

Chlorination of Taurine by Human Neutrophils: *EVIDENCE FOR HYPOCHLOROUS ACID GENERATION*

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The model hydrogen peroxide-myeloperoxidase-chloride system is capable of generating the powerful oxidant hypochlorous acid, which can be quantitated by trapping the generated species with the β -amino acid, taurine. The resultant stable product, taurine chloramine, can be quantitated by its ability to oxidize the sulfhydryl compound, 5-thio-2-nitro-benzoic acid to the disulfide, 5,5'-dithiobis(2-nitroben-zoic acid) or to oxidize iodide to iodine. Using this system, purified myeloperoxidase in the presence of chloride and taurine converted stoichiometric quantities of hydrogen peroxide to taurine chloramine. Chloramine generation was absolutely dependent on hydrogen peroxide, myeloperoxidase, and chloride and could be inhibited by catalase, myeloperoxidase inhibitors, or chloride-free conditions. In the presence of taurine, intact human neutrophils stimulated with either phorbol myristate acetate or opsonized zymosan particles generated a stable species capable of oxidizing 5-thio-2-nitrobenzoic acid or iodide. Resting cells did not form this species. The oxidant formed by the stimulated neutrophils was identified as taurine chloramine by both ultraviolet spectrophotometry and electrophoresis. Taurine chloramine formation by the neutrophil was dependent on the taurine concentration, time, and cell number. Neutrophil-dependent chloramine generation was inhibited by catalase, the myeloperoxidase inhibitors, azide, cyanide, or aminotriazole and by chloride-free conditions, but not by superoxide dismutase or hydroxyl radical scavengers. Thus, it appears that stimulated human neutrophils can utilize the hydrogen peroxide-myeloperoxidase-chloride system to generate taurine chloramine. Based on the demonstrated ability [...]

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EVIDENCE FOR HYPOCHLOROUS ACID GENERATION

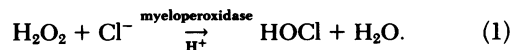
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ABSTRACT The model hydrogen peroxide-myeloperoxidase-chloride system is capable of generating the powerful oxidant hypochlorous acid, which can be quantitated by trapping the generated species with the β -amino acid, taurine. The resultant stable product, taurine chloramine, can be quantitated by its ability to oxidize the sulfhydryl compound, 5-thio-2-nitrobenzoic acid to the disulfide, 5,5'-dithiobis(2-nitrobenzoic acid) or to oxidize iodide to iodine. Using this system, purified myeloperoxidase in the presence of chloride and taurine converted stoichiometric quantities of hydrogen peroxide to taurine chloramine. Chloramine generation was absolutely dependent on hydrogen peroxide, myeloperoxidase, and chloride and could be inhibited by catalase, myeloperoxidase inhibitors, or chloride-free conditions. In the presence of taurine, intact human neutrophils stimulated with either phorbol myristate acetate or opsonized zymosan particles generated a stable species capable of oxidizing 5-thio-2-nitrobenzoic acid or iodide. Resting cells did not form this species. The oxidant formed by the stimulated neutrophils was identified as taurine chloramine by both ultraviolet spectrophotometry and electrophoresis. Taurine chloramine formation by the neutrophil was dependent on the taurine concentration, time, and cell number. Neutrophil-dependent chloramine generation was inhibited by catalase, the myeloperoxidase inhibitors, azide, cyanide, or aminotriazole and by chloride-free conditions, but not by superoxide dismutase or hydroxyl radical scavengers. Thus, it appears that stimulated human neutrophils can utilize the hydrogen peroxide-myeloperoxidase-chloride system to generate taurine chloramine. Based on the demonstrated ability of the myeloperoxidase

system to generate free hypochlorous acid we conclude that neutrophils chlorinate taurine by producing this powerful oxidant. The biologic reactivity and cytotoxic potential of hypochlorous acid and its chloramine derivatives suggest that these oxidants play an important role in the inflammatory response and host defense.

INTRODUCTION

Human neutrophils can utilize the H_2O_2 -myeloperoxidase-halide system to generate a powerful oxidant capable of destroying a variety of microorganisms and mammalian cell targets (1-4). Chloride, bromide, iodide, or thiocyanate may all be effective substrates in the myeloperoxidase system but, based on their *in vivo* concentrations, Cl^- is thought to be the preferred halide (1, 4). In the presence of Cl^- , the final mediator of myeloperoxidase-dependent events has been debated but increasing evidence indicates that hypochlorous acid (HOCl) is the responsible toxic species (5-10). In 1976, Harrison and Schultz (10) demonstrated that a model system consisting of H_2O_2 , purified myeloperoxidase, and Cl^- generated HOCl (Eq. 1).



Unlike other peroxidases that may oxidize acceptor molecules via the formation of a chlorinating intermediate (11), myeloperoxidase preferentially catalyzes the peroxidation of the chloride ion to free HOCl (10).

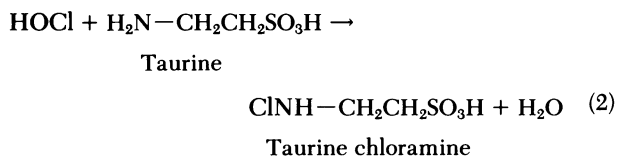
Recently, we have demonstrated that human neutrophils stimulated with phorbol myristate acetate (PMA)¹ destroyed cell targets by a myeloperoxidase-dependent process that could be inhibited by a variety of compounds known to scavenge HOCl (8, 9). Al-

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¹Abbreviations used in this paper: DTNB, 5-5'-dithiobis(2-nitrobenzoic acid); PMA, phorbol myristate acetate; TNB, 5-thio-2-nitrobenzoic acid.

though these data indicated that intact neutrophils have the potential to generate a species with characteristics similar, if not identical, with HOCl, the rates and quantities of the chlorinating species produced are unknown. Attempts to directly measure the generation of chloride peroxidation products by intact cells are hampered by the short half-life of this oxidant in biological systems containing multiple potential substrates. To quantitatively detect chlorinating species generated by neutrophils, we have taken advantage of the ability of HOCl to rapidly react with the β -amino acid taurine to form the stable oxidant, taurine chloramine (12-14) (Eq. 2).



In this manner we have examined the ability of intact, human neutrophils to utilize the H_2O_2 -myeloperoxidase- Cl^- system to generate a chloride peroxidation product capable of oxidizing taurine to its chloramine.

METHODS

Cell preparation. Neutrophils were obtained from the venous blood of healthy volunteers. Neutrophil preparations were isolated by Ficoll-Hypaque density centrifugation followed by dextran sedimentation (15). The preparations contained >96% neutrophils and 2-3% eosinophils. Cells were suspended in Dulbecco's phosphate-buffered saline (Gibco Laboratories, Grand Island Biological Co., Grand Island, NY; pH 7.4 with 1 mg/ml glucose). In some experiments Dulbecco's buffer was replaced with a 0.1 M NaCl-0.029 M sodium phosphate buffer with 1.5 mM MgSO_4 (pH 7.4) or a halide-free 0.1 M phosphate buffer with 1.5 mM MgSO_4 (16).

Assay systems

Myeloperoxidase system. Varying concentrations of H_2O_2 were incubated alone or with purified myeloperoxidase in the presence or absence of taurine (Sigma Chemical Co., St. Louis, MO) in Dulbecco's buffer (pH 7.4) in final volume of 1 ml at 37°C. In some experiments the Dulbecco's buffer was replaced with the chloride or chloride-free phosphate buffer. The H_2O_2 concentration was calculated from the absorbance at 230 nm assuming an extinction coefficient of $81 \text{ M}^{-1} \text{ cm}^{-1}$ (17). Purified canine myeloperoxidase was prepared (18) and kindly provided by Dr. J. Schultz (Papanicolaou Cancer Research Institute, Miami, FL). Myeloperoxidase activity was assayed by the *o*-dianisidine method (19) and activity expressed in international units. At the end of the incubation period, 50 μg of catalase (type C-40, 12,000 U/mg, Sigma Chemical Co.) were added to reduce residual H_2O_2 and duplicate samples assayed for taurine chloramine formation.

Neutrophil system. Neutrophils were incubated with varying concentrations of taurine for indicated periods of time at 37°C in the described buffers (pH 7.4) in a final volume of 1 ml. Neutrophils were stimulated to generate oxygen metabolites and release lysosomal enzymes by the

addition of PMA (8, 9) (Consolidated Midland Corp., Fortsetter, NY) or opsonized zymosan particles (Sigma Chemical Co.) prepared as previously described (15). At the end of the incubation period, catalase (50 μg) was added and duplicate samples assayed for taurine chloramine formation as described below.

Other additions to the myeloperoxidase or neutrophil systems included bovine superoxide dismutase (2,700 U/mg, Sigma Chemical Co.), bovine catalase (80,000 U/mg, Worthington Biochemical Corp., Freehold, NJ), sodium azide, sodium cyanide (Fisher Scientific Co., Pittsburgh, PA), ethanol, mannitol or 3-amino-1,2,4-triazole (aminotriazole, Sigma Chemical Co.). Superoxide dismutase was assayed according to the method of McCord and Fridovich (20). Catalase was washed over an XM-100A ultrafiltration membrane (Amicon Corp., Lexington, MA) and assayed before use (19).

Taurine chloramine quantitation

The ability of HOCl to react with taurine was confirmed by adding a known amount of reagent NaOCl (Fisher Scientific Co.) to taurine. The OCl^- concentration was calculated from its absorbance at 292 nm at pH 12.0 assuming an extinction coefficient of $350 \text{ M}^{-1} \text{ cm}^{-1}$ (21). Fig. 1 shows the UV spectrum of a solution of OCl^-/HOCl ($\text{pK}_a \sim 7.4$)

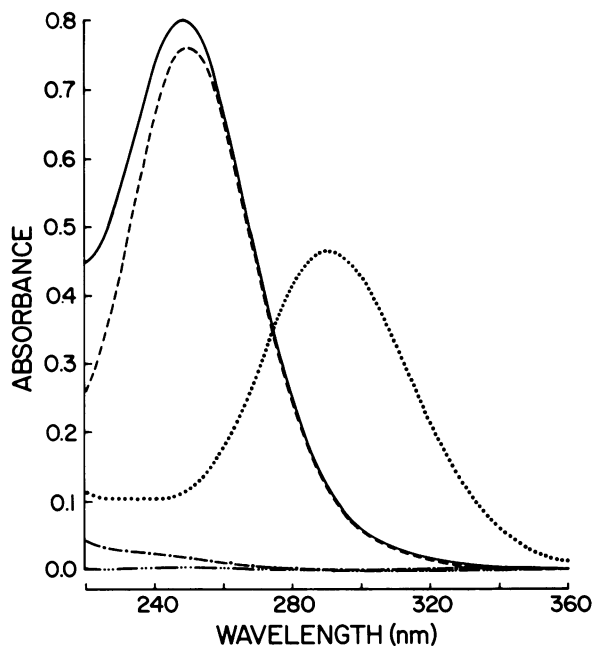


FIGURE 1 The UV absorption spectrum of synthesized taurine chloramine. Samples were incubated in Dulbecco's buffer (pH 7.4) for 5 min at 25°C in a final volume of 1.5 ml and the spectra recorded. After the samples were scanned, 3.5 μmol of H_2O_2 in a volume of 10 μl were added to the OCl^-/HOCl or the taurine- OCl^-/HOCl mixtures, incubated for 5 min at 25°C and rescanned. (a) 15 μmol of taurine alone (.....); (b) 3.0 μmol of a solution of OCl^-/HOCl alone (.....); (c) 15 μmol of taurine and 3.0 μmol of OCl^-/HOCl (----); (d) OCl^-/HOCl plus H_2O_2 (-.-.-.-); (e) taurine chloramine plus H_2O_2 (—).

(21) in Dulbecco's buffer (pH 7.4). Taurine does not absorb in this region. If a solution of OCl^-/HOCl (referred to as HOCl) was added to an excess of taurine a new compound was formed (Fig. 1). The resultant spectrum with a λ maximum of ~ 250 nm is characteristic of monochloramines (12, 13). Under these conditions, dichloramine formation (λ maximum 300 nm) was not detected (14). To rule out the formation of a mixture of taurine chloramine and HOCl, H_2O_2 was added to the system. H_2O_2 can reduce HOCl but does not react with taurine chloramine (7). As expected, the addition of H_2O_2 abolished the HOCl spectrum but had no effect on the taurine chloramine concentration (Fig. 1). The slight increase in the absorbance of the H_2O_2 -HOCl or H_2O_2 -taurine chloramine spectrum is due to the absorbance of H_2O_2 alone. The taurine chloramine spectrum did not change after a 4-h incubation at 37°C . Thus, under our conditions, HOCl can react with taurine to form the stable chloramine. Although taurine chloramine can be quantitated directly by its absorbance at 250 nm, the system is not sufficiently sensitive for the neutrophil studies because of the low extinction coefficient of chloramine ($\sim 398 \text{ M}^{-1} \text{ cm}^{-1}$) (14).

Because 1 mol taurine chloramine retains 2 mol oxidizing equivalents (7), its concentration can be sensitively determined by the ability to oxidize 2 mol of the sulfhydryl compound 5-thio-2-nitrobenzoic acid (TNB) to 1 mol of the disulfide, 5,5'-dithiobis(2-nitrobenzoic acid) (DTNB) or 2 mol of I^- to I_2 (7, 14). TNB was prepared by reducing DTNB (Sigma Chemical Co.) (7) and its concentration determined by its absorbance at 412 nm assuming an extinction coefficient of $1.36 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1}$ (22). The I_2 concentration was determined spectrophotometrically in the presence of excess I^- as I_3^- ($\epsilon = 2.29 \times 10^4 \text{ M}^{-1} \text{ cm}^{-1}$) (23).

In experiments with the model myeloperoxidase system or intact neutrophils, samples were assayed for taurine chloramine formation at the end of the incubation period by adding either TNB (0.45 mM) or KI (20 mM) to the entire mixture. The samples were immediately centrifuged (500 g for 5 min) and the absorbance recorded at 412 or 350 nm, respectively. In selected neutrophil experiments, the supernatant and cell button were analyzed separately for chloramine content. Catalase was added to all systems before the addition of TNB or KI due to the ability of high concentrations of H_2O_2 to slowly oxidize TNB or I^- . The addition of catalase did not interfere with either of the assays.

High voltage paper electrophoresis

In selected experiments the presence of taurine chloramine formation was examined by high voltage paper electrophoresis. Supernates were removed from the test mixtures and spotted on Whatman No. 3MM paper (Whatman Ltd., England). Electrophoresis runs were carried out for 30 min in a borate buffer (pH 9.0) at 3,000 V and a current of 150 mA. Taurine was visualized with ninhydrin spray (Sigma Chemical Co.) and the taurine chloramine was visualized with a 2% KI solution in acetone (13).

Hydrogen peroxide generation

H_2O_2 generation was determined in the presence of 1 mM azide by the method of Thurman et al. (24).

RESULTS

Taurine chloramine generation by the H_2O_2 -myeloperoxidase- Cl^- system. Under ideal conditions, the

addition of known amounts of H_2O_2 to purified myeloperoxidase and Cl^- should result in the formation of an equimolar concentration of HOCl. If the experiment is performed in the presence of an excess of taurine, then the HOCl should be trapped as the chloramine. Fig. 2 illustrates the relationship between H_2O_2 added and taurine chloramine detected in a cell-free myeloperoxidase system. Under the conditions used, $\sim 100\%$ of the H_2O_2 was converted to chloramine. The presence of taurine chloramine was confirmed by the UV absorption spectrum of the sample. As described in Methods, the chloramine had an absorption maximum at ~ 250 nm and was resistant to H_2O_2 (data not shown). The amount of taurine chloramine formed was also dependent on the taurine concentration. As little as 0.1 mM taurine (100 nmol/ml) scavenged almost all of the HOCl (data not shown). If 100 nmol of H_2O_2 were added to myeloperoxidase and Cl^- in the absence of taurine, incubated for 60 min and then taurine added, <5 nmol of chloramine could be detected. Although the enzyme does not require taurine to generate HOCl, myeloperoxidase is rapidly inactivated by this oxidant (6, 10) and any residual HOCl would be reduced by H_2O_2 (7). Thus, it appears that the myeloperoxidase system can utilize H_2O_2 to generate HOCl which, in the presence of taurine, leads to the formation of chloramine.

Requirements for taurine chloramine formation by the model myeloperoxidase system. Elimination of H_2O_2 , myeloperoxidase, or Cl^- completely inhibited the formation of taurine chloramine (Table I). As ex-

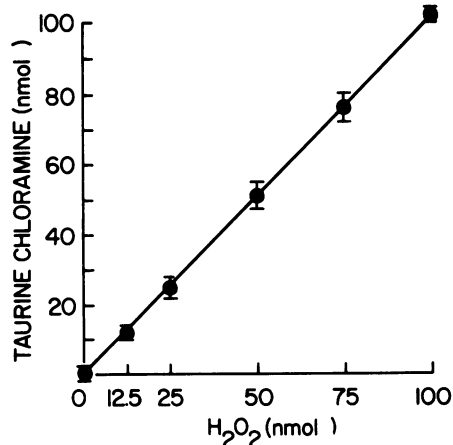


FIGURE 2 The formation of taurine chloramine by a model H_2O_2 -myeloperoxidase- Cl^- system. Known amounts of H_2O_2 were added to 16 mU of purified myeloperoxidase in Dulbecco's buffer (pH 7.4) in a final volume of 1 ml in the presence of 5 mM taurine. Samples were incubated for 60 min at 37°C and assayed by the TNB method. Results are expressed as the mean ± 1 SD of four experiments.

TABLE I
Taurine Chloramine Generation by the H_2O_2 -
Myeloperoxidase- Cl^- System

Additive*	Taurine chloramine formed nmol†
$H_2O_2 + Cl^-$	0.0±0.0
$H_2O_2 +$ myeloperoxidase (Cl^- free)	0.0±0.0
Complete system	102.8±1.4
Complete system + catalase (50 μ g/ml)	4.6±2.1
Complete system + heat-inactivated catalase (50 μ g/ml)	101.8±1.2
Complete system + superoxide dismutase (10 μ g/ml)	101.3±1.5
Complete system + azide (1 mM)	2.7±0.6
Complete system + cyanide (1 mM)	5.0±1.6
Complete system + aminotriazole (10 mM)	0.0±0.0
Complete system + ethanol (40 mM)	100.9±2.7
Complete system + mannitol (40 mM)	99.0±4.6

* The complete system consisted of 100 nmol H_2O_2 , 16 mU of myeloperoxidase, and 5 mM taurine incubated for 60 min in Dulbecco's phosphate-buffered saline at 37°C in a final volume of 1 ml. In the Cl^- free experiments a 0.1 M phosphate buffer was used. The 0.1 M NaCl-phosphate buffer gave results identical to those obtained with the Dulbecco's buffer.

† The taurine chloramine concentration was determined as described by the TNB method. Results are expressed at the mean±1 SD of four experiments.

pected, the addition of catalase (degrades H_2O_2) markedly inhibited chloramine formation, while neither superoxide dismutase (dismutates $O_2^{\cdot -}$ to H_2O_2) nor heat-inactivated catalase had any effect (Table 1). Azide, cyanide, and aminotriazole, three classic myeloperoxidase inhibitors, all blocked taurine chloramine formation, and the OH^{\cdot} scavengers, ethanol or mannitol, were without an inhibitory effect. Thus, as predicted, taurine chloramine formation was dependent on H_2O_2 , functional myeloperoxidase, and Cl^- .

Taurine chloramine formation by intact neutrophils. If intact neutrophils can utilize the H_2O_2 -myeloperoxidase- Cl^- system to generate HOCl or a species of similar reactivity we would predict that, in the presence of taurine, stimulated neutrophils should produce taurine chloramine. Fig. 3 demonstrates that a mixture of 2.5×10^6 PMA-stimulated neutrophils incubated with taurine for 2 h accumulated a species capable of oxidizing TNB to DTNB. Resting cells did not oxidize any of the TNB added. Addition of a reducing agent (cysteine) resulted in the recovery of >98% of the added TNB. The presence of taurine chloramine was confirmed in scaled up experiments by UV spectrophotometry. Fig. 4 illustrates the typical absorption maximum of taurine chloramine in the supernatant of

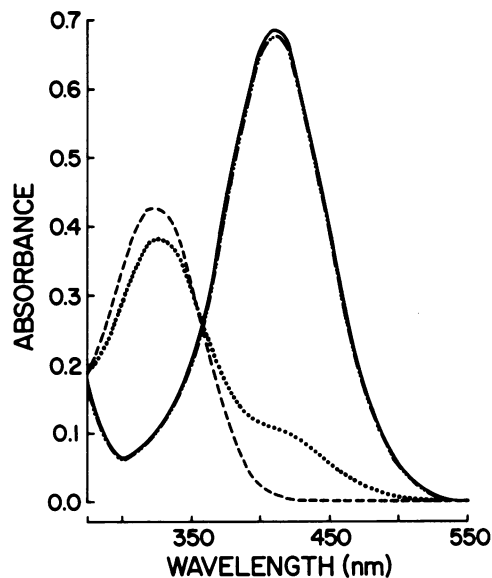


FIGURE 3 Optical spectra of TNB oxidized by PMA-stimulated neutrophils in the presence of taurine. 2.5×10^6 PMA-stimulated neutrophils were incubated with 15 mM taurine for 2 h at 37°C in 1 ml of Dulbecco's buffer (pH 7.4). At the end of the incubation period catalase was added followed by TNB (450 nmol). The mixture was centrifuged, diluted, and immediately scanned. After the spectrum was recorded, a small amount of solid cysteine was added (to reduce DTNB to TNB) and the sample scanned. (a) 225 nmol of DTNB diluted as above, λ maximum \sim 325 nm (-----); (b) 450 nmol of TNB, λ maximum 412 nm (—); (c) 450 nmol of TNB incubated with cell mixture ($\cdot \cdot \cdot \cdot \cdot$); (d) TNB-cell mixture following addition of excess cysteine (- $\cdot \cdot \cdot$ -). Note that the cells stimulated in the presence of taurine contained a species that decreased the absorption peak of the TNB at 412 nm and increased the absorption peak at 325 nm suggesting a mixture of DTNB and TNB. The addition of cysteine to this mixture ablated the 325-nm peak and regenerated the 412-nm maximum.

20×10^6 PMA-stimulated neutrophils after a 2-h incubation in the presence of taurine. No peak was observed in this region with resting cells (Fig. 4). As expected, the addition of H_2O_2 did not reduce the chloramine (data not shown). Identical results were obtained with opsonized zymosan-stimulated neutrophils. The presence of taurine chloramine was also reinforced by high voltage paper electrophoresis. In this system, taurine standards migrated 5.5 cm, while synthesized taurine chloramine migrated 17.0 cm (toward the anode), and could be visualized by its ability to oxidize I^- to I_2 . Electrophoresis of supernates obtained from PMA- or zymosan-stimulated neutrophils revealed an I^- reactive material that comigrated with synthesized taurine chloramine (i.e., 17 cm toward the anode). Thus, we conclude that intact, stimulated neutrophils can generate taurine chloramine.

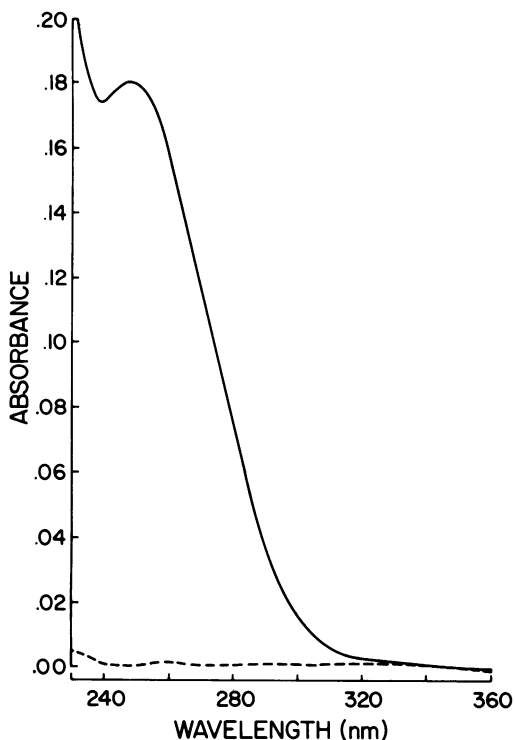


FIGURE 4 Optical spectra of oxidant generated by PMA-stimulated neutrophils in the presence of taurine. 20×10^6 neutrophils were incubated with or without 15 mM taurine in the presence or absence of PMA (100 ng/ml) for 2 h at 37°C in 2 ml of Dulbecco's buffer (pH 7.4). At the end of the incubation period the samples were centrifuged, diluted, and scanned. The reference cuvettes contained resting or stimulated cells in the absence of taurine. (a) Resting cells vs. resting cells with taurine (-----) and (b) stimulated cells vs. stimulated cells with taurine (—). Stimulated cells released material that absorbed strongly below 240 nm masking the descending shoulder of the chloramine.

Quantitation of taurine chloramine generation by neutrophils. Both PMA- or zymosan-stimulated neutrophils generated increasing amounts of taurine chloramine as the taurine concentration was increased until a plateau was reached at ~ 15 mM taurine (Fig. 5). The dose of PMA (30 ng/ml) and zymosan (1.25 mg/ml) used gave maximal results. Resting cells did not generate detectable amounts of taurine chloramine. In addition, 15 mM taurine did not depress the neutrophils' respiratory burst as assessed by O_2 consumption, $\text{O}_2^{\cdot -}$ generation (data not shown), or by H_2O_2 release from azide. 2.5×10^5 PMA- or zymosan-stimulated neutrophils in the presence of 1 mM azide generated 39.7 ± 4.6 and 32.0 ± 8.3 nmol $\text{H}_2\text{O}_2/\text{h}$ alone while producing 43.0 ± 3.7 and 32.0 ± 10.0 nmol/h, respectively, in the presence of 15 mM taurine (mean \pm 1 SD, $n = 3$).

Taurine chloramine generation was dependent on

the length of the incubation period and was essentially complete after a 2-h incubation (Fig. 6A). The relationship between neutrophils number and taurine chloramine formation is illustrated in Fig. 6B. In 15 experiments 2×10^6 PMA-stimulated neutrophils generated 126.4 ± 13.0 nmol chloramine during a 2-h incubation, while zymosan-stimulated cells produced 108.0 ± 11.1 (mean \pm 1 SD). In three paired experiments, chloramine concentrations determined by the TNB method varied $<5\%$ from those obtained with the KI technique. PMA-stimulated cells generated 129.7 ± 11.1 nmol of taurine chloramine by the TNB method and 123.5 ± 9.7 nmol with the KI method, while zymosan-stimulated cells formed 119.1 ± 8.5 nmol and 116.7 ± 8.7 nmol by the TNB and KI methods, respectively. Although chloramine determinations were routinely made on the entire incubation mixture, separate analysis of the supernatant and the cell button revealed that $\sim 10\%$ of the oxidant remained associated with the neutrophil.

These results indicate that neutrophils can produce significant amounts of taurine chloramine, but the quantitation assumes that the generated chloramine is stable in the complex cell system. To assess the stability of taurine chloramine in the neutrophil system, we examined our ability to recover 50 nmol of exogenous taurine chloramine from mixtures of both resting and PMA- or zymosan-stimulated cells. After a 2-h incubation, $>95\%$ of the exogenous taurine chloramine could be recovered from either the resting or stimulated cell mixtures. Thus, this quantitative technique appears to accurately reflect the total amount of taurine chloramine generated.

Requirements for taurine chloramine formation by neutrophils. If taurine chloramine generation by the stimulated neutrophil is mediated by HOCl or a similar species, we would expect a requirement for H_2O_2 , myeloperoxidase, and Cl^- . As shown in Table II, catalase significantly inhibited taurine chloramine generation by 2×10^6 PMA- or zymosan-stimulated neutrophils. The inhibitory effect of catalase increased to 90 and 65% with 1×10^6 PMA- or zymosan-stimulated cells, respectively (PMA control 53.8 ± 9.3 ; $n = 5$, zymosan control 53.7 ± 10.4 ; $n = 3$). Heat-inactivated catalase had no inhibitory effect with either stimulus (Table II). The addition of superoxide dismutase did not inhibit chloramine generation by the neutrophils with either stimulus. The heme-enzyme inhibitors, azide, cyanide, and aminotriazole were each capable of inhibiting chloramine formation, while neither ethanol nor mannitol had any inhibitory effect (Table II).

Because endogenous myeloperoxidase was apparently required for chloramine formation, we examined the effect of exogenous peroxidase in the neutrophil

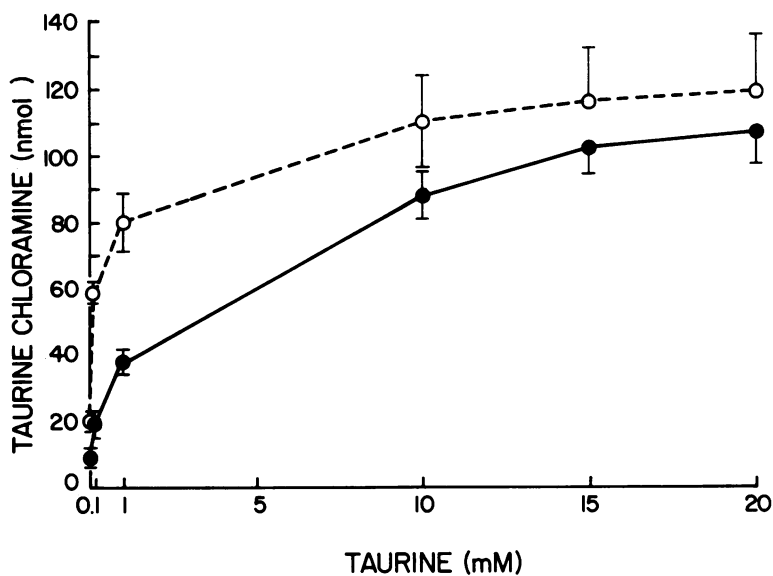


FIGURE 5 The effect of taurine concentration on the formation of taurine chloramine by neutrophils. 2×10^6 neutrophils were incubated with varying concentrations of taurine for 2 h at 37°C in 1 ml of Dulbecco's buffer (pH 7.4). Neutrophils were incubated in the presence of 30 ng/ml of PMA (-○-○-) or 1.25 mg/ml of opsonized zymosan (-●-●-●-). Taurine formation was quantitated by the TNB method. Results are expressed as the mean \pm 1 SD of three experiments.

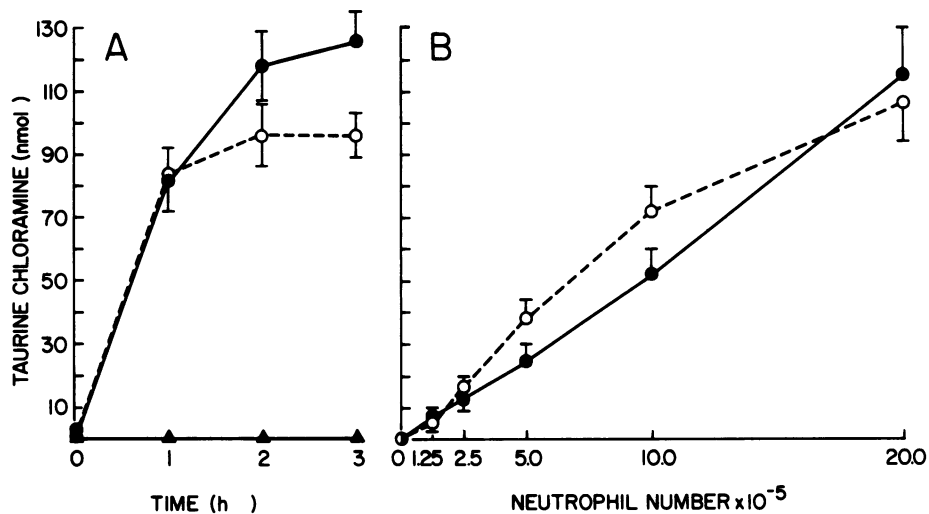


FIGURE 6(A) Taurine chloramine formation by neutrophils as a function of time. 2×10^6 neutrophils were incubated with 15 mM taurine for varying periods of time at 37°C in 1 ml of Dulbecco's buffer (pH 7.4). Neutrophils were incubated in the absence of a stimulus (-▲-▲-) and in the presence of 30 ng/ml of PMA (-○-○-) or 1.25 mg/ml of opsonized zymosan (-●-●-●-). Taurine chloramine formation was quantitated by the TNB method. Results are expressed as the mean \pm 1 SD of five experiments. (B) Taurine chloramine formation as a function of neutrophil number. Varying numbers of neutrophils were incubated with 15 mM taurine for 2 h at 37°C in 1 ml of Dulbecco's buffer (pH 7.4). Neutrophils were stimulated with either 30 ng/ml of PMA (-○-○-) or 1.25 mg/ml of opsonized zymosan particles (-●-●-●-). Taurine chloramine formation was quantitated by the TNB method. Results are expressed as the mean \pm 1 SD of five experiments.

TABLE II
Characteristics of Taurine Chloramine Generation
by Neutrophils

Additive*	Percentage of control†	
	PMA	Zymosan
Catalase (40 µg/ml)	16.7±2.7	51.2±3.8
Heat-inactivated catalase (40 µg/ml)	100.0	96.5
Superoxide dismutase (10 µg/ml)	105.8±3.4	113.6±11.7
Azide‡ (0.1 mM)	3.4±4.4	2.6±4.2
Azide (10 µM)	3.0±3.6	6.3±4.0
Azide (1 µM)	37.7±1.5	76.4±7.5
Azide (0.1 µM)	87.7±5.9	97.3±3.1
Cyanide§ (0.5 mM)	1.2±1.1	3.4±4.8
Cyanide (0.1 mM)	20.3±6.7	33.7±4.5
Cyanide (10 µM)	95.3±4.2	98.0±3.5
Aminotriazole (10.0 mM)	0.0±0.0	2.8±3.4
Ethanol (40 mM)	106.1±4.8	93.4±4.0
Mannitol (40 mM)	103.9±7.2	105.4±5.0
Chloride-free system	8.0	8.7
Chloride-free + H ₂ O ₂ (100 nmol)	8.4	7.2
Chloride-free + myeloperoxidase (16 mU)	10.0	3.5
Chloride-free + H ₂ O ₂ + myeloperoxidase	12.0	7.6

* The complete system consisted of 2×10^6 neutrophils incubated with 15 mM taurine in Dulbecco's buffer (pH 7.4) in a final volume of 1 ml at 37°C for 2 h. The cells were stimulated with either PMA (30 ng/ml) or zymosan (1.25 mg/ml). Neutrophils were preincubated with azide, cyanide, or aminotriazole for 5 min at room temperature before addition of the stimulus. The chloride-free experiments were performed in the 0.1 M sodium phosphate buffer with MgSO₄ (pH 7.4).

† Results are expressed as the mean percentage of control \pm 1 SD of six paired experiments with catalase, superoxide dismutase, azide, cyanide and aminotriazole. Heat-inactivated catalase was studied in two-paired experiments. The control taurine chloramine generation for PMA- and zymosan-stimulated cells were 115.1±15.3 and 98.2±20.4, respectively. The results for the Cl⁻-free experiments are expressed as the mean of two-paired experiments. The controls were performed in a 0.1 M NaCl-phosphate buffer and the values for 2×10^6 PMA- and zymosan-stimulated neutrophils were 122.0 and 85.5 nmol taurine chloramine, respectively.

‡ If the azide concentration was increased to 1 mM, the TNB oxidation was inhibited only 50%. Further studies revealed that in the presence of 1 mM azide, high concentrations of H₂O₂ were formed capable of mediating TNB or I⁻ oxidation. In contrast, higher concentrations of cyanide did not increase TNB oxidation. However, cyanide along with other nucleophiles can directly reduce DTNB (25). In this system the addition of \leq 1 mM cyanide reduced $<$ 5 nmol of DTNB.

system. PMA is a poor stimulus for myeloperoxidase release (26, 27) and exogenous enzyme might be expected to enhance chloramine generation. Indeed, in three experiments the addition of 16 mU of purified myeloperoxidase enhanced chloramine generation by 2×10^6 PMA-stimulated neutrophils from 124.0±8.0 nmol to 178.0±4.0 nmol during a 2-h incubation. Zymosan-stimulated neutrophils were not affected by the addition of myeloperoxidase (119.1±3.5 vs. 116.7±3.7 nmol taurine chloramine; $n = 3$).

A possible requirement for Cl⁻ was examined by performing the neutrophil experiments in a Cl⁻-free buffer (Table II). In the absence of Cl⁻ chloramine generation was significantly inhibited. Although the functional activity of the neutrophil in a Cl⁻-free system is questionable, neither the addition of exogenous H₂O₂ nor myeloperoxidase alone or in combination stimulated chloramine generation. Thus, it appears that both PMA- or zymosan-stimulated neutrophils utilize the H₂O₂-myeloperoxidase-Cl⁻ system to generate taurine chloramine.

DISCUSSION

Neutrophils can generate H₂O₂ and release myeloperoxidase into a Cl⁻ containing environment to form a powerful oxidizing agent capable of attacking microorganisms, mammalian cells (1, 4), and a variety of soluble inflammatory mediators (for recent examples, see references 28 and 29). In the presence of H₂O₂ and Cl⁻, purified myeloperoxidase can continuously generate free HOCl via a mechanism that does not require the presence of organic chloride acceptors (10). Apparently, the enzyme system does not directly chlorinate substrates nor does it generate other chloride oxidation products. In solution, HOCl may be in equilibrium with other oxychloro compounds, but HOCl and its anion OCl⁻ are the predominant species under physiologic conditions (30).

Both HOCl and OCl⁻ are powerful oxidants capable of reacting with amines, amino acids, sulfhydryls, disulfides, thioethers, aromatics, and a variety of potentially important biological substrates (30-32). Because of this reactivity, steady-state concentrations of HOCl and OCl⁻ would be difficult to detect in complex biological systems. In this study, we chose taurine as a model compound capable of reacting with HOCl/OCl⁻ at diffusion controlled rates to form a stable and quantifiable chloramine derivative (12-14). The applicability of this technique was initially examined in a model H₂O₂-myeloperoxidase-Cl⁻ system wherein H₂O₂ was stoichiometrically oxidized to a product capable of chlorinating taurine. Based on the known ability of the myeloperoxidase system to generate free

HOCl (10), it would seem that this oxidant was responsible for taurine chloramine formation. Thus, it appeared that taurine might allow us to trap and quantify chloride peroxidation products generated by the intact neutrophil.

In the presence of taurine, PMA- or zymosan-stimulated neutrophils generated large quantities of a species capable of oxidizing TNB to DTNB or I^- to I_2 . Both UV spectrophotometry and high voltage paper electrophoresis helped confirm the formation of taurine chloramine. Although monochloramines can be oxidized to dichloramines (13, 14, 30), we found no evidence for the generation of this oxidant by the neutrophil in the presence of a large excess of taurine. These data indicate that neutrophils can form an oxidant capable of attacking the amine group of taurine to form the N-Cl derivative.

Chloramine generation by the neutrophils plateaued as the concentration of exogenous taurine reached 15 mM. The optimal concentration of taurine was significantly higher than that used in the model myeloperoxidase system and probably reflects the increased number of potential substrates in the more complex neutrophil system. Similarly, at low taurine concentrations, increased amounts of chloramine were detected with PMA as the stimulant as opposed to zymosan particles. Higher concentrations of taurine might be required to scavenge the oxidant with zymosan because the chloride peroxidation product can react with the opsonized particle.² Alternatively, zymosan is internalized and high taurine concentrations could favor an increased intravacuolar concentration of the scavenger. It is interesting to note that even in the absence of exogenous taurine, small amounts of a stable oxidant could be detected with either stimulant (Fig. 5). Whether this represents the formation of an endogenous chloramine or some other stable oxidant remains to be determined.

Analysis of the oxidant distribution in the supernatant and cell button revealed that the majority of the chloramine was in the extracellular milieu, but we cannot be certain that this necessarily reflects extracellular chloramine generation. Either the chloride peroxidation products or the chloramine itself might be generated within the vacuole and then diffuse out of the cell. At present, the amounts of taurine available within the vacuolar space to scavenge the oxidant are unknown. Alternatively, neutrophils do release H_2O_2 and myeloperoxidase extracellularly (1, 4), and we might be detecting only those chloramines generated outside the cell. In other studies, myeloperoxidase-

mediated iodination products generated by neutrophils are also preferentially localized in the extracellular rather than the intracellular environment (1, 16).

Studies designed to determine the requirements for taurine chloramine generation by the neutrophil clearly indicated a pivotal role for H_2O_2 , myeloperoxidase, and Cl^- . First, the ability of the cell to generate the chloramine was inhibited in the presence of catalase. Although catalase inhibited taurine chloramine generation by either PMA- or zymosan-stimulated cells, it was more effective in the PMA system. PMA-stimulated cells release large amounts of oxygen metabolites extracellularly but only small quantities of myeloperoxidase (26, 27), while zymosan-stimulated neutrophils generate oxygen metabolites and release large amounts of myeloperoxidase into the phagocytic vacuole. Thus, in the presence of PMA, catalase may not have to gain access into the small vacuolar space to inhibit chloramine formation and, in addition, would be competing for H_2O_2 with a lower concentration of myeloperoxidase. Second, a role for myeloperoxidase was underlined by the ability of azide, cyanide, and aminotriazole to inhibit taurine chloramine generation. Although cyanide and aminotriazole can directly inhibit myeloperoxidase, they can also react with HOCl (30). It should be noted that a variety of peroxidase inhibitors seem to mediate their effects by interacting with the activated halide rather than directly with the enzyme (33, 34). Finally, a Cl^- requirement was based on the inhibitory effects of Cl^- -free conditions on chloramine formation. These results are not surprising since the oxidant being measured is itself a chlorinated derivative. Although neutrophils incubated in Cl^- -free buffers may have altered functions, neither exogenous H_2O_2 nor myeloperoxidase stimulated chloramine generation. Thus, it appears that intact neutrophils can utilize the H_2O_2 -myeloperoxidase- Cl^- system to generate an oxidant capable of chlorinating taurine. Based on the demonstrated ability of the myeloperoxidase system to peroxidize Cl^- to HOCl, it would seem that this is the most likely species responsible for chloramine formation. Although this conclusion is indirect, it should be reemphasized that myeloperoxidase does not directly chlorinate organic Cl^- acceptors and has not been demonstrated to generate any other chloride oxidation products (10, 11).

HOCl is a powerful oxidant capable of reacting with a host of biologically important molecules (30–32). We have previously shown that a human tumor cell line is sensitive to low concentrations of HOCl (8) and Thomas (17) has demonstrated its strong bactericidal potential. Indeed, neutrophils are capable of incorporating chloride into insoluble fractions of phagocytosed bacteria (35). The reactivity of HOCl also suggests

² S. J. Weiss. Unpublished observations.

potential control mechanisms for limiting its production. HOCl generation by neutrophils would be dependent on the quantity of H₂O₂ produced, the rate of H₂O₂ catabolism by endogenous catalase or the glutathione redox system, the amount of myeloperoxidase released, and the availability of potential substrates for HOCl. If sufficient myeloperoxidase is released but there are no available substrates for HOCl, either myeloperoxidase (6, 10) or possibly NADPH oxidase itself (36) could be rapidly inactivated. In our system, 2 × 10⁶ zymosan-stimulated neutrophils generated ~260 nmol of H₂O₂ (based on values obtained for 2.5 × 10⁵ cells) compared with ~100 nmol of taurine chloramine. We conclude that ~40% of the detected H₂O₂ was utilized by the myeloperoxidase system to generate HOCl. Although the utilization of H₂O₂ by myeloperoxidase may be maximized in our system via the maintenance of high concentrations of taurine, chloramines are oxidants that may be capable of inactivating myeloperoxidase (37). In the presence of an acceptor molecule that does not form an oxidant, even more effective utilization of H₂O₂ by myeloperoxidase might be attained.

In this study, taurine was used to trap and quantitate HOCl, but these results also suggest that neutrophils may use amines present at inflammatory sites to generate stable chloramine derivatives. Chloramines can oxidize sulfhydryl containing compounds, thioethers,³ iodide, and may slowly hydrolyze to regenerate HOCl (38). Indeed, Thomas (38) has recently demonstrated the bactericidal potential of a variety of chloramines. Thus, neutrophils could also utilize the myeloperoxidase system to generate long-lived oxidants capable of mediating inflammatory effects long after the termination of the respiratory burst. Based on the demonstrated reactivity of HOCl and its chloramine derivatives, these neutrophil-derived oxidants should play an important role in host defense and the inflammatory response.

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