



**Safety Data Sheet** (previously Material Safety Data Sheet) –  
**Escitalopram Tablets**

As per 29 CFR 1910.1200(b)(6)(vii), “Any drug, as that term is defined in the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq.), when it is in solid, final form for direct administration to the patient (e.g., tablets or pills); drugs which are packaged by the chemical manufacturer for sale to consumers in a retail establishment (e.g., over-the-counter drugs); and drugs intended for personal consumption by employees while in the workplace (e.g., first aid supplies)”, are exempted from the requirements of the Hazard Communication Standard.

Cadista is a solid, oral dosage form (tablets and capsules) manufacturer / distributor, therefore finished pharmaceutical products manufactured/distributed by Cadista are covered under this exemption. Therefore, Material Safety Data Sheets are not available.

All Cadista finished products are labeled in compliance with the requirements of the Food and Drug Administration (FDA) and must be used in the prescribed manner. Each package of the finished pharmaceutical product is supplied with a package insert/insert (approved labeling) which provides necessary drug safety information. Either an approved labeling or the drug information in the Physician’s Desk Reference (PDR) may be considered a MSDS for the purposes of compliance with the standard.

If you need any further information, please contact Cadista Customer Service at 1-800-313-4623.

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**Operations Center**  
**Jubilant Cadista Pharmaceuticals Inc.**  
207 Kiley Drive  
Salisbury, MD 21801-2249  
Main Number: 410-860-8500  
Fax: 410-860-8719  
Website: [www.cadista.com](http://www.cadista.com)

**Sales & Marketing Office**  
**Jubilant Cadista Pharmaceuticals Inc.**  
790 Township Line Road, Suite 175  
Yardley, PA 19067  
Main Number: 410-860-8500  
Fax: 215-443-9646  
Website: [www.cadista.com](http://www.cadista.com)