



PRESS RELEASE

SIGMA-ALDRICH LIFE SCIENCE & TECHNOLOGY SITE IN WUXI, PR CHINA, RECEIVES EXCiPACT GMP CERTIFICATION AS PHARMACEUTICAL EXCIPIENT SUPPLIERS

Brussels, 14 May 2019

EXCiPACT asbl is pleased to announce that the **Sigma-Aldrich Life Science & Technology site in Wuxi, PR China** has recently been awarded an EXCiPACT GMP Certificate from **Bureau Veritas**, one of EXCiPACT's internationally-recognised Certification Bodies. The Certificate demonstrates that this site manufactures pharmaceutical excipients according to the EXCiPACT Good Manufacturing Practice (GMP) Certification Standard. Its scope covers the testing and packaging for organic and inorganic salts, organic and inorganic acids and bases used as pharmaceutical excipients.

Both **Bureau Veritas** and their auditors had to undergo a rigorous assessment process in order to be EXCiPACT Registered. Both have satisfied the additional EXCiPACT competency requirements set out in the EXCiPACT annex to the ISO/IEC 17021-1:2015 standard. These include completion of the 2-day EXCiPACT Auditor Training Course, post-course written examination and independently witnessed audit to verify their competency to the required standard. **Bureau Veritas** also had to have their auditor's report verified by their independent certification board prior to issuing the certificate.

For full details of all sites that have been EXCiPACT certified to date in 17 Countries (Canada, China, Belgium, France, Germany, India, Israel, Italy, Japan, Romania, Saudi Arabia, Singapore, Spain, Switzerland, The Netherlands, UK and USA), see <http://www.excipact.org/certificate-holders.html>.

EU and U.S. pharmaceutical regulations require drug manufacturers to conduct either their own or commission third-party physical risk-assessment audits of all their starting material suppliers to demonstrate GMP and/or GDP compliance thus increasing the audit burden. However, using GMP and GDP standards designed specifically for pharmaceutical excipients, the voluntary, independent, high quality, EXCiPACT Certification Scheme helps pharmaceutical excipient users and their suppliers to demonstrate their commitment to GMP/GDP, to reduce their audit burden, to save costs and to assure product quality.

Notes for the Editor

EXCiPACT asbl provides management oversight for the voluntary, high quality international EXCiPACT Certification Scheme that provides for independent 3rd party GMP and/or GDP certification of manufacturers, suppliers and distributors of pharmaceutical excipients worldwide. The Scheme ensures patient safety through supplier quality while minimising the overall costs for assessing the excipient supply chain. It has been operational since early 2013 since when there has been considerable interest among pharmaceutical excipient suppliers, their customers and regulators. For further information see www.excipact.org or contact info@excipact.org

EXCiPACT asbl, Rue du Luxembourg 16B, B-1000 Brussels, Brussels Capital Region, Belgium

info@excipact.org www.excipact.org VAT: BE0545 771 884

The French-speaking Business Court of Brussels Capital Region is competent for dispute resolution and litigation

EXCiPACT is a registered trademark