

**K812332 KERMATH L/C ARM**Sep 16, 1981  
29 days to decisionK812332 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k812332/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Aug 18, 1981
Decision date	Sep 16, 1981
Days to decision	29 days
Third-party review	No

**APPLICANT**

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Company	<b>Kermath Mfg. Corp.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1987

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Device record: <https://www.510kdatabase.net/k812332/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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