

Certification of Substances Department

**Certificate of suitability**  
**No. R1-CEP 2006-272-Rev 04**

1 *Name of the substance:*

2 **GEMCITABINE HYDROCHLORIDE**

3 *Name of holder:*

4 **SCINOPHARM TAIWAN, LTD.**

5 No. 1, Nan-Ke 8th Road

6 Taiwan-74144 Shan-Hua, Tainan

7 *Site(s) of production:*

8 **SEE ANNEX 1**

**Notice**

**NOT FOR REGISTRATION PURPOSES**

**For filing purposes please contact ScinoPharm Taiwan  
to obtain a complete "controlled copy" of this CEP.**

**ScinoPharm Taiwan- Regulatory Technical Services  
(SPT.RTS@scinopharm.com.tw)**

18 Isopropanol

not more than 0.5%

19 In the last steps of the synthesis water is used as solvent.

20 A risk management summary for elemental impurities has been provided.

21 The re-test period of the substance is 5 years if stored in double polyethylene bags, placed in  
22 either a polyethylene bottle or drum.

23 The holder of the certificate has declared the absence of use of material of human or animal  
24 origin in the manufacture of the substance.

25 The submitted dossier must be updated after any significant change that may alter the quality,  
26 safety or efficacy of the substance.

27 Manufacture of the substance shall take place in accordance with the Good Manufacturing Practice  
28 and in accordance with the dossier submitted.


Address: 7 Allée Kastner, CS 30026

F-67081 Strasbourg (France)

Tel: +33 (0) 3 88 41 30 30 – Fax: +33 (0) 3 88 41 27 71 - e-mail: cep@edqm.eu

Internet: <http://www.edqm.eu>

- 29 Failure to comply with these provisions will render this certificate void.
- 30 This certificate is renewed from **4 July 2013** according to the provisions of Resolution AP-CSP  
31 (07) 1, and of Directive 2001/83/EC and Directive 2001/82/EC and any subsequent amendment,  
32 and the related guidelines.
- 33 This certificate has three annexes, the first of 1 page, the second of 3 pages and the third of  
34 2 pages.
- 35 This certificate has:  
36 lines.



On behalf of the  
Director of EDQM



Strasbourg, 14 December 2018

DECLARATION OF ACCESS *(to be filled in by the certificate holder under their own responsibility)*

**SCINOPHARM TAIWAN, LTD.** as holder of the certificate of suitability

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The holder also certifies that no significant changes to the operations as described in the CEP dossier have been made since the granting of this version of the certificate.

Date and Signature *(of the CEP holder)*:

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