FDA 510(k) Premarket Notification

K780248 CRONEX QC-1 PROCESSOR

CLEARED

Mar 2, 1978
17 days to decision

K780248 · Product code: **IXW** · Radiology
Source: https://www.510kdatabase.net/k780248/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Processor, Radiographic-film, Automatic (IXW)
Date received	Feb 13, 1978
Decision date	Mar 2, 1978
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**E.I. Dupont DE Nemours & Co., Inc.**
Location	Mchenry, IL, US
510(k) history	253 submissions · 252 cleared · 1976-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k780248/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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