

K800223 PATIENT POSITIONING SYSTEM 1D OR 2DMar 3, 1980
28 days to decisionK800223 · Product code: **IWE** · Radiology
Source: <https://www.510kdatabase.net/k800223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	Feb 4, 1980
Decision date	Mar 3, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Laboratoire National D'essais
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k800223/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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