MARKET ACCESS TO CHINA



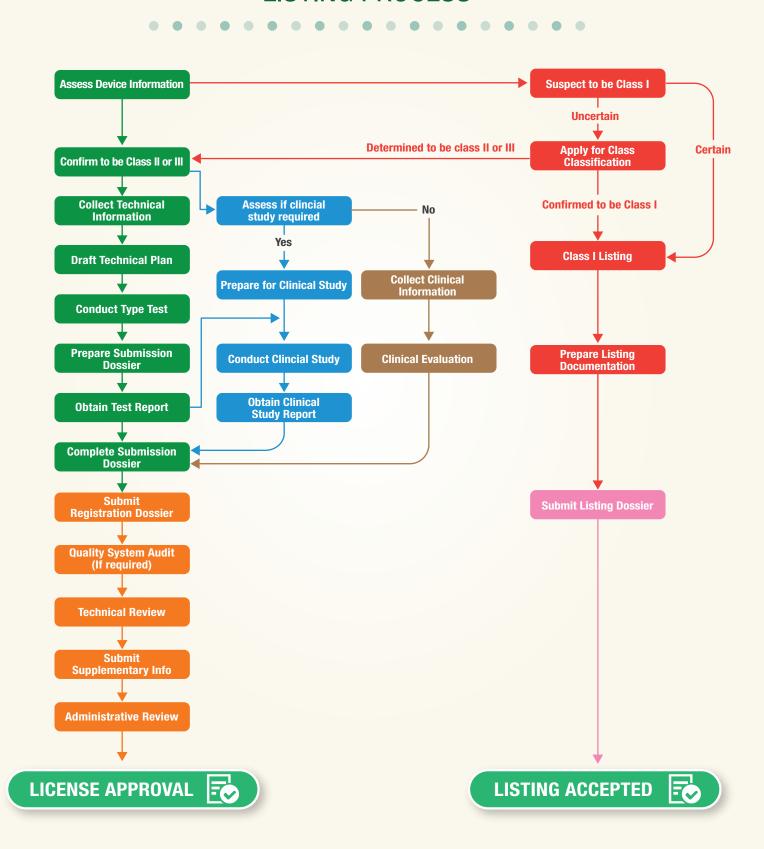




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LISTING PROCESS



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DOCUMENTS REQUIRED

Administrative Information

- Model List
- Quality System Certificate
- Authorization Letter

Device Information

- Product Descriptions
- Intended uses/indications
- Contraindications/Warnings
- Device History

Non-clinical Test Data

- Risk Management File
- Essential Requirement Checklist
- Complied Standards
- Bench Test Reports

Label Information

- Packing Material Artworks
- Pack Insert or Product Manual
- Information for Doctor
- Information for Patient

Vee Care provides a complete set of regulatory and compliance solutions for medical manufacturers.

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HONG KONG

17/F, Chung Pont Com. Bldg. 300 Hennessy Road, Wanchai, Hong Kong

Off: (852) 2893-3118 Fax: (852) 2893-0863

TAIWAN

8/F-1, No. 80, Zhouzi Street, Neihu District, Taipei City 114, Taiwan Off: (886) 02-2758-8892

Room 3501-1A13, East Tower, Fortune Plaza, No.116, Ti Yu Dong Road, Tianhe District, Guangdong, China

Off: (86) 20-8981-5060

USA

4610 Highland Drive Salt Lake City, UT 84117, USA Off: (1) 801-201-167

AUSTRALIA

Level 25, 100 Mount Street, North Sydney, NSW 2060, Australia Off: (61) 2-9959-2400

UK

Unit 808, 54 Bloomfield Avenue, Belfast, BT5 5AD, Northern Ireland, United Kingdom

Off: (44) 743-439-9728