

**K880505 ECG/RESPIRATION/ARRHYTHMIA MODULE FOR RM-301**Dec 27, 1988  
326 days to decisionK880505 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k880505/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Feb 5, 1988
Decision date	Dec 27, 1988
Days to decision	326 days
Third-party review	No

**APPLICANT**

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Company	<b>Ppg Industries, Inc.</b>
Location	Pleasantville, NY, US
Contact	STEVE BRODY
Website	<a href="http://corporate.ppg.com/Home.aspx">http://corporate.ppg.com/Home.aspx</a>
510(k) history	20 submissions · 20 cleared · 1987-1992

PPG Industries, Inc. is a global manufacturer of paints, coatings, and specialty products headquartered in Pittsburgh, Pennsylvania, with a manufacturing facility in Pleasantville, US. The company operates in more than 50 countries and reported net sales of \$15.9 billion in 2025. PPG Industries received FDA 510(k) clearances from total submissions, with all submissions resulting in clearance. The company's regulatory activity focused on cardiovascular devices, including physiological monitoring systems, recording equipment, and transducer catheters. Clearances span from 1...

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