

## The Effect of $U_3O_8$ Additon on the Sinterability of CANDU-Type $UO_2$ Pellet

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### 1. Introduction

A  $UO_2$  scrap rate, in general, is approximately 5% in the manufacturing process of CANDU  $UO_2$  pellets. The scrap is oxidized into  $U_3O_8$  powder under an air atmosphere at constant temperature for recycling.

If pellets, which consist of  $UO_2$  and  $U_3O_8$ , are sintered in a  $H_2$  atmosphere, the component  $U_3O_8$  acts as the pore, and the sintered density of the pellet decreases. In other words,  $U_3O_8$  is a density controller as well as a pore-former.

In this study, the effects of an additional amount of  $U_3O_8$  powder, formed according to the oxidation temperatures, on the sinterability of CANDU-type  $UO_2$  pellet were investigated.

### 2. Methods and Results

CANDU-type pellets (97%TD,  $\sim 8\mu m$ ) were oxidized with 3 different temperature conditions (350, 450 and 550 °C) for 4 hours under an air atmosphere in a product-scale oxidation furnace. The results of XRD analysis showed that all oxidized powders formed according to the oxidation temperature were confirmed as  $U_3O_8$ . (Fig. 1)

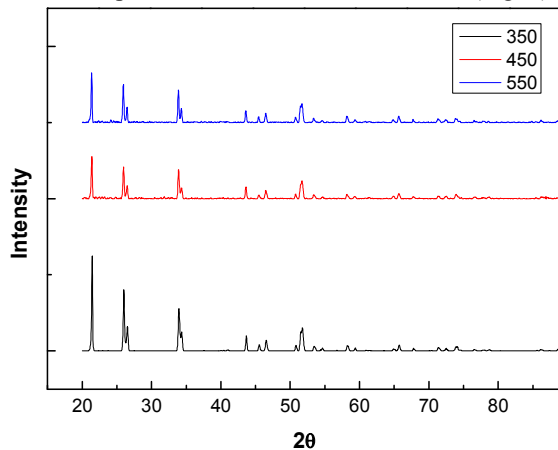


Fig. 1. XRD pattern according to the oxidation temperature

Fig. 2 shows the specific surface area ( $m^2/g$ ) of  $U_3O_8$  powder formed according the oxidation temperature. As shown in Fig. 2, the specific surface area of  $U_3O_8$  powder decreases as the oxidation temperature increases. If the

oxidation temperature is increasing, the strength of grain boundary of  $UO_2$  scrap pellet is weaker than that of the grain. Accordingly, the oxygen diffuses rapidly along the grain boundary. The 2-3 crystal grains become one particle during high oxidation temperature. Therefore, the oxidation temperature increases, the particle size of the  $U_3O_8$  powder increases. The specific surface area of the powder then decreases. [1]

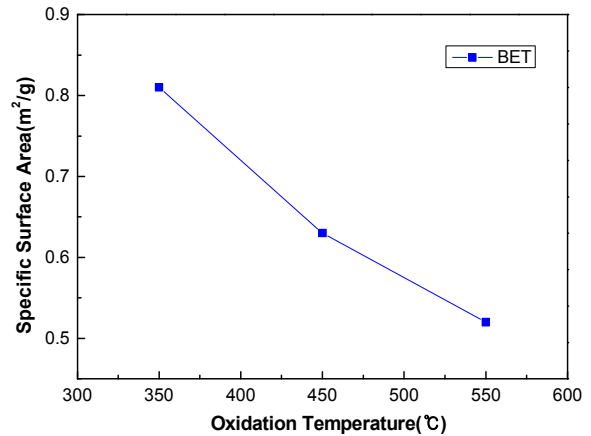


Fig. 2. Specific surface area of  $U_3O_8$  powder according to the oxidation temperature

$U_3O_8$  powder, which is formed at each oxidation temperature, was added to ex-ADU natural  $UO_2$  powder as 5 conditions (0, 2.5, 5, 7.5 and 10wt%). Next, these powders were mixed for 30 min. in a tubular mixer with 0.19wt% Acrawax for a lubricant.

The mixed powders were compacted with the density of 5.67g/cc, such as a density of product-scale. The green pellets were sintered at 1610 °C for 6 hours in a  $H_2$  atmosphere. The sintered density of the pellet was measured by an immersion method.

Fig. 3 shows the sintered density according to an addition amount of  $U_3O_8$  powder that formed at the each oxidation temperature.

As shown in Fig. 3, the sintered density of  $UO_2$  pellet decreases with increase in the additional amount of  $U_3O_8$  powder. However,  $U_3O_8$  powder formed at the lower oxidation temperature is added into the  $UO_2$  powder under the same addition, the drop width of the sintered density of the pellets appeared to be small.

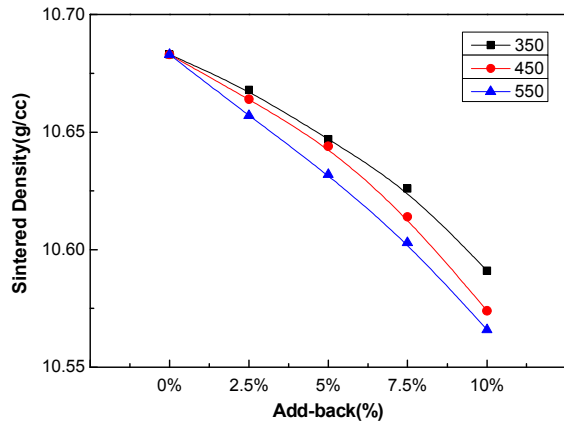


Fig. 3. Sintered density according to addition amount of  $U_3O_8$

Fig. 4 shows the relationship between specific surface area of  $U_3O_8$  powder and sintered density drop with a 1wt% addition of  $U_3O_8$  powder.

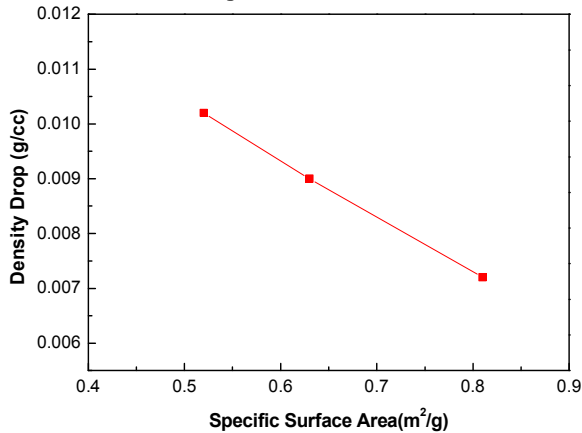


Fig. 4. Density drop according to the specific surface area of  $U_3O_8$  powder

As shown in Fig. 4, the density drop decreases with an increase in the specific surface area of  $U_3O_8$  powder.

It is thought that the specific surface area of  $U_3O_8$  is an important factor to control the sintered density of the pellet for recycle of the  $UO_2$  scrap. [2]

### 3. Conclusion

- The result of XRD analysis showed that oxidized powder that was oxidized at 350, 450 and 550 °C, was confirmed as  $U_3O_8$ .
- Increasing of the additional amount of  $U_3O_8$  powder causes decreasing of the sintered density of  $UO_2$  pellet.
- Decreasing of the oxidation temperature of  $U_3O_8$  powder causes increasing of specific surface area of  $U_3O_8$  powder and decreasing of the density drop of  $UO_2$  pellet.

### References

- [1] S.H. Na et al., Proc. of the Korean Radioactive Waste Society, Autumn 2011, Jeju, 9(2) 2011, pp.163-164
- [2] C.M. Yang et al., Transactions of the Korean Nuclear Society Spring Meeting, Jeju, Korea, May 17-18, 2012