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Marvin J. Hirn, Esquire Greenbaum, Doll, Matthews and Boone 3300 First Mational Tower Louisville, Kentucky 40202

Dear Mr. Hirn:

This is in further reference to your letter dated June 13. 1974 concerning the Commission's jurisdiction over wheelchairs. By letter dated September 10, 1974 (enclosed), the Assistant General Counsel for the Food and Drug Division of the Department of Health, Education and Welfare advised the Commission that wheelchairs are classified as a "device" under the Federal Food, Drug, and Cosmetic Act (FFDCA). In view of section 3(a)(1)(H) of the Consumer Product Safety Act (15 U.S.C. 2052(a)(1)(H)) which excludes from the Commission's jurisdiction, "devices" as that term is defined in section 201(h) of the FFDCA, it is our conclusion that the Commission does not have jurisdiction to regulate wheelchairs. Rather, jurisdiction over that product lies with the Food and Drug Administration.

Sincerely,

Original signed by

David Schmelmaltzer

Assistant

General Counsel

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DEPARTMENT OF HEALTH, EDUCATION, AND WELFARE OFFICE OF THE SECRETARY ROCKVILLE, MD. 20852

OFFICE OF THE GENERAL COUNSEL

September 10, 1974

Michael A. Brown, Esq. General Counsel Consumer Product Safety Commission Washington, D. C. 20207

Dear Mike:

This is in response to your letter of August 19, 1974, in which you inquire whether the Food and Drug Administration regulates wheelchairs as a "device" under the Federal Food, Drug, and Cosmetic Act.

FDA does classify wheelchairs as a device under the Act. Wheelchairs are included within the device inventory that has already been established, and are presently undergoing classification by the General Hospital and Personal Use Device Classification Panel to determine whether they should be subject to pre-market clearance, standardization, or simple market surveillance.

Sincerelly yours

Peter Barton Hutt Assistant General Counsel Food and Drug Division