

K813123 EPC FILTERNov 27, 1981
23 days to decisionK813123 · Product code: **DRR** · CardiovascularSource: <https://www.510kdatabase.net/k813123/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Amplifier And Signal Conditioner, Biopotential (DRR)
Date received	Nov 4, 1981
Decision date	Nov 27, 1981
Days to decision	23 days
Third-party review	No

APPLICANT

Company	Codman & Shurtleff, Inc.
Location	Mchenry, IL, US
510(k) history	152 submissions · 151 cleared · 1976-2020

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Device record: <https://www.510kdatabase.net/k813123/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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