

K812354 KONTRON PMS 7000Sep 25, 1981
37 days to decisionK812354 · Product code: **DQK** · Cardiovascular
Source: <https://www.510kdatabase.net/k812354/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Aug 19, 1981
Decision date	Sep 25, 1981
Days to decision	37 days
Third-party review	No

APPLICANT

Company	Kontron Instruments, Inc.
Location	Mchenry, IL, US
510(k) history	57 submissions · 57 cleared · 1981-1993

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Device record: <https://www.510kdatabase.net/k812354/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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