

K800091 COMBISON 100-MODEL 100 & 110Feb 11, 1980
28 days to decisionK800091 · Product code: **IYO** · Radiology
Source: <https://www.510kdatabase.net/k800091/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Imaging, Pulsed Echo, Ultrasonic (IYO)
Date received	Jan 14, 1980
Decision date	Feb 11, 1980
Days to decision	28 days
Third-party review	No

APPLICANT

Company	Kt Medical, Inc.
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1980-1980

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k800091/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026