



international excipients  
certification

## NEWS RELEASE

### **NANJING DUOYUAN BIOCHEMISTRY CO. LTD. CHINESE SITE RECEIVES EXCiPACT GMP CERTIFICATION AS PHARMACEUTICAL EXCIPIENT SUPPLIERS**

***Brussels, 26 July 2016***

EXCiPACT asbl is delighted to announce that the Nanjing Duoyuan Biochemistry Co. Ltd Chinese site has recently been awarded an EXCiPACT Certificate from SGS, one of EXCiPACT's internationally-recognised Certification Bodies.

The Certificate demonstrates that the **Nanjing Duoyuan Biochemistry Co. Ltd site in Nanjing City, Jiangsu Province in PR China**, manufactures pharmaceutical excipients according to the EXCiPACT Good Manufacturing Practice (GMP) Certification Standard. Its scope covers manufacturing of pharmaceutical grade Lanolin. For full details of all sites certified to date in Canada, China, Belgium, France, Germany, The Netherlands, India, Israel, Saudi Arabia, Singapore, Spain, Switzerland, UK and USA, see <http://www.excipact.org/certification/certificates/>

Both SGS and their auditors had to undergo a rigorous assessment process in order to be EXCiPACT Registered. This required the successful completion of the EXCiPACT Training Programme and post-course examination followed by an independently witnessed audit to verify that their competency was to the required standard. SGS also had to have their auditor's report verified by an independent certification board prior to issuing the certificate.

EU and U.S. pharmaceutical regulations require drug manufacturers to conduct either their own or commission 3<sup>rd</sup> party physical audits of all their starting material suppliers to demonstrate GMP and/or GDP compliance thus increasing the audit burden. Using GMP and GDP standards designed for excipients, the independent, high quality 3<sup>rd</sup> Party EXCiPACT Certification Scheme is already helping excipient users and suppliers to reduce their audit burden, save costs and assure quality.

#### Notes for the Editor

EXCiPACT asbl provides management oversight for a high quality, voluntary international scheme that provides for independent 3<sup>rd</sup> party certification of manufacturers, suppliers and distributors of pharmaceutical excipients worldwide. The Scheme will ensure patient safety through supplier quality while minimising the overall costs for assessing the excipient supply chain. It was launched in January 2012 since when there has been considerable interest among pharmaceutical excipient suppliers, customers and regulators. For further information see [www.excipact.org](http://www.excipact.org) or contact [info@excipact.org](mailto:info@excipact.org). **EXCiPACT is a registered trademark.**

minimize risks, maximize benefits

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