

K830878 BINOCULAR LOUPEApr 6, 1983
19 days to decisionK830878 · Product code: **HJH** · Ophthalmic
Source: <https://www.510kdatabase.net/k830878/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Loupe, Binocular, Low Power (HJH)
Date received	Mar 18, 1983
Decision date	Apr 6, 1983
Days to decision	19 days
Third-party review	No

APPLICANT

Company	Almore Intl., Inc.
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1980-1986

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Device record: <https://www.510kdatabase.net/k830878/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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