

**K802248 DISARP ORIGINAL**Sep 26, 1980  
10 days to decisionK802248 · Product code: **GAO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k802248/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Suture, Nonabsorbable (GAO)
Date received	Sep 16, 1980
Decision date	Sep 26, 1980
Days to decision	10 days
Third-party review	No

**APPLICANT**

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Company	<b>Unoplast A/S</b>
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1980-1990

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

Device record: <https://www.510kdatabase.net/k802248/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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