FDA 510(k) Premarket Notification

CLEARED

K073159 TRAUSON BONE SCREW, MODELS HA 3.5 AND 4.5

Jul 23, 2008
258 days to decision

K073159 · Product code: **HWC** · Orthopedic
Source: https://www.510kdatabase.net/k073159/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Nov 8, 2007
Decision date	Jul 23, 2008
Days to decision	258 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Trauson (Jiangsu) Medical Instrument Co., Ltd.**
Location	Shanghai, CN
Contact	Diana Hong
510(k) history	4 submissions · 4 cleared · 2008-2009

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k073159/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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