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# Greenhouse Gas and Air Pollutant Emissions from Composting

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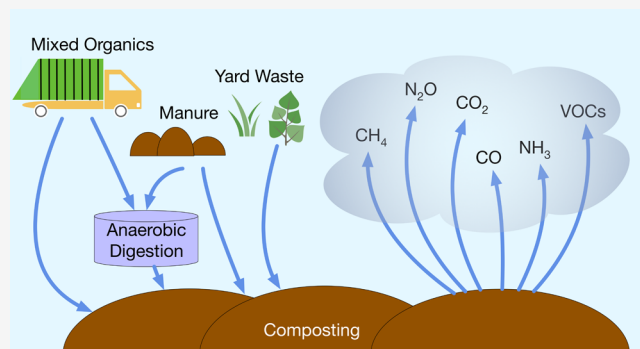
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**ABSTRACT:** Composting can divert organic waste from landfills, reduce landfill methane emissions, and recycle nutrients back to soils. However, the composting process is also a source of greenhouse gas and air pollutant emissions. Researchers, regulators, and policy decision-makers all rely on emissions estimates to develop local emissions inventories and weigh competing waste diversion options, yet reported emission factors are difficult to interpret and highly variable. This review explores the impacts of waste characteristics, pretreatment processes, and composting conditions on CO<sub>2</sub>, CH<sub>4</sub>, N<sub>2</sub>O, NH<sub>3</sub>, and VOC emissions by critically reviewing and analyzing 388 emission factors from 46 studies. The values reported to date suggest that CH<sub>4</sub> is the single largest contributor to 100-year global warming potential (GWP<sub>100</sub>) for yard waste composting, comprising approximately 80% of the total GWP<sub>100</sub>. For nitrogen-rich wastes including manure, mixed municipal organic waste, and wastewater treatment sludge, N<sub>2</sub>O is the largest contributor to GWP<sub>100</sub>, accounting for half to as much as 90% of the total GWP<sub>100</sub>. If waste is anaerobically digested prior to composting, N<sub>2</sub>O, NH<sub>3</sub>, and VOC emissions tend to decrease relative to composting the untreated waste. Effective pile management and aeration are key to minimizing CH<sub>4</sub> emissions. However, forced aeration can increase NH<sub>3</sub> emissions in some cases.



**KEYWORDS:** Greenhouse Gases, Composting, Air Quality, Ammonia, Methane, Anaerobic Digestion

## 1. IMPORTANCE OF COMPOSTING EMISSIONS

Composting is an essential part of any strategy to divert organic waste and reduce fugitive methane (CH<sub>4</sub>) emissions from landfills.<sup>1,2</sup> In the United States (U.S.), 6–9% of total municipal solid waste is currently composted, although as much as 34% could be composted if all food and yard waste were diverted from landfills.<sup>3,4</sup> Composting can treat organic waste directly or treat solids remaining after organic waste has undergone anaerobic digestion (AD), ultimately reducing the total mass of waste through aerobic biochemical decomposition and yielding soil amendments for agricultural or landscaping applications. Most reported values for mass loss during composting on a dry basis fall in the range of 10–60%.<sup>5–8</sup> The motivations for composting are (1) avoidance of fugitive CH<sub>4</sub> emissions associated with the anaerobic decomposition that occurs in solid waste landfills and manure storage lagoons, (2) the diversion of organic waste from landfills, and (3) generation of compost that is free of harmful pathogens and ready for use in agricultural applications or for erosion control.<sup>9,10</sup> Although avoiding CH<sub>4</sub> emissions from landfills is one of the motivations for composting organic waste, the composting process itself emits greenhouse gases (GHGs) and air pollutants, and these emissions are still not well understood.<sup>11</sup>

Gaseous emissions from the composting of organic waste have impacts on both climate change and air quality. The

GHG emissions are directly relevant to policy. For example, life-cycle GHG emissions from bioenergy production routes—some of which incorporate composting of residual solids—must be thoroughly documented in the U.S., as they are tied to the Renewable Fuel Standard Renewable Identification Numbers (RINs) and California’s Low Carbon Fuel Standard (LCFS) Carbon Intensity (CI) scores, both of which carry substantial monetary value.<sup>12</sup>

Non-GHG air pollutant emissions from composting facilities affect local and regional air quality and, as a result, human health in surrounding communities.<sup>13</sup> Emissions of ammonia (NH<sub>3</sub>) and volatile organic compounds (VOCs) from composting are of particular concern because they are precursors of secondary fine particulate matter (PM<sub>2.5</sub>), which is the primary driver of air pollution-related health impacts.<sup>14,15</sup> A detailed description of NH<sub>3</sub> emissions and PM<sub>2.5</sub> formation is provided in the [Supporting Information](#). VOCs are also precursors to tropospheric ozone formation which impacts human health and sensitive vegetation and

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ecosystems.<sup>14</sup> Lastly, VOCs and NH<sub>3</sub> have low odor detection thresholds and can cause a public nuisance for surrounding communities. Odorous pollutants can impact permitting for new facilities, particularly in nonattainment areas in the U.S. where ambient air pollutant concentrations exceed the National Ambient Air Quality Standards. Our review covers NH<sub>3</sub> and VOC emissions but does not include other odorous compounds such as noncarbon-containing volatile sulfur compounds (e.g., hydrogen sulfide). Because of the additional environmental and human health impacts from non-GHG emissions, it is essential to balance ambitious landfill diversion goals with local air quality and odor concerns associated with operating composting facilities.<sup>16</sup>

Despite the importance of GHG and air pollutant emissions from composting, available data can be difficult to interpret and use for policy implementation. The California Air Resources Board released a recommended methodology for estimating composting emissions in 2015, but the method was only applicable to mixtures with at least 85% green waste and a maximum of 15% food waste, biosolids, or manure.<sup>17</sup> The degree to which specific composting practices and incoming waste composition affect emissions per unit of composted material is not well understood. Furthermore, researchers incorporating composting emissions into life-cycle assessments (LCAs) are often not experts in different measurement techniques and the degree to which measurement methods affect the accuracy of empirical data. This knowledge gap makes prioritization of emissions mitigation strategies and scenario planning for zero-waste policies challenging.

The purpose of this review is to evaluate data available in the scientific literature on air emissions from composting operations, discuss the merits and trade-offs of measurement strategies employed in past studies, and provide guidance for researchers and decision-makers who seek to integrate composting emission factors into policy and environmental impact studies.

Regarding GHGs, we mainly focus on CH<sub>4</sub> and nitrous oxide (N<sub>2</sub>O) emissions because they are the primary drivers of net climate forcing impacts from composting.<sup>11,18</sup> Although contemporary carbon emitted as CO<sub>2</sub> during composting is not thought to have a net climate impact, we also include limited data on CO<sub>2</sub> emissions results, reported separately from CH<sub>4</sub> and N<sub>2</sub>O, in the Supporting Information (Figure S1). This review also includes data on NH<sub>3</sub> and VOCs because of their importance for air quality and air pollution-related human health impacts.<sup>15</sup> Empirical emission values collected from the literature were differentiated based on the type of material being composted, measurement methods used in the study, and the management strategies employed during the composting process, with the goal of developing more representative and material-specific recommendations for composting emission factor ranges.

Prior reviews have explored some dimensions of this topic but fall short of providing recommended ranges for emission factors that can be used in future LCAs and policy-making. For example, Amlinger et al. (2008) primarily focused on their own measured results for CH<sub>4</sub>, N<sub>2</sub>O, and NH<sub>3</sub> but included a review of prior results to inform the development of a helpful, mostly qualitative table summarizing the effects of different compost management strategies on emissions and the mechanisms behind those effects.<sup>19</sup> Brown et al. (2008) reviewed a broader set of literature values on GHG emissions associated with different alternatives for disposing of/treating

organic waste, including landfilling and anaerobic digestion, in comparison to composting.<sup>20</sup> Lou and Nair (2009) compared GHG emissions from composting and landfilling organic waste, concluding that landfilling results in higher GHG emissions as compared to composting, a conclusion that reflects broad consensus in the research community.<sup>21</sup> Pardo et al. (2015) conducted a meta analysis of 50 studies to establish the relative impacts of different management strategies, such as forced aeration versus turning and the addition of bulking agents.<sup>22</sup> Pardo et al. (2015) considered the same raw feedstocks as those included in this review and focused on the relative impact of different operational practices and conditions but did not establish emission factors per tonne of waste composted. Bong et al. (2017) and Sayara and Sánchez (2021) provide more qualitative reviews of composting GHG emissions and discuss GHG mitigation strategies.<sup>23,24</sup> Sayara and Sánchez (2021) summarize research regarding the impact of composting practices and feedstock characteristics on emissions, while Bong et al. (2017) focus more specifically on the variability of scope definition and inventory analysis in published LCAs of composting. Neither review provides recommended emission factors.

Although this review focuses on gaseous emissions during the composting process itself, excluding truck transport and combustion of fuels to operate equipment, composting emission factors are more meaningful in a broader context, where each end-to-end process for managing organic waste can be compared. There are two main competing routes of relevance: (1) composting followed by land application of finished compost and (2) landfilling untreated organic waste. The use phase for finished compost is essential to include in life-cycle emissions inventories; applying compost can reduce the need for synthetic fertilizers and, in some cases, increase the net primary productivity on degraded lands.<sup>25</sup> The comparison between composting emissions and landfilling organic waste is another important topic, and this has been explored more thoroughly in prior reviews, although gaps in empirical data remain.<sup>21,26</sup> In the [Supporting Information](#), we provide an overview of the state of knowledge related to how compost application and landfilling organic waste affect net GHG emissions.

## 2. THE ROLE OF COMPOSTING IN ORGANIC WASTE MANAGEMENT

Organic wastes that can be composted include the entire organic fraction of municipal solid waste (OFMSW, which includes a variety of organic waste types), food waste, yard waste, sewage sludge, manure, and digestates (residual solids remaining after AD). The most commonly composted material is source-separated yard waste. According to the U.S. Environmental Protection Agency (EPA), approximately 0.4% of food waste and 63% of yard waste are currently composted in the U.S. While technically compostable, paper waste is more commonly recycled unless it is soiled or otherwise unsuitable,<sup>27</sup> so we have excluded it from this review. Solid digestate can be directly applied to agricultural land as a fertilizer amendment, but there are typically seasonal limitations on this practice due to nutrient runoff concerns in some states, so AD facilities may send digestate to composting facilities during part or all of the year.<sup>15,28–31</sup>

The wastes processed at composting facilities vary in moisture content, the carbon-to-nitrogen ratio (C:N), pH, volatile solids (VS) content, and other characteristics that lead

to varying rates of aerobic decomposition and emissions to the atmosphere. VS refers to the part of compostable materials that is combusted at 550 °C in the presence of air after 2 h and can be a proxy for the fraction of biodegradable material.<sup>32</sup> The composting process itself involves a diverse microbial community, in which the relative abundance and activity level of different microbes shift over time. Because levels of aeration and the composition of organic matter will vary, there is also heterogeneity across a given pile or windrow. The multistage composting process begins with the mesophilic phase, in which mesophilic microbes break down easily degradable compounds until the generated waste heat increases the temperature to 40 °C, which inhibits their growth.<sup>33</sup> It is during the mesophilic phase that nitrifying and denitrifying bacteria produce N<sub>2</sub>O.<sup>33</sup> Above 40 °C, the thermophilic microbes begin to dominate, and the increased activity of methanogens results in greater CH<sub>4</sub> emissions.<sup>33</sup> At this point, reaching temperatures above 55 °C is desirable because this kills most human and plant pathogens; however, aeration is necessary to prevent the pile from exceeding 65 °C, the threshold where most microbes are killed and the rate of decomposition decreases.<sup>34</sup> After the thermophilic phase, the compost cools and undergoes a curing and maturation stage, during which slow decomposition continues as mesophilic microbes become dominant again.

Composting operations are designed to facilitate this natural process, and practices at different facilities are distinguished by the manner in which material is stored and aerated, either in windrows or vessels and with manual, passive, or forced aeration. With in-vessel composting, material is contained in a series of containers or concrete bunkers, in which the temperature and air flow are controlled. This approach requires less land area than windrow composting and can be more efficient with proper management but is a more expensive method.<sup>35</sup> In industrial-scale windrow composting operations, material is placed in rows of long and narrow piles called windrows. These windrows can either be left uncovered or can be enclosed by plastic sheeting or within bags. The dimensions of these piles are typically 2–6 m wide and 1–3 m in height, which is large enough to maintain thermophilic composting conditions while also ensuring adequate aeration.<sup>36,37</sup> There are several methods of aeration used to ensure the aerobic conditions required for composting. One method is to periodically turn uncovered compost piles manually or mechanically. Alternatively, static piles, either uncovered or enclosed by plastic sheeting, are aerated by natural, passive, or forced means. Natural aeration strictly relies on diffusion for air flow through the pile, but this approach can be inhibited by high moisture content material that reduces air space and increases the likelihood of conditions in the pile becoming anaerobic.<sup>38</sup> Passively aerated piles include perforated pipes to promote air circulation that is driven by thermal gradients. Forced aeration similarly uses perforated pipes but includes a positively or negatively pressurized pump to either push or pull air through the composting pile on prescribed cycles to control temperature and optimize the composting process. In negative aeration, the air drawn from piles may be treated with a biofilter to control odor and VOCs.<sup>39,40</sup> Naturally and passively aerated piles compost at a slower rate, whereas the controlled forced aeration or turning of piles results in shorter composting cycles.<sup>41</sup>

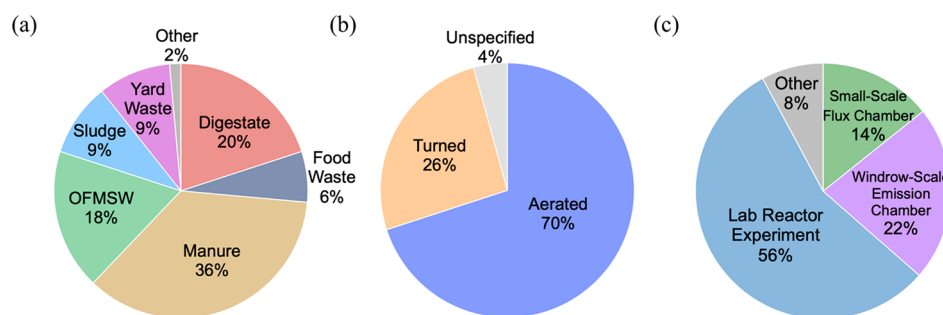
### 3. UNDERSTANDING AND MEASURING EMISSIONS FROM COMPOSTING

**3.1. Overview of Key Emission Sources.** By mass, CO<sub>2</sub> is the dominant compound emitted to the atmosphere during composting operations.<sup>33</sup> During each stage of composting, some carbon present in the organic material is oxidized to CO<sub>2</sub>. Because this CO<sub>2</sub> production is a natural part of organic decomposition and the carbon present in most compost feedstocks is biogenic (part of the contemporary carbon cycle, in contrast to fossil carbon), these emissions are considered to be climate-neutral.<sup>33,42,43</sup> Emissions of other air quality and climate-relevant pollutants vary, depending on factors like oxygen availability, temperature, and moisture content. Under anaerobic conditions, decomposition occurs more slowly; and methanogens create greater quantities of CH<sub>4</sub>, while emitted CO<sub>2</sub> decreases. Localized areas of anaerobic decomposition in composting operations are inevitable, but turning and aeration can minimize CH<sub>4</sub> emissions. Methanotrophs play an important role in consuming CH<sub>4</sub> that may be produced in localized anaerobic regions of the pile or windrow; one study suggested that 46–98% of CH<sub>4</sub> produced during composting operations is consumed by methanotrophs before it can escape to the atmosphere.<sup>44</sup> Carbon monoxide (CO) formation is well documented, but the mechanisms are still not fully understood by the scientific community.<sup>45</sup> CO in composting environments can be formed through thermochemical processes, stimulated by heat and ultraviolet radiation, and CO can also be produced and consumed by microbes.<sup>45–47</sup>

Nitrogen cycling in composting operations involves numerous direct and indirect processes, but an understanding of the basic mechanisms is important, given the relevance of N<sub>2</sub>O and NH<sub>3</sub> emissions to the climate and human health. Biological removal of nitrogen involves nitrification and denitrification and ultimately results in N<sub>2</sub>O emissions.<sup>48</sup> NH<sub>3</sub> is produced as microbes consume peptides and amino acids present in protein-rich waste. Nitrification is a two-step process in which microbes oxidize NH<sub>3</sub> to nitrite (NO<sub>2</sub><sup>-</sup>) and subsequently oxidize NO<sub>2</sub><sup>-</sup> to nitrate (NO<sub>3</sub><sup>-</sup>). A fraction of the NO<sub>2</sub><sup>-</sup> formed will be converted to nitric oxide (NO) and eventually N<sub>2</sub>O by ammonia oxidizing bacteria, rather than forming NO<sub>3</sub><sup>-</sup>. During denitrification, microbes anaerobically convert NO<sub>3</sub><sup>-</sup> back to NO<sub>2</sub><sup>-</sup>, then to NO, and ultimately to N<sub>2</sub>O, most (but not all) of which is ultimately converted to nitrogen gas (N<sub>2</sub>). NH<sub>3</sub> can also be directly emitted to the atmosphere, particularly from well-aerated piles where it escapes before microbes are able to oxidize it. NH<sub>3</sub> emissions from compost increase with increasing aeration, lower C:N ratios, higher temperatures, and higher pH.<sup>49</sup> The conditions for reducing NH<sub>3</sub> volatilization, unfortunately, can be counter to the optimal microbial conditions for fast and efficient composting.<sup>50</sup>

**3.2. Emissions Measurement Methods.** Many methods are used to measure emission rates from composting, and each has advantages and disadvantages to consider when interpreting and using the empirical data. Emissions can be characterized in controlled laboratory experiments or with in situ field measurements. Sampling can be conducted continuously in the field with pollutant analyzers or intermittently by collecting discrete samples of emitted gas into canisters or bags that are later analyzed in the laboratory. Pollutant concentrations can be measured at a single point or integrated across the composting pile. The trade-off in





**Figure 1.** Breakdown of the 140 composting scenarios and study methods associated with the reported emission factors collected for analysis. Emission factors are categorized by (a) composted material, (b) aeration method, and (c) measurement method.

temporal and spatial resolution between these sampling approaches depends on the sampling conditions and objectives of the study.

Laboratory experiments have been used to approximate the composting process under controlled conditions in reactors that are typically ~10–200 L in volume.<sup>51–64</sup> Lab experiments allow for a better understanding and characterization of how specific environmental conditions, such as temperature, pH, moisture content, and material, affect pollutant emissions than can often be attained with field measurements. However, the smaller lab-scale and experimental conditions may not be representative of the real-world. These emission factors should be used with caution or ideally validated against field measurements for similar materials and conditions.

Measurements can be made in the field as relatively controlled experiments of pilot- or full-scale test windrows that are maintained separately from normal operations.<sup>19,42,65–70</sup> In situ sampling of full-scale commercial windrows operating under normal composting conditions is also common.<sup>11,71–75</sup> Ideally, field measurements of emissions would be fully integrated over the windrow or pile surface, over the full duration of the composting cycle, and without disrupting normal composting conditions. This ideal measurement approach is not practical under many sampling scenarios, however, given researcher resources and environmental/operational conditions. As such, many sampling methods have been used for field measurements, including but not limited to flux chambers, gas probes, wind tunnels, open emission chambers, tracer releases, inverse dispersion analysis, micrometeorological mass balance, and high-density spot sampling. Each approach has its limitations, as described below. These emission measurements can also be accompanied by intermittent measurements and/or continuous monitoring of conditions in the windrow, which is important for developing a deeper understanding of the mechanisms driving emissions over time and space. Detailed descriptions of each measurement method and their impacts on reported emission factors are included in the [Supporting Information](#).

**3.3. Characterization of Composting Emissions Studies in the Current Literature.** We conducted a survey of peer-reviewed studies that report CO<sub>2</sub>, CH<sub>4</sub>, N<sub>2</sub>O, NH<sub>3</sub>, and/or VOC emissions from composting, keeping track of feedstock type and composting conditions. Where possible, we converted reported emission factors to units of kilograms of pollutant emitted per kilogram of wet (sometimes referred to as fresh or green) feedstock material composted. Composting emission factors are most commonly reported in terms of wet material because this is practical for commercial operations and general material flow tracking. However, these values should be

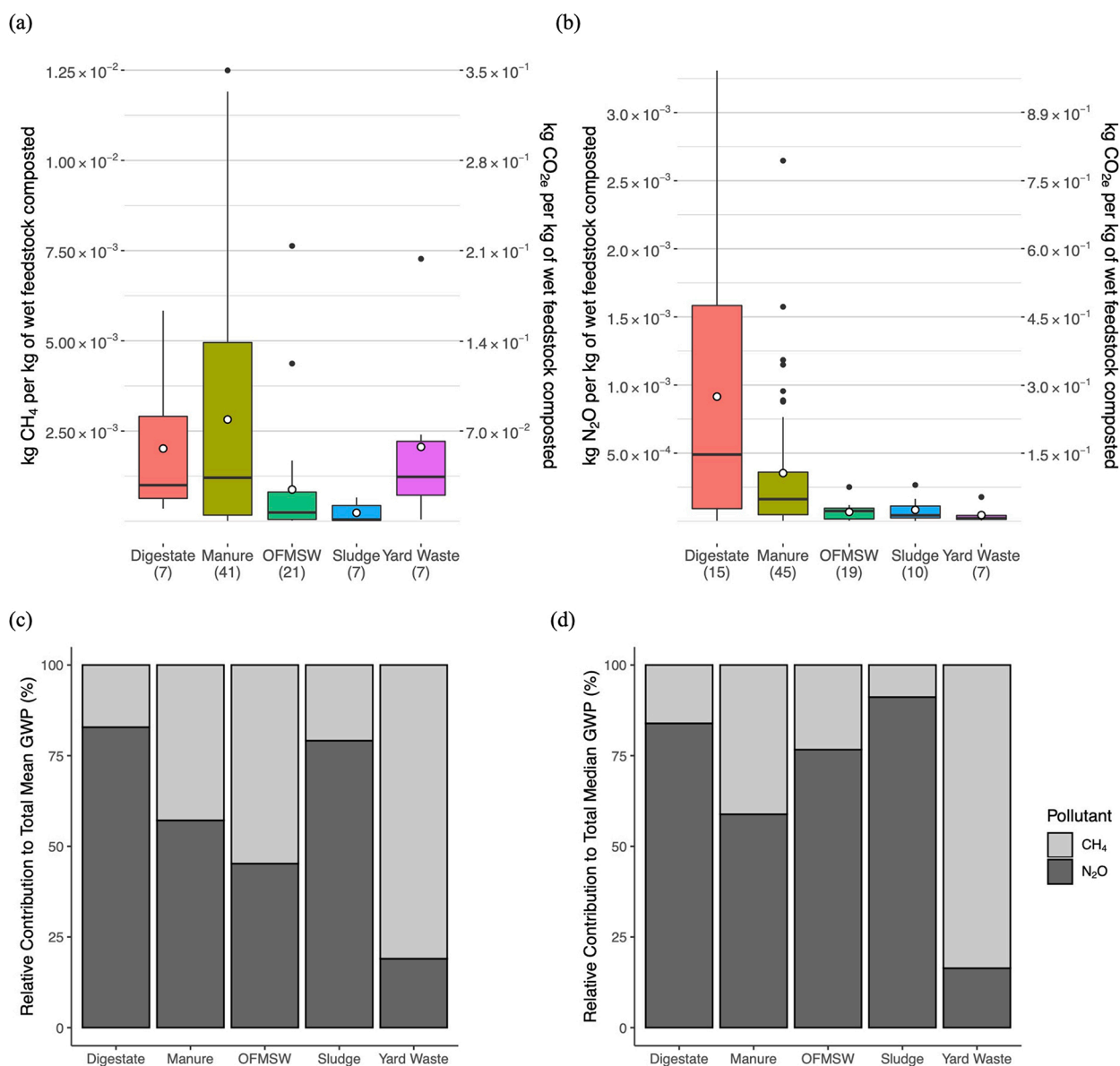
converted to a per-dry-mass basis for use in carbon or nitrogen balance modeling since water makes up a significant portion of composting feedstocks (36–85%, [Table S1](#)). We do not provide emission factors on a dry basis because several studies did not provide sufficient data on moisture content to calculate these conversions.

Studies that did not give enough information to calculate reasonable emission factors and secondary sources that did not provide original data were excluded from our review. In one case, we excluded 6 measured emission factors from further analysis because the authors acknowledged that two of their small-scale measurement methods, a static flux chamber method and a funnel method, significantly underestimated GHG emissions.<sup>76</sup> In total, 388 emission factors from 46 studies reporting emission measurements were considered in the survey, corresponding to 140 composting scenarios ([Table S1](#)).

A majority of currently available research on composting emissions is focused on manure composting. Therefore, manure composting comprises most of the emissions observations across all pollutant types ([Figure 1](#) and [Table S1](#)). The literature survey does include an extensive accounting of available literature on GHG composting emissions from food waste, OFMSW, yard waste, and anaerobically digested materials. Emissions from the composting of solid digestate are particularly understudied, and given the importance of these emissions for regulatory decision-making in waste-to-energy pathways, this topic requires further research. Of the collected data, most emission factors are associated with forced aeration ([Figure 1](#)). This is not necessarily the most common industry practice but is more easily replicated in lab-based studies ([Figure 1](#)). Most commercial composting operations involve outdoor windrows that can be turned or forcibly aerated, are not equipped with effective emission control systems, and allow all fugitive emissions to be released to the atmosphere. Alternatively, in-vessel or fully enclosed composting facilities can more easily be equipped with scrubbers and biofilters to reduce atmospheric emissions.<sup>77</sup>

## 4. COMPOSTING EMISSION FACTORS BY SOURCE MATERIAL AND MANAGEMENT PRACTICES

**4.1. Greenhouse Gas Emissions.** Establishing definitive, broadly applicable GHG emission factors for composting organic wastes is difficult because emissions vary due to a number of factors beyond feedstock (waste) type. These include the following: local climatic conditions at the composting site; composting method and duration; aeration method and frequency; use of a bulking agent intended to provide structure to piles/windrows and facilitate aeration; and



**Figure 2.** Distributions of (a) CH<sub>4</sub> and (b) N<sub>2</sub>O emission factors for composting reported in the literature and relative contribution to total GWP<sub>100</sub> based on (c) mean values and (d) median values. The sample size (*n*) of data points contributing to each boxplot is indicated in the *x*-axis labels for parts (a) and (b); the first value refers to the sample size of CH<sub>4</sub> emission factors, and the second value refers to that of N<sub>2</sub>O emission factors. Parts (a) and (b) have two *y*-axes: the left axis indicates the per-tonne mass of the specified pollutant emitted (exact values), and the right axis shows the CO<sub>2</sub>-equivalent emission factor (rounded values), so that CH<sub>4</sub> and N<sub>2</sub>O emissions can be compared with respect to GWP<sub>100</sub>. The mean values for the boxplot data are indicated by the open point symbols, while outliers are shown as closed circles.

the feedstocks' VS content, C:N ratio, moisture content, and pH. In this section, we differentiate previously published emission factors based on source material and management practices to elucidate the impact of these variables on GHG emissions. Our discussion of GHG emissions from composting is focused on CH<sub>4</sub> and N<sub>2</sub>O, as these gases are most likely to drive net changes in radiative forcing from composting operations. Biogenic CO<sub>2</sub>, by contrast, is not included in our GHG footprint calculations because it is part of the contemporary carbon cycle and will be resequenced during plant regrowth.<sup>20,33,42,43,78</sup> However, depending on how a particular researcher or practitioner chooses to account for carbon flows, it may be important to account for CO<sub>2</sub>. Further information and emission factor distributions for CO<sub>2</sub> are discussed in the [Supporting Information](#).

**4.1.1. Variation by Feedstock Type.** GHG emission factors by feedstock type are presented in [Figure 2](#) and [Table 1](#). [Figure 2](#) shows the distributions of CH<sub>4</sub> and N<sub>2</sub>O emission factors by feedstock type (manure, OFMSW, sludge, and yard waste) and for digestate. The distribution for digestate includes data across all original feedstocks to allow for a general comparison to raw material composting; the effect of AD as a pretreatment to composting is discussed in more detail in [Section 4.1.3](#). We grouped together studies examining OFMSW, household waste, kitchen waste, and food waste because of ambiguous distinctions between these feedstocks. If yard waste is collected separately and paper/paperboard is recycled, the remaining OFMSW will be primarily composed of food, food-soiled paper products, and other paper products that cannot be recycled.<sup>15</sup> However, in that case, a composter processing this

Table 1. Summary of GHG Emission Factor Data for Composting Raw Materials<sup>a</sup>

feedstock	pollutant	emission factor				sample size
		kg pollutant/kg of wet feedstock		kg CO <sub>2e</sub> /kg of wet feedstock		
		mean	median	mean	median	
manure	CH <sub>4</sub>	2.82 × 10 <sup>-3</sup>	1.21 × 10 <sup>-3</sup>	7.90 × 10 <sup>-2</sup>	3.39 × 10 <sup>-2</sup>	41
	N <sub>2</sub> O	3.54 × 10 <sup>-4</sup>	1.62 × 10 <sup>-4</sup>	1.05 × 10 <sup>-1</sup>	4.83 × 10 <sup>-2</sup>	45
	CO <sub>2</sub>	1.40 × 10 <sup>-1</sup>	1.47 × 10 <sup>-1</sup>	1.40 × 10 <sup>-1</sup>	1.47 × 10 <sup>-1</sup>	30
OFMSW	CH <sub>4</sub>	8.79 × 10 <sup>-4</sup>	2.43 × 10 <sup>-4</sup>	2.46 × 10 <sup>-2</sup>	6.80 × 10 <sup>-3</sup>	21
	N <sub>2</sub> O	6.80 × 10 <sup>-5</sup>	7.50 × 10 <sup>-5</sup>	2.03 × 10 <sup>-2</sup>	2.24 × 10 <sup>-2</sup>	19
	CO <sub>2</sub>	5.63 × 10 <sup>-2</sup>	4.30 × 10 <sup>-2</sup>	5.63 × 10 <sup>-2</sup>	4.30 × 10 <sup>-2</sup>	3
sludge	CH <sub>4</sub>	2.34 × 10 <sup>-4</sup>	4.50 × 10 <sup>-5</sup>	6.55 × 10 <sup>-3</sup>	1.26 × 10 <sup>-3</sup>	7
	N <sub>2</sub> O	8.36 × 10 <sup>-5</sup>	4.36 × 10 <sup>-5</sup>	2.49 × 10 <sup>-2</sup>	1.3 × 10 <sup>-2</sup>	7
	CO <sub>2</sub>	1.75 × 10 <sup>-2</sup>	1.75 × 10 <sup>-2</sup>	1.75 × 10 <sup>-2</sup>	1.75 × 10 <sup>-2</sup>	2
yard waste	CH <sub>4</sub>	2.06 × 10 <sup>-3</sup>	1.23 × 10 <sup>-3</sup>	5.77 × 10 <sup>-2</sup>	3.44 × 10 <sup>-2</sup>	7
	N <sub>2</sub> O	4.54 × 10 <sup>-5</sup>	2.27 × 10 <sup>-5</sup>	1.35 × 10 <sup>-2</sup>	6.76 × 10 <sup>-3</sup>	7
	CO <sub>2</sub>	1.71 × 10 <sup>-1</sup>	1.56 × 10 <sup>-1</sup>	1.71 × 10 <sup>-1</sup>	1.56 × 10 <sup>-1</sup>	4

<sup>a</sup>Digestate is excluded in this table because of variation in the original raw feedstock materials.

high-moisture food waste-dominated material will likely need to add a bulking agent, such as wood chips, sawdust, dry leaves, shredded paper/cardboard, or other materials that are very similar to yard waste and/or paper and paperboard. Therefore, the final material that is composted in all of these studies is likely to be similar regardless of whether yard waste and/or paper/paperboard in the original waste stream are diverted for other uses.

Table 1 provides mean and median emission factors by feedstock type, which can be useful for researchers and LCA practitioners who must approximate composting emissions as part of their analyses of waste management or waste-to-energy systems. It is important to note that assembling results from all prior field- or lab-based research may not provide a representative sample of real-world composting operations. For example, the majority of emissions data for composting manure came from studies examining either beef cattle, dairy cattle, or swine manure. In almost all studies considered here, composting operations for yard waste involved open, turned windrows. Most surveyed studies of sludge composting emissions were lab-based, involving closed reactors and forced aeration, and only examined wastewater treatment sludge.

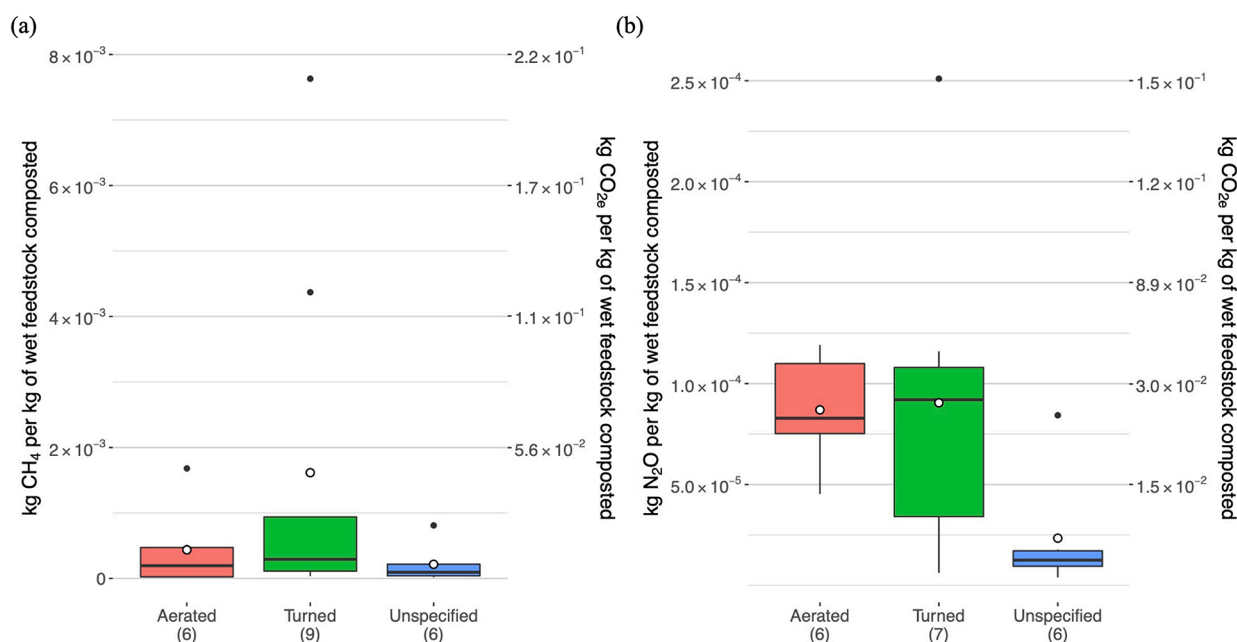
Based on 100-year global warming potential (GWP<sub>100</sub>) values for N<sub>2</sub>O and CH<sub>4</sub> (GWP<sub>100</sub> is equal to 298 and 28, respectively.), the median emission values for sludge, digestate, and OFMSW suggest that N<sub>2</sub>O is the largest contributor to total CO<sub>2</sub>-equivalent (CO<sub>2e</sub>) emissions, while CH<sub>4</sub> emissions are higher on a CO<sub>2e</sub> basis for yard waste (Figure 2). Manure composting resulted in the highest total GWP<sub>100</sub>, with a roughly even split between CH<sub>4</sub> and N<sub>2</sub>O on a CO<sub>2e</sub> basis (Figure 2, Table 1). Pardo et al. (2015) similarly found that composting manure resulted in the highest CH<sub>4</sub> and N<sub>2</sub>O emissions when comparing across different feedstocks.<sup>22</sup>

**4.1.2. Impact of Feedstock Characteristics.** In addition to the type of feedstock (e.g., manure, food waste), measurable characteristics including moisture content, VS content, and pH play a role in determining emissions. We attempted linear regressions using the ordinary least-squares method and more robust regressions using M-estimation to assess the relationship between each feedstock characteristic listed in Table S1 (% bulking agent, VS content, C:N ratio, moisture content, and pH) and each emission factor. The observed relationships were not statistically significant, even when controlling for

feedstock type or measurement methods. However, a few general trends emerged, consistently offering residual standard error less than 0.01 with varying degrees of freedom. The data collected in the literature survey suggests a positive correlation between moisture content and CH<sub>4</sub> emissions, and this holds true when controlling for feedstock type, which is supported by results from Pardo et al. (2015).<sup>22</sup> There is limited data that may suggest a negative correlation between moisture content and N<sub>2</sub>O emissions ( $n = 84$ ) and a positive correlation between VS content and N<sub>2</sub>O emissions ( $n = 22$ ), but further study is required to support any definitive conclusions. The collected data does not support a correlation between the C:N ratio and GHG emissions, and the impact of pH on overall emissions is likely negligible. Contrary to the results of our literature survey and analysis, Jiang et al. (2011) found in a lab-based study of swine manure composting that moisture content did not significantly impact CH<sub>4</sub> emissions, the C:N ratio was negatively correlated with CH<sub>4</sub> emissions, and neither moisture content nor the C:N ratio had an impact on N<sub>2</sub>O emissions.<sup>53</sup>

**4.1.3. Impact of Anaerobic Digestion Prior to Composting.** The literature on GHG emissions from the composting of digestate is limited, making it difficult to draw conclusions about the impact of AD as a strategy for pretreating organic waste prior to composting. However, even with the limited data available, there are some basic relationships that can be used to approximate differences in composting emissions between post-AD material and untreated material.

Li et al. (2018) provide one of the only studies that directly compares emissions from the composting of post-AD digestate to the same undigested material as a control.<sup>63</sup> In this lab-based experimental study, Li et al. composted raw, untreated feedstock—a mixture of manure and agricultural residues—as well as feedstock that first underwent AD for varied digestion times. Corn stover was added to ensure a similar bulk density across all samples during composting. CH<sub>4</sub> emissions during composting increased relative to the nondigested control treatment when the feedstock material underwent AD for only 15 days but decreased when the digestion time was 30 or 45 days. Without additional data on the microbiomes and volatile solids content in these composting experiments, it is only possible to speculate as to why shorter AD residence times caused elevated CH<sub>4</sub> emissions during subsequent composting.



**Figure 3.** Distributions of (a)  $\text{CH}_4$  and (b)  $\text{N}_2\text{O}$  emission factors for OFMSW composting based on the aeration method. The sample size ( $n$ ) of data points contributing to each boxplot is indicated in the  $x$ -axis labels. Each figure has two  $y$ -axes: the left axis indicates the per-tonne mass of the specified pollutant emitted (exact values), and the right axis shows the  $\text{CO}_2\text{-e}$  equivalent emission factor (rounded values). The mean values for the boxplot data are indicated by the open point symbols, while outliers are shown as closed circles.

It is possible that insufficient residence times during AD may allow digestate to be “seeded” with methanogens.<sup>63</sup> It is also possible that insufficient AD residence times result in higher concentrations of intermediate products from the hydrolysis, acidogenesis, and acetogenesis stages of AD in the final digestate. Additional studies and data would be required to support the development of feedstock-specific composting emission factors for post-AD materials.

Li et al. (2018) offer more conclusive results regarding the impact of AD on  $\text{N}_2\text{O}$  emissions from composting, which can be the primary contributor to total  $\text{GWP}_{100}$  from composting (Figure 2).<sup>63</sup> Piles pretreated with AD had 57–81% lower  $\text{N}_2\text{O}$  emissions relative to the nondigested control. Longer digestion times resulted in further reductions in  $\text{N}_2\text{O}$  emissions. Li et al. reported an average VS reduction during AD of 61% (individual VS reduction data for each batch treated with AD was not reported). For perspective, the mean  $\text{N}_2\text{O}$  emission factor reported by Li et al. for post-AD manure is 69% lower than the mean  $\text{N}_2\text{O}$  emission factor for composting untreated manure in Table 1.

In addition to Li et al. (2018) study, seven other studies measured GHG emissions from composting digested materials but did not include controls (identical untreated materials).<sup>11,54,61,62,64,71,77,79</sup> For instance, Colón et al. (2012) included a comparison of in-vessel composting with and without AD pretreatment, finding that  $\text{N}_2\text{O}$  emissions were 53% lower and  $\text{CH}_4$  emissions were  $\sim 7$  times higher for OFMSW treated with AD relative to raw OFMSW.<sup>77</sup> These results support the assertion that AD can reduce  $\text{N}_2\text{O}$  emissions from composting, but it is important to note that Colón et al. observed real-world operations at facilities with similar but not identical OFMSW feedstocks. Maulini-Duran et al. (2013) compared two different types of wastewater treatment sludge: (1) sludge sent directly from a wastewater treatment facility to composting without undergoing AD and (2) sludge at a separate facility, treated with AD and

subsequently sent to composting.<sup>54</sup> They found that  $\text{CH}_4$  and  $\text{N}_2\text{O}$  emissions were respectively 60 and  $>100$  times higher for the post-AD material. However, there was not a proper control in this study because the source material originated from entirely different facilities. The  $\text{N}_2\text{O}$  emission factors reported by Maulini-Duran et al. for composting post-AD sludge aligned better with  $\text{N}_2\text{O}$  emissions reported by several other studies for composting raw, untreated sludge.<sup>19,60</sup>

Preble et al. (2020) measured emissions at a commercial-scale composting facility that processed digestate remaining after dry (high-solids) AD of OFMSW and calculated GHG emission factors per unit of incoming material.<sup>11</sup> This study did not include a control comparison to untreated OFMSW. However, it is notable that their reported  $\text{N}_2\text{O}$  emission factor is approximately 80% lower than mean and median  $\text{N}_2\text{O}$  emission factors for composting untreated OFMSW shown in Table 1. Conversely, Preble et al. report a  $\text{CH}_4$  emission factor that is  $\sim 5$  times higher than the mean value and  $\sim 18$  times higher than the median value for untreated OFMSW (Table 1). Although Preble et al. did not directly measure VS reduction during the dry AD process, the EPA WARM uses a VS reduction of 75% during AD of municipal food waste, a reasonable proxy for OFMSW.<sup>80</sup>

Like Preble et al. (2020), Beylot et al. (2015) studied the emissions from composting post-AD OFMSW and observed  $\text{N}_2\text{O}$  emissions that were 75% lower than what is reported by Preble et al. and per-tonne  $\text{CH}_4$  emissions that were  $\sim 30\%$  higher.<sup>79</sup> Zeng et al. (2016) conducted a series of lab-based trials to assess nitrogen emissions from composting digested OFMSW under a variety of conditions, including varied bulking agents, feedstock mixing ratios, and initial moisture content, and found  $\text{N}_2\text{O}$  emissions ranging from  $5.6 \times 10^{-4}$  to  $3.3 \times 10^{-3}$  kg per tonne of wet feedstock.<sup>61</sup> This range is higher than what has been reported from field measurements of both pre- and post-AD OFMSW composting. While lab-based experiments can be useful for comparing a range of



Table 2. Summary of NH<sub>3</sub> and VOC Emission Factor Data for Composting Raw Materials and Digestate

feedstock	NH <sub>3</sub> emission factors (kg NH <sub>3</sub> /kg of wet feedstock)			VOC emission factors (kg VOC/kg of wet feedstock)		
	mean	median	sample size	mean	median	sample size
manure	$2.04 \times 10^{-3}$	$1.64 \times 10^{-3}$	44	$6.06 \times 10^{-5}$	$6.06 \times 10^{-5}$	2
OFMSW	$1.03 \times 10^{-3}$	$2.79 \times 10^{-4}$	29	$1.71 \times 10^{-3}$	$3.60 \times 10^{-4}$	13
sludge	$7.70 \times 10^{-4}$	$3.27 \times 10^{-4}$	13	$1.77 \times 10^{-4}$	$1.80 \times 10^{-4}$	3
yard waste	$8.91 \times 10^{-5}$	$2.50 \times 10^{-5}$	5	$5.23 \times 10^{-4}$	$4.62 \times 10^{-4}$	4
digestate	$5.50 \times 10^{-4}$	$6.22 \times 10^{-5}$	25	$1.16 \times 10^{-4}$	$3.72 \times 10^{-5}$	11

materials and conditions under controlled conditions, we advise against relying on these values to represent commercial composting conditions.

Based on the limited data available on emissions from pre- and post-AD organic waste, the question is whether there is a defensible method for approximating differences in composting emissions in the absence of reliable measured data. The studies reviewed here suggest that treating waste with AD, thereby lowering its VS content, can subsequently reduce N<sub>2</sub>O emissions during composting relative to the alternative approach of sending untreated material straight to composting without AD. For researchers and practitioners who must approximate emission factors for composting digestate, it may be appropriate to select a measured emission factor for composting raw materials and apply a reduction factor equivalent to the estimated VS reduction. For example, by applying the lowest observed reduction in N<sub>2</sub>O emissions (57%) from directly comparable emission measurements in Li et al. (2018) to the mean values in Table 1, we estimate that composting digested OFMSW emits  $3.9 \times 10^{-5}$  kg of N<sub>2</sub>O per wet tonne and digested manure emits  $2.0 \times 10^{-4}$  kg of N<sub>2</sub>O per wet tonne.<sup>63</sup> In practice, longer AD residence times and greater reductions in VS may lead to further reductions in N<sub>2</sub>O emissions during composting.

Unfortunately, approximating differences in CH<sub>4</sub> emissions may be more challenging than estimating N<sub>2</sub>O. Li et al. (2018) found that changes in CH<sub>4</sub> emissions were dependent on AD residence time, with shorter residence times translating to elevated CH<sub>4</sub> emissions. The increase in CH<sub>4</sub> emissions when comparing the data for digested OFMSW from Preble et al. (2020) and Beylot et al. (2015) to mean or median values reported in Table 1 may be driven by management practices and/or the fact that the material was anaerobically digested. Highly degradable feedstocks, like manure, OFMSW, and digestate, can create oxygen-depleted zones in compost piles that are compacted and/or not sufficiently aerated, thereby increasing CH<sub>4</sub> production.<sup>51,54</sup> Because of variability in pile management and lack of detailed reporting on these practices, it is likely safest to assume that AD has no effect on CH<sub>4</sub> emissions during composting, provided AD residence times are not below industry standard practices.

**4.1.4. Impact of Composting Methods.** As noted previously, different methods for managing compost piles are likely to impact emissions, particularly if some are more effective than others at maintaining aerobic conditions. With regards to composting methods, our analysis focuses on how turning or forced aeration impacts GHG emissions. Using OFMSW as an example, Figure 3 demonstrates the differences in distributions of both CH<sub>4</sub> and N<sub>2</sub>O emission factors when grouping by the aeration method. The median CH<sub>4</sub> emission factor was ~1.5 times higher when the primary aeration method was turning versus forced aeration, and the mean value was nearly 4 times higher. This is supported by a meta-analysis

from Pardo et al. (2015), which found turning to be associated with higher GHG emissions.<sup>22</sup> An important caveat is that this trend may be related to the relatively high number of lab-based studies among those involving forced aeration. Because it is easier to control conditions and maintain proper aeration in laboratory settings that often use enclosed compost reactors, these results may not accurately reflect emissions in industrial scale composting.

As would be expected, several studies have confirmed through measurements that CH<sub>4</sub> emissions decrease with higher aeration rates, but these studies were less consistent in their findings regarding the impact of aeration on N<sub>2</sub>O emissions.<sup>53,58,60,81</sup> Unlike the CH<sub>4</sub> emission factor distributions, the N<sub>2</sub>O emission factor distributions do not diverge significantly based on aeration methods (Figure 3). For instance, the average N<sub>2</sub>O emission factor for composting with turning is only 4% greater than that for composting with forced aeration. There is not sufficient evidence to suggest that the aeration method has a significant impact on N<sub>2</sub>O emissions from composting.

**4.2. Ammonia Emissions.** Table 2 presents the feedstock-specific mean and median NH<sub>3</sub> emission factors from our literature survey. Boxplot visualizations of the NH<sub>3</sub> data are provided in the Supporting Information (Figure S2). As is the case with GHG emissions, the highest average NH<sub>3</sub> emission factor is associated with manure, followed by OFMSW. Composting yard waste emits the least NH<sub>3</sub>. As discussed earlier, NH<sub>3</sub> is a product of microbial decomposition of proteins in the composted waste, and a fraction of that nitrogen will ultimately be emitted as N<sub>2</sub>O. Elevated NH<sub>3</sub> and N<sub>2</sub>O emissions can simply indicate that a protein-rich feedstock is being decomposed through the nitrification and denitrification processes, although these emissions can also be sensitive to compost management methods.<sup>60</sup>

According to Andraskar et al. (2021),<sup>82</sup> maintaining aerobic conditions is imperative for controlling NH<sub>3</sub> and other odorous emissions because many of these compounds are produced from anaerobic processes.<sup>82</sup> Bulking agents can increase porosity to facilitate better aeration; Zhang et al. (2021) found that composting kitchen waste emitted 62% more NH<sub>3</sub> than composting a mixture of 85% kitchen and 15% garden waste.<sup>83</sup> Shao et al. (2014) observed a similar effect on NH<sub>3</sub> emissions as the bulking agent-to-substrate ratio increased.<sup>84</sup> In terms of operational methods, there appear to be trade-offs in NH<sub>3</sub> emission rates when forced aeration is used to maintain aerobic conditions. Several studies have found that intermittent aeration at lower rates reduced NH<sub>3</sub> emissions during swine manure composting.<sup>53,60,81</sup> This is generally supported by other experimental studies that have observed increases in NH<sub>3</sub> emissions when forced aeration increases.<sup>49,53,60</sup> In addition to managing aeration, composters can also use microbial inoculation to control NH<sub>3</sub> emissions.<sup>82,85</sup> Chen et al. (2022) measured an ~20%

reduction in  $\text{NH}_3$  emissions when composting sewage sludge with a compound bacterial consortium relative to the control.<sup>85</sup>

Unlike for  $\text{N}_2\text{O}$ , there is not consistent evidence to suggest whether treating waste with AD prior to composting increases or decreases  $\text{NH}_3$  emissions. A study by Smet et al. (1999) measured odors from OFMSW composting, AD of OFMSW, and digested OFMSW composting and found that AD pretreatment reduced  $\text{NH}_3$  emissions by 73%.<sup>86</sup> Even when including emissions during AD, composting raw OFMSW still emitted 72% more  $\text{NH}_3$  than combined AD and composting. Maulini-Duran et al. (2013) observed a 98% decrease in  $\text{NH}_3$  emissions when comparing raw sludge composting to digested sludge composting; however, as noted earlier, this study did not include a proper experimental control, as the material came from two entirely different facilities.<sup>54</sup> Rincón et al. (2019) compared emissions from 15 different feedstocks, including 5 different digestates, and found that on average and a wet mass basis,  $\text{NH}_3$  emissions were 87% lower for digested materials compared to raw feedstocks.<sup>87</sup> Like Maulini-Duran et al., Rincón et al. did not include proper experimental controls since the feedstocks all came from different sites and the digestates were not derived from the same material in the raw feedstocks. In contrast, Colón et al. (2012) found that  $\text{NH}_3$  emissions from OFMSW composting roughly doubled with AD pretreatment.<sup>77</sup> Similarly, Li et al. (2018) observed an increase of up to 40% in  $\text{NH}_3$  emissions from manure composting when materials first underwent AD.<sup>63</sup> The mean  $\text{NH}_3$  emission factor for composting post-AD materials (including OFMSW, manure, and sludge) is lower than that for raw OFMSW or manure but on the same order of magnitude as the mean value for composting sludge (Table 2). Generally, composting untreated yard waste appears to emit less  $\text{NH}_3$  than composting digestates, but there is no available emissions data on composting digested yard waste.

**4.3. VOC Emissions.** Of all the compounds discussed in this review, VOCs are the least commonly reported, and although individual compounds may have differing effects on local odor concerns and air quality, VOC emissions are typically summed and reported as a total mass. VOCs include ketones, alcohols, terpenes, and other carbon compounds that can participate in atmospheric reactions with the exception of carbon monoxide, carbon dioxide, carbonic acid, metallic carbides or carbonates, and ammonium carbonate as defined by the EPA.<sup>88</sup>  $\text{CH}_4$  is a VOC, although it is often reported separately because of its relevance for climate forcing; remaining VOCs are reported as non-methane VOCs. All of the values reported in this section and in Table 2 exclude  $\text{CH}_4$ . The surveyed literature includes a total of 33 VOC emission factors, 11 of which are for digestate composting. The summary data broken down by feedstock type is provided in Table 2, but the sample sizes are limited ( $n < 5$  for all raw feedstocks except OFMSW), so those emission factors should be used with caution. Further research is required to establish accurate distributions of these feedstock-specific emission factors. Without controlling for feedstock type, the mean emission factor is  $8.14 \times 10^{-4}$ , and the median is  $1.06 \times 10^{-3}$  kg VOC per kg of wet feedstock for composting.<sup>11,54,72,77,84,86,87,89,90</sup> If pressed to assume a nonzero value, researchers and practitioners may choose to use a median or mean value that excludes digestate. For nondigestate feedstocks (including sludge, OFMSW, and yard waste), the

mean emission factor is  $1.18 \times 10^{-3}$ , and the median is  $2.1 \times 10^{-4}$  kg VOC per kg of wet feedstock ( $n = 21$ ).

All of the surveyed VOC emission factors for digestate composting are at least an order of magnitude lower than the average for nondigestate composting, suggesting AD may reduce VOC emissions from composting. Most of the surveyed literature provides evidence to support this conclusion with the exception of Colón et al. (2012) which measured higher VOC emissions from composting post-AD OFMSW compared to raw OFMSW.<sup>77</sup> Smet et al. (1999) compared the VOC emissions from composting and AD of OFMSW, as they did with  $\text{NH}_3$ , and the results show a 99% reduction in VOC emissions from composting when AD pretreatment was used. Expanding the system boundary, the combined AD and composting scenario had 63% fewer VOC emissions than direct OFMSW composting.<sup>86</sup> Maulini-Duran et al. (2013) observed a decrease in VOC emissions from composting post-AD sludge relative to sludge that was not treated with AD.<sup>54</sup> More recently, Rincón et al. (2019) compared odorous emissions from each raw feedstock type and digestate type with the exception of digested yard waste for which they had no data.<sup>87</sup> On average, VOC emissions from digestate composting are 94% lower than those from raw material composting. When controlling for feedstock type, the VOC emission factors for composting digested materials are consistently lower than their raw counterparts, but as stated before, Rincón et al. did not include ideal experimental controls.<sup>87</sup> Beyond AD, other technology options exist to specifically target and reduce VOC emissions from composting; these include but are not limited to pretreatment techniques, incineration, biotrickling filters, bioscrubber technology, and membrane bioreactors.<sup>82,91</sup>

## 5. CONCLUSIONS

Properly accounting for composting emissions, and for organic waste management-related emissions in general, in an LCA can be exceptionally challenging. There are still large gaps in the empirical data available for the range of materials that can be composted and the key greenhouse gases and air pollutants. More fundamentally, there is a limited scientific understanding of the complex microbial communities that break down plant matter, and emissions estimates are likely to evolve as scientists gain an improved understanding of the complex chemical and biological mechanisms at work in these environments. However, by analyzing data reported across the literature and disaggregating emission factors based on pile management strategies and starting material, basic patterns emerge that can inform best-estimates for use in future analyses.

Our findings suggest that  $\text{N}_2\text{O}$  is typically the dominant contributor to the  $\text{GWP}_{100}$  of direct emissions from composting operations in properly aerated piles/windrows, assuming biogenic  $\text{CO}_2$  emissions do not have a net climate impact. When controlling for feedstock type,  $\text{N}_2\text{O}$  accounts for 45–79% based on mean values and 59–91% based on median values of total GHG emissions on a  $\text{GWP}_{100}$  basis. Yard waste is a notable exception where GHG emissions are dominated by  $\text{CH}_4$  (80% of  $\text{GWP}_{100}$  based on mean values or 83% based on median values), likely because of its high C:N ratio compared to other waste types such as food waste and manure. Among observed feedstock types,  $\text{N}_2\text{O}$  emissions appear to be highest for manure and lowest for yard waste and may be influenced by initial VS content.  $\text{N}_2\text{O}$  emissions seem to be impacted by whether the incoming material was previously processed in an

AD facility, and this paper outlines a suggested method for adjusting the estimated N<sub>2</sub>O emission factor based on VS reductions during AD. Conversely, CH<sub>4</sub> appears to be primarily related to whether the pile or windrow is adequately aerated. The impact of VS reduction, through AD or otherwise, prior to composting did not appear to have a substantial impact on CH<sub>4</sub> emissions, although direct comparisons in the empirical data are extremely limited and warrant further study. Assuming CH<sub>4</sub> emissions are heavily influenced by pile management, reducing GHG emissions from well-managed, properly aerated compost piles may require more focus on the composition and quality of feedstock materials to reduce N<sub>2</sub>O emissions.

Regarding NH<sub>3</sub> and VOCs, the available data suggests that treating waste with AD prior to composting may reduce these emissions, but more measurements are required to definitively support this conclusion. This uncertainty is echoed by inconsistent results and disagreement in current scientific literature, emphasizing the need for further research in this area. Reducing these emissions is a key part of improving air quality in local and surrounding communities not only because of odor concerns but also because both NH<sub>3</sub> and VOC contribute to atmospheric formation of PM<sub>2.5</sub>, which has significant human health impacts. Therefore, though AD does not have a clear benefit with respect to limiting GHG emissions from composting, it can still play a role in effective organic waste management because of its potential to reduce other harmful emissions.

## ■ ASSOCIATED CONTENT

### SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.est.2c05846>.

Relevance of ammonia emissions; detailed descriptions of measurement methods; carbon dioxide emissions from composting; boxplot distribution of carbon dioxide emission factors for composting, Figure S1; discussion of net emissions and offsets from compost application to soil; comparison of compost emissions to landfill emissions; and boxplot distribution of ammonia emission factors for composting, Figure S2 (PDF)

Table S1 including emission factors (XLSX)

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## Notes

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