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Title: COMPARISON OF DIFFERENT APPROACHES FOR THE ESTIMATION OF ODOUR EMISSIONS FROM LANDFILL SURFACES

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Abstract: The study addresses the problem of the assessment of odour emissions from landfills. The aim of the project is the research concerning how to quantify odour emissions from landfill surfaces, since there is no evidence of a widely accepted method to evaluate odour emissions from this particular kind of source. Seven different methods have been developed and investigated; these methods can be seen as based on three distinct approaches to the problem, both experimental and computational. The first approach provides to use models for the estimation the landfill gas production, whereby the second and the third approach are based on direct measurement campaigns on the landfill surface, for the determination either of the methane or for the direct measurement of the odour concentration. The methods have then been compared in terms of resulting odour impact by application of a specific atmospheric dispersion model.

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4 **Comparison of different approaches for the estimation of odour**  
5 **emissions from landfill surfaces**

6 Lucernoni, L. Capelli, S. Sironi  
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12 The present work addresses the issue of assessment of odour emissions from landfills, an aspect that in  
13 recent years has become a very important matter, for researchers and managers as well as for the general  
14 public. The aim is to shed some light on the quantification of odour emissions from landfill surfaces, since  
15 there is no evidence of a widely accepted methodology. The methods that are here presented can be seen as  
16 based on three distinct approaches to the problem, both experimental and computational. The first approach  
17 entails the use of specific models for the estimation the landfill gas production, whereby the second and the  
18 third approach are based on direct measurement campaigns on the landfill surface, for the determination  
19 either of the methane or of the odour concentration. The methods have then been compared in terms of  
20 resulting odour impact by application of a specific atmospheric dispersion model.

21  
22 As novelty of this work and its contributions to the field of olfactometry and odour dispersion modelling,  
23 please consider our paper “Comparison of different approaches for the estimation of odour emissions from  
24 landfill surfaces” for publication on Waste Management Journal.

25  
26 The paper was presented during the Sardinia Symposium 2015 and resulted a selected paper considered for  
27 publication on Waste Management Journal.

28  
29 We hope this paper will be of interest for the readers of this journal.  
30  
31

32 Best regards,

33  
34 Laura Capelli and Federico Lucernoni  
35  
36

1 COMPARISON OF DIFFERENT APPROACHES FOR THE ESTIMATION OF  
2 ODOUR EMISSIONS FROM LANDFILL SURFACES

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7 ABSTRACT: The study addresses the problem of the assessment of odour emissions from  
8 landfills. The aim of the project is the research concerning how to quantify odour emissions from  
9 landfill surfaces, since there is no evidence of a widely accepted method to evaluate odour  
10 emissions from this particular kind of source. Seven different methods have been developed and  
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16 compared in terms of resulting odour impact by application of a specific atmospheric dispersion  
17 model.

18 Keywords: odour impact assessment; dispersion modelling; surface sampling; wind tunnel;  
19 satatic hood; flux chamber<sup>1</sup>.

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LFG: landfill gas  
OER: Odour Emission Rate  
SOER: Specific Odour Emission Rate

## 20 **1. INTRODUCTION**

21 Olfactory pollution is the immission of polluting compounds in the atmosphere that, although if  
22 not directly dangerous for the human health, are nonetheless characterized by intense and/or  
23 unpleasant smell. Typical examples are gaseous emissions coming from landfill surfaces (Ying et  
24 al., 2012), intensive farming, etc. This kind of pollution is one of the most significant causes of  
25 environmental discomfort since it lowers the quality of the environment and it may lead to  
26 psychophysiological disorders and a general worsened life quality (Palmiotto et al., 2014). In  
27 Italy there are no comprehensive regulations concerning the problem of odoriferous emissions. In  
28 order to devise proper strategies to reduce the nuisance related to odour emissions it is necessary  
29 to have specific methods for univocal odour assessment (Balling et al., 1980; Hobson, 1995;  
30 Stordeur, 1981), debunking the common belief that odour characterization is more art than  
31 science (Koe, 1989; Jiang, 1996). While in the scientific community there is a satisfactory  
32 agreement regarding the analytical techniques – i.e. dynamic olfactometry for the determination  
33 of odour concentration (Sironi et al., 2014) – odour sampling is a quite more debated task,  
34 especially if diffused emissions or area sources are concerned. The evaluation of odour emissions  
35 from landfills is even more complicated, due to the specific characteristics of this kind of source,  
36 which is surely not an active area source, but neither properly a passive area source, as the  
37 landfill surface is typically crossed by a – low – flux, and there is currently no widely accepted  
38 method for odour assessment on landfills. However, landfills have always been considered as  
39 important sources of unpleasant odours, for this reason the development of specific methods for  
40 the assessment of odours emissions or the definition of specific odour emission factors would be  
41 of great interest both for environmental authorities, as well as for landfill managers and  
42 operators.

43 This is the reason why, in the present work, different methods for the evaluation of odour  
44 emissions from landfills have been inspected: the 7 methods developed are all retraceable to one

45 of three distinct approaches to the matter.

46 The first approach – that comprises methods 1 and 2 - entails the usage of specific software for  
47 the quantification of the landfill gas (LFG) production: the software used in the first method is  
48 the US-EPA LandGEM, while the second method exploited a similar software specifically  
49 developed for the present project with some improved features with respect to the US-EPA  
50 LandGEM.

51 The second approach – that includes the third method – relies on the direct measurement of CH<sub>4</sub>  
52 on the landfill surface, which involved the necessity to define a sampling methodology tailored  
53 for this peculiar type of source.

54 The third approach – that comprises methods 4, 5, 6 and 7 – involves the direct measurement of  
55 the odour concentration at the source. In the fourth method the concentration was considered  
56 independent from the wind speed, while in methods 5, 6, 7 the landfill surface was treated as a  
57 liquid area source, thereby considering the concentration as a function of the wind speed on the  
58 surface (Sironi et al., 2005).

59 Since the first two approaches evaluated are based on the quantification of methane emissions,  
60 the Odour Emission Rate (OER) needs to be obtained indirectly by multiplying the emitted gas  
61 flow rate by the LFG odour concentration. The odour concentration ( $c_{od}$ ) of the LFG emitted  
62 through the landfill surface was estimated by means of a correlation investigated between  
63  $c_{od}$  and  $c_{CH_4}$ .

64 The OER values obtained with the seven described methods were then used as inputs to run the  
65 atmospheric dispersion model CALPUFF, which allowed to visualize the odour impacts of the  
66 studied landfill resulting from the different emission scenarios considered. The impacts have then  
67 been compared with one another in order to make some consideration about the different models'  
68 accuracy and reliability.

69 **2. MATERIALS AND METHODS**

70 **2.1 The Studied Site**

71 The chosen landfill is located in Northern Italy, and it is classified as “Landfill for Non-  
72 Hazardous Municipal Solid Waste Disposal”.

73 The site was chosen because it is object of repeated olfactometric monitoring campaigns by the  
74 Olfactometric Laboratory of Politecnico di Milano since several years: this allows to have access  
75 to a great amount of emission, olfactometric and meteorological data, as well as a consolidated  
76 experience about the landfill odour impact gained throughout the years, both crucial aspects for  
77 the present work. The landfill is operational since the early 1990s and has a waste processing  
78 capacity of several millions cubic meters, making it one of the biggest landfills in Northern Italy.  
79 The site is subdivided in six allotments: now only one allotment is still active, all the others are  
80 closed.

81 In order to be able to apply the dispersion model to the emissions coming from a landfill, the  
82 whole area (about 250'000 m<sup>2</sup>) needs to be considered as an emission source. For modelling  
83 purposes the landfill surface was divided in a reasonable number of emissive cells having smaller  
84 dimensions: 40 emissive cells were considered, a number of sources that is in line also with the  
85 suggestions of the User's Manual of the CALPUFF software (Scire et al., 2000). The geometry  
86 of the cells was kept as simple as possible (simple shapes like squares and triangles).

87 In order to glean the emission scenarios relying on an experimental approach it was also  
88 necessary to carry out tailored campaigns in the field. The campaigns spanned from 1/3/2014 to  
89 30/10/2014 and interested mainly allotment 1, which is the oldest one.

90 The software initially used to assess the methane production is the US-EPA LandGEM  
91 (Alexander et al., 2005); hereafter another newer software was adopted, similar to the default  
92 LandGEM but with some reasoned modifications. This will be discussed in the following  
93 sections.

94 **2.2 Modifications To The LandGEM Software**

95 One of the novel aspects of the present study regarded the modification and improvement of the  
96 US-EPA LandGEM software aiming to optimize its performances. The main improvement  
97 concerns the input mode of the initial parameters of the model which the software is very  
98 sensitive to. More in detail, the biggest flaw in the default model is in the input procedure of the  
99 fundamental parameters of the model, since besides the waste inflow that can be considered  
100 changing year by year, the other parameters, i.e. the methane generation rate ( $k$ ) and the methane  
101 generation potential ( $L_0$ ), are kept constant for the whole simulation (Alexander et al., 2005).  
102 This aspect makes it impossible to account for the possible variations that may occur as a  
103 consequence of a change in the processed MSW quality, given that both ( $k$ ) and ( $L_0$ ) show a  
104 strong dependence on the bio-degradability of the considered waste.

105 The many regulations addressing the problem of landfill management have defined the “classes”  
106 of wastes treated there and this classification has been known to change over time. In the end it is  
107 the class of the waste that affects the most the two input parameters ( $k$ ) and ( $L_0$ ): at this point it is  
108 evident that assuming them unchangeable is an unacceptable approximation. Thus, the  
109 innovative idea is the possibility of considering these two key parameters as well varying year by  
110 year. The modified software maintains the same Graphical User Interface (GUI) in MS Excel as  
111 the standard US-EPA LandGEM and allows to consider the parameters ( $k$ ) and ( $L_0$ ) changing  
112 yearly.

113 **2.3 Design And Development Of Specific Sampling Procedures**

114 It is necessary to clarify the particular nature of the landfill source typology. This kind of source  
115 is an area source but it cannot be defined neither active nor passive, referring to the classification  
116 made in the german guideline VDI 3880 (VDI, 2011). In fact, even if not presenting a significant  
117 outflow – typical of the active sources – it has a distinct non-zero emission flow that cannot be

118 neglected. This is why it is necessary to develop specific sampling devices and procedures for  
119 this unique type of source.

120 The first method adopted for the direct measure of methane emissions on the considered landfill  
121 surface was the so-called Static Hood device (SH) that was specifically realized starting from a  
122 design found in the scientific literature that was deemed appropriate (Rachor et al., 2013). The  
123 mentioned work was picked as reference since it is a rare example of a study that involved tests  
124 and experiments in the field that were repeatable and that were performed over a rather long  
125 period (i.e. 2 years). The hood geometry proposed in that research paper is cylindrical, 50 cm  
126 high with a base section of 0.12 m<sup>2</sup>. Within the research group further considerations were  
127 pondered concerning the geometry of the hood. In particular, about the height of the chamber two  
128 considerations were made, concerning on one hand molecular diffusion, the most significant  
129 phenomenon in the process, and on the other hand the CH<sub>4</sub> ascension velocity, which depends on  
130 the LFG outflow from the landfill surface, and was hypothesized as monodimensional  
131 (perpendicular to the surface).

132 Since both processes are rather slow, it was concluded that it was necessary to have a shorter  
133 hood, minimizing its height in order to have a homogeneous methane concentration profile in the  
134 hood. The chamber assembled at Politecnico is 10 cm high, allowing both an easy placement of  
135 the hood on the surface and an accurate measure; the base section is 0.25 m<sup>2</sup> and the material  
136 used is steel, providing the required robustness to withstand the stresses involved in the laying of  
137 the device. The chamber is provided with lateral flanges to avoid excessive plunging in the soil.  
138 The hood is equipped with an open tube that connects the interior to the exterior of the hood, in  
139 order to keep the internal pressure equal to the external pressure, thus avoiding overpressures that  
140 may affect the LFG emission from the surface portion covered by the hood.

141 The emitted methane flow can be evaluated according to the following mass balance:

142 
$$Q_{CH_4} = \frac{V_{hood}}{S_{hood}} * \frac{\partial c_{CH_4}}{\partial t} \quad (1)$$



143 Once the CH<sub>4</sub> flow rate is known, it is possible to calculate the LFG emission by considering the  
144 LFG composition (i.e., the CH<sub>4</sub> content). A specific procedure for sampling was designed,  
145 providing that after each measurement the hood is lifted and ventilated, in order to minimize  
146 LFG build-up in the hood and perturbations on the emission source. Afterwards, the hood is  
147 positioned on the same spot and the next measure is performed, restarting the time count from  
148 time zero.

149 After the found experimental correlation was proved to agree with the theoretical expectations,  
150 the sampling time for measurements with the static hood was defined in 10 minutes. The hood  
151 developed is portrayed in Fig.1.

152

153 Figure 1

154 Another device was involved in the study, a Flux Chamber (FC) similar to the one designed by  
155 the US EPA (Klenbusch, 1986) to be used in fluxed experiments. The hood realized at  
156 Politecnico is hemispherical with a 50 cm diameter and a 32 dm<sup>3</sup> volume. The emitted methane  
157 flow in this case can be calculated as shown in Eq.2 where the fluxes are expressed in dm<sup>3</sup>/h:

$$158 \quad F_{air} + F_{CH_4} = F_{out} \quad (2)$$

159 This sampling procedure is the one used for collecting samples to be analysed by means of  
160 olfactometry. The main reason behind is that odour samples have typical volumes of 6 dm<sup>3</sup> and  
161 considering a static hood with a volume of 25 dm<sup>3</sup> this would mean a huge perturbation of the  
162 system that would affect the results; the inconvenience can be overcome operating with a Flux  
163 Chamber where the volume inside the hood is constantly recirculated thanks to the continuous air  
164 flow. Also in this case different tests were performed: several experiments were conducted  
165 adopting an air flow rate ranging between 100 dm<sup>3</sup>/h and 300 dm<sup>3</sup>/h, thereby verifying that the  
166 measured concentration is inversely proportional to the flow rate. Then, a specific sampling

167 procedure for the measurement of both ( $c_{od}$ ) and ( $c_{CH_4}$ ) with the Flux Chamber was defined.  
168 The neutral air flow rate was defined in 200 dm<sup>3</sup>/h while the sampling time in 12 minutes. The  
169 Flux Chamber is depicted in Fig.2:

170

171 Figure 2.

172 Another device that was used in the present work was the Wind Tunnel (WT), which was  
173 designed at Politecnico in a previous project for odour sampling from passive area source  
174 (Capelli et al., 2009), with the aim of making the flow fluxed in the hood as uniform as possible,  
175 in order to realistically simulate the action of the wind on the surface (Jiang et al., 1995). Even if  
176 this sampling system was validated and codified only for liquid area sources – while the studies  
177 to validate it also for solid area sources are still ongoing (Capelli et al., 2012) – it was decided to  
178 use it for the determination of the odour emissions from the landfill surface since it represents the  
179 only sampling system available that has been validated for passive area sources. However, it is  
180 known that for area sources characterized by a very low emissivity, it tends to overestimate the  
181 actual emission (Frechen et al., 2004; VDI, 2011). Previous studies on the Wind Tunnel led to  
182 the definition of the optimal sweep air flow rate, found to be 2500 dm<sup>3</sup>/h (Capelli et al., 2009). At  
183 the outlet of the Wind Tunnel an odour sample is collected in a specific Nalophan<sup>TM</sup> bag by  
184 means of a vacuum pump. From the odour concentration it is possible to evaluate the Specific  
185 Odour Emission Rate (SOER) expressed in [ou<sub>E</sub>/s/m<sup>2</sup>] according to Eq.3:

186 
$$SOER = \frac{c_{od} * Q_{air}}{A_{WT}} \quad (3)$$

187 With:  $c_{od}$ = odour concentration [ou<sub>E</sub>/m<sup>3</sup>]

188  $Q_{air}$ = sweep air flow rate [m<sup>3</sup>/s]

189  $A_{WT}$ = base section of the WT [m<sup>2</sup>]

190

191 The OER is then obtained by multiplying the SOER by the site surface. The outlet odour  
192 concentration, the SOER and the OER, i.e. the amount of odorous compounds that migrate to the  
193 gas phase as a consequence of forced convection, are a function of the defined sweep flow rate.  
194 More in detail, considering the motion regime laminar, assuming that the mass transfer  
195 phenomena can be described by the laws of Prandtl's Fluid-Dynamic Boundary Layer Theory, it  
196 is possible to deduce that both SOER and OER will be proportional to the square root of the  
197 sweep air velocity as shown in Eq.4, where  $\alpha$  is a proportionality constant and the hypothesis of  
198 laminar motion regime holds true:

$$199 \quad SOER, OER = \alpha v^{\frac{1}{2}} \quad (4)$$

200 This means that the OER data obtained with the WT will need to be recalculated at the correct  
201 wind speed for each hour of the simulation time domain, before using them as inputs for the  
202 CALPUFF dispersion model (Jiang et al., 2001; Sohn et al., 2003); this operation can be done  
203 according to Eq.5:

$$204 \quad OER_{v_2} = OER_{v_1} \left( \frac{v_2}{v_1} \right)^{\frac{1}{2}} \quad (5)$$

205 In Eq.5  $v_1$  is the air speed during the sampling conditions – in our case corresponding to 0.035  
206 m/s – while  $v_2$  is the wind speed at a specific hour of the time domain of the simulation.

## 207 **2.4 Evaluation Of The Emitted LFG Odour Concentration**

208 In order to be able to assess the odour emissions associated with the LFG emissions from the  
209 landfill surface, it is necessary to know the effluent flow rate and its odour concentration,  
210 required for OER calculation. The determination of the odour concentration of the LFG emitted  
211 from the landfill surface is a quite complicated task: in literature there are some odour  
212 concentration values relevant to pure LFG (Sironi et al., 2005), but these values overestimate the  
213 concentration of the LFG emitted through the landfill surface, as they do not consider the effects  
214 of odour concentration reduction that occur while the gas crosses the landfill surface cover

215 material (Capanema et al., 2014). For this reason, it was decided to try to estimate the LFG odour  
216 concentration by investigating the correlation between the CH<sub>4</sub> concentration (in ppm) and the  
217 odour concentration (in ou<sub>E</sub>/m<sup>3</sup>). For this purpose, specific sampling campaigns at the landfill  
218 were conducted, where the sampling method involved the Flux Chamber both for the assessment  
219 of CH<sub>4</sub> concentration and for the retrieval of odour samples for olfactometric analysis. The  
220 resulting values underwent a screening process: in particular the odour concentration values  
221 below 40 ou<sub>E</sub>/m<sup>3</sup> were neglected due to the detection limits that are characteristic of the  
222 olfactometric analysis. The remaining data were then plotted with methane concentration on the  
223 axis of abscissae and the odour concentration on the axis of ordinates.

224 The resulting correlation is here reported:

$$225 \qquad y = 0.6907x + 83.026 \qquad (6)$$

226 In Eq.6 (x) is ( $c_{CH_4}$ ) in ppm while (y) is ( $c_{od}$ ) in ou<sub>E</sub>/m<sup>3</sup>. Since in this case the CH<sub>4</sub> concentration  
227 in LFG is assumed to be 50%, i.e. 500000 ppm, the resulting odour concentration for the LFG  
228 turned out to be 345000 ou<sub>E</sub>/m<sup>3</sup>.

## 229 **2.5 Optimization Of The Model For The Assessment Of The Wind Speed Depending** 230 **Emission Rate**

231 When it is necessary to recalculate OER as a function of the wind speed, it is necessary to define  
232 which wind speed has to be considered. In this sense, it was decided to investigate what  
233 differences were observed when using the wind speed recalculated at source level (at 2 m)  
234 instead of the one recorded by the meteorological station (at 10 m). Two possible laws were  
235 considered for wind speed recalculation: a Power Law and a Logarithmic Law. The resulting  
236 models and impacts were then compared with one another:

- 237 - in the first case the OER recalculation considered the wind speed recorded at 10 m;
- 238 - in the second case the OER recalculation considered the wind speed at 2 m recalculated

239 with a Power Law model that is obtained empirically starting from a known velocity at  
 240 certain height, the height corresponding to the desired wind velocity and a so-called  
 241 Hellman's parameter that depends on terrain and stability class. This can be viewed  
 242 formulaically in Eq.7. The model is suggested in the heights range 30-300 m (Cook,  
 243 1997);

$$244 \quad v_w^{h1} = v_w^{h2} * \left(\frac{h1}{h2}\right)^\alpha \quad (7)$$

245 - in the third case the OER recalculation considered the wind speed at 2 m evaluated with a  
 246 Logarithmic Law obtained as a consequence of fluid-dynamic considerations concerning  
 247 the inertial sub-layer, considering that both the "Law of the Wall" and the "Velocity  
 248 Defect Law" hold true, that produces the wind profile expressed by Eq.8 (Drew et al.,  
 249 2013; Tieleman, 2008):

$$250 \quad v_w^h = \frac{u_*}{k} \left[ \ln\left(\frac{h}{z}\right) + 5.75 \left(\frac{h}{H}\right) - 1.88 \left(\frac{h}{H}\right)^2 - 1.33 \left(\frac{h}{H}\right)^3 + 0.25 \left(\frac{h}{H}\right)^4 \right] \quad (8)$$

252 where H is the neutral boundary layer that can be calculated according to Eq.9:

$$253 \quad H = \frac{u_*}{Bf} \quad (9)$$

255 *f* is the Coriolis parameter, B is an empirical constant that can be assumed equal to 6.

256

### 257 **3. RESULTS AND DISCUSSION**

#### 258 **3.1 Determination Of The Emitted LFG Flow Rate With LandGEM, normal and modified**

259 It is necessary to underline that the LandGEM provides the LFG production rate, not the LFG  
 260 emission, which is the quantity required for the OER calculation. First, it is necessary to input the

261 parameters for the US EPA software. According to the regulations in force since 1993 and until  
262 2013, the value of the production rate (k) during the landfill operation years was determined.  
263 Then the arithmetical mean of these values was picked as the value to be used in the simulations  
264 involving the standard US EPA software. The obtained value is reported in Eq.10:

$$265 \quad k = 0.038 \text{ y}^{-1} \quad (10)$$

266 Analogously, also for the biogas generation potential ( $L_0$ ) the mean value was used for running  
267 the program; the resulting value is shown in Eq.11:

$$268 \quad L_0 = 135 \text{ m}^3/\text{Mg} \quad (11)$$

269 The  $\text{CH}_4$  concentration in the LFG was considered equal to 500'000 ppm, which is the average  
270 value relevant to the site under investigation. It is possible to run the LandGEM software and  
271 obtain the flow rate of the  $\text{CH}_4$  produced by the landfill for the year 2014, reported in Eq.12:

$$272 \quad Q_{LFG} = 3.545 * 10^7 \text{ m}^3/\text{y} \quad (12)$$

273 Then, knowing the LFG flow rate collected by the collection system, which in the studied case is  
274 equal to 2200  $\text{m}^3/\text{h}$ , the LFG emission can be then evaluated as shown in Eq.13:

$$275 \quad Q_{emitted} = Q_{tot} - Q_{collect} = 1872.77 \text{ m}^3/\text{h} \quad (13)$$

276 The landfill OER is then calculated considering the LFG odour concentration, which was  
277 considered equal to 345'000  $\text{ou}_E/\text{m}^3$  (Eq. 6) as shown in Eq.14:

$$278 \quad OER = Q_{emitted} * c_{od} = 179473 \text{ ou}_E/\text{s} \quad (14)$$

279 It is also possible to obtain the SOER of the landfill, by dividing the OER by the surface of the  
280 site as expressed in Eq.15:

$$281 \quad SOER = \frac{OER}{A_{tot}} = 0.88 \text{ ou}_E/(\text{s} * \text{m}^2) \quad (15)$$

282 From Eq.15 it is possible to determine the OER for each emissive cell, multiplying the SOER by  
283 each cell surface. The outcome is the input for the CALPUFF dispersion model. The obtained

284 emission data are used as input for the CALPUFF atmospheric dispersion model, producing the  
285 impact depicted in Fig.3.

286 The odour impact in Fig.3 is represented on a 4 km by 4 km map centred on the landfill, the  
287 shown iso-concentration lines represent the 98<sup>o</sup> percentile of the hourly averaged odour  
288 concentrations considering a peak-to-mean ratio of 2.3; this means that the mean values are  
289 multiplied by a factor that accounts for peak oscillations around the mean value of concentration  
290 over 60 minutes (Schauberger et al., 2012). The isopleths represented in the impact map refer to  
291 the range of concentration between 1 and 10 ou<sub>E</sub>/m<sup>3</sup>.

292 These parameters are a commonly accepted choice and are those suggested by the guidelines  
293 issued by Regione Lombardia (D.g.r. n. IX/3018, 2012), which is the only regulation addressing  
294 the problem of olfactory pollution available in Italy. Looking at Fig.3 it is possible to observe a  
295 significant impact since the isopleths cover almost the entire map. This result is likely to be an  
296 over-estimation of the real impact since the many years of odour monitoring campaigns  
297 conducted by the Olfactometric Laboratory of Politecnico at this exact site never highlighted a  
298 critical situation such as the one depicted in Fig.3.

299 Regarding the application of the modified LandGEM, considering the average rainfall of the site  
300 and the influence that the nature of the waste has on the LandGEM parameters (k) and (L<sub>0</sub>), their  
301 values were defined for each year of landfill operation. The biogas amount produced in 2014,  
302 computed with the modified software is reported in Eq.16:

$$303 \quad Q_{tot} = 3.427 * 10^7 \text{ m}^3/y \quad (16)$$

304 Then it was possible to evaluate the actually emitted flow rate considering the same collected  
305 flow of 2200 m<sup>3</sup>/h. Then the OER and SOER can be assessed as in the previous case. The  
306 calculations are shown in Eq.17, Eq.18 and Eq.19:

$$307 \quad Q_{emitted} = Q_{tot} - Q_{collect} = 1712.58 \text{ m}^3/h \quad (17)$$

$$308 \quad OER = Q_{emitted} * c_{od} = 164122 \text{ ou}_E/s \quad (18)$$

309

)

310

$$SOER = \frac{OER}{A_{tot}} = 0.80 \text{ ou}_E / (s * m^2) \quad (19)$$

311

From Eq.19 it is possible to determine the OER for each emissive cell, multiplying the SOER by

312

each cell surface. The outcome is the input for the CALPUFF dispersion model (Fig. 3, right).

313

314

Figure 3.

315

In Fig.3 it is possible to see the impact resulting from this method. Even if the OER is reduced

316

with respect to the first method, the impact shown in Fig.6 is still overestimated. The new model

317

with modified LandGEM shows an improvement since the impact over-estimation is reduced;

318

but the main criticism relevant to this approach is that both models produce an estimate of the LFG

319

production, not the emission and this is conceptually a step back with respect to the

320

characterization of the LFG emissions from landfill surfaces.

321

### 3.2 Experimental Measure Of The CH<sub>4</sub> Concentration On The Landfill Surface

322

The results here presented refer to CH<sub>4</sub> concentration measures conducted in the field in the year

323

2014 adopting a sampling methodology that involved the use of a Flux Chamber with a neutral

324

air flow rate of 200 dm<sup>3</sup>/h. After evaluating the mean of all measured values, which turned out to

325

be equal to 47 ppm, it was possible to determine the LFG flow rate by means of a mass balance

326

written for the Flux Chamber. Assuming as usual that the methane concentration in the pure

327

biogas is 50% v/v, it is possible to compute the emitted biogas flow rate as shown in Eq.20:

328

$$Q_{LFG} = \frac{Q_{CH_4}}{c_{CH_4}^{theo,pure}} = 0.003 \frac{m^3}{s} \quad (10)$$

329

Using the correlation of Eq.6 the odour concentration of the biogas can be evaluated – equal to

330

345'000 ou<sub>E</sub>/m<sup>3</sup> – and consequently also the OER, as shown in Eq.21:



331 
$$OER_{landfill} = c_{od} * Q_{LFG} = 1023.5 \frac{ou_E}{s} \quad (21)$$

332 The SOER is then calculated according to Eq.22, from which it is possible to determine the OER  
 333 for each emissive cell. The outcome is the input for the CALPUFF dispersion model which  
 334 provides the impact depicted in Fig.4 (left).

335 
$$SOER_{landfill} = \frac{OER_{landfill}}{A_{tot}} = 0.005 \frac{ou_E}{s \cdot m^2} \quad (22)$$

336 Just looking at the OER and SOER values obtained with this method it is possible to foresee that  
 337 the resulting impact will be much less significant than in the previous cases, and looking at Fig.4  
 338 makes it visually evident. It is important to highlight that in all impact maps the concentration  
 339 range represented is 1-10  $ou_E/m^3$ , but in this particular case since the maximum value is lower  
 340 than 1  $ou_E/m^3$ , the selected range is 0.1-1  $ou_E/m^3$ . A very low emission such as the one obtained  
 341 in this scenario can be explained since the sampling campaigns have been conducted only on  
 342 allotment 1, the oldest one that is closed since 1994 and this probably led to an under-estimation  
 343 of the real impact of the landfill, which is still operational.

344

345 Figure 4.

346 **3.4 Experimental Measure Of The Odour Concentration On The Landfill Surface**

347 Adopting the same sampling methodology used in method 3 (Flux Chamber), in this case 11 air  
 348 samples were collected from the site that underwent an olfactometric analysis in order to  
 349 determine their odour concentration. The geometric mean of the odour concentrations was  
 350 calculated and turned out to be 135  $ou_E/m^3$ . The SOER was obtained as shown in Eq.23:

351 
$$SOER = \frac{c_{od} \left[ \frac{ou_E}{m^3} \right] * Q_{air} \left[ \frac{m^3}{s} \right]}{A_{hood} [m^2]} = \frac{135 * 200}{1000 * 3600 * 0.25} = 0.03 \frac{ou_E}{m^2 \cdot s} \quad (23)$$

352 Also in this situation both the OER and the resulting odour impact are very low, as can be seen in

353 Fig.4 (right). Methods 3 and 4 produce relatively coherent results, thus confirming the hypothesis  
354 of a correlation existing between odour concentration and CH<sub>4</sub> concentration. Even if both  
355 methods under-estimate the real emissions, because relying on data gathered only from allotment  
356 1, the approaches involving experimental campaigns in the field with the Flux Chamber seem to  
357 be the most reliable tools for the assessment of odour emissions from landfill surfaces.

### 358 **3.5 Assessment Of The OER As A Function Of The Wind Speed**

359 In this scenario the landfill is assumed as a completely passive area source; thus conducting  
360 sampling by means of a Wind Tunnel. In this case 15 samples were collected and analysed,  
361 giving an average odour concentration of 218 ou<sub>E</sub>/m<sup>3</sup>. Then the SOER was evaluated according  
362 to Eq.24:

$$363 \quad SOER = \frac{c_{od} * Q_{air}}{A_{WT}} = 1.21 \frac{ou_E}{m^2s} \quad (24)$$

364 In this case, the OER obtained needs to be pre-processed before being used in the dispersion  
365 model; it is necessary to recalculate it as a function of the actual wind velocity at each hour of the  
366 simulation time domain according to Eq.5.

367 In method 5 the wind speed v<sub>2</sub> is the velocity recorded at a meteorological station nearby, at a  
368 height of 10 m. The velocity v<sub>1</sub> is the velocity inside the Wind Tunnel, i.e. 3.5 cm/s. The  
369 obtained OER values are used as input of the dispersion model producing the impact shown in  
370 Fig.5 (left). It is possible to observe that the isopleths cover the whole area of the map. The odour  
371 impact is very high and, as discussed for methods 1 and 2, it is an overestimation with respect to  
372 the real situation.

373 In the last two scenarios – methods 6 and 7 – as previously mentioned, the wind speed used for  
374 the OER recalculation is the one evaluated at the source level, i.e. 2 m, with tailored wind profile  
375 models exploiting a Power Law and a Logarithmic Law, respectively. The resulting odour  
376 impacts obtained with the dispersion model CALPUFF can be observed in Fig.5 (centre and

377 right).

378 Figure 5.

379 It is worth underlying that methods 5, 6, 7 all share a common flaw, given by the fact that they  
380 consider the landfill surface as a passive area source, and therefore the emissions as a function of  
381 the wind speed over the emitting surface. It is likely that the LFG emissions from landfill  
382 surfaces have a dependence on the meteorological parameters that may affect the phenomenon,  
383 but the assumption that there is a well-defined dependence on the wind speed – which is true for  
384 passive area sources – for this source typology, is groundless since in this case the driving  
385 phenomenon of the whole process is not forced convection.

### 386 **3.6 Comparison Of The OER Obtained With The Different Methods**

387 In order to compare the different approaches adopted, Tab.1 summarizes the OER values  
388 obtained with methods 2, 4 and 7. These three were picked since – among the proposed  
389 approaches – they look like the best LandGEM-based method, the best Flux-Chamber-based  
390 method and the best Wind-Tunnel-based method respectively. More in detail, Tab.1 highlights  
391 how the OER order of magnitude changes from one method to the next one. Relying on a careful  
392 consideration and on the direct experience, it is believed that the most reliable methods are  
393 methods 3 and 4: they are experimental methods involving direct measurement campaigns in the  
394 field with a tailored sampling system adopting a specifically designed Flux Chamber device.

## 395 **4. CONCLUSIONS**

396 The purpose of the present work was the investigation of specific methods for the quantification  
397 of odour emissions from landfill surfaces since up to date there are no codified and universally  
398 accepted methodologies to address this peculiar problem. In facts, landfills represent a particular

399 type of area sources that cannot be considered nor fully active nor fully passive.

400 Table 1. Estimated OER comparison for methods 2, 4 and 7.

Cell Name	Cell UTM East Coordinates [m]	Cell UTM North Coordinates [m]	OER Mod. 2 [ou <sub>E</sub> /s]	OER Mod. 4 [ou <sub>E</sub> /s]	OER Mod. 7 [ou <sub>E</sub> /s]
A01	493808 E	5056844 N	5156.4	192	14532.3
A02	493881 E	5056871 N	5156.4	192	14532.3
A03	493953 E	5056899 N	5156.4	192	14532.3
A04	494029 E	5056930 N	5156.4	192	14532.3
A05	494095 E	5056955 N	5156.4	192	14532.3
A06	493774 E	5056904 N	5156.4	192	14532.3
A07	493845 E	5056932 N	5156.4	192	14532.3
A08	493917 E	5056962 N	5156.4	192	14532.3
A09	493993 E	5056991 N	5156.4	192	14532.3
A10	494060 E	5057016 N	5156.4	192	14532.3
A11	493733 E	5056973 N	5156.4	192	14532.3
A12	493805 E	5057001 N	5156.4	192	14532.3
A13	493877 E	5057029 N	5156.4	192	14532.3
A14	493552 E	5057059 N	5156.4	192	14532.3
A15	494016 E	5057092 N	5156.4	192	14532.3
A16	493692 E	5057041 N	5156.4	192	14532.3
A17	493764 E	5057069 N	5156.4	192	14532.3
A18	493837 E	5057097 N	5156.4	192	14532.3
A19	493911 E	5057128 N	5156.4	192	14532.3
A20	493650 E	5057109 N	5156.4	192	14532.3
B01	493724 E	5057138 N	2922.2	108.8	8235.7
C01	493796 E	5057166 N	3931.7	146.4	11080.9
D01	493870 E	5057196 N	4816.4	179.3	13574.1
E01	493921 E	5057225 N	3113.2	115.9	8773.9
F01	493609 E	5057177 N	1728.2	64.4	4870.6
G01	493682 E	5057206 N	4146	154.4	11684.9
H01	493756 E	5057236 N	4146	154.4	11684.9
I01	493830 E	5057266 N	4413.5	164.3	12438.8
J01	493538 E	5057150 N	3741.6	139.3	10545
K01	493558 E	5057108 N	1423.6	53	4012.3
L01	493590 E	5057055 N	3029.4	112.8	8537.8
M01	493622 E	5057001 N	1482.5	55.2	4178.1
N01	493964 E	5057166 N	513.2	19.1	1446.4
O01	493674 E	5056935 N	1031.3	38.4	2906.5
P01	493526 E	5057177 N	1218.2	45.4	3433.3
Q01	493644 E	5057246 N	4060.6	151.2	11444.2
Q02	493715 E	5056895 N	4060.6	151.2	11444.2

Q03	493529 E	5057202 N	4060.6	151.2	11444.2
Q04	493748 E	5057281 N	4060.6	151.2	11444.2
R01	493881 E	5057279 N	3096.2	115.3	8726.2

401

402 Different methods for the evaluation of odour emissions from landfill surfaces were developed  
403 and investigated. At first the LandGEM-based method was considered and a new possibility was  
404 explored modifying the default US-EPA software. The improved software designed allows to  
405 consider the two most significant input parameters of the LandGEM model variable yearly  
406 according to a set of aspects such as waste quality and composition. The new software shows an  
407 improvement with respect to the standard version of LandGEM, reducing the over-estimation by  
408 10%. This improvement does not fix the underlying flaws of the concept of using a model for the  
409 estimation of the LFG generation for environmental emission assessment purposes (Capelli et al.,  
410 2014).

411 Another novel feature of the research is the optimization of the OER recalculation procedure  
412 involved in the methods relying on measurements with the Wind Tunnel. The changes also in  
413 this case led to an improvement of the previous situation but ultimately proved that anyhow the  
414 application of the Wind Tunnel to landfills, i.e. their assimilation to passive area sources, even  
415 though they characterized by a minimal yet non-negligible emissive flow which is not affected  
416 by forced convection, leads to an over-estimation of the OER.

417 Another aspect of the work entailed the estimation of the odour concentration of the LFG emitted  
418 through the landfill surface by means of specific olfactometric campaigns aiming to correlate  
419 odour and CH<sub>4</sub> concentration of the emitted LFG. The found correlation even though not  
420 exceptional is satisfactory, but further studies will be required in order to increase data  
421 robustness. In the future probably the samples should be collected not only in allotment 1  
422 (closed), but in all allotments including the still active one.

423 Finally, the most significant result of the present work was the clear statement that experimental  
424 campaigns on site are fundamental for these kind of evaluations. Moreover, two sampling

425 systems, i.e. Static Hood and Flux Chamber, were designed and tested, and the optimal sampling  
426 procedures were defined for the two devices; this led to the determination of a precise chain of  
427 actions for both the Static Hood and the Flux Chamber measurements.

428 Actually, it is here necessary to stress that the proposed emission models will have to be further  
429 compared and weighed by means of specific validation measurement campaigns in the field  
430 involving for instance field inspections, questioning and/or electronic noses (Capelli et al., 2013).  
431 Moreover, future investigations and experimental campaigns will need to span on larger temporal  
432 scales in order to be able to highlight other aspects affecting emissions such as seasonal effects  
433 and dependence on meteorological conditions.

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533

534 FIGURES

535

536 Figure 1:



537

538

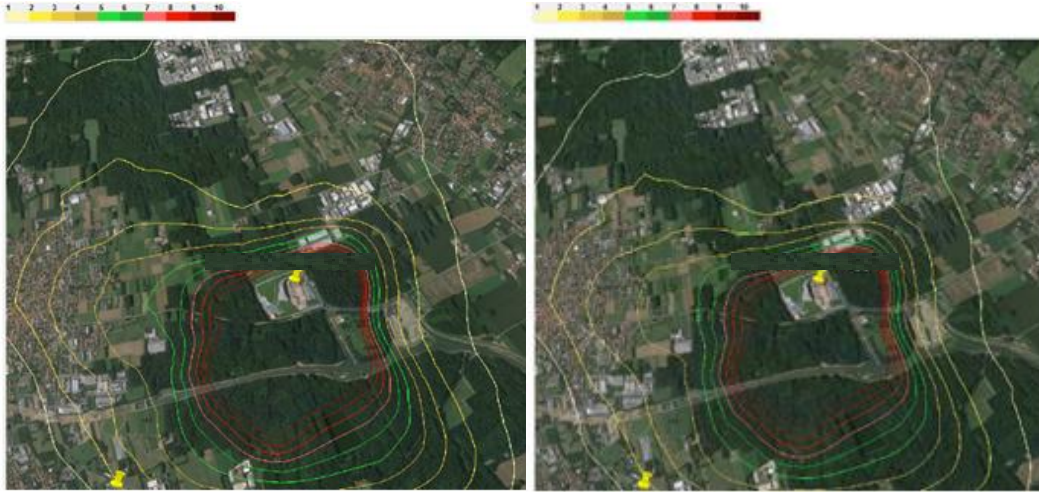
539 Figure 2:



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542 Figure 3:



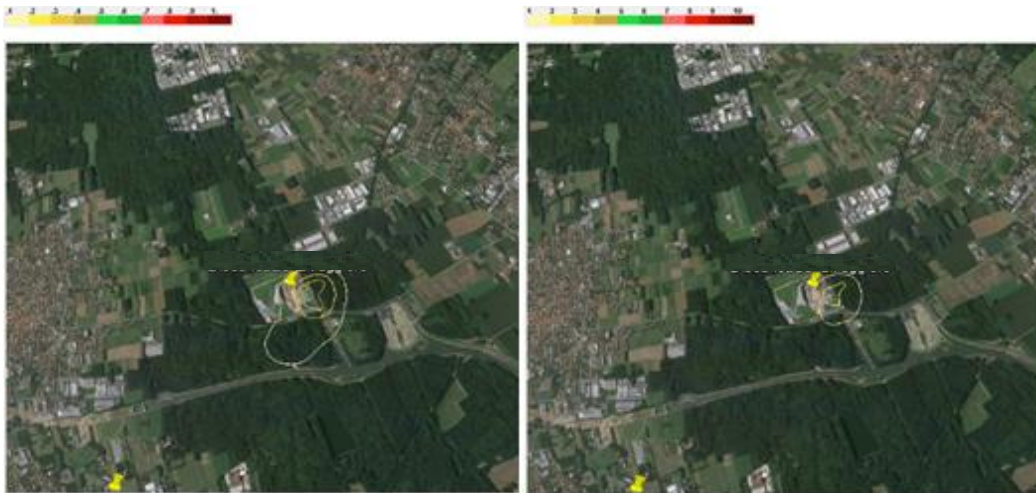
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Figure 4:



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Figure 5:



550

1 Highlights for the article (<85 characters/highlight):

2

- 3
- Odour emission assessment from landfills is important for landfill operation and management.
- 4
- Different methods for the evaluation of odour emissions from landfills were inspected.
- 5
- A new LandGEM-like software was developed for LFG generation assessment.
- 6
- One main outcome is that experimental campaigns on site are fundamental for emission assessment.
- 7
- Two sampling systems were designed and tested and the optimal sampling procedures were defined.

8

9

1 Figure captions:

2

3 Figure 1. Static hood: picture of the static hood used for static sampling; design based on the ones  
4 proposed by the UK EA and by Rachor.

5 Figure 2. Flux chamber: picture of the flux chamber used for fluxed sampling; design based on the  
6 US EPA guidelines.

7 Figure 3. Odour dispersion modelling results A: Odour impacts predicted with methods 1 (left) and  
8 2 (right).

9 Figure 4. Odour dispersion modelling results B: Odour impacts predicted with methods 3 (left) and  
10 4 (right).

11 Figure 5. Odour dispersion modelling results C: Odour impacts predicted with methods 5 (left), 6  
12 (centre) and 7 (right).

13