

**K810795 INSTAFIL**Jun 12, 1981  
81 days to decisionK810795 · Product code: **KPE** · Chemistry  
Source: <https://www.510kdatabase.net/k810795/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, I.v. (KPE)
Date received	Mar 23, 1981
Decision date	Jun 12, 1981
Days to decision	81 days
Third-party review	No

**APPLICANT**

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Company	<b>Instafil, Inc.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k810795/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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