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Mometasone Furoate Cream

DEFINITION

Mometasone Furoate Cream is Mometasone Furoate in a suitable cream base. It contains NLT 90.0% and NMT 110.0% of the labeled amount of mometasone furoate ($C_{27}H_{30}Cl_2O_6$).

IDENTIFICATION

- A. The retention time of the major peak of the *Sample solution* corresponds to that of the *Standard solution*, both relative to the internal standard, as obtained in the Assay.
- B. THIN-LAYER CHROMATOGRAPHIC IDENTIFICATION TEST (201).

Standard solution: 0.2 mg/mL of [USP Mometasone Furoate RS](#) in acetonitrile

Sample solution: 0.2 mg/mL of mometasone furoate from Cream in acetonitrile

Developing solvent system: Chloroform and ethyl acetate (3:1)

Acceptance criteria: The R_F value of the principal spot of the *Sample solution* corresponds to that of the *Standard solution*.

ASSAY

• PROCEDURE

[NOTE—Protect from light.]

Diluent A: Tetrahydrofuran and glacial acetic acid (100:1)

Diluent B: Acetonitrile, water, and glacial acetic acid (50:50:1)

Solution A: Water

Solution B: Acetonitrile

Mobile phase: See [Table 1](#).

Table 1

Time (min)	Solution A (%)	Solution B (%)
0	70	30
2	70	30
45	45	55
46	70	30
50	70	30

Internal standard solution: 1.4 mg/mL of diethyl phthalate in acetonitrile

Standard stock solution: 0.2 mg/mL of [USP Mometasone Furoate RS](#) in *Diluent A*

Standard solution: 0.05 mg/mL of mometasone furoate and 0.35 mg/mL of diethyl phthalate from equal quantities of the *Standard stock solution* and the *Internal standard solution*, in *Diluent B*

Sample solution: Transfer a portion of Cream, equivalent to 1.0 mg of mometasone furoate, to a 50-mL, screw-capped centrifuge tube. Add 5.0 mL of *Diluent A* and a few glass beads, and mix on a vortex mixer. Add 5.0 mL of *Internal standard solution*, and mix. Add 10.0 mL of *Diluent B*, mix on a vortex mixer for 1 min, and centrifuge for 10 min. Pass the aqueous phase through a polypropylene filter of 0.2- μ m pore size, discarding the first 1–2 mL of filtrate.

Chromatographic system

(See [Chromatography \(621\), System Suitability](#).)**Mode:** LC**Detector:** UV 254 nm**Column:** 4.6-mm × 25-cm; 5-μm packing L60**Flow rate:** 2 mL/min**Injection size:** 20 μL**System suitability****Sample:** Standard solution

[NOTE—The relative retention times for diethyl phthalate and mometasone furoate are 0.4 and 1.0, respectively.]

Suitability requirements**Tailing factor:** NMT 1.5 for the mometasone furoate peak**Relative standard deviation:** NMT 2.0%**Analysis****Samples:** Standard solution and Sample solutionCalculate the percentage of mometasone furoate ($C_{27}H_{30}Cl_2O_6$) in the portion of Cream taken:

$$\text{Result} = (R_U/R_S) \times (C_S/C_U) \times 100$$

 R_U = ratio of the mometasone furoate peak response to the diethyl phthalate peak response from the Sample solution R_S = ratio of the mometasone furoate peak response to the diethyl phthalate peak response from the Standard solution C_S = concentration of [USP Mometasone Furoate RS](#) in the Standard solution (mg/mL) C_U = nominal concentration of mometasone furoate in the Sample solution (mg/mL)**Acceptance criteria:** 90.0%–110.0%**PERFORMANCE TESTS**

- [MINIMUM FILL \(755\)](#): Meets the requirements

IMPURITIES

- [ORGANIC IMPURITIES](#)

[NOTE—Protect from light.]

Diluent A, Solution A, Solution B, Mobile phase, and Standard stock solution: Prepare as directed in the Assay.**Diluent C:** Acetonitrile, water, and glacial acetic acid (30:70:1)**System suitability solution:** 0.1 μg/mL of [USP Mometasone Furoate RS](#) from Standard stock solution in Diluent C**Blank solution:** Diluent C and Diluent A (3:1)**Sample solution:** Transfer a portion of Cream, equivalent to 2.0 mg of mometasone furoate, to a 50-mL, screw-capped centrifuge tube. Add 5.0 mL of Diluent A and a few glass beads, and mix on a vortex mixer. Add 15.0 mL of Diluent C, and mix. Centrifuge for 10 min. Pass the aqueous phase through a 0.2-μm polypropylene filter, discarding the first 1–2 mL of filtrate.**Chromatographic system**(See [Chromatography \(621\), System Suitability](#).)**Mode:** LC**Detector:** UV 254 nm**Column:** 4.6-mm × 25-cm; 5-μm packing L60**Column temperature:** 25 ± 5°**Flow rate:** 2 mL/min**Injection size:** 50 μL**System suitability****Sample:** System suitability solution**Suitability requirements****Relative standard deviation:** NMT 10%**Analysis****Samples:** System suitability solution, Blank solution, and Sample solution

[NOTE—Exclude any peak areas less than those from the System suitability solution. Also exclude any peaks with the same retention time as that observed in the Blank solution. Any peaks having a relative retention time of about 1.04 or 1.13 are controlled in the Mometasone Furoate monograph and, therefore, are not included in the total specified and unspecified impurities limit.]

Calculate the percentage of each impurity in the portion of Cream taken:

$$\text{Result} = (r_U/r_T) \times 100$$

r_U = peak response of each impurity from the *Sample solution*

r_T = sum of all the peak responses from the *Sample solution*

Acceptance criteria: See [Table 2](#).

Table 2

Name	Relative Retention Time	Acceptance Criteria, NMT (%)
9 α -Chloro-11 β ,17,21-trihydroxy-16 α -methylpregna-1,4-diene-3,20-dione 17-(2-furoate)	0.56	0.1
9 α ,21-Dichloro-11 β ,17-dihydroxy-16 α -methylpregna-1,4-diene-3,20-dione	0.73	0.1
21-Chloro-17-hydroxy-16 α -methylpregna-1,4-diene-3,11,20-trione 17-(2-furoate)	0.88	0.1
21-Chloro-9 β ,11 β -epoxy-17-hydroxy-16 α -methylpregna-1,4-diene-3,20-dione 17-(2-furoate)	0.94	1.0
Mometasone furoate	1.0	—
Unspecified individual impurity	—	0.2
Total specified and unspecified impurities	—	1.0

SPECIFIC TESTS

- [MICROBIAL ENUMERATION TESTS \(61\)](#) and [TESTS FOR SPECIFIED MICROORGANISMS \(62\)](#): It meets the requirements of the tests for absence of *Staphylococcus aureus*, *Pseudomonas aeruginosa*, *Escherichia coli*, and *Salmonella* species.

ADDITIONAL REQUIREMENTS

- **PACKAGING AND STORAGE:** Preserve in well-closed containers, and store at controlled room temperature.
- [USP REFERENCE STANDARDS \(11\)](#)
[USP Mometasone Furoate RS](#)

Auxiliary Information - Please [check for your question in the FAQs](#) before contacting USP.

Topic/Question	Contact	Expert Committee
MOMETASONE FUROATE CREAM	Documentary Standards Support	SM22020 Small Molecules 2
REFERENCE STANDARD SUPPORT	RS Technical Services RSTECH@usp.org	SM22020 Small Molecules 2

Chromatographic Database Information: [Chromatographic Database](#)

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