

**K822420 REPLAM HYDROXYAPATATITE**Sep 28, 1982  
47 days to decisionK822420 · Product code: **FMF** · General Hospital  
Source: <https://www.510kdatabase.net/k822420/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Aug 12, 1982
Decision date	Sep 28, 1982
Days to decision	47 days
Third-party review	No

**APPLICANT**

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Company	<b>Replam Corp.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1983

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Device record: <https://www.510kdatabase.net/k822420/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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