

## Fabrication and characterization of $UAl_x$ intermetallic compounds

Ji Min Nam, Moon Soo Sim, Ho Jin Ryu\*, Se Jung Jang, Jong Man Park  
Korea Atomic Energy Research Institute: 1045 Daedeok-daero, Yuseong-gu, Daejeon 305-353, Republic of Korea  
\*Corresponding author: hjryu@kaeri.re.kr

### 1. Introduction

Currently, uranium aluminum alloys have been used as dispersion fuel in research reactors [1] and U-Al dispersion targets for  $^{99}Mo$  medical radioisotope production [2]. One of the conventional manufacturing processes of the U-Al dispersion fuels and targets is the grinding and crushing of cast  $UAl_2$  ingot by mechanical methods. Also, produced powder was mixed with Al [3]. However, it is complicated and inefficient to fabricate U-Al. Therefore, KAERI has produced U-Al powder with varying Al content using a centrifugal atomization method [4].

In this study, U-Al alloy and  $UAl_x$  intermetallic compound powders were produced by a centrifugal atomization method. The atomized powders were characterized by X-ray diffraction, SEM, EDX, and density measurements.

### 2. Experimental procedures

U-1wt%Al, U-10wt%Al and U-20wt%Al alloy powders were fabricated in this experiment. Superheated molten uranium-aluminum was fed through a small nozzle onto a graphite disk spinning at about 30,000 rpm, and liquid alloy droplets were then spread from the disk by a centrifugal force and cooled in an argon atmosphere. Fig. 1 shows the fabrication steps for the atomized U-Al powders. This method has advantages such as a single-step process, short processing time, a high production yield rate, high purity with less defects, and spherical formation for easy dispersion plate fabrication.

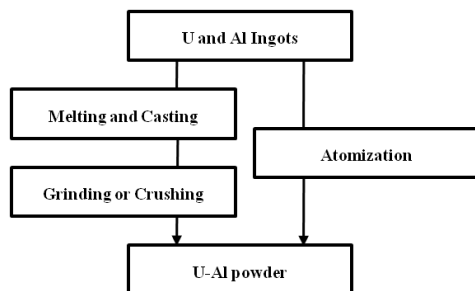


Fig. 1. Step for the fabrication of powders.

The microstructures of U-Al powders were observed by scanning electron microscopy (SEM). X-ray diffraction and Energy dispersive x-ray spectroscopy (EDX) experiments were performed to characterize the  $UAl_x$ .

### 3. Results and discussion

The differences in powder microstructures in accordance with aluminum content were compared, as shown in Fig. 2(a-c). Typical solidification microstructures were developed in rapidly solidified spherical powders. Fig. 2(a) shows the spherical morphology of the atomized U-1wt%Al particles and a cross-section image. The fine grain structure is evident. The boundaries of U and  $UAl_2$  were observed in U-10wt%Al, as shown in Fig. 2(b). Fig. 2(c) shows the equiaxed zone starts from one part of a liquid droplet, and then a columnar zone develops along the thermal flow direction. Columnar dendrite growth of  $UAl_2$  was observed in U-20wt%Al.

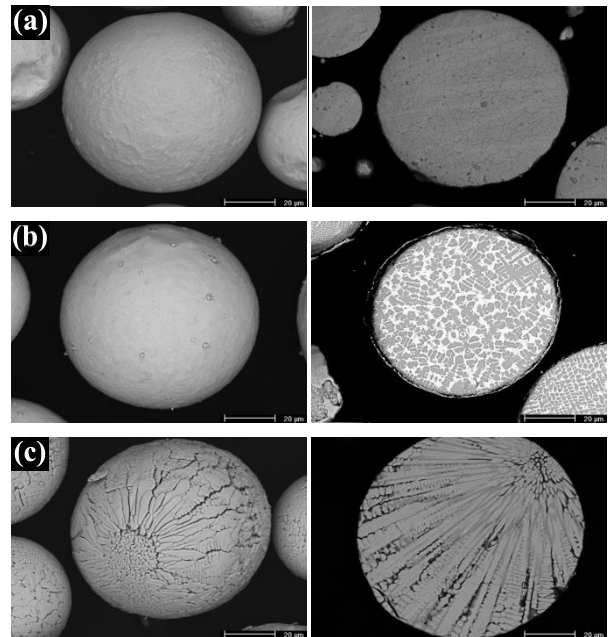
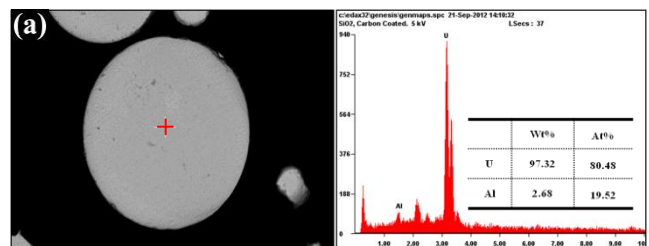


Fig. 2. Spherical morphologies and cross-section SEM images of atomized particles : (a) U-1wt%Al, (b) U-10wt%Al, (c) U-20wt%Al

EDX analyses, as in Fig. 3, show that U-10wt%Al is composed of uranium and  $UAl_2$ , and U-20wt%Al consists of  $UAl_2$  and  $UAl_3$ .



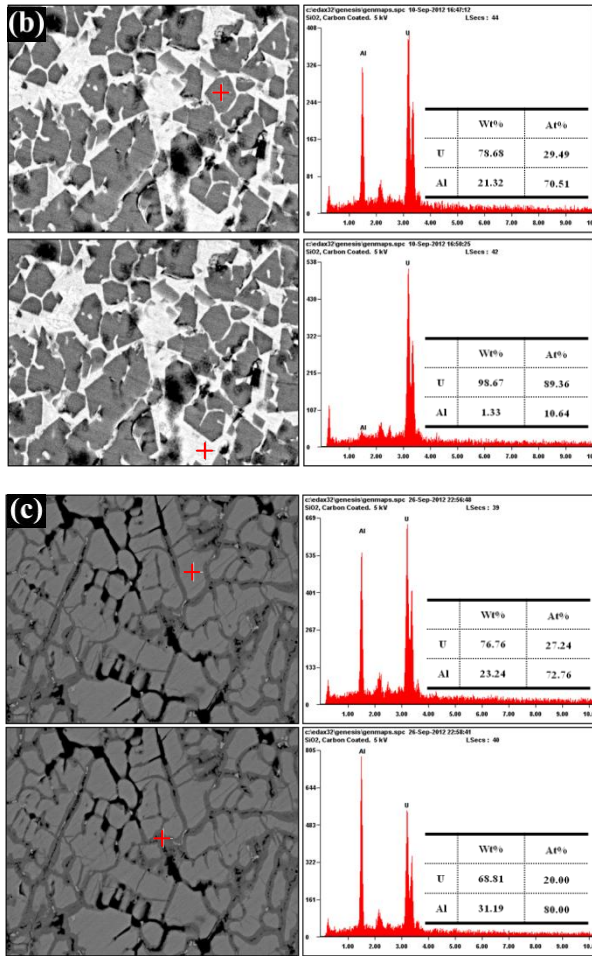


Fig. 3. Cross-section SEM images and EDX spectra of atomized particles : (a) U-1wt%Al, (b) U-10wt%Al, (c) U-20wt%Al

XRD patterns of the particles are shown in Fig. 4. It was observed that the compounds of U-Al were composed of  $UAl_2$  and  $UAl_3$  in U-10wt%Al and U-20wt%Al. However, a  $UAl_3$  pattern could not be found in U-1wt%Al.

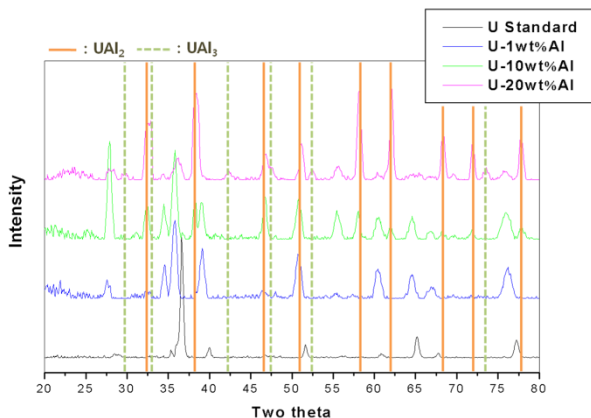


Fig. 4. X-ray diffraction patterns of U and U-Al powders.

#### 4. Conclusions

1. Atomized U-Al powders with compositions of U-1wt%Al, U-10wt%Al, U-20wt%Al were fabricated.
2. XRD analyses identified  $UAl_2$  and  $UAl_3$  intermetallic compounds formed in the atomized particles.

#### ACKNOWLEDGMENTS

This study was carried out as a National Nuclear R&D Program sponsored by Ministry of Education, Science and Technology.

#### REFERENCES

- [1] Farhan Muhammad, et al., Annals of Nuclear Energy 36 (2009) 998.
- [2] Nuclear Energy Agency, The supply of medical radioisotopes (2010).
- [3] Kanwar Liaqat Ali, et al., Nuclear Engineering and Design 255 (2013) 77.
- [4] Chang Kyu Kim, et al., Nuclear Engineering and Technology 39 (2007) 617.