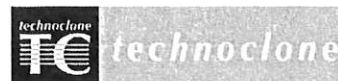


CERTIFICATE OF ANALYSIS

Technoview LMWH CON L



Temp.: TF0326.07

Attachment 17 DQ0H00.03

3F51C00.02

REF 5090042-RUO
 LOT 3F51C00.02
 2027-04-30

| Analytical Procedures | | Requirement | Result | |
|---|----------------------------------|---|------------|----|
| visual inspection | Cap color | red | confirmed | OK |
| | Vial type | clear glass vial | confirmed | OK |
| | Vial label | according to packaging SOP | confirmed | OK |
| | Vial barcode | according to packaging SOP | confirmed | OK |
| | Box label | according to packaging SOP | confirmed | OK |
| | Box barcode | according to packaging SOP | confirmed | OK |
| | Package insert | present | confirmed | OK |
| | Batch table | present | confirmed | OK |
| | Appearance before reconstitution | buff colored plug of lyophilized material | confirmed | OK |
| | Appearance after reconstitution | amber colored liquid | confirmed | OK |
| Virology | HBsAg / HBsAg | Negative | Negative | OK |
| | HIV-Ak / HIV-Ab | Negative | Negative | |
| | HCV-AK / HCV-Ab | Negative | Negative | |
| The testing methods applied were FDA-approved or CE marked. | | | | |
| Recovery of control | | 0.32 - 0.59 IU/mL | 0.42 IU/mL | OK |

Prepared by

Maylis RAUSCHER

16. SEP. 2025

Date

Approved by

Michael KAFKA

16. SEP. 2025

Date