Sandoz Inc.
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SANDOZ

Dear Valued Customer,

All Sandoz Inc. finished products are labeled in compliance with the requirements of the Food and Drug Administration (FDA) and must be used in the prescribed manner. Within each package of finished pharmaceutical products is a Package Insert which will provide all necessary information; this information is also available in the Physician's Desk Reference (PDR).

Pharmaceutical products which are solids, e.g., tablets and capsules, and are in final form for direct administration to patients, are exempted from the requirements of the Hazard Communication Standard (29 CFR 1910:1200). Either an approved package insert or the drug information in the PDR will be considered an MSDS for purposes of compliance with the Standard. This exemption does not include finished pharmaceutical products in non-solid or non-final form. Accordingly, Sandoz Inc. will furnish MSDSs for products supplied in bulk, ampoules, syringes, or vials, and as creams, liquids, lotions, ointments, or transdermal systems.

If you have any questions regarding a specific pharmaceutical product or component, please do not hesitate to call 1-800-525-8747.

Sincerely,

Sandoz Inc.