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Absorbent structure and absorbent article comprising the absorbent structure

57	ABSTRACT (NOT MORE THAN 150 WORDS)
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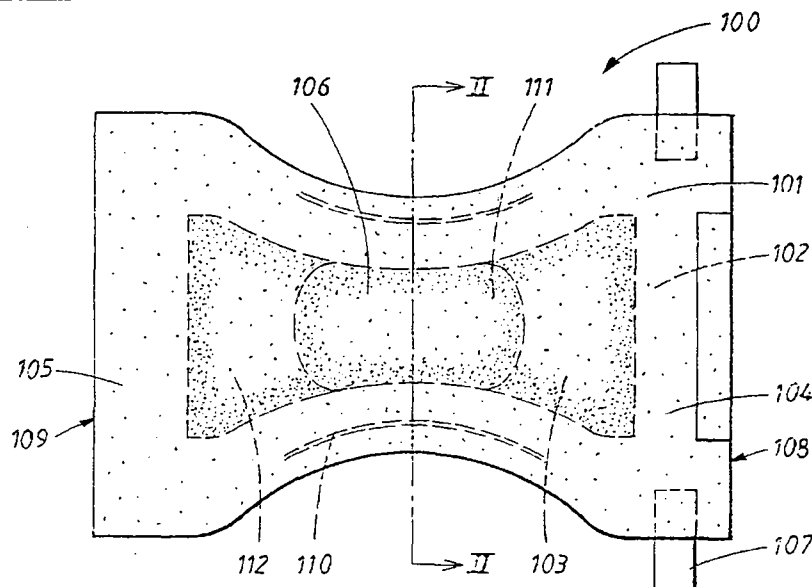
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(54) Title: ABSORBENT STRUCTURE AND ABSORBENT ARTICLE COMPRISING THE ABSORBENT STRUCTURE



Abstract: The invention relates to an absorbent structure (103; 303) for use in a diaper (100; 300), an incontinence pad, a sanitary towel or the like, which absorbent structure (103; 303) has at least one absorbent layer (112; 311; 312) comprising fluff pulp and superabsorbent particles. The average absorption capacity per superabsorbent particle in the absorbent layer (112; 311; 312) is greater than 8.0 mg sodium chloride solution, and the number of superabsorbent particles per cm³ of the absorbent layer (112; 311; 312) is smaller than 1100. The invention also relates to a diaper (100; 300), an incontinence pad, a sanitary towel or the like, which comprises an absorbent structure (103; 303) as above.

ABSORBENT STRUCTURE AND ABSORBENT ARTICLE COMPRISING THE ABSORBENT STRUCTURE

TECHNICAL FIELD

- 5 The present invention relates to an absorbent structure for use in an absorbent article such as a diaper, an incontinence pad, a sanitary towel or the like, which absorbent structure has at least one absorbent layer comprising fluff pulp and superabsorbent particles.

10 BACKGROUND ART

- An absorbent structure for disposable absorbent articles such as diapers, incontinence pads and sanitary towels is usually constructed from one or more layers of hydrophilic fibres, for example cellulose fluff pulp. In order to obtain high absorption capacity and also a high liquid-retaining capacity when
15 the article is subjected to external loading, the absorbent structure usually contains superabsorbent particles, which are polymers with the ability to absorb many times their own weight of water or body fluid. The effectiveness of the superabsorbent depends on many factors such as, for example, the physical shape of the superabsorbent particles. Other examples of properties
20 which influence the functioning of the superabsorbent are absorption rate, gel strength and liquid-retaining capacity.

- The absorbent structure can also contain other components, for example in order to improve its liquid-spreading properties or increase its cohesive
25 capacity and ability to withstand deformation during use.

- It is of considerable importance that the absorbent article is capable of rapidly receiving and absorbing large quantities of liquid. It is also of considerable importance that the total absorption capacity of the article can be utilized. In
30 order for it to be possible to utilize the total absorption capacity of the article, it is essential that the liquid can be spread from the wetting area to other parts of the absorbent structure.

One problem, above all for diapers and incontinence pads which are intended to receive and absorb relatively large quantities of liquid, is that there is a risk of them leaking before their total absorption capacity is fully utilized. One cause of leakage is that the absorbent structure, in particular
5 when repeated wetting takes place, has an impaired ability to receive and absorb large quantities of liquid rapidly. A major cause of it being difficult for the absorbent structure to function satisfactorily when repeated wetting takes place, that is to say a second wetting and subsequent wettings, is that the superabsorbent material in a swollen state can block the pores in the porous
10 fibrous structure and thus interfere with the transport of liquid from the wet area out to other parts of the absorbent structure. This phenomenon is referred to as gel blocking and results in the total absorption capacity of the absorbent structure not being utilized optimally. It also leads to an increased risk of leakage.

15

The problem of gel blocking increases when the proportion of superabsorbent material in an absorbent structure is high. In order to obtain an article which is discreet and comfortable to wear, however, it is an advantage to have a thin article which contains a relatively high proportion of
20 superabsorbent material.

DISCLOSURE OF INVENTION

The problem of gel blocking during use of thin absorbent articles having a relatively high content of superabsorbent material has been reduced by
25 means of the present invention.

An absorbent structure according to the invention is characterized mainly in that the average absorption capacity per superabsorbent particle in the absorbent layer is greater than 8.0 mg and in that the number of
30 superabsorbent particles per cm^3 of the absorbent layer is smaller than 1100. The absorption capacity is measured using 0.9% by weight sodium chloride solution.

By limiting the number of superabsorbent particles per unit of volume, it has been found that it is possible to maintain a fibrous network with a pore structure which can transport liquid in the absorbent structure even after the structure has been subjected to a first wetting. It has also been found that, with a limited number of superabsorbent particles per unit of volume, it is essential that the average absorption capacity per superabsorbent particle is greater than 8.0 mg. The advantage of such an absorbent structure is that the risk of gel blocking decreases at the same time as it is possible to obtain a thin absorbent structure.

10

According to one embodiment, the average absorption capacity per superabsorbent particle in the absorbent layer is greater than 9.5 mg. The absorption capacity is measured using 0.9% by weight sodium chloride solution. Furthermore, the number of superabsorbent particles per cm³ of the absorbent layer is smaller than 600.

15

According to another embodiment, the average absorption capacity per superabsorbent particle in the absorbent layer is greater than 14.0 mg. The absorption capacity is measured using 0.9% by weight sodium chloride solution. In such an embodiment, the number of superabsorbent particles per cm³ of the absorbent layer is smaller than 450.

20

According to one embodiment, the superabsorbent particles have a particle size which is greater than 600 µm. The superabsorbent particles are preferably polyacrylate-based. In order to obtain a high absorption capacity, it is also possible to change the morphology of the superabsorbent particles. An example of superabsorbent particles with a changed morphological structure is microporous superabsorbent particles. A high absorption capacity can also be obtained by means of a special chemical composition of the superabsorbent particles.

25

30

The superabsorbent particles can be surface cross-linked or have a gradually increasing cross-linking towards the surface of the particles. A surface cross-linked superabsorbent is cross-linked in two different steps. First, the polymer is cross-linked so that a homogeneous cross-linked gel is formed. In cases
5 where polymerization and cross-linking do not result in particles being formed simultaneously, particles are produced in a following process step. In another following process step, the formed particles are cross-linked in the second step, but then only partly. The additional cross-linking can be effected so that there is a higher cross-linker content next to the surface of the particle
10 compared with the centre of the particle. In this way, a more firmly cross-linked particle shell is produced, which surrounds a particle core with a lower degree of cross-linking.

Superabsorbents with a low degree of cross-linking provide a high absorption
15 capacity. However, a problem with such superabsorbents is that, in a swollen state, they are soft and sticky, which results in the risk of gel blocking in the absorbent structure already being high at a low superabsorbent material content. Superabsorbents with a high degree of cross-linking keep their shape better in a swollen state and do not stick to the same great extent
20 either. However, a problem with a superabsorbent with a high degree of cross-linking is that it has a considerably lower absorption capacity. So, by surface cross-linking the superabsorbent, or alternatively creating a cross-linking gradient so that the particle surface is cross-linked more firmly than the inner particle core, a superabsorbent is obtained which both has high
25 absorption capacity and essentially maintains its shape in a swollen state.

According to one embodiment of an absorbent structure according to the invention, the average distance between centres of the superabsorbent particles in the absorbent layer in a dry state is greater than 700
30 micrometres, more preferably greater than 1000 micrometres and even better greater than 1200 micrometres. The average centre-centre distance (l_{cc}) of the superabsorbent particles is obtained using the following equation:

$$l_{cc} = (1/n)^{1/3}$$

n = number of superabsorbent particles per unit of volume of material

According to another embodiment, the density of the absorbent layer is greater than 0.12 g/cm³, more preferably greater than 0.17 g/cm³ and even better greater than 0.25 g/cm³. The absorbent layer can moreover comprise bonding means, such as bonding fibres for example. Examples of bonding fibres are synthetic fibres made of polyolefin. In order to function as bonding fibres, the fibres are heated to their melting point, the fibres being bonded to the material in the absorbent layer. Bonding fibres made of bicomponent fibres are common. If bicomponent fibres are used as bonding fibres, one component is melted while the other component is intact, that is to say does not melt but instead maintains the structure of the fibre.

The invention also relates to an absorbent article such as a diaper, an incontinence pad, a sanitary towel or the like, which comprises an absorbent structure according to any one of the embodiments described.

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 shows a diaper according to the invention, seen from the side which is intended to lie against the wearer during use;

Figure 2 shows a cross section along the line II-II through the diaper shown in Figure 1;

25

Figure 3 shows a cross section of an alternative embodiment of an absorbent article according to the invention.

DETAILED DESCRIPTION OF THE FIGURES

30 The diaper 100 shown in Figure 1 comprises a liquid-permeable surface layer 101, a backing layer 102, which is at least essentially liquid-impermeable,

and an absorbent structure 103 enclosed between the liquid-permeable surface layer 101 and the backing layer 102.

5 The diaper is intended to surround the lower part of the abdomen of a wearer like a pair of absorbent underpants. To this end, it is shaped with a rear portion 104 and a front portion 105, and a narrower crotch portion 106 which is located between the front portion 105 and the rear portion 104 and is intended during use to be arranged in the crotch of the wearer between the legs of the latter. In order that it is possible for the diaper to be fastened
10 together in the desired pants-shape, tape tabs 107 are arranged close to the rear waist edge 108 of the diaper. During use, the tape tabs 107 are fastened to the front portion 105 of the diaper, close to the front waist edge 109, so that the diaper is held together around the waist of the wearer. Other fastening devices are of course also possible, such as hook and loop
15 fastening for example.

The diaper 100 according to Figure 1 also comprises pretensioned elastic means 110 which can consist of elastic bands, thread-covered elastic threads, elastic foam or another suitable material. For the sake of simplicity,
20 the elastic means 110 have in Figure 1 been shown in the stretched state. As soon as stretching stops, however, they contract and form elastic leg-bands of the diaper.

The liquid-permeable surface layer 101 is, for example, a nonwoven material
25 or a perforated film, or a laminate thereof. Examples of polymers from which the liquid-permeable surface layer 101 can be made are polyethylene, polypropylene, polyester or copolymers thereof. In order that the liquid-permeable surface layer 101 will allow the discharged body fluid to pass through rapidly, it is common for the surface layer to be surfactant-coated
30 and/or perforated. Another suitable material for use as the liquid-permeable surface layer is a layer of continuous fibres which are interconnected in a spot, line or patch bonding pattern but are otherwise on the whole not

bonded to one another. The backing layer 102 is, for example, a plastic film, which is preferably breathable, a hydrophobic nonwoven layer or a laminate thereof.

5 The absorbent structure 103 of the diaper 100 is constructed from an upper liquid-receiving layer 111 and a lower liquid-distribution and storage layer 112. The lower liquid-distribution and storage layer 112 has a greater extent in the plane of the article than the upper liquid-receiving layer 111. The upper receiving layer 111 is to be capable of rapidly receiving large quantities of
10 liquid in a short time, that is to say have a high instantaneous liquid absorption capacity, while the lower distribution and storage layer 112 is to have a high wicking capacity and high storage capacity and to be capable of draining liquid from the receiving layer 111. The lower distribution and storage layer 112 in the absorbent structure 103 consists of an absorbent
15 layer according to the invention. The lower liquid-distribution and storage layer 112 therefore also comprises superabsorbent particles in addition to cellulose fluff pulp. The average absorption capacity per superabsorbent particle in the liquid-distribution and storage layer 112 is greater than 8.0 mg sodium chloride solution. Furthermore, the number of superabsorbent
20 particles per cm^3 of the liquid-spreading and storage layer 112 is smaller than 1100. According to one embodiment, the average absorption capacity per superabsorbent particle in the liquid-distribution and storage layer 112 is greater than 9.5 mg, and the number of superabsorbent particles per cm^3 of the liquid-distribution and storage layer is smaller than 600. According to
25 another example, the average absorption capacity per superabsorbent particle in the liquid-distribution and storage layer 112 is greater than 14.0 mg, and the number of superabsorbent particles per cm^3 of the liquid-distribution and storage layer is smaller than 450. The absorption capacity is measured throughout using 0.9% by weight sodium chloride solution.

30

The average distance between centres of the superabsorbent particles in the liquid-distribution and storage layer 112 in a dry state is, for example, greater

than 700 micrometres, preferably greater than 1000 micrometres and even better greater than 1200 micrometres. The density of the absorbent structure in the liquid-distribution and storage layer 112 is, for example, greater than 0.12 g/cm³, preferably greater than 0.17 g/cm³ and even better greater than 0.25 g/cm³.

Suitable materials for use as the receiving layer 111 include, for example, an open nonwoven layer made of synthetic or natural fibres. A difference in properties between the liquid-distribution and storage layer 112 and the receiving layer 111 can be brought about by, for example, the liquid-distribution and storage layer 112 being compressed more firmly than the receiving layer 111. A firmly compressed fibrous structure with high density spreads the liquid better than a corresponding fibrous structure with lower density, which, because of its larger pore size, has a higher instantaneous liquid absorption capacity but lower wicking capacity. Differences in absorption properties between the two layers can also be brought about by means of different fibrous structures with different properties. Accordingly, cellulose fluff pulp produced in a conventional chemical way, chemical pulp (CP), has higher liquid-wicking capacity compared with pulp produced in a mechanical or chemithermomechanical way. Therefore, cellulose fluff pulp produced in a conventional chemical way, chemical pulp (CP), is an example of a suitable material for the liquid-distribution and storage layer 112, and pulp produced in a mechanical or chemithermomechanical way is an example of a material for the receiving layer 111. A fibrous structure containing chemically stiffened cellulose fibres also has a higher instantaneous liquid absorption capacity but lower spreading capacity than conventional chemical pulp and is therefore an example of a material for the receiving layer 111. Another suitable material for use as the receiving layer 111 is a superabsorbent foam, for example a polyacrylate-based foam. A polyacrylate-based foam is produced by a solution which consists of at least monomer, cross-linker, initiator and surfactant being saturated and pressurized with carbon dioxide in a vessel while being stirred. When the

solution is removed from the vessel through a nozzle, the solution expands and a foamed structure is obtained. The foamed structure is then locked by polymerization and cross-linking being initiated by, for example, UV radiation. Finally, the material is compressed and dried. On wetting, such a
5 superabsorbent foam expands greatly, which results in it being capable of receiving a large quantity of liquid in a short time. Such a receiving layer can consist of, for example, a continuous layer, which is positioned at least in the crotch portion of the article, or alternatively of a number of strips with hollow spaces between the strips. The receiving layer can also consist of a fibrous
10 layer with superabsorbent particles or a superabsorbent coating bonded to the fibrous layer.

In order to reduce the occurrence of undesirable bacterial growth and problems with odour, the absorbent structure 103 and/or the liquid-permeable
15 surface layer 101 can comprise bacteria-inhibiting and/or odour-inhibiting substances. An example of a bacteria-inhibiting and odour-inhibiting substance is a superabsorbent material which has a lower pH than a conventional superabsorbent. A superabsorbent material with a lower pH than a conventional superabsorbent has a lower degree of neutralization than
20 a conventional superabsorbent, the degree of neutralization being, for example, between 20 and 60%. The superabsorbent particles according to the invention can be, for example, a superabsorbent with a degree of neutralization between 20 and 60%.

25 Figure 2 shows a cross section along the line II-II through the diaper 100 shown in Figure 1. The diaper 100 shown in Figure 2 therefore has a liquid-permeable surface layer 101, a backing layer 102, and an absorbent structure 103 enclosed between the liquid-permeable surface layer 101 and the backing layer 102.

30

The absorbent structure 103 of the diaper is constructed from an upper liquid-receiving layer 111 and a lower liquid-distribution and storage layer

112. The lower liquid-distribution and storage layer 112 in the absorbent structure 103 consists of an absorbent layer according to the invention. The liquid-distribution and storage layer 112 therefore also comprises superabsorbent particles in addition to cellulose fluff pulp. The average
5 absorption capacity per superabsorbent particle in the liquid-spreading and storage layer 112 is greater than 8.0 mg sodium chloride solution, and the number of superabsorbent particles per cm^3 of the liquid-spreading and storage layer 112 is smaller than 1100.

10 Figure 3 shows a cross section of an alternative embodiment of an absorbent article according to the invention. The diaper 300 shown in Figure 3 is essentially constructed in the same way as the diaper in Figure 2. The diaper 300 therefore has a liquid-permeable surface layer 301, a backing layer 302, and an absorbent structure 303 enclosed between the liquid-permeable
15 surface layer 301 and the backing layer 302.

The absorbent structure 303 of the diaper is constructed from an upper liquid-receiving layer 311 and a lower liquid-distribution and storage layer 312. Both the upper liquid-receiving layer 311 and the lower liquid-distribution
20 and storage layer 312 in the absorbent structure 303 consist of absorbent layers according to the invention. The upper liquid-receiving layer 311 and the lower liquid-distribution and storage layer 312 therefore consist of fibrous structures which comprise superabsorbent particles, the average absorption capacity per superabsorbent particle in both layers 311, 312 being greater
25 than 8.0 mg sodium chloride solution. Furthermore, the number of superabsorbent particles per cm^3 of the absorbent structure in both layers is smaller than 1100. In the absorbent structure 303, both the upper liquid-receiving layer 311 and the lower liquid-distribution and storage layer 312 therefore consist of absorbent layers according to the invention.

30

The invention is of course not limited to the illustrative embodiments above but can of course be applied to other embodiments within the scope of the

patent claims. The invention therefore also comprises incontinence pads, pant diapers, sanitary towels, panty liners and the like. The invention also includes belt-supported diapers.

5 It is furthermore possible, for example, for the whole of the absorbent structure to consist of only one absorbent layer, in which case the whole absorbent structure consists of an absorbent layer according to the invention. According to another example, the absorbent structure can consist of a multilayer structure where the upper liquid-receiving layer consists of an
10 absorbent layer according to the invention. The liquid-receiving layer therefore comprises fibres and superabsorbent particles, the average absorption capacity for the superabsorbent particles in the liquid-receiving layer being greater than 8.0 mg sodium chloride solution (% by weight of sodium chloride is 0.9%), and the number of superabsorbent particles per
15 cm³ of the liquid-receiving layer being smaller than 1100. A cellulose fluff pulp with superabsorbent material of conventional type, for example, is used as the liquid-distribution and storage layer. It is also possible for the liquid-distribution and storage layer to consist of several different layers, at least one of the layers containing superabsorbent material and, for example, one
20 layer consisting of pure pulp in order to obtain good liquid distribution. Different layers can moreover have a difference in concentration of superabsorbent material, a liquid-distribution and storage layer with a gradually increasing/decreasing content of superabsorbent material being obtained. It is also possible for the liquid-distribution and storage layer to
25 consist of or comprise a layer made of a superabsorbent foam.

It is furthermore also possible for the absorbent structure to include one or more tissue layers or types of material or component other than those described above. The design of the layers can also vary. For example, one or
30 more layers in the absorbent structure can have cut-outs, that is to say cavities. The cut-outs extend, for example, in the longitudinal direction of the

absorption structure. It is of course also possible to have other physical designs of cut-out.

Example 1 – Determining volume and density of the absorbent layer.

- 5 When measuring the volume (cm^3) of the absorbent layer in an absorbent article, the absorbent layer is separated from the rest of the material in the article. If the absorbent structure has several different absorbent layers with mutually different properties, the various absorbent layers are also separated from one another, after which volume and density are measured for each
- 10 absorbent layer.

The absorbent layer is then weighed, and the thickness of the absorbent layer is measured. When measuring the thickness, use is made of a thickness gauge which has a circular foot with a diameter of 80 mm. The foot

15 is to exert a pressure of 0.5 kPa on the absorbent layer. The thickness is measured at five different points, which are distributed uniformly over the surface of the absorbent layer. The average value from these five measuring points represents the thickness of the absorbent layer in the volume calculation. The area of the absorbent layer is then measured, the volume

20 being obtained by multiplying the thickness by the area. The density of the absorbent layer is then obtained by dividing the weight of the absorbent layer by the volume.

Example 2 – Determining the number of superabsorbent particles per unit of

25 volume and measuring the absorption capacity of the superabsorbent particles.

The example is based on an absorbent layer which contains the superabsorbent particles. In this connection, it is also described how the

30 superabsorbent particles are to be separated from the pulp structure. It is of utmost importance that no material is lost in the handling described below. Ensure therefore that necessary measures are taken to avoid loss.

The absorbent layer is first separated from the rest of the material in the article. The superabsorbent particles are then separated from the fluff pulp in the absorbent layer by finely dividing the layer, that is to say tearing it into small pieces, and then shaking the superabsorbent particles out of the pulp structure. It is also possible to use an apparatus for separating the superabsorbent particles from the pulp structure. If an apparatus is used for separating the superabsorbent particles from the pulp structure, however, it is a condition that the superabsorbent particles are not damaged mechanically.

5 The moisture content of the superabsorbent particles is to be less than 5.0%. All indications in the present invention relate to superabsorbent particles with a moisture content of less than 5.0%. The moisture content is determined according to the method ISO 17190-4 "Determination of moisture content by mass loss upon heating". If the moisture content exceeds 5.0%, the

10 superabsorbent is dried at 60°C until the moisture content is less than 5.0%.

15

Particles with a diameter smaller than 150 µm are then separated. All indications of the number of superabsorbent particles in the present invention relate to particles with a diameter of 150 µm or greater. Particles with a diameter smaller than 150 µm are therefore not included in the expression "superabsorbent particles" according to the invention. In order to separate particles smaller than 150 µm, use is made of apparatus described in ISO 17190-3 "Determination of particle size distribution by sieve fractionation". Particles smaller than 150 µm are sieved out. The remaining particles are

20 then weighed. This weight therefore constitutes the total weight of the superabsorbent particles.

25

In order to calculate the number of superabsorbent particles per unit of volume, the superabsorbent particles are divided up into smaller portions. To divide the superabsorbent particles up into smaller portions, use is made of a

30 "Rotary Sample Divider – laborette 27" from Fritsch GmbH Laborgerätebau

or similar apparatus. Each portion is assumed to have a representative particle size distribution. Three of these portions are then weighed, and the number of particles in these three portions is counted manually. Each portion weighed 0.1 gram, so altogether the number of superabsorbent particles in
5 0.3 gram was counted. The weight of the samples is to be +/- 10% of the given values. The measuring accuracy is to be +/- 0.005 gram. The average weight of the individual superabsorbent particles is then calculated by dividing the weight of the sample (roughly 0.3 gram) by the number of manually counted particles. By then dividing the total weight of the
10 superabsorbent particles by the average weight of the superabsorbent particles, the total number of superabsorbent particles in the absorbent layer is obtained. In order finally to obtain the number of superabsorbent particles per cm^3 of the absorbent layer, the total number of superabsorbent particles is divided by the volume of the absorbent layer.

15

The absorption capacity of the superabsorbent particles is then measured according to ISO 17190-6 "Gravimetric determination of fluid retention after centrifugation". The absorption capacity is measured on three other portions. All portions have a representative particle size distribution, and the
20 absorption capacity per particle can therefore be calculated by dividing the measured absorption capacity by the number of particles previously counted manually.

Liquid used for measurement is 0.9% by weight sodium chloride solution.

25

Superabsorbent materials tested are three different size fractions of particulate polyacrylate-based superabsorbent from BASF with the designation Hysorb C 7100 and two different size fractions of particulate polyacrylate-based superabsorbent from Dow with the designation Drytech
30 S230R. The average particle size of the superabsorbent particles from BASF with the designation Hysorb C 7100 was in the first case a normal particle-size distribution, that is to say the measurement was performed on the whole

particle fraction in the commercially available grade; in the second case the particle size was between 600 μm and 710 μm , and in the third case the average particle size was between 710 μm and 850 μm . The average particle size of the superabsorbent particles from Dow with the designation Drytech S230R was in one case a normal particle size distribution, that is to say the measurement was performed on the whole particle fraction in the commercially available grade, and the average particle size in the other case was greater than 600 μm .

10 The superabsorbent with the designation Hysorb C 7100 with a normal particle size distribution is called A below.

The superabsorbent with the designation Hysorb C 7100 with a particle size between 600 μm and 710 μm is called B below.

15

The superabsorbent with the designation Hysorb C 7100 with a particle size between 710 μm and 850 μm is called C below.

20 The superabsorbent with the designation Drytech S230R with a normal particle size distribution is called D below.

The superabsorbent Drytech S230R with a particle size greater than 600 μm is called E below.

25 Result

	Superabsorbent	Abs. cap. (g/g)	Abs. cap/particle (mg/particle)
	A	35.6	1.4
	B	38.0	7.4
	C	37.2	9.5
30	D	33.6	1.7
	E	36.5	8.4

It can be seen from the result that the average absorption capacity per particle for superabsorbent C and E is greater than 8.0 mg sodium chloride solution, while the average absorption capacity per particle for superabsorbent A, B and D is less than 8.0 mg sodium chloride solution.

5

Example 3 – Measuring admission time in absorbent structure.

Measurement of admission time on a first, a second, a third and a fourth measurement was performed for five different absorbent structures. The
10 absorbent structures contained 50% by weight superabsorbent and 50% by weight chemical fluff pulp. The chemical fluff pulp was manufactured by Weyerhaeuser and is called NB 416.

Superabsorbent material in absorbent structure 1 was superabsorbent A, that
15 is to say Hysorb C 7100 with a normal particle size distribution.

Superabsorbent material in absorbent structure 2 was superabsorbent B, that is to say Hysorb C 7100 with a particle size between 600 μm and 710 μm .

20 Superabsorbent material in absorbent structure 3 was superabsorbent C, that is to say Hysorb C 7100 with a particle size between 710 μm and 850 μm .

Superabsorbent material in absorbent structure 4 was superabsorbent D, that is to say Drytech S230R with a normal particle size distribution.

25

Superabsorbent material in absorbent structure 5 was superabsorbent E, that is to say Drytech S230R with a particle size greater than 600 μm .

The absorbent structures 1, 2 and 3 had a density which was 0.25 g/cm^3 , a
30 weight per unit area which was 600 g/m^2 and an area which was 10x28 cm.

The absorbent structures 4 and 5 had a density which was 0.25 g/cm^3 , a weight per unit area which was 600 g/m^2 and an area which was $10 \times 40 \text{ cm}$.

For measurement, the absorbent structure was placed on a foam mattress of the tempur type. The absorbent structure was then subjected to a load of 0.64 kPa and four doses of 80 ml each of sodium chloride solution (0.9% by weight) were added. The time between the liquid doses was 10 minutes. The time for the liquid to be admitted into the absorbent structure was measured. The admission time was measured in seconds.

10

Result

	Abs. struct. 1 (s)	Abs. struct. 2 (s)	Abs. struct. 3 (s)
1 st wetting	115	103	110
2 nd wetting	210	149	146
15 3 rd wetting	312	230	217
4 th wetting	354	275	259

	Abs. struct. 4 (s)	Abs. struct. 5 (s)
1 st wetting	67	83
20 2 nd wetting	80	69
3 rd wetting	122	109
4 th wetting	160	132

The result shows that of the absorbent structures which contained superabsorbent Hysorb C 7100, that is to say absorbent structures 1-3, absorbent structure 3 has the fastest admission time on repeated wetting. Absorbent structure 3 contained superabsorbent particles with an average absorption capacity which is greater than 8.0 mg.

Of the absorbent structures which contained Drytech S230R, that is to say absorbent structures 4-5, absorbent structure 5 has the fastest admission time on repeated wetting. Absorbent structure 5 contained superabsorbent

particles with an average absorption capacity which is greater than 8.0 mg, while absorbent structure 4 contained superabsorbent particles with an average absorption capacity which is lower than 8.0 mg.

5 Example 4 – Measuring liquid distribution in absorbent structure.

Measurement of liquid distribution after a first, a second, a third and a fourth wetting was performed for absorbent structure 4 and absorbent structure 5. The liquid distribution was measured after each wetting immediately before
10 the next liquid dose was added. The liquid distribution was measured in cm.

Result

	Abs. struct. 4 (cm)	Abs. struct. 5 (cm)
1 st wetting	20	25
15 2 nd wetting	22	26
3 rd wetting	29	34
4 th wetting	34	39

The result shows that absorbent structure 5 which contained superabsorbent
20 particles with an average absorption capacity which is greater than 8.0 mg spreads the liquid further than absorbent structure 4 which contained superabsorbent particles with an average absorption capacity which is lower than 8.0 mg.

25 Example 5 – Measuring rewet.

Measurement of rewet after the fourth wetting was performed for absorbent structure 4 and absorbent structure 5. Measurement of rewet was started 10 minutes after the fourth dose of liquid had been applied. After 10 minutes, 15
30 pieces of filter paper and a weight (5 kPa) were placed on the wetting point of the absorbent structure. After 15 seconds, the weight was removed, and the bundle of filter paper was weighed. The rewet was calculated by subtracting

the dry weight of the filter paper from the wet weight. The rewet was measured in grams of sodium chloride solution (0.9% by weight).

Result

- 5 Absorbent structure 4: 9.5 grams
- Absorbent structure 5: 7.9 grams

10 The result shows that absorbent structure 5 has lower rewet than absorbent structure 4. The result therefore shows that the absorbent structure which contained superabsorbent particles with an average absorption capacity which is greater than 8.0 mg has lower rewet than the absorbent structure which contained superabsorbent particles with an average absorption capacity which is lower than 8.0 mg.

CLAIMS

- 5 1. Absorbent structure for use in a diaper, an incontinence pad, a sanitary towel or the like, which absorbent structure has at least one absorbent layer comprising fluff pulp and superabsorbent particles, characterized in that the average absorption capacity per superabsorbent particle in the absorbent layer is greater than 8.0 mg sodium chloride solution and in that the number of superabsorbent particles per cm³ of the absorbent layer is smaller than 1100.
- 10
2. Absorbent structure according to Claim 1, characterized in that the average absorption capacity per superabsorbent particle in the absorbent layer is greater than 9.5 mg sodium chloride solution and in that the number of superabsorbent particles per cm³ of the absorbent layer is smaller than 600.
- 15
3. Absorbent structure according to Claim 2, characterized in that the average absorption capacity per superabsorbent particle in the absorbent layer is greater than 14.0 mg sodium chloride solution and in that the number of superabsorbent particles per cm³ of the absorbent layer is smaller than 450.
- 20
4. Absorbent structure according to any one of Claims 1 – 3, characterized in that the superabsorbent particles have a particle size which is greater than 600 µm.
- 25
5. Absorbent structure according to any one of Claims 1 – 4, characterized in that the average distance between centres of the superabsorbent particles in the absorbent layer in a dry state is greater than 700 µm.
- 30
6. Absorbent structure according to Claim 5, characterized in that the average distance between centres of the superabsorbent particles in the absorbent layer in a dry state is greater than 1000 µm.
- 35
7. Absorbent structure according to Claim 6, characterized in that the average distance between centres of the superabsorbent particles in the absorbent layer in a dry state is greater than 1200 µm.

8. Absorbent structure according to any one of Claims 1-7, characterized in that the density of the absorbent layer in a dry state is greater than 0.12 g/cm^3 .

5 9. Absorbent structure according to Claim 8, characterized in that the density of the absorbent layer in a dry state is greater than 0.17 g/cm^3 .

10 10. Absorbent structure according to Claim 9, characterized in that the density of the absorbent layer in a dry state is greater than 0.25 g/cm^3 .

11. Absorbent structure according to any one of the preceding claims, characterized in that the absorbent layer comprises bonding means.

12. Absorbent structure according to any one of the preceding claims, characterized in that the superabsorbent particles are surface cross-linked.

15

13. Absorbent article such as a diaper, an incontinence pad, a sanitary towel or the like, characterized in that it comprises an absorbent structure according to any one of Claims 1 -12.

20 14. An absorbent structure according to any one of Claims 1 to 12, substantially as herein described with reference to and as illustrated in any of the examples and accompanying drawings.

25 15. An absorbent article according to Claim 13, substantially as herein described with reference to and as illustrated in any of the examples and accompanying drawings.