Determination of Sr-90 and Pb-210 in freshwater fish in Austria



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A method for the determination of ⁹⁰Sr and ²¹⁰Pb in freshwater fish was developed. The determinations were conducted within a project on behalf of the Federal Ministry of Health. The aim of this project was to get an overview of natural radionuclides and artificial radionuclides in wild caught freshwater fish in different lakes in Austria.

For sampling the Neusiedler See in Burgenland, two lakes in Styria the Grundlsee and the Toplitz See and the Zeller See in Salzburg were chosen. Chub (leuciscus cephalus), pike (esox lucius), perch (perca fluviatilis), carp (cyprinus carpio), catfish (silurus glanis), pike-perch (sander lucioperca) and burbot (lota lota) were analysed.

The fish sample was ashed and dissolved. After ammonium oxalate precipitation and destruction of the oxalate 90 Sr and 210 Pb were separated with strontium specific extraction columns (Eichrom Industries, Inc. / TrisKem International). 90 Sr and 210 Pb were measured with Hisafe 3 and Quantulus 1220^{IM} . For the determination of the chemical recovery first the initial strontium and lead concentration in the sample was measured and then a Sr(NO₃)₂ and Pb(NO₃)₂ carrier solution was added. The strontium and lead concentrations were measured with ICP-MS. First different elution solutions were tested to elute the 210 Pb from the column. Then the whole procedure was tested with the IAEA-414 fish reference material. The sample amount for the determination of the wild caught fish varied between 190 g and 2 kg. 5.5 g Sr-spec® resin were used for column preparation.

Festing different elution solutions for F	ъ-210
chemical recovery of the column for	r Pb [%]
6 M HCI	4
Water	12
0.1 M Ammonium oxalate	75

Determination of ²¹⁰Pb in IAEA-414 fish

25 g of reference material were ashed and	analysed for ²¹⁰ Pb.
The chemical recovery for Sr and for Pb wa	s 83,3% and 79,6% respectively.
Laboratory value ²¹⁰ Pb [Bq/kg]	2.08 ± 0.32
Reference sheet information value [Bq/kg]	2.1 (1.8 – 2.5 (95% confidence interval))

Sample Preparation:
ashing at 500°C
\downarrow
Addition of Sr-carrier and Pb-carrier solution
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digestion with conc. HNO_3 and H_2O_2
\downarrow
precipitation with ammonium oxalate
\downarrow
destruction of the oxalate with conc. HNO ₃
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dissolving in 8 M HNO ₃ (n-Oct. saturated)
Ļ
\rightarrow Sr and Pb separation

Sr and Pb separation with Sr-spec:
loading the sample on the column
elution of the 90 Y with 8 M HNO3
washing with 3 M HNO $_3$ + 0.05 M Oxalic acid
washing with 3 M HNO ₃
elution of ⁹⁰ Sr with 0.05 M HNO ₃ \rightarrow \rightarrow LSC (⁹⁰ Sr) + ICP-MS (Sr-carrier)
\downarrow elution of ^{210}Pb with 0,1 M ammonium oxalate \rightarrow LSC ($^{210}\text{Pb})$ + ICP-MS (Pb-carrier)

Results:

For elution of ²¹⁰Pb the most adequate solution tested was the 0.1 M ammonium oxalate solution. The separation method is a fast and suitable method for determining ⁹⁰Sr and ²¹⁰Pb in the same sample aliquot with just one column (Sr-spec®) separation.

The Figures below show the results of our measurements. The median chemical recovery for ⁹⁰Sr and ²¹⁰Pb was 76% and 70% respectively. The sample with the highest ²¹⁰Pb and ⁹⁰Sr activity concentration was perch from Grundlsee. This was a sample from combined small fish which still contained all the fishbone. From the other samples most of the fishbone were taken out. The lower limit of detection for ⁹⁰Sr and ²¹⁰Pb varied with the amount of sample used and the chemical recovery. For 1 kg fresh weight and the median chemical recovery for ⁹⁰Sr and ²¹⁰Pb was 6.9 mBq/kg_{fresh weight} and 7.4 mBq/kg_{fresh weight} respectively.

