

**K933778 WHITMORE ANKLE X-RAY EXTENSION (AXE)**Dec 13, 1993  
132 days to decisionK933778 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k933778/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Aug 3, 1993
Decision date	Dec 13, 1993
Days to decision	132 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Whitmore Ent., Inc.</b>
Location	San Antonio, TX, US
Contact	EARL HARMSSEN
510(k) history	2 submissions · 2 cleared · 1993-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933778/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 20, 2026