

K820730 MICRO CISE DIAMOND SCALPELApr 9, 1982
23 days to decisionK820730 · Product code: **GDX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k820730/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Scalpel, One-piece (GDX)
Date received	Mar 17, 1982
Decision date	Apr 9, 1982
Days to decision	23 days
Third-party review	No

APPLICANT

Company	CooperVision, Inc.
Location	Southampton, GB
Website	https://www.coopervision.com
510(k) history	97 submissions · 94 cleared · 1978-2024

CooperVision, Inc. is a contact lens manufacturer based in Southampton, GB. The company specializes in ophthalmic devices for vision correction. CooperVision has received FDA 510(k) clearances from total submissions since its first clearance in 1978. Ophthalmic devices represent 88% of the company's regulatory submissions. The company remains active, with its latest FDA 510(k) clearance in 2024. Recent cleared devices include daily disposable contact lenses in spheric, toric, and multifocal designs. Notable product families include MyDay, Clariti 1 day, Biofinity, and Ava...

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