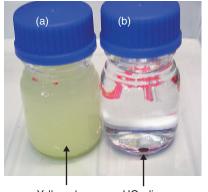
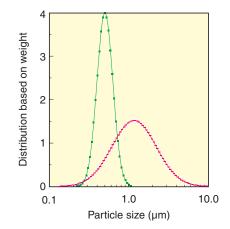
1–13 How do Microscopic Particles of Debris Change in Water? — To Understand the Chemical Effects of Hydrogen Peroxide and Boric Acid —



Yellow slurry UO₂ disc

Fig.1-29 Yellow slurry of UO₄·4H₂O and a UO₂ disc in water

(a) Yellow slurry of UO₄·4H₂O given by the immersion of a UO₂ disc in 10wt% H_2O_2 aqueous solution, (b) UO₂ disc immersed in pure water over a period corresponding to (a). UO₂ discs as well as powders are fully oxidized and pulverized into microscopic particles by H_2O_2 .



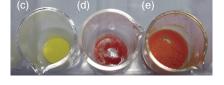


Fig.1-30 Particle size distribution of UO_{4} ·4H₂O

particle size distribution of the UO_{4} ·4H₂O given by the immersion of UO_2 in 10wt% H₂O₂ aqueous solution (pH 5.6), the distribution after hydrochloric acid (HCl) has been added to the slurry (pH 2.8). As the acidity of the aqueous solution is stronger, the particles are flocculated; consequently, the particle size becomes larger. This property means that the slurry is mesocolloidal. Fig.1-31 Uranium trioxide hydrate (c) Uranium trioxide dihydrate (UO₃·2H₂O), (d) uranium trioxide monohydrate (UO₃·H₂O), and (e) UO₃. UO₃·2H₂O are formed by the moistureabsorption of UO₃; in contrast, a perfect dehydration of UO₃·2H₂O cannot be accomplished by heating at 300 °C. Under this condition, UO₃·H₂O is formed.

During the accident at the TEPCO's Fukushima Daiichi NPS, the nuclear fuels melted together with the structural materials of the nuclear reactors and became debris. It is supposed that a portion of the debris may have dropped into the cooling water. In such a quenching condition, the debris could be pulverized into small particles. In addition, the particle size would become smaller by oxidation. Small powders of the debris will also be produced by cutting under the removal operations of the debris from the nuclear reactors. However, the chemical behaviors of the small particles of the debris in water are not understood.

Therefore, to investigate the chemical change and pulverization of debris immersed in cooling water for a long time, uranium dioxide (UO₂), a component of the debris, was immersed in pure water as well as sodium chloride (NaCl), hydrogen peroxide (H₂O₂), and boric acid (H₃BO₃) aqueous solutions for a month at 50 °C. As shown in Fig.1-29, UO₂ reacted perfectly with H₂O₂; consequently, yellow particles of uranium peroxide tetrahydrate (UO₄·4H₂O) were formed. It was found that the average particle size was smaller than 1 μ m, as shown in Fig.1-30, and that the slurry showed mesocolloidal properties. In contrast, UO₂ did not show notable chemical changes in the other aqueous solutions.

The chemical bonding of the UO₂ and H₃BO₃ was investigated by infrared spectroscopy. From the result, H₃BO₃ was predicted not to accompany the debris under removal operations from the H₃BO₃ aqueous solution. The influence of heating temperature and humidity on the hydration number of uranium peroxide hydrates was investigated. It was found that although UO₄·4H₂O changed to orange anhydrous uranium trioxide (UO₃) under heating at 300 °C, UO₃ absorbed the moisture and easily suffered the hydration. Please see Fig.1-31.

This topic includes a part of the results of "Establishment of basic technology for decommissioning and safety of nuclear reactors for power generation" in FY 2013 that was entrusted to IRID from METI and of the subsidy program "Project of Decommissioning and Contaminated Water Management" in the FY 2013 supplementary budget and conducted by JAEA as a member of IRID.

Reference

Otobe, H. et al., Corrosion of Uranium and Plutonium Dioxides in Aqueous Solutions, Proceedings of 2014 Nuclear Plant Chemistry Conference (NPC 2014), Sapporo, Japan, 2014, paper 10173, 11p., in USB Flash Drive.