

K840305 CATEGORY 1 CIRCULATORY SYSTEM DEVICEApr 23, 1984
90 days to decisionK840305 · Product code: **DWS** · CardiovascularSource: <https://www.510kdatabase.net/k840305/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instruments, Surgical, Cardiovascular (DWS)
Date received	Jan 24, 1984
Decision date	Apr 23, 1984
Days to decision	90 days
Third-party review	No

APPLICANT

Company	Imm Enterprises , Ltd.
Location	Walker, MI, US
510(k) history	52 submissions · 52 cleared · 1983-1984

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Device record: <https://www.510kdatabase.net/k840305/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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