

Certification of Substances Department

**Certificate of suitability**  
**No. R0-CEP 2022-010 - Rev 00**

1 *Name of the substance:*

2 **MAGNESIUM HYDROXIDE**

3 *Name of holder:*

4 **ANHUI POLY PHARM. CO., LTD.**

5 No.58 Xiahong Road

6 Hi-tech Industrial Development Zone

7 China-246 000 Anqing, Anhui Province

8 *Site(s) of production:*

9 **SEE ANNEX 1**

10 After examination of the information provided on the manufacturing method and subsequent  
11 processes (including purification) for this substance on the site(s) of production listed in annex, we  
12 certify that the quality of the substance is suitably controlled by the current version of the  
13 monograph **MAGNESIUM HYDROXIDE** no. 39 of the European Pharmacopoeia, current edition  
14 including supplements, only if it is supplemented by the test(s) mentioned below.

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

16 A risk management summary for elemental impurities has been provided. (Annex 2)

17 The re-test period of the substance is 18 months if stored in double polyethylene bags, in a  
18 triple laminated aluminium bag, placed in a fibre drum.

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

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

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


25 Failure to comply with these provisions will render this certificate void.

26 This certificate is granted within the framework of the procedure established by the European  
27 Pharmacopoeia Commission [Resolution AP-CSP (07) 1] for a period of five years starting from  
28 **26 July 2022**. Moreover, it is granted according to the provisions of Directive 2001/83/EC and  
29 Directive 2001/82/EC and any subsequent amendment, and the related guidelines.

30 This certificate has two annexes, the first of 1 page and the second of 2 pages.

31 This certificate has:

32 lines.



On behalf of the  
Director of EDQM

Strasbourg, 26 July 2022

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

**ANHUI POLY PHARM. CO., LTD.** , as holder of the certificate of suitability

**R0-CEP 2022-010 - Rev 00 for Magnesium hydroxide**

hereby authorises .....

*(name of the pharmaceutical company)*

to use the above-mentioned certificate of suitability in support of their application(s) for the following  
Marketing Authorisation(s): *(name of product(s) and marketing number(s), if known)*

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)*:

**Certification of Substances Department**

## **Annex 1: Site(s) of production for R0-CEP 2022-010 - Rev 00**

### **Production of Magnesium hydroxide:**

ANHUI POLY PHARM. CO., LTD.  
No.58 Xiahong Road  
Hi-tech Industrial Development Zone  
China-246 000 Anqing, Anhui Province



### Risk assessment results of elemental impurities in API

Intended route of administration/Use of the substance: Oral				
Elements	Class	Intentionally added?	Considered in risk management?	Conclusion
Cd	1	No	Yes	Absent*
Pb	1	No	Yes	Absent
As	1	No	Yes	Absent
Hg	1	No	Yes	Absent
Co	2A	No	Yes	Absent
V	2A	No	Yes	Absent
Ni	2A	No	Yes	Absent
Tl	2B	No	No	N/A
Au	2B	No	No	N/A
Pd	2B	No	No	N/A
Ir	2B	No	No	N/A
Os	2B	No	No	N/A
Rh	2B	No	No	N/A
Ru	2B	No	No	N/A
Se	2B	No	No	N/A
Ag	2B	No	No	N/A
Pt	2B	No	No	N/A

Intended route of administration/Use of the substance: Oral				
Elements	Class	Intentionally added?	Considered in risk management?	Conclusion
Li	3	No	No	N/A
Sb	3	No	No	N/A
Ba	3	No	No	N/A
Mo	3	No	Yes	Absent
Cu	3	No	Yes	Absent
Sn	3	No	No	N/A
Cr	3	No	Yes	Absent

\* Absent: less than 30% ICH Q3D Option 1 limit.