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CA 2710784 A1 2009/07/09

(21) **2 710 784**

**(12) DEMANDE DE BREVET CANADIEN
CANADIAN PATENT APPLICATION**

(13) A1

(86) Date de dépôt PCT/PCT Filing Date: 2008/12/20
(87) Date publication PCT/PCT Publication Date: 2009/07/09
(85) Entrée phase nationale/National Entry: 2010/06/18
(86) N° demande PCT/PCT Application No.: US 2008/087869
(87) N° publication PCT/PCT Publication No.: 2009/086224
(30) Priorité/Priority: 2007/12/21 (US12/004,719)

(51) Cl.Int./Int.Cl. *D21H 13/10* (2006.01),
D21H 13/20 (2006.01), *D21H 15/00* (2006.01)

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(54) Titre : PAPIERS CONTENANT DES FIBRIDES DERIVES DE DIAMINODIPHENYLSULFONE
(54) Title: PAPERS CONTAINING FIBRIDS DERIVED FROM DIAMINO DIPHENYL SULFONE

(57) Abrégé/Abstract:

This invention relates to papers made with fibrils containing a polymer or copolymer derived from a monomer selected from the group consisting of 4,4'diaminodiphenyl sulfone, 3,3'diaminodiphenyl sulfone, and mixtures thereof. Such papers have high thermal stability and accept ink more readily than papers made solely with aramid fibrils.



(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
9 July 2009 (09.07.2009)(10) International Publication Number
WO 2009/086224 A3(51) International Patent Classification:
D21H 13/10 (2006.01) **D21H 15/00** (2006.01)
D21H 13/20 (2006.01)

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(21) International Application Number:
PCT/US2008/087869(22) International Filing Date:
20 December 2008 (20.12.2008)(25) Filing Language:
English(26) Publication Language:
English(30) Priority Data:
12/004,719 21 December 2007 (21.12.2007) US(71) Applicant (for all designated States except US): **E. I. DU PONT DE NEMOURS AND COMPANY** [US/US]; 1007 Market Street, Wilmington, Delaware 19898 (US).

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(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, NO, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report (Art. 21(3))
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments (Rule 48.2(h))

(88) Date of publication of the international search report:
20 August 2009

WO 2009/086224 A3

(54) Title: PAPERS CONTAINING FIBRIDS DERIVED FROM DIAMINO DIPHENYL SULFONE

(57) Abstract: This invention relates to papers made with fibrids containing a polymer or copolymer derived from a monomer selected from the group consisting of 4,4'diaminodiphenyl sulfone, 3,3'diaminodiphenyl sulfone, and mixtures thereof. Such papers have high thermal stability and accept ink more readily than papers made solely with aramid fibrids.

TITLE OF INVENTION

5 PAPERS CONTAINING FIBRIDS DERIVED FROM
DIAMINO DIPHENYL SULFONE

Background of the Invention

10 1. **Field of the Invention.**

This invention relates to papers made with fibrils containing a polymer or copolymer derived from a monomer selected from the group consisting of 4,4'diaminodiphenyl sulfone, 3,3'diaminodiphenyl sulfone, and mixtures thereof.

15 Such papers have high thermal stability and accept ink more readily than papers made solely with aramid fibrils.

2. **Description of Related Art.**

20 Papers made from high performance materials have been developed to provide papers with improved strength and/or thermal stability. Aramid paper, for example, is synthetic paper composed of aromatic polyamides. Because of its heat and flame resistance, electrical insulating properties, toughness and flexibility, the paper has been used as electrical insulation material and a base for aircraft honeycombs. Of 25 these materials, Nomex® of DuPont (U.S.A.) is manufactured by mixing poly(metaphenylene isophthalamide) floc and fibrils in water and then subjecting the mixed slurry to papermaking process to make formed paper followed by hot calendering of the formed paper. This paper is known to have excellent electrical insulation properties and with strength and toughness, which remains high even at 30 high temperatures.

Generally such aramid papers are difficult to color and print; for some applications aramid papers are coated to provide a better surface for printing of bar codes and other indicia. This requires an additional step after paper manufacture and the resulting waste that is generated by an additional manufacturing step. Therefore, 35 there is an ongoing need for high performance papers with improved properties, particularly papers that will accept ink or color more readily than high performance papers such as known aramid papers.

Brief Summary of the Invention

This invention relates to a highly printable thermally stable paper comprising non-granular, fibrous or film-like polymer fibrils comprising a polymer or copolymer derived from an amine monomer selected from the group consisting of 4,4'-diaminodiphenyl sulfone, 3,3'-diaminodiphenyl sulfone, and mixtures thereof, the fibrils having an average maximum dimension of 0.1 to 1 mm, a ratio of maximum to minimum dimension of 5:1 to 10:1, and a thickness of no more than 2 microns; and at least one high performance floc selected from the group of para-aramid, meta-aramid, carbon, glass, and mixtures thereof, the floc having a length of from 2 to 25 mm. In various embodiments, this invention also relates to heat resistant tags and labels, wrapped wires and conductors, laminate structures, honeycomb structures, and electrical devices comprising this highly printable thermally stable paper. (As employed herein "film-like" means "film".)

This invention also relates to a process for making thermally stable paper comprising the steps of:

a) forming an aqueous dispersion of 10 to 95 parts by weight polymer fibrils comprising a polymer or copolymer derived from an amine monomer selected from the group consisting of 4,4'-diaminodiphenyl sulfone, 3,3'-diaminodiphenyl sulfone, and mixtures thereof and 90 to 5 parts by weight of at least one high performance floc selected from the group of para-aramid, meta-aramid, carbon, glass, liquid crystalline polyester, polyphenylene sulfide, polyether-ketone-ketone, polyether-ether-ketone, polyoxadiazole, polybenzazole, and mixtures thereof, based on the total weight of the floc and fibrils;

b) blending the dispersion to form a slurry,
c) draining the aqueous liquid from the slurry to yield a wet paper composition, and
d) drying the wet paper composition to make a formed paper.

If desired, the process includes the additional step of consolidating the formed paper under heat and pressure to make a calendered paper.

Detailed Description of the Invention

This invention relates to the use of polymer fibrils containing a polymer or copolymer derived from a monomer selected from the group consisting of 4,4'-diaminodiphenyl sulfone, 3,3'-diaminodiphenyl sulfone, and mixtures thereof in

papers for improved printability without sacrificing thermal stability of the paper. Such polymers have [SO₂] linkages that help promote printability of the paper.

The term “fibrils” as used herein, means a very finely-divided polymer product of small, filmy or irregular fibrous shape particles. There are essentially two types of fibrils; “filmy” fibrils and “fibrous shape” or “stringy” fibrils. Filmy fibrils are essentially two-dimensional particles having a length and width on the order of 100 to 1000 micrometers and a thickness of 0.1 to 1 micrometer. Fibrous shape or stringy fibrils usually have length of up to 2-3 mm, a width of 10 to 50 microns, and a thickness of 0.1 to 1 micrometer. Fibrils are made by streaming a polymer solution into a coagulating bath of liquid that is immiscible with the solvent of the solution. The stream of polymer solution is subjected to strenuous shearing forces and turbulence as the polymer is coagulated. The predominant shape of the fibrils is determined by the type of polymer and the particular processing conditions during their coagulation.

Preferably, fibrils have a melting point or decomposition point above 320°C. Fibrils are not fibers, but they are fibrous in that they have fiber-like regions connected by webs. In one embodiment, fibrils have an aspect ratio of 5:1 to 10:1. In another embodiment, fibrils are used wet in a never-dried state and can be deposited as a binder physically entwined about other ingredients or components of a paper. The fibrils can be prepared by any method including using a fibrilating apparatus of the type disclosed in U.S. Patent No. 3,018,091 where a polymer solution is precipitated and sheared in a single step. Fibrils can also be made via the processes disclosed in U.S. Patent Nos. 2,988,782 and 2,999,788.

The fibrils comprise a polymer or copolymer derived from an amine monomer selected from the group consisting of 4,4'-diaminodiphenyl sulfone, 3,3'-diaminodiphenyl sulfone, and mixtures thereof. Such polymers and copolymers generally having the structure:



wherein Ar1 and Ar2 are any unsubstituted or substituted six-membered aromatic group of carbon atoms and Ar1 and Ar2 can be the same or different. In some preferred embodiments Ar1 and Ar2 are the same. Still more preferably, the six-membered aromatic group of carbon atoms has meta- or para-oriented linkages versus the SO₂ group. This monomer or multiple monomers having this general structure are

reacted with an acid monomer in a compatible solvent to create a polymer. Useful acids monomers generally have the structure of



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wherein Ar3 is any unsubstituted or substituted aromatic ring structure and can be the same or different from Ar1 and/or Ar2. In some preferred embodiments Ar3 is a six-membered aromatic group of carbon atoms. Still more preferably, the six-membered aromatic group of carbon atoms has meta- or para-oriented linkages. In some 10 preferred embodiments Ar1 and Ar2 are the same and Ar3 is different from both Ar1 and Ar2. For example, Ar1 and Ar2 can be both benzene rings having meta-oriented linkages while Ar3 can be a benzene ring having para-oriented linkages. Examples of useful monomers include terephthaloyl chloride, isophthaloyl chloride, and the like. In some preferred embodiments, the acid is terephthaloyl chloride or its mixture with 15 isophthaloyl chloride and the amine monomer is 4,4'diaminodiphenyl sulfone. In some other preferred embodiments, the amine monomer is a mixture of 4,4'diaminodiphenyl sulfone and 3,3'diaminodiphenyl sulfone in a weight ratio of 3:1, which creates a fibrid made from a copolymer having both sulfone monomers.

In still another preferred embodiment, the fibrils contain a copolymer, the 20 copolymer having both repeat units derived from sulfone amine monomer and an amine monomer derived from paraphenylene diamine and/or metaphenylene diamine. In some preferred embodiments the sulfone amide repeat units are present in a weight ratio of 3:1 to other amide repeat units. In some embodiments, at least 80 mole percent of the amine monomers is a sulfone amine monomer or a mixture of sulfone 25 amine monomers. For convenience, herein the abbreviation "PSA" will be used to represent all of the entire classes of fibers made with polymer or copolymer derived from sulfone monomers as previously described.

In one embodiment, the polymer and copolymer derived from a sulfone monomer can preferably be made via polycondensation of one or more types of 30 diamine monomer with one or more types of chloride monomers in a dialkyl amide solvent suchs as N-methyl pyrrolidone, dimethyl acetamide, or mixtures thereof. In some embodiments of the polymerizations of this type an inorganic salt such as lithium chloride or calcium chloride is also present. If desired the polymer can be isolated by precipitation with non-solvent such as water, neutralized, washed, and

dried. The polymer can also be made via interfacial polymerization which produces polymer powder directly that can then be dissolved in a solvent for fiber production.

Specific methods of making PSA fibers or copolymers containing sulfone amine monomers are disclosed in Chinese Patent Publication 1389604A to Wang et al. This reference discloses a fiber known as polysulfonamide fiber (PSA) made by spinning a copolymer solution formed from a mixture of 50 to 95 weight percent 4,4'diaminodiphenyl sulfone and 5 to 50 weight percent 3,3'diaminodiphenyl sulfone copolymerized with equimolar amounts of terephthaloyl chloride in dimethylacetamide. Chinese Patent Publication 1631941A to Chen et al. also discloses a method of preparing a PSA copolymer spinning solution formed from a mixture of 4,4'diaminodiphenyl sulfone and 3,3'diaminodiphenyl sulfone in a mass ratio of from 10:90 to 90:10 copolymerized with equimolar amounts of terephthaloyl chloride in dimethylacetamide. Still another method of producing copolymers is disclosed in United States Patent No. 4,169,932 to Sokolov et al. This reference discloses preparation of poly(paraphenylene) terephthalamide (PPD-T) copolymers using tertiary amines to increase the rate of polycondensation. This patent also discloses the PPD-T copolymer can be made by replacing 5 to 50 mole percent of the paraphenylene diamine (PPD) by another aromatic diamine such as 4,4'diaminodiphenyl sulfone.

In one embodiment, a portion of the PSA fibrils can be replaced by another, second, non-granular, fibrous or film-like polymer binder. Such binders include fibrils made from another polymer or copolymer. In a preferred embodiment the polymer binder is selected from the group of meta-aramid fibrils, para-aramid fibrils, and mixtures thereof. The preferred meta-aramid fibrils are poly(metaphenylene isophthalamide) fibrils.

In one embodiment, it is believed that up to about 80 weight percent of the PSA fibrils can be replaced with MPD-I fibrils with good result. However, in a preferred embodiment, 20 to 50 weight percent of the PSA fibrils are replaced with MPD-I fibrils. It is believed the improved dyeability and printability of the paper due to the additional polysulfone groups provided by the PSA fibrils is retained even with only 20 weight percent PSA fibrils in the paper.

If desired, the fibrils in the paper can be filled with different fillers including carbon black, graphite, and mineral powders. In a preferred embodiment the filled

fibrils are PSA fibrils. Method of filling fibrils with carbon black or graphite is described, for example, in United States Patent No. 5,482,773 to Bair.

The PSA fibrils are combined with at least one high performance floc selected from the group of para-aramid, meta-aramid, carbon, glass, liquid crystalline polyester, polyphenylene sulfide, polyether-ketone-ketone, polyether-ether-ketone, polyoxadiazole, polybenzazole, and mixtures thereof.

By "floc" is meant fibers having a length of 2 to 25 millimeters, preferably 3 to 7 millimeters and a diameter of 3 to 20 micrometers, preferably 5 to 14 micrometers. If the floc length is less than 3 millimeters, the paper strength is severely reduced, and if the floc length is more than 25 millimeters, it is difficult to form a uniform paper web by a typical wet-laid method. If the floc diameter is less than 5 micrometers, it can be difficult to commercially produce with adequate uniformity and reproducibility, and if the floc diameter is more than 20 micrometers, it is difficult to form uniform paper of light to medium basis weights. Floc is generally made by cutting continuous spun filaments into specific-length pieces.

The high performance floc includes flocs of para-aramid, meta-aramid, carbon, glass, liquid crystalline polyester, polyphenylene sulfide, polyether-ketone-ketone, polyether-ether-ketone, polyoxadiazole polybenzazole, and mixtures thereof.

By aramid is meant a polyamide wherein at least 85% of the amide (-CONH-) linkages are attached directly to two aromatic rings. A para-aramid is such a polyamide that contains a para configuration or para-oriented linkages in the polymer chain, while meta-aramid is such a polyamide that contains a meta configuration or meta-oriented linkages in the polymer chain. Additives can be used with the aramid and, in fact, it has been found that up to as much as 10 percent, by weight, of other polymeric material can be blended with the aramid or that copolymers can be used having as much as 10 percent of other diamine substituted for the diamine of the aramid or as much as 10 percent of other diacid chloride substituted for the diacid chloride of the aramid. In some embodiments, the preferred para-aramid is poly(paraphenylene terephthalamide). Methods for making para-aramid fibers useful are generally disclosed in, for example, United States Patent Nos. 3,869,430; 3,869,429; and 3,767,756. Various forms of such aromatic polyamide organic fibers are sold under the trademarks of Kevlar® and Twaron® by respectively, E. I. du Pont de Nemours and Company, of Wilmington, Delaware; and Teijin, Ltd, of Japan. Also, fibers based on copoly(p-phenylene/3,4'-diphenyl ether terephthalamide) are defined

as para-aramid fibers as used herein. One commercially available version of these fibers is known as Technora® fiber also available from Teijin, Ltd.

In some embodiments, the preferred meta-aramids are poly(meta-phenylene isophthalamide)(MPD-I) and its copolymers. One such meta-aramid floc is Nomex® aramid fiber available from E. I. du Pont de Nemours and Company of Wilmington, DE, however, meta-aramid fibers are available in various styles under the trademarks Conex®, available from Teijin Ltd. of Tokyo, Japan,; Apyeil®, available from Unitika, Ltd. of Osaka, Japan; New Star® Meta-aramid, available from Yantai Spandex Co. Ltd, of Shandong Province, China; and Chinfunex® Aramid 1313 available from Guangdong Charming Chemical Co. Ltd., of Xinhui in Guangdong, China. Meta-aramid fibers are inherently flame resistant and can be spun by dry or wet spinning using any number of processes; however, U.S. Patent Nos. 3,063,966; 3,227,793; 3,287,324; 3,414,645; and 5,667,743 are illustrative of useful methods for making aramid fibers that could be used.

Additives can be used with the aramid and, in fact it has been found that up to as much as 10 percent, by weight, of other polymeric material can be blended with the aramid or that copolymers can be used having as much as 10 percent of other diamine substituted for the diamine of the aramid or as much as 10 percent of other diacid chloride substituted for the diacid chloride of the aramid.

Commercially available carbon fibers include Tenax® fibers available from Toho Tenax America, Inc, and commercially available glass fibers include borosilicate glass microfiber type 253 sold by Johns Manville Co. Useful commercially available liquid crystal polyester fibers include Vectran® HS fiber available from Swicofil AG Textile Services. Polyphenylene sulfide fiber has good

heat resistance, chemical resistance, and hydrolysis resistance. At least 90% of the constituent units of these fibers are of a polymer or copolymer having phenylene sulfide structural units of -(C₆H₄-S)-. Polyphenylene sulfide fiber is sold under the tradenames Ryton® by American Fibers and Fabrics, Toray PPS® by Toray Industries Inc., Fortron® by Kureha Chemical Industry Co. and Procon® by Toyobo Co.

Polyether-ketone-ketone and polyether-ether-ketone fibers include Zyex® PEEK and Zyex® PEK fibers available from Zyex Ltd. (UK). Polyoxadiazole fibers also have good heat resistance and are disclosed in, for example, U. S. Patent No. 4,202,962 to Bach and the Encyclopedia of Polymer Science and Engineering, Vol 12, p. 322-339 (John Wiley & Sons, New York, 1988). In some embodiments the

polyoxadiazole fiber contains polyarylene-1,3,4-oxadiazole polymer, polyarylene-1,2,4-oxadiazole polymer, or mixtures thereof. In some preferred embodiments, the polyoxadiazole fiber contains polyparaphhenylene-1,3,4-oxadiazole polymer. Suitable polyoxadiazole fibers are known commercially under various tradenames such as

5 Oxalon®, Arselon®, Arselon-C® and Arselon-S® fiber. Useful commercially available polybenzazole fibers include Zylon® PBO-AS (Poly(p-phenylene-2,6-benzobisoxazole) fiber, Zylon® PBO-HM (Poly(p-phenylene-2,6-benzobisoxazole)) fiber, available from Toyobo, Japan.

In some preferred embodiments the high performance floc has a high modulus.

10 As used herein high modulus fibers are those having a tensile or Young's modulus of 600 grams per denier (550 grams per dtex) or greater. High modulus of the floc provides stiffness and also can provide improved dimensional stability to the paper that can translate to the final applications of the paper. In a preferred embodiment, the Young's modulus of the fiber is 900 grams per denier (820 grams per dtex) or greater.

15 In the preferred embodiment, the fiber tenacity is at least 21 grams per denier (19 grams per dtex) and its elongation is at least 2% so as to provide a high level of mechanical properties to the final application of the paper.

In a preferred embodiment the high modulus floc is heat resistant fiber. By "heat resistant fiber" it is meant that the fiber preferably retains 90 percent of its fiber weight when heated in air to 500° C at a rate of 20 degrees Celsius per minute. Such fiber is normally flame resistant, meaning the fiber or a fabric made from the fiber has a Limiting Oxygen Index (LOI) such that the fiber or fabric will not support a flame in air, the preferred LOI range being about 26 and higher. The preferred heat resistant fiber is para-aramid fiber, particularly poly(paraphenylene terephthalamide) fiber.

25 In one embodiment, the fibrils are combined with at least one high performance floc and at least one other floc. In one preferred embodiment, the at least one other floc is a floc that contains a polymer or copolymer derived from a monomer selected from the group consisting of 4,4'diaminodiphenyl sulfone, 3,3'diaminodiphenyl sulfone, and mixtures thereof.

30 The fibrils and the floc are combined to form a thermally stable paper. As employed herein the term paper is employed in its normal meaning and it can be prepared using conventional paper-making processes and equipment and processes. The fibrous material, i.e. fibrils and floc can be slurried together to form a mix which is converted to paper such as on a Fourdrinier machine or by hand on a handsheet

mold containing a forming screen. Reference may be made to Gross USP 3,756,908 and Hesler et al. USP 5,026,456 for processes of forming fibers into papers. If desired, once paper is formed it is calendered between two heated calendering rolls with the high temperature and pressure from the rolls increasing the bond strength of 5 the paper. Calendering also provides the paper with a smooth surface for printing. Several plies with the same or different compositions can be combined together into the final paper structure during forming and/or calendering. In one embodiment, the paper has a weight ratio of fibrils to floc in the paper composition of from 95:5 to 10:90. In one preferred embodiment, the paper has a weight ratio of fibrils to floc in 10 the paper composition of from 60:40 to 10:90.

In one embodiment, the formed paper has a density of about 0.1 to 0.5 grams per cubic centimeter. In some embodiments the thickness of the formed paper ranges from about 0.002 to 0.015 inches. The thickness of the calendered paper is dependent upon the end use or desired properties and in some embodiments is typically from 15 0.001 to 0.005 mils (25 to 130 micrometers) thick. In some embodiments, the basis weight of the paper is from 0.5 to 6 ounces per square yard (15 to 200 grams per square meter).

Additional ingredients such as fillers for the adjustment of paper conductivity and other properties, pigments, antioxidants, etc in powder or fibrous form can be 20 added to the paper composition of this invention. If desired, an inhibitor can be added to the paper to provide resistance to oxidative degradation at elevated temperatures. Preferred inhibitors are oxides, hydroxides and nitrates of bismuth. An especially effective inhibitor is a hydroxide and nitrate of bismuth. One desired method of 25 incorporating such fillers into the papers is by first incorporating the fillers into the fibrils during fibril formation. Other methods of incorporating additional ingredients into the paper include adding such components to the slurry during paper forming, spraying the surface of the formed paper with the ingredients and other conventional techniques.

When PSA fibrils are incorporated as binders in papers, the sulfone groups in 30 the PSA fibrils provide improved sites for accepting printing ink on the surface of the papers over papers having, for example, only MPD-I fibrils as binders.

In one embodiment the thermally stable paper can be made using a process comprising the steps of:

- a) forming an aqueous dispersion of 10 to 95 parts by weight polymer fibrils comprising a polymer or copolymer derived from an amine monomer selected from the group consisting of 4,4'diaminodiphenyl sulfone, 3,3'diaminodiphenyl sulfone, and mixtures thereof, and 90 to 5 parts by weight of at least one high performance floc selected from the group of para-aramid, meta-aramid, carbon, glass, liquid crystalline polyester, polyphenylene sulfide, polyether-ketone-ketone, polyether-ether-ketone, polyoxadiazole, polybenzazole, and mixtures thereof, based on the total weight of the floc and fibrils;
- 5 b) blending the dispersion to form a slurry,
- c) draining the aqueous liquid from the slurry to yield a wet paper composition, and
- 10 d) drying the wet paper composition to make a formed paper.

In another embodiment, the floc is a mixture of flocs further comprising at least one floc containing a polymer or copolymer derived from a monomer selected from the group consisting of 4,4'diaminodiphenyl sulfone, 3,3'diaminodiphenyl sulfone, and mixtures thereof.

The paper can be formed on equipment of any scale from laboratory screens to commercial-sized papermaking machinery, such as a Fourdrinier or inclined wire machines. The general process involves making a dispersion of the fibrils and floc, and optionally additional ingredients such as fillers, in an aqueous liquid, draining the liquid from the dispersion to yield a wet composition and drying the wet paper composition.

The dispersion can be made either by dispersing the floc in the aqueous liquid and then adding the fibrils or by dispersing the fibrils in the liquid and then adding the fibers. The dispersion can also be made by combining a floc-containing dispersion with a fiber-containing dispersion. The concentration of floc in the dispersion can range from 0.01 to 1.0 weight percent based on the total weight of the dispersion. The concentration of a fibrils in the dispersion can be up to 20 weight percent based on the total weight of solids.

In some embodiments, a portion of the PSA fibrils the aqueous dispersion can be replaced by another, second, non-granular, fibrous or film-like polymer binder. Such binders include fibrils made from another polymer or copolymer. In a preferred embodiment the polymer binder is selected from the group of meta-aramid fibrils,

para-aramid fibrids, and mixtures thereof. The preferred meta-aramid fibrids are poly(metaphenylene isophthalamide) fibrids.

In one preferred embodiment, dye or pigment is included in the aqueous dispersion to make a colored paper. Any dye or pigment compatible with the final application of the paper and that is adequately bound to the sulfone groups in the paper can be used. In one preferred embodiment, the dye or pigment is added in an amount that results in the desired coloration in the final paper. The preferred dyes and pigments can withstand the calendering process, that is, temperatures of 250 degrees Celsius or greater; in some especially preferred embodiments the dyes and pigments can withstand temperatures of 310 degrees Celsius or greater.

The aqueous liquid of the dispersion is generally water, but may include various other materials such as pH-adjusting materials, forming aids, surfactants, defoamers and the like. The aqueous liquid is usually drained from the dispersion by conducting the dispersion onto a screen or other perforated support, retaining the dispersed solids and then passing the liquid to yield a wet paper composition. The wet composition, once formed on the support, is usually further dewatered by vacuum or other pressure forces and further dried by evaporating the remaining liquid.

A next step, which can be performed if higher density and strength are desired, is calendering one or more layers of the paper in the nip of metal-metal, metal-composite, or composite-composite rolls. Alternatively, one or more layers of the paper can be compressed in a platen press at a pressure, temperature and time, which are optimal for a particular composition and final application. Also, heat-treatment as an independent step before, after or instead of calendering or compressing, can be conducted if strengthening or some other property modification is desired without or in addition to densification.

The paper is useful as printable material for high temperature tags, labels, and security papers. The paper can also be used as a component in materials such as printed wiring boards; or where dielectric properties are useful, such as electrical insulating material for use in motors, transformers and other power equipment. In these applications, the paper can be used by itself or in laminate structures either with or without impregnating resins, as desired. In another embodiment, the paper is used as an electrical insulative wrapping for wires and conductors. The wire or conductor can be totally wrapped, such a spiral overlapping wrapping of the wire or conductor, or can wrap only a part or one or more sides of the conductor as in the case of square

conductors. The amount of wrapping is dictated by the application and if desired multiple layers of the paper can be used in the wrapping. In another embodiment, the paper can also be used as a component in structural materials such as core structures or honeycombs. For example, one or more layers of the paper may be used as the 5 primarily material for forming the cells of a honeycomb structure. Alternatively, one or more layers of the paper may be used in the sheets for covering or facing the honeycomb cells or other core materials. Preferably, these papers and/or structures are impregnated with a resin such as a phenolic, epoxy, polyimide or other resin. However, in some instances the paper may be useful without any resin impregnation.

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Test Methods

Thickness and Basis Weight (Grammage) were determined for papers of this 15 invention in accordance with ASTM D 374 and ASTM D 646 correspondingly. At thickness measurements, method E with pressure on specimen of about 172 kPa was used.

Density (Apparent Density) of papers was determined in accordance with ASTM D 202.

20 Tensile Strength and Elongation were determined for papers of this invention on an Instron-type testing machine using test specimens 2.54 cm wide and a gage length of 18 cm in accordance with ASTM D 828.

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Example 1

Fibrils from a copolymer of 4, 4'diaminodiphenyl sulfone and 3, 3'diaminodiphenyl sulfone were prepared as follows. A 10% solution of a copolymer of 4, 4'diaminodiphenyl sulfone and 3, 3'diaminodiphenyl sulfone in DMAc was 30 precipitated in a water bath at high shear stress using a Waring blender. The precipitate was then washed with water and dispersed in the same blender with water for 10 minutes to form fibrils. The fibrils had a freeness of about 450 ml Shopper-Riegler.

A water slurry of these fibrils containing 2.0 grams (dry weight) of the solids 35 was placed together with 2 grams of poly(metaphenylene isophthalamide) floc in a laboratory mixer (British pulp evaluation apparatus) with about 1600 g of water and

agitated for 3 minutes, forming a 50/50 percent by weight mixture of fibrils and floc. The poly(metaphenylene isophthalamide) floc had a linear density of 0.22 tex (2.0 denier) and length of 0.64 cm.

The dispersion was then poured, with 8 liters of water, into an approximately 5 21 x 21 cm handsheet mold and a wet-laid sheet was formed. The sheet was placed between two pieces of blotting paper, hand couched with a rolling pin and dried in a handsheet dryer at 190°C to make formed paper. After drying, the formed paper was calendered in the metal-metal nip at temperature of 300 C and linear pressure of about 3000 N/cm.

10 The final calendered paper had a basis weight of 83.4 g/m², a thickness of 0.094 mm, a density of 0.89 g/cm³, a tensile strength of 26.0 N/cm, and an elongation of 3.22%. This paper is printed without prior coating to provide a printed label or tag.

Example 2

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Example 1 was repeated to make first formed and then calendered paper, however the 50/50 slurry blend of fibrils and floc contained 1.7 grams (dry weight) of fibrils and 1.7 grams of poly(paraphenylene terephthalamide) floc. The 20 poly(paraphenylene terephthalamide) floc had a linear density 0.17 tex (1.5 denier) and length of 0.64 cm. The final calendered paper had a basis weight of 71.9 g/m², a thickness of 0.079 mm, a density of 0.91 g/cm³, a tensile strength of 23.3 N/cm, and an elongation of 1.90%. This paper is printed without prior coating to provide a printed label or tag.

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Example 3

The process of Example 1 is repeated to make first formed and then calendered paper with the addition of 2 grams of the Basacryl Red GL dye, available 30 from BASF Wyandotte Corp., Charlotte, N.C., is added to the 1600 grams of water slurry. The fibrils accept the red dye and a colored paper is made.

Example 4

35 Example 1 is repeated to make first formed and then calendered paper except that 10 weight percent of the poly(metaphenylene isophthalamide) MPD-I floc is replaced with floc made from a copolymer derived from 4,4'diaminodiphenyl sulfone

and 3,3'diaminodiphenyl sulfone amine monomers(~70:30 ratio) PSA. The PSA floc has the same cut length as the MPD-I floc. The final floc mixture has a composition of 80% MPD-I floc, 10% PET floc, and 10% PSA floc. The final calendered paper is printed without prior coating to provide a printed label or tag.

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Example 5

Example 1 is repeated to make first formed and then calendered paper except that in the aqueous dispersion 20 weight percent of the PSA fibrils are replaced with 10 MPD-I fibrils. The final calendered paper is printed without prior coating to provide a printed label or tag.

CLAIMS

5 **What is claimed is:**

1. A highly printable thermally stable paper, comprising:
10 a) non-granular, fibrous or film-like polymer fibrils comprising a polymer or copolymer derived from an amine monomer selected from the group consisting of 4,4'-diaminodiphenyl sulfone, 3,3'-diaminodiphenyl sulfone, and mixtures thereof, the fibrils having an average maximum dimension of 0.1 to 1 mm, a ratio of maximum to minimum dimension of 5:1 to 10:1, and a thickness of no more than 2 microns; and

15 b) at least one high performance floc selected from the group of para-aramid, meta-aramid, carbon, glass, liquid crystalline polyester, polyphenylene sulfide, polyether-ketone-ketone, polyether-ether-ketone, polyoxadiazole, polybenzazole, and mixtures thereof, the floc having a length of from 2 to 25 mm;

20 wherein, the weight ratio of fibrils to floc in the paper composition is from 95:5 to 10:90.

2. The paper of claim 1, further comprising:

25 c) at least one floc containing a polymer or copolymer derived from a monomer selected from the group consisting of 4,4'-diaminodiphenyl sulfone, 3,3'-diaminodiphenyl sulfone, and mixtures thereof.

3. The paper of claim 1 wherein the meta-aramid fiber is poly (metaphenylene isophthalamide) fiber.

4. The paper of claim 1, further comprising a second non-granular, fibrous or film-like polymer binder.

35 5. The paper of claim 4 wherein the polymer binder is selected from the group of meta-aramid fibrils, para-aramid fibrils, and mixtures thereof.

6. The paper of claim 5 wherein the meta-aramid is poly (metaphenylene isophthalamide).

7. A heat resistant tag or label, or security paper comprising the paper of claim 1.

8. A wire or conductor wrapped with the paper of claim 1.

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9. A laminate structure comprising the paper of claim 1.

10. A honeycomb structure comprising the paper of claim 1.

10 11. An electrical device comprising the paper of claim 1.

12. A process for making thermally stable formed paper comprising the steps of:

15 a) forming an aqueous dispersion of 10 to 95 parts by weight polymer fibrils comprising a polymer or copolymer derived from an amine monomer selected from the group consisting of 4,4'diaminodiphenyl sulfone, 3,3'diaminodiphenyl sulfone, and mixtures thereof, and 90 to 5 parts by weight of at least one high performance floc selected from the group of para-aramid, meta-aramid, carbon, glass, liquid crystalline polyester, polyphenylene sulfide, polyether-ketone-ketone, polyether-ether-ketone, polyoxadiazole, polybenzazole, and mixtures thereof, based on the total weight of the floc and fibrils;

20 b) blending the dispersion to form a slurry,

c) draining the aqueous liquid from the slurry to yield a wet paper composition, and

d) drying the wet paper composition to make a formed paper.

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13. The process of claim 12 wherein the water is drained from the slurry via a screen or wire belt.

14. The process of claim 12 further comprising at least one floc containing a

30 polymer or copolymer derived from a monomer selected from the group consisting of 4,4'diaminodiphenyl sulfone, 3,3'diaminodiphenyl sulfone, and mixtures thereof.

15. The process of claim 12 further comprising calendering the formed paper with heat and pressure.