

**K902334 MONITOR, MODEL 2817**Jun 26, 1990  
33 days to decisionK902334 · Product code: **DXJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k902334/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	May 24, 1990
Decision date	Jun 26, 1990
Days to decision	33 days
Third-party review	No

**APPLICANT**

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Company	<b>Esaote Biomedica Spa</b>
Location	Cambridge, MA, US
Contact	BARRY SALL
510(k) history	4 submissions · 4 cleared · 1990-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902334/>; 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 21, 2026