methanol, and dry in a desiccator (in vacuum, silica gel) for 5 hours: the crystals melt between 112°C and 118°C.

**Optical rotation**  
$[\alpha]_D^20 = -23 - 27^\circ$ (after drying, 0.25 g, chloroform, 25 mL, 200 mm).

**Melting point**  
203 – 209°C

**Loss on drying**  
Not more than 0.5% (0.5 g, in vacuum, silica gel, 4 hours).

**Residue on ignition**  
Not more than 0.1% (0.5 g).

**Assay**  
Weigh accurately about 0.2 g of Norethisterone, previously dried, dissolve in 40 mL of tetrahydrofuran, add 10 mL of a solution of silver nitrate (1 in 20), and titrate with 0.1 mol/L sodium hydroxide VS (potentiometric titration). Perform a blank determination, and make any necessary correction.

Each mL of 0.1 mol/L sodium hydroxide VS = 29.843 mg of $\text{C}_9\text{H}_{10}\text{NO}_2$

**Containers and storage**  
Containers—Tight containers. Storage—Light-resistant.

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**Norfloxacin**

ノルフロキサシン

\[\text{C}_{16}\text{H}_{18}\text{FN}_3\text{O}_3: 319.33}\]

1-Ethyl-6-fluoro-1,4-dihydro-4-oxo-7-(piperazin-1-yl)quinoline-3-carboxylic acid  [70458-96-7]

Norfloxacin, when dried, contains not less than 99.0% of $\text{C}_{16}\text{H}_{18}\text{FN}_3\text{O}_3$.

**Description**  
Norfloxacin occurs as a white to pale yellow crystalline powder.

It is freely soluble in acetic acid (100), slightly soluble in ethanol (99.5) and in acetone, very slightly soluble in methanol, and practically insoluble in water.

It dissolves in dilute hydrochloric acid TS and in sodium hydroxide TS.

It is hygroscopic.

It is gradually colored by light.

**Identification** (1)  
Dissolve 0.01 g of Norfloxacin in a solution of sodium hydroxide (1 in 250) to make 100 mL. To 5 mL of this solution add a solution of sodium hydroxide (1 in 250) to make 100 mL. Determine the absorption spectrum of this solution as directed under the Ultraviolet-visible Spectrophotometry, and compare the spectrum with the Reference Spectrum: both spectra exhibit similar intensities of absorption at the same wavelength.

(2)  
Dissolve a suitable amount of Norfloxacin in a suitable amount of acetone, evaporate the acetone under reduced pressure, and dry the residue. Determine the infrared absorption spectrum of the residue so obtained as directed in the potassium bromide disk method under the Infrared Spectrophotometry, and compare the spectrum with the Reference Spectrum: both spectra exhibit similar intensities of absorption at the same wave numbers.

**Purity** (1)  
Sulfate—Dissolve 1.0 g of Norfloxacin in 7 mL of 0.5 mol/L sodium hydroxide TS and 23 mL of water, and add 1 drop of phenolphthalein TS. Add gradually diluted hydrochloric acid (1 in 3) to this solution until the red color disappears, then add 0.5 mL of dilute hydrochloric acid, and cool in ice for 30 minutes. Filter through a glass filter (G4), and wash the residue with 10 mL of water. Combine the filtrate and the washing, and add 1 mL of dilute hydrochloric acid and water to make 50 mL. Perform the test using this solution as the test solution. Prepare the control solution as follows. To 0.50 mL of 0.005 mol/L sulfuric acid VS add 7 mL of 0.5 mol/L sodium hydroxide TS and 1 drop of phenolphthalein TS, add diluted hydrochloric acid (1 in 3) until the red color disappears, then add 1.5 mL of dilute hydrochloric acid, 1 or 2 drops of bromophenol blue TS and water to make 50 mL (not more than 0.024%).

(2)  
Heavy metals—Proceed with 2.0 g of Norfloxacin according to Method 2, and perform the test. Prepare the control solution with 3.0 mL of Standard Lead Solution (not more than 15 ppm).

(3)  
Arsenic—Prepare the test solution with 1.0 g of Norfloxacin according to Method 3, and perform the test using Apparatus B (not more than 2 ppm).

(4)  
Related substances—Conduct this procedure without exposure to light, using light-resistant vessels. Dissolve 0.10 g of Norfloxacin in 50 mL of a mixture of methanol and acetone (1:1), and use this solution as the sample solution. Pipet 1 mL of the sample solution, add a mixture of methanol and acetone (1:1) to make exactly 100 mL. Pipet 2 mL of this solution, add a mixture of methanol and acetone (1:1) to make exactly 10 mL, and use this solution as the standard solution. Perform the test with these solutions as directed under the Thin-layer Chromatography. Spot 20 µL each of the sample solution and the standard solution on a plate of silica gel with fluorescent indicator for thin-layer chromatography (5 – 7 µm in particle diameter). Develop with a mixture of methanol, chloroform, toluene, diethylamine and water (20:20:10:7:4) to a distance of about 9 cm, and air-dry the plate. Examine under ultraviolet light (main wavelength: 254 nm and 366 nm): the number of the spot other than the principal spot from the sample solution is not more than 2 and they are not more intense than the spot from the standard solution.

**Loss on drying**  
Not more than 1.0% (1 g, 105°C, 2 hours).

**Residue on ignition**  
Not more than 0.10% (1 g).

**Assay**  
Weigh accurately about 0.5 g of Norfloxacin, previously dried, dissolve in 50 mL of acetic acid (100), and titrate with 0.1 mol/L perchloric acid VS (potentiometric titration). Perform a blank determination, and make any necessary correction.

Each mL of 0.1 mol/L perchloric acid VS = 31.933 mg of $\text{C}_{16}\text{H}_{18}\text{FN}_3\text{O}_3$

**Containers and storage**  
Containers—Tight containers. Storage—Light-resistant.