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

Certificate of Class I

Awarded to

Megagen Implant Co., Ltd.
472, Hanjanggun-ro, Jain-myeon, Gyeongsan-si, South Korea, 712-852

Listing Number: E220672
Registration Number: 3005554774

This is to certify that Auto-Max™ has been registered with the U.S. FDA and it is safe for marketing in the United States of America.

Indications for Use:	Prior to dental implant surgery, it is required to make sufficient bone in height and width. Especially dental implant surgery on the maxilla and mandible need sufficient bone in vertical height to hold dental implant in place. Therefore, GBR(Guided Bone Regeneration) helps having enough bone quantity to place dental implant and increases initial stability using Auto-Max™.
Product Class:	Accessories, Implant, Dental, Endosseous
Product Code:	NDP
Trade Name:	Auto-Max™
Regulation Number:	21 CFR 872.3980
Regulatory Class:	Class I
FDA Registered Date:	Dec 17, 2012

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

KoDent

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