 3). In the study by Cydzik-Kwiatkowska et al. (2022) [55], AGS fermentation enabled a biogas yield of 400 mL/gVS, 420 mL/gVS and 455 mL/gVS after 0.5, 4.0 and 8.0 min of ultrasonic pretreatment, respectively. The biogas yield in raw AGS was 375 mL/gVS [55]. Kazimierowicz et al. (2023) [47] achieved biogas and CH₄ production of 476 \pm 20 mL/gVS and 341 \pm 13 mL/gVS, respectively, under mesophilic conditions by digesting AGS after digestion with solidified CO₂ at an optimal ratio of solidified CO₂ to AGS of 0.3 [47]. Similar results were obtained under thermophilic conditions [56]. At the same ratio of solidified CO₂ to AGS, 482 \pm 21 mL/gVS biogas and 337 \pm 14 mL/gVS CH₄ were obtained [56]. In studies investigating the effects of the thermal hydrolysis process (THP) on the solubilisation of the main organic substances of SS and the effectiveness of the subsequent biochemical methane potential (BMP) tests under mesophilic conditions (35 °C), the results were between 940–1070 mL/gVS CH₄, depending on the variant [57].

Significantly higher AD effects than in S1V1 were observed in S1V2 and S1V3, where the proportion of WF in VS was 8% and 15%, respectively (p < 0.5). However, it should be noted that no significant differences were observed between the results in these two variants (p > 0.5). Biogas production was 823.1 \pm 33.4 mL/gVS in S1V2 and 838.3 \pm 32.3 mL/gVS in S1V3 (Table 3). There were also no differences in CH₄ content, which was between 63.2 \pm 1.3% and 63.5 \pm 1.1% (p > 0.5) (Table 3). The kinetics of the process were also very similar, with an r-value of 0.22 1/day for both variants (Figure 3). A significant improvement in AD efficiency was observed in S1V4, where the proportion of VS from WF was increased to 25% (p < 0.5). The biogas production in this variant was 925.8 \pm 36.8 mL/gVS, with k = 222.0 mL/day and r = 0.24 1/day (Table 3, Figure 3). The CH₄ content fluctuated near the level of 65.0 \pm 1.2%, resulting in unit production of 602.0 \pm 23.2 mLCH₄/gVS at a rate (k) of 138.5 mLCH₄/day (Table 3, Figure 3). The average amount of biogas produced in S1V4 was 17.85% higher than in S1V1, but CH₄ production increased on average by 19.85% thanks to the use of AGS and WF in AC-D under mesophilic conditions.

AC-D of SS with WF under mesophilic conditions was analysed by Grosser and Neczaj (2018) [58], who observed an increase in the efficiency of biogas production at a proportion of more than 10% WF. The highest biogas production value of 27.5 L/L·d was achieved with 52% WF addition, which was three times higher than with SS mono-digestion. Increasing the WF addition above 54% led to a drastic decrease in biogas production to 1.2 L/L·d. At the same time, a decrease in CH₄ yield was also observed. The highest CH₄ production values between 404 and 448 L/kgVS were recorded at 34% WF. The biogas yield for this

variant ranged from 536 to 589 L/kgVS [58]. Davidsson et al. (2008) [31] found that the addition of 10 to 30% WF increased CH₄ production by 9 to 12% and reached values in the range of 295–344 L/kgVS. The CH₄ concentration in the biogas ranged from 66 to 69% [31]. Martínez et al. (2016) achieved a cumulative CH₄ production of 700 mL/gVS under mesophilic conditions in the AC-D of SS and butchery fat waste as co-substrate [59].



Figure 3. Kinetics of biogas and CH_4 production in stage 1 of the experiment. (r—production rate (mL/day); k—production rate constant (1/day)).

Table 3. Production efficiency and composition of the biogas depending on the variant in stage 1.

| | Production Efficiency | | | | | | Main Components of Biogas | |
|---------|-----------------------------------|----------------|----------------|-----------------|------------------|------------------|---------------------------|---------------------|
| Variant | | Biogas | | | Methane | | | icitis of Diogus |
| | mL/gFM | mL/gTS | mL/gVS | mL/gFM | mL/gTS | mL/gVS | CH4 [%] | CO ₂ [%] |
| S1V1 | $\textbf{27.23} \pm \textbf{1.1}$ | 554.5 ± 22.3 | 760.5 ± 29.3 | 17.28 ± 0.6 | 351.8 ± 15.5 | 482.5 ± 22.6 | 63.4 ± 1.1 | 36.1 ± 1.3 |
| S1V2 | 62.1 ± 1.6 | 697.5 ± 31.3 | 823.1 ± 33.4 | 39.21 ± 1.3 | 440.6 ± 21.6 | 519.9 ± 18.8 | 63.2 ± 1.3 | 30.1 ± 1.1 |
| S1V3 | 96.8 ± 2.8 | 748.5 ± 33.5 | 838.3 ± 32.3 | 61.49 ± 1.6 | 475.5 ± 20.5 | 532.5 ± 21.1 | 63.5 ± 1.1 | 29.5 ± 1.3 |
| S1V4 | 128.5 ± 2.5 | 826.6 ± 35.4 | 925.8 ± 36.8 | 98.37 ± 1.5 | 537.5 ± 22.4 | 602.0 ± 23.2 | 65.0 ± 1.2 | 28.9 ± 1.2 |

The increase in AD efficiency achieved was less than the previously published research results suggested. For this reason, and in view of the relative ease with which such a technological solution could be used in practice, tests were carried out under thermophilic conditions. By using thermophilic AGS fermentation at a temperature of 55 °C (S2V1), 835.2 \pm 30.3 mL/gVS of biogas and 575.9 \pm 24.6 mL/gVS of methane were obtained (Figure 4, Table 4). These values were 8.94% and 16.16% higher than the values achieved in S1V1. A significantly higher efficiency of CH₄ production was caused by an increase in the proportion of this component in the gas mixture to 68.9 \pm 1.2% (p < 0.5) (Table 4). An even greater improvement in the technological effects achieved was observed in the variants in which AGS was supplemented with WF. In S2V2, where the proportion of VS from WF was 8%, 1118.5 \pm 39.3 mL/g biogas VS was obtained at the end of the process, with a CH₄ content of 69.0 \pm 1.3% (Table 4). Compared to S1V2, the amount of biogas increased by 26.4% and that of methane by 32.7% (p < 0.5). The kinetics of the biogas

production process were characterised by a rate (r) of 279.5 mL/day and a rate constant (k) of 0.25 1/day (Figure 4). Similar proportions of SS and WF in the AD process were used by Silvestre et al. (2014) [29], who obtained 0.26 L/L·d CH₄ without the addition of WF, 0.345 L/L·d CH₄ with 8% WF and 0.575 L/L·d CH₄ with 27% WF. At a WF content of 29%, a decrease in CH₄ production to a value of 0.545 L/L·d was observed [29]. Davidsson et al. (2008) [31] also used similar ratios of the substrates mentioned. SS AD od SS without WF yielded 271 L/kgVS CH₄. The addition of 10% WF led to an increase in CH₄ production to 295–308 L/kgVS. With 30% WF, however, CH₄ production increased to 344 L/kgVS [31].



Figure 4. Kinetics of biogas and CH_4 production in stage 2 of the experiment. (r—production rate (mL/day); k—production rate constant (1/day)).

| Table 4. Production efficiency and composition of the biogas depending on the variant in stag | ge. | 2 |
|--|-----|---|
|--|-----|---|

| | Production Efficiency | | | | | Main Components of Biogas | | |
|---------|-----------------------|-----------------|-----------------|-----------------|----------------|---------------------------|--------------|---------------------|
| Variant | | Biogas | | | Methane | | | lents of blogas |
| | mL/gFM | mL/gTS | mL/gVS | mL/gFM | mL/gTS | mL/gVS | CH4 [%] | CO ₂ [%] |
| S2V1 | 29.90 ± 1.2 | 609.0 ± 24.4 | 835.2 ± 30.3 | 20.62 ± 0.8 | 419.9 ± 19.6 | 575.9 ± 24.6 | 68.9 ± 1.2 | 30.6 ± 1.1 |
| S2V2 | 81.3 ± 1.9 | 1016.8 ± 36.2 | 1118.5 ± 39.3 | 126.2 ± 2.1 | 702.1 ± 25.8 | 772.3 ± 24.1 | 69.0 ± 1.3 | 30.0 ± 1.3 |
| S2V3 | 130.0 ± 2.6 | 1083.2 ± 32.3 | 1278.2 ± 40.2 | 67.07 ± 1.4 | 753.7 ± 29.6 | 889.4 ± 29.7 | 69.6 ± 1.3 | 29.1 ± 1.2 |
| S2V4 | 148.8 ± 2.1 | 1150.4 ± 31.1 | 1288.4 ± 36.4 | 103.4 ± 1.7 | 799.9 ± 28.7 | 895.8 ± 30.1 | 69.5 ± 1.2 | 29.2 ± 1.1 |

Increasing the VS proportion of WF to 15% led to a significant improvement in AD efficiency. The amount of biogas produced was $1278.2 \pm 40.2 \text{ mL/gVS}$ (r = 319.5 mL/day, k = 0.26 1/day) and the CH₄ content was $889.4 \pm 29.7 \text{ mL/gVS}$ (r = 222.2 mL/day, k = 0.25 1/day) (Figure 4, Table 4). The CH₄ content of the biogas was $69.6 \pm 1.3\%$ (Table 4). Compared to S1V3, the biogas yield increased by 34.4% and the CH₄ content by 40.1% (p < 0.5) (Table 4). Increasing the addition of WF to the substrate mixture in S2V4 had no significant effect on the biogas and CH₄ yield (p > 0.5). The observed values did not differ significantly from those in S2V3 (p > 0.5). The amount of biogas produced was $1288.4 \pm 36.4 \text{ mL/gVS}$ (Figure 4, Table 4). The CH₄ content was lower than in S2V3 and was $69.5 \pm 1.2\%$ (Table 4). The values characterising the kinetics of AD were also similar to those recorded in S2V3 (Figure 4). Silvestre et al. (2014) [29] compared the mesophilic

and thermophilic AC-D effects of SS with the addition of WF. The highest CH₄ yield was obtained with a WF content of 27%. In this case, mesophilic AD proved to be more effective and yielded 0.575 L/L·d CH₄. However, 0.4 L/L·d CH₄ was obtained by thermophilic fermentation. Increasing the WF content to 37–39% led to a decrease in AD efficiency [29]. In the mesophilic and thermophilic AD of SS and fats, oils and grease, the CH₄ yield during single-stage fermentation was 473 mL/gVS and 551 mL/gVS, respectively. After the initial hydrolysis, significantly higher values were recorded, reaching 1040 mL/gVS and 1083 mL/gVS at 35 °C and 52 °C, respectively [60].

The increase in biogas and CH₄ production for the variants V1–V4 during mesophilic fermentation in S1 was strongly positively correlated with the C/N ratio, and the coefficients of determination were $R^2 = 0.9301$ (Figure 5A) and $R^2 = 0.9256$ (Figure 5B), respectively. A strong positive correlation between these parameters was also observed in S2, as evidenced by the coefficients of determination of $R^2 = 0.8067$ for biogas (Figure 5A) and $R^2 = 0.8121$ for CH₄ (Figure 5B). Similar relationships were observed between VS concentration and biogas and CH₄ production, with very strong positive correlations in S1 (Figure 5C) and strong positive correlations in S2 (Figure 5D).



Figure 5. Correlations between biogas production (**A**) and CH_4 (**B**) and the C/N ratio and between biogas production (**C**) and CH_4 (**D**) and the VS concentration.

4. Conclusions

The pre-hydrodynamically cavitated AGS used in the tests was mineralised, as confirmed by low concentrations of indicators of organic compound content, including VS and TOC. This had a direct impact on the very low C/N ratio, which is significantly different from the values considered optimal for proper AD. A justified technological measure was to supplement the AGS biomass with another substrate rich in organic compounds and capable of AD. The addition of WF to the substrate composition had a positive effect on the properties of the substrate. The concentration of organic matter was significantly increased, and the C/N ratio improved, which led to an increase in the efficiency of AD.

Under mesophilic fermentation conditions, the highest technological effects in terms of biogas and CH_4 production were observed when the proportion of VS from WF was 25%. In this variant, the average amount of biogas produced increased by 17.85% and CH_4 by 19.85% compared to the fermentation of the AGS monosubstrate.

The highest technological effects were observed when the mixture of AGS and WF was subjected to AD under thermophilic conditions. In the variant in which the proportion of WF in the VS was 15%, $1278.2 \pm 40.2 \text{ mL/gVS}$ biogas and $889.4 \pm 29.7 \text{ mL/gVS}$ CH₄ were obtained. The CH₄ content of the biogas was $69.6 \pm 1.3\%$. The increase in biogas yield compared to pure AD in AGS was 34.4%, and that CH₄ was 40.1%. Increasing the addition of WF to the substrate mixture to 25% VS no longer had a significant effect on increasing the efficiency of AD.

Very strong positive correlations ($R^2 > 0.9$) were observed between biogas and CH₄ production and the C/N ratio and VS concentration during mesophilic fermentation. Strong positive correlations ($R^2 > 0.8$) were observed between these parameters during thermophilic fermentation.

Research proves that the enrichment of AGS with a high content of organic compounds substrate is fully justified technologically justified. Considering the properties of AGS, both in terms of the structure and complexity of the granules as well as the chemical composition, this can be considered a necessary procedure that determines the correct course of methane fermentation and, consequently, the appropriate mineralisation and stabilisation of AGS. It should be emphasised that AC-D can be, and in many cases is, used in existing plants without the need for significant structural or technological changes so that the results obtained can be quickly put into practice. However, it should be emphasised that the investigations presented were carried out on a laboratory scale in respirometric batch reactors. In order to obtain more detailed results, the experimental work should be continued in reactors with a continuous operation whose operational and technological parameters (organic loading rate, hydraulic retention time, substrate dosing method, degree of mixing, and digester geometry) correspond to real operating conditions. Only the results obtained after long-term use of the AGS and WF mixture under conditions similar to large-scale operation can form the basis for establishing a reliable and credible energy and economic balance and conducting a life cycle assessment of the entire technology.

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