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SOFT ABSORBENT TISSUE PAPER
WITH HIGH TEMPORARY WET STRENGTH

FIELD OF THE INVENTION

This invention relates to tissue paper webs. More particularly, it relates to soft, absorbent tissue paper webs which can be used in sanitary tissue, facial tissue products, and paper napkins.

BACKGROUND OF THE INVENTION

Paper webs or sheets, sometimes called tissue or paper tissue webs or sheets, find extensive use in modern society. Such items as paper towels, napkins, and facial tissues are staple items of commerce. It has long been recognized that three important physical attributes of these products are their softness; their absorbency, particularly their absorbency for aqueous systems; and their strength, particularly their strength when wet. Research and development efforts have been directed to the improvement of each of these attributes without deleteriously affecting the others as well as to the improvement of two or three attributes simultaneously.

Softness is the tactile sensation perceived by the consumer as he/she holds a particular product, rubs it across his/her skin, or crumples it within his/her hand. This tactile sensation is a combination of several physical properties. One of the more important physical properties related to softness is generally considered by those skilled in the art to be the stiffness of the paper web from which the product is made. Stiffness, in turn, is usually considered to be directly dependent on the dry tensile strength of the web.

Strength is the ability of the product, and its constituent webs, to maintain physical integrity and to resist tearing, bursting, and shredding under use conditions, particularly when wet.

Absorbency is the measure of the ability of a product, and its constituent webs, to absorb quantities of liquid, particularly aqueous solutions or dispersions. Overall absorbency as perceived by the human consumer is generally considered to be a combination of the total

quantity of liquid a given mass of tissue paper will absorb at saturation as well as the rate at which the mass absorbs the liquid.

The use of wet strength resins to enhance the strength of a paper web is widely known. For example, Westfelt described a number of such materials and discussed their chemistry in *Cellulose Chemistry and Technology*, Volume 13, at pages 813-825 (1979).

Freimark et al. in U.S. Pat. No. 3,755,220 issued August 28, 1973 mention that certain chemical additives known as debonding agents interfere with the natural fiber-to-fiber bonding that occurs during sheet formation in papermaking processes. This reduction in bonding leads to a softer, or less harsh, sheet of paper. Freimark et al. go on to teach the use of wet strength resins to enhance the wet strength of the sheet in conjunction with the use of debonding agents to off-set undesirable effects of the wet strength resin. These debonding agents do reduce dry tensile strength, but there is also generally a reduction in wet tensile strength.

Shaw, in U.S. Pat. No. 3,821,068, issued June 28, 1974, also teaches that chemical debonders can be used to reduce the stiffness, and thus enhance the softness, of a tissue paper web.

Chemical debonding agents have been disclosed in various references such as U.S. Pat. No. 3,554,862, issued to Hervey et al. on January 12, 1971. These materials include quaternary ammonium salts such as trimethylcocoammonium chloride, trimethyloleylammonium chloride, dimethyldi(hydrogenated-tallow)ammonium chloride and trimethylstearyl-ammonium chloride.

Emanuelsson et al., in U.S. Pat. No. 4,144,122, issued March 13, 1979, teach the use of complex quaternary ammonium compounds such as bis(alkoxy-(2-hydroxy)-propylene) quaternary ammonium chlorides to soften webs. These authors also attempt to overcome any decrease in absorbency caused by the debonders through the use of nonionic surfactants such as ethylene oxide and propylene oxide adducts of fatty alcohols.

Armak Company, of Chicago, Illinois, in their bulletin 76-17 (1977) disclose that the use of dimethyldi(hydrogenated-tallow)ammonium chloride in combination with fatty acid esters of polyoxyethylene glycols may impart both softness and absorbency to tissue paper webs.

One exemplary result of research directed toward improved paper webs is described in U.S. Pat. No. 3,301,746, issued to Sanford and Sisson on January 31, 1967. Despite the high quality of paper webs made

by the process described in this patent, and despite the commercial success of products formed from these webs, research efforts directed to finding improved products have continued.

For example, Becker et al. in U.S. Pat. No. 4,158,594, issued January 19, 1979, describe a method they contend will form a strong, soft, fibrous sheet. More specifically, they teach that the strength of a tissue paper web (which may have been softened by the addition of chemical debonding agents) can be enhanced by adhering, during processing, one surface of the web to a creping surface in a fine patterned arrangement by a bonding material (such as an acrylic latex rubber emulsion, a water soluble resin, or an elastomeric bonding material) which has been adhered to one surface of the web and to the creping surface in the fine patterned arrangement, and creping the web from the creping surface to form a sheet material.

It is an object of this invention to provide a process for making soft, absorbent tissue paper webs with high temporary wet strength.

It is a further object of this invention to provide soft, absorbent tissue paper sheets with high temporary wet strength.

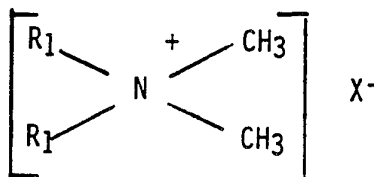
It is a still further object of this invention to provide soft, absorbent paper towel products with high permanent wet strength.

These and other objects are obtained using the present invention, as will become readily apparent from a reading of the following disclosure.

SUMMARY OF THE INVENTION

The present invention provides soft, absorbent tissue paper webs having high wet strength, and a process for making the webs. Briefly, the tissue paper webs comprise:

- (a) papermaking fibers;
- (b) from about 0.01% to about 2.0% by weight of a quaternary ammonium compound having the formula



wherein each R₁ substituent is a C₁₂-C₁₈ aliphatic hydrocarbon radical, and X⁻ is a compatible anion;

- (c) from about 0.01% to about 2.0% by weight of a polyhydroxy plasticizer; and
- (d) from about 0.01% to about 3.0% by weight of a water-soluble temporary wet strength resin.

Examples of quaternary ammonium compounds suitable for use in the present invention include the well-known dialkyldimethylammonium salts such as ditallowdimethylammonium chloride, ditallowdimethylammonium methylsulfate, di(hydrogenated tallow)dimethylammonium chloride; with di(hydrogenated tallow)dimethylammonium methylsulfate being preferred.

Examples of polyhydroxy plasticizers useful in the present invention include glycerol and polyethylene glycols having a molecular weight of from about 200 to about 2000, with polyethylene glycols having a molecular weight of from about 200 to about 600 being preferred.

The temporary wet strength resins useful in the present invention include all those commonly used in papermaking. Examples of preferred temporary wet strength resins include cationic starch-based resins and the cationic polymers described in U.S. Pat. No. 4,981,557, Bjorkquist, issued January 1, 1991.

A particularly preferred tissue paper embodiment of the present invention comprises from about 0.01% to about 0.5% by weight of the quaternary ammonium compound, from about 0.01% to about 0.5% by weight of the polyhydroxy plasticizer, and from about 0.1% to about 1.5% by weight of the water-soluble temporary wet strength resin, all quantities of these additives being on a dry fiber weight basis of the tissue paper.

Briefly, the process for making the tissue webs of the present invention comprises the steps of forming a papermaking furnish from the aforementioned components, deposition of the papermaking furnish onto a foraminous surface such as a Fourdrinier wire, and removal of the water from the deposited furnish.

All percentages, ratios and proportions herein are by weight unless otherwise specified.

The present invention is described in more detail below.

DETAILED DESCRIPTION OF THE INVENTION

While this specification concludes with claims particularly pointing out and distinctly claiming the subject matter regarded as the invention, it is believed that the invention can be better understood

from a reading of the following detailed description and of the appended example.

As used herein, the terms tissue paper web, paper web, web, and paper sheet all refer to sheets of paper made by a process comprising the steps of forming an aqueous papermaking furnish, depositing this furnish on a foraminous surface, such as a Fourdrinier wire, and removing the water from the furnish as by gravity or vacuum-assisted drainage, with or without pressing, and by evaporation.

As used herein, an aqueous papermaking furnish is an aqueous slurry of papermaking fibers and the chemicals described hereinafter.

The first step in the process of this invention is the forming of an aqueous papermaking furnish. The furnish comprises papermaking fibers (hereinafter sometimes referred to as wood pulp), at least one wet strength resin, at least one quaternary ammonium and at least one polyhydroxy plasticizer, all of which will be hereinafter described.

It is anticipated that wood pulp in all its varieties will normally comprise the papermaking fibers used in this invention. However, other cellulosic fibrous pulps, such as cotton linters, bagasse, rayon, etc., can be used and none are disclaimed. Wood pulps useful herein include chemical pulps such as Kraft, sulfite and sulfate pulps as well as mechanical pulps including for example, ground wood, thermomechanical pulps and chemically modified thermomechanical pulp (CTMP). Pulps derived from both deciduous and coniferous trees can be used. Also applicable to the present invention are fibers derived from recycled paper, which may contain any or all of the above categories as well as other non-fibrous materials such as fillers and adhesives used to facilitate the original papermaking. Preferably, the papermaking fibers used in this invention comprise Kraft pulp derived from northern softwoods.

Wet Strength Resins

The present invention contains as an essential component from about 0.01% to about 3.0%, more preferably from about 0.1% to about 1.5% by weight, on a dry fiber weight basis, of a water-soluble temporary wet strength resin.

Wet strength resins useful herein can be of several types. Generally, those resins which have previously found and which will hereafter find utility in the papermaking art are useful herein.

Numerous examples are shown in the aforementioned paper by Westfelt, incorporated herein by reference.

In the usual case, the wet strength resins are water-soluble, cationic materials. That is to say, the resins are water-soluble at the time they are added to the papermaking furnish. It is quite possible, and even to be expected, that subsequent events such as cross-linking will render the resins insoluble in water. Further, some resins are soluble only under specific conditions, such as over a limited pH range.

Wet strength resins are generally believed to undergo a cross-linking or other curing reactions after they have been deposited on, within, or among the papermaking fibers. Cross-linking—or curing does not normally occur so long as substantial amounts of water are present.

Of particular utility are the various polyamide-epichlorohydrin resins. These materials are low molecular weight polymers provided with reactive functional groups such as amino, epoxy, and azetidinium groups. The patent literature is replete with descriptions of processes for making such materials. U.S. Pat. No. 3,700,623, issued to Keim on October 24, 1972 and U.S. Pat. No. 3,772,076, issued to Keim on November 13, 1973 are examples of such patents and both are incorporated herein by reference.

Polyamide-epichlorohydrin resins sold under the trademarks Kymene 557H and Kymene 2064 by Hercules Incorporated of Wilmington, Delaware, are particularly useful in this invention. These resins are generally described in the aforementioned patents to Keim.

Base-activated polyamide-epichlorohydrin resins useful in the present invention are sold under the Santo Res trademark, such as Santo Res 31, by Monsanto Company of St. Louis, Missouri. These types of materials are generally described in U.S. Pat. Nos. 3,855,158 issued to Petrovich on December 17, 1974; 3,899,388 issued to Petrovich on August 12, 1975; 4,129,528 issued to Petrovich on December 12, 1978; 4,147,586 issued to Petrovich on April 3, 1979; and 4,222,921 issued to Van Eenam on September 16, 1980, all incorporated herein by reference.

Other water-soluble cationic resins useful herein are the polyacrylamide resins such as those sold under the Parez trademark, such as Parez 63INC, by American Cyanamid Company of Stamford, Connecticut. These materials are generally described in U.S. Pat. Nos. 3,556,932 issued to Coscia et al. on January 19, 1971; and 3,556,933 issued to

Williams et al. on January 19, 1971, all incorporated herein by reference.

Other types of water-soluble resins useful in the present invention include acrylic emulsions and anionic styrene-butadiene latexes. Numerous examples of these types of resins are provided in U.S. Patent 3,844,880, Meisel, Jr. et al., issued October 29, 1974, incorporated herein by reference.

Still other water-soluble cationic resins finding utility in this invention are the urea formaldehyde and melamine formaldehyde resins. These polyfunctional, reactive polymers have molecular weights on the order of a few thousand. The more common functional groups include nitrogen containing groups such as amino groups and methylol groups attached to nitrogen.

Although less preferred, polyethylenimine type resins find utility in the present invention.

More complete descriptions of the aforementioned water-soluble resins, including their manufacture, can be found in TAPPI Monograph Series No. 29, Wet Strength In Paper and Paperboard, Technical Association of the Pulp and Paper Industry (New York; 1965), incorporated herein by reference.

The above-mentioned wet strength additives typically result in paper products with permanent wet strength, i.e., paper which when placed in an aqueous medium retains a substantial portion of its initial wet strength over time. However, permanent wet strength in some types of paper products can be an unnecessary and undesirable property. Paper products such as toilet tissues, etc., are generally disposed of after brief periods of use into septic systems and the like. Clogging of these systems can result if the paper product permanently retains its hydrolysis-resistant strength properties.

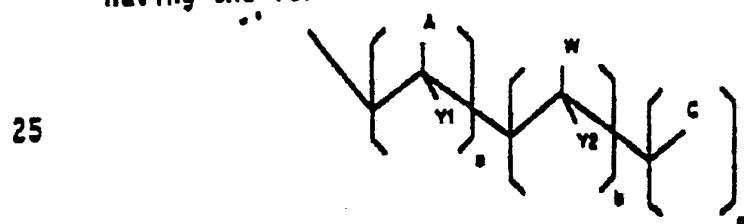
More recently, manufacturers have added temporary wet strength additives to paper products for which wet strength is sufficient for the intended use, but which then decays upon soaking in water. Decay of the wet strength facilitates flow of the paper product through septic systems. As used herein, the term "temporary wet strength resin" refers to a resin that allows the tissue paper, when placed in an aqueous medium, to lose a majority of its initial wet strength in a short period of time, e.g., two minutes or less, more preferably, 30 seconds or less.

Examples of suitable temporary wet strength resins include modified starch temporary wet strength agents such as National Starch 78-0080,

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10 Examples of suitable temporary wet strength resins include modified starch temporary wet strength agents such as National Starch 78-0080, marketed by the National Starch and Chemical Corporation (New York, New York). This type of wet strength agent can be made by reacting dimethoxyethyl-N-methyl-chloroacetamide with cationic starch polymers. Modified starch temporary wet strength agents are also described in U.S. Pat. No. 4,675,394, Solarek, et al., issued June 23, 1987, and incorporated herein by reference.

15 Preferred temporary wet strength resins include those described in U.S. Pat. No. 4,981,557, Bjorkquist, issued January 1, 1991, and incorporated herein by reference. The temporary wet strength resins described in U.S. Pat. No. 4,981,557 comprise a polymer characterized by the substantially complete absence of nucleophilic functionalities and having the formula:



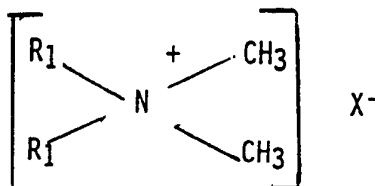
wherein: A is



35 and X is $-\text{O}-$, $-\text{NCH}_3-$, and R is a substituted or unsubstituted aliphatic groups; Y₁ and Y₂ are independently $-\text{H}$, $-\text{CH}_3$ or a halogen; W is a nonnucleophilic, water-soluble nitrogen heterocyclic moiety; C is a cationic monomeric unit; the mole percent of a is from about 30% to about 70%, the mole percent of b is from about 30%

Quaternary Ammonium Compound

The present invention contains as an essential component from about 0.01% to about 2.0%, more preferably from about 0.01% to about 0.5% by weight, on a dry fiber weight basis, of a quaternary ammonium compound having the formula:



In the structure noted above each R_1 is an aliphatic hydrocarbon radical selected from the group consisting of alkyl having from about 12 to about 18 carbon atoms, coconut and tallow. X^- is a compatible anion, such as an halide (e.g., chloride or bromide) or methylsulfate. Preferably, X^- is methylsulfate.

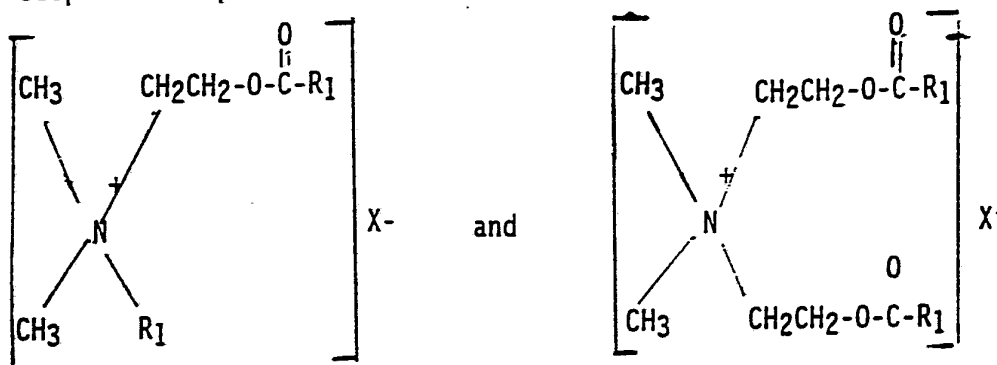
As used above, "coconut" refers to the alkyl and alkylene moieties derived from coconut oil. It is recognized that coconut oil is a naturally occurring mixture having, as do all naturally occurring materials, a range of compositions. Coconut oil contains primarily fatty acids (from which the alkyl and alkylene moieties of the quaternary ammonium salts are derived) having from 12 to 16 carbon atoms, although fatty acids having fewer and more carbon atoms are also present. Swern, Ed. in Bailey's Industrial Oil and Fat Products, Third Edition, John Wiley and Sons (New York 1964) in Table 6.5, suggests that coconut oil typically has from about 65 to 82% by weight of its fatty acids in the 12 to 16 carbon atoms range with about 8% of the total fatty acid content being present as unsaturated molecules. The principle unsaturated fatty acid in coconut oil is oleic acid. Synthetic as well as naturally occurring "coconut" mixtures fall within the scope of this invention.

Tallow, as is coconut, is a naturally occurring material having a variable composition. Table 6.13 in the above-identified reference edited by Swern indicates that typically 78% or more of the fatty acids of tallow contain 16 or 18 carbon atoms. Typically, half of the fatty acids present in tallow are unsaturated, primarily in the form of oleic acid. Synthetic as well as natural "tallows" fall within the scope of the present invention.

Preferably, each R_1 is C_{16} - C_{18} alkyl, most preferably each R_1 is straight-chain C_{18} alkyl.

Examples of quaternary ammonium compounds suitable for use in the present invention include the well-known dialkyldimethylammonium salts such as ditallowdimethylammonium chloride, ditallowdimethylammonium methylsulfate, di(hydrogenated tallow) dimethylammonium chloride; with di(hydrogenated tallow)dimethylammonium methylsulfate being preferred. This particular material is available commercially from Sherex Chemical Company Inc. of Dublin, Ohio under the tradename "Varisoft^R 137".

Biodegradable mono and di-ester variations of the quaternary ammonium compound can also be used, and are meant to fall within the scope of the present invention. These compounds have the formula:



with R₁ and X⁻ as defined above.

Polyhydroxy Plasticizer

The present invention contains as an essential component from 0.01% to about 2.0%, more preferably from about 0.01% to about 0.5% by weight, on a dry fiber weight basis, of a polyhydroxy plasticizer.

Examples of polyhydroxy plasticizers useful in the present invention include glycerol and polyethylene glycols having a molecular weight of from about 200 to about 2000, with polyethylene glycols having a molecular weight of from about 200 to about 600 being preferred.

A particularly preferred polyhydroxy plasticizer is polyethylene glycol having a molecular weight of about 400. This material is available commercially from the Union Carbide Company of Danbury, Connecticut under the tradename "PEG-400".

Optional Ingredients

Other chemicals commonly used in papermaking can be added to the papermaking furnish so long as they do not significantly and adversely affect the softening, absorbency, and wet strength enhancing actions of the three required chemicals.

For example, surfactants may be used to treat the tissue paper webs of the present invention. The level of surfactant, if used, is preferably from about 0.01% to about 2.0% by weight, based on the dry fiber weight of the tissue paper. The surfactants preferably have alkyl chains with eight or more carbon atoms. Exemplary anionic surfactants are linear alkyl sulfonates, and alkylbenzene sulfonates. Exemplary nonionic surfactants are alkylglycosides including alkylglycoside esters such as Crodesta™ SL-40 which is available from Croda, Inc. (New York, NY); alkylglycoside ethers as described in U.S. Patent 4,011,389, issued to W. K. Langdon, et al. on March 8, 1977; and alkylpolyethoxylated esters such as Pegosperse™ 200 ML available from Glyco Chemicals, Inc. (Greenwich, CT) and IGEPAL RC-520 available from Rhone Poulenc Corporation (Cranbury, N.J.).

Other types of chemicals which may be added include dry strength additives to increase the tensile strength of the tissue webs. Examples of dry strength additives include cationic polymers from the ACCO chemical family such as ACCO 771 and ACCO 514. The level of dry strength additive, if used, is preferably from about 0.01% to about 1.0%, by weight, based on the dry fiber weight of the tissue paper.

The above listings of additional chemical additives is intended to be merely exemplary in nature, and are not meant to limit the scope of the invention.

The papermaking furnish can be readily formed or prepared by mixing techniques and equipment well known to those skilled in the papermaking art.

The three types of chemical ingredients described above i.e. quaternary ammonium compounds, polyhydroxy plasticizers, and water soluble temporary wet strength resins are preferably added to the aqueous slurry of papermaking fibers, or furnish in the wet end of the papermaking machine at some suitable point ahead of the Fourdrinier wire or sheet forming stage. However, applications of the above chemical ingredients subsequent to formation of a wet tissue web and prior to drying of the web to completion will also provide significant softness, absorbency, and wet strength benefits and are expressly included within the scope of the present invention.

It has been discovered that the chemical ingredients are more effective when the quaternary ammonium compound and the polyhydroxy plasticizer are first pre-mixed together before being added to the papermaking furnish. A preferred method, as will be described in

greater detail hereinafter in Example 1, consists of first heating the polyhydroxy plasticizer to a temperature of about 150°F, and then adding the quaternary ammonium compound to the hot plasticizer to form a fluidized "melt". Preferably, the molar ratio of the quaternary ammonium compound to the plasticizer is about 1 to 1, although this ratio will vary depending upon the molecular weight of the particular plasticizer and/or quaternary ammonium compound used. The quaternary ammonium compound and polyhydroxy plasticizer melt is then diluted to the desired concentration, and mixed to form an aqueous solution containing a vesicle suspension of the quaternary ammonium compound/polyhydroxy plasticizer mixture which is then added to the papermaking furnish.

Without being bound by theory, it is believed that the plasticizer enhances the flexibility of the cellulosic fibers, improves the absorbency of the fibers, and acts to stabilize the quaternary ammonium compound in the aqueous solution. Separately, the temporary wet strength resins are also diluted to the appropriate concentration and added to the papermaking furnish. The quaternary ammonium/polyhydroxy plasticizer chemical softening composition acts to make the paper product soft and absorbent, while the temporary wet strength resin insures that the resulting paper product also has high temporary wet strength. In other words, the present invention makes it possible to not only improve both the softness and absorbency rate of the tissue webs, but also provides a high level of temporary wet strength.

The second step in the process of this invention is the depositing of the papermaking furnish on a porous surface and the third is the removing of the water from the furnish so deposited. Techniques and equipment which can be used to accomplish these two processing steps will be readily apparent to those skilled in the papermaking art.

The present invention is applicable to tissue paper in general, including but not limited to conventionally felt-pressed tissue paper; pattern densified tissue paper such as exemplified in the aforementioned U.S. Patent by Sanford-Sisson and its progeny; and high bulk, uncompacted tissue paper such as exemplified by U.S. Patent 3,812,000, Salvucci, Jr., issued May 21, 1974. The tissue paper may be of a homogenous or multilayered construction; and tissue paper products made therefrom may be of a single-ply or multi-ply construction. The tissue paper preferably has a basis weight of between 10 g/m² and about 65 g/m², and density of about 0.60 g/cc or less. Preferably, basis weight

will be below about 35 g/m² or less; and density will be about 0.30 g/cc or less. Most preferably, density will be between 0.04 g/cc and about 0.20 g/cc.

Conventionally pressed tissue paper and methods for making such paper are known in the art. Such paper is typically made by depositing papermaking furnish on a foraminous forming wire. This forming wire is often referred to in the art as a Fourdrinier wire. Once the furnish is deposited on the forming wire, it is referred to as a web. The web is dewatered by pressing the web and drying at elevated temperature. The particular techniques and typical equipment for making webs according to the process just described are well known to those skilled in the art. In a typical process, a low consistency pulp furnish is provided in a pressurized headbox. The headbox has an opening for delivering a thin deposit of pulp furnish onto the Fourdrinier wire to form a wet web. The web is then typically dewatered to a fiber consistency of between about 7% and about 25% (total web weight basis) by vacuum dewatering and further dried by pressing operations wherein the web is subjected to pressure developed by opposing mechanical members, for example, cylindrical rolls. The dewatered web is then further pressed and dried by a stream drum apparatus known in the art as a Yankee dryer. Pressure can be developed at the Yankee dryer by mechanical means such as an opposing cylindrical drum pressing against the web. Multiple Yankee dryer drums may be employed, whereby additional pressing is optionally incurred between the drums. The tissue paper structures which are formed are referred to hereinafter as conventional, pressed, tissue paper structures. Such sheets are considered to be compacted since the web is subjected to substantial mechanical compressional forces while the fibers are moist and are then dried while in a compressed state.

Pattern densified tissue paper is characterized by having a relatively high bulk field of relatively low fiber density and an array of densified zones of relatively high fiber density. The high bulk field is alternatively characterized as a field of pillow regions. The densified zones are alternatively referred to as knuckle regions. The densified zones may be discretely spaced within the high bulk field or may be interconnected, either fully or partially, within the high bulk field. Preferred processes for making pattern densified tissue webs are disclosed in U.S. Patent No. 3,301,746, issued to Sanford and Sisson on January 31, 1967, U.S. Patent No. 3,974,025, issued to Peter G. Ayers on August 10, 1976, and U.S. Patent No. 4,191,609, issued to Paul D.

Trokhan on March 4, 1980, and U.S. Patent 4,637,859, issued to Paul D. Trokhan on January 20, 1987; all of which are incorporated herein by reference.

In general, pattern densified webs are preferably prepared by depositing a papermaking furnish on a foraminous forming wire such as a Fourdrinier wire to form a wet web and then juxtaposing the web against an array of supports. The web is pressed against the array of supports, thereby resulting in densified zones in the web at the locations geographically corresponding to the points of contact between the array of supports and the wet web. The remainder of the web not compressed during this operation is referred to as the high bulk field. This high bulk field can be further dedensified by application of fluid pressure, such as with a vacuum type device or a blow-through dryer, or by mechanically pressing the web against the array of supports. The web is dewatered, and optionally predried, in such a manner so as to substantially avoid compression of the high bulk field. This is preferably accomplished by fluid pressure, such as with a vacuum type device or blow-through dryer, or alternately by mechanically pressing the web against an array of supports wherein the high bulk field is not compressed. The operations of dewatering, optional predrying and formation of the densified zones may be integrated or partially integrated to reduce the total number of processing steps performed. Subsequent to formation of the densified zones, dewatering, and optional predrying, the web is dried to completion, preferably still avoiding mechanical pressing. Preferably, from about 8% to about 55% of the tissue paper surface comprises densified knuckles having a relative density of at least 125% of the density of the high bulk field.

The array of supports is preferably an imprinting carrier fabric having a patterned displacement of knuckles which operate as the array of supports which facilitate the formation of the densified zones upon application of pressure. The pattern of knuckles constitutes the array of supports previously referred to. Imprinting carrier fabrics are disclosed in U.S. Patent No. 3,301,746, Sanford and Sisson, issued January 31, 1967, U.S. Patent No. 3,821,068, Salvucci, Jr. et al., issued May 21, 1974, U.S. Patent No. 3,974,025, Ayers, issued August 10, 1976, U.S. Patent No. 3,573,164, Friedberg et al., issued March 30, 1971, U.S. Patent No. 3,473,576, Amneus, issued October 21, 1969, U.S. Patent No. 4,239,065, Trokhan, issued December 16, 1980, and U.S. Patent

No. 4,528,239, Trokhan, issued July 9, 1985, all of which are incorporated herein by reference.

Preferably, the furnish is first formed into a wet web on a foraminous forming carrier, such as a Fourdrinier wire. The web is dewatered and transferred to an imprinting fabric. The furnish may alternately be initially deposited on a foraminous supporting carrier which also operates as an imprinting fabric. Once formed, the wet web is dewatered and, preferably, thermally predried to a selected fiber consistency of between about 40% and about 80%. Dewatering is preferably performed with suction boxes or other vacuum devices or with blow-through dryers. The knuckle imprint of the imprinting fabric is impressed in the web as discussed above, prior to drying the web to completion. One method for accomplishing this is through application of mechanical pressure. This can be done, for example, by pressing a nip roll which supports the imprinting fabric against the face of a drying drum, such as a Yankee dryer, wherein the web is disposed between the nip roll and drying drum. Also, preferably, the web is molded against the imprinting fabric prior to completion of drying by application of fluid pressure with a vacuum device such as a suction box, or with a blow-through dryer. Fluid pressure may be applied to induce impression of densified zones during initial dewatering, in a separate, subsequent process stage, or a combination thereof.

Uncompacted, nonpattern-densified tissue paper structures are described in U.S. Patent No. 3,812,000 issued to Joseph L. Salvucci, Jr. and Peter N. Yiannos on May 21, 1974 and U.S. Patent No. 4,208,459, issued to Henry E. Becker, Albert L. McConnell, and Richard Schutte on June 17, 1980, both of which are incorporated herein by reference. In general, uncompacted, nonpattern-densified tissue paper structures are prepared by depositing a papermaking furnish on a foraminous forming wire such as a Fourdrinier wire to form a wet web, draining the web and removing additional water without mechanical compression until the web has a fiber consistency of at least 80%, and creping the web. Water is removed from the web by vacuum dewatering and thermal drying. The resulting structure is a soft but weak high bulk sheet of relatively uncompacted fibers. Bonding material is preferably applied to portions of the web prior to creping.

Compacted non-pattern-densified tissue structures are commonly known in the art as conventional tissue structures. In general, compacted, non-pattern-densified tissue paper structures are prepared by

depositing a papermaking furnish on a foraminous wire such as a Fourdrinier wire to form a wet web, draining the web and removing additional water with the aid of a uniform mechanical compaction (pressing) until the web has a consistency of 25-50%, transferring the web to a thermal dryer such as a Yankee and creping the web. Overall, water is removed from the web by vacuum, mechanical pressing and thermal means. The resulting structure is strong and generally of singular density, but very low in bulk, absorbency and in softness.

The tissue paper web of this invention can be used in any application where soft, absorbent tissue paper webs with high temporary wet strength are required. One particularly advantageous use of the tissue paper web of this invention is in sanitary tissue products.

Analysis of the amount of treatment chemicals herein retained on tissue paper webs can be performed by any method accepted in the applicable art. For example, the level of the quaternary ammonium compound, such as DTMAMS, retained by the tissue paper can be determined by solvent extraction of the DTMAMS by an organic solvent followed by an anionic/cationic titration using Dimidium Bromide as indicator; the level of the polyhydroxy plasticizer, such as PEG-400, can be determined by extraction in an organic solvent followed by gas chromatography to determine the level of PEG-400 in the extract; the level of temporary wet strength resin such as a temporary wet strength resin with a nitrogen moiety (e.g., as described in U.S. Patent 4,981,557, D. W. Bjorkquist issued January 1, 1991) resin can be determined by subtraction from the total nitrogen level obtained via the Nitrogen Analyzer, the amount of quaternary ammonium compound level, determined by the above titration method. These methods are exemplary, and are not meant to exclude other methods which may be useful for determining levels of particular components retained by the tissue paper.

Hydrophilicity of tissue paper refers, in general, to the propensity of the tissue paper to be wetted with water. Hydrophilicity of tissue paper may be somewhat quantified by determining the period of time required for dry tissue paper to become completely wetted with water. This period of time is referred to as "wetting time." In order to provide a consistent and repeatable test for wetting time, the following procedure may be used for wetting time determinations: first, a conditioned sample unit sheet (the environmental conditions for testing of paper samples are $23 \pm 1^\circ\text{C}$ and $50 \pm 2\% \text{RH}$. as specified in TAPPI

Method T 402), approximately 4-3/8 inch x 4-3/4 inch (about 11.1 cm x 12 cm) of tissue paper structure is provided; second, the sheet is folded into four (4) juxtaposed quarters, and then crumpled into a ball approximately 0.75 inches (about 1.9 cm) to about 1 inch (about 2.5 cm) in diameter; third, the balled sheet is placed on the surface of a body of distilled water at $23 \pm 1^\circ\text{C}$ and a timer is simultaneously started; fourth, the timer is stopped and read when wetting of the balled sheet is completed. Complete wetting is observed visually.

The preferred hydrophilicity of tissue paper depends upon its intended end use. It is desirable for tissue paper used in a variety of applications, e.g., toilet paper, to completely wet in a relatively short period of time to prevent clogging once the toilet is flushed. Preferably, wetting time is 2 minutes or less. More preferably, wetting time is 30 seconds or less. Most preferably, wetting time is 10 seconds or less.

Hydrophilicity characters of tissue paper embodiments of the present invention may, of course, be determined immediately after manufacture. However, substantial increases in hydrophobicity may occur during the first two weeks after the tissue paper is made: i.e., after the paper has aged two (2) weeks following its manufacture. Thus, the above stated wetting times are preferably measured at the end of such two week period. Accordingly, wetting times measured at the end of a two week aging period at room temperature are referred to as "two week wetting times."

The density of tissue paper, as that term is used herein, is the average density calculated as the basis weight of that paper divided by the caliper, with the appropriate unit conversions incorporated therein. Caliper of the tissue paper, as used herein, is the thickness of the paper when subjected to a compressive load of 95 g/in² (14.7 g/cm²).

The following example illustrates the practice of the present invention but is not intended to be limiting thereof.

EXAMPLE 1

The purpose of this example is to illustrate one method that can be used to make soft, absorbent and high temporary wet strength tissue fibrous structure treated with a mixture of Dihydrogenated Tallow Dimethyl Ammonium Methyl Sulfate (DTDMAMS) and a polyhydroxy plasticizer (PEG-400) in the presence of a temporary wet strength resin in accordance with the present invention.

A pilot scale Fourdrinier papermaking machine is used in the practice of the present invention. First, a 1% solution of the chemical softener composition containing DTMAMS and PEG-400 is prepared according to the following procedure: 1. An equivalent molar concentration of DTMAMS and PEG-400 is weighed; 2. PEG is heated up to about 150°F; 3. DTMAMS is dissolved into PEG to form a melted solution; 4. Shear stress is applied to form a homogeneous mixture of DTMAMS in PEG; 5. The dilution water is heated up to about 150°F; 6. The melted mixture of DTMAMS/PEG-400 is diluted to a 1% solution; and 7. Shear stress is applied to form an aqueous solution containing a vesicle suspension of the DTMAMS/PEG-400 mixture.

Second, a 3% by weight aqueous slurry of NSK is made up in a conventional re-pulper. The NSK slurry is refined gently and a 2% solution of the temporary wet strength resin (as described in U.S. Pat. No. 4,981,557, D. W. Bjorkquist issued January 1, 1991) is added to the NSK stock pipe at a rate of 0.75% by weight of the dry fibers. The absorption of the temporary wet strength resin onto NSK fibers is enhanced via an in-line mixer. The NSK slurry is diluted to about 0.2% consistency at the fan pump.

Third, a 3% by weight aqueous slurry of Eucalyptus fibers is made up in a conventional re-pulper. A 1% solution of the chemical softener mixture is added to the Eucalyptus stock pipe before the stock pump at a rate of 0.2% by weight of the dry fibers. The absorption of the chemical softener mixture to CTMP can be enhanced via an in-line mixer. The Eucalyptus slurry is diluted to about 0.2% consistency at the fan pump.

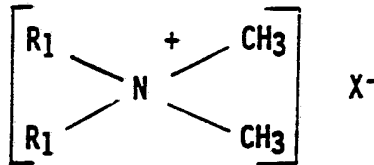
The treated furnish mixture (30% of NSK/70% of Eucalyptus) is blended in the head box and deposited onto a Fourdrinier photopolymer wire to form an embryonic web. Dewatering occurs through the photo-polymer wire and is assisted by a deflector and vacuum boxes. The photo-polymer wire has 400 discontinuous Linear Idaho cells per square inch, 70 percent of open areas and 2 mils of photo-polymer depth. The embryonic wet web is transferred from the photo-polymer wire, at a fiber consistency of about 15% at the point of transfer, to a photo-polymer belt having 711 Linear Idaho cells per square inch, 36 percent of knuckle areas and 8 mils of photo-polymer depth. Further de-watering is accomplished by vacuum assisted drainage until the web has a fiber consistency of about 28%. The patterned web is pre-dried by air blow-through to a fiber consistency of about 65% by weight. The web is

then adhered to the surface of a Yankee dryer with a sprayed creping adhesive comprising 0.25% aqueous solution of Polyvinyl Alcohol (PVA). The fiber consistency is increased to an estimated 99% before the dry creping the web with a doctor blade. The doctor blade has a bevel angle of about 24 degrees and is positioned with respect to the Yankee dryer to provide an impact angle of about 83 degrees; the Yankee dryer is operated at about 800 fpm (feet per minute) (about 244 meters per minute). The dry web is formed into roll at a speed of 700 fpm (214 meters per minute).

Two plies of the web are formed into tissue paper products and laminating together using conventional ply bonding techniques well known in the papermaking industry. The tissue paper has about 23 lbs./1000 sq. ft. basis weight, contains about 0.05% of DTMAMS, 0.05% PEG-400, and about 0.5% of the temporary wet strength resin. Importantly, the resulting tissue paper is soft, absorbent and has high temporary wet strength.

What is claimed is:

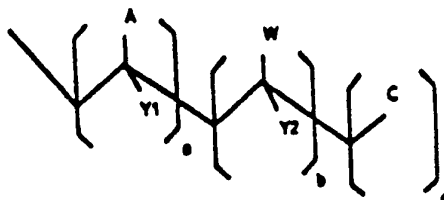
1. A strong, soft, absorbent tissue paper web characterized in that it comprises:
 - (a) papermaking fibers;
 - (b) from 0.01% to 2.0%, preferably from 0.01% to 0.5% by weight of a quaternary ammonium compound having the formula



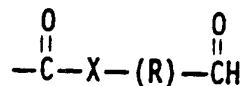
wherein each R_1 substituent is a C_{12} - C_{18} aliphatic hydrocarbon radical, preferably C_{16} - C_{18} alkyl and X^- is a compatible anion;

- (c) from 0.01% to 2.0%, preferably from 0.01% to 0.5% by weight of a polyhydroxy plasticizer; and
 - (d) from 0.01% to 3.0%, preferably from 0.1% to 1.5% by weight of a water-soluble temporary wet strength resin.
2. The paper web of Claim 1 wherein said polyhydroxy plasticizer is selected from the group consisting of glycerol and polyethylene glycols having a molecular weight from 200 to 2000, preferably a polyethylene glycol having a molecular weight from 200 to 600.
3. The paper web of Claim 1 or 2 wherein X^- is a halogen or methyl sulfate, preferably X^- is methyl sulfate.

4. The paper web of any of Claims 1-3 wherein said water-soluble temporary wet strength resin comprises a polymer characterized by the substantially complete absence of nucleophilic functionalities and having the formula:



wherein: A is

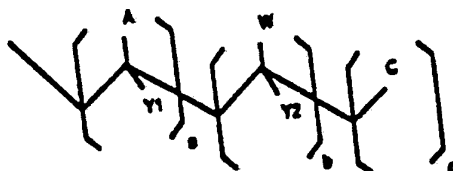


and X is --O--, --NCH₃--, and R is a substituted or unsubstituted aliphatic groups; Y₁ and Y₂ are independently --H, --CH₃ or a halogen; W is a nonnucleophilic, water-soluble nitrogen heterocyclic moiety; C is a cationic monomeric unit; the mole percent of a is from 30% to 70%, the mole percent of b is from 30% to 70%, and the mole percent of c is from 1% to 40%; and said polymer has an average molecular weight of between 30,000 and 200,000.

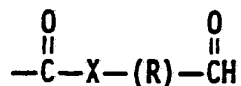
5. The paper web of any of Claims 1-3 wherein said water-soluble temporary wet strength resin is a cationic starch-based resin.
6. The paper web of any of Claims 1-5 wherein said polyhydroxy plasticizer is a polyethylene glycol having a molecular weight from 200 to 600.
7. The tissue paper of any of Claims 1-6 wherein said quaternary ammonium compound is di(hydrogenated tallow)dimethylammonium and wherein X⁻ is methyl sulfate.

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8. The paper web of Claim 6 or 7 wherein said water-soluble temporary wet strength resin comprises a polymer characterized by the substantially complete absence of nucleophilic functionalities and having the formula:



wherein: A is



and X is --O--, --NCH₃--, and R is a substituted or unsubstituted aliphatic groups; Y₁ and Y₂ are independently --H, --CH₃ or a halogen; W is a nonnucleophilic, water-soluble nitrogen heterocyclic moiety; C is a cationic monomeric unit; the mole percent of a is from 30% to 70%, the mole percent of b is from 30% to 70%, and the mole percent of c is from 1% to 40%; and said polymer has an average molecular weight of between 30,000 and 200,000.

9. The paper web of Claim 4 or 8 wherein said water-soluble temporary wet strength resin further comprises a cationic starch-based temporary wet strength resin.

INTERNATIONAL SEARCH REPORT

PCT/US 92/08898

International Application No

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) ⁶		
According to International Patent Classification (IPC) or to both National Classification and IPC		
Int.Cl. 5 D21H17/06; D21H17/07; D21H17/45		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁷		
Classification System	Classification Symbols	
Int.Cl. 5	D21H	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁸		
III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹		
Category ^o	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
X	<p>DATABASE WPIL Section Ch, Week 8833, Derwent Publications Ltd., London, GB; Class A97, AN 88-231810 & JP,A,63 165 597 (OJI PAPER KK ET AL.) 8 July 1988 see abstract</p> <p style="text-align: center;">---</p>	1, 3, 7
P, X	<p>DATABASE WPIL Section Ch, Week 9221, Derwent Publications Ltd., London, GB; Class E16, AN 92-170428 & JP,A,4 100 995 (NIPPON OILS & FATS) 2 April 1992 see abstract</p> <p style="text-align: center;">---</p> <p style="text-align: right;">-/--</p>	1-3
<p>^o Special categories of cited documents :¹⁰</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"I" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"&" document member of the same patent family</p>		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
25 FEBRUARY 1993	10. 03. 93	
International Searching Authority	Signature of Authorized Officer	
EUROPEAN PATENT OFFICE	SONGY Odile	

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		Relevant to Claim No.
Category ^o	Citation of Document, with indication, where appropriate, of the relevant passages	
A	EP,A,0 049 924 (THE PROCTER & GAMBLE COMPANY) 21 April 1982 see the whole document ---	1-9
A	WO,A,8 804 704 (WEYERHAEUSER COMPANY) 30 June 1988 see page 4, line 5 - page 5, line 16; table 1 ---	2,6
A	EP,A,0 350 276 (THE PROCTER & GAMBLE COMPANY) 10 January 1990 & US,A,4 981 557 cited in the application -----	4-8

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

US 9208898
SA 66040

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report.
The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information. 25/02/93

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP-A-0049924	21-04-82	US-A- 4351699	28-09-82
		AU-A- 7633481	22-04-82
		JP-A- 57136426	23-08-82
		US-A- 4441962	10-04-84

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		JP-A- 2124911	14-05-90
		US-A- 5138002	11-08-92
