

## Aluminum analysis in biological standardized reference materials

Y Katoh, T Sato<sup>1)</sup>, Y Yamamoto<sup>2)</sup>, Y Gotoh and K Yamamoto

Faculty of Health Sciences, School of Radiology, Tokyo Metropolitan University 7-2-10, Higashi-Ogu, Arakawa-ku, Tokyo 116-8551, Japan

\*1) Tokyo Metropolitan Institute Neuroscience  
2-6, Musashidai, Fuchu-shi, Tokyo 183-8526, Japan

\*2) Department of Legal Medicine, School of Medicine, Shiga University of Medical Science  
Seta-tsukiwa-cyo, Otsu-chi, Siga 520-2192, Japan

### Abstract

We have investigated the multi-elemental abundances in biological materials by Instrumental Neutron Activation Analysis (INAA). The application of INAA for the Al determination has some problems: 1) the biological material includes P (several  $10^3$  to a few  $10^4$  ppm) and Si (several  $10^2$  to a few  $10^3$  ppm). 2) P and Si interfere with the Al determination due to  $^{31}\text{P}(n,\alpha)^{28}\text{Al}$  and  $^{28}\text{Si}(n,p)^{28}\text{Al}$  reactions.

In this study, P in biological standardized reference materials (SRMs) was determined by Liquid Scintillation Counter as a nuclear method. After being irradiated, the SRMs and comparative standards were treated using the ordinary nitric acid method. The samples were measured recurrently for a long-term. The determination of  $^{35}\text{S}$  count rate in samples must be done post-160 day post-irradiation. Subtracting the decay-corrected  $^{35}\text{S}$  count rate of post-160 day irradiation from that of the post-30 day post-irradiation,  $^{32}\text{P}$  count rate in sample is capable of determining.