



U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION



Certificate of Class I

Awarded to

Megagen Implant Co., Ltd.

377-2 Gyocheon-Ri, Jain-Myeon, Gyeongsan-Si, South Korea

Listing Number: D061501

Registration Number: 3005554774

This is to certify that Sinus Lift Kit has been registered with the U.S. FDA and it is safe for marketing in the United States of America.

Indications for Use: Sinus Lift Kit is intended to perform sinus membrane lift in general dentistry and oral surgery procedures. The device includes surgical curette, elevator, retractor and plugger which are hand-held devices.

Product Class: Elevator, Surgical, Dental

Product Code: EMJ

Subsequent Code: EMK, ELA, EKR, EIG

Trade Name: Sinus Lift Kit

Regulation Number: 21 CFR 872.4565

Regulatory Class: Class I

FDA Registered Date: November 14, 2008

Presented by: Official Correspondent, Consultant and U.S. FDA Designated Agent



Kodent Inc.

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