

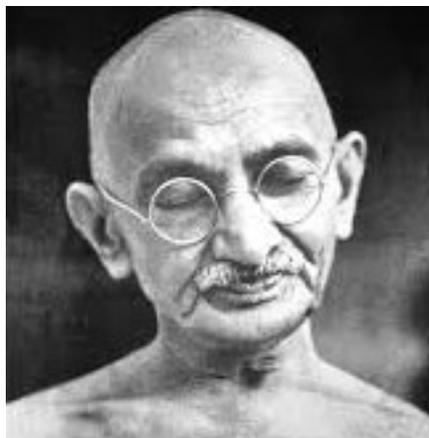
INVERSE DISPERSION MODELLING FOR A QUICK SCAN SERVICE TO ASSESS FUGITIVE EMISSIONS FROM LANDFILLS

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An ounce of
practice is worth
more than tons
of preaching.

—Mahatma Gandhi

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Context

- CH₄ produced by landfills
- GHG MRV and mitigation

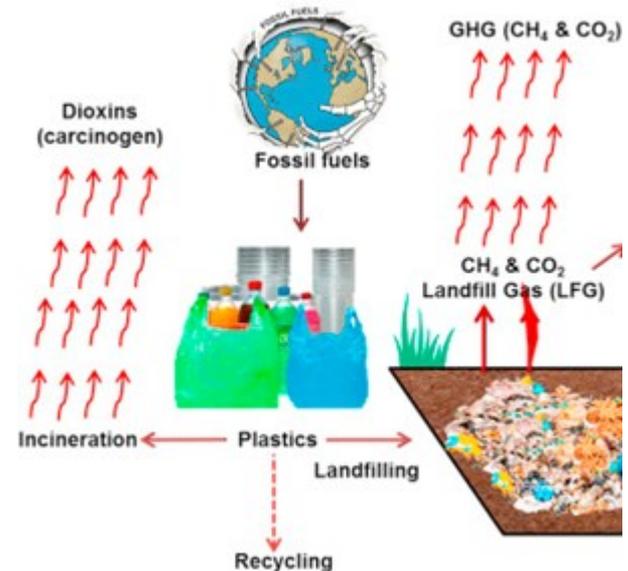
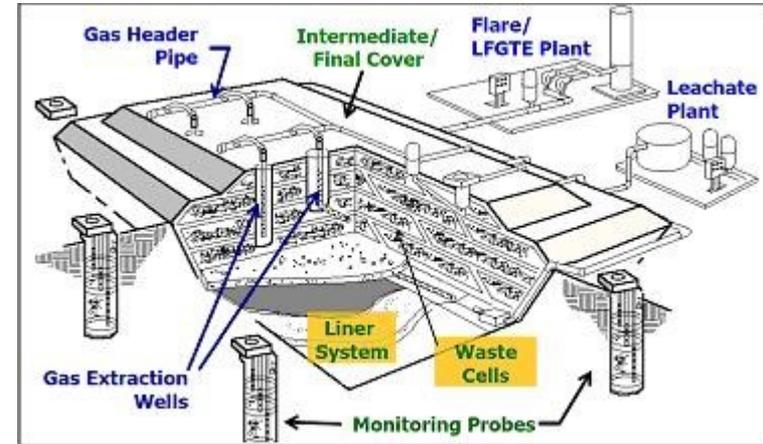
Inverse modeling approach

- Experimental site
- Measurement protocol
- Multi-sources back-plume approach
- AMIS adaptive Bayesian approach

Dual tracer method

- Methodology
- Measurement protocol
- Results

Conclusion



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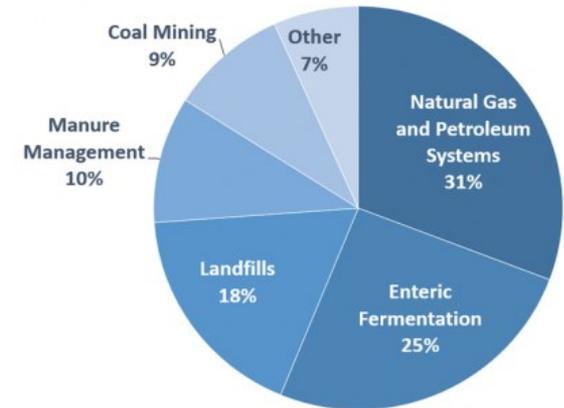
Conclusion

1 kg of CH₄ has the same impact on global warming than releasing 28 kg of CO₂ on a 100 years scale (IPCC 2013).

CH₄

- 14% of the world GHG emissions
- 8.6 % of the European Union GHG emissions in 2010.
 - 595 MtCO₂eq in 1990
 - 405 MtCO₂eq. in 2010 (a reduction of more than 30%).
 - 49% Agriculture & 32% Waste

2015 U.S. Methane Emissions, By Source



Source US-EPA Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2015.

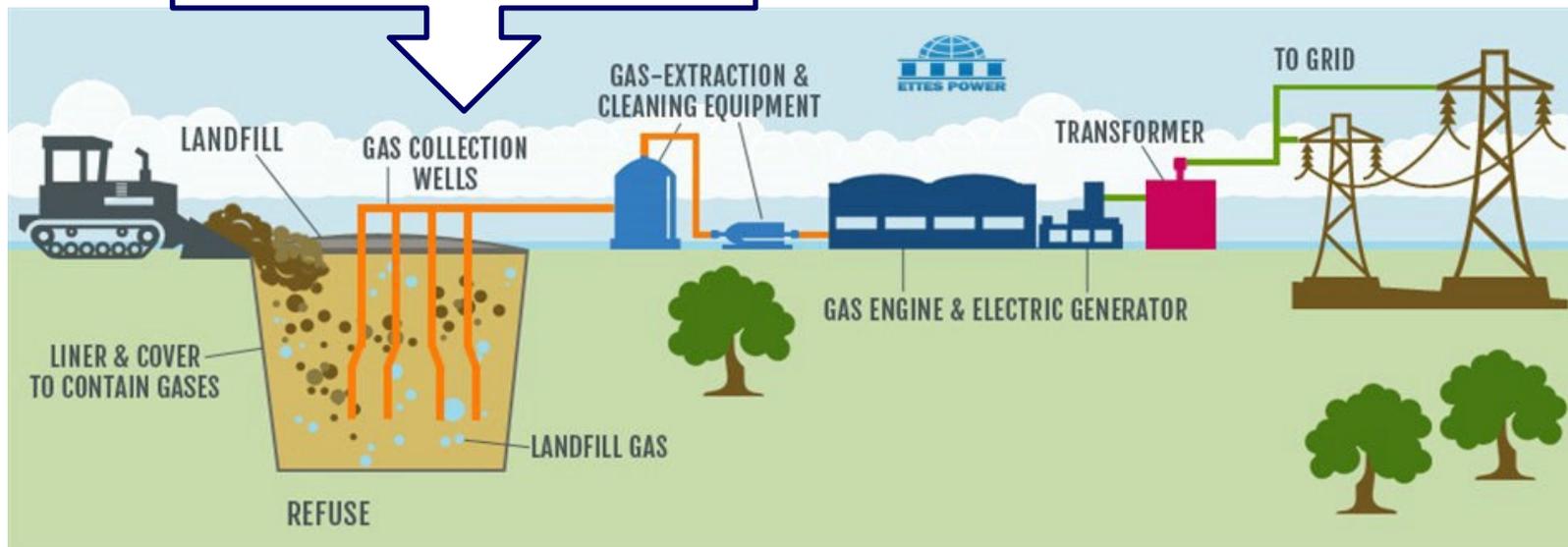
Sources : <http://www.eea.europa.eu/data-and-maps/data/data-viewers/greenhouse-gases-viewer>

Landfills : >15% of the total emission of CH₄

Need of a routine “quick scan” service

- **GHG MRV**
- **Checking collect and countermeasure efficiency**
- **Optimizing the biogas collect and its valorisation**

- **METHANE: ~50%**
- **CO₂ : 35%**
- **Other gases : 15%**



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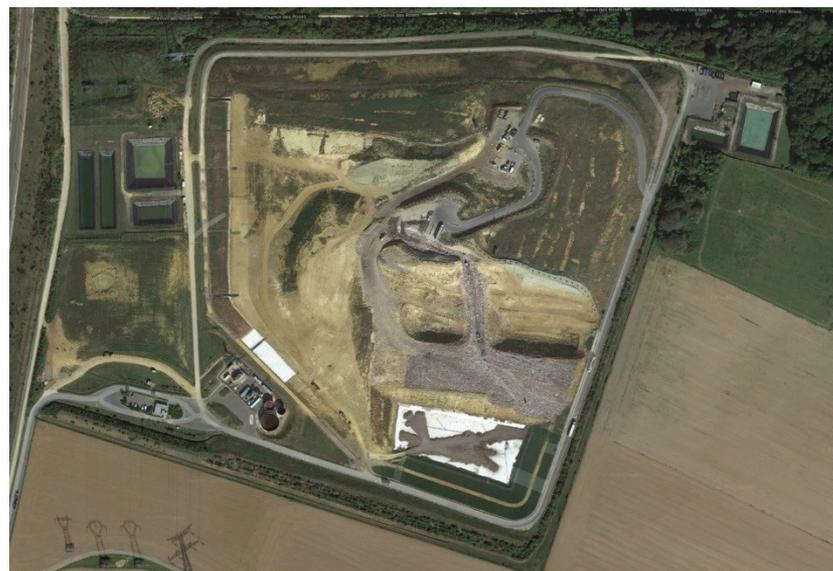
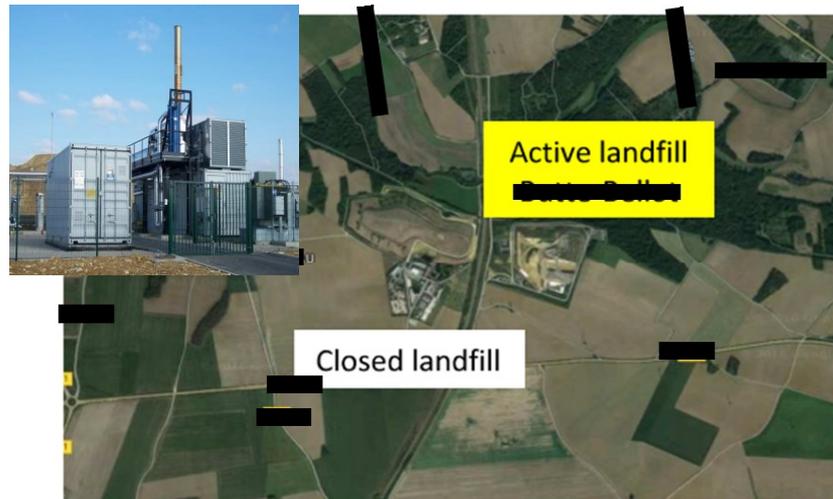
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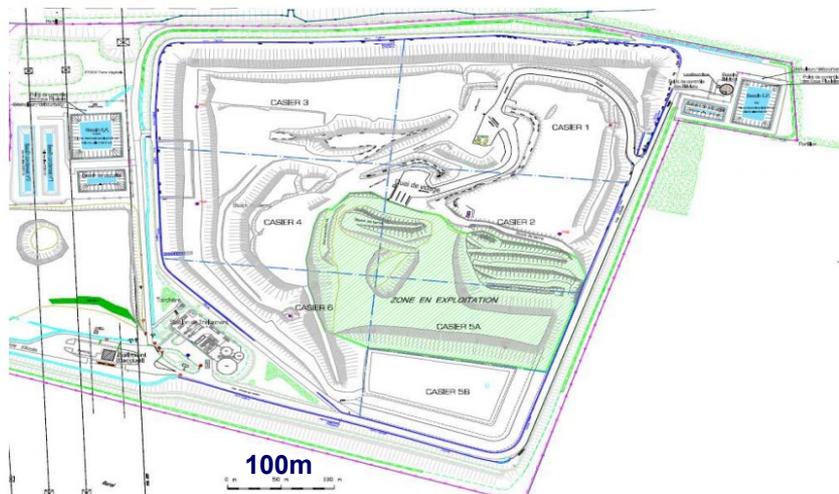
Domestic /urban waste landfill

Rural site - Flat natural terrain

- Old one closed (1974- 2004)
- Active one from 2005
 - ~ 250 000 t / year
 - 30 ha x H=20m ; 2.8 M m³ /3.1 M t
- Biogas 4.5 M Nm³ collected
- 9.5 Gwh_e ~900 toe (tep) saved



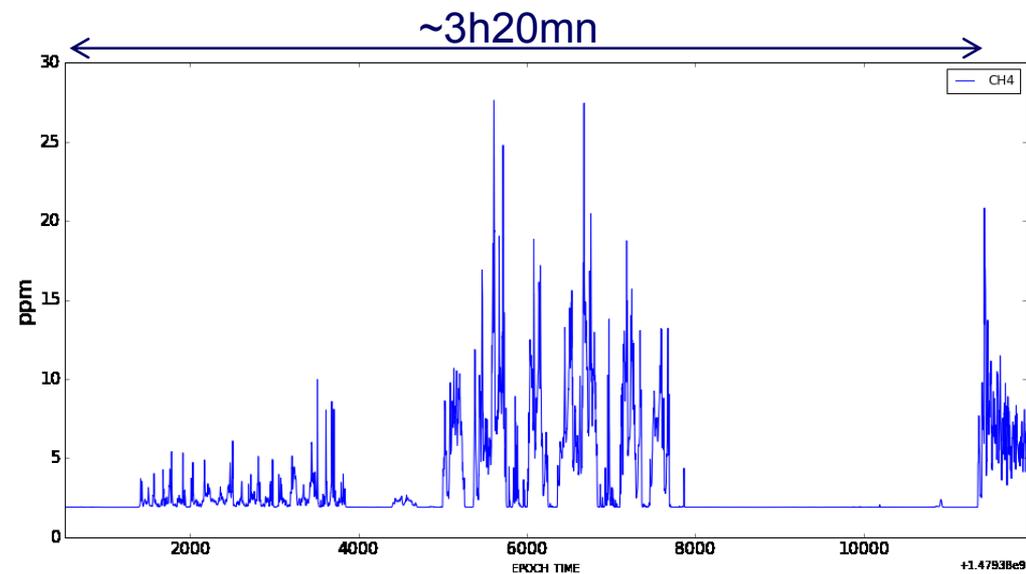
Présentation de l'Installation



CH₄ measurement (dt = 1s)

- Picarro analyser (G2203)
- Cavity Ring Down Spectroscopy
- PC +GPS
- Power supply

Meteorological ground station



Base on linearity between sources and receptors and steady state assumption : n area sources, m receptors

$$\begin{bmatrix} A_{11} & \dots & A_{1n} \\ \vdots & \ddots & \vdots \\ A_{m1} & \dots & A_{mn} \end{bmatrix} \begin{bmatrix} Q_1 \\ \vdots \\ Q_n \end{bmatrix} + \begin{bmatrix} B_1 \\ \vdots \\ B_m \end{bmatrix} = \begin{bmatrix} C_1 \\ \vdots \\ C_m \end{bmatrix}$$

- A_{ij} is the atmospheric transfer coefficient from source i to measurement j
- Q_j is the emission rate of the source j from 1 to n
- C_i is the measured concentration at receptor i from 1 to m
- B_i is the background at receptor i from 1 to m =>

Finding Q_j required $[A_{ij}]$ (pseudo)inversion

- Quick means n and m not to big
- $[A_{ij}]$ well conditioned

The “Condition Number” of matrix A , $\kappa(A)$, is an indicator of how the error on concentration may be amplified when source rates are obtained

$$\kappa(A) = \frac{M_{max}}{M_{min}} = \|A\|_M \cdot \|A^{-1}\|_M$$

- Considering Euclidian distance

$$\|x\|_E = \sqrt{\sum_1^N x_i^2}$$

- How much the matrix $A(n,m)$ can stretch vectors ?
 - $M_{max} = \max\left(\frac{\|Ax\|}{\|x\|}\right) = \|A\|_M$ Considering “max distance”
- how much the matrix A can shrink vectors ?
 - $M_{min} = \min\left(\frac{\|Ax\|}{\|x\|}\right) = \frac{1}{\|A^{-1}\|_M}$

We need to solve a system $AQ = C$

- C is corrected from background
- $\|C\| \leq M_{max} \|Q\|$

$A(Q + \delta Q) = C + \delta C$ or $A\delta Q = \delta C$

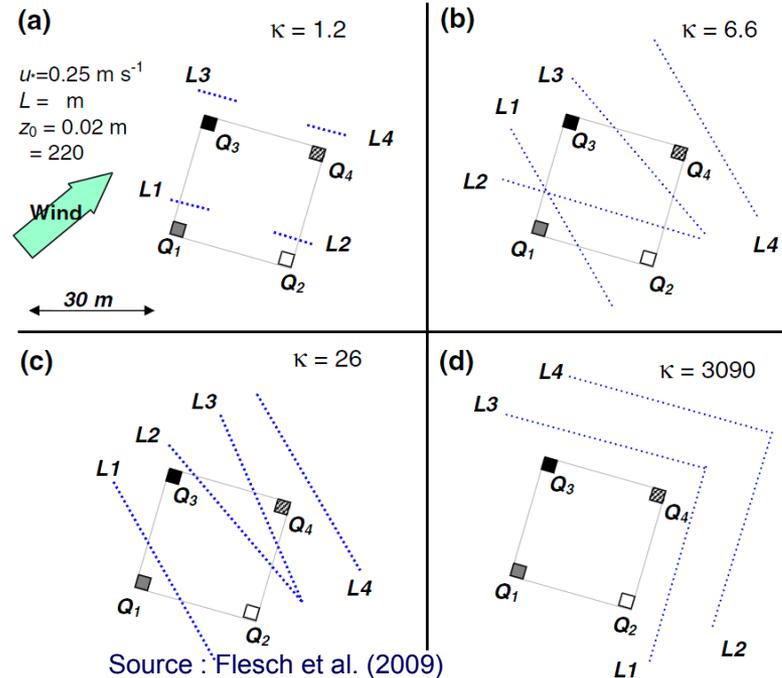
- $\|\delta C\| \geq M_{min} \|\delta Q\|$

$$\|C\| \cdot M_{min} \cdot \|\delta Q\| \leq M_{max} \|Q\| \|\delta C\|$$

$$\frac{\|\delta Q\|}{\|Q\|} \leq \kappa(A) \left(\frac{\|\delta C\|}{\|C\|} \right)$$

If κ is greater than ~ 20

→ We cannot separate the different sources

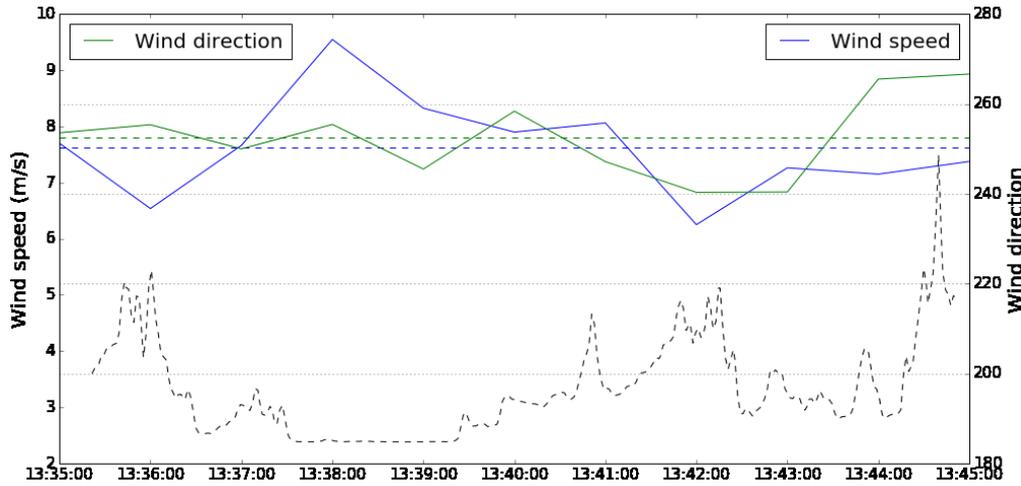


Optimisation of pseudo-sensors (matrix C)

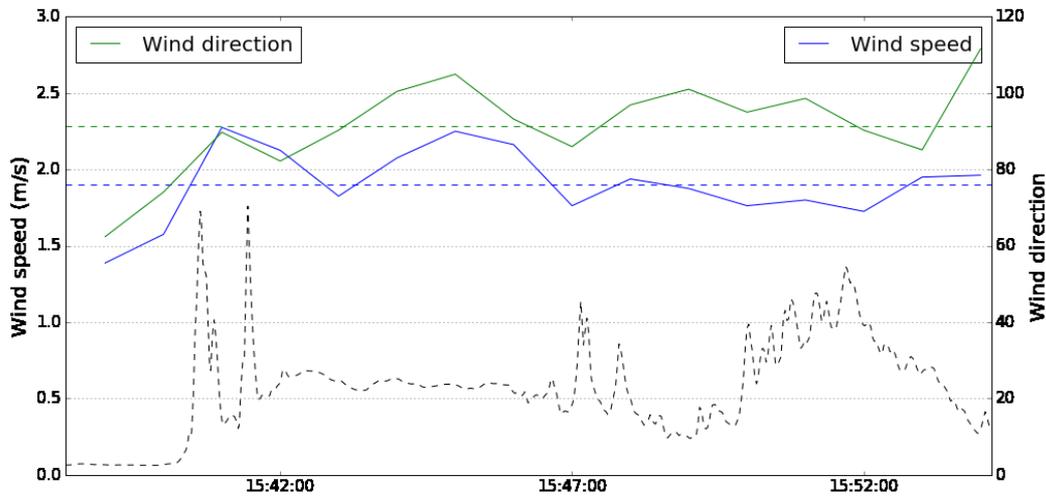
HARMO18



17 November 2016

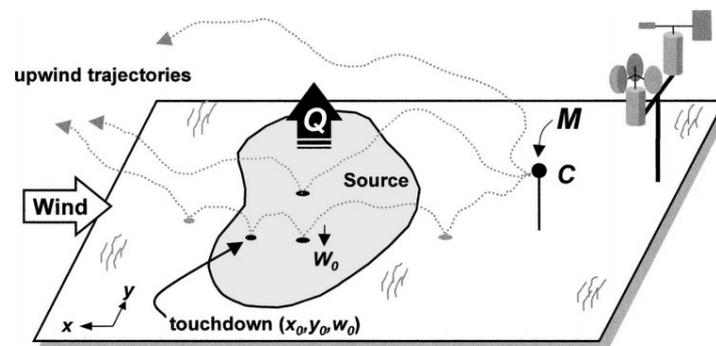
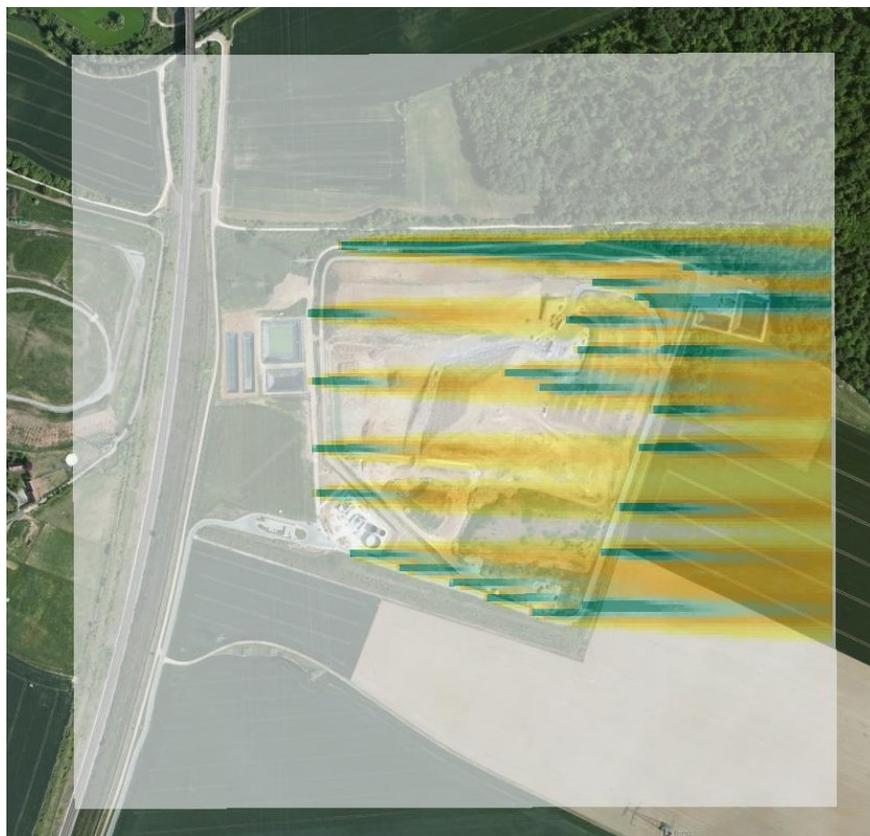


5 December 2016



Atmospheric transfert coef. (matrix A)

Matrix A is computed using back-plume stochastic lagrangian model (MSS in ARIA View Package)



Horizontal grid resolution	5 meters
Domain dimension	1 kilometer x 1 kilometers
Number of vertical levels	21
Lowest / Highest vertical level	3 / 600 meters
Number of receptors	27 (for the 17 th November) 29 (for the 5 th December)
Number of sources	4 (for the 17 th November) 5 (for the 5 th December)
Computation of source-receptor matrix	Each 5 minutes
Estimation of emission rates	Each 15 minutes

17 November 2017

Before mitigation actions

The global emission for the site: **3.85 t/day**



5 December 2017

After mitigation actions

The global emission for the site : **2.92 t/day**



Find the minimum of a cost function

The *Adaptive Multiple Importance Sampling* (AMIS) algorithm is based on an sampling scheme, where a target distribution (namely the posterior distribution) is approximated by weighted samples from a proposition distribution

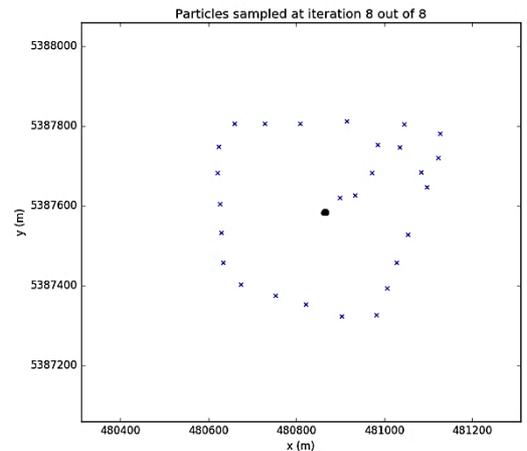
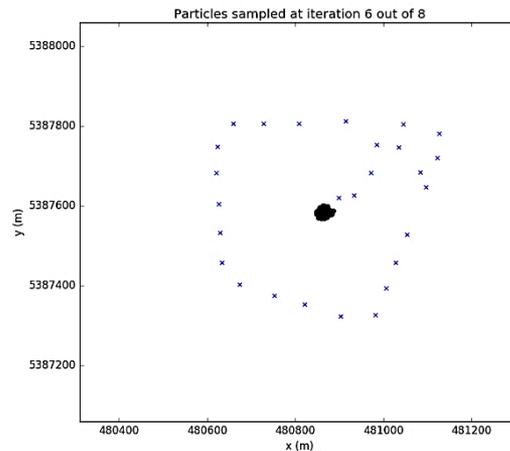
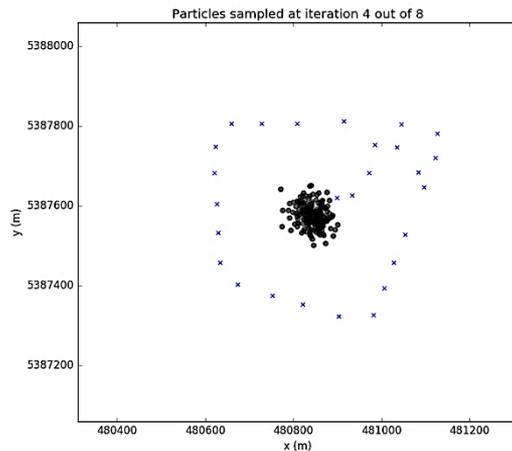
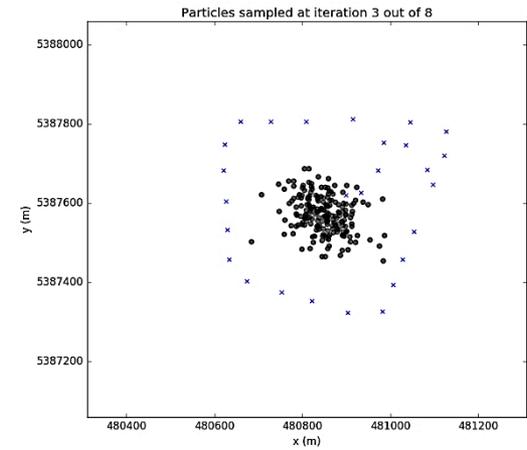
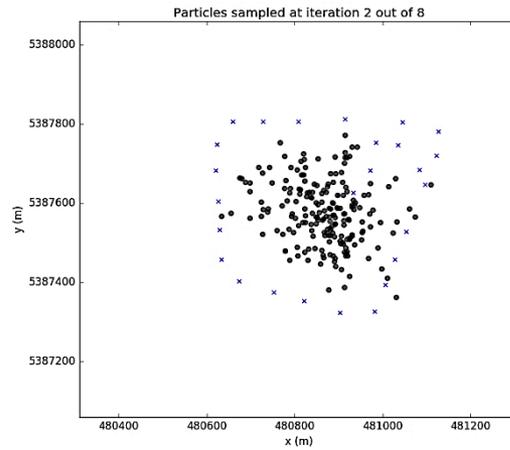
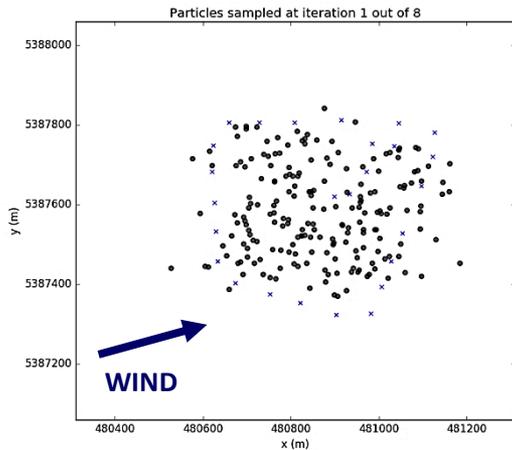
The “*Adaptive*” algorithm improves the standard importance sampling procedure by:

- Allowing the update of the proposal distribution, which can be chosen as a flexible combination of well-known kernels (e.g. a multivariate Gaussian mixture)
- Optimally recycling the importance weights at each iteration to fully exploit the full available information and accelerate the convergence

Current version of the software: a unique point source → here 4 or 5 area sources !



Test of the software “as is” on the 17 Nov 2017

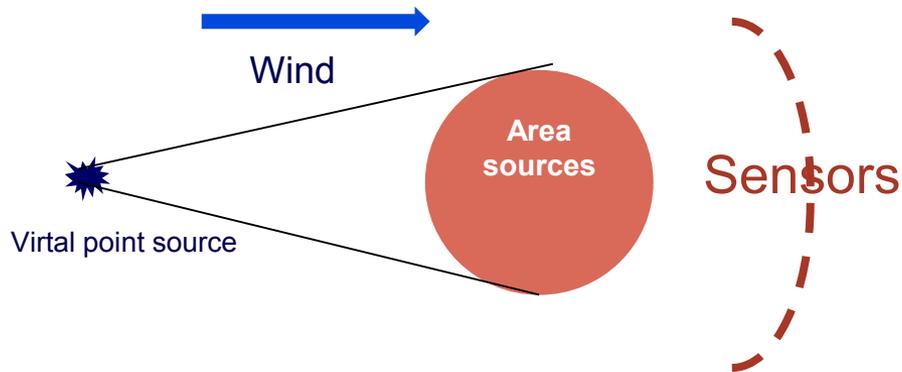


Surprise : good order of magnitude !

- Back-plume method : 3.85 tons/day
- AMIS method : 3.46 tons/day

What we should expect with a single source

- A virtual point source upwind
- If in the area source → a large underestimation



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Experimental set

- Dual gas is acetylene C_2H_2
- A C_2H_2 bottle and suitable flowmeter
- Same mobile C_2H_2/CH_4 PICARRO (G2203) installed in car
- Additional measurements with G2201Piacarro CDRS (carbon isotopes apportionment)



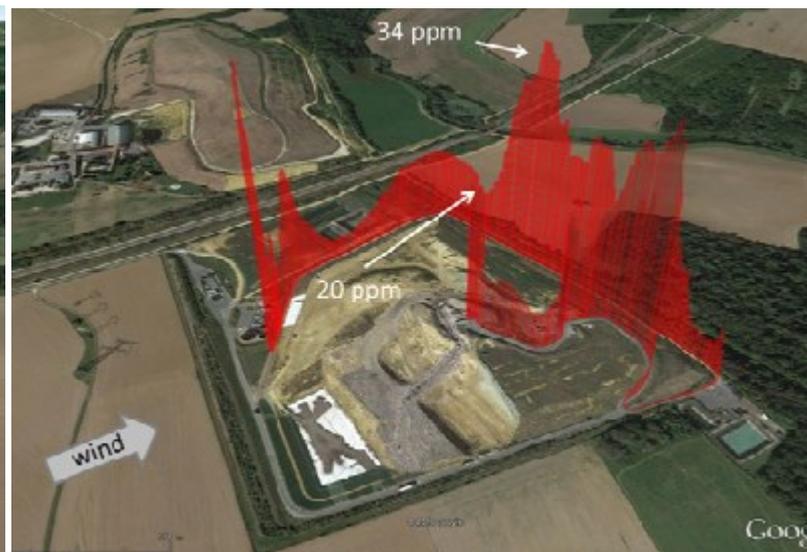
LSCE

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& DE L'ENVIRONNEMENT

Principle

- Release a known gas (here C_2H_2) in the same place of the unknown emission (here CH_4)
- $$Q_{CH_4} = Q_{C_2H_2} \frac{A_{CH_4} M_{CH_4}}{A_{C_2H_2} M_{C_2H_2}}$$
 - Q_{CH_4} targeted emission
 - $Q_{C_2H_2}$ is the know release of acetylene (flowmeter + the bottle is weighted before and after the release (kg/h))
 - $\frac{A_{CH_4}}{A_{C_2H_2}}$ ratio of area peaks for the two gases (Picarro measurements)
 - $\frac{M_{CH_4}}{M_{C_2H_2}}$ ratio of molar mass

13 Sept 2016 Preliminary survey (before any mitigation actions)



- Identifying pics and order of magnitude
- Optimization of the C₂H₂ release
- Optimization of ambient concentration measurement protocol

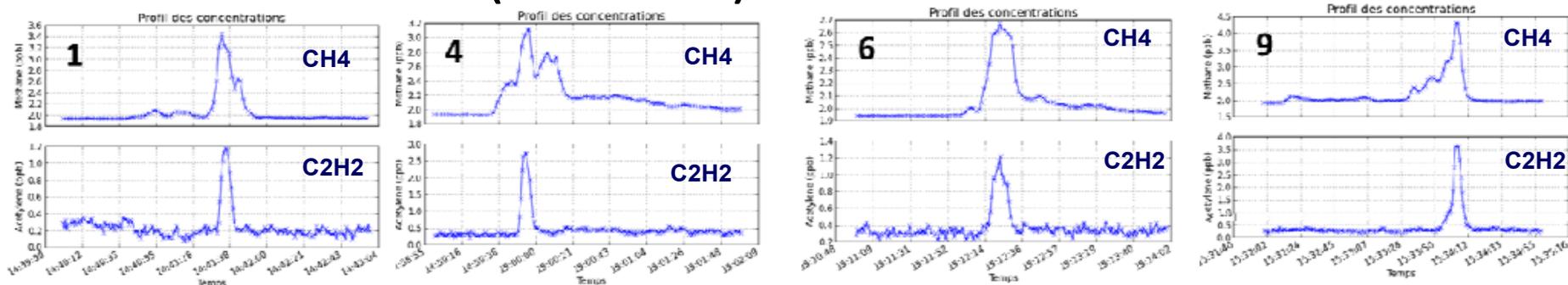
Variable (unit)	Typical value (during measurement)	Range of value (min/max)
Temperature (°C)	28	23/31
Air Pressure (hPa)	1011	1009/1013
Wind speed (m/s)	5	2.7/12.1
Wind direction (degrees) (coming from the direction)	SE	ESE/SSE
Rain (mm/3h)	0	0/1.4



19 Sept 2016 Dual gas tracer release

Variable (unit)	Typical value (during measurement)	Range of value (min/max)
Temperature (°C)	20	15/20
Air Pressure (hPa)	1018	1018/1020
Wind speed (m/s)	5	3.1/6.7
Wind direction (degrees) (coming from the direction)	NNW	NNW/WNW
Rain (mm/3h)	0.2	0/0.3

Time series downwind (14 transects)



Results

N° Pic	1	2	3	4	5	6	7	8	9	10	11	12	13	14
Emissions de CH ₄ (kg/j)	5533	6005	7823	6925	4375	5177	3314	5278	6688	5697	4528	3485	6920	7497
Moyenne (kg/j)	5813 ± 1318													

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Application to a mitigation policy

	Units	Reverse Modelling (1)	Dual Tracer (2)	((1)- (2)) / (2)
Before Mitigation (dual tracer, no reverse modelling)	t CH ₄ emitted/d	5.8	5.8 ± 1.3	
After phase I (North west of site : biogas network improvement)	t CH ₄ emitted/d	3.85	3.6 ± 1,3	- 6,5%
After phase II (cover and biogas network improvement)	t CH ₄ emitted/d	2.92	2 ± 0,5	-31%

On progress

- **In 2017: installation of an additional cogeneration engine corresponding to the amount of CH₄ recovered and it is working!**
- **Test on an other site in Netherland (with TNO)**

To HARMO18 Organizers !

For your attention !

For Partners of the WASTE MITI 2 consortium



For Climate KIC Contribution

