

Natural Environment Research Council Institute of Geological Sciences

Mineral Reconnaissance Programme Report

A report prepared for the Department of Industry

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Report 44

Reconnaissance geochemical maps of parts of south Devon and Cornwall

INSTITUTE OF GEOLOGICAL SCIENCES

Natural Environment Research Council

Mineral Reconnaissance Programme

Report No. 44

Reconnaissance geochemical maps of parts of south Devon and Cornwall

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Bibliographical reference

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This brief report accompanies the following 1:50 000 scale geochemical maps, which are placed on open file at the offices of the Institute of Geological Sciences at London, Keyworth, Exeter, Leeds and Edinburgh.

- 1 Chromium in stream sediments, Camelford and Padstow area*
- 2 Chromium in stream sediments, Launceston and Okehampton area*
- 3 Chromium in stream sediments, Truro, Bodmin and Liskeard area*
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- 28 Zinc in stream sediments, South Devon and the Tamar Estuary area

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- 90 Cerium in panning concentrates, Launceston and Okehampton area.

- 91 Cerium in panning concentrates, Truro, Bodmin and Liskeard area.
- 92 Cerium in panning concentrates, South Devon and the Tamar Estuary area.

* The Camelford and Padstow area is bounded by National Grid lines 180 and 230 E and 70 and 100 N.

The Launceston and Okehampton area is bounded by National Grid lines ${}^{2}30$ and ${}^{2}90 E$ and ${}^{\circ}70$ and ${}^{1}00 N$. The Truro, Bodmin and Liskeard area is bounded by National Grid lines ${}^{1}70$ and ${}^{2}30 E$ and ${}^{\circ}40$ and ${}^{\circ}70 N$. The South Devon and the Tamar Estuary area is bounded by National Grid lines ${}^{2}30$ and ${}^{2}90 E$ and ${}^{\circ}40$ and ${}^{\circ}70 N$.

INTRODUCTION

A geochemical drainage survey of South Devon and Cornwall was conducted between 1970 and 1976. Efforts were made to maintain a consistent sampling method, but because of improvements in analytical techniques, and changes in instrumentation, there is an unavoidable variation in absolute values from area to area. For this reason the raw data are presented using an expanding polygon display which emphasises local variations in element concentration.

The results contain many anomalous concentrations of elements of potential economic value but few of these are likely to be of significance, considering the high level of contamination, from past mine workings and from refuse, which exists in the region. Efforts were made to avoid samples badly contaminated collecting by agricultural or domestic refuse, or at locations immediately downstream of roads. It was considered that, where the drainage was contaminated by mine waste, benefit could be gained from the sample analysis as a check on the records of ore components from that mine. It will be noticed that, in several cases, the presence of ore elements not mentioned in the records is thus brought to light. As one purpose of this survey was to determine if there was any significant mineralisation outside the established mining areas of South Devon and Cornwall, samples were not collected from the most intensively mined areas such as the southern edge of the St Austell Granite.

SAMPLING METHODS

Drainage sampling sites were selected, using 1:25 000-scale Ordnance Survey maps, to cover as many streams up to third order as possible, allowing for sites likely to be contaminated. The samples were collected from points in the stream profile where some natural concentration of heavy minerals was likely to have taken place, and clear of dilution by locally derived material from collapsed banks. The sediment was sieved through a 1/10-inch mesh and the fine fraction split into

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two portions of approximately 2 kg each. One portion was taken as the sediment sample, for drying and further sieving through a 150 μ m mesh, and the other was washed to remove clay minerals and organic matter and then panned until about 50 g of concentrate remained. After drying, a 12 g subsample of this was mixed with 4 g of elvacite for X-ray fluorescence analysis.

ANALYTICAL METHODS

Sediment samples

Atomic absorbtion spectrophotometry: copper, lead, zinc (and silver).

Optical emission spectrometry: iron, manganese, nickel, cobalt, tin, barium, zirconium, chromium, molybdenum (vanadium, niobium, boron and beryllium).

Delayed neutron measurement: uranium. Colorimetry: (tungsten and arsenic).

Concentrate samples

X-ray fluorescence spectrometry: cerium, barium, antimony, tin, copper, lead, zinc, calcium, nickel, iron, manganese, titanium (and arsenic).

Elements shown in parentheses were determined for certain samples, but are not included in the accompanying geochemical maps. Listings are available on application to the Officer in Charge, MMAGU, Institute of Geological Sciences, Keyworth, Nottingham.

PRESENTATION

The analytical and locational data were copied onto punched cards and processed on the dual IBM 360/195 computer at S.R.C. Rutherford Laboratory, using the G-EXEC program package. Summary statistics were calculated for each element, and values exceeding five times the geometric mean were truncated to a convenient round figure at about that value (with the exception of tin where partial classification was found to be necessary). The data were plotted using GPLOT software on a CIL drum plotter, and the plots superimposed on Ordnance Survey 1:50 000 base maps for copying. The limitations of accuracy of the plotter and of the determination of National Grid References by Knox Protractor combine to locate any point to within about 2 mm of its true position on the map.

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