

K090839 MODIFICATION TO SYNTEC-TAICHUN NON-STERILE BONE PLATE AND SCREW IMPLANTS

Apr 24, 2009
28 days to decision

K090839 · Product code: **HRS** · Orthopedic
Source: <https://www.510kdatabase.net/k090839/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Plate, Fixation, Bone (HRS)
Date received	Mar 27, 2009
Decision date	Apr 24, 2009
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Syntec Scientific Corp.
Location	Taipei City, TW
Contact	CAROL CHANG
510(k) history	13 submissions · 13 cleared · 1998-2012

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k090839/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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