

TIMOLOL MALEATO (EUR. PH.)

| | | |
|------------------------------|--------------------------------|---------------------|
| PRODUCT CODE: 009675 | CAS Nº: 26921-17-5 | ANALYSIS Nº: 141/24 |
| MANUFACTURER BATCH: 22010575 | CERTIFICATE ID: 42.354 | |
| SUPPLIER BATCH: 233502 | MANUFACTURING DATE: 27/08/2022 | |
| METAPH BATCH: 0260524 | EXPIRY DATE: 27/08/2027 | |

| ATTRIBUTES | SHOULD BE | IS |
|---------------------------|--|------------------------------|
| Appearance | White or almost white, crystalline powder or colourless crystals | White crystalline powder (*) |
| Solubility | Soluble in water and in ethanol (96 %) | Complies |
| Melting point | About 199 °C | 199 °C (*) |
| Identification A | Complies | Complies (*) |
| Identification B | Complies | Complies (*) |
| Specific optical rotation | -6.2 / -5.7 | -5.9 (*) |
| Appearance of solution | Clear solution and not more intensely coloured than ref. sol. B8 | Complies |
| pH | 3.8 - 4.3 | 4.0 |
| Enantiomeric purity | | |
| Impurity A | =< 1.0 % | 0.03 % |
| Related substances | | |
| Impurity B | =< 0.2 % | < 0.05 % |
| Impurity C | =< 0.2 % | Not Detected |
| Impurity D | =< 0.2 % | < 0.05 % |
| Impurity E | =< 0.2 % | Not Detected |
| Impurity F | =< 0.2 % | < 0.05 % |
| Unspecified impurities | =< 0.10 % | < 0.05 % |
| Total impurities | =< 0.4 % | < 0.05 % |
| Loss on drying | =< 0.5 % | 0.1 % (*) |
| Sulfated ash | =< 0.1 % | < 0.1 % (*) |
| Assay | 98.5 - 101.0 % | 99.1 % (*) |

COMPLIES WITH

European Pharmacopoeia 11.0

REMARKS

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

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

(*) Data controlled by Metapharmaceutical Industrial SL.

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.

Analysis date: 17/06/2024
 Signature: Albert Sanchez Lopez (QP)
 Conclusion: Complies
 Original certificate available upon request

Manufacturer: 40000581



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STORAGE

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

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