U.S. PHARMACOPEIA

| Search USP29 | Go |
|--------------|----|

Sodium Lauryl Sulfate

Sulfuric acid monododecyl ester sodium salt.

Sodium monododecyl sulfate [151-21-3].

» Sodium Lauryl Sulfate is a mixture of sodium alkyl sulfates consisting chiefly of sodium lauryl sulfate [CH₃(CH₂)₁₀CH₂OSO₃Na]. The combined content of sodium chloride and sodium sulfate is not more than 8.0 percent.

Packaging and storage— Preserve in well-closed containers.

Identification—

A: Ignite about 500 mg at 800° until the carbon is consumed: the residue dissolved in 10 mL of water responds to the tests for <u>Sodium</u> (191).

B: A solution (1 in 10) after acidification with hydrochloric acid and gentle boiling for 20 minutes, responds to the tests for <u>Sulfate</u> (191).

Alkalinity— Dissolve 1.0 g in 100 mL of water, add phenol red TS, and titrate with 0.10 N hydrochloric acid: not more than 0.60 mL is required for neutralization.

Heavy metals, Method II (231): 0.002%.

Organic volatile impurities, *Method IV* (467): meets the requirements.

Sodium chloride— Dissolve about 5 g, accurately weighed, in about 50 mL of water. Neutralize the solution with 0.8 N nitric acid, using litmus paper as the indicator, add 2 mL of potassium chromate TS, and titrate with 0.1 N silver nitrate VS. Each mL of 0.1 N silver nitrate is equivalent to 5.844 mg of NaCl.

Sodium sulfate—

Lead nitrate solution— Dissolve 33.1 g of lead nitrate in water to make 1000 mL.

Procedure— Transfer about 1 g of Sodium Lauryl Sulfate, accurately weighed, to a 250-mL beaker, add 35 mL of water, and warm to dissolve. To the warm solution add 2.0 mL of 1 N nitric acid, mix, and add 50 mL of alcohol. Heat the solution to boiling, and slowly add 10 mL of Lead nitrate solution, with stirring. Cover the beaker, simmer for 5 minutes, and allow to settle. If the supernatant is hazy, allow to stand for 10 minutes, heat to boiling, and allow to settle. When the solution is almost to a boiling point, decant as much liquid as possible through a 9-cm filter paper (Whatman No. 41 or equivalent). Wash four times by decantation, each time using 50 mL of 50 percent alcohol, and bring the mixture to a boil. Finally, transfer the filter paper to the original beaker, and immediately add 30 mL of water, 20.0 mL of 0.05 M edetate disodium VS, and 1 mL of ammonia-ammonium chloride buffer TS. Warm to dissolve the precipitate, add 0.2 mL of eriochrome black TS and titrate with 0.05 M zinc sulfate VS. Each mL of 0.05 M edetate disodium is equivalent to 7.102 mg of Na₂SO₄.

Unsulfated alcohols— Dissolve about 10 g, accurately weighed, in 100 mL of water, and add 100 mL of alcohol. Transfer the solution to a separator, and extract with three 50-mL portions of solvent hexane. If an emulsion forms, sodium chloride may be added to promote separation of the two layers. Wash the combined solvent hexane extracts with three 50-mL portions of water, and dry with anhydrous sodium sulfate. Filter the solvent hexane extract into a tared beaker, evaporate on a steam bath until the odor of solvent hexane no longer is perceptible, dry the residue at 105 of for 30 minutes, cool, and weigh. The weight of the residue is not more than 4.0% of the weight of the Sodium Lauryl Sulfate taken.

Total alcohols— Transfer about 5 g, accurately weighed, to an 800-mL Kjeldahl flask, and add 150 mL of hydrochloric acid, and a few boiling chips. Attach a reflux condenser to the Kjeldahl flask, heat carefully to avoid excessive frothing, and then boil for about 4 hours. Cool the flask, rinse the condenser with ether, collecting the ether in the flask, and transfer the contents to a 500-mL separator, rinsing the flask twice with ether and adding the washings to the separator. Extract the solution with two 75-mL portions of ether, evaporate the combined ether extracts in a tared beaker on a steam bath, dry the residue at 105 of 500 minutes, cool, and weigh. The residue represents the total alcohols, and is not less than 59.0% of the weight of Sodium Lauryl Sulfate taken.

Residual solvents (467): meets the requirements. (Official January 1, 2007)

Auxiliary Information— Staff Liaison: Catherine Sheehan, B.Sc., Scientist

Expert Committee: (EM105) Excipient Monographs 1

USP29-NF24 Page 3424

Phone Number: 1-301-816-8262