证券代码：000028 、200028　　证券简称：一致药业、一致B　　公告编号：2011-14

深圳一致药业股份有限公司

关于属下深圳致君制药有限公司

无菌头孢制剂生产获得欧盟GMP认证的公告

本公司及董事保证信息披露的内容真实、准确、完整，没有虚假记载、误导性陈述或重大遗漏。

本公司全资子公司深圳致君制药有限公司无菌头孢制剂生产于2011年2月接受了瑞典药品管理局EU-GMP符合性检查，并于近日收到瑞典药品管理局颁发的欧盟认可的GMP认证证书，证书编号为24:2010/512180（依照欧盟指令2001/83/EC第111（5）号法案检查及颁发），证书有效期三年。获得欧盟认证将给致君制药无菌头孢制剂在国内和欧洲等国际市场的销售带来积极影响。

特此公告。

深圳一致药业股份有限公司董事会

二〇一一年五月十三日

附：认证证书

LÄKEMEDELSVERKET
MEDICAL PRODUCTS AGENCY

Drug Inspectorate

Date: May 3, 2011

Dnr: 24:2010/512180

To whom it may concern

CERTIFICATE OF GMP COMPLIANCE OF A MANUFACTURER [1]

Issued following an inspection in accordance with Art. 111(5) of Directive 2001/83/EC

The competent authority of Sweden confirms that:

The manufacturer: **Shenzhen Zhijun Pharmaceutical Co., Ltd.**
Hi-Tech Zone GuanLan, Baoan Area
Shenzhen, Guandong, PRC

has been inspected during February 22-25, 2011 for compliance to EU-GMP.

The inspection was a general GMP inspection and covered the manufacturing of medicinal products in the form of cephalosporin powder for injection by aseptic filling.

From the knowledge gained during inspection, it is considered that the manufacturer complies with the principles and guidelines of Good Manufacturing Practice requirements referred to in; "The principles and guidelines of Good Manufacturing Practice" laid down in Directive 91/356 as amended by 2003/94/EC[1].

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than three years have elapsed since the date of that inspection.

The authenticity of this certificate may be verified with the issuing authority.

On behalf of the Medical Products Agency, Sweden

LÄKEMEDELSVERKET · MEDICAL PRODUCTS AGENCY ·

Tor Gråberg
Chief Pharmaceutical Inspector

Dr Bruno Barbieri
Principal Pharmaceutical Inspector

[1]These requirements fulfil the GMP recommendations of WHO.

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