

**K833283 RX ORY PENTRON C & B**Nov 28, 1983  
67 days to decisionK833283 · Product code: **EIS** · DentalSource: <https://www.510kdatabase.net/k833283/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Remover, Crown (EIS)
Date received	Sep 22, 1983
Decision date	Nov 28, 1983
Days to decision	67 days
Third-party review	No

**APPLICANT**

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Company	<b>Genesis Industries, Inc.</b>
Location	Walker, MI, US
510(k) history	17 submissions · 17 cleared · 1983-2000

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

Device record: <https://www.510kdatabase.net/k833283/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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