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ASIA EDITION

MET appoints a new GMP Manager

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Global combination device testing laboratory, **Medical Engineering Technologies** (MET), has newly appointed Kevin Claris as Manager for Good Manufacturing Practice and ISO 17025 documentation.

Kevin has 30 years' experience of working with cGMP environments. He has previously pursued his career within both Pfizer and Mylan. This work has mostly been in manufacturing environments. As well as having a great deal of experience in the implementation of quality systems, Kevin has been involved in the development of bronchodilators and their associated MHRA and FDA submissions.

Kevin has a particular interest in training and will be working with the MET technical and quality teams to further develop our

training programmes

Our laboratories specialise in design validation testing of drug delivery devices. Contact our team to find out more about performance testing, drug stability and compatibility, extractables and leachables.

For more than 20 years MET has successfully delivered design validation testing to medical device producers in many countries across: Africa, Asia, Australasia, Europe and the Americas.