

**K951901 MASTERSCREEN PFT BODY**Jul 31, 1995  
98 days to decisionK951901 · Product code: **JEH** · AnesthesiologySource: <https://www.510kdatabase.net/k951901/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plethysmograph, Volume (JEH)
Date received	Apr 24, 1995
Decision date	Jul 31, 1995
Days to decision	98 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Erich Jaeger GmbH &amp; Co. KG</b>
Location	Wuerzburg, Germany, DE
Contact	DETLEF GROTHEER
510(k) history	4 submissions · 4 cleared · 1995-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951901/> 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 25, 2026