

**K914309 THE GENESIS BED FROM HILL-ROM, MODEL 355
SERIES**Apr 20, 1992
208 days to decisionK914309 · Product code: **HDD** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k914309/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Obstetrical, Ac-powered (and Accessories) (HDD)
Date received	Sep 25, 1991
Decision date	Apr 20, 1992
Days to decision	208 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Hill-Rom, Inc.
Location	Batesville, TN, US
Contact	FORD WILDER
Website	http://www.hill-rom.com
510(k) history	33 submissions · 33 cleared · 1986-2021

Hill-Rom, Inc. is a global medical device manufacturer specializing in hospital care equipment and patient handling solutions. The company operates with a manufacturing facility in Batesville, Indiana, and serves healthcare providers across multiple care settings including hospitals, surgical centers, and home healthcare environments. Hill-Rom has received FDA 510(k) clearances from total submissions since its first clearance in 1986. The company's regulatory portfolio focuses on General Hospital devices, including patient beds, stretchers, warming systems, and fluid mana...

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