Long-Chain Alkanoylcholines, a New Category of Soft Antimicrobial Agents That Are Enzymatically Degradable

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A new category of amphiphilic hydrolyzable quaternary ammonium compounds with rapid and high levels of antimicrobial activity was studied. The compounds, alkanoylcholines with hydrocarbon chains of 10 to 14 carbon atoms, are hydrolyzed by butyrylcholine esterase, which is present in human serum and mucosal membranes. The hydrolysis products are common components of human metabolism. Alkanoylcholines were tested and found to be active against gram-negative and gram-positive bacteria as well as yeasts. The microbicidal activities of the alkanoylcholines were comparable to the activities of the stable quaternary ammonium compounds of corresponding chain length and increased with an increasing number of carbon atoms. The compounds were also found to be hydrolyzed by enzymes present in certain microorganisms. The degradation was achieved after reaching the microbicidal effect.

Amphiphilic quaternary ammonium compounds are surface-active substances that are known for their membrane-disruptive effects and antimicrobial activities (10, 14). These compounds exhibit rapid activity against a broad range of microorganisms: gram-positive and gram-negative bacteria, fungi, and certain viruses. However, the high degree of affinity of these agents for biological membranes in general results in low selectivity, which can lead to damage to mammalian cells if exposure times are sufficiently long.

The side effects of this type of compound can be minimized by introducing a cleavable moiety into the molecule, such as an ester bond. This will give compounds that are more labile and that can be subject to hydrolytic degradation. The hydrolysis step should preferably lead not only to a nonactive compound but also to degradation products that are significantly less toxic than the original substance. Compounds based on this type of concept have been introduced as “soft” drugs. Soft drugs are defined as biologically active, therapeutically useful compounds characterized by a controllable degradation to nontoxic moieties after they achieve their therapeutic effect. This concept can be applied to limit the toxicities of antimicrobial agents, and the first soft drug to be introduced was actually an antimicrobial agent (2).

The work in our laboratory relating to soft antimicrobial agents has been centered around cationic amphiphilic compounds which degrade into natural products of human metabolism. In line with this work, we have earlier introduced long-chain alkyl betaine esters as soft antimicrobial agents (6, 12). These compounds were shown to have good microbicidal activity and are degraded to betaine and long-chain alcohols. The rate of hydrolysis of the compounds is pH dependent and was found to be highly influenced by the amphiphilic nature of the substances (22, 23).

In order to expand the field of soft antimicrobial agents to include enzymatically degradable compounds, we initiated studies with a series of alkanoylcholines with various hydrocarbon chain lengths (Fig. 1) (4). Our interest in these compounds was due to their potential degradation by hydrolytic enzymes. The hydrolysis products of long-chain alkanoylcholines, choline and fatty acids, are common human metabolites, and their metabolic pathways have been well investigated (13, 18).

Butyrylcholine esterase (BChE; EC 3.1.1.8), which is found in mammalian serum and mucosal membranes, is known to catalyze the hydrolysis of short-chain alkanoylcholines (19). Early studies of BChE indicated that butyrylcholine is the substrate with optimal activity and that long-chain alkanoylcholines are not hydrolyzed by the enzyme (9). We found, however, that the long-chain alkanoylcholines are hydrolyzed in vitro but that the reaction does not follow Michaelis-Menten kinetics and shows deviation from linearity caused by inhibition of the reaction at high substrate concentrations (4). We showed that the inhibition could be a result of conformational changes of the protein due to the amphiphilic nature of the substrates. Furthermore, the formation of mixed micelles between the substrate and the fatty acids formed during the hydrolysis reaction will lower the monomer concentration of the substrate, which could also partially explain the inhibition.

The in vitro hydrolysis studies were complemented with an investigation on the hydrolysis of alkanoylcholines in a biological environment, the rat intestine (5). These studies showed that alkanoylcholines are readily hydrolyzed in vivo and that the rate of hydrolysis decreases with increasing hydrocarbon chain length. Furthermore, the hydrolysis follows the occurrence of BChE through the rat intestinal tract (11, 20), indicating that BChE is responsible for the degradation of the amphiphilic alkanoylcholines.

The stabilities of the alkanoylcholines could also be influenced by hydrolytic enzymes of microbial origin, but the presence of BChE in microorganisms has not been reported previously.

This report describes the microbicidal activities of long-chain alkanoylcholines against a range of bacteria and yeasts as well as the hydrolytic effects on the substances of enzymes present in the microorganisms.

MATERIALS AND METHODS

Compounds investigated. The alkanoylcholine bromides were prepared as follows. Fatty acid chlorides (C10 to C14) were dissolved in dry diethyl ether, and
the solution was added dropwise under stirring to a solution of N,N-dimethylaminoethanol in dry diethyl ether. The precipitates, 2-(N,N-dimethylamino)ethyl ester hydrochlorides, were filtered off and recrystallized from warm ethyl acetate. The products were further used for the syntheses of both labeled and unlabeled alkanoylcholines.

The aminooxy ether hydrochloride was dissolved in water, and a saturated solution of NaOH was added to liberate the free base. The solution was then extracted twice with diethyl ether, and the organic phase was dried with MgSO₄. An appropriate amount of [³H]methyl iodide dissolved in toluene was added, and the solution, in a thoroughly closed vial, was left for 2 h at room temperature. An excess of methyl bromide was then added, and the mixture was left overnight. The precipitate was filtered off and washed with diethyl ether. The radiochemical purity of the products was verified by high-pressure liquid chromatography (HPLC) with on-line beta-radiation counting detection by calculating the areas of the peaks obtained in the chromatograms (24).

For the syntheses of unlabeled alkanoylcholines, the extraction of free amino ester was performed with ethyl acetate. Only the excess of methyl bromide was added, and the mixture was kept at 37°C for 1 h and then at room temperature overnight.

A more detailed description of the synthesis can be found in reference 3.

Electrospray ionization mass spectrometric analysis of the compounds gave spectra containing only the M⁺ molecular ion (258, 286, and 314 for C₁₀, C₁₂, and C₁₄, respectively). However, increasing the accelerating voltage led to collision-induced dissociation and a certain extent of fragmentation. For C₁₀, an increase in the voltage from 30 to 100 V gave a spectrum with the molecular ion (258 m/z) and major fragmentation ions at 199 [(CH₃)₂NCH₂CH₂OC(CH₃)₃]⁺, 85 [(CH₃)₂NCH₂CH₂]⁺, 71 [(CH₃)₂NCH₂CH₃]⁺, 57 [(CH₃)₂NCH₂H]⁺, and 43 [(CH₃NCH)⁺].

Instrumentation and chromatography. The liquid scintillation counter used in the study was an LKB 1217 instrument (LKB Wallac, Turku, Finland). The radiochromatography system was composed of an LKB model 2150/2152 dual pump gradient delivery system, a 100-µl loop, a 150-by-4.6-mm (inner diameter) analytical column packed with 5-µm Kromasil C₈ (Eka Nobel, Surte, Sweden), and a Radiomatic A-250 radioactivity monitor. The detector was run in the homogeneous mode and was equipped with a 2.5-ml liquid scintillation cell.

The scintillator fluid/mobile phase ratio was 4:1. The mobile phases consisted of acetone/toluene-water mixtures adjusted to pH 3.0 with hydrochloric acid, with a flow rate of 1 ml/min.

The electrospray ionization mass spectrometer used for structure confirmation was a Trio 2000 instrument (Fison Instruments, Altrincham, United Kingdom). Samples were dissolved in acetonitrile-water (1:1) and were introduced into the ion source at a flow rate of 10 µl/min.

Microorganisms and culture conditions. The antimicrobial activity was tested on Salmonella typhimurium 395 MS (serogroup B, S form) and 395 MR10 (R mutant, chemotype Rd), which have been described in detail elsewhere (7), and on Salmonella braenderup CCUG 12644 (serogroup C1); Salmonella infantis CCUG 12615 (serogroup C1); Escherichia coli CCUG 11412 and CCUG 11320; Streptococcus agalactiae CCUG 4208, CCUG 4209, and CCUG 4210; Staphylococcus saprophyticus CCUG 18736; Campylobacter jejuni (C. col) CCUG 11701 and CCUG 11702; and Lactobacillus casei CCUG 21451 (Culture Collection, University of Göteborg, Göteborg, Sweden). Six Candida albicans and four Candida glabrata isolates were obtained from recently cultured clinical specimens. All bacteria and yeasts were maintained on agar slants in the cold.

The solid medium used for S. typhimurium, E. coli, and S. saprophyticus (group I) was nutrient agar, for S. agalactiae, L. casei, and C. jejuni (C. col) (group II) the solid medium was blood agar, and for C. albicans and C. glabrata (group III) the solid medium was Sabouraud agar. The liquid media were nutrient broth, brain heart infusion broth, and Sabouraud broth, respectively.

All microorganisms were grown in 20 ml of broth, groups I and III were grown on a rotary shaker at 37°C for 18 h, S. agalactiae and L. casei were grown in a CO₂ atmosphere at 37°C for 48 h, and C. jejuni (C. col) was grown in a microaerophilic atmosphere (5% O₂, 10% CO₂, 85% N₂) at 42°C for 48 h.

Microbiological activities of alkanoylcholines. The susceptibilities of the microorganisms to the alkanoylcholines were determined by two different techniques. In both techniques, 20 ml of the bacteria and yeast cultures were harvested by centrifugation and were washed once with phosphate-buffered saline solution. The sediments were resuspended in 2 ml of 0.1 M phosphate buffer (pH 8.0). The yeast suspensions were diluted to obtain an optical density of 0.7 absorbance units at 610 nm, and 40 ml of the cell suspension was then centrifuged and resuspended in 2 ml of 0.1 M phosphate buffer (pH 8.0). The microorganisms were exposed to serial dilutions of the esters in 0.1 M phosphate buffer at pH 8 and 37°C for 10 min.

The first of the techniques used the spiral plater system (Spiral Systems, Cincinnati, Ohio). The samples (1 ml per vial) were diluted 100-fold with Letheen broth (Thiotone peptone [BBL Microbiology Systems, Cockeysville, Md.], 10.0 g; beef extract, 5.0 g; lecithin, 0.7 g; polyborate 80, 5.0 g; sodium chloride, 5.0 g; distilled water, 1,000 ml). This broth inactivates quaternary ammonium salts and is recommended as a neutralizing diluent for the evaluation of disinfectants containing cationic surfactants (1). The suspensions and dilutions were then plated onto nutrient agar.

The other method used a microtiter plate technique for the rapid determination of minimum microbicidal concentrations. Aliquots of 5 µl were taken from each well (0.2 ml per well) and were directly transferred onto nutrient agar plates after the 10-min exposure. All agar plates were incubated for 1 to 3 days in the appropriate atmosphere and at the appropriate temperature for the microorganism tested.

Uptake and hydrolysis of alkanoylcholines. Undiluted cell suspensions of the organisms (100 µl) were added to 900 µl of preincubated 0.1 M phosphate buffer (pH 8.0) which contained different concentrations of radiolabeled tetradecanoylcholine. After mixing, the suspensions were incubated at 37°C for 10 min, after which the cells were pelleted by centrifugation at 12,500 rpm (Microfuge E; Beckman) for 5 min. The radioactivity of the supernatant was measured either after separation by HPLC with on-line detection or with a liquid scintillation counter.

RESULTS

Antimicrobial effects of alkanoylcholines as a function of chain length. To determine the influence of the chain length of alkanoylcholines on the bactericidal effect, we tested a series of esters with 10 to 14 carbon atoms in the alkyl chain, decanoylcholine (C₁₀), dodecanoylcholine (C₁₂), and tetradecanoylcholine (C₁₄), against S. typhimurium 395 MR10 and S. infantis 12615 using the spiral plater technique (Fig. 2 and 3). In 0.1 M sodium phosphate buffer at pH 8.0 and 37°C, C₁₄ showed the highest and C₁₀ showed the lowest level of activity. C₁₂ was only slightly less effective than C₁₄. There was no significant difference in the antibacterial effects on the two Salmonella strains.

We tested the same series of alkanoylcholines against C. albicans and C. glabrata. The yeasts were exposed to different concentrations of the esters at 37°C. The killing effects of the alkanoylcholines against both Candida strains increased, as expected, with an increase in the chain length of the ester (Fig. 4 and 5). The effect of the stable quaternary ammonium com-
pound hexadecyltrimethylammonium bromide (CTAB) on C. albicans was tested as a comparison with those of the hydrolysable cationic surfactants. CTAB was slightly more effective than C_{14}.

**Determination of MBCs.** The microtiter plate technique was used to determine MBCs and minimal fungicidal concentrations (MFCs) (Table 1). Low MBCs were found with C_{14} for both gram-positive and gram-negative bacteria. The only deviation was found with *S. saprophyticus*. C_{10} and C_{12} were only tested on *Salmonella* spp., and the activity of C_{12} was found to be in the same range as that of C_{14}.

MFCs were determined for *C. albicans*, *C. glabrata*, *C. parapsilosis*, and *C. krusei*. The killing effect was similar for all species and increased with an increase in the alkyl chain length of the choline esters.
Degradation of tetradecanoylcholine in the presence of bacteria and yeasts. A survey of the capacity of different bacteria and yeasts to degrade C14 was performed by exposing the microorganisms to 0.4 mM [3H]tetradecanoylcholine at 37°C for 10 min. The suspensions were then centrifuged, and the radioactivity in the supernatant was measured in a scintillation counter. Assuming that most of the nonhydrolyzed alkanoylcholine was bound to the cells, enhanced radioactivity in the supernatant would indicate that C14 was hydrolyzed and that choline was released. As can be seen in Table 2, the highest level of activity in the supernatant was found with the Salmonella species tested (67 to 79%) and with the C. glabrata isolates (69 to 100%). In contrast, S. agalactiae gave only 6% activity in the supernatant.

In order to ascertain the contribution of C14 to the radioactivity in the supernatant, the activities of the different components were determined after separation by HPLC. The differences observed between the strains (Table 3) were analogous to those recorded in Table 2. The most effective hydrolysis of C14 was found with C. glabrata, which gave 0.33 mM choline after 10 min from 0.4 mM ester (Table 3). The four Salmonella spp. showed hydrolysis that was almost as good as that of C. glabrata, e.g., 0.29 mM choline with S. infantis.

### DISCUSSION

It has been established that cationic surfactants interact with the lipid bilayer structures of microbial cell membranes. The cationic polar head group of this class of compounds interacts with negative charges on the surface of the membrane, while the hydrocarbon chain is intercalated into the hydrophobic interior (14). The series of alkanoylcholines studied by us show differences in their antimicrobial effects, depending on the length of the hydrocarbon chain, since the choline part of the compounds is identical throughout the series. The highest degree of bactericidal and fungicidal activity was found for tetradecanoylcholine (C14). The decyl derivative (C10) lacked activity at concentrations of less than 2 mM for the bacteria and 5 mM for the yeasts under the experimental conditions used (Fig. 2 to 5). The differences in activity between C12 and C14 were largest against C. glabrata and C. albicans, less difference was observed against S. typhimurium, and the antimicrobial effects of both compounds were almost the same against S. infantis. These results are in good agreement with those of studies of the nonhydrolyzable amphiphilic ammonium compounds (8). In that study, optimal activity against bacterial strains and yeasts was found with chain lengths of 14 and 16, respectively. This can be explained by differences in the envelope and phospholipid compositions of the microorganisms, which affect interactions between the compounds studied and the cell membrane.

The results presented above are also in good agreement with those found with the long-chain alkyl betainates previously developed in our laboratory (12). Furthermore, the concentrations of both groups of hydrolyzable compounds necessary to give antimicrobial activity are comparable to those of the corresponding stable quaternary ammonium antimicrobial agents such as CTAB (Fig. 4) (12). As can be seen, CTAB shows a slightly greater antimicrobial effect than C14. However, since the antimicrobial effect is dependent to a large extent on the number of carbon atoms in the hydrophobic chain, substances with similar chain lengths should be compared. The motivation for comparing C14 with CTAB (CTAB has 16 carbon atoms in the hydrophobic chain) is that for both the betaine and choline esters the critical micelle concentration is similar to that of a stable quaternary ammonium compound with two extra CH2 groups, i.e., the critical micelle concentration of C14 is comparable to that of CTAB (4, 17). This would indicate that the ester function will be infused into the hydrophobic interior of the micelle. That this comparison of CTAB and C14 will also hold true in a system such as a bacterial membrane, however, cannot be taken for granted. Studies of transport kinetics in liposome lecithin membranes indicates that the flux of SO4 2− across the membrane caused by the introduction of stable
quaternary ammonium compounds with 12 carbon atoms in the hydrophobic chain is similar to that for a betaine ester with the same number of carbon atoms (16). Furthermore, it is known that the stable quaternary ammonium compounds show maximum antimicrobial activity with chain lengths of 14 to 16 carbon atoms (8). We have earlier shown that for betaine esters the maximum activity is also found with chain lengths of 16 to 18 carbon atoms (12). Thus, for the alkanoylcholines a more appropriate comparison might have been CTAB and a C₁₆ ester. However, for the alkanoylcholines, we have only tested chain lengths of up to 14 carbon atoms because of the negative effect on enzymatic hydrolysis by the more hydrophobic esters (4).

Tetradecanoylcholine, C₁₄, being the most effective ester, was chosen as the test substance against a representative selection of different bacteria and yeasts (Table 1). The MBC obtained showed that the antimicrobial effects of hydrolyzable quaternary ammonium compounds are excellent against gram-negative as well as gram-positive bacteria and are comparable to those of stable quaternary ammonium compounds. The MBC of C₁₄ ranges between 0.4 and 0.8 mM. The low MBC (0.2 mM) for S. typhimurium 395 R10 may be due to the deep rough character of the outer membrane of the bacteria, making it both hydrophobic and accessible to surface-active agents (15). The deviation observed with S. saprophyticus (3.2 mM) is probably due to the fact that staphylococci grow in clusters, which impedes the accessibility of the antimicrobial agent to the individual cells and implies that in these cases there is more than one viable cell per CFU. Kinetic studies of biocide killing of cluster-forming bacteria show that plots of the natural logarithm of survivors versus time are convex, i.e., high survival rates with short exposure times (21). This multiple-hit phenomenon, together with the obstructed access of the antimicrobial agent to the individual cell, would be expected to give lower killing rates with the short exposure times used in the present study. However, if the exposure time is sufficiently long, the effect of C₁₄ should be as good as that against the other microorganisms tested. The MBC of C₁₄ against different Candida species is in the range of 0.2 to 0.8 mM. However, the antimicrobial effect is more chain length dependent with yeasts than with bacteria.

The primary idea of using alkanoylcholines as soft antimicrobial agents was that the compounds are readily degraded by BChE, which is present in mammalian tissues and serum (4). However, during the studies of the antimicrobial effect, it was found that some of the bacteria and yeasts investigated also catalyzed the hydrolysis of alkanoylcholines. Despite this, the killing effect of the compounds in 10-min exposures was not affected conspicuously. Thus, C. glabrata cells were killed by C₁₄ as well as catalyzed the hydrolysis of the compound. Similar results were found with the different Salmonella spp., while the other microorganisms tested showed less or almost no hydrolysis capacity (Table 2).

The hydrolysis results in Table 3 show that the amount of choline released increases with the amount of added choline ester up to a plateau level. In the case of E. coli, however, no choline seems to be generated above a certain ester concentration. This could be due to a denaturation of the enzyme by excess ester. A similar effect was previously observed in vitro with BChE (4).

In conclusion, the long-chain alkanoylcholines show rapid antimicrobial activity against a wide range of microorganisms. At the same time, the esters are enzymatically hydrolyzable to nonactive components not only by BChE in mammalian systems, as shown earlier, but also by enzyme systems present in the microorganisms themselves. These substances are thought to have a good potential to replace the more toxic stable quaternary ammonium compounds.

ACKNOWLEDGMENT

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REFERENCES


TABLE 3. Concentration of free choline in the supernatant after hydrolysis of C₁₄ by different microorganisms

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<th>Organism</th>
<th>OD&lt;sub&gt;610&lt;/sub&gt;&lt;sup&gt;a&lt;/sup&gt;</th>
<th>Concen (mM)</th>
<th>Choline concn (mM)</th>
<th>Ester concn (mM)</th>
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* The results were obtained after exposure of different concentrations of [³H]tetradecanoylcholine to bacteria and yeasts at 37°C for 10 min. The radioactivity was measured after separation by HPLC and on-line detection.
* OD<sub>610</sub>, optical density at 610 nm, measured after diluting bacteria 1/5 and yeasts 1/10 in buffer.

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<sup>b</sup> OD<sub>610</sub>, optical density at 610 nm, measured after diluting bacteria 1/5 and yeasts 1/10 in buffer.


