

**K091964 CREST GLIDE CLINICAL GUM PROTECTION DENTAL FLOSS**Sep 24, 2009  
85 days to decisionK091964 · Product code: JES · Dental  
Source: <https://www.510kdatabase.net/k091964/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Floss, Dental (JES)
Date received	Jul 1, 2009
Decision date	Sep 24, 2009
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Procter &amp; Gamble Co.</b>
Location	Cincinnati, OH, US
Contact	VICTORIA P SCHOFIELD
Website	<a href="http://www.pg.com/">http://www.pg.com/</a>
510(k) history	23 submissions · 23 cleared · 1988-2014

Procter & Gamble Co. is a consumer health and personal care company headquartered in Cincinnati, US. The company develops and markets a broad range of health and wellness products globally. Procter & Gamble has received FDA 510(k) clearances from total submissions, with no denied submissions on record. The company's regulatory focus has centered on Obstetrics & Gynecology devices, which represent the dominant category of its submissions. FDA 510(k) clearances span from 1988 to 2014, establishing a historical regulatory record in feminine care and oral health device categories.

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091964/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 6, 2026