

K063303 UNIVERSAL LOCKING SYSTEM, 2.7MM PLATES & SCREWSNov 22, 2006
21 days to decisionK063303 · Product code: **HRS** · Orthopedic
Source: <https://www.510kdatabase.net/k063303/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Plate, Fixation, Bone (HRS)
Date received	Nov 1, 2006
Decision date	Nov 22, 2006
Days to decision	21 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Zimmer, Inc.
Location	Warsaw, IN, US
Contact	ANTHONY FRANCALANCIA
Website	https://www.zimmerbiomet.com
510(k) history	373 submissions · 352 cleared · 1976-2026

Zimmer, Inc. is a leading orthopedic medical device manufacturer based in Warsaw, US. The company specializes in innovative surgical implants and trauma solutions. Zimmer, Inc. maintains a strong FDA 510(k) regulatory record with cleared devices from total submissions since 1976. Orthopedic devices represent approximately 90% of the company's submission portfolio. The company remains actively engaged in product development, with the latest FDA 510(k) clearance in 2026. Recent cleared devices reflect the company's focus on joint reconstruction and trauma fixation. Notable ...

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Device record: <https://www.510kdatabase.net/k063303/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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