FDA 510(k) Premarket Notification

K072871 MS SYSTEM (PROVISIONAL)

CLEARED

Jan 10, 2008
93 days to decision

K072871 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k072871/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Oct 9, 2007
Decision date	Jan 10, 2008
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Osstem Implant Co., Ltd.**
Location	Busan, KR
Contact	CATHRYN N CAMBRIA
Website	https://www.osstem.com
510(k) history	68 submissions · 68 cleared · 2006-2026

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k072871/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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