

K822305 ORTHO-LEVELAug 20, 1982
24 days to decisionK822305 · 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

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Device record: <https://www.510kdatabase.net/k822305/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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