



# Net-zero carbon condition in wastewater treatment plants: A systematic review of mitigation strategies and challenges

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## ABSTRACT

The wastewater sector accounts for up to 7 and 10% of anthropogenic CH<sub>4</sub> and N<sub>2</sub>O emissions, respectively. Nowadays wastewater treatment plants are going through a paradigm shift to approach a net-zero carbon condition. Numerous ongoing measures have taken place to identify the sources of greenhouse gases and minimize the carbon footprint. This paper systematically reviews all known practices leading towards net-zero carbon wastewater treatment. The greenhouse gas emissions from the wastewater sector are identified and carbon footprint quantification tools, such as reliable models and emission factors are compared. The direct process emissions can contribute to over 60% of the carbon footprint in wastewater treatment plants, while around 30% of the carbon footprint is due to energy-related indirect emissions. Therefore, greenhouse gas mitigation via process optimization and energy usage in wastewater treatment plants are comprehensively described. The implantation of novel nitrogen removal processes can reduce both greenhouse gas emissions and energy consumption. Other techniques such as source separation systems can potentially allow mitigation of N<sub>2</sub>O emissions by 60% while avoiding energy-intensive nitrogen fertilizer production. Nutrient recovery methods are another approach which offer negative value for the net carbon footprint. Recovering N<sub>2</sub>O for energy production is a promising method which can lead to both direct and indirect carbon footprint reductions. Ultimately, to achieve full decarbonization any remaining emissions need to be offset, including carbon footprint of chemicals usage and transportation.

## 1. Introduction

The biological wastewater treatment process has historically been focused on efficient removing organic pollutants and nutrients to reduce the impact of wastewater on the aquatic environment. However, the wastewater sector is now undergoing a paradigm shift towards an integrated operation which is focused on resource recovery and net-zero carbon condition [1]. It is estimated that wastewater treatment plants (WWTPs) are responsible for nearly 5% of the global non-CO<sub>2</sub> greenhouse gas (GHG) emissions and this is projected to increase by 22% by 2030 [2,3].

GHG emissions from WWTPs are classified as either direct or indirect. The direct, non-biogenic GHG emissions, termed also scope 1 emissions, take place during wastewater and sludge treatment processes

[4]. The direct CO<sub>2</sub> emissions are considered carbon neutral due to their biogenic nature, i.e., produced by the biodegradation of organic compounds in wastewater. This approach is in contrast with other sectors, such as transportation and energy, where the fossil-based CO<sub>2</sub> is the major contributor to GHG emissions. On the other hand, non-CO<sub>2</sub> direct emissions, including methane (CH<sub>4</sub>) and nitrous oxide (N<sub>2</sub>O), are of a significant concern due to a high global warming potential (GWP) of those gases. It is widely acknowledged that the wastewater treatment sector significantly contributes to CH<sub>4</sub> and N<sub>2</sub>O emissions, estimated at 7–10% for each gas [5,6]. However, these percentages can vary depending on site-specific factors, such as treatment technologies, plant size, and regional differences in wastewater management practices. Based on the latest Intergovernmental Panel on Climate Change (IPCC) report [7], the GWP in a 100 years horizon is 27.2, 29.8 and 273 for non-fossil origin CH<sub>4</sub>, fossil-origin CH<sub>4</sub> and N<sub>2</sub>O, respectively.

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**Nomenclature and abbreviations**

Item	Description		
AD	Anaerobic Digestion	HRT	Hydraulic Retention Time
ADM	Anaerobic Digestion Model	IEA	International Energy Agency
ANAMMOX	Anaerobic Ammonia Oxidation	IPCC	Intergovernmental Panel on Climate Change
AnAOB	Anaerobic Ammonia Oxidizing Bacteria	ISO	International Organization for Standardization
AOB	Ammonia Oxidizing Bacteria	LCA	Life Cycle Assessment
ASM	Activated Sludge Model	ML	Machine Learning
ASMN	Activated Sludge Model for Nitrogen	MLR	Mixed Liquor Recirculation
BEAM	Biosolids Emissions Assessment Model	N	Nitrogen
BNR	Biological Nutrient Removal	N <sub>2</sub>	Dinitrogen
BOD	Biochemical Oxygen Demand	N <sub>2</sub> O	Nitrous Oxide
BW	Black Water	N <sub>2</sub> OR	Nitrous Oxide Reductase
CANDO	Coupled Aerobic-Anoxic Nitrous Decomposition Operation	NH <sub>2</sub> OH	Hydroxylamine
CE	Circular Economy	NH <sub>4</sub> <sup>+</sup>	Ammonium
CF	Carbon Footprint	NO	Nitric Oxide
CH <sub>4</sub>	Methane	NO <sub>2</sub> <sup>-</sup>	Nitrite
CHP	Combined Heat and Power	OFMSW	Organic Fraction of Municipal Solid Waste
CO <sub>2e</sub>	Carbon Dioxide Equivalent	P	Phosphorus
COD	Chemical Oxygen Demand	PANDA	Partial Nitrification, Denitrification and Anaerobic Ammonia Oxidation
CWA	Clean Water Act	PD	Partial Denitrification
DAMO	Denitrifying Anaerobic Methane Oxidation	PD/A	Partial Denitrification-Anammox
DAMO/A	Denitrifying Anaerobic Methane Oxidation-Anammox	PE	Population Equivalent
DEFRA	Department for Environment, Food and Rural Affairs	PN/A	Partial Nitrification-Anammox
DNRA	Dissimilatory Nitrate Reduction to Ammonium	SBR	Sequencing Batch Reactor
DO	Dissolved Oxygen	SCENA	Shortcut Enhanced Nutrient Abatement
DW	Dry Waste	SCND	Shortcut Nitrification-Denitrification
ECAM	Energy Performance and Carbon Emissions Assessment and Monitoring	SHARON	Single Reactor System for High Activity Ammonium Removal over Nitrite
EF	Emission Factor	SNAD	Simultaneous Partial Nitrification-Anammox and Denitrification
EPA	Environmental Protection Agency	SRT	Solid Retention Time
EU	European Union	TCD	Thermal Conductivity Detector
FOG	Fat-Oil-Grease	TKN	Total Kjeldahl Nitrogen
FTIR	Fourier Transformed Infrared Spectroscopy	TN	Total Nitrogen
FU	Functional Unit	TSS	Total Suspended Solids
FW	Food Waste	UK	United Kingdom
GC	Gas Chromatography	US	United States
GEA	German Environmental Agency	VS	Volatile Solid
GHG	Greenhouse Gas	WEF	Water Environmental Federation
GW	Grey Water	WFD	Water Framework Directive
GWP	Global Warming Potential	WRRF	Water Resource Recovery Facility
		WWTP	Wastewater Treatment Plant

The indirect GHG emissions, referred to as scope 2 emissions, are due to energy and electricity consumption by WWTPs. The remaining indirect emissions, referred to as scope 3 emissions, are associated with transportation and production of chemicals outside of the plant [2]. The total sum of GHG emissions from all the scopes is known as the carbon footprint (CF) and reported based on the carbon dioxide equivalent (CO<sub>2e</sub>) unit [8].

In comparison with the transportation and energy sectors, the wastewater sector has fewer studies on net-zero carbon pathways [1]. However, many countries are beginning to move towards low-carbon operation and undertaking actions to decarbonize municipal water management [9]. Net-zero targets are included in near-term policies and plans, and the discussion on the net-zero condition has been evolving recently in line with the targets of the Paris Agreement [10].

Metcalf et al. [11] outlined criteria for a net-zero transition in wastewater treatment, including the overall energy balance, process-related GHG emissions, chemical consumption and sludge disposal. Therefore, the net-zero pathways for wastewater sector is more diverse and complex compared to other sectors. A multi-criteria approach that considers both wastewater reclamation and other

sustainability goals need to be defined specifically for the wastewater sector [1]. The net-zero carbon condition refers to a prospective WWTP which aims to offset certain CF that is sum of scope 1, 2 and 3 emissions in wastewater treatment and thus attain net-zero GHG emission. The net-zero carbon WWTP employs novel strategies to mitigate N<sub>2</sub>O and CH<sub>4</sub> emissions via process modification. Furthermore, this plant needs to mitigate CF via energy optimization as well as chemicals and transportation optimization. The number of scientific papers on the topic has increased rapidly in the last decade (Fig. 1), confirming growing attention from the scientific community.

This review aims to provide comprehensive insights into effectively reducing CF and achieving a net-zero carbon condition in municipal WWTPs. The paper covers crucial aspects, including relevant regulations, GHG sources, and quantification methods and tools for CF assessment. Process optimization, energy usage, and reduction and offsets of indirect emissions are discussed to minimize CF. Practices for net-zero carbon wastewater treatment and a multi-criteria approach to decarbonize WWTPs are thoroughly reviewed with an environmental protection, resource recovery, and reducing impact. GHG emissions in WWTPs, particularly N<sub>2</sub>O and CH<sub>4</sub>, and their sources are explored.

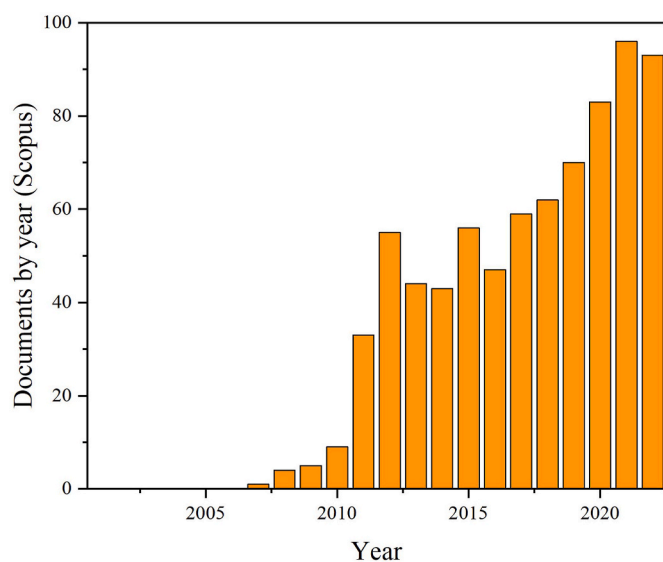


Fig. 1. Number of publications covering CF of WWTPs documented in the Scopus database since 2007 (analysis based on the keywords: CF and wastewater).

Methods for estimating the emissions and effective mitigation strategies are also discussed. Energy optimization in WWTPs is addressed, emphasizing reduced consumption and increased production through co-digestion and biogas utilization. The review explores the optimization of chemical dosages and transportation to minimize emissions. Additionally, the study investigates the potential of recovering  $N_2O$  from wastewater as an energy source. The findings of study carry significant implications, supporting climate change mitigation, environmental protection, and sustainable wastewater treatment with a reduced CF. The review underscores the need for comprehensive strategies to achieve net-zero carbon conditions in municipal WWTPs.

## 2. Evolving wastewater regulations and recommendations towards a net-zero carbon condition

Fig. 2 presents a summary of historical regulations and



Fig. 2. Summary of significant wastewater management regulations/recommendations.

recommendations in the wastewater sector and outlines the recent trends that would affect the sustainability of wastewater treatment. For a long time, the regulations focused on protecting the aquatic environment, while only recently the efficient use/recovery of resources and reducing the impact on the entire environment have been postulated.

### 2.1. Protection of the aquatic environment

Based on the Public Health Act established in United Kingdom (UK) in 1875, the authorities were obliged to provide clean water, dispose and reuse all sewage. A century later (1972), the Environmental Protection Agency of United States (US EPA) published its definition of the minimum standards for secondary treatment, including major parameters, such as total suspended solids (TSS) and biochemical oxygen demand (BOD). A few of US EPA regulations brought changes in planning and designing WWTPs, including the Clean Water Act (CWA) (1972) that defined the minimum standards for discharging wastewater. Further bio-solids regulations were set in 1993 for pathogen and heavy metal content as well as safe handling and use of bio-solids [11].

Meanwhile, across the European Union (EU), the first directive concerning municipal wastewater treatment (91/271/EEC) was adopted in 1991 to protect the aquatic environment from unfavorable effects of municipal wastewater discharge. In 1998, another directive (98/15/EC) clarified the requirements in relation to the discharges from municipal WWTPs to areas which are potentially subject to eutrophication [12].

In the beginning of 21<sup>st</sup> century, the EU Water Framework Directive (WFD) was adopted, establishing a framework for water policy (2000/60/EC). The WFD established water quality standards and discharge controls to mitigate the impact of human activities on surface water quality [13]. In 2015, the United Nations set a global target recommendation through its sustainable development goals to reduce untreated wastewater by half by the year 2030 [14].

### 2.2. Efficient reuse and recovery of resources

In 2011, Water Environmental Federation (WEF) stated “WWTPs are not waste disposal facilities, but rather water resource recovery facilities that produce clean water, recover nutrients (such as phosphorus (P) and nitrogen (N)), and have the potential to reduce the nation’s dependence upon fossil fuel through the production and use of renewable energy” [15]. In 2012, WEF officially started using the novel term of water resource recovery facility (WRRF), instead of WWTP [16].

In 2017, according to the new German Environmental Agency regulation (GEA) [17], sewage sludge must be recycled to recover P in plants with population equivalent (PE) over 50,000. The regulation is aimed at closing the P cycle and replacing over 50% of the imported P.

### 2.3. Reducing impact on the entire environment

The landmark Paris Agreement [18] aimed at limiting the global warming by prohibiting the temperature rise <2 °C above the pre-industrial levels, while IPCC [7] encouraged to follow even more ambitious goal leveled at 1.5 °C. In order to achieve this milestone, countries are aiming to achieve net-zero carbon condition by 2050. In 2019, the European Green Deal was introduced, which re-evaluated the EU’s commitment to the net-zero concept with the introduction of the ‘Fit for 55’ package. The package outlines the path towards carbon-neutral transition, including short-term targets for 2030, such as a minimum reduction of 55% in GHG emissions compared to the 1990 level [19].

According to the International Energy Agency (IEA), developing and deploying clean energy technologies is crucial for achieving net-zero carbon conditions [20]. In practice, achieving net-zero globally will require a two-pronged approach, i.e. removing carbon from the atmosphere and restricting human-produced emissions. Based on the IPCC [7] definition, the net-zero carbon entails balancing anthropogenic GHG

emissions with their intentional removal from the atmosphere. Although the proposed net-zero pathways can be found in various sectors, no framework for the wastewater sector has been complied yet.

Municipal WWTPs can be a significant element of circular economy (CE) due to incorporation of resource recovery and energy production, while minimizing the environmental impact on the aquatic environment. Salminen et al. [21] presented a concept of a water-smart CE which would decrease energy and water loss and recover valuable resources and reuse wastewater. The authors suggested taxation of water abstraction and tax relief on the recycled materials and instruments targeting pollutants in wastewater.

To proactively respond to climate change and reduce GHG emissions, China committed in 2020 to peak carbon emissions by 2030 and achieve carbon neutrality by 2060, known as the “dual carbon” goal [22]. According to the European Environment Agency, the EU’s GHG emissions fell by 23.2% since 1990. The EU’s global contribution declined from 15% to 8% between 1990 and 2018. Despite this, the EU aims to cut carbon emissions by 55% by 2030 (compared to 1990 level) and achieve carbon neutrality by 2050 as per their new policy announcement [23]. Under the Paris Agreement, Australia also set its target to reduce GHG emissions in 2030 by 26–28% from 2005 level and achieve net-zero emission by 2050 [24]. Germany and the UK led 50% of the EU’s GHG emission reduction from 1990 to 2019. Romania, France, Italy, Poland, and the Czech Republic contributed 33% of the EU’s total reduction during that period. Over 29 years, the UK and Germany were responsible for nearly half of the overall net reduction in the EU-KP. In 2019, Germany, Spain, and Poland played a significant role in the EU-KP’s net GHG reduction, accounting for over half of the total absolute reduction [25]. In the same year, the Netherlands’ GHG emissions dropped by 3.2% compared to 2018, with total emissions approximately 18% lower than the 1990 levels [26].

### 3. Identification of sources of GHG emissions in wwtpds

The largest contribution to the total CF in WWTPs comes from scope 1 CF, which is due to the direct fugitive emissions from wastewater treatment and anaerobic digestion (AD) processes. A few studies which are presented in Table 1 [4,27,28] reported that the direct emissions contributed to over 60% of the total CF.

The indirect energy-related emissions, which are caused by the electricity purchased from the grid, contributed on average to the total CF with a lower share of approximately 20%. Those emissions play a minor role especially in the countries, such as Finland and Austria, that have a high share of renewable energy sources [27,28]. In contrast, Hu et al. [29] analyzed the CF of WWTPs in China and found a lower share of 29% for the direct emissions, while 45% of the total CF was related to

sludge handling. This high share resulted from a large number of studied WWTPs performing sludge incineration and landfilling. In the case of plants without on-site electricity production (via e.g., biogas production through AD and energy co-generation), a dominant percentage share can also be attributed to the indirect emissions [30]. The indirect emissions related to chemicals consumed at the plant and transportation showed a marginal effect on the CF of WWTPs [4].

Kadam et al. [31] presented a discussion on achieving carbon neutrality in municipal WWTPs based on feasible technical aspects. Their proposed plant incorporates various advanced primary treatment techniques, leading to efficiently organic recovery and reducing the oxygen demand for oxidation of organic compounds. Consequently, biogas production during biological processes significantly increases. The optimized approach also involves converting concentrated primary sludge to biomethane through anaerobic digestion. Additionally, H<sub>2</sub> gas derived from N upgrades plays a crucial role in enhancing biomethane quality by reducing CO<sub>2</sub> content in the biogas. These findings demonstrate the potential of municipal WWTPs to achieve high process efficiency and energy utilization. To achieve a net-zero carbon condition, a hybrid system integrating both GHG emission reduction solutions and energy recovery is suggested [32]. Specific solutions for the three scopes of emissions are discussed in Sections 5 to 7.

Wu et al. [33] performed a comprehensive CF analysis of different wastewater treatment configurations. The authors reported that scope 1 emissions accounted for 23–83% of the total CF in different configuration scenarios. Scope 1 emission could be the primary contributor to CF for configurations with anaerobic processes. This is due to the reduction in scope 2 and scope 3 emissions resulting from energy recovery and reduced sludge production. Recent studies by Yao et al. [34] reported N<sub>2</sub>O emissions alone to exceed 50% of the total CF based on site-specific monitoring of BNR WWTPs employing such configurations as A/O, anaerobic-anoxic-oxic (A<sub>2</sub>O), and SBR. The contribution of scope 2 emissions is also highly variable and can range from 14 to 68% of the total CF of WWTP (Table 1). Scope 2 emissions could be minimized by reducing the energy consumption, but most wastewater treatment configurations with relatively low GHG emissions involve energy recovery, which offsets the scope 2 emissions. Scope 3 emissions play a minor role in the overall GHG emissions in WWTPs, contributing only 1–13% to the overall CF.

#### 3.1. WWTP emission hotspots

Fig. 3 shows the potential sources of different GHGs that are emitted directly and indirectly. The preliminary stage of wastewater treatment is known as a potential source of CH<sub>4</sub> emissions. Anaerobic conditions often prevail in sewer systems, and intense CH<sub>4</sub> fluxes have been

**Table 1**  
CF distribution in WWTPs - overview.

Reference -number of studied plants	Contribution to the total CF (%)										
	Scope 1					Scope 2	Scope 3				
	Bioreactor	AD	Recipient	Sludge handling	Total			Grit and screening handling	Chemicals	Sludge handling (3rd parties)	Transport
Parravicini et al. [27] – 2 WWTPs	42.9	26.3	9.1	–	78.4	15.5	–	4.2	–	1.7	5.8
Hu et al. [29] – 344 WWTPs	29	–	–	–	29	26	–	–	45	–	–
Maktabifard et al. [30] – 6 WWTPs	58	3	1	10	72	26	1	1	–	–	2
Awaitey [28] – 4 WWTPs	52	1.5	2	11	66.5	18	0.5	8	–	7	15.5
Min	29	3	0	0	29	15.5	0	0	0	0	0
Max	58	26.3	9.1	11	78.4	26	1	8	45	7	15.5
Average	45.5	10.3	4	10.5	61.5	21.4	0.8	4.4	15	2.9	17.1
Deviation	10.9	11.3	3.6	0.5	19.2	4.7	0.3	2.8	21.2	2.9	16.8

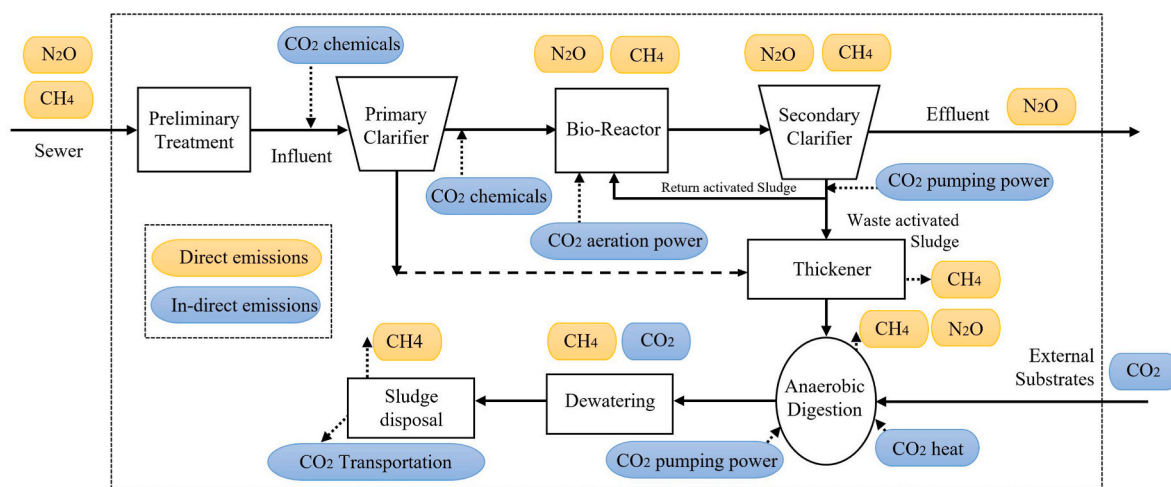


Fig. 3. Sources of the direct and indirect GHG emissions in WWTPs.

reported at the headworks of the WWTP, especially in aerated grit chambers. Other potential CH<sub>4</sub>-forming spots are primary clarifiers, and then anaerobic compartments in the biological reactors. CH<sub>4</sub> can subsequently be stripped in the aerobic compartments of the reactors [35]. Due to the leakage of gases from internal facilities in the plant, biogas production installations may become significant CH<sub>4</sub> sources. Aerobically stabilized sludge, when stored for several months, can also contribute to CH<sub>4</sub> emissions due to the fact that anaerobic conditions start to prevail in the long-term perspective [36]. The effluent BOD load can contribute to further direct CH<sub>4</sub> emissions from the natural processes in recipients [27].

The GHG emissions from the preliminary treatment and sludge treatment stages do not attract as much attention as the emissions from the biological stage employing biological nutrient removal (BNR). Bio-reactors have been recognized as the main source of GHG emissions in WWTPs. Campos et al. [37] reported that, the dominant N<sub>2</sub>O production

occurs in the bioreactors (90%) while for CH<sub>4</sub> emissions the sludge line with AD was responsible for over 70% and the remaining portion originates from bioreactors. N<sub>2</sub>O produced under favorable conditions (either anoxic or aerobic) is emitted via either saturation-induced liquid-gas transfer or stripped to the air via aeration [38]. The effluent N load also becomes the source of direct N<sub>2</sub>O emissions in the recipients.

### 3.2. N<sub>2</sub>O emissions

WWTPs are considered significant anthropogenic sources of atmospheric N<sub>2</sub>O, contributing 3–10% of the total emissions [5]. The latest IPCC report [7] shows a progressive increase in the atmospheric N<sub>2</sub>O concentration, which has been levelled at the rate of 0.95 ppb/year for the last decade (2010–2019). The report emphasizes that wastewater treatment is the fourth largest sector responsible for N<sub>2</sub>O emissions. The N<sub>2</sub>O emissions from this sector increased from 0.2 Tg N/year to 0.35 Tg

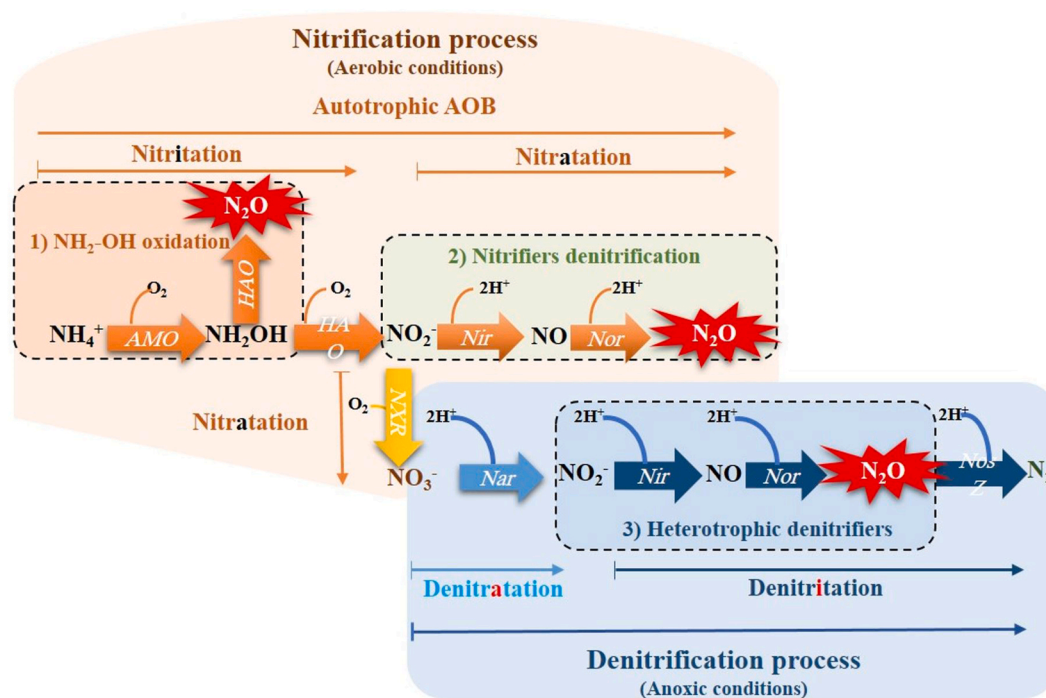


Fig. 4. Biological pathways of N<sub>2</sub>O production and sink in the bioreactors (responsible genes: AMO: ammonia monooxygenase; HAO: hydroxylamine oxidoreductase; NXR: nitrite oxidoreductase; Nar: nitrite oxidoreductase; Nir: nitrite reductase; Nor: nitrite reductase; NosZ: Nitrous oxide reductase).

N/year between the 1980s and 2010s.

The  $N_2O$  emissions occur dynamically and normally remain beyond the control of the plant operators. Even small changes, such as dissolved oxygen (DO) variations, can significantly affect the liquid  $N_2O$  production by microorganisms [3].

### 3.2.1. Identification of the dominant pathways

$N_2O$  production pathways involve three microbiological reactions, which require either aerobic or anoxic conditions (Fig. 4). In two pathways mediated by ammonia oxidizing bacteria (AOB),  $N_2O$  can be an intermediate product of hydroxylamine ( $NH_2OH$ ) oxidation and the final product of autotrophic denitrification. Moreover,  $N_2O$  is an intermediate product of denitrification by heterotrophs. If that process is not disturbed,  $N_2O$  is further reduced to dinitrogen ( $N_2$ ) in the final step of denitrification and this pathway can thus be a sink of  $N_2O$  [39]. Based on a literature review of  $N_2O$  emissions from WWTPs, Vasilaki et al. [39] formulated four main operational factors responsible for liquid  $N_2O$  production in the conventional nitrification-denitrification systems, including (i) low DO conditions, nitrite ( $NO_2^-$ ) accumulation and changes in the ammonium ( $NH_4^+$ ) concentration in aerobic compartments, (ii) low chemical oxygen demand (COD) to N ratio and  $NO_2^-$  accumulation in anoxic bioreactors, (iii) alternation of the anoxic to aerobic environments in intermittent compartments, (iv) sudden fluctuations (shocks) in the processes conditions.

Diverse contributing factors and dynamic reactions can occur simultaneously in the bioreactors, which are beyond operational control in full-scale WWTPs. Furthermore, heterotrophic denitrification can consume liquid  $N_2O$  produced via the different pathways. Regardless of the various production and consumption pathways of liquid  $N_2O$ , the stripping process should be considered as the process leading to  $N_2O$  emissions.

### 3.2.2. Additional sources of $N_2O$ emissions in WWTPs

Although the direct  $N_2O$  process emissions released from bioreactors has the highest contribution to the total CF of WWTPs, a minor fraction of  $N_2O$  is also emitted from other units of the plant. Hwang et al. [40] performed  $N_2O$  measurements in different units of a WWTP, including the primary clarifier and secondary clarifier, as well as the sludge thickener and anaerobic digester. Small quantities of  $N_2O$  (0.012 g  $N_2O$ /kg TN) were produced in the digester, accounting for less than 1% of the total  $N_2O$  emissions. Regarding the primary and secondary clarifier,  $N_2O$  emission factors (EFs) of 0.22–0.26 g  $N_2O$ /kg TN were reported based on the full-scale measurements of Hwang et al. (2016). For comparison, Solís et al. [41] applied an EF of 0.01 kg  $N_2O$ -N/kg TN to predict the direct  $N_2O$  emissions from sludge storage, assuming uncovered storage throughout the year. Another study by Caniani et al. [42] found interesting results in the disinfection unit, observing high  $N_2O$  EF of 0.008 kg  $CO_{2e}$ /kg COD, due to the interaction between disinfecting agent and  $NH_2OH$ .

### 3.3. $CH_4$ emissions

The latest IPCC [7] report shows a progressive increase in the atmospheric concentration of  $CH_4$ , which has been levelled at the high rate of 9.3 ppb/year in the last decade. On a global scale, the wastewater sector accounts for approximately 5–7% of anthropogenic  $CH_4$  emissions, ranking fifth after livestock (32%), oil and gas (25%), landfills (13%), and coal mining (11%) [43]. In the United States,  $CH_4$  emissions from wastewater treatment increased from 10% to 14% between 1990 and 2019 [44]. A recent study by Song et al. [45] utilized the updated data from municipal WWTPs in US and estimated  $10.9 \pm 7.0$  MMT  $CO_{2e}$ /year of annual  $CH_4$  emissions which is about the double amount of emissions estimated by IPCC [36] (4.3–6.1 MMT  $CO_{2e}$ /year).

$CH_4$  is mainly produced during the anaerobic degradation of organic matter in bioreactors and digesters. The effectively utilized  $CH_4$  can assist the plant to move towards energy neutrality and indirectly offset

$CO_2$  emissions. On the other hand,  $CH_4$  would increase the plant CF when emitted through incomplete combustion or leakage [46].

Studies at a regional level are usually based on  $CH_4$  EFs [47]. Case studies of  $CH_4$  emissions at the plant level are less frequent in the literature [48]. Zhao et al. [49] found that the  $CH_4$  emissions from municipal WWTPs in China were over three times higher compared to the US plants. Although an empirical approach is highly uncertain, it is still a suitable way to estimate  $CH_4$  emissions from WWTPs. Zhang et al. [6] concluded that it would be crucial to perform extensive measurements under various treatment processes, scales, and locations to obtain more accurate  $CH_4$  emission predictions.

A full-scale study of Ribera-Guardia et al. [50] reported the long-term  $CH_4$  emissions from aerobic compartments of a plug-flow bioreactor. The peak  $CH_4$  emissions were found in the first aerobic zone, while decreasing towards the end of the bioreactor. The authors assumed that  $CH_4$  was produced under anaerobic conditions in the initial stages of the plant and then stripped in the aerobic compartment of bioreactors. The highest liquid  $CH_4$  concentrations were detected in the plant influent (0.55 mg  $CH_4$ /l) and reject water from the anaerobic digesters (0.52 mg  $CH_4$ /l).

Hwang et al. [30] determined the  $CH_4$  EFs in different units of a municipal WWTP in South Korea. The sludge thickening process produced the largest  $CH_4$  emissions (2.09 g  $CH_4$ /kg BOD) considerably more than aeration basin (0.72 g  $CH_4$ /kg BOD), primary clarifier (0.26 g  $CH_4$ /kg BOD) and secondary clarifier (0.67 g  $CH_4$ /kg BOD).  $CH_4$  was primarily released from the digesters with a high  $CH_4$  EF of 227 g  $CH_4$ /kg BOD. Wei et al. [51] analyzed four scenarios of sludge treatment and disposal. The highest contribution to  $CH_4$  emissions was incineration (45.1%), followed by sanitary landfills (23%), land utilization (17.7%), and building materials (14.2%).

In a plant-wide model, implemented by Solís et al. [52],  $CH_4$  emissions were calculated by a modified Anaerobic Digestion Model No. 1 (ADM1) [53]. Fugitive emissions were included as 2.7% of the produced biogas which was un-combusted in combined heat and power (CHP) units or slipped from anaerobic digester. The remaining amount of biogas was assumed to be completely combusted and  $CH_4$  was fully converted to  $CO_2$  while generating heat and electricity. It was assumed that  $CH_4$  from the digester effluent was fully stripped to the atmosphere. Furthermore, the direct emissions due to sludge storage were estimated at 8.7 kg  $CH_4$  per ton of volatile solids (VS). Overall, for the studied WWTP with the flow rate of 21,000  $m^3$ /d, the CF was 19,000 kg  $CO_{2e}$ /d and  $CH_4$  emissions contributed 5.8% to the total CF.

## 4. Estimation methods of GHG emissions

### 4.1. GHG emissions inventories

**European inventory:** The annual EU GHG inventory 1990–2019 report [54] contains the  $CH_4$  and  $N_2O$  emissions from the wastewater sector. Both  $N_2O$  and  $CH_4$  emissions account for 0.6% of the total EU GHG emissions in 2019 with 35.5 Mt  $CO_{2e}$  (domestic wastewater) and 9.3 Mt  $CO_{2e}$  (industrial wastewater). Between 1990 and 2019, the total emissions from wastewater treatment in EU decreased by 43.7%. Due to the application of new wastewater treatment processes,  $CH_4$  and  $N_2O$  emissions from domestic wastewater decreased by approximately 50% and 17%, respectively [54].

**US inventory:** According to the US GHG Emissions and Sinks 1990–2020 Inventory Report [55], the wastewater treatment sector significantly contributed to the total  $N_2O$  country-wide emissions. It was the second largest anthropogenic source of  $N_2O$  emissions in 2020, levelled at 23.5 Mt  $CO_{2e}$ .

The wastewater sector contributed to 5.5% of  $N_2O$  fluxes, which resulted in 0.4% of the total GHG emission in the US. Comparing to 1990, a 41.8% increase was reported due to the country's growing population and a peaking protein consumption. The wastewater sector was ranked seventh of  $CH_4$  emission sources with its 18.3 Mt  $CO_{2e}$

fluxes. During the period of 1990–2002, CH<sub>4</sub> emissions remained stable, but a decreasing trend was observed later due to a reduction in the use of on-site septic systems [55].

**Australian inventory:** In the quarterly updated national inventory GHG (Australia's Greenhouse Accounts [56]), the levels of ozone layer depleting gases are given for different sectors, including wastewater handling. In 2010, wastewater treatment contributed to the total Australia's emissions in 0.51%, reflecting 0.8 Mt CO<sub>2e</sub>. After a decade, the emissions increased to 0.9 Mt CO<sub>2e</sub> constituting 0.72% of the total GHG emissions in the continent.

**Chinese inventory:** The most recent Chinese national GHG report from 2014 revealed that N<sub>2</sub>O emissions from WWTPs were 1.1 Mt of N<sub>2</sub>O, accounting for 5.6% of the national N<sub>2</sub>O emissions. CH<sub>4</sub> emissions from wastewater treatment were 2.7 Mt of CH<sub>4</sub>, accounting for 4.9% of China's total CH<sub>4</sub> emissions [6].

#### 4.2. Direct measurements

There are various methods on *in-situ* measurements of GHG emissions that can be applied in WWTPs. In hermetic facilities, the direct GHG emissions can be analyzed continuously in the ventilation air by combining the concentration results with the airflow parameters [35]. In the case of open-to-air plants, collection of flux gas requires a dedicated sampling technique. In order to aggregate samples in open-surface tanks, different gas hoods and flux chambers are installed in multiple measurement points to address spatial and temporal variability along the wastewater train. In both cases, liquid N<sub>2</sub>O concentrations should be simultaneously examined [57] and they need to be complemented by the analysis of conventional qualitative and quantitative operating parameters. The analytical techniques for measuring GHG concentration in collected flux-gas samples include Fourier Transformed Infrared Spectroscopy (FTIR) or Gas Chromatography with a Thermal Conductivity Detector (GC TCD).

Duan et al. [58] provided a critical review on using isotope technology for direct N<sub>2</sub>O measurements, including both natural abundance and labelled isotope approaches. However, the authors emphasized that the accuracy and reliability of these techniques requires further improvement.

In order to reliably validate N<sub>2</sub>O emission models, data from full-scale measurement campaigns are essential. However, the number of plants which measure N<sub>2</sub>O emissions is currently very limited [59]. The long-term (yearly) N<sub>2</sub>O monitoring data are available in the literature for Viikinmäki WWTP in Helsinki [60] and Kralingseveer WWTP [35].

#### 4.3. CF calculation tools

General requirements that are dedicated to CF calculation can be found in International Organization for Standardization (ISO) 14,064 standard [61]. The IPCC (2006) with 2019 refinement for the national GHG inventories [36] is a frequently applied guidance note for the estimation of GHG emissions from WWTPs.

Life cycle assessment (LCA) is a broad technique to systematically evaluate multiple environmental impacts associated with municipal water cycle infrastructure, including wastewater collection and treatment [62]. The CF is part of LCA that quantifies a negative impact on the climate change. The LCA can be an accurate tool for calculating the indirect GHG emissions including high level of details in inventory data [63]. The direct GHG releases to air from the wastewater and sludge lines of WWTPs are incorporated via additive calculation procedures, often supported by secondary data estimations sourced from, either empirical or mechanistic models as well as *in-situ* measurement campaigns [63].

Several tools are available specifically for the CF assessment of WWTPs, usually as open-source files or applications. These are designed to deliver CF results based on the user's data input comprising operational parameters of the analyzed WWTP or its part. Table 2 shows an

**Table 2**

Overview of the CF assessment tools for WWTPs.

Tool name (if available)	Country of origin	Year	Methodology	Open source
1 Quantifying the GHG emissions of WWTPs (as part of a thesis project)	Netherlands	2009	ISO	YES - source code for MatLab
2 Biosolids Emissions Assessment Model (BEAM)	Canada	2010–2011	IPCC, WRI	YES - Excel
3 Worldbank Organization - Sustainable Urban Energy and Emissions Planning Toolkits - Energy Balance and GHG Inventory Spreadsheet	USA	2011–2012	IPCC	YES - Excel
4 Waste Sector GHG Protocol Calculation Tool	France	2013	GHG Protocol	YES - Excel
5 Energy Performance and Carbon Emissions Assessment and Monitoring (ECAM)	Germany	2015	IPCC	YES - online
6 GESTABoues for wastewater treatment sludge management	France	2016	Own - based on literature	YES - after free registration
7 C-FOOT-CTRL	UK/Greece	2019	Own - based on literature	NO
8 Queensland Water GHG Calculator	Australia	2010–2019	National Greenhouse and Energy Reporting Standard	YES - excel
9 Calculation of the CF from Swedish WWTPs (SVU 12–120)	Sweden	2012–2021	Own - based on literature	YES - excel

overview of CF calculation tools specifically dedicated to WWTPs. The Carbon Footprint Calculation Tool (CFCT) [64] has been utilized frequently in the literature [4,30] which is an adjustable spreadsheet developed originally for Swedish case studies [65] and has been updated since then.

When planning the CF calculation for a WWTP, two main approaches for setting boundary conditions of the CF analysis can be applied. The first method covers a full life cycle of a WWTP within the construction, operation and demolition stages [66]. Another method would be an in-depth analysis of the operation stage with detailed considerations of process emissions [4].

Overlooking the construction and demolition phases of WWTPs and supporting buildings is a common simplification in LCA. The impacts of these phases are usually considered insignificant compared to the emissions from the operational stage [67]. The inclusion of new aspects in WWTPs, such as energy and resource recovery, would broaden the

scope of carbon accounting. Therefore, Li et al. [46] proposed a system expansion approach, while assuming that the products recovered from wastewater, such as struvite, vivianite, biodiesel, bioplastics, biochar and protein, should substitute similar products on the market. This would result in offsetting carbon emissions from the system.

#### 4.4. Empirical EFs

Simple empirical models are the most common approach to estimate N<sub>2</sub>O emissions from wastewater and N<sub>2</sub>O EFs are advantageous for providing a better insight into the WWTPs CF. It is important to note that the specific functional units (FUs) used to report N<sub>2</sub>O EF might vary in the literature. The influent total Kjeldahl nitrogen (TKN) load was used most frequently, but the total removed N load or influent NH<sub>4</sub> load have occasionally been reported in the literature as well (Table 3).

The latest IPCC guidelines [7] recommended 1.6% kg N<sub>2</sub>O/kg influent N-load as the EF based on the results of various monitoring campaigns. The reported N<sub>2</sub>O EF for other processes, such as nitrite “shunt” or deammonification, are typically higher than full nitrification/denitrification. For example, Gustavsson and La Cour Jansen [68] estimated 6% kg N<sub>2</sub>O/kg NH<sub>4</sub> removed, EF for the nitrification process, while the EF for partial nitrification anammox process was 3% kg N<sub>2</sub>O/kg N removed [69].

Continuous, long-term monitoring campaigns can substantially improve the N<sub>2</sub>O EF accuracy [70]. A high inconsistency in the annual EFs (0.1–8% of the N load) was found after monitoring fourteen

**Table 3**  
Measured and estimated N<sub>2</sub>O EFs from WWTPs in different studies.

Reference	N <sub>2</sub> O EF benchmark			Type of study
	kg N <sub>2</sub> O/kg TKN	kg N <sub>2</sub> O/kg N <sub>removed</sub>	kg N <sub>2</sub> O/kg NH <sub>4</sub> removed	
Solis et al. [41]	0.55%			Practical modelling
Li et al. [73]				Practical modelling
Maktabifard et al. [59]	0.94%			Practical modelling
Zaborowska et al. [74]	1.6%			Practical modelling
Domingo-Félez and Smets [75]			1.2–4.6%	Theoretical modelling
Massara et al. [76]	1–11%			Theoretical modelling
Pocquet et al. [77]			1%	Theoretical modelling
Ni et al. [78]	0.69–3.5%			Practical modelling
Foley et al. [79]		1.57%		LCA
Gruber et al. [80]	1–2.4%			Long term monitoring campaign
Sun et al. [81]	0.2–1.6%			Sampling measurements
Kosonen et al. [60]	1.9%			Long term monitoring campaign
Wang et al. [82]	0.095–3.44%			Online measurements
Hwang et al. [40]	1.606%			Sampling measurements
Duan et al. [83]	0.58%			Long term monitoring campaign
IPCC [36]	1.6%			Inventory report
EEA [84]	0.035%			Inventory report
NGER [85]	0.7–0.8%	1%		Inventory report
WERF [86]			0.01–2.5%	Inventory report
IPCC [87]			0.32–0.5%	Inventory report

full-scale WWTPs in Switzerland, but the EFs were strongly correlated with the effluent NO<sub>2</sub><sup>-</sup> concentrations. The authors proposed a national EF calculated from the weighted (weights based on the fraction of N loading in the country-scale) EFs of carbon removal (EF: 0.1–8% depending on the expected variability in plant performance), nitrification only (EF: 1.8%), and full N removal (EF: 0.9%). This approach allowed to estimate country-specific N<sub>2</sub>O emissions from WWTPs. In Switzerland, the average EFs and total annual N<sub>2</sub>O emissions ranged between 0.9 and 3.6% and 410–1690 ton N<sub>2</sub>O (corresponding to 0.3–1.4% of the total GHGs), respectively [70].

The fixed EF approach neglects the impact of operational conditions and wastewater characteristics on N<sub>2</sub>O emissions. Valkova et al. [3] suggested that the N<sub>2</sub>O EF by IPCC overestimates N<sub>2</sub>O emissions for the plants with the high efficiency of total nitrogen (TN) removal. The authors proposed that the annual direct N<sub>2</sub>O emissions can be predicted based on the annual average TN removal efficiency. In ten Austrian WWTPs with high N removal efficiency (83–92%), the N<sub>2</sub>O EF of 0.12% ± 0.1% kg N<sub>2</sub>O/kg TKN was found. The N<sub>2</sub>O EFs do not capture emission dynamics, potentially leading to overestimation or underestimation of emissions [71]. To improve the accuracy of N<sub>2</sub>O emissions and CF assessment in process optimization and mitigation strategies for individual WWTPs, local measurements and mathematical models are preferred over fixed EFs. Table 3 shows significant variations in N<sub>2</sub>O EFs depending on the estimation methodology, plant location, and study conditions. The scale of estimation, from global inventories through individual facilities, also impacts the results. Nayeb et al. [72] found high uncertainty in the CF estimation using national-scale inventories. Maktabifard et al. [30] demonstrated over 50% uncertainty in total CF estimation when considering a wide range of N<sub>2</sub>O EFs from literature. High uncertainties were especially observed in the facilities dominated by the direct emissions.

#### 4.5. Mathematical models

Mechanistic models are considered powerful tools for determining liquid N<sub>2</sub>O production. These models can help identify the dominant N<sub>2</sub>O production pathways and mitigation strategies. In comparison with empirical models, the use of mechanistic models may significantly reduce the uncertainty of the total CF results, as the CF results are highly sensitive to N<sub>2</sub>O emissions [59].

Single-pathway (AOB denitrification/NH<sub>2</sub>OH oxidation) and two-pathway models have been suggested for N<sub>2</sub>O production by AOBs, while heterotrophic denitrification could be modelled as three or four-step process [88]. These models are added as extensions to existing activated sludge models (ASMs). For example, Kim et al. [89] combined the ASM for Nitrogen (ASMN) [90] and ADM1 [53] in the ASMN for GHGs (ASMN\_G). Blomberg et al. [91] extended the ASM3 [92] with a dynamic N<sub>2</sub>O prediction and implemented the model in a full-scale WWTP in Finland. Zaborowska et al. [93] extended the ASM2d (ASM2d-N<sub>2</sub>O) and included both N and P removal processes to model a full-scale WWTP in Poland. Research attention has primarily focused on modelling N<sub>2</sub>O emissions, but efforts are being made to develop models considering CO<sub>2</sub> and CH<sub>4</sub> emissions from wastewater (ASMs) and sludge treatment (ADM). Mannina et al. [94] proposed a simple mass balance-based model with N<sub>2</sub>O from nitrification and denitrification. A modified Benchmark Simulation Model No. 2 (BSM2) incorporated GHG emissions were considered (N<sub>2</sub>O and CO<sub>2</sub> from AS, CO<sub>2</sub> and CH<sub>4</sub> from AD) along with the indirect emissions from the electricity and chemicals [95,96]. These models investigated the influence of different operational scenarios on the effluent quality, operational cost and GHG emissions. The DO control in the aerobic zone was a commonly studied strategy for GHG mitigation strategy. Other model-based studies considered the influent COD/N ratio [94], TSS control in the primary clarifier and CEPT [95,97], MLR ratio [96] and sidestream deammonification [74]. The advantage of the modelling approach was the possibility to take into account the processes interactions and identification of



trade-offs between conflicting objectives [98]. In the approach proposed by Arnell et al. [97], the plant-wide model was combined with LCA to account for the global environmental impact due to external resource use.

The models of  $N_2O$  production/consumption should be integrated with a stripping model to predict  $N_2O$  emission. Baresel et al. [99] initially applied the saturation-induced liquid-gas transfer and the stripping  $N_2O$  emission model in the anoxic and aerobic compartments, respectively. These models were adopted in several subsequent studies [59,91,93].

The inclusion of all the known  $N_2O$  production pathways may result in over parameterized complex models. These models require more calibration efforts due to a higher number of coefficients. Moreover, calibration of the models by using short-term measurement data would reduce the accuracy and increase the uncertainty of the model [59].

Although mechanistic models have explicit advantages in determining  $N_2O$  mitigation strategies for WWTPs, their practical use in GHG inventories is still limited. Another emerging mathematical model is Machine Learning (ML), which can transform WWTP data into knowledge. By training ML models with outputs from ASMs, the prediction capabilities can be improved. These trained ML models are highly effective for online monitoring and providing decision support and are widely used in industry. Limitations of ML approaches may arise from data scarcity for learning and testing. To address this, Mehrani et al. [100] proposed a hybrid model combining a mechanistic model with ML to simulate  $N_2O$  production in a sequencing batch reactor. Szelag et al. [101] examined various ML models for  $N_2O$  emission prediction, using global sensitivity analysis to select a ML model as an alternative to the mechanistic model.

## 5. Mitigation of GHG emissions via process modification

From the CF perspective of WWTPs, the direct emissions from the treatment processes usually play the most significant role. This chapter presents  $N_2O$  mitigation strategies in order to reach a net-zero condition and explores the implementation of innovative removal processes, source separation, and nutrient recovery for decarbonizing WWTPs.

### 5.1. Approaches to $N_2O$ mitigation strategy

$N_2O$  mitigation strategies are mainly focused on [83]: (i) Optimizing aeration mode and DO set-point, (ii) Preventing DO gradients via mixing optimization, (iii) Avoiding  $NH_4^+$  peaks, (iv) Preventing  $NO_2^-$  accumulation, (v) Ensuring complete denitrification by supplying sufficient carbon source or/and increasing hydrolysis in the primary clarifiers.

Duan et al. [83] evaluated  $N_2O$  production by adopting a multi-pathway mechanistic model. The employed multi-criteria strategy in an Australian plant, which was focused on reducing DO levels, resulted in 35% mitigation of  $N_2O$  emissions with the EF reduced from 0.89 to 0.58% of the influent TN load. This could mainly be associated with the declines in  $N_2O$  generated via the  $NH_2OH$  oxidation pathway. Based on lower DO levels simultaneous nitrification and denitrification was encouraged and thus nitrite accumulation was minimized, leading to less  $N_2O$  production via AOB denitrification pathway. Another study by Mampaey et al. [102] reported a decline of over 50% in  $N_2O$  emissions when one-stage granular SHARON (partial nitrification) reactor cycles were reduced by 1 h, which also falls in the mitigation category of optimizing aeration.

One of the most effective ways to recognize  $N_2O$  mitigation strategies is testing different operational modes [39]. Rodriguez-Caballero et al. [103] suggested that the optimum control strategy in sequencing batch reactors (SBRs) would be shortening aerobic-anoxic cycles (20 min of aeration with a short anoxic phase).

### 5.2. Novel processes for N removal

Nitrogen removal systems, which are based on anaerobic ammonia oxidation (anammox), can be categorized into partial nitrification-anammox (PN/A), simultaneous PN/A and denitrification (SNAD), partial denitrification-anammox (PD/A), and denitrifying anaerobic methane oxidation (DAMO)-anammox (DAMO/A) systems (Fig. 5). PN/A and SNAD can effectively treat ammonia-rich wastewater, while reducing energy consumption and sludge production compared to the traditional nitrification-denitrification processes. In contrast, PD/A and DAMO/A are used to treat  $NO_3^-$ -rich wastewater, with a focus on reducing energy consumption (no oxygen demand and reducing C/N ratio to  $< 3$ ) and achieving sustainable removal efficiencies [104]. Therefore, anammox-based systems are becoming a promising technology in WWTPs for energy neutrality [105].

There are three main pathways of  $N_2O$  production in anammox-based systems, including  $NH_2OH$  oxidation ( $(NH_4^+ \rightarrow NH_2OH \rightarrow N_2O)$ ) and autotrophic denitrification both mediated by AOB ( $NO_2^- \rightarrow NO \rightarrow N_2O$  or  $NH_2OH \rightarrow N_2O$  or  $NH_2OH + NO \rightarrow N_2O$ ), as well as heterotrophic denitrification ( $NO_3^- \rightarrow NO_2^- \rightarrow NO \rightarrow N_2O$ ) mediated by diverse groups of denitrifiers. The  $N_2O$  consumption ( $N_2O \rightarrow N_2$ ) is a unique  $N_2O$  degradation (nosZ-dominated) catalyzed by denitrifiers [106]. Other microorganisms present in anammox-based systems, such as anaerobic ammonia oxidizing bacteria (AnAOB), dissimilatory nitrate reduction to ammonia (DNRA) heterotrophs and DAMO autotrophs do not contribute to  $N_2O$  emission in anammox systems [107–109].

**PN/A.** The reported overall  $N_2O$  emissions in PN/A systems are highly variable (dependent on the scale) and range from 0.08% to 6.6% of N-load [5,110]. The mitigation of  $N_2O$  emission in PN/A systems can be achieved by adjusting various operation parameters, such as  $NO_2^-/NH_4^+$  and C/N ratios, aeration strategies, DO set points, temperatures, and pH [111,112].

$N_2O$  emission is dominated by  $NH_2OH$  oxidation pathway at low DO levels ( $< 0.5$  mg  $O_2/L$ ), low inorganic carbon, high pH or low  $NO_2^-$  concentration, while AOB denitrification pathway dominates at higher inorganic carbon, lower pH or higher  $NO_2^-$  concentration [5]. Properly selecting intermittent aeration strategies can lead to frequent shifts in the predominating pathways of  $N_2O$  production and help mitigate  $N_2O$  emissions [113]. Beier et al. [114] aimed to predict the production of  $N_2O$  in a side-stream system using a novel modelling approach. During the periods when the system was aerated, the average EF was achieved 0.05 mg  $N_2O/mgNH_4^+$ . During the anoxic phases, the production factor was determined to be 33%. The overall EF of oxidized  $NH_4^+$  was found to be 27%, with denitrification being the primary source of  $N_2O$  production.

A high  $NO_2^-/NH_4^+$  ratio can result in an increased  $N_2O$  production [115], but a low  $NO_2^-$  and high  $NH_4^+$  loading is expected to minimize  $N_2O$  production rates [116].

**SNAD.**  $N_2O$  is produced through the oxidation of  $NH_2OH$  by AOB and reduction of  $NO_2^-$  to NO, then to  $N_2O$  under aerobic conditions [117].  $N_2O$  production is also linked to heterotrophic denitrification under anoxic conditions [118]. The  $N_2O$  production in SNAD systems was found to be slightly lower (by 4%) compared to PN/A systems due to lower  $NO_2^-$  accumulation [82]. This reduction may be due to the consumption of  $NO_2^-$  and/or NO during anammox metabolism [119]. Moreover, complete heterotrophic denitrification by the NosZ enzyme was identified as a promising strategy for mitigating  $N_2O$  emissions [120]. For comparison,  $N_2O$  emission was mitigated by 80% and 70% in a SNAD biofilter compared to traditional nitrification and nitrification-denitrification biofilters [117].

**PD/A.** In PD/A systems, diverse microbial communities compete for  $NO_2^-$ . The presence of AnAOB and heterotrophic denitrifying/Dissimilatory Nitrate Reduction to Ammonium (DNRA) in this system is crucial for  $N_2O$  mitigation.  $N_2O$  is not produced during anammox metabolism and should not be produced by heterotrophs catalyzing partial denitrification (PD) ( $NO_3^- \rightarrow NO_2^-$ ) (Fig. 5) [121]. Low  $N_2O$  emissions were

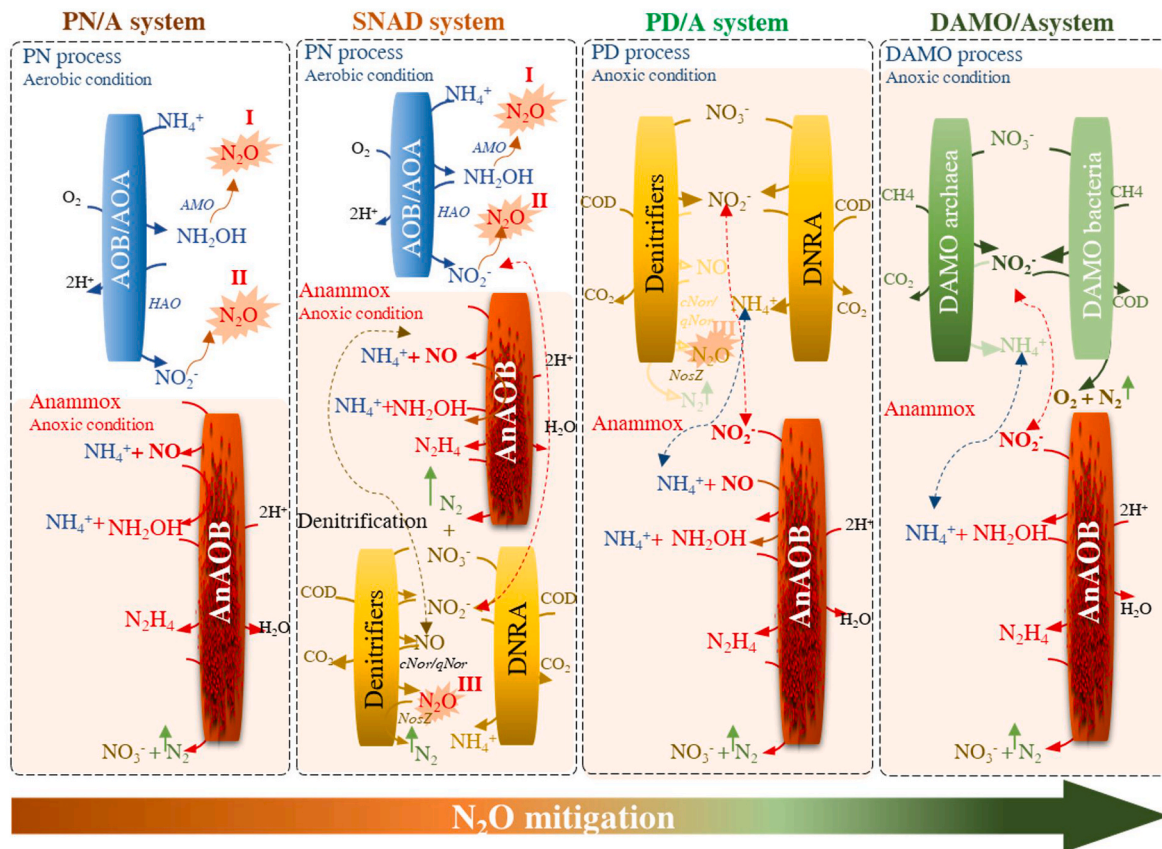


Fig. 5. Anammox based N removal systems.

observed in PD/A systems either with or without COD addition. Mitigation of  $\text{N}_2\text{O}$  was successful ( $\text{EF} = 0.002\%$ ) with low levels of COD addition and remained low during long-term PD/A performance [122]. However, adding COD resulted in approximately 50% reduction in  $\text{N}_2\text{O}$  emissions compared to the period without COD addition, indicating further reduction through the activity of *NosZ* enzyme in denitrifiers [122]. The  $\text{N}_2\text{O}$  conversion to  $\text{N}_2$  gas was comparable to  $\text{N}_2\text{O}$  production from  $\text{NO}_2^-$  by heterotrophs under COD-limited conditions [123]. Thus, PD/A and DAMO/A have gained a significant attention for applications in WWTP as an alternative process to nitrification-denitrification and pursuing efficient N removal in a sustainable way. The most important challenge for a stable PD/A performance is a balance in  $\text{NO}_2^-$  by the activities of coexisting denitrifiers and AnaAOB [123]. Moreover, low process temperatures and low influent C/N ratios could limit the startup of PD/A systems [121]. In contrast, heterotrophs can produce  $\text{N}_2\text{O}$  when PD is not successfully performed to only produce  $\text{NO}_2^-$  during unbalanced COD addition [124]. For example, 30% of total  $\text{N}_2\text{O}$  was observed to be produced in the AnaAOB-dominated anoxic zone, potentially by heterotrophic denitrification or other unknown pathways [125]. Proper control of denitrifiers through adjusting COD addition (proper C/N ratio and COD type) appears to be crucial for  $\text{N}_2\text{O}$  mitigation in PD/A systems [126]. Comprehensive research should be taken for a deep understanding PD/A microbial metabolism and microbial growth rates, which are needed to improve the efficiencies of N and COD removal rates in WWTPs in the future.

**DAMO/A.** DAMO microorganisms can be classified as DAMO archaea and DAMO bacteria. DAMO-archaea can reduce  $\text{NO}_3^-$  to  $\text{NO}_2^-$  (substrate for AnaAOB) by  $\text{CH}_4$  oxidation under anoxic conditions, leading to  $\text{N}_2\text{O}$  and  $\text{CH}_4$  mitigation. In contrast, DAMO-bacteria perform full denitrification ( $\text{NO}_3^- \rightarrow \text{N}_2$ ) by utilizing  $\text{CH}_4$  as an electron donor [127]. Combining DAMO microorganisms and AnaAOB have many advantages: i)  $\text{CH}_4$  could be decreased to 15% in the effluent wastewater

[128]; ii) minimal amounts of  $\text{N}_2\text{O}$  were emitted [129]; iii) excellent nitrogen removal efficiency (99%) was achieved and cost savings were 49% [130]. The DAMO/A systems exhibit negligible  $\text{N}_2\text{O}$  emissions due to the absence of  $\text{N}_2\text{O}$ -generating enzymes in DAMO archaea and AnaAOB, while DAMO bacteria effectively remove excess  $\text{NO}_2^-$  [104]. Thus, DAMO/A has significantly attracted attention for applying it in WWTP as an alternative process to nitrification-denitrification and realizing optimal N removal efficiency and environmental sustainability. However, according to Samedo et al. [131], *Thauera linaloolentis* strain 47 Lol<sup>T</sup> contains *NosZ*, which converts  $\text{N}_2\text{O}$  to  $\text{N}_2$ . However, Mania et al. [132] found that when  $\text{NO}_3^-$  levels are high, the regulation of *NosZ* activity and  $\text{N}_2\text{O}$  reduction may not be possible. Consequently, elevated  $\text{NO}_3^-$  levels can significantly contribute to environmental impact and result in higher net  $\text{N}_2\text{O}$  emissions [131]. More research is still needed to understand DAMO/A microbial metabolism.

### 5.3. Source separation systems

In source separation systems, black water (BW), grey water (GW) and food waste (FW) are separated from municipal wastewater and solid waste as shown in Fig. 6. Separating the different kinds of wastewater at source can potentially increase the treatment efficiency, nutrient recovery and biogas yield. Kjerstadius et al. [133] compared the CF of conventional systems and a source separation system in Swedish WWTPs. The LCA showed that the source separation system could decrease the overall CF by 34 kg  $\text{CO}_2\text{e}/\text{PE}/\text{year}$ , primarily due to an increasing the biogas production.

Other studies, e.g. Remy [134], were in-line with the findings of Kjerstadius et al. [133] regarding the reduction of the net CF after source separation. In contrast, Thibodeau [135] reported a CF of 65 kg  $\text{CO}_2\text{e}/\text{PE}/\text{year}$  for a source separation scenario, which was higher by 12 kg  $\text{CO}_2\text{e}/\text{PE}/\text{year}$  in comparison with the conventional system. The wet

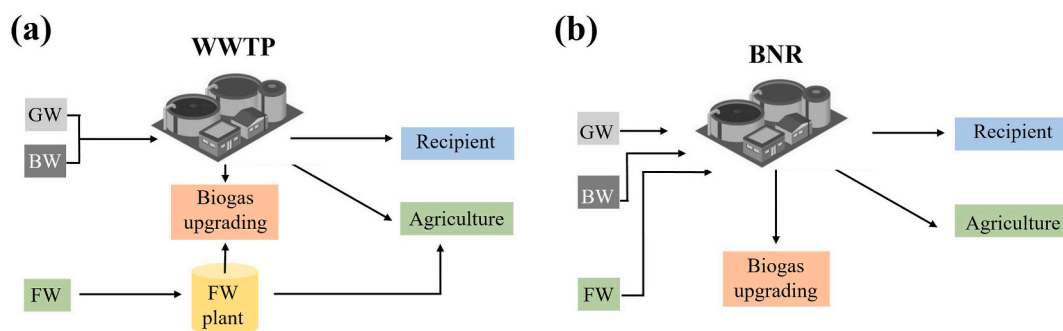


Fig. 6. (A) Conventional systems vs (B) source separation system of wastewater treatment (adopted from Kjerstadius et al. [133]).

fraction of BW was entirely returned to agriculture, leading to an increase in the CF resulting from transportation. However, the increased CF could be balanced out by selling bio-products as fertilizer.

Another benefit of source separation could be an increased recycle of nutrients to farmlands. The range of excess return of nutrients (N and P) was 1.7–4.3 kg N/PE/year and 0.06–0.54 kg P/PE/year, respectively [133–135]. The increased recovery of nutrients and utilizing them on farmlands could partially offset CF [4] and further decrease the indirect CF of WWTPs. Besson et al. [136] performed an LCA study and compared the environmental impacts of a central WWTP with source separation systems. Their results showed that for maximizing nutrient recovery and reducing GHG emissions, urine and BW separation are more favorable scenarios than a centralized WWTP. The separation allowed for mitigating  $N_2O$  emissions by 60% and avoiding energy-intensive N fertilizer production (Habor-Bosch process), which emits a large amount of GHGs (8.6 kg  $CO_{2e}/kg$  N) [136].

In another study, Tian et al. [137] utilized urine wastewater derived from a source separation system to pretreat WAS for AD. The authors observed a significant improvement in cumulative methane production, showing a 23% increase after pretreatment with urine wastewater at a volumetric proportion of 1:8. The presence of urea and hydrolyzed free ammonia in urine wastewater had a synergistic effect on sludge disintegration, leading to a reduction in sludge volume and increased availability of substrates for biological treatment processes. These findings highlight the potential of urine wastewater for pretreatment to enhance the efficiency of AD processes and improve biogas production from WAS. Badeti et al. [138] conducted a theoretical investigation of the impact of urine diversion on the biological treatment processes at a decentralized WWTP. Their simulations showed that 33% of aeration energy could be saved with 90% urine diversion. Furthermore, direct  $N_2O$  and  $CO_2$  emissions in the treatment processes could be reduced by 98% and 25%, respectively, while indirect GHG emissions could be reduced by 20%. Source separating sanitation was considered in a new city district of Hiedanranta, Finland, Lehtoranta [139] considered source separating sanitation and applied LCA to compare it with a conventional sanitation system. The study revealed that with a separating system, 3 to 10 times more N could be recovered compared to the conventional system. Based on their environmental performance, source separating systems with improved nutrient recovery showed reduced climate and eutrophication impacts.

#### 5.4. Systems with nutrient recovery

##### 5.4.1. P recovery from wastewater

The P load in urban wastewater could theoretically replace up to 50% of the mineral P fertilizer used in agriculture. Moreover, P mining from the phosphate rock is associated with environmental concerns, such as air pollution and water contamination [140]. Amann et al. [141] used LCA to evaluate 18 P recovery technologies in terms of the energy demand and GWP. The analysis showed that a wide-range of GHG emissions can be estimated through the implementation of different P

recovery strategies. The recovery from the liquid phase reduces the net GHG emissions. For the recovery from sludge, high GHG emissions were attributed to the processes which were already practiced in the full-scale. The recovery from sewage sludge ash showed the highest potential for an efficient P recycling. This method could annually decrease the cumulative energy demand and GWP by 0.096% and 0.1% per PE, respectively [141].

Another LCA study by Duan et al. [142] analyzed and compared the environmental impacts of different P removal and recovery methodologies. For chemical P removal and recovery pathways, the net specific CF was 44.5 kg  $CO_{2e}/kg$   $P_{removed}$  and  $-3.76$  kg  $CO_{2e}/kg$   $P_{recovered}$ , respectively. The negative value indicated the potential savings of CF due to P recovery. In another review, Zhao et al. [143] explored potential pathways for P recovery in WWTPs with a special focus on the A-2B-centered process, which involves an anaerobic fixed bed reactor to capture COD for improving energy efficiency. Simultaneously, this process allows for P recovery through further treatment with P crystallization. The A-2B-centered process exhibited the lowest specific energy consumption and GHG emissions when considering the amount of P recovered, indicating its promising approach in terms of both energy efficiency and environmental impact. The management of P resources in WWTPs encompasses multifaceted concerns related to environmental protection, energy efficiency, and GHG emissions. By exploring and implementing efficient recovery paths like the A-2B-centered process, WWTPs can take significant steps towards sustainable P management.

##### 5.4.2. N recovery from wastewater

Recovery of N as a nutrient would decrease both direct environmental impacts and energy consumption. Nitrification-denitrification could be energy intensive and without any additional benefits aside from meeting the effluent standards. N recovery allows for the simultaneous treatment of wastewater, while consuming less energy and recovering resources. Beckinghausen et al. [144] performed a comprehensive review of approximately 50 N recovery techniques. The authors focused on techniques that could produce fertilizers. The highlighted technique was permeable membrane, which consumed smaller amount of energy (1–1.2 kWh/kg N). Moreover, it showed the potential to recover  $NH_4^+$  from wastewater and the market value of the recovered product (ammonium sulphate). Other techniques, such as vacuum membrane distillation, can recover large amounts of energy in the form of heat (27 kWh/kg N), but the process is highly energy demanding (60 kWh/kg N). These techniques require further investigation to evaluate the costs of energy, as the amount of recovered heat highly effects the overall energy balance [145]. Innovative technologies, such as membranes, sorbents, electrodialysis, bioelectrochemical process, and even microalgae, are used to enhance nutrient recovery from wastewater, moving beyond conventional crystallization methods. Moreover, at the full-scale scale, some physicochemical methods, such as air stripping and struvite formation, are employed to recover P and N from side-streams [146].

## 6. Mitigation of GHG emissions via energy usage optimization

Wastewater treatment has a high potential to decarbonize and move towards net-zero carbon condition, since water reclamation from wastewater is considered to be the most energy-intensive water infrastructure [1].

### 6.1. CF due to energy consumption

Energy efficiency has received growing attention as the number of WWTPs is growing globally and the effluent quality criteria become more strict [147]. The water industry is estimated to account for 2.1% in the US and 0.8% in the EU of the total electricity consumption [148]. Around 23 MMT CO<sub>2e</sub> was generated during electricity consumption, which accounts for 37.7% of the entire CO<sub>2e</sub> release in the water industry [149]. Aeration, pumping, and heating are the main energy consumers in WWTPs [149]. In a typical WWTP, aeration can account for up to 60% of all electricity usage, followed by sludge treatment and return activated sludge (RAS) recirculation from secondary clarifiers [150].

Decarbonization of the energy sector can lead to a reduction in the indirect emissions in WWTPs. Coal-fired electricity production is being replaced by natural gas and more countries are substituting fossil fuels with renewable resources, such as wind, solar, hydropower, and nuclear power [149]. Hydropower and nuclear power can be seen as the cleanest energy source with a maximum carbon intensity of 40 g CO<sub>2e</sub>/kWh and 110 g CO<sub>2e</sub>/kWh respectively, while coal is the most carbon-intensive energy source (Table 4) with the maximum carbon intensity of 1689 g CO<sub>2e</sub>/kWh [151]. Currently, about 88% of the grid energy is satisfied by fossil fuels. Over the next 20 years, it is planned to increase the usage of renewable energy by 25%, which would decrease the CF related to the energy mix by up to 42%.

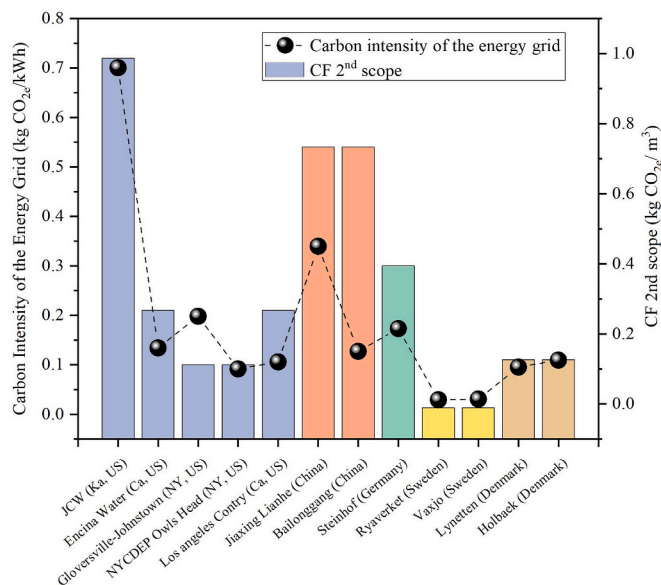
Delre et al. [132] evaluated the effect of electricity usage on the CF of seven different WWTPs located in Denmark and Sweden. In the Danish WWTPs, where the energy mix mostly consists of fossil fuels, energy consumption accounted for up to 29% of the CF. In the Swedish WWTPs, where nuclear and hydropower account for most of the energy mix, the CF-related energy consumption was as low as 3%. In the US, the fuel mix varies substantially by the state, resulting in energy carbon intensity ranging from 0.12 kg CO<sub>2e</sub>/kWh in Idaho to 1.15 kg CO<sub>2e</sub>/kWh in Washington, DC [82]. As shown in Fig. 7, the Midwest and Southwest regions contribute heavily to CF. This is due to the heavy reliance on fossil fuels in these regions. In the Midwest, 66.5% of the energy mix is satisfied by coal, while only 6% of the energy mix is from renewable energy sources. In contrast, in the Southwest regions, natural gas and coal make up 45% and 22% of the energy mix. On the other hand, approximately 50% of the energy mix in some regions is fueled by renewable energy, and thus the specific GHG emissions were found as low as 0.38 kg CO<sub>2e</sub>/m<sup>3</sup><sub>wastewater</sub> [148].

Although the energy mix has a noticeable effect on the indirect CF of WWTPs, other factors also play an important role. For example, the Owls Head WWTP (New York, US) has a substantially lower CF related to energy compared to the Gloversville-Johnstown WWTP (New York, US), even though the energy mix is similar in those plants. This is because Owls Head WWTP uses chlorine as a disinfectant, while Gloversville-

**Table 4**

EFs for different sources of electric energy [151].

Energy Source	min (g CO <sub>2e</sub> /kWh)	max (g CO <sub>2e</sub> /kWh)
Coal	675	1689
Oil	510	1170
Natural gas	290	930
Biogas	50	700
Hydropower	3	40
Nuclear power	4	110



**Fig. 7.** Comparison between carbon intensity of the energy in different countries and scope 2 emissions of selected WWTPs [82,152].

Johnstown WWTP uses ultraviolet (UV) for effluent disinfection, which requires electricity consumption [82].

### 6.2. Reducing energy consumption

Minimizing energy consumption can play a pivotal role in the CF reduction. There are two major approaches to decrease the power use in WWTPs: (i) reduction through operational changes and (ii) application of innovative, energy-efficient wastewater treatment processes [153]. Table 5 shows the energy mix and CF reduction under different scenarios in several case studies.

The DO, solids retention time (SRT), and mixed liquor recirculation (MLR) are crucial operational parameters in the bioreactor, which highly determine the operation of the aeration system and electricity consumption. Energy consumption could be decreased by 7–9% through the adjustment of SRT, DO, and MLR [154]. Mamais et al. [150] also obtained a 4500 MWh/year energy reduction by optimizing the SRT and DO concentration in a WWTP. Borzooei et al. [159] concluded that theoretically a reduction of up to 5000 MWh/year in energy consumption would be possible through operational changes in the largest Italian WWTP (2.1 million PE).

The advanced equipment also brings benefits to the CF. Li et al. [157] installed a TN online monitoring equipment at both influent and effluent, with a carbon source dosing system. As a consequence, the TN treatment efficiency increased by 1%, while the energy consumption and carbon emission decreased by 5.6% and 12.7%, respectively. Li et al. [157] updated the traditional blower to magnetic suspension centrifugal blowers, resulting in a 15–24% decrease in the aeration energy and a 4.6–7.7% decrease in the CF. Additionally, Ke and Lu [160] noted that the energy efficiency of the magnetic suspension blower could be improved by 20%.

ML is one of the novel techniques for reducing the energy demand of WWTPs. Cao and Yang [161] developed a model (OS-LEM) to control DO supply of aerobic bioreactors, which can reduce the energy consumption by 40% in comparison with the conventional on/off DO control. Similarly, Ramli et al. [158] used ML to build a prediction model to reduce the energy consumption of WWTPs by forecasting it one month ahead of time, which estimated a 2.2% reduction in energy consumption.

**Table 5**  
Reported energy reductions in WWTPs under different scenarios.

Reference	Scenario	GHG emission related to energy (kgCO <sub>2e</sub> /m <sup>3</sup> )	Reduction in Energy Demand		Total GHG emission decrease (%)	Remarks
			(MWh/year)	%		
Mamais et al. [150]	Reduction in SRT and DO	–	4500	–	–	Psytalia (Greece) WWTP in Greece was analyzed.
Daskiran et al. [154]	Reduction in SRT and MLR	0.1	3510	–	0%	Emission rates matched the active scenario in the WWTP
	Reduction in SRT and DO	0.06	3530	–	–15%	The emission rates were about 0.17 kgCO <sub>2</sub> /m <sup>3</sup> more than the active scenario in the WWTP
	Reduction in MLR and DO	0.05	4661	–	–9%	The emission rates were about 0.1 kgCO <sub>2</sub> /m <sup>3</sup> more than the active scenario in the WWTP
Borzoeei et al. [155]	Reduction in SRT	–	5000	–	–	In Castiglione Torinese WWTP (Italy) different scenarios where SRT was optimized were tested
Longo et al. [147]	Minimizing energy consumption of pumps, motors, and other electro-mechanic devices	–	–	10–20%	–	WWTP in Missouri Southeast, USA
Torregrossa et al. [156]	optimizing pumping, blowers, and AD	–	–	50–80%	–	Various WWTPs in Europe
Li et al. [157]	Upgrading to a magnetic suspension blower	–	–	15–24%	4.6–7.7%	14 WWTPs in North-Central China were examined
	Accurate aeration system modification	–	–	6.3%	10.37%	The implementations were studied on two WWTP in China
	Intelligent precise aeration system for sewage treatment	–	–	5.3%	10.93%	The implementations were studied on a WWTP in China
Ramli et al. [158]	Optimizing pumping, blowers, and AD	–	–	3%	–	WWTP in Peninsulay (Malaysia)

### 6.3. Increasing energy production inside WWTPs

#### 6.3.1. Co-digestion

Co-digestion is proposed as a straightforward approach for improving macronutrient balance, correcting moisture, and/or diluting inhibitory or toxic compounds [162], which possessed a positive effect on energy production and CF reduction. Most energy neutral plants located in developed countries adopt biogas recovery through co-digestion [163]. Koch et al. [164] reported more than double energy recovery when adding a ratio of 10% FW as a co-substrate in a WWTP. The biogas output increased by 17% by co-digesting 7% of fat with the mixed sewage sludge [30]. Sarpong and Gude [165] reported that the utilization of fat-oil-grease (FOG) for co-digestion with low-strength wastewater can increase energy production by 0.08 kWh/m<sup>3</sup>, while the total required energy of the studied plant was reported as 0.32 kWh/m<sup>3</sup>. Wickham et al. [166] demonstrated a biogas production increase of up to 191% using carbonated soft drinks as a co-substrate. An increase of 81% in biogas production was achieved by adding 3% of glycerol as a co-substrate [167]. Similarly, biogas production was

increased by up to 209% by using grease trap water as a co-substrate [168]. Many plants, employing co-digestion, have reached 100% energy neutrality and even become net energy positive, including Zurich Wedholzli WWTP-Switzerland (42 GWh/year), Point Loma WWTP-US (193 GWh/year), Grevesmuhlen WWTP-Germany (193 GWh/year), and Sheboygan Regional WWTP-US (32 GWh/year) [169]. Table 6 provides the potential increase in the biogas production rate for various co-substrates.

#### 6.3.2. Biogas utilization and upgrading

The produced biogas can be utilized internally for valorization in combustion engines to supply electricity and heat generation. Most WWTPs that produce biogas through AD in developing countries do not have thermal energy recovery or electricity production, thus the biogas is flared and wasted [163]. A decade ago, the USEPA estimated that if 544 US WWTPs with AD utilize CHP, the reduction in energy consumption would be equivalent to the emissions from 430,000 cars [173]. The Psytalia WWTP deployed CHP and revealed the lowest annual energy consumption among 10 studied WWTPs in Greece, at 15 kWh/PE

**Table 6**  
Biogas production rate increase after using different co-substrates.

Co-substrate	Condition	Energy production without co-substrate (kWh/d)	Energy Production with co-substrate (kWh/d)	Biogas Production without co-substrate (L/gVS/day)	Biogas Production with co-substrate (L/gVS/day)	Reference
Microalgae	–	774	1189	–	–	[170]
Cheese whey	–	774	1531	–	–	
The organic fraction of municipal solid waste (OFMSW)	Volumetric ratio 75:25;	7688	8418	–	–	[171]
	HRT 20 d					
Dry waste (DW)	Volumetric ratio 80:20;	677	10,639	–	–	
	Hydraulic Retention Time (HRT) 16 d					
OFMSW + DW	Volumetric ratio 68:23:9;	6037	8972	–	–	
	HRT 18 d					
Grease (G)	TS: G; 50:50	–	–	0.48	0.78	[172]
Septage (SP)	TS:SP; 90:10	–	–	0.48	0.46	
Whey (W)	TS: W; 70:30	–	–	0.48	0.75	

[150].

The produced biogas can be upgraded to biomethane, which has the same CH<sub>4</sub> quality as natural gas and could be utilized as a fuel for vehicles or pumped into the grid [174]. Biogas upgrading, involving CO<sub>2</sub> removal, employs gas sorption and separation techniques. Physical and chemical scrubbing are effective but energy-intensive, while pressure swing adsorption is less energy-intensive, but more complex. Membrane separation technologies are environmentally friendly with a relatively low energy consumption, whereas cryogenic separation demands high energy for cooling [175]. Following the technological advances in biogas upgrading, a transition away from CHP and towards upgrading biogas to biomethane has been trending [176]. Biogas is already being converted to biomethane in 15 EU countries and 10 of those countries are injecting biomethane into the natural gas grid. The countries with the highest number of biomethane plants include Germany (185 plants), UK (80 plants), and Sweden (61 plants), while the other countries are lagging behind [174].

To address GHG release from biogas upgrading, emerging technologies focus on CO<sub>2</sub> utilization through methanation, combining CO<sub>2</sub> with H<sub>2</sub> to produce CH<sub>4</sub> via chemical and biological processes. Implementing this technology requires investment and operational costs, especially for H<sub>2</sub> production through electrolysis. The biomethanation process efficiency is a limitation, but benefits include carbon neutrality and continuous fuel delivery for grid balance [175].

Nguyen et al. [169] compared the CHP and biomethane for biogas utilization from WWTPs. It was concluded that the CHP was effective with a negative CF, while the biogas upgrading to biomethane resulted in a higher CF due to a high energy input during the upgrading process. Ravina and Genon [177] also concluded that CHP of biogas with the thermal energy utilization produces the lowest CO<sub>2e</sub> (−0.277tCO<sub>2</sub>/t biogas). The biomethane upgrade (−0.13 tCO<sub>2</sub>/t biogas) to the Italy's national grid enabled to store biomethane at a lower cost for further utilization. Thus, upgrading biogas to biomethane could be a more sustainable scenario than on-site biogas consumption in CHP units, especially if the gas energy content is not fully cogenerated and the produced thermal energy is not capitalized. Regarding the CH<sub>4</sub> loss during upgrading, a sustainable system should reduce CH<sub>4</sub> slip below 4%, with a target value of 0.05% [177].

## 7. Mitigation of GHG emissions via chemicals and transportation optimization

To achieve full decarbonization and the net-zero carbon condition in WWTPs, both upstream and downstream, indirect emissions are ought to be reduced. Ambiguous GHG emissions from WWTPs are primarily related to chemical agent usage and transportation of by-products (grit, screenings, and sludge) [28].

### 7.1. Reducing chemical consumption

Chemicals are used at different stages of wastewater treatment and sludge management. To minimize their contribution to the CF, not only consumption cuts can be implemented, but also some sustainability criteria introduced to the purchase procedure may result in reducing the indirect GHG emissions. Factors linked to the usage of chemical agents in the context of upstream indirect emissions are leveled at 1964 kg CO<sub>2e</sub>/kg for inorganic agents and 1909 kg CO<sub>2e</sub>/kg for organic agents (Ecoinvent database 3.6).

The inorganic agents are commonly used to support enhanced biological P removal. Metal salts, such as aluminum (III) sulphate, iron (III) chloride, and lime in the form of Ca(OH)<sub>2</sub> or CaO<sub>2</sub>, are added to trigger precipitation. The use of lime compounds results in a pH increase, which requires the addition of acid chemicals to adjust the pH level of the WWTP's effluent. Moreover, both metal salts and calcium agents result in an increase in sludge volume, leading to higher GHG emissions associated with sludge disposal. It is preferable to use recycled inorganic

chemical agents instead of new resources in order to avoid emissions related to the extraction and manufacturing processes of raw materials. Organic agents and polymers are regularly applied for flocculation, coagulation, and dewatering processes. Recycling could effectively reduce the indirect GHG fluxes. Instead of synthetic, polymers of bio-based origin can be applied as a more environmentally friendly choice [178,179].

The application of chemical agents influences the entire wastewater treatment process and its stages [180]. The enhancement of primary sedimentation via chemically forced flocculation contributes to a higher chemical consumption in the overall GHG balance. Ultimately, it becomes an advantage due to the biogas and energy production bust. In contrast, the chemical agents used to support dewatering processes may become a target in lowering CO<sub>2e</sub> indirect emissions via overall chemical agent consumption cuts at WWTPs.

### 7.2. Reducing transport-related CF

Conveying activities and services, such as upstream transportation of chemical agents and downstream transport of wastewater treatment by-products (grit, screenings, and sludge), contribute to the indirect emissions of WWTPs. It is often challenging to implement a transport-related emission reduction plan based on switching to more environmentally-friendly means of transportation, such as rail transport, or shortening the transportation distance. Therefore, more advanced reduction scenarios are sought.

According to the Department for Environment, Food and Rural Affairs (DEFRA) [181] EF database, the amount of GHG emitted to the atmosphere depends on (1) engine standard, which translates into the type of fuel combusted, (2) size of the vehicle – rigid, and (3) the laden. Emission reduction can be achieved via a detailed transportation plan, including optimization of laden for each material transported, adaptation of the vehicle rigid type (elimination of oversizing), and optimization of fuel type engines.

Transport-related concerns resulted in the promotion of various biofuels: especially bioethanol from sugar and starch crops and biodiesel from oilseed crops [182]. Triggered by the Renewable Energy Directive I and II [183,184], the process of replacing fossil-origin gasoline and diesel with biofuels presents at least 65% GHG savings in comparison with the conventional fuels.

## 8. Recovering N<sub>2</sub>O from wastewater as an energy source

Recovering N<sub>2</sub>O as an energy source would be a win-win strategy. The process would be a step-forward towards net-zero operation by offering both the possibility of energy generation and mitigation of N<sub>2</sub>O emissions. There are limited studies available in the literature that investigate N<sub>2</sub>O recovery, which has also been being used as a powerful oxidant for energy generation. Wu et al. [185] suggested that applying Coupled Aerobic-Anoxic Nitrous Decomposition Operation (CANDO) might be one of the bioprocesses with a high potential to achieve energy neutral wastewater treatment paradigm. The process involves three steps: i) partial nitrification of NH<sub>4</sub><sup>+</sup> to NO<sub>2</sub><sup>-</sup>; ii) partial anoxic reduction of NO<sub>2</sub><sup>-</sup> to N<sub>2</sub>O; and 3) N<sub>2</sub>O conversion to N<sub>2</sub> with energy recovery by either catalytic decomposition or co-combustion of N<sub>2</sub>O with CH<sub>4</sub>. Yu et al. [186] documented a novel process that recovers N<sub>2</sub>O by inhibiting the activity of the nitrous-oxide reductase (N<sub>2</sub>OR) in denitrifying bacteria. N<sub>2</sub>O recovery was demonstrated in batch experiments in which NO was the only electron-acceptor. N<sub>2</sub>O recovery level in their study reached approximately 70% of the total gas production.

The most investigated N<sub>2</sub>O recovery process is CANDO, which has been tested with various configurations and real wastewater [187]. However, achieving stable long-term operation requires further experiments. Promising alternatives include autotrophic photoelectrotrophic denitrification, sulfur-driven denitrification, and hydrogenotrophic denitrification [188]. These autotrophic processes are currently

immature technologies and need intensive investigations. N<sub>2</sub>O recovery from high-strength wastewater is more favorable than low-strength (mainstream) wastewater due to a higher energy and economic potential. Recently, studies such as Deng et al. [189] and Huo et al. [190] have developed reliable models to simulate N<sub>2</sub>O recovery from Fe(II) EDTA-NO. The simulation results of Huo et al. [190] suggested that with a S/N mass ratio of around 1, high-purity N<sub>2</sub>O could be more rapidly and efficiently recovered from Fe(II)EDTA-NO (N<sub>2</sub>O recovery efficiency reached up to 85%). Deng et al. [189] highlighted that sufficiently increasing the headspace volume of the reactor was considered an ideal strategy to obtain ideal N<sub>2</sub>O production of 86.6%. Duan et al. [188] critically reviewed the progress of research on N<sub>2</sub>O recovery and counted (i) unstable nitrification, (ii) the low energy potential and (iii) environmental risks associated with intentional N<sub>2</sub>O emissions in mainstream treatment, as technical and economic challenges.

## 9. Challenges and limitations while implementing net-zero carbon practices

Both direct (fugitive, process-related) and indirect (due to energy and chemicals consumption, transportation) GHG emissions have some potential for reduction. However, implementing mitigation strategies for the net-zero carbon concept in WWTPs comes with recognized limitations and trade-offs. Effluent quality, operational cost and GHG emissions are potentially conflicting objectives [95,97,191]. For example, energy savings in the AS bioreactor, achieved by decreasing aeration intensity, may result in increased N<sub>2</sub>O emissions [192]. Co-digestion of external substrates reduces energy-related emissions and increases process-related GHG emissions [30]. Those studies proposed either separate evaluation criteria or integrated plant performance indicators. LCA complements decision-making to emphasize environmental impact [97]. In a full-scale WWTP modelling study [74], three sustainability criteria were considered: effluent TN concentration, overall energy balance and GHG footprint. Investigating the DO set-point in the aerobic zone and MLR rate showed improvements in GHG emissions and effluent quality. Technological upgrades, such as CEPT and deammonification in reject water treatment, enhanced energy neutrality and reduced GHG footprint, but did not improve effluent quality simultaneously.

## 10. Conclusions

Several steps may be considered when moving WWTPs towards net-zero carbon conditions. The direct emissions from treatment processes can contribute to more than 60% of the CF in WWTPs, while energy consumption with over 30% contribution to the CF has also a high potential for decarbonization of WWTPs. The remaining emissions are of lesser importance and can be offset by optimizing chemical usage and transportation. The first major step towards net-zero carbon condition is identification of emission hotspots and quantification of those emissions via either measurements, reliable models or EFs. The second step is GHG mitigation via process modification and energy usage optimization. The implementation of novel N removal processes such as PD/A and DAMO/A, could reduce GHG emissions and energy consumption while ensuring reliable N removal efficiencies. Other techniques such as source separation systems could potentially allow mitigation of N<sub>2</sub>O emissions by 60% while avoiding energy-intensive N fertilizer production. Nutrient recovery methods are another approach which offered negative value for the net CF. Permeable membrane N recovery offered 1 kWh/kg N of energy savings and P recovery led to -3.76 kg CO<sub>2e</sub>/kg P<sub>recovered</sub> CF savings. Upgrading biogas to biomethane could be a more sustainable scenario than on-site biogas consumption in CHP units, especially if the thermal energy is not capitalized. Recovering and utilizing N<sub>2</sub>O for energy production is a promising method which leads to both direct and indirect CF reductions.

## Declaration of competing interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper

## Data availability

No data was used for the research described in the article.

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