

K081390 BIOACTIF OSTEOTRANS INTERFENCE SCREWAug 12, 2008
85 days to decisionK081390 · Product code: **HWC** · Orthopedic
Source: <https://www.510kdatabase.net/k081390/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	May 19, 2008
Decision date	Aug 12, 2008
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Takiron Co., Ltd.
Location	Himeji, Hyogo, JP
Contact	KUNIHURO HATA
510(k) history	6 submissions · 6 cleared · 2007-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k081390/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 19, 2026