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## Lepu Medical Technology (Beijing) Co., Ltd. Reminding Announcement on the CE Certification of the Three Sirolimus-eluting Stent Systems

The company and all members of the Board of Directors hereby guarantee that the information presented in this Announcement is true, accurate and complete with no false or misleading statement or material omission. The English version of this Announcement is for reference only.

Lepu Medical Technology (Beijing) Co., Ltd. (hereinafter brief as "Lepu Medical") continues to accelerate the implementation of its international development strategy. This announcement is hereby given that the three Sirolimus-eluting Stent Systems: Partner<sup>®</sup> Sirolimus-eluting Coronary Stent System, Nano+<sup>TM</sup> Polymer-free Sirolimus-eluting Coronary Stent System and GuReater<sup>®</sup> CoCr Sirolimus-eluting Coronary Stent System which are independently developed by Lepu Medical acquired EU (European Union) CE Certification recently. The CE certificates were officially issued by the EU Notified Body after review and approval.

Under the increasingly severe situation of new EU regulations and supervision, the success of CE certification of Lepu Medical's core products shows that all technical specifications of the products can continuously meet the requirements of the latest EU regulations and standards. The CE certification strengthened the market competitiveness of the series of drug-eluting stent products. Under the national medical insurance drug and equipment centralized procurement policy, Lepu Medical has accelerated the adjustment of its core product sales strategy, which will speed up the company's pace of opening up the international market, and fully implement the company's globalization. This strategy lays a solid foundation for the company's overall business growth, and at the same time brings more choices and benefits to cardiovascular patients worldwide.

> The Board of Directors Lepu Medical Technology (Beijing) Co., Ltd. November 10, 2020