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## **LifeTech Scientific Corporation**

**先健科技公司**

*(Incorporated in the Cayman Islands with limited liability)*

**(Stock Code: 1302)**

### **VOLUNTARY ANNOUNCEMENT**

#### **The first enrollment for confirmatory clinical trials of IBS<sup>®</sup> Sirolimus-Eluting Iron Bioresorbable Coronary Scaffold System had been completed in China**

This announcement is made by LifeTech Scientific Corporation (the “**Company**”, together with its subsidiaries, the “**Group**”) on a voluntary basis to provide its shareholders and potential investors with information in relation to the latest business and new product development of the Group.

The board (the “**Board**”) of directors (the “**Directors**”) of the Company is pleased to announce that on 10 March 2022, the first enrollment for confirmatory clinical trials of the Group’s self-developed IBS<sup>®</sup> Sirolimus-Eluting Iron Bioresorbable Coronary Scaffold System (“**IBS<sup>®</sup> Coronary Scaffold**” or the “**Product**”) had been completed in Fuwai Yunnan Cardiovascular Hospital in China.

IBS<sup>®</sup> Coronary Scaffold is the world’s first fully degradable iron-based absorbable coronary scaffold. The scaffold backbone is made of high-strength and high-plasticity pure nitrided iron tube, which has ultrathin strut thickness and excellent mechanical properties. Innovative materials research and unique technical path enable the Product to retain the advantages of a permanent metal coronary stent, such as full range of specifications, superior mechanical properties, good biocompatibility, and simple operations, as well as having the characteristic of being fully absorbable. IBS<sup>®</sup> Coronary Scaffold begins to degrade after completing the effective support of blood vessels (3-6 months after implantation), and safely enters into the final phase of degradation process in about 2 years. The Product will eventually be absorbed harmlessly by human tissues, thus effectively avoiding a series of long-term prognostic problems that may be caused by a permanent coronary stent.

The completion of the first enrollment for the IBS<sup>®</sup> Coronary Scaffold's confirmatory clinical trials in China is a major milestone in the Company's research and development of iron-based bioabsorbable material. With the steady advancement of follow-up clinical trials, there will be more evidence-based medical evidences to further confirm the safety and effectiveness of the Product. After being successfully marketed, IBS<sup>®</sup> Coronary Scaffold will bring unprecedented treatment for patients with coronary heart disease in China, which lays a solid foundation for the Product to enter into the global market simultaneously.

By order of the Board  
**LifeTech Scientific Corporation**  
**XIE Yuehui**  
*Executive Director, Chairman and  
Chief Executive Officer*

Hong Kong,  
14 March 2022

*As at the date of this announcement, the Board comprises Mr. XIE Yuehui and Mr. LIU Jianxiong being executive Directors; Mr. JIANG Feng being non-executive Director; and Mr. LIANG Hsien Tse Joseph, Mr. WANG Wansong and Mr. ZHOU Luming being independent non-executive Directors.*