

**K810538 PLASTICEPH**Apr 29, 1981  
62 days to decisionK810538 · Product code: **EAG** · Radiology  
Source: <https://www.510kdatabase.net/k810538/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cephalometer (EAG)
Date received	Feb 26, 1981
Decision date	Apr 29, 1981
Days to decision	62 days
Third-party review	No

**APPLICANT**

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Company	<b>Morningstar Dental Co.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k810538/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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