

**K883299 SUTURE RETENTION DEVICE**Aug 23, 1988  
19 days to decisionK883299 · Product code: **KGS** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k883299/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Retention Device, Suture (KGS)
Date received	Aug 4, 1988
Decision date	Aug 23, 1988
Days to decision	19 days
Third-party review	No

**APPLICANT**

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Company	<b>Edwards Orthopaedics Div., Baxter Healthcare Corp.</b>
Location	Santa Ana, CA, US
Contact	RONALD L DIECK
510(k) history	2 submissions · 2 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883299/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 18, 2026