

REVIEW OF LITERATURE ON RUTHENIUM BEHAVIOR IN NUCLEAR POWER PLANT SEVERE ACCIDENTS

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During a hypothetical severe accident in a pressurized water reactor (PWR), fission products (FPs) are released from the nuclear fuel and may reach the reactor containment building. Among the FPs, ruthenium is of particular interest due to its ability to form volatile oxide compounds in highly oxidizing conditions. In addition, ruthenium is a very hazardous compound because it is chemically toxic and also because of its radiotoxicity.

The topic of ruthenium is examined in terms of nuclear safety issues. A review of the literature regarding ruthenium oxides properties, gaseous and aqueous chemistry is compiled. The study focuses on ruthenium tetroxide (RuO_4), which is highly reactive and volatile and is the most likely gaseous chemical form under the conditions prevailing in the containment. The interactions between ruthenium oxides and containment surfaces, which could be most important in overall ruthenium behavior, are also discussed. Finally, an evaluation of the possible revolatilization phenomena of ruthenium adsorbed on PWR containment surfaces or dissolved in the sump under superoxidizing conditions (radiolysis) is also presented. In this case, ruthenium dioxide (RuO_2) must also be considered.

Knowledge of all these phenomena is required to accurately predict ruthenium behavior and to make a best-estimate assessment of the potential ruthenium source term.

aerosol form through the reactor coolant system (RCS), partially condensed onto steel surfaces, and reach the containment. Some fission products, still in gas or aerosol form, may leave the containment via different leaks and will lead to radioactive contamination of the environment: This is called source term.

Among all types of severe accidents, some happen in a highly oxidizing atmosphere due to possible contact between degraded fuel and air. The different possible air ingress pathways have been studied,^{1,2} and three main possibilities have been identified.

The first one corresponds to air ingress during refueling operations. During this process, the reactor vessel head is removed and the fuel is immersed in water. In the case of possible water loss, the refueling cavity could be emptied and fuel would come into direct contact with the air.¹

The second possibility is air ingress following lower-head reactor pressure vessel melt-through due to molten nuclear fuel (called corium) interaction with steel.² Indeed, subsequent to this phenomenon, a gas circulation is created between the reactor pit, lower-head vessel break, and the primary system.

The last possibility is air ingress directly through a break in the primary or secondary coolant system.¹

Other scenarios leading to contact between fuel and air are possible, e.g., a spent-fuel storage pool water-draining accident or a fuel-handling accident.

This technical note reviews the literature regarding ruthenium oxides gaseous and aqueous chemistry in the containment during an oxidizing severe accident. This study focuses particularly on the ruthenium tetroxide chemistry.

I. BACKGROUND

A severe accident would happen in a nuclear reactor if loss of coolant due to an initiating event occurs together with partial or total failure of the safeguard systems, leading to irreversible core dewatering. After the nuclear fuel degradation, fission products released from fuel are transported in gas, vapor, or

II. RUTHENIUM ISSUE

II.A. Does Ruthenium Represent a Real Issue for Nuclear Safety?

The study of ruthenium by Institut de Radioprotection et de Sûreté Nucléaire as well as other organizations devoted to nuclear safety research in France and other countries is motivated by several factors.

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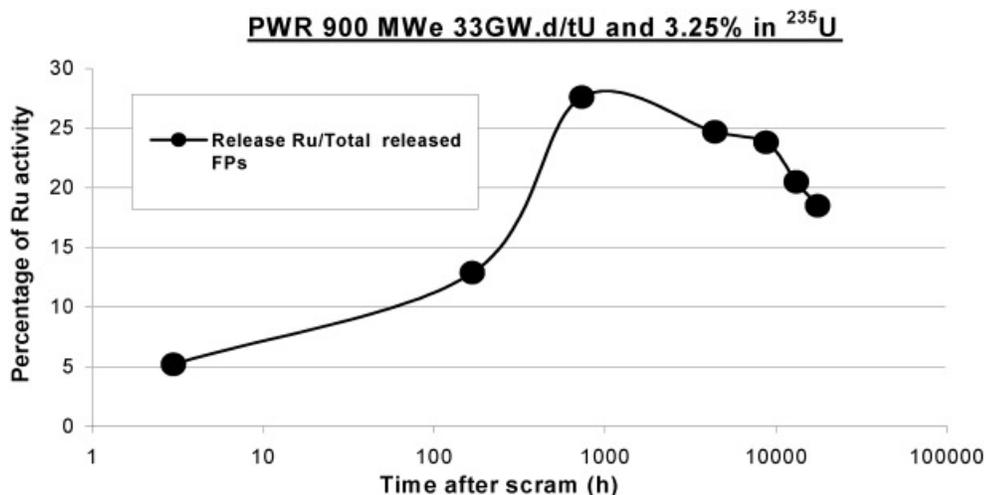


Fig. 1. Percentage of ruthenium activity compared to the total activity of all other fission products.

First, the quantity of ruthenium formed increases with fuel burnup. With identical burnup, it increases with fuel enrichment in ²³⁵U; moreover, it is greater in mixed-oxide fuel than in UO₂ fuel. As a consequence, the amount of ruthenium formed during fuel life will continuously rise in the future, because utilities tend to optimize fuel performances.

Second, ruthenium is a high-specific-activity fission product. The graph in Fig. 1 shows the evolution, during the first 2 yr after an emergency scram, of the important part of ruthenium activity with respect to the total activity of all the fission products released (¹³¹I, ¹³³Xe, etc.)^{3,4}; it has been assumed that 50% of the bundle inventory is released.

Finally, ruthenium has, in addition to its chemical toxicity, which was already known through toxicological studies,⁵⁻⁷ a high radiotoxicity, essentially through its isotopes ¹⁰⁶Ru ($T_{1/2} = 369$ days) and ¹⁰³Ru ($T_{1/2} = 39.3$ days). It is therefore an important short- and middle-term radiocontaminant.

In radiological protection, four groups of radionuclides were defined,⁸ and ¹⁰⁶Ru was ranked in group 2, high radiotoxicity, in the same way as ¹³¹I, and ¹⁰³Ru in group 3, moderate radiotoxicity, as ¹³⁷Cs.

The danger of ruthenium particles is their specific activity and the high associated risk of irradiation and internal contamination (inhalation, ingestion). For example, an 8- μ m-diam particle that has a 1500-Bq activity [calculated for RBMK fuel (Chernobyl) with an average burnup] may induce >1.6 mGy/cm²·h to skin (external irradiation).⁹

Under the action of oxidizing agents, which may be produced either from sump water radiolysis or from air radiolysis, ruthenium metal or dioxide could be oxidized up to +VIII oxidation state. Its volatile compounds (essentially RuO₄) may be present in the containment in the case of a severe accident and could represent a high potential risk for nuclear safety.

II.B. Ruthenium Release from Degraded Fuel and Transport in RCS

Metal ruthenium is considered as a moderately volatile species; then the fraction emitted, from UO₂ fuel heated to

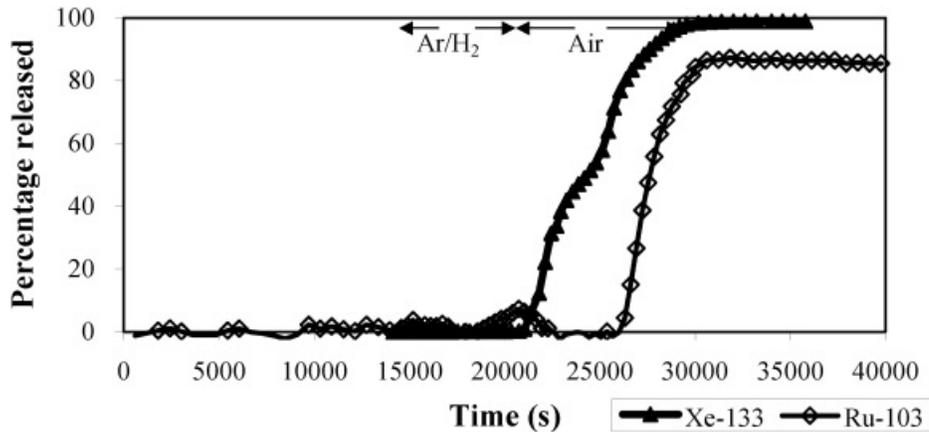
2573 K, varies between 1 and 10% (Ref. 10). Indeed, according to the Ellingham diagram, to obtain ruthenium oxides, it is necessary to have preliminarily oxidized most of the other elements, especially uranium, plutonium, and molybdenum. The presence of volatile ruthenium species is possible only if the fuel is subject to strong oxidation, such as the one resulting from air ingress in the reactor pressure vessel. Affinity for oxygen is in decreasing order: Zr > UO₂ > Ru. So oxygen availability for ruthenium depends strongly on the UO₂ matrix oxidation state.¹ As indicated by Hunt et al.,¹¹ some elements can become oxidized (ruthenium extreme case) only if UO₂ has previously been oxidized in UO_{2+x}. Oxygen potential is of prime importance for the ruthenium release rate.

Ruthenium release from the fuel matrix (or simulant) under air atmosphere has been highlighted by various experiments.

Atomic Energy of Canada Limited carried out some tests that consisted of heating pieces of irradiated fuel pellets under different atmospheres. The fuel burnup of the test called H02 was 10 GWd/t U. This test was performed in an Ar/H₂ atmosphere and next in air. The maximum temperature reached was 2163 K. Figure 2 shows that as soon as an oxidizing atmosphere is present (transition from Ar/H₂ mixture to air atmosphere), ruthenium release increases drastically to reach a value close to that of ¹³³Xe, which is a very volatile compound released at nearly 100%. It is worth noticing the presence of an induction time (~5000 s) assumed to correspond to the time necessary for oxidation of the UO₂ matrix by air; the next ruthenium release is extremely quick.¹²

In the framework of the PHEBUS-RUSET program, in 2002 the Hungarian research organization AEKI studied the ruthenium release at high temperature in an oxidizing environment. These tests confirmed the presence of a gaseous type of ruthenium.¹³

Once released out of the fuel matrix, ruthenium can be transported through the RCS under various physicochemical forms: ruthenium trioxide [RuO₃(g)], ruthenium tetroxide [RuO₄(g)], RuO₂ aerosol (ruthenium dioxide), and ruthenium mixed aerosols (Cs₂RuO₄...) before reaching the containment via the RCS break.



Xenon and Ruthenium release percentages as functions of time for HCE3 test H02 (sheathed Darlington fuel, air, 1890°C).

Fig. 2. Variation of ruthenium and xenon releases according to the nature of gas flow.¹²

The British technology company AEA-Technology and VTT Processes (Finland) have studied ruthenium transport. It has been observed that the RuO_4 partial pressure measured is far higher than the predicted thermodynamic equilibrium calculation, so ruthenium tetroxide decomposition may have been limited.^{14,15} Similar results were obtained in other experiments.¹⁶

Interesting information can also be drawn from the Chernobyl accident even if this accident is not strictly a pressurized water reactor (PWR) severe accident; nevertheless, a strong ruthenium release occurred. Indeed, total releases of ^{103}Ru were higher than those of ^{137}Cs , 9.3×10^{16} versus 4.7×10^{16} Bq (Ref. 17), corresponding to $\sim 2.9\%$ of the ruthenium initial inventory,¹⁸ so it confirms the possibility that ruthenium can be released outside the fuel matrix.

Many radioactive particles emitted contain ruthenium, essentially in metal form.¹⁷ Various authors have tried to explain the origin of this metal form, but assumptions vary from one author to another. Nevertheless, Broda et al.¹⁹ have formulated a pertinent hypothesis: They postulated that some radioactive compounds, emitted in gas form, precipitated and condensed when they reached the airborne plume. Ruthenium and molybdenum volatile oxides, formed due to heating from graphite combustion, reduced or decomposed at high temperature, are probably the main compounds of such precipitates. This scenario has been subsequently suggested as the origin of the very high purity of radioactive ruthenium and of the fact that many particles have been found incorporated in large fragments of nonradioactive material. In addition, the formation of volatile compounds is also confirmed, because the condensed ruthenium particles showed a constant $^{103}\text{Ru}/^{106}\text{Ru}$ isotopic ratio.

This is only possible if they come from an initial gaseous release form because this ratio depends on burnup, so variable according to the ruthenium location in the fuel matrix.

In the release of solid particles from fuel, this ratio could not be identical, it would change according to the fragments analyzed.²⁰

All these experimental observations indicate that ruthenium can be released from fuel in large quantities according to conditions and next transported through the RCS. But then, how do emitted ruthenium compounds behave in the containment?

III.C. Ruthenium Behavior in the Reactor Containment

Ruthenium behavior is directly linked to the nature of the ruthenium compound. The $\text{RuO}_3(\text{g})$ is unlikely to reach the containment because it is thermodynamically unstable in the temperature range of the containment severe-accident conditions.^{21,22} In the gas phase, mixed aerosols are not the major risk from a source-term point of view, because after aerosol settling in the aqueous phase (sump), ruthenates will be formed. The two essential species are RuO_4 , the most stable gaseous form in containment conditions, and the little soluble aerosol RuO_2 , which can have an impact in terms of potential revolatilization under RuO_4 form.

In the aqueous phase, ionic compounds of ruthenium, essentially $\text{RuO}_4^-/\text{RuO}_4^{2-}$, have to be taken into account because they can be involved in reactions leading to the formation of either gaseous RuO_4 or RuO_2 .

The potential reactivity of ruthenium onto surfaces is important. Therefore, the important surfaces present inside the PWRs' containment have to be considered in terms of reactivity; these surfaces are made of painted steel and stainless steel surfaces (Types 316L and 304L) (Ref. 23). Ruthenium tetroxide seems to have a certain affinity for iron oxides^{22,24} and for organic compounds.^{25,26} Reactions involving RuO_4 with the internal surfaces of the containment can be expected to occur, especially as the developed total surface is very large—several thousands of square meters.

During a severe accident, there is a high dose rate in the containment in the gaseous and aqueous phases resulting from the large amount of fission products present in the containment. Radiolytic reactions occur in the atmosphere and the sump leading to the formation of oxidants that might oxidize ruthenium up to a +VIII oxidation state.

III. GENERAL PROPERTIES CONCERNING RUTHENIUM OXIDES

Although it is an iron homologue, ruthenium has neither the same reactivity nor the same chemistry. Ruthenium chemistry is extremely complex; many compounds are known—inorganic as well as organic forms. Its external electronic structure is $4d^75s^1$, which gives it nine stable oxidation states lying between 0 and +VIII, and even a $-II$ oxidation state, $[\text{Ru}(\text{CO})_4^{2-}]$ (Ref. 25). Our paper is limited to compounds that could be involved in nuclear safety issues.

With alkali metals and alkali earth metals present in large quantity as fission products, ruthenium can form mixed compounds, such as Cs_2RuO_4 , SrRuO_4 , BaRuO_3 , etc.

Ruthenium halides can also be quoted: RuF_6 to RuF_3 , RuCl_3 , RuCl_2 , RuI_3 , RuI_2, \dots (Ref. 27). The last two compounds are of particular interest, because iodine is a very hazardous fission product.²⁸ In aqueous phase, oxidation states +VI, +VII, and +VIII give essentially tetrahedral oxocomplex ruthenate ions (RuO_4^{2-}) and perruthenate ions (RuO_4^-), as well as ruthenium tetroxide and its hydrolyzed form, H_2RuO_5 .

The possible existence of ruthenium hydroxides and ruthenium oxyhydroxides is mentioned in the literature even if the data are scarce.

Besides metal ruthenium, ruthenium aerosols (type Cs_2RuO_4), and possibly ruthenium hydroxides, ruthenium oxides are the main compounds supposed to be formed during a severe accident. The four main ruthenium oxides are RuO , RuO_2 , RuO_3 , and RuO_4 (Ref. 29).

Table I lists all the ruthenium compounds that may be concerned in this topic; they are classified according to their oxidation states. Letters (g), (aq), and (c) refer, respectively, to gaseous, aqueous, and condensed phases.

The main characteristics of ruthenium oxide compounds are the following:

1. *RuO*: This oxide is only gaseous. According to Schäfer et al., when the temperature is >1273 K, the three oxides RuO_4 , RuO_3 , and RuO_2 are decomposed into RuO (Ref. 30). How-

ever, various authors, such as Rard³¹ and Bell²¹ have doubts concerning its existence. After experimental works, Eichler et al.²⁹ claim that RuO monoxide only exists when temperatures are >1900 K. Enthalpy of formation has been established by Norman et al.³² at 1900 K.

2. *RuO₂*: Ruthenium dioxide is the most stable ruthenium oxide. It appears under brown-black powder with a tetragonal rutile structure, type TiO_2 . One means of production is to roast Ru_{metal} in an oxygen stream in a very high temperature muffle furnace²⁵; another is to dehydrate $\text{Ru}(\text{OH})_3$ hydroxide at 723 K (Ref. 33). By heating RuO_2 under an oxygen flow, volatilization takes place, indicating the formation of higher oxidation-state oxides. These oxides, which are thermodynamically unstable, decompose on cooling to form anhydrous RuO_2 blue crystals. This phenomenon has been experimentally confirmed by Rard.³¹ In anhydrous form, dioxide is insoluble, but a hydrated form exists ($\text{RuO}_2 \cdot x\text{H}_2\text{O}$ with $2 < x < 3$), so its dissolution becomes possible³³ even if this is quite difficult. RuO_2 is also assumed to exist in gaseous form but solely at very high temperatures, >2000 K (Ref. 34), as in the reactor core in severe-accident conditions.

3. *RuO₃*: Very little information is available in the literature concerning this oxide. Its existence in gas form was questioned for a long time, but Schäfer et al.³⁰ and Bell and Tagami²¹ in 1963 proved its existence. According to these authors, the partial pressures of ruthenium trioxide and ruthenium tetroxide are equal in a ($\text{RuO}_2 + \text{O}_2$) environment at 1 atm and at 1327 K. RuO_3 predominates in the gaseous phase above this temperature up to 2000 K, and RuO_4 , below this temperature (1327 K). Eichler et al.²⁹ indicate that ruthenium trioxide in a gaseous phase can condense in dioxide form. Moreover, $\text{RuO}_3(\text{g})$ could also be transformed in solid RuO_3 form³⁵ even if this hypothesis is not accepted by all the scientific community.

4. *RuO₄*: Ruthenium tetroxide can exist in a gaseous state or in a condensed state. The properties of this compound are extensively detailed later. An important characteristic is that

TABLE I
Possible Ruthenium Oxidation States Discussed in This Paper

Oxidation State	Type of Compound	Name	Physical State
+8	RuO_4 (associated with $\text{H}_2\text{RuO}_5/\text{HRuO}_5^-$)	Ruthenium tetroxide	(c), (aq), (g)
+7	RuO_4^-	Perruthenate ion	(aq)
+6	RuO_4^{2-} RuO_3	Ruthenate ion Ruthenium trioxide	(aq) (g)
+5	$\text{Ru}_2\text{O}_5, 2\text{H}_2\text{O}$	Hydrated pentavalent ruthenium oxide	(c)
+4	RuO_2 $\text{RuO}_2 \cdot 2\text{H}_2\text{O}$ $\text{Ru}(\text{OH})_2^{2+}/\text{Ru}_4(\text{OH})_{12}^{4+}$ $\text{Ru}(\text{OH})_4$	Ruthenium dioxide Hydrated ruthenium dioxide Ruthenium monomer/tetramer Ruthenium hydroxide	(c) (c) and (aq) (aq) (aq)
+3	$\text{Ru}^{3+}/\text{Ru}(\text{OH})_3$	Trivalent cation and its hydroxide	(aq)
+2	Ru^{2+}	Divalent cation	(aq)
0	Ruthenium	Metal ruthenium	(c)

this oxide is very volatile with very toxic vapors. To illustrate this point, the liquid-vapor equilibrium constant of ruthenium tetroxide has been calculated with GEMINI 1 (Ref. 36) and the thermodynamic databank associated COACH:



The constant of this reaction represents the RuO_4 vapor pressure in the standard reference state.

IV. RUTHENIUM OXIDE CHEMISTRY IN SEVERE-ACCIDENT CONDITIONS

IV.A. Ruthenium Reactivity in Gas Phase

From a nuclear safety point of view, ruthenium compounds can be split into two categories, the gaseous forms and the aerosol forms, amounts of which are very much predominant. Possible gaseous forms can only be RuO_3 and RuO_4 . However, the maximal temperature in the containment in the case of a severe accident will very likely remain below 413 K, a value not compatible with the thermodynamic stability of trioxide,^{21,22} as already mentioned.

Figure 3, obtained by means of the thermodynamic code COACH-GEMINI 1 (Ref. 36), indicates clearly that RuO_3 decomposes at ~ 1500 K to form RuO_2 and RuO_4 .

Even if thermodynamic equilibrium calculations also show $\text{RuO}_4(\text{g})$ instability at low temperature and all the more in the temperature range of the containment (313 to 413 K), several arguments favor a more complete study of the reactivity associated with this species.

First, recent experimental data^{13,14} tend to prove that RuO_4 would be formed in the RCS in a nonnegligible amount. Second, RuO_4 has a very high saturation vapor pressure, so its volatility makes it hazardous for the population. Third, in the reactor containment conditions (oxidizing medium), this high oxidation state (+ VIII) could be stabilized.

Concerning aerosols, there are mixed ruthenium aerosols and ruthenium dioxide $\text{RuO}_2(\text{s})$. Mixed aerosols are water soluble, and ruthenium dioxide, on first analysis, does not represent a major danger in this form, except in the case of early containment failure, but it can be a possible source of $\text{RuO}_4(\text{g})$ formation by oxidation of aerosol deposits on the walls. To sum up, to be able to make an accurate estimation of the ruthenium source term, a better understanding of tetroxide and dioxide behavior is needed.

IV.A.1. Ruthenium Tetroxide

For several authors,^{24,37} ruthenium tetroxide appears in two forms in solid state. The first condensed state is brownish globules, which are little soluble in water and seem to constitute a crystalline state. They are produced by solidification of liquid tetroxide below 300 K. The second state corresponds to very unstable yellow needles, which are obtained by condensation of tetroxide in the vapor state. These needles melt at ~ 298 K. This form is more soluble in water and produces a golden yellow solution; its solubility in water is estimated at 17.1 g/l at 273 K and 20.3 g/l at 293 K (Ref. 26). It is also very soluble in tetrachloromethane and produces an orange red solution. Its sublimation can be observed at 280 K with a pressure of ~ 4 mbar (Ref. 24). This crystalline form can be obtained with sublimation and next condensation of the brownish form.

However, a study by Nikol'skii³⁸ focused on determining the melting point, vapor pressure, and solubility in water refutes this hypothesis of ruthenium tetroxide polymorphism.

Tetroxide vapors are yellow and toxic and smell strongly like ozone. They are reactive, particularly with organic materials and reducing agents. Sensitivity to photolysis has also been demonstrated.³⁵ Indeed, in a dark place in dry atmosphere and in a sealed tube, ruthenium tetroxide can be stored for several years. On the other hand, with light and humidity, its decomposition happens quickly.

The duration was not indicated. RuO_4 is also supposed to decompose in an explosive way into RuO_2 when temperatures

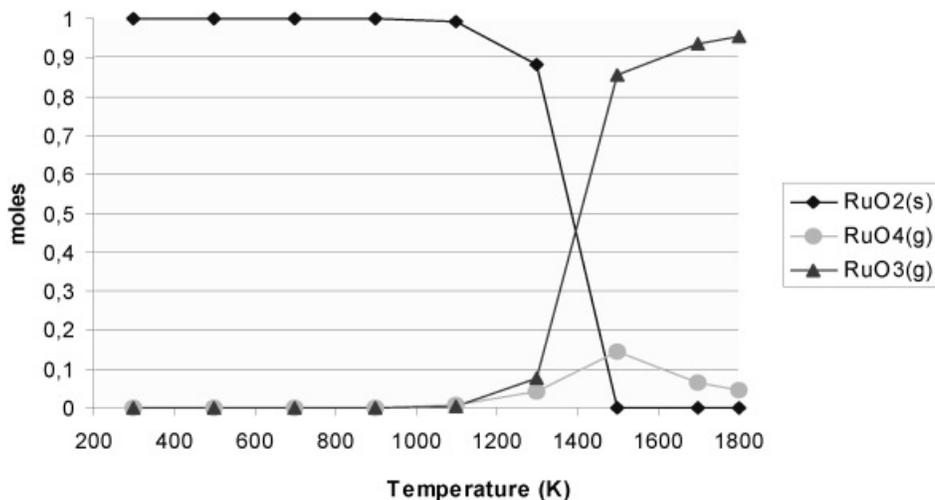


Fig. 3. Thermodynamic stability of $\text{RuO}_4(\text{g})$, $\text{RuO}_3(\text{g})$, and $\text{RuO}_2(\text{s})$ as a function of temperature for, respectively, 1 mol of ruthenium, 200 mol of H_2O , and 200 mol of O_2 (15% M).

are near 381 K (Ref. 39). This last point is not mentioned in any other paper; nevertheless, it is worth noticing that only these authors have manipulated ~ 100 g of tetroxide, which represents a huge amount—never handled again.

IV.A.1.a. Thermal Stability of RuO₄(g). RuO₂ stability and RuO₄ volatility can be easily explained with covalent bonds. Binding energy between tetroxide atoms is low because they are perfectly saturated, and the structure is symmetrical, so this compound can be easily melted, sublimated, and volatilized.

Few experimental studies on ruthenium tetroxide stability in gaseous phase have been performed. Ruthenium tetroxide decomposition mechanisms and the associated kinetics are still unknown. According to Leudet,⁴⁰ tetroxide decomposition at low temperatures takes place according to the following reaction:



The equilibrium constant is provided by the following relation:

$$\ln K = \frac{14880}{T} - 11.53 + 1.135 \ln T \quad (T \text{ in K}) \quad .$$

The RuO₄(g) stability rises with temperature. The equilibrium constant calculated (7×10^{19} at 293 K and 2×10^{15} at 373 K) indicates an almost complete decomposition if equilibrium is reached.

Ortner⁴¹ introduced gaseous tetroxide in a sealed tube (Carius tube) maintained at a constant temperature for some time and then rapidly cooled to stop the reaction, but the precise boundary experimental conditions were not provided.

They noticed that in the presence of wet air, decomposition happens only after an induction time, the duration of which depends on steam concentration and temperature. But, after adding a drying agent, the induction time has been shortened. No hypothesis was formulated.

The decomposition seems to be initiated by ruthenium deposition on glass walls; this would indicate an autocatalytic reaction, all the more that in introducing RuO₂ into the tube, induction time was also reduced. The decomposition kinetic was studied between 383 and 413 K with an RuO₄ initial pressure of ~ 0.3 atm in the presence of 25 to 150 mg of RuO₂. The determined kinetic rate law is of first order with respect to RuO₄. An Arrhenius law expressed the rate constant:

$$k \text{ (min}^{-1}\text{)} = 7.12 \times 10^{14} \exp(-128 \times 10^3/(\text{room temperature})) \\ (T \text{ in K}) \quad .$$

Corresponding reaction half-lives calculated at different temperatures are reported in Table II.

The kinetics of the RuO₄ decomposition is very slow at room temperature, which may justify the existence of RuO₄(g) even if it is not thermodynamically stable. The temperature range ($\Delta T = 30$ K), where the kinetic data were determined, is limited, and above 373 K, extrapolation is uncertain. Moreover, the possible explosive decomposition of tetroxide at temperatures near 381 K noticed by Sainte Claire Deville and Debray³⁹ is not consistent with this study even if the possible explanation is that the amount of the tetroxide involved in these tests was weaker. The influence of the presence of metal ruthenium in powder form on the decomposition rate has also

TABLE II
Ruthenium Tetroxide Decomposition Kinetic Constant
Between 293 and 423 K (Ref. 41)

Temperature (K)	k (min ⁻¹)	Calculated Half-Life (h)
293	1.3×10^{-8}	8.9×10^5
323	1.6×10^{-6}	7220
373	9.6×10^{-4}	12
423	1.2×10^{-1}	9.6×10^{-2}

been studied. The results obtained for different concentrations of tetroxide indicate that 54% of RuO₄ is decomposed after 3 h at 383 K, and after 2 h at 413 K, it is 100%. Nevertheless, the lack of information about the precise conditions of experimentation limits the interest of these results.

At the end of the 19th century, Debray and Joly⁴² had already pointed out the impact of water steam on ruthenium tetroxide stability. Indeed, RuO₄ can react with water steam to form a black coating of lower oxide. This black amorphous precipitate, dried in an oven at 383 K, consists of Ru₂O₅·2H₂O, and even after drying, it remains partially hydrated. Tetroxide decomposition, which is slow in the dark, is accelerated with sunlight, especially with ultraviolet rays. According to Wöhler et al.,⁴³ dioxide hydrates would be formed.

Ortins de Bettencourt and Jouan²⁴ have also investigated RuO₄ decomposition kinetics. Even though they had great experimental difficulties, particularly in tetroxide generation and its transfer into the reactor vessel, they confirmed that the reaction is of first order.

More recently, Zimmerman et al.³⁵ have again shown interest in RuO₄ decomposition and the resulting products. Their study is based on irradiation of RuO₄(g) by a mercury arc lamp or by a photolytic flash. Photolytic decomposition was observed in measuring the absorbance decrease of RuO₄.

IV.A.1.b. RuO₄(g) Reactivity Under Radiation. The formation of air radiolysis compounds has been extensively studied.^{44–46} Willis and Boyd⁴⁷ published all the radiolytic yield values (G values) of primary compounds coming from air radiolysis, as well as the reactions between these different species. The set of thermal and radiolytic reactions can be solved by the Facsimile software⁴⁸ in order to predict the gas mixture composition as a function of boundary conditions. By way of illustration, the main compounds generated by radiolysis are shown in Fig. 4 for prototypical conditions (10 kGy/h dose rate, 373 K, steam mass fraction = 30%, and no potential sorption of radiolysis compounds on surfaces).

The main radiolytic compounds present in the containment in the case of a severe accident are NO₂, N₂O, and O₃, which are oxidants.

The ruthenium tetroxide concentration in containment atmosphere is quite difficult to evaluate in a severe accident; it will nevertheless be extremely low with respect to atmospheric compounds. It is then legitimate to assume that most of the energy released by ionizing radiations is distributed between N₂, O₂, and H₂O molecules, and the influence of radiation on ruthenium species will be due to chemical reactions with air radiolysis compounds rather than to direct radiations. The

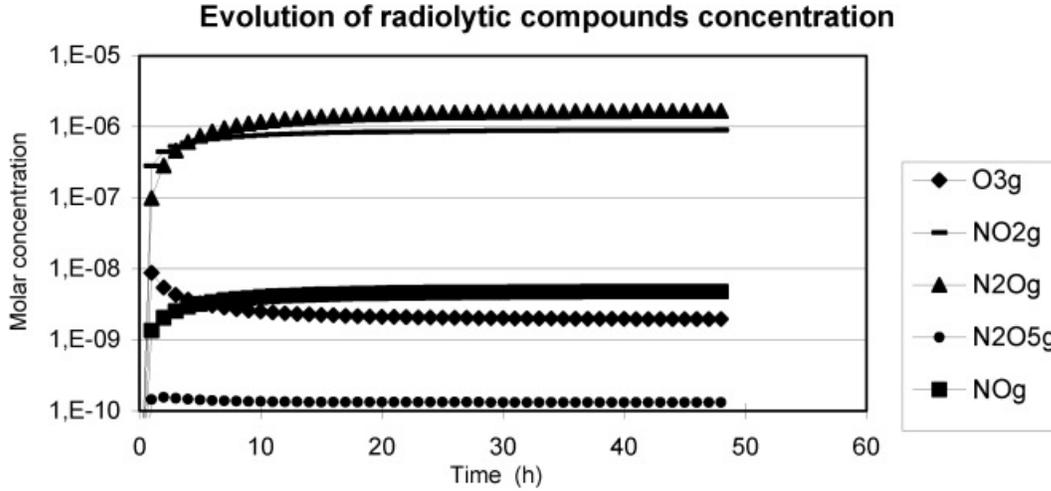


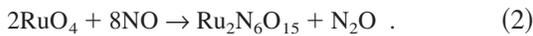
Fig. 4. Air radiolysis compounds concentration profiles as a function of time (DR = 10 kg/h at 373 K, steam mass fraction = 30%).

$\text{RuO}_4(\text{g})$ stability under radiolysis is legitimately questionable. The few experimental tests from literature are summarized in this paper.

Igarashi et al.⁴⁹ have studied NO_x action on $\text{RuO}_4(\text{g})$ stability. The addition of NO and NO_2 facilitates ruthenium absorption in water. This effect is even stronger in the case of monoxide. Addition of NO_2 with an NO_2/RuO_4 molar ratio of ~ 189 reduces the proportion of tetroxide in the gas phase by a factor of 168, and the addition of NO with a 22 ratio, reduces it by 1180.

This phenomenon could be due to tetroxide transformation into a nitrosylruthenium complex, which is less volatile and more soluble in water.

In 1955, Fletcher et al.⁵⁰ demonstrated the formation of a species of binuclear ruthenium. This species is obtained via the reaction



In the presence of water, the final compound is not only in $\text{Ru}_2\text{N}_6\text{O}_{15}$ form; aquo-nitrato- RuNO^{3+} forms are also generated.⁵¹

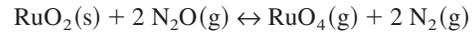
IV.A.2. Thermal and Radiolytic Reactivity of Ruthenium Dioxide

If $\text{RuO}_2(\text{c})$ reactivity with air is reasonably limited because of its thermodynamic stability, a possible release of $\text{RuO}_4(\text{g})$ at 600 K has been shown experimentally¹³ coming from the $\text{RuO}_2(\text{c})$ deposit; the RuO_4 partial pressure measured is $\sim 10^{-6}$ bar. Maas and Longo⁵² also suspected the possibility of reoxidizing ruthenium dioxide with air at low temperatures.

From a theoretical point of view, the equilibrium constant calculated³⁶ for the reaction $\text{RuO}_2(\text{s}) + \text{O}_2(\text{g}) \leftrightarrow \text{RuO}_4(\text{g})$ is equal to 5×10^{-17} at 373 K. For an oxygen partial pressure of ~ 0.2 bar, the resulting tetroxide concentration is $\sim 3 \times 10^{-19}$ mol/l. This reaction cannot form tetroxide in great proportion. The RuO_2 reactivity seems to be limited with oxygen, but for radiolysis products it is no longer the case.

Reactivity with ozone, $\text{RuO}_2(\text{s}) + \frac{2}{3}\text{O}_3(\text{g}) \leftrightarrow \text{RuO}_4(\text{g})$, is much higher, indeed $k = 0.2$ at 373 K. For prototypical ozone concentration, estimated at $\sim 10^{-8}$ mol/l or 3×10^{-7} bar, the tetroxide partial pressure obtained is $\sim 2 \times 10^{-7}$ bar, or 6×10^{-9} mol/l in terms of concentration.

Reactivity with nitrous oxide is also enhanced, as indicated by the following reaction:



$$k = 2.6 \times 10^{14} \text{ at } 373 \text{ K} \quad .$$

According to the air radiolysis model, for an N_2O concentration of $\sim 10^{-6}$ mol/l, corresponding to 3×10^{-5} bar, the consumption of all $\text{N}_2\text{O}(\text{g})$ will lead to the production of RuO_4 with a 1.5×10^{-5} bar partial pressure once equilibrium has been reached.

Another reaction between ruthenium dioxide and nitrogen dioxide is also possible by the following reaction:



$$k = 0.2 \text{ at } 373 \text{ K} \quad .$$

If the NO_2 concentration is identical to that of N_2O , equilibrium is not moved much to $\text{RuO}_4(\text{g})$ formation. For a NO_2 partial pressure equal to 10^{-6} mol/l, P_{RuO_4} should not be $> 10^{-10}$ bar when equilibrium is reached. This reaction does not constitute a potential source of gaseous tetroxide formation.

Even if RuO_2 and RuO_4 are the species of main interest, ruthenium hydroxides or oxyhydroxide species in a gaseous state can also exist. Unfortunately, very few data in literature are available on this subject, insofar as no experimental evidence has been published yet. Only a few authors have mentioned them because they have demonstrated their thermodynamic stabilities in the gaseous phase. Two compounds might exist, RuO_3OH and RuOH . In our conditions, only $\text{RuO}_3(\text{OH})$ could come up because it would be stable when temperatures are low. Its partial

TABLE III
RuO₄ Solubility in Water as a Function of Temperature²⁵

	Temperature (K)									
	273	283	288	293	298	303	313	323	343	348
Solubility (g/ℓ)	17.12	18.53	19.63	20.33	21.62	21.13	20.81	20.88	21.41	22.49

pressure is supposed to be close to 10⁻⁷ bar at 500 K according to the FACT thermodynamic code.⁵³

In the absence of references, all previous equilibrium constants were calculated with COACH, the database associated with GEMINI 1 (Ref. 36).

The study of previous reactions associated with the few experimental results highlights that RuO₂ deposits could be oxidized to a significant extent leading to gaseous ruthenium tetroxide formation with partial pressures that may reach 10⁻⁷ to 10⁻⁵ bar in the containment.

IV.B. Ruthenium Thermal Reactivity in Aqueous Phase

A severe accident leads to the formation of an aqueous phase in the containment (with a minimum volume of ~200 m³ corresponding to the water volume of the RCS for a French 900-MW(electric) PWR, for example).

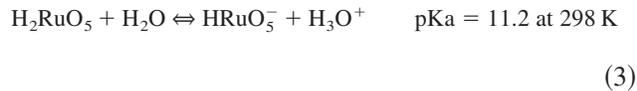
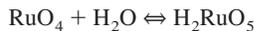
Ruthenium behavior in aqueous solution has to be considered for an accurate source-term evaluation. Indeed, the aqueous reactions could either trap ruthenium in solution through stabilization or produce volatile ruthenium tetroxide.

Possible forms in solution are mainly RuO₄ transferred from the gaseous phase, RuO₂ coming from the soluble aerosol settling, and ruthenate and perruthenate coming from the settling of mixed aerosols. Sump pH may vary from 4 to 10, according to the availability of the spray system.

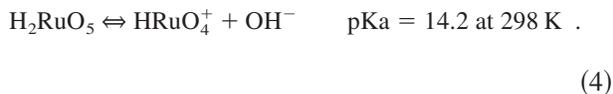
IV.B.1. Behavior of Dissolved Tetroxide

Ruthenium tetroxide is a soluble molecule in water, as reported in Table III. According to the pH solution, it can be present in H₂RuO₅ acid form, known as hyperruthenic, and it is an amphoteric species. Martin⁵⁴ suggested that RuO₄ could act as a weak acid and as a weak base, as shown with the following equilibria:

weak acid:



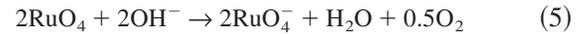
weak base:



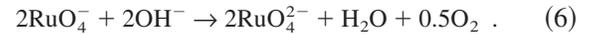
Zoubov and Pourbaix⁵⁵ have specified that tetroxide decomposes rapidly with light, forming a black deposit of

RuO₂·2H₂O hydrated dioxide. Hydrochloric acid reduces RuO₄ in tetra- and trivalent complex forms, becoming oxidized into chlorine.

Analyses of pH-potential scale³³ and that of redox potential values, indicated in Table IV, show that in the pH range of interest (4 to 10), tetroxide is not stable at 298 K in aqueous solutions without oxidizing and complexing agents. Reduction to hydrated ruthenium dioxide or into ruthenium metal occurs. In a sodium hydroxide solution, tetroxide is reduced in perruthenate (Ru-VII) and ruthenate (Ru-VI) ions²⁶:



and



Nowogrocki and Tridot⁵⁶ have studied the Eq. (6) reaction in a RuO₄⁻ concentration range of ~10⁻² mol/ℓ, and the following kinetic rate law was determined:

$$\frac{-d[\text{RuO}_4^-]}{dt} = k [\text{RuO}_4^-]^2 \cdot [\text{OH}^-]^3$$

with $k = 20 \ell^4/\text{mol}^4 \cdot \text{h}$ (at room temperature) .

Nikitina et al.⁵⁷ have also studied the kinetics of perruthenate reduction into alkaline solution but in much lower concentration ranges (10⁻⁴/10⁻⁶ mol/ℓ). With constant ionic strength, the kinetic rate law of RuO₄⁻ reduction was established:

TABLE IV
Redox Potential in Acid and Alkaline Medium Calculated with a 10⁻¹⁰ mol/ℓ Reference Concentration in Ruthenium³³

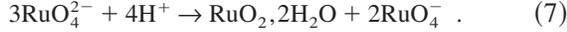
Oxidizing/Reducing Agent	E° (V)	E° (V) - k pH (4)	E° (V) - k pH (10)
RuO ₄ (aq)/RuO ₄ ²⁻	1.59	1.59	1.59
RuO ₄ (aq)/RuO ₄ ⁻	1.00	1.00	1.00
RuO ₄ ²⁻ /RuO ₂	2.01	1.24	0.53
RuO ₄ ⁻ /RuO ₂	1.53	1.02	0.55
RuO ₄ (aq)/RuO ₂	1.40	1.02	0.65
RuO ₂ /Ru ²⁺	1.12	0.94	0.23
RuO ₂ /Ru	0.79	0.55	0.19
O ₂ /H ₂ O	1.23	0.99	0.64

$$\frac{-d[\text{RuO}_4^-]}{dt} = k_1 [\text{RuO}_4^-]^2 \cdot [\text{OH}^-] + k_2 [\text{RuO}_4^-]$$

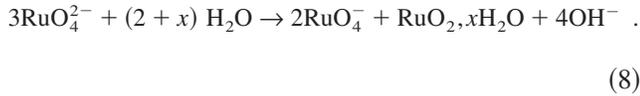
with $k_1 = 0.41 \pm 0.08 \ell^2/\text{mol}^2 \cdot \text{s}$ and $k_2 = (1.2 \pm 0.3) \times 10^{-4} \text{ s}^{-1}$ at 293 K.

These two kinetic rate laws provide very significant differences in the perruthenate disappearance rate.

As shown on the pH-potential scale, at room temperature, orange solutions of ruthenate are stable in a very alkaline medium. If the pH is sufficiently decreased, below 12, ruthenate disproportionates into green perruthenate and $\text{RuO}_2 \cdot 2\text{H}_2\text{O}$, as indicated in Eq. (7):



This ruthenate disproportionation reaction was checked experimentally by Connick and Hurley,⁵⁸ who worked with quite low ruthenium concentrations, $\sim 10^{-5} \text{ mol}/\ell$. However, they suggested the following general reaction:



The precipitate obtained was analyzed with X-ray diffraction (before and after heating at 413 K and next at 973 K), and the results confirm the $\text{RuO}_2 \cdot 2\text{H}_2\text{O}$ form.

For pH lower than 12, perruthenate can now be reduced into hydrated dioxide as follows:

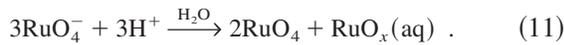


Nevertheless, the degree of hydration of the ruthenium dioxide is still uncertain. Zoubov and Pourbaix⁵⁵ are inclined to favor $\text{RuO}_2 \cdot 2\text{H}_2\text{O}$ formation, while other authors, like Nowogrocki and Tridot⁵⁶ or Rard,³¹ proposed the generic $\text{Ru}_2\text{O}_5 \cdot x\text{H}_2\text{O}$ compound formation instead.

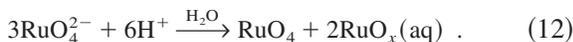
Zoubov and Pourbaix⁵⁵ indicated that when pH is lower than 7.5, ruthenate ion decomposition can lead to the formation of RuO_4 or H_2RuO_5 (hyperruthenic acid). This statement was also confirmed by Nowogrocki and Tridot,⁵⁹ who performed ruthenate and perruthenate solution acidification tests by making measurements with spectrophotometry, pH-metry, and conductimetry. Ionic equilibria and redox equilibria make it possible to confirm Zoubov's results. They suggest a reactional mechanism that is slightly different from the previous ones, in particular due to the formation of a pentavalent ruthenium hydrated oxide instead of hydrated dioxide, such as



and



It can be summarized as



Nowogrocki and Tridot⁵⁶ demonstrated the possible formation of volatile ruthenium tetroxide from a 0.5 N sodium

TABLE V
Stability Domain of Ruthenium Oxides in Aqueous Solution as a Function of pH

	pH		
	4	7	10
Stable species	$\text{RuO}_2(\text{c})$	$\text{RuO}_2(\text{c})$ RuO_4^-	RuO_4^- RuO_4^{2-}

hydroxide solution containing perruthenate ions in the presence of an oxidizing hypochlorite agent.

This information is important in terms of nuclear safety because these boundary conditions are probable. Ruthenium ionic species [(+VII) and (+VI)] can be present in acid or in alkaline sump; oxidizing agents (due to radiolytic compounds) can also be there. To sum up, the aqueous phase could be a potential source of volatile ruthenium. Table V recapitulates the stability domain of the main ruthenium compounds as a function of pH conditions.

IV.B.2. RuO_4 Volatility

In studying RuO_4 partition coefficients between the organic phase and the aqueous phase for different acidities, Martin⁵⁴ noticed that ruthenium tetroxide in water and in CCl_4 made an ideal mixture (for dissolved RuO_4 concentration lower than 0.2 mol/l; RuO_4 vapor pressure in equilibrium with CCl_4 solution at 303 K obeys Henry's law).

Nikol'skii³⁸ measured RuO_4 saturation vapor pressure between 1 and 364 K. Two empirical equations were recommended to calculate P_{SatRuO_4} as a function of temperature (lower or higher than 298 K corresponding to the tetroxide melting point). Sublimation, vaporization, and fusion enthalpies were also calculated, as well as theoretical melting (298 K) and boiling points (364 K):

1. For $273 \text{ K} < T < 298 \text{ K}$:

$$\log P = \frac{2880.3}{T} + 10.673 \quad (T \text{ in K and } P \text{ in mm Hg}) \quad .$$

The uncertainty associated with P is $\sim 5\%$.

2. For $298 \text{ K} < T < 364 \text{ K}$:

$$\log P = \frac{-2847.9}{T} + 1.74 \log T - 0.008406T + 8.737$$

$$(T \text{ in K and } P \text{ in mm Hg}) \quad .$$

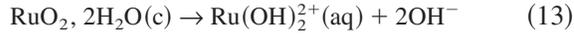
The uncertainty associated with P is $\sim 2\%$.

In a comparison between Nikol'skii's values and those of the COACH thermodynamic database,³⁶ 300 and 360 K, respectively, are obtained as well as 1.5×10^3 and $2.5 \times 10^4 \text{ Pa}$ with Nikol'skii relationships and 4.9×10^2 and 8.8×10^3 with the COACH database. Uncertainties concerning vapor pressure are quite significant.

IV.B.3. Soluble Ruthenium Dioxide

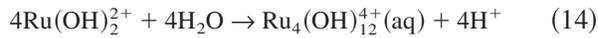
Ruthenium dioxide in its hydrated form, $\text{RuO}_2 \cdot x\text{H}_2\text{O}$ with $2 < x < 3$, may be soluble in the aqueous phase even if solubility turns out to be difficult, especially in a noncomplexing acid medium.³³

Solubility is due to the formation of the following specific compounds (monomer or polymer):



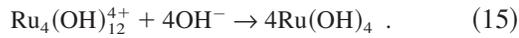
$$K_S = [\text{Ru}(\text{OH})_2^{2+}(\text{aq})][\text{OH}^-]^2 = 8.5 \times 10^{-28}$$

and



$$K_S = 7 \times 10^{-46} .$$

There is a slow polymerization of the Ru(IV) monomer, which makes thermodynamically stable tetramer.^{60,61} This species was observed experimentally by Bremard et al.⁶²; they also showed that this tetramer can precipitate into an hydroxide according to Eq. (15):



Equation (15) is purely a formality for ion precipitate because it is probably hydrated.

Finally, even if according to the pH-potential scale, $\text{RuO}_2 \cdot 2\text{H}_2\text{O}$ is insoluble in noncomplexing acid solutions, it can be dissolved partially in mono- or polynuclear forms. It could even appear in a hexavalent state in ruthenate form in the presence of oxygenated alkaline solutions.⁵⁵

IV.B.4. Ruthenium Hydroxides

$\text{Ru}(\text{OH})_3$ hydroxide is stable in aqueous phase whatever the pH and for a certain range of potential. Otherwise, it is likely to be oxidized into hydrated dioxide $\text{RuO}_2 \cdot 2\text{H}_2\text{O}$ or $\text{Ru}(\text{OH})_4$ even into RuO_4 ; it can also be easily reduced into elementary ruthenium.⁵⁵

IV.B.5. Soluble Ruthenium Aerosols

When soluble ruthenium aerosols are dissolved, they liberate either ruthenate or perruthenate anions. As a consequence, the chemistry involved is the same as already described for these compounds.

IV.C. Ruthenium Reactivity Under Radiolysis in Aqueous Phase

The presence of radionuclides in sump in severe-accident conditions will produce a large dose rate (up to 10 kGy/h) in this phase and will then lead to aqueous solution radiolysis phenomena. As the main compound in this phase is water, we can easily go back to a water radiolysis issue. Sodium hydroxide introduced via the containment spray system does not question this hypothesis nor the initial presence of boric acid (H_3BO_3) in the primary system. Water radiolysis produces molecular, ionic, and radical species, which may be either oxidizing or reducing agents. Redox reactions involving ruthenium dissolved species into solution may take place.

Ruthenium can react with radiolytic compounds. The most numerous free radicals are $e^-_{(\text{aq})}$ and OH^\bullet ; H^\bullet is present to a lesser extent.⁶³

Hydrated electron is a powerful reducing agent, while OH^\bullet is a strong oxidizing agent. Rate constants for radical reactions are often high, $\sim 10^9$ to 10^{10} $\ell/\text{mol} \cdot \text{s}$, close to the diffusion limit.

The analyses of redox potentials of $\text{O}_2/\text{O}_2^{\bullet -}$ (-0.33 V) and $(\text{O}_2 + 2\text{H}^+)/\text{H}_2\text{O}_2$ (1 V) show that $\text{O}_2^{\bullet -}$ can be either an oxidizing or reducing agent. It is a weaker oxidizing agent than OH^\bullet but able to reduce metallic ions ($\text{HO}_2^{\bullet -}$, its conjugated acid, is stronger than $\text{O}_2^{\bullet -}$). In an acid medium when there is no reactant, $\text{O}_2^{\bullet -}$ reacts preferentially with $\text{HO}_2^{\bullet -}$ to produce hydrogen peroxide.

Generally, H_2O_2 is not very reactive, but its action should not be omitted; it is both an oxidizing and a reducing agent. Indeed, concerning iodine, for example, H_2O_2 is able to reduce I_2 into I^- . Finally, the hydrogen radical H^\bullet , which corresponds to conjugated acid of aqueous electron, is also a strong reducing agent.

For the ruthenium safety issue, our interest is limited to the potential role of an oxidizing agent in ruthenium behavior. The redox potential reported in Table VI (Refs. 64 and 65) shows us that the presence of hydroxyl radical OH^\bullet could stabilize $\text{RuO}_4(\text{aq})$ due to the very high redox potential of the $\text{OH}^\bullet/\text{H}_2\text{O}$ coupling, which is also true for the $\text{H}_2\text{O}_2/\text{H}_2\text{O}$ coupling.

So in severe-accident conditions, ruthenium tetroxide could be stable in the sump. Table VII is an update of Table V, taking into account the radiolytic conditions as well as the thermal conditions.

Haïssinsky and Dran⁶⁶ have performed ruthenate ions radiolysis experiments in sodium hydroxide aqueous solution. The radioactive source was ^{60}Co with an activity of 1700 Ci (maximum dose rate, 4.8 kGy/h). Sodium ruthenate was reduced into Ru-IV hydroxide precipitate. Nevertheless, perruthenate ions, RuO_4^- , are supposed to be formed transiently due to the following oxidation reaction:



Ruthenate reduction yield falls when its concentration rises, which is directly attributed to the reaction with radical-ions $\text{O}^{\bullet -}$. Ruthenate disappearance and Ru(IV) hydroxide formation increase in aerated solution and at low concentration in

TABLE VI
Redox Potential as a Function of pH, Calculated with a Reference Concentration of 10^{-10} mol/ ℓ in Ruthenium³³

Oxidizing/Reducing Agent	E° (V)	E° (V) - k pH (4)	E° (V) - k pH (10)
$\text{RuO}_4(\text{aq})/\text{RuO}_4^{2-}$	1.59	1.59	1.59
$\text{RuO}_4(\text{aq})/\text{RuO}_4^-$	1.00	1.00	1.00
$\text{RuO}_4^{2-}/\text{RuO}_2$	2.01	1.24	0.53
$\text{RuO}_4^-/\text{RuO}_2$	1.53	1.02	0.55
$\text{RuO}_4(\text{aq})/\text{RuO}_2$	1.40	1.02	0.65
$\text{O}_2/\text{H}_2\text{O}$	1.23	0.99	0.64
$\text{OH}^\bullet/\text{H}_2\text{O}$	2.74	2.50	2.15
$\text{H}_2\text{O}_2/\text{H}_2\text{O}$	1.78	1.54	1.18

TABLE VII
Stability Range of Ruthenium Oxides as a Function of pH
Containing Oxidizing Agents Formed by Water Radiolysis

	pH		
	4	7	10
Stable species	RuO ₂ (c) RuO ₄	RuO ₂ (c) RuO ₄ ²⁻ /RuO ₄ ⁻ RuO ₄	RuO ₄ ²⁻ RuO ₄ ⁻

Ru(VI) ($\sim 10^{-4} M$), but the reduction is not complete even for long irradiations (100 h, corresponding to 480 kGy).

No specific study of reactions between ruthenium compounds and hydrogen peroxide (H₂O₂) is available in literature.

V. INTERACTIONS BETWEEN RUTHENIUM OXIDES AND CONTAINMENT SURFACES

In a French 900-MW(electric) PWR, the internal wall of the reactor containment building is covered with a 6-mm-thick metal skin made of steel,⁶⁷ which ensures leaktightness. This metallic skin is painted to improve leaktightness and protect against corrosion but also for easier decontamination in case of incidents or accidents. The paint that is most often used in the containment is a water-based polyamide epoxy type. Its chemical structure was established in Aujollet's doctoral thesis.⁶⁸ In a 1300- to 1450-MW(electric) PWR, there is no metallic skin, but the concrete wall is also painted in the same way.

Inside the containment, many other elements are made of steel but not painted, e.g., the steam generators and all the hoopings that maintain the heat insulator. Most of these surfaces are made of Types 316L and 304L stainless steel.²³

V.A. Reactivity of RuO₄ with Surfaces in the Gaseous Phase

No experimental study of the interaction between gaseous ruthenium tetroxide and paints (epoxy) has been carried out yet. However, it is well known that RuO₄(g) is very reactive with organic compounds.²⁵⁻²⁷ Tetroxide is used in organic chemistry as an oxidizing agent. Table VIII lists the various organic

TABLE VIII
Organic Compounds Able to Be Oxidized by RuO₄

Substrates	Oxidation Products
Primary alcohols	Aldehydes, carboxylic acids
Secondary alcohols	Ketones
Aldehydes	Carboxylic acids
Alkenes	Aldehydes, carboxylic acids
Alkynes	α -diketones, carboxylic acids
Amides	Imides
Ethers	Esters
Sulphides	Sulphones

compounds that can be oxidized by RuO₄. A strong reactivity of tetroxide vapors with the painted surfaces of the containment can be expected.

On the other hand, ruthenium tetroxide affinity for iron oxides has already been studied.^{52,69} Besides, ruthenium was also found on structure material particles (iron and others) after the Chernobyl accident. Kashparov et al.⁷⁰ think that the origin of these particles is linked to the following process: First, ruthenium seems to have been oxidized on the fuel particle surface; second, RuO₄ seems to have been vaporized, and these vapors seem to have condensed onto aerosol particles formed by structural materials.

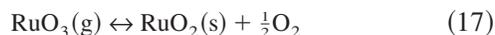
Maas and Longo⁵² mentioned that ruthenium trapping by iron oxides is very efficient, at $\sim 99.9\%$. The following deposition mechanism was proposed: RuO₄(g) \rightarrow RuO₂(s) + O₂(g) [Eq. (1)].

The large specific surface of iron oxides and the variability of iron oxidation states are supposed to promote the reduction of RuO₄ onto the surface. Even if RuO₂ is often considered as fixed irreversibly by iron oxides, thereby avoiding any later revolatilization, the deposition mechanism is not really understood. No existence of mixed ruthenium/iron oxides has yet been mentioned.

Holdoway⁶⁹ performed various tests in order to study ruthenium volatilization and RuO₂ deposition phenomena. One of them consisted of heating ruthenium dioxide that had been previously activated [¹⁰³RuO₂(s)] up to 1323 K in an O₂ stream and then following the vaporized ruthenium dioxide progressively deposited along the stainless steel thermal-gradient tube located downstream of the furnace.

The measured deposits are distributed as shown in Table IX. The results obtained clearly indicate the existence of a volatile ruthenium compound, even at low temperature, contrary to thermodynamic predictions. The most probable candidate is ruthenium tetroxide.

Deposition reactions considered by Holdoway are the following:



and

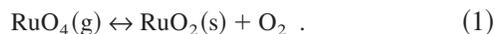


TABLE IX
Test Results on ¹⁰³RuO₂ Deposits on
Stainless Steel Surfaces⁶⁹

Sample	Temperature (K)	¹⁰³ RuO ₂ Deposited (gamma-count/min)
1	323	1 139
2	324	1 474
3	328	1 625
4	332	2 063
5	339	2 695
6	353	2 936
7	277	4 843
8	413	16 092
9	465	23 068

Apparently, ruthenium deposits on steel are not uniform, because deposits take place primarily on already deposited RuO₂ and not on still available surface sites. The presence of RuO₂ would then catalyze RuO₄(g) condensation. Moreover, this self-catalytic reduction phenomenon is also mentioned by various authors.^{24,35,52}

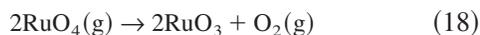
Ortins de Bettencourt and Jouan²⁴ have observed that the nature of the ruthenium deposits varies as a function of the temperature deposition. At ~773 K, ruthenium is deposited in an irreversible way, but at 423 K, the crystals obtained disappear completely when the tube is rinsed. The latter are assumed to be hydrated RuO₂ or Ru₂O₅, whose existence has already been mentioned by Debray and Joly.⁴²

Eichler et al.²⁹ also studied adsorption of ruthenium tetroxide onto stainless steel substrates. During thermochromatography tests carried out between 1500 and 300 K, a large proportion of the ruthenium was volatilized under an oxygen stream, and some of it settled in the column at ~800 K; a weak part, probably in RuO₄ form, was able to reach the active charcoal trap set at the outlet at room temperature. When quartz powder was added, ruthenium volatility was strongly reduced, and the fraction that reached the charcoal trap was drastically reduced, near zero. Furthermore, a second deposit area, at ~400 K, occurred. The authors interpreted the deposit phenomena with the same reactions as Holdoway [Eqs. (17) and (1)]. The RuO₂(s) is then in equilibrium either with RuO₃ or with RuO₄, according to temperature and oxygen partial pressure, as Schäfer et al.³⁰ and Bell and Tagami²¹ have already assumed. Finally, Eichler et al.²⁹ attributed the low-temperature deposit area at ~400 K to the formation of RuO₃(s) in non-equilibrium conditions, even if its existence has not been established. The RuO₃(g) would no longer be in equilibrium with RuO₂(s), as expected at high temperatures, but gaseous ruthenium trioxide would condense directly.

Another possible adsorption reaction at quite a low temperature can be RuO₄(g) → RuO₃(ads) + ½O₂, whose enthalpy of dissociation reaction has been calculated as -19.3 ± 11 kJ/mol.

The existence of ruthenium trioxide in solid form was also mentioned by Kim and Winograd,⁷¹ who provide its X-ray photoelectron spectroscopy (XPS) spectrum with binding energy values for Ru 3d5/2 and O1s. They found a surface layer corresponding to RuO₃ on anhydrous RuO₂(s).

As mentioned previously, Zimmerman et al.³⁵ focused their studies on RuO₄ photochemical decomposition and the resulting products formed. A thin brown layer of RuO₃(s) for irradiating photons with a 438.5-nm wavelength was observed. According to the authors, the only energetically possible elementary process would be



in which RuO₃ could be either gaseous ($\Delta G_{298}^\circ = +197$ kJ/mol and $\Delta H_{298}^\circ = +264$ kJ/mol) or solid.

Sakurai et al.⁷² studied RuO₄(g) interactions with various surfaces, including stainless steels, at room temperature and low pressure. Contrary to the already quoted hypothesis concerning RuO₄ decomposition into (RuO₂ + O₂), they were not able to obtain any experimental evidence about dioxide formation, even if the black color of the deposit is consistent with RuO₂. First, no oxygen release was experimentally measured. Second, the spectrum from X-ray diffraction did not corre-

spond to that of RuO₂ and was different from the well-characterized spectra of the oxides of the different metals concerned. Third, they even observed ruthenium deposits on gold, a priori well known to be nonoxidizable. These facts meant they could refute the assumption of metal oxidation by ruthenium tetroxide, reduced to RuO₂, which led to formation of metal oxides. To support this assumption, no presence of ruthenium mixed oxides with atoms of the various substratum involved was detected by X-ray diffraction (XRD) and XPS analyses. No atom of the metal substratum on the deposit surface was also observed, not even in the subsurface down to a depth of ~20 Å. Moreover, temperatures of ~1273 K are generally necessary to form mixed oxides,⁷³ but these tests were carried out at room temperature.

Hydrolysis of this deposit turned out to be efficient and led to hydrogen peroxide release, measured with spectrophotometric and iodometric methods. Some of the hydrolyzed compounds were able to cover the initial ruthenium black deposit and protect it from any later hydrolysis. The oxidation state of the ruthenium deposit was determined to +IV by performing XPS and infrared analysis. According to all these elements, it was concluded that the deposit was not RuO₂ but a ruthenium tetroxide form modified being in contact with the metal. In fact, a RuO₄ molecule is supposed to be linked to another one via peroxide Ru-O-O-Ru bonds. In that case, the black deposit observed would consist of (RuO₄)_n polymerized with peroxide bonds.

The hydrolysis reaction may correspond to ruthenium hydroxide formation:



Using XRD spectral analysis, Cains and Barnes⁷⁴ have also characterized ruthenium deposits formed from RuO₄(g), at temperatures of ~423 K and above. Ruthenium dioxide in crystal form was determined without any bonds with ferrous metal. The XPS analysis of Ru 3p and 3d levels confirmed it.

Supposing ruthenium tetroxide can be adsorbed easily on iron substrates, the information published in literature is not consistent concerning the adsorption mechanism involved and the deposited products resulting from the ruthenium-iron oxide interaction. If a pseudoconsensus about the ruthenium dioxide presence in the deposits exist, others species might be expected.

V.B. Reactivity of Dissolved Species with Immersed Surfaces

Immersed surfaces in sump contain substrates that are potentially reactive toward ruthenium species dissolved in this aqueous phase. Nevertheless, the total surface is quite small compared with total surface in the atmosphere. Most of the immersed steel or concrete surfaces are painted, while a small section of steel surface is not painted. No data were found in the literature on paint/dissolved ruthenium interactions, but reactivity to stainless steel has already been broached.

Carlson⁷⁵ studied the deposit of dissolved ruthenium onto stainless steel-etched plates for 6 h in buffer solutions at pH 4, 7, and 9 at 333 K. The pH increases ruthenium adsorption on stainless steel. Whatever the buffer employed, the fraction deposited after 24 h (or 168 h) is always low, the maximum is reached at pH 9, with ~10%, while it is only 1 and 0.5% at pH 7 and 4, respectively. At pH 9, equilibrium is not reached after 24 h; the kinetics of adsorption is slow. The temperature

effect is significant; indeed, the deposited ruthenium fraction gradually rises with temperature over the 6-h test. Ruthenium deposit did not appear homogeneous on sample plates; it depends strongly on the nature of the metal surface and on the presence, or not, of available adsorption sites. Finally, tests for redissolution were not conclusive, the desorbed fraction is very weak, and ruthenium is probably chemisorbed.

Belloni et al.⁷⁶ carried out experiments on the adsorption of ruthenium on different surfaces, such as platinum, glass, silver, etc. Unlike some other fission products, ruthenium adsorption does not follow a Langmuir isotherm, but the kinetics obeys Henry's law (linear isotherm). Moreover, ruthenium adsorption increases with temperature^{75,76} and is barely reversible.

Finally, we point to the existence of an American patent⁷⁷ that reveals inhibition of stainless steel corrosion when there is ruthenium in acid solution in contact with steel surfaces. This may be due to surface covering with RuO₂, which would then have a covering property and such adsorption on steel would prevent any corrosion phenomenon. This mechanism is very probable if a parallel is drawn between ruthenium and technetium. Indeed, chemical properties of RuO₂ are close to those of TcO₂, but pertechnetate ions, TcO₄⁻, even in very low concentration, are well known³³ to have electrolytic iron and mild steel corrosion inhibitive properties due to the formation of an insoluble TcO₂ film on their surface.

To sum up, ruthenium species are expected to be reactive on immersed surfaces with steel or with paint, and the trapping seems to be almost irreversible.

VI. CONCLUSION

Basically, the ruthenium chemistry is very complex and all the more so in severe-accident conditions where radiolytic phenomena are involved.

Ruthenium tetroxide is the most volatile ruthenium oxide in the conditions prevailing in the containment, with potentially large consequences on the source term, so understanding its behavior is important, because for accidents involving highly oxidizing conditions such as air ingress, its presence in the containment reactor is suspected.

From a thermodynamic point of view, ruthenium tetroxide is not stable at ~373 K, but kinetic effect seems to occur and could allow its existence for a while, promoted by the presence of air radiolysis products in the atmosphere resulting in rather oxidizing conditions. Experimental data tend to prove that RuO₄ remains stable at room temperature for a certain period of time.

Even if there are some discrepancies in literature concerning the adsorption mechanisms involved and the deposited products resulting from the ruthenium–stainless steel surfaces interaction, it is obvious that tetroxide is also very reactive to steel and paints. This is true for immersed surfaces as well.

In the sump, the oxidizing agents formed by water radiolysis could oxidize ruthenate and perruthenate ions in ruthenium tetroxide.

Concerning ruthenium dioxide, which is the main compound of ruthenium expected to reach the containment in aerosol form, it is generally not soluble and it will be distributed in the sump after settling or will be deposited onto walls in the atmosphere. Nevertheless, considering redox potential, it can be expected that a fraction of the ruthenium dioxide deposits

could be oxidized by ozone, which is a product of air radiolysis. Unfortunately, no data were found in literature concerning this potential reaction, which could greatly impact the ruthenium source term.

For all the aforementioned reasons, it is of primary importance to obtain experimental data on ruthenium behavior in severe-accident containment conditions, and more precisely on ruthenium tetroxide stability in the gaseous phase, and on possible formation of this oxide from ruthenium deposits or from ruthenate and perruthenate anions.

In the near future, experiments will be carried out to explore this field to better quantify the ruthenium source term and obtain data on kinetics concerning all these potentially relevant reactions.

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