

**K770180 SUB. OF DIACETONE ACRYLAMIDE**Feb 15, 1977  
19 days to decisionK770180 · Product code: **FPX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k770180/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Strip, Adhesive, Closure, Skin (FPX)
Date received	Jan 27, 1977
Decision date	Feb 15, 1977
Days to decision	19 days
Third-party review	No

**APPLICANT**

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Company	<b>Johnson &amp; Johnson Professionals, Inc.</b>
Location	Raynham, MA, US
Website	<a href="https://www.jnj.com">https://www.jnj.com</a>
510(k) history	206 submissions · 184 cleared · 1976-2000

Johnson & Johnson Professionals, Inc. is a medical device company based in Raynham, Massachusetts. The company specializes in surgical and orthopedic devices. The company has received FDA 510(k) clearances from total submissions between 1976 and 2000. Orthopedic devices and neurosurgical instruments represent core product categories. Notable cleared devices include hip and elbow prostheses, programmable valve systems, and aneurysm clips. The company is inactive and represents a historical regulatory record with no submissions in more than two decades. Explore the complete...

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