

Certificate of Analysis

Product name: **Fludrocortisoni acetat micr.**

Batch number / Weight: **23J19-B01-234562 / 1 G**

Analysed according to: **Ph.Eur.11.3**

Number of analysis / Inspection Code **23J19-B01 / INS-23-8422**

Reference Code / No.: **V01457 / 2142AM0 B0022223**

| Tests | Requirement | Result | Unit | Standard remark |
|-------------------------------|--|---------|------|-----------------|
| Appearance | (almost) white, crystalline powder | Conform | | |
| Particle size | 99% < 25 µm / 90% <10 µm | Conform | | |
| Identification A | Conform | Conform | | |
| Identification B | Conform | Conform | | |
| Specific optical rotation | +148 - +156 | +150,7 | | |
| Related substances | Conform | Conform | | |
| Any impurity | <=1,0 | 0,08 | % | |
| Total impurities | <=1,5 | 0,13 | % | |
| Loss on drying | <=1,0 | 0,05 | % | |
| Metallic residues | CHMP/ICH/353369/2013 | Conform | | Data producer |
| Residual solvents | CPMP/ICH/82 260/06 | Conform | | Data producer |
| TSE/BSE-statement: | No contamination with TSE/BSE-risk materials | Conform | | Data producer |
| Assay Fludrocortisone acetate | 97,0 - 103,0 | 100,31 | %m/m | |

All the manufacturing stages of this batch have been carried out in full compliance with the GMP requirements of the EU and with the national legal requirements.

Analysis performed by the authorized lab Fagron PL lab

Release:
Ewelina Gadzinowska
Qualified Person

12-01-2024

Expiration: 18-05-2027

Conclusion: APPROVED

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fagron.pl Fagron Services Northern Europe Sp. z o.o.
ul. Armii Krajowej 3
32-540 Trzebinia, Poland
e-mail: fsne@fagron.pl

fagron.pl

Fagron sp. z o.o.
ul. Pasternik 26, 31-354 Kraków, Poland
tel.: +48 12 3343 512
e-mail: biuro@fagron.pl