

K073307 MAX-FACE MAXILLOFACIAL OSTEOSYNTHESIS SYSTEM

Oct 29, 2008
341 days to decision

K073307 · Product code: JEY · Dental
Source: <https://www.510kdatabase.net/k073307/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	Nov 23, 2007
Decision date	Oct 29, 2008
Days to decision	341 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bio-Technology USA, Inc.
Location	Austin, TX, US
Contact	IAN GORDON
510(k) history	1 submissions · 1 cleared · 2008-2008

510k Database - www.510kdatabase.net
 Device record: <https://www.510kdatabase.net/k073307/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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