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PATENT SPECIFICATION



Application Date: April 23, 1936. No. 14356/36. (Divided out of No. 478,942).

Complete Specification Left: May 24, 1937.

Complete Specification Accepted: Jan. 24, 1938.

PROVISIONAL SPECIFICATION

Improvements in and relating to Colour Forming Developers and Processes of Colour Development

We, Kodak Limited, a Company registered under the Laws of Great Britain, of Kodak House, Kingsway, London, W.C.2, do hereby declare the 5 nature of this invention which has been communicated to us by Eastman Kodak Company, a Company organised under the Laws of the State of New Jersey, United States of America, of 343, State 10 Street, Rochester, New York, United States of America, to be as follows:—

This invention relates to improvements

This invention relates to improvements in colour forming developers and in processes of colour development for use 15 in connection with colour photography.

It is known that coloured photographic images may be formed by using a developer which forms a coloured compound on development. The coloured 20 compound thus formed is deposited adjacent to the silver grains of the silver image during the development. It is also known that a coloured image may be formed by adding to certain developer 25 solutions a compound which couples, during development, with the oxidation product of the developing agent and forms a colouring substance which is likewise deposited adjacent to the silver 30 grains of the silver image during development. Such a compound, which is employed in conjunction with a developing agent for the silver and which couples with the oxidation product thereof during 35 development, is referred to herein as a colour coupler.

The present invention concerns new or improved colour forming developers comprising an aromatic amino compound 40 serving as the developing agent and a colour coupler as hereinafter defined and also includes a new or improved colour development process which consists in developing a reducible silver salt image 45 in a photographic element with the aid of an aromatic amino compound in presence of a colour coupler as hereinafter defined.

The silver can be removed from the 50 image after colour development leaving a clear transparent dye image.

The invention also includes a photographic element having at least one layer containing a clear transparent image composed essentially of the product 55 resulting from the coupling in situ during development of a colour coupler as hereinafter defined with the oxidation product of an aromatic amino developing

When a silver halide emulsion containing a latent photographic image is deevloped the silver halide is reduced to metallic silver and the developing agent is oxidized. The aromatic diamino compounds which have been used as developing agents form, on oxidation, products which will couple with colour couplers during development to form dyes. If such colour couplers are added to the 70 developer solution, or incorporated in the emulsion layer, the dye which is thus formed by coupling during development is deposited in the gelatine or other silver halide carrier adjacent to the metallic 75 silver grain. It is desirable that the dyes thus formed should not readily wander from the place of formation. It is accordingly desirable that they should be insoluble in water. They are not 80 physically attached to the silver grain and therefore the silver may be subse-quently bleached out of the carrier layer leaving a pure dye image.

Numerous substances have hitherto 85 been employed or proposed as colour couplers among which may be mentioned phenols, naphthols, cresols, nitrophenylacetonitriles and acetoacetic esters. has not, however, always been possible 90 among those hitherto available to select one which exhibits all the desired combination of properties required for any specific case. In colour-developing a gelatino-silver halide emulsion layer it is 95 necessary to select a colour coupler which will give just the desired shade in conjunction with the colours which are produced in other layers. It is moreover important to employ a colour coupler 100 which gives a coloured compound which is resistant to the normal processing baths

employed, although it may often be desirable to have one which gives a colour 5 processing.

employed according to the present invention are suitable in carrying out the which can be destroyed and/or removed if desired during some step in the processing. Many of the colour couplers 427,472, 427,516, 427,518, 427,520, 440,089 and 447,092.

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Ethyl-β-(1-benzthiazolylhydrazone) butyrate

CH2-COOC2H3

Ethyl p-nitrophenylacetate

3. C₂H₅OOC—CH₂—COOC₂H₅

Ethyl malonate

4.

Dihydro-2-hydroxyquinoxaline

OH

15

1:4-Dihydroxyisoquinoline (4-Hydroxyisocarbostyril)

5

Rhodanine

The colours given by the colour couplers named above when used in conjunction with p-aminodiethylaniline, 20 for example, as developing agent are

1. Magenta 2. Yellow 3. Yellow 4. Blue

5. Yellow-brown 20 The aromatic amino compounds which may be used as developing agents in the present invention include the mono- di-, and tri-amino aryl compounds. Among 30 the monoamino compounds may be mentioned the aminophenols and aminocresols and their halogen substituted derivatives as well as the amino-naphthols. The developing agents usually used are the diamino compounds such as para-phenylene diamine and its substitution 35 diamino compounds products. These developing agents may

be substituted in the amino groups or in the ring or in both, forming compounds such as the alkyl phenylene diamines. 40 toluylene-diamines, alkyl-toluylene amines and aminodiphenylamines. These compounds are usually kept in the salt form such as hydrochloride or sulphate since these are more stable than the 45 amines themselves. As examples of developing agents of this class, there may be mentioned diethyl para-phenylene diamine, mono-methyl para-phenylene diamine, dimethyl para-phenylene diamine 50 and ortho-amino-diethylaniline.

The present invention may be utilised in the formation of coloured photographic images on plates or papers as well as on films and the dyes will be formed 55 when gelatine or other carrier for the silver halide is used. The plates, films or papers may have differently sensi-

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tized emulsions of the mixed grain type convert it into a developable silver salt or superimposed on one side or on both sides of the support. The dyes formed may be decolourized by an oxidizing 5 agent such as chromic acid and colourless soluble compounds thereby formed. The bleaching of the dye in this manner need not destroy the silver image but may

image which can in turn be coloured, 10 bleached and recoloured a number of times.

Dated this 22nd day of May, 1937.
W. P. THOMPSON & CO.,
12, Church Street, Liverpool, 1, Chartered Patent Agents.

COMPLETE SPECIFICATION

Improvements in and relating to Colour Forming Developers and Processes of Colour Development

We, Kodak Limited, a Company registered under the Laws of Great
15 Britain, of Kodak House, Kingsway,
London, W.C.2, do hereby declare the nature of this invention which has been communicated to us by Eastman Kodak Company, a Company organised under 20 the Laws of the State of New Jersey, United States of America, of 343, State Street, Rochester, New York, United States of America, and in what manner the same is to be performed, to be parties the same is to be performed, to be particu-25 larly described and ascertained in and by the following statement:

This invention relates to improvements in colour forming developers and in processes of colour development for use 30 in connection with colour photography.

It is known that coloured photographic

images may be formed by using a developer which forms a coloured compound on development. The coloured 35 compound thus formed is deposited adjacent to the silver grains of the silver image during the development. It is also known that a coloured image may be formed by adding to certain developer 40 solutions or by incorporating in the gelatino-silver halide emulsion before or after exposure a compound which couples, during development, with the oxidation product of the developing agent and 45 forms a colouring substance which is likewise deposited adjacent to the silver grains of the silver image during development. Such a compound, which is employed in conjunction with a

couples with the oxidation product thereof during development, is referred to herein as a colour coupler. The present invention concerns new or 55 improved colour forming developers comprising an aromatic amino compound

50 developing agent for the silver and which

serving, as the developing agent and a colour coupler as hereinafter defined and also includes a new or improved colour 60 development process which consists in developing a reducible silver salt image in a photographic element with the aid of an aromatic amino compound in presence of a colour coupler as hereinafter defined as well as the colour photographic 65. elements resulting therefrom. It also includes photographic sensitive elements having such a colour coupler incorporated in one or more emulsion layers.

The silver can be removed from the 70 image after colour development leaving a clear transparent dye image.

The invention also includes a photographic element having at least one layer containing a clear transparent 75 image composed essentially of the product resulting from the coupling in situ, during development of a developable silver salt, of a colour coupler as hereinafter defined with the oxidation product 80 of an aromatic amino developer agent.

When a silver halide emulsion containing a latent photographic image is developed, the silver halide is reduced to metallic silver and the developing agent 85 is oxidised. The aromatic diamino compounds which have been used as developing agents form, on oxidation, products which will couple with colour couplers during development to form dyes. such colour couplers are added to the developer solution, or incorporated in the emulsion layer, the dye which is thus formed by coupling during development is deposited in the gelatine or other silver 95 halide carrier adjacent to the metallic silver grain. It is desirable that the dyes thus formed should not readily wander from the place of formation. I is accordingly, desirable that they should 100 be insoluble in water. They are probably not physically attached to the silver grain. The silver may be subsequently bleached out of the carrier layer leaving a pure dye image.

Numerous substances have hitherto been employed or proposed as colour couplers among which may be mentioned phenols naphthols, cresols, nitrophenylacetonitriles and acetoacetic esters. It 110 has not, however, always been possible among those hitherto available to select one which exhibits all the desired combination of properties required for

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any specific case. In colour-developing a gelatino-silver halide emulsion layer it is necessary to select a colour coupler which will give just the desired shade

5 in conjunction with the colours which are produced in other layers. It is moreover important to employ a colour coupler which gives a coloured compound which is resistant to the normal processing 10 baths employed, although it may often be desirable to have one which gives a

colour which can be destroyed and/or removed if desired during some step in the processing. Many of the colour couplers employed according to the 15 present invention are suitable in carrying out the processing described in our prior patents Nos. 427,472, 427,516, 427,518, 427,520, 440,032, 440,089, and 447,092.

According to the present invention the 20 substance employed as a colour coupler is

one of the following:

Ethyl-\$-(1-benzthiazolylhydrazone) butyrate

2.
$$CH_2$$
— $COOC_2H_5$

Ethyl p-nitrophenylacetate

3. C₂H₅OOC—CH₂—COOC₂H₅

Ethyl malonate

Dihydro-2-hydroxyquinoxaline

1:4-Dihydroxyisoquinoline (4-Hydroxyisocarbostyril)

Rhodanine

These compounds, when present during 30 the development of a silver salt with an aromatic amino developing agent, couple with the oxidation product of such developing agent forming a dye simultaneously with the formation of the silver 35 image.

Ethyl β -(1-benzthiazolyl hydrazone) butyrate was prepared by the action of The

1-benzthiazolylhydrazine on ethyl acetoacetate in acetic acid solution. 40 product was recrystallized from dilute alcohol and melted at 108-109° C.

Dihydro - 2 - hydroxyquinoxaline was

prepared in the following manner: 300 g. of o-phenylenediamine and 105 g. zinc dust are mixed thoroughly in a 45 flask and 270 g. of chloroacetic acid is added and mixed uniformly with the other reactants. In two or three minutes, a very energetic reaction takes place. When it has subsided, the resulting 50 gummy mass is heated on a steam bath for one hour. The reaction mixture is extracted with boiling water as long as an extract, upon being chilled deposits a crop of the product. The dihydro-2-55 hydroxyquinoxaline after recrystalliza-

	tion from water, melts at 133-135° C.
	The aromatic amino compounds which
	may be used as developing agents in the
	present invention include the mono- di-,
=	and tri-amino aryl compounds. Among
Ð	the measure and compounds. Among
	the monoamino compounds may be men-
	tioned the aminophenols and amino-
	cresols and their halogen substituted
	derivatives as well as the amino-
10	naphthols. The developing agents
	usually used are the diamino compounds
	such as para-phenylene diamine and its
	substitution products. These developing
	agents may be substituted in the amino
15	group or in the ring or in both, forming
TO	sompounds such as the all-
	compounds such as the alkyl phenylene
	diamines, toluylene-diamines, alkyl-
	toluylene diamines and aminodiphenyl-
	_ •
	A .
	Diethyl para phenylenediamine -
	Sodium sulphite

amines. These compounds are usually kept in the salt form such as hydro- 20 chloride or sulphate since these are more stable than the amines themselves. As examples of developing agents of this class, there may be mentioned diethyl para-phenylene diamine, mono-methyl 25 para-phenylene diamine, dimethyl paraphenylene diamine and ortho-aminodiethylaniline.

As would be expected from the behaviour of known colour couplers the 30 shade of the colour obtained by coupling generally varies in accordance with the developing agent selected.

EXAMPLE. A developing formula which may be 35 used is the following:

			\mathbf{A}										
	Diethyl	para	$_{ m pheny}$	lenedia	$_{ m mine}$	-	-	_	-	_	_	-	1 gram
40	Sodium	sulph	11te -	-	-	-	-	-	-	_			$0.5~\mathrm{gram}$
	Sodium	carbo	nate	-	-	-	-	-	-	-	-	-	20 grams
	Water	-	-	-	-	-	-	-	-	-	-	-	1 l $itre$
			В										
	Colour coupler Water miscible solvent, such as acetone. For use, B is added to A.			The	mol	molecular equivalent of the developing ag						ping agent.	
45				-	-	-	-	-	-	-	-	50 cc.	

The developing agent and the proportions of the ingredients used in the above formula may, of course, be varied. 50 Solvents other than acetone, such as alcohols, may also be used.

The colours formed by the compounds of the present invention on coupling with the oxidization product of the developer

> 1. Magenta 2. Yellow 3. Yellow 4. Blue

60

5. Yellow-brown Although we have described our invention with particular reference to the use of the colour coupler in the developing solution itself, our invention is in no way 65 limited to this method. As an alternative method, the colour coupler may be incorporated in the photographic layer before development, and either before or after exposure. It may be absorbed upon 70 the sensitive silver halide grains.

The present invention may be utilised in the formation of coloured photographic images on plates or papers as well as on films employing gelatine or other carrier 75 for the silver halide. The plates, films or papers may have differently sensitized emulsions of the mixed grain type or superimposed on one side or on both sides of the support. The dyes formed may be 80 decolourized by an oxidizing agent such

as chromic acid and colourless soluble compounds thereby formed. The bleaching of the dye in this manner need not destroy the silver image but may convert it into a developable silver salt image 85 which can in turn be coloured, bleached and recoloured a number of times.

Having now particularly described and ascertained the nature of the said invention and in what manner the same is to 90 be performed, as communicated to us by our foreign correspondents, we declare that what we claim is:-

1. A colour developer comprising an aromatic amino developing agent and a 95 colour coupler consisting of one of the following compounds, namely, Ethyl- β -(1 - benzthiazolylhydrazone) butyrate, Ethyl p - nitrophenyl - acetate, Ethyl malonate, Dihydro - 2 - hydroxyquin- 100 malonate, Dihydro - 2 - hydroxyquinoxaline, 1:4-Dihydroxyisoquinoline (4-Hydroxyisocarbostyril), Rhodanine.

2. A colour forming developer as claimed in Claim 1, in which the developing agent is an aromatic diamino 105 compound.

3. A colour forming developer as claimed in claim 2, in which the arematic diamino compound is an alkyl substituted phenylene diamine.

4. A process of colour development which includes developing a reducible silver salt image with an aromatic amino developing agent in presence of a colour

coupler as defined in claim 1.

5. A process of colour development which includes developing a reducible silver salt image with a colour forming 5 developer as claimed in any of claims 1

to 3.

6. A photographic element having a layer containing a clear transparent dye image composed essentially of the product 10 resulting from the coupling in situ, during development of a developable silver salt, of a colour coupler as defined in claim 1 with the oxidation product of an aromatic amino developing agent and 15 subsequent removal of metallic silver.

7. A sensitive photographic element having a colour coupler as defined in claim 1 incorporated in one or more emulsion layers.

8. In the production of colour photographic records, especially multi-layer records, the employment in conjunction with aromatic amino developing agents of colour couplers of the nature herein defined

Dated this 21st day of May, 1937. W. P. THOMPSON & CO., 12. Church Street, Liverpool, Chartered Patent Agents.

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ERRATUM

SPECIFICATION No. 478,934.

In the heading on page 1, for "No. 14356/36." read "No. 14356/37."

THE PATENT OFFICE, March 4th, 1938.