FDA 510(k) Premarket Notification

K822305 ORTHO-LEVEL

CLEARED

Aug 20, 1982
24 days to decision

K822305 · Product code: **HTH** · Orthopedic
Source: https://www.510kdatabase.net/k822305/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Protractor (HTH)
Date received	Jul 27, 1982
Decision date	Aug 20, 1982
Days to decision	24 days
Third-party review	No

APPLICANT

Company	**Frontier Products**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1983