FDA 510(k) Premarket Notification

# K791364 QUINTON MODEL 630 AND 636 ECG DATA CART

**CLEARED**

Aug 3, 1979
24 days to decision

K791364 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k791364/

## SUBMISSION DETAILS

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Electrocardiograph (DPS) |
| Date received | Jul 10, 1979 |
| Decision date | Aug 3, 1979 |
| Days to decision | 24 days |
| Third-party review | No |

## APPLICANT

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

**510k Database** - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k791364/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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