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(54) Titre : COMPOSITIONS DE LESSIVE AVEC UN COLORANT DE THIAZOLIUM
(54) Title: LAUNDRY CARE COMPOSITIONS WITH THIAZOLIUM DYE

(57) **Abrégé/Abstract:**

The present invention relates to thiazolium dyes, laundry care compositions comprising one or more thiazolium dyes, processes of making such dyes and laundry care compositions and methods of using same. The dyes, compositions and methods of the present invention are advantageous in providing improved hueing of fabric, including whitening of white fabric, while avoiding significant build up of bluing dyes on the fabric.



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(54) **Title:** LAUNDRY CARE COMPOSITIONS WITH THIAZOLIUM DYE

(57) **Abstract:** The present invention relates to thiazolium dyes, laundry care compositions comprising one or more thiazolium dyes, processes of making such dyes and laundry care compositions and methods of using same. The dyes, compositions and methods of the present invention are advantageous in providing improved hueing of fabric, including whitening of white fabric, while avoiding significant build up of bluing dyes on the fabric.

WO 2007/087252 A1

LAUNDRY CARE COMPOSITIONS WITH THIAZOLIUM DYE

CROSS-REFERENCES TO RELATED APPLICATIONS

This application claims priority under 35 U.S.C. § 119(e) to U.S. Provisional

5 Application Serial No. 60/761,428 filed January 23, 2006.

FIELD OF THE INVENTION

The present invention relates to thiazolium dyes, laundry care compositions comprising one or more thiazolium dyes, processes of making such dyes and laundry care
10 compositions and methods of using same.

BACKGROUND OF THE INVENTION

Fabrics, typically lighter colored fabrics such as white fabrics, that are worn and/or laundered typically discolor. For example, white fabrics which are repeatedly
15 laundered can exhibit a yellowing in color appearance which causes the fabric to look older and worn. In an effort to overcome such fabric discoloration, certain laundry detergent products include a hueing or bluing dye which attaches to fabric during the laundry wash and/or rinse cycle. Unfortunately, such hueing or bluing dye typically tends to accumulate on the fabric, thus giving the fabric an undesirable bluish tint. As a result,
20 a chlorine treatment is generally employed to reduce the aforementioned accumulation of bluing dyes. While a chlorine treatment can be effective, it is an additional, inconvenient step in the laundry process. Additionally, a chlorine treatment is costly and harsh on fabrics - contributing to increased fabric degradation. Accordingly, a need exists for improved laundry care products which can counter the undesirable discoloration of
25 fabrics, including the yellowing of white fabrics.

SUMMARY OF THE INVENTION

The present invention relates to thiazolium dyes, laundry care compositions comprising one or more thiazolium dyes, processes of making such dyes and laundry care
30 compositions and methods of using same. The dyes, compositions and methods of the present invention are advantageous in providing improved hueing of fabric, including whitening of white fabric, while avoiding significant build up of bluing dyes on the fabric.

DETAILED DESCRIPTION OF THE INVENTION

Definitions

As used herein, the term "laundry care composition" includes, unless otherwise indicated, granular, powder, liquid, gel, paste, bar form and/or flake type washing agents and/or fabric treatment compositions.

As used herein, the term "fabric treatment composition" includes, unless otherwise indicated, fabric softening compositions, fabric enhancing compositions, fabric freshening compositions and combinations thereof. Such compositions may be, but need not be rinse added compositions.

As used herein, the articles including "the", "a" and "an" when used in a claim, are understood to mean one or more of what is claimed or described.

As used herein, the terms "include", "includes" and "including" are meant to be non-limiting.

As used herein, the term polyether is defined as at least two repeating ether units that are chemically bound via the ethers' oxygen atoms. Such polyethers may be derived from materials including but not limited to ethylene oxide, propylene oxide, butylene oxide, hexylene oxide, glycidol, epichlorohydrin, pentanerythritol, glucose or combinations thereof.

As used herein capped polyether means a polyether that terminates in an alkyl or aryl moiety, including but not limited to a moiety selected from methyl, ethyl, butyl, isopropyl, tertiary butyl, amyl, benzyl, pentyl, and acetyl moieties.

As used herein "EO" stands for an ethylene oxide moiety.

As used herein "PO" stands for a propylene oxide moiety.

The test methods disclosed in the Test Methods Section of the present application should be used to determine the respective values of the parameters of Applicants' inventions.

Unless otherwise noted, all component or composition levels are in reference to the active portion of that component or composition, and are exclusive of impurities, for example, residual solvents or by-products, which may be present in commercially available sources of such components or compositions.

All percentages and ratios are calculated by weight unless otherwise indicated. All percentages and ratios are calculated based on the total composition unless otherwise indicated.

It should be understood that every maximum numerical limitation given throughout this specification includes every lower numerical limitation, as if such lower numerical limitations were expressly written herein. Every minimum numerical limitation given throughout this specification will include every higher numerical limitation, as if such higher numerical limitations were expressly written herein. Every numerical range given throughout this specification will include every narrower numerical range that falls within such broader numerical range, as if such narrower numerical ranges were all expressly written herein.

The citation of any document is not to be construed as an admission that it is prior art with respect to the present invention.

Laundry Care Compositions

In one aspect, a laundry care composition that may comprise a laundry care ingredient and a suitable thiazolium dye is disclosed. Suitable thiazolium dyes include thiazolium dyes that exhibit good tinting efficiency during a laundry wash cycle without exhibiting excessive undesirable build up after laundering. Thus, undesirable bluing after repeated washings with the detergent compositions of the invention is avoided and costly and harsh chlorine treatments are unnecessary. Suitable thiazolium dyes include those thiazolium dyes that are described under the heading "Suitable Thiazolium Dyes" of the present specification.

In one aspect, the laundry care compositions disclosed in the present specification can employ the thiazolium dyes disclosed in the present specification as detailed by Formulae V through VIII of the present specification.

In one aspect suitable thiazolium dyes include thiazolium dye molecules numbers 1- 80 as detailed in Tables 1 and 2 of the present specification.

In one aspect, suitable thiazolium dyes include thiazolium dye molecules numbers 1, 4, 5, 7, 8, 12, 13, 15, 16, 17, 21, 24, 25, 26, 30, 31, 33, 36, 38, 40, 45 and 48 as detailed in Tables 1 and 2 of the present specification.

In one aspect, suitable thiazolium dyes include thiazolium dye molecules numbers 12, 13, 15, 16, 24, 25, 26, 30, 31, 33, 36, 38, 40, 45 and 48 as detailed in Tables 1 and 2 of the present specification.

In one aspect, the laundry care compositions disclosed in the present specification can employ combinations of any of the suitable thiazolium dyes disclosed in the present specification.

In one aspect, the laundry care compositions disclosed in the present specification can employ a non-hueing dye in combination with the thiazolium dye. The non-hueing dye may be selected from non-hueing dyes disclosed in U.S. Patent Application 2005/028820 A1, U.S. Patent 4,137,243, U.S. Patent 4,601,725 and U.S. Patent 4,871,371. While not being bound by theory, it is believed that the combination of both a thiazolium dye and a non-hueing dye allows for flexibility to color blend to a desired hue.

In one aspect, the laundry care compositions disclosed in the present specification can employ a non-hueing dye, that may be non-substantive in nature, in combination with the thiazolium dye. The combination of both a thiazolium dye and a non-hueing dye can allow customization of product color and fabric tint. In one aspect, Acid Blue 7 may be employed as a non-hueing, non-tinting dye.

In one aspect, any of the components, including the suitable thiazolium dyes, may be employed in the laundry care compositions in an encapsulated form. Such encapsulates may comprise one or more of such components.

In one aspect a laundry care compositions comprising a thiazolium dye and a laundry care ingredient and having a hueing efficiency of greater than 10 but less than 40, from about 15 to about 35, or even from about 15 to about 30 and a wash removability of from about 30% to about 85%, from about 40% to about 85%, from about 50% to about 85% are disclosed.

Suitable laundry care ingredients include, but are not limited to, those materials described in the present specification as useful aspects of the present invention, including adjunct materials as described in the present specification.

Liquid, Laundry Detergent Compositions

In one aspect, the laundry care compositions disclosed herein, may take the form of liquid, laundry detergent compositions. In one aspect, such compositions may be a heavy duty liquid composition. Such compositions may comprise a sufficient amount of a surfactant to provide the desired level of one or more cleaning properties, typically by weight of the total composition, from about 5% to about 90%, from about 5% to about 70% or even from about 5% to about 40% and a sufficient of suitable thiazolium dye that is described under the heading "Suitable Thiazolium Dyes" of the present specification, to

provide a tinting effect to fabric washed in a solution containing the detergent, typically by weight of the total composition, from about 0.0001% to about 0.05%, or even from about 0.001% to about 0.01%.

The liquid detergent compositions comprise an aqueous, non-surface active liquid carrier. Generally, the amount of the aqueous, non-surface active liquid carrier employed in the compositions herein will be effective to solubilize, suspend or disperse the composition components. For example, the compositions may comprise, by weight, from about 5% to about 90%, from about 10% to about 70%, or even from about 20% to about 70% of an aqueous, non-surface active liquid carrier.

The most cost effective type of aqueous, non-surface active liquid carrier may be water. Accordingly, the aqueous, non-surface active liquid carrier component may be generally mostly, if not completely, water. While other types of water-miscible liquids, such alkanols, diols, other polyols, ethers, amines, and the like, have been conventionally been added to liquid detergent compositions as co-solvents or stabilizers, for purposes of the present invention, the utilization of such water-miscible liquids may be minimized to hold down composition cost. Accordingly, the aqueous liquid carrier component of the liquid detergent products herein will generally comprise water present in concentrations ranging from about 5% to about 90%, or even from about 20% to about 70%, by weight of the composition.

The liquid detergent compositions herein may take the form of an aqueous solution or uniform dispersion or suspension of surfactant, thiazolium dye, and certain optional other ingredients, some of which may normally be in solid form, that have been combined with the normally liquid components of the composition, such as the liquid alcohol ethoxylate nonionic, the aqueous liquid carrier, and any other normally liquid optional ingredients. Such a solution, dispersion or suspension will be acceptably phase stable and will typically have a viscosity which ranges from about 100 to 600 cps, more preferably from about 150 to 400 cps. For purposes of this invention, viscosity is measured with a Brookfield LVDV-II+ viscometer apparatus using a #21 spindle.

Suitable surfactants may be anionic, nonionic, cationic, zwitterionic and/or amphoteric surfactants. In one aspect, the detergent composition comprises anionic surfactant, nonionic surfactant, or mixtures thereof.

Suitable anionic surfactants may be any of the conventional anionic surfactant types typically used in liquid detergent products. Such surfactants include the alkyl

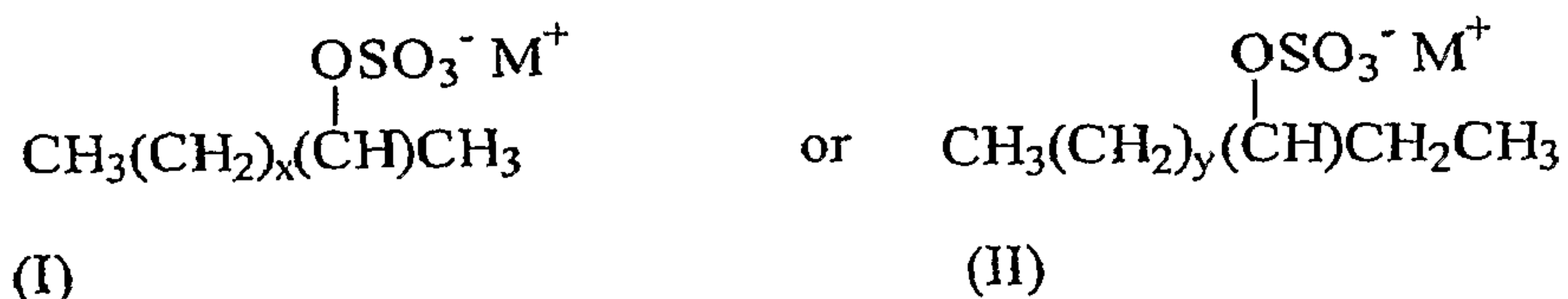
benzene sulfonic acids and their salts as well as alkoxyated or non-alkoxyated alkyl sulfate materials.

Exemplary anionic surfactants are the alkali metal salts of C₁₀₋₁₆ alkyl benzene sulfonic acids, preferably C₁₁₋₁₄ alkyl benzene sulfonic acids. In one aspect, the alkyl group is linear. Such linear alkyl benzene sulfonates are known as "LAS". Such surfactants and their preparation are described for example in U.S. Patents 2,220,099 and 2,477,383. Especially preferred are the sodium and potassium linear straight chain alkylbenzene sulfonates in which the average number of carbon atoms in the alkyl group is from about 11 to 14. Sodium C₁₁-C₁₄, e.g., C₁₂, LAS is a specific example of such surfactants.

Another exemplary type of anionic surfactant comprises ethoxylated alkyl sulfate surfactants. Such materials, also known as alkyl ether sulfates or alkyl polyethoxylate sulfates, are those which correspond to the formula: R'-O-(C₂H₄O)_n-SO₃M wherein R' is a C₈-C₂₀ alkyl group, n is from about 1 to 20, and M is a salt-forming cation. In a specific embodiment, R' is C₁₀-C₁₈ alkyl, n is from about 1 to 15, and M is sodium, potassium, ammonium, alkylammonium, or alkanolammonium. In more specific embodiments, R' is a C₁₂-C₁₆, n is from about 1 to 6 and M is sodium.

The alkyl ether sulfates will generally be used in the form of mixtures comprising varying R' chain lengths and varying degrees of ethoxylation. Frequently such mixtures will inevitably also contain some non-ethoxylated alkyl sulfate materials, i.e., surfactants of the above ethoxylated alkyl sulfate formula wherein n=0. Non-ethoxylated alkyl sulfates may also be added separately to the compositions of this invention and used as or in any anionic surfactant component which may be present. Specific examples of non-alkoxyated, e.g., non-ethoxylated, alkyl ether sulfate surfactants are those produced by the sulfation of higher C₈-C₂₀ fatty alcohols. Conventional primary alkyl sulfate surfactants have the general formula: ROSO₃⁻M⁺ wherein R is typically a linear C₈-C₂₀ hydrocarbyl group, which may be straight chain or branched chain, and M is a water-solubilizing cation. In specific embodiments, R is a C₁₀-C₁₅ alkyl, and M is alkali metal, more specifically R is C₁₂-C₁₄ and M is sodium.

Specific, nonlimiting examples of anionic surfactants useful herein include: a) C₁₁-C₁₈ alkyl benzene sulfonates (LAS); b) C₁₀-C₂₀ primary, branched-chain and random alkyl sulfates (AS); c) C₁₀-C₁₈ secondary (2,3) alkyl sulfates having formulae (I) and (II):



wherein M in formulae (I) and (II) is hydrogen or a cation which provides charge neutrality, and all M units, whether associated with a surfactant or adjunct ingredient, can either be a hydrogen atom or a cation depending upon the form isolated by the artisan or the relative pH of the system wherein the compound is used, with non-limiting examples of preferred cations including sodium, potassium, ammonium, and mixtures thereof, and x is an integer of at least about 7, preferably at least about 9, and y is an integer of at least 8, preferably at least about 9; d) C₁₀-C₁₈ alkyl alkoxy sulfates (AE_xS) wherein preferably x is from 1-30; e) C₁₀-C₁₈ alkyl alkoxy carboxylates preferably comprising 1-5 ethoxy units; f) mid-chain branched alkyl sulfates as discussed in US 6,020,303 and US 6,060,443; g) mid-chain branched alkyl alkoxy sulfates as discussed in US 6,008,181 and US 6,020,303; h) modified alkylbenzene sulfonate (MLAS) as discussed in WO 99/05243, WO 99/05242, WO 99/05244, WO 99/05082, WO 99/05084, WO 99/05241, WO 99/07656, WO 00/23549, and WO 00/23548.; i) methyl ester sulfonate (MES); and j) alpha-olefin sulfonate (AOS).

Suitable nonionic surfactants useful herein can comprise any of the conventional nonionic surfactant types typically used in liquid detergent products. These include alkoxylated fatty alcohols and amine oxide surfactants. Preferred for use in the liquid detergent products herein are those nonionic surfactants which are normally liquid.

Suitable nonionic surfactants for use herein include the alcohol alkoxyate nonionic surfactants. Alcohol alkoxyates are materials which correspond to the general formula: R¹(C_mH_{2m}O)_nOH wherein R¹ is a C₈ - C₁₆ alkyl group, m is from 2 to 4, and n ranges from about 2 to 12. Preferably R¹ is an alkyl group, which may be primary or secondary, that contains from about 9 to 15 carbon atoms, more preferably from about 10 to 14 carbon atoms. In one embodiment, the alkoxylated fatty alcohols will also be ethoxylated materials that contain from about 2 to 12 ethylene oxide moieties per molecule, more preferably from about 3 to 10 ethylene oxide moieties per molecule.

The alkoxylated fatty alcohol materials useful in the liquid detergent compositions herein will frequently have a hydrophilic-lipophilic balance (HLB) which ranges from about 3 to 17. More preferably, the HLB of this material will range from about 6 to 15,

most preferably from about 8 to 15. Alkoxylated fatty alcohol nonionic surfactants have been marketed under the tradename Neodol® by the Shell Chemical Company.

Another suitable type of nonionic surfactant useful herein comprises the amine oxide surfactants. Amine oxides are materials which are often referred to in the art as “semi-polar” nonionics. Amine oxides have the formula:
 $R(EO)_x(PO)_y(BO)_zN(O)(CH_2R')_2 \cdot qH_2O$. In this formula, R is a relatively long-chain hydrocarbyl moiety which can be saturated or unsaturated, linear or branched, and can contain from 8 to 20, preferably from 10 to 16 carbon atoms, and is more preferably C₁₂-C₁₆ primary alkyl. R' is a short-chain moiety, preferably selected from hydrogen, methyl and -CH₂OH. When x+y+z is different from 0, EO is ethyleneoxy, PO is propyleneoxy and BO is butyleneoxy. Amine oxide surfactants are illustrated by C₁₂₋₁₄ alkyldimethyl amine oxide.

Non-limiting examples of nonionic surfactants include: a) C₁₂-C₁₈ alkyl ethoxylates, such as, NEODOL® nonionic surfactants from Shell; b) C₆-C₁₂ alkyl phenol alkoxyates wherein the alkoxyate units are a mixture of ethyleneoxy and propyleneoxy units; c) C₁₂-C₁₈ alcohol and C₆-C₁₂ alkyl phenol condensates with ethylene oxide/propylene oxide block polymers such as Pluronic® from BASF; d) C₁₄-C₂₂ mid-chain branched alcohols, BA, as discussed in US 6,150,322; e) C₁₄-C₂₂ mid-chain branched alkyl alkoxyates, BAE_x, wherein x 1-30, as discussed in US 6,153,577, US 6,020,303 and US 6,093,856; f) Alkylpolysaccharides as discussed in U.S. 4,565,647 Llenado, issued January 26, 1986; specifically alkylpolyglycosides as discussed in US 4,483,780 and US 4,483,779; g) Polyhydroxy fatty acid amides as discussed in US 5,332,528, WO 92/06162, WO 93/19146, WO 93/19038, and WO 94/09099; and h) ether capped poly(oxyalkylated) alcohol surfactants as discussed in US 6,482,994 and WO 01/42408.

In the laundry detergent compositions herein, the deterative surfactant component may comprise combinations of anionic and nonionic surfactant materials. When this is the case, the weight ratio of anionic to nonionic will typically range from 10:90 to 90:10, more typically from 30:70 to 70:30.

Cationic surfactants are well known in the art and non-limiting examples of these include quaternary ammonium surfactants, which can have up to 26 carbon atoms. Additional examples include a) alkoxyate quaternary ammonium (AQA) surfactants as discussed in US 6,136,769; b) dimethyl hydroxyethyl quaternary ammonium as discussed in US 6,004,922; c) polyamine cationic surfactants as discussed in WO 98/35002, WO

98/35003, WO 98/35004, WO 98/35005, and WO 98/35006; d) cationic ester surfactants as discussed in US Patents Nos. 4,228,042, 4,239,660 4,260,529 and US 6,022,844; and e) amino surfactants as discussed in US 6,221,825 and WO 00/47708, specifically amido propyldimethyl amine (APA).

5 Non-limiting examples of zwitterionic surfactants include: derivatives of secondary and tertiary amines, derivatives of heterocyclic secondary and tertiary amines, or derivatives of quaternary ammonium, quaternary phosphonium or tertiary sulfonium compounds. See U.S. Patent No. 3,929,678 to Laughlin et al., issued December 30, 1975 at column 19, line 38 through column 22, line 48, for examples of zwitterionic
10 surfactants; betaine, including alkyl dimethyl betaine and cocodimethyl amidopropyl betaine, C₈ to C₁₈ (preferably C₁₂ to C₁₈) amine oxides and sulfo and hydroxy betaines, such as N-alkyl-N,N-dimethylammino-1-propane sulfonate where the alkyl group can be C₈ to C₁₈, preferably C₁₀ to C₁₄.

Non-limiting examples of ampholytic surfactants include: aliphatic derivatives of
15 secondary or tertiary amines, or aliphatic derivatives of heterocyclic secondary and tertiary amines in which the aliphatic radical can be straight- or branched-chain. One of the aliphatic substituents contains at least about 8 carbon atoms, typically from about 8 to about 18 carbon atoms, and at least one contains an anionic water-solubilizing group, e.g. carboxy, sulfonate, sulfate. See U.S. Patent No. 3,929,678 to Laughlin et al., issued
20 December 30, 1975 at column 19, lines 18-35, for examples of ampholytic surfactants.

Granular Laundry Detergent Compositions

In one aspect, the laundry care compositions disclosed herein, may take the form of granular, laundry detergent compositions. Such compositions may comprise a
25 sufficient of suitable thiazolium dye that is described under the heading "Suitable Thiazolium Dyes" of the present specification, to provide a tinting effect to fabric washed in a solution containing the detergent, typically by weight of the total composition, from about 0.0001% to about 0.05%, or even from about 0.001% to about 0.01%.

Granular detergent compositions of the present invention may include any number
30 of conventional detergent ingredients. For example, the surfactant system of the detergent composition may include anionic, nonionic, zwitterionic, ampholytic and cationic classes and compatible mixtures thereof. Detergent surfactants for granular compositions are described in U.S. Patent 3,664,961, Norris, issued May 23, 1972, and in U.S. Patent 3,919,678, Laughlin et al., issued December 30, 1975. Cationic surfactants

include those described in U.S. Patent 4,222,905, Cockrell, issued September 16, 1980, and in U.S. Patent 4,239,659, Murphy, issued December 16, 1980.

Nonlimiting examples of surfactant systems include the conventional C₁₁-C₁₈ alkyl benzene sulfonates ("LAS") and primary, branched-chain and random C₁₀-C₂₀ alkyl sulfates ("AS"), the C₁₀-C₁₈ secondary (2,3) alkyl sulfates of the formula
5 $\text{CH}_3(\text{CH}_2)_x(\text{CHOSO}_3^- \text{M}^+) \text{CH}_3$ and $\text{CH}_3(\text{CH}_2)_y(\text{CHOSO}_3^- \text{M}^+) \text{CH}_2\text{CH}_3$ where x and (y + 1) are integers of at least about 7, preferably at least about 9, and M is a water-solubilizing cation, especially sodium, unsaturated sulfates such as oleyl sulfate, the C₁₀-C₁₈ alkyl alkoxy sulfates ("AE_xS"; especially EO 1-7 ethoxy sulfates), C₁₀-C₁₈ alkyl
10 alkoxy carboxylates (especially the EO 1-5 ethoxycarboxylates), the C₁₀-C₁₈ glycerol ethers, the C₁₀-C₁₈ alkyl polyglycosides and their corresponding sulfated polyglycosides, and C₁₂-C₁₈ alpha-sulfonated fatty acid esters. If desired, the conventional nonionic and amphoteric surfactants such as the C₁₂-C₁₈ alkyl ethoxylates ("AE") including the so-called narrow peaked alkyl ethoxylates and C₆-C₁₂ alkyl phenol alkoxyates (especially
15 ethoxylates and mixed ethoxy/propoxy), C₁₂-C₁₈ betaines and sulfobetaines ("sultaines"), C₁₀-C₁₈ amine oxides, and the like, can also be included in the surfactant system. The C₁₀-C₁₈ N-alkyl polyhydroxy fatty acid amides can also be used. See WO 9,206,154. Other sugar-derived surfactants include the N-alkoxy polyhydroxy fatty acid amides, such as C₁₀-C₁₈ N-(3-methoxypropyl) glucamide. The N-propyl through N-hexyl C₁₂-C₁₈
20 glucamides can be used for low sudsing. C₁₀-C₂₀ conventional soaps may also be used. If high sudsing is desired, the branched-chain C₁₀-C₁₆ soaps may be used. Mixtures of anionic and nonionic surfactants are especially useful. Other conventional useful surfactants are listed in standard texts.

The detergent composition can, and preferably does, include a detergent builder.
25 Builders are generally selected from the various water-soluble, alkali metal, ammonium or substituted ammonium phosphates, polyphosphates, phosphonates, polyphosphonates, carbonates, silicates, borates, polyhydroxy sulfonates, polyacetates, carboxylates, and polycarboxylates. Preferred are the alkali metal, especially sodium, salts of the above. Preferred for use herein are the phosphates, carbonates, silicates, C₁₀-C₁₈ fatty acids,
30 polycarboxylates, and mixtures thereof. More preferred are sodium tripolyphosphate, tetrasodium pyrophosphate, citrate, tartrate mono- and di-succinates, sodium silicate, and mixtures thereof.

11

Specific examples of inorganic phosphate builders are sodium and potassium tripolyphosphate, pyrophosphate, polymeric metaphosphate having a degree of polymerization of from about 6 to 21, and orthophosphates. Examples of polyphosphonate builders are the sodium and potassium salts of ethylene diphosphonic acid, the sodium and potassium salts of ethane 1-hydroxy-1, 1-diphosphonic acid and the sodium and potassium salts of ethane, 1,1,2-triphosphonic acid. Other phosphorus builder compounds are disclosed in U.S. Patents 3,159,581; 3,213,030; 3,422,021; 3,422,137; 3,400,176 and 3,400,148. Examples of nonphosphorus, inorganic builders are sodium and potassium carbonate, bicarbonate, sesquicarbonate, tetraborate decahydrate, and silicates having a weight ratio of SiO_2 to alkali metal oxide of from about 0.5 to about 4.0, preferably from about 1.0 to about 2.4. Water-soluble, nonphosphorus organic builders useful herein include the various alkali metal, ammonium and substituted ammonium polyacetates, carboxylates, polycarboxylates and polyhydroxy sulfonates. Examples of polyacetate and polycarboxylate builders are the sodium, potassium, lithium, ammonium and substituted ammonium salts of ethylene diamine tetraacetic acid, nitrilotriacetic acid, oxydisuccinic acid, mellitic acid, benzene polycarboxylic acids, and citric acid.

Polymeric polycarboxylate builders are set forth in U.S. Patent 3,308,067, Diehl, issued March 7, 1967. Such materials include the water-soluble salts of homo- and copolymers of aliphatic carboxylic acids such as maleic acid, itaconic acid, mesaconic acid, fumaric acid, aconitic acid, citraconic acid and methylenemalonic acid. Some of these materials are useful as the water-soluble anionic polymer as hereinafter described, but only if in intimate admixture with the nonsoap anionic surfactant. Other suitable polycarboxylates for use herein are the polyacetal carboxylates described in U.S. Patent 4,144,226, issued March 13, 1979 to Crutchfield et al., and U.S. Patent 4,246,495, issued March 27, 1979 to Crutchfield et al.

Water-soluble silicate solids represented by the formula $\text{SiO}_2 \cdot \text{M}_2\text{O}$, M being an alkali metal, and having a $\text{SiO}_2:\text{M}_2\text{O}$ weight ratio of from about 0.5 to about 4.0, are useful salts in the detergent granules of the invention at levels of from about 2% to about 15% on an anhydrous weight basis. Anhydrous or hydrated particulate silicate can be utilized, as well.

Any number of additional ingredients can also be included as components in the granular detergent composition. These include other detergency builders, bleaches,

bleach activators, suds boosters or suds suppressors, anti-tarnish and anti-corrosion agents, soil suspending agents, soil release agents, germicides, pH adjusting agents, nonbuilder alkalinity sources, chelating agents, smectite clays, enzymes, enzyme-stabilizing agents and perfumes. See U.S. Patent 3,936,537, issued February 3, 1976 to Baskerville, Jr. et al.

Bleaching agents and activators are described in U.S. Patent 4,412,934, Chung et al., issued November 1, 1983, and in U.S. Patent 4,483,781, Hartman, issued November 20, 1984. Chelating agents are also described in U.S. Patent 4,663,071, Bush et al., from Column 17, line 54 through Column 18, line 68. Suds modifiers are also optional ingredients and are described in U.S. Patents 3,933,672, issued January 20, 1976 to Bartoletta et al., and 4,136,045, issued January 23, 1979 to Gault et al. Suitable smectite clays for use herein are described in U.S. Patent 4,762,645, Tucker et al., issued August 9, 1988, Column 6, line 3 through Column 7, line 24. Suitable additional detergency builders for use herein are enumerated in the Baskerville patent, Column 13, line 54 through Column 16, line 16, and in U.S. Patent 4,663,071, Bush et al., issued May 5, 1987.

Rinse Added Fabric Conditioning Compositions

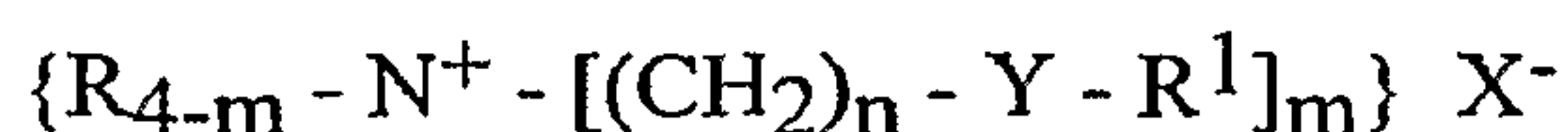
In one aspect, the laundry care compositions disclosed herein, may take the form of rinse added fabric conditioning compositions. Such compositions may comprise a fabric softening active and a sufficient amount of suitable thiazolium dye, that is described under the heading "Suitable Thiazolium Dyes" of the present specification, to provide a tinting effect to fabric treated by the composition, typically from about 0.00001 wt. % (0.1 ppm) to about 1 wt. % (10,000 ppm), or even from about 0.0003 wt. % (3 ppm) to about 0.03 wt. % (300 ppm) based on total rinse added fabric conditioning composition weight. In another specific embodiment, the compositions are rinse added fabric conditioning compositions. Examples of typical rinse added conditioning composition can be found in U.S. Provisional Patent Application Serial No. 60/687582 filed on October 8, 2004.

In one embodiment of the invention, the fabric softening active (hereinafter "FSA") is a quaternary ammonium compound suitable for softening fabric in a rinse step. In one embodiment, the FSA is formed from a reaction product of a fatty acid and an aminoalcohol obtaining mixtures of mono-, di-, and, in one embodiment, triester compounds. In another embodiment, the FSA comprises one or more softener quaternary

ammonium compounds such, but not limited to, as a monoalkylquaternary ammonium compound, a diamido quaternary compound and a diester quaternary ammonium compound, or a combination thereof.

In one aspect of the invention, the FSA comprises a diester quaternary ammonium (hereinafter "DQA") compound composition. In certain embodiments of the present invention, the DQA compounds compositions also encompasses a description of diamido FSAs and FSAs with mixed amido and ester linkages as well as the aforementioned diester linkages, all herein referred to as DQA.

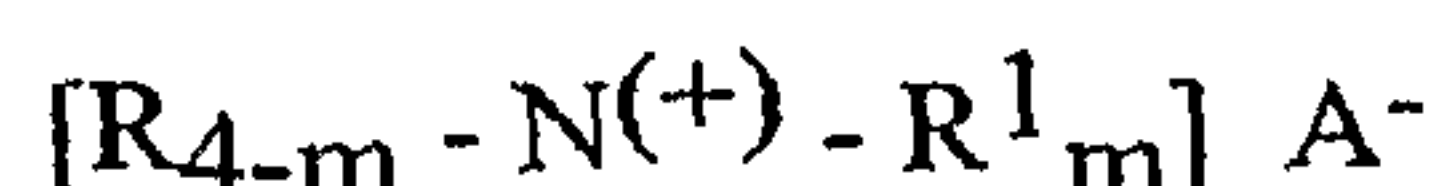
A first type of DQA ("DQA (1)") suitable as a FSA in the present CFSC includes a compound comprising the formula:



wherein each R substituent is either hydrogen, a short chain C₁-C₆, preferably C₁-C₃ alkyl or hydroxyalkyl group, e.g., methyl (most preferred), ethyl, propyl, hydroxyethyl, and the like, poly (C₂-C₃ alkoxy), preferably polyethoxy, group, benzyl, or mixtures thereof; each m is 2 or 3; each n is from 1 to about 4, preferably 2; each Y is -O-(O)C-, -C(O)-O-, -NR-C(O)-, or -C(O)-NR- and it is acceptable for each Y to be the same or different; the sum of carbons in each R¹, plus one when Y is -O-(O)C- or -NR-C(O)-, is C₁₂-C₂₂, preferably C₁₄-C₂₀, with each R¹ being a hydrocarbyl, or substituted hydrocarbyl group; it is acceptable for R¹ to be unsaturated or saturated and branched or linear and preferably it is linear; it is acceptable for each R¹ to be the same or different and preferably these are the same; and X⁻ can be any softener-compatible anion, preferably, chloride, bromide, methylsulfate, ethylsulfate, sulfate, phosphate, and nitrate, more preferably chloride or methyl sulfate. Preferred DQA compounds are typically made by reacting alkanolamines such as MDEA (methyldiethanolamine) and TEA (triethanolamine) with fatty acids. Some materials that typically result from such reactions include N,N-di(acyl-oxyethyl)-N,N-dimethylammonium chloride or N,N-di(acyl-oxyethyl)-N,N-methylhydroxyethylammonium methylsulfate wherein the acyl group is derived from animal fats, unsaturated, and polyunsaturated, fatty acids, e.g., tallow, hardened tallow, oleic acid, and/or partially hydrogenated fatty acids, derived from vegetable oils and/or partially hydrogenated vegetable oils, such as, canola oil, safflower oil, peanut oil, sunflower oil, corn oil, soybean oil, tall oil, rice bran oil, palm oil, etc. Non-limiting examples of suitable fatty acids are listed in US 5,759,990 at column 4, lines 45-66. In

one embodiment the FSA comprises other actives in addition to DQA (1) or DQA. In yet another embodiment, the FSA comprises only DQA (1) or DQA and is free or essentially free of any other quaternary ammonium compounds or other actives. In yet another embodiment, the FSA comprises the precursor amine that is used to produce the DQA.

5 In another aspect of the invention, the FSA comprises a compound, identified as DTTMAC comprising the formula:



wherein each m is 2 or 3, each R^1 is a C_6 - C_{22} , preferably C_{14} - C_{20} , but no more than one being less than about C_{12} and then the other is at least about 16, hydrocarbyl, or substituted hydrocarbyl substituent, preferably C_{10} - C_{20} alkyl or alkenyl (unsaturated alkyl, including polyunsaturated alkyl, also referred to sometimes as "alkylene"), most preferably C_{12} - C_{18} alkyl or alkenyl, and branch or unbranched. In one embodiment, the Iodine Value (IV) of the FSA is from about 1 to 70; each R is H or a short chain C_1 - C_6 , preferably C_1 - C_3 alkyl or hydroxyalkyl group, e.g., methyl (most preferred), ethyl, propyl, hydroxyethyl, and the like, benzyl, or $(R^2 O)_{2-4}H$ where each R^2 is a C_1 - C_6 alkylene group; and A^{-} is a softener compatible anion, preferably, chloride, bromide, methylsulfate, ethylsulfate, sulfate, phosphate, or nitrate; more preferably chloride or methyl sulfate. Examples of these FSAs include dialkyldimethylammonium salts and dialkylenedimethylammonium salts such as ditallowdimethylammonium and ditallowdimethylammonium methylsulfate. Examples of commercially available dialkylenedimethylammonium salts usable in the present invention are di-hydrogenated tallow dimethyl ammonium chloride and ditallowdimethyl ammonium chloride available from Degussa under the trade names Adogen[®] 442 and Adogen[®] 470 respectively. In one embodiment the FSA comprises other actives in addition to DTTMAC. In yet another embodiment, the FSA comprises only compounds of the DTTMAC and is free or essentially free of any other quaternary ammonium compounds or other actives.

In one embodiment, the FSA comprises an FSA described in U.S. Pat. Pub. No. 2004/0204337 A1, published Oct. 14, 2004 to Corona et al., from paragraphs 30 – 79.

In another embodiment, the FSA is one described in U.S. Pat. Pub. No. 2004/0229769 A1, published Nov. 18, 2005, to Smith et al., on paragraphs 26 – 31; or U.S. Pat. No. 6,494,920, at column 1, line 51 *et seq.* detailing an "esterquat" or a quaternized fatty acid triethanolamine ester salt.

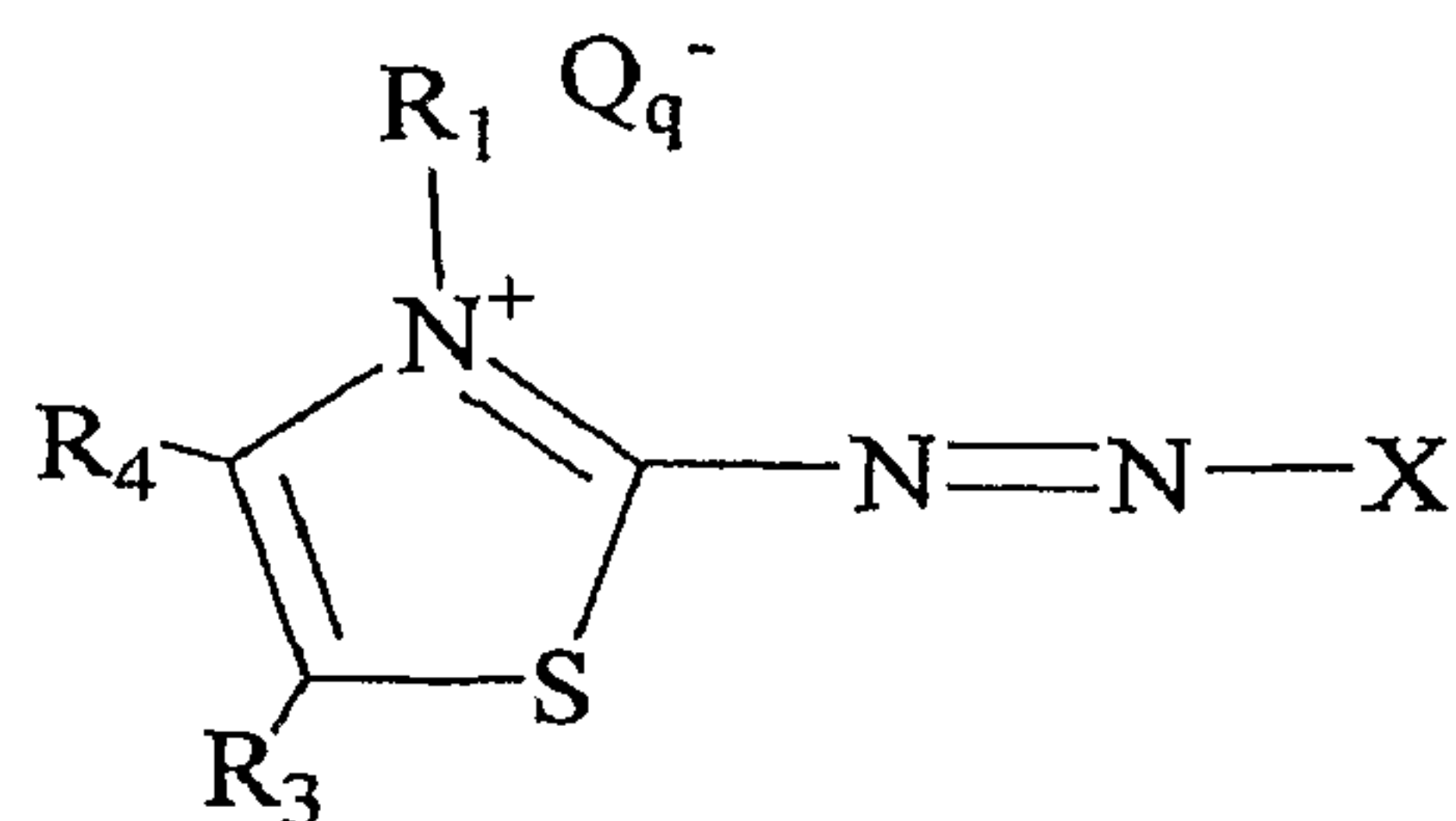
In one embodiment, the FSA is chosen from at least one of the following: ditallowoyloxyethyl dimethyl ammonium chloride, dihydrogenated-tallowoyloxyethyl dimethyl ammonium chloride, ditallow dimethyl ammonium chloride, ditallowoyloxyethyl dimethyl ammonium methyl sulfate, dihydrogenated-tallowoyloxyethyl dimethyl ammonium chloride, dihydrogenated-tallowoyloxyethyl dimethyl ammonium chloride, or combinations thereof.

In one embodiment, the FSA may also include amide containing compound compositions. Examples of diamide comprising compounds may include but not limited to methyl-bis(tallowamidoethyl)-2-hydroxyethylammonium methyl sulfate (available from Degussa under the trade names Varisoft 110 and Varisoft 222). An example of an amide-ester containing compound is N-[3-(stearoylamino)propyl]-N-[2-(stearoyloxy)ethoxy)ethyl]]-N-methylamine.

Another specific embodiment of the invention provides for a rinse added fabric care composition further comprising a cationic starch. Cationic starches are disclosed in US 2004/0204337 A1. In one embodiment, the fabric care composition comprises from about 0.1% to about 7% of cationic starch by weight of the laundry care composition. In one embodiment, the cationic starch is HCP401 from National Starch.

Suitable Thiazolium Dyes

Suitable thiazolium dyes include azo dyes that may have Formula (I) below:



Formula I

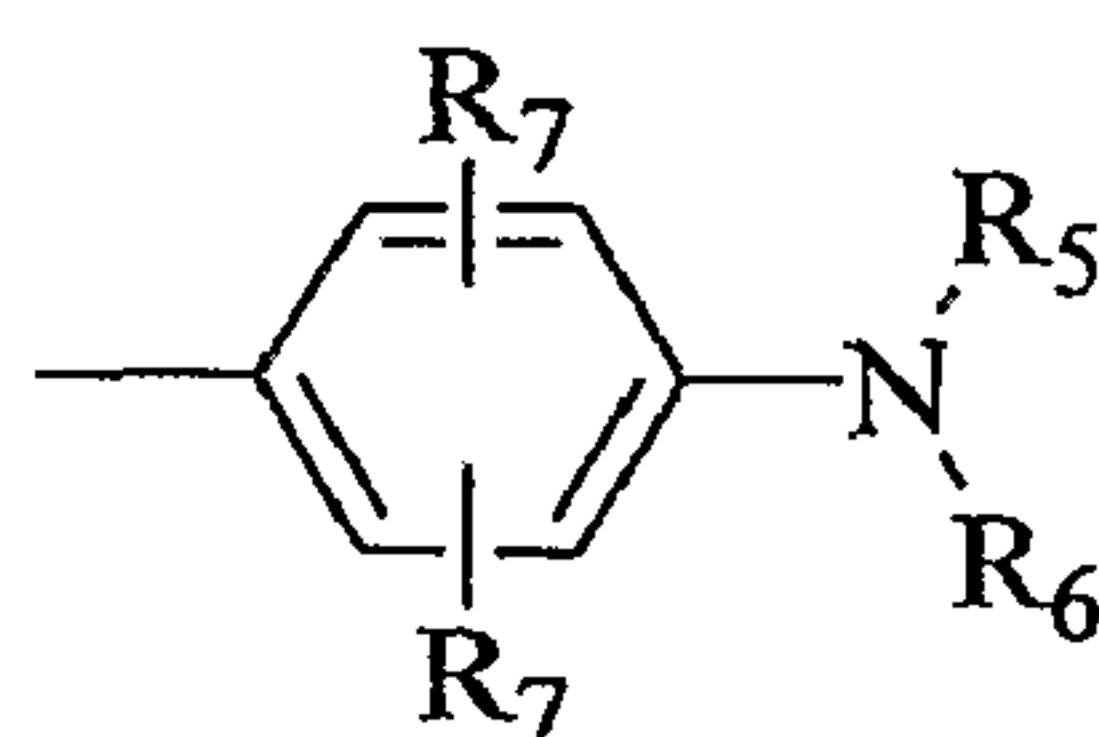
wherein:

R₃ and R₄ may be identical or different and, independently of one another, are hydrogen, a saturated or unsaturated (C₁-C₂₂)-alkyl group, a (C₁-C₂₂)-alkyl group substituted by a halogen atom, a hydroxy-(C₂-C₂₂)-alkyl group optionally interrupted by oxygen, a polyether group derived from ethylene oxide, propylene oxide or butylene oxide, an amino-(C₁-C₂₂)-alkyl group,

16

a substituted or unsubstituted phenyl group or a benzyl group, a (C₁-C₂₂)-alkyl group terminated in sulfonate, sulfate, or carboxylate, or the radical groups R₃ and R₄, together with the remaining molecule, can form a heterocyclic or carbocyclic, saturated or unsaturated, substituted or unsubstituted ring system optionally substituted by halogen, sulfate, sulfonate, phosphate, nitrate, and carboxylate;

X may be a radical group of the phenol series or a heterocyclic radical group or aniline series or m-toluidine series that may have Formula II below;



Formula II

wherein:

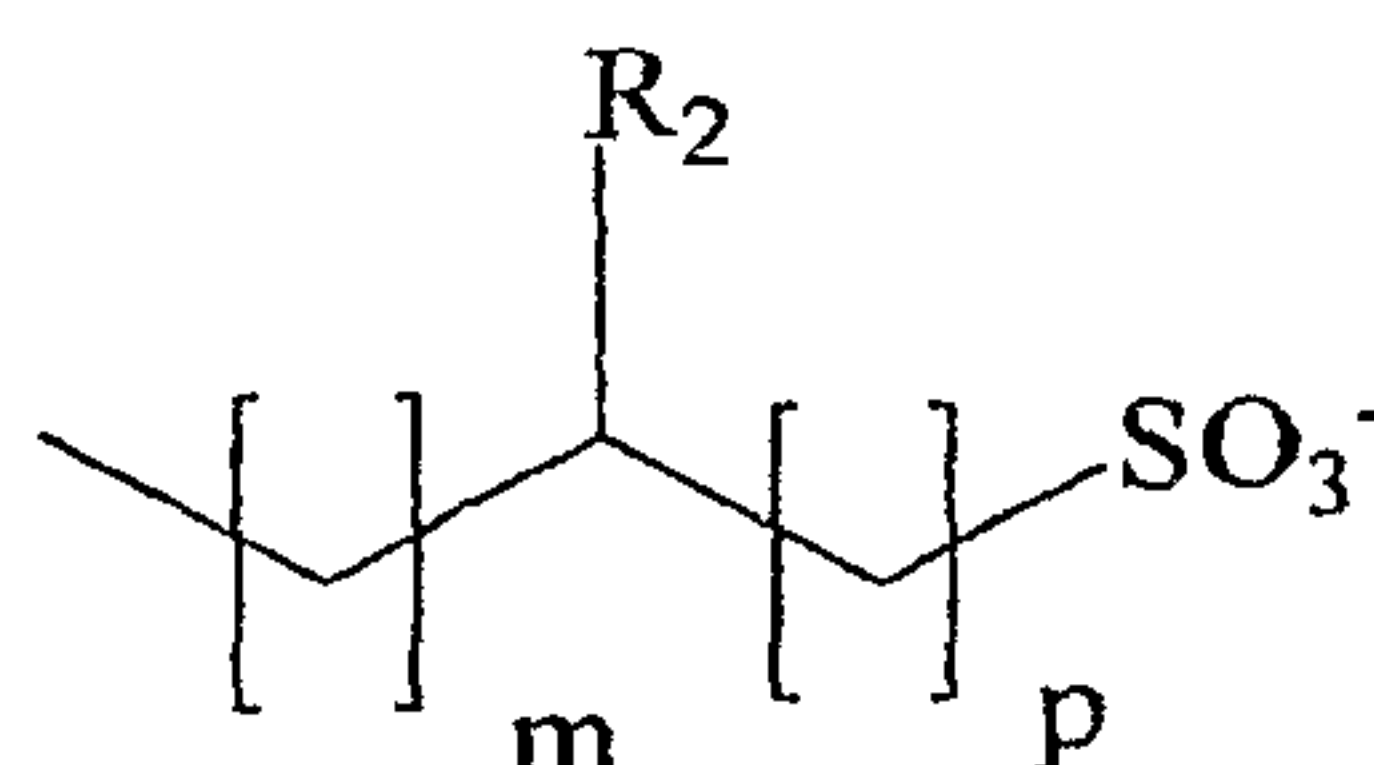
R₅ and R₆ may be identical or different and, independently of one another, are a straight or branched saturated or unsaturated (C₁-C₂₂)-alkyl group, a (C₁-C₂₂)-alkyl ether group, a hydroxy-(C₂-C₂₂)-alkyl group optionally interrupted by oxygen, a polyether group derived from ethylene oxide, propylene oxide, butylene oxide, glycidyl or combinations thereof, an amino-(C₁-C₂₂)-alkyl group, a substituted or unsubstituted phenyl group or a benzyl group, a linear or branched (C₁-C₂₂)-alkyl group terminated in a linear or branched (C₁-C₂₂)-alkyl, hydroxyl, acetate, sulfonate, sulfate, or carboxylate, group or R₅ and R₆ or R₅ and R₇ or R₆ and R₇, together with the nitrogen atom, form a 5-membered to 6-membered ring system, which may comprise a further heteroatom; or R₅ and R₆ or R₅ and R₇ or R₆ and R₇, form with a carbon atom of the benzene ring an optionally oxygen-containing or nitrogen containing five or six-membered heterocycle which may be substituted with one or more (C₁-C₂₂)-alkyl group;

17

R_7 may be identical or different and, independently of one another, are hydrogen, a halogen atom, a saturated or unsaturated (C_1-C_{22}) -alkyl group, a (C_1-C_{22}) -alkyl ether group, a hydroxyl group, a hydroxy- (C_1-C_{22}) -alkyl group, a (C_1-C_{22}) -alkoxy group, a cyano group, a nitro group, an amino group, a (C_1-C_{22}) -alkylamino group, a (C_1-C_{22}) -dialkylamino group, a carboxylic acid group, a $C(O)O-(C_1-C_{22})$ -alkyl group, a substituted or unsubstituted $C(O)O$ -phenyl group;

Q^- may be an anion that balances the overall charge of the compound of Formula I, and the index q may be either 0 or 1. Suitable anions include chloro, bromo, methosulfate, tetrafluoroborate, and acetate anions.

R_1 may be a (C_1-C_{22}) -alkyl, an alkyl aromatic or an alkyl sulfonate radical having Formula (III) below;



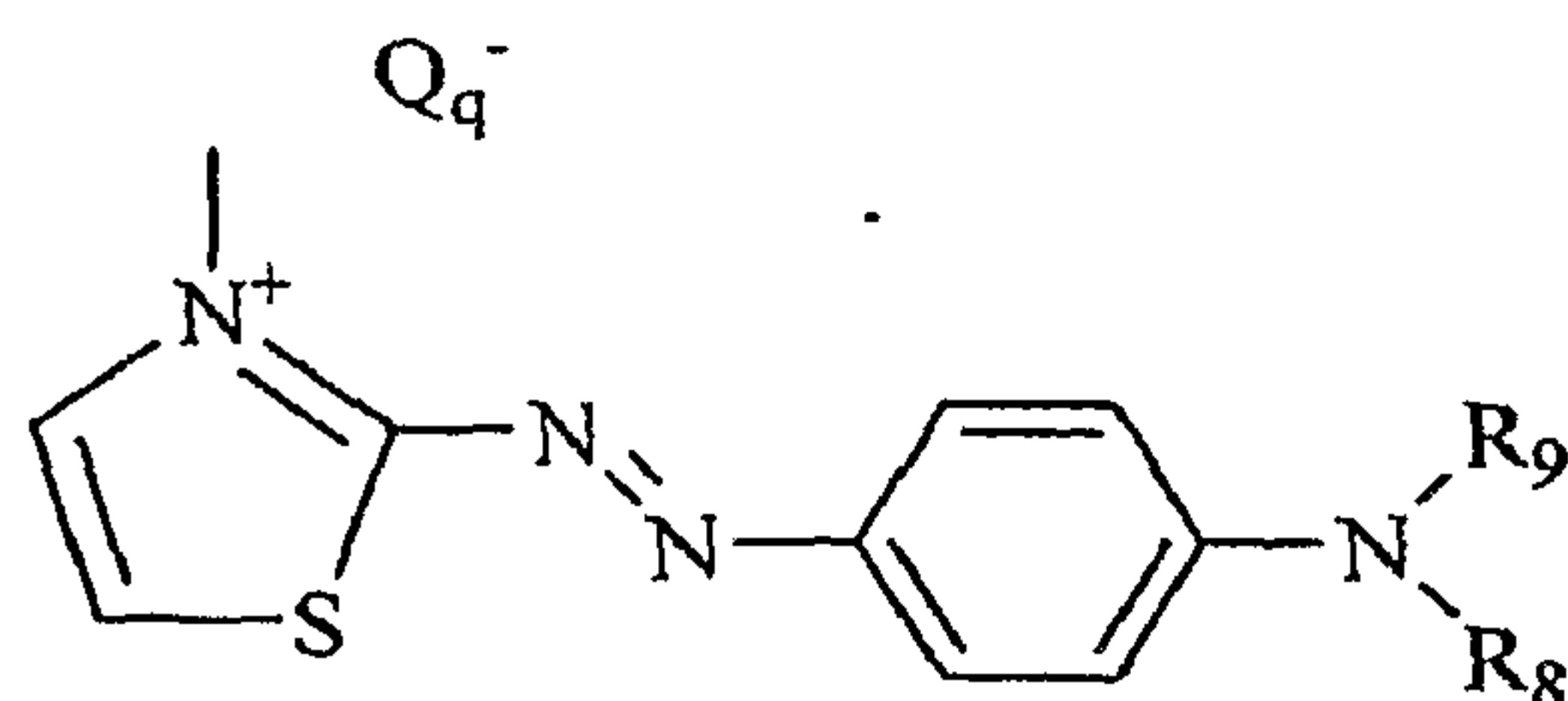
Formula III

wherein

R_2 is hydrogen, methyl, ethyl, propyl, acetate or a hydroxyl group; m and p are integers from 0 to $(n-1)$, n is an integer from 1 to 6 and $m+p = (n-1)$;

with the proviso that the heterocycle of the Formula (I) comprises at least two and at most three heteroatoms, where the heterocycle has at most one sulfur atom;

In one aspect, a suitable thiazolium dye may have Formula IV below:

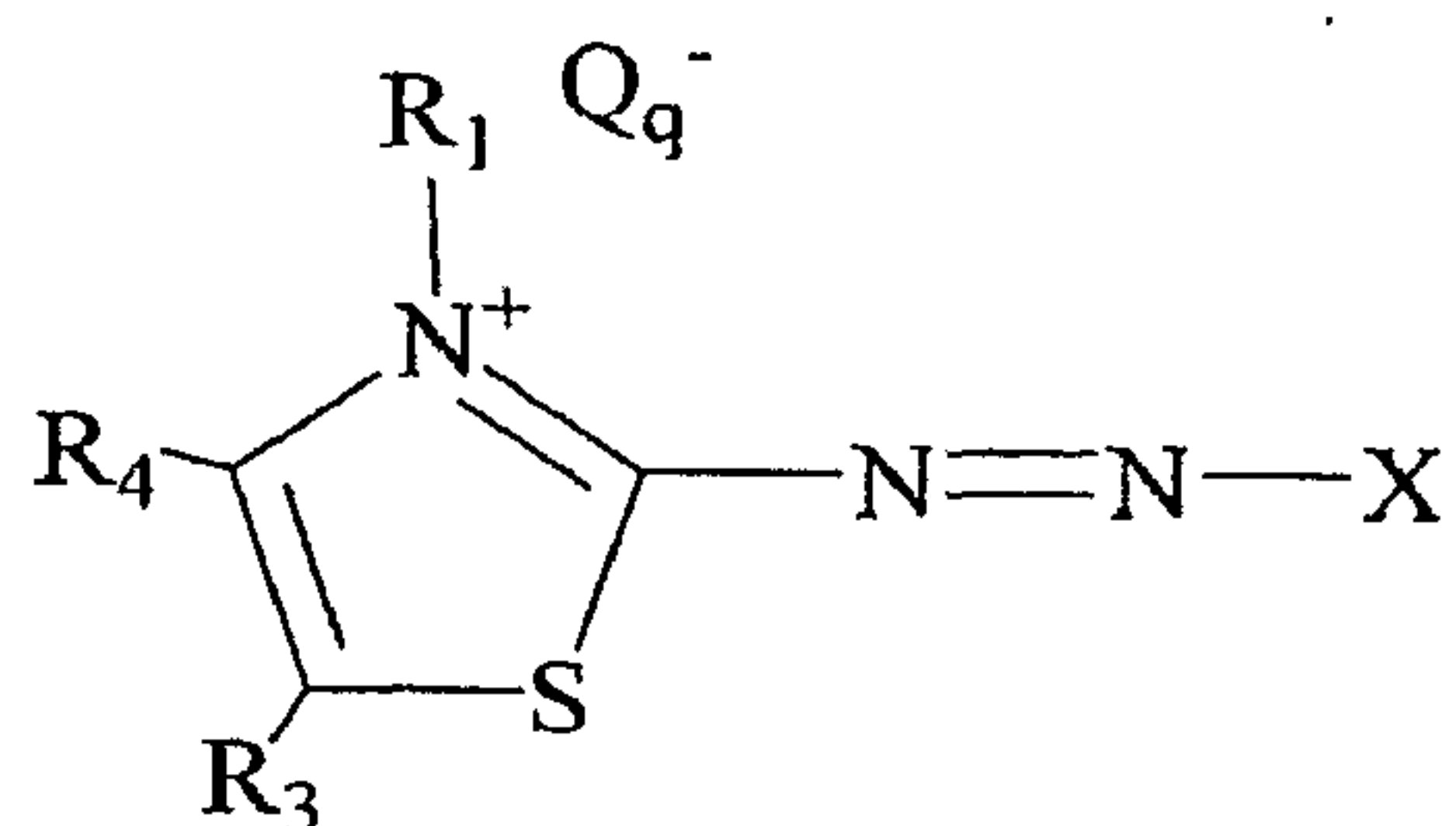


Formula IV

wherein R_8 and R_9 may be identical or different and, independently of one another, may be a saturated or unsaturated (C_1-C_{22}) -alkyl group, a (C_1-C_{22}) -alkyl group, a hydroxy- $(C_2-$

C₂₂)-alkyl group optionally interrupted by oxygen, a polyether group derived from ethylene oxide, propylene oxide or butylene oxide, an amino-(C₁-C₂₂)-alkyl group, a substituted or unsubstituted phenyl group or a benzyl group, a (C₁-C₂₂)-alkyl group terminated in sulfonate, sulfate, or carboxylate, or R₈ and R₉, together with the nitrogen atom, may form a 5-membered to 6-membered ring system, which may comprise a further heteroatom; or R₈ or R₉ may form, with a carbon atom of the benzene ring, an optionally oxygen-containing or nitrogen containing five or six-membered heterocycle which may be substituted with one or more (C₁-C₂₂)-alkyl groups, and mixtures thereof, and R₁₀ is hydrogen or methyl. For Formula IV, Q⁻ is as described for Formula I above.

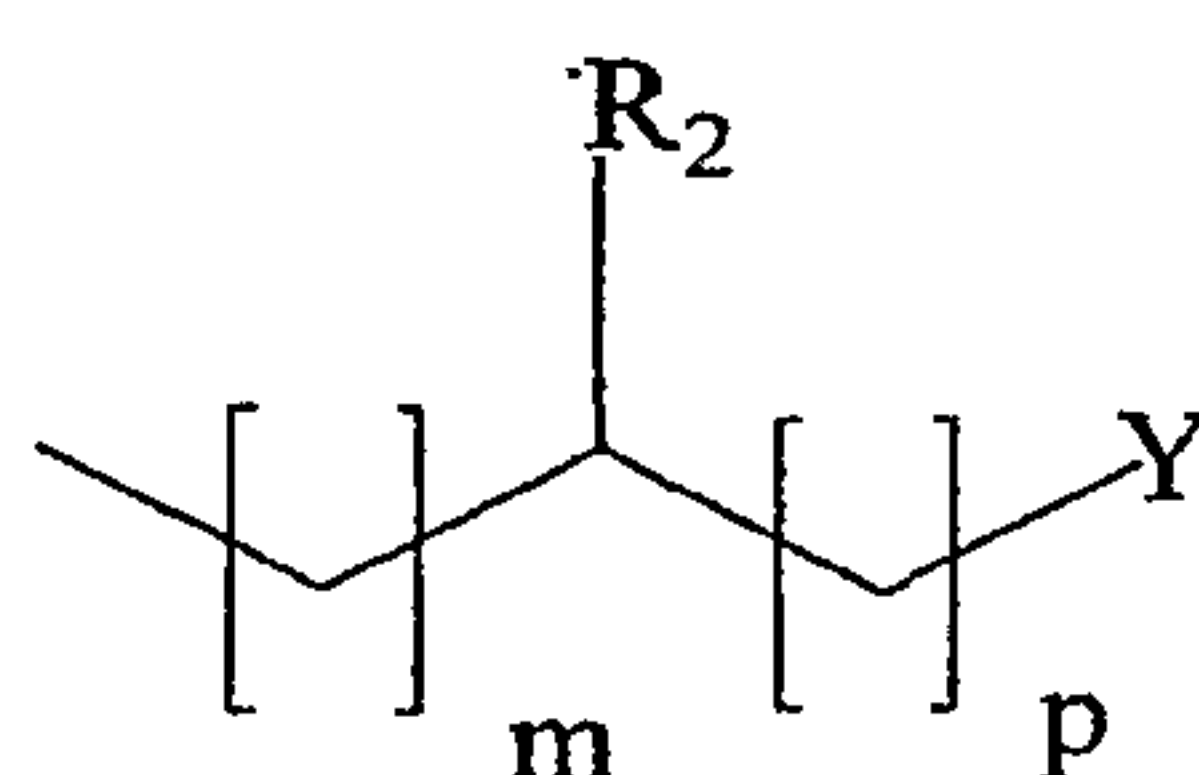
In one aspect, suitable thiazolium dyes may have Formula (V);



Formula V

wherein:

- a.) R₁ may be selected from a branched or unbranched (C₁-C₂₂)-alkyl moiety, an aromatic alkyl moiety, a polyalkylene oxide moiety, or a moiety having Formula (VI) below;



Formula VI

wherein

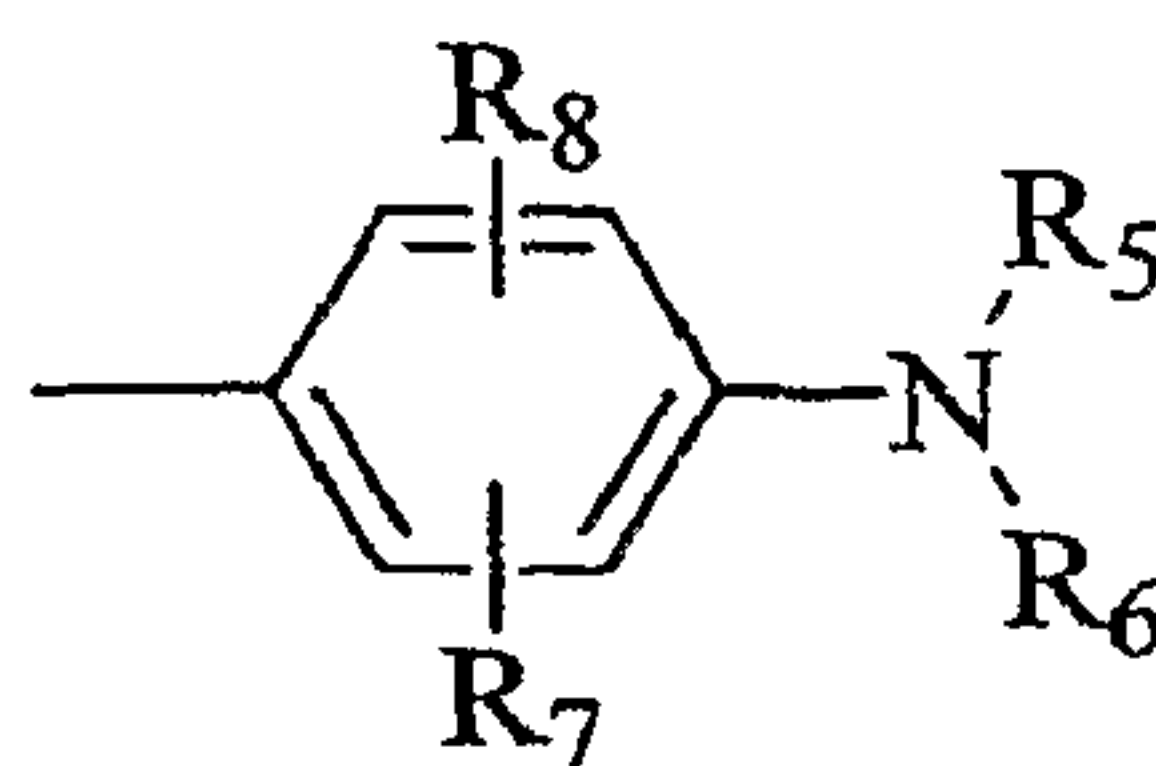
- (i) R₂ may be selected from hydrogen, methyl, ethyl, propyl, acetate or a hydroxyl moiety; m and p may be, independently, integers from 0 to (n-1), with the proviso that n is an integer from 1 to 6 and m+p = (n-1)
- (ii) Y may be selected from a hydroxyl, sulfonate, sulfate, carboxylate or acetate moiety;

- b.) R₃ and R₄:

19

- i.) may be independently selected from hydrogen; a saturated or unsaturated (C₁-C₂₂)-alkyl moiety; a hydroxy-(C₂-C₂₂)-alkyl moiety; a hydroxy-(C₂-C₂₂)-alkyl moiety comprising, in addition to the hydroxyl oxygen, an oxygen atom; a polyether moiety; an amino-(C₁-C₂₂)-alkyl moiety; a substituted or unsubstituted phenyl moiety; a substituted or unsubstituted benzyl moiety; a (C₁-C₂₂)-alkyl moiety terminated in sulfonate, sulfate, acetate, or carboxylate; or
- ii.) when taken together may form a saturated or unsaturated heterocyclic or carbocyclic moiety; or
- iii.) when taken together may form a saturated or unsaturated heterocyclic or carbocyclic moiety substituted by, sulfate, sulfonate, phosphate, nitrate, and carboxylate;

c.) X may be moiety having Formula VII below;



Formula VII

wherein:

i.) R₅ and R₆:

- (a) may be independently selected from hydrogen; a saturated or unsaturated (C₁-C₂₂)-alkyl moiety; a hydroxy-(C₂-C₂₂)-alkyl moiety; a hydroxy-(C₂-C₂₂)-alkyl moiety comprising, in addition to the hydroxyl oxygen, an oxygen atom; a capped or uncapped polyether moiety; an amino-(C₁-C₂₂)-alkyl moiety; a substituted or unsubstituted phenyl moiety; a substituted or unsubstituted benzyl moiety; a (C₁-C₂₂)-alkyl moiety comprising a terminating C₁-C₄ alkyl ether, sulfonate, sulfate, acetate or carboxylate moiety; a thiazole moiety or

- (b) when taken together may form a saturated or unsaturated heterocyclic moiety; or

(c) when taken together form a saturated or unsaturated heterocyclic moiety substituted by one or more, alkoxyate, sulfate, sulfonate, phosphate, nitrate, and/or carboxylate moieties;

(d) when taken together with R_7 , R_8 , or R_7 and R_8 form one or more saturated or unsaturated heterocyclic moieties, optionally substituted by one or more alkoxyate, sulfate, sulfonate, phosphate, nitrate, and/or carboxylate moieties; or

(e) when taken together form a thiazole moiety;

ii.) R_7 and R_8 may be independently selected from hydrogen or a saturated or unsaturated alkyl moiety;

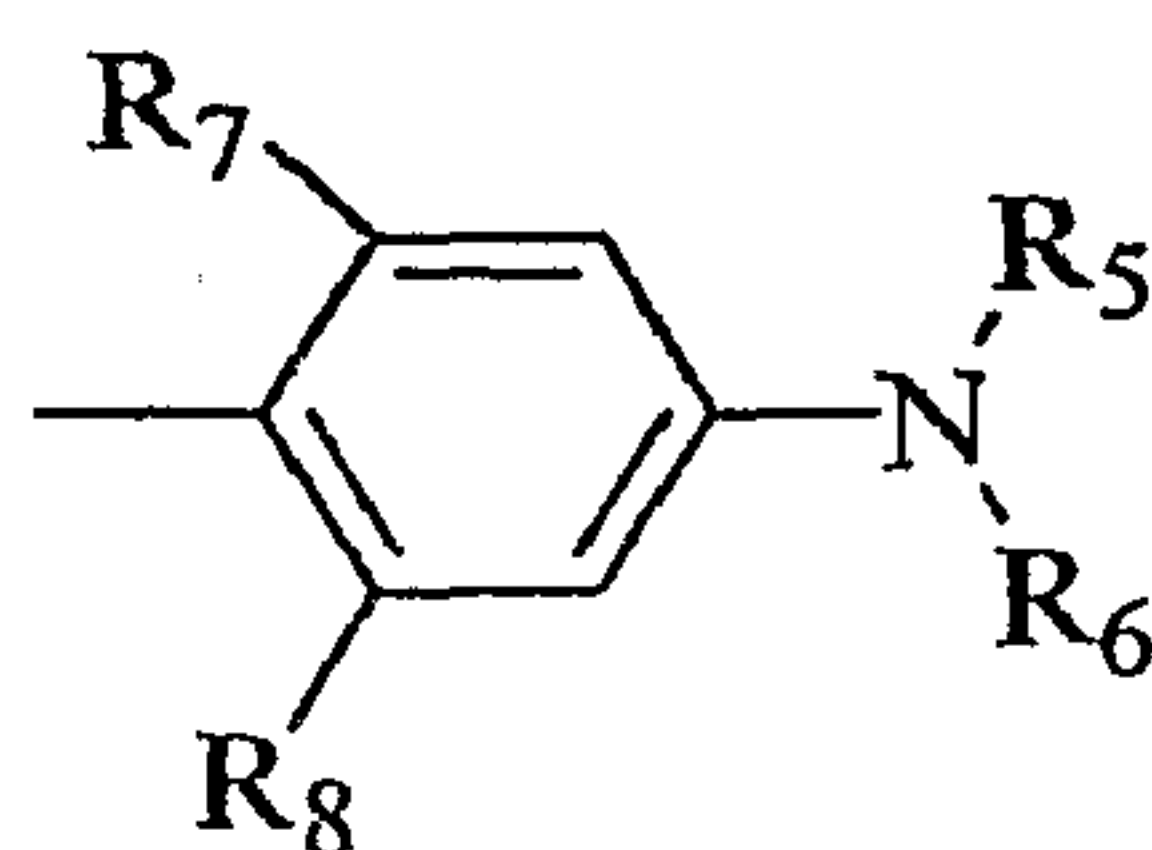
d.) Q^- may be an anion that balances the overall charge of the compound of Formula I, and the index q is 0 or 1. Suitable anions include chloro, bromo, methosulfate, tetrafluoroborate, and acetate anions.

In one aspect, for Formula V:

a.) R_1 may be a methyl moiety;

b.) R_3 and R_4 may be hydrogen; and

c.) X may have Formula VIII below:



Formula VIII

wherein

(i) R_5 and R_6 may be as defined for Formula VII above;

(ii) R_7 may be hydrogen or a methyl moiety; and

(iii) R_8 may be hydrogen.

In one aspect, for Formula VII R_5 and R_6 each comprise, independently, from 1 to 20 alkylene oxide units and, independently, a moiety selected from the group consisting of: styrene oxide, glycidyl methyl ether, isobutyl glycidyl ether, isopropylglycidyl ether, t-butyl glycidyl ether, 2-ethylhexylglycidyl ether, or glycidylhexadecyl ether.

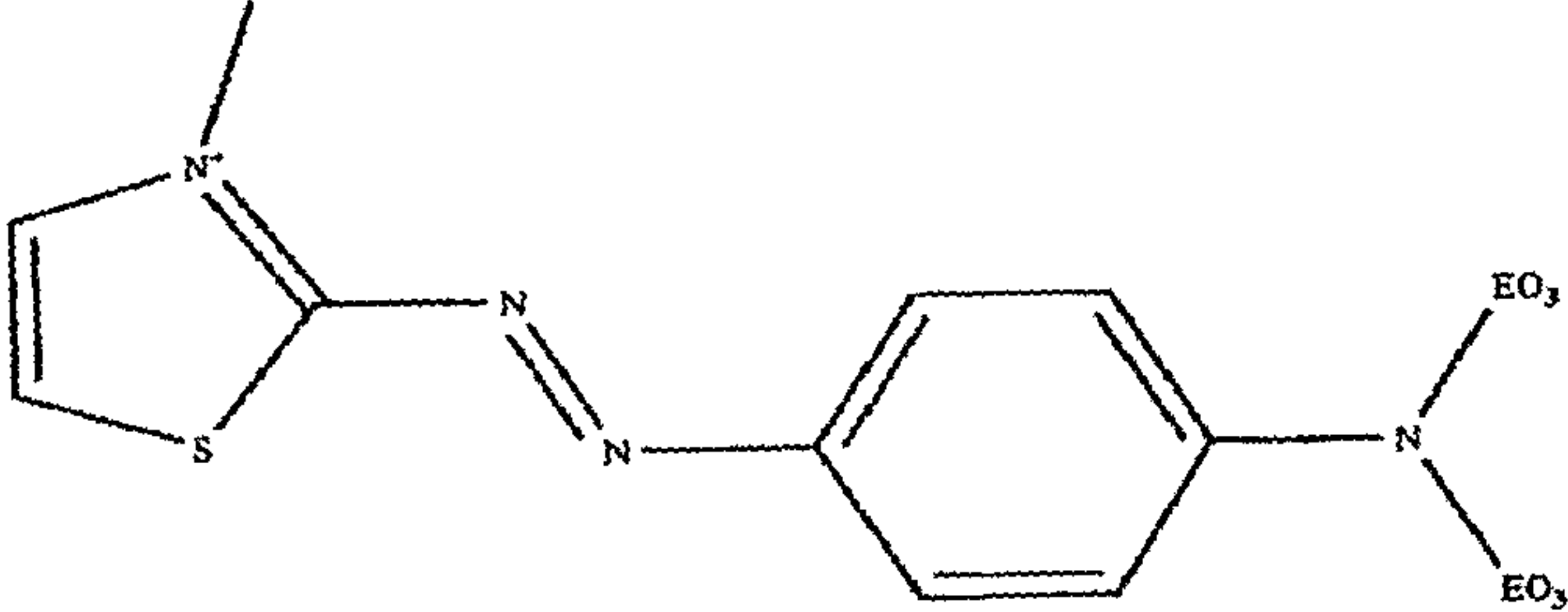
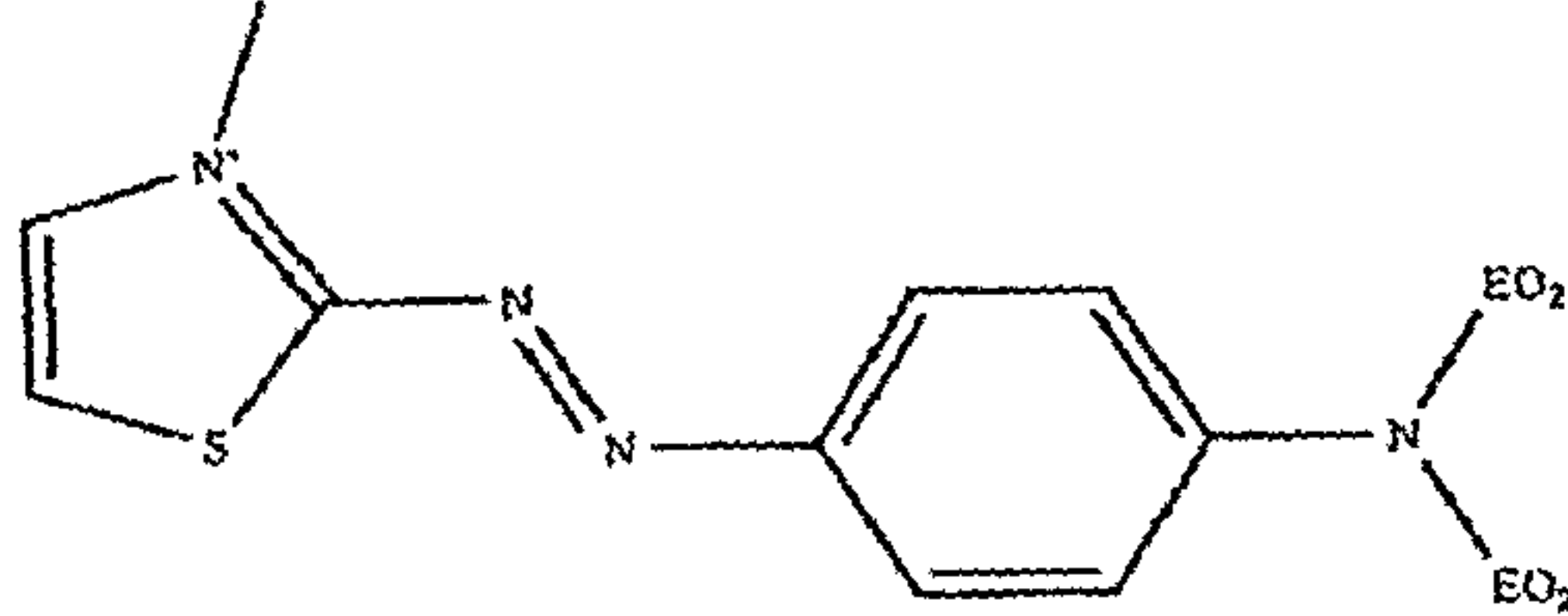
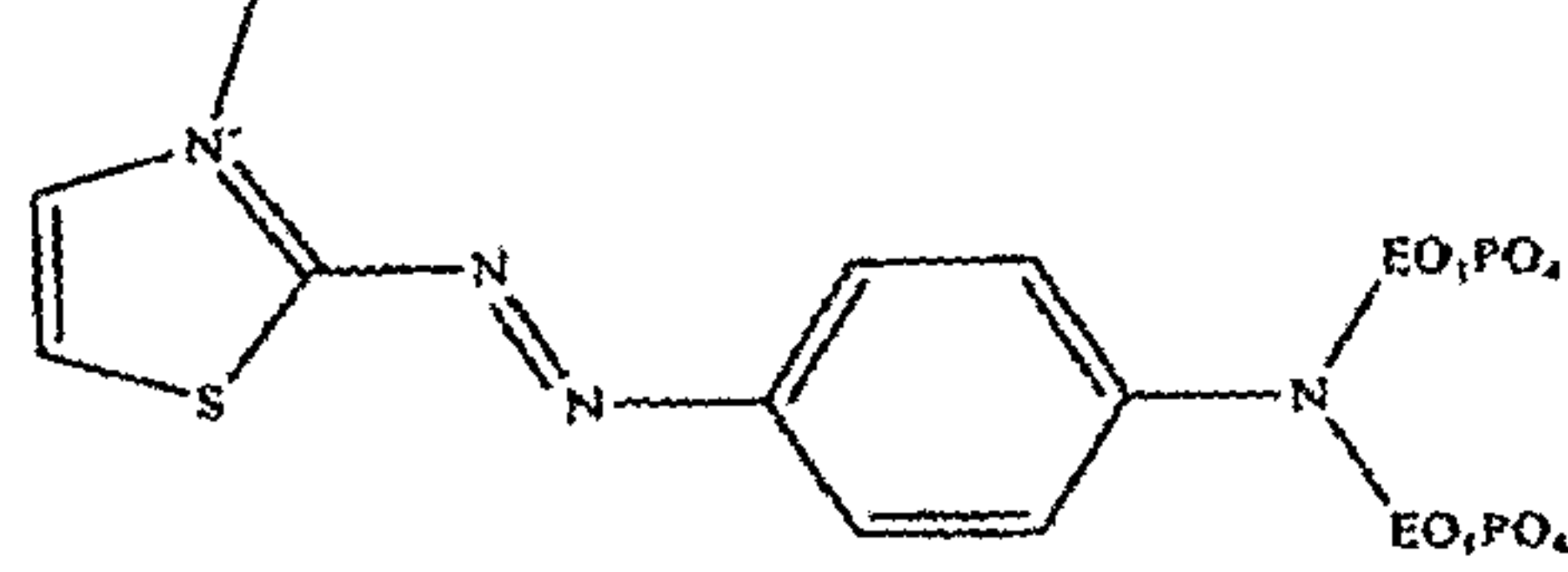
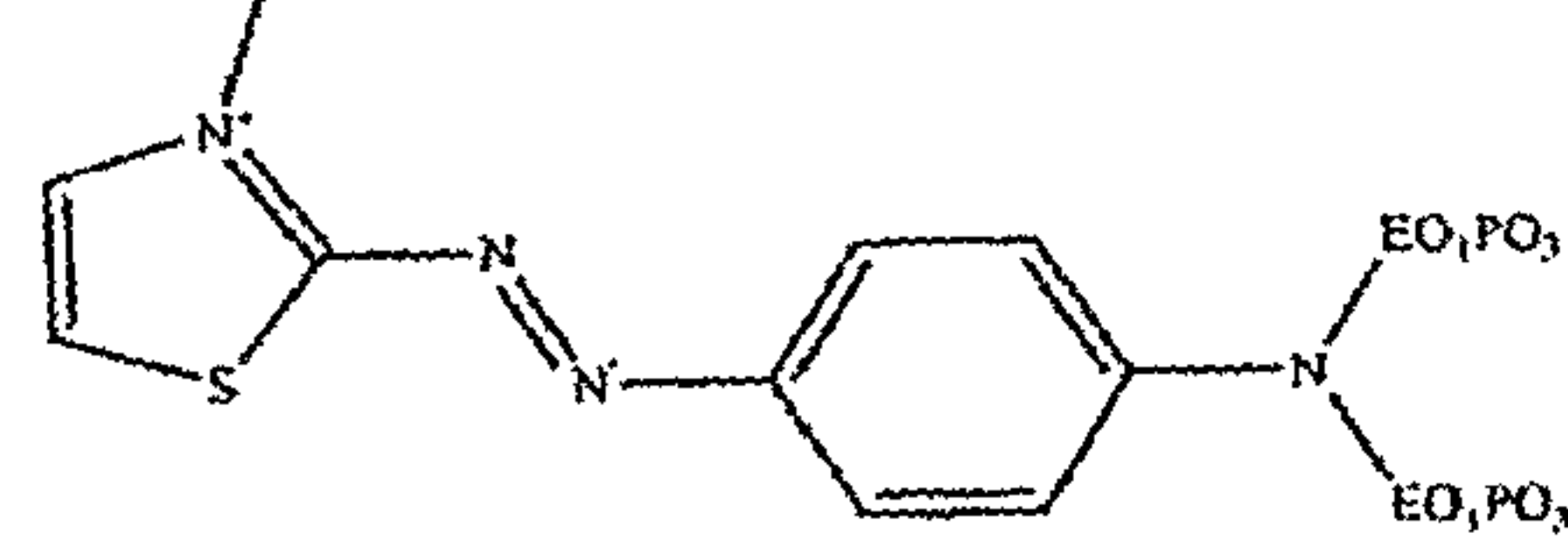
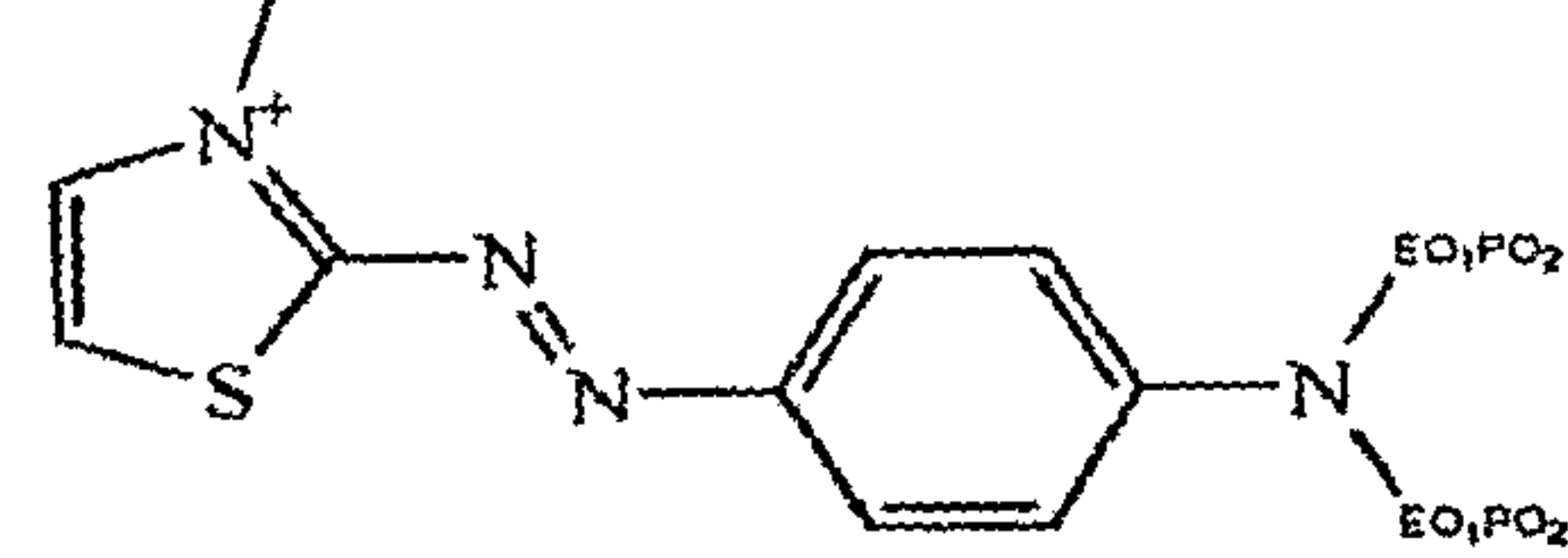
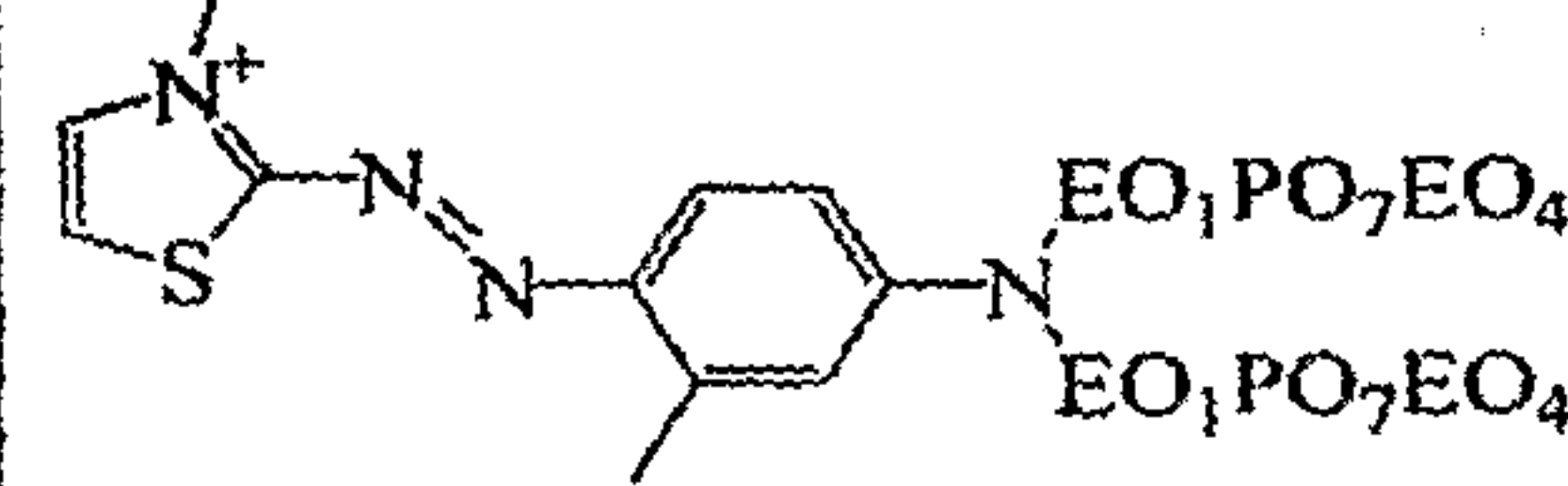
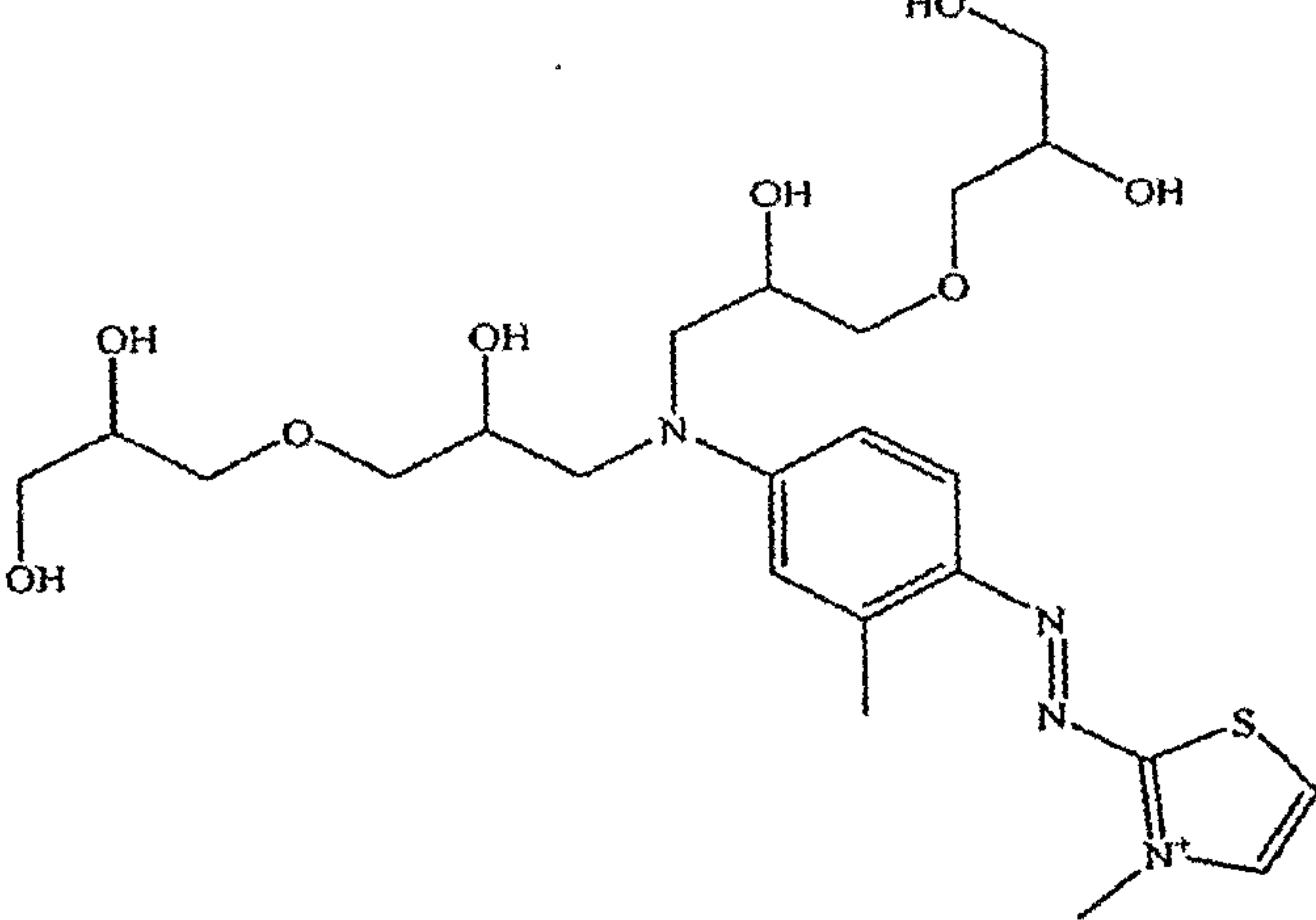
In one aspect, suitable thiazolium dyes are set forth in Table 1 below and are defined as Table 1 Thiazolium Dyes. The chemical names, as determined by ChemFinder software Level:Pro; Version 9.0 available from CambridgeSoft, Cambridge, Massachusetts, U.S.A., for such dyes are respectively provided in Table 2 below. Such dyes are associated, as needed to balance the molecule's charge, with an anion Q⁻. Such anion is not shown in the structures below but for the purposes of the present specification is assumed to be present as required. Such anion is as described above for Formula (I).

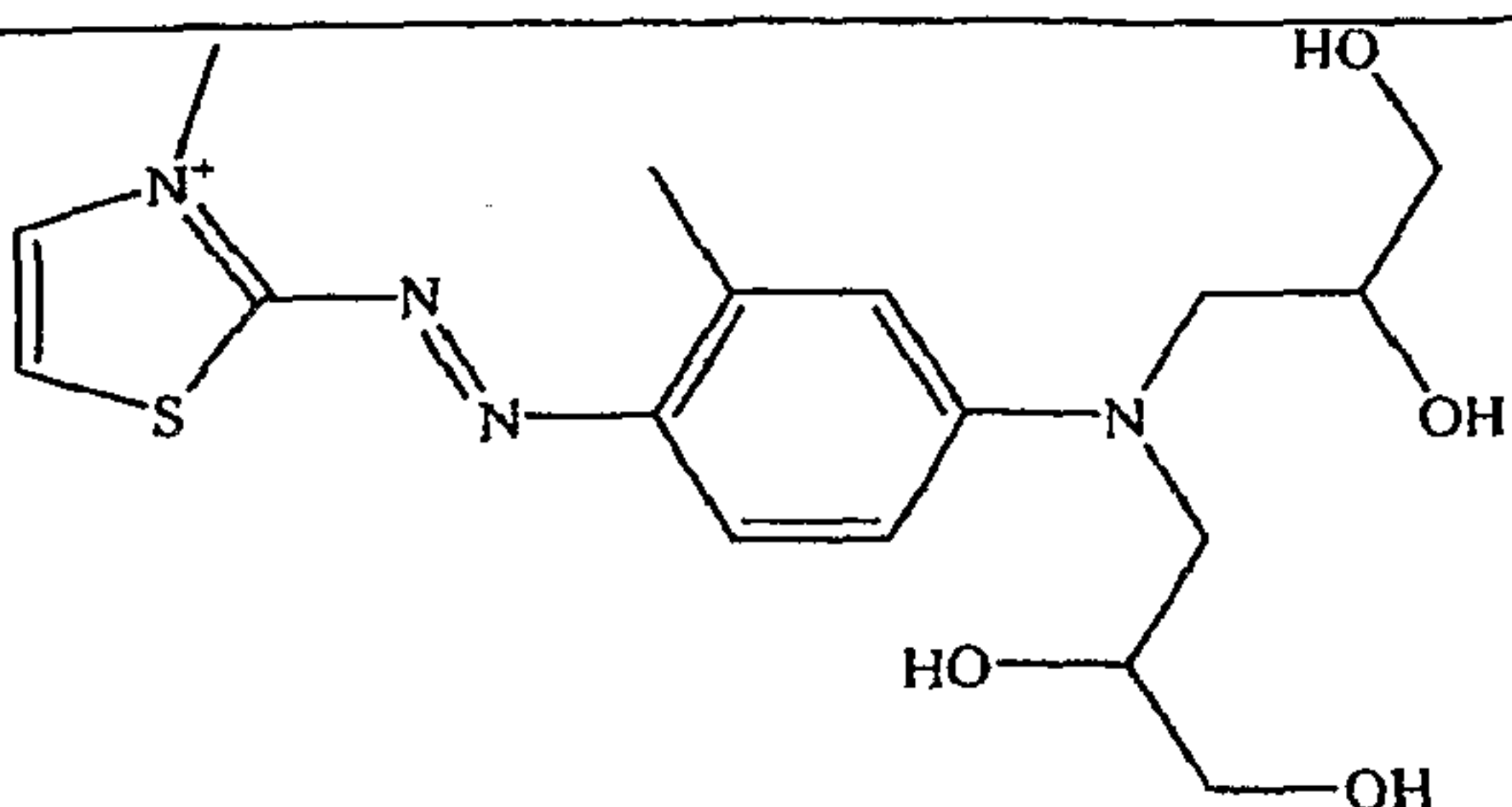
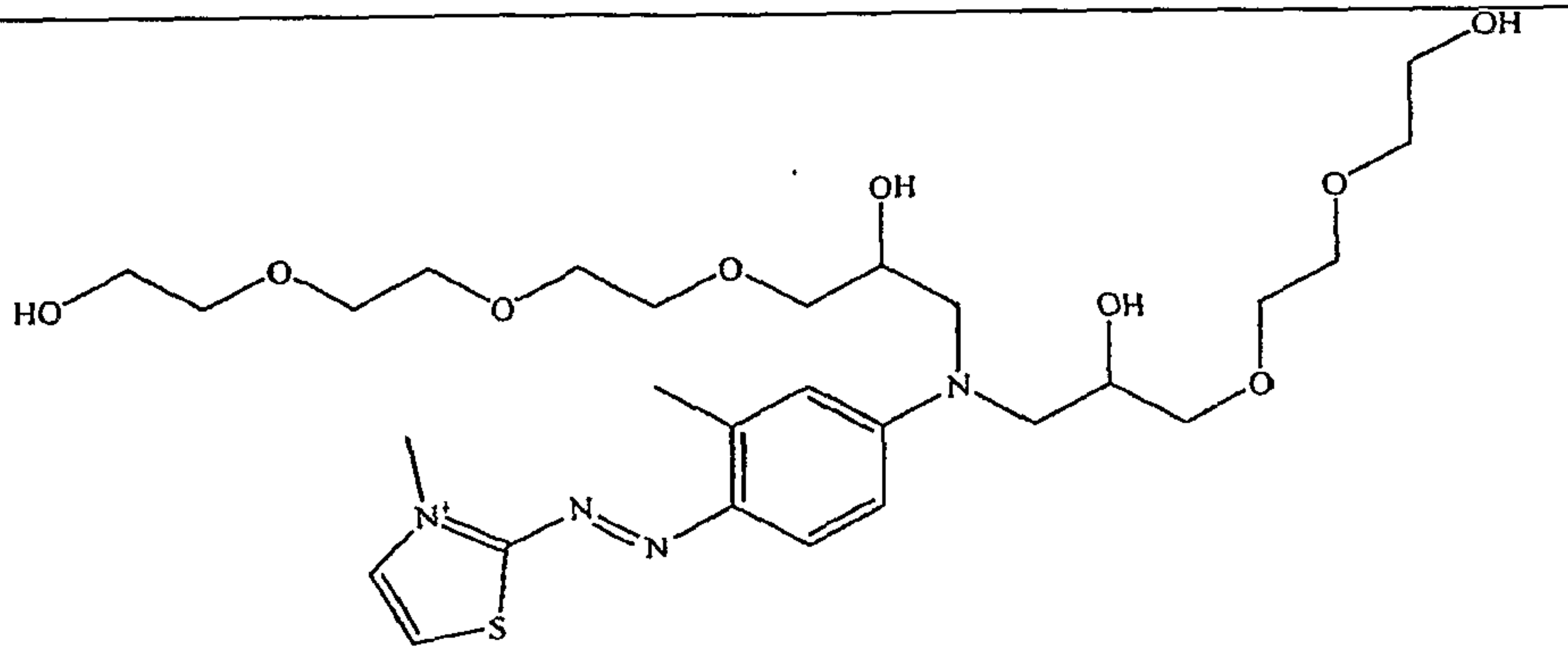
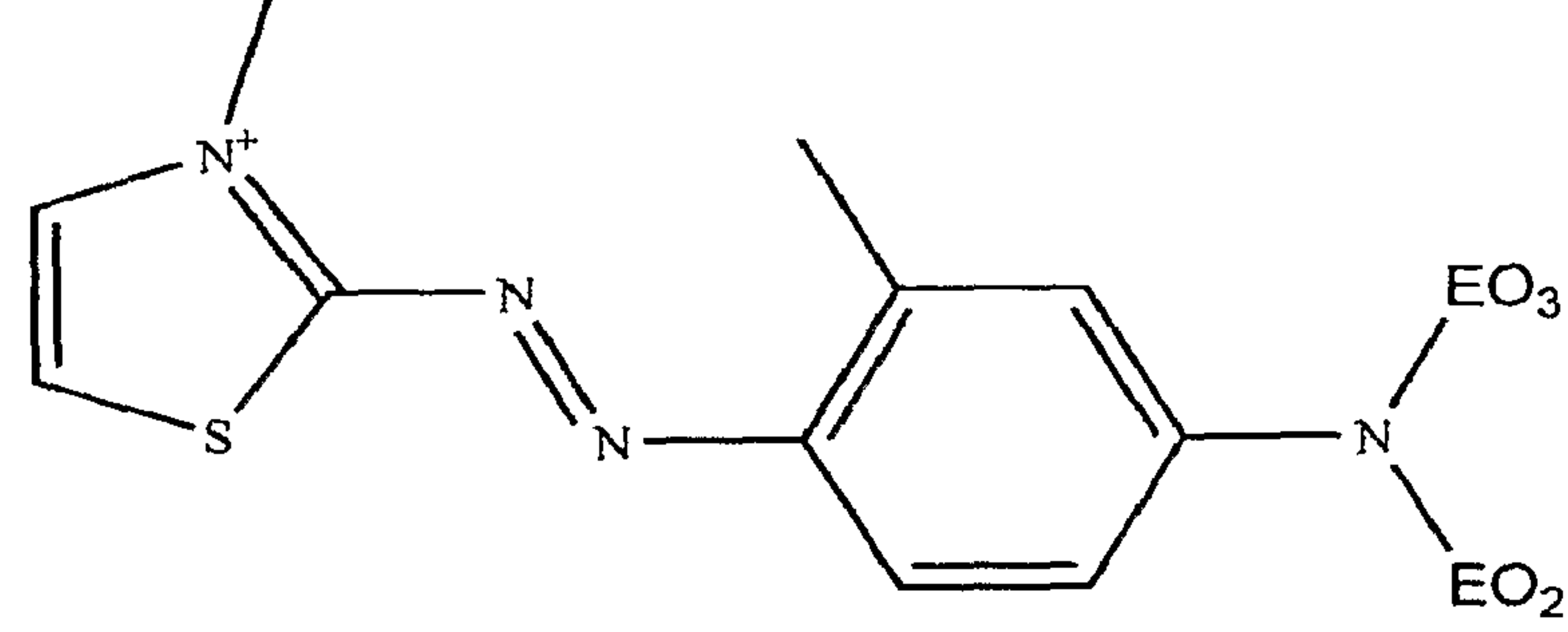
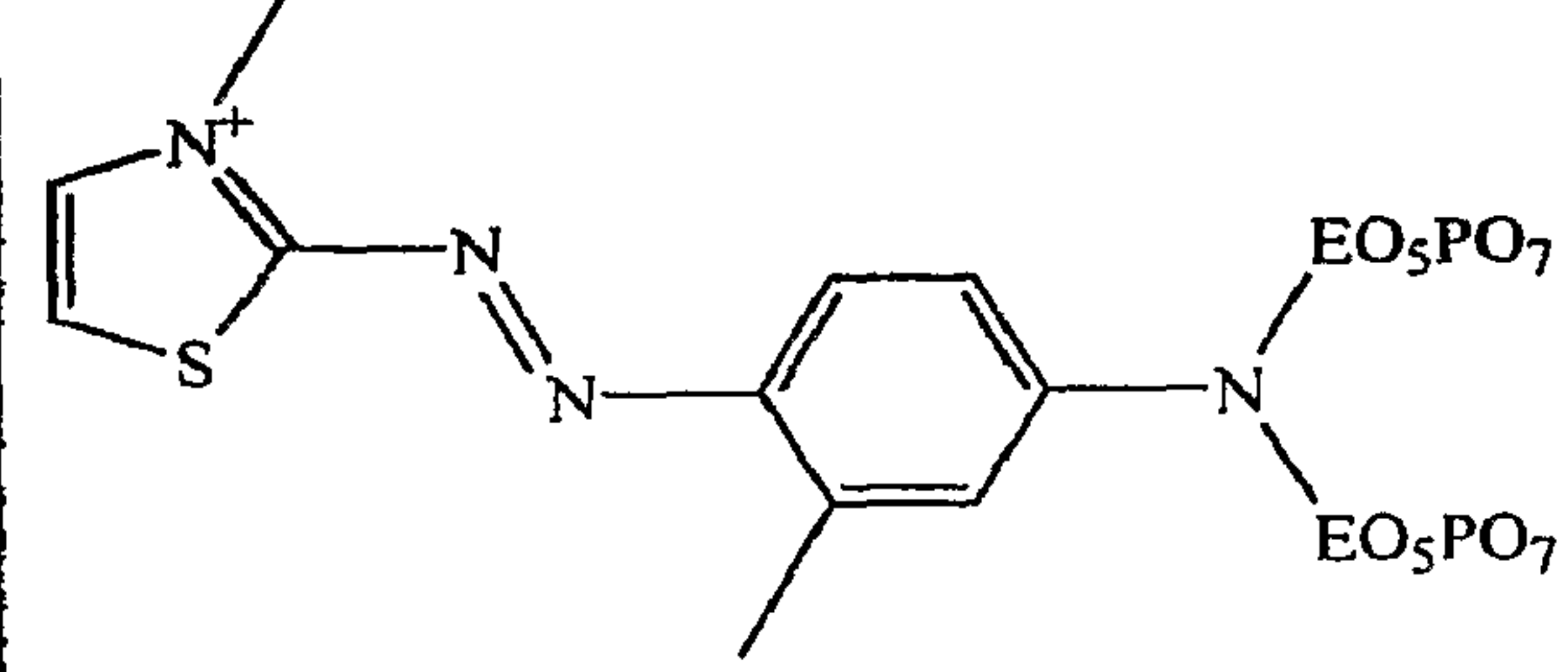
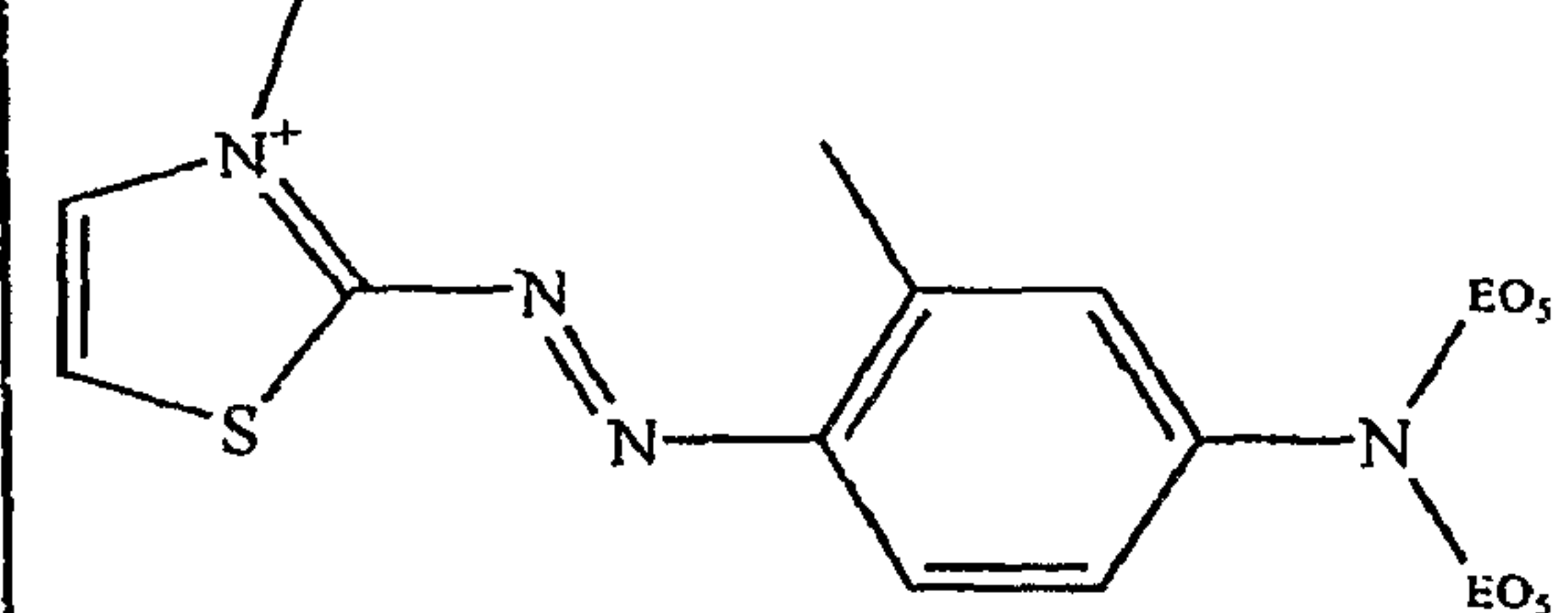
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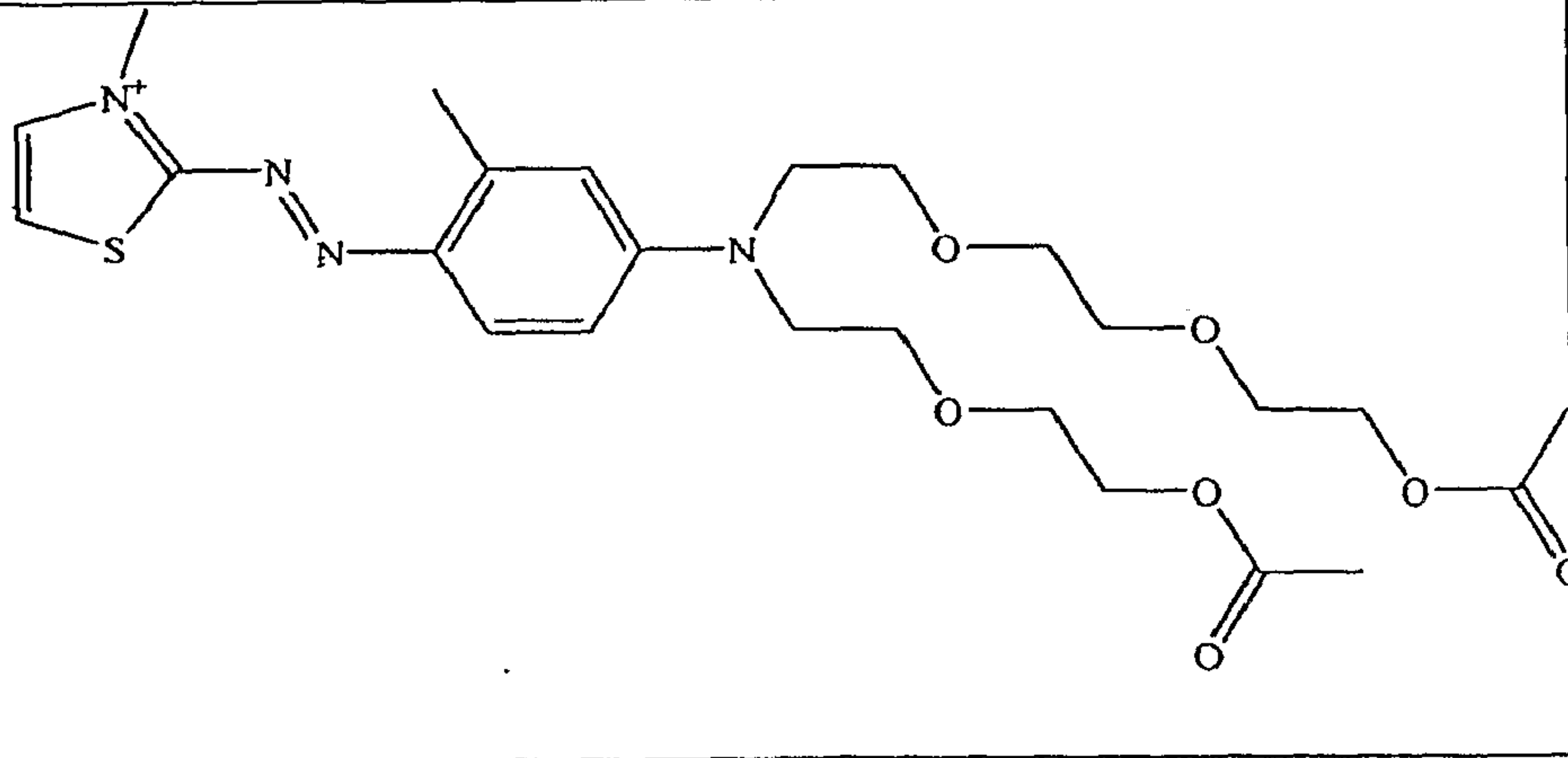
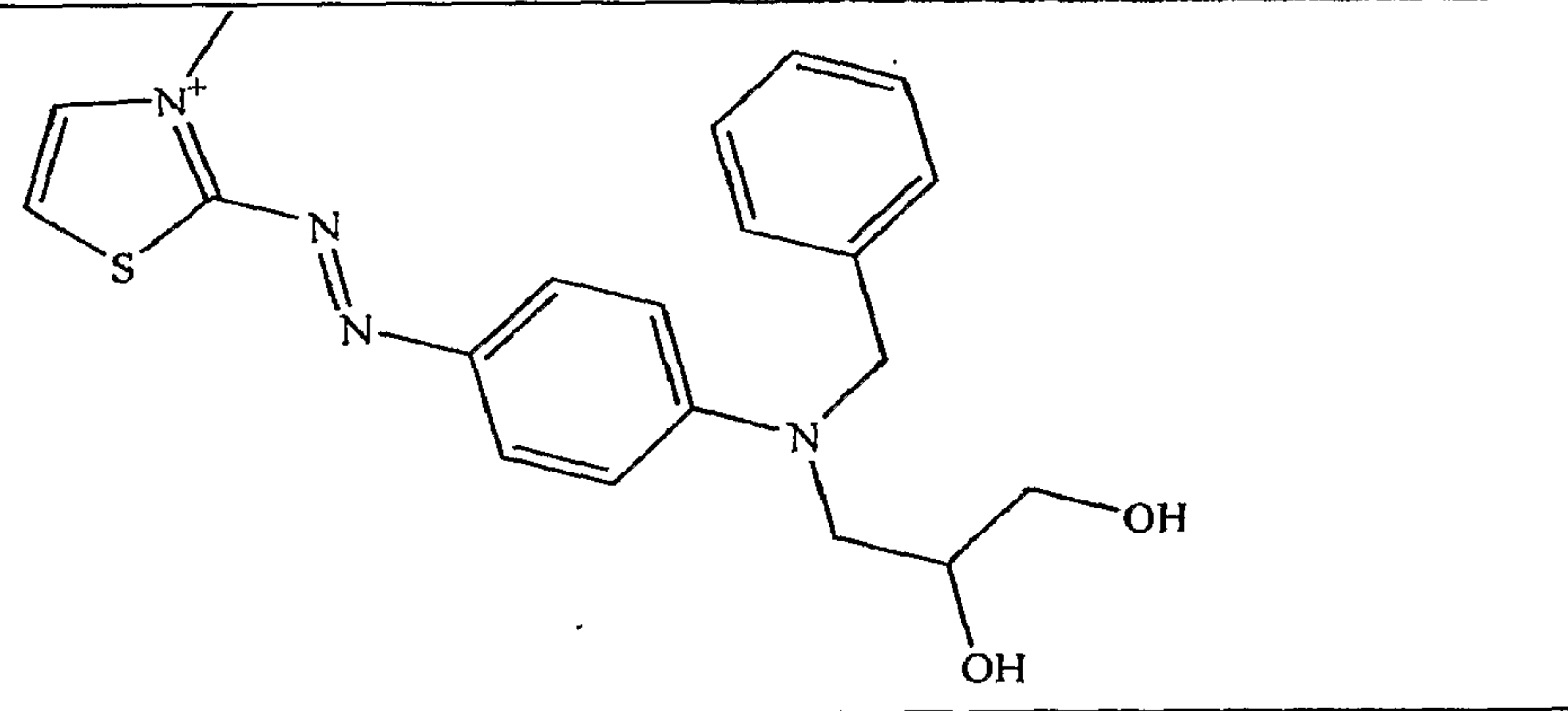
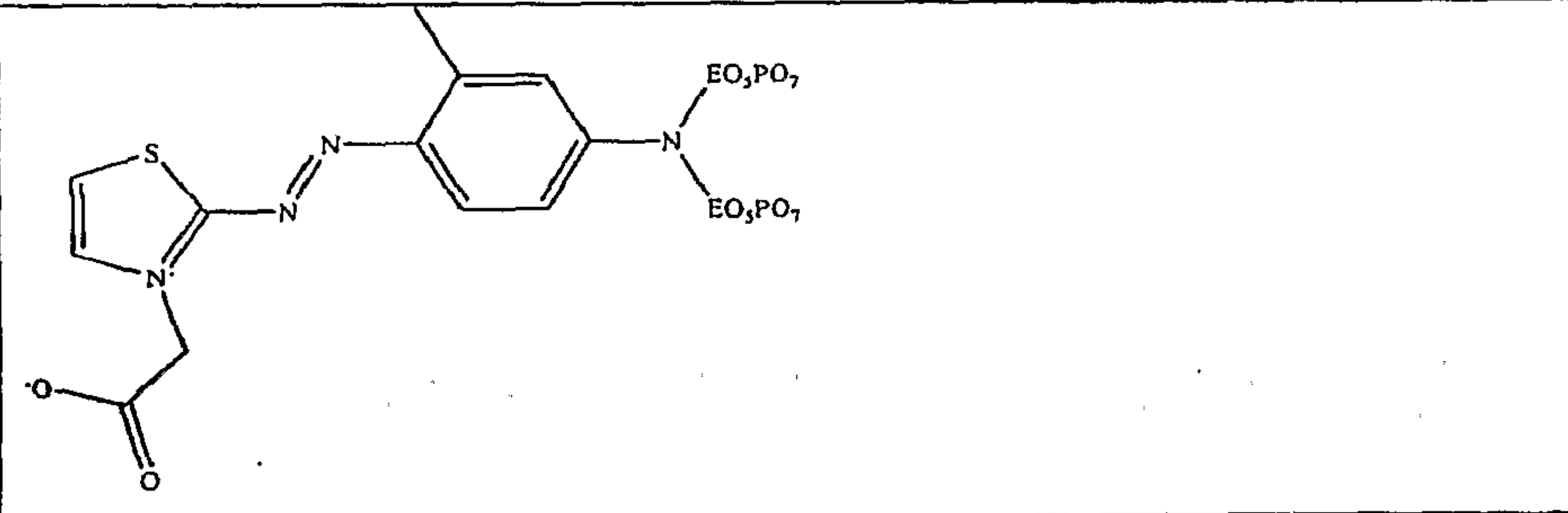
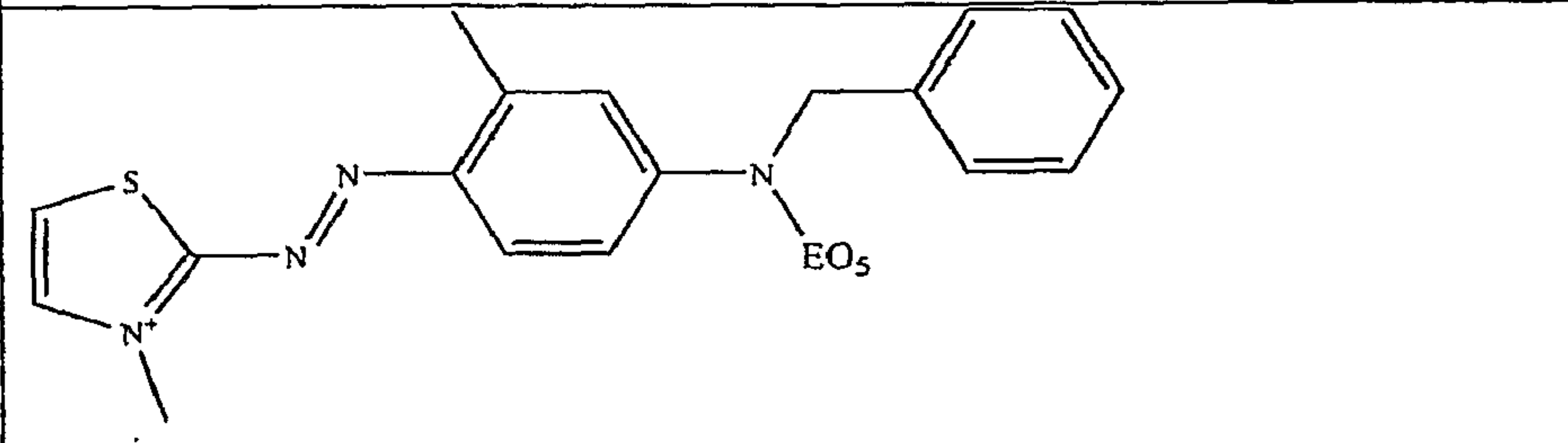
Table 1

No.	Structure
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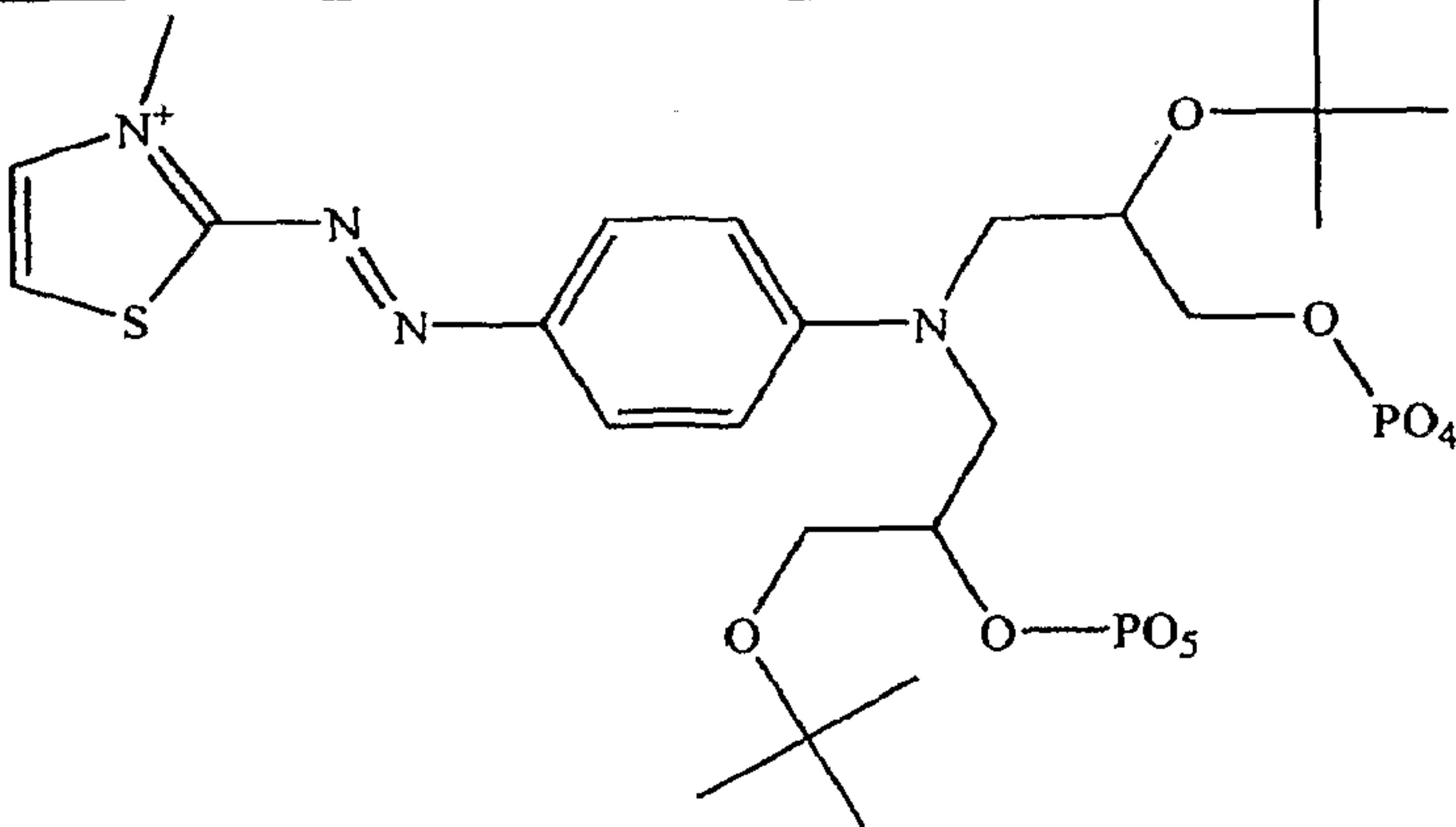
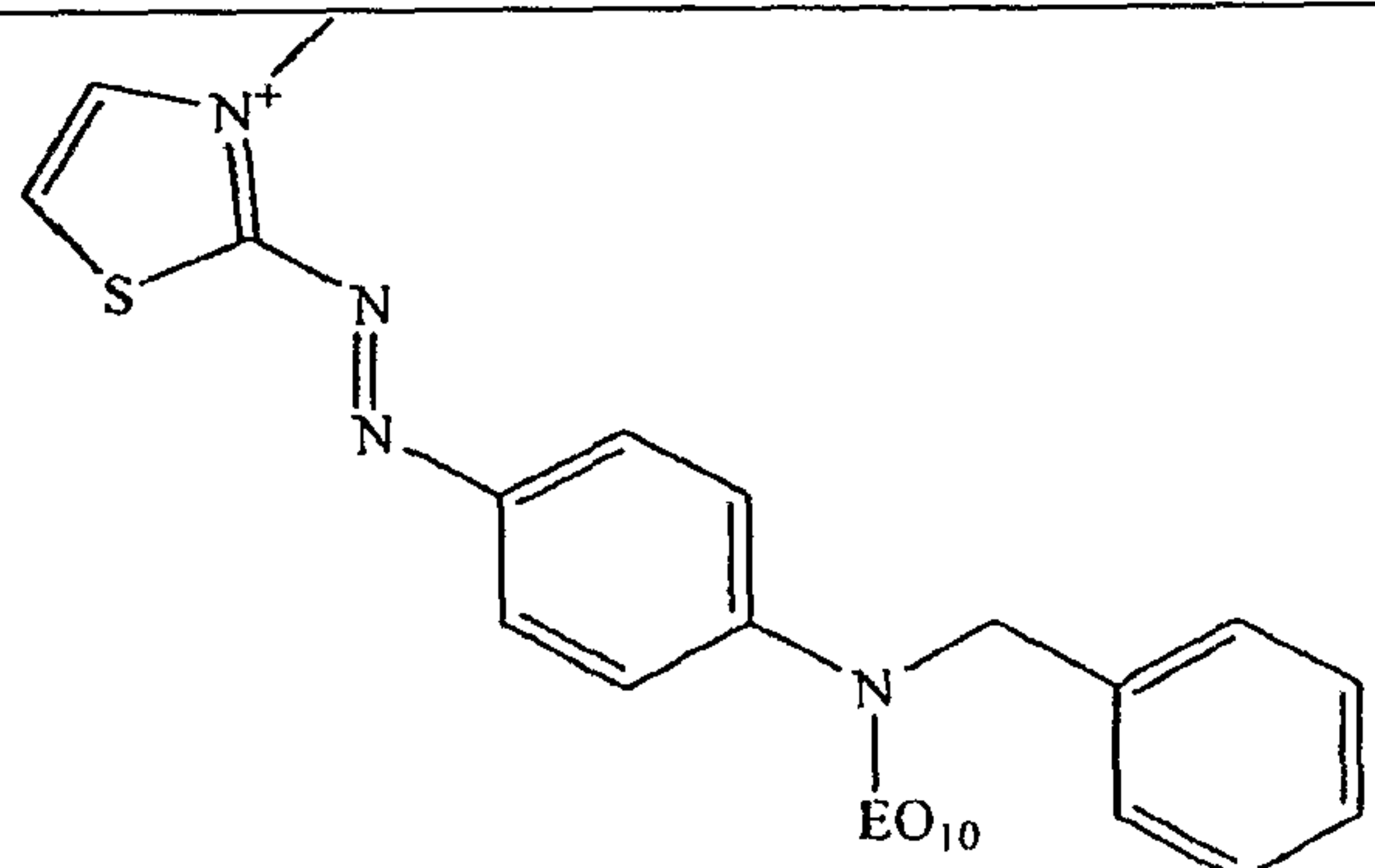
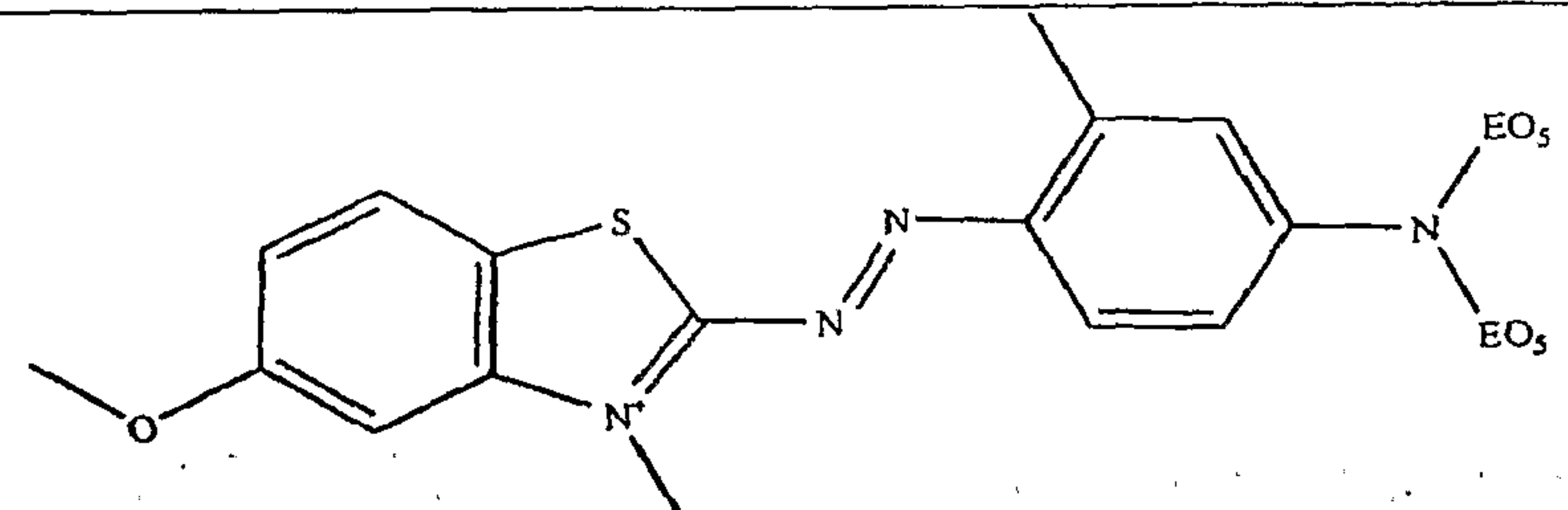
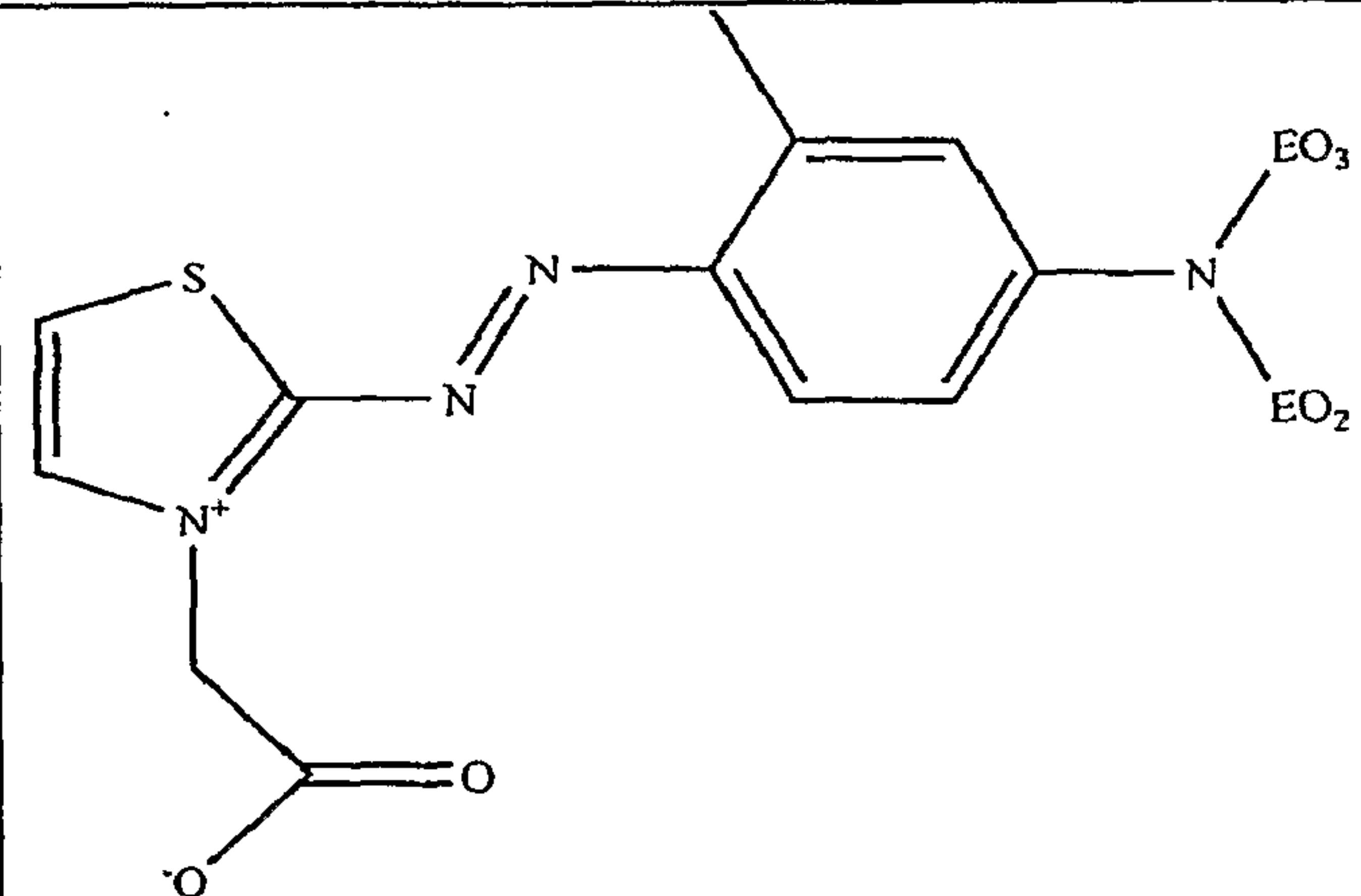
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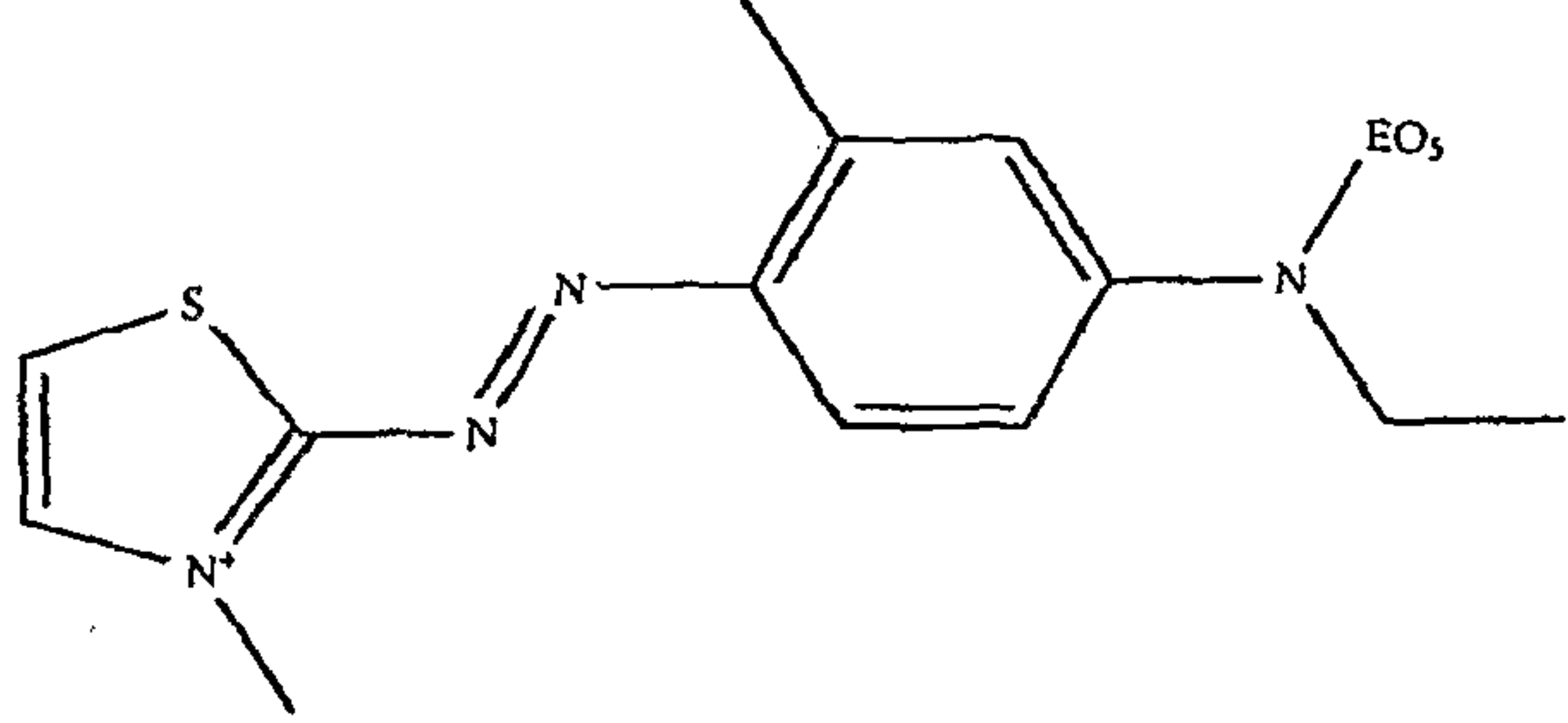
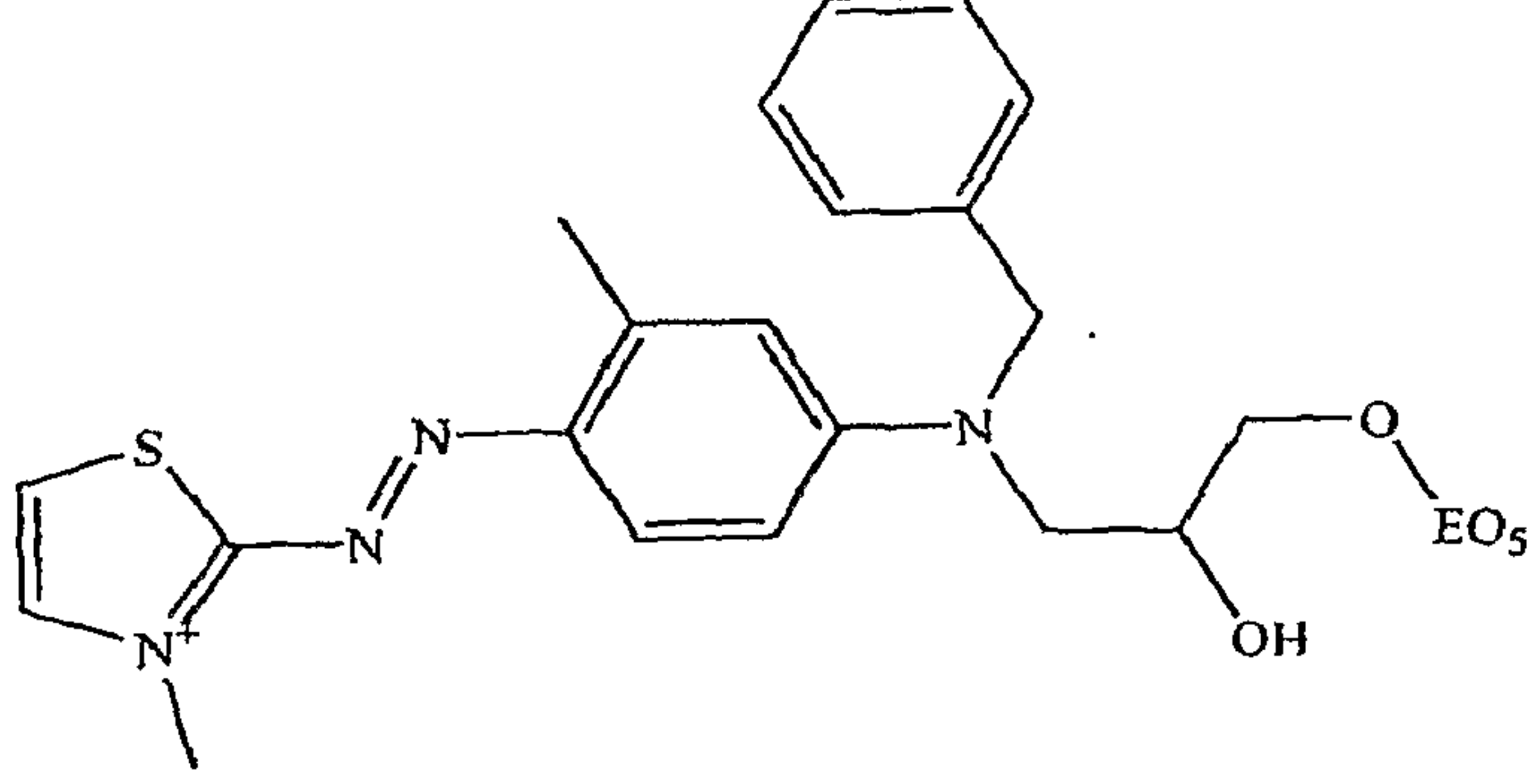
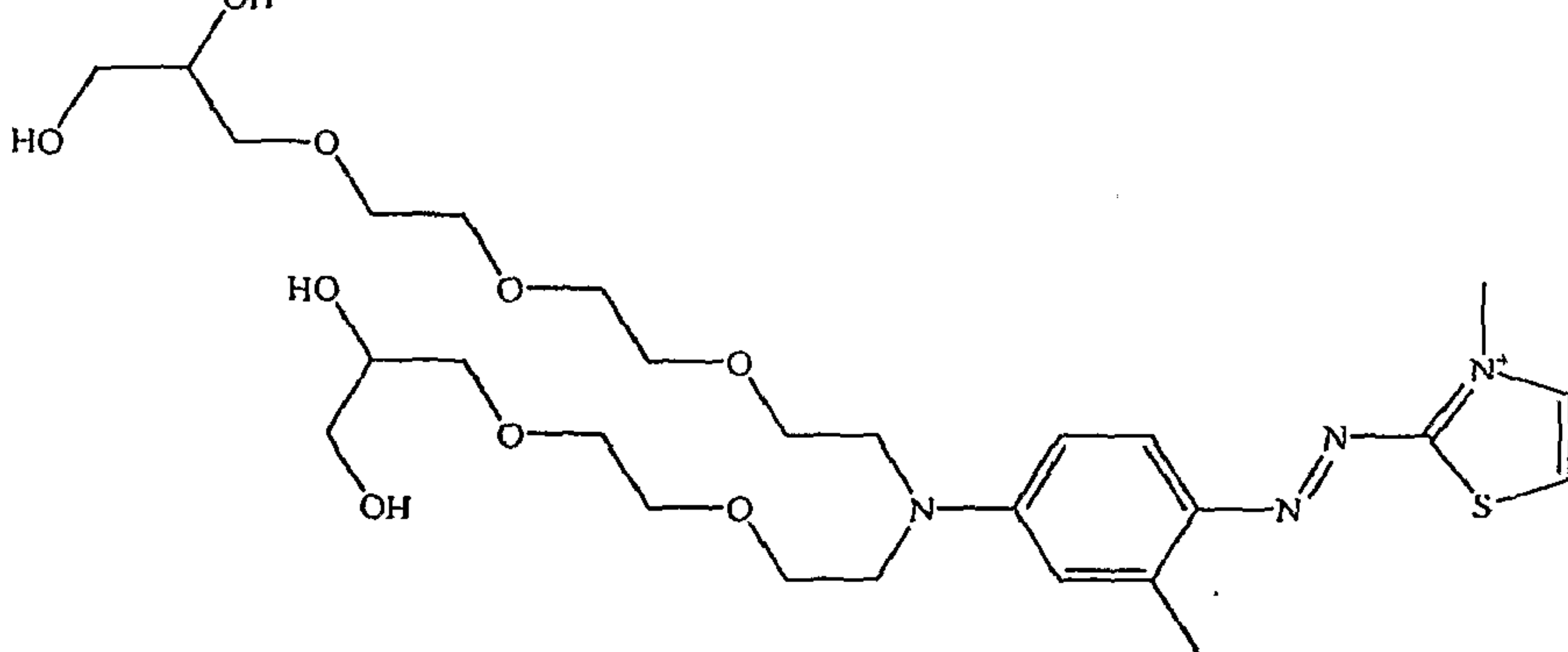
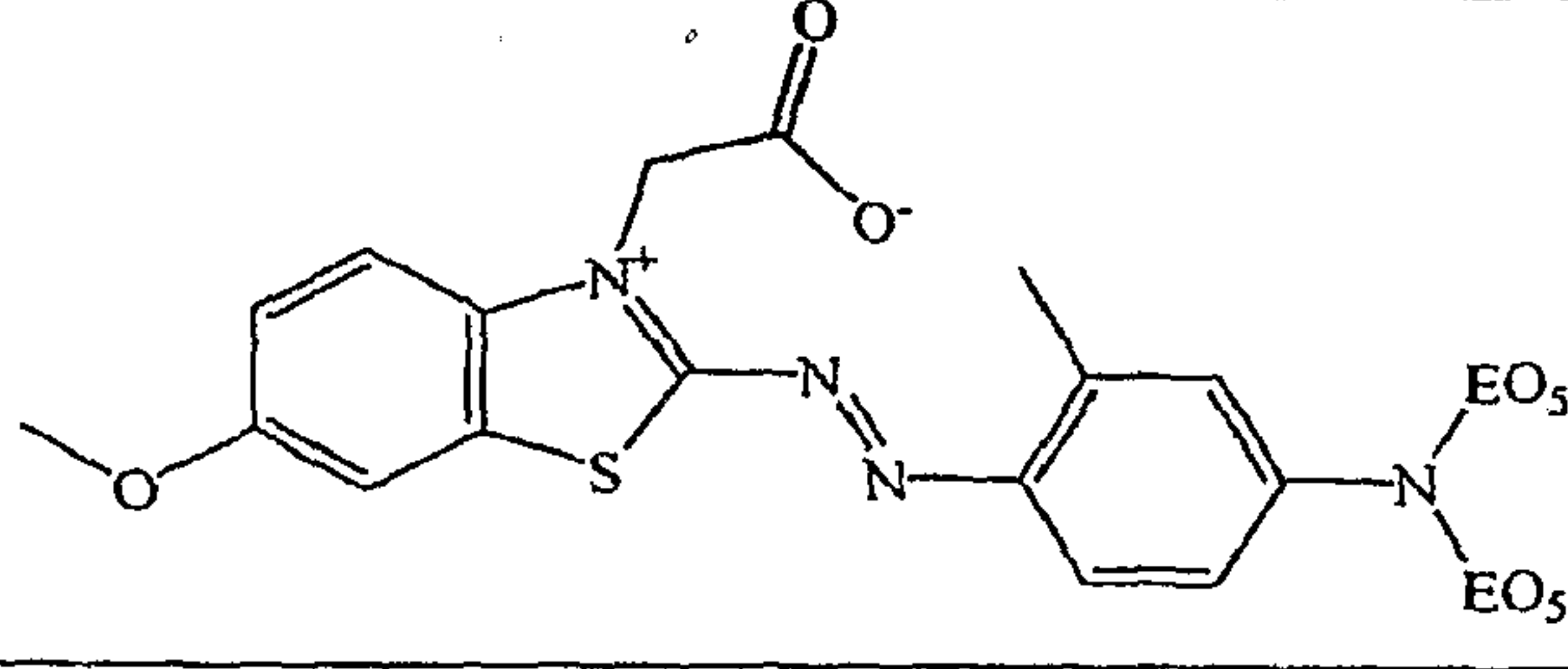
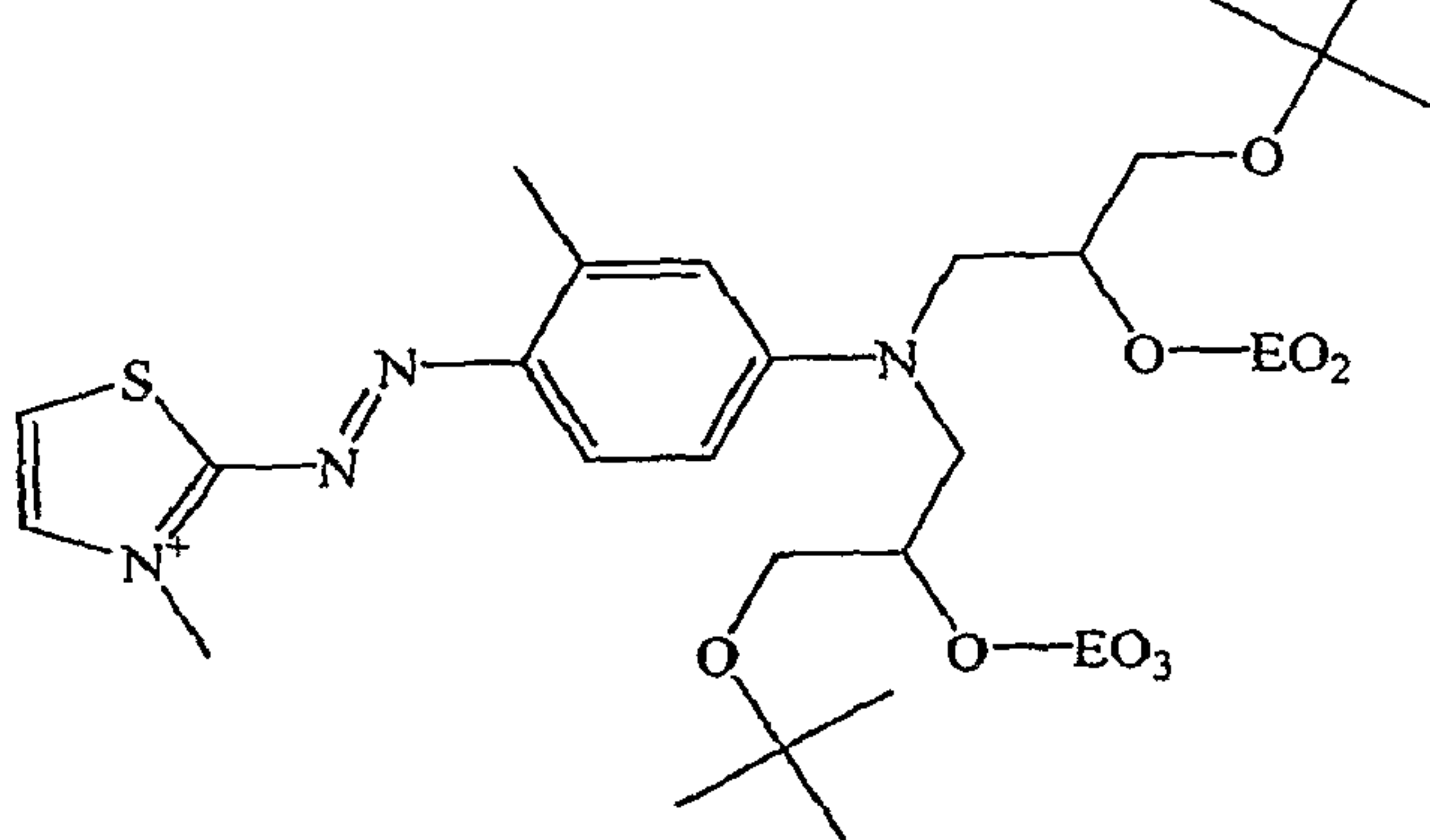
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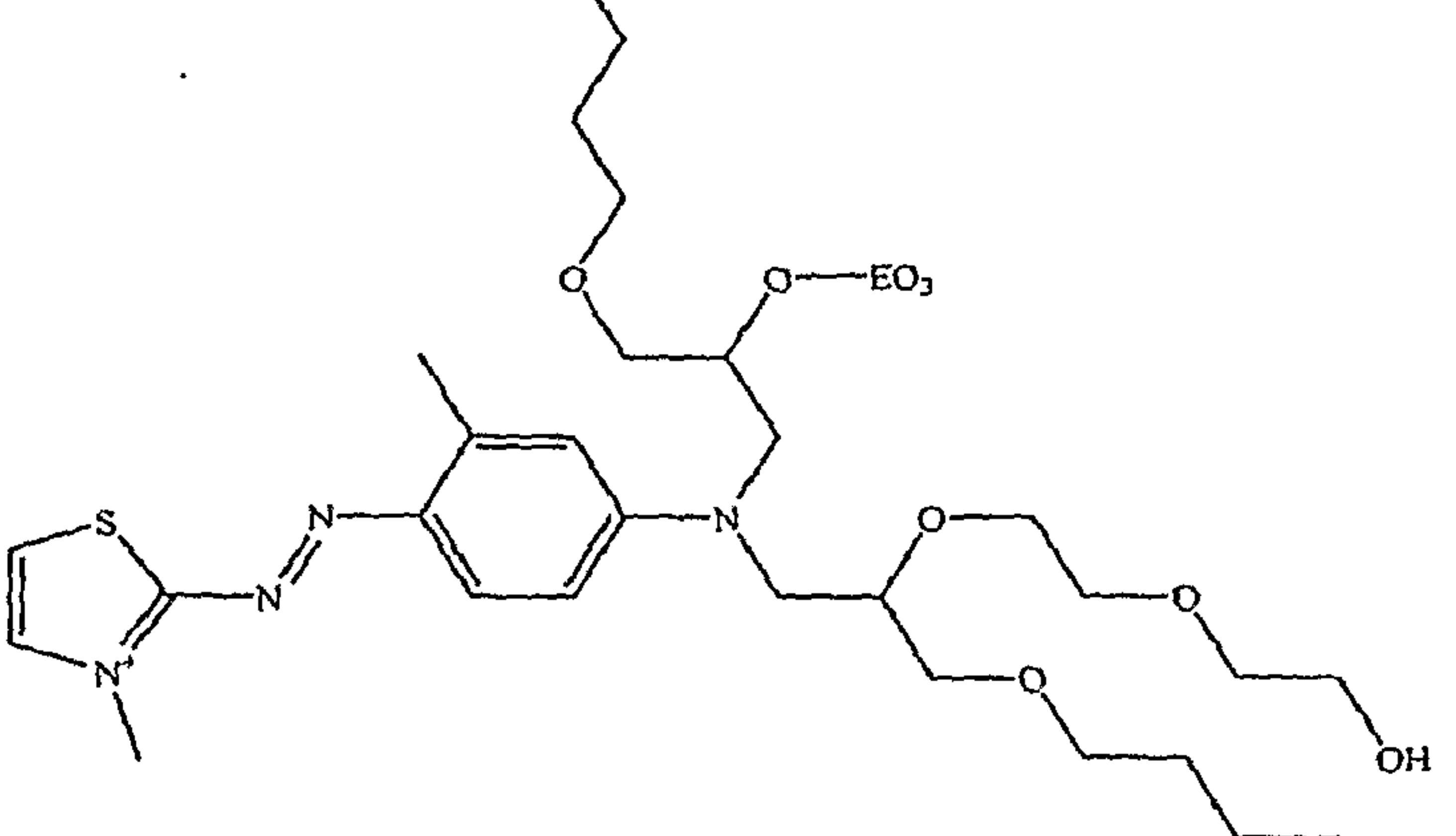
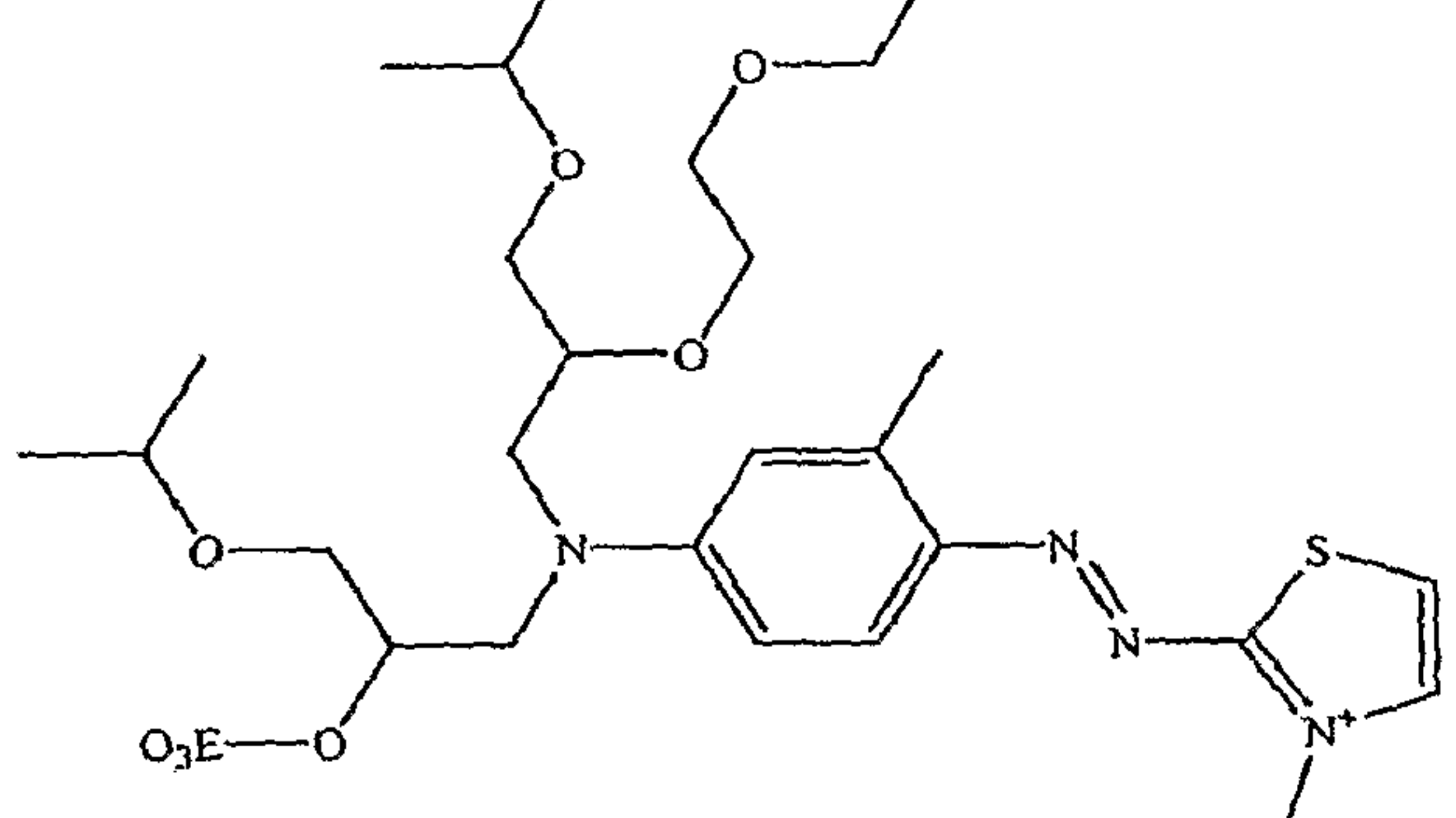
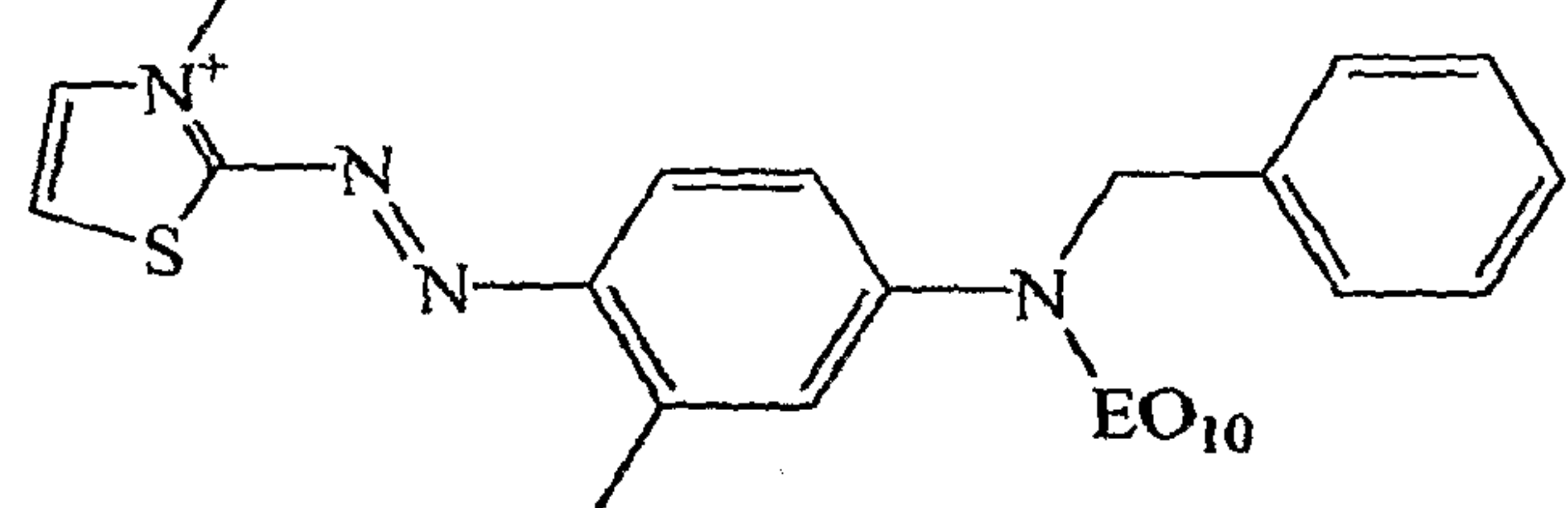
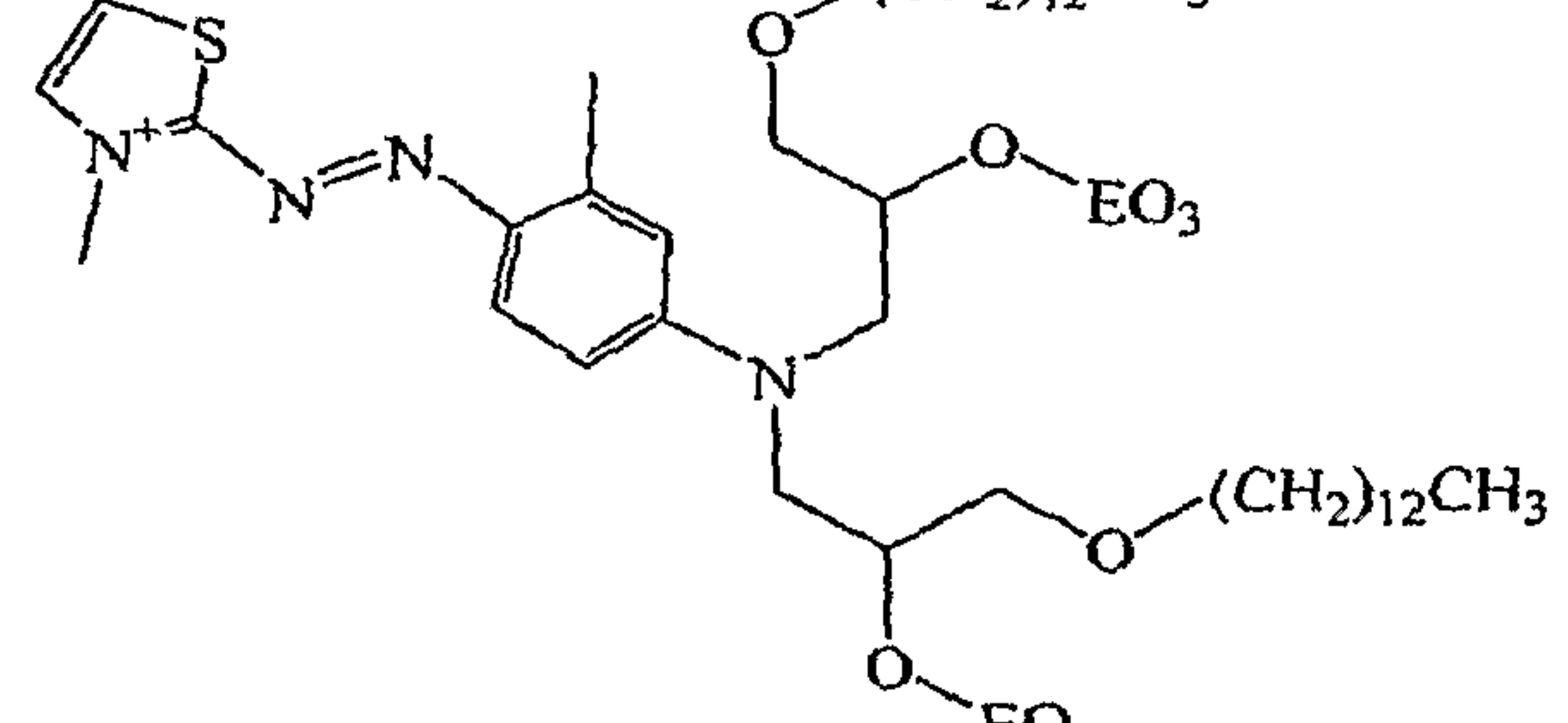
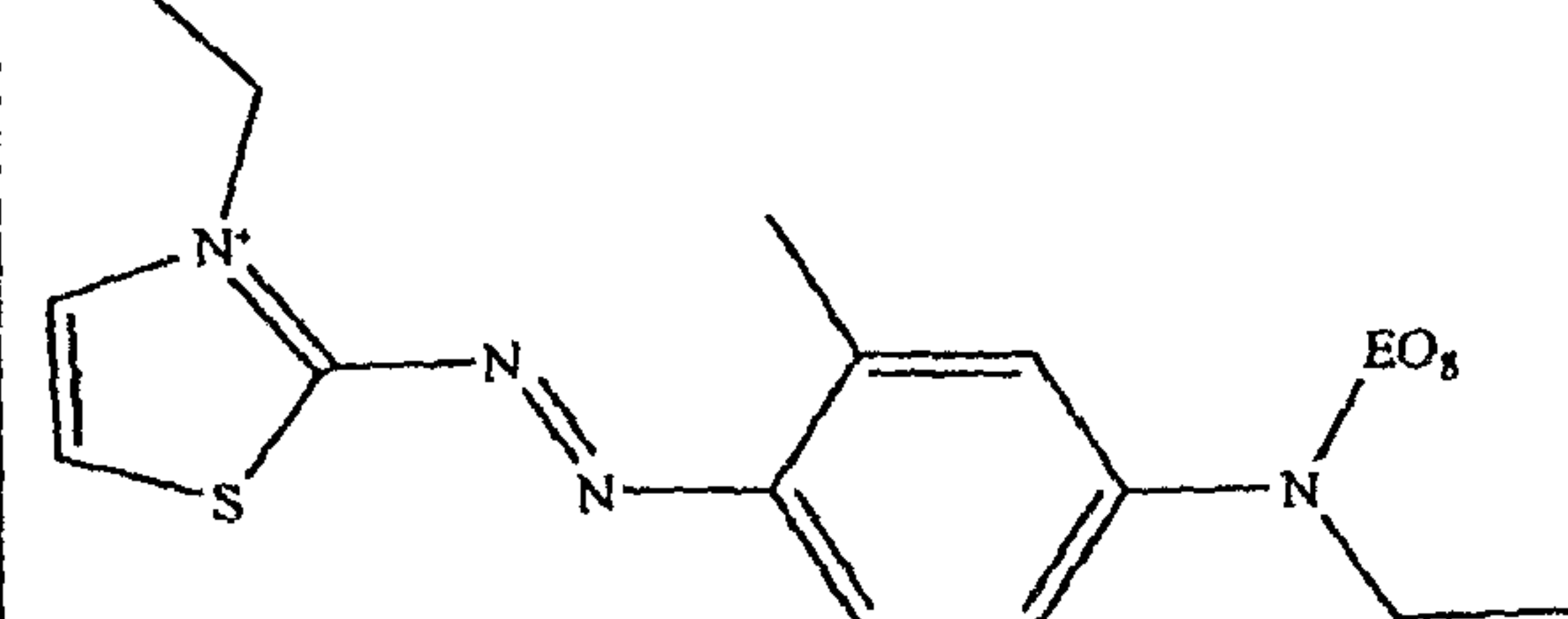
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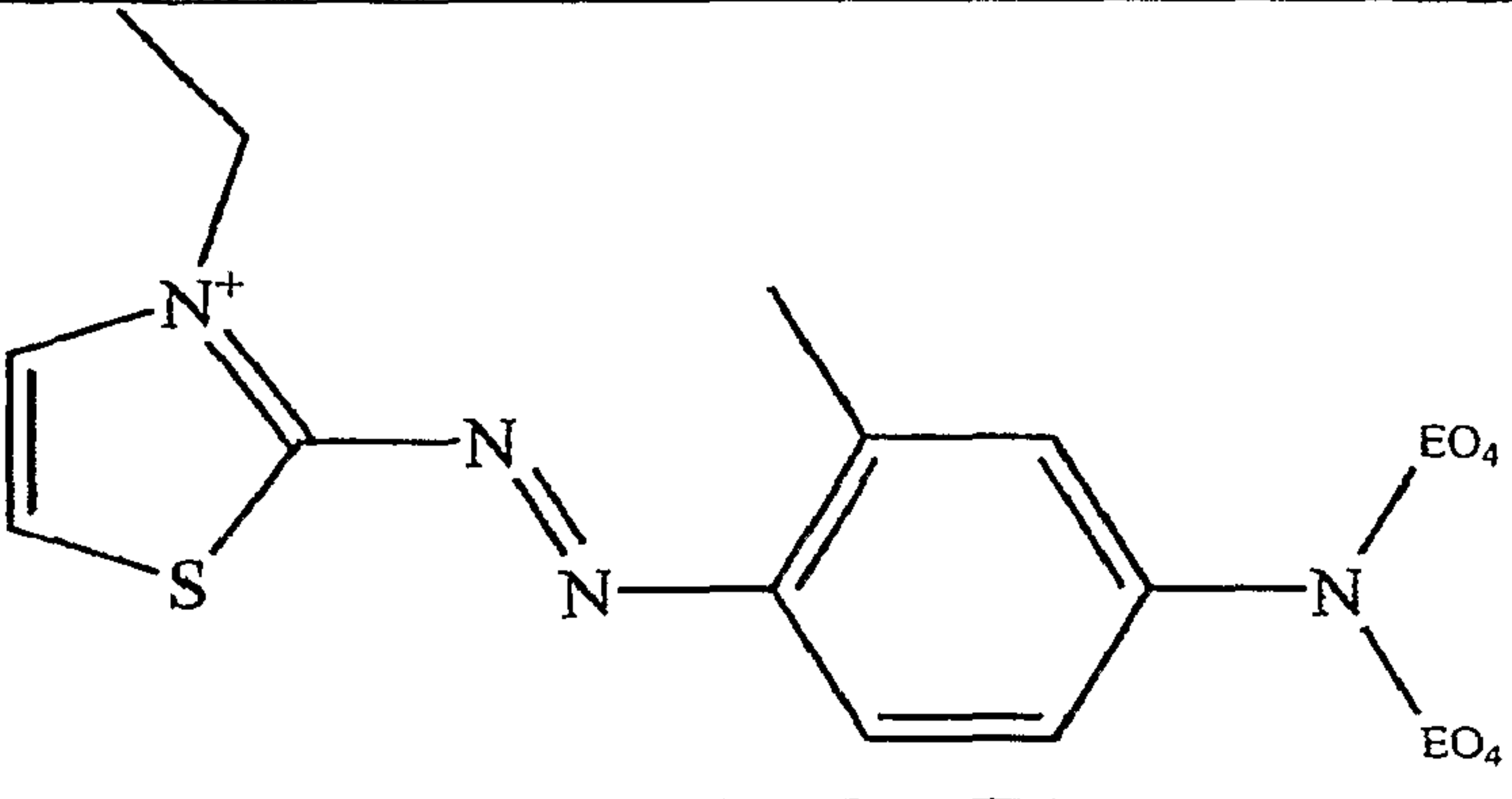
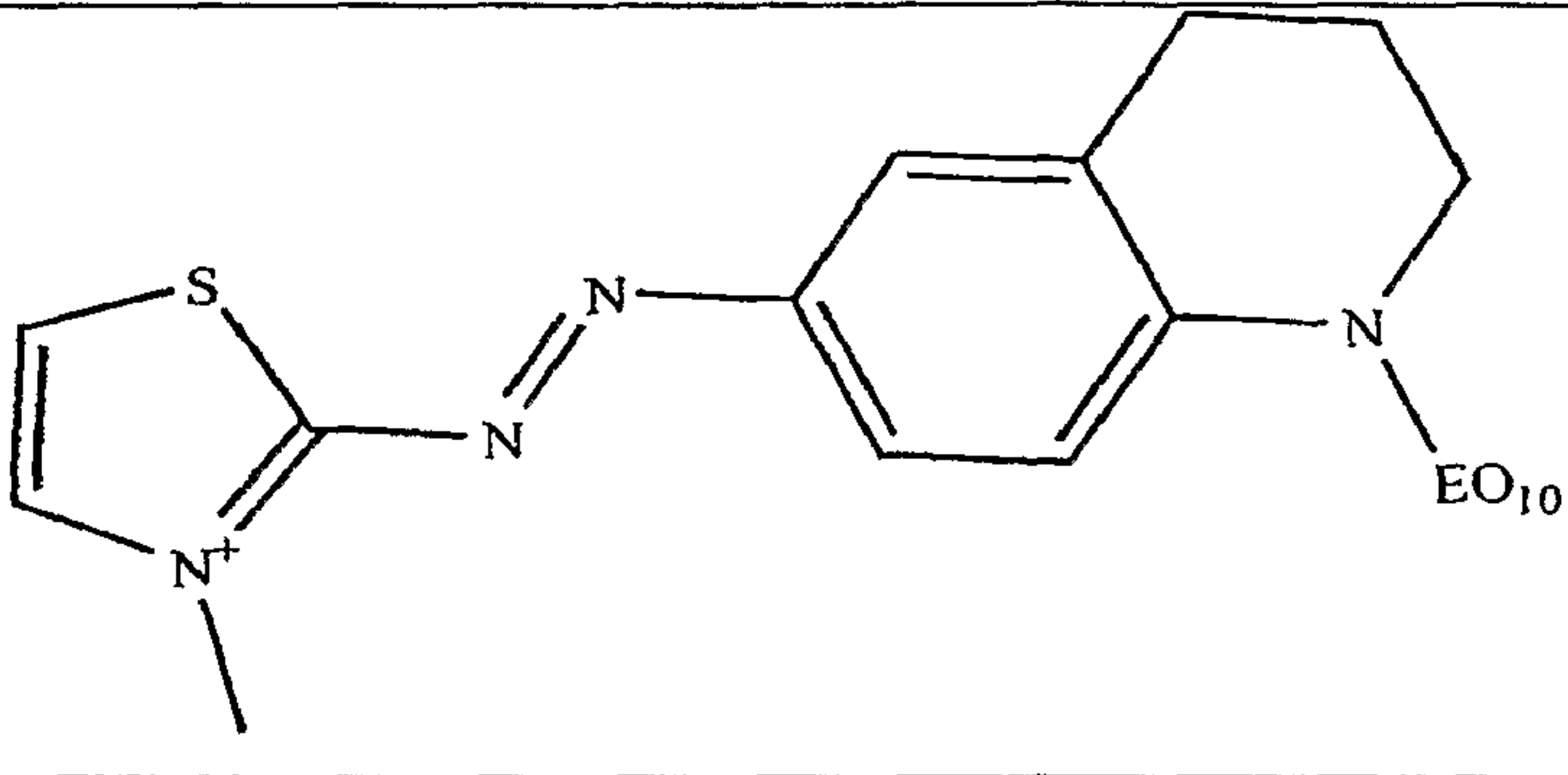
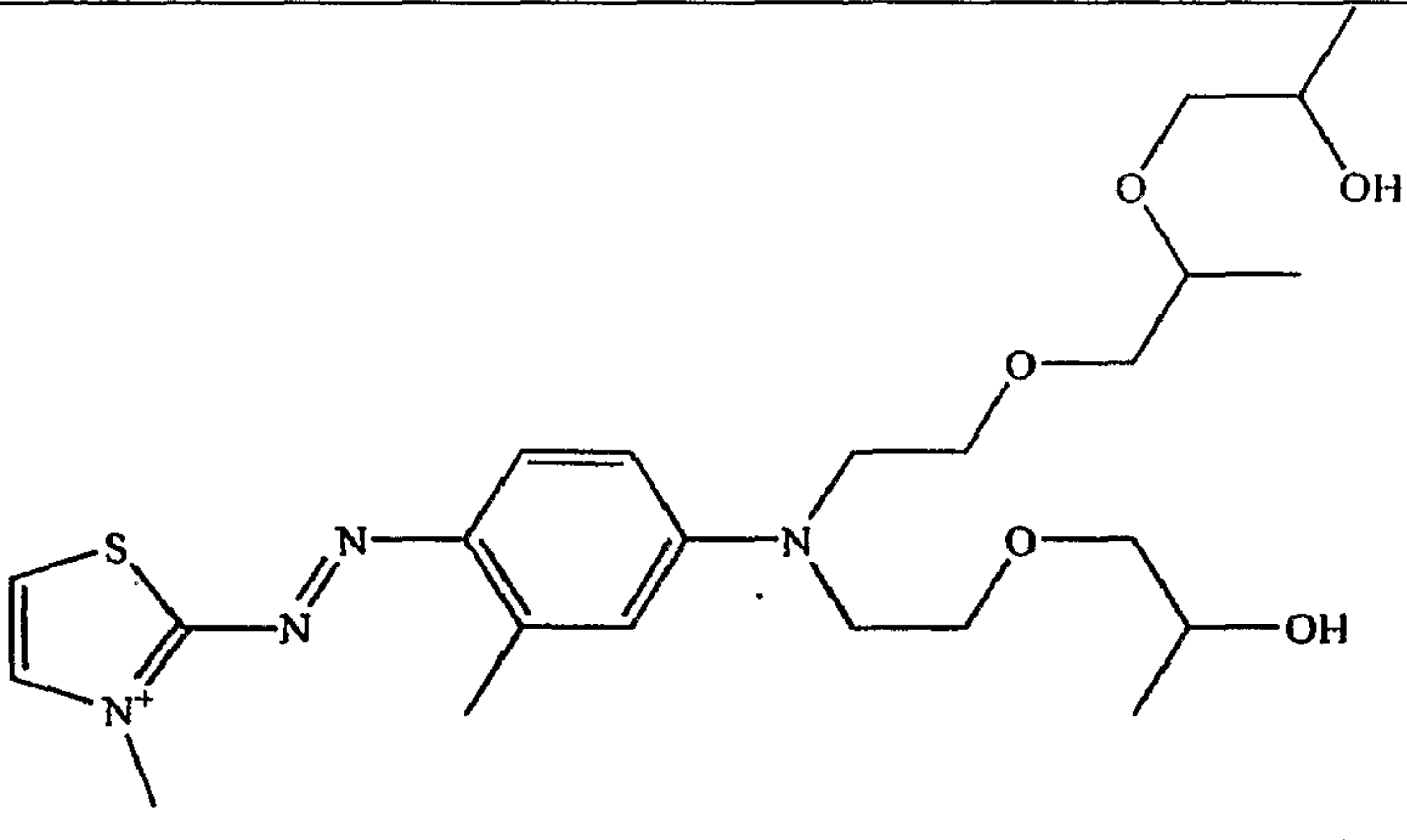
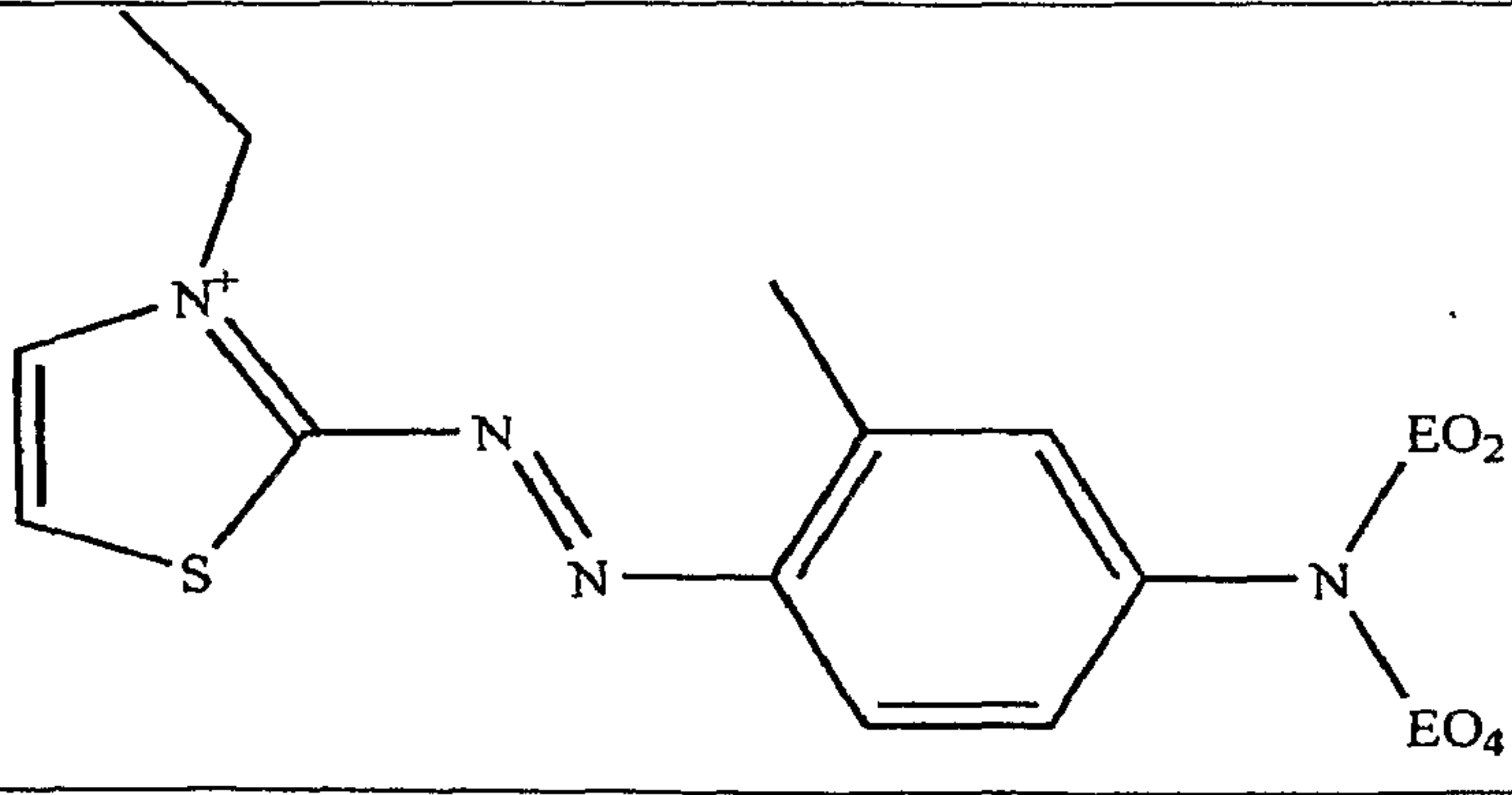
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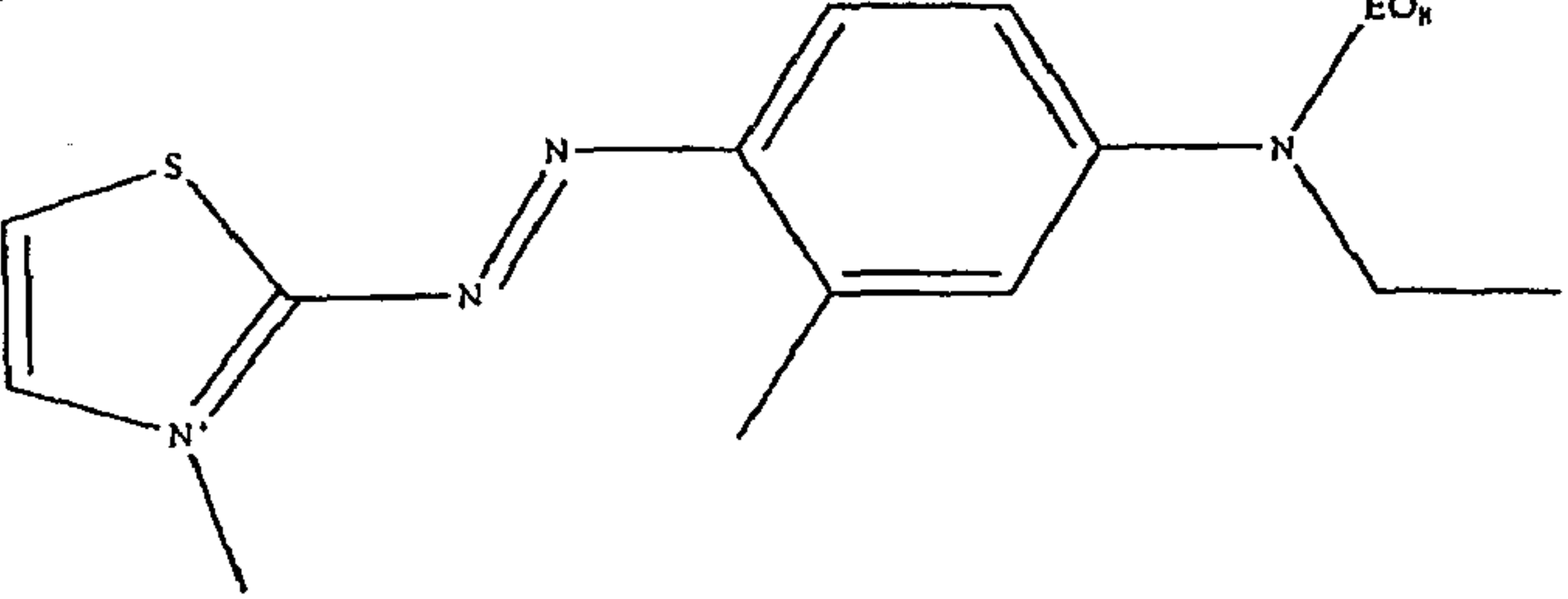
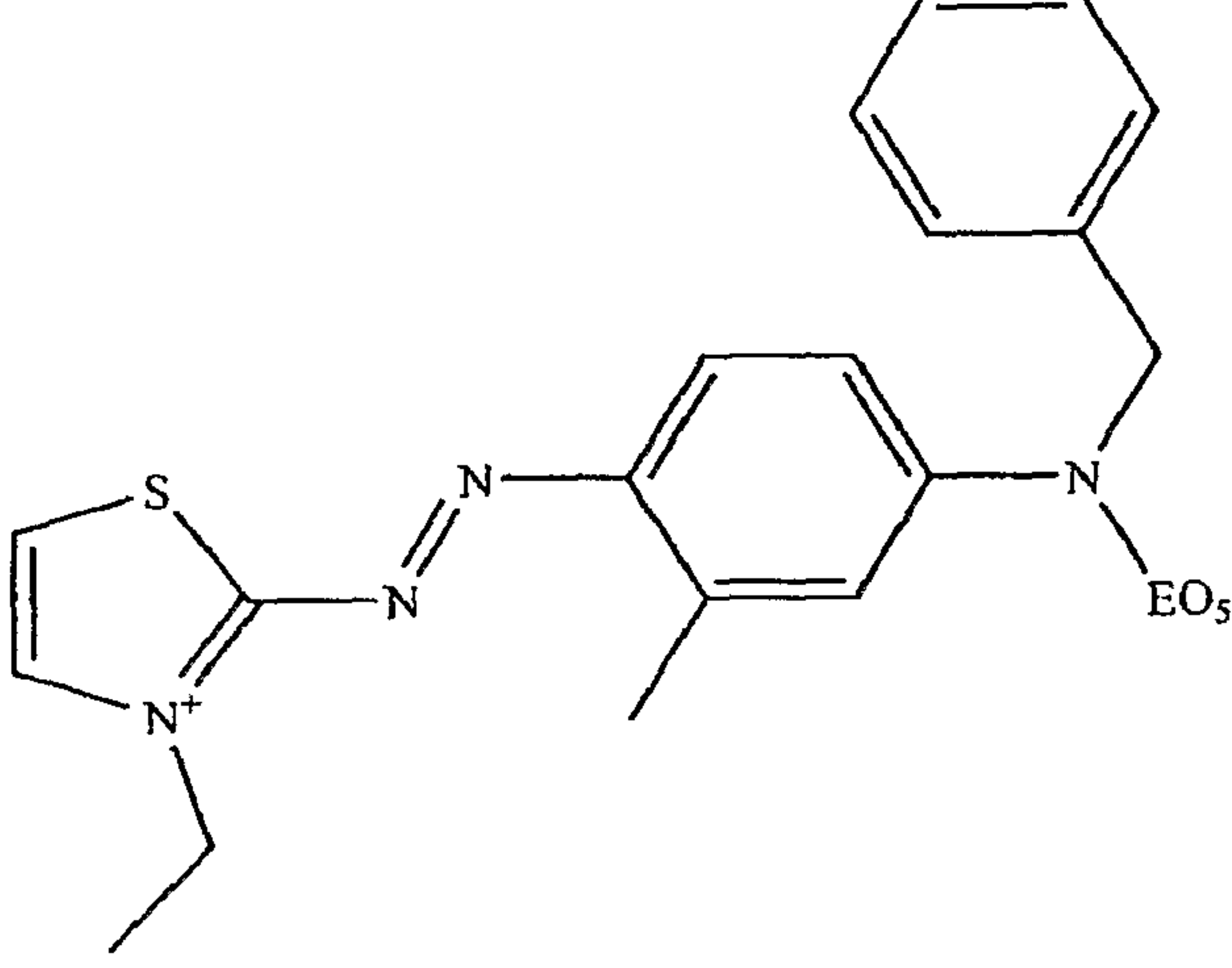
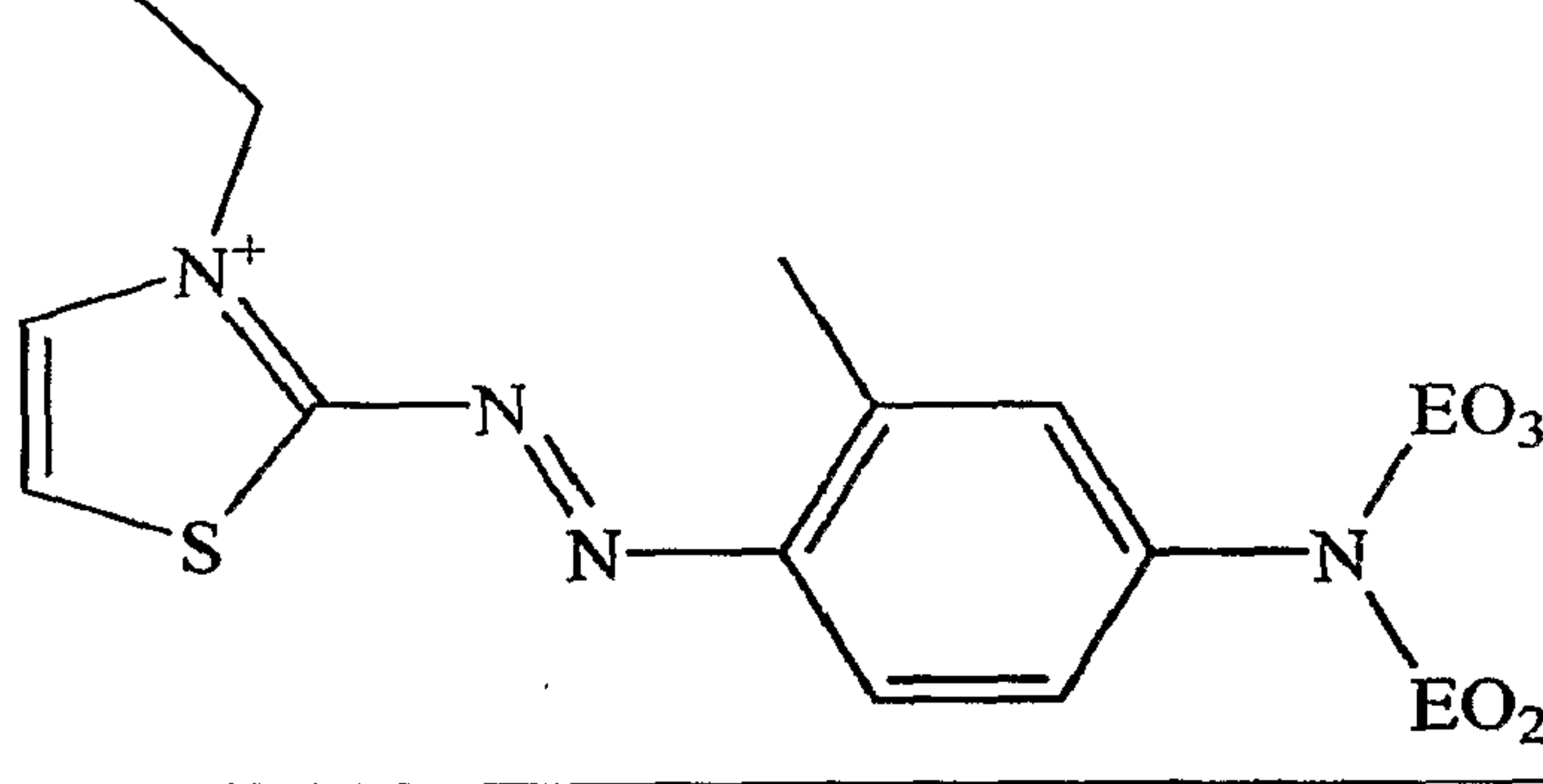
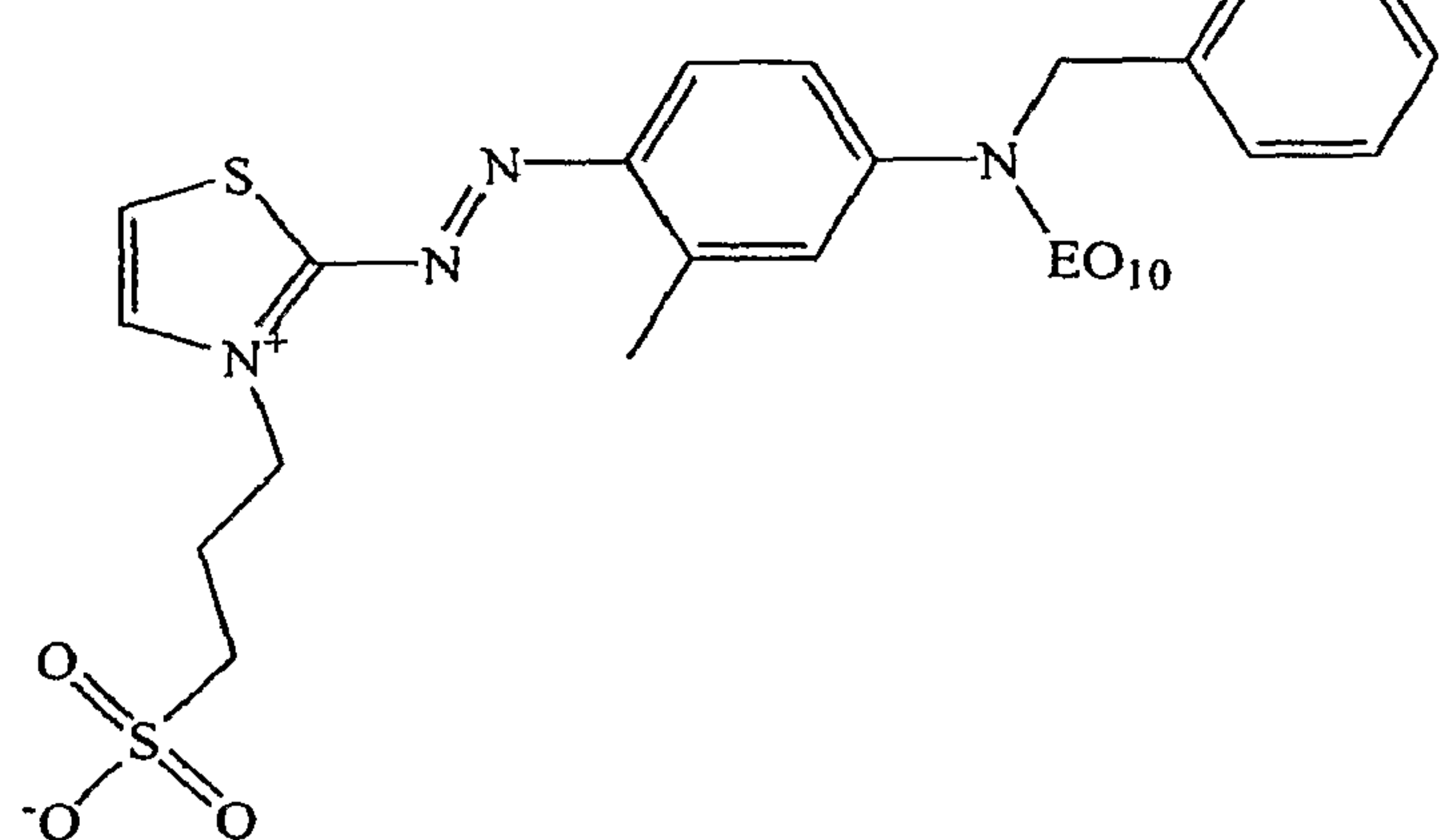
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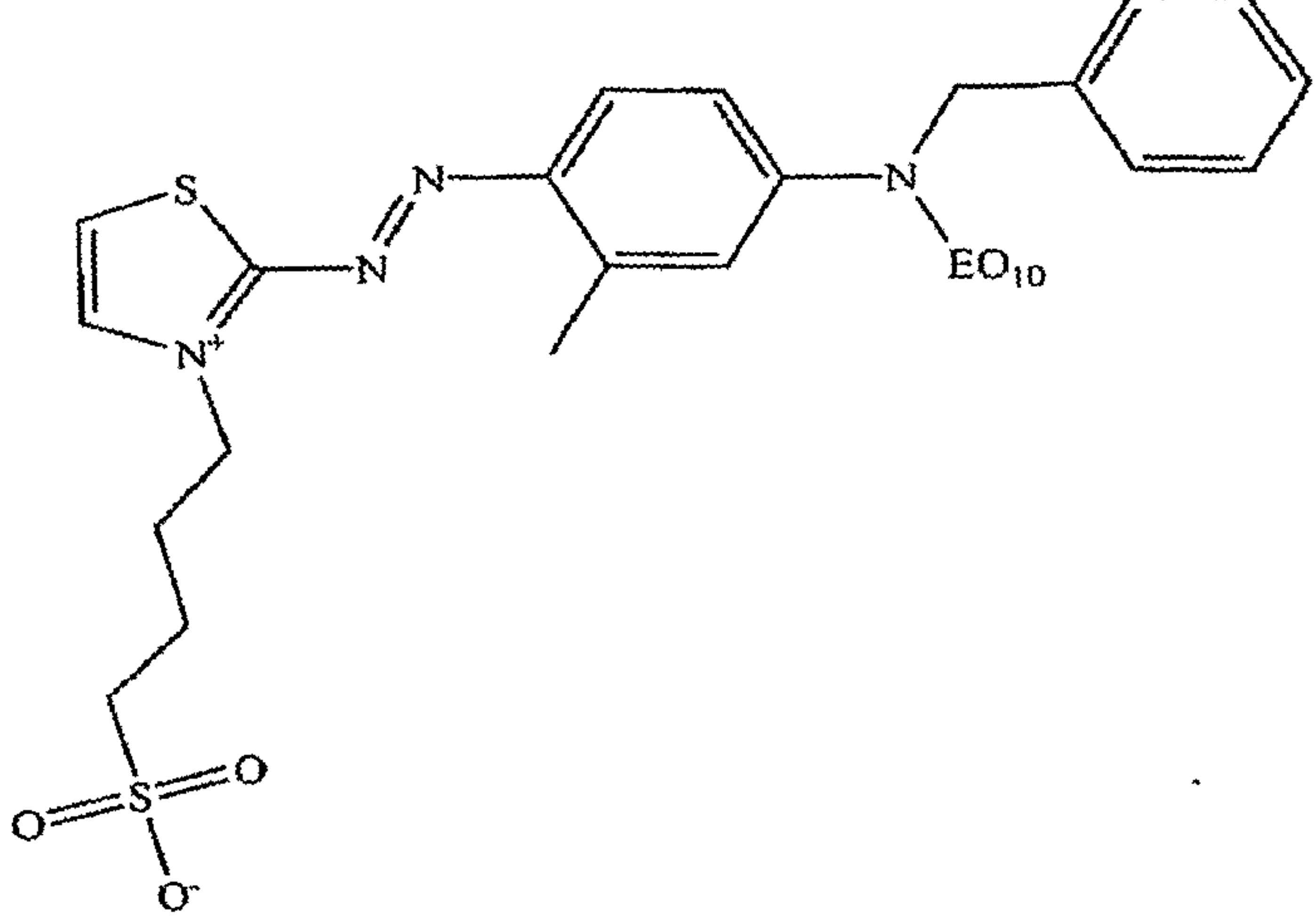
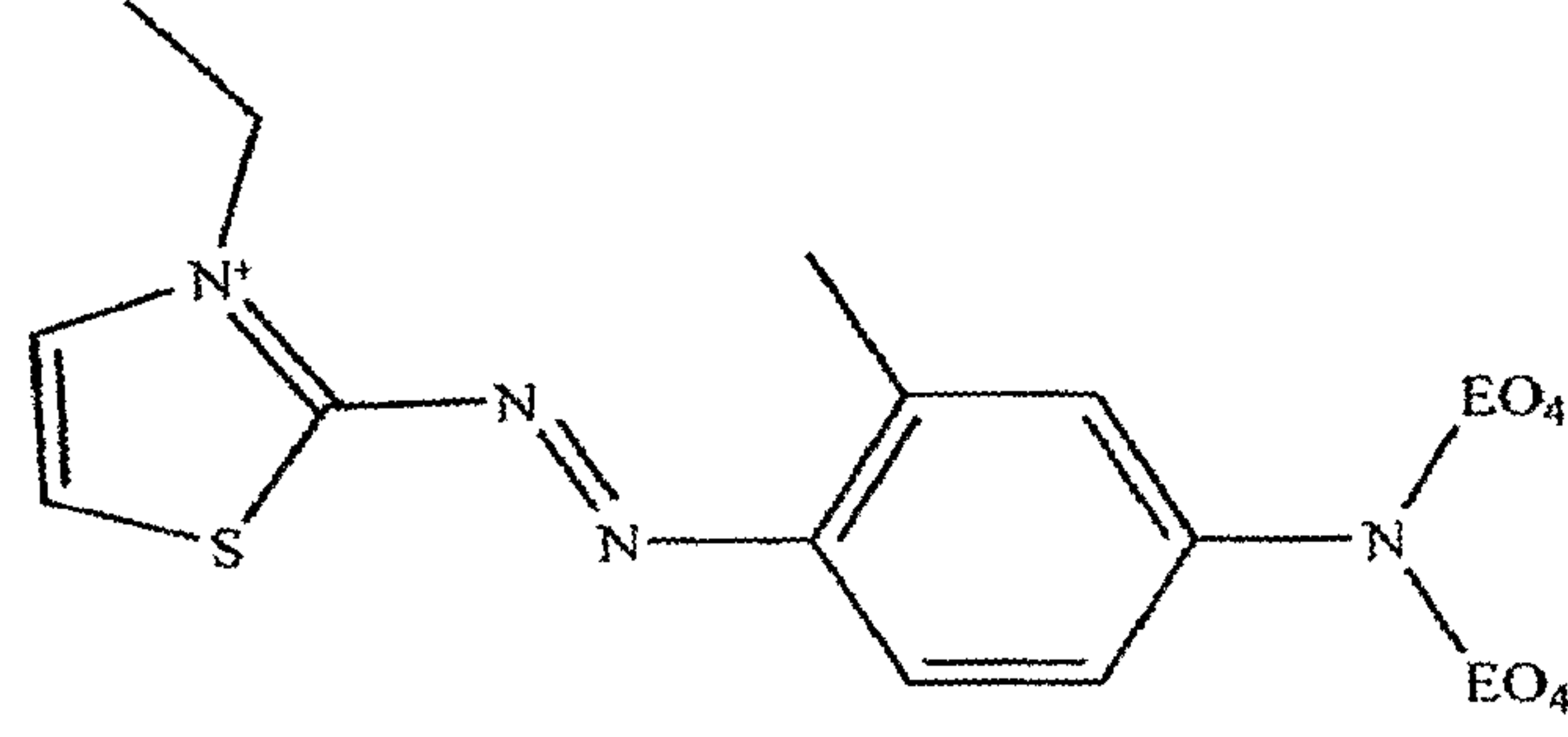
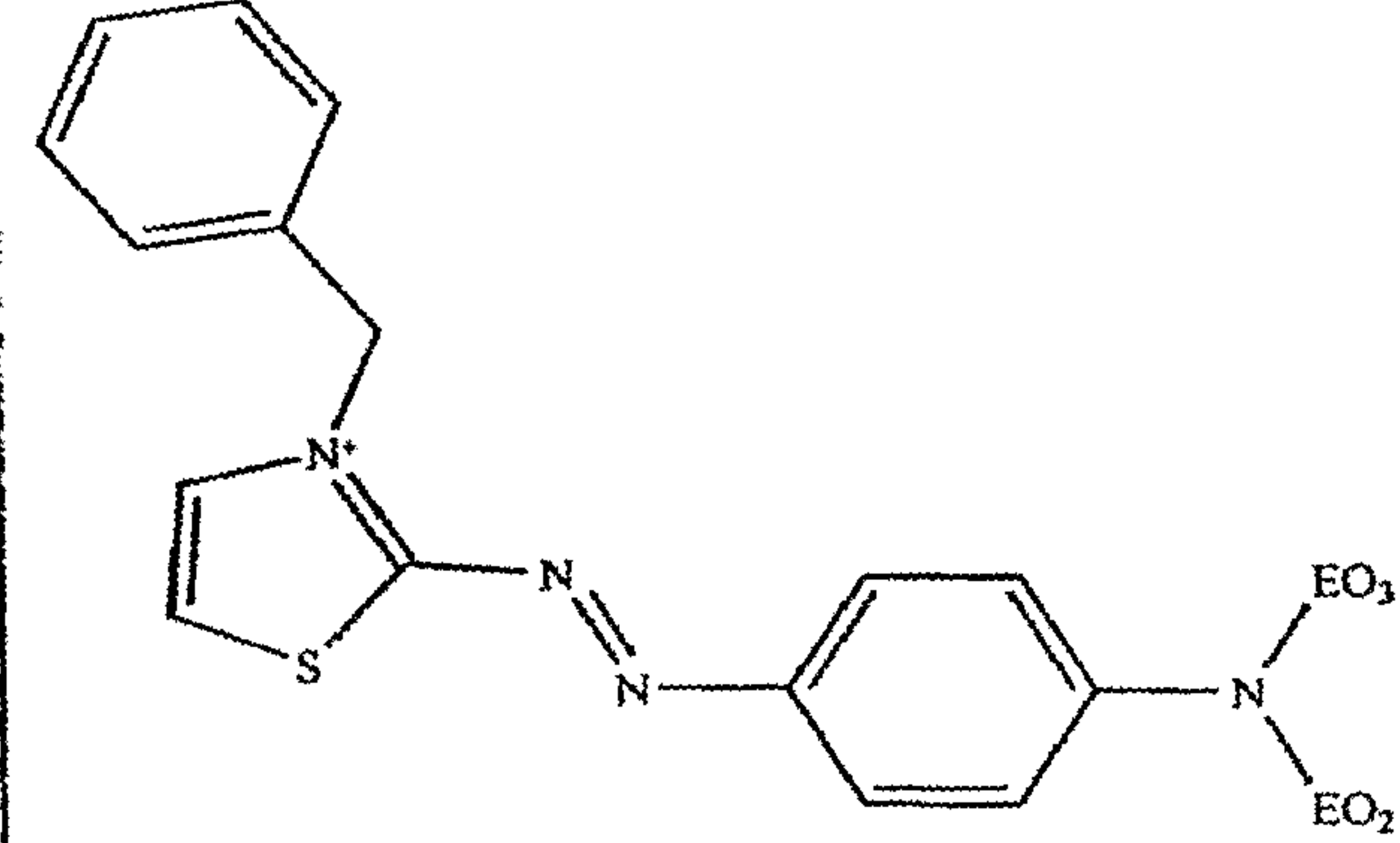
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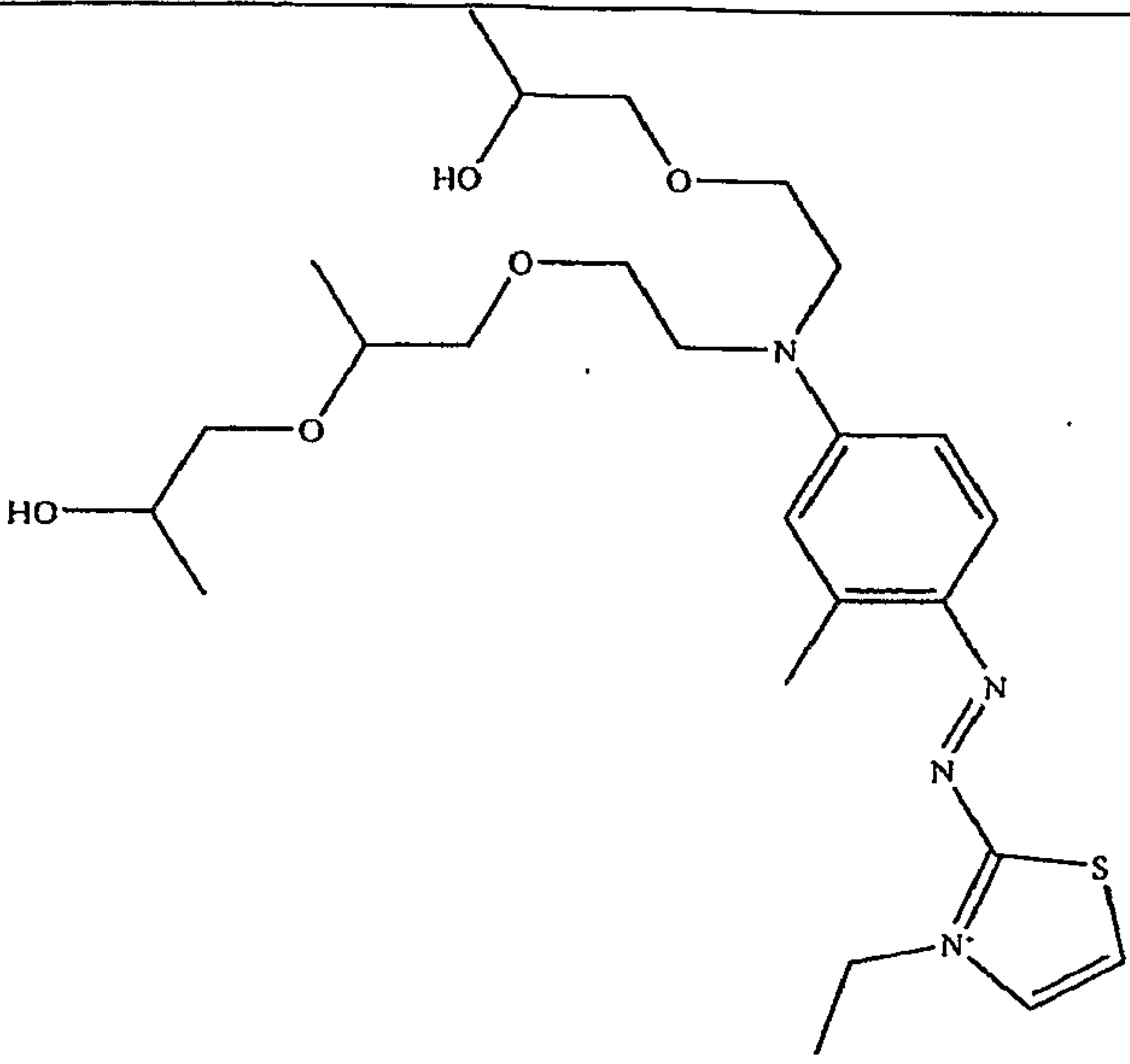
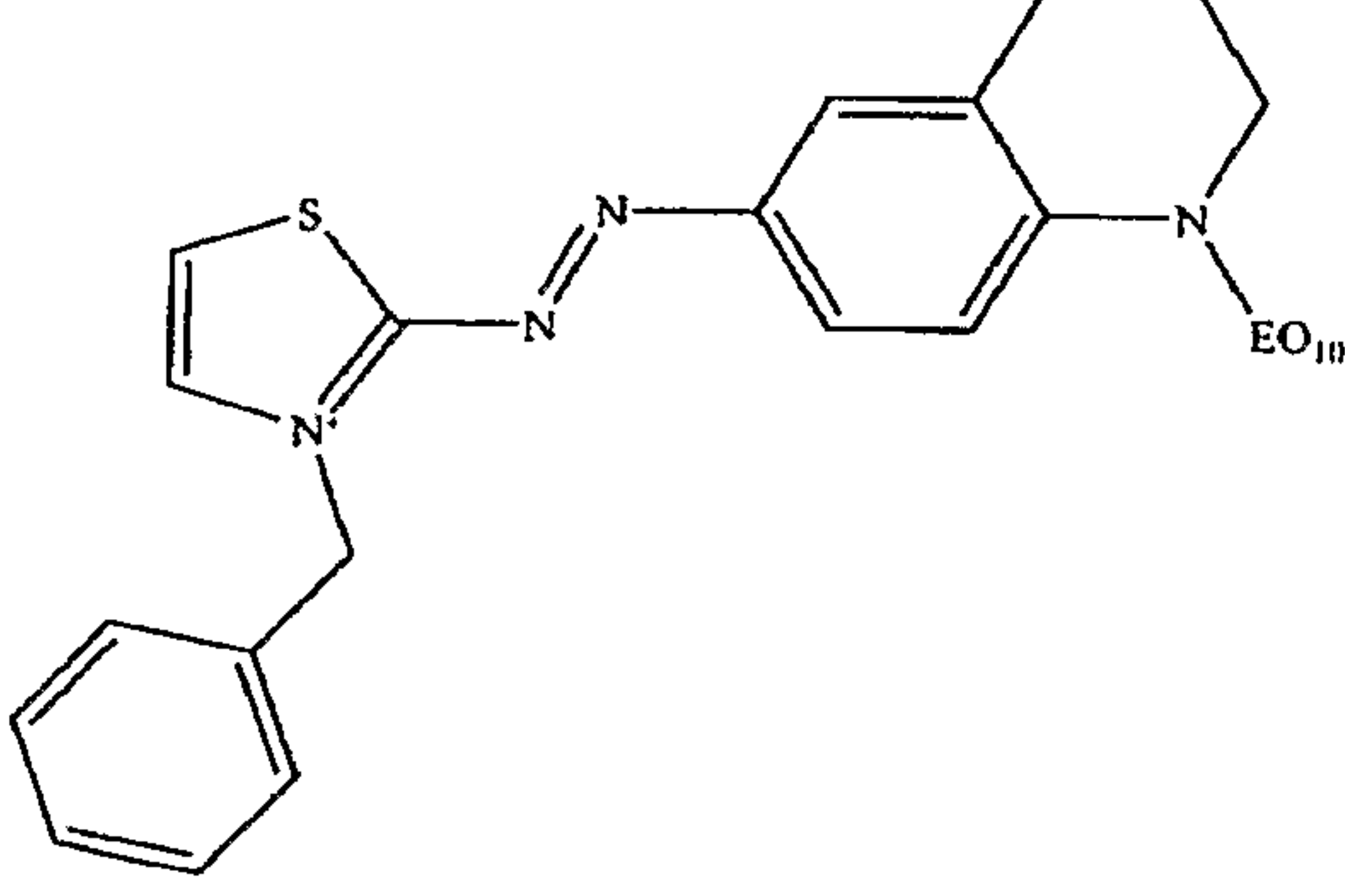
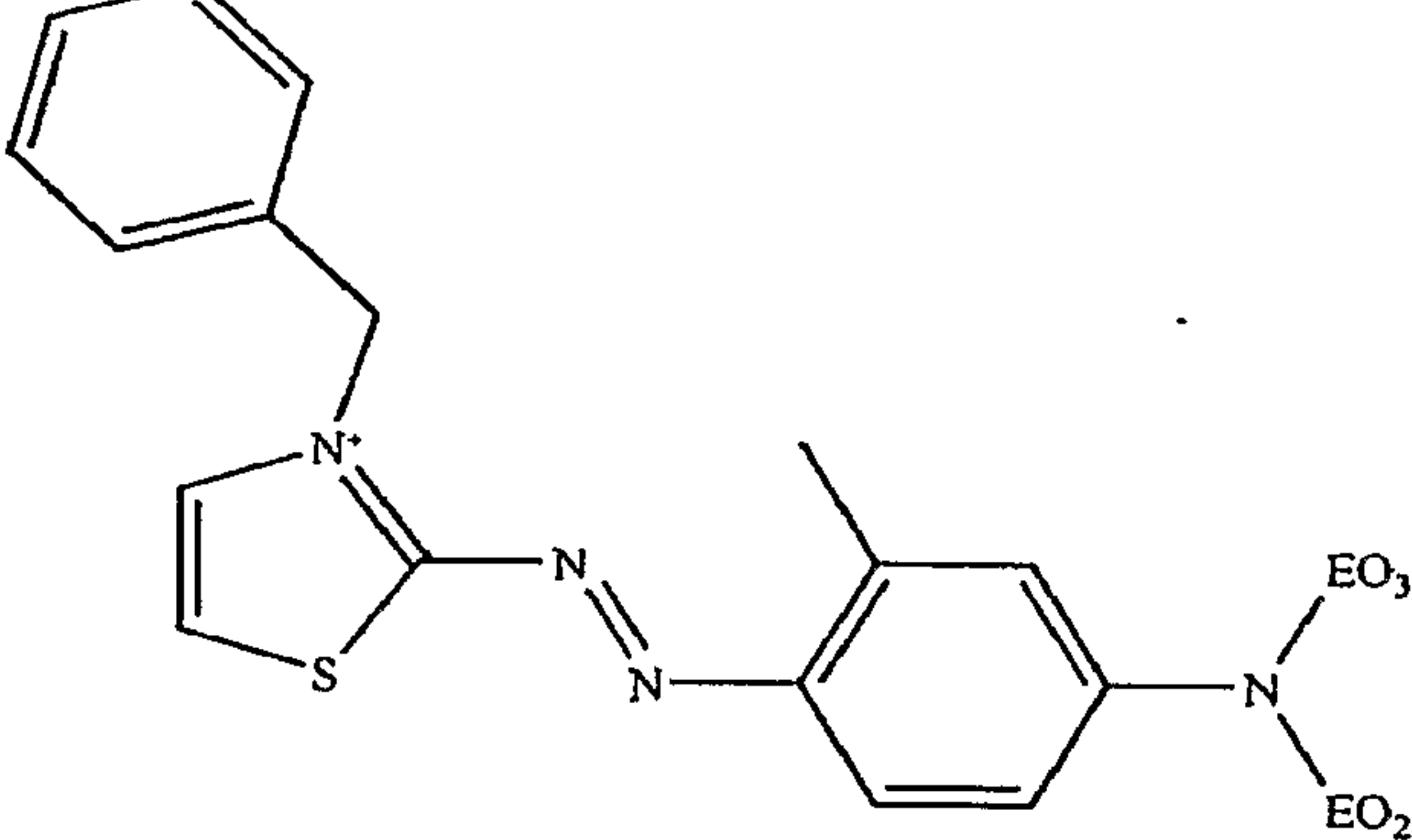
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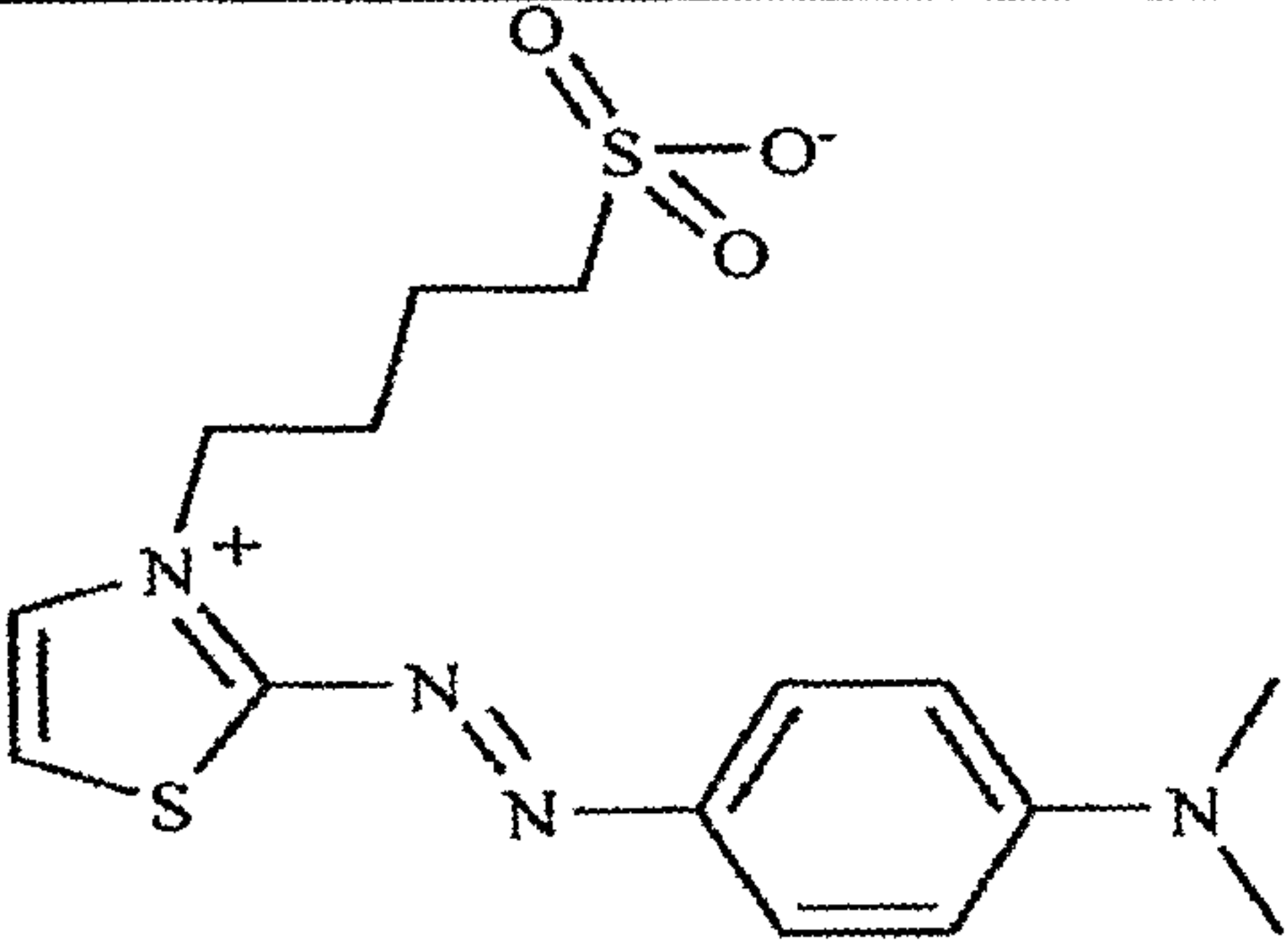
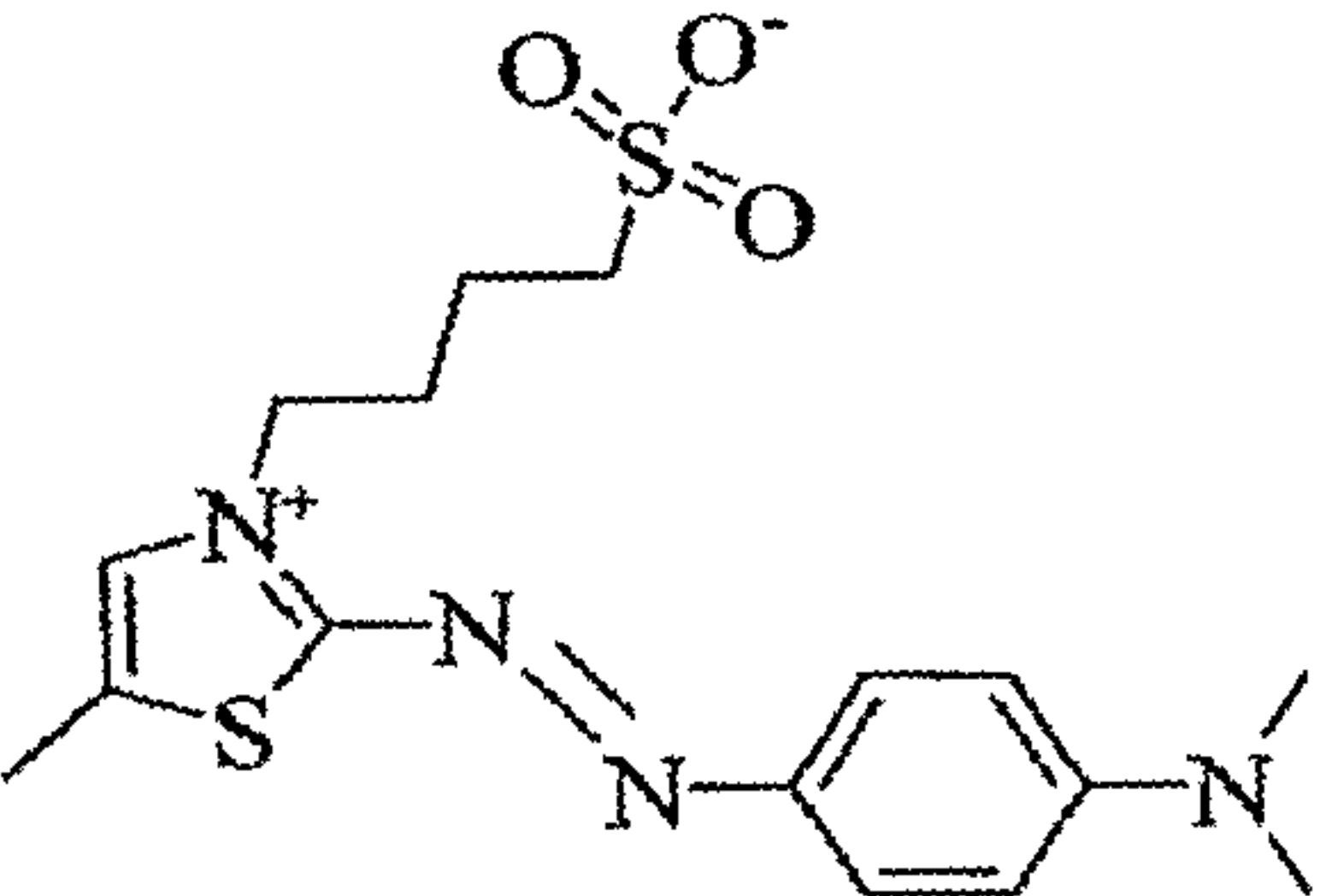
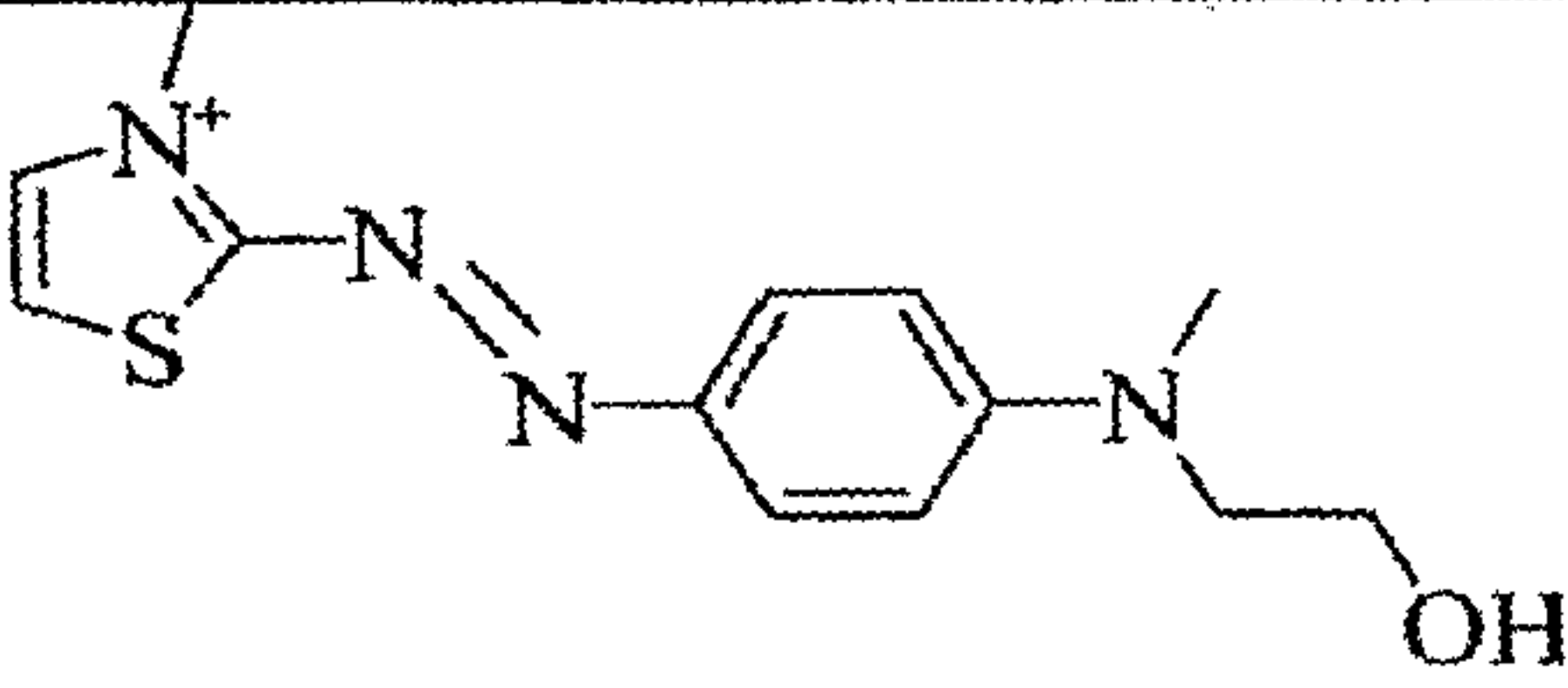
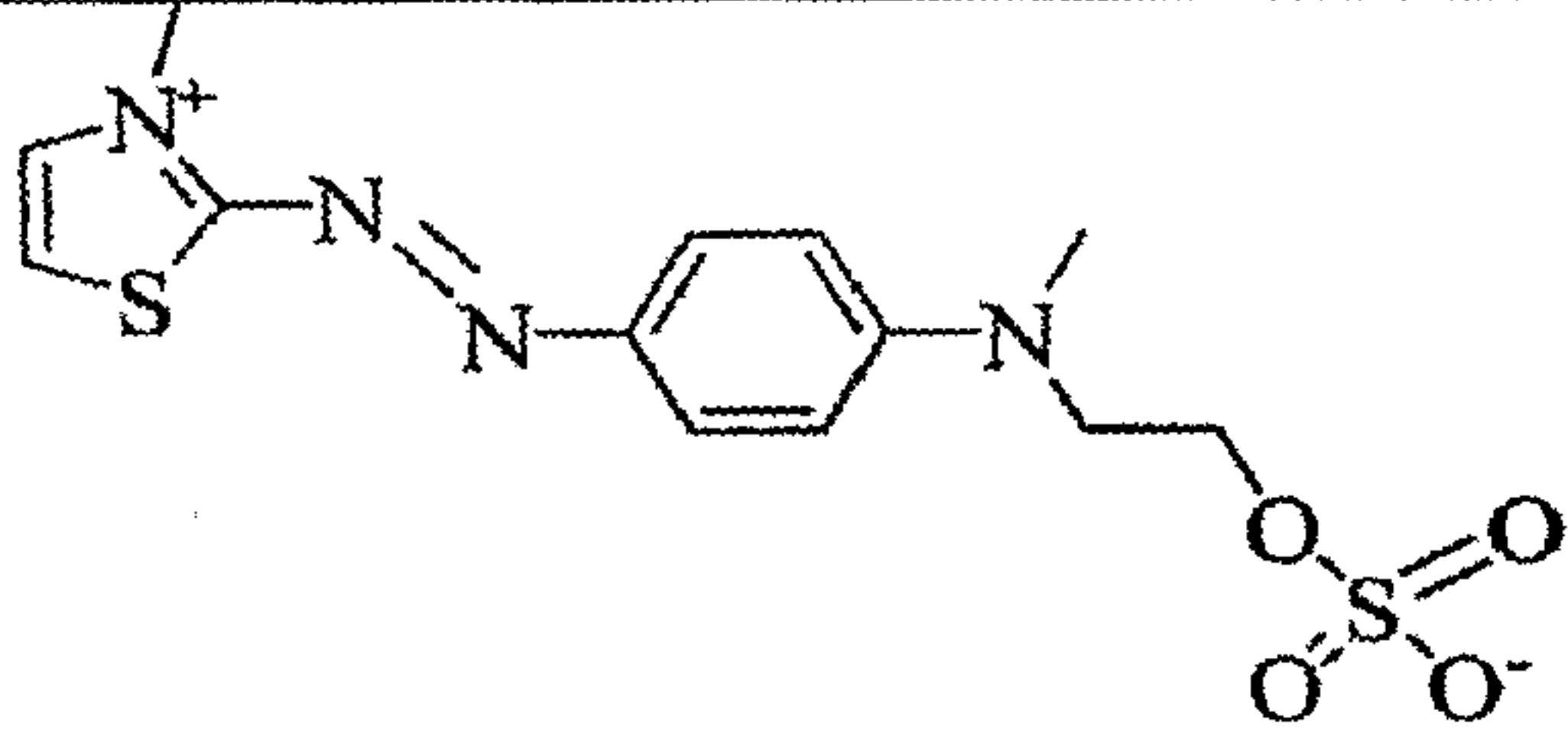
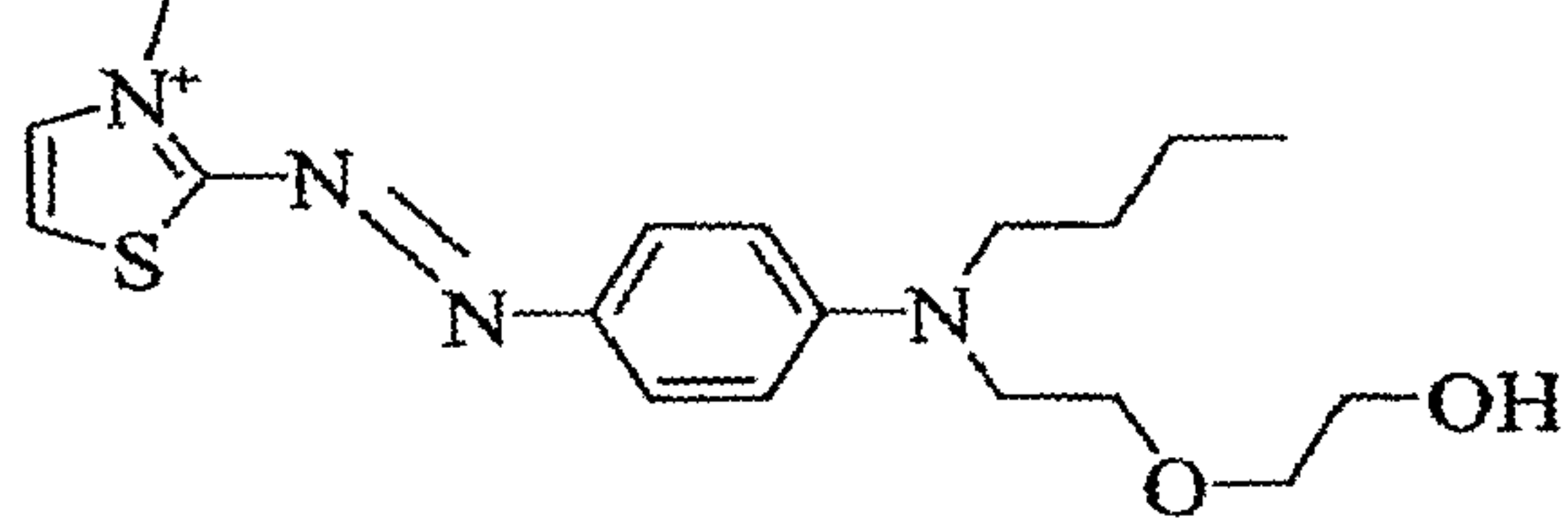
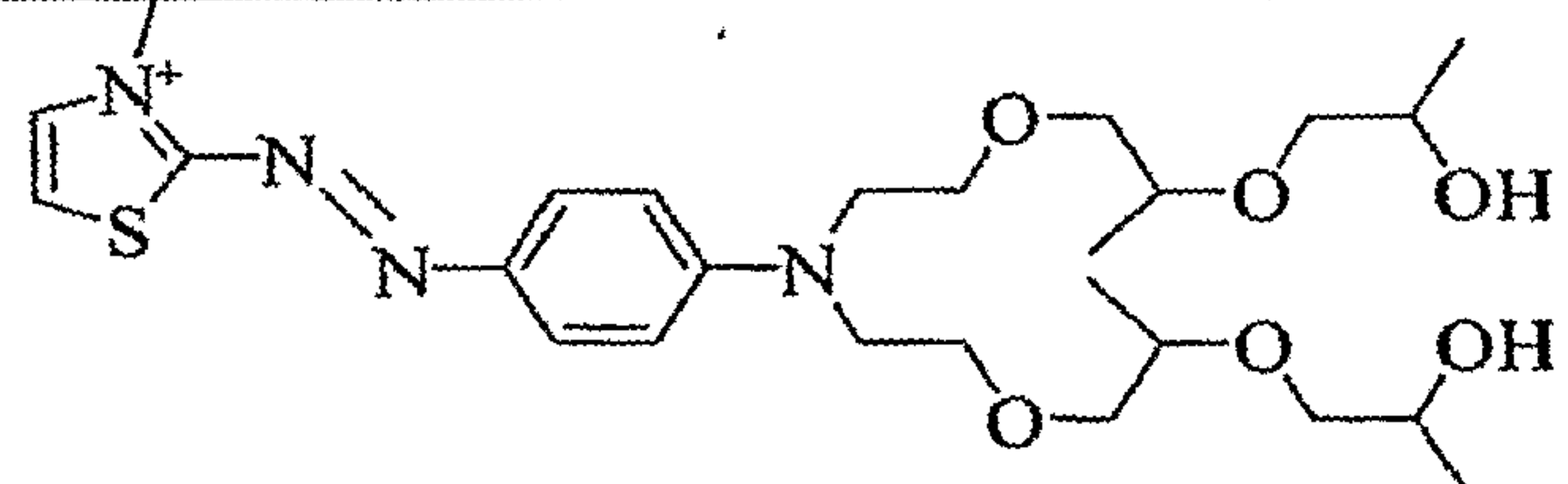
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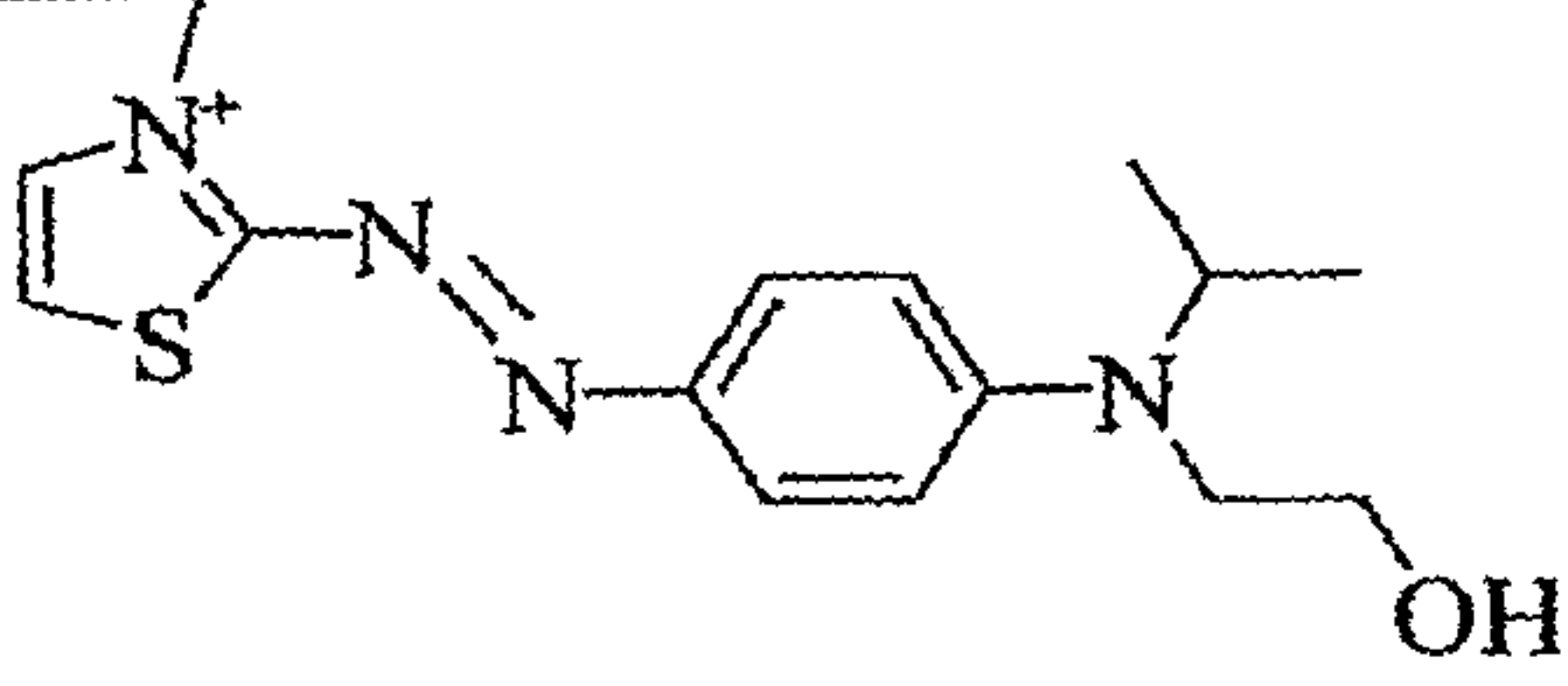
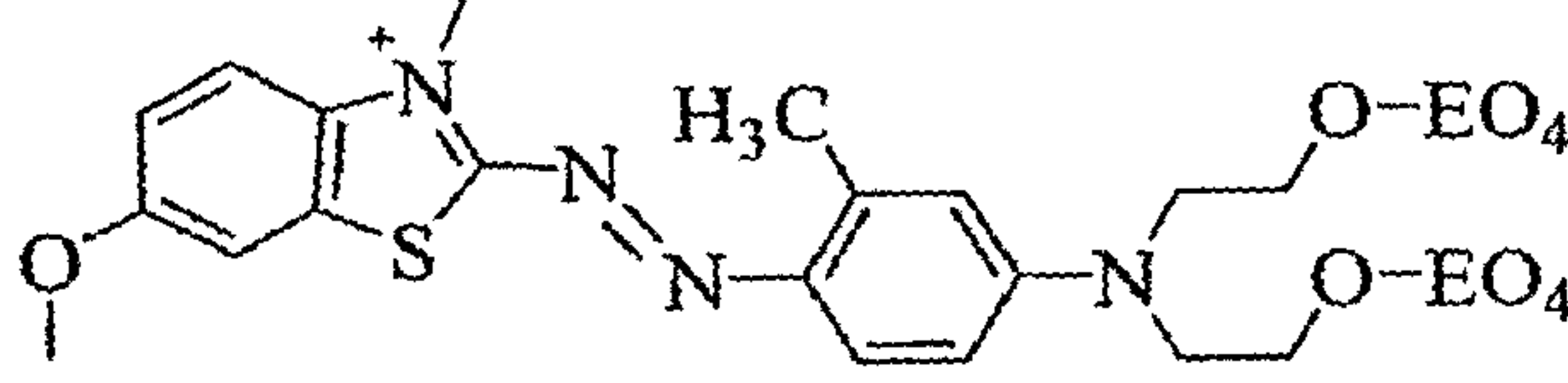
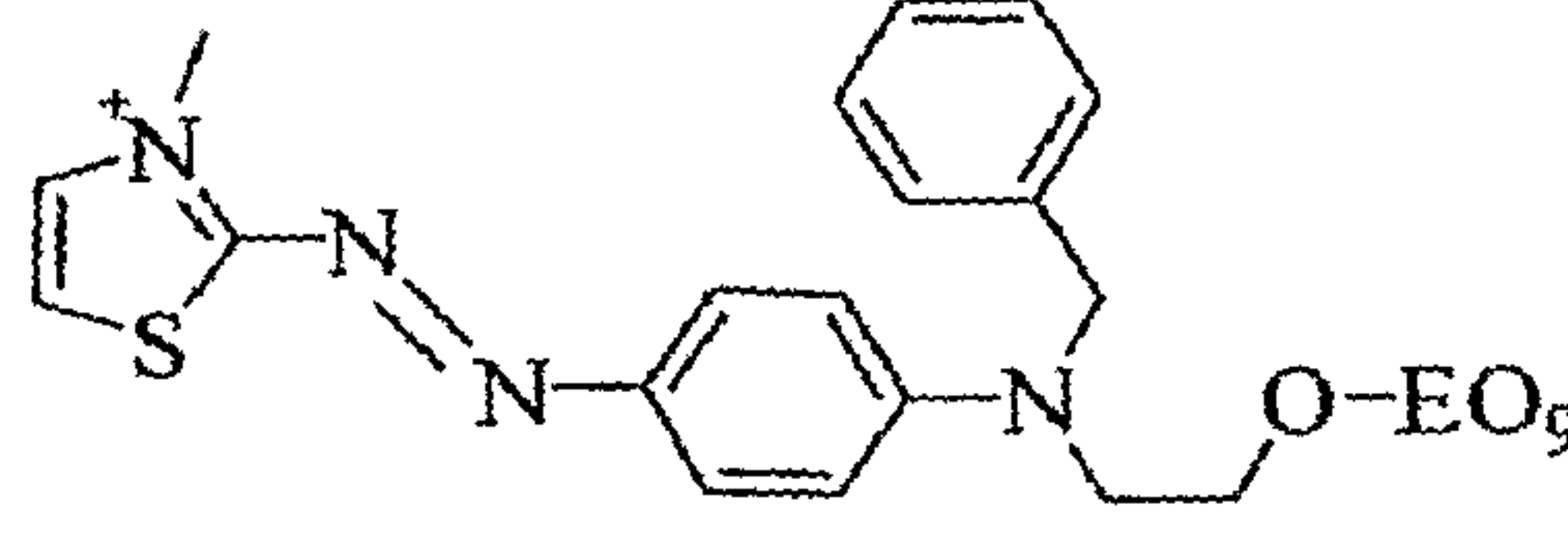
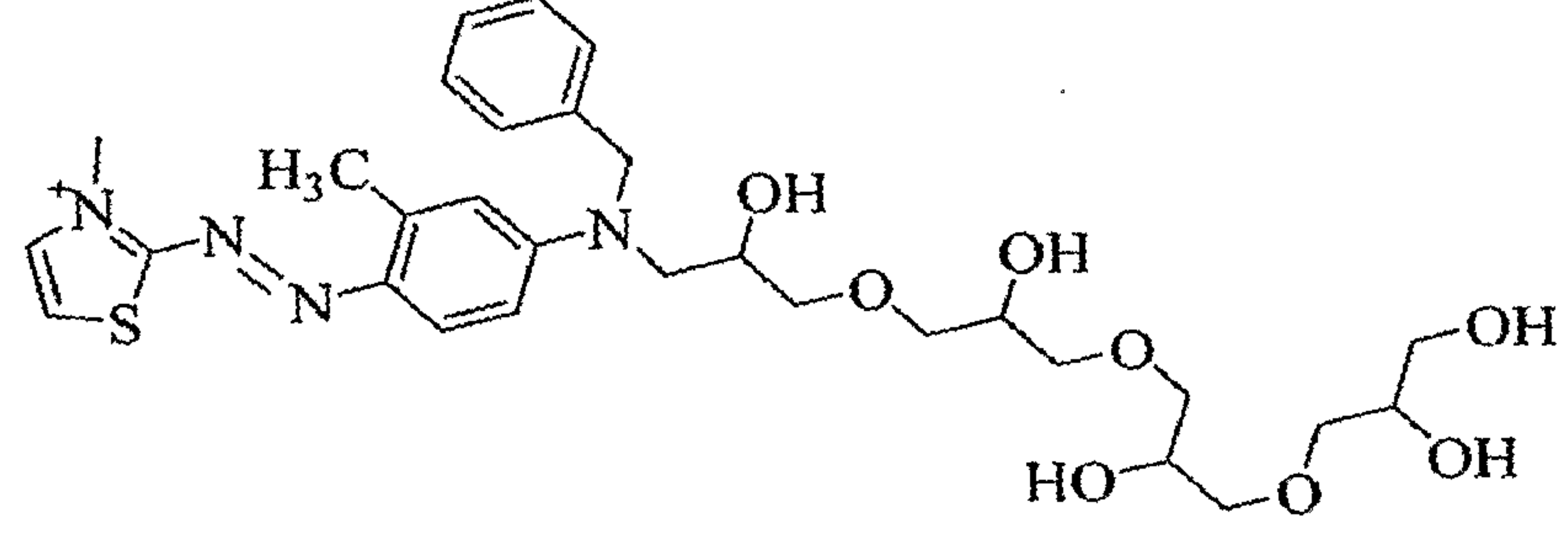
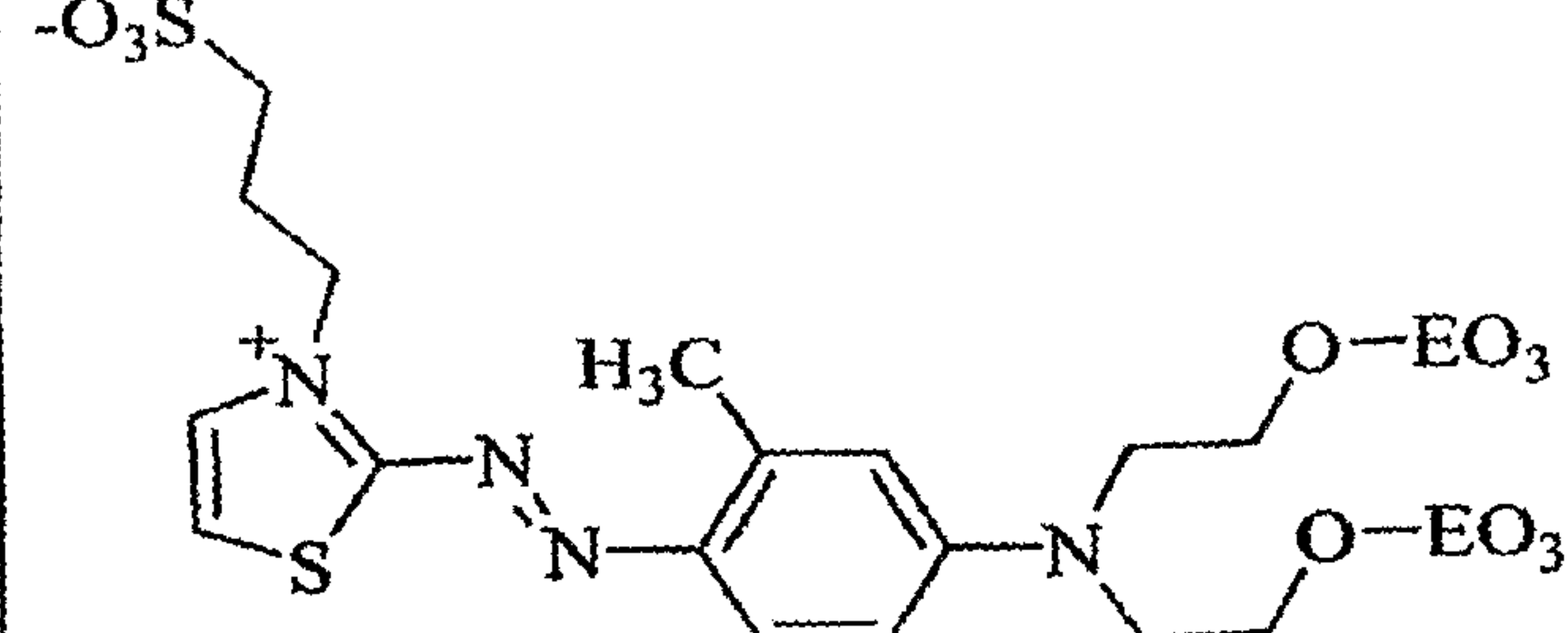
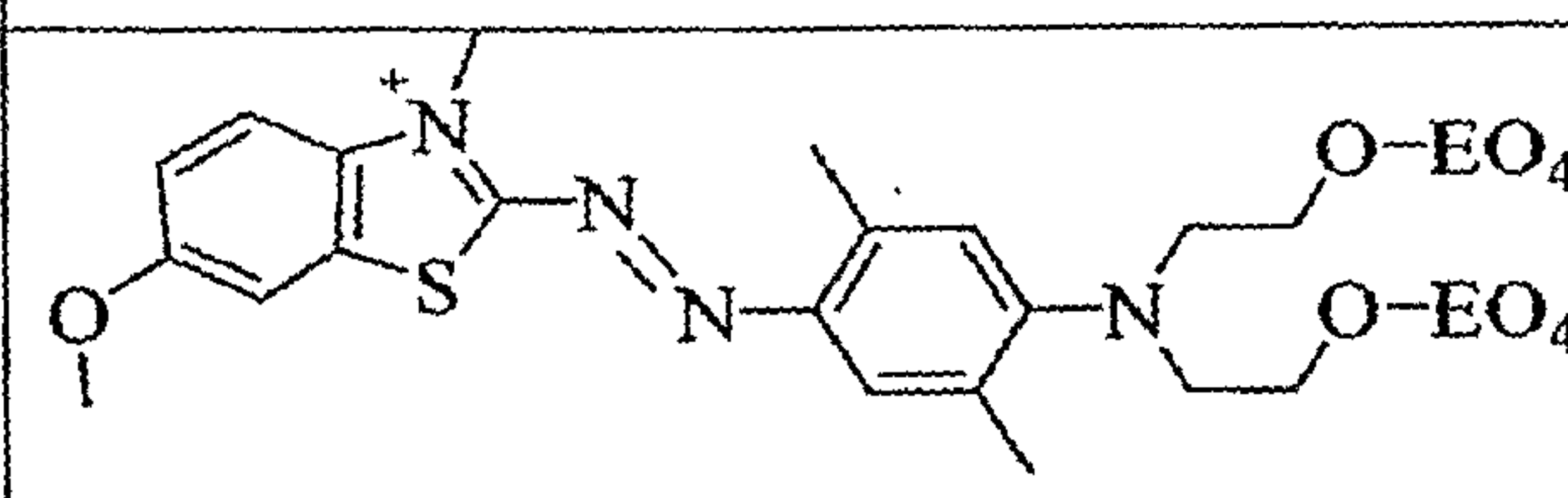
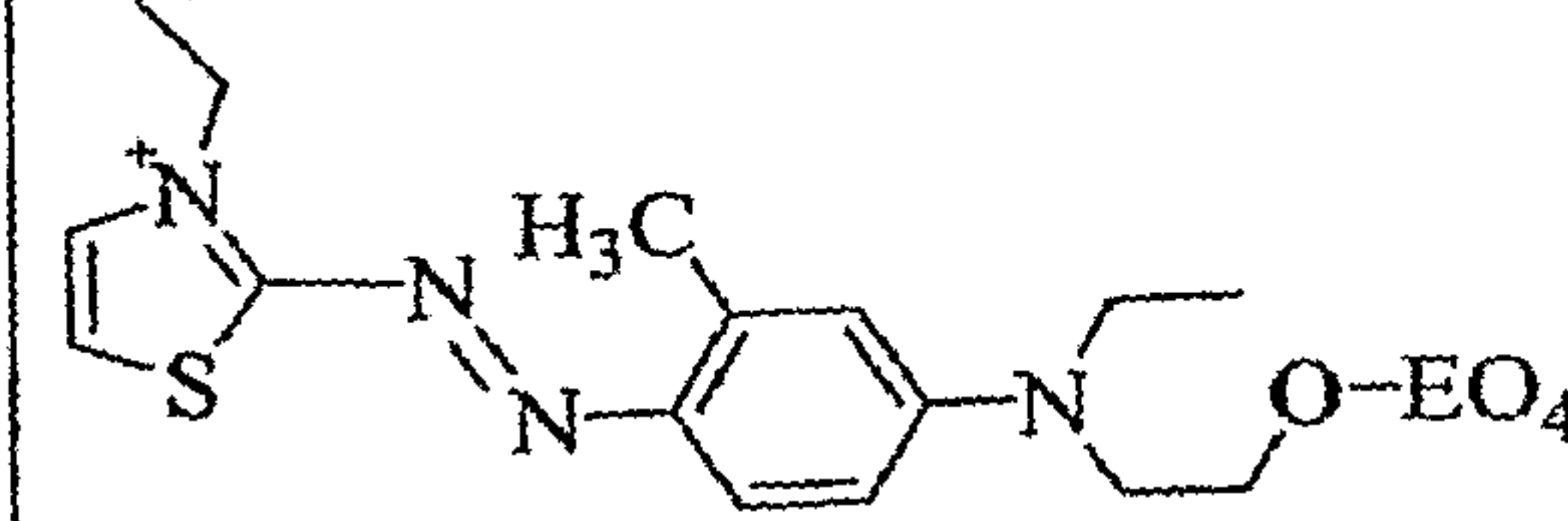
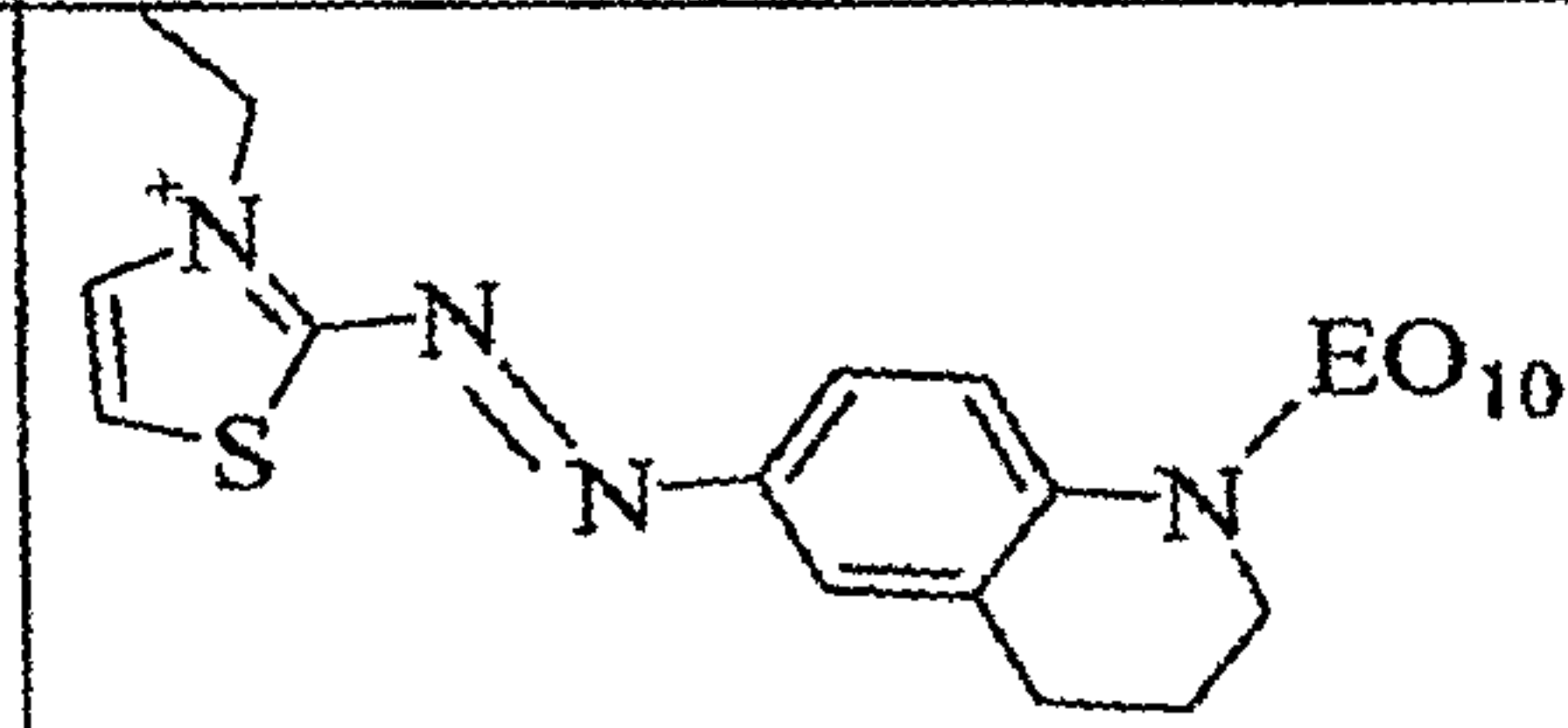
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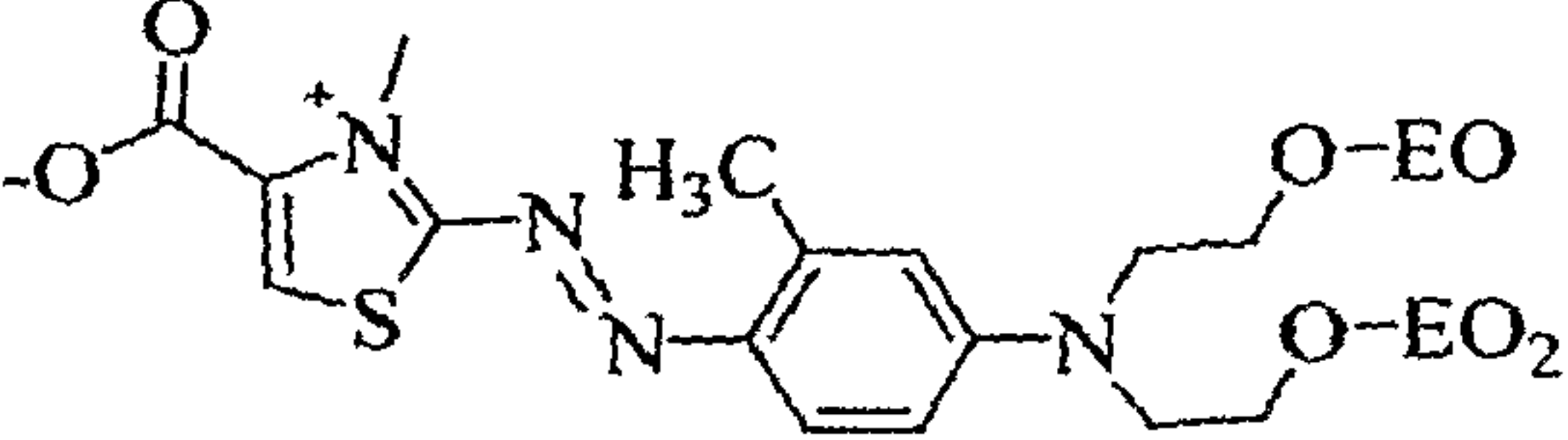
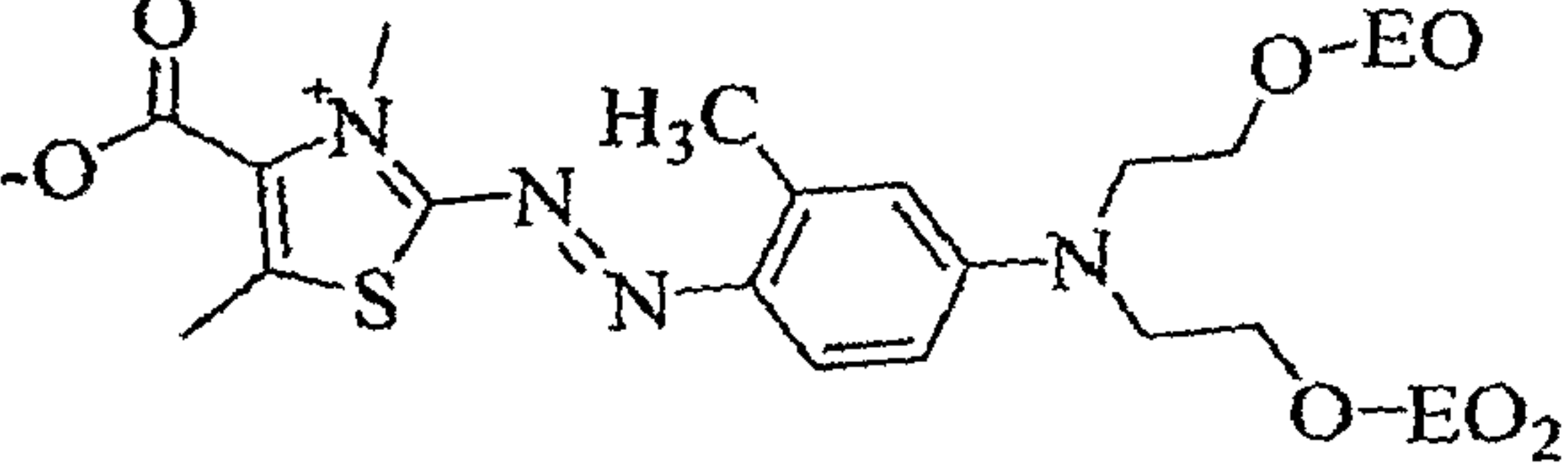
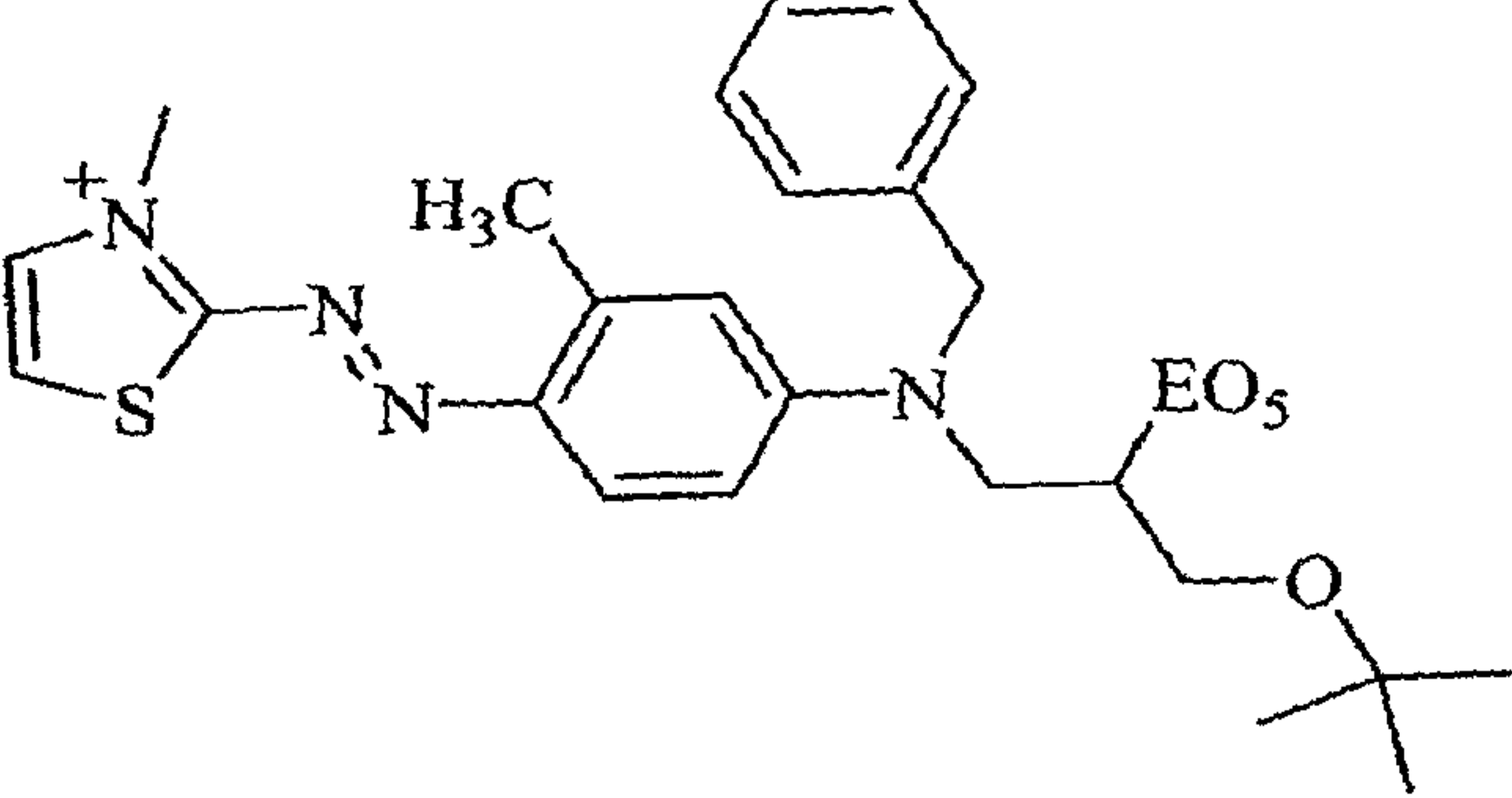
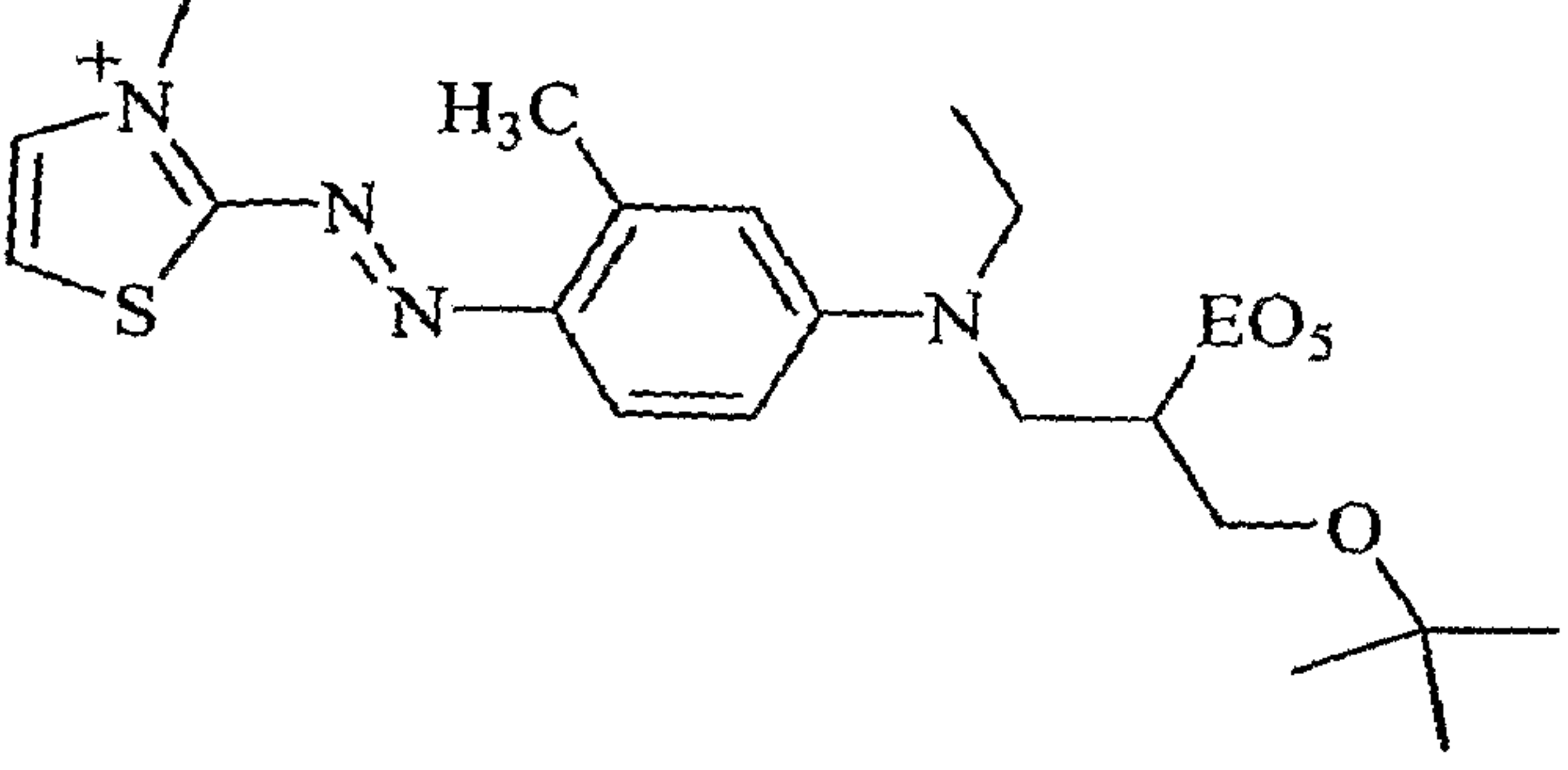
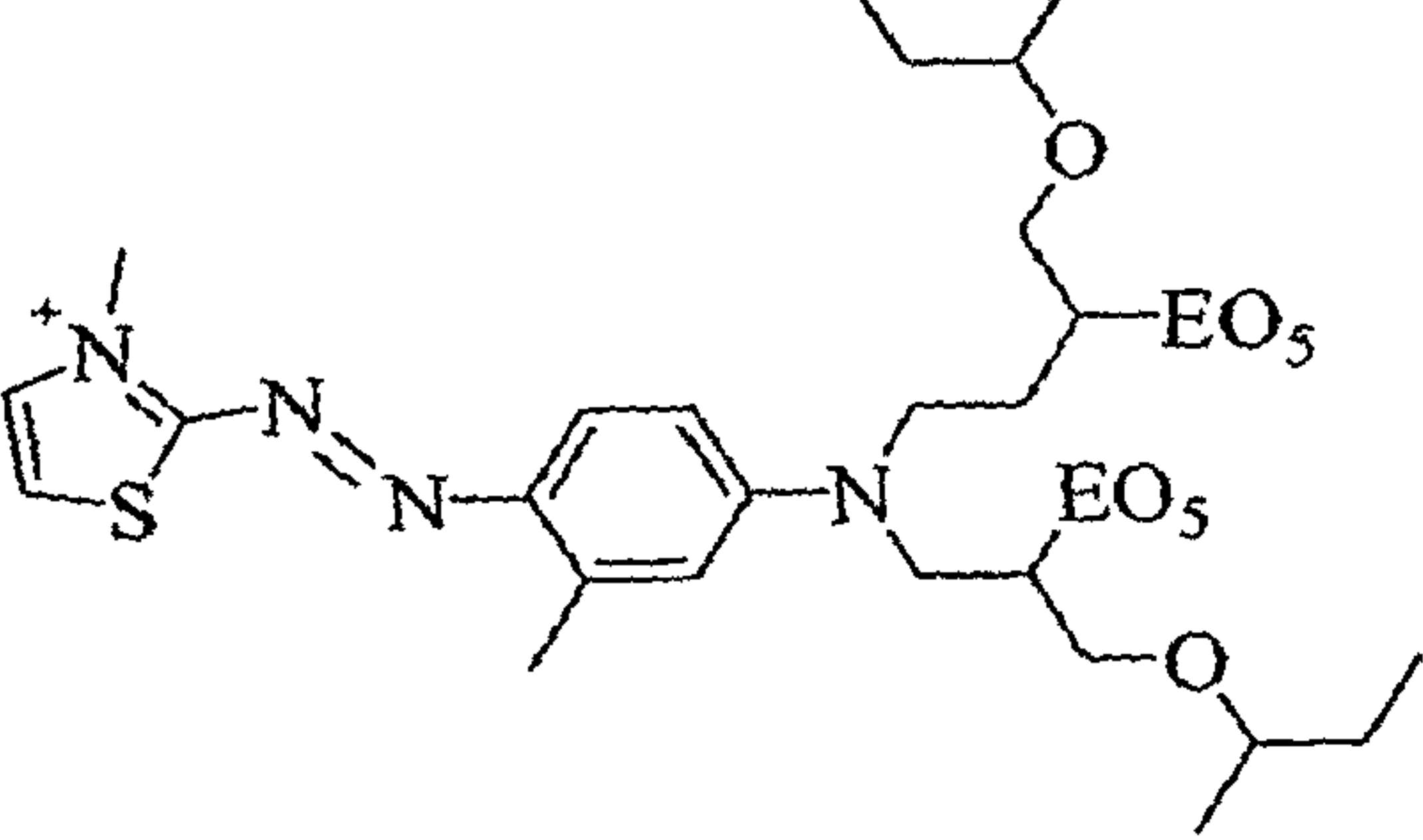
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Table 2

No.	Name
1	(E)-2-((4-(benzyl(methyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
2	(E)-2-((4-(dimethylamino)phenyl)diazenyl)-3-methylthiazol-3-ium
3	(E)-2-((4-(bis(2-hydroxyethyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
4	(E)-2-((4-(bis(2-(2-(2-hydroxyethoxy)ethoxy)ethyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
5	(E)-2-((4-(bis(2-(2-hydroxyethoxy)ethyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
6	(E)-2-((4-(bis(14-hydroxy-5,8,11-trimethyl-3,6,9,12-tetraoxapentadecyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
7	(E)-2-((4-(bis(2-(2-(2-(2-hydroxypropoxy)propoxy)propoxy)ethyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
8	(E)-2-((4-(bis(2-(2-(2-hydroxypropoxy)propoxy)ethyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
9	(E)-2-((4-(bis(35-hydroxy-5,8,11,14,17,20,23-heptamethyl-3,6,9,12,15,18,21,24,27,30,33-undecaoxapentatriacontyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
10	(E)-2-((4-(bis(3-(2,3-dihydroxypropoxy)-2-hydroxypropyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
11	(E)-2-((4-(bis(2,3-dihydroxypropyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
12	(E)-2-((4-((2-hydroxy-3-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)propyl)(2-hydroxy-3-(2-(2-hydroxyethoxy)ethoxy)propyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
13	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
14	(E)-2-((4-(bis(35-hydroxy-17,20,23,26,29,32-hexamethyl-3,6,9,12,15,18,21,24,27,30,33-undecaoxahexatriacontyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium

15	(E)-2-((4-(bis(14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
16	(E)-2-((4-((2-(2-(2-acetoxyethoxy)ethoxy)ethyl)(2-(2-acetoxyethoxy)ethyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
17	(E)-2-((4-(benzyl(2,3-dihydroxypropyl)amino)phenyl)diazanyl)-3-methylthiazol-3-ium
18	(E)-2-(2-((4-(bis(35-hydroxy-17,20,23,26,29,32-hexamethyl-3,6,9,12,15,18,21,24,27,30,33-undecaoxaheptatriacontyl)amino)-2-methylphenyl)diazanyl)thiazol-3-ium-3-yl)acetate
19	(E)-2-((4-(benzyl(14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
20	(E)-2-((4-((2-tert-butoxy-15-hydroxy-6,9,12-trimethyl-4,7,10,13-tetraoxahexadecyl)(2-(tert-butoxymethyl)-17-hydroxy-5,8,11,14-tetramethyl-3,6,9,12,15-pentaoxaoctadecyl)amino)phenyl)diazanyl)-3-methylthiazol-3-ium
21	(E)-2-((4-(benzyl(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)amino)phenyl)diazanyl)-3-methylthiazol-3-ium
22	(E)-2-((4-(bis(14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2-methylphenyl)diazanyl)-5-methoxy-3-methylbenzo[d]thiazol-3-ium
23	(E)-2-(2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazanyl)thiazol-3-ium-3-yl)acetate
24	(E)-2-((4-(ethyl(14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
25	(E)-2-((4-(benzyl(1,17-dihydroxy-3,6,9,12,15-pentaoxaoctadecan-18-yl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
26	(E)-2-((4-((2-(2-(2-(2,3-dihydroxypropoxy)ethoxy)ethoxy)ethyl)(2-(2-(2,3-dihydroxypropoxy)ethoxy)ethyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
27	(E)-2-(2-((4-(bis(14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2-methylphenyl)diazanyl)-6-methoxybenzo[d]thiazol-3-ium-3-yl)acetate
28	(E)-2-((4-((3-tert-butoxy-2-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)propyl)(3-

	tert-butoxy-2-(2-(2-hydroxyethoxy)ethoxy)propyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
29	(E)-2-((4-((3-butoxy-2-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)propyl)(3-butoxy-2-(2-(2-hydroxyethoxy)ethoxy)propyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
30	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)-3-isopropoxypropyl)(2-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)-3-isopropoxypropyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
31	(E)-2-((4-(benzyl(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
32	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)-3-(tridecyloxy)propyl)(2-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)-3-(tridecyloxy)propyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
33	(E)-3-ethyl-2-((4-(ethyl(23-hydroxy-3,6,9,12,15,18,21-heptaoxatricosyl)amino)-2-methylphenyl)diazanyl)thiazol-3-ium
34	(E)-2-((4-(bis(2-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)ethyl)amino)-2-methylphenyl)diazanyl)-3-ethylthiazol-3-ium
35	(E)-2-((1-(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)-1,2,3,4-tetrahydroquinolin-6-yl)diazanyl)-3-methylthiazol-3-ium
36	(E)-2-((4-((2-(2-hydroxypropoxy)ethyl)(2-(2-(2-hydroxypropoxy)propoxy)ethyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
37	(E)-2-((4-(bis(2-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)ethyl)amino)-2-methylphenyl)diazanyl)-3-ethylthiazol-3-ium
38	(E)-2-((4-(ethyl(23-hydroxy-3,6,9,12,15,18,21-heptaoxatricosyl)amino)-2-methylphenyl)diazanyl)-3-methylthiazol-3-ium
39	(E)-2-((4-(benzyl(14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2-methylphenyl)diazanyl)-3-ethylthiazol-3-ium
40	(E)-3-ethyl-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazanyl)thiazol-3-ium
41	(E)-3-(2-((4-(benzyl(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)amino)-2-methylphenyl)diazanyl)thiazol-3-ium-3-yl)propane-1-sulfonate

42	(E)-4-(2-((4-(benzyl(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)amino)-2-methylphenyl)diazenyl)thiazol-3-ium-3-yl)butane-1-sulfonate
43	(E)-2-((4-(bis(2-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-3-ethylthiazol-3-ium
44	(E)-3-benzyl-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)phenyl)diazenyl)thiazol-3-ium
45	(E)-3-ethyl-2-((4-((2-(2-hydroxypropoxy)ethyl)(2-(2-(2-hydroxypropoxy)propoxy)ethyl)amino)-2-methylphenyl)diazenyl)thiazol-3-ium
46	(E)-3-benzyl-2-((1-(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)-1,2,3,4-tetrahydroquinolin-6-yl)diazenyl)thiazol-3-ium
47	(E)-3-benzyl-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)thiazol-3-ium
48	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)propyl)(2-(2-hydroxyethoxy)propyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
49	(E)-2-((4-(benzyl(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
50	(E)-2-((4-(bis(14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2,6-dimethylphenyl)diazenyl)-3-methylthiazol-3-ium
51	(E)-2-((4-((4-(17-hydroxy-3,6,9,12,15-pentaoxaheptadecyloxy)-3-methoxybenzyl)(methyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
52	(E)-2-((1-(1-hydroxy-2,5,8,11,14,17,20,23,26-nonaoxaoctacosan-28-yl)-1,2,3,4-tetrahydroquinolin-6-yl)diazenyl)-3-methylthiazol-3-ium
53	(E)-4-(2-((4-(dimethylamino)phenyl)diazenyl)thiazol-3-ium-3-yl)butane-1-sulfonate
54	(E)-4-(2-((4-(dimethylamino)phenyl)diazenyl)-5-methylthiazol-3-ium-3-yl)butane-1-sulfonate
55	(E)-2-((4-((2-hydroxyethyl)(methyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
56	(E)-2-(methyl(4-((3-methylthiazol-3-ium-2-yl)diazenyl)phenyl)amino)ethyl

	sulfate
57	(E)-2-((4-(butyl(2-(2-hydroxyethoxy)ethyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
58	(E)-2-((4-(bis(2-(2-(2-hydroxypropoxy)propoxy)ethyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
59	(E)-2-((4-((2-hydroxyethyl)(isopropyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
60	(E)-2-((4-((14-hydroxy-3,6,9,12-tetraoxatetradecyl)(1-hydroxy-3,6,9,13-tetraoxapentadecan-15-yl)amino)-2-methylphenyl)diazenyl)-6-methoxy-3-methylbenzo[d]thiazol-3-ium
61	(E)-2-((4-(benzyl(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
62	(E)-2-((4-(benzyl(3-(3-(3-(2,3-dihydroxypropoxy)-2-hydroxypropoxy)-2-hydroxypropoxy)-2-hydroxypropyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
63	(E)-3-(2-((4-(bis(2-(2-(2-(2-hydroxyethoxy)ethoxy)ethoxy)ethyl)amino)-2-methylphenyl)diazenyl)thiazol-3-ium-3-yl)propane-1-sulfonate
64	(E)-2-((4-(bis(14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2,5-dimethylphenyl)diazenyl)-6-methoxy-3-methylbenzo[d]thiazol-3-ium
65	(E)-3-ethyl-2-((4-(ethyl(14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2-methylphenyl)diazenyl)thiazol-3-ium
66	(E)-3-ethyl-2-((1-(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)-1,2,3,4-tetrahydroquinolin-6-yl)diazenyl)thiazol-3-ium
67	(E)-3-ethyl-2-((1-(29-hydroxy-3,6,9,12,15,18,21,24,27-nonaoxanonacosyl)-1,2,3,4-tetrahydroquinolin-6-yl)diazenyl)thiazol-3-ium
68	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-3,5-dimethylthiazol-3-ium
69	(E)-3-ethyl-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-5-methylthiazol-3-ium
70	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-

	hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-3,5-dimethylthiazol-3-ium
71	(E)-3-ethyl-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-5-methylthiazol-3-ium
72	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-3,5-dimethylthiazol-3-ium
73	2-((E)-(4-((14-hydroxy-3,6,9,12-tetraoxatetradecyl)(17-hydroxy-3-(4-((E)-thiazol-2-yl)diazenyl)phenyl)-6,9,12,15-tetraoxa-3-azaheptadecyl)amino)phenyl)diazenyl)-3-methylthiazol-3-ium
74	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-3-methyl-5-nitrothiazol-3-ium
75	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-3-methyl-5-nitrothiazol-3-ium
76	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium-4-carboxylate
77	(E)-2-((4-((2-(2-(2-hydroxyethoxy)ethoxy)ethyl)(2-(2-hydroxyethoxy)ethyl)amino)-2-methylphenyl)diazenyl)-3,5-dimethylthiazol-3-ium-4-carboxylate
78	(E)-2-((4-(benzyl(2-(tert-butoxymethyl)-17-hydroxy-3,6,9,12,15-pentaoxaheptadecyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium
79	(E)-2-((4-((2-(tert-butoxymethyl)-14-hydroxy-3,6,9,12-tetraoxatetradecyl)(ethyl)amino)-2-hydroxyphenyl)diazenyl)-3-methylthiazol-3-ium
80	(E)-2-((4-((13-(sec-butoxymethyl)-1-hydroxy-3,6,9,12-tetraoxapentadecan-15-yl)(2-(sec-butoxymethyl)-14-hydroxy-3,6,9,12-tetraoxatetradecyl)amino)-2-methylphenyl)diazenyl)-3-methylthiazol-3-ium

In one aspect, suitable thiazolium dyes include thiazolium dye molecules numbers 1, 4, 5, 7, 8, 12, 13, 15, 16, 17, 21, 24, 25, 26, 30, 31, 33, 36, 38, 40, 45 and 48 as detailed in Tables 1 and 2 of the present specification.

In one aspect, suitable thiazolium dyes include thiazolium dye molecules numbers
5 12, 13, 15, 16, 24, 25, 26, 30, 31, 33, 36, 38, 40, 45 and 48 as detailed in Tables 1 and 2 of the present specification.

The suitable thiazolium dyes disclosed herein may be made by procedures known in the art and/or in accordance with the examples of the present specification.

Adjunct Materials

10 While not essential for the purposes of the present invention, the non-limiting list of adjuncts illustrated hereinafter are suitable for use in the laundry care compositions and may be desirably incorporated in certain embodiments of the invention, for example to assist or enhance performance, for treatment of the substrate to be cleaned, or to modify the aesthetics of the composition as is the case with perfumes, colorants, dyes or the like.
15 It is understood that such adjuncts are in addition to the components that were previously listed for any particular embodiment. The total amount of such adjuncts may range from about 0.1% to about 50%, or even from about 1% to about 30%, by weight of the laundry care composition.

The precise nature of these additional components, and levels of incorporation thereof, will depend on the physical form of the composition and the nature of the operation for which it is to be used. Suitable adjunct materials include, but are not limited to, polymers, for example cationic polymers, surfactants, builders, chelating agents, dye transfer inhibiting agents, dispersants, enzymes, and enzyme stabilizers, catalytic materials, bleach activators, polymeric dispersing agents, clay soil removal/anti-redeposition agents, brighteners, suds suppressors, dyes, additional perfume and perfume delivery systems, structure elasticizing agents, fabric softeners, carriers, hydrotropes, processing aids and/or pigments. In addition to the disclosure below, suitable examples of such other adjuncts and levels of use are found in U.S. Patent Nos. 5,576,282, 6,306,812 B1 and 6,326,348 B1.

As stated, the adjunct ingredients are not essential to Applicants' cleaning and laundry care compositions. Thus, certain embodiments of Applicants' compositions do not contain one or more of the following adjunct materials: bleach activators, surfactants, builders, chelating agents, dye transfer inhibiting agents, dispersants, enzymes, and

enzyme stabilizers, catalytic metal complexes, polymeric dispersing agents, clay and soil removal/anti-redeposition agents, brighteners, suds suppressors, dyes, additional perfumes and perfume delivery systems, structure elasticizing agents, fabric softeners, carriers, hydrotropes, processing aids and/or pigments. However, when one or more adjuncts are present, such one or more adjuncts may be present as detailed below:

Surfactants - The compositions according to the present invention can comprise a surfactant or surfactant system wherein the surfactant can be selected from nonionic and/or anionic and/or cationic surfactants and/or ampholytic and/or zwitterionic and/or semi-polar nonionic surfactants. The surfactant is typically present at a level of from about 0.1%, from about 1%, or even from about 5% by weight of the cleaning compositions to about 99.9%, to about 80%, to about 35%, or even to about 30% by weight of the cleaning compositions.

Builders - The compositions of the present invention can comprise one or more detergent builders or builder systems. When present, the compositions will typically comprise at least about 1% builder, or from about 5% or 10% to about 80%, 50%, or even 30% by weight, of said builder. Builders include, but are not limited to, the alkali metal, ammonium and alkanolammonium salts of polyphosphates, alkali metal silicates, alkaline earth and alkali metal carbonates, aluminosilicate builders polycarboxylate compounds, ether hydroxypolycarboxylates, copolymers of maleic anhydride with ethylene or vinyl methyl ether, 1,3,5-trihydroxybenzene-2,4,6-trisulphonic acid, and carboxymethyloxysuccinic acid, the various alkali metal, ammonium and substituted ammonium salts of polyacetic acids such as ethylenediamine tetraacetic acid and nitrilotriacetic acid, as well as polycarboxylates such as mellitic acid, succinic acid, oxydisuccinic acid, polymaleic acid, benzene 1,3,5-tricarboxylic acid, carboxymethyloxysuccinic acid, and soluble salts thereof.

Chelating Agents - The compositions herein may also optionally contain one or more copper, iron and/or manganese chelating agents. If utilized, chelating agents will generally comprise from about 0.1% by weight of the compositions herein to about 15%, or even from about 3.0% to about 15% by weight of the compositions herein.

Dye Transfer Inhibiting Agents - The compositions of the present invention may also include one or more dye transfer inhibiting agents. Suitable polymeric dye transfer inhibiting agents include, but are not limited to, polyvinylpyrrolidone polymers, polyamine N-oxide polymers, copolymers of N-vinylpyrrolidone and N-vinylimidazole, polyvinylloxazolidones and polyvinylimidazoles or mixtures thereof. When present in the

compositions herein, the dye transfer inhibiting agents are present at levels from about 0.0001%, from about 0.01%, from about 0.05% by weight of the cleaning compositions to about 10%, about 2%, or even about 1% by weight of the cleaning compositions.

Dispersants - The compositions of the present invention can also contain dispersants. Suitable water-soluble organic materials are the homo- or co-polymeric acids or their salts, in which the polycarboxylic acid may comprise at least two carboxyl radicals separated from each other by not more than two carbon atoms.

Enzymes - The compositions can comprise one or more detergent enzymes which provide cleaning performance and/or fabric care benefits. Examples of suitable enzymes include, but are not limited to, hemicellulases, peroxidases, proteases, cellulases, xylanases, lipases, phospholipases, esterases, cutinases, pectinases, keratanases, reductases, oxidases, phenoloxidases, lipoxygenases, ligninases, pullulanases, tannases, pentosanases, malanases, β -glucanases, arabinosidases, hyaluronidase, chondroitinase, laccase, and amylases, or mixtures thereof. A typical combination is a cocktail of conventional applicable enzymes like protease, lipase, cutinase and/or cellulase in conjunction with amylase.

Enzyme Stabilizers - Enzymes for use in compositions, for example, detergents can be stabilized by various techniques. The enzymes employed herein can be stabilized by the presence of water-soluble sources of calcium and/or magnesium ions in the finished compositions that provide such ions to the enzymes.

Catalytic Metal Complexes - Applicants' compositions may include catalytic metal complexes. One type of metal-containing bleach catalyst is a catalyst system comprising a transition metal cation of defined bleach catalytic activity, such as copper, iron, titanium, ruthenium, tungsten, molybdenum, or manganese cations, an auxiliary metal cation having little or no bleach catalytic activity, such as zinc or aluminum cations, and a sequester having defined stability constants for the catalytic and auxiliary metal cations, particularly ethylenediaminetetraacetic acid, ethylenediaminetetra (methylenephosphonic acid) and water-soluble salts thereof. Such catalysts are disclosed in U.S. patent 4,430,243.

If desired, the compositions herein can be catalyzed by means of a manganese compound. Such compounds and levels of use are well known in the art and include, for example, the manganese-based catalysts disclosed in U.S. patent 5,576,282.

Cobalt bleach catalysts useful herein are known, and are described, for example, in U.S. patents 5,597,936 and 5,595,967. Such cobalt catalysts are readily prepared by known procedures, such as taught for example in U.S. patents 5,597,936, and 5,595,967.

Compositions herein may also suitably include a transition metal complex of a macropolycyclic rigid ligand - abbreviated as "MRL". As a practical matter, and not by way of limitation, the compositions and cleaning processes herein can be adjusted to provide on the order of at least one part per hundred million of the benefit agent MRL species in the aqueous washing medium, and may provide from about 0.005 ppm to about 25 ppm, from about 0.05 ppm to about 10 ppm, or even from about 0.1 ppm to about 5 ppm, of the MRL in the wash liquor.

Preferred transition-metals in the instant transition-metal bleach catalyst include manganese, iron and chromium. Preferred MRL's herein are a special type of ultra-rigid ligand that is cross-bridged such as 5,12-diethyl-1,5,8,12-tetraazabicyclo[6.6.2]hexadecane.

Suitable transition metal MRLs are readily prepared by known procedures, such as taught for example in WO 00/32601, and U.S. patent 6,225,464.

Processes of Making Laundry Care Compositions

The laundry care compositions of the present invention can be formulated into any suitable form and prepared by any process chosen by the formulator, non-limiting
5 examples of which are described in U.S. 5,879,584; U.S. 5,691,297; U.S. 5,574,005; U.S. 5,569,645; U.S. 5,565,422; U.S. 5,516,448; U.S. 5,489,392; and U.S. 5,486,303.

In one aspect, the liquid detergent compositions disclosed herein may be prepared by combining the components thereof in any convenient order and by mixing, e.g., agitating, the resulting component combination to form a phase stable liquid detergent
10 composition. In one aspect, a liquid matrix is formed containing at least a major proportion, or even substantially all, of the liquid components, e.g., nonionic surfactant, the non-surface active liquid carriers and other optional liquid components, with the liquid components being thoroughly admixed by imparting shear agitation to this liquid combination. For example, rapid stirring with a mechanical stirrer may usefully be
15 employed. While shear agitation is maintained, substantially all of any anionic surfactant and the solid ingredients can be added. Agitation of the mixture is continued, and if necessary, can be increased at this point to form a solution or a uniform dispersion of insoluble solid phase particulates within the liquid phase. After some or all of the solid-

form materials have been added to this agitated mixture, particles of any enzyme material to be included, e.g., enzyme prills, are incorporated. As a variation of the composition preparation procedure described above, one or more of the solid components may be added to the agitated mixture as a solution or slurry of particles premixed with a minor portion of one or more of the liquid components. After addition of all of the composition components, agitation of the mixture is continued for a period of time sufficient to form compositions having the requisite viscosity and phase stability characteristics. Frequently this will involve agitation for a period of from about 30 to 60 minutes.

In another aspect of producing liquid detergents, the thiazolium dye is first combined with one or more liquid components to form a thiazolium dye premix, and this thiazolium dye premix is added to a composition formulation containing a substantial portion, for example more than 50% by weight, more than 70% by weight, or even more than 90% by weight, of the balance of components of the laundry detergent composition. For example, in the methodology described above, both the thiazolium dye premix and the enzyme component are added at a final stage of component additions. In another aspect, the thiazolium dye is encapsulated prior to addition to the detergent composition, the encapsulated dye is suspended in a structured liquid, and the suspension is added to a composition formulation containing a substantial portion of the balance of components of the laundry detergent composition.

Various techniques for forming detergent compositions in such solid forms are well known in the art and may be used herein. In one aspect, when the laundry care composition is in the form of a granular particle, the thiazolium dye is provided in particulate form, optionally including additional but not all components of the laundry detergent composition. The thiazolium dye particulate is combined with one or more additional particulates containing a balance of components of the laundry detergent composition. Further, the thiazolium dye, optionally including additional but not all components of the laundry detergent composition may be provided in an encapsulated form, and the thiazolium dye encapsulate is combined with particulates containing a substantial balance of components of the laundry detergent composition.

Methods of Using Laundry Care Compositions

The laundry care compositions disclosed in the present specification may be used to clean or treat a fabric. Typically at least a portion of the fabric is contacted with an embodiment of the aforementioned laundry care compositions, in neat form or diluted in a

liquor, for example, a wash liquor and then the fabric may be optionally washed and/or rinsed. In one aspect, a fabric is optionally washed and/or rinsed, contacted with an embodiment of the aforementioned laundry care compositions and then optionally washed and/or rinsed. For purposes of the present invention, washing includes but is not limited to, scrubbing, and mechanical agitation. The fabric may comprise most any fabric capable of being laundered or treated.

The laundry care compositions disclosed in the present specification can be used to form aqueous washing solutions for use in the laundering of fabrics. Generally, an effective amount of such compositions is added to water, preferably in a conventional fabric laundering automatic washing machine, to form such aqueous laundering solutions. The aqueous washing solution so formed is then contacted, preferably under agitation, with the fabrics to be laundered therewith. An effective amount of the laundry care composition, such as the liquid detergent compositions disclosed in the present specification, may be added to water to form aqueous laundering solutions that may comprise from about 500 to about 7,000 ppm or even from about 1,000 to about 3,000 ppm of laundry care composition.

In one aspect, one or more of the thiazolium dyes disclosed in the present specification may be provided, for example via a laundry care composition, such that during the wash cycle and or rinse cycle the concentration of such one or more dyes may be from about 0.5 parts per billion (ppb) to about 5 part per million (ppm), from about 1 ppb to about 600 ppb, from about 5 ppb to about 300 ppb, or even from about 10 ppb to about 100 ppb of thiazolium dye. In one aspect such concentrations may be achieved during the washing cycle, and/or rinse cycle, of a 17 gallon automatic laundry washing machine.

In one aspect, the laundry care compositions may be employed as a laundry additive, a pre-treatment composition and/or a post-treatment composition.

TEST METHODS

I. Method for Determining of Hueing Efficiency for Detergents

- a.) Two 25 cm x 25 cm fabric swatches of 16 oz white cotton interlock knit fabric (270 g/square meter, brightened with UvitexTM BNB fluorescent whitening agent, from Test Fabrics. P.O. Box 26, Weston, PA, 18643), are obtained.

- b.) Prepare two one liter aliquots of tap water containing 1.55 g of AATCC standard heavy duty liquid (HDL) test detergent as set forth in Table 3.
- c.) Add a sufficient amount the dye to be tested to one of the aliquots from Step b.) above to produce an aqueous solution absorbance of 1 AU.
- 5 d.) Wash one swatch from a.) above in one of the aliquots of water containing 1.55 g of AATCC standard heavy duty liquid (HDL) test detergent and wash the other swatch in the other aliquot. Such washing step should be conducted for 30 minutes at room temperature with agitation. After such washing step separately rinse the swatches and dry the swatches.
- 10 e.) After rinsing and drying each swatch, the hueing efficiency, DE^*_{eff} , of the dye is assessed by determining the L^* , a^* , and b^* value measurements of each swatch using a Hunter LabScan XE reflectance spectrophotometer with D65 illumination, 10° observer and UV filter excluded. The hueing efficiency of the dye is then calculated using the following equation:

$$DE^*_{eff} = ((L^*_c - L^*_s)^2 + (a^*_c - a^*_s)^2 + (b^*_c - b^*_s)^2)^{1/2}$$

wherein the subscripts c and s respectively refer to the L^* , a^* , and b^* values measured for the control, i.e., the fabric sample washed in detergent with no dye, and the fabric sample washed in detergent containing the dye to be screened.

25 II. Method for Determining Wash Removability

- a.) Prepare two separate 150 ml aliquots of HDL detergent solution set forth in Table 1, according to AATCC Test Method 61-2003, Test 2A and containing 1.55 g/ liter of the AATCC HDL formula in distilled water.
- 30 b.) A 15 cm x 5 cm sample of each fabric swatch from the Method for Determining of Hueing Efficiency For Detergents described above is washed in a Launderometer for 45 minutes at 49°C in 150 ml of a the HDL detergent solution prepared according to Step II. a.) above.
- c.) The samples are rinsed with separate aliquots of rinse water and air dried in the dark, the amount of residual coloration is assessed by measuring the

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DE*_{res} of the dye is assessed by determining the L*, a*, and b* value measurements of each swatch using a Hunter LabScan XE reflectance spectrophotometer with D65 illumination, 10° observer and UV filter excluded. The hueing efficiency of the dye is then calculated using the following equation:

$$DE^*_{res} = ((L^*_c - L^*_s)^2 + (a^*_c - a^*_s)^2 + (b^*_c - b^*_s)^2)^{1/2}$$

wherein the subscripts c and s respectively refer to the L*, a*, and b* values measured for the control, i.e., the fabric sample initially washed in detergent with no dye, and the fabric sample initially washed in detergent containing the dye to be screened. The wash removal value for the dye is then calculated according to the formula: % removal = 100 x (1 – DE*_{res}/DE*_{eff}).

Table 3

Ingredient	weight percent
C11.8 linear alkylbenzene sulfonic acid	12.00
Neodol 23-9	8.00
citric acid	1.20
C12-14 fatty acid	4.00
sodium hydroxide ¹	2.65
ethanolamine	0.13
borax	1.00
DTPA ²	0.30
1,2-propanediol	8.00
brightener 15	0.04
water	balance

¹ formula pH adjusted to 8.5

² diethylenetriaminepentaacetic acid, pentasodium salt

EXAMPLES

The following examples illustrate the compositions of the present invention but are not necessarily meant to limit or otherwise define the scope of the invention herein.

Example 1

5 The following liquid formulas are within the scope of the present invention.

Ingredient	1a	1b	1c	1d	1e	1f ⁵
	wt %	wt %	wt %	wt %	wt %	wt %
sodium alkyl ether sulfate	14.4%	14.4%		9.2%	5.4%	
linear alkylbenzene	4.4%	4.4%	12.2%	5.7%	1.3%	22.0%
sulfonic acid						
alkyl ethoxylate	2.2%	2.2%	8.8%	8.1%	3.4%	18.0%
amine oxide	0.7%	0.7%	1.5%			
citric acid	2.0%	2.0%	3.4%	1.9%	1.0%	1.6%
fatty acid	3.0%	3.0%	8.3%			16.0%
protease	1.0%	1.0%	0.7%	1.0%		2.5%
amylase	0.2%	0.2%	0.2%			0.3%
lipase				0.2%		
borax	1.5%	1.5%	2.4%	2.9%		
calcium and sodium						
formate	0.2%	0.2%				
formic acid						1.1%
amine ethoxylate polymers	1.8%	1.8%	2.1%			3.2%
sodium polyacrylate					0.2%	
sodium polyacrylate						
copolymer				0.6%		
DTPA ¹	0.1%	0.1%				0.9%
DTPMP ²			0.3%			
EDTA ³					0.1%	
fluorescent whitening						
agent	0.15%	0.15%	0.2%	0.12%	0.12%	0.2%
ethanol	2.5%	2.5%	1.4%	1.5%		
propanediol	6.6%	6.6%	4.9%	4.0%		15.7%
sorbitol				4.0%		

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ethanolamine	1.5%	1.5%	0.8%	0.1%		11.0%
sodium hydroxide	3.0%	3.0%	4.9%	1.9%	1.0%	
sodium cumene sulfonate			2.0%			
silicone suds suppressor			0.01%			
perfume	0.3%	0.3%	0.7%	0.3%	0.4%	0.6%
Compound 16 of Table 1	0.005%	0.005%				
Compound 24 of Table 1			0.005%			
Compound 13 of Table 1				0.008%		
Compound 36 of Table 1					0.008%	
Compound 21 of Table 1						0.015%
Liquitint Aqua AS ⁴		0.005%				
opacifier ⁶		0.5%				
water	<u>balance</u>	<u>balance</u>	<u>balance</u>	<u>balance</u>	<u>balance</u>	<u>balance</u>
	100.0%	100.0%	100.0%	100.0%	100.0%	100.0%

¹ diethylenetriaminepentaacetic acid, sodium salt² diethylenetriaminepentakis(methylenephosphonic acid), sodium salt³ ethylenediaminetetraacetic acid, sodium salt⁴ a non-tinting dye used to adjust formula color5 ⁵ compact formula, packaged as a unitized dose in polyvinyl alcohol film⁶ Acusol OP 301

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Example 2

The following granular detergent formulas are within the scope of the present invention.

Ingredient	2a	2b	2c	2d
	wt %	wt %	wt %	wt %
Na linear alkylbenzene sulfonate	3.4%	3.3%	11.0%	3.4%
Na alkylsulfate	4.0%	4.1%		4.0%
Na alkyl sulfate (branched)	9.4%	9.6%		9.4%
alkyl ethoxylate			3.5%	
type A zeolite	37.4%	35.4%	26.8%	37.4%
sodium carbonate	22.3%	22.5%	35.9%	22.3%
sodium sulfate	1.0%		18.8%	1.0%
sodium silicate			2.2%	
protease	0.1%	0.2%		0.1%
sodium polyacrylate	1.0%	1.2%	0.7%	1.0%
carboxymethylcellulose			0.1%	
PEG 600		0.5%		
PEG 4000		2.2%		
DTPA	0.7%	0.6%		0.7%
fluorescent whitening agent	0.1%	0.1%	0.1%	0.1%
sodium percarbonate		5.0%		
sodium nonanoyloxybenzenesulfonate		5.3%		
silicone suds suppressor	0.02%	0.02%		0.02%
perfume	0.3%	0.3%	0.2%	0.3%
Compound 15 of Table 1	0.015% ¹			
Compound 48 of Table 1		0.017% ²		
Compound 38 of Table 1			0.017% ³	
Compound 33 of Table 1				0.02% ⁴
water and miscellaneous	balance	balance	balance	balance

¹ formulated as a particle containing 0.5% dye, 99.5% PEG 4000

² formulated as a layered particle containing 2% dye according to US 2006 252667 A1

5 ³ formulated as a particle containing 0.5% dye according to US 4,990,280

⁴ formulated as a particle containing 0.5% dye with zeolite

Example 3

The following rinse added fabric conditioning formulas are within the scope of the present invention.

Ingredients	3a	3b	3c	3d
Fabric Softening Active ^a	13.70%	13.70%	13.70%	13.70%
Ethanol	2.14%	2.14%	2.14%	2.14%
Cationic Starch ^b	2.17%	2.17%	2.17%	2.17%
Perfume	1.45%	1.45%	1.45%	1.45%
Phase Stabilizing Polymer ^c	0.21%	0.21%	0.21%	0.21%
Calcium Chloride	0.147%	0.147%	0.147%	0.147%
DTPA ^d	0.007%	0.007%	0.007%	0.007%
Preservative ^e	5 ppm	5 ppm	5 ppm	5 ppm
Antifoam ^f	0.015%	0.015%	0.015%	0.015%
Compound 45 of Table 1	30 ppm			15 ppm
Compound 25 of Table 1		30 ppm		
Compound 30 of Table 1			30 ppm	15 ppm
Tinopal CBS-X ^g	0.2	0.2	0.2	0.2
Ethoquad C/25 ^h	0.26	0.26	0.26	0.26
Ammonium Chloride	0.1%	0.1%	0.1%	0.1%
Hydrochloric Acid	<u>0.012 %</u>	<u>0.012 %</u>	<u>0.012 %</u>	<u>0.012 %</u>
Deionized Water	Balance	Balance	Balance	Balance

^a N,N-di(tallowoxyloxyethyl)-N,N-dimethylammonium chloride.

^b Cationic starch based on common maize starch or potato starch, containing 25% to 95% amylose and a degree of substitution of from 0.02 to 0.09, and having a viscosity measured as Water Fluidity having a value from 50 to 84.

^c Copolymer of ethylene oxide and terephthalate having the formula described in US 5,574,179 at col.15, lines 1-5, wherein each X is methyl, each n is 40, u is 4, each R¹ is essentially 1,4-phenylene moieties, each R² is essentially ethylene, 1,2-propylene moieties, or mixtures thereof.

^d Diethylenetriaminepentaacetic acid.

^e KATHON[®] CG available from Rohm and Haas Co.

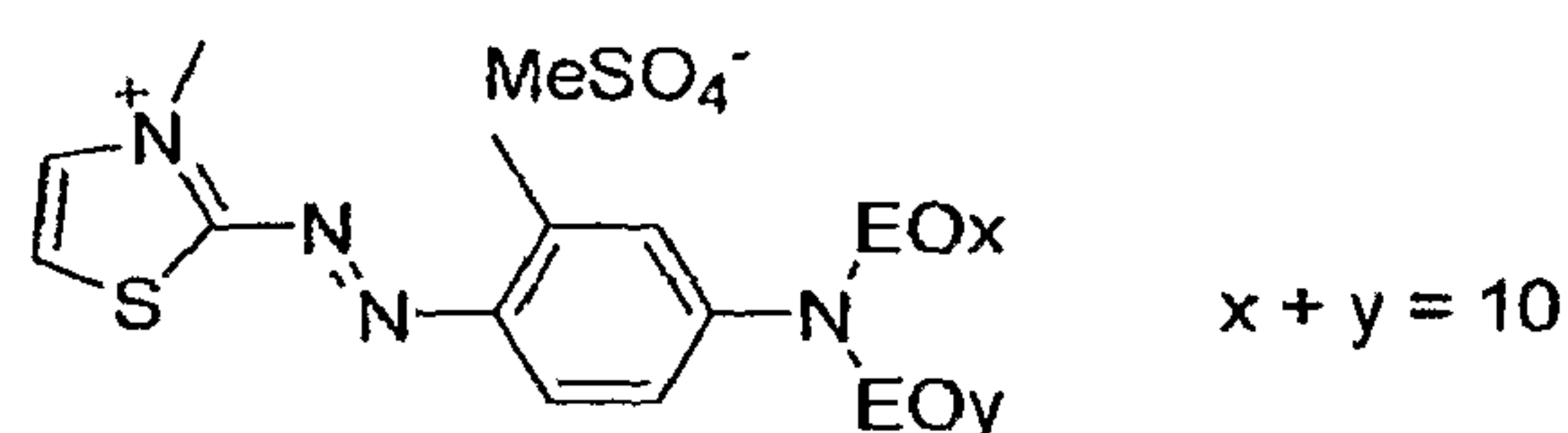
^f Silicone antifoam agent available from Dow Corning Corp. under the trade name DC2310.

5 ^g Disodium 4,4'-bis-(2-sulfostyryl) biphenyl, available from Ciba Specialty Chemicals.

^h Cocomethyl ethoxylated [15] ammonium chloride, available from Akzo Nobel

Example 4

Synthesis of mtol-10EO methylthiazolium



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Five hundred and forty-nine grams of 85% phosphoric acid, 75 grams of 98% sulfuric acid and 9 drops of 2-ethyl hexanol defoamer are added to a 100 milliliter three necked flask equipped with a thermometer, cooling bath, and mechanical stirrer. The mixture is cooled and 30.9 grams of 2-aminothiazole is added to the flask. The mixture is further cooled to below 0°C after which 105 grams of 40% nitrosyl sulfuric acid are added while the temperature is maintained below 5°C. After three hours the mixture gives a positive nitrite test and 25 grams of sulfamic acid are added slowly while the temperature is kept below 5°C. A negative nitrite test is evident after one hour.

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A 2000 milliliter beaker is charged with 190 grams 10 EO m-toluidine intermediate, 200 grams of water, 200 grams of ice and 12 grams of urea. The mixture is cooled to 0°C. The diazo solution is added dropwise to the beaker over about 30 minutes, while maintaining the temperature below 10°C. The resulting mixture is stirred for several hours and allowed to stand overnight, after which 780 grams of 50% sodium hydroxide is added to neutralize excess acid to a pH of about 7 while the temperature is kept below 20°C. The bottom salt layer is removed and the product is washed with 200 milliliters of a 10% sodium sulfate solution. The aqueous layer is removed and the desired product is obtained as an orange liquid (240 grams, 70% actives).

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One hundred grams of the orange liquid from above and 28.40 grams of dimethyl sulfate are placed into a 500 milliliter flask equipped with a reflux condenser, thermometer, heating mantle and mechanical stirrer. The reaction mixture is heated to

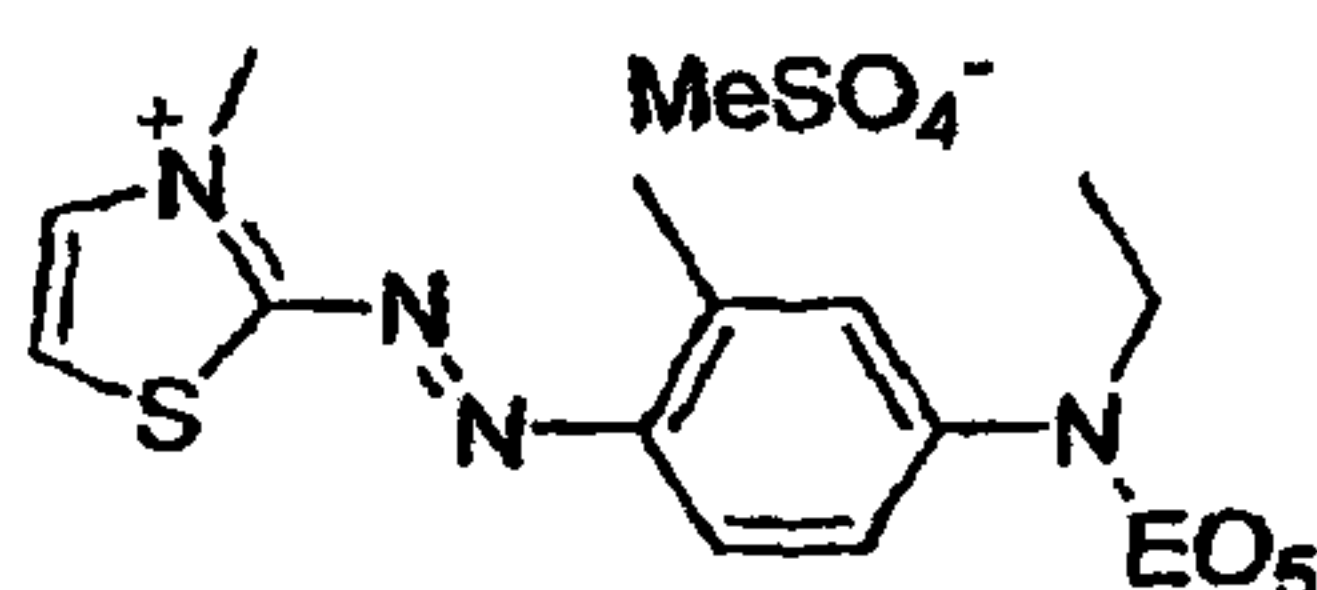
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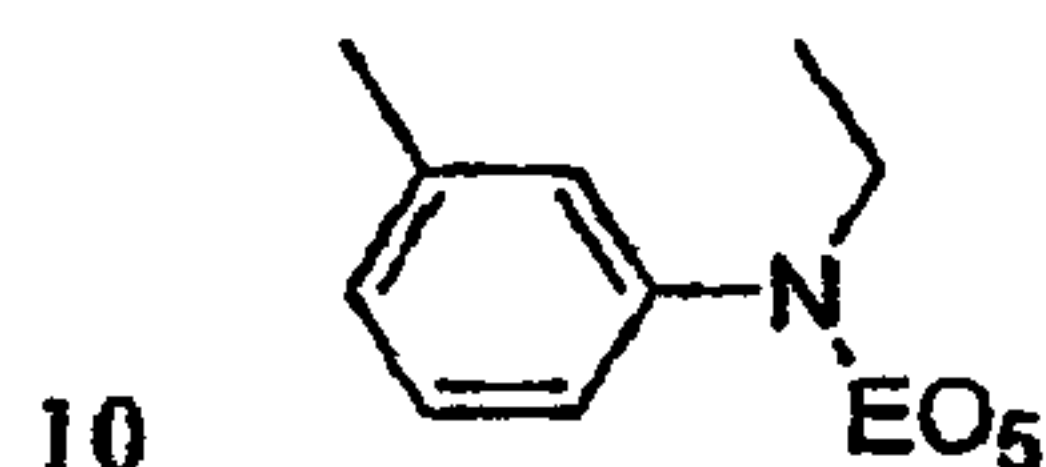
70°C for two hours. The reaction is cooled and the pH is adjusted to 7 with 10 grams of 20% ammonium hydroxide and is used without further purification.

5 Example 5

The procedure of Example 4 is used to make N-ethyl-mtol-5EO

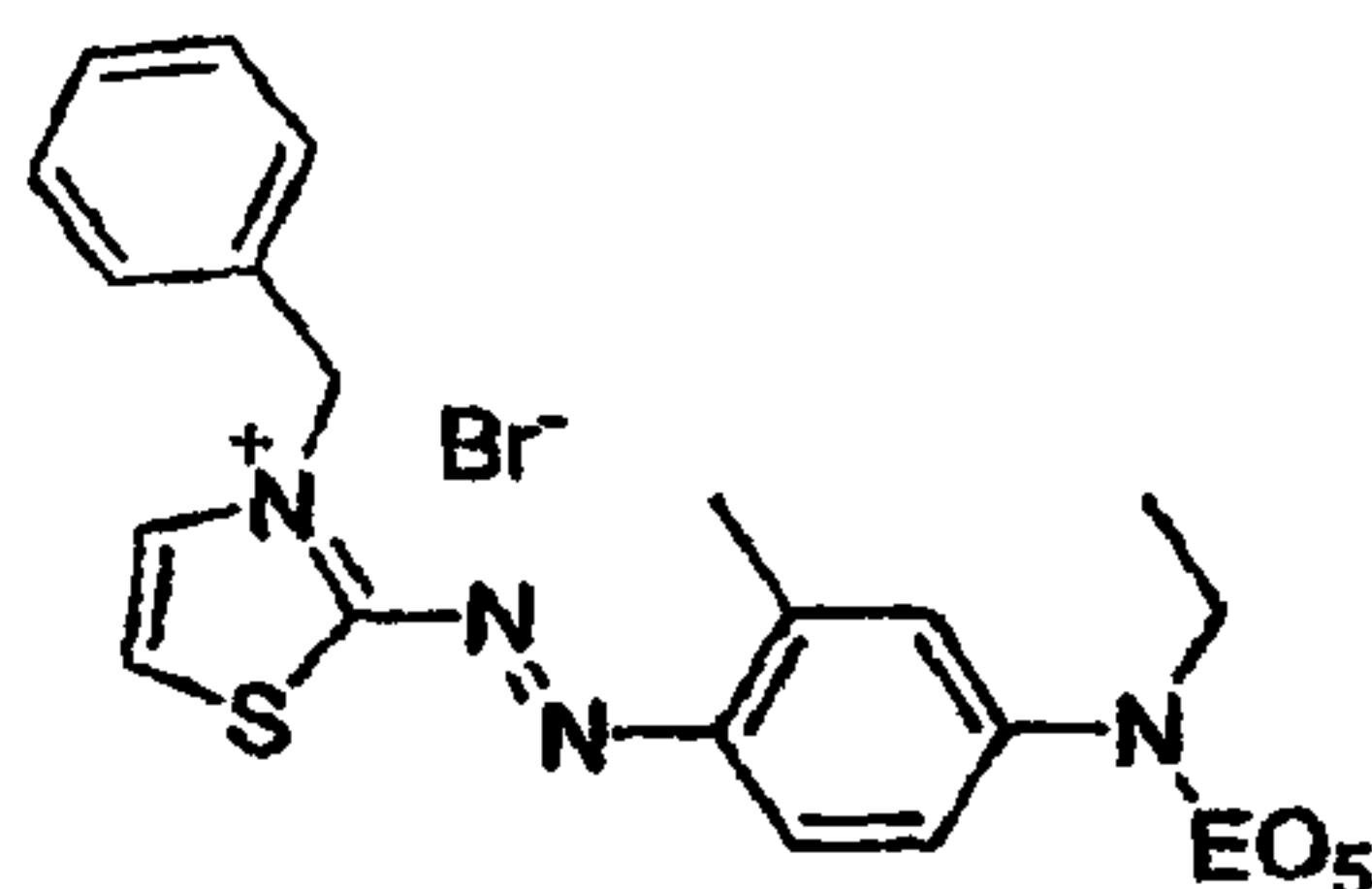


with the difference being the use of the following m-toluidine intermediate:



Example 6

The procedure of Example 5, with the noted changes, is used to make:



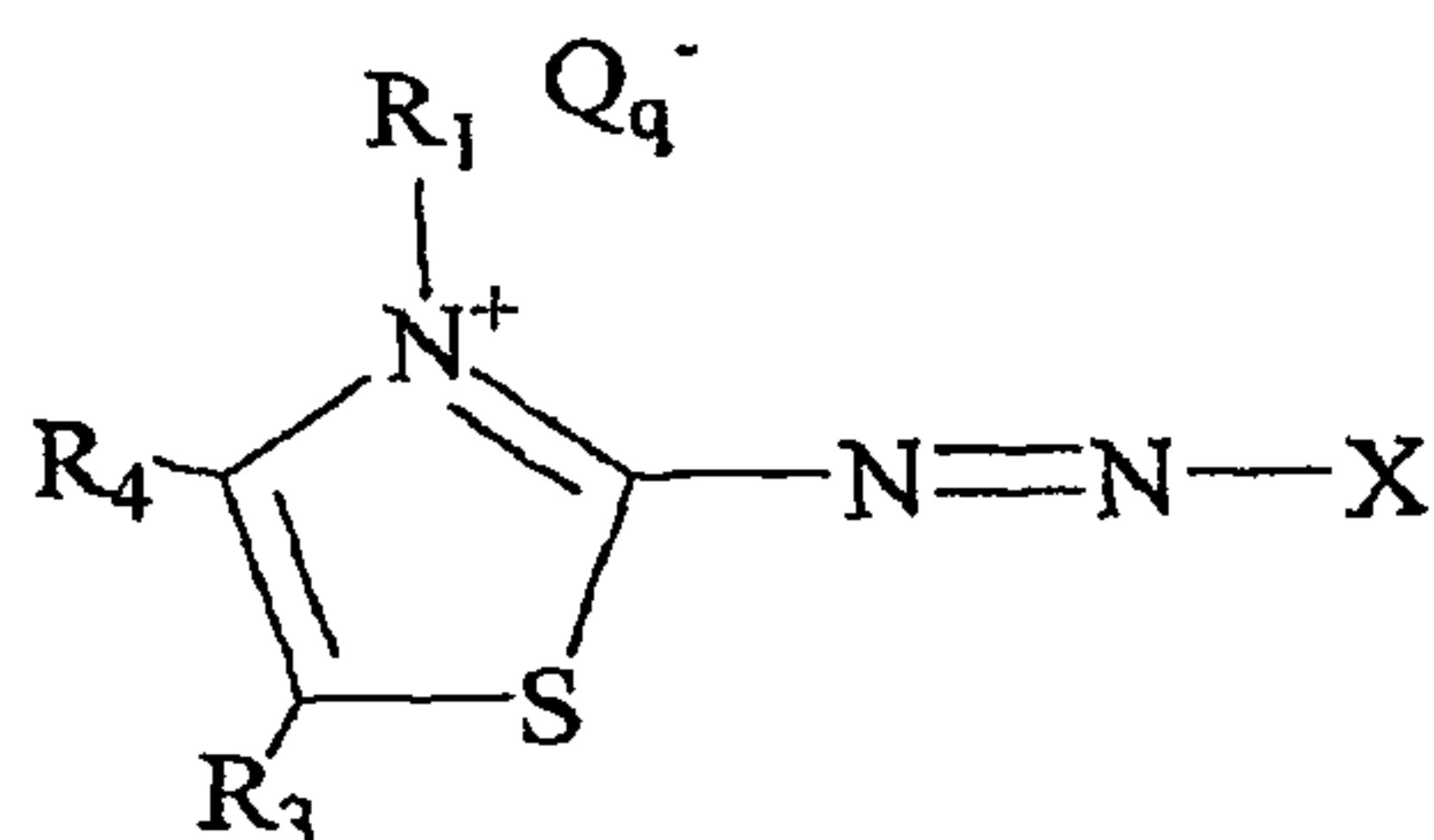
15 Twenty grams of the orange liquid per Example 5, as obtained via Example 4, and nine grams of benzyl bromide are placed into a 250 milliliter flask equipped with a reflux condenser, thermometer, heating mantle and mechanical stirrer. The reaction mixture is heated to 70°C for two hours. The reaction is cooled and the pH is adjusted to 7 with 4 grams of 50% sodium hydroxide and is used without further purification.

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While particular embodiments of the present invention have been illustrated and described, it would be obvious to those skilled in the art that various other changes and modifications can be made.

What is claimed is:

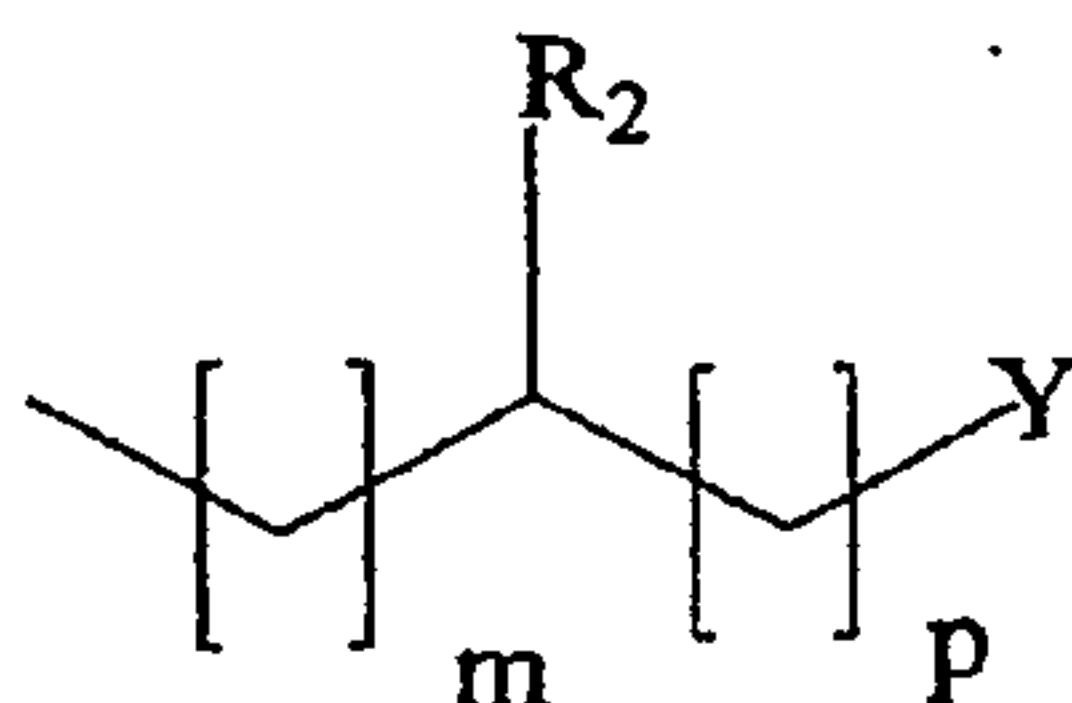
1. A whitening agent which is a thiazolium dye of the general Formula (V);



Formula V

wherein:

- 5 a.) R_1 is a branched or unbranched (C_1 - C_{22})-alkyl moiety, an aromatic alkyl moiety, a polyalkylene oxide moiety, or a moiety having Formula (VI) below;



Formula VI

wherein

- 10 (i) R_2 is hydrogen, methyl, ethyl, propyl, acetate or a hydroxyl moiety; m and p are, independently, integers from 0 to $(n-1)$, with the proviso that n is an integer from 1 to 6 and $m+p = (n-1)$
- 15 (ii) Y is a hydroxyl, sulfonate, sulfate, carboxylate or acetate moiety;

b.) R_3 and R_4 :

- 20 i.) are independently hydrogen; a (C_1 - C_{22})-alkyl moiety or (C_2 - C_{22})-alkenyl moiety; a hydroxy-(C_2 - C_{22})-alkyl moiety; a hydroxy-(C_2 - C_{22})-alkyl moiety comprising, in addition to the hydroxyl oxygen, an oxygen atom; a polyether moiety; an amino-(C_1 - C_{22})-alkyl moiety; a substituted or unsubstituted phenyl moiety; a substituted or unsubstituted

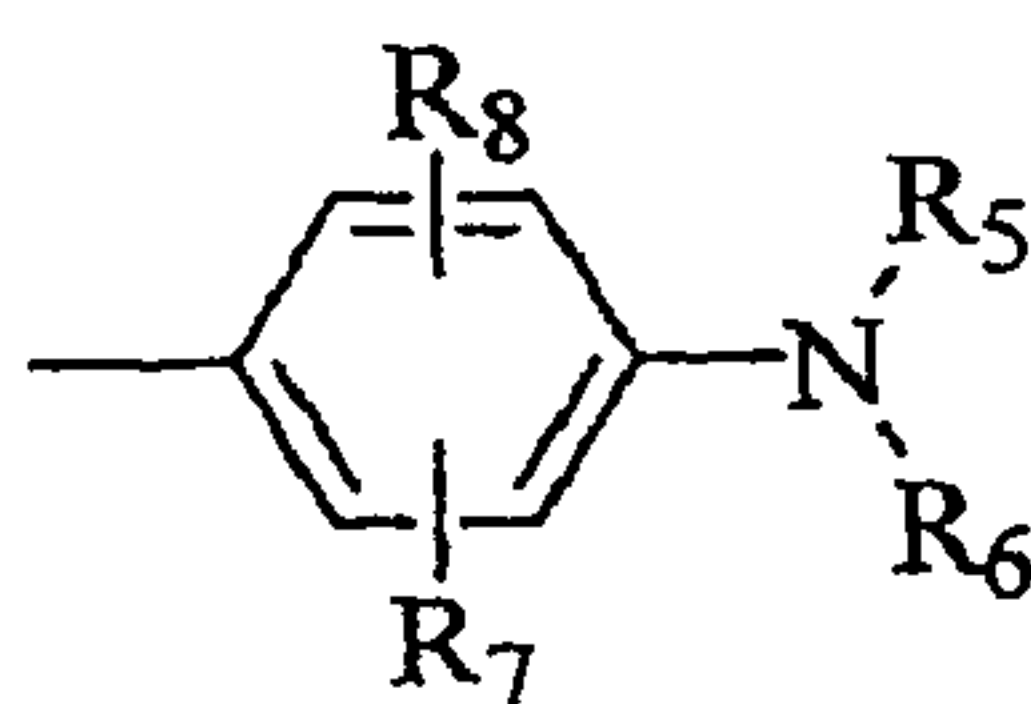
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benzyl moiety; a (C₁-C₂₂)-alkyl moiety terminated in sulfonate, sulfate, acetate, or carboxylate; or

ii.) when taken together form a saturated or unsaturated heterocyclic or carbocyclic moiety; or

iii.) when taken together form a saturated or unsaturated heterocyclic or carbocyclic moiety substituted by, sulfate, sulfonate, phosphate, nitrate, or carboxylate;

c.) X is moiety having Formula VII below;



Formula VII

wherein:

i.) R₅ and R₆:

(a) are independently hydrogen; a (C₁-C₂₂)-alkyl

moiety or (C₂-C₂₂)-alkenyl moiety; a hydroxy-

(C₂-C₂₂)-alkyl moiety; a hydroxy-(C₂-C₂₂)-alkyl moiety comprising, in addition to the hydroxyl oxygen, an

oxygen atom; a capped or uncapped polyether moiety;

an amino-(C₁-C₂₂)-alkyl moiety; a substituted or

unsubstituted phenyl moiety; a substituted or

unsubstituted benzyl moiety; a (C₁-C₂₂)-alkyl moiety

comprising a terminating C₁-C₄ alkyl ether, sulfonate,

sulfate, acetate or carboxylate moiety; a thiazole moiety

or

(b) when taken together form a saturated or unsaturated

heterocyclic moiety; or

(c) when taken together form a saturated or unsaturated

heterocyclic moiety substituted by one or more,

alkoxylate, sulfate, sulfonate, phosphate, nitrate, or

carboxylate moieties; or

(d) when taken together with R₇, R₈, or R₇ and R₈ form one

or more saturated or unsaturated heterocyclic moieties,

59

optionally substituted by one or more alkoxyate, sulfate, sulfonate, phosphate, nitrate, or carboxylate moieties; or

(e) when taken together form a thiazole moiety;

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ii.) R_7 and R_8 are independently hydrogen or an alkyl moiety or alkenyl moiety;

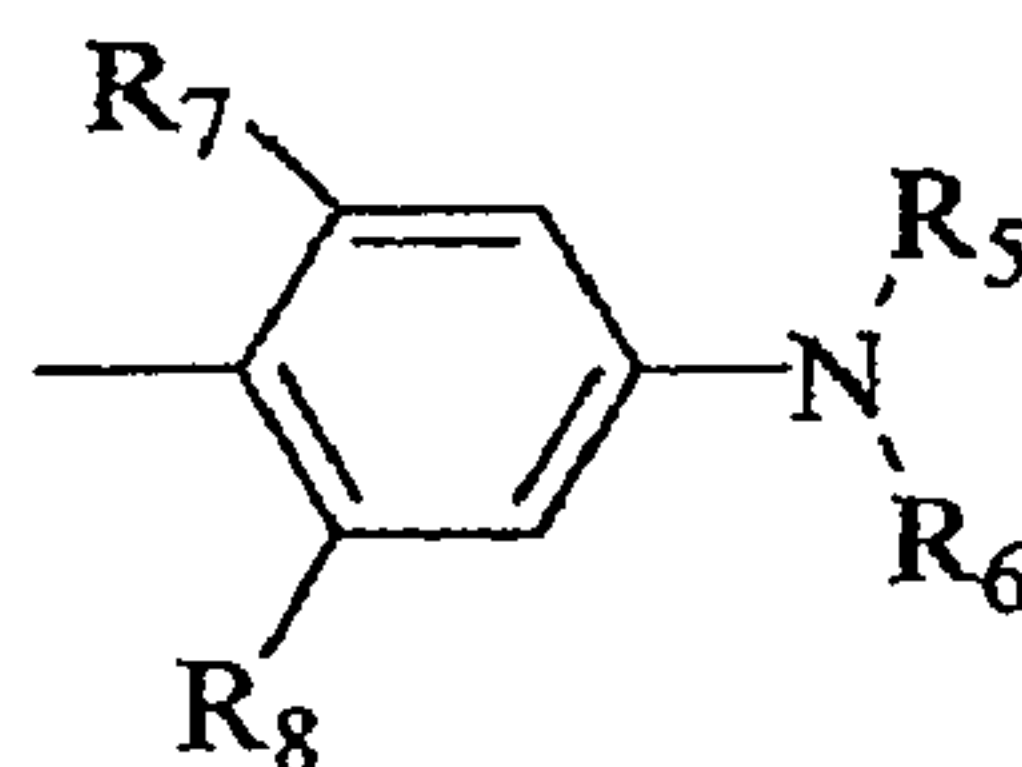
d.) Q^- is an anion that balances the overall charge of the compound of Formula V, and the index q is 0 or 1.

2. The whitening agent of Claim 1, wherein:

a.) R_1 is a methyl moiety;

b.) R_3 and R_4 are hydrogen; and

c.) X has Formula VIII below:



Formula VIII

wherein

(i) R_5 and R_6 are as defined in Claim 1;

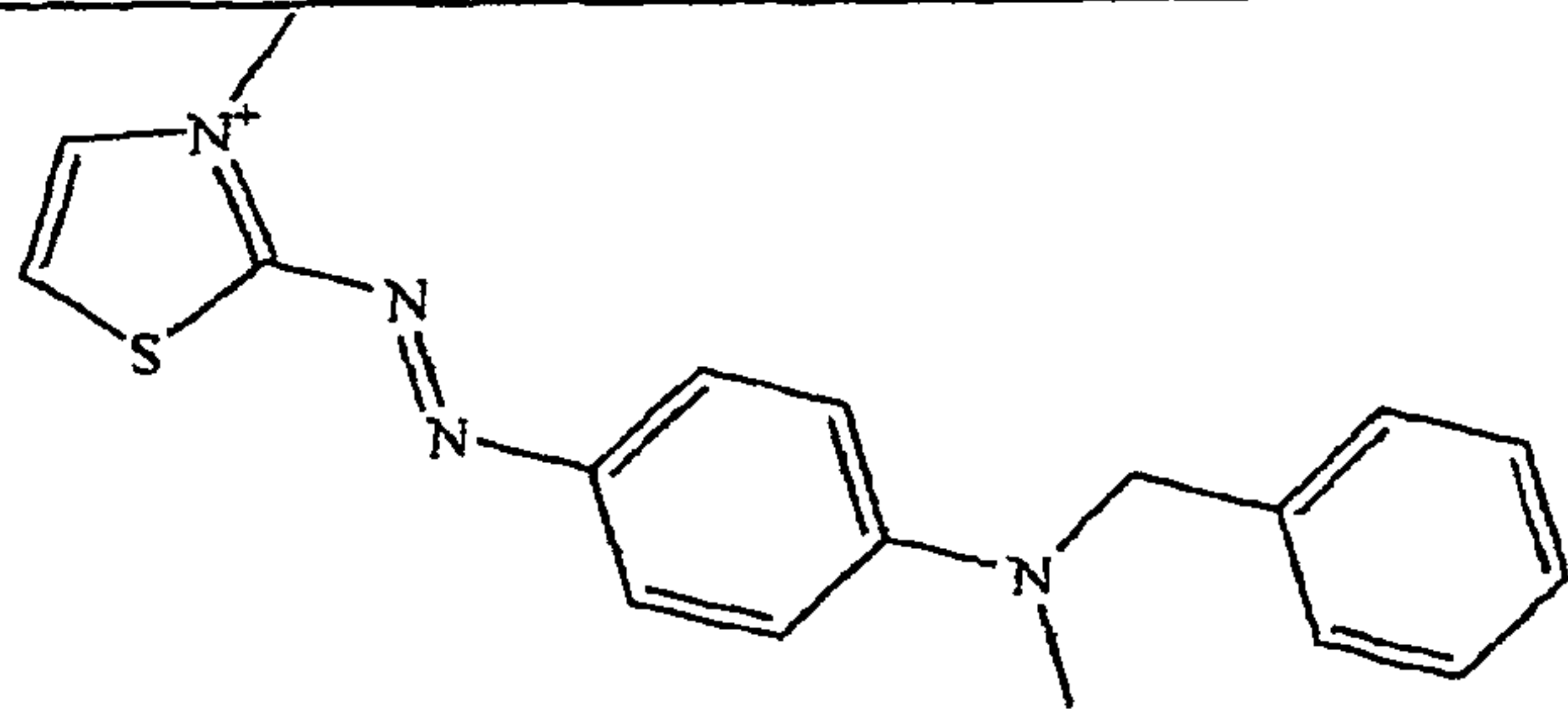
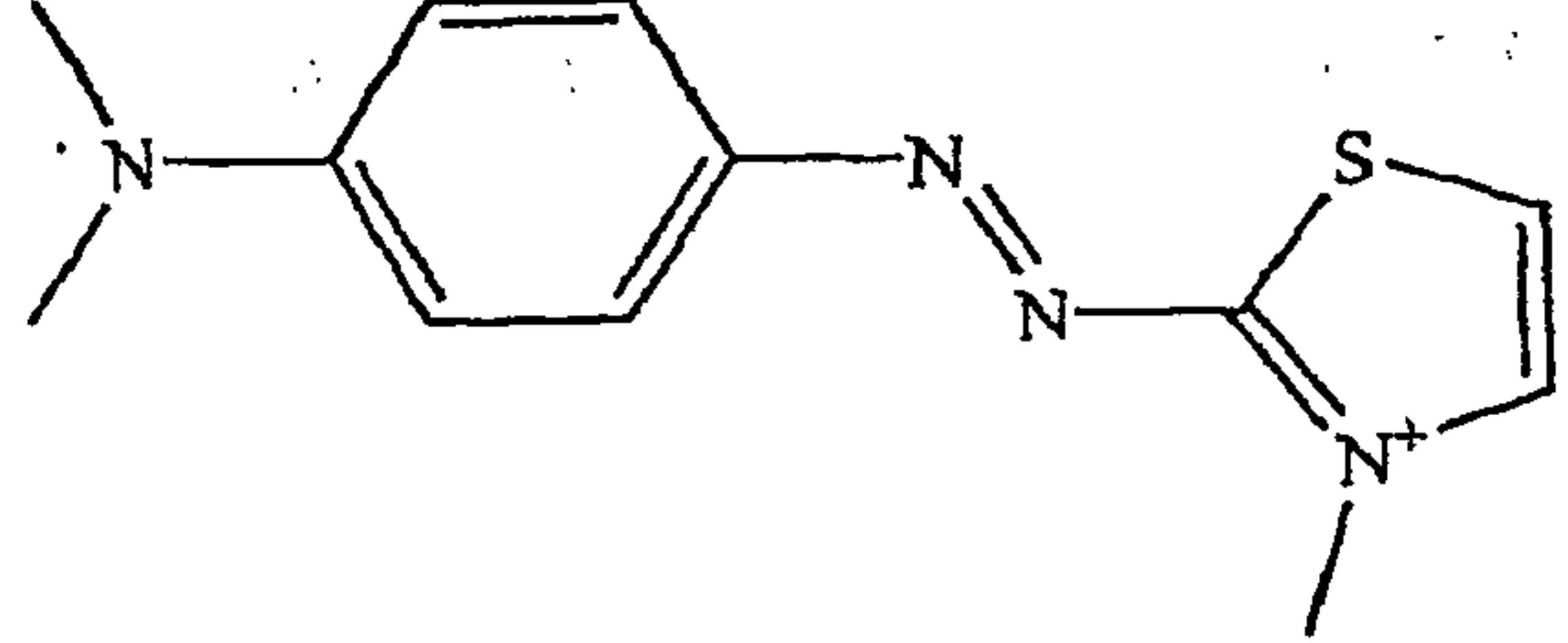
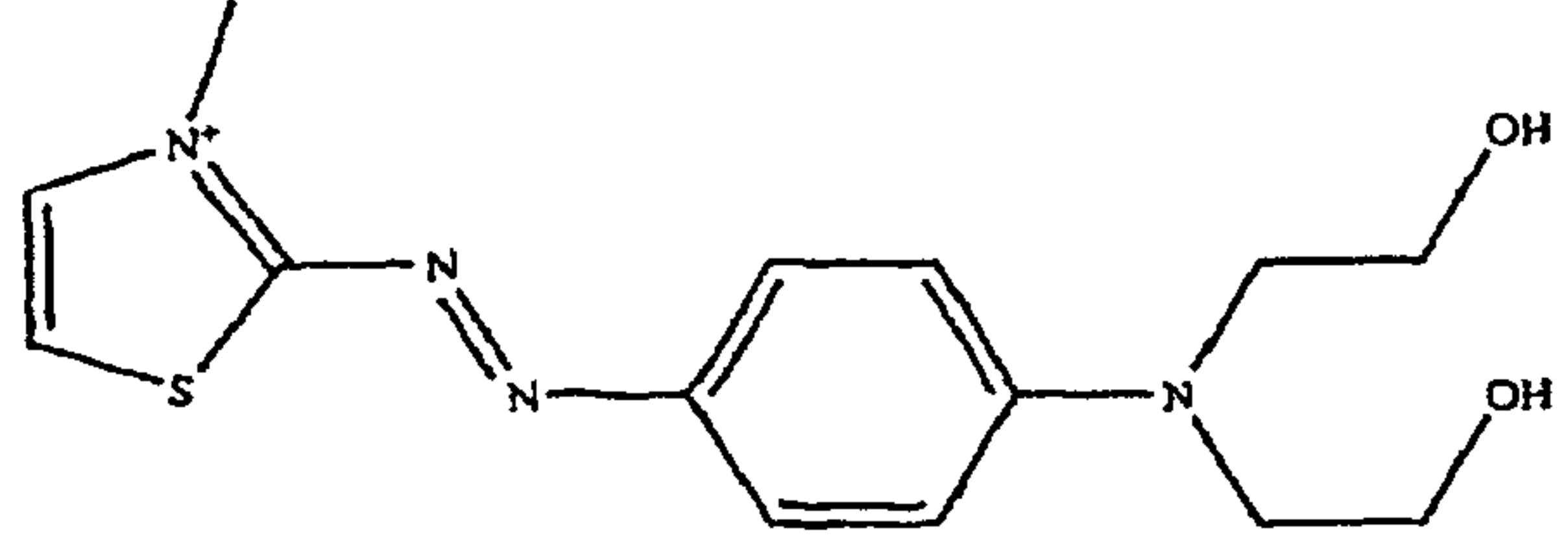
(ii) R_7 is hydrogen or a methyl moiety; and

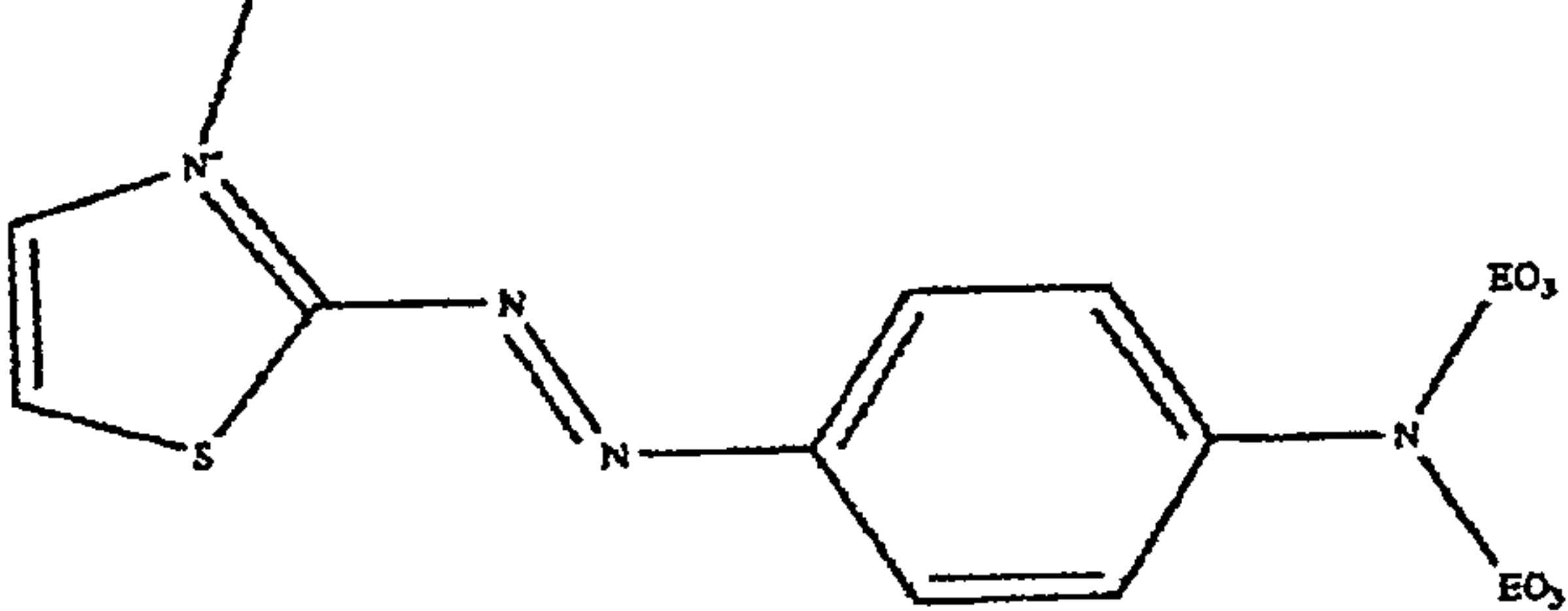
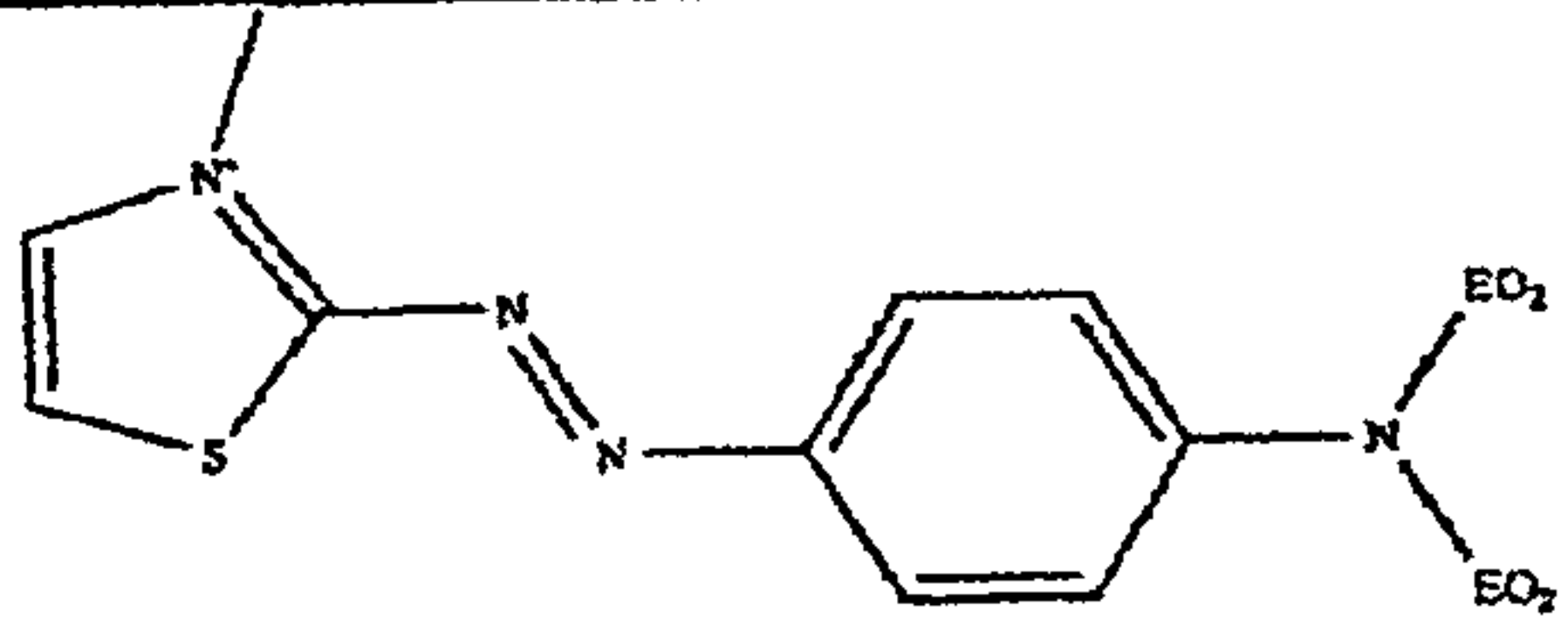
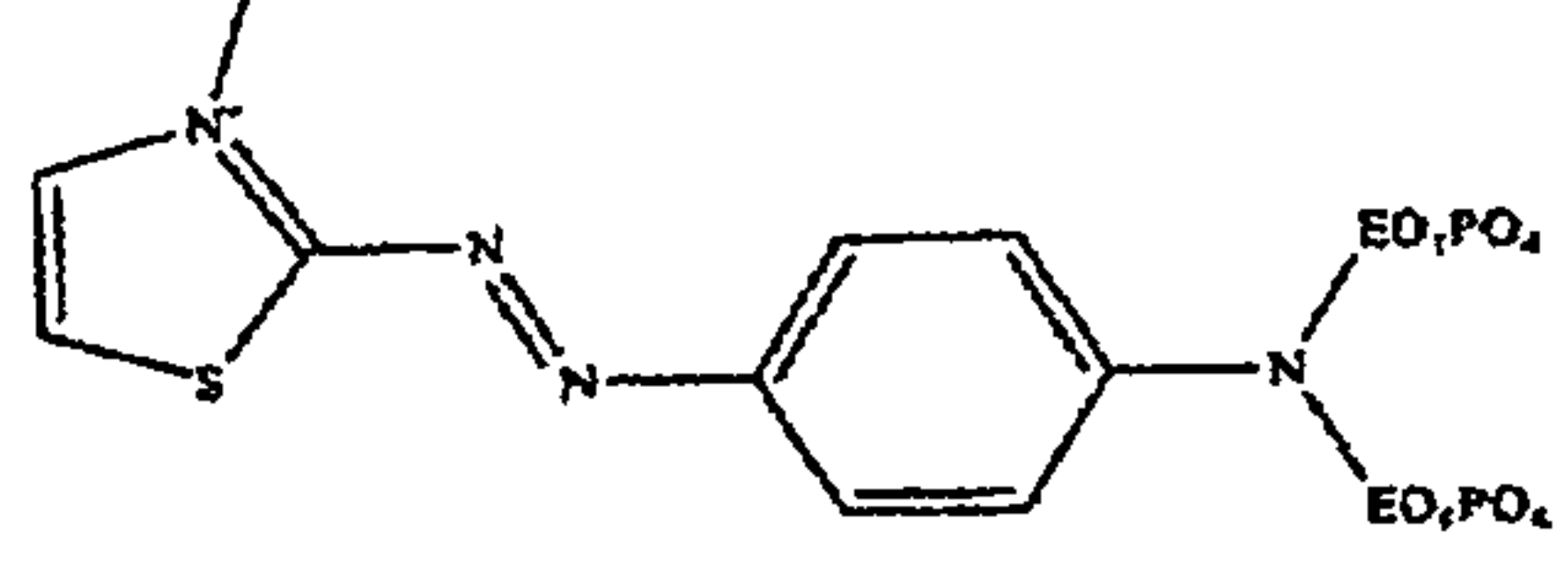
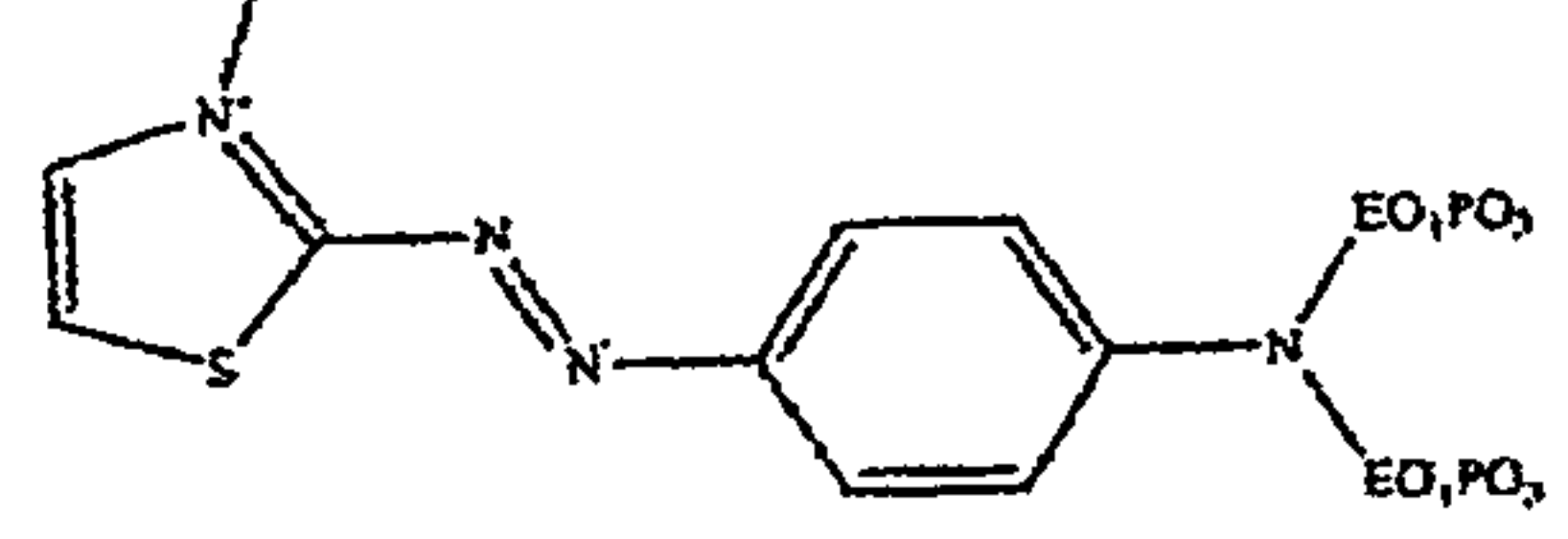
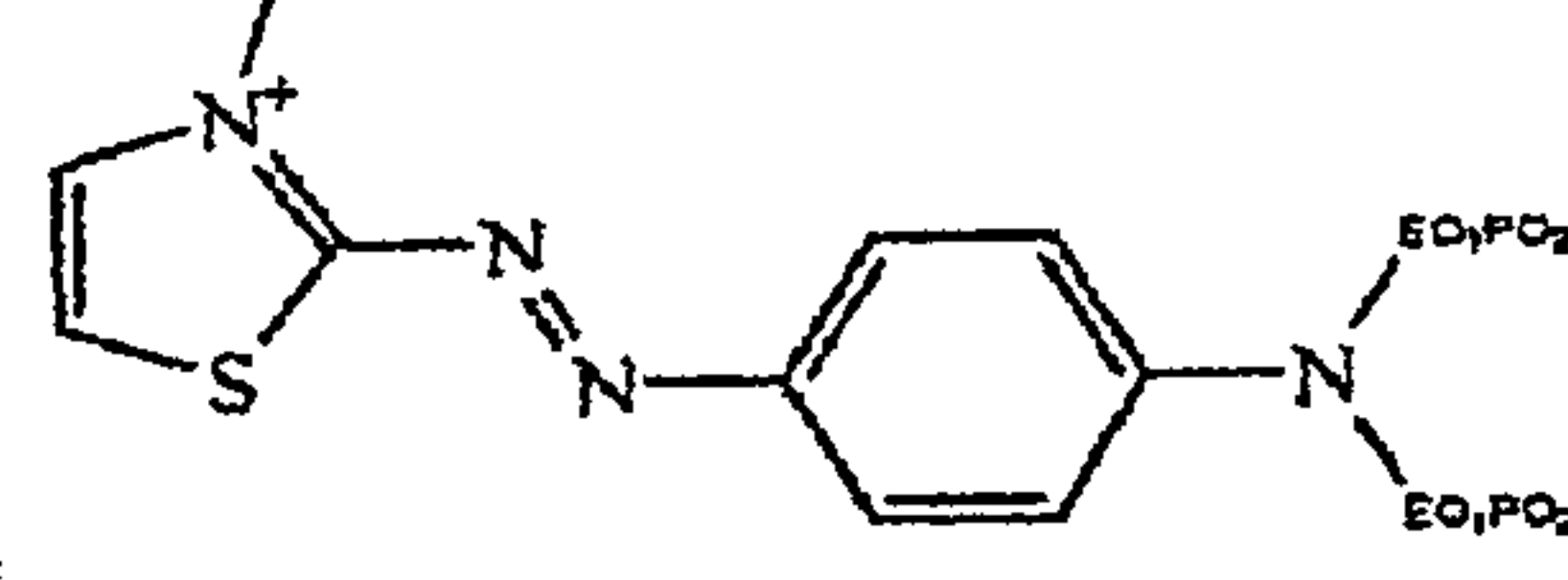
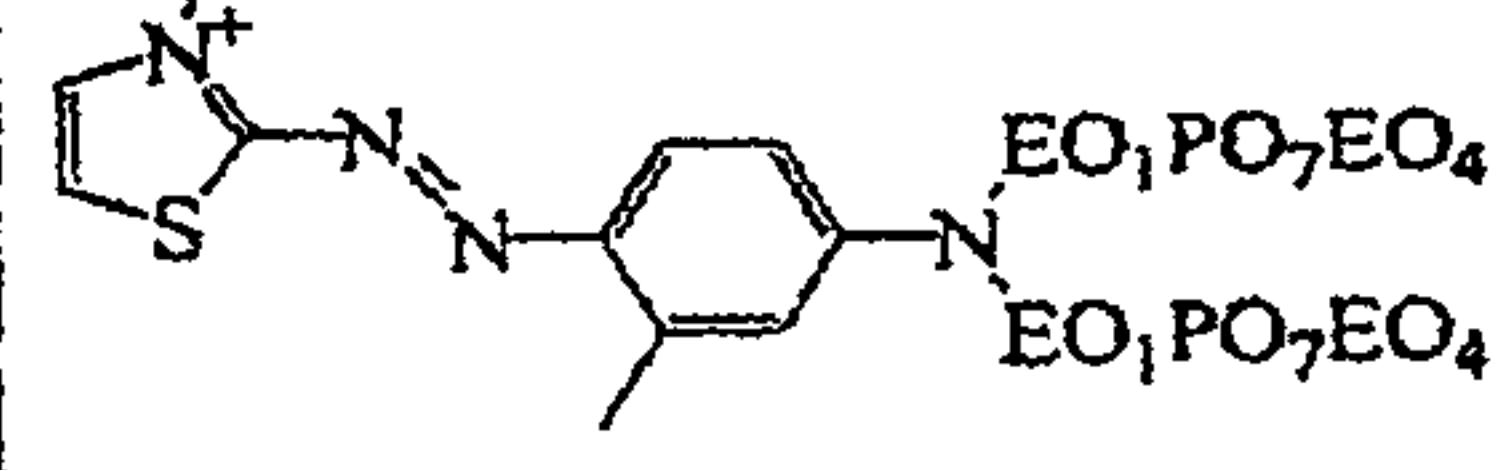
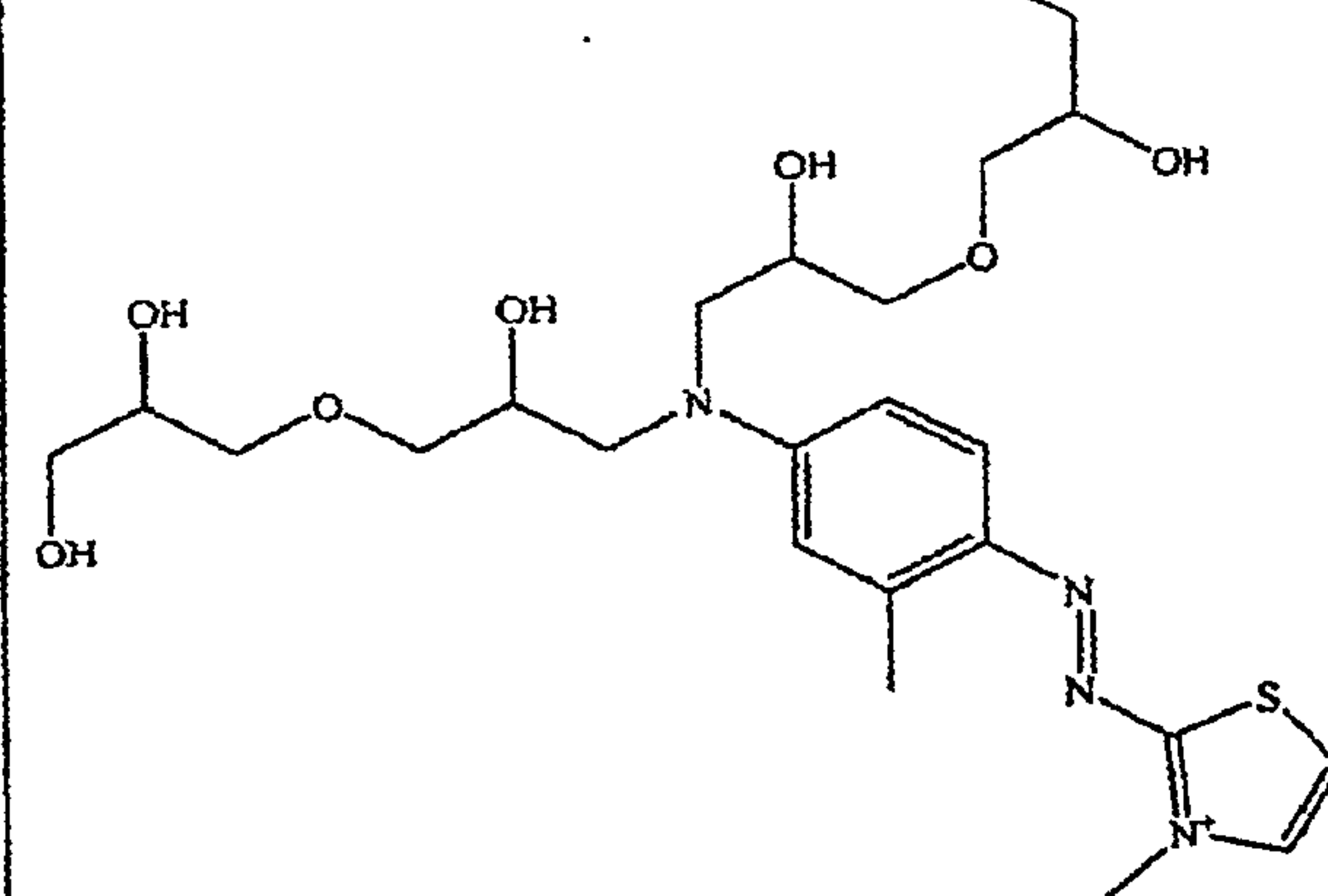
(iii) R_8 is hydrogen.

3. The whitening agent of Claim 1, wherein R_5 and R_6 each is, independently, from 1 to 20 alkylene oxide units and, independently, a moiety selected from the group consisting of: styrene oxide, glycidyl methyl ether, isobutyl glycidyl ether, isopropylglycidyl ether, t-butyl glycidyl ether, 2-ethylhexylglycidyl ether, and glycidylhexadecyl ether.

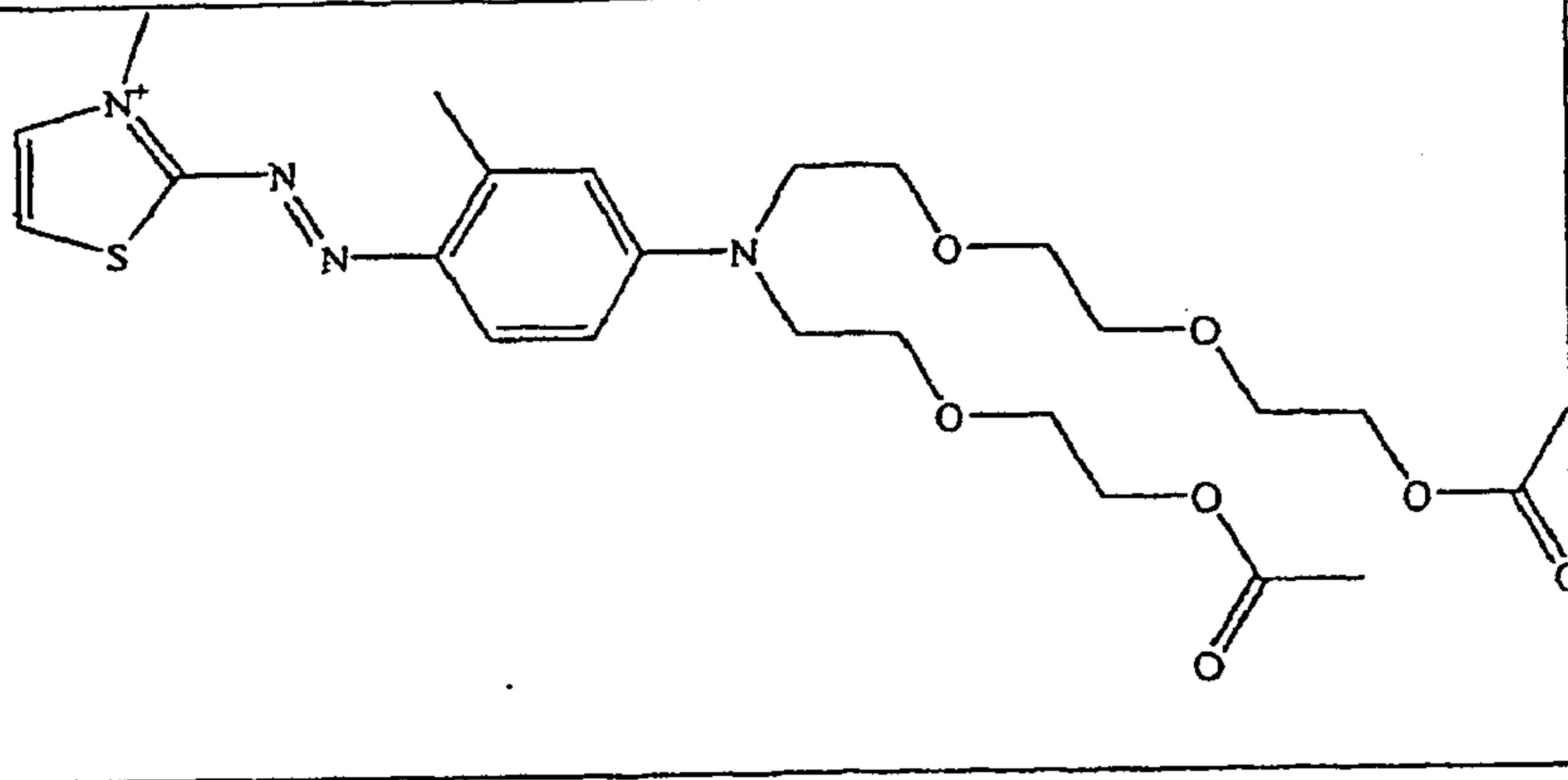
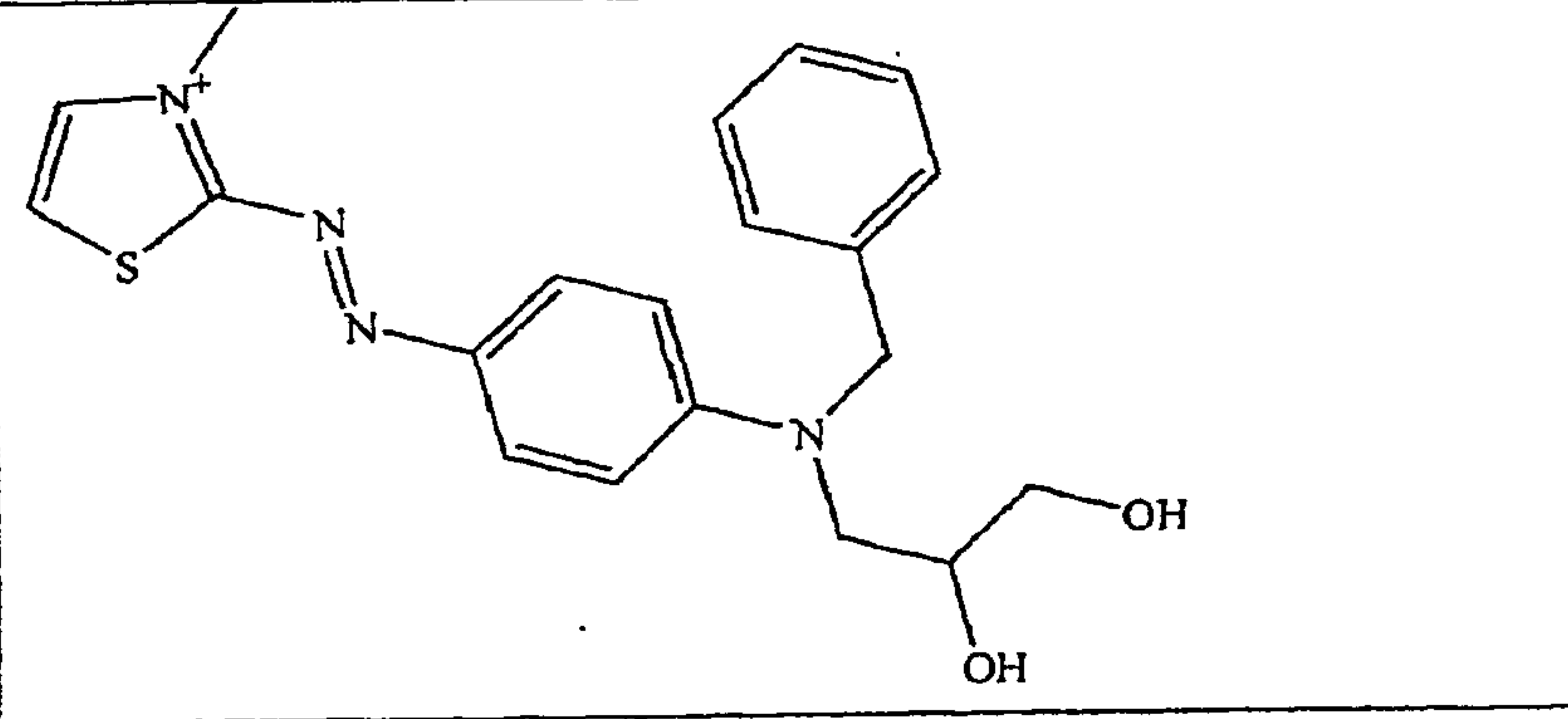
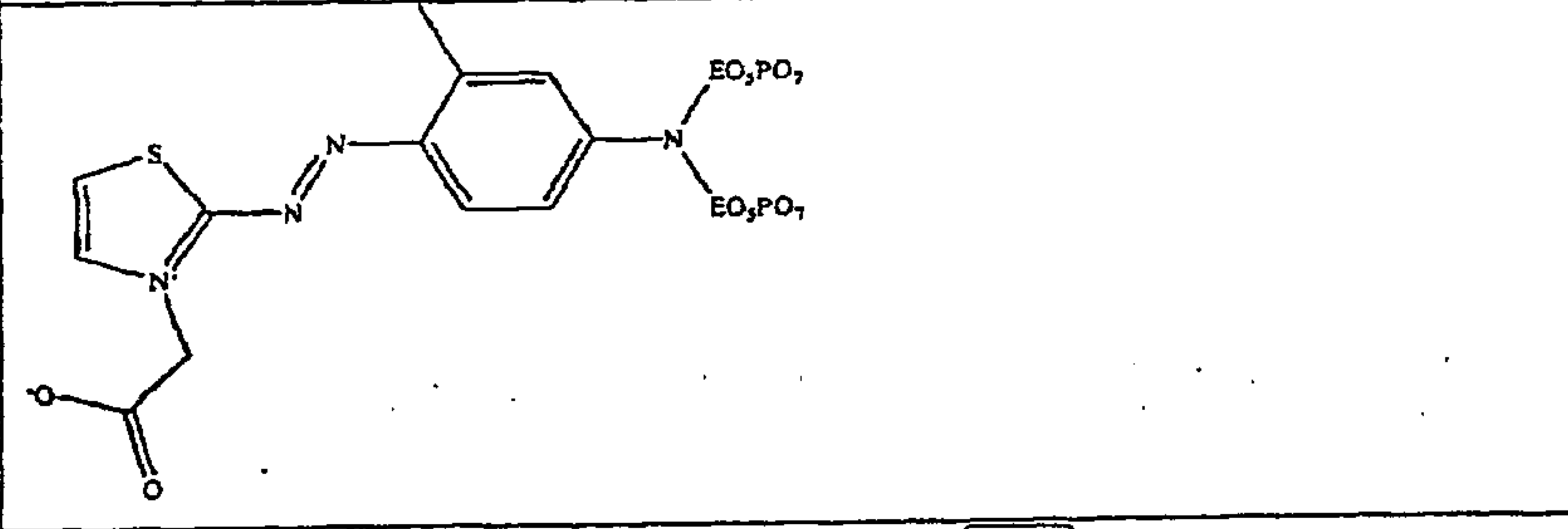
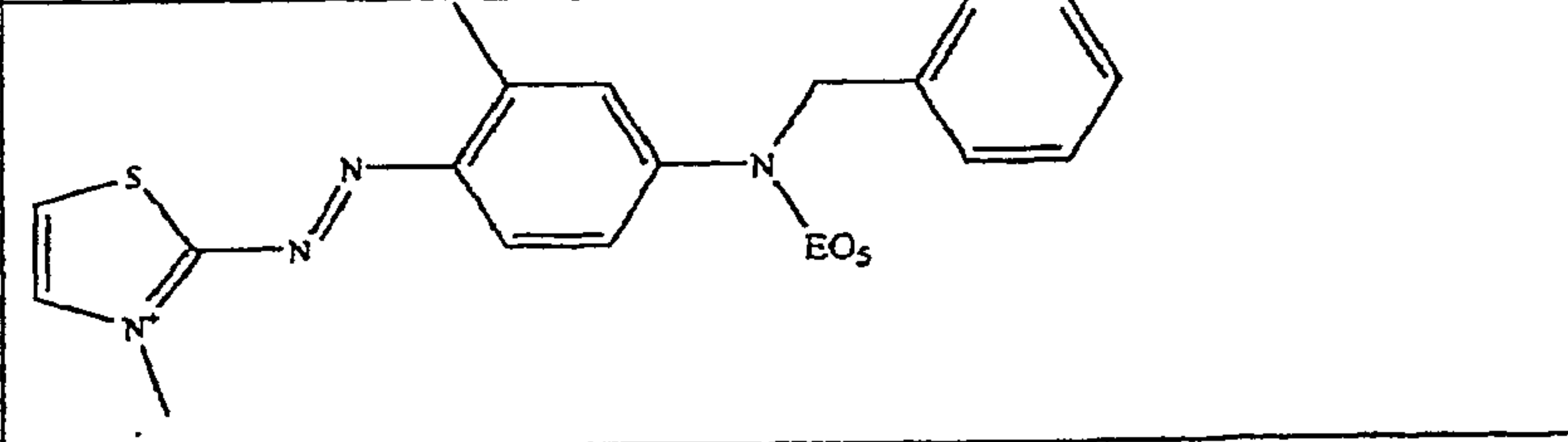
4. The whitening agent of Claim 1, wherein the thiazolium dye is one of Table 1
Thiazolium Dyes 1 through 80 or mixtures thereof, wherein Table 1 is as follows:

Table 1

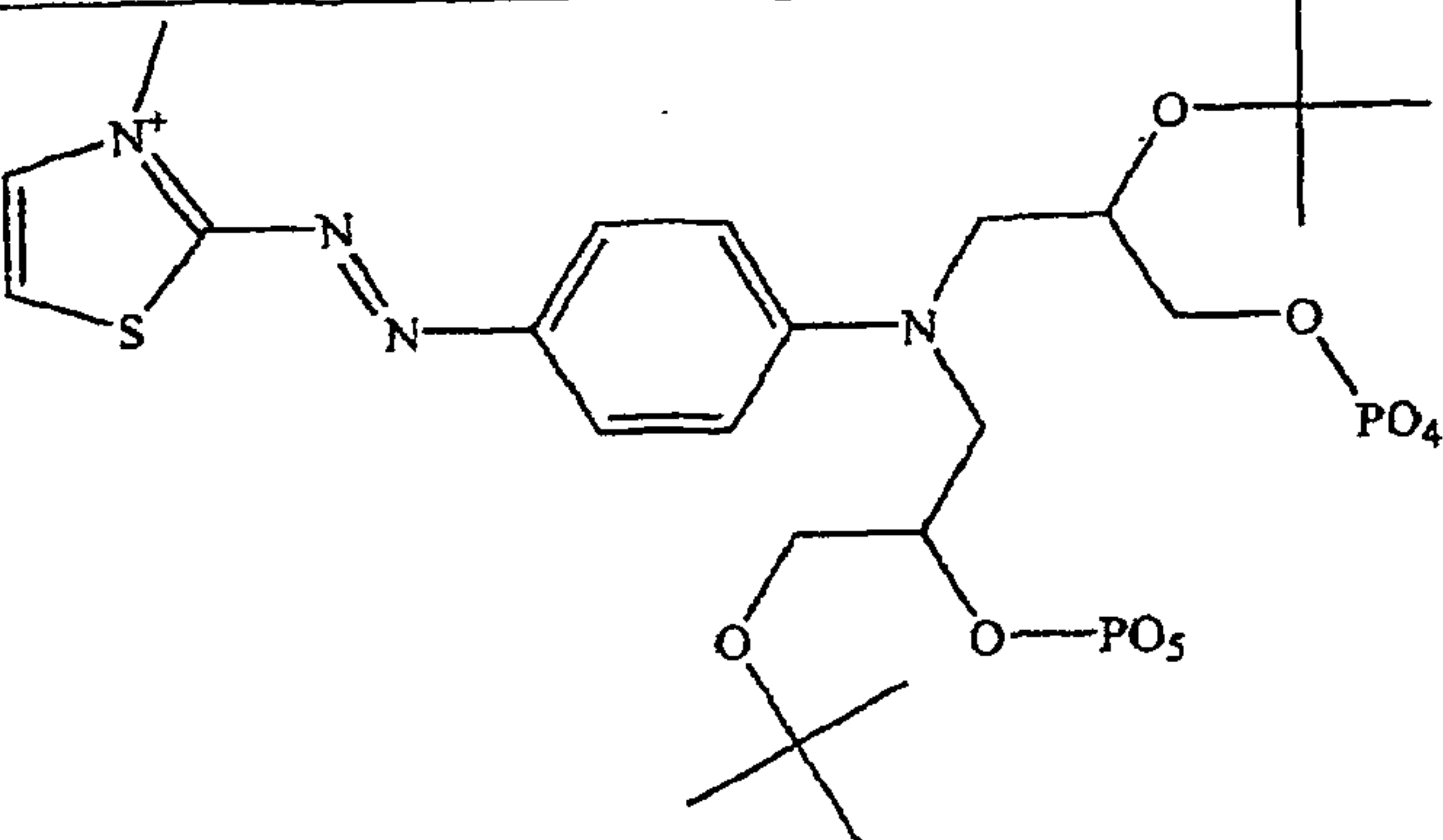
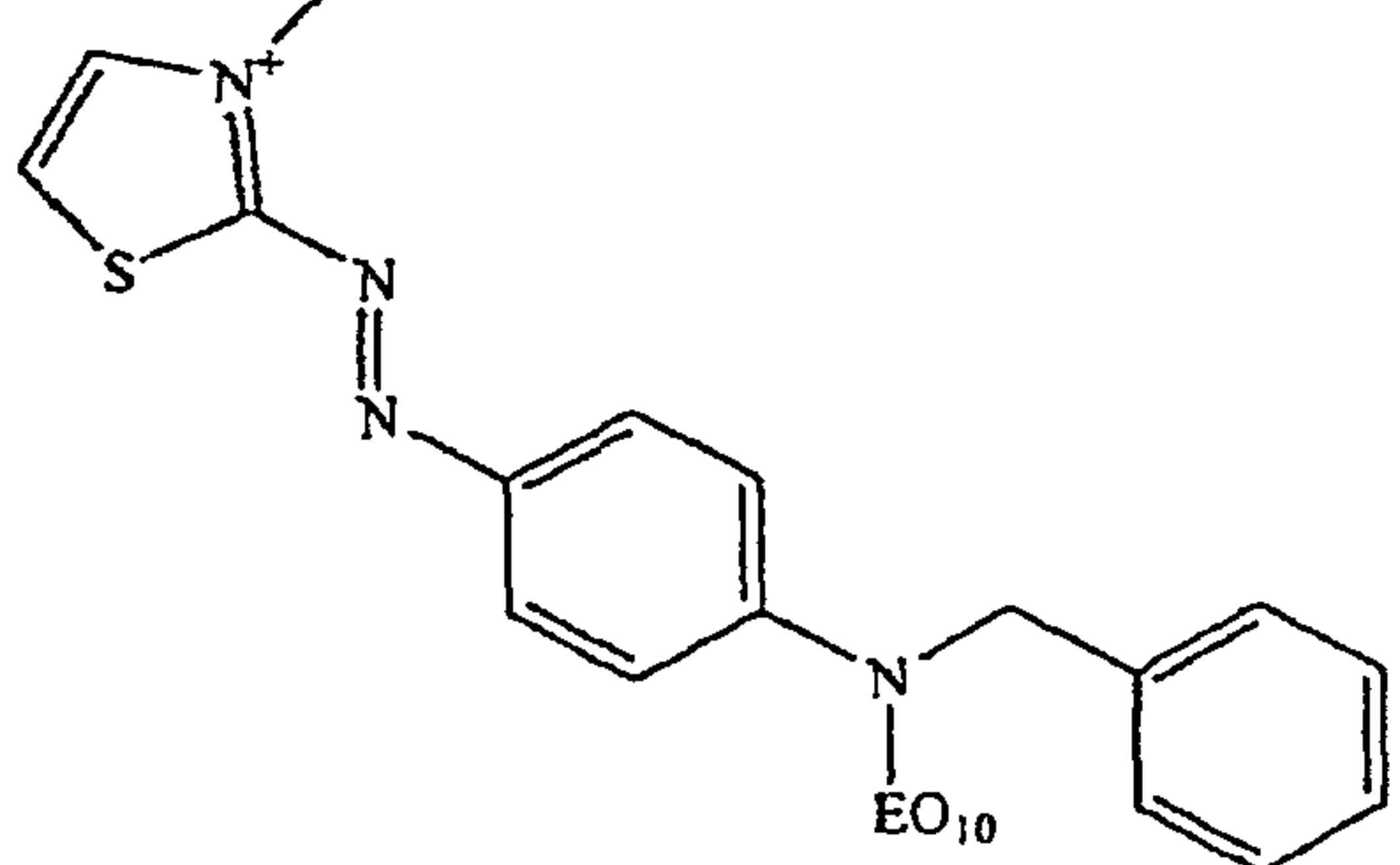
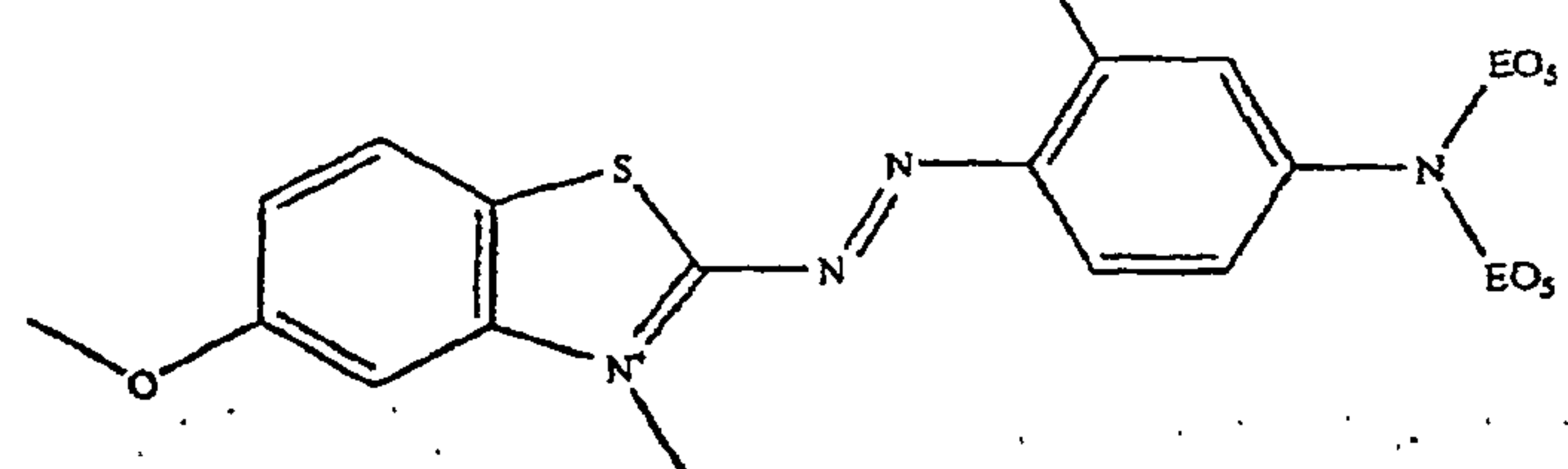
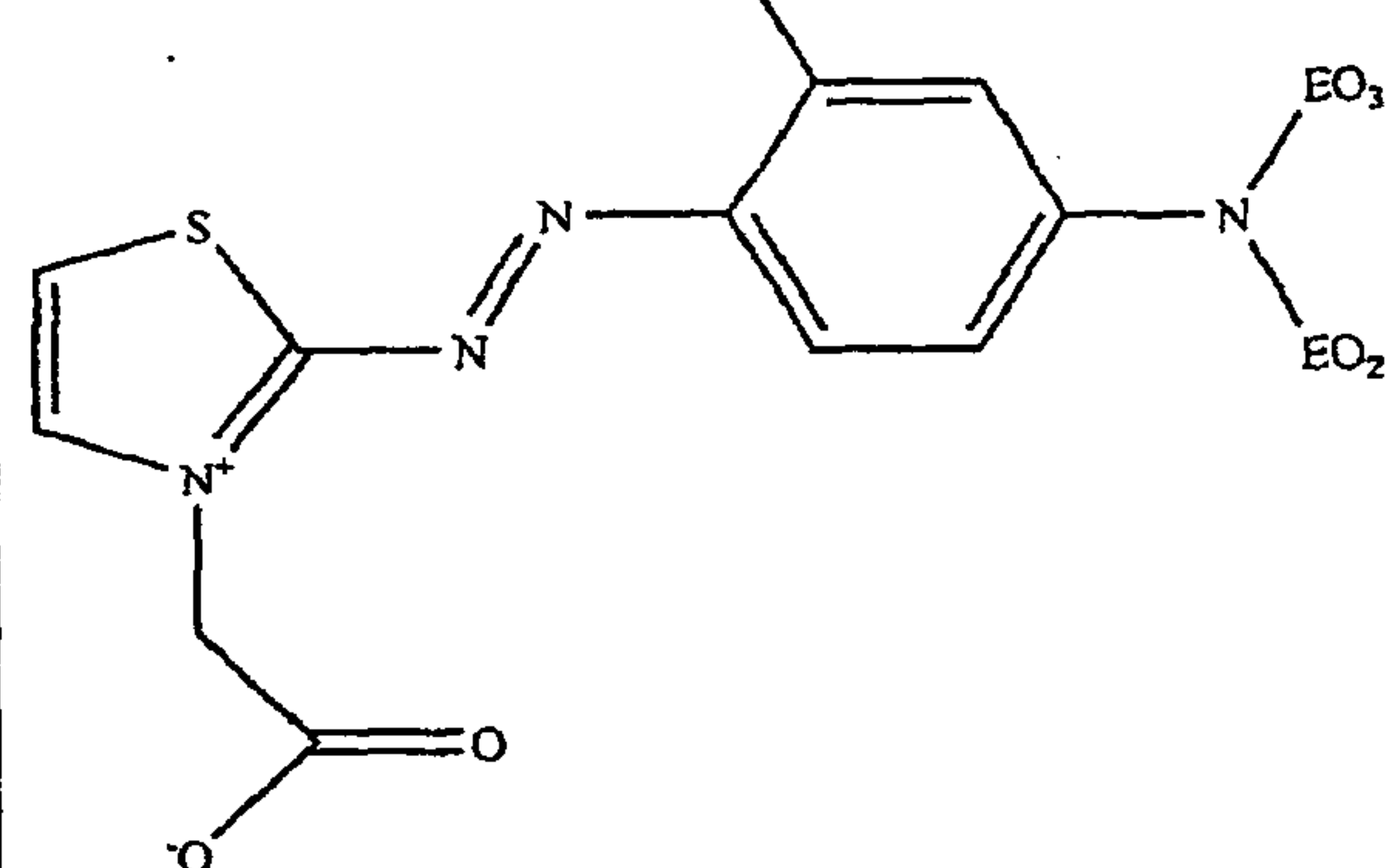
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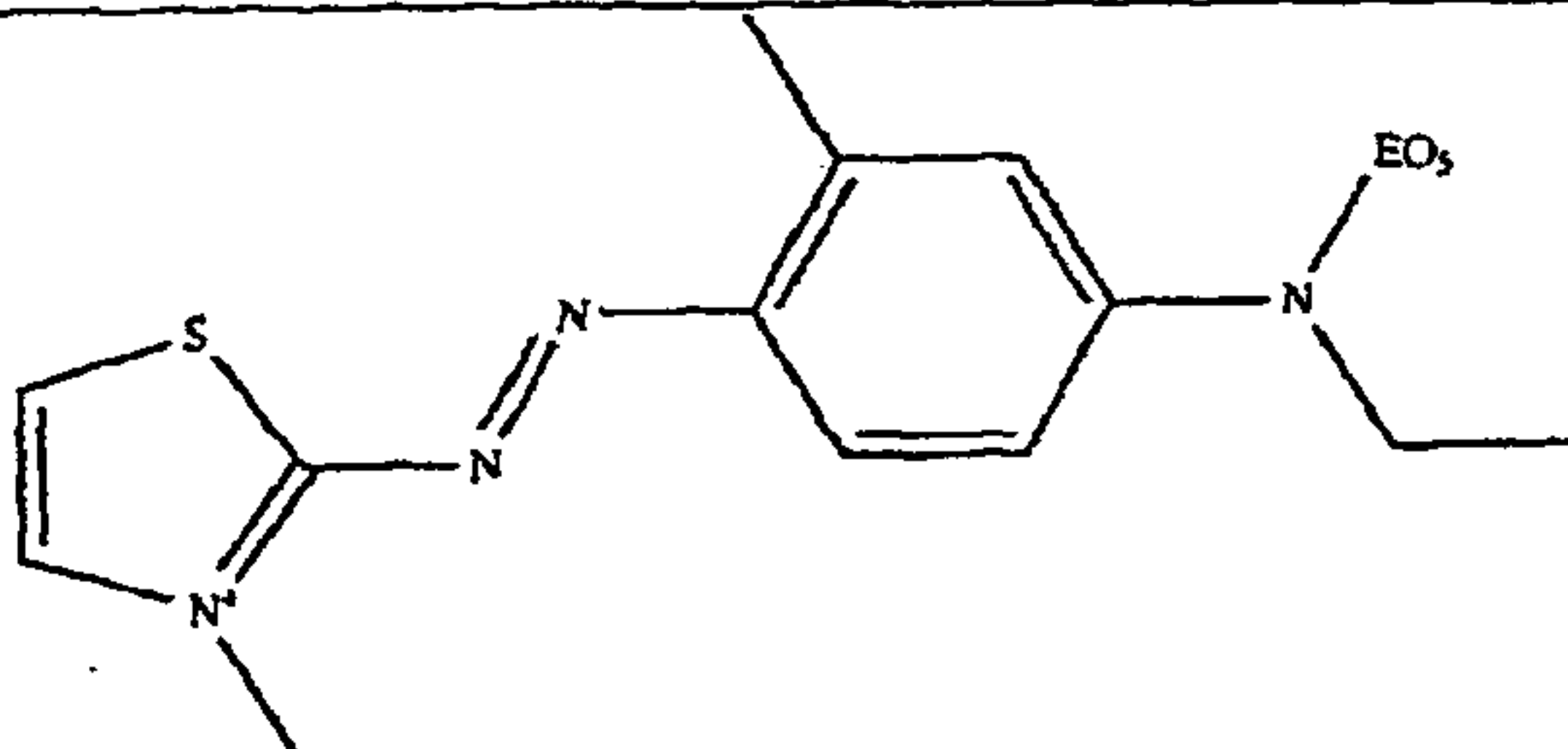
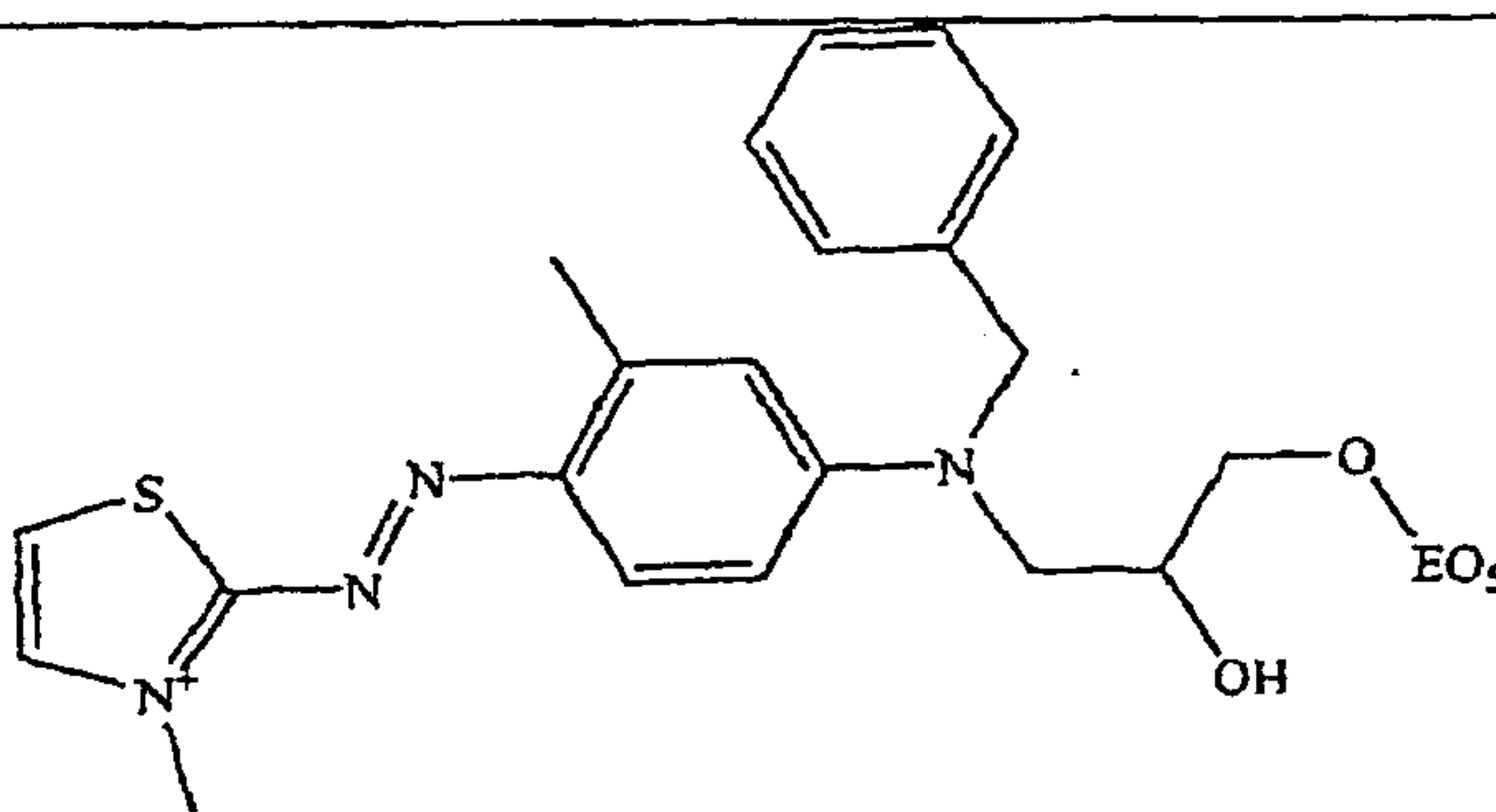
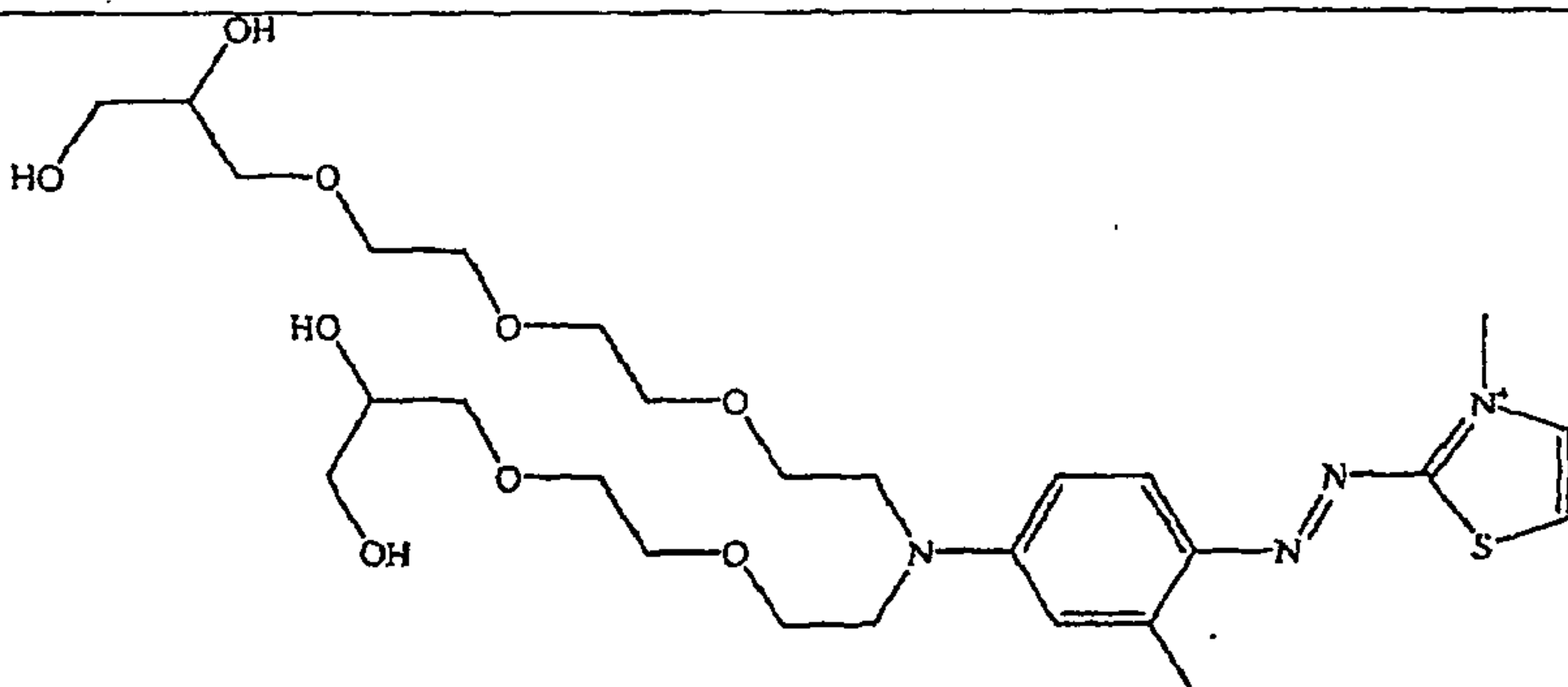
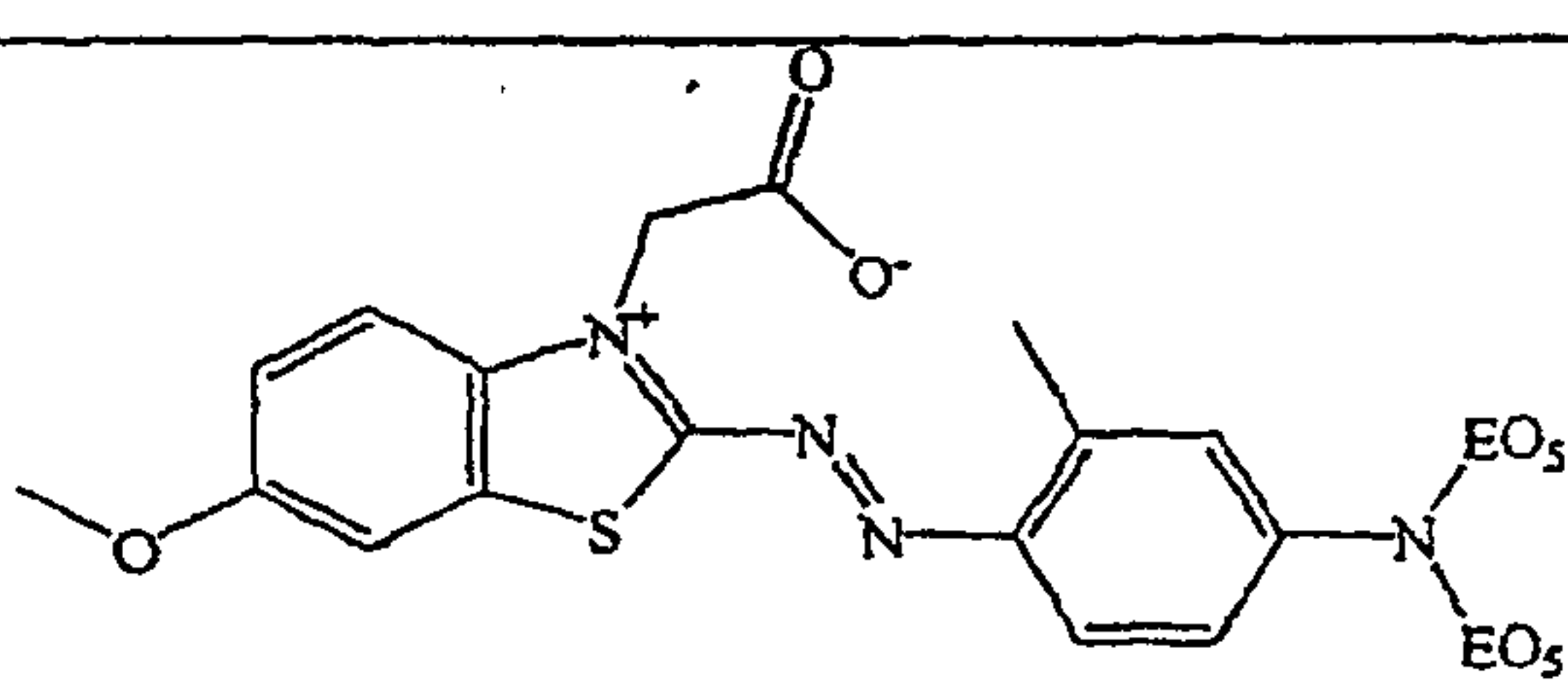
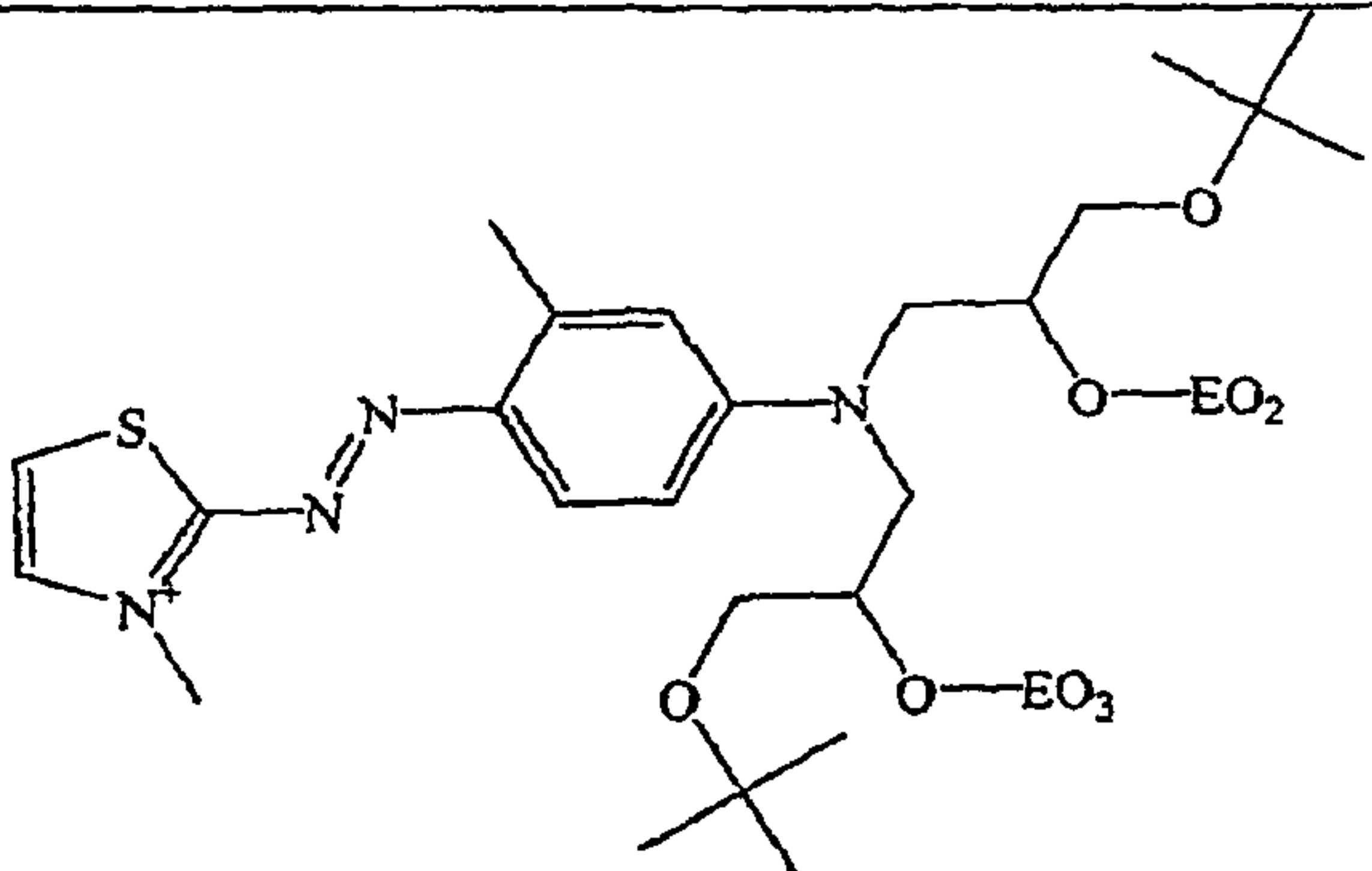
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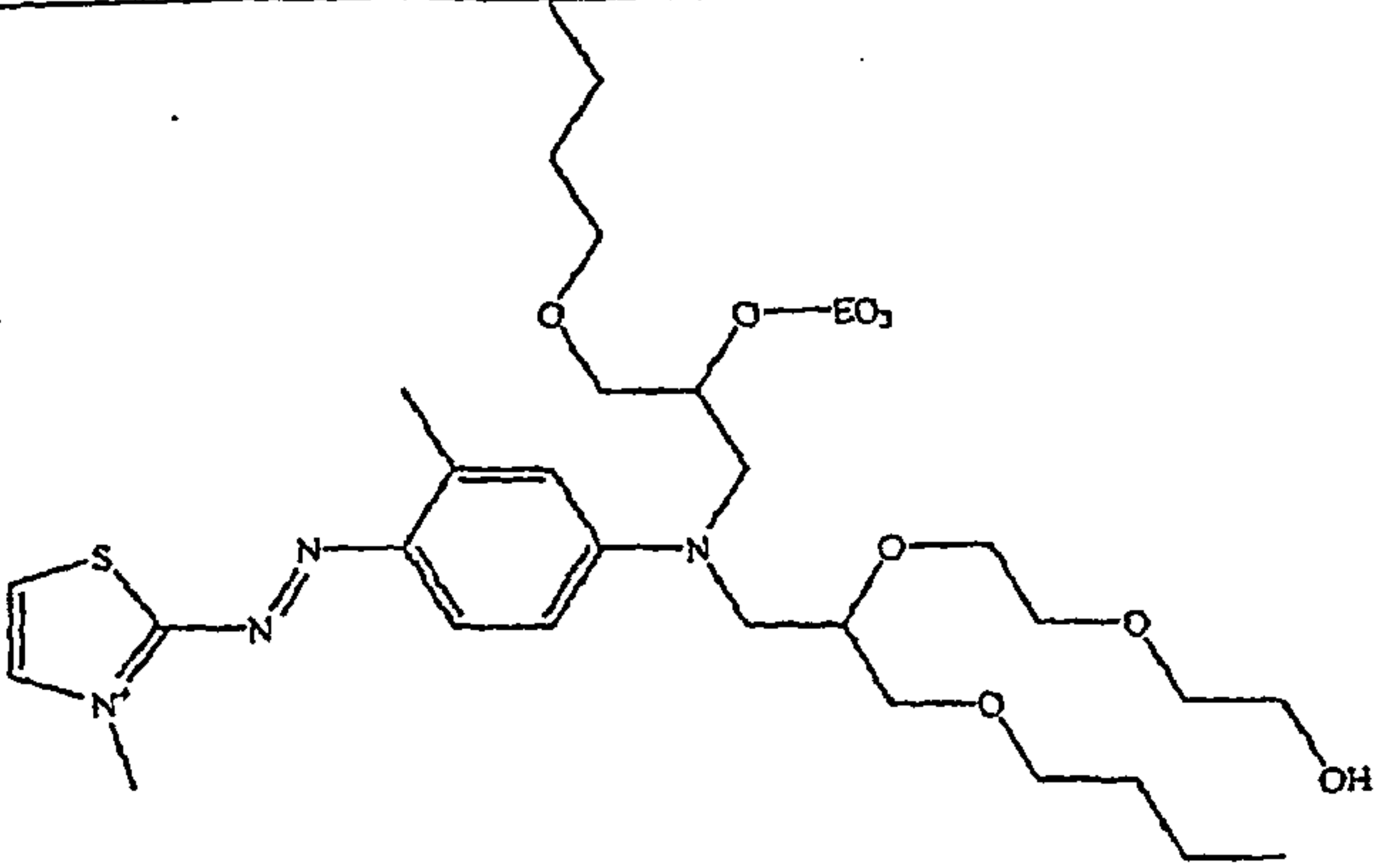
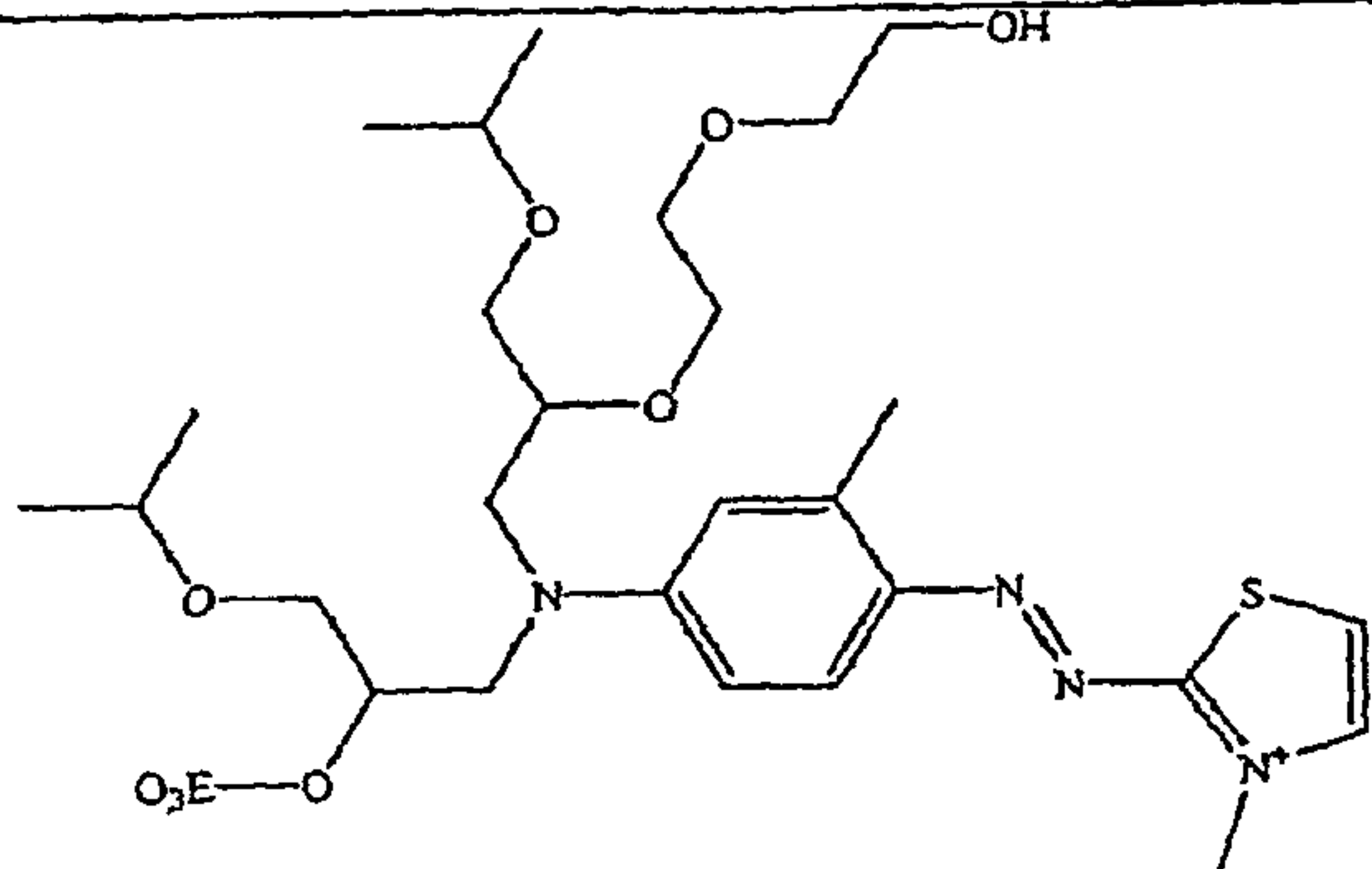
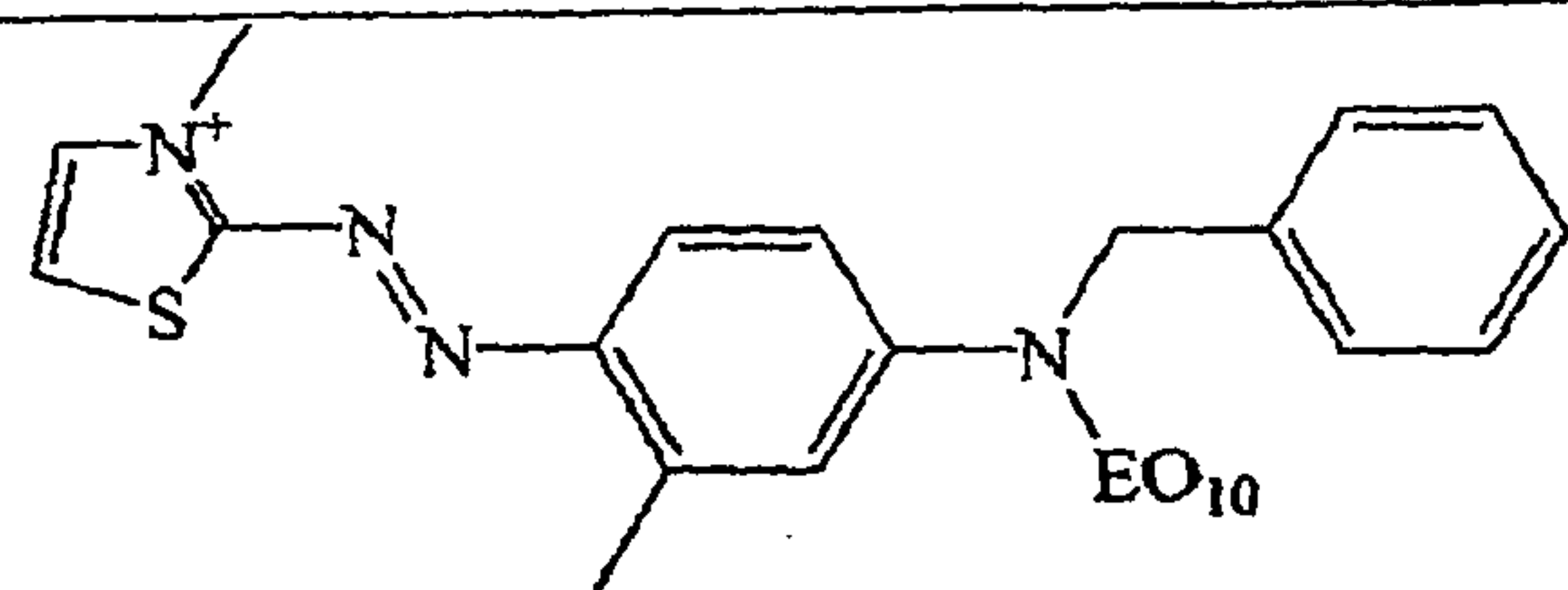
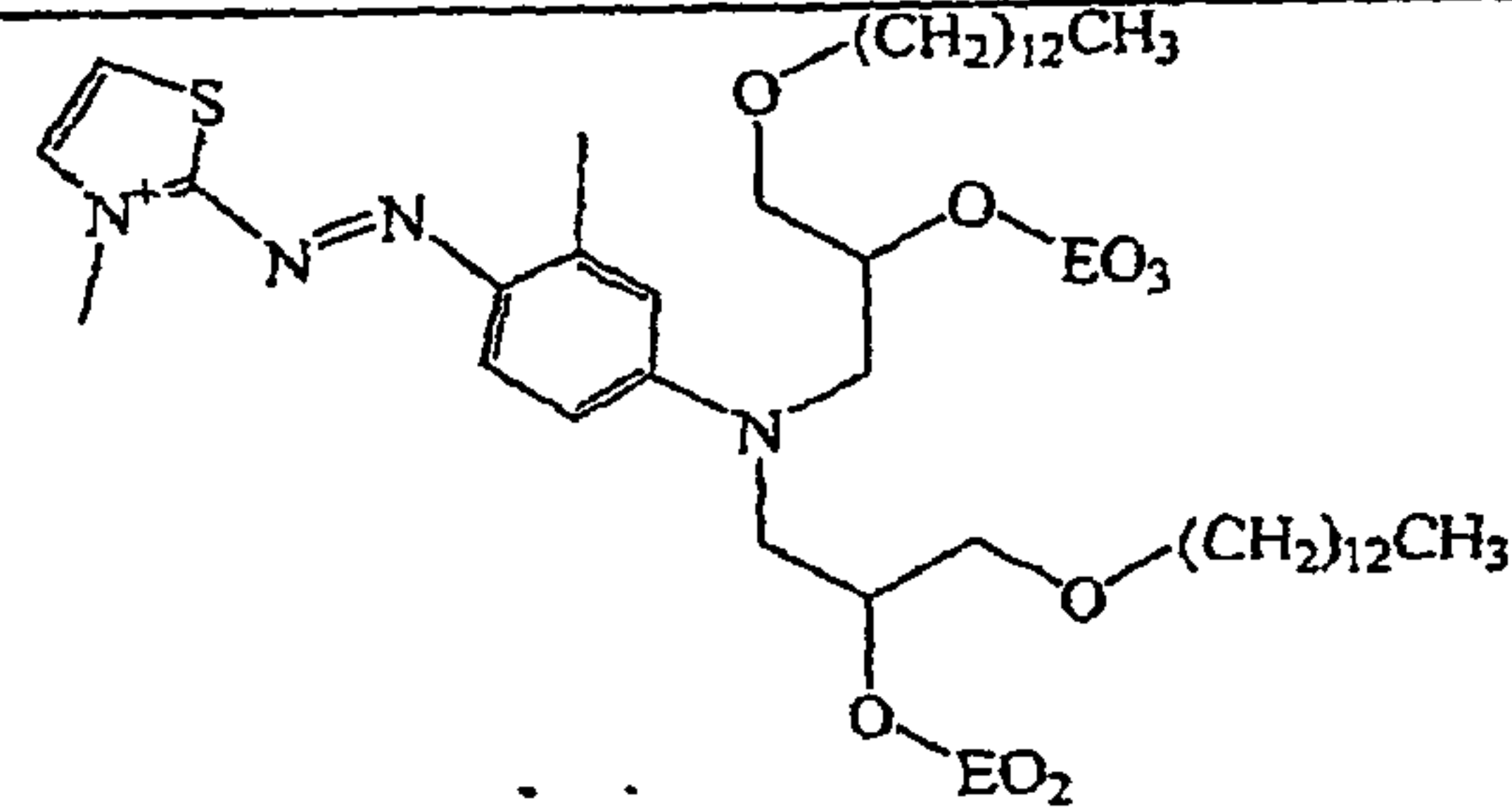
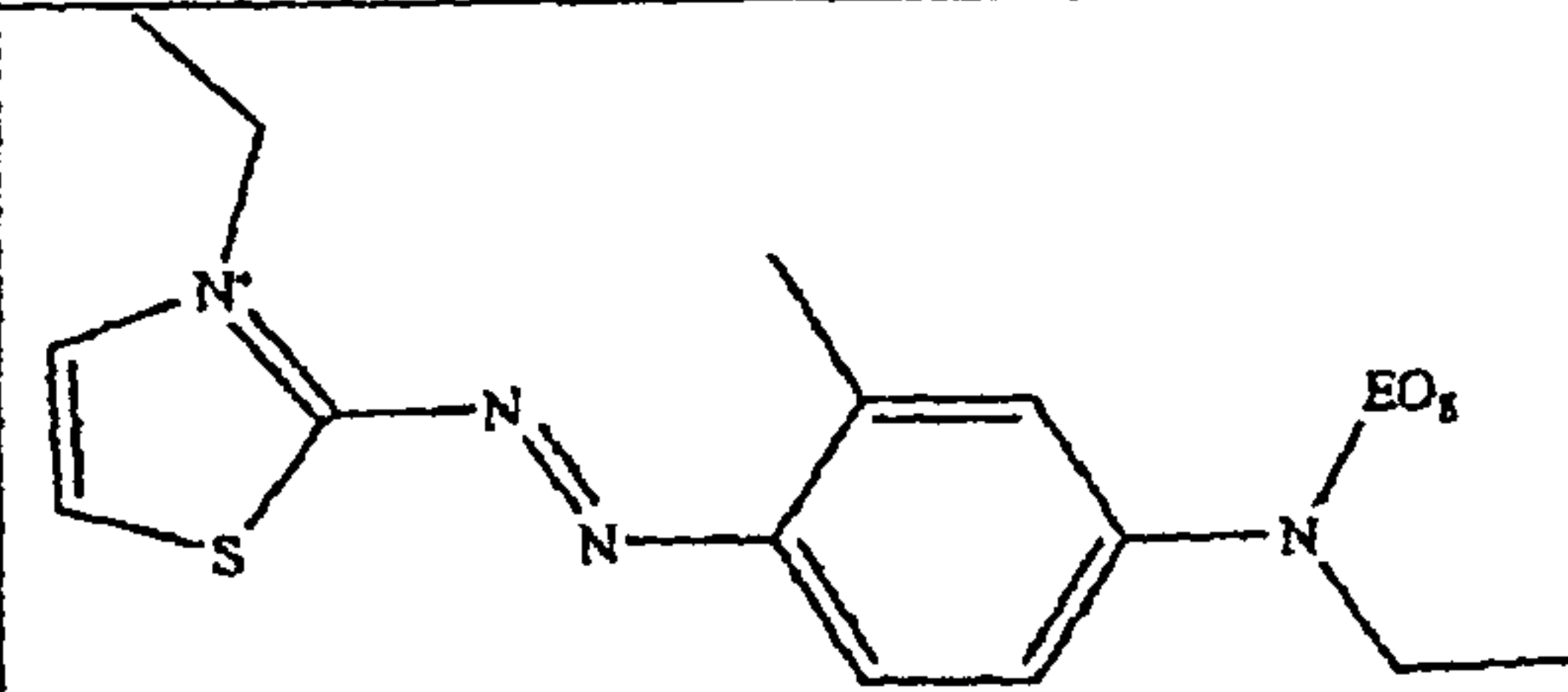
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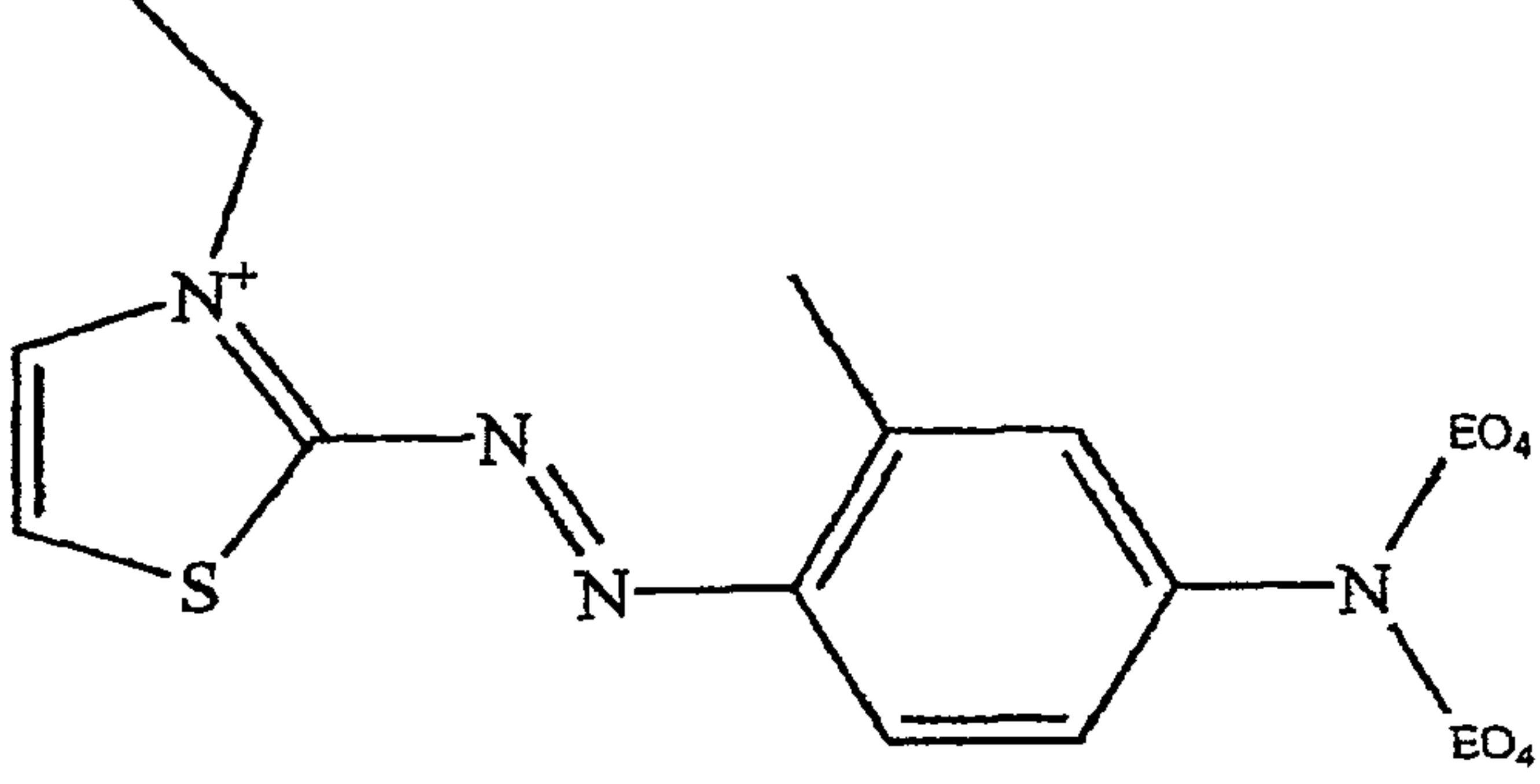
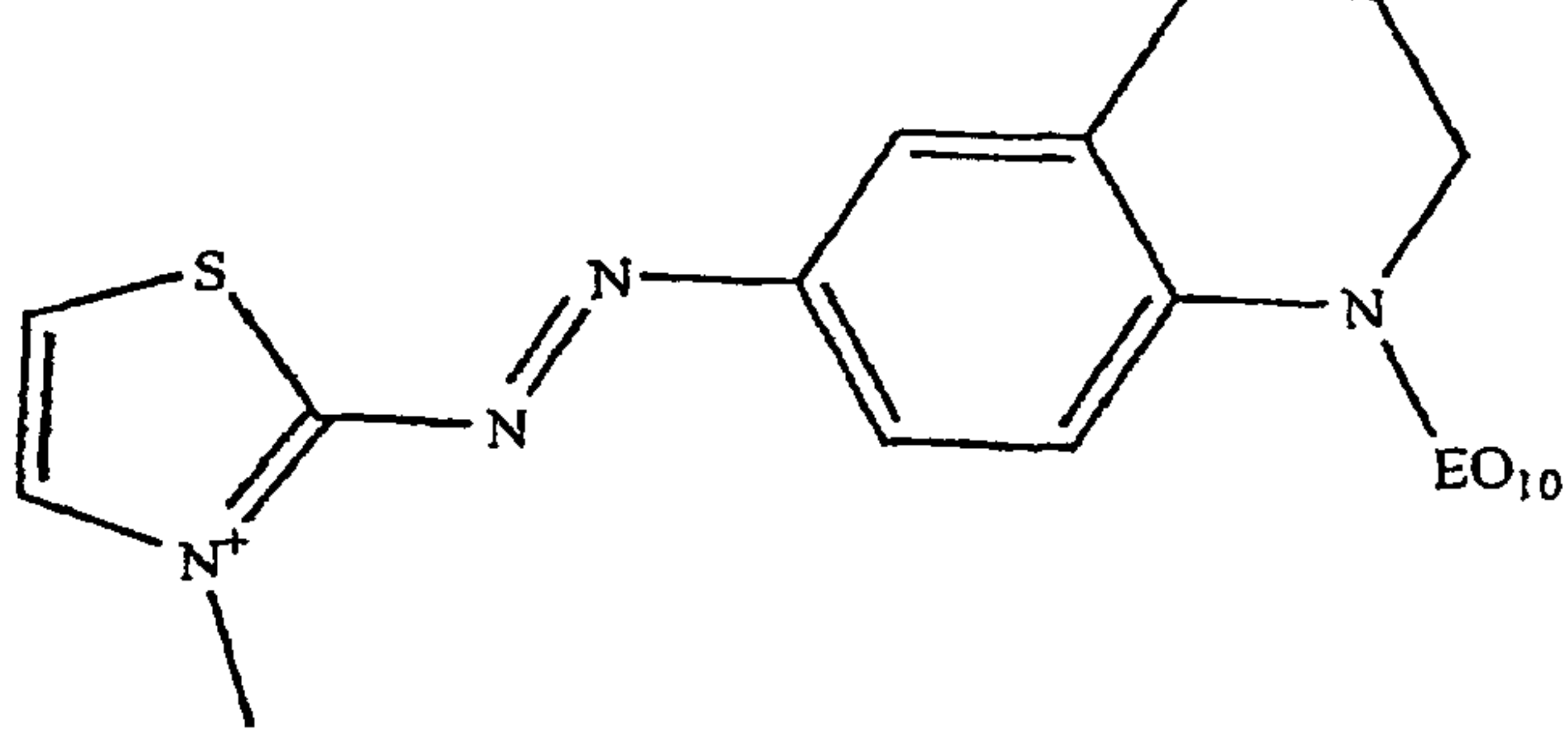
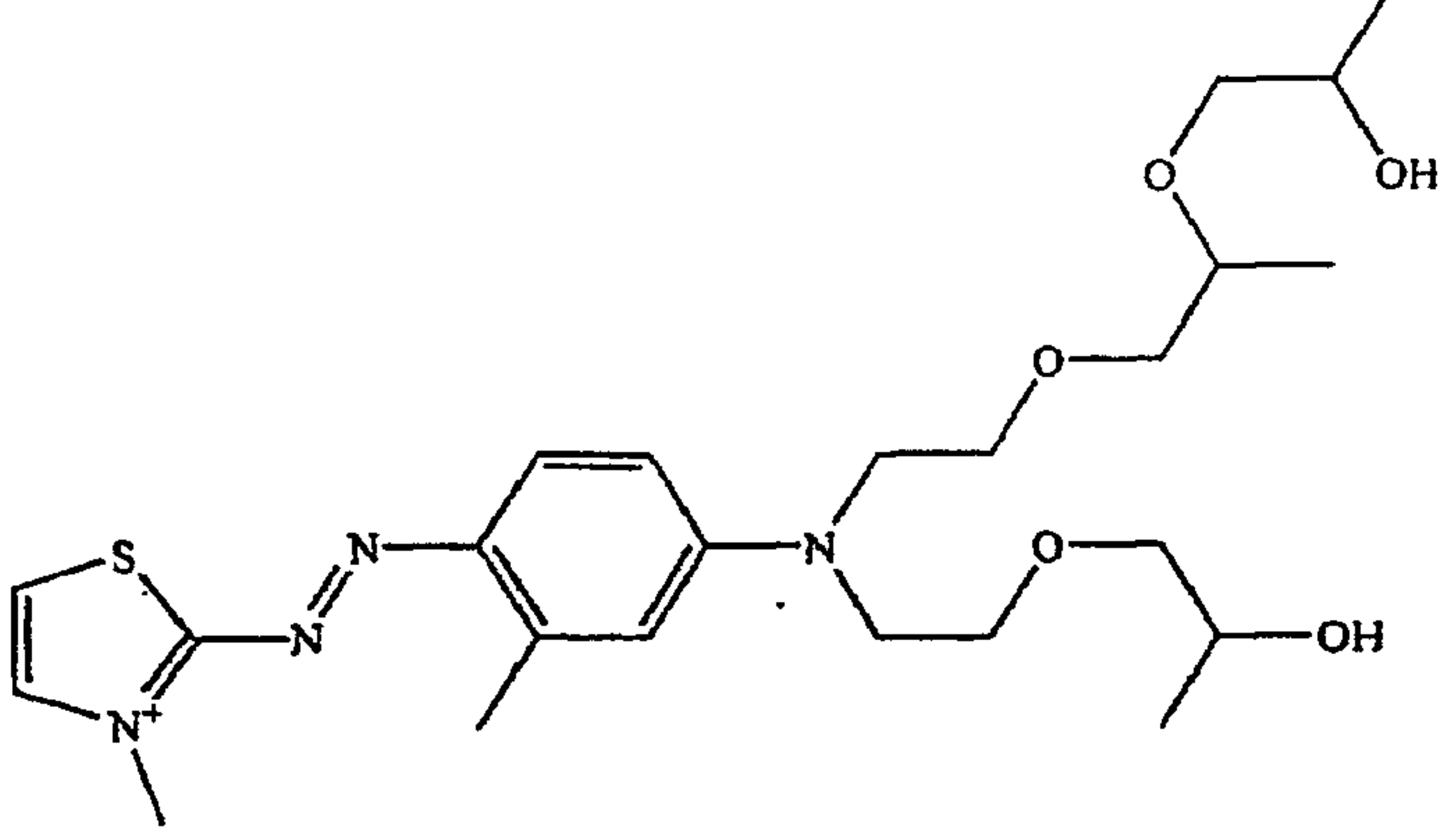
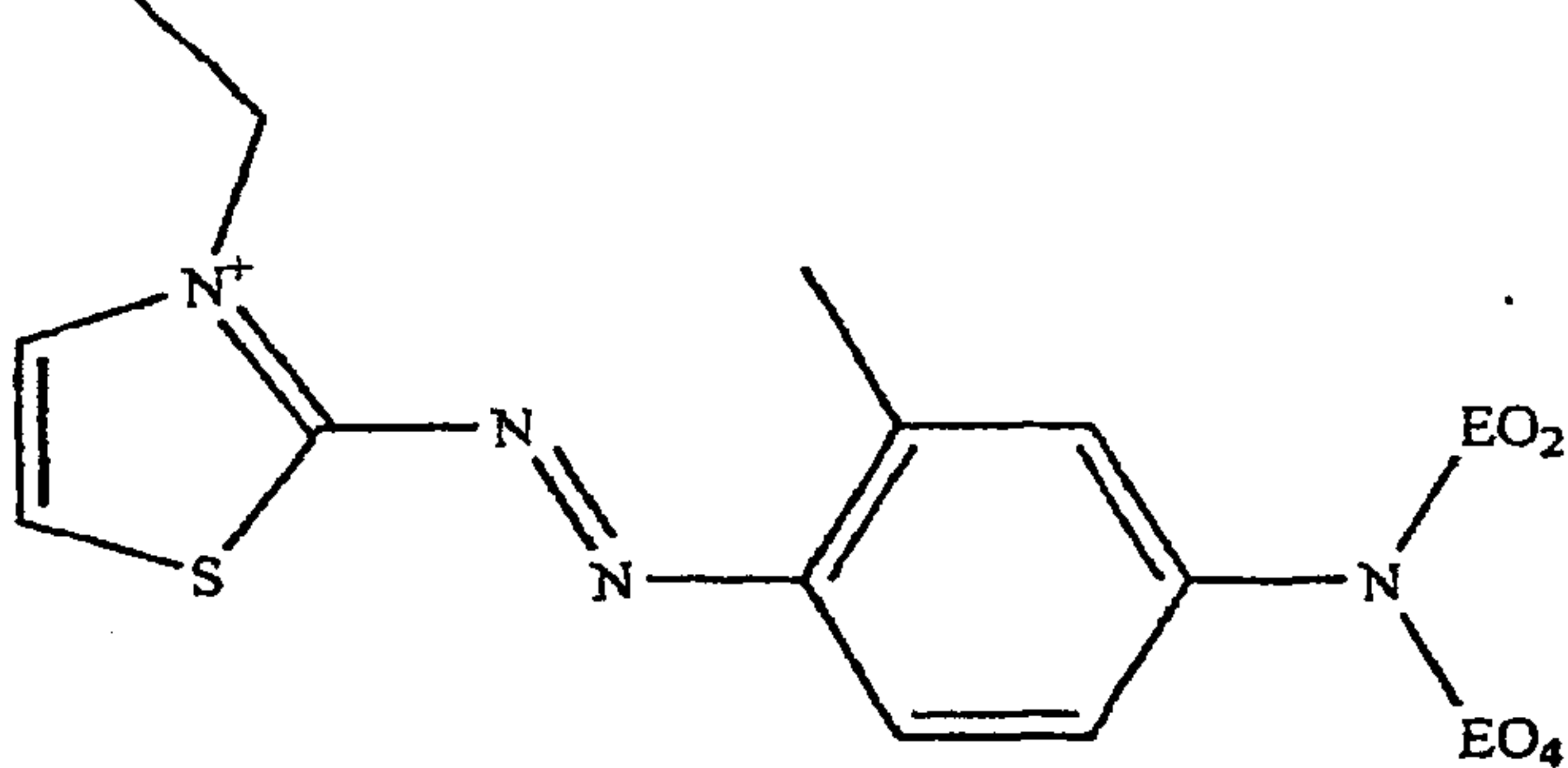
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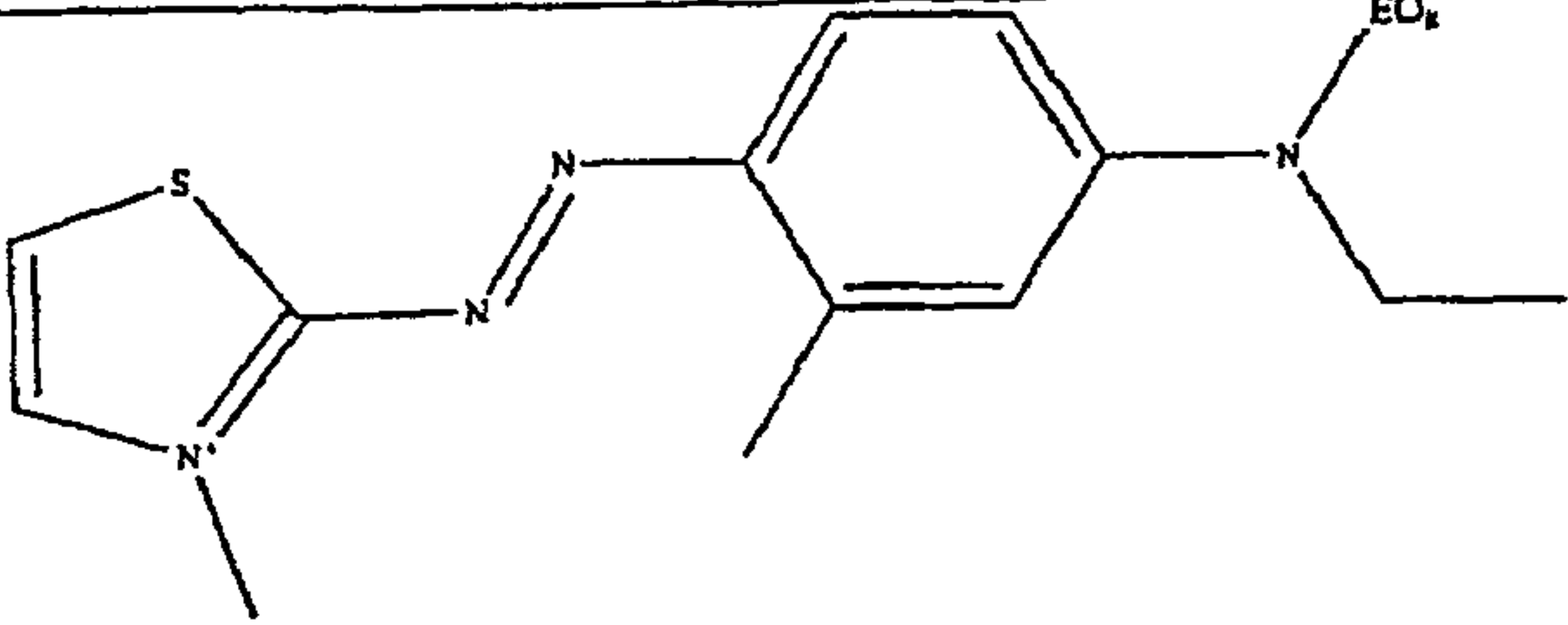
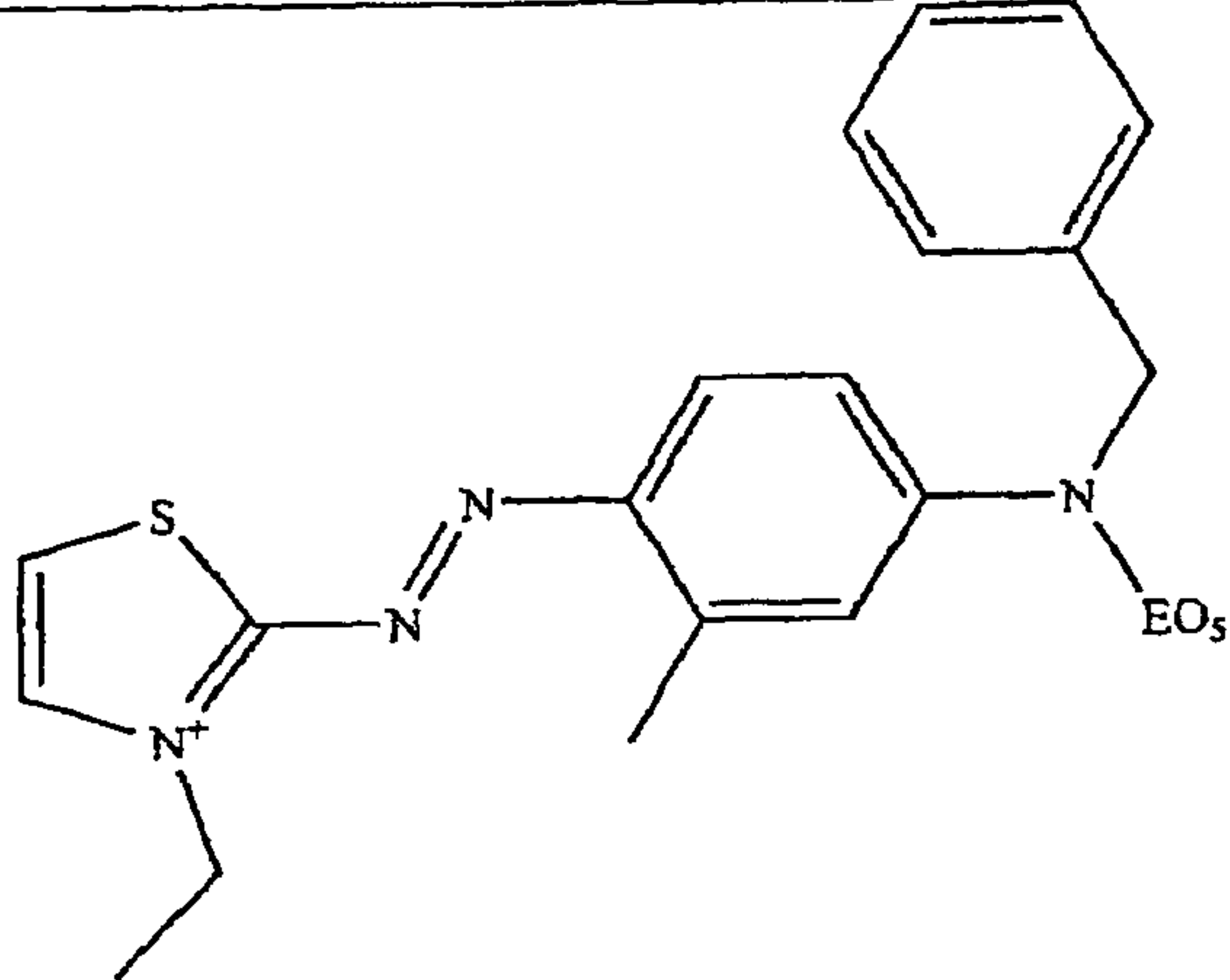
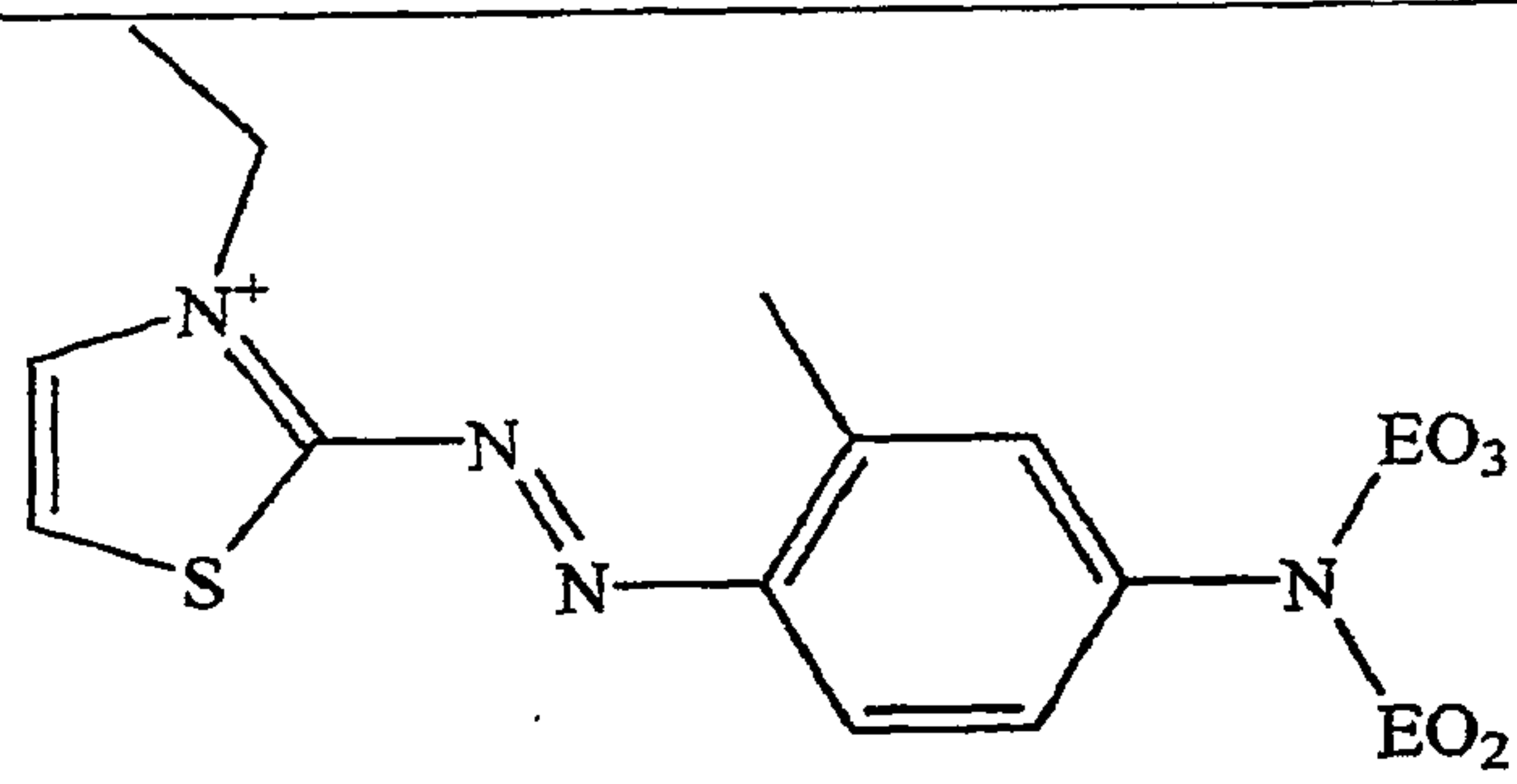
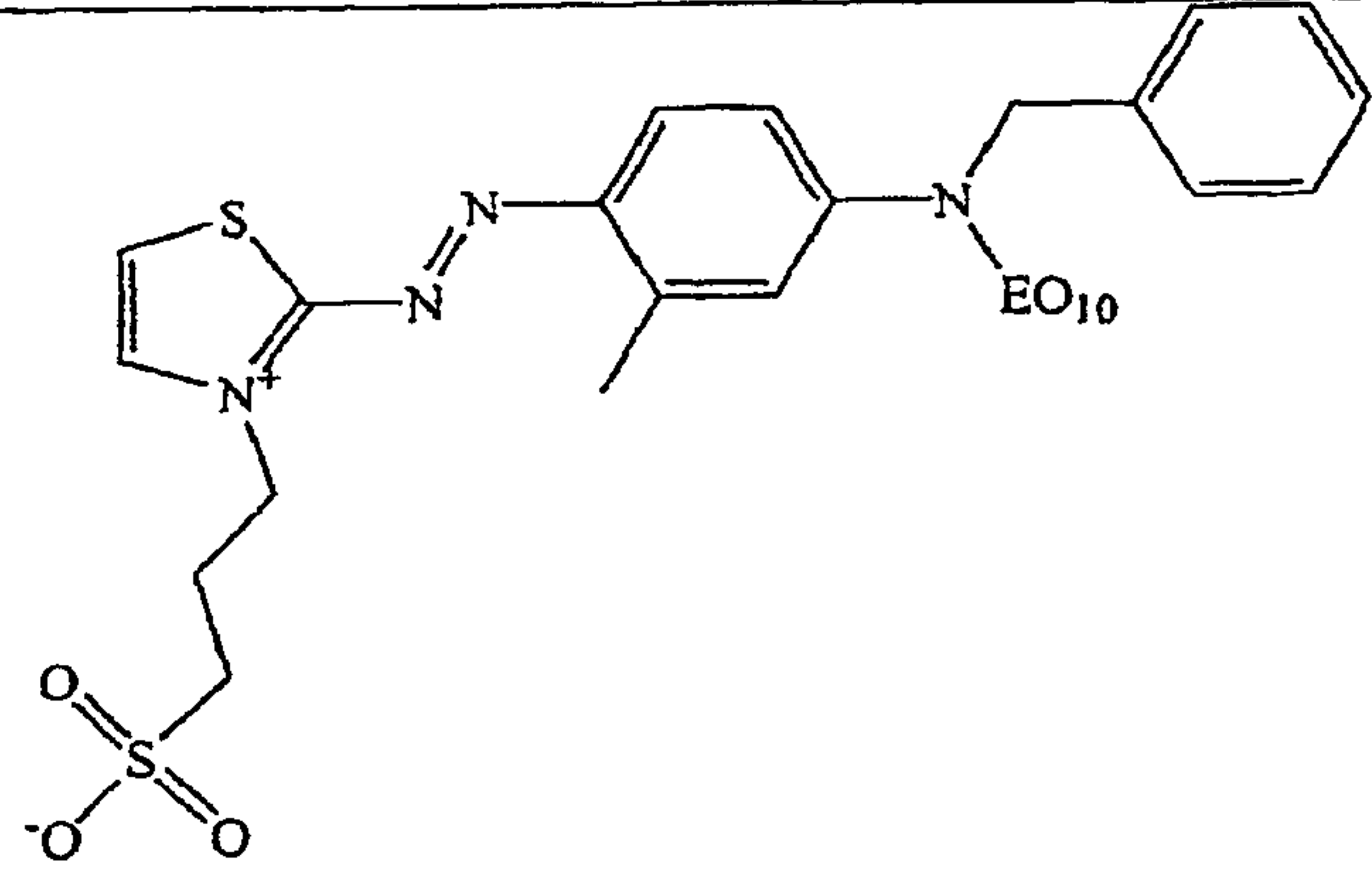
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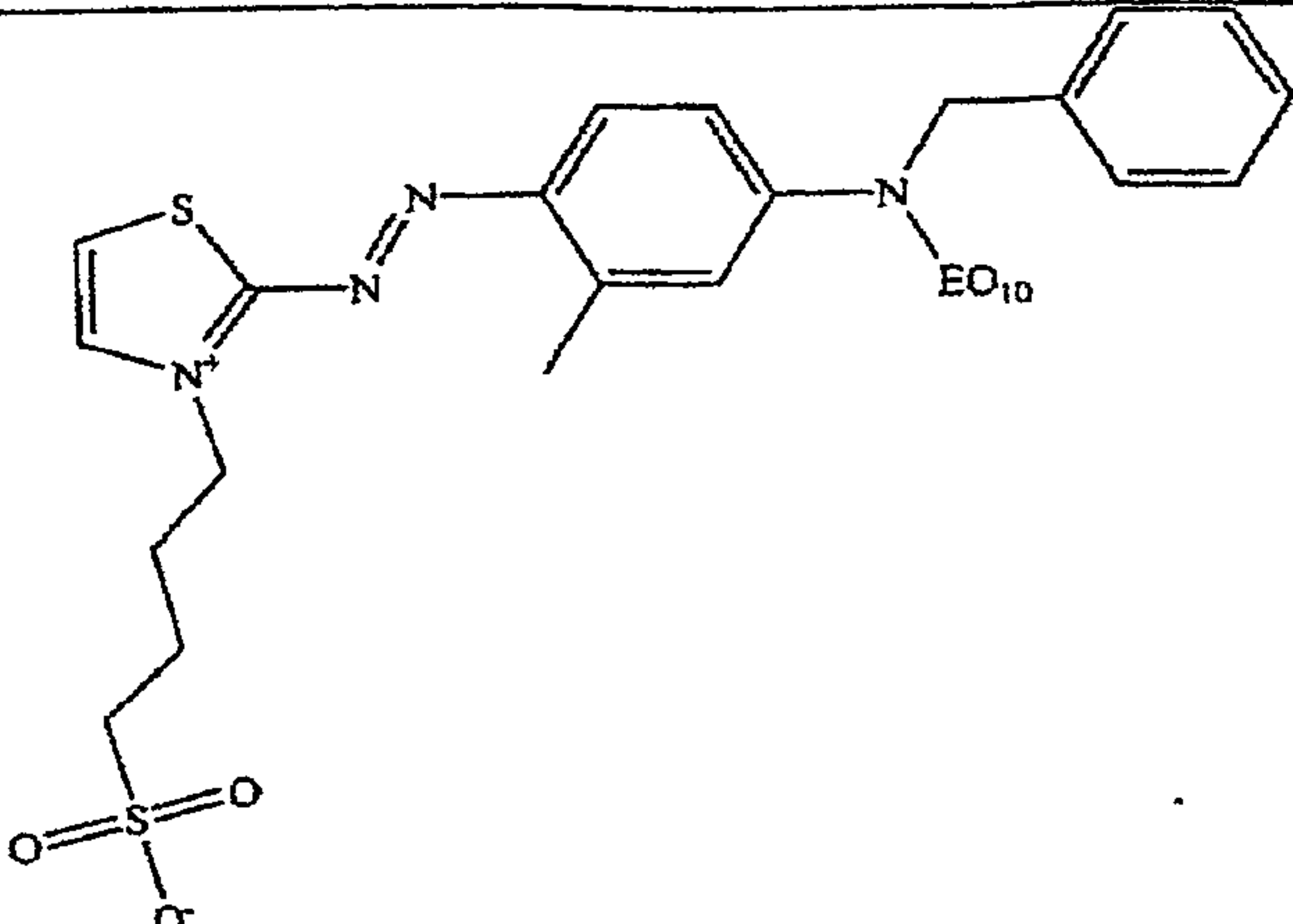
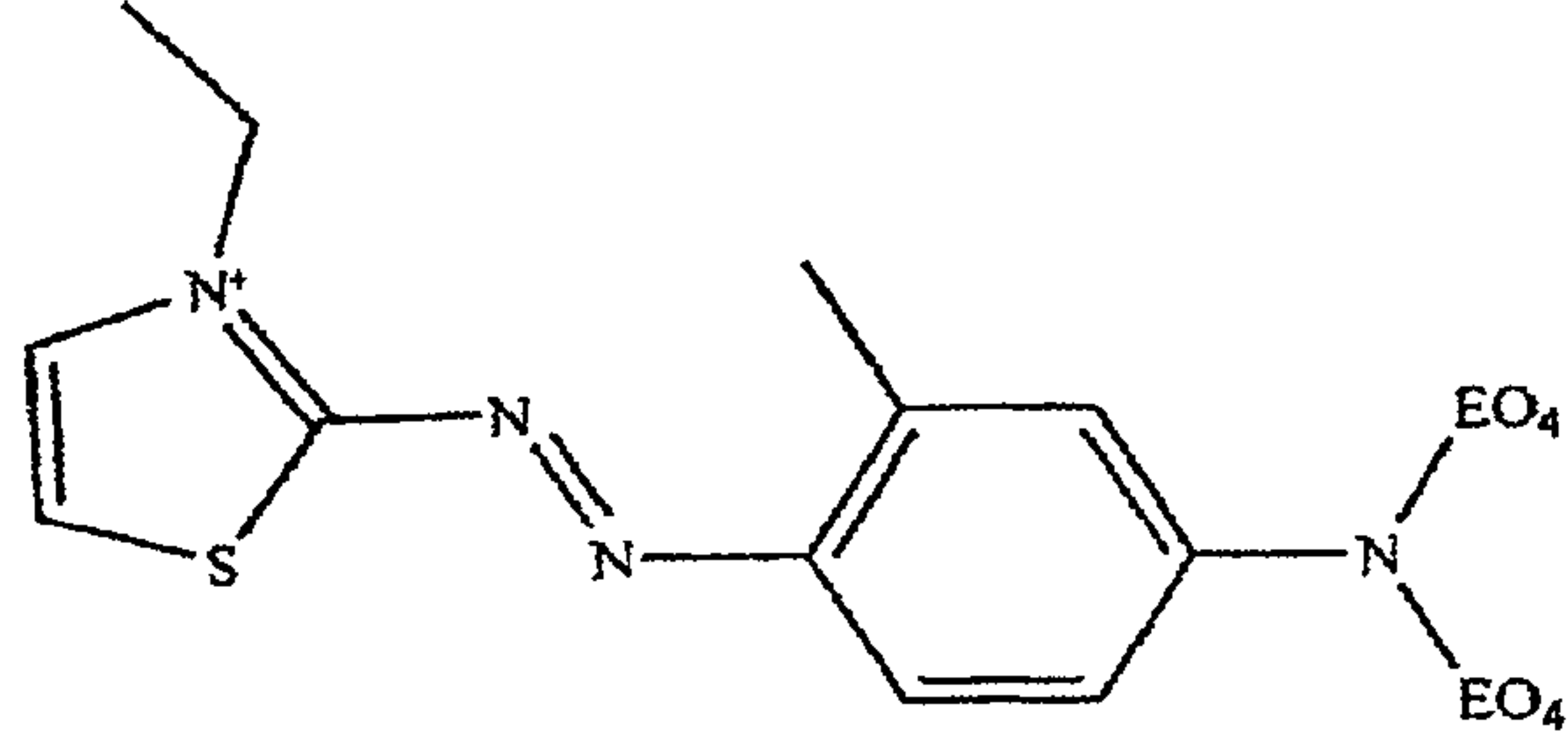
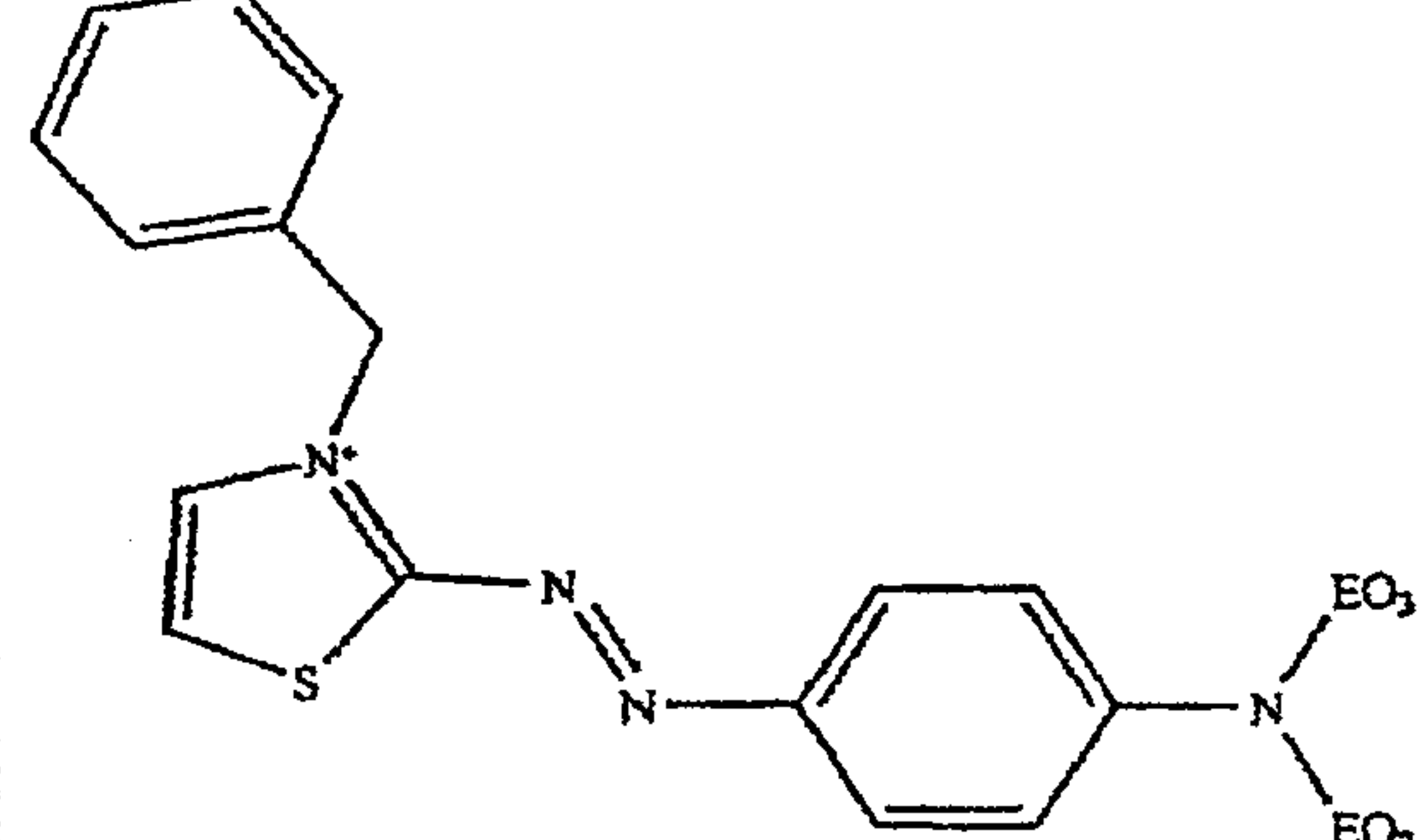
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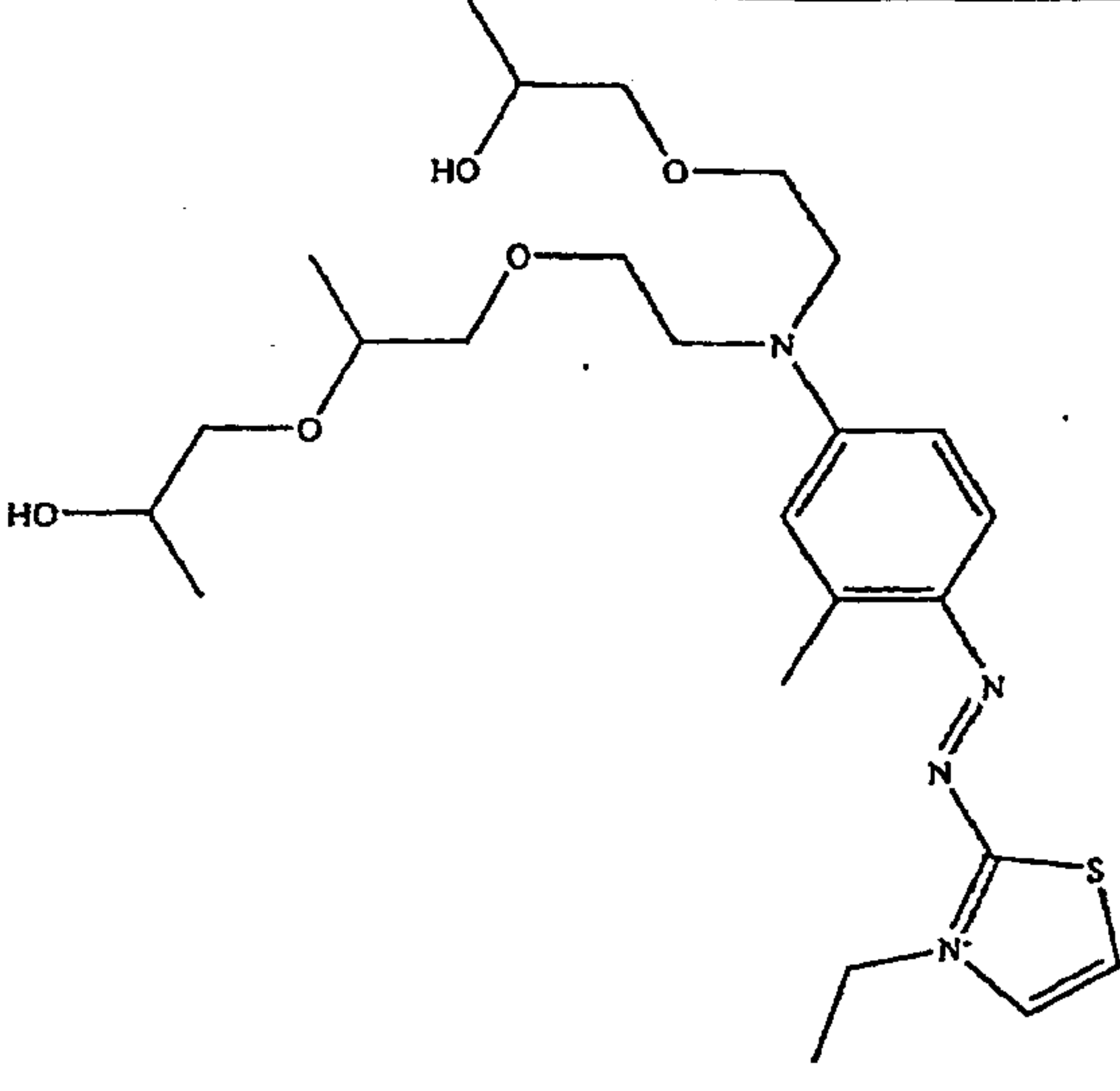
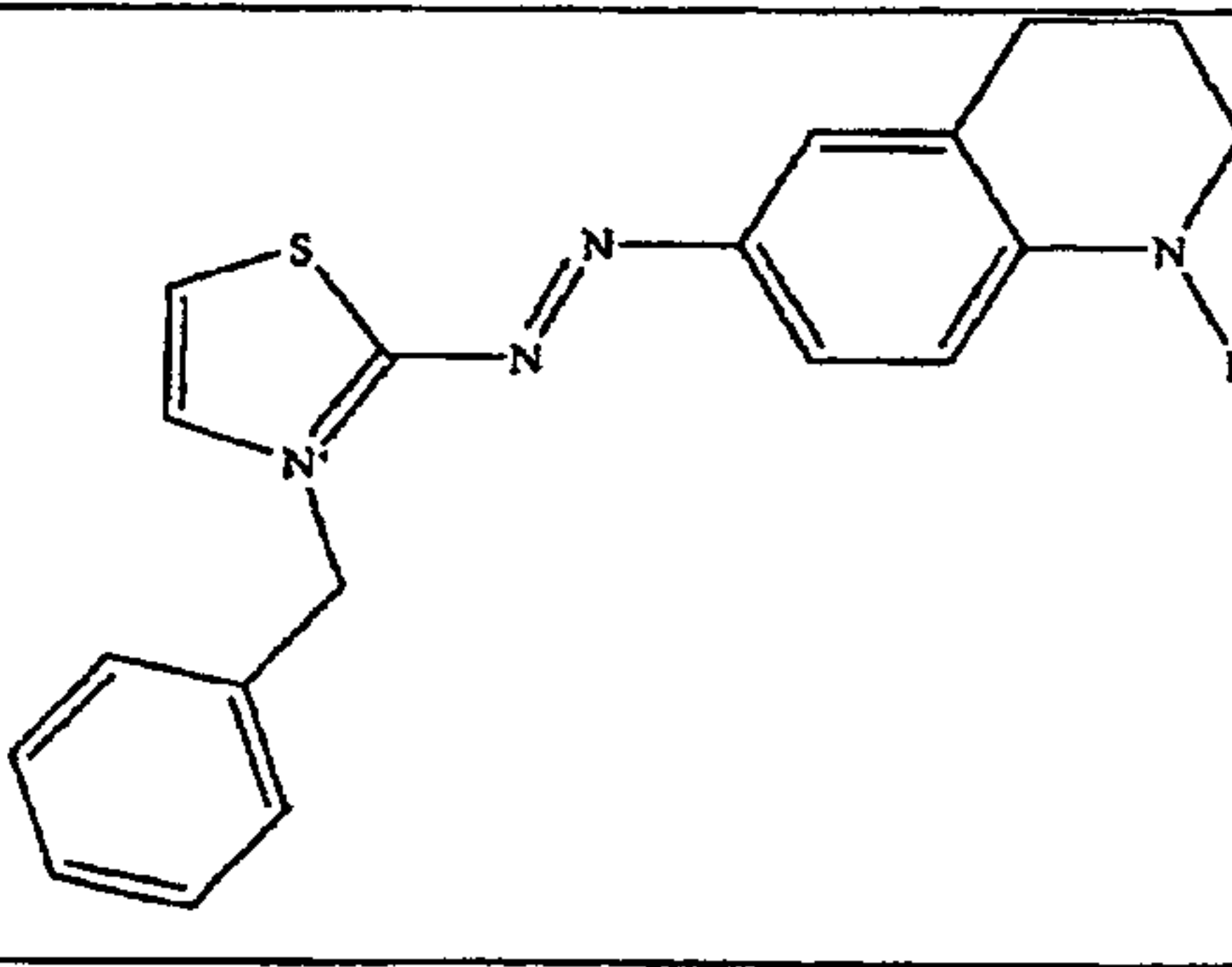
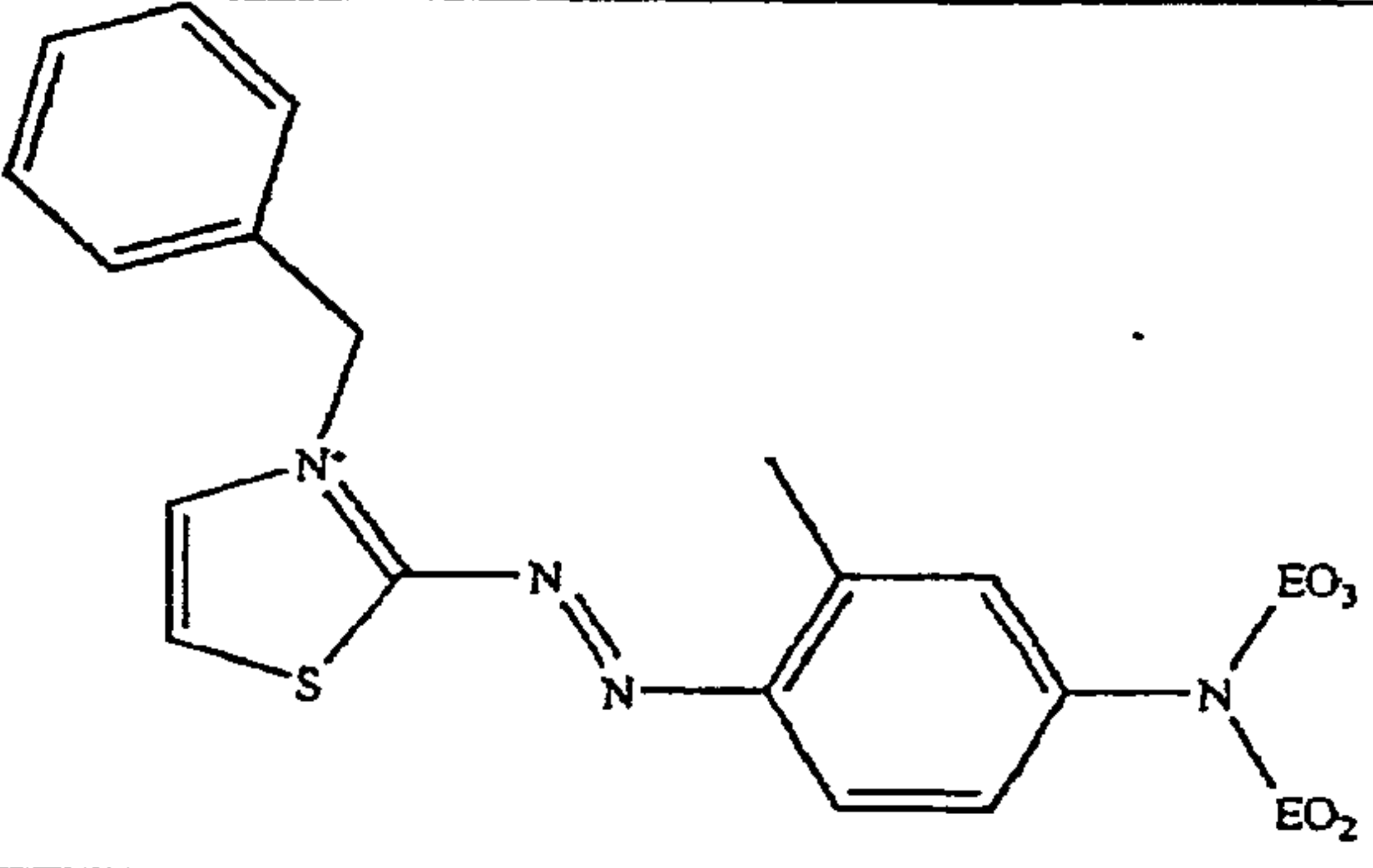
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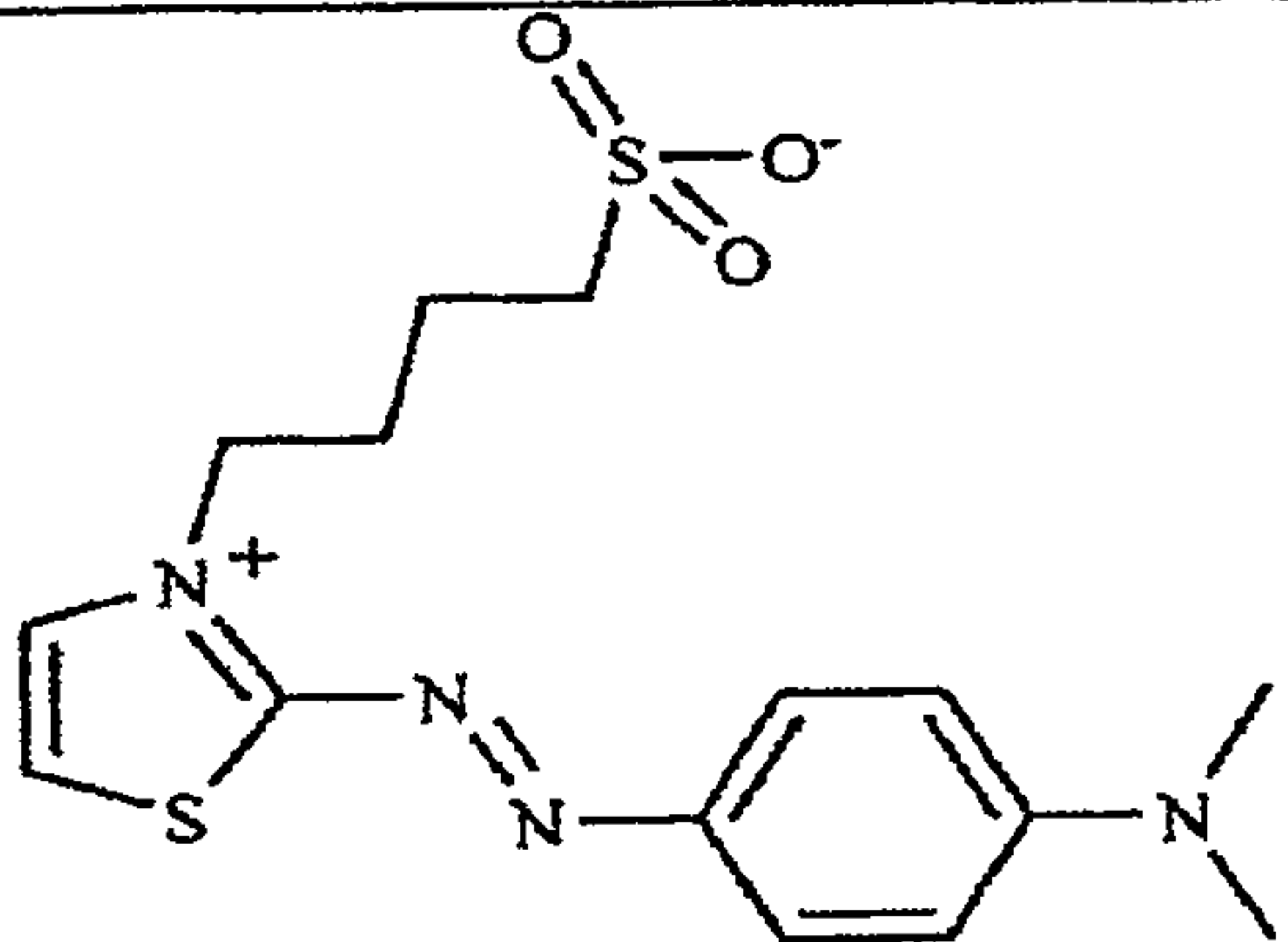
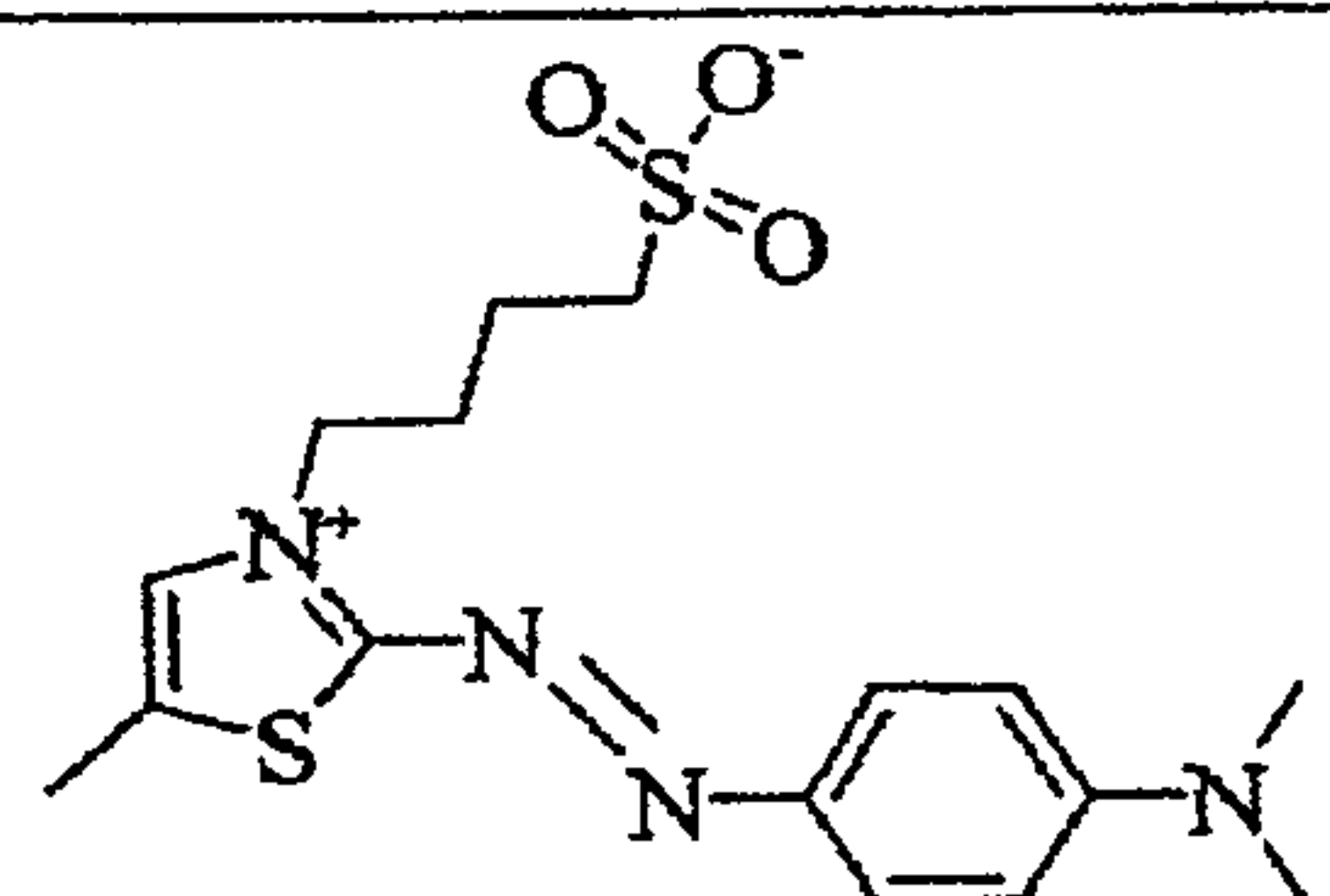
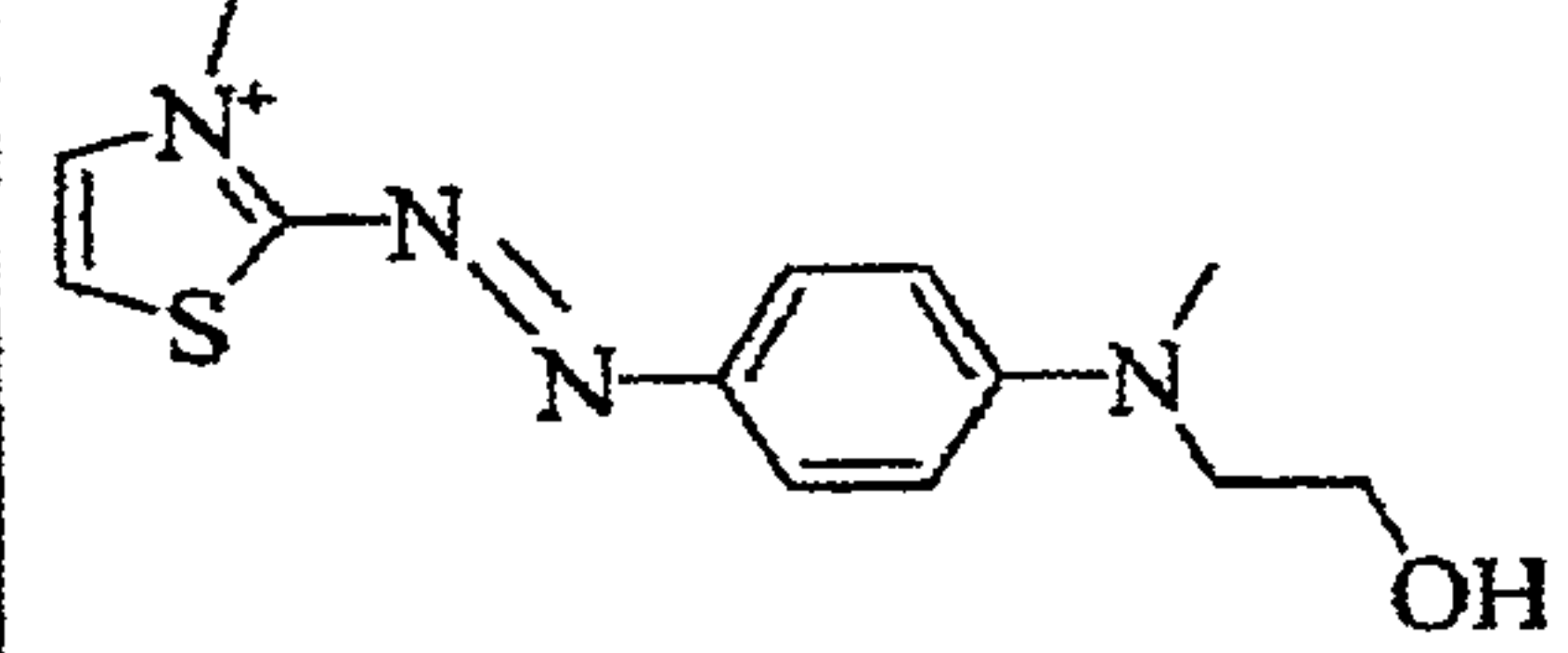
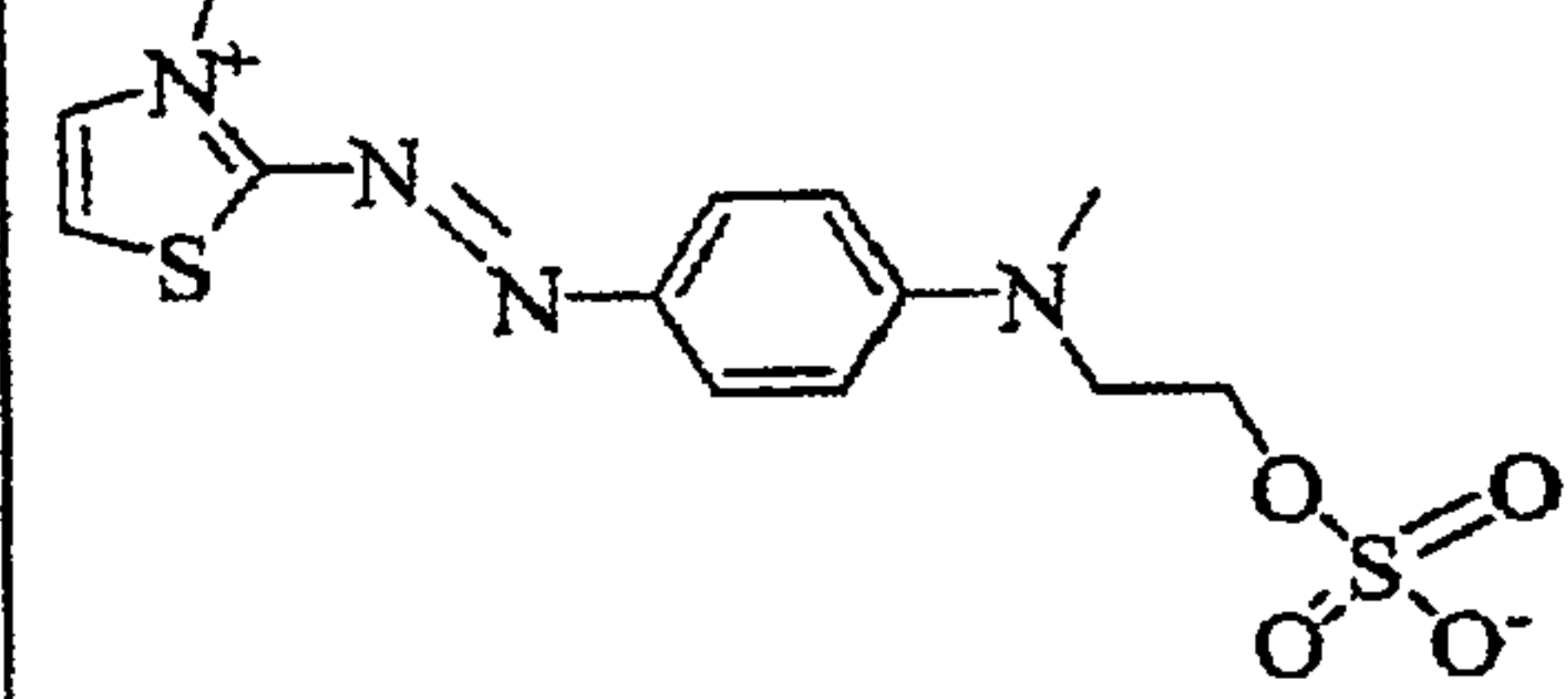
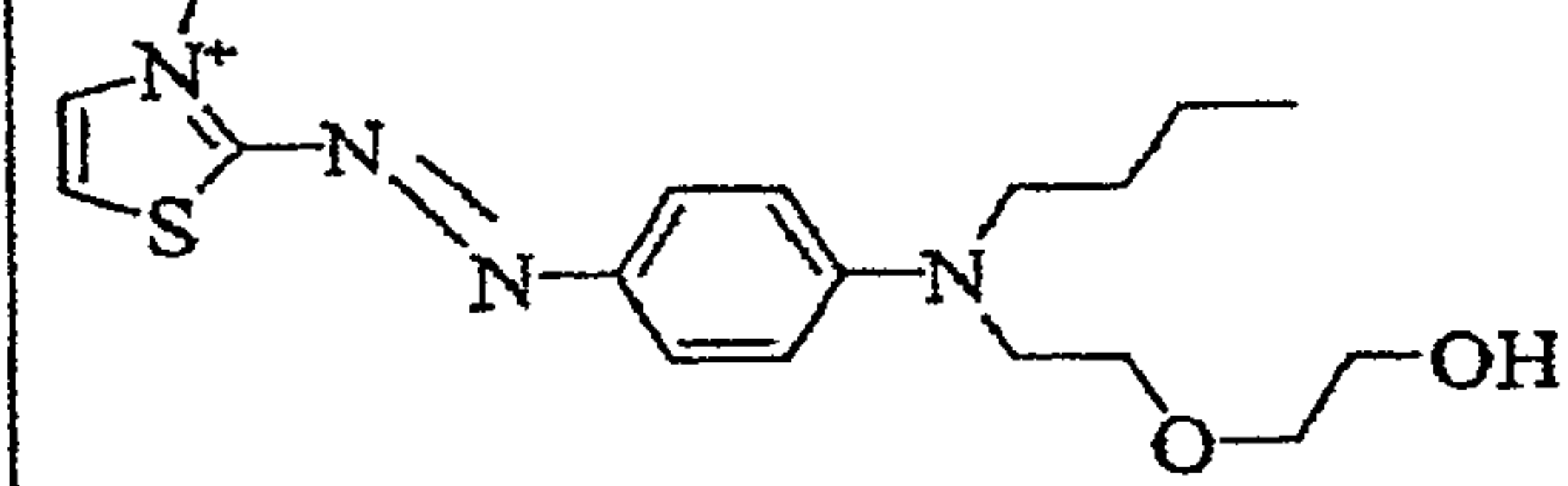
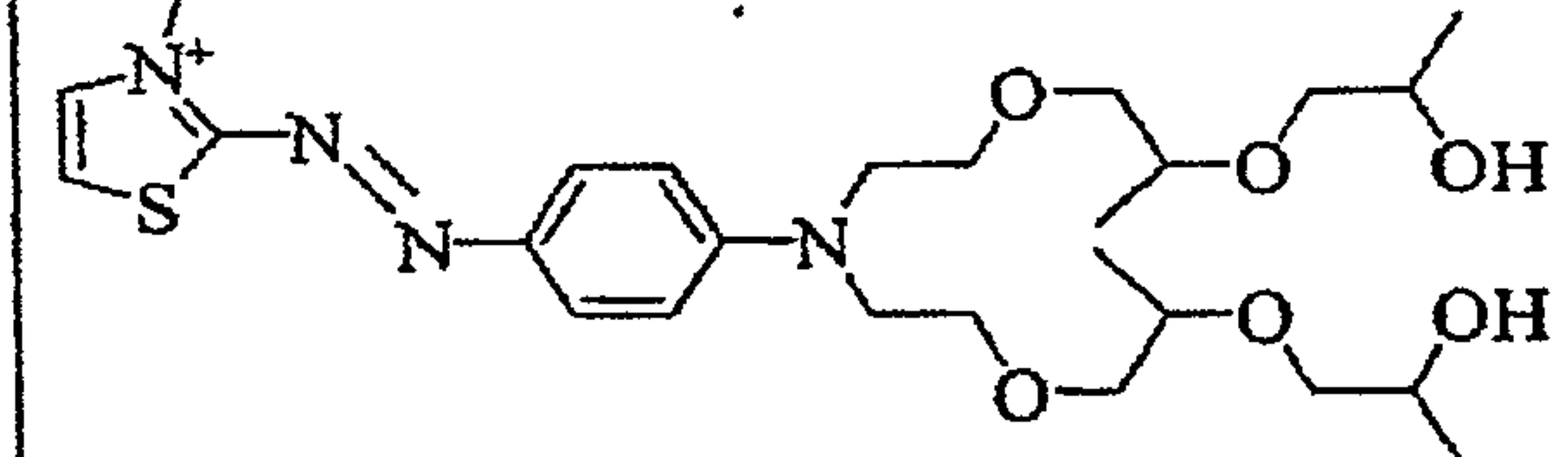
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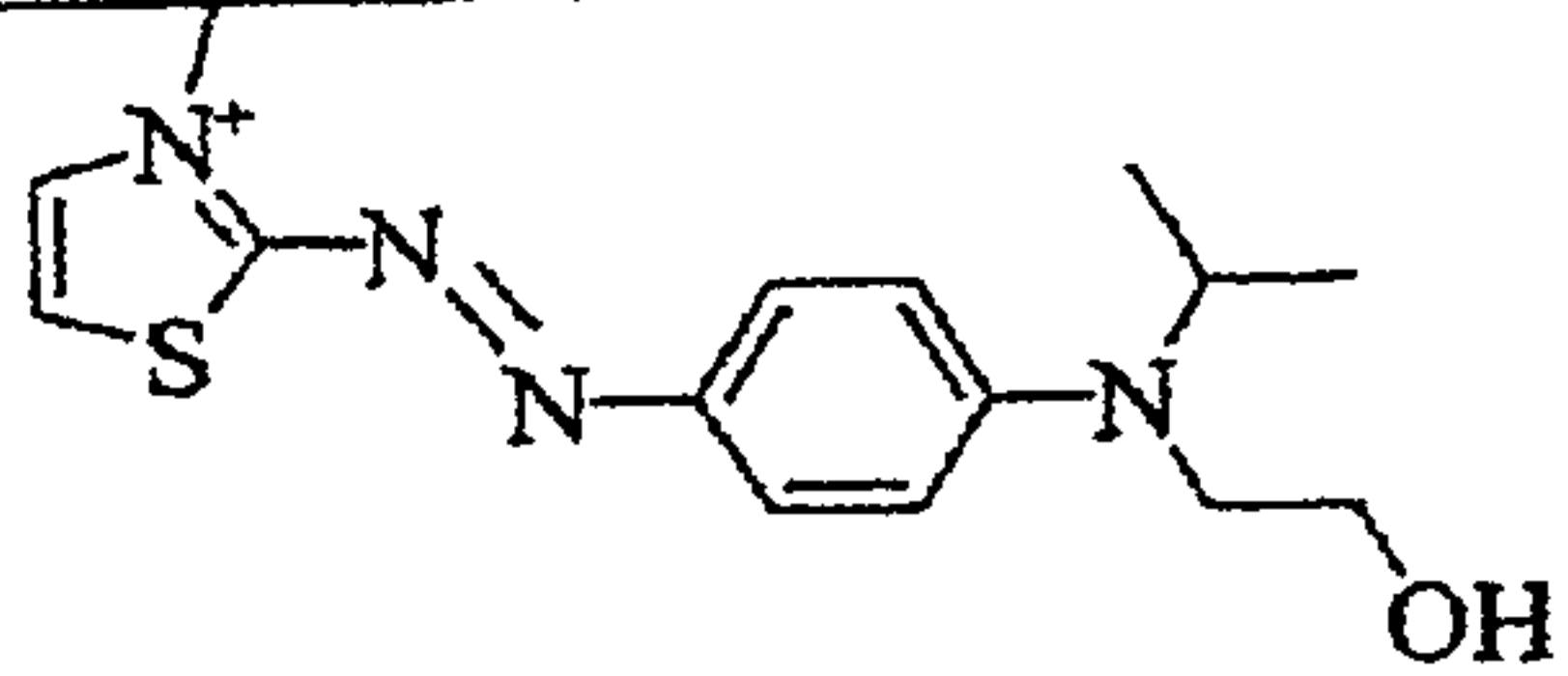
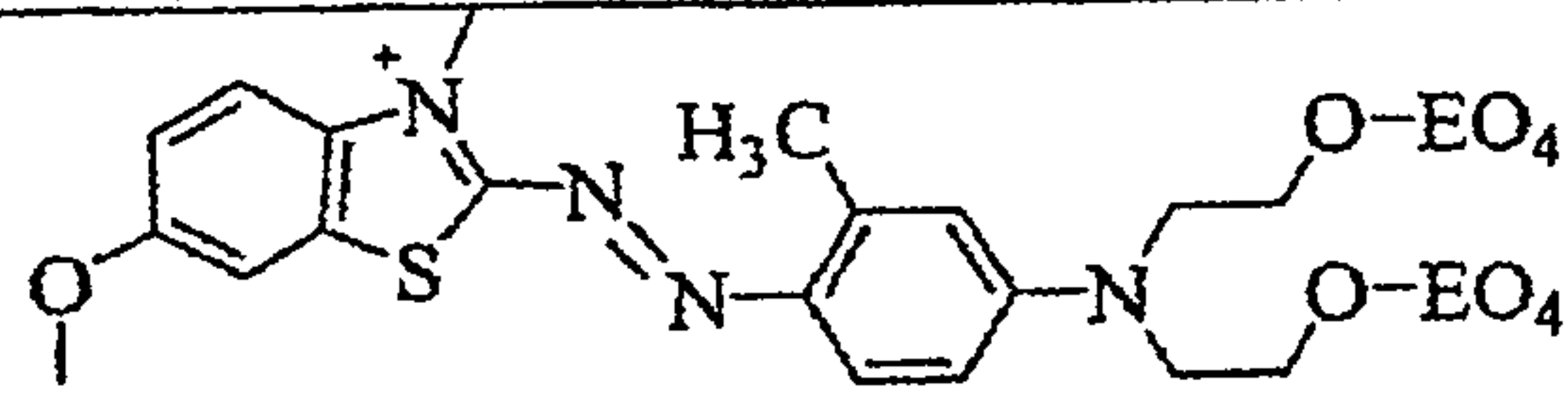
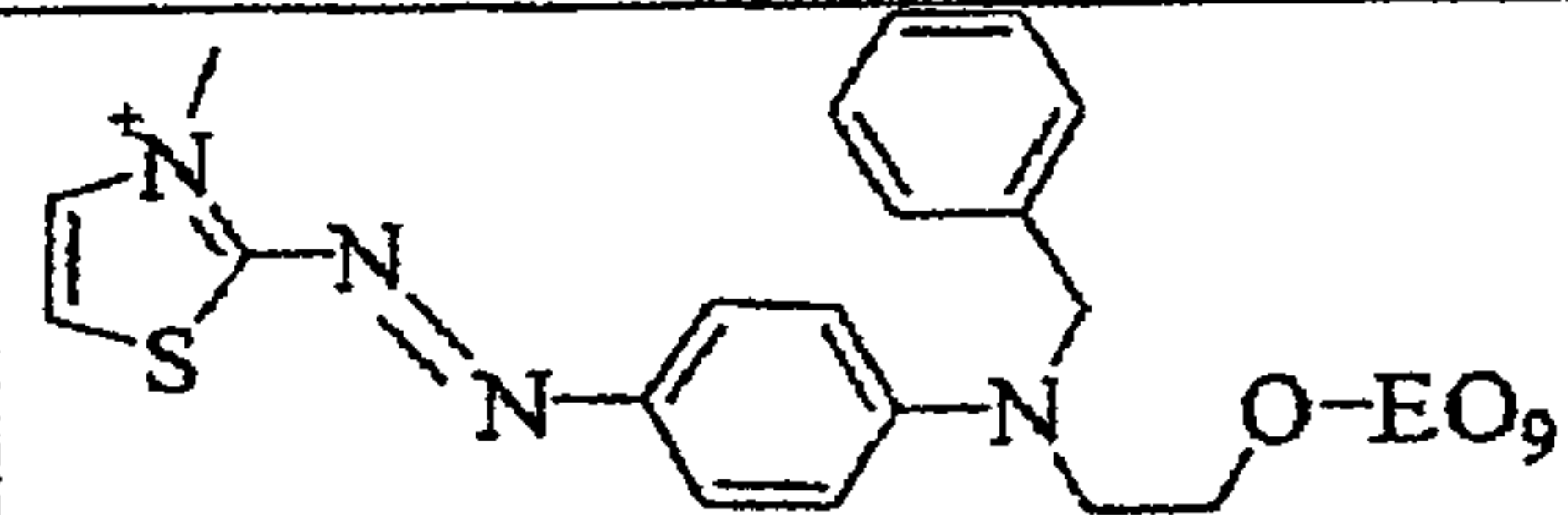
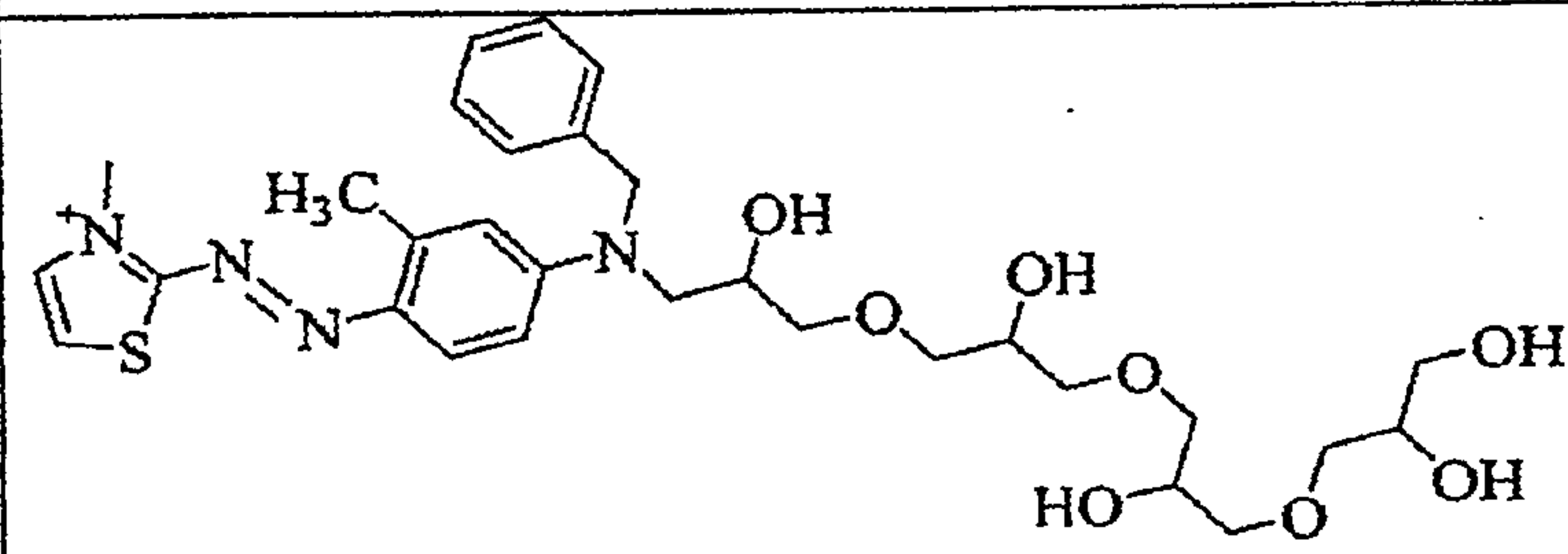
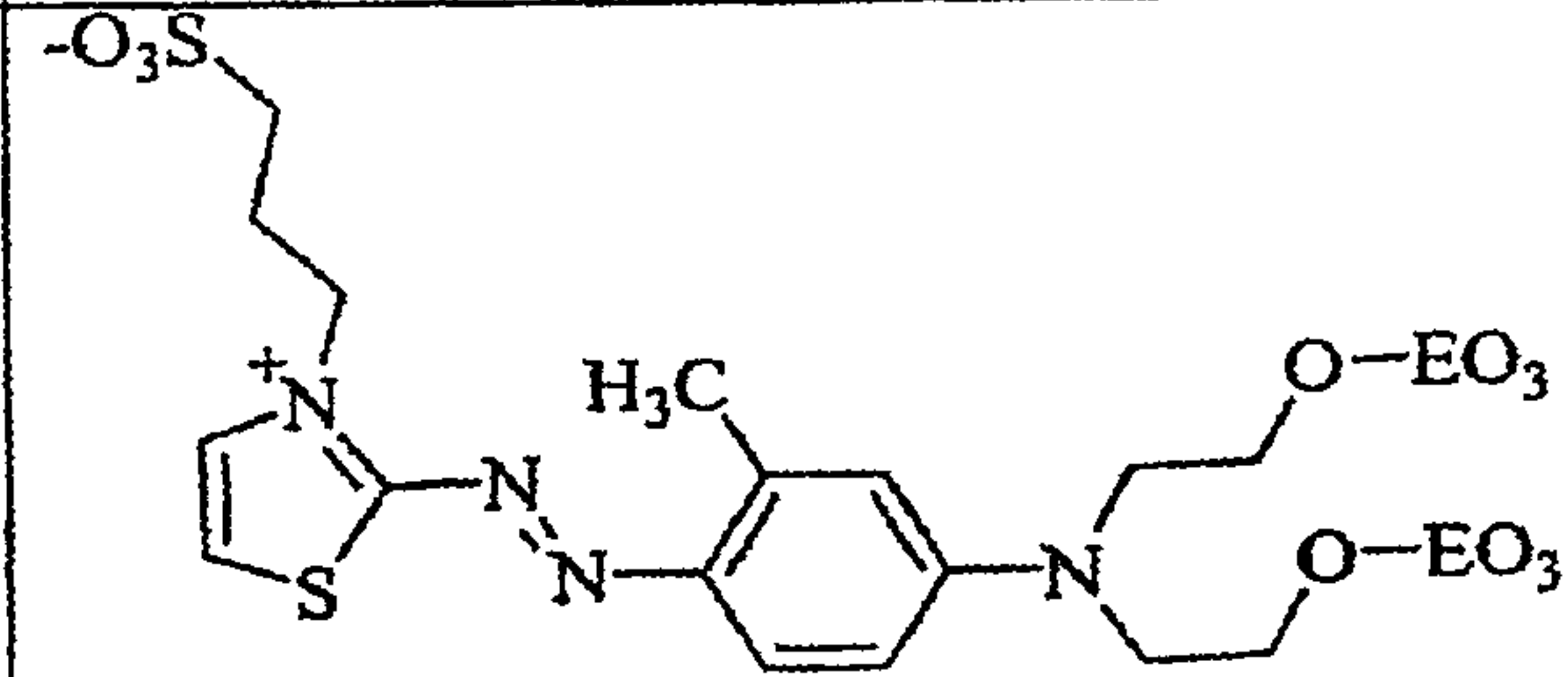
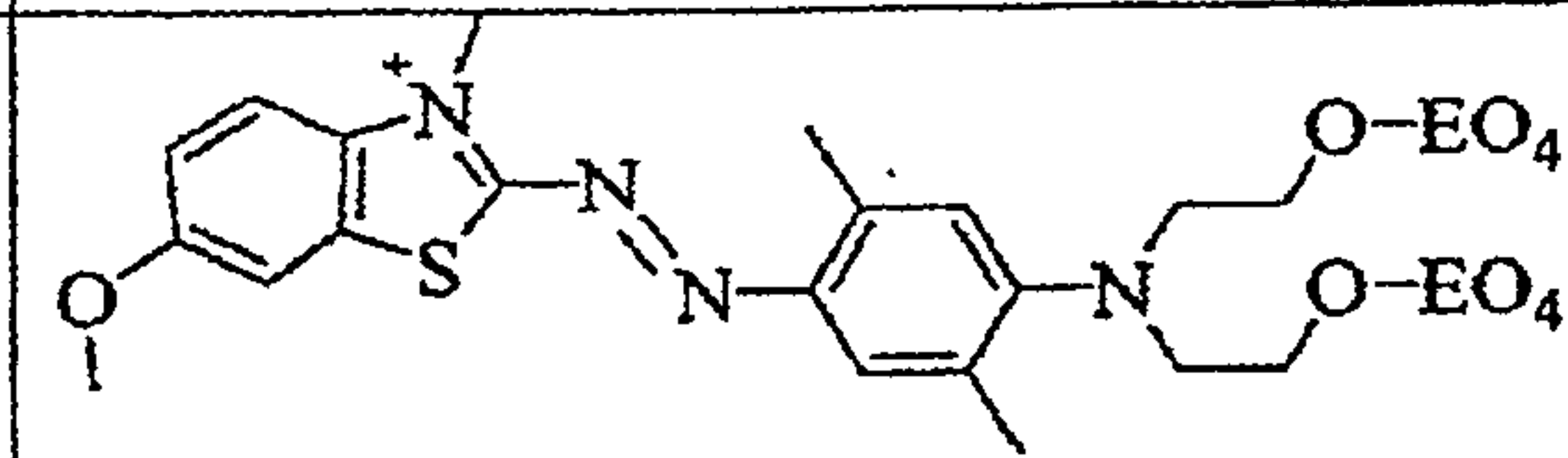
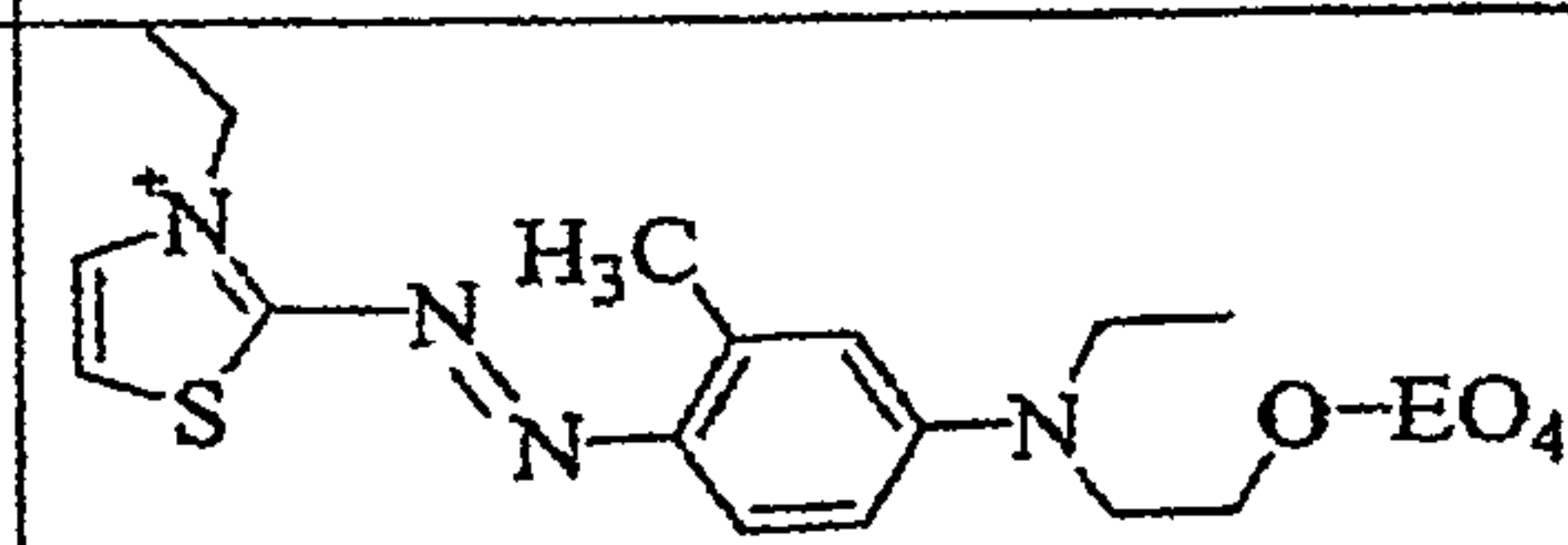
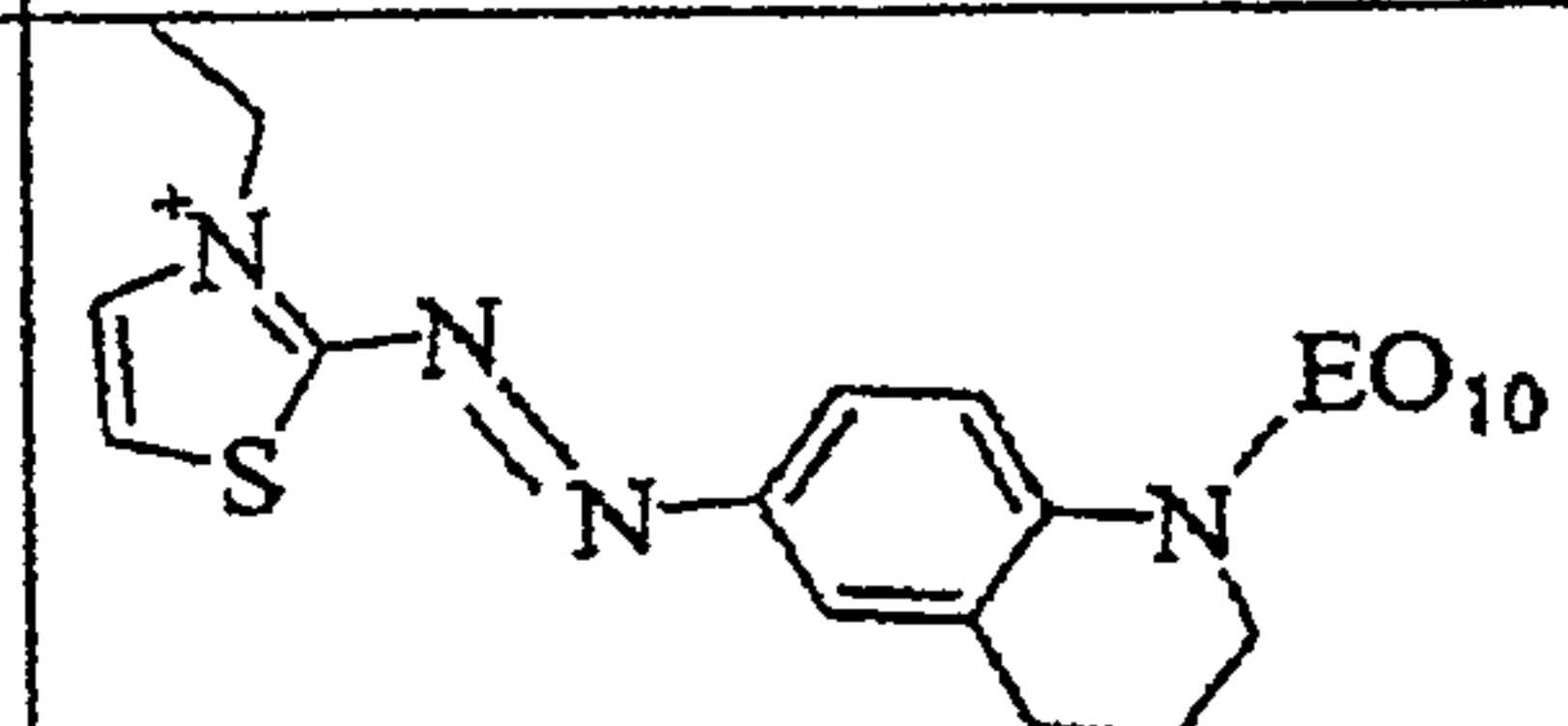
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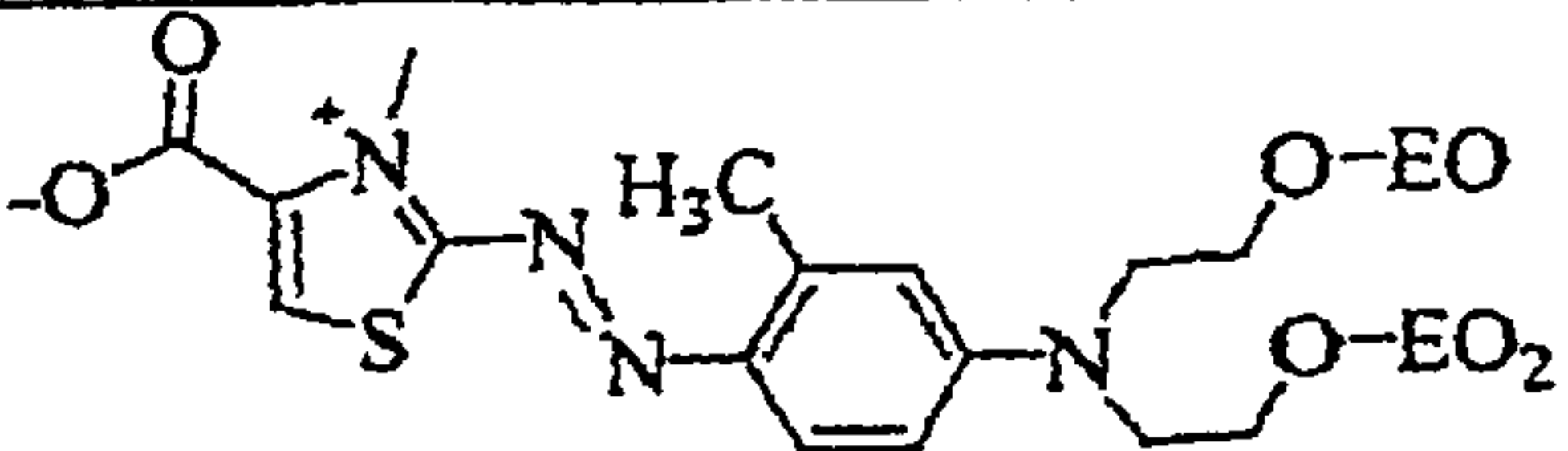
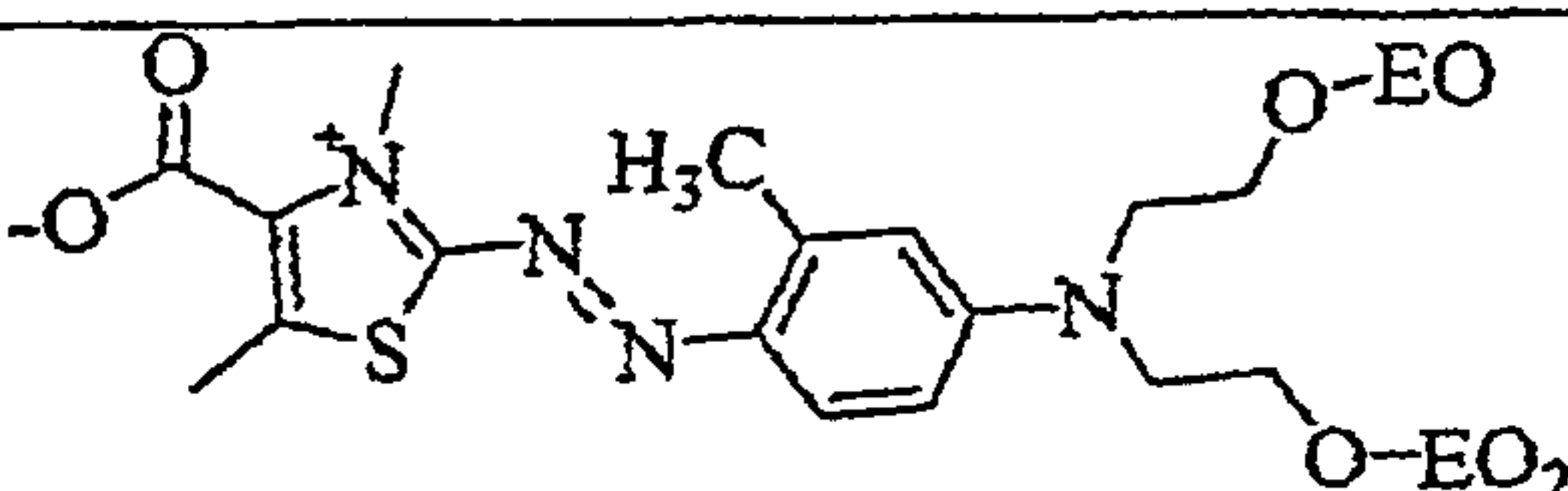
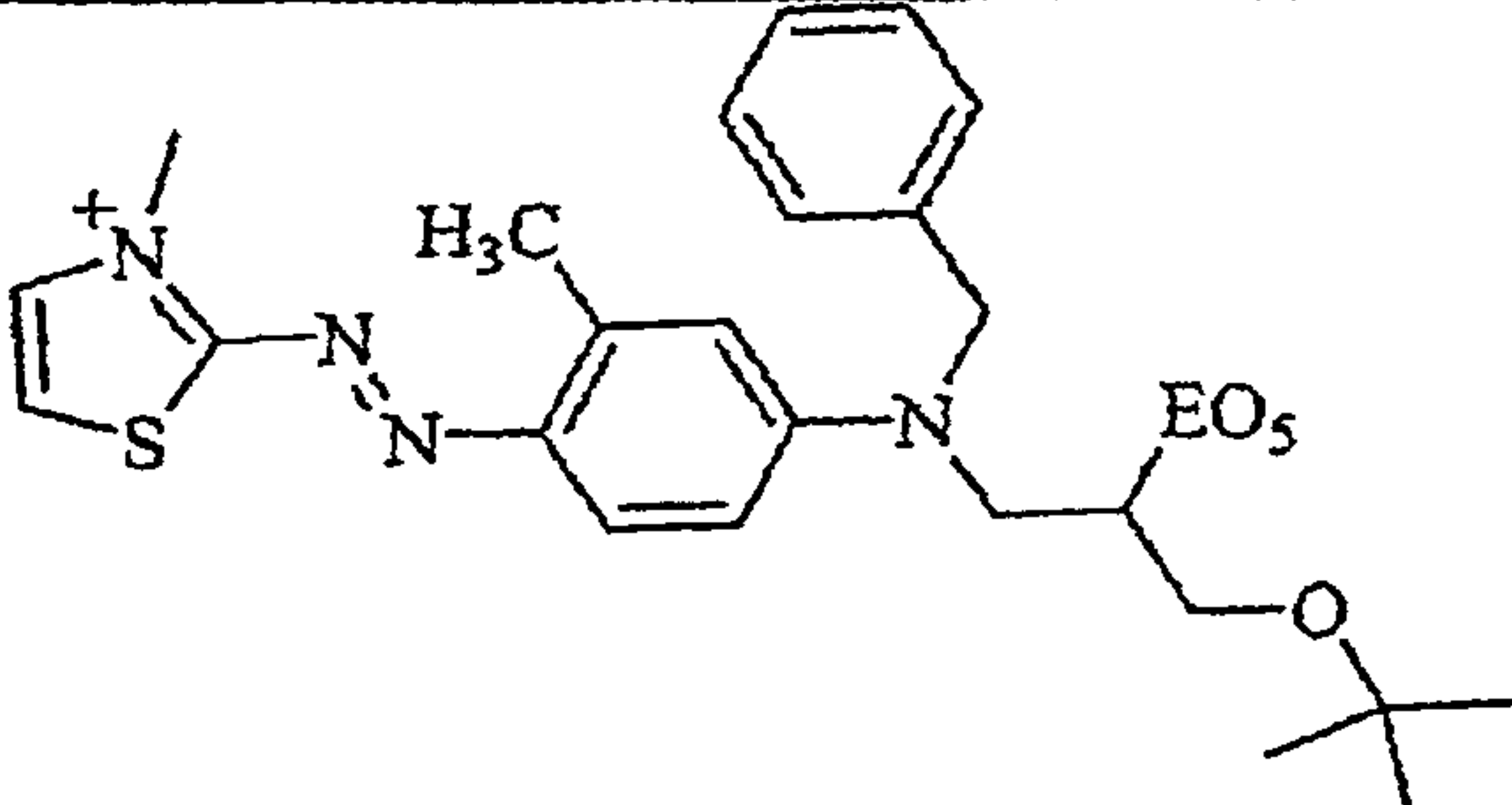
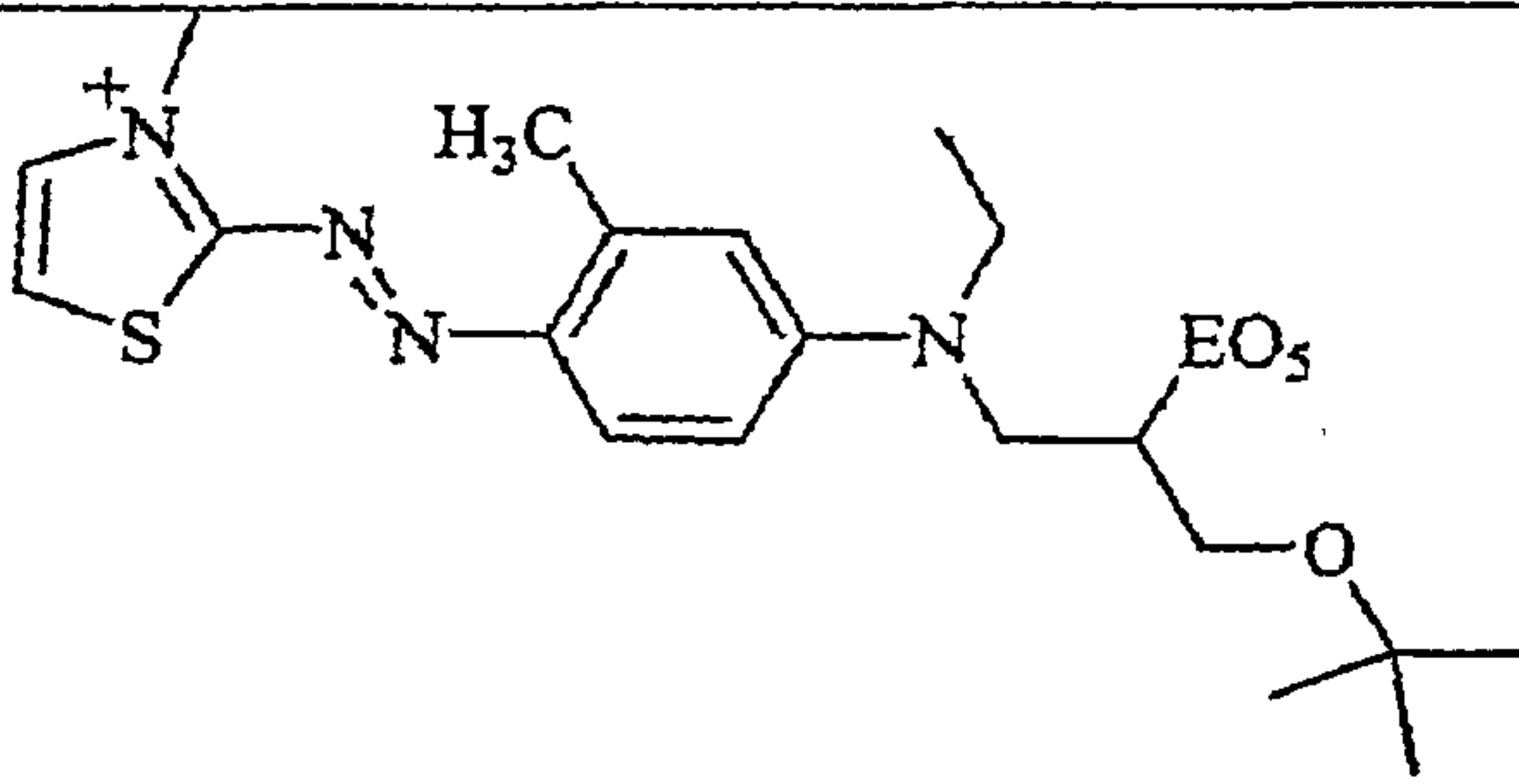
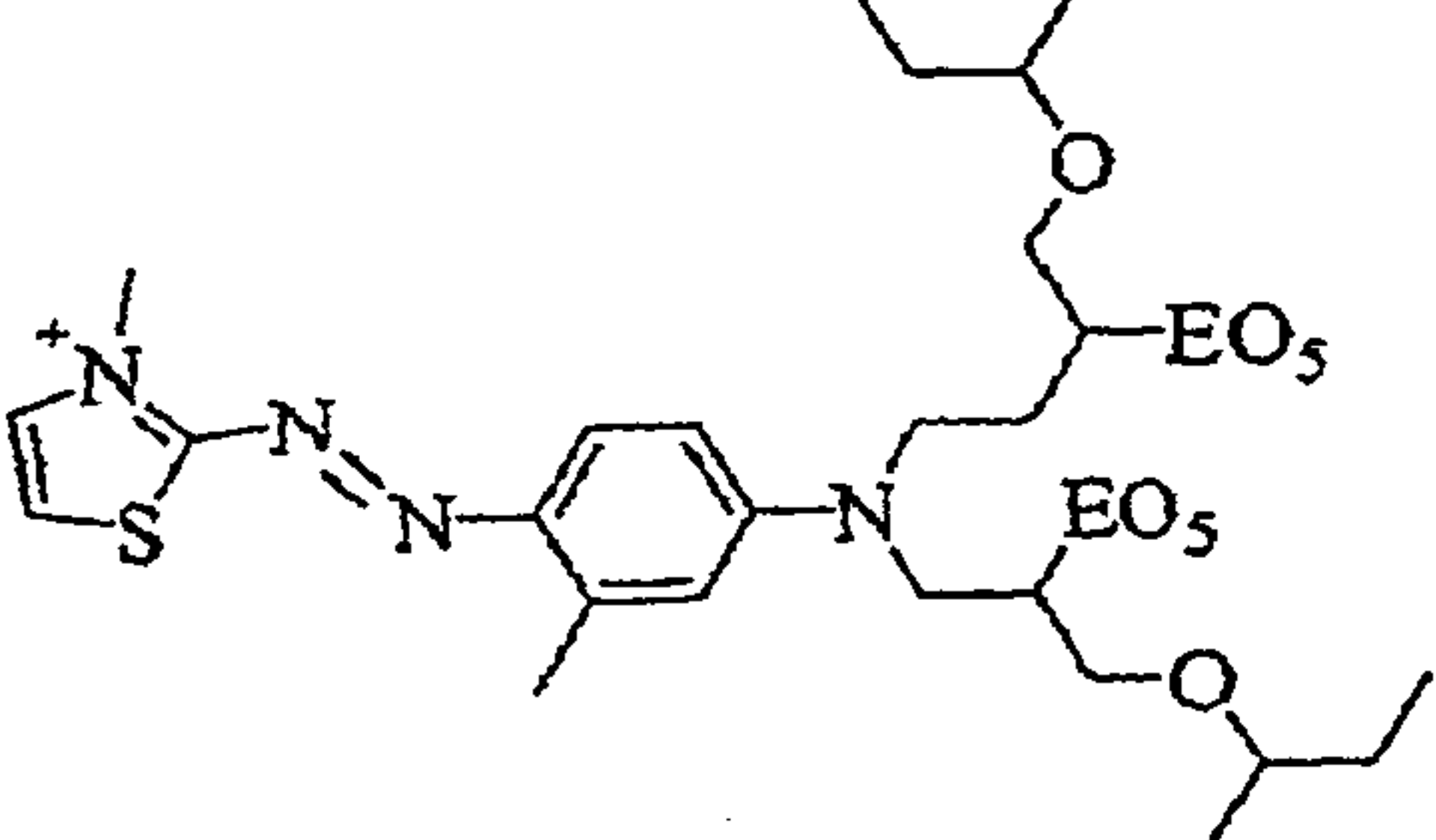
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5. The whitening agent of Claim 4, wherein the thiazolium dye is one of Table 1 Thiazolium Dyes 1, 4, 5, 7, 8, 12, 13, 15, 16, 17, 21, 24, 25, 26, 30, 31, 33, 36, 38, 40, 45, 48 or mixtures thereof.
6. The whitening agent of Claim 5, wherein the thiazolium dye is one of Table 1 Thiazolium Dyes 12, 13, 15, 16, 24, 25, 26, 30, 31, 33, 36, 38, 40, 45, 48 or mixtures thereof.