

**K193637 MRH ALFA Radiographic System, MRH II Radiographic System, MRH IIE Radiographic System**Jan 28, 2020  
32 days to decisionK193637 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k193637/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, X-ray, Stationary (KPR)
Date received	Dec 27, 2019
Decision date	Jan 28, 2020
Days to decision	32 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Compania Mexicana DE Radiologia Cgr, S.A. DE C.V.</b>
Location	Denton, TX, US
Contact	Tonatiuh Monroy Soberon
510(k) history	4 submissions · 4 cleared · 2013-2020

**REGULATORY CONSULTANT**

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Consulting firm	<b>Emergo Global Consulting, LLC</b>
Contact	Stuart R. Goldman

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k193637/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated June 18, 2026