FDA 510(k) Premarket Notification

K782072 QUINTON STATUS 1000

CLEARED

Feb 23, 1979
72 days to decision

K782072 · Product code: **DQK** · Cardiovascular
Source: https://www.510kdatabase.net/k782072/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Programmable (DQK)
Date received	Dec 13, 1978
Decision date	Feb 23, 1979
Days to decision	72 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003