would increase the asymmetry between the pyrrole and pyridine nitrogens and thus increase the quadrupole splitting. The very extensive π system of the phthalocyanine ring allows for even greater delocalization of electrons from pyrrole nitrogen π orbitals, and it may be that it is this characteristic of the molecule, rather than the axial asymmetry of the pyridine ligands, which causes the large quadrupole splitting. Mössbauer experiments on iron(II) phthalocyanine with axially symmetric ligands (like CO) would show which of these two explanations for the large quadrupole splitting is correct.

Acknowledgment.—We wish to thank Professor A. J. Bearden for his advice concerning the initiation of these experiments and the use of his laboratory and Mössbauer spectrometer. The support of Professor Martin Kamen and the advice of Professor C. L. Perrin concerning preparation of these compounds is also appreciated. Grants-in-aid to Professor Bearden (National Science Foundation GB 5458) and Professor Kamen (National Institute of Health HO-01262 and TRE-1045-02) supported this work.

| IBM WATSON LABORATORY | Thomas H. Moss |
|----------------------------|--------------------|
| AT COLUMBIA UNIVERSITY | |
| NEW YORK, NEW YORK 10025 | |
| DEPARTMENT OF CHEMISTRY | ARTHUR B. ROBINSON |
| UNIVERSITY OF CALIFORNIA | |
| at San Diego | |
| LA JOLLA, CALIFORNIA | |
| Received February 23, 1968 | |
| | |

Monodentate Ethylenediamine Complexes of Chromium(III)

Sir:

A recent paper by Childers, *et al.*,¹ reported the successful chromatographic isolation of the pentaaquo(2-

aminoethylammonium)chromium(III) cation, Cr(enH)- $(OH_2)_5^{4+}$, supposedly the first isolation of a complex of chromium(III) with a monodentate ethylenediamine ligand. A chromium(III) complex containing monodentate ethylenediamine may have been prepared more than half a century ago by the founder of coordination chemistry, Alfred Werner.

In an unpublished, but complete, typewritten manuscript dated June 1916, Werner described the preparation, properties, and constitution of a series of chromium complexes which apparently still remains unreported in the published literature. He formulated the parent compound as $H[Cr(en)_2Cl_4]$ and postulated a coordination number of 8 for chromium in this compound in analogy with its heavier congeners molybdenum and tungsten. We have been unable to prepare these compounds according to Werner's directions, but, on the basis of preliminary reflection spectra of Werner's samples (courtesy of Anorganisch-Chemisches Institut der Universität Zürich), Jørgensen² has tentatively identified the chloride of the series as 1,2,3-[Cr(en)-(enH)Cl₃]Cl, a compound of hexacoordinated chromium containing both bidentate and monodentate ethylenediamine.

Werner's original paper has recently been published both in English³ and German.⁴ Small samples of Werner's compounds are available from the author.

(1) R. F. Childers, K. G. Vander Zyl, D. A. House, R. G. Hughes, and C. S. Garner, *Inorg. Chem.*, 7, 749 (1968).

(2) C. K. Jørgensen in "Werner Centennial," G. B. Kauffman, Symposium Chairman, Advances in Chemistry Series, No. 62, American Chemical Society, Washington, D. C., 1967, pp 163, 164.

(3) G. B. Kauffman, Chymia, **12**, 217, 221 (1967).

(4) G. B. Kauffman, Naturwissenschaften, 54, 573 (1967).

Department of Chemistry George B. Kauffman California State College at Fresno Fresno, California 93726

RECEIVED APRIL 22, 1968