

**K770941 APPLIC SYSTEM ACTIVATOR, ESPE**Jun 2, 1977  
10 days to decisionK770941 · Product code: **EID** · DentalSource: <https://www.510kdatabase.net/k770941/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Restorative And Impression Material (EID)
Date received	May 23, 1977
Decision date	Jun 2, 1977
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Espe GmbH (Us)</b>
Location	Mchenry, IL, US
510(k) history	59 submissions · 57 cleared · 1977-1993

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

Device record: <https://www.510kdatabase.net/k770941/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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