

UNITED ARAB EMIRATES

MINISTRY OF HEALTH

DEPARTMENT OF DRUG CONTROL



دولة الإمارات العربية المتحدة  
وزارة الصحة

## MANUFACTURING SITE REGISTRATION CERTIFICATE

It is certified that the following Manufacturing Site has been registered in the UAE Ministry of Health, in accordance with the Article 65 of the Federal Law No. 3 of 1984.

Certificate #	097	Reg. Date	05/07/2005
Registration #	GRC/141/05	Reg. Expiry Date	04/07/2010
Committee Meeting No.	24/2005	MTG. Date:	05/07/2005
Manufacturing Site Name	ICN POLFA RZESZÓW S.A		
Address	2 Przemyslowa Str., 35-959 RZESZÓW, POLAND		
Line(s) of Production & Other Activities, Registered for			
<b>Non Sterile Dosage Forms: Tablets, Capsules, Coated Tablets, Syrup and Suspension</b>			
Manufacturing Site for Product Classes	<b>CONVENTIONAL</b>		
<ol style="list-style-type: none"><li>1. The evidence(s) for GMP for the above mentioned activity(s) is/are acceptable.</li><li>2. Failure to provide current acceptable GMP or any equivalent evidence prior to the expiry / as &amp; when requested could result in refusal to receive product registration dossier or removal of the affected company and its product(s) from the register, according to the case.</li><li>3. Registration as Manufacturing Site makes it eligible to involve in the registered activities in respect to the products to be registered in the U.A.E.</li><li>4. The Manufacturing Site should apply for minor variation as and when an amendment is needed in registration status.</li><li>5. This registration applies only to the above site name and address.</li></ol>			
Applicant for registering the site	ICN POLFA RZESZÓW S.A 2 Przemyslowa Str., 35-959 RZESZÓW, POLAND		



Director of Drug Control Department

Issued on: 07/02/2007

SYED CO.DOC E

AED 1000/
DRUG REGISTRATION CERTIFICATE
FEE RECEIPT NO.: 31196211
DATE: 18/2/2007
TECHNICAL AFFAIRS SEC. DRUG REGISTRATION & PRICING UNIT M.O.H., ABU DHABI - U.A.E.