

IRSN

INSTITUT
DE RADIOPROTECTION
ET DE SÛRETÉ NUCLÉAIRE

Faire avancer la sûreté nucléaire

Source term assessment of a nuclear release: Inverse modelling method and application examples

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IRSN is the French technical safety organization

Role of IRSN in case of a Radiological Emergency

- Assess risk induced by accidental situation
- Provide technical expertise to public Authorities

Task

- Evaluation of the reactors state, releases to the environment (diagnosis/prognosis)
- Evaluation of the radiological consequences (doses and depositions, diagnosis/prognosis), all spatial scales

Development of operational tools

Previous events

- ❑ Main nuclear accidents: Chernobyl, Fukushima
 - ❑ Minor events: radionuclide detection by monitoring system (iodine detection 2011-2012, cesium detection 2013, forest fires in areas contaminated by the Chernobyl accident...)
- **Need to develop a tool to assess atmospheric releases by using environmental observations**
- ❑ Source location
 - ❑ Source term (ST: temporal evolution of the release rate + distribution between radionuclides)

Operational method - source term assessment

ATMOSPHERIC TRANSPORT MODEL

Gaussian puff – Eulerian - Lagrangian

+

OBSERVATIONS

Air concentration

Ambient gamma dose rate

STEP1 : DEFINE THE A PRIORI INFORMATION

Source location - First guess

Gathered measurements (spatial distribution, temporal resolution, order of magnitude)

Isotopic composition

→ Reduce the number of parameters + limit the solution space

IF AIR CONCENTRATION OBSERVATIONS

IF DOSE RATE OBSERVATIONS

STEP2 : IDENTIFY THE POTENTIAL RELEASE PERIODS

Extraction of the plume component

Inverse modelling

→ Reduce the number of parameters + limit the solution space

STEP3 : ESTIMATE THE RELEASE RATES

carry out for the all period

carry out for the periods identified during Step 2

Cost function $J(\sigma) = \|\mu - H\sigma\|^2 + \lambda \|\sigma - \sigma_b\|^2$

Cost function $J(\sigma) = \|\mu - H\sigma\|^2 + \lambda \|\sigma - \sigma_b\|^2 + \sum_{i=1}^{n_{isotope}-1} r_i^2$

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Iodine detection by air sampling monitoring stations in Europe Oct-Nov 2011

□ Description of the event

- 9/11/2011 Iodine detection on several stations in Europe (7 countries) - Source location and

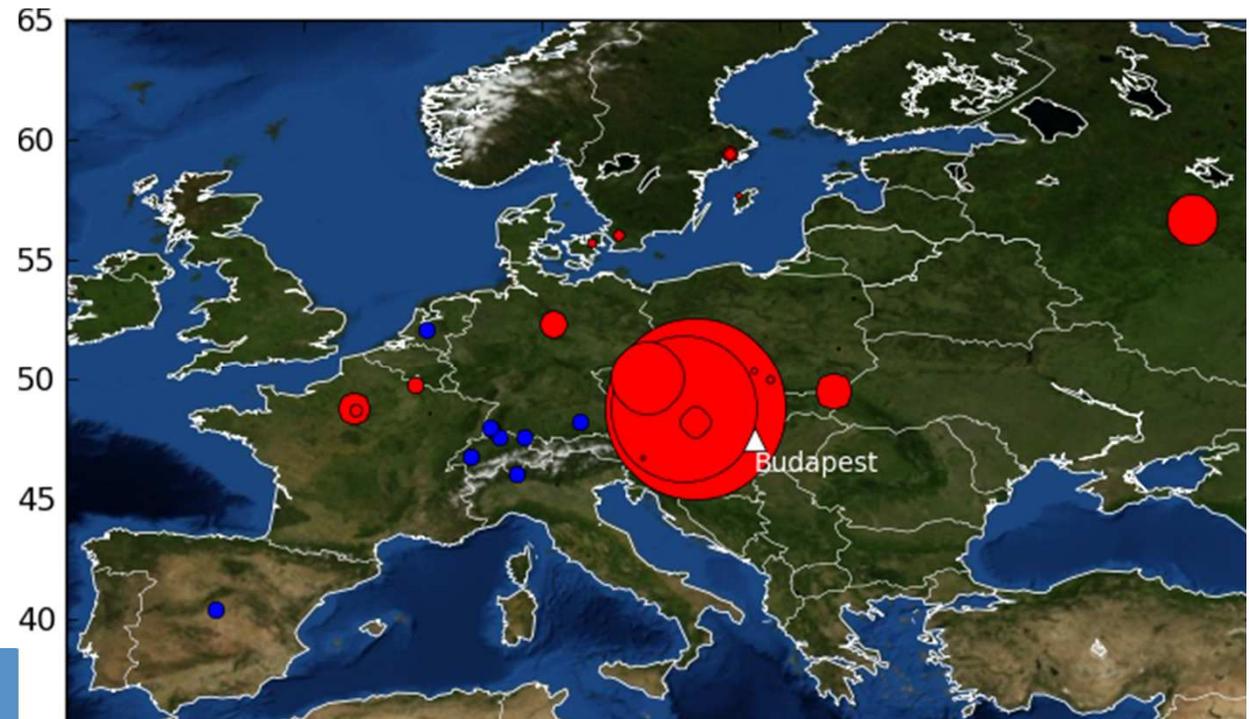
Use of the IRSN's inverse modelling tool

- Source location: to narrow the list of potential candidates
- Source term: to assess the source term

- 17/11/2011 IAEA's confirmation of the source location + provision of a source term assessment – iodine measured in 9 countries in Europe)

□ Measurements

Averaged over 1 – 7 days periods	
Czech Republic	27 $\mu\text{Bq}/\text{m}^3$
E of Austria	65 $\mu\text{Bq}/\text{m}^3$
E of Germany	14 $\mu\text{Bq}/\text{m}^3$
Poland	13 $\mu\text{Bq}/\text{m}^3$
Slovakia	16 $\mu\text{Bq}/\text{m}^3$
Sweden	5 $\mu\text{Bq}/\text{m}^3$
Hungary	87 $\mu\text{Bq}/\text{m}^3$
Ukraine	7 $\mu\text{Bq}/\text{m}^3$
N-NE of France	5 $\mu\text{Bq}/\text{m}^3$



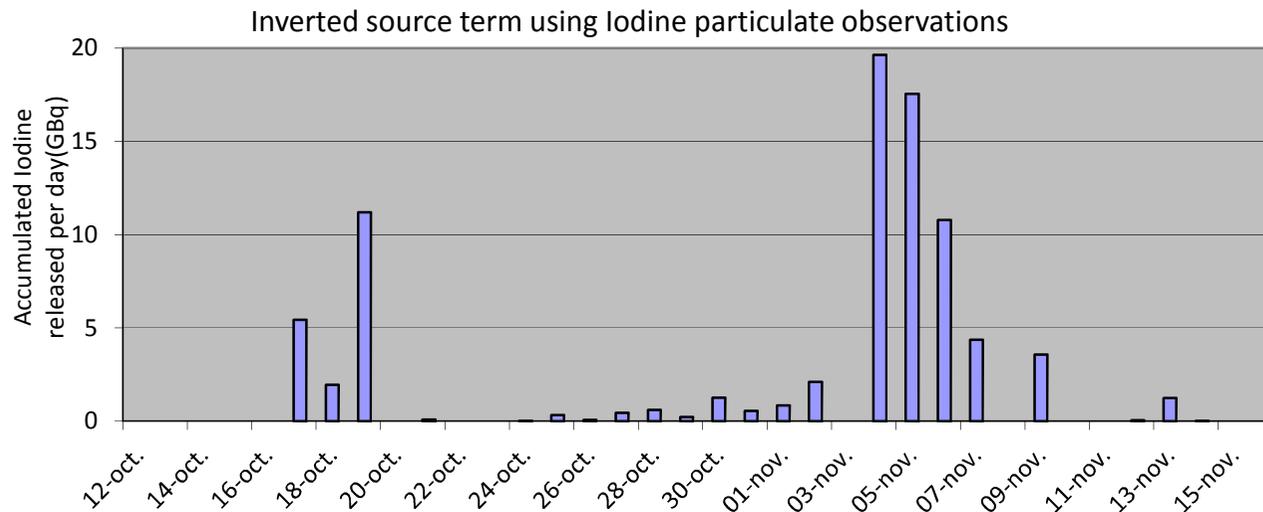
1- Application to air concentration observations

ATM	IdX (C3X platform)	Eulerian model	No chemistry of iodine
Meteorological data	ECMWF	Spatial resolution ~10 km	
Air concentration observations	41 Stations	59 Observations (>0: 17-oct – 10-nov 2011)	Time Averaged (1 to 7 days)

Step 1: Define the a priori information

- A priori source term unknown ($\sigma_b = 0$)
- “Shorten” the observation period (Consider only periods when iodine may have been observed in the stations location)
- Time scale of the inverted source term : daily

Step 3: Source term assessment

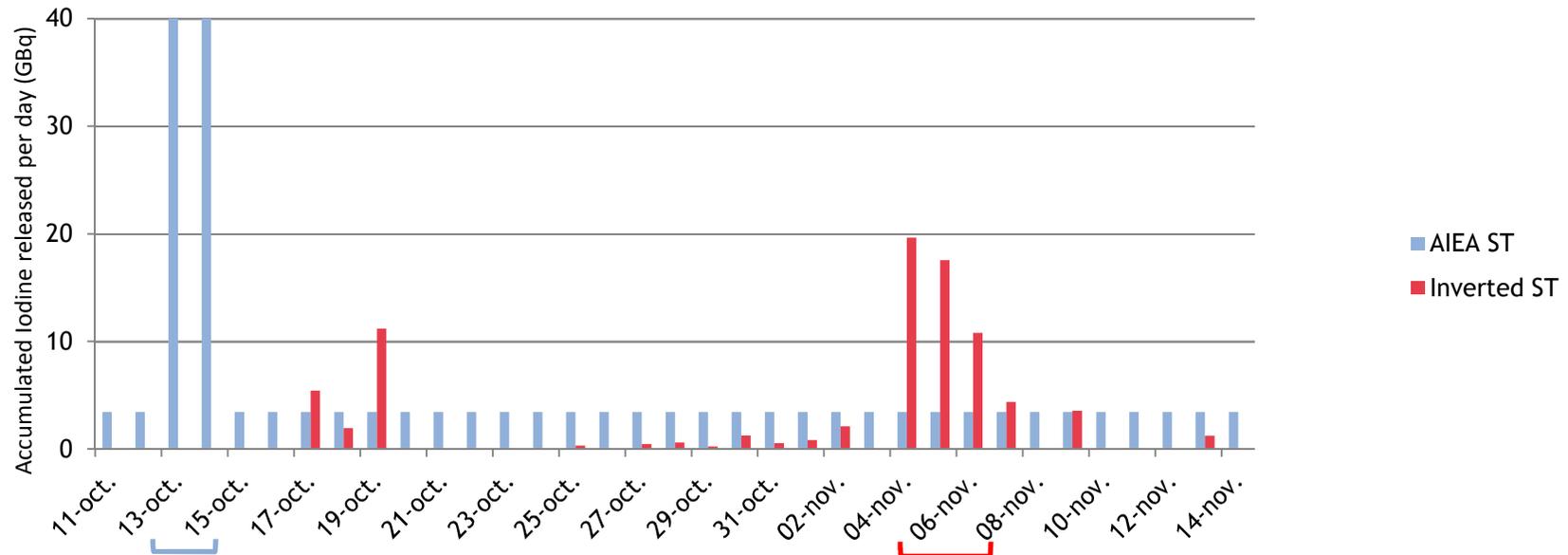


1- Application to air concentration observations

Validation of the source term: Comparison to other source term (IAEA source term)

IAEA ST:

Between 8/09/2011 – 16/11/2011 : 342 GBq released
with a maximum rate of 108 GBq/48 hours in October 12-14



Releases transported toward the south then west – they are not observed by the available monitoring stations.
→ Releases not observed: the Inversion method cannot reconstruct those events

Releases responsible for the iodine observed in France and Scandinavia. They explain the maximum concentration values observed in Austria

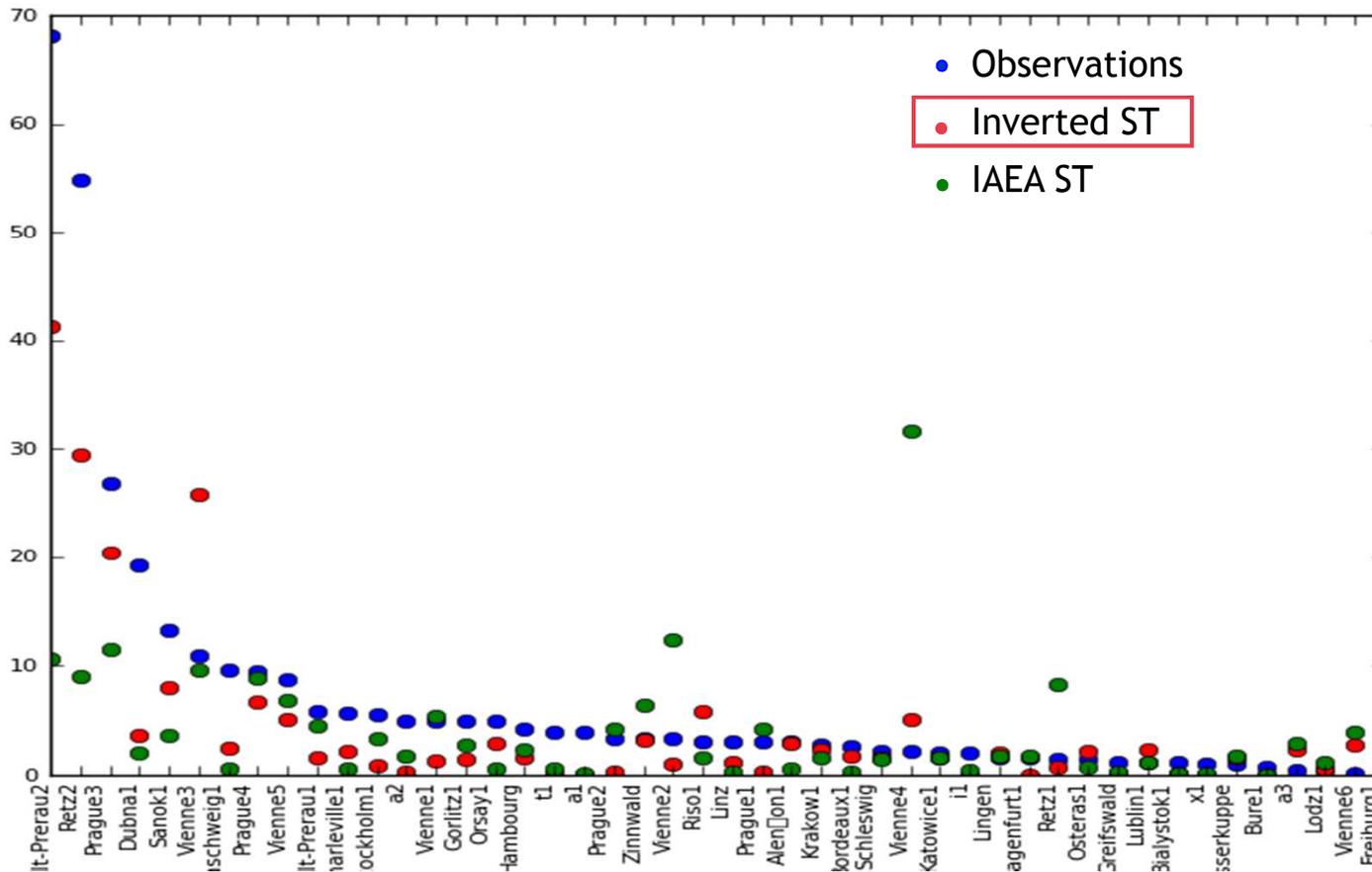
No inconsistencies between the 2 ST.

Release rates were not constant

1- Application to air concentration observations

Validation of the source term: model to data comparisons

Direct simulation using the inverted source term and comparisons with observations

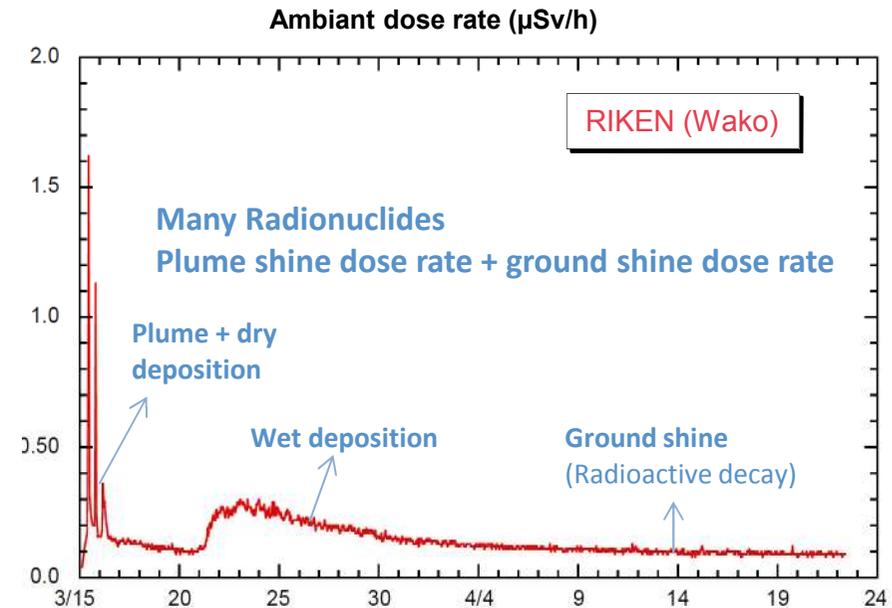
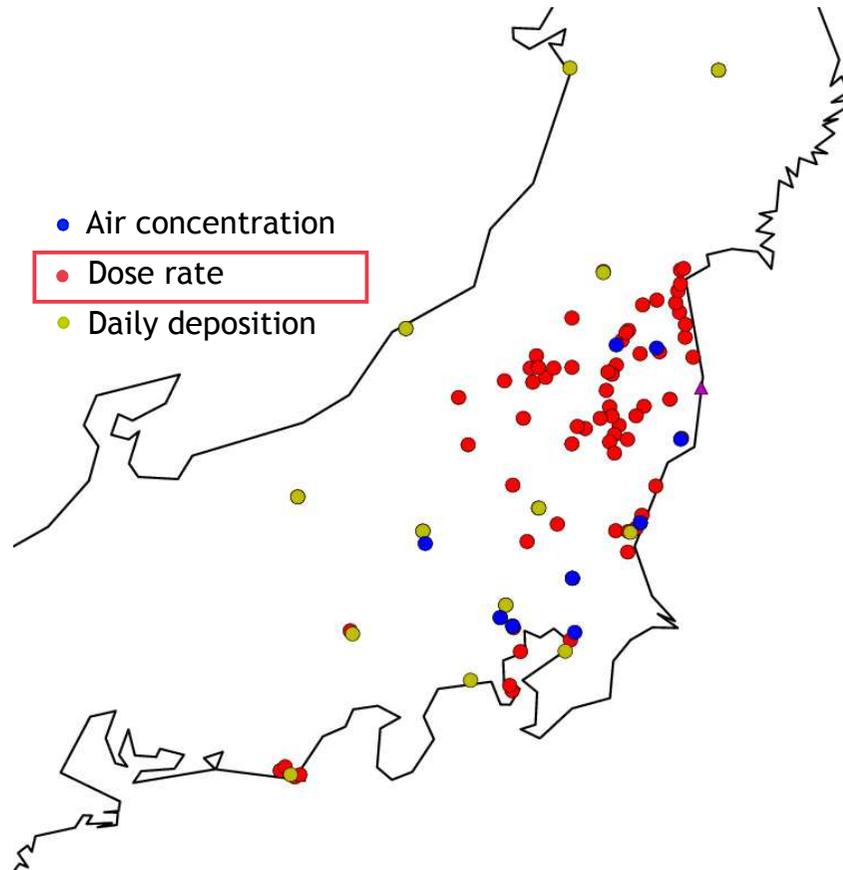


Better model to data agreement with the inverted ST.

Relatively good agreement between model and observations: inverted source term is realistic

Fukushima accident - *Saunier et al., ACP 2013*

ATM	IdX (C3X platform) Eulerian	No chemistry of iodine + Radioactive decay and filiation	Dry deposition $v_{\text{dep}} = 2 \cdot 10^{-3} \text{ cm/s}$	Wet deposition $\Lambda_s = \alpha p_o$ $\alpha = 5 \cdot 10^{-5}$	Vertical diffusion Troen and Mahrt scheme
Met. Data	ECMWF	Spatial resolution 0.125°	Temporal resolution 1h	11/03/2011 9h - 27/03/2011 18h	
Observations	Dose rate	Temporal frequency : 10min – 1h		57 Stations	



Difficult to use (Not a model parameter & Aggregate all the gamma radiations emitted by any radionuclides present on the ground and in the air)

Step 1: Define the a priori information

□ A priori source term unknown ($\sigma_b = 0$)

□ Time scale of the inverted source term : 1h

□ Isotopic composition of the ST :

The dose rate signal is mainly due to 8 radionuclides: ^{134}Cs , ^{136}Cs , ^{137}Cs , $^{137\text{m}}\text{Ba}$, ^{132}Te , ^{132}I , ^{131}I , ^{133}Xe with secular equilibrium and constant ratio ($^{134}\text{Cs}/^{137}\text{Cs}$) hypothesis, it leads to 5 radionuclides : ^{134}Cs , ^{136}Cs , ^{132}Te , ^{131}I and ^{133}Xe

□ Define isotopic soft constraints

Radionuclides released in proportions that depends on their physicochemical properties + the core inventory

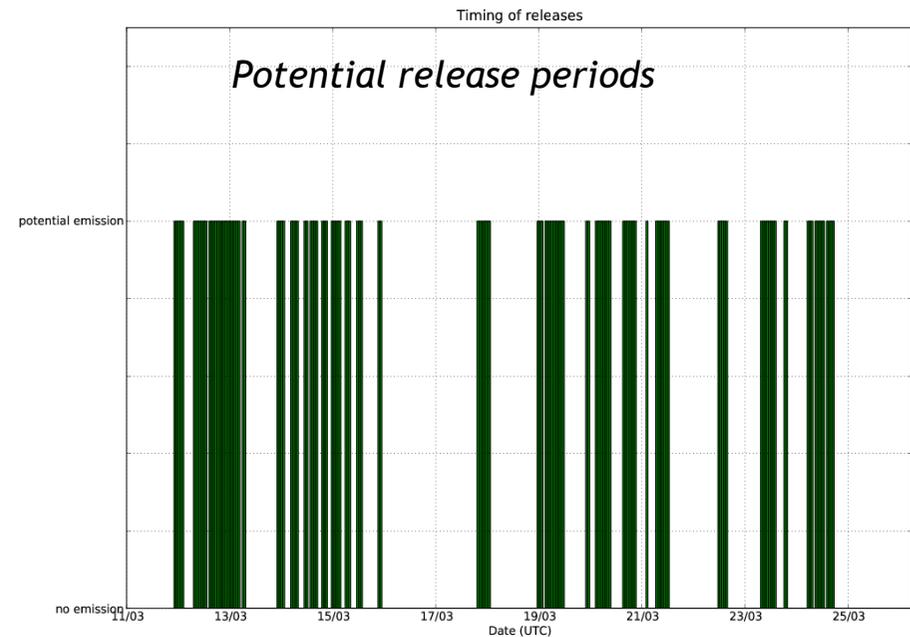
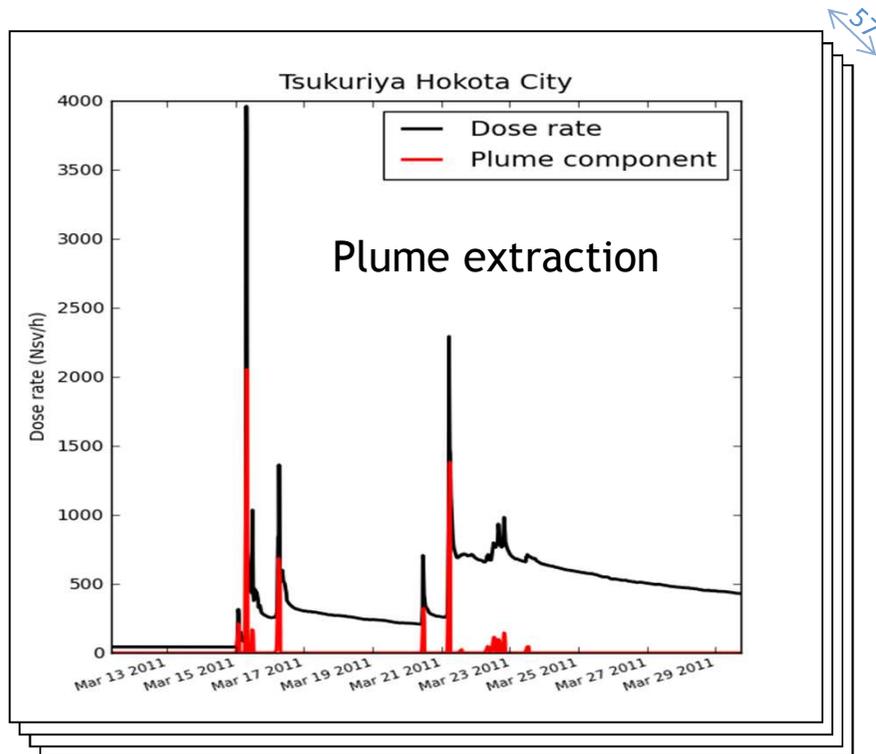
$$0.6 \leq \frac{\sigma_{^{132}\text{Te}}}{\sigma_{^{134}\text{Cs}}} \leq 16$$

$$2 \leq \frac{\sigma_{^{131}\text{I}}}{\sigma_{^{134}\text{Cs}}} \leq 100$$

$$0.1 \leq \frac{\sigma_{^{133}\text{Xe}}}{\sigma_{^{134}\text{Cs}}} \leq 10000$$

$$0.1 \leq \frac{\sigma_{^{136}\text{Cs}}}{\sigma_{^{134}\text{Cs}}} \leq 0.5$$

Step 2: Identify the potential release periods

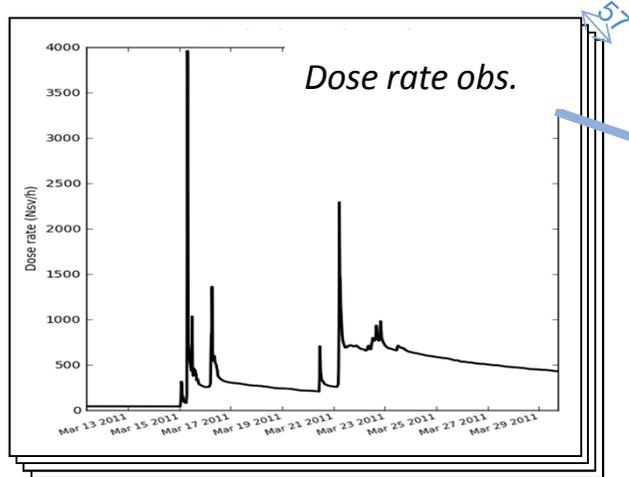


→
Inverse modelling

Cost function

$$J(\sigma) = \|\mu - H\sigma\|^2 + \lambda \|\sigma - \sigma_b\|^2$$

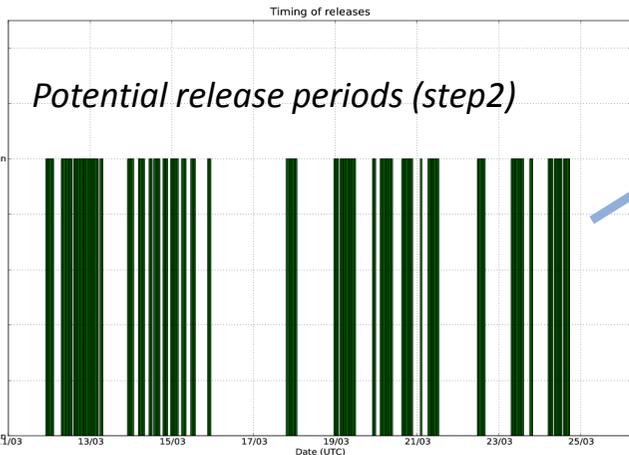
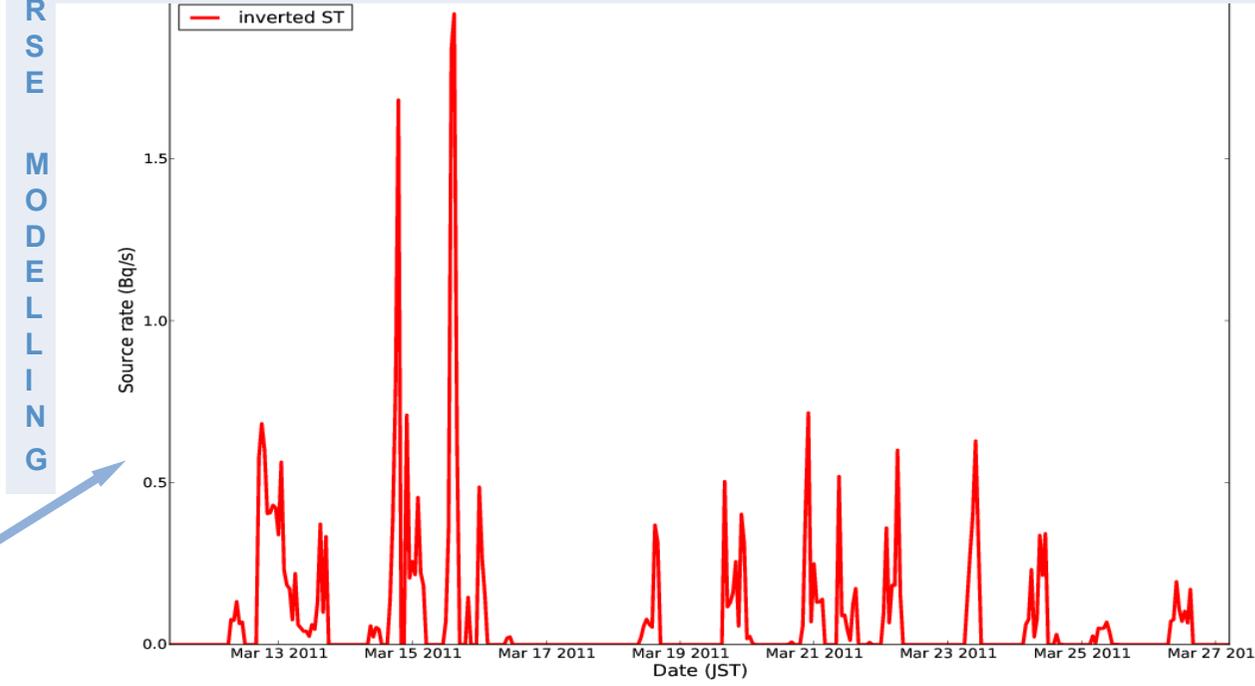
Step 3: Estimate the release rates



INVERSE MODELLING

Cost function

$$J(\sigma) = \|\mu - H\sigma\|^2 + \lambda \|\sigma - \sigma_b\|^2 + \sum_{i=1}^{n_{isotope}-1} r_i^2$$



Validation - Comparisons to other ST

Source Term (PBq)	¹³³ Xe	¹³¹ I	¹³² I	¹³⁷ Cs	¹³⁶ Cs
Inverted ST	12100	103	35.5	15.5	3.7
Mathieu et al. (2012)	5950	197	56.4	20.6	9.8
Winiarek et al. (2012)	-	190-380	-	12-19	-
Terada et al. (2012)	-	150	-	13	-
Stohl et al. (2012a)	13400-20000	-	-	23.3-50.1	-
TEPCO (2012)	500	500	-	10	-

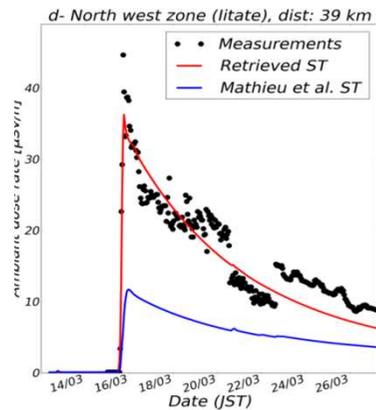
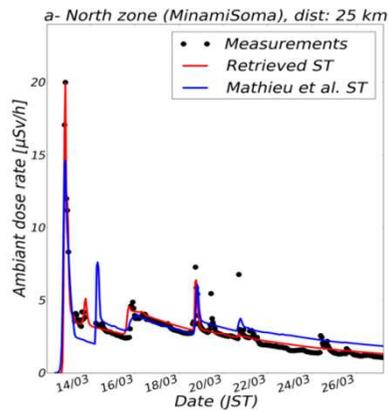
Inverted source term is consistent with the other estimations.

Underestimation in iodine and cesium in comparison to Mathieu et al ST (several events are not identified by inversion).

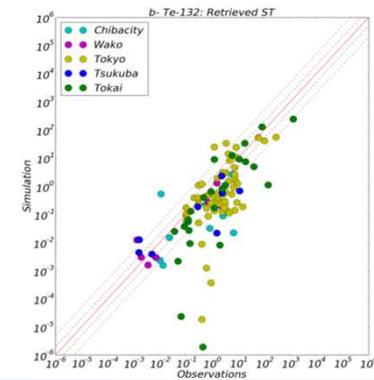
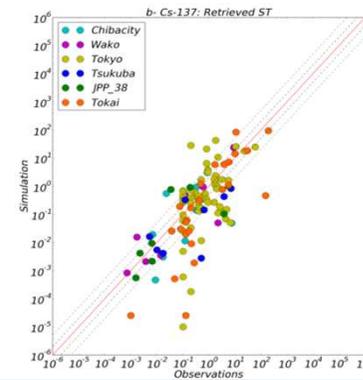
Amount of noble gases is similar to Stohl et al estimation: probably overestimated

Validation – Model to data comparisons

Dose Rate	Inverted ST	Mathieu et al. ST
Fac2 (%)	79.8	60.0
Bias	0.42	0.59



Air Concentration	Inverted Fac5 (%)	Mathieu Fac5 (%)
^{136}Cs	52,3	35,4
^{137}Cs	58,2	47,0
^{131}I	57,1	31,4
^{132}Te	53,7	40,1



Inverted quantities are consistent with the other estimations.

Model-to-data comparisons show a relatively good agreement.

Assessment of the isotopic composition need to be improved.

Releases transported toward the ocean (and not detected by the dose rate network) are not reconstructed).

Model uncertainties (Met data, deposition process...).

Reliable inversed modelling method to assess the source term

- ❑ Efficient operational tool, perfectly suited to crisis management (Air concentration or Gamma dose rate measurements).
- ❑ Can be used with various ATM (Eulerian, Gaussian, Lagrangian).

What are the next steps ?

- ❑ Simultaneous reconstruction of release events detected close to the source location as well as those detected far away.
- ❑ Improve the reconstruction of the isotopic composition by using all together air concentration, deposition and dose rate observations.

What are the challenges ?

- ❑ How can we take observations with greatly diverging orders of magnitude into account?
Inverse modelling tool (cost function) is designed to strengthen the higher values.
Application to the Fukushima case: assess the releases transported toward the Pacific ocean as well as those detected on Japan territory.
- ❑ How can we use all together different kind of observations?
Application to the Fukushima case: better assess the isotopic composition + assess the releases transported toward the Pacific ocean

Thank you for your attention