

SUBSTITUTED PYRIDINE-2-ONES AND SUBSTITUTED PYRIDINE-2-THIONES AS BIOCIDES

FIELD OF THE INVENTION

This invention relates to novel 1-hydroxy-6-substituted pyridine-2-ones and 2-thiones, a process for their preparation, and their use as biocides. These compounds exhibit good biocidal activity, particularly antifungal activity.

BACKGROUND OF THE INVENTION

Compounds exhibiting biocidal activity are well known in the art. For example, pyrithione salts, such as zinc pyrithione, are known to provide excellent biocidal activity, including broad spectrum anti-bacterial and anti-fungal activity. There are many uses for these pyrithiones. By way of illustration, U.S. Pat. No. 4,818,436 discloses the use of pyrithiones in metalworking fluids, U.S. Pat. No. 4,401,770 discloses urethane shoe inserts having antimicrobial activity; and U.S. Pat. No. 4,935,061 discloses their use in paints.

Despite the excellent biocidal (particularly fungicidal) activity attributable to pyrithione salts, these compounds do have drawbacks for certain applications, most notably limited solubility in certain organic solvents and aqueous media. Accordingly, new compounds exhibiting excellent biocidal activity, but also exhibiting good solubility in organic solvents would be useful to the biocides manufacturing community.

One compound that exhibits good biocidal efficacy and solubility in shampoo formulations is 1-hydroxy-4-methyl-6-(2,4,4-trimethylpentyl)-2-pyridone. Unfortunately, this compound is more expensive to manufacture than otherwise might be desired. This compound and related compounds are disclosed in U.S. Pat. No. 3,883,545; their use in combating dandruff is described in U.S. Pat. No. 4,185,106; and their use in an antidandruff shampoo is described in U.S. Pat. No. 4,711,775. U.S. Pat. No. 4,916,228 discloses the preparation of a broad class of 1-hydroxy-2-pyridones by a process comprising reacting a pyrone having a specified structural formula with a hydroxylammonium salt in the presence of at least one alkali metal carbonate and/or hydrogen carbonate.

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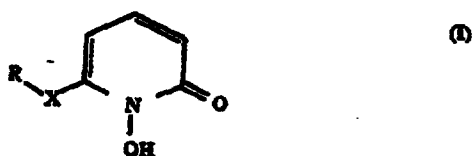
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A technical journal publication entitled "Quantitative Structure-Activity Analysis in a Series of Antimycotically Active N-Hydroxypyridones", Journal of Medicinal Chemistry, 1974, Vol. 17, No. 7, p. 753, describes a variety of 1-hydroxy-substituted-2-pyridones, including those having para-substitutions of $-\text{SCH}_2\text{C}_6\text{H}_4\text{Cl}$ (compound 15 in Table II thereof) and $-\text{OCH}_2\text{C}_6\text{H}_4\text{Cl}$ (compound 16 in Table II thereof), as having antimycotic activity. However, the method of preparation of these compounds is not described in this technical publication, and it is believed that these compounds were made from 6-chloro-2-pyrone which is not commercially available.

Other new biocides providing desired solubility characteristics, as well as new processes for their preparation, would be highly desired by the biocides manufacturing community.

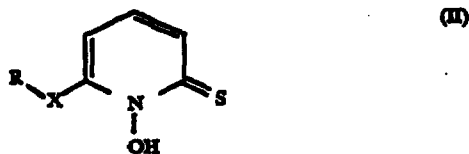
SUMMARY OF THE INVENTION

In one aspect, the present invention relates to a compound of the formula:



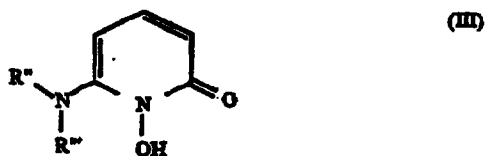
wherein X is an oxygen or sulfur moiety and R is a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms, with the proviso that R is other than chlorobenzyl.

In another aspect, the present invention relates to a compound of the formula:



wherein Y is an oxygen, sulfur, or NR' moiety wherein N is nitrogen and R and R' are independently each a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms.

In yet another aspect, the present invention relates to a compound of the formula:



wherein R'' and R''' are independently either hydrogen or a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms, with the proviso that either R or R' is a hydrocarbon radical.

Preferred hydrocarbon radicals for the R group in the above formulae (I), (II), and (III) are aliphatic hydrocarbons having between 3 and 15 carbons, more preferably a straight chain hydrocarbon having between 5 and 10 carbons. The term "substituted hydrocarbon" is intended to include substituents such as halogen such as chloro, iodo fluoro or bromo, alkoxy such as methoxy, ethoxy, propoxy or butoxy, nitro, thio, amino combinations thereof, and the like. Illustrative hydrocarbon groups include n-octyl, 2,4,4-trimethylpentyl, 3,5,5-trimethylhexyl, combinations thereof, and the like.

In still another aspect, the present invention relates to a process for producing compound of Formulae (I) or (II) wherein X or Y is oxygen which comprises the steps of:

- (a) reacting 2,6-dichloropyridine N-oxide, a hydroxy compound containing between 1 and 20 carbon atoms (such as an alcohol such as n-octanol or a phenol compound such as 4-methylphenol, preferably selected from the group consisting of alkyl, arylalkyl and aryl compounds containing at least one hydroxyl group, and combinations thereof) and a base (such as sodium hydroxide or potassium hydroxide), optionally in the presence of water or an organic solvent, at an elevated temperature to produce a corresponding 2-chloro-6-substituted-pyridine N-oxide and
- (b) reacting said 2-chloro-6-substituted-pyridine N-oxide with a base to produce the corresponding 1-hydroxy-6-substituted-pyridine-2-one.

In yet another aspect, the present invention relates to a process for producing a compound of Formula (II) wherein Y is nitrogen or a compound of Formula (III) which comprises the steps of:

- (a) reacting 2,6-dichloropyridine N-oxide, a primary or secondary amine containing between 1 and 20 carbon atoms (such as dimethanolamine or triethanolamine) and a base (such as sodium hydroxide or potassium hydroxide), optionally in the presence of water or an organic solvent, at an elevated temperature to produce a corresponding 2-chloro-6-substituted-pyridine N-oxide and
- (b) reacting said 2-chloro-6-substituted-pyridine N-oxide with a sulfur-containing base to produce the corresponding 1-hydroxy-6-substituted-pyridine-2-thione.

In still another aspect, this invention relates to a process for producing the compound of Formula (I) or (II), wherein X or Y is sulfur, which comprises the following steps:

- (a) reacting a 2,6-dichloro pyridine N-oxide, a thiol compound having between 1 and 20 carbon atoms, (for example alkyl, arylalkyl and aryl thiol compounds, and combinations thereof) and base (such as sodium hydroxide or potassium hydroxide) in an organic solvent at an elevated temperature to produce a corresponding 2-chloro-6-substituted-pyridine N-oxide and
- (b) reacting said 2-chloro-6-substituted-pyridine N-oxide with a base to produce the corresponding 1-hydroxy-6-substituted pyridine-2-one (i.e., the compound of formula (I)) or 1-hydroxy-6-substituted pyridine-2-thione (i.e., the compound of formula (II)).

The above processes can be carried out sequentially or simultaneously in a single step. Surfactants and/or phase transfer catalysts are optionally employed to facilitate the step (a) reaction in any of these processes.

In yet another aspect, the present invention relates to an antimicrobial composition comprising a functional component selected from the group consisting of paints, adhesives, coatings, elastomers, sealants, shampoos, skin care medicaments and metalworking fluid plus an antimicrobially effective amount of a compound represented by the above Formula (I) (II), or (III), or a salt of the compound of Formula (I) (II), or (III). Illustrative salts include the amine salts, alkali metal and alkaline earth metal salts and the like.

In still another aspect, the invention relates to a method for inhibiting the growth of microorganisms by contacting said microorganisms with a composition containing an antimicrobial effective amount of a one of the above-described 1-hydroxy-6-substituted pyridine-2-ones, 1-hydroxy-6-substituted pyridine-2-thiones, a salt thereof, or a combination thereof, of this invention and at least one component selected from the group consisting of soaps, shampoos, skin care medicaments, cosmetics, and other polymer compounds such as adhesives, coatings, elastomers, sealants and paints.

These and other aspects will become apparent upon reading the following detailed description of the invention.

DETAILED DESCRIPTION OF THE INVENTION

It has now been surprisingly discovered that a straightforward process can be employed to produce 1-hydroxy-6-substituted-2-pyridones or 1-hydroxy-6-substituted-2-thiones, utilizing readily available starting materials. These compounds provide excellent antimicrobial efficacy and are inexpensive to produce.

The novel 1-hydroxy-6-substituted pyridine-2-one and 1-hydroxy-6-substituted pyridine-2-thione compounds of this invention are shown formulae (I), (II) and (III) given above.

The reactions of the processes of this invention are suitably conducted at atmospheric pressure, although higher or lower pressures may be used if desired. Suitable reaction temperatures for steps (a) and (b) range between a temperature of about 25°C. to about 150°C., preferably employing a reflux temperature of between about 70°C. and about 90°C. Total reaction time for the process of this invention can vary over a wide range, but is advantageously between 30 minutes and 5 hours for step (a) and between thirty minutes and 5 hours for step (b).

The reactions of the processes of this invention are suitably carried out in the presence of a base and an organic solvent. Suitable bases include alkali metal hydroxides such as sodium hydroxide or potassium hydroxide, alkaline earth metal hydroxides such as calcium or magnesium hydroxide, pyridine, triethylamine and other tertiary amine bases, potassium carbonate, "DABCO" amine catalyst (1,4-diazabicyclo(2.2.2)octane), "DBU" (1,8-diazabicyclo(5.4.0)undec-7-ene), "DBN" (1,5-diazabicyclo(4.3.0)non-5-ene, t-butyltetramethyl-guanidine, combinations thereof, and the like. Preferred bases for step

(a) are anhydrous, granular or powdered sodium or potassium hydroxide, and preferred bases for step (b) are concentrated aqueous solutions of sodium or potassium hydroxide having a concentration of base of between about 10% and about 90% by weight based upon the weight of the solution. Preferably, the base is employed in at least an amount equal to the number of moles for each reactant, and a molar excess of base relative to each reactant can be employed as desired.

The reactions of the processes of this invention are suitably carried out in the presence of a solvent. Suitable organic solvents include, for example, ether, and acetone, methylene chloride, benzene, toluene, pyridine, tetrahydrofuran ("THF"), acetonitrile, dimethylsulfoxide ("DMSO"), dimethylformamide ("DMF"), and combinations thereof. Although the preferred solvent is acetonitrile, it is more preferred to run the reactions neat. As another alternative, a reactive solvent can be employed, if desired, for example by using an organic alcohol, e.g. isopropanol, or n-octanol, in an amount sufficient to provide the desired reactant and solvent characteristics.

The molar ratio of reactants for the processes of this invention can vary over a wide range, but is preferably between 10:1 and 1:10, more preferably between 2:1 and 1:2, most preferably about 1:1.

The antimicrobial compositions of this invention suitably comprise an antimicrobially effective amount of a compound as defined by the structural formulae given herein above, and combinations thereof, and at least one component selected from the group consisting of soaps, shampoos, skin care medicaments, cosmetics and paints. The antimicrobial compound can be employed as is or in the form of a salt, such as an amine salt, such as the monoethanolamine ("MEA") salt, and the use of the salt form is preferred. By the term "antimicrobial effective amount" is meant an amount sufficient to impart to the compositions resistance against microbial attack by fungi and/or bacteria. Preferably the antimicrobial compounds are employed in the composition in a total amount of between about 0.01 and about 10 weight percent, more preferably between about 0.01 and about 5 weight percent, based upon the total weight of the composition. A particularly preferred use for the biocidal compounds of the present invention is in personal care products such as in shampoos and other hair care products, as well as in paint, paint bases and polymer compositions such as adhesive coatings, sealants, elastomers, acids and the like.

With respect to paints improved organic solubility and biocidal efficacy associated with the compounds of the present invention are expected to provide advantages when used in a wide variety of paints, including indoor and outdoor household paints, industrial and commercial paints and in particular marine paints for use, for example, on ships hulls.

Typically, a paint composition will contain a resin, a pigment and various optional additives such as thickening agent(s), wetting agents and the like, as is well known in the art. The resin is preferably selected from the group consisting of vinyl, alkyl, epoxy, acrylic, polyurethane and polyester resins, and combinations thereof. The resin is preferably employed in an amount of between about 20% and about 80% based upon the weight of the paint or paint base.

In addition, the paint composition of the present invention contains optional additional additives which have a favorable influence on the viscosity, the wetting power and the dispersibility, as well as on the stability to freezing and electrolytes and on the foaming properties. If a marine paint is being fabricated, the paint preferably contains a co-biocide such as cuprous oxide or copper thio-cyanate, a swelling agent to cause the paint to gradually "slough off" in its marine environment, thereby causing renewed biocidal efficacy of newly exposed biocide at the surface of the paint in contact with the water medium of the marine environment. Illustrative swelling agents are naturally-occurring or synthetic clays, such as kaolin, montmorillonite (bentonite), clay mica (muscovite), and chlorite (hectorite), and the like. In addition to clays, other swelling agents, including natural or synthetic polymers, such as that commercially available as POLYMERGEL, have been found to be useful in the compositions of the present invention to provide the desired "sloughing off". Swelling agents can be used singly or in combination. The total amount of optional additives is preferably no greater than 20% by weight, more preferably between about 1% and about 5% by weight, based upon the total weight of the paint composition.

Illustrative thickening agents include cellulose derivatives, for example methyl, hydroxyethyl, hydroxypropyl and carboxymethyl cellulose, poly(vinyl alcohol), poly(vinylpyrrolidone), poly(ethylene-glycol), salts of poly(acrylic acid) and salts of acrylic acid/acrylamide copolymers.

Suitable wetting and dispersing agents include sodium polyphosphate, salts of low-molecular-weight poly(acrylic acid), salts of poly(ethane-sulfonic acid), salts of poly(vinyl-phosphonic acid), salts of poly(maleic acid) and salts of copolymers of maleic acid with ethylene, 1-olefins with 3 to 18 carbon atoms and/or styrene.

In order to increase the stability to freezing and electrolytes there may be added to the paint composition various monomer 1,2-diols, for example glycol, propylene-glycol-(1,2), and butylene-glycol-(1,2) or polymers thereof, or ethoxylated compounds, for example reaction products of ethylene oxide with long-chain alkanols, amines, carboxylic acids, carboxylic acid amides, alkyd phenols, poly(propylene-glycol) or poly(butylene-glycol). The minimum temperature of film formation (white point) of the paint composition may be reduced by adding solvents, such as ethylene-glycol, butyl-glycol, ethyl-glycol acetate, ethyl-diglycol acetate, butyl-diglycol acetate, benzene or alkylated aromatic hydrocarbons. As defoaming agents there are suitable for example poly(propylene-glycol) and polysiloxanes.

The compounds of the present invention have many desirable attributes. They possess good antimicrobial activity and are compatible with components of conventional soaps, shampoos, skincare medicaments, plastics and other polymer compositions and the like. These polymers are also non-volatile, hydrolytically-stable, thermally-stable and may be soluble in water and organic solvents. Furthermore, they form no undesirable colors in typical personal care items. Still further, they are expected to be favorably cost competitive with known antimicrobial additives used in conventional skin care formulations.

The following examples are intended to illustrate, but in no way limit the scope of, the present invention. In the examples, the term "g" denotes grams, "mol" denotes moles, and all percents are given on a weight basis unless otherwise specified.

EXAMPLES

Example 1

Preparation of 1-Hydroxy-6-isopropoxyloxy-pyridine-2(1H)-ones

The 0.82 g (0.0050 moles) of 2,6-dichlopyridine N-oxide and 0.78 g (0.013 moles) of isopropanol was reacted with 0.210 g (0.0050 moles) of ground sodium

hydroxide in 8.2 ml of DMSO at 80°C. overnight to give 2-chloro-6-isopropoxy-pyridine N-oxide. The excess of isopropanol was removed in vacuo and 0.800 g (0.020 mole) of ground sodium hydroxide was reacted at 3 hours at 100°C. to give 1-hydroxy-6-isopropoxy-pyridine-2(1H)-one. After cooling, it was added 74 ml of water and was adjusted with 6N HCl to pH 5. It was extracted with ethyl acetate (3x25 ml). The ethyl acetate extract was washed with water (2x10 ml) and dried over sodium sulfate. The solution was saturated with sodium chloride and was extracted with ethyl acetate (3x25 ml). The ethyl acetate extract was washed with water (2x10 ml) and dried over sodium sulfate. The combined ethyl acetate extracts were filtered and stripped in vacuo to give 0.50 g (59%). It was recrystallized from 5 ml of ethyl acetate, 5 ml of hexanes, and activated carbon: mp 119°-120°C.

Example 2

Preparation of 1-Hydroxy-6-(2,4,4-trimethylpentoxy)pyridine-2(1H)-one

The 0.82 g (0.0050 moles) of 2,6-dichloropyridine N-oxide and 0.664 g (98%) (0.0050 moles) of 2,4,4-trimethyl-1-pentanol was reacted with 0.211 g (0.0050 moles) of ground sodium hydroxide in 8.2 ml of DMSO at 100°C. overnight to give 2-chloro-6-(2,4,4-trimethylpentoxy)pyridine N-oxide. It was reacted with 0.600 g (0.015 moles) of ground sodium hydroxide at 100°C. for 4.5 hours to give 1-hydroxy-6-(2,4,4-trimethylpentoxy)pyridine-2(1H)-one. After cooling, it was added 74 ml of water and was adjusted with 6N HCl to pH 3. It was extracted with ethyl acetate (3x25 ml). The combined ethyl acetate extract was washed with a saturated sodium chloride solution (2x10 ml) and dried over sodium sulfate. The ethyl acetate extract was filtered and stripped in vacuo to give 1.08 g (90%). It was recrystallized from ethyl acetate and hexanes: mp 133.5°-134°C.

Example 3

Preparation of 1-Hydroxy-6-(3,5,5-trimethylhexyloxy)pyridine-2(1H)-one

The 0.82 g (0.0050 moles) of 2,6-dichloropyridine N-oxide and 0.80 g (90%) (0.0050 moles) of 3,5,5-trimethyl-1-hexanol was reacted with 0.206 g (0.0050 moles) of ground sodium hydroxide in 8.2 ml of DMSO at 80°C. for 8.5 hours to give 2-chloro-6-(3,5,5-trimethylhexyloxy)pyridine N-oxide. It was reacted with 0.600 g (0.015 moles) of ground sodium hydroxide at 80°C. for 4 hours to give 1-hydroxy-6-(3,5,5-trimethylhexyloxy)pyridine-2(1H)-one. After cooling, it was added 74 ml of water and was adjusted with 6N HCL to pH 3. The precipitate was filtered and washed with water to give 0.577 g (45%). It was recrystallized from ethyl acetate and hexanes: mp 130.5°-131°C.

Example 4

Preparation of 1-Hydroxy-6-octyloxy-pyridine-2(1H)-one

The 0.82 g (0.0050 moles) of 2,6-dichloropyridine N-oxide and 0.658 g (99%) (0.0050 moles) of 1-octanol was reacted with 0.200 g (0.0050 moles) of ground sodium hydroxide in 8.2 ml of DMSO at 80°C. for 4.5 hours to give 2-chloro-6-octyloxy-pyridine N-oxide. It was reacted with 0.600 g (0.015 moles) of ground sodium hydroxide at 80°C. for 2.5 hours to give 1-hydroxy-6-octyloxy-pyridine-2(1H)-one. After cooling, it was added 74 ml of water and was adjusted with 6N HCL to pH 3. The precipitate was filtered and washed with water to give 0.577 g (48%). It was recrystallized from ethanol and hexanes: mp 119°-120°C.

Example 5

Preparation of 1-Hydroxy-6-octylthiopyridine-2(1H)-one

The 2.01 g (0.0122 moles) of 2,6-dichloropyridine N-oxide and 2.10 g (85%) (0.0122 moles) of 1-octylmercaptan was reacted with 0.488 g (0.0122 moles) of ground sodium hydroxide in 8.2 ml of DMSO at 80°C. for 6 hours to give 2-chloro-6-octylthiopyridine N-oxide. It was reacted with 1.47 g (0.0367 moles) of ground sodium hydroxide at 80°C. for 3 hours to give 1-hydroxy-6-octylthiopyridine-2(1H)-one. After cooling, it was added 180 ml of water

and was adjusted with 6N HCL to pH 3. The precipitate was filtered, washed with water and washed with 150 ml of petroleum ether to give 1.34 g (43%).

Example 6

Preparation of 1-Hydroxy-6-octylsulfonylpyridine-2(1H)-one

The 0.204 g (0.00080 moles) of 1-hydroxy-6-octylthiopyridine-2(1H)-one was reacted with 0.36 ml (0.0032 moles) of 30% hydrogen peroxide in 2.0 ml of glacial acetic acid for overnight at room temperature to give 1-hydroxy-6-octylsulfonylpyridine-2(1H)-one. It was stripped in vacuo and was then dissolved with ethanol and stripped in vacuo three times to give 0.22 g. It was recrystallized from acetone and petroleum ether.

Example 7

Preparation of 1-Hydroxy-6-thiophenylpyridine-2(1H)-one

The 1.64 g (0.010 moles) of 2,6-dichloropyridine N-oxide and 1.11 g (99%) (0.010 moles) of thiophenol was reacted with 0.400 g (0.010 moles) of ground sodium hydroxide of 16.4 ml of DMSO at 80°C. for 5 hours to give 2-chloro-6-thiophenylpyridine N-oxide. It was reacted with 1.20 g (0.030 moles) of ground sodium hydroxide at 80°C. for 2.5 hours to give 1-hydroxy-6-thiophenylpyridine-2(1H)-one. After cooling, it was added 148 ml of water and was adjusted with 6N HCl to pH 3. The precipitate was filtered, washed with water and washed with hexanes to give 1.64 g (75%). It was recrystallized from ethanol: mp 180°-181°C.

Example 8

Preparation of 1-Hydroxy-6-thiophenylsulfonylpyridine-2(1H)-one

The 1.00 g (0.00456 moles) of 1-hydroxy-6-thiophenylpyridine-2(1H)-one reacted with 1.30 ml (0.0114 moles) of 30% hydrogen peroxide in 5 ml of glacial acetic acid at room temperature overnight. The precipitate was dissolved with more 5 ml of glacial

acetic acid and was then added 1.30 ml (0.0114 moles) of 30% hydrogen peroxide at room temperature over the weekend. The precipitate was filtered and washed with hexanes to give 0.624 g. The acetic acid and the hexanes were stripped in vacuo to give 0.45 g. The combined product gave 1.14 g (93%).

Example 9

Preparation of 1-Hydroxy-6-phenyloxypyridine-2(1H)-one

The 0.82 g (0.0050 moles) of 2,6-dichloropyridine N-oxide and 0.47 g (0.0050 moles) of phenol was reacted with 0.200 g (0.0050 moles) of ground sodium hydroxide in 8.2 ml of DMSO at 80°C. for 6 hours to give 2-chloro-6-phenyloxypyridine N-oxide. It was reacted with 0.600 g (0.015 moles) of ground sodium hydroxide at 80°C. for 3.5 hours to give 1-hydroxy-6-phenyloxypyridine-2(1H)-one. After cooling, it was added 74 ml of water and was adjusted with 6N HCl to pH 3. The water solution was extracted with dichloromethane (2x25 ml) and was dried over sodium sulfate. The dichloromethane solution was stripped in vacuo to give 1.29 g which was purified with Flash Chromatography(9:1 ethyl acetate:methanol):0.82 (65%).

Example 10

Preparation of 1-hydroxy-6-octylaminopyridine-2(1H)-ones

To 1.53 g(0.008moles) of 2,6-dichloropyridine in 13.3 ml of toluene was added 1.26 g) 0.0162 moles) 1-octylamine and refluxed for 3 hours. The reaction content was cooled to room-temperature, washed several times with water to yield 2.02 g of 2-chloro-6-octylaminopyridine-N-oxide as a yellow liquid.

The yellow liquid refluxed with 2.7 g of potassium tert-butoxide in tert-butyl alcohol overnight. Water was added to the reaction flask and a precipitate was formed upon adjustment of pH to 3. The precipitate filtered and washed to yield 1.36 g of product.

Example 11

Preparation of 1-hydroxy-6-octyloxy-pyridine-2(1H)-thiones

2-Chloro-6-octyloxy-pyridine-N-oxide was prepared according to Example 4 and subsequent reaction with NaSH in acetonitrile and extraction with methylene chloride yielded the product as white powder.

Example 12

Preparation of 1-hydroxy-6-octylamino-pyridine-2(1H)-thiones

A reaction mixture of 0.52 g of 2-chloro-6-octylamino-pyridine-N-oxide (prepared as described in Example 10), 5 ml DMSO and 0.55 ml of 46% solution of NaSH was heated at 80 degrees for two hours. The reaction was diluted with water, pH was adjusted to three, and the product was extracted with methylene chloride.

Example 13

Proposed example for preparation of 1-hydroxy-6-octylthio-pyridine-2(1H)-thiones

2-Chloro-6-octylthio-pyridine-N-oxide is prepared according to the method described in Example 5, and subsequent reaction with aqueous solution of NaSH yields the titled product.

Determination of the Minimum Inhibitory Concentrations (MIC's)
for Antimicrobial Compounds of this Invention

Solutions of the experimental compounds in dimethyl sulfoxide were serially diluted in nutrient broth (Tryptic Soy Broth for bacteria and Sabouraud Dextrose Broth for fungi) in microliter plates. Equal volumes of a broth suspension of bacteria (10^6 CFU/ml) or fungi (10^5 cells or spores/ml) were added to each dilution, and the plates were incubated at 37°C. (bacteria and yeast) or 28°C. (molds). Bacteria, yeast and molds were incubated two, five and seven days respectively before determining the highest inhibitory dilution.

TABLE I

Compound	MIC (ppm)					
	<u>MIC of the 1-hydroxy-6-substituted-2-pyridones</u>					
	A	B	C	D	E	F
Example 1	256	≥1024	128	512	512	512
Example 2	64	64	8	32	8	16
Example 3	32	32	2	2	16	2
Example 4	32	16	4	2	32	4
Example 5	256	16	8	32	8	nt
Example 6	512	64	128	256	256	nt
Example 7	64	128	16	64	64	nt
Example 8	256	8	1024	1024	1024	nt
Example 9	256	512	32	256	512	nt
Example 10	32	32	8	16	32	16
Example 11	16	16	4	4	8	16
Example 12	256	64	8	16	64	64
OCTOPIROX® biocide	128	64	2	2	2	16-32

A = Escherichia coli

B = Staphylococcus aureus

C = Candida albicans

D = Fusarium sp.

E = Aspergillus niger

F = Aureobasidium pullulans

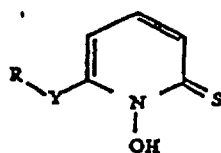
"nt" denotes not tested

The results provided in Table 1 show that the compounds of the present invention provide MIC's that are sometimes better than those provided by a commercial biocide, OCTOPIROX® biocide, a product of Hoechst Company. Note that the compounds of Examples 3, 4 and 11 provide particularly excellent MIC results.

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What is claimed is:

1. A compound of the formula:



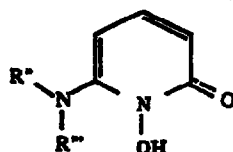
wherein Y is a sulfur, or NR' moiety, wherein N is nitrogen and R and R' are independently each a substituted or unsubstituted hydrocarbon radical having from 1 to 20 carbon atoms.

2. The compound of claim 1 wherein R or R' is an aliphatic or aromatic hydrocarbon having from 2 to 18 carbon atoms, with the proviso that said aromatic hydrocarbon have at least 6 carbon atoms.

3. The compound of claim 1 wherein R or R' is a straight chain hydrocarbon having from 5 to 10 carbons.

4. The compound of claim 1 wherein R or R' is an n-octyl moiety.

5. A compound of the formula:

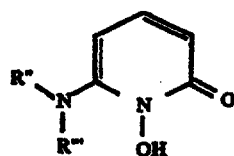


wherein R'' and R''' are independently either hydrogen or a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms.

6. The compound of claim 5 wherein R'' or R''' is an aliphatic or aromatic hydrocarbon having from 2 to 18 carbon atoms, with the proviso that said aromatic hydrocarbon have at least 6 carbon atoms.

7. The compound of claim 5 wherein R'' or R''' is a straight chain hydrocarbon having from 5 to 10 carbons.

8. A compound of the formula:



wherein R'' and R''' are independently either hydrogen or a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms, and wherein R'' or R''' is an n-octyl moiety.



9. A process for producing the compound of claim 1, wherein Y in the formula of claim 1 is oxygen, which comprises the steps of:

- (a) reacting 2,6-dichloropyridine N-oxide, a hydroxy compound containing from 1 to 20 carbon atoms and a base, optionally in the presence of water or an organic solvent, to produce a corresponding 2-chloro-6-substituted-pyridine N-oxide, and
- (b) reacting said 2-chloro-6-substituted-pyridine N-oxide with a sulfur-containing base to produce the corresponding 1-hydroxy-6-substituted-pyridine-2-thione,

steps (a) and (b) being carried out at a temperature independently selected from about 25 to about 150 degrees Centigrade.

10. A process for producing a compound of claim 1, wherein Y in the formula of claim 1 is NR' which comprises the steps of:

- (a) reacting 2,6-dichloropyridine N-oxide, a primary or secondary amine containing from 1 to 20 carbon atoms and a base, optionally in the presence of water or an organic solvent, at an elevated temperature to produce a corresponding 2-chloro-6-substituted-pyridine N-oxide, and
- (b) reacting said 2-chloro-6-substituted-pyridine N-oxide with a sulfide or hydrosulfide salt to produce the corresponding 1-hydroxy-6-substituted-pyridine-2-thione,

steps (a) and (b) being carried out at a temperature independently selected from about 25 to about 150 degrees Centigrade.

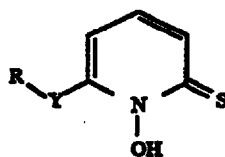
11. A process for producing a compound of claim 1, wherein Y in the formula of claim 1 is sulfur, which comprises the steps of:

- (a) reacting a 2,6-dichloropyridine N-oxide, a thiol compound having from 1 to 20 carbon atoms, and base in an organic solvent at an elevated temperature to produce a corresponding 2-chloro-6-substituted-pyridine N-oxide, and
- (b) reacting said 2-chloro-6-substituted-pyridine N-oxide with a sulfide or hydrosulfide salt to produce the corresponding 1-hydroxy-6-substituted-2-pyridine-2-thione,



steps (a) and (b) being carried out at a temperature independently selected from about 25 to about 150 degrees Centigrade.

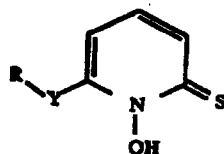
12. A process for producing a compound of claim 5 which comprises the steps of:
- (a) reacting 2,6-dichloropyridine N-oxide, a primary or secondary amine containing from 1 to 20 carbon atoms and a base, optionally in the presence of water or an organic solvent, at an elevated temperature to produce a corresponding 2-chloro-6-substituted-pyridine N-oxide, and
 - (b) reacting said 2-chloro-6-substituted-pyridine N-oxide with a base to produce the corresponding 1-hydroxy-6-substituted-pyridine-2-one,
- steps (a) and (b) being carried out at a temperature independently selected from about 25 to about 150 degrees Centigrade.
13. An antimicrobial composition comprising a functional product selected from the group consisting of shampoos or skin care medicaments further comprising an antimicrobially effective amount of a compound of the formula:



wherein Y is an oxygen, sulfur, or NR' moiety, wherein N is nitrogen and R and R' are each a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms and conventional antimicrobial additives.

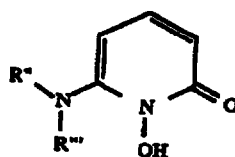
14. The composition of claim 13 wherein R or R' in said formula is an aliphatic or aromatic hydrocarbon having from 3 to 18 carbons.
15. The composition of claim 13 wherein R or R' in said formula is a straight chain hydrocarbon having from 5 to 10 carbons.
16. The compound of claim 1 wherein R or R' in said formula is an n-octyl moiety.
17. A method for inhibiting the growth of microorganisms by contacting said microorganisms on skin or scalp with a shampoo or skin care composition comprising an antimicrobial effective amount of a compound of the formula:





wherein Y is an oxygen, sulfur, or NR' moiety, wherein N is nitrogen and R and R' are each a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms.

18. An antimicrobial composition comprising a functional product selected from the group consisting of shampoos or skin care medicaments, further comprising an antimicrobially effective amount of a compound of the formula:



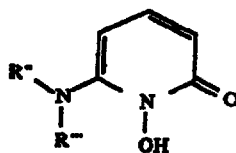
wherein R'' and R''' are independently either hydrogen or a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms and conventional antimicrobial additives.

19. The composition of claim 18 wherein R'' or R''' in said formula is an aliphatic or aromatic hydrocarbon having from 3 to 18 carbons.

20. The composition of claim 18 wherein R'' or R''' in said formula is a straight chain hydrocarbon having from 5 to 10 carbons.

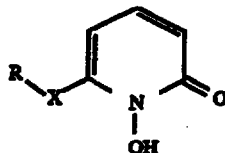
21. The composition of claim 18 wherein R'' or R''' in said formula is an n-octyl moiety.

22. A method for inhibiting the growth of microorganisms by contacting said microorganisms or skin or scalp with a shampoo or skin care composition comprising an antimicrobial effective amount of a compound of the formula:



wherein R'' and R''' are either hydrogen or a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms and conventional antimicrobial additives.

23. An antimicrobial composition comprising a functional product selected from the group consisting of shampoos or skin care medicaments and further comprising an antimicrobially effective amount of a compound of the formula:



wherein X is an oxygen or sulfur moiety and R is a substituted or unsubstituted hydrocarbon radical having between 1 and 20 carbon atoms, with the proviso that R is other than chlorobenzyl.

