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1.19 VALIDATION OF DRY DEPOSITION MODELS FOR SUBMICRONIC AND MICRONIC AEROSOLS

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INTRODUCTION

The atmosphere is a major transfer path of pollutants released in gaseous form or in aerosols form from an industry to the land and/or sea environment and, consequently, to humans. In order to estimate the impacts of an atmospheric release on human and on the environment, it is necessary to assess the dispersion and deposition (wet and dry) of these pollutants. Dry deposition was studied from experimental campaigns carried out *in situ* and in laboratory which allowed understanding globally the phenomena governing it. However, it still remains uncertainties as for the assessment of dry deposition velocity, in particular for submicronic and micronic aerosols. The dry deposition velocity depends on numerous factors such as micro-meteorological conditions, pollutant and substrate properties. This implies that dry deposition velocity cannot be accurately assessed without *in situ* measurements. Generally, a conservative value of $5 \cdot 10^{-3} \text{ m}\cdot\text{s}^{-1}$ is used in operational models due to a lack of specific knowledge of the site being studied.

On May 18th, 2001, the spent fuel reprocessing plant of COGEMA La Hague released into the atmosphere, Ruthenium and Rhodium-106 (¹⁰⁶Ru-Rh) aerosols which marked the near environment of the site. Following this release, grass was sampled by different laboratories (GRNC, 2002) (Crabol and Maro, 2001) and the results obtained were analyzed (Maro *et al.*, 2002).

The purpose of this document is to show the results of ¹⁰⁶Ru-Rh measurements carried out on grass, and the interpretation made regarding the operational deposition models.

EQUIPMENT AND METHOD

On May 18th, 2001, a failure in a gas-treatment line of the spent fuel reprocessing plant of COGEMA La Hague resulted in a ¹⁰⁶Ru and ¹⁰⁶Rh release into the atmosphere. The activity released during 1 hour was estimated to $4.5 \cdot 10^9 \text{ Bq}$ (COGEMA, 2001). ¹⁰⁶Ru and ¹⁰⁶Rh are beta- and gamma-emitting radioelements. ¹⁰⁶Ru and ¹⁰⁶Rh are in radioactive equilibrium with respectively radioactive time periods of 372.6 days and 30 s. Following this release, grass was sampled by COGEMA, ACRO, OPRI and IRSN (GRNC, 2002) (Crabol and Maro, 2001) (Maro *et al.*, 2002) under wind of the facility in the wind direction 320°. Grass was sampled on both sides of the average wind direction between 700 m and 6,700 m from the discharge point (figure 1).

Grass samples were then dried to 60°C then ¹⁰⁶Ru-Rh concentrations were measured using gamma spectrometry in the Roule Mountain underground laboratory at Cherbourg (French Navy) with a very low background noise deeply improving measures reliability.

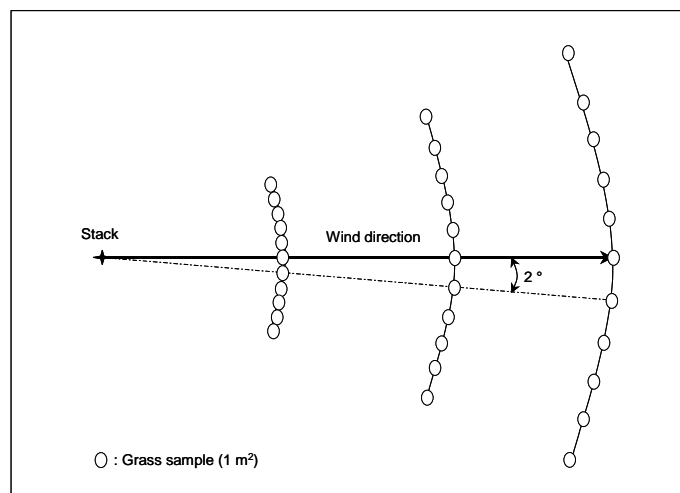


Figure 1. Position of sample points regarding the discharge point and the average wind direction

RESULTS AND DISCUSSION

Results of $^{106}\text{Ru-Rh}$ measurements in grass samples

Measurement results are displayed in figure 2 depending on the distance from the discharge point (UP2-800 stack).

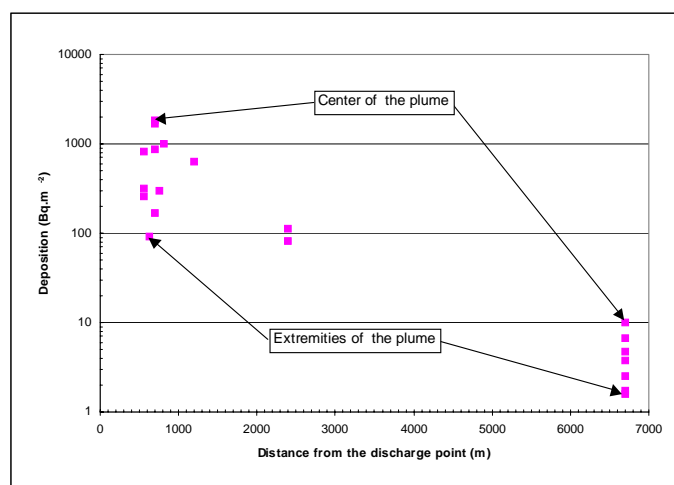


Figure 2. Evolution of $^{106}\text{Ru-Rh}$ deposition in environment depending on the distance from the discharge point

These results show a marking of the grass, up to a few thousands of Bq.m^{-2} of $^{106}\text{Ru-Rh}$, at 700 m from the discharge stack, in the wind direction. This released activity reduces by 10 (100 Bq.m^{-2}) at 2,400 m and by 100 at least (10 Bq.m^{-2}) at 6,700 m.

Comparison of measurement results with calculation results using a constant dry deposition velocity of $5 \cdot 10^{-3} \text{ m.s}^{-1}$ generally used for operational models

An atmospheric plume may progressively deplete in the wind direction, as aerosols stick to surfaces, such as leaves and soil. The released aerosol quantity may be determined from a term called dry deposition velocity. This parameter of which dimension is a velocity (m.s^{-1}), is the quotient of the dry deposition flux density on the soil ($\text{Bq.m}^{-2}.\text{s}^{-1}$) and the atmospheric concentration at the soil level (Bq.m^{-3}). The dry deposition velocity values vary with the

atmospheric stability, the wind velocity and the surface condition, but also the aerosol granulometry. The most used hypothesis for operational atmospheric dispersion models, is a constant dry deposition velocity for a given physico-chemical form. With no precise data on the physical characteristics of the released aerosols, the typical value of this deposition velocity is $5 \cdot 10^{-3} \text{ m.s}^{-1}$.

Basing on the Atmospheric Transfert Coefficients (ATC) measured at 700 m from the IRSN atmospheric dispersion study campaigns (experimental ATC) (Maro *et al.*, 1999) and the $^{106}\text{Ru-Rh}$ quantity released, assessed from $^{106}\text{Ru-Rh}$ measurements of COGEMA La Hague, the dry deposition velocity of aerosols may be estimated to $5.7 \cdot 10^{-2} \text{ m.s}^{-1}$ (Figure 3).

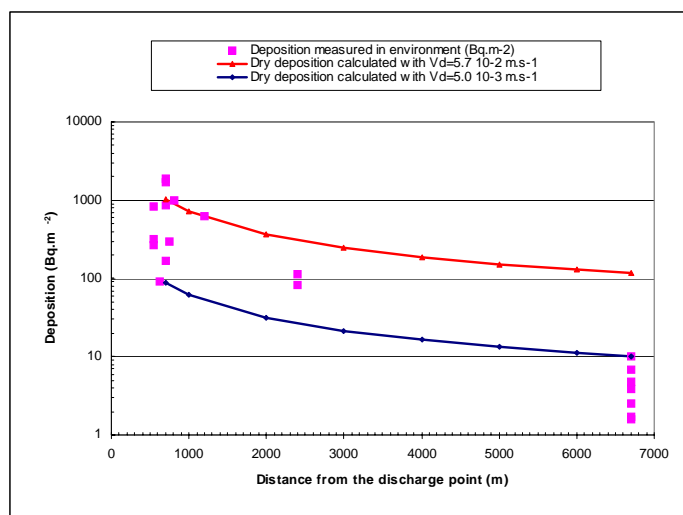


Figure 3. Comparison between $^{106}\text{Ru-Rh}$ deposition in environment with deposition calculated with dry deposition velocities of $5 \cdot 10^{-3} \text{ m.s}^{-1}$ and $5.7 \cdot 10^{-2} \text{ m.s}^{-1}$

This deposition velocity is higher (factor of 10) than the deposition velocity of $5 \cdot 10^{-3} \text{ m.s}^{-1}$, generally used for operational models. An hypothesis to explain this significant value of the deposition velocity would be the non incorporation of the electrostatic field in the deposition velocity assessment for very low granulometry aerosols (Gensdarmes, 2002). Another hypothesis is that the significant value of deposition velocity may also result from the presence of strong local turbulence which may increase the deposition velocity of fine aerosols. Besides, during this incident, it is likely that ruthenium was released in gaseous form, RuO_4 , then was progressively reduced in the form of RuO_2 aerosols. One may think that this reduction of RuO_4 into RuO_2 in the facility resulted from homogeneous nucleation which produced ultrafine aerosols (a few nanometers).

However, for this deposition velocity of $5.7 \cdot 10^{-2} \text{ m.s}^{-1}$ determined from measurements at 700 m from the discharge point, the activity settled in $^{106}\text{Ru-Rh}$ should be 120 Bq.m^{-2} at 6,700 m, whereas the activity measured in environment at this distance is about 5 Bq.m^{-2} (Figure 3). We tried to determine if this deviation between measurements and model might be explained by taking into account the coagulation phenomenon of $^{106}\text{Ru-Rh}$ aerosols on atmospheric aerosols in the atmospheric dispersion modelling.

Comparison of measurement results with calculation results taking into account the coagulation phenomenon of $^{106}\text{Ru-Rh}$ aerosols on atmospheric aerosols

In order to take into account the aerosol coagulation phenomenon (Boulaud and Renoux, 1998) in the modelling, we considered a population of atmospheric aerosol with R_1 radius comprised between 10^{-2} and $1 \mu\text{m}$ and a population of ultra-fine $^{106}\text{Ru-Rh}$ aerosols with

R_2 radius comprised between 1 and 10 nanometers. The concentration of atmospheric aerosols N_1 is considered as equal to 5,000 particles per cm^3 (measuring campaign in La Hague dated March 11th, 2002). The coagulation phenomenon of R_1 radius particles with R_2 radius particles is represented by a coagulation coefficient noted K_{12} . K_{12} is therefore the total number of collisions, within one cm^3 and per s , between R_1 radius particles and R_2 radius particles, for a concentration of 1 particle per cm^3 of each type. To follow the evolution of the number of N_2 particles with R_2 radius regarding the population of the number of N_1 particles with R_1 radius, resolve the following differential equation (1):

$$\frac{dN_2}{dt} = -\frac{1}{2} \cdot K_{12} \cdot N_1 \cdot N_2 \quad (1)$$

By integration, we obtain equation 2:

$$N_2 = N_{02} \cdot \text{Exp}\left(-\frac{1}{2} \cdot K_{12} \cdot N_1 \cdot t\right) \quad (2)$$

with N_{02} : initial concentration of R_2 radius particles.

The N_2 population would have reduced by half after a $T_{1/2} = -\frac{2 \cdot \text{LN}(0.5)}{K_{12} \cdot N_1}$ (equation 3) time period.

The initial deposition velocities for atmospheric particles and $^{106}\text{Ru-Rh}$ particles are respectively $1 \cdot 10^{-4}$ and $5,7 \cdot 10^{-2} \text{ m}\cdot\text{s}^{-1}$. Then, from measurements performed in environment, we determined, by mathematic adjustment (figure 4), to which coagulation coefficient (K_{12}) would correspond this deposition evolution depending on the distance regarding the discharge point.

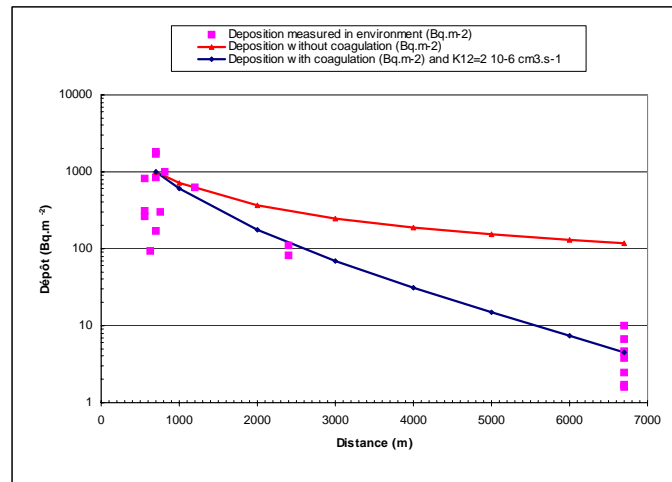


Figure 4. Comparison between $^{106}\text{Ru-Rh}$ deposition in environment and deposition calculated without and with coagulation phenomenon aerosols

The best adjustment between measurements and modelling is determined for a K_{12} value of $2 \cdot 10^{-6} \text{ cm}^3 \cdot \text{s}^{-1}$. For example, a K_{12} coagulation coefficient of $2 \cdot 10^{-6} \text{ cm}^3 \cdot \text{s}^{-1}$ may be obtained for atmospheric aerosols with $0.15 \mu\text{m}$ radius and $^{106}\text{Ru-Rh}$ aerosols with 1 nm radius or an atmospheric aerosols with $0.55 \mu\text{m}$ radius and $^{106}\text{Ru-Rh}$ aerosols with 2 nm radius (table 1). Therefore, it can be noted that $^{106}\text{Ru-Rh}$ aerosols with nanometric radius and atmospheric aerosols for which radius corresponds to the accumulation mode ($0.1 - 0.2 \mu\text{m}$), the evolution of the ruthenium deposition, following the incident of May 18th, 2002, may be explained. The coagulation phenomenon will increase the size of $^{106}\text{Ru-Rh}$ aerosols initially released and therefore reduce the dry deposition velocity of them. Besides, for aerosols of which radius is comprised between a few nanometers and some thousands of nanometers, the deposition

velocity is inversely proportional to the particle radius. Particles thus stuck to the atmospheric aerosol may then be conducted over long distances as their deposition velocity is very low.

Table 1. Values of atmospheric aerosol radius and ¹⁰⁶Ru-Rh aerosol radius determined from sampling campaign carried out in La Hague

Atmospheric release of ¹⁰⁶ Ru-Rh (Bq)	Dry deposition velocity of the atmospheric aerosols (m.s ⁻¹)	Dry deposition velocity of the ¹⁰⁶ Ru-Rh aerosols (m.s ⁻¹)	Coagulation coefficient K ₁₂ .10 ¹⁰ (cm ³ .s ⁻¹)	Time period T _{1/2} (s)	Radius of the atmospheric aerosols (µm)	Radius of the ¹⁰⁶ Ru-Rh aerosols (µm)
4,5 10 ⁹	1 10 ⁻⁴	5.7 10 ⁻²	20000	140	0.15	1 10 ⁻³
4,5 10 ⁹	1 10 ⁻⁴	5.7 10 ⁻²	20000	140	0.55	2 10 ⁻³

CONCLUSION

This study shows that the consideration of the aerosol coagulation phenomenon in the modelling of the atmospheric dispersion and deposition allows explaining the ¹⁰⁶Ru-Rh deposition measured in environment following the incident of May 18th in the COGEMA La Hague facility. To do so, one must take into account the evolution of two different aerosol size distributions and dry deposition velocities. Moreover, the consideration in the modelling of local phenomena (electric field, turbulence...) should allow explaining the significant value of the dry deposition velocity (5,7 10⁻² m.s⁻¹) observed in near field (700 m). Lastly, these various hypotheses shall be invalidated or validated using field experiments carried out producing fluorescein monodispersed aerosols as tracer. This technique is being developed by IRSN.

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