AMS ¹⁴C CHRONOLOGICAL STUDY ON HOLOCENE ACTIVITIES OF ACTIVE FAULTS IN JAPAN

T. NAKAMURA,¹ M. OKAMURA,² K. SHIMAZAKI,³ T. NAKATA,⁴ N. CHIDA,⁵ Y. SUZUKI,⁶ M. OKUNO⁷ and A. IKEDA¹

Slipping of active faults generates earthquakes. If we can detect traces of ancient slips of active faults, we can correlate them with earthquakes that occurred in the past. The scale of a slip correlates with the time interval between the slip and the previous one during which strain accumulated in the fault system. Thus it is very important to examine the history of fault activities in the late Quaternary, to provide a risk estimation of the next slip time for prevention of disasters. To establish chronology of past slips of active faults, AMS ¹⁴C dating of macro fossil, charcoal and soil organic material samples collected around the fault is very promising, because the method requires carbon samples of very small size.

Some active faults were discovered in a shallow sea area. They provided us an ideal field to study behavior of a fault during the Holocene, because sea sediments deposited continuously, by recording seismic events of the faults in them. The following procedures are applied to two submarine active fault systems located in the northwestern part of Beppu Bay, eastern Kyushu and off the coast of the town of Futami in the northwestern Shikoku, Japan: 1) surveying an exact line of a submarine fault and continuation of sediment layers on both sides of it by using a high-resolution single-channel seismic profiler; 2) collecting sediment samples on both sides of the fault by using a piston core sampler; 3) analyzing core samples for magnetic susceptibility, micropaleontology and lithology; 4) correlating exactly each sediment layer of the cores on both sides of the fault to recognize several seismic events by relative vertical displacements of relevant layers between the two core sediments; 5) establishing chronology of the events by AMS ¹⁴C dating of mainly shell fossil samples selected from the sediments.

We also applied AMS ¹⁴C dating to establish the chronology of the Nojima fault, located in Awajishima, Hyogo Prefecture, which is one of the faults that generated the 1995 Hyogoken Nanbu Earthquake that occurred on 17 January 1995 and destroyed the city of Kobe. Preliminary chronological results are: 1) the Nojima fault has repeated the displacements a few times in the last 2000 yr; 2) a slip definitely occurred along the Nojima fault *ca*. 2000 yr ago; 3) several types of deformation, including an open crack, were likely to be formed close to the fault, associated with an earthquake that occurred 400 yr ago, that is probably assigned as the 1596 Keicho Earthquake.

¹Dating and Materials Research Center, Nagoya University, Japan ²Faculty of Science, Kochi University, Japan ³Earthquake Research Institute, University of Tokyo, Japan ⁴Department of Geography, Hiroshima University, Japan ⁵Faculty of Education, Oita University, Japan ⁶Faculty of Literature, Aichi Prefectural University, Japan ⁷Graduate School of Human Informatics, Nagoya University, Japan

ENVIRONMENTAL ⁹⁰Sr MEASUREMENTS

M. PAUL,¹ D. BERKOVITS,¹ H. FELDSTEIN,² L. DEWAYNE CECIL³ and S. VOGT⁴

 90 Sr (T=28.5 yr) is a long-lived anthropogenic radionuclide produced in fission. The current and historical distribution of 90 Sr in the environment is useful for the prediction of the behavior of this nuclide released into the environment. The rapid radiochemical detection of 90 Sr in environmental

samples poses non-trivial problems mainly because its pure beta activity is masked by a shorterlived strontium isotope ⁸⁹Sr, also a fission product. AMS offers a direct, fast and sensitive measurement of ⁹⁰Sr concentration by atom counting, the major detection problem being in this case the interference of stable isobaric ⁹⁰Zr impurities. The ⁹⁰Y isobaric daughter reaches secular equilibrium where 90 Y/ 90 Sr = 2 × 10⁻⁴ and is insignificant in atom counting. AMS measurements of 90 Sr were performed with the Rehovot 14UD Pelletron accelerator at a terminal voltage of 11 MV using our standard detection system. Samples of SrH₂ were prepared by reduction of SrO to Sr and conversion to hydride in a hydrogen chamber at 400 deg C. SrH₃⁻ are the preferred ions owing to high beam intensity and low Coulomb explosion effects. The ions were identified by measuring time-of-flight, total energy and three independent energy loss signals in an ionization chamber filled with 15.7 torr of isobutane. The measurements are normalized to a known standard of 90 Sr/Sr = 4.3 × 10⁻¹¹. For this standard 400 90 Sr¹¹⁺ ions were counted in 13 min. A ratio of 90 Sr/Sr = 2 × 10⁻¹³ was measured for a blank sample (3 counts in 33 min), probably due to the vicinity of the intense group of ⁹⁰Zr ions in the spectra. This sensitivity corresponds in practice to a ⁹⁰Sr specific activity of 0.2 pCi/L in environmental samples, a factor of *ca*. 5 better than typically achieved in standard beta counting methods. Environmental samples are being prepared and studied with our AMS system.

Work was supported in part by the International Atomic Energy Commission (Vienna) and the Department of Energy (USA).

¹Racah Institute of Physics, Hebrew University, Jerusalem 91904 Israel
²Soreq NRC, Yavne, Israel
³U.S. Geological Survey, Idaho Falls, Idaho 83403 USA
⁴Department of Chemistry, Purdue University, West Lafayette, Indiana 47907 USA

EXPANDED SMALL SAMPLE CAPABILITIES AT NOSAMS: PREPARATION AND ANALYSIS OF CONVENTIONAL GRAPHITE TARGETS CONTAINING $\lesssim 0.15~\mu g$ CARBON

A. PEARSON, A. P. McNICHOL, R. J. SCHNEIDER and K. F. VON REDEN

Woods Hole Oceanographic Institution, Woods Hole, Massachusetts 02543 USA

Eighty small AMS samples containing between 15 and 150 μ g carbon were prepared by reduction of CO₂ over cobalt metal at 625 ± 25°C in the presence of excess H₂. Analysis of the graphitization procedure shows that optimal reduction conditions favor temperatures below 608°C, and success rates improve with larger sample sizes. AMS results for the 41 OxI and OxII standards found that 84% of targets containing ≥ 20 μ g carbon had "good" (accurate to within 2% of "true" and imprecision of ~2%) performance, while only 20% of targets containing <20 μ g carbon were classified "good". Mean fm results are 1.051 ± 0.015 (HOxI) and 1.364 ± 0.015 (HOXII), for an overall imprecision of ±15‰. The graphitization blank is insignificant—equivalent to ~0.01 mmol modern carbon.

Detailed analysis of the raw AMS data suggests that machine efficiency issues are critical below the 100 μ g threshold and must be addressed to achieve high quality results on small samples. Sample ¹²C current output (I12) scales linearly with sample size. The low currents associated with small samples in turn affect the isotopic ratios ¹⁴C/¹²C and δ^{13} C. The depression in ¹⁴C/¹²C ratio is especially significant and is not believed to be the result of an unquantified "dead" carbon blank. While application of a size-calibrated model correction factor to all calculations is possible, the use of size-matching standards is recommended as the preferable option for analysis of sub-milligram samples.