and orange cell emitted a fluorescence longer than 475 nm and 530 nm, respectively. However, no fluorescence was detected from the signet ring cell, amoebocyte, granular amoebocyte or lymphocyte (fig.).

In contrast to earlier results 3-5, we have verified that the signet ring cell is the vanadocyte in A. ahodori after fractionation of cells on a Ficoll density gradient, ESR spectrometry and neutron activation analysis of vanadium⁹. Our conclusions are supported by evidence obtained by X-ray micro-analysis 10-12.

If the tunichrome is involved in the accumulation of vanadium ions in ascidian blood cells, it would seem necessary that the vanadocyte (the signet ring cell) contain the tunichrome. However, no fluorescence due to the tunichrome was detected in the vanadocytes from A. ahodori.

We have extracted a vanadium-binding substance called vanadobin from the blood cells of A. sydneiensis samea. This substance is colorless and can maintain the vanadium ion in the vanadyl form (VO (IV)) even under aerobic conditions. Moreover, this substance has an affinity for exogenous vanadium ions (V) and contains a reducing sugar 13. Taking all the above data into account, we suggest that it is not the tunichrome but rather the vanadobin that is the substance involved in the accumulation of vanadium ions from seawater in ascidian blood cells.

de Vincentiis and his co-worker first noted the emission of fluorescence from ascidian blood cells and the follicle and test cells of ascidian eggs ^{14,15}. Recently, it has been reconfirmed, in a detailed study, that ascidian eggs emit fluorescence from the myoplasmic region of the cytoplasm 16. It will be of interest to determine the substance(s) from which such autonomous fluorescence is derived.

Acknowledgments. We are grateful to Prof M. Yoshida and M. Yamamoto and their staff of the Ushimade Marine Biological Station, Okayama University, for supplying the animals, and to Prof. M. Sugai of Toyama University for providing help in the use of MMSP. Our gratitude is also expressed to Prof O. Nikaido and F. Suzuki of Kanazawa University and Prof. A. Shima of the University of Tokyo for their invaluable suggestions at the beginning of this study. This work was supported in part by a grant-in-aid from the Ministry of Education, Science and Culture, Japan, to H.M. (No. 6088009) and to H.M. and H.S. (Nos. 61030035 and 62540540).

- Present address: Faculty of Pharmaceutical Sciences, University of Tokushima, Sho-machi 1, Tokushima 770 (Japan).
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0014-4754/88/100906-02\$1.50 + 0.20/0

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Announcements

Hildegard Doerenkamp and Gerhard Zbinden Foundation for Realistic Animal Protection in Scientific Research Scientific Award 1988

A prize of DM 50000.- will be awarded for outstanding scientific contributions leading to the reduction of animal use in biomedical research. The specific topic for the year 1988 is: "Reduction of animal use in biomedical research by computer modelling.'

Preference will be given to applications leading to a reduction of the use of large animals (dogs, cats, monkeys). Research in pharmacokinetics and drug metabolism is included in the topic.

The applications may consist of published or unpublished reports on computer use in all areas of biomedical research, provided that they are directly relevant to the topic of this

Computer programs for simulation of animal experiments in teaching and research are also acceptable. No special application forms are required. The jury reserves the right to split the prize among not more than three applicants. Languages: English, German, French.

Deadline for submission is December 31, 1988. Applications should be sent to: Prof. G. Zbinden, Institute of Toxicology, Schorenstraße 16, CH-8603 Schwerzenbach/Switzerland.

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