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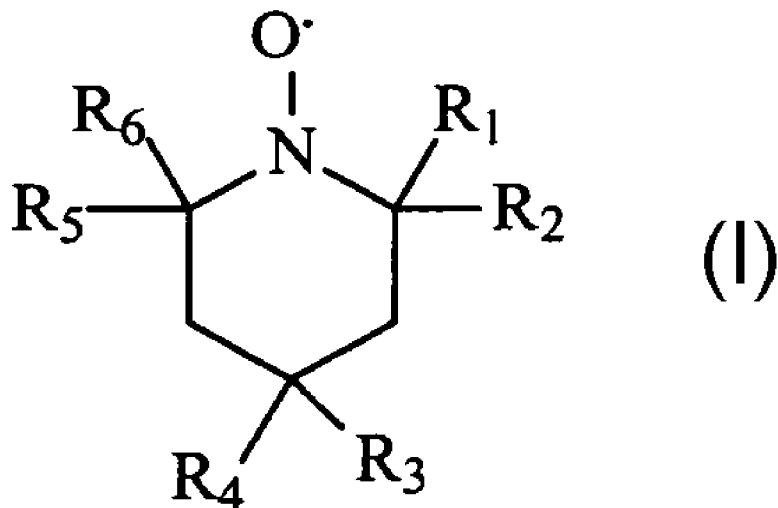
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(54) Title: HEAT TRANSFERABLE MATERIAL FOR IMPROVED IMAGE STABILITY



(57) Abstract: A heat transferable material includes a heat transferable polymeric binder and a light stabilizer that is an N-oxyl radical derived from a hindered amine, the N-oxyl radical having the following formula, (I) wherein R₁, R₂, R₅, and R₆ are each independently selected from a straight or branched C₁-C₆ alkyl, and R₃ and R₄ are each independently selected from H, OH, OR, COOH, or COOR, wherein R is a straight or branched C₁-C₆ alkyl or alkene, and having a molecular weight of 600 or less, is described. The heat transferable material can be in one or more sections or patches on a thermal donor element to provide a protective overcoat material. Optionally, a patch in the donor element can also include a dye. The heat transferable material provides better image stability and improved iridescence when applied to a thermal, inkjet, electrophotographic, or silver halide receiver.

HEAT TRANSFERABLE MATERIAL
FOR IMPROVED IMAGE STABILITY

FIELD OF THE INVENTION

5 The present invention relates to a heat transferable material including a heat transferable polymeric binder and an N-oxyl radical that is derived from a hindered amine, and that behaves as a light stabilizer that provides improved image stability and reduced iridescence when applied to a receiver.

10

BACKGROUND OF THE INVENTION

There are many ways of forming an image. Images can be formed through thermal transfer of dyes, inkjet applications, electrophotographic reproduction, and silver halide image development. Also known is that all such images are susceptible to environmental factors, particularly light fade. Thermal, 15 inkjet, and electrophotographic images also can suffer from iridescence problems, which are unsightly to the viewer. Typically, iridescence is caused by the interaction between the materials on the receiver and any materials applied to the receiver in forming the image.

20 To form any printed image, the image is either chemically developed from film, or developed from an electronic signal generated from either a digital capture device, or scanning of a film. For thermal, inkjet, and electrophotographic prints, electronic signals indicating appropriate colors are used to produce cyan, magenta and yellow color signals. These signals are then transmitted to a printer where colored material is transferred to a receiver. A color 25 hard copy is thus obtained that corresponds to the original image.

Thermal, ink jet, and electrophotographic prints are susceptible to retransfer of colorants to adjacent surfaces and to discoloration by fingerprints because the colorants remain at the surface of the receiver. Heat can be used to drive the colorants deeper into the receiver. Application of a protective overcoat 30 on these types of prints, as well as silver halide prints, is also known, and effectively reduces retransfer and discoloration by adding a protective polymeric layer over the image.

The protective overcoat can also provide improved light stability to the underlying imaging colorants, including dyes. The most common approach is to filter out UV radiation since it is known that UV radiation is detrimental to the underlying colorants. Improved image stability can be achieved with the addition

5 of a UV absorbing dye in a protective overcoat, as described in U.S. Patent 4,522,881. This approach has practical limitations on the amount of UV radiation that can be absorbed because there is a practical limitation on the thickness of the protective overcoat as well as the concentration of the UV absorbing dye that can be applied.

10 Improved image stability can also be achieved by incorporating light stabilizers in close proximity to the colorants within the receiver. Light stabilizers can be added to the receiver during manufacture by aqueous or solvent coating or thermal extrusion of materials incorporating the light stabilizers. If thermal extrusion is used, only light stabilizers with very high thermal stabilities
15 can be used due to the temperatures of extrusion, typically 250°C or higher. The light stabilizers must be incorporated within the receiver in such a manner that they will react with the colorants when applied to the receiver.

U.S. Patent 5,332,713 discloses a transferable protection overcoat on a donor element for transfer to a thermal print. The transferable protection
20 overcoat comprises poly(vinyl formal), poly(vinyl benzal) or poly(vinyl acetal) containing at least 5 mole % hydroxyl. The overcoat provides inferior gloss and iridescence performance due to refractive index mismatch with the dye receiving layer.

U.S. Patent 5,387,573 discloses a protective overcoat including
25 particles in an amount of up to 75% of the thickness of the heat transferable protective overcoat. Although the particles reduce the iridescence problems, the particles lower the gloss of the imaged print.

U.S. Patent 5,670,449 discloses the use of elastomeric beads in a protective overcoat for better raw-stock keeping, but the gloss performance of
30 these protective overcoats is not optimum.

U.S. Patent 6,942,956 discloses a protective overcoat comprising a gloss-enhancing agent, and a mixture of inorganic and organic particles. In some

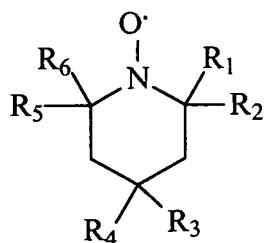
embodiments, the protection layer contains from 5% to 60% by weight inorganic particles, from 25% to 80% by weight polymeric binder and from 5% to 60% by weight of organic particles, and an effective amount of at least one gloss-enhancing compound. The gloss enhancing compound consists of an organic 5 molecule that is essentially colorless, does not scatter light, is substantially not absorbing of light at a wavelength from 400 to 800 nm, and has a maximum absorption at a wavelength less than 400 nm. The inorganic particles, e.g. silica, are required to provide smooth protective overcoat tear-off, but these degrade 10 gloss and are detrimental to the gravure coating quality. The organic particles are required to reduce iridescence, but these reduce gloss. The gloss improvement 15 provided is not adequate.

U.S. Patent 7,301,012 and 7,384,138 disclose the use of hindered amine light stabilizers (HALS) in a receiver to provide image dye stability.

There remains a need for a heat transferable protective overcoat 15 that provides greater image stability, reduced iridescence, and can be manufactured at low cost.

SUMMARY OF THE INVENTION

The present invention relates to a heat transferable donor element 20 comprising a polymeric support, the support having at least one portion thereof coated with a heat transferable material comprising a polymeric binder and a light stabilizer that is an N-oxyl radical derived from a hindered amine, the N-oxyl radical having the following formula:



25 wherein R₁, R₂, R₅, and R₆ are each independently selected from a straight or branched C₁-C₆ alkyl or alkene, and R₃ and R₄ are each independently selected from H, OH, OR, COOH, or COOR, wherein R is a straight or branched C₁-C₆ alkyl or alkene, and having a molecular weight of 600 or less.

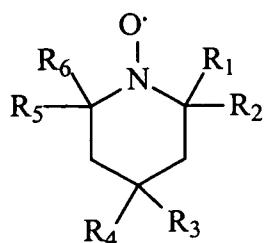
This invention also provides an assemblage for transferring material imagewise to a receiver element from the donor element of this invention. The heat transferable material can be in one or more protective overcoat patches. Suitable receiver elements for transfer of a protective overcoat 5 include any colorant containing material, including, for example, an inkjet receiver, a thermal receiver, an electrophotographic receiver, or a silver halide print.

The heat transferable donor element of this invention containing a transferable polymeric binder and a transferable N-oxyl radical light stabilizer 10 provides the advantages of reducing light fade, reducing iridescence, and lowering costs for image production by reducing or eliminating the need for UV absorbing materials. Other advantages will be apparent upon review of this document in full.

15

DETAILED DESCRIPTION OF THE INVENTION

The invention relates to a heat transferable donor element for use with receivers for thermal, inkjet, and electrophotographic printing, as well as silver halide prints. A heat transferable material is present on at least a portion of the heat transferable donor element, wherein the donor element has a support and 20 disposed on at least one side of the support, a heat transferable polymeric binder and a light stabilizer that is an N-oxyl radical that is derived from a hindered amine. Sometimes this N-oxyl radical is known in the art as a “hindered amine light stabilizer” (HALS). The N-oxyl radical has a molecular weight of at least 140 and less than 600 and generally has the following formula:



25

wherein R₁, R₂, R₅, and R₆ are each independently selected from a straight or branched C₁-C₆ alkyl or alkene, and R₃ and R₄ are each independently selected

from H, OH, OR, COOH, or COOR, wherein R is a straight or branched C1-C6 alkyl or alkene.

According to various embodiments, R₃ and R₄ can each separately be chosen from CH₂CH₃, CH₃, or H. For example, R₃ and R₄ can be both

5 hydrogen. According to various embodiments, R₁, R₂, R₅, and R₆ can each independently be chosen from CH₂CH₃, CH₃, or H. Also, R₁, R₂, R₅, and R₆ can each independently be chosen from CH₃ or H, and typically each is CH₃. A useful compound is available commercially as TEMPO from Evonik/Degussa.

It is noted that the light stabilizer is an oxyl radical, and is a singlet 10 oxygen quencher. It is present in an active form. When present in a heat transferable material on a donor element, the light stabilizer is transferred like a colorless dye from the donor element or to a receiver element upon printing. That is, the light stabilizer migrates upon heating from the heat transferable donor element to the receiver element. For this reason, an N-oxyl radical light stabilizer 15 with a low molecular weight as noted above is desired so that it can more easily transfer between the donor element and receiver element. Similarly, side chains for R₁-R₆ with less steric hindrance are useful to enable migration.

It has been observed that the N-oxyl radical appears to react to the presence of plasticizer and bind thereto. The presence of a plasticizer in the 20 receiver layer to which the N-oxyl radical is being transferred appears to bind the N-oxyl radical and prevent any retransfer or further migration into the other receiver layers. Presence of plasticizer in the heat transferable material including the N-oxyl radical can hamper transfer of light stabilizer to the receiver element, instead mordanting the HALS in the patch on the donor element. It is desirable to 25 have little or no plasticizer present in the heat transferable material including the N-oxyl radical, for example, an amount of plasticizer of 5% or less by weight of the heat transferable material, typically 3% or less by weight, more typically 0 to 2% by weight. In many embodiments, there is no plasticizer in the heat transferable material.

30 Any material can be used as the support for the donor element of the invention provided it is dimensionally stable and can withstand the heat of thermal transfer, for example from a thermal printing head. Suitable materials can

include, for example, polyesters such as poly(ethylene terephthalate); polyamides; polycarbonates; glassine paper; condenser paper; cellulose esters such as cellulose acetate; fluorine polymers such as poly(vinylidene fluoride) or poly(tetrafluoroethylene-co-hexafluoropropylene); polyethers such as

5 polyoxymethylene; polyacetals; polyolefins such as polystyrene, polyethylene, polypropylene or methylpentene polymers; and polyimides such as polyimide amides and polyetherimides. The support can have a thickness of from 2 to 30 μm , although thicker or thinner supports could be used for specific applications.

10 According to certain embodiments where a high gloss image is desired, the support can have a surface roughness, Ra, of 18 nm or less on the side of the support on which the heat transferable material is provided.

The heat transferable material can be provided in one or more sections, or patches, on the donor element, or a single heat transferable material can coat the length of the donor element. The donor element can be provided as 15 sheets or rolls of any desired width and length suitable for the intended thermal transfer apparatus. The patches on a donor element can be the same or different, and can be in a repeating pattern if desired. The patches provide a protective overcoat. A donor element can also include one or more colored dye patches followed by a protective overcoat patch, or a single color patch followed by a 20 protective overcoat patch. The sequence can be repeated, if desired. An exemplary sequence commonly used in thermal dye diffusion printing is a repeat of yellow, magenta, cyan, and protective overcoat patches. The present invention is directed to the protective overcoat patches that can be used solely in the donor element or as protective overcoat patches along with one or more color patches.

25 A UV absorber can be present in amounts of 20% or less by weight, or 5% or less in the heat transferable material. In some instances no UV absorber is present.

For the protective overcoat patches, the heat transferable material comprises, besides the N-oxyl radical light stabilizer, one or more heat 30 transferable polymeric binders. Any known heat transferable polymeric binder can be used. For example, the present invention can include the use of a heat transferable polymeric binder blend for use in heat transferable material such as a

polyvinyl acetal resin blended with a polystyrene/allyl alcohol copolymer incorporating one or more heat transferable N-oxyl radical light stabilizers derived from the hindered amines that results in improved image stability of the resulting dye diffusion thermal transfer prints. This resin blend may be used in the

5 protective overcoat layer or patch. The protective overcoat layer is generally included in final prints with enhanced optical properties. The protective overcoat layer provides better refractive index matching with the underlying dye receiving layer. In addition to its use in dye diffusion thermal transfer systems, the layer may be used in applications such as, for example, a thermal transfer layer applied

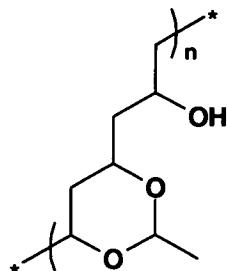
10 to an inkjet receiver.

The application of a protective overcoat layer or patch can eliminate concerns commonly encountered with dye diffusion thermal transfer prints. In addition, the protective overcoat layer or patch can also provide improved dye stability by acting as a barrier to UV light and pollutant gases such as ozone and nitrogen dioxide. If desired, the protective overcoat layer or patch may also provide the prints with a glossy surface, comparable to that obtained from glossy silver-halide photographic prints, with no iridescence. For example, the gloss, when measured at 20°, is at least 50 when transferred at a line time of 1 ms, and at least 45 when transferred at a line time of 0.5 ms. The protective overcoat layer desirably results in a smooth tear-off between transferred and non-transferred protective overcoat. For improvements in manufacturing, it is desirable that the materials utilized in the protective overcoat layer or patch enable use of low-cost solvents.

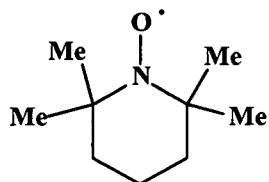
In particulate, the heat transferable donor element of this invention

25 is a protective overcoat layer patch (or protective material) on a thermal print provided by uniform application of heat using a thermal head. The protective overcoat layer, which may also be referred to as a protective overcoat or protective overcoat patch, can include at least one poly(vinyl acetal) resin of the following Formula I:

Formula I



wherein n is from 10 to 100. The average molecular weight can be in the range of from 4,000 to 100,000, for example from 15,000 to 80,000. Optionally, the 5 protective overcoat layer can also include at least one styrene/allyl alcohol copolymer resin, such as Lyondell SAA-100. As noted above, the protective overcoat layer also includes an N-oxyl radical light stabilizer, for example defined by the following Formula III, known as TEMPO:



10 In one embodiment of the invention, the protective overcoat layer is the only layer on the donor element and can be used in conjunction with a dye donor element that contains the heat transferable image dyes.

For example, in many embodiments, the heat transferable polymers used in the protective overcoat layer comprise a polyvinyl acetal resin blended 15 with a polystyrene/allyl alcohol resin. When used in a thermal transfer system, the refractive index of the poly(vinyl acetal) resin blended with a polystyrene/allyl alcohol resin can be closely matched to the refractive index of the dye receiving layer to alleviate the low gloss of current protective overcoat, resulting from the mismatch in refractive index between dye receiving layer and the protective 20 overcoat layer. Typically, refractive index is in the range of from 1.50 to 1.65, more typically in the range of from 1.54 to 1.65. Use of the poly(vinyl acetal) resin blended with a polystyrene/allyl alcohol resin gives gloss values 10-15 units higher than the gloss values achieved with a poly(vinyl acetal) protective

overcoat, as measured with a BYK- Gardner micro-TRI-gloss® meter (see experimental section).

In one embodiment, the protective overcoat layer may contain crosslinked elastomeric organic beads. The beads can have a glass transition 5 temperature (Tg) of 45°C or less, for example, 10°C or less. The elastomeric beads can be made from an acrylic polymer or copolymer, such as butyl-, ethyl-, propyl-, hexyl-, 2-ethylhexyl-, 2-chloroethyl-, 4-chlorobutyl- or 2-ethoxyethyl- acrylate or methacrylate; acrylic acid; methacrylic acid; hydroxyethyl acrylate; a styrenic copolymer, such as styrene-butadiene, styrene-acrylonitrile-butadiene, 10 styrene-isoprene, or hydrogenated styrene-butadiene; or mixtures thereof. The elastomeric beads can be crosslinked with various crosslinking agents, which can be part of the elastomeric copolymer, such as but not limited to divinylbenzene; ethylene glycol diacrylate; 1,4-cyclohexylene-bis(oxyethyl) dimethacrylate; 1,4-cyclohexylene-bis(oxypropyl) diacrylate; 1,4-cyclohexylene-bis(oxypropyl) 15 dimethacrylate; and ethylene glycol dimethacrylate. The elastomeric beads can have from 1 to 40%, for example, from 5 to 40%, by weight of a crosslinking agent. The elastomeric microbeads may be employed in any amount effective for the intended purpose. In general, good results have been obtained at a coverage of from 2 to 25 mg/m². The elastomeric microbeads generally have a particle size of 20 from 4 µm to 10 µm. At these levels, the beads are not detrimental to gloss, and are beneficial for finishing operations involving web-transport and spool winding.

The elastomeric beads may be crosslinked with various crosslinking agents, which may also be part of the elastomeric copolymer, such as divinylbenzene; ethylene glycol diacrylate; 1,4-cyclohexylene- bis(oxyethyl) 25 dimethacrylate; 1,4-cyclohexylene-bis(oxypropyl) diacrylate; 1,4-cyclohexylene-bis(oxypropyl) dimethacrylate; and ethylene glycol diacrylate.

The glass transition temperatures can be determined by the method of differential scanning calorimetry (DSC) at a scanning rate of 20°C/minute and the onset in the change in heat capacity was taken as the Tg.

30 Following are examples of typical elastomeric microbeads that may be employed in the invention:

Bead 1) poly(butyl acrylate-co-divinylbenzene) (80:20 mole ratio) having a nominal diameter of approximately 4 μm and a Tg of approximately -31°C.

Bead 2) poly(styrene-co-butyl acrylate-co-divinylbenzene) (40:40:20 mole ratio) having a nominal diameter of approximately 4 μm and a Tg of approximately 45°C.

Bead 3) poly(ethyl acrylate-co-ethylene glycol diacrylate) (90:10 mole ratio) having a nominal diameter of approximately 5 μm and a Tg of approximately -22°C.

10 Bead 4) poly(2-ethylhexyl acrylate-co-styrene-co- divinylbenzene) (45:40:15 mole ratio) having a nominal diameter of approximately 5 μm and a Tg of approximately 20°C.

Bead 5) poly[2-chloroethylacrylate-co-1,4-cyclohexylene- bis(oxypropyl) diacrylate] (80:20 mole ratio) having a nominal diameter of approximately 7 μm and a Tg of approximately -10°C.

15 Bead 6) poly(butyl methacrylate-co-hydroxyethyl-acrylate-co- divinylbenzene)(65:10:25 mole ratio) having a nominal diameter of approximately 6 μm and a Tg of approximately 29°C.

20 Bead 7) poly(styrene-co-butadiene-co-divinylbenzene)(40:50:10 mole ratio) having a nominal diameter of approximately 8 μm and a Tg of approximately -55°C.

Bead 8) poly(styrene-co-2-ethoxyethyl acrylate-co-ethylene glycol diacrylate)(20:45:35 mole ratio) having a nominal diameter of approximately 4 μm and a Tg of approximately -5°C.

25 Bead 9) poly(styrene-co-hexyl acrylate-co- divinylbenzene)(10:70:20 mole ratio) having a nominal diameter of approximately 4 μm and a Tg of approximately -15°C.

30 Non-heat transferable polymeric binders may also be present in the donor element but they are not transferred during thermal printing. Such polymeric binders are well known in the art and include but are not limited to, thermoplastic resins, for example, acrylic resins, such as poly(methyl methacrylate), poly(ethyl methacrylate), poly(butyl acrylate), vinyl resins, such as poly(vinyl acetate), vinyl chloride-vinyl acetate copolymer, poly(vinyl alcohol),

poly(vinyl butyral), and cellulose derivatives, such as ethyl cellulose, nitrocellulose, and cellulose acetate, and thermosetting resins, for example, unsaturated polyester resins, polyester resins, polyurethane resins, and aminoalkyd resins, in known amounts.

5 The heat transferable protective overcoat layer formulation may be formed by dissolving or dispersing the various resins, the light stabilizer, and optional beads in a suitable solvent, such as a mixture of toluene and n-butanol. An ultraviolet (UV) absorber may also be included in the formulation. While any known UV absorber may be used, the useful material is TINUVIN® 460 (Ciba).

10 While any known N-oxyl radical may be used, a useful compound is 1,1,5,5-tetramethylpentamethylene nitroxide (TEMPO).

Although not required for this invention, inorganic particles or organic beads other than the crosslinked elastomeric beads may be added.

15 The formulation is coated onto the support sheet, for example, by gravure printing, screen printing, or reverse coating using a gravure plate, and drying the coating. The formulation is generally applied to provide a dry coverage of at least 0.03 g/m² to 1.7 g/m² to obtain a dried layer of less than 1 µm. Thicker coatings can be applied if desired, for example in the 2-3 g/m² range.

20 In one embodiment of the invention, the protective overcoat layer contains, from 20% to 45% by weight poly(vinyl acetal) binder, typically from 35 to 45% from 20% to 50% by weight polystyrene/allyl alcohol polymeric binder, typically from 40 to 50% %, from 0.50 to 3.0%, typically from 1.0 to 2.0% by weight of the N-oxyl radical light stabilizer, from 0 to 30%, typically from 3 to 15% by weight of the UV absorbing compound, and from 0.5% to 4% crosslinked elastomeric beads, typically from 1.0 to 3.0%.

25 In practice, yellow, magenta and cyan dyes are thermally transferred from a dye donor element to form an image on the dye receiving element or sheet. The thermal head is then used to transfer a clear protective overcoat layer from a clear patch on the dye donor element or from a separate donor element, onto the dye imaged receiving sheet by uniform application of heat. The clear protective overcoat layer adheres to the print and is released from the donor support in the area where heat is applied.

An adhesive layer may be provided on the surface of the heat transferable protective overcoat layer to improve transferability and adhesion to the receiver surface. The adhesive layer may be formed of any conventional pressure-sensitive adhesive or heat sensitive adhesive having a glass transition 5 temperature (Tg) of from 40 to 80°C. To maintain high gloss and freedom from iridescence, material selection is determined by refractive index matching requirements.

The protective overcoat layer may be provided on the substrate sheet through a peel layer. The provision of the peel layer permits the overcoat 10 layer to be more easily transferred from the thermal transfer sheet onto the receiver. The peel layer may comprise, for example, waxes, such as microcrystalline wax, carnauba wax, paraffin wax, Fischer-Tropsh wax, various types of low-molecular weight polyethylene, Japan wax, beeswax, spermaceti, insect wax, wool wax, shellac wax, candelilla wax, petrolatum, partially 15 modified wax, fatty esters, and fatty amides, and thermoplastic resins, such as silicone wax, silicone resin, fluororesin, acrylic resin, polyester resin, polyurethane resin, cellulose resin, vinyl chloride-vinyl acetate copolymer, and nitrocellulose. Further, the peel layer may comprise a binder resin and a releasable material. Binder resins usable herein include thermoplastic resins, for 20 example, acrylic resins, such as poly(methyl methacrylate), poly(ethyl methacrylate), poly(butyl acrylate), vinyl resins, such as poly(vinyl acetate), vinyl chloride-vinyl acetate copolymer, poly(vinyl alcohol), poly(vinyl butyral), and cellulose derivatives, such as ethyl cellulose, nitrocellulose, and cellulose acetate, and thermosetting resins, for example, unsaturated polyester resins, polyester 25 resins, polyurethane resins, and aminoalkyd resins. Releasable materials include but are not limited to waxes, silicone wax, silicone resins, melamine resins, fluororesins, fine powders of talc or silica, and lubricants such as surfactants or metal soaps.

The peel layer may be formed by dissolving or dispersing the 30 materials in a suitable solvent to prepare a coating liquid for a peel layer, coating the coating liquid onto a substrate sheet by gravure printing, screen printing,

reverse coating using a gravure plate or other means, and drying the coating. The coverage is generally from 0.1 to 10 g/m² on a dry basis.

The donor element of the present invention may be used in sheet form or in a continuous roll or ribbon.

5 In some embodiments of the invention, the donor element comprises a poly(ethylene terephthalate) support coated with sequential repeating areas of yellow, cyan and magenta dye, and the inventive protection overcoat layer. The process steps are sequentially performed for each color to obtain a three-color dye transfer image with a protection layer on top.

10 The donor layer can include beads. The beads can have a particle size of from 0.5 to 20 μm , typically from 2.0 to 15 μm . The beads can act as spacer beads under the compression force of a wound up donor roll, improving raw stock keeping of the donor roll by reducing the material transferred from the donor layer to the slipping layer, as measured by the change in sensitometry under
15 accelerated aging conditions, or the appearance of unwanted dye in the protective overcoat layer, or from the backside of the donor element, for example, a slipping layer, to the donor layer. The use of the beads can result in reduced mottle and improved image quality. The beads can be employed in any amount effective for the intended purpose. In general, good results have been obtained at a coverage of
20 from 0.003 to 0.20 g/m². Beads suitable for the donor layer can also be used in the slip layer.

The beads in the donor layer can be crosslinked, elastomeric beads. The beads in the donor layer can be hard polymeric beads. Suitable beads can include divinylbenzene beads, beads of polystyrene crosslinked with at least 20
25 wt. % divinylbenzene, and beads of poly(methyl methacrylate) crosslinked with at least 20 wt. % divinylbenzene, ethylene glycol dimethacrylate, 1,4-cyclohexylene-bis(oxyethyl) dimethacrylate, 1,4-cyclohexylene-bis(oxypropyl) dimethacrylate, or other crosslinking monomers known to those familiar with the art.

30 Useful elastomeric microbeads have a lower Tg and are compressed under the weight of the thermal print head during printing, thereby allowing better contact between the donor and dye receiver elements. When microbeads having a high Tg are used, the microbeads are too rigid and prevent

intimate contact between the donor and dye receiver during printing, resulting in image mottle and poor image quality. The improved dye donor element/dye receiver element contact achievable with the low Tg elastomeric microbeads results in reduced mottle and improved image quality. The crosslinked

5 elastomeric beads employed in the invention have a Tg of 45°C or less, or typically 10° C or less.

The donor layer of the donor element can be formed or coated on a support. The donor layer composition can be dissolved in a solvent for coating purposes. The donor layer can be formed or coated on the support by techniques 10 such as, but not limited to, a gravure process, spin-coating, solvent-coating, extrusion coating, or other methods known to practitioners in the art.

The protective overcoat layer of the donor element may be coated on the support or printed thereon by a printing technique such as a gravure process.

15 A slipping layer may be used on the back side of the heat transferable donor element of the invention to prevent the printing head from sticking to the donor element. Such a slipping layer would comprise either a solid or liquid lubricating material or mixtures thereof, with or without a polymeric binder or a surface-active agent. Useful lubricating materials include oils or semi-crystalline organic solids that melt below 100°C such as poly(vinyl stearate), beeswax, perfluorinated alkyl ester polyethers, poly-caprolactone, silicone oil, poly(tetrafluoroethylene), carbowax, poly(ethylene glycols), or any of those materials disclosed in U.S. Patents 4,717,711; 4,717,712; 4,737,485; and 20 4,738,950. Suitable polymeric binders for the slipping layer include poly(vinyl alcohol-co-butyral), poly(vinyl alcohol-co-acetal), polystyrene, poly(vinyl acetate), cellulose acetate butyrate, cellulose acetate propionate, cellulose acetate, or ethyl cellulose.

25 For example, the slipping layer formulation most desired for resistive head thermal media incorporates a synergistic combination of lubricants from a friction perspective and in terms of headwear or print head buildup. This slip layer is disclosed in U.S. Patent 7,078,366. Primarily, it includes a donor element for thermal dye transfer comprising a support having on one side thereof

a heat transferable layer and on the other side a slipping layer comprising a material comprising a maleic anhydride polyethylene graft copolymer and at least one other hydrocarbon wax. There can be a protective overcoat layer on one side of the donor element and on the other side a slipping layer comprising a

5 lubricating material. The lubricating material comprises a solid polymer derived from a polyolefin and an ethylenically unsaturated carboxylic acid or ester or anhydride thereof; and at least one wax. The polymer may be an alpha-olefin maleic anhydride copolymer, a maleic anhydride polyethylene graft copolymer, a copolymer of an alpha-olefin and isopropyl maleate. The polyolefin is derived

10 from an alpha-olefin containing between two to eight carbon atoms, preferably where the alpha-olefin is ethylene and/or propylene. The ethylenically unsaturated carboxylic acids are those having between 3 to 12 carbon atoms. The ethylenically unsaturated carboxylic acid, ester or anhydride may be, for example, maleic acid, ethylmaleic acid, propylmaleic acid, isopropyl maleic acid, fumaric acid, methylenemalonic acid, glutaconic acid, itaconic acid, methylitaconic acid, mesacomic acid, citraconic acid, or a mixture thereof, as well as corresponding esters, anhydrides or mixtures of such acids, esters and anhydrides. The other wax can be an olefinic wax, a saturated hydrocarbon polymer, a linear low molecular weight polyethylene, a branched hydrocarbon with a number average molecular

15 weight of no more than 10,000 and a melting point or softening point of no more than 120°C, or a synthetic wax comprising a saturated or unsaturated hydrocarbon. The other wax may be selected from, for example, a mineral wax, a vegetable wax, an animal wax or a synthetic wax that is a saturated or unsaturated hydrocarbon polymer. The ratio of the first wax to the other wax is 5:1 to 1:10.

20 Typically, the slipping layer comprises at least three different waxes, the polymer derived from the polyolefin and the ethylenically unsaturated carboxylic acid or ester or anhydride thereof, a highly branched alpha-olefin polymer, and at least one other wax. This slipping layer formulation for resistive head thermal media incorporates a synergistic combination of lubricants from a friction perspective

25 and in terms of headwear buildup. Additional benefits include preventing or reducing folds, especially when used with relatively fast printers, for example at 4 milliseconds or less per line. A still further benefit is the prevention of retransfer

of dye from the dye donor during production. Finally, the slip layer is capable of being coated at high speed.

The amount of lubricating material used in the slip layer is dependent, at least in part, upon the type of lubricating material, but can be in the 5 range of from 0.001 to 2 g/m², although less or more lubricating material can be used as needed. If a polymeric binder is used, the lubricating material can be present in a range of from 0.1 to 50 weight %, typically from 0.5 to 40 weight %, of the polymeric binder. In one embodiment, the slipping layer comprises from 10 to 80 percent by weight of the polymer derived from a polyolefin and an 10 ethylenically unsaturated carboxylic acid or ester or anhydride thereof; from 10 to 80 percent by weight of the highly branched α -olefin polymer, and from 10 to 80 percent by weight of a substantially linear wax, based on the total weight of the three waxes.

Any binder may also be used in the slipping layer provided it will 15 be useful for the intended effect. In some embodiments, polymeric thermoplastic binders are employed. Examples of such materials include, for example, poly(styrene-co-acrylonitrile) (70/30 wt. ratio); poly(vinyl alcohol-co-butyral) (available commercially as Butvar[®] 76.RTM. by Monsanto Corp.); poly(vinyl alcohol-co-acetal); poly(vinyl alcohol-co-benzal); polystyrene; poly(vinyl acetate); cellulose acetate butyrate; cellulose acetate propionate; cellulose acetate; ethyl cellulose; cellulose triacetate; poly(methyl methacrylate); and copolymers 20 of methyl methacrylate. In another embodiment, the thermoplastic binder is cellulose acetate propionate or polyvinyl acetal.

The amount of the optional binder employed in the slipping layer 25 of the invention is not critical such as from 0.1 to 2 g/m².

The dye receiving element that is used with the donor element of the present invention usually comprises a support having thereon a dye image receiving layer. The support for the image receiving layer may be transparent or reflective. The support may be a transparent film such as a poly(ether sulfone), a 30 polyimide, a cellulose ester such as cellulose acetate, a poly(vinyl alcohol-co-acetal) or a poly(ethylene terephthalate). Opaque, reflective supports can include plain paper, coated paper, synthetic paper, photographic paper support, melt-

extrusion-coated paper, and laminated paper, such as biaxially oriented support laminates. Biaxially oriented support laminates suitable for use as receivers are described in U.S. Patents 5,853,965; 5,866,282; 5,874,205; 5,888,643; 5,888,681; 5,888,683; and 5,888,714. Biaxially oriented supports can include a paper base

5 and a biaxially oriented polyolefin sheet, for example, polypropylene, laminated to one or both sides of the paper base. The support can be a reflective paper, for example, baryta- coated paper, white polyester (polyester with white pigment incorporated therein), an ivory paper, a condenser paper, or a synthetic paper, for example, DuPont Tyvek® by E.I. DuPont de Nemours and Company (Wilmington, DE). The support can be employed at any desired thickness, for example, from 10 µm to 1000 µm. Exemplary supports for the dye image- receiving layer are disclosed in U.S. Patents 5,244, 861 and 5,928,990 and EP 671,281. Other suitable supports as known to practitioners in the art can also be used. According to various embodiments, the support can be a composite or laminate structure

10 comprising a base layer and one or more additional layers. The base layer can comprise more than one material, for example, a combination of one or more of a microvoided layer, a foamed layer, a layer with hollow particles, a nonvoided layer, a synthetic paper, a natural paper, and a polymer. The dye image-receiving layer may comprise, for example, a polycarbonate, a polyurethane, a polyester, poly(styrene-co-acrylonitrile), polycaprolactone, vinyl-series resins, such as halogenated polymers (for example, polyvinyl chloride and polyvinylidene chloride), poly(vinyl acetate), ethylene-vinyl acetate copolymer, vinyl chloride-vinyl acetate copolymer, or mixtures thereof. Latex polymers may be used in the dye image-receiving layer. The latex polymer may be a dispersion in which

15 hydrophobic polymers comprising a monomer unit of, for example, water-insoluble vinyl chloride dispersed as fine particles in a water-soluble dispersion medium. The dispersed state may be one in which polymer is emulsified in a dispersion medium, one in which polymer underwent emulsion polymerization, one in which polymer underwent micelle dispersion, one in which polymer

20 molecules partially have a hydrophilic structure, or the like. For such latex polymers it is desirable to prepare the dye image-receiving layer by applying an aqueous type coating solution and then drying it. Exemplary aqueous coating

25

formats are disclosed in U.S. Patent Application Publication 2008/0254241. The dye image-receiving layer may be present in any amount that is effective for the intended purpose. In general, good results have been obtained at a concentration of from 1 to 5 g/m².

5 Additional polymeric layers can be present between the support and the dye image-receiving layer. The additional layers can provide coloring, adhesion, antistatic properties, act as a dye-barrier, act as a dye mordant layer, or a combination thereof. For example, a polyolefin such as polyethylene or polypropylene can be present. White pigments such as titanium dioxide, zinc oxide, and the like can be added to the polymeric layer to provide reflectivity.

10 A subbing layer can be used over the polymeric layer in order to improve adhesion to the dye image-receiving layer. This can be called an adhesive or tie layer. Exemplary subbing layers are disclosed in U.S. Patents 4,748,150, 4,965,238, 4,965,239, and 4,965,241. An antistatic layer as known to 15 practitioners in the art can also be used in the receiver element. The receiver element can also include a backing layer. Suitable examples of backing layers include those disclosed in U.S. Patents 5,011,814 and 5,096,875.

20 The dye image-receiving layer, or an overcoat layer thereon, can contain a release agent, for example, a silicone or fluorine based compound, as is conventional in the art. Various exemplary release agents are disclosed, for example, in U.S. Patents 4,820,687 and 4,695,286.

 The receiver element can also include stick preventative agents, as described for the donor element. The receiver element and dye donor element can include the same stick preventative agent.

25 The dye image-receiving layer can be formed on the support by any method known to practitioners in the art, including but not limited to printing, solution coating, dip coating, and extrusion coating. When the dye image-receiving layer is extruded, the process can include (a) forming a melt comprising a thermoplastic material; (b) extruding or coextruding the melt as a single-layer 30 film or a layer of a composite (multilayer or laminate) film; and (c) applying the extruded film to the support for the receiver element. Exemplary extruded

receiving layer formats are disclosed in U.S. Patents 7,125,611, 7,091,157, 7,005,406, 6,893,592, and 6,897,183.

The donor element of this invention can also include a stick preventative agent to reduce or eliminate sticking between the donor element and the receiver element during printing. The stick preventative agent can be present in any layer of the donor element, so long as the stick preventative agent is capable of diffusing through the layers of the donor element to the heat transferable layer, or transferring from the slip layer to the heat transferable layer. For example, the stick preventative agent can be present in one or more patches of the heat transferable layer, in the support, in an adhesive layer, in a dye-barrier layer, in a slip layer, or in a combination thereof. According to various embodiments, the stick preventative agent can be in the slip layer, the heat transferable layer, or both. The stick preventative agent can be in one or more patches of that. If more than one dye patch is present in the heat transferable layer, the stick preventative agent can be present in the last patch of the layer to be printed, such as the cyan layer. However, the dye patches and protective overcoat patches can be in any order. The stick preventative agent can be a silicone- or siloxane-containing polymer. Suitable polymers can include graft copolymers, block polymers, copolymers, and polymer blends or mixtures. Suitable stick preventative agents are described, for example, in U.S. Patent 7,067,457.

Release agents as known to practitioners in the art can also be added to the dye donor element, for example, to the dye donor layer, the slip layer, or both. Suitable release agents can include, for example, those described in U.S. Patents 4,740,496 and 5,763,358.

Thermal printing heads, which can be used to transfer heat transferable materials from the donor elements of the invention, are available commercially. There can be employed, for example, a Fujitsu Thermal Head FTP-040 MCSOO1, a TDK Thermal Head LV5416 or a Rohm Thermal Head KE 2008-F3.

A thermal transfer assemblage of the invention comprises

- (a) a heat transferable donor element, and
- (b) a dye receiving element,

the dye receiving element being in a superposed relationship with the heat transferable donor element so that the heat transferable layer of that donor element is in contact with the dye image-receiving layer of the receiving element.

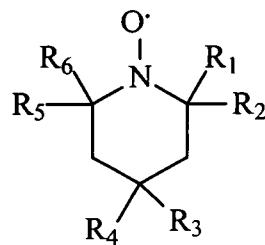
5 The assemblage comprising these two elements may be pre-assembled as an integral unit. This may be done by temporarily adhering the two elements together at their margins. After transfer, the dye receiving element is then peeled apart to reveal the dye transfer image.

When a three-color image is to be obtained, the assemblage is formed on three occasions during the time when heat is applied by the thermal 10 printing head. After the first dye is transferred, the elements are peeled apart. A second dye donor element (or another area of the donor element with a different dye area) is then brought in register with the dye receiving element and the process is repeated. The third color is obtained in the same manner. Finally, a protective overcoat layer is applied on top.

15 When a protective overcoat material is applied, it can be patterned to provide a matte or glossy finish by varying thickness, line time, print energy, or some combination thereof. Further, expandable or pre-expanded beads can be used in a laminate or protective overcoat layer to affect a gloss or matte finish depending on the amount and size of the beads. Overcoats, whether patterned or 20 not, can be provided on any colorant containing material, for example but not limited to, printed ink jet, thermal, or electrophotographic receivers, or silver halide prints.

The following represent at least some of the embodiments of this invention:

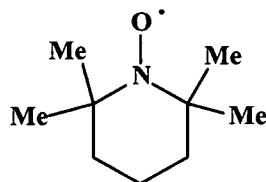
25 Embodiment 1: A heat transferable donor element comprising a polymeric support, the support having at least one portion thereof coated with a heat transferable material comprising a heat transferable polymeric binder and a light stabilizer that is an N-oxyl radical that is derived from a hindered amine, the N-oxyl radical having the following formula:



wherein R₁, R₂, R₅, and R₆ are each independently selected from a straight or branched C₁-C₆ alkyl or alkene, and R₃ and R₄ are each independently selected from H, OH, OR, COOH, or COOR, wherein R is a straight or branched C₁-C₆ alkyl or alkene, and having a molecular weight of 600 or less.

5 Embodiment 2: The element of embodiment 1 comprising at least one protective overcoat patch.

Embodiment 3: The element of embodiment 1 or 2 wherein the N-oxyl radical light stabilizer is:

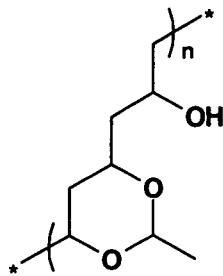


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Embodiment 4: The element of any of embodiments 1 to 3 wherein the heat transferable material further comprises a UV absorbing material in the amount of 20% by weight or less.

15 Embodiment 5: The element of any of embodiments 1 to 4 wherein the heat transferable material further comprises a plasticizer in the amount of 5% by weight or less.

Embodiment 6: The element of any of embodiments 1 to 5 wherein the heat transferable material further comprises at least one resin selected from Formula I, styrene/allyl alcohol copolymer, and the combination thereof, wherein Formula I
20 is



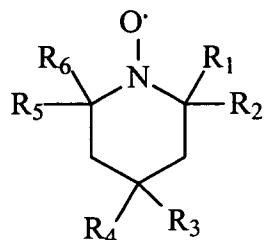
wherein n is from 10-100.

Embodiment 7: The element of embodiment 6 wherein the heat transferable material comprises from 40% to 90% by weight of the resin of

5 Formula I, and from 2 to 20% of a UV absorbing material.

Embodiment 8: The element of any of embodiments 1 to 7 further comprising an adhesive layer on the surface of the heat transferable material.

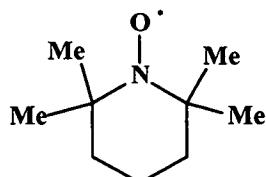
Embodiment 9: A heat transferable overcoat material comprising a heat transferable polymeric binder and an N-oxyl radical light stabilizer of the following formula:



wherein R₁, R₂, R₅, and R₆ are each independently selected from a straight or branched C₁-C₆ alkyl or alkene, and R₃ and R₄ are each independently selected from H, OH, OR, COOH, or COOR, wherein R is a straight or branched C₁-C₆

15 alkyl or alkene, and having a molecular weight of 600 or less.

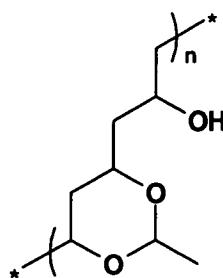
Embodiment 10: The overcoat material of embodiment 9 wherein the N-oxyl radical light stabilizer is:



Embodiment 11: The overcoat material of embodiment 9 or 10 further comprising a UV absorbing material in the amount of 20% or less.

Embodiment 12: The overcoat material of any embodiments 9 to 11 further comprising a plasticizer in the amount of 0 to 2%.

5 **Embodiment 13:** The overcoat material of any of embodiments 9 to 12 further comprising at least one resin selected from Formula I, styrene/allyl alcohol copolymer, and the combination thereof, wherein Formula I is:



wherein n is from 10-100.

10 **Embodiment 14:** The overcoat material of embodiment 13 wherein the heat transferable material comprises from 40% to 90% by weight of the resin of Formula I, and from 2 to 20% of a UV absorbing material.

Embodiment 15: A donor element comprising a polymeric support and the overcoat material of any of embodiments 9 to 12.

15 **Embodiment 16:** A method of coating a receiver material with a protective overcoat material, comprising:

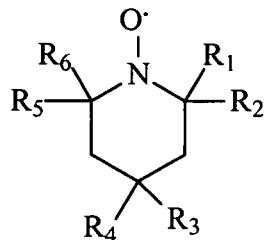
contacting the donor element of embodiment 15 with a receiver element;

20 applying heat or pressure sufficient to transfer the protective overcoat material from the donor element to the receiver element.

Embodiment 17: The method of embodiment 16 wherein the receiver element is selected from an inkjet receiver, a thermal receiver, an electrophotographic receiver, or a silver halide print.

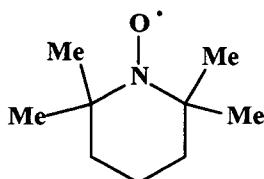
25 **Embodiment 18:** A thermal transfer assemblage comprising a receiver element in contact with at least a portion of a heat transferable donor element, wherein the donor element comprises a polymeric support at least one portion thereof coated with a heat transferable material comprising a heat transferable

polymeric binder and a light stabilizer that is an N-oxyl radical that is derived from a hindered amine, the N-oxyl radical having the following formula:



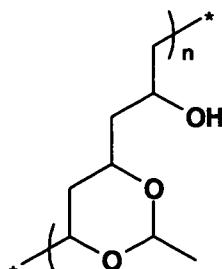
wherein R₁, R₂, R₅, and R₆ are each independently selected from a straight or branched C₁-C₆ alkyl or alkene, and R₃ and R₄ are each independently selected from H, OH, OR, COOH, or COOR, wherein R is a straight or branched C₁-C₆ alkyl or alkene, and having a molecular weight of 600 or less.

Embodiment 19: The assemblage of embodiment 18 wherein the light stabilizer in the donor element is:



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Embodiment 20: The assemblage of embodiment 18 or 19 wherein the heat transferable material of the donor element further comprises at least one resin selected from Formula I, styrene/allyl alcohol copolymer, and the combination thereof, wherein Formula I is



15

wherein n is from 10-100.

Embodiment 21: The assemblage of embodiment 20 wherein the heat transferable material of the donor element comprises from 40% 90% by weight of the resin of Formula I, and from 2 to 20% of a UV absorbing material.

Embodiment 22: The assemblage of any of embodiments 18 to 21 wherein the donor element comprises two or more patches of a heat transferable material, wherein at least one patch includes a dye and at least one patch comprises a protective overcoat material.

5 Embodiment 23: The assemblage of any of embodiments 18 to 22 wherein the receiver element is selected from an inkjet receiver, a thermal receiver, an electrophotographic receiver, or a silver halide print.

EXAMPLES

10 The following examples are provided to illustrate the invention.

Receiving Elements:

Thermal Receiver R-1 was used throughout these experiments, having an overall thickness of about 220 μm and a thermal dye receiving layer thickness of about 3 μm . R-1 was prepared by melt extruding the tie layer and 15 dye receiving layer onto the paper support, resulting in the following structure:

Co-extruded polyester-polycarbonate-silicone dye receiving layer
PELESTAT [®] 300 (Sanyo Chemical Industries, Ltd.) tie layer
Microvoided composite film OPPalyte [®] 350 K18 (ExxonMobil)
Pigmented polyethylene
Cellulose Paper
Polyethylene
Polypropylene film

KODAK Professional EKTATHERM ribbon, catalogue # 106-7347, was used in a KODAK Thermal Photo Printer, model number 6850, with receiver R-1 to 20 produce multiple, identical test target prints whose records were composed of neutral, monochrome, and bi-chromes consisting of two colors. Each record was arranged in a 15 step incremental density change from minimum density (Dmin) to maximum density (Dmax). The control or experimental protective overcoat patch on the donor ribbon was then transferred onto a test target print. The 25 protective overcoats were laminated with a transfer line-time of 0.8 ms.

Control Donor Element C-1

KODAK Professional EKTATHERM ribbon, catalogue # 106-7347, was used in a KODAK Thermal Photo Printer, model number 6850.

The protective overcoat of the donor elements were prepared by

5 coating on the back side of a 4.5 μm poly(ethylene terephthalate) support:

- 1) a subbing layer of titanium alkoxide, Tyzor TBT[®], (DuPont Corp.) (0.13 g/m²) from a n-propyl acetate and n-butyl alcohol solvent mixture (85/15), and
- 2) a slipping layer containing 0.02 g/m² Polywax[®] 400, 0.02 g/m²

10 Vybar[®] 103, 0.02 g/m² Ceramer 1608, all from Baker-Petrolite Corp., and 0.38 g/m² poly(vinyl acetal) binder KS-1 from Sekisui Co, coated from a 75/20/5 solvent mixture of toluene, methanol, and cyclopentanone.

15 A transferable protective overcoat layer of poly(vinyl acetal) KS-10 at a laydown of 0.63 g/m², IPA-ST (Nippon) at a laydown of 0.46 g/m², and UV Absorber TINUVIN[®] 460 (Ciba Specialties Co.), was coated on the front side of the donor element at a laydown of 0.11 g/m², and contained 4 μm poly(divinylbenzene) beads at a laydown of 0.03g/m². The materials were coated from solvent 3-pentanone.

20 **Invention Examples 1-20:**

Invention Example 1 was prepared as Control Donor Element C-1 except the front side of the element was prepared in the following manner:

To a magnetically stirred 16 oz. clear jar with threaded cap containing 226 g toluene and 25 g n-butanol was added 19.8 g polyvinyl acetal

25 KS-10 from Sekisui. The mixture was stirred at room temperature until a solution was obtained. Then, 22.5 g of styrene/allyl alcohol copolymer (Lyondell SAA-100) were added and the mixture was stirred at room temperature until a solution was obtained. Further, 4.37 g of Tinuin[®] 460 (Ciba) were added and the mixture stirred at room temperature until a solution was obtained. Additionally, 0.52 g of

30 TEMPO (Evonik/Degussa) was added and the mixture stirred at room temperature until a solution was obtained. Then, 1.30 g of four (4) μm poly(divinylbenzene) beads were added and the mixture was stirred for 24 hours. The resulting mixture

was coated on the front side of the donor element to give the TEMPO and Tinuvin® 460 levels shown in TABLE 1 below.

Invention Examples 2-6 were produced as in Invention Example 1 but in the absence of Tinuvin® 460, and with increasing amounts of TEMPO of 0 5 up to 0.0215 g/m² in increments of 0.0054 g/m².

Invention Examples 7-20 were produced as in Invention Example 1 but with Tinuvin® 460 in increasing amounts of 0.045, 0.090, and 0.180 g/m², and with increasing amounts of TEMPO (0 to 0.0215 g/m²) in increments of 0.0054 g/m².

10

Light Fade Test Method

Test target Status A densities were measured with an X-Rite Transmission/Reflection Densitometer model 820 from X-Rite Incorporated.

The test targets were subjected to 50 Klux high intensity daylight 15 using a xenon light source at room temperature. Test target dye densities were read at 1.0 and Delta density changes from start densities were calculated and reported as a Delta density. A lower absolute number indicates less change from the original sample, and therefore a better result (for example, -0.20 is better than -0.40, having less color change). TABLES 1-4 show the results of shifts in blue 20 and red at the end of a 28 day fade period. TABLES 5 and 6 show the results of shifts in blue and red at the end of a 21 day fade period.

TABLE 1

Donor Element	TEMPO (g/m²)	Tinuvin® 460 (g/m²)	Delta Blue of Red	Delta Red of Blue
Control C-1	0	0.0900	-0.40	-0.39
Invention 1	0.0108	0.0900	-0.26	-0.24
Invention 2	0	0	-0.47	-0.56
Invention 3	0.0054	0	-0.34	-0.43
Invention 4	0.0108	0	-0.29	-0.37
Invention 5	0.0162	0	-0.28	-0.35
Invention 6	0.0216	0	-0.27	-0.33

The data in TABLE 1 show the improvement in light fade by use of TEMPO as compared to Tinuvin® 460.

TABLE 2

Donor Element	TEMPO (g/m ²)	Tinuvin® 460 (g/m ²)	Delta Blue of Red	Delta Red of Blue
Control C-1	0	0.0900	-0.40	-0.39
Invention 7	0	0.0450	-0.40	-0.46
Invention 8	0.0054	0.0450	-0.29	-0.32
Invention 9	0.0108	0.0450	-0.27	-0.29
Invention 10	0.0162	0.0450	-0.27	-0.28
Invention 11	0.0216	0.0450	-0.25	-0.24

5

The data in TABLE 2 demonstrate how the addition of TEMPO to the protective overcoat allows for the coverage reduction of Tinuvin® 460 from 0.0900 g/m² to one half the coverage, 0.0450 g/m².

10

TABLE 3

Donor Element	TEMPO (g/m ²)	Tinuvin® 460 (g/m ²)	Delta Blue of Red	Delta Red of Blue
Control C-1	0	0.0900	-0.40	-0.39
Invention 12	0	0.0900	-0.37	-0.40
Invention 13	0.0054	0.0900	-0.28	-0.27
Invention 14	0.0108	0.0900	-0.26	-0.24
Invention 15	0.0162	0.0900	-0.25	-0.24
Invention 16	0.0216	0.0900	-0.24	-0.21

The data in TABLE 3 demonstrate the improvement in image stability achieved by the addition of TEMPO where the coverage of Tinuvin® 460 is equal to that in C-1.

TABLE 4

Donor Element	TEMPO (g/m ²)	Tinuvin [®] 460 (g/m ²)	Delta Blue of Red	Delta Red of Blue
Control C-1	0	0.0900	-0.40	-0.39
Invention 17	0	0.1800	-0.33	-0.34
Invention 18	0.0054	0.1800	-0.27	-0.22
Invention 19	0.0108	0.1800	-0.24	-0.19
Invention 20	0.0162	0.1800	-0.23	-0.17

The data in TABLE 4 demonstrate the image stability improvement resulting from increased solubility of Tinuvin[®] 460 in a solvent mixture.

5

Comparative Example 1:

The donor element of this Example was prepared as described for Invention Example 1, but substituting Tinuvin[®] 123 for TEMPO. The data in TABLE 5 demonstrate the importance of low molecular weight HALS. Tinuvin[®]

10 123, a commercially available hindered amine light stabilizer with the structure shown below, has a large molecular weight (737 MW) relative to TEMPO (156 MW). Also, Tinuvin[®] 123 exists as the alkyl oxy not the nitroxyl radical of TEMPO. Tinuvin[®] 123 is available from Ciba [bis(1-octyloxy-2,2,6,6-tetramethyl-4-piperidyl) sebacate] and has the following structure:

15

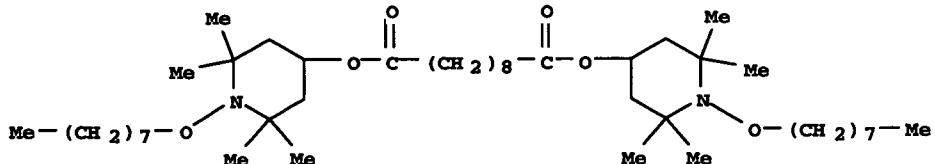


TABLE 5

Donor Element	Tinuvin [®] 123 (g/m ²)	Tinuvin [®] 460 (g/m ²)	Delta Blue of Red
Control C-1	0	0.0900	-0.51
Comparative Example 1	0.0108	0.0900	-0.53

Comparative Iridescence

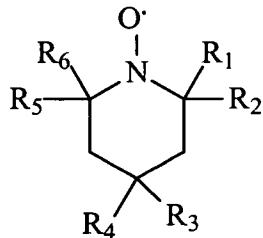
After lamination of the protective overcoat donor elements to the imaged receiver, samples were visually assessed for iridescence on a scale of 0 5 (none) to 5 (severe). TABLE 7 below shows data for samples laminated at 0.8 ms line-time.

TABLE 7

Donor Element		Iridescence
Control C-1		3
Invention 1		0
Invention 2		0
Invention 3		0
Invention 4		0
Invention 5		0
Invention 6		0
Invention 7		0
Invention 8		0

CLAIMS:

1. A heat transferable donor element comprising a polymeric support, the support having at least one portion thereof coated with a heat transferable material comprising a heat transferable polymeric binder and a light stabilizer that is an N-oxyl radical that is derived from a hindered amine, the N-oxyl radical having the following formula:

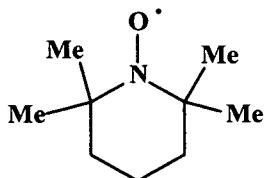


wherein R₁, R₂, R₅, and R₆ are each independently selected from a straight or branched C₁-C₆ alkyl or alkene, and R₃ and R₄ are each independently selected

10 from H, OH, OR, COOH, or COOR, wherein R is a straight or branched C₁-C₆ alkyl or alkene, and having a molecular weight of 600 or less.

2. The element of claim 1 comprising at least one protective overcoat patch.

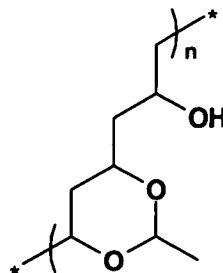
15 3. The element of claim 1 or 2 wherein the N-oxyl radical light stabilizer is:



4. The element of any of claims 1 to 3 wherein the heat transferable 20 material further comprises a UV absorbing material in the amount of 20% by weight or less.

5. The element of any of claims 1 to 4 wherein the heat transferable material further comprises a plasticizer in the amount of 5% by weight or less.

6. The element of any of claims 1 to 5 wherein the heat transferable material further comprises at least one resin selected from Formula I, styrene/allyl alcohol copolymer, and the combination thereof, wherein Formula I is

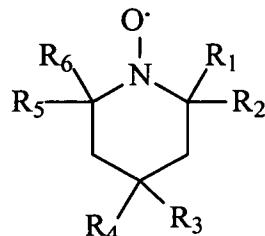


5 wherein n is from 10-100.

7. The element of claim 6 wherein the heat transferable material comprises from 40% to 90% by weight of the resin of Formula I, and from 2 to 20% of a UV absorbing material.

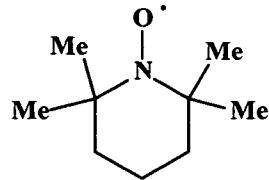
10

8. A heat transferable overcoat material comprising a heat transferable polymeric binder and a light stabilizer that is an N-oxyl radical that is derived from a hindered amine, the N-oxyl radical having the following formula:



15 wherein R₁, R₂, R₅, and R₆ are each independently selected from a straight or branched C₁-C₆ alkyl or alkene, and R₃ and R₄ are each independently selected from H, OH, OR, COOH, or COOR, wherein R is a straight or branched C₁-C₆ alkyl or alkene, and having a molecular weight of 600 or less.

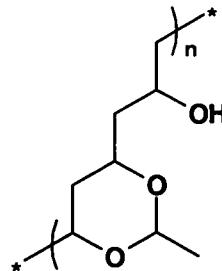
9. The overcoat material of claim 9 wherein the N-oxyl radical light stabilizer is:



5 10. The overcoat material of claim 8 or 9 further comprising a UV absorbing material in the amount of 20% or less.

11. The overcoat material of any of claims 8 to 10 further comprising a plasticizer in the amount of 0 to 2%.

10 12. The overcoat material of any of claims 8 to 11 further comprising at least one resin selected from Formula I, styrene/allyl alcohol copolymer, and the combination thereof, wherein Formula I is:



15 wherein n is from 10-100.

13. A method of coating a receiver element with a protective overcoat material, comprising:

20 contacting the donor element of any of claims 1 to 7 with a receiver element;

applying heat or pressure sufficient to transfer the protective overcoat material from the donor element to the receiver element.

14. A thermal transfer assemblage comprising a receiver element in contact with at least a portion of the donor element of any of Claims 1 to 7.
- 5 15. The assemblage of claim 14 wherein the donor element comprises two or more patches of a heat transferable material, wherein at least one patch includes a dye and at least one patch includes a protective overcoat material.

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2010/000465

A. CLASSIFICATION OF SUBJECT MATTER INV. B41M7/00 B41M5/42 B41M5/392 ADD.		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) B41M		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 2005/064320 A1 (SIMPSON WILLIAM H [US] ET AL) 24 March 2005 (2005-03-24) paragraphs [0008], [0021] – [0031], [0051], [0065] – [0069], [0081] – [0085] -----	1-15
A	US 5 658 850 A (TANIGUCHI KEISHI [JP] ET AL) 19 August 1997 (1997-08-19) column 8, lines 12-25 column 9, lines 19-24 -----	1-15
A	US 4 756 963 A (YAMAMOTO MAYUMI [JP] ET AL) 12 July 1988 (1988-07-12) column 3, lines 38-50 column 4, line 59 – column 5, line 34 -----	1-15
A	US 2002/158960 A1 (KASPERCHIK VLADEK P [US] ET AL) 31 October 2002 (2002-10-31) paragraphs [0032] – [0034], [0037] ----- -/-	1-15
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C.		<input checked="" type="checkbox"/> See patent family annex.
<p>* Special categories of cited documents :</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>"L" 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 when the document is taken alone</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>		
Date of the actual completion of the international search	Date of mailing of the international search report	
11 June 2010	14/07/2010	
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Patosuo, Susanna	

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2010/000465

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 6 942 956 B2 (SIMPSON WILLIAM H [US] ET AL) 13 September 2005 (2005-09-13) cited in the application column 3, lines 12-24 column 6, line 60 - column 7, lines 25, 43 -----	1-15

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2010/000465

Patent document cited in search report	Publication date	Patent family member(s)			Publication date
US 2005064320	A1	24-03-2005	JP WO	2007506582 T 2005032842 A1	22-03-2007 14-04-2005
US 5658850	A	19-08-1997	NONE		
US 4756963	A	12-07-1988	JP JP JP	1928846 C 6053436 B 62059076 A	12-05-1995 20-07-1994 14-03-1987
US 2002158960	A1	31-10-2002	US	2004026020 A1	12-02-2004
US 6942956	B2	13-09-2005	EP US WO	1680281 A1 2005064319 A1 2005032841 A1	19-07-2006 24-03-2005 14-04-2005