

**K032156 ACCUFIX RADIOLUCENT PELVIS AND BELLY
BOARDS, ACCUFIX CANTILEVER HEAD BOARD WITH
SHOULDER DEPRESSION, ACCUFIX RADIOLUCENT**

Sep 2, 2003
50 days to decision

K032156 · Product code: **IYE** · Radiology
Source: <https://www.510kdatabase.net/k032156/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jul 14, 2003
Decision date	Sep 2, 2003
Days to decision	50 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Wfr/Aquaplast Corp.
Location	Wyckoff, NJ, US
Contact	J. DAMON KIRK
510(k) history	8 submissions · 8 cleared · 1994-2012

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k032156/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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