



| TETRACAINA HCL (EUR. PH.) | | | | |
|---------------------------|-----------------|--------------------------------|--|--|
| PRODUCT CODE: 1703 | CAS Nº: 136-47- | -0 ANALYSIS Nº: 209/25 | | |
| MANUFACTURER BATCH: | 003240901 | CERTIFICATE ID: 47.313 | | |
| SUPPLIER BATCH: | | MANUFACTURING DATE: 09/02/2025 | | |
| МЕТАРН ВАТСН: | 0050725 | RETEST DATE: 31/01/2028 | | |

| ATTRIBUTES | SHOULD BE | IS |
|------------------------|--|------------------------------|
| Appearance | White or almost white, slightly hygroscopic, | White, slightly hygroscopic, |
| | crystalline powder | crystalline powder |
| Solubility | Freely soluble in water, soluble in ethanol (96 %) | Complies (*) |
| Melting range | 134 - 150 °C | 148-149 °C (*) |
| Identification A | Complies | Complies |
| Identification D | Complies | Complies |
| Appearance of solution | Clear and colourless | Complies |
| рН | 4.5 - 6.5 | 5.8 |
| Related substances | | |
| Impurity A | =< 0.05 % | Not Detected |
| Impurity B | =< 0.15 % | Not Detected |
| Impurity C | =< 0.15 % | Not Detected |
| Unspecified impurities | =< 0.10 % | 0.07 % (RT = 24.03 min) |
| Total impurities | =< 0.5 % | 0.07 % |
| Loss on drying | =< 1.0 % | 0.04 % |
| Sulfated ash | =< 0.1 % | 0.08 % |
| Assay | 99.0 % - 101.0 % | 100.5 % |
| COMPLIES WITH | | |

COMPLIES WITH

European Pharmacopoeia 11.0

REMARKS

Product fully analyzed within the EU, in compliance with the currents regulations "EudraLex - Volume 4 - Good Manufacturing Practice (GMP) guidelines" Part-II.

Tetracaine Hydrochloride is subjected to the requirements of the ICH Q3D "Elemental Impurities" guideline and the requirements of guides EMA/CHMP/ICH/82260/2006.

All data controlled by Metapharmaceutical Industrial SL.

 $(\ensuremath{^*})$ Data adapted from the manufacturer's certificate of analysis.

Absence of N-nitrosamines impurities has been ensured after a risk evaluation according to ICH Q9, ICH M7 and in accordance with guidelines EMA/428592/2019 Rev 2 and EMA/189634/2019.

Certificates of residual solvents, allergens, non-GMO and BSE-TSE, among others, are available upon request.

All methods of analysis are validated by official pharmacopoeias or are validated by internal methods of the manufacturer, which can be obtained at specific request. The above information does not exempt from the obligation to identify the product before use.

STORAGE

Store in a cool place. Keep the container tightly closed in a dry and well-ventilated place.

Analysis date: 26/08/2025

Signature: Albert Sánchez López (QP)

Conclusion: Complies

Original certificate available upon request

Manufacturer: 40001510

SHANDONG CHENGHUI SHUANGDA

PHARMACEUTICAL CO.,LTD.

Economic Development Zone of Pingyuan County, 253100, Dezhou City, Shandong

2073 03 FNG

Page 1 of 1