

K070292 DUROM HIP RESURFACING SYSTEM, FEMORAL COMPONENTSApr 26, 2007
85 days to decisionK070292 · Product code: **KXA** · Orthopedic
Source: <https://www.510kdatabase.net/k070292/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Femoral, Resurfacing (KXA)
Date received	Jan 31, 2007
Decision date	Apr 26, 2007
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Zimmer GmbH
Location	Warsaw, IN, US
Contact	ANTHONY FRANCALANCIA
510(k) history	43 submissions · 43 cleared · 2004-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k070292/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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