§7. Corrosion Characteristic of AlN, Y<sub>2</sub>O<sub>3</sub>, Er<sub>2</sub>O<sub>3</sub> and Al<sub>2</sub>O<sub>3</sub> in Flinak for Molten Salt Blanket System

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Regarding the heat exchanger for liquid breeder blanket system using molten salts, tritium barrier and anticorrosion barrier play important roles. To develop the heat exchanger, it is essential to optimize chemical stability of the barrier material against molten salts. In the present study, the corrosion of AlN,  $Y_2O_3$   $Er_2O_3$  and  $Al_2O_3$  in Flinak was examined to consider possibility as barrier materials.

The static corrosion tests on these ceramic specimens were performed in high purity Flinak (46.5LiF-11.5NaF-42KF mol%) for 230 and 1010 hours. The specimens and Flinak were heated in SS316L crucibles at 600°C in Ar atmosphere. Weight change by the corrosion test was measured. Surface analysis of the specimens was carried out by SEM and XPS. Elution of elements from the specimens was determined based on chemical analysis of the Flinak after the corrosion test using ICP. With the corrosion test, the compatibility of the ceramics in molten Flinak was also considered based on the standard Gibbs energy for formation. According to the estimation as shown table 1, AlN is expected to be stable in Flinak. On the other hand,  $Er_2O_3$ ,  $Y_2O_3$  and  $Al_2O_3$  are expected to be unstable.

Figure 1 shows SEM images and weigh change before and after the corrosion test. While AlN rarely changed in structure and weight, the structure of  $Er_2O_3$ ,  $Y_2O_3$  and  $Al_2O_3$  drastically changed in weigh and structure through the corrosion test.

The rough structure at the surface of the AlN specimen before the corrosion test had changed to slightly smooth after the corrosion test. According to surface analysis by XPS measurement, substitution from nitrogen to oxygen near surface was clarified after the corrosion test. This means that  $Al_2O_3$  was formed at the surface. In Flinak after the corrosion tests, Fe and Cr from the SS316L crucibles were detected. When Al was dissolved into molten Flinak in the corrosion process, Fe and Cr would be oxidized as a redox reaction.

The weight loss in the  $Er_2O_3$  specimens after corrosion test was large with breaking. It includes loss of fragments due to breaking during the corrosion test and Flinak removing process by water rinse. The initial  $Er_2O_3$ surface consisted of round shape grains. After the corrosion test, the grains became rough and formed porous and interstitial structure. According to XPS measurement, fluorine was detected on the specimen surface after the corrosion test. Er dissolution was also detected in the Flinak. These suggest that the surface of  $Er_2O_3$  specimens was corroded via the fluoridation process.

The initial  $Y_2O_3$  specimens consisted of large grains appeared over several micrometers. After the corrosion test, the surface became more porous and interstitial structure than that before the corrosion test. Although the  $Y_2O_3$  specimen gained in weight after 230 hour corrosion test, this weight gain would be due to residual Flinak and water remained in the pores of the specimens. The fluorine was also detected on the surface. These suggest that the surface could be corroded via fluoridation process.

Although initial  $Al_2O_3$  consisted of grains like gathering pine cones, after corrosion test, the structure was drastically changed to be rough structure. In the  $Al_2O_3$ specimens after corrosion test, fluorine, lithium, sodium and potassium were detected. The weight increased after the corrosion test. The weight gain of the  $Al_2O_3$  specimens would also be due to retaining Flinak and/or water in the porous structure.

These results mean that  $Er_2O_3$ ,  $Y_2O_3$  and  $Al_2O_3$ dissolved into Flinak and that the corrosion is not negligible and that the AlN specimens seem to possess corrosion resistance against Flinak. For details, the slight weight loss was detected. Reaction of AlN with moisture forms Al(OH)<sub>3</sub> and ammonia.<sup>1)</sup> And the hydroxide could change to Al<sub>2</sub>O<sub>3</sub> and/or its hydrate. After the corrosion test, the substitution of nitrogen to oxygen was detected at the surface. The oxygen rich layer such Al(OH)<sub>3</sub> or Al<sub>2</sub>O<sub>3</sub> at the specimen surface would promote the corrosion behavior.

In conclusion, it was found that these corrosion behaviors agreed with the prediction based on thermodynamic evaluation. AlN indicates compatibility in molten Flinak. This suggests that AlN is suitable for surface coating of the structural materials for blanket system using molten salt like a Flinak.<sup>2</sup>

Table 1 Gibbs energy for reaction of ceramics material in Flinak at  $600^{\circ}$ C

		Possible R	eaction	∆ Gr [J/K.mo	ol] Thermodynami Stability	с	
AIN	P1 11.0	$AIN + 3LiF = AIF_3 + Li_3N$		+2.015	Stable		
	Fluoridation	$A1N + 3KF = A1F_3$	$+3K + 1/2N_2$	-148.607	Unstable		
		$A1N + 3H_2O =$		-318.676	Unstable	Unutuhla	
	Oxidation	NH3 + 1/2Al2O3.3(H2O)		(-325.219 at 27	°C) Unstable	Unstable	
		AIN + 3/4O <sub>2</sub> - 1/2Al <sub>2</sub> O <sub>3</sub> + 1/2N <sub>2</sub>		-51.351	Unstable	Unstable	
$Al_2O_3$		$Al_2O_3+6LiF = 2AlF_3+3Li_2O$		-23.293	Unstable		
		$Al_2O_3+6KF - 2AlF_3+3K_2O$		-18.280	Unstable		
Y <sub>2</sub> O <sub>3</sub>		$Y_2O_3 + 6LiF - 2YF_3 + 3Li_2O$		-2.227	Unstable	Unstable	
		$\mathbf{Y}_2\mathbf{O}_3 + \mathbf{6KF} = \mathbf{2YF}_3 + \mathbf{3K}_2\mathbf{O}$		+2.786 at 597	°C Stable over 527	°C	
	Fluoridation			negative under :	527°C Unstable under 52	7°€	
Er <sub>2</sub> O <sub>3</sub> ZrO <sub>4</sub>		$Er_2O_3 + 6LiF = 2ErF_3 + 3Li_2O$		-24.365	Unstable		
		$Er_2O_3 + 6KF = 2ErF_3 + 3K_2O$		-19.355	Unstable		
		$ZrO_4+4LiF = ZrF_4 + 2Li_2O$		-16.484	Unstable		
$210_4$		ZrO <sub>4</sub> +4KF – Z	$rF_4 + 2K_2O$	-13.139	Unstable		
Corrosion test 230h initial	AIN Rough surf		5 m	Y <sub>2</sub> O <sub>3</sub> ugh porcus ructure	Al <sub>2</sub> O <sub>3</sub> Pine cone like grains	5	
	-0.233g/	m <sup>2</sup> -44	ig/m²	+195g/m <sup>2</sup>	+981g/m <sup>2</sup>		
HOTOL	-0.0921 Smooth st	-	g/m² structure Ra	-58.9g/m <sup>2</sup>	+74.0g/m <sup>2</sup>		
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					5000		

Fig. 1 SEM images and weight change by corrosion test.

1) Krnel, K. et al.: J. Euro. Ceram. Soc., **21** (2001) 2075

2) Watanabe, T. et al.: J. Plasma Physics and Research, in press