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(54) Title: CHROMOGENIC BLACK-AND-WHITE PHOTOGRAPHIC IMAGING SYSTEMS

(57) Abstract

The invention is accomplished by forming balanced cyan, magenta, and yellow coupler and emulsion mixes. There is at least one layer in which silver halide emulsion has been sensitized to blue light or silver halide emulsion sensitive to green light. Regardless of the color sensitivity of the silver halide layer that contains silver contains a mix of cyan, magenta and yellow dye-forming couplers. Further, in order to have a black-and-white image that has a lightness such as observed by the human eye in a scene, it is preferred that the ratios of red sensitive emulsion to green sensitive emulsion to blue sensitive emulsion in the photographic element is about 2:3:1.

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Title of the Invention

CHROMOGENIC BLACK-AND-WHITE PHOTOGRAPHIC IMAGING SYSTEMS

Cross-Reference to Related Application

This is a continuation-in-part of our earlier filed application U.S. Serial No. 810,311 filed on December 19, 1991.

Field of the Invention

This invention relates to the formulation of a photographic system which produces black-and-white images using a combination of cyan, magenta, and yellow dyes. The dyes are formed during a color development step from a mixture of cyan, magenta, and yellow dye forming couplers which, when allowed to react with oxidized color developer, form a neutral image.

Background of the Invention

Black-and-white images formed in a photographic process are generally produced by developing silver halide in a black-and-white developer to form a silver image. A black-and-white developer, such as hydroquinone, is commonly used to reduce the exposed silver halide to silver metal. The undeveloped silver halide is removed from the print by 'fixing' with aqueous sodium thiosulfate. The silver metal remaining in the print represents the image.

In the photographic industry, a photofinisher who wishes to produce both black-and-white and color pictures or prints must have separate processing systems; one for color and one for black-and-white, as the two systems are not compatible. It would, therefore, be advantageous for the photofinisher to have one process capable of producing either black-and-white or color materials.

U.S. 4,348,474 - Scheerer discloses a system wherein black-and-white images are formed by the use of one emulsion that is treated with three sensitizing dyes.

U.S. 2,186,736 - Schneider discloses the use of several color components in one layer for a black-and-white image formation.

U.S. 2,592,514 - Harsh discloses a color film in which couplers forming more than one color are present in the same layer of the color film.

There have been commercialized products that have formed black-and-white images by the use of pan sensitized emulsions which contain three spectral sensitizing dyes, color dye forming couplers and one emulsion. These pan-sensitive emulsions are sometimes coated in a fast and a slow layer to form images after exposure and development of the couplers. While the above products are somewhat successful, they do not achieve a neutral image. Additionally, the tone reproduction of such materials is severely limited by the contrast range of the emulsion.

Problem to be Solved by the Invention

There is a need for high quality black-and-white photographic products that are suitable for development in color imaging systems. Further, there is a desire that silver be recovered from the photographic print used to form the black-and-white images rather than being a part of the image and, therefore, not recoverable.

Summary of the Invention

It is an object of the invention to provide black-and-white images that are developable in color processes.

It is an object of the invention to provide black-and-white images of improved tone scale by the use of color couplers and conventional color processing.

The invention is generally accomplished by providing a photographic element for forming neutral images comprising at least one layer of balanced cyan, magenta, and yellow dye-forming couplers with silver halide grains comprising at least one of blue sensitized silver halide grains and green sensitized silver halide grains.

In a preferred embodiment, there is formed a photographic element comprising at least one layer of balanced cyan, magenta, and yellow dye-forming couplers with blue sensitized silver halide grains, and at least one layer of balanced cyan, magenta, and yellow dye-forming couplers with green sensitized silver halide grains.

In another preferred embodiment, there is formed a photographic element comprising at least one layer of balanced cyan, magenta, and yellow dye-forming couplers and blue sensitized silver halide grains, and green sensitized silver halide grains.

There is also provided a method of forming neutral black-and-white images by developing the above elements of the invention.

In a preferred embodiment, the invention is generally accomplished by forming multilayers consisting of mixtures of coupler dispersions and emulsions. In such a multilayer, there are at least two layers such that there is at least one layer in which silver halide emulsion has been sensitized to blue light, one silver halide emulsion layer sensitive to red light, and one silver halide layer emulsion sensitive to green light. Regardless of the color sensitivity of the silver halide, each emulsion containing layer also contains a mixture of cyan, magenta, and yellow dye-forming couplers.

Further, in order to have a black-and-white image that reproduces the lightness ratios in a scene such as observed by the human eye, it is preferred that the contrast ratios of red sensitive emulsion to green

sensitive emulsion to blue sensitive emulsion in the photographic element be about 2:3:1. These contrast ratios have been found to be similar to the relative response of the eye to color lightness changes. This contrast ratio can, however, be changed to meet customer preferences, as well as to increase the range of contrasts that can be effected when black and white or color negatives varying in contrast are to be printed. In like manner, the overall contrast and tone reproduction can be customized by adjusting the silver halide and color coupler coverages such as is typically done with present color and black and white photographic film and paper systems.

Advantageous Effect of the Invention

The invention has numerous advantages over the prior art. By the formation of an accurate black-and-white reproduction of a color exposure, the photographic products of the invention eliminate the need for a separate processing system in order to form black-and-white photographs. The black-and-white photographic system of the invention further allows the recovery of substantially all of the silver from a black-and-white photographic image. Another advantage of the system is that the reproduction of lightness ratios and tone is more accurate than any other system using color couplers to form black-and-white images. Another advantage is that if the lightness and tone of the black-and-white image are desired to be changed, this can be accomplished by the use of conventional color filters during printing of the negative. A photographic print formed in accordance with the invention will respond to changes in filtration of colored light during printing in a manner that allows ready adjustment of tone and lightness. This advantage is not available in other black-and-white photographic systems where pan or ortho sensitive emulsion systems are employed.

Brief Description of the Drawings

Figure 1 shows the lightness reproduction vs. chroma of 30 different standard colors and neutral densities when Kodak Panalure M™ black and white paper is compared to Kodak Supra™ color paper when printed with the identical negative. In each case, the tail of the arrow represents the lightness and chroma values obtained with the color paper, while the head of the arrow and the length of the arrow illustrates the change in chroma and lightness when the negative is printed onto Kodak Panalure M black and white paper.

Figure 2 illustrates the comparison when Kodak Polycontrast III™ black and white paper is used in place of the Kodak Panalure M paper.

Figure 3 illustrates the changes in lightness obtained when a black and white printing paper such as that described in this invention is used in place of the Kodak Panalure M paper.

Detailed Description of the Invention

To make a black and white image using a mixture of dyes formed from couplers, it is necessary to balance the ratio of the couplers in the imaging layer so that after exposure and color development, the resultant image is neutral and lacks any specific color bias. There may, however, be photographic market requirements whereby the color of the desired reproduction may not be neutral. For example, to accurately reproduce the tone of a 'sepia-toned' print, it would be necessary to alter the ratios of the couplers in the dispersion or the ratios of the dispersions in the emulsion layer in such a way that the preferred "sepia-toned" color balance is obtained. This process can be easily done using simple mathematical models. Also, many 'black and white papers' based upon silver halide systems which are presently in the market place are known not to produce a neutral image.

Depending upon the formulation of the silver halide material and the nature of the black and white development process, a wide variety of shades of green, red, yellow or brown can be produced. Each having its own unique characteristic color and photographic application.

The phrase "balanced cyan, magenta, and yellow dye-forming couplers" means that the couplers are balanced to provide a generally neutral image. This neutral balanced image would preferably for most uses be black and white. It is also possible in accordance with the invention technique to balance to give a sepia tone or slightly bluish tone to the image but still have a generally neutral image.

The invention utilizes an oil-in-water dispersion containing a mixture of cyan, magenta, and yellow dye-forming couplers. It should also be obvious that separate dispersions containing cyan, magenta and yellow dye forming couplers could be used. In addition, other dispersion addenda such as coupler solvent, auxiliary coupler solvent and/or dye stabilizers can be added. Dispersion addenda such as latex polymers or hydrophobic polymers may also be added. The aqueous phase of the dispersion is composed of gelatin, a surfactant, and water. The composition of the oil phase portion of the dispersion is adjusted so that when processed in a color developing bath, a neutral image is formed whose density varies only in proportion to the amount of silver developed in the process. In the instance where separate coupler dispersions are used, the appropriate ratios of each dispersion are added to the layer so that after exposure and development a neutral image is formed.

Once prepared, the coupler dispersion may in one embodiment be coated in a multilayer format much like that used in conventional color film or paper. There are two major differences, however; the first difference is

that the same neutral dye-forming coupler dispersion is coated in each emulsion containing layer. Thus, regardless of whether the element is exposed to red, green, or blue light, a neutral image is formed during color development in proportion to the amount of silver development. After color development, the developed and undeveloped silver are removed from the element by bleaching and fixing, or more simply, blixing (bleach-fixing).

The second difference is that the ratios of sensitized silver halide in the element are adjusted so that the lightness of the object being reproduced in the original scene is more accurately reproduced. This effect is obtained by coating the spectrally sensitized silver halide layers in amounts which correspond to the eye's relative sensitivity to light. It is generally agreed that the eye's response the red, green and blue light is in the ratio of about 2:3:1. Higher numbers indicate greater sensitivity. Therefore, in the invention element, the ratios of the amount of red sensitive emulsion to the amount of green sensitive emulsion to the amount of blue sensitive emulsion is also preferred to be about 2:3:1. However, this ratio can be adjusted to any ratio depending upon the needs and requirements of the photographic system. For example, films designed for X-ray applications which are currently coated on a blue support might choose to enhance the visual process of contrast discrimination by using ratios of 2:2:1 or 3:2:1.

In the second embodiment of the invention, the oil in water dispersion containing a mixture of cyan, magenta, and yellow dye-forming couplers is coated in a layer that contains silver halide grains sensitized to more than one color. In this embodiment the silver halide grains are a mixture of grains sensitized to be sensitive to different light colors. Preferably, the silver halide emulsion contains blue sensitized, green

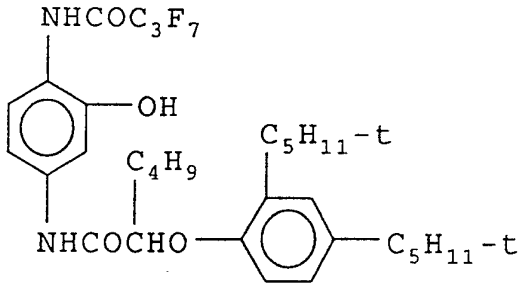
sensitized, and red sensitized silver halide grains. An element may only contain the blue sensitized and green sensitized silver halide grains to form an ortho sensitized element.

The invention may be performed with the materials conventionally utilized in color papers. As known, such papers comprise couplers for forming yellow, cyan, and magenta dyes. It is most common to use predominantly silver chloride emulsions with color paper, as they are suitable for fast processing. It should be apparent that other photographic systems may require the use of emulsions other than silver chloride. Such systems may in fact require silver chlorobromide, silver bromide, silver bromoiodide or silver chlorobromoiodide. The emulsions are sensitized to light in the wavelength to be absorbed by the particular layer where they are present. For instance, silver halide grains in the yellow layer will be most sensitive to blue light, and silver halide grains in the magenta layer will be most sensitive to green light. The use of sensitizing dyes to provide such emulsions is well known. Reference is made to *Research Disclosure* #308119, published December, 1989 for a description of emulsion formation, sensitizing dyes, antifoggant and stabilizers, couplers, hardeners, coating aids, and other conventional materials for use in silver halide image formation. The invention is considered to be able to be practiced with any of the known materials for use in color silver halide photography. Further, it is anticipated that the technique will be satisfactory for use with future materials using silver halide and dye-forming couplers that form yellow, cyan, and magenta dyes.

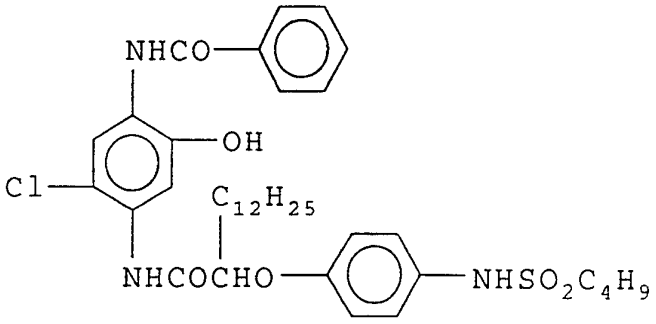
The following is a listing of preferred cyan, magenta, and yellow couplers for the invention. Following the couplers are the structures of three preferred hydroquinones for the invention.

Cyan Couplers

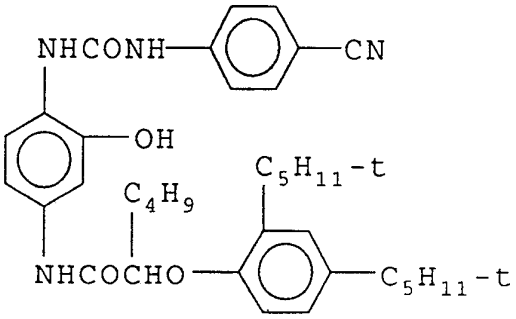
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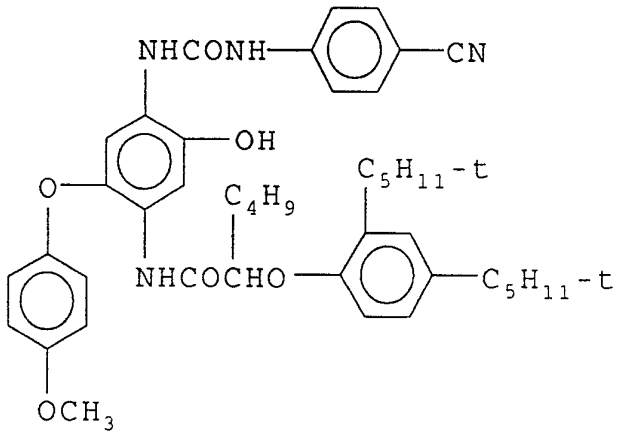
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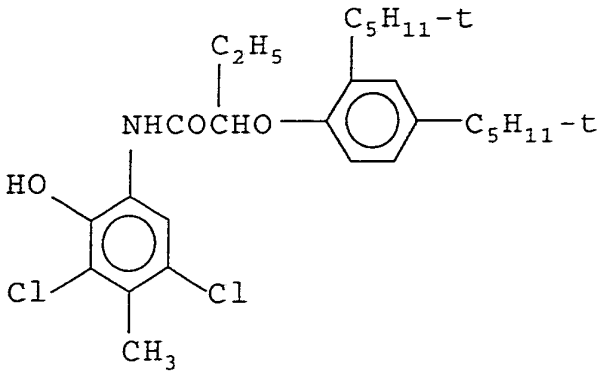
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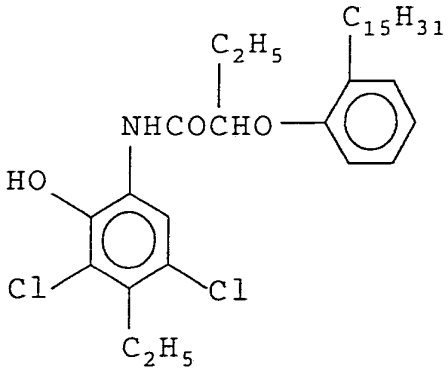
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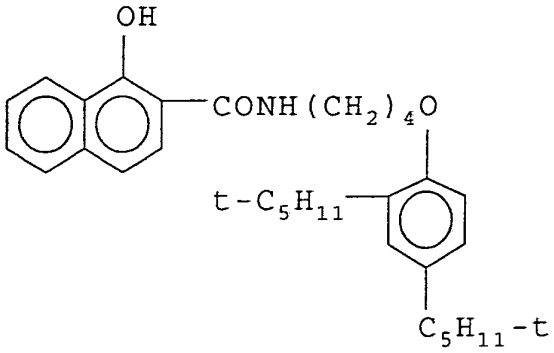
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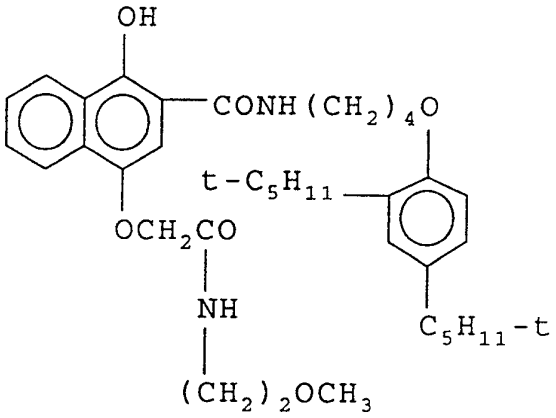
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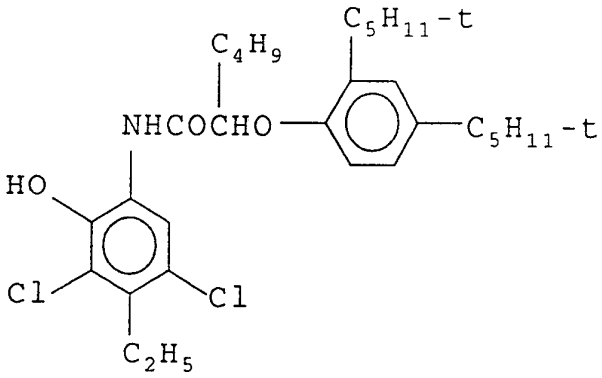
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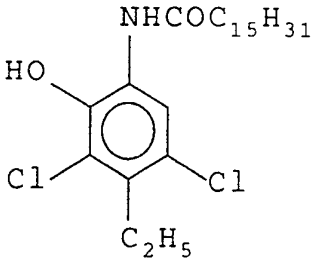
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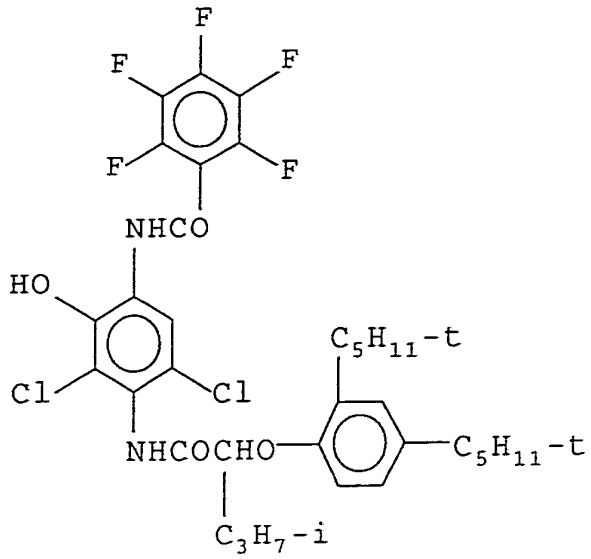
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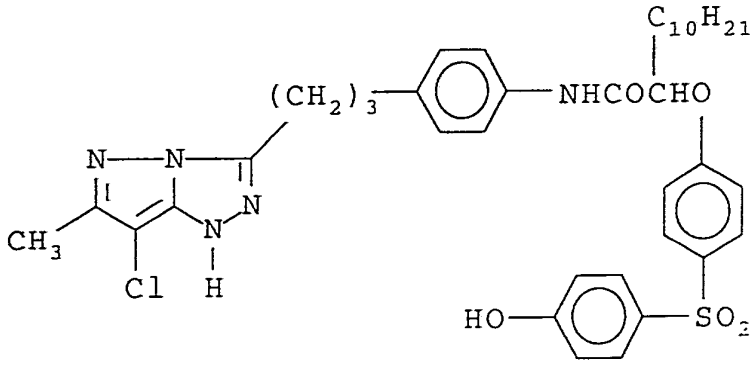


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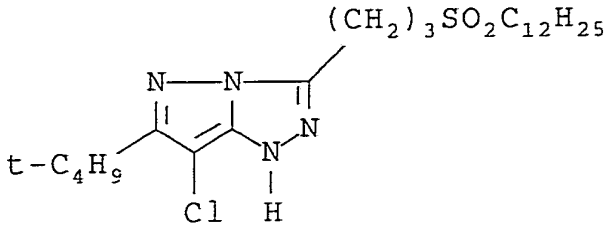


Magenta Couplers

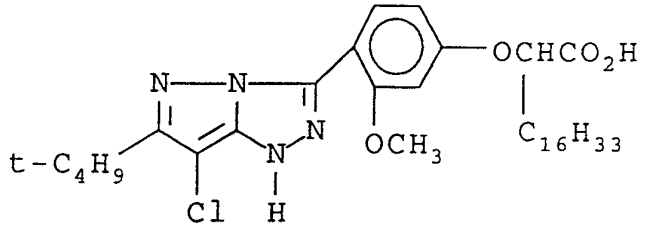
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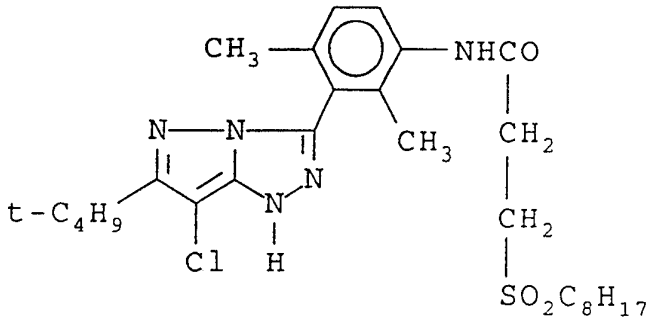
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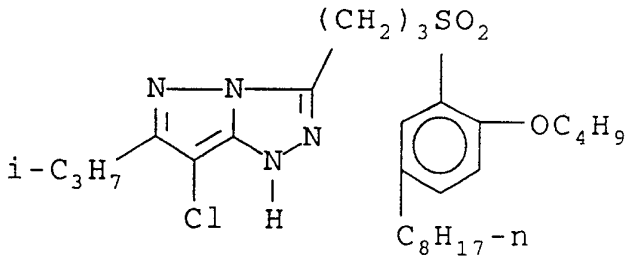
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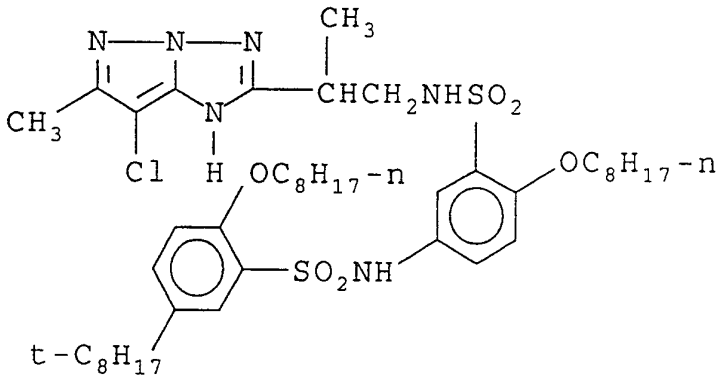
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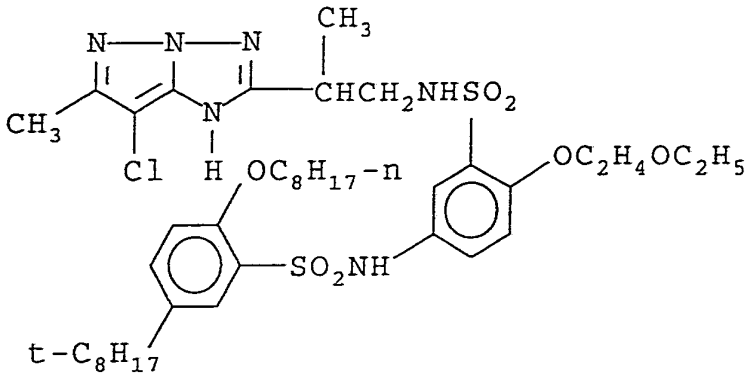
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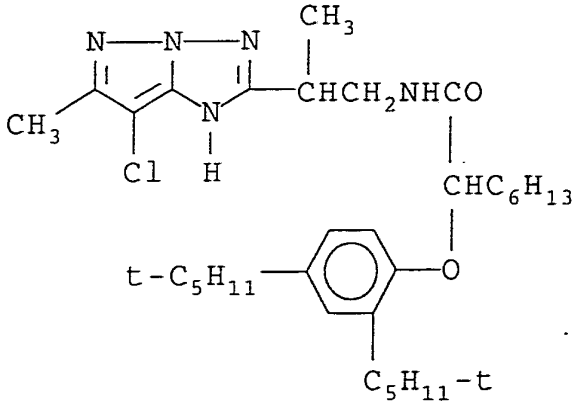
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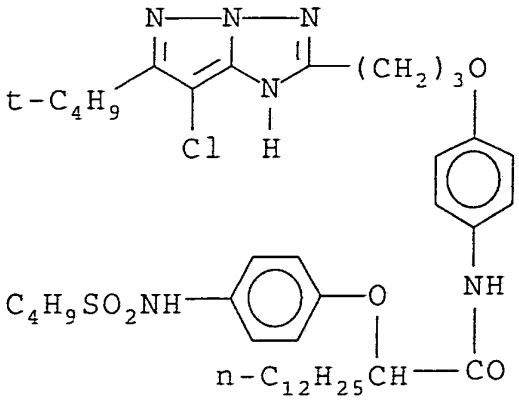
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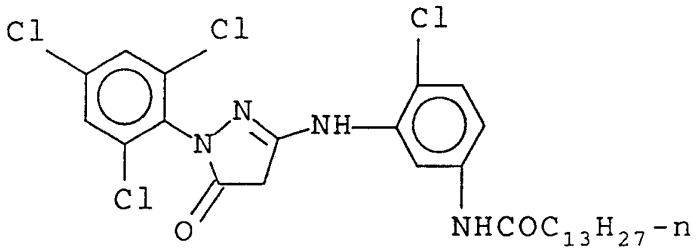
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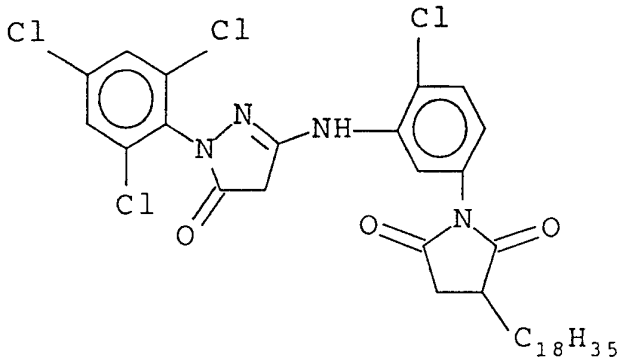
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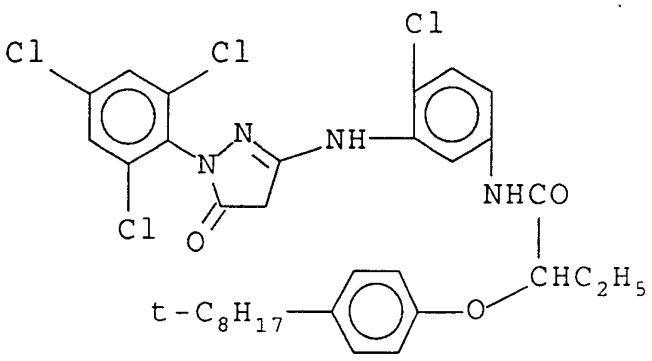

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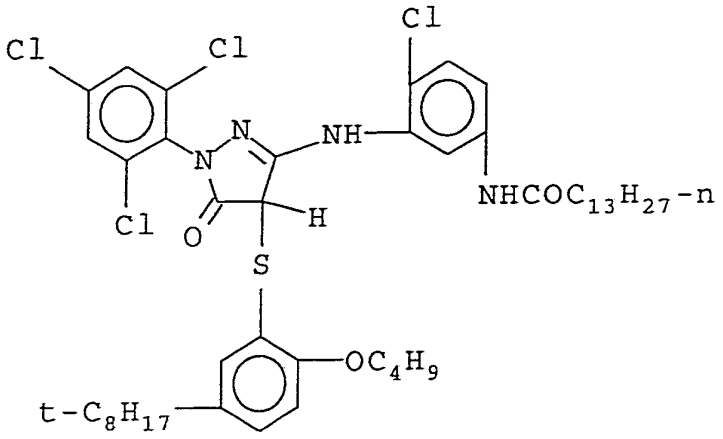

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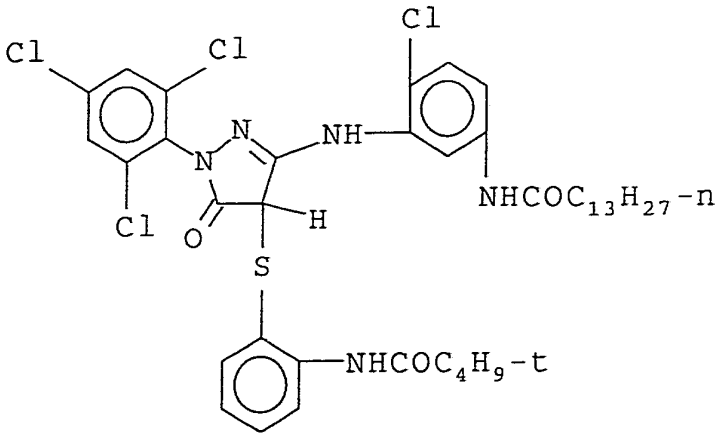
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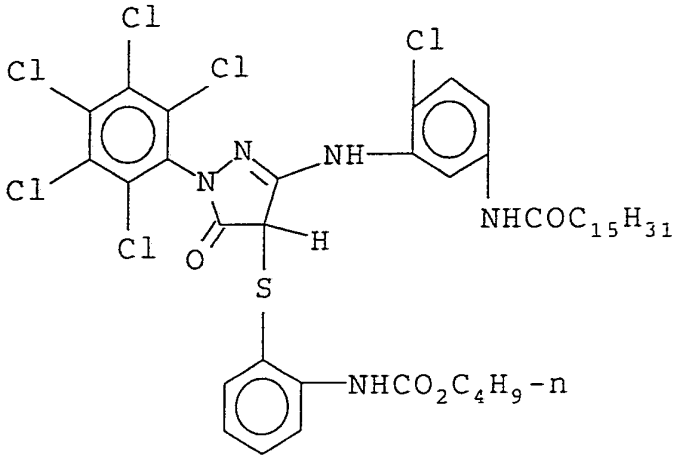
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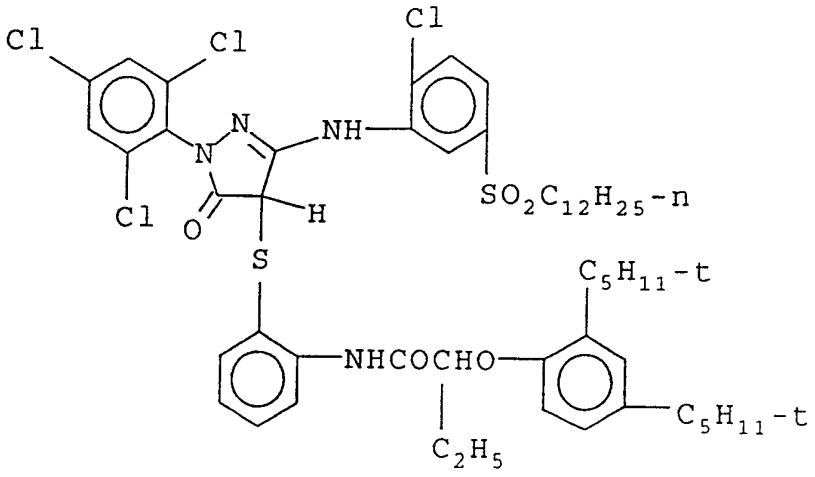
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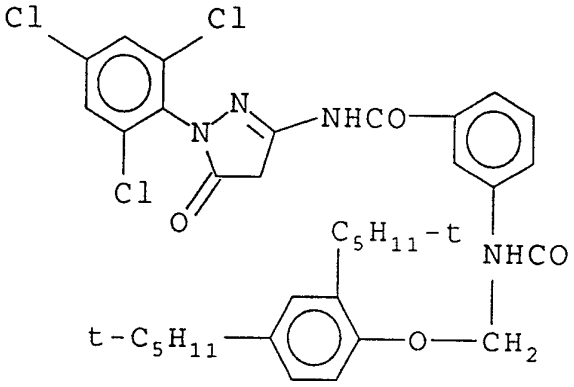
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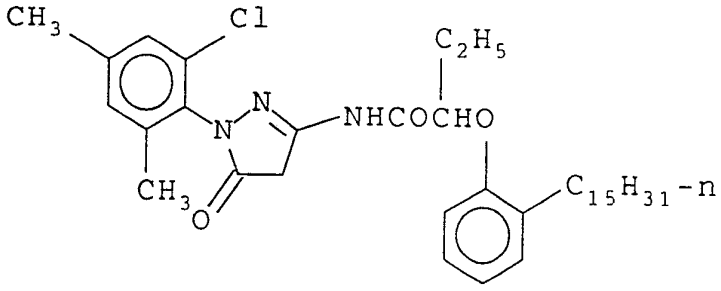
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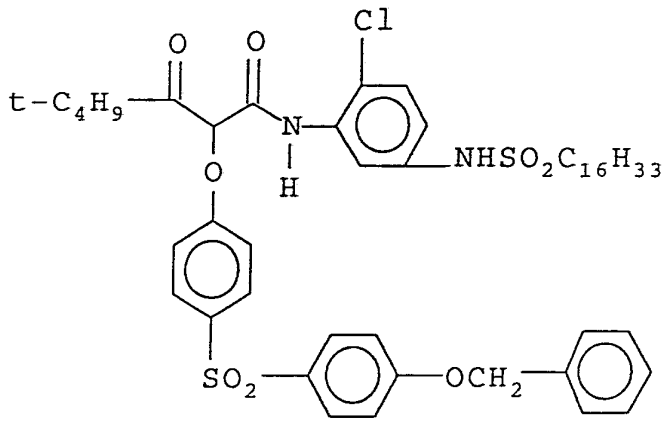


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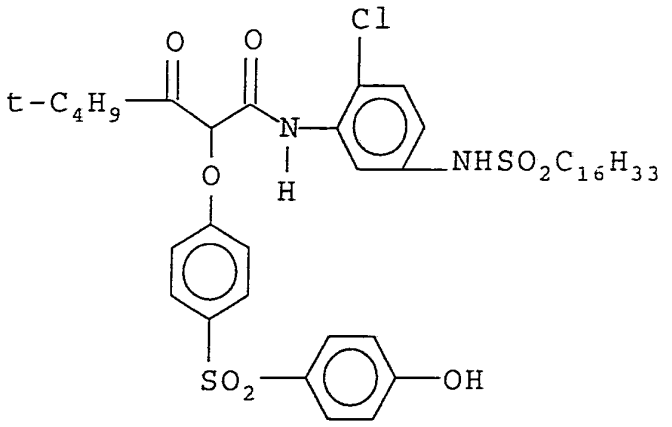


Yellow Couplers

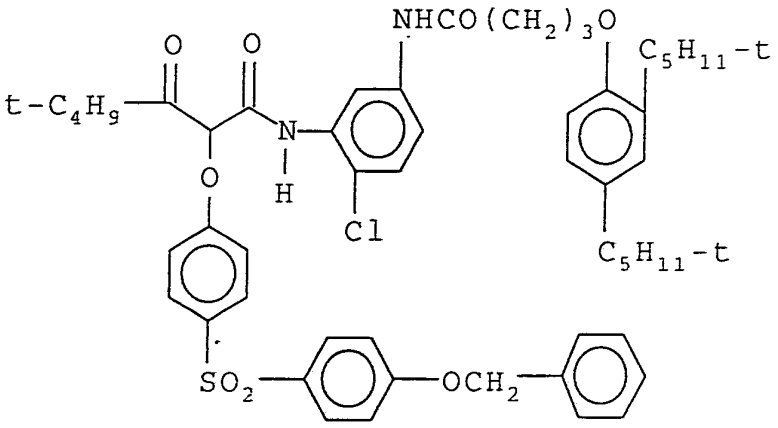
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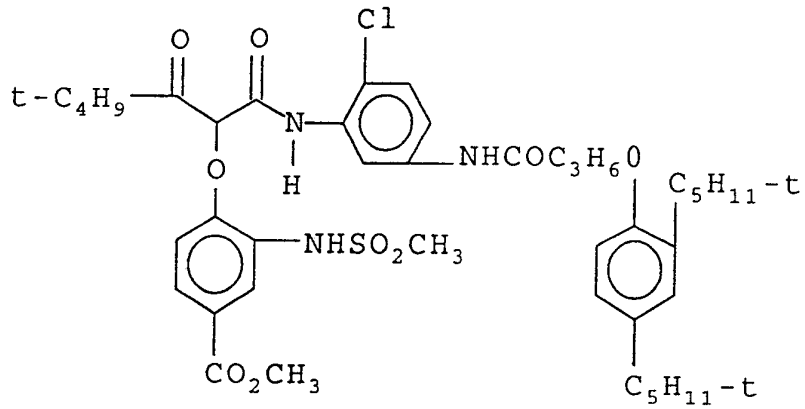
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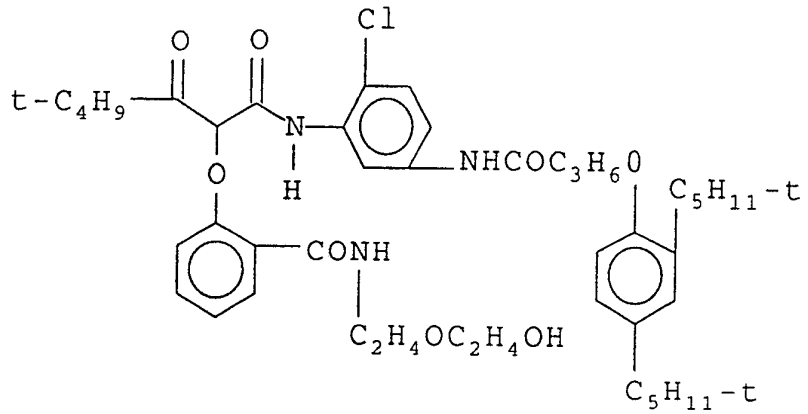
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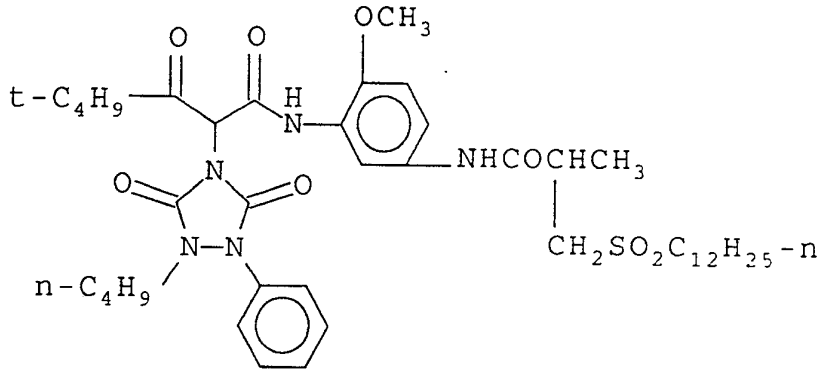
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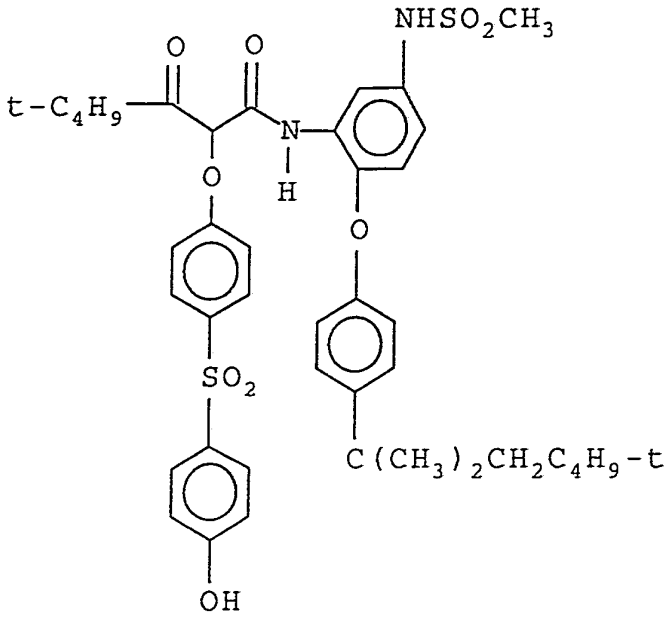
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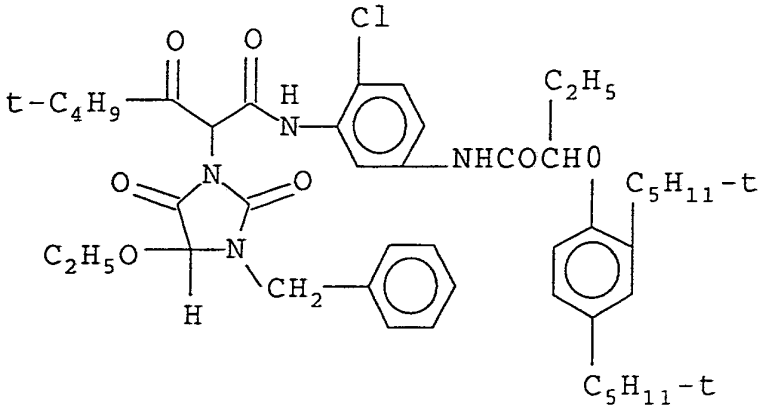
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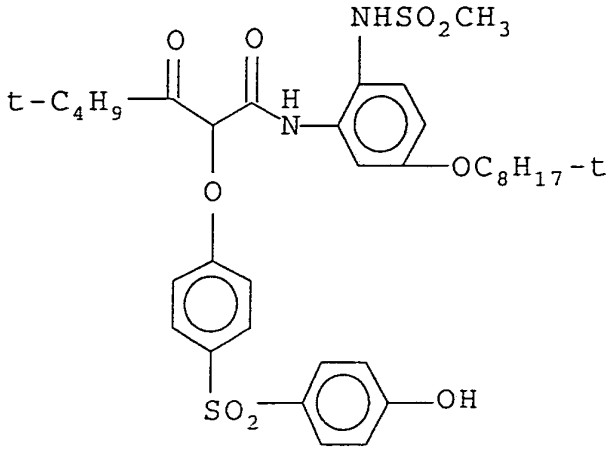
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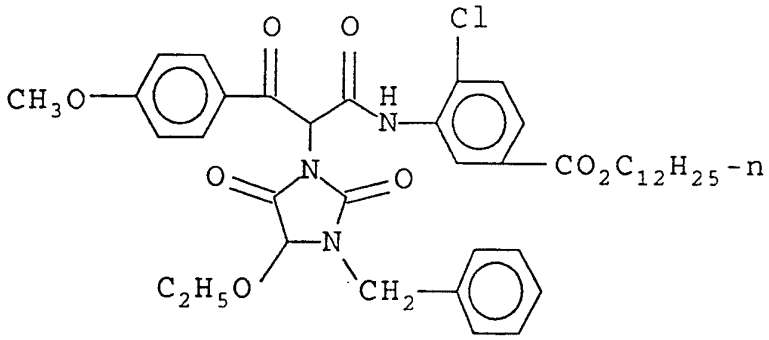
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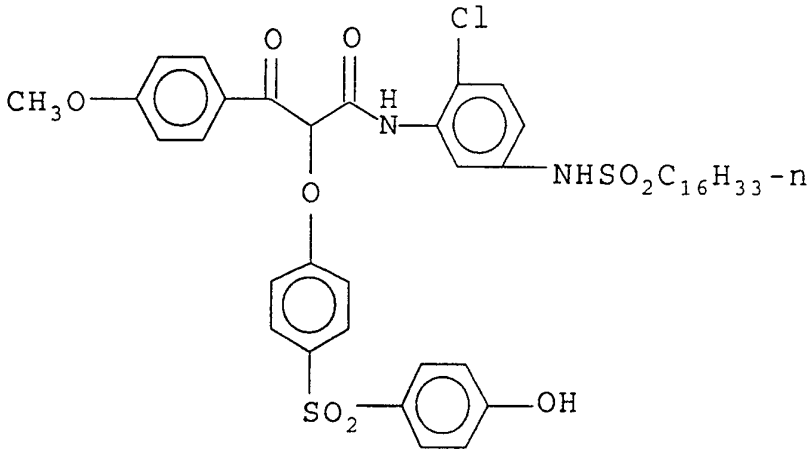
C10



C11



C12

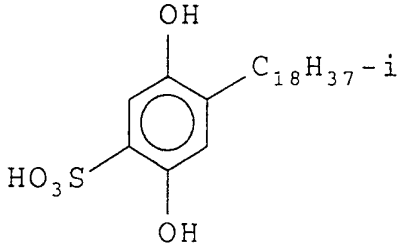


Hydroquinones

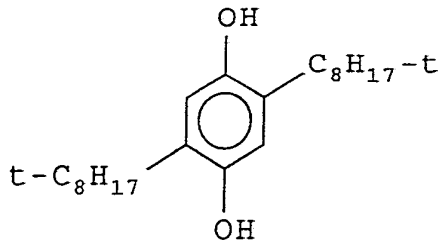
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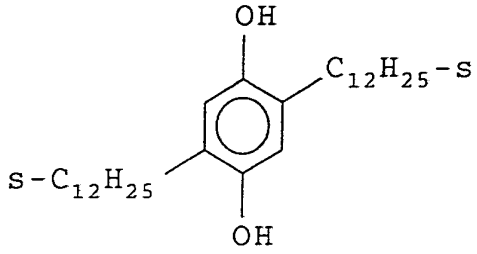
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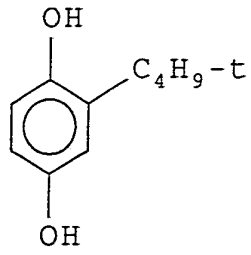
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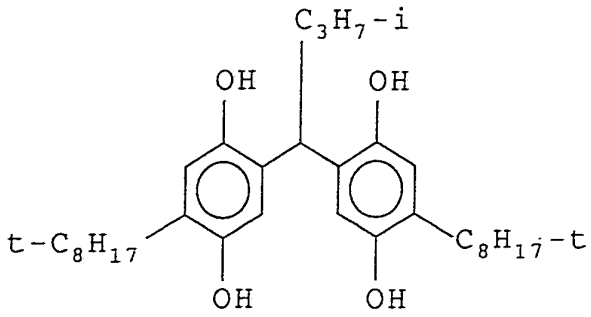
H4



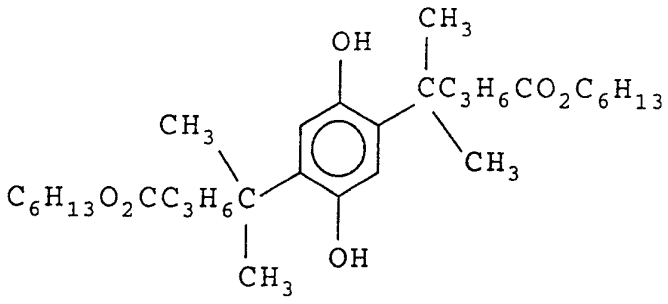
H5



H6



H7



The examples below are intended to be illustrative and not exhaustive of the invention:

Practice of the Invention

The following example is used to illustrate the practice of this invention:

Example 1: Formulation of a neutral image forming dispersion

The oil phase of the dispersion formula is composed of a mixture of:

Cyan coupler A	50.0 grams
Magenta coupler B	37.1 grams
Yellow coupler C	65.6 grams
Coupler solvent D	62.6 grams
Auxiliary solvent E	78.5 grams

The aqueous phase of the dispersion is composed of a mixture of:

Gelatin	120.0 grams
Alkanol-XC Surfactant	12.0 grams
Water	1574.2 grams
Total	2000.0 grams

Procedure:

1) The materials used in the oil phase are combined and heated to 125C with stirring until dissolution occurs.

2) After dissolution occurs, the hot oil phase is quickly added to the preheated (70°C) aqueous phase mixture with stirring.

3) The mixture is then passed through a colloid mill, collected, then rapidly chilled until the dispersion is set.

The above preferred formula was derived from a statistically designed central composit mixture experiment in which the ratio of the components of the

oil phase were systematically varied in different proportions. In all, 27 different formulations, including replicates of the center point were made and tested. Once the coupler dispersions were prepared, they were mixed with additional gelatin, water, and silver halide and coated on a resin coated paper support in a single layer format. The following sample format describes the structure:

Coating 1

Overcoat	1076. mg gel/m^2 spreading aids gel hardener
Emulsion	710. mg/m2 red sensitive silver chloride 1615. mg gel/m^2 377. mg cyan coupler/m^2 280. mg magenta coupler/m^2 495. mg yellow coupler/m^2 (Coupler coverages based upon dispersion formulation above.)
Gel sub	3229. mg gel/m^2
Support	resin coated paper

Once coated, the samples were given a step exposure on a KODAK Model 1B sensitometer. Specifically, they were exposed for 0.1 seconds to a 3000°K tungsten light source through a 0-3 density step wedge. After exposure, the samples were processed through a standard Process RA-4 color process. The process consists of a 45-second developer, a 45-second bleach-fix, and a 90-second wash. The coated paper samples were subsequently dried with hot air.

The RA-4 Color Developer consists of:

Triethanol amine	12.41 g/l
Phorwite REU	2.30
Lithium polystyrene sulfonate (30%)	0.30
N,N-diethylhydroxylamine (85%)	5.40
Lithium sulfate	2.70
Kodak color developer CD-3	5.00
DEQUEST 2010 (60%)	1.16
Potassium carbonate	21.16
Potassium bicarbonate	2.79
Potassium chloride	1.60
Potassium bromide	7.00 mg/l@
Water	to make 1 liter@
pH @ 80F is 10.04 +/- 0.05	

The RA-4 Bleach-fix consists of:

Ammonium thiosulfate (56.5%)	127.40 g/l
Sodium metabisulfite	10.00
Glacial acetic acid	10.20
Ammonium ferric EDTA (44%)	110.40
Water	to make 1 liter@
pH @ 80F is 5.5 +/- 0.10	

Processing the exposed paper samples is done with the developer and bleach-fix temperatures adjusted to 95°F. Washing is performed with tap water at 90°F.

Testing for Neutrality:

After the developed samples are dried, the densities of each step are measured to determine which exposure step produced a status A density nearest 1.0. Once determined, that exposure step has its visible absorption spectra measured. The spectrophotometric data is then converted to colorimetric data, and the corresponding a*, b*, C* and L* values are calculated.

These color coordinates are well-known metrics in the CIE System (International Commission on

Illumination), and their derivation is discussed at length in many texts on color science. One such example is PRINCIPLES OF COLOR TECHNOLOGY, 2nd Edition, authored by Fred W. Billmeyer, Jr. and Max Saltzman, which is published by John Wiley and Sons, New York with pages 25-66 being of particular interest.

Accordingly, the metrics, a* and b* are measures of the color of an object. The value of a* are generally thought of as a measure of the amount of redness or greenness of an object. An object with a positive value for a* is increasingly red while a negative value for a* indicates the degree of greenness in the object. Likewise for b*, a positive value indicates more yellowness; while negative b* values indicate increasing blueness.

An objects lightness or darkness is measured using the term L*. An L* value of 100 indicates that the object is perfectly white; while an L* value of 0 indicates that the object is perfectly black. Values of L* between 0 and 100 indicate intermediate lightness. Chroma, or color saturation, is calculated as C* using the equation described below. In black and white imaging systems it has little meaning since it is derived from the a* and b* terms which are both near zero indicating no color. In black and white photographs, C* is also near zero. For an object to be rendered neutral, it should have a very small values for a* and b*. In fact, the closer that a* and b* are to zero, the lesser amount of color in the object and the more neutral appearing the object will be rendered. In the examples below, L* describes the lightness of the sample patch and does not relate to the color of the patch.

In our colorimetric calculations, we assume a color temperature of D5500 and have used the 1931 CIE 2 degree standard observer.

In the coated example described above, the colorimetry results for the test patch produced an a*

value of -1.15 and a b* of 1.23 with a lightness L* of 38.2. Both the a* and b* values are close to zero indicating neutrality was achieved. As added confirmation, the sample patch when viewed under the appropriate D5500 illumination appeared visually neutral with no evidence of a color bias.

The calculation of the colorimetric values for a*, b*, L* and C* is straightforward and is discussed in detail in the reference noted above. However, to further clarify their calculation, the following derivations are given for the four terms.

First it is necessary to know the spectral power distribution of the light source used as the viewing illuminant. For these calculations we have chosen a daylight source with a color temperature of 5500^{O} Kelvin (D5500). The spectral power distribution, P(lambda), of this source is well known and we have chosen to use the spectral range of 340nm to 800nm in 10nm increments. Secondly, it is necessary to know the reflectance spectrum, R(lambda), of the object being observed. These data are conveniently obtained by using any commercial reflectance spectrophotometer. The wavelength range measured is 340nm to 800nm. Lastly, it is necessary to know the 1931 CIE 2^{O} standard observer functions x(lambda), y(lambda) and z(lambda). These values are also conveniently obtained over the necessary wavelength range from the reference text cited above.

Once the above values are obtained, they are multiplied together as a function of wavelength and the individual values for x(lambda), y(lambda) and z(lambda) are summed to obtain the tristimulus values X, Y and Z. Once obtained, the tristimulus values are used to calculate the colorimetric values of a*, b*, L* and C* using the equations given below:

$$a^* = 500[(X/X_n)^{1/3} - (Y/Y_n)^{1/3}]$$

$$b^* = 200[(Y/Y_n)^{1/3} - (Z/Z_n)^{1/3}]$$

$$L^* = 116(Y/Y_n)^{1/3} - 16$$

$$C^* = (a^{*2} + b^{*2})^{1/2}$$

In these equations, X_n, Y_n and Z_n represent the tristimulus values for the reference white.

Other metrics are also useful when evaluating the tone scale reproduction of various photographic systems. For example, when comparing the color reproduction of an object when reproduced by two different color photographic systems, it is often convenient to describe the tonal difference as delt-E*, which is defined as:

$$\text{delt-E}^* = [(\text{delt-L}^*)^2 + (\text{delt-a}^*)^2 + (\text{delt-b}^*)^2]^{1/2}$$

This equation is known as the color difference equation and was defined by the CIE in 1976. Examination of the equation shows that color difference can be determined by calculating the square root of the sums of the squares of the differences of L*, a* and b* of the two objects.

To further the analogy, it is often desirable to assess the overall tone reproduction of a black and white print which was used to reproduce a scene containing various and assorted colors and compare its tone reproduction to that of a color photograph of the same original scene. One method of comparing two photographic reproductions for tonal accuracy is to use the color difference equation described above.

To assess the difference, it is first necessary to photograph the scene. In our case, a test scene is prepared which contains a neutral step wedge exposure and

assorted other colors which include red, green, blue, cyan, magenta, yellow and other color shades. The color negative used to photograph the scene is developed and subsequently used to print the scene onto the black and white papers being compared. Two commercially available black and white printing papers are Kodak Polycontrast III paper and Kodak Panalure M paper. The Polycontrast III paper is an ortho sensitive paper. This means that it is spectrally sensitive in the blue and green regions of the visible spectrum. In contrast, the Kodak Panalure M paper is pan sensitive. This means that it has sensitivity to red, green and blue light. The third black and white paper to be tested is the paper described below in the multilayer invention.

To make the comparison, the papers described above and a Kodak Ektacolor Supra™ color paper are printed onto with the color negative containing the test scene. The color print is developed in the standard Kodak RA-4 Color Development Process. The Kodak Polycontrast III and the Kodak Panalure M prints are developed in their respective recommended black and white processes. The chromogenic black and white paper of the invention is developed in the same Kodak RA-4 Color Development Process as is the color print above. After printing each paper and matching the density of one of the neutral patches (at approximately 0.8 density), each color and neutral patch of each print has its visible absorbtion spectra determined. These absorption spectra are then converted to their corresponding tristiumlus values and, hence, their colorimetric values are described using the equations above.

To assess the overall colorimetric reproduction error, a comparison can be made between the Kodak Ektacolor Supra Color Paper print relative to the Kodak Polycontrast III print, the Kodak Panalure M print and the chromogenic black and white print from the invention described below. To assess the overall error in tonal

reproduction it is convenient to sum the delt-E*'s for each of the 30 patches of each print. Clearly, the print which has the lowest delt-E* sum has the smallest error in tonal reproduction relative to the color print. The following table shows the accumulated delt-E*'s for the black and white prints described above:

Table 1

Print Material	delt-E*
Kodak Polycontrast III	755.7
Kodak Panalure M	688.3
Invention	633.7

The values for delt-E* listed in Table 1 include in their calculation, the relative changes in a* and b* for each test patch. Since it is expected that a* and b* will approach zero in a black and white print the values for delt-E* will be expectedly large. To better clarify the tonal issue of these reproductions, it is simplier to compare the absolute differences in lightness of each patch by comparing and defining a new colorimetric term delt-Z* which is simply the absolute value of the differences in lightness between two test patches:

$$\text{delt-Z}^* = | L^*_1 - L^*_2 |$$

If we then sum the delt-Z* values for each of the test patches for each of the prints using the Kodak Ektacolor Supra Color Print as the reference print, we can see then in Table 2, below, that the invention retains the lower overall error in lightness reproduction when compared to either of the other black and white print materials.

Table 2

Print Material	delt-E*
Kodak Polycontrast III	312.6
Kodak Panalure M	182.0
Invention	107,8

Graphical information for these data are presented in figures 1, 2 and 3. These graphs show the relative changes in chroma C* and lightness L* for each of the 30 different test patches in each test print when comparing the tonal reproduction of the Kodak Polycontrast III, Kodak Panalure M and the invention when compared to Kodak Ektacolor Supra Color Paper.

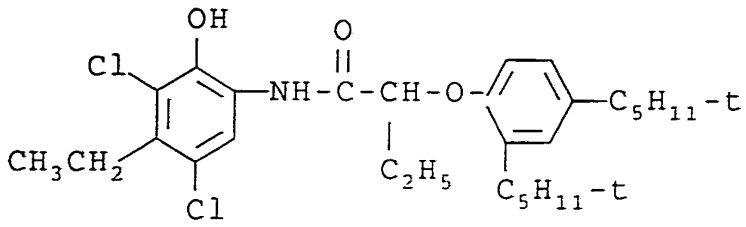
Formulation of a multilayer example in color paper format:

Layer	Composition	mg/m^2
Overcoat	Gelatin	1346.
UV absorbing layer	Tinuvin 326 Tinuvin 328 Oxidized developer scavenger (F) Gelatin	64.6 366. 64.6 656.
Red sensitive layer	Red sensitive silver Cyan coupler (A) Magenta coupler (B) Yellow coupler (C) (From the neutral dye forming dispersion) Gelatin	322. 280. 208. 368. 1184.
UV absorbing layer	Tinuvin 326 Tinuvin 328 Oxidized developer scavenger (F) Gelatin	64.6 366. 64.6 656.
Green sensitive layer	Green sensitive silver Cyan coupler (A) Magenta coupler (B) Yellow coupler (C) (From the neutral dye forming dispersion) Gelatin	407. 421. 313. 553. 1184.
Interlayer	Oxidized developer scavenger (F) Gelatin	93.6 753.
Blue sensitive layer	Blue sensitive silver Cyan coupler (A) Magenta coupler (B) Yellow coupler (C) (From the neutral dye forming dispersion) Gelatin	80.7 140. 104. 184. 1507.
Standard color paper support (resin coated)		

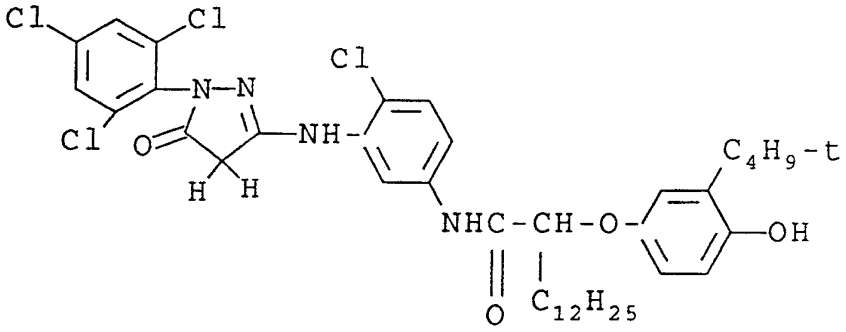
The film structure illustrated above is similar to current conventional color paper. However, it is considered that for the black-and-white paper of the invention, the Interlayer and middle UV absorbing layer would preferably not be present as there is no need to prevent cross contamination of oxidized developer between layers.

All three emulsions used in the multilayer element are silver chloro-bromide (99:1).

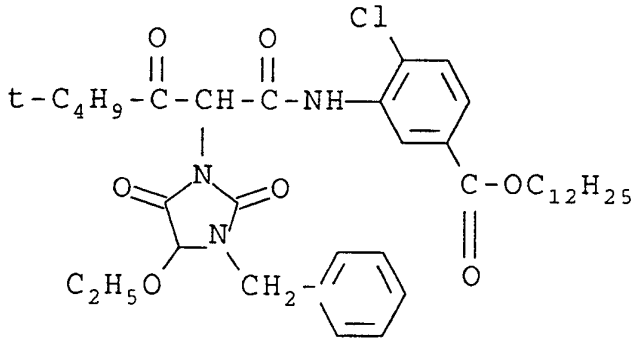
This coating was made on a conventional film forming machine. After coating it was exposed and processed as described earlier. The colorimetric data was also obtained as described earlier. The results of the analysis show that at status A densities of 1.03 red, 1.04 green and 1.02 blue, the corresponding a* and b* values are 0.18 and 0.09 respectively. The related L* value is 38.5. In addition to the extremely low a* and b* values, the exposed patches appeared to be visually neutral.



A

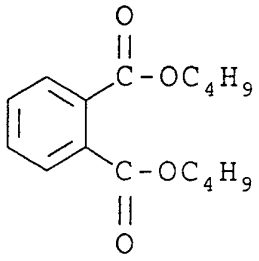


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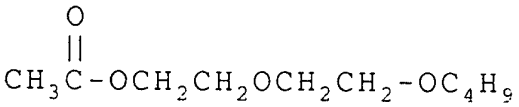


C

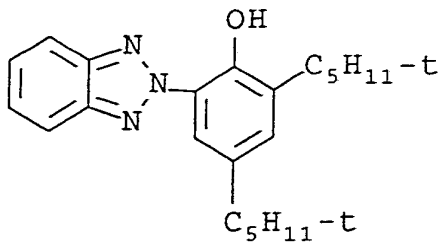
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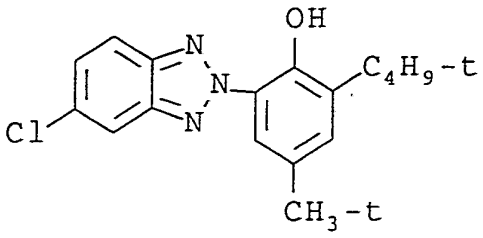
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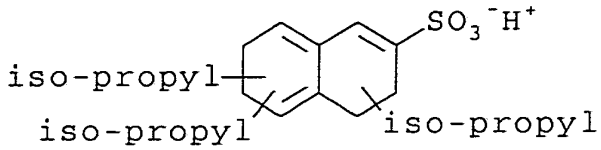
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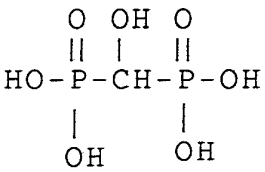
Tinuvin 328



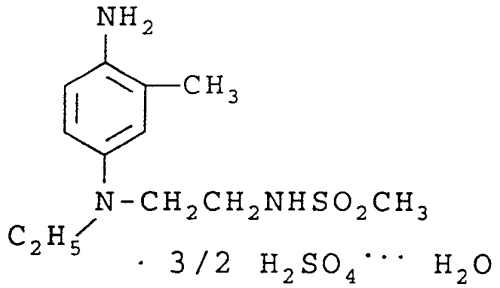
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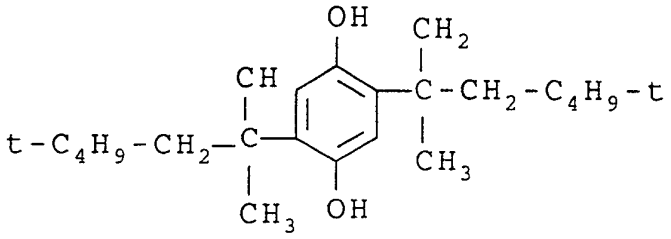
Alkanol-XC



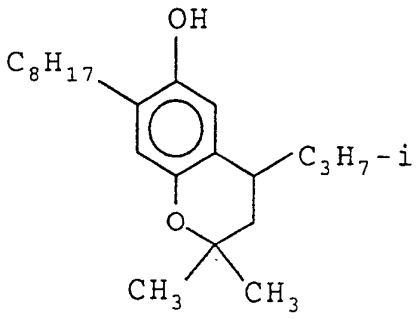
Dequest 2010



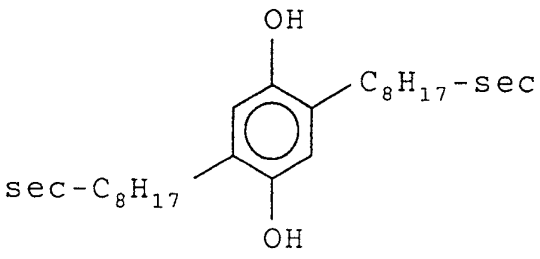
CD3



F



L



N

Layer Structure 1: Multilayer Format (Example 1)

Layer	Material	Coverage (mg/M2)
Overcoat	gelatin	1345.0
	N	21.5
	D	64.6
	Alkanol-XC	21.5
UV Absorber	gelatin	1398.8
	Tinuvin 326	113.0
	Tinuvin 328	640.2
	N	75.2
Red Sensitive Layer	gelatin	1990.6
	Red sensitive silver halide	351.7
	Cyan coupler (A)	478.3
	Magenta coupler (B)	252.2
	Yellow coupler (C)	562.5
	Stabilizer (L)	187.0
	Coupler Solvent (D)	562.0
	Aux. Solvent (E)	530.9
Green Sensitive Layer	gelatin	2152.0
	Green sensitive silver halide	187.7
	Cyan coupler (A)	340.0
	Magenta coupler (B)	178.2
	Yellow coupler (C)	399.8
	Stabilizer (L)	132.9
	Coupler Solvent (D)	399.5
	Aux. Solvent (E)	377.4
Blue Sensitive Layer	gelatin	1506.4
	Blue sensitive silver halide	70.4
	Cyan coupler (A)	175.7
	Magenta coupler (B)	92.7
	Yellow coupler (C)	206.6
	Stabilizer (L)	68.7
	Coupler Solvent (D)	206.5
	Aux. Solvent (E)	195.0
Resin Coated Support		

Layer Structure 2: Multilayer Format (Example 2)

Layer	Material	Coverage (mg/M2)
Overcoat	gelatin	1345.0
	N	21.5
	D	64.6
	Alkanol-XC	21.5
UV Absorber	gelatin	1398.8
	Tinuvin 326	113.0
	Tinuvin 328	640.2
	N	75.2
Red, Green, and Blue Sensitive Layer	gelatin	3509.5
	Red sensitive silver halide (optional)	351.7
	Green sensitive silver halide	187.7
	Blue sensitive silver halide	70.4
	Cyan coupler (A)	997.6
	Magenta coupler (B)	526.1
	Yellow coupler (C)	1173.1
	Stabilizer (L)	390.0
	Coupler Solvent (D)	1172.1
	Aux. Solvent (E)	1107.3
Resin Coated Support		

Layer Structure 1 shown above, gives the structure and composition of the photographic element referred to in Example 1. Layer Structure 2 shown above shows the structure of Example 2 in which all three spectrally sensitized emulsions are contained in a single emulsion layer. This element would be referred to as being 'pansensitized'. It is formed and processed as in Example 1 to produce a pleasing black and white print.

Examples 3-7: Blended Yellow Emulsions for Improved Exposure Latitude

It was found that in order to accurately reproduce the lightnesses of objects in the element described herein when using color negatives, that the amount of blue sensitive emulsion needed was comparatively smaller relative to the amounts of red or green sensitive emulsion.

However, because the emulsions used in these elements are monodisperse, they produce only a narrow range of exposure latitude when processed. This deficiency is even more exaggerated when the coverage of the emulsion is significantly reduced for reasons such as the lightness reproduction referred to above.

To correct for this deficiency, it is possible to design an emulsion which is more disperse, thus improving the desired exposure latitude. This polydisperse emulsion when used in place of the mondisperse emulsion would increase the relative exposure latitude of the element resulting in black and white prints having more perceivable detail in tones where the polydisperse emulsion was exposed.

An alternative approach to this correct for this deficiency is to blend two or more emulsions of different grain sizes or sensitivities which have similar spectral sensitizations. If one chooses to develop two emulsions of similar sensitizations but different grain sizes, there will be a difference in their sensitivity towards light. The larger grain will be more sensitive and the smaller grain less sensitive. Since these two or more emulsions will inherently have different sensitivities because of their respective grain size differences, the blend of at least two such emulsions will result in an effectively increased exposure latitude.

An example of this technology was achieved by blending two, monodisperse yellow emulsions of different grain sizes. Table 1 given below describes the emulsion grains used to achieve this improvement in exposure latitude. The choice of the relative sizes of the emulsions is based upon known metrics for estimating relative sensitivity differences between emulsion grain sizes where the grain shapes are similar.

In the example shown below, the calculated sensitivity difference between the first and second grain is 0.23 log exposure. This difference in sensitivity was judged to be sufficient to increase the relative exposure latitude of the element without being so great as to not be exposed due to insufficient sensitivity or so large as to not be significantly different from the first emulsion in sensitivity.

TABLE 1

Example No.	First Emulsion Cubic Edge Length	Second Emulsion Cubic Edge Length	Ratio of Emulsions
3	0.60μ	0.46μ	1.00:0.00
4	0.60μ	0.46μ	0.75:0.25
5	0.60μ	0.46μ	0.50:0.50
6	0.60μ	0.46μ	0.25:0.75
7	0.60μ	0.46μ	0.00:1.00

The elements described above were prepared according to the description and formulas shown in the multilayer element described in Example 1, Layer Structure 1. The blended emulsions described above as Examples 3-7 were substituted for the single blue sensitive emulsion. The blended pair of emulsions were coated at the same coverage as the single emulsion. The materials prepared were exposed with a color negative which prior had been exposed to a variety of colored test

objects of various lightnesses. The examples were then processed through the standard **Kodak Ektacolor RA-4 process**. The resulting prints were then judged for improved exposure latitude by experienced observers.

The results of the judging indicate that the print showing the greatest improvement in exposure latitude is when the blend ratio of the first and second emulsions is 0.50 to 0.50 as in example print number 3. The prints exhibiting the least amount of exposure latitude were Examples 3 and 7. These observations are totally consistent with the premise that the exposure latitude of a sensitized layer can be increased by combining emulsions of different sensitivity. This advantage of mixed emulsions is also considered to result if the single layer structure of Example 2 is used with blended emulsions.

Examples 8-10: Use of Alternate Yellow Dye Forming Couplers for Improved Light Stability

An observed deficiency in the performance of the element described in Example 1 is the imbalance in the relative fade rates of the three image dyes towards light. If the rate of fade of the three dyes are not the same, the print will develop an undesirable coloration after prolonged exposure to light. In the first example, previously described, prolonged exposure of the print to light results in a change from a neutral appearing image towards a 'bluish' image. The change in image hue is caused by a loss of yellow image dye at a rate greater the loss of cyan or magenta image dyes.

We have found that some yellow dye forming couplers are less susceptible towards light fade than others. The data presented below in Table 2 shows the comparison fades of coupler C with couplers C5 and C8, two alternate yellow dye forming couplers. These two couplers were dispersed in an oil phase containing a cyan

coupler A and a magenta coupler B along with coupler solvent and an auxiliary solvent at a level equimolar to coupler C.

TABLE 2

Change in Red, Green and Blue Density From Initial Density 1.0 After 2 Weeks' Exposure to 50 klux Daylight

Example	Yellow Coupler	Delta Red	Delta Green	Delta Blue
8	(C)	-0.10	-.0.09	-0.22
9	(C5)	-0.15	-0.11	-0.16
10	(C8)	-0.09	-0.06	-0.10

Examples 11-15

Using a modified dispersion formulation in which the stabilizer, (L), was omitted from the oil phase, other yellow couplers were compared to (C) to determine their light stability. The results of these experiment are given below in Table 3.

TABLE 3

Change in Blue Density from Initial Density 1.0 After 2 Weeks' Exposure to 50 klux Daylight

Example	Yellow Coupler	Delta Blue
11	(C)	-0.50
12	(C2)	-0.38
13	(C5)	-0.33
14	(C3)	-0.36
15	(C6)	-0.38

These dispersions were prepared by replacing the (C) coupler in the chromogenic dispersion with an equimolar amount of each of the other yellow couplers. The dispersion formula used and method of preparation is given in Example 1.

The data in Tables 2 and 3 clearly show that the light stability of the yellow component in the chromogenic dispersion is significantly improved by replacing coupler (C) with any of these preferred couplers.

Layer Structure 3 below describes the coating format in which each of these Examples 11-15 was prepared.

Layer Structure 3

Single Layer Coating Structure

Layer	Description	Coverage (mg/M2)
Overcoat	Gelatin	1076
UV Absorber	Gelatin	1334
	Tinuvin 326	129
	Tinuvin 328	732
Emulsion/Dispersion	Gelatin	1614
	Red Sensitive Silver Halide	699
	Cyan Coupler (A)	323
	Magenta Coupler (B)	172
	Yellow Coupler (varies)	380
	Coupler Solvent (D)	380
	Aux. Coupler Solvent (E)	358
Underlayer	Gelatin	3228
Resin Coated Paper Support		

Example 16: Use of Polyester Supports for Diverse Applications

Coating upon resin coated paper stock produces a photographic element suitable for direct reflection viewing. Resin coated paper stock represents only one type of material upon which this type of sensitized material may be coated.

We have produced an example of materials in Example 1 in which the support is a clear 0.18 mm polyester base. This material is suitable for backlit display viewing where the display box contains a light and a translucent diffuser.

Polyester base sheet support thickness can be varied to meet the need of the particular application. Typically, supports of this nature are about 0.10 to 0.18 mm of thickness. Other clear supports such as acetate or cellulose nitrate may also be used.

Other examples of applications where a clear support would be necessary are X-ray films, motion picture films, motion picture intermediate films, slide films, etc.

Example 17

In some applications, the print display box does not contain a translucent diffuser over the light. For applications such as these, a light diffusing layer is coated between the clear photographic support of Example 16 and the sensitized layers of the element. The diffusing layer consists of a mixture of gelatin and titanium dioxide particles whose thickness and concentration are adjusted to provide a minimum density without becoming so translucent as to reveal the back lighting of the display box.

Example 18

A second type of support, commercially available from DuPont, is known as Melinex™. This material is a diffuse reflective support comprised of a voided polyester filled with barium sulfate particles. It was sensitized with the element described above in Example 1 and produced a black and white print.

Other support types could have included films made from polyvinyl acetate, polyvinyl chloride, polyethylene, polycarbonate, polystrene, cellulose nitrate, etc. For completeness, biaxially oriented and voided polypropylene or polyethylene supports as well as paper fiber supports which include materials known to act as water vapor or oxygen barrier layers. In the instance of oxygen barriers, it is well known that polyvinyl alcohol is highly effective.

Color Development Process Modification Requirement For Examples 16-18 Elements:

When sensitizing a support which is transparent, it is necessary to increase the coverage of the sensitizing materials such as the coupler dispersions and the silver halide. Normally it is required to approximately double the coverage of the emulsions and dispersions compared to a reflective support in order to achieve a sufficiently high enough Dmax and contrast. This requires that approximately twice as much silver halide be developed than in the standard color development processing step.

We have learned that in the standard 45 second color development step, that insufficient color developer can diffuse into the element in time to react with the developing silver halide. The solution to this problem is to increase the development time (and also the bleach-fix time) to approximately twice the normal time. With this minor process modification, contrast and D_{max}

densities are substantially higher than with the standard color development time and are judged adequate for direct transmission viewing.

Example 19: Development of an Exceptionally Low Silver Halide Containing Paper

An example of an element has been produced wherein the amounts of silver halide have been substantially reduced. In this example the amounts of red, green and blue silver halide coated are shown in Table 4 below compared to the levels used in Example 1.

TABLE 4

Comparison of Silver Halide Levels in Standard RA4 Color Development Process vs. a Development-Amplifier Process

	Example 1 RA-4 Process	Developer-Amplifier Process
	Nominal Silver Coverage (mg/M2)	Nominal Silver Coverage (mg/M2)
Red	352.	86.7
Green	188.	34.1
Blue	70.4	15.5

The key to the substantial reduction in the amount of silver halide coated, is the used of the 'developer-amplifier' solution in place of the standard color developer bath. The developer-amplifier solution contains hydrogen peroxide. The presence of the hydrogen peroxide allows the re-use of the developed silver halide by acting as an electron transfer agent between the developed silver and the color developer, thus resulting in a substantially reduced need for developed silver.

The formula used for the developer-amplifier solution is given below in Table 5.

TABLE 5

Developer-Amplifier Solution Composition

Component	Concentration g/liter
Kodak Color Developer CD-3 (Methane sulfonamide, N-[2-(4-amino-3-methylphenyl)-ethylamino]ethyl]-sulfate)	3.50
Hydrogen peroxide	50.0 cc of a 3.0% aqueous solution
Potassium bromide	0.001
Potassium chloride	0.50
Potassium carbonate	25.0
Kodak anti-calcium #5 (Dequest 2010) (1-hydroxyethyl-1,1-diphosphonic acid 60%)	0.60
Kodak anti-calcium #8 (diethylenetriamine-pentaacetic acid penta sodium salt 40%)	2.0 cc
N,N-diethylhydroxyl amine (85%)	4.0 cc
Adjust to pH = 10.3 with potassium hydroxide or sulfuric acid	

TABLE 6

Processing Sequence and Times

Process Step	Time
Develop-amplify	45 seconds
Bleach-fix	45 seconds
Wash	90 seconds

Processing temperature is 90°F

Examples 20-23: Addition of an Antioxidant to the Dispersion Oil Phase to Reduce Printout

It is widely known that upon exposure to light photographs fade. The degree of fade is proportional to the type of photographic image (silver vs. pigmented vs. dye), the intensity of light and the wavelength of light. In some photographic images, areas where there is no image (known as D-min) are also discolored. This discoloration is predominantly due the formation of a yellow stain. The yellow stain is known as 'print-out'. Various additives and treatments are usually employed to reduce or minimize this stain build-up upon prolonged exposure to light.

Among the more effective stain preventers used in color photographs using dye images are the class of antioxidants known as hydroquinones. Inclusion of a hydroquinone into the oil phase of the coupler dispersion serves to act as an antioxidant and significantly reduces the yellow stain build-up upon prolonged exposure to light.

We have found that inclusion of such a hydroquinone into the dispersion oil phase significantly reduces the formation of print-out during exposure to light. Table 7 below shows the reduction in print-out obtained when samples of the photographic element formed in Example 1 are exposed to 2 weeks of 50 klux sunshine.

TABLE 7

Printout Reduction as A Function of Antioxidant Level

Example	Weight Fraction of Antioxidant in the Dispersion	Print-out After 2 Weeks of 50 Klux Sunshine Exposure Change in Blue D_{min} Density
20	0.000	0.08
21	0.013	0.05
22	0.026	0.03
23	0.052	0.00

In the Examples 20-23 above, the hydroquinone chosen is 1,4-benzenediol, 2,5-di-sec-dodecyl. The data in Table 7 clearly shows how the inclusion of the hydroquinone antioxidant reduces the amount of printout formed during exposure to light. It should also be noted that hydroquinones other than the example shown above would be equally as effective.

Examples 24-30: Addition of a Hydrophobic Polymer to Reduce the Sensitivity to Leuco Dye Formation in the Bleach Fix

It is widely known that cyan dyes formed in the chromogenic development process have electrolytic reduction potentials low enough to make the dye susceptible to leuco cyan dye formation if the oxidation potential of the bleach-fix (blix) falls below a certain minimum threshhold. The susceptibility of the dye to reduction is a function of the concentration of ferrous ion in the blix, the pH of the blix and the hydrophobicity of the dispersion in which the cyan dye was formed.

Leuco cyan dye formation usually is not a problem in the development process, unless the blix becomes exhausted and is under-replenished. When the blix is under-replenished, the amount of ferrous ion generated as the ferric ion oxidizes the developed silver to form silver bromide, increases to a level where significant leuco cyan dye formation may occur. This leuco cyan dye formation is undesirable since it results in the apparent reduction of cyan contrast in the print which affects color reproduction.

In the instance of a chromogenically formed neutral print such as described in this application, the result is an image which appears red in color since it is lacking in the desired cyan dye amount.

In order to reduce this sensitivity to leuco cyan dye formation, we have found that the addition of a hydrophobic polymer, such as poly-t-butyl acrylamide, to the oil phase of the chromogenic dispersion, results in a balanced neutral dye image which is substantially reduced in leuco cyan dye sensitivity.

Table 8, shown below, gives the dispersion formulations used to test the invention:

Table 8

Measurement of the Leuco Cyan Dye Sensitivity of the Chromogenic Dispersion

	Couplers Used in Chromogenic Dispersion				Change in Cyan Dye Density
Example	Cyan	Magenta	Yellow	Polymer Ratio[1]	from Dr=1.0
24	A	B	C8	0.44	-0.093
25	A	B	C5	0.44	-0.057
26	A	B	C	0.44	-0.054
27	A	B	C8	0.66	-0.099
28	A	B	C5	0.66	-0.071
29	A	B	C	0.66	-0.034
30	A5	B	C5	0.44	-0.031
Reference	A	B	C	0.00	-0.131

[1]Ratio of poly-t-butyl acrylamide polymer to yellow coupler in the chromogenic dispersion.

It is seen in Table 8 that the inclusion of the polymer, poly-t-butylacrylamide to a variety of dispersions effectively and surprisingly, reduces the sensitivity of the processed coated to the formation of leuco cyan dye.

Description of the Procedure Used to Perform the Leuco Cyan Dye Test:

The chromogenic dispersion was prepared as described earlier. Where required, the polymer, poly-t-butyl acrylamide, is added to the oil phase of the

dispersion in the ratios described in the Table 8 above. Once the dispersion was prepared, it was coated, using standard coating techniques onto resin coated color paper stock with an amount of silver halide.

After the coating is prepared, it is exposed and processed as described earlier. Using an X-rite densitometer, the area of the coating having a status A density of 1.0 is determined and marked. The coating is then immersed in a reducing bath of ferrous sulfate for 5 minutes. After rinsing in distilled water and drying, the density of the sample is re-measured and the change in density from Dr=1.0 calculated by difference from the original sample.

A ferrous sulfate bath, such as that described below, simulates the chemistry of an exhausted bleach-fix bath where the bleach (ferric ion) has been reduced by developed silver to ferrous ion, which is known as a powerful reducing agent.

The ferrous sulfate bath used to treat the samples is prepared by dissolving 41.8g of tetra-sodium-ethylenediaminetetraacetic acid (EDTA) in 1.0 L of distilled water, then adjusting the pH to 4.0 with a 10% solution of nitric acid. The solution is then de-oxygenated by bubbling nitrogen through it for 15 minutes. Then 15.2g of ferrous sulfate (heptahydrate) is added with stirring. The pH of the solution is then raised to 5.0 with dilute ammonium hydroxide.

It should be readily apparent that this invention is not limited to the application described above. Listed below, are examples of other photographic systems where the ability to produce a black-and-white image which does not contain silver are of utility and advantageous:

1) Producing black-and-white motion picture print film. Producing black-and-white motion picture print film on an acetate or polyester base that prints from color negative products can be printed onto using current

motion picture printing technology. There currently is not a black-and-white motion picture print film on the market. This would allow contrast manipulation in the printing stage rather than the processing stage as is currently done, by regulating the ratio of the red, green, and blue sensitive layers independently. It would also allow the color timers the possibility of making scene-to-scene contrast changes in the printing stage similar to the way they currently make scene-to-scene color timing. This is not possible currently.

2) Producing black-and-white motion picture intermediate film. It would be extremely useful in the special effects industry and would provide much greater flexibility in creating burn-in mattes, and holdout mattes.

3) Producing a black-and-white display film for backlighting applications.

4) Producing X-ray film from which all the silver is recovered.

5) Producing a masking film.

6) Producing a motion picture sound track film.

7) A black-and-white layer incorporated into a color negative film in a very slow emulsion layer for detail enhancement.

8) Increasing exposure latitude in a color print by using the neutral image forming dispersion in a layer adjacent to one or more color layers such as described in U.S. Patent 4,946,765 (P. Hahm, Eastman Kodak Co.).

These and other variations of the invention are within the scope of the invention disclosed herein.

The invention has been described in detail with particular reference to preferred embodiments thereof, but it will be understood that variations and modifications can be effected within the spirit and scope of the invention.

CLAIMS:

1. A photographic element for forming neutral images comprising at least one layer of balanced cyan, magenta, and yellow dye-forming couplers with silver halide grains comprising at least one of blue sensitized silver halide grains and green sensitized silver halide grains.

2. The element of Claim 1 wherein said at least one layer comprises at least two layers, one layer comprising blue sensitized silver halide and one comprising green sensitized silver halide grains and each layer further comprising balanced cyan, magenta and yellow dye-forming couplers.

3. The element of Claim 2 wherein said element further comprises a layer comprising balanced cyan, magenta, and yellow dye-forming couplers and red sensitized silver halide grains.

4. The element of Claim 1 wherein said element comprises one layer comprising balanced cyan, magenta and yellow dye-forming couplers and blue sensitized and green sensitized silver halide grains.

5. The element of Claim 3 further comprising red sensitized silver halide grains.

6. The element of Claim 1 wherein said element forms a black-and-white neutral image when developed.

7. The photographic element of Claim 1 wherein said sensitized silver is coated in ratios corresponding to eye response.

8. The photographic element of Claim 1 wherein said silver halide is in the ratio of two red sensitized,

to three green sensitized, to one blue sensitized silver halide.

9. The element of Claim 1 wherein said element comprises a photographic paper.

10. The element of Claim 1 wherein each of said at least one layers is neutral balanced.

11. The element of Claim 1 comprising a motion picture film.

12. The element of Claim 1 comprising an X-ray film.

13. The element of Claim 1 comprising a* and b* values of about 0.

14. A method of forming a black-and-white image comprising providing a photographic element comprising at least one layer of neutral balanced cyan, magenta, and yellow dye-forming couplers with silver halide grains comprising at least red sensitized silver halide grains and green sensitized silver halide grains.

15. The method of Claim 1 wherein said at least one layer comprises at least two layers, one layer comprising blue sensitized silver halide and one comprising green sensitized silver halide grains and each layer further comprising balanced cyan, magenta and yellow dye-forming couplers.

16. The method of Claim 2 wherein said element further comprises a layer comprising balanced cyan, magenta, and yellow dye-forming couplers and red sensitized silver halide grains.

17. The method of Claim 1 wherein said element comprises one layer comprising balanced cyan, magenta and yellow dye-forming couplers and blue sensitized and green sensitized silver halide grains.

18. The method of Claim 3 further comprising red sensitized silver halide grains.

19. The method of Claim 13 wherein said element forms a black-and-white neutral image when developed.

20. The method of Claim 13 wherein said sensitized silver is coated in ratios corresponding to eye response.

21. The method of Claim 13 wherein said silver halide comprises the weight ratio of two red sensitized, three green sensitized, and one blue sensitized silver halide grains.

22. A photographic element comprising, at least one layer of balanced cyan, magenta, and yellow dye-forming couplers with blue sensitized silver halide grains, and at least one layer of balanced cyan, magenta, and yellow dye-forming couplers with green sensitized silver halide grains.

23. The element of Claim 22 wherein further including at least one layer of balanced cyan magenta and yellow dye forming couplers with red sensitized silver halide grains.

24. The element of Claim 22 wherein said element forms a black-and-white neutral image when developed.

25. The photographic element of Claim 23 wherein said sensitized silver is coated in ratios corresponding to eye response.

26. The photographic element of Claim 23 wherein said silver halide is in the ratio of two red sensitized, to three green sensitized, to one blue sensitized silver halide.

27. The element of Claim 22 wherein the silver halide is selected from at least one member of the group consisting of silver chloride, silver chlorobromide, silver bromide, silver bromoiodide or silver chlorobromoiodide.

28. The element of Claim 22 wherein said element comprises a photographic paper.

29. The element of Claim 23 wherein each of said at least one layers is neutral balanced.

30. The element of Claim 23 comprising a* and b* values of about 0.

31. A method of forming a black-and-white image comprising providing a photographic element comprising at least one layer of neutral balanced cyan, magenta, and yellow dye-forming couplers with red sensitized silver halide grains, at least one layer of neutral balanced cyan, magenta, and yellow dye-forming couplers with blue sensitized silver halide grains, and at least one layer of neutral balanced cyan, magenta, and yellow dye-forming couplers with green sensitized silver halide grains, exposing said element to a color image and developing said element.

32. The method of Claim 31 wherein said element forms a black-and-white neutral image when developed.

33. The method of Claim 31 wherein said sensitized silver is coated in ratios corresponding to eye response.

34. The method of Claim 32 wherein said silver halide is in the weight ratio of two red sensitized, three green sensitized, and one blue sensitized silver halide grains.

35. A photographic element comprising, at least one layer of balanced cyan, magenta, and yellow dye-forming couplers and blue sensitized silver halide grains, and green sensitized silver halide grains.

36. The element of Claim 35 wherein said at least one layer further includes red sensitized silver halide grains.

37. The element of Claim 35 wherein said element forms a black-and-white neutral image when developed.

38. The photographic element of Claim 36 wherein said sensitized silver is coated in ratios corresponding to eye response.

39. The photographic element of Claim 36 wherein said silver halide is in the ratio of two red sensitized, to three green sensitized, to one blue sensitized silver halide.

40. The element of Claim 35 wherein the silver halide is selected from at least one member of the group

consisting of silver chloride, silver chlorobromide, silver bromide, silver bromoiodide or silver chlorobromoiodide.

41. The element of Claim 35 wherein said element comprises a photographic paper.

42. The element of Claim 35 wherein said at least one layer is neutral balanced.

43. The element of Claim 35 comprising a* and b* values of about 0.

44. A method of forming a black-and-white image comprising providing a photographic element comprising at least one layer of neutral balanced cyan, magenta, and yellow dye-forming couplers with red sensitized silver halide grains, blue sensitized silver halide grains, and green sensitized silver halide grains, and exposing said element to a color image and developing said element.

45. The method of Claim 44 wherein said element forms a black-and-white neutral image when developed.

46. The method of Claim 44 wherein said sensitized silver is coated in ratios corresponding to eye response.

47. The method of Claim 46 wherein said silver halide is in the weight ratio of two red sensitized, three green sensitized, and one blue sensitized silver halide grains.

FIG. 1

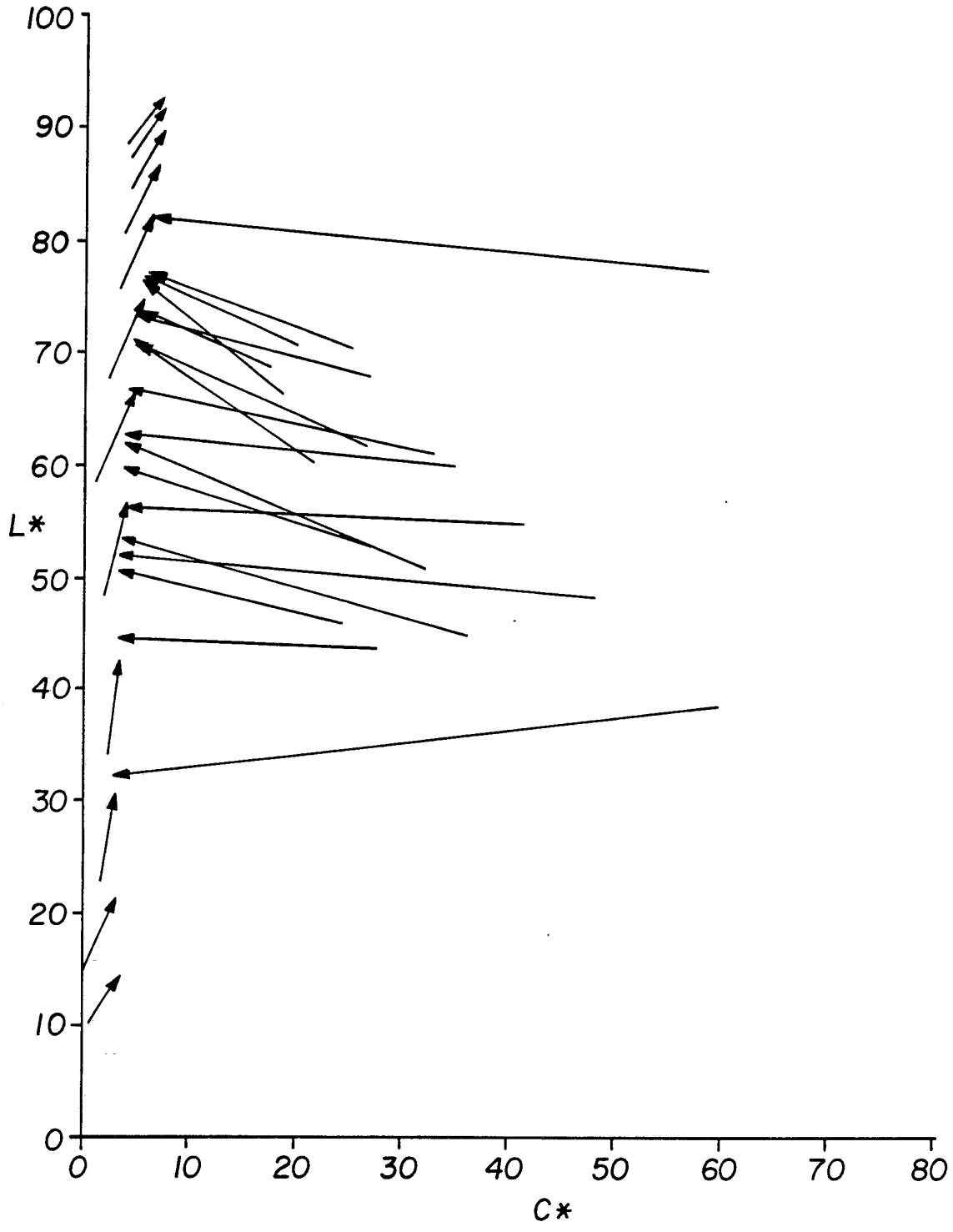


FIG. 2

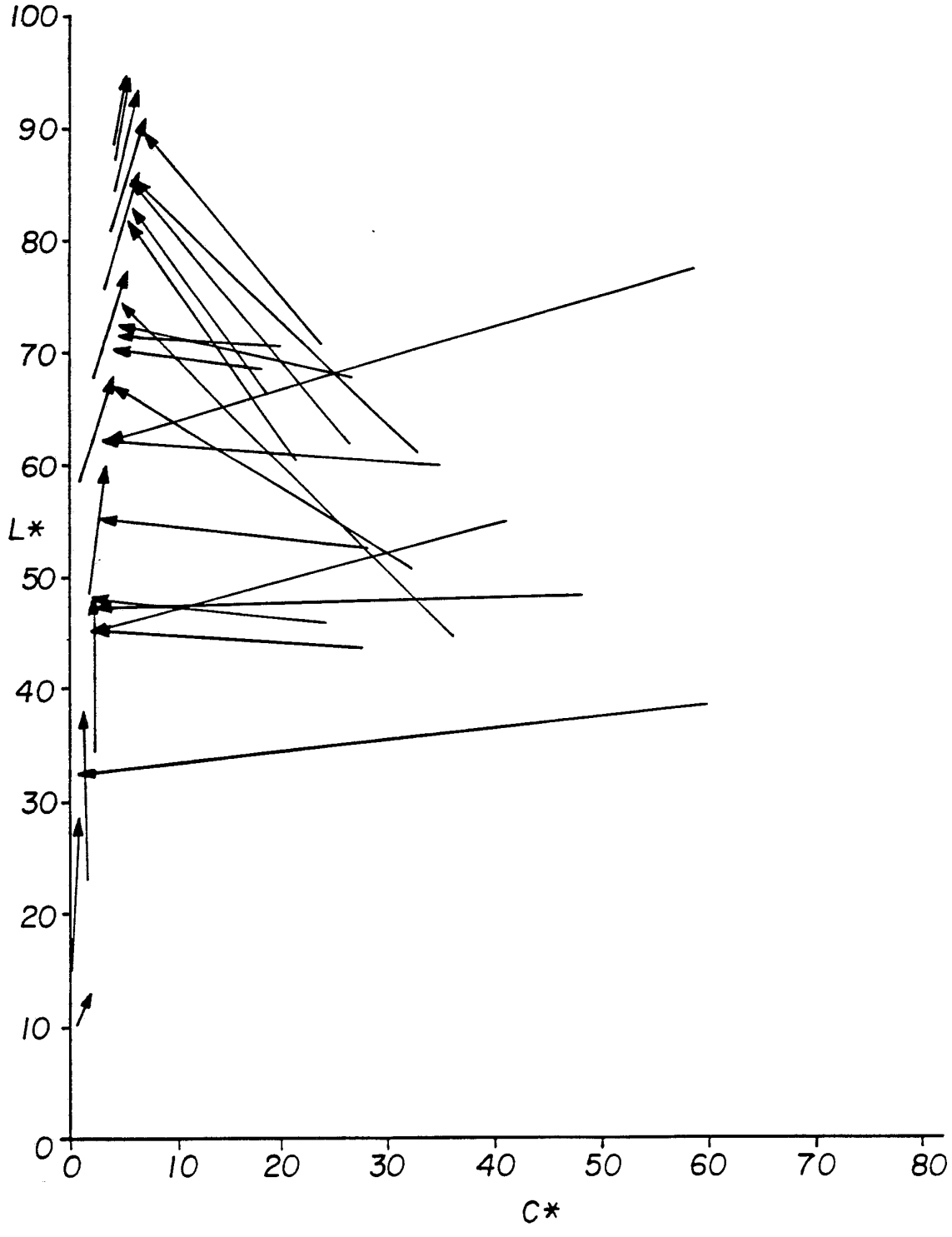
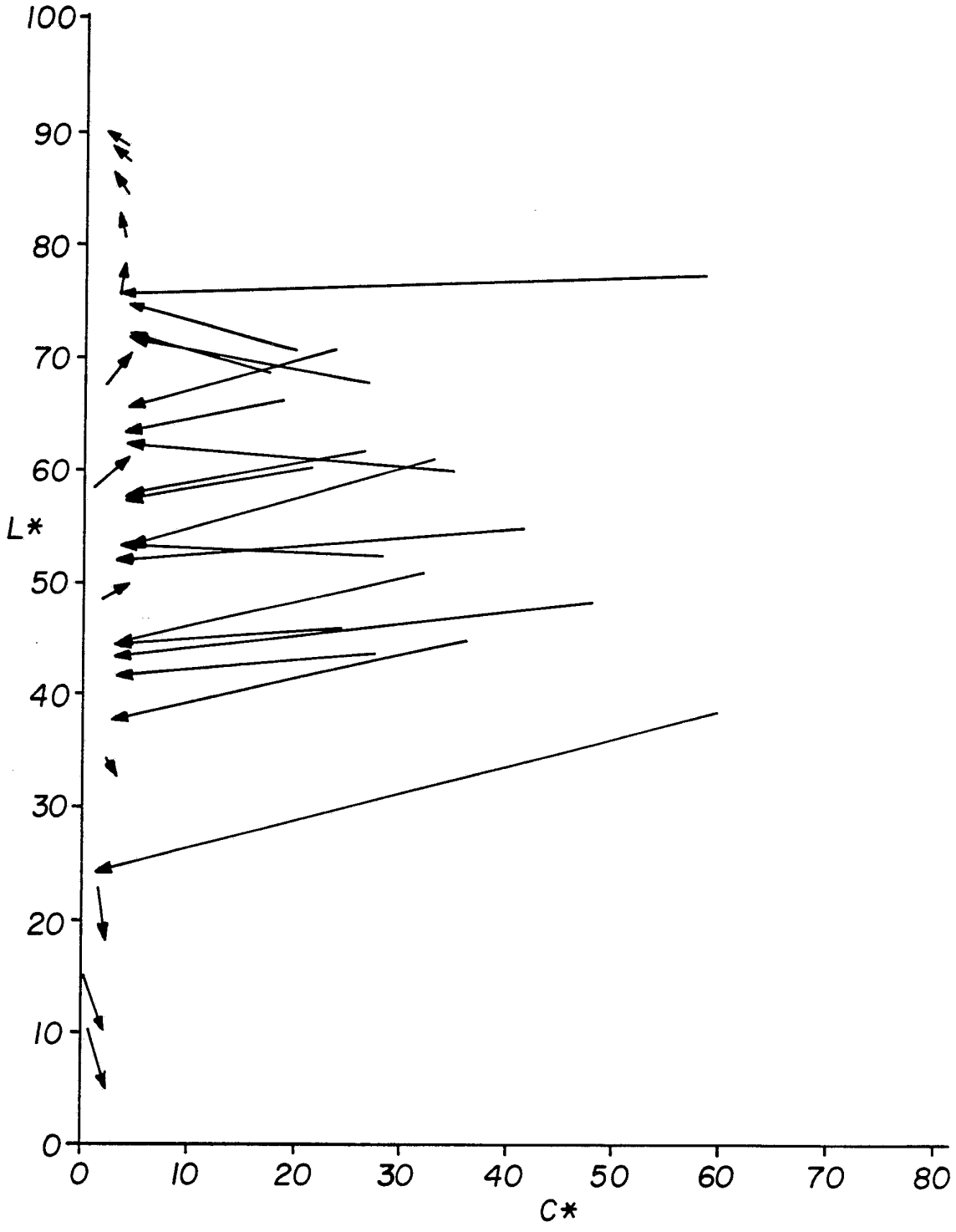


FIG. 3



INTERNATIONAL SEARCH REPORT

International Application No PCT/US 92/10704

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all)[6]

According to International Patent Classification (IPC) or to both National Classification and IPC

Int.Cl. 5 G03C7/30

II. FIELDS SEARCHED

Minimum Documentation Searched[7]

Classification System	Classification Symbols
Int.Cl. 5	G03C

Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched[8]

III. DOCUMENTS CONSIDERED TO BE RELEVANT[9]

Category °	Citation of Document,[11] with indication, where appropriate, of the relevant passages[12]	Relevant to Claim No.[13]
X	DE,A,743 341 (I. G. FARBENINDUSTRIE) 23 December 1943 see the whole document	1,4,6,9, 10,14, 17, 35-37, 40-42, 44-45
Y	PATENT ABSTRACTS OF JAPAN vol. 015, no. 301 (P-1233)31 July 1991 & JP,A,31 05 341 (KONICA) 2 May 1991 see abstract	1-47
Y	PATENT ABSTRACTS OF JAPAN vol. 015, no. 305 (P-1234)5 August 1991 & JP,A,31 07 144 (KONICA) 7 May 1991 see abstract	1-47

-/--

° Special categories of cited documents :[10]

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"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)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"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

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"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.

"&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report
29 MARCH 1993	22 APR 93
International Searching Authority	**Signature of Authorized Officer**
EUROPEAN PATENT OFFICE	PHILOSOPH L.

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)

Category °	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No.
Y	GB,A,1 160 525 (EASTMAN KODAK COMPANY) 6 August 1969 see the whole document ---	1-47
A	EP,A,0 321 401 (ILFORD) 21 June 1989 ---	
A	EP,A,0 025 775 (CIBA-GEIGY) 25 March 1981 -----	

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

US 9210704
SA 68306

This annex lists the patent family members relating to the patent documents cited in the above-mentioned international search report. The members are as contained in the European Patent Office EDP file on
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Patent document cited in search report	Publication date	Patent family member(s)	Publication date
DE-A-743341		BE-A- 419447	
		DE-C- 750971	
		FR-A- 816328	
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