

K770574 CRONEX QUANTA III INTENSIFYING SCREENSApr 5, 1977
8 days to decisionK770574 · Product code: **EAM** · Radiology
Source: <https://www.510kdatabase.net/k770574/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screen, Intensifying, Radiographic (EAM)
Date received	Mar 28, 1977
Decision date	Apr 5, 1977
Days to decision	8 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

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Device record: <https://www.510kdatabase.net/k770574/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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