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**Urine-absorbing aids for  
incontinence — Polyacrylate  
superabsorbent powders —**

**Part 6:**

**Test method for determination of  
the fluid retention capacity in saline  
solution by gravimetric measurement  
following centrifugation**

*Aides pour absorption d'urine — Méthodes d'essai pour caractériser  
les matériaux absorbants à base de polymères —*

*Partie 6: Détermination gravimétrique de la capacité de rétention de  
fluides en solution saline après centrifugation*

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## Foreword

ISO (the International Organization for Standardization) is a worldwide federation of national standards bodies (ISO member bodies). The work of preparing International Standards is normally carried out through ISO technical committees. Each member body interested in a subject for which a technical committee has been established has the right to be represented on that committee. International organizations, governmental and non-governmental, in liaison with ISO, also take part in the work. ISO collaborates closely with the International Electrotechnical Commission (IEC) on all matters of electrotechnical standardization.

The procedures used to develop this document and those intended for its further maintenance are described in the ISO/IEC Directives, Part 1. In particular, the different approval criteria needed for the different types of ISO documents should be noted. This document was drafted in accordance with the editorial rules of the ISO/IEC Directives, Part 2 (see [www.iso.org/directives](http://www.iso.org/directives)).

Attention is drawn to the possibility that some of the elements of this document may be the subject of patent rights. ISO shall not be held responsible for identifying any or all such patent rights. Details of any patent rights identified during the development of the document will be in the Introduction and/or on the ISO list of patent declarations received (see [www.iso.org/patents](http://www.iso.org/patents)).

Any trade name used in this document is information given for the convenience of users and does not constitute an endorsement.

For an explanation of the voluntary nature of standards, the meaning of ISO specific terms and expressions related to conformity assessment, as well as information about ISO's adherence to the World Trade Organization (WTO) principles in the Technical Barriers to Trade (TBT) see [www.iso.org/iso/foreword.html](http://www.iso.org/iso/foreword.html).

This document was prepared by Technical Committee ISO/TC 173, *Assistive products*, Subcommittee SC 3, *Aids for ostomy and incontinence*.

This second edition cancels and replaces the first edition (ISO 17190-6:2001), which has been technically revised. The main changes compared to the previous edition are as follows:

- full text review and new laboratory analysis with statistical evaluation;
- sample weighing simplified;
- request for duplication removed.

A list of all parts in the ISO 17190 series can be found on the ISO website.

Any feedback or questions on this document should be directed to the user's national standards body. A complete listing of these bodies can be found at [www.iso.org/members.html](http://www.iso.org/members.html).

# Urine-absorbing aids for incontinence — Polyacrylate superabsorbent powders —

## Part 6:

## Test method for determination of the fluid retention capacity in saline solution by gravimetric measurement following centrifugation

**WARNING** — - This document does not claim to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this document to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use. It is expected that the person performing this test has been fully trained in all aspects of this procedure.

### 1 Scope

This document provides a test method for the determination of the fluid retention capacity of polyacrylate superabsorbent powders in saline solution, following centrifugation.

### 2 Normative references

The following documents are referred to in the text in such a way that some or all of their content constitutes requirements of this document. For dated references, only the edition cited applies. For undated references, the latest edition of the referenced document (including any amendments) applies.

ISO 187, *Paper, board and pulps — Standard atmosphere for conditioning and testing and procedure for monitoring the atmosphere and conditioning of samples*

ISO 3696, *Water for analytical laboratory use — Specification and test methods*

### 3 Terms and definitions

For the purposes of this document, the following terms and definitions apply.

ISO and IEC maintain terminological databases for use in standardization at the following addresses:

- ISO Online browsing platform: available at <https://www.iso.org/obp>
- IEC Electropedia: available at <http://www.electropedia.org/>

#### 3.1

##### **sample**

product or portion of a product taken from a production lot for testing purposes and identifiable and traceable back to its origin

#### 3.2

##### **specimen**

specific portion of the identified *sample* (3.1) upon which a test is performed

## 4 Principle

The sample is weighed and placed in a bag. The bag is submerged in the fluid to be absorbed and afterwards centrifuged for a specified time, at a specified centrifugal force, to determine the amount of fluid retained.

## 5 Reagents and materials

### 5.1 Water.

Grade 1 water in accordance with ISO 3696, with the exception that the conductivity can be as high as 30  $\mu\text{S}/\text{cm}$ .

### 5.2 Sodium chloride solution.

**5.2.1** 0,9 % mass fraction of sodium chloride solution in water. Weigh  $(9,00 \pm 0,01)$  g of sodium chloride into a 1 l beaker and add  $(991,0 \pm 0,1)$  g of deionized water (grade 3). Stir until dissolved.

**5.2.2** The conductivity of the solution should be checked prior to each use using properly calibrated measuring equipment. The expected conductivity of a 0,9 % saline solution is of the order of 1600 S/m at 25 °C. Each testing lab shall determine the correct conductivity for the conditions obtaining in the lab.

It is also recommended that the temperature of the solution be maintained at  $(23 \pm 2)$  °C for the duration of the test. As this matches the required laboratory temperature it is not necessary to record the solution temperature.

### 5.3 Nonwoven bag.

NOTE Bags described in this document are often referred to as “teabags”.

The bag has the external dimensions of  $(60 \times 40)$  mm<sup>2</sup> to  $(60 \times 85)$  mm<sup>2</sup> and made of non-apertured heat-sealable nonwoven. One example of a suitable specification for the nonwoven is the following:

- Mass per unit area:  $(16,5 \pm 1,5)$  g/m<sup>2</sup>;
- Thermoplastic fibre content:  $(4,0 \pm 0,8)$  g/m<sup>2</sup>;
- Web tensile strength in cross direction:  $(70 \pm 12)$  N/m;
- Air permeability (4 plies) –  $2,30 \pm 0,50$  l.min<sup>-1</sup>.cm<sup>-2</sup> at a pressure drop of 124 Pa.

Normally the teabag paper is supplied as a roll, for example 120 mm wide. This shall be stored flat as storing in an upright position (as a wheel) will compress the paper at bottom of the roll and affect its characteristics.

The teabag is made by cutting sections, for example 60 mm wide, which will provide teabags  $60 \times 60$  mm<sup>2</sup> when folded and sealed. In this example, the paper may be folded in half and sealed along two sides ready for the addition of superabsorbent powder and the sealing of the final side prior to running the test.

It is recommended to have teabags made in bulk and delivered ready-made for use in the lab.

In any case, teabags should be stored in cool, dry conditions. For best practice storing teabags in a desiccator prior to use is also recommended.