- (2) Water-soluble substances—To 5 g of Purified Absorbent Cotton add 500 mL of water, and boil gently for 30 minutes, while adding water to maintain the original volume. Pour the extract through a funnel into another vessel, transfer the cotton to the funnel, and press out the water absorbed therein with a glass rod. Wash the cotton with two 150-mL portions of hot water, pressing the cotton after each washing. Filter the combined extracts and washings. Evaporate to concentrate the filtrate, transfer to a weighing bottle, and dry at 105°C to constant mass: the amount of the residue is not more than 14.0 mg. Perform a blank determination, and make any necessary correction.
- (3) Dyes—Digest 10 g of Purified Absorbent Cotton with 100 mL of ethanol (95), press out, and transfer 50 mL of the extracts to a Nessler tube. Observe downward: a yellow color develops, but neither a blue nor a green color develops.
- (4) Fluorescent whitening agents—Irradiate Purified Absorbent Cotton under ultraviolet rays in a dark place: no fluorescence is perceptible on the surface.
- (5) Submersion rate—Prepare a test basket, weighing 3.0 g, form copper wire 0.44 mm in diameter in the form of a cylinder 50.0 mm in diameter and 80.0 mm in depth, with spaces of 20 mm between the wires. Place 5 g of Purified Absorbent Cotton in the basket, hold the basket on its side 12 mm above the surface of water between 24°C and 26°C, and drop the basket gently into the water, which is 200 mm deep: the time required for complete submersion is not more than 8 seconds.
- (6) Absorbency—Leave the submerged basket at the bottom of the water in (5) as it is for 3 minutes. Lift the basket gently from the water, keeping its side horizontal, and allow to drain for 1 minute on the wire gauze of a sieve No. 10 in the same horizontal position. Then place in a beaker and weigh: the mass of water absorbed is not less than 100.0 g.
- (7) Other filaments—Dip 1.0 g of Purified Absorbent Cotton in 0.5 mol/L iodine TS for 1 minute, and wash well with water: no colored filament is found.
- (8) Neps and adhering impurities—Spread evenly about 1 g of Purified Absorbent Cotton between two 10 cm-square, colorless, transparent plates, and examine neps and adhering impurities (fragments of rinds and seeds): the total number of the fragments more than 2.5 mm in diameter is not more than 5.
- (9) Short fibers—Take 0.10 g of Purified Absorbent Cotton, separate the fibers into two groups, one consisting of fibers not exceeding 6.0 mm in length (short fibers) and the other consisting of fibers exceeding 6.0 mm in length, weigh both groups, and determine the content (%) of the short fibers: not more than 10%.

Percentage (%) of the short fibers
$$= \frac{W_2}{W_1 + W_2} \times 100$$

W₁: Mass of the group of fibers exceeding 6.0 mm in length

 W_2 : Mass of the group of fibers not exceeding 6.0 mm in length

Total ash Not more than 0.25% (5 g, proceed as directed in the Total ash under Crude Drugs).

Containers and storage Containers—Well-closed containers.

Sterile Absorbent Cotton

滅菌脱脂綿

Sterile Absorbent Cotton is sterilized Absorbent Cotton.

Description Apply the Description under Absorbent Cotton

Purity Proceed as directed in the Purity under Absorbent Cotton.

Total ash Proceed as directed in the Total ash under Absorbent Cotton.

Sterility Take Sterile Absorbent Cotton from package abacterially under an aseptic circumstances, sample about 0.5 g of it (whole content in the case of less than 0.5 g) evenly from 5 different parts around the center portion, put the samples in a test tube of 25 mm × 200 mm containing 60 mL each of fluid thioglycollate medium I for the Sterility Test for the growth of bacteria and fungi: it meets the requirements of the Sterility Test. In the case of the test for the growth of fungi, a 200-mL Erlenmeyer flask can also be used. In this connection, perform an efficient test of the medium under a condition without the samples: the medium supports the substantial growth of the incubated microorganisms.

Sample number used in the Sterility Test is indicated in the following table.

Number of products of the same kind sterilized simultaneously	Number of products used for test		
Not more than 100	4		
100 to not more than 500	10		
Not less than 500	20		

Containers and storage Containers—Tight containers impervious to any microbe.

Sterile Purified Absorbent Cotton

滅菌精製脱脂綿

Sterile Purified Absorbent Cotton is sterilized Purified Absorbent Cotton.

Description Apply the Description under Purified Absorbent Cotton.

Purity Proceed as directed in the Purity under Purified Absorbent Cotton.

Total ash Proceed as directed in the Total ash under Purified Absorbent Cotton.

Sterility Take Sterile Purified Absorbent Cotton from package abacterially under an aseptic circumstances, sample about 0.5 g of it (whole content in the case of less than 0.5 g) evenly from 5 different parts around the center portion, put

the samples in a test tube of $25 \text{ mm} \times 200 \text{ mm}$ containing 60 mL each of fluid thioglycollate medium I for the Sterility Test for the growth of bacteria and fungi: it meets the requirements of the Sterility Test. In the case of the test for the growth of fungi, a 200-mL Erlenmeyer flask can also be used. In this connection, perform an efficient test of the medium under a condition without the samples: the medium supports the substantial growth of the incubated microorganisms.

Sample number used in the Sterility Test is indicated in the following table.

Number of products of the same kind sterilized simultaneously	Number of products used for test		
Not more than 100			
100 to not more than 500	10		
Not less than 500	20		

Containers and storage Containers—Tight containers impervious to any microbe.

Absorbent Gauze

ガーゼ

Absorbent Gauze consists of non-fatty and well-bleached cotton cloth of plain weave using pure cotton threads obtained from hairs of the seed of *Gossypium hirsutum* Linné or other species of the same genus (*Malvaceae*). The amount of Absorbent Gauze is expressed in terms of its type, length and width.

Description Absorbent Gauze occurs as white cotton cloth. It is odorless and tasteless.

- Purity (1) Water-soluble substances—Place 20 g of Absorbent Gauze in 500 mL of water, and boil gently for 15 minutes, while adding water to maintain the original volume. Pour the extract through a funnel into a 1000-mL flask, transfer the Absorbent Gauze to the funnel, press out the water absorbed therein with a glass rod, and wash Absorbent Gauze with two 250-mL portions of boiling water, pressing after each washing. Combine the extract and the washings, filter, and add water to make 1000 mL. Transfer 400 mL of the filtrate to a beaker, evaporate to concentrate, and place the residue in a weighing bottle. Wash the beaker with a small amount of water, combine the washings with the residue in the weighing bottle, and dry at 105°C to constant mass: the mass of the residue is not more than 20.0 mg. Perform a blank determination, and make any necessary correction.
- (2) Acid or alkali—To 200 mL of the extract obtained in (1), add 5 drops of phenolphthalein TS: no red color develops. To 200 mL of the sample solution add 2 drops of methyl orange TS: no red color develops.
- (3) Dextrin or starch—To 200 mL of the extract obtained in (1) add 2 drops of iodine TS: no red-purple to blue color develops.
- (4) Dyes—Digest 10 g of Absorbent Gauze with 80 mL of ethanol (95), press out, and transfer 50 mL of the extracts to a Nessler tube. Observe downward: a yellow color de-

velops, but neither a blue nor a green color develops.

- (5) Fluorescent whitening agents—Irradiate Absorbent Gauze with ultraviolet rays in a dark place: no fluorescence is perceptible on the surface.
- (6) Submersion rate—Prepare a test basket from copper wire 0.4 mm in diameter (No. 26 wire) in the form of a cylinder 50.0 mm in diameter and 80.0 mm in depth, with spaces of 20 mm between the wires, the basket weighing 3.0 g. Place 10 g of Absorbent Gauze evenly in the basket, hold the basket on its side 12 mm above the surface of water between 24°C and 26°C, and drop the basket gently into the water, which is 200 mm deep: the time required for complete submersion is not more than 8 seconds.
- (7) Other filaments—Dip 1.0 g of Absorbent Gauze in 0.5 mol/L iodine TS for 1 minute, and wash well with water: no colored filament is found.

Texture The texture requirements of Absorbent Gauze are given in the following table.

Туре	Threads per 1 cm (number)			Tolerance in average counts of	Standard	Standard	
	Warp		Filling		threads per 1 cm	width (cm)	mass (g)
	Average	Tolerance	Average	Tolerance	× 1 cm		
I	12	±1	12	±1	24 + 2	30 + 0.5 - 1.0	width 30 cm length 100 cm 10.3 + 8%
II	12	± 1	12	± 1	24 ^{+ 2} _{- 1}	91.4 + 1.5	width 91.4 cm length 30 cm 8.7 + 8%
III	11	± 1	9	± 1	20 + 2	91.4 + 1.5	width 91.4 cm length 30 cm 7.6 + 8%
IV	9	± 1	8	± 1	17 + 2	91.4 + 1.5	width 91.4 cm length 30 cm 6.1 + 8%

Length: Place Absorbent Gauze on a flat plate, eliminate the unnatural creases or tensions, and measure the full length at the center line: the length is not less than 95% of the labeled length. When it has closely woven parts at both edges in the direction of the length, measure the full length. When it has no closely woven parts, measure only the net.

Width: Place Absorbent Gauze on a flat plate, smooth out the unnatural creases or tensions, and measure the full width at more than 3 different locations: the average of these measurements is not less than 80% and not more than 120% of the labeled width which is not more than 5 cm, not more than -1.0 cm and not less than +0.5 cm of the labeled width which is between 5 cm and 30 cm, and within ± 1.5 cm of the labeled width which is above 30 cm. When it has closely woven parts at both edges in the direction of the width, measure the full width. When it has no closely woven parts, measure only the net.

Number of threads: Prepare a frame of 1 cm \times 1 cm, and set the thread to the edge of the frame. Count the integral number of the threads in the frame and average the results of more than 3 counts. Except the closely woven parts.

Mass: Fold Absorbent Gauze into about a 10-cm square, allow to stand at ordinary temperature for 4 hours in a desiccator, previously saturated with the vapor above a saturated solution of sodium nitrite, and weigh. For pieces in various