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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-µm 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 solution*

Calculate 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 $\alpha$ -Chloro-11 $\beta$ ,17,21-trihydroxy-16 $\alpha$ -methylpregna-1,4-diene-3,20-dione 17-(2-furoate)	0.56	0.1
9 $\alpha$ ,21-Dichloro-11 $\beta$ ,17-dihydroxy-16 $\alpha$ -methylpregna-1,4-diene-3,20-dione	0.73	0.1
21-Chloro-17-hydroxy-16 $\alpha$ - methylpregna-1,4-diene-3,11,20-trione 17-(2-furoate)	0.88	0.1
21-Chloro-9 $\beta$ ,11 $\beta$ -epoxy-17-hydroxy-16 $\alpha$ -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	<a href="#">Documentary Standards Support</a>	SM22020 Small Molecules 2
REFERENCE STANDARD SUPPORT	RS Technical Services <a href="mailto:RSTECH@usp.org">RSTECH@usp.org</a>	SM22020 Small Molecules 2

**Chromatographic Database Information:** [Chromatographic Database](#)

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