

K895208 ZYMED MODEL 1610Dec 12, 1989
117 days to decisionK895208 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k895208/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 17, 1989
Decision date	Dec 12, 1989
Days to decision	117 days
Third-party review	No

APPLICANT

Company	Agilent Technologies
Location	Walker, MI, US
Contact	JOSE R GONZALEZ
Website	http://www.agilent.com
510(k) history	13 submissions · 13 cleared · 1982-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895208/> Data sourced from FDA 510(k) public records (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. - Generated May 25, 2026