

K810265 VISIO-BANDFeb 23, 1981
21 days to decisionK810265 · Product code: **KLE** · DentalSource: <https://www.510kdatabase.net/k810265/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Agent, Tooth Bonding, Resin (KLE)
Date received	Feb 2, 1981
Decision date	Feb 23, 1981
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Espe-Premier Sales Co.
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k810265/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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