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Research Article

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The Effect of Tuftsin on the Nitrous Blue Tetrazolium Reduction of Normal Human Polymorphonuclear Leukocytes

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ABSTRACT The influence of Tuftsin, the synthetic phagocytosis-stimulating tetrapeptide (L-threonyl-L-lysyl-L-prolyl-L-arginine), on the nitrous blue tetrazolium (NBT) reduction by human polymorphonuclear leukocytes was investigated. It was found that this substance increases the NBT reduction by approximately as much as endotoxin. Other tetrapeptides do not share this property. When Tuftsin analogs are added to the cell suspension and incubated, they prevent the action of both Tuftsin and endotoxin but not of methylene blue. When washed of the analogs, the cells regain the property to be activated by both Tuftsin and endotoxin. It appears that methylene blue on one hand and Tuftsin and endotoxin on the other hand have different sites for their actions. We suggest that whereas methylene blue diffuses into the cell and acts directly upon the hexosemonophosphate shunt activation, Tuftsin and endotoxin appear to act on the cell membrane binding to specific receptors. By treating the cells with Tuftsin analogs, we probably block these receptors.

INTRODUCTION

Fidalgo and Najjar demonstrated in 1967 (1) the presence in dog sera of a specific "leukophilic" gamma globulin that can bind to autologous leukocytes and stimulates their phagocytic activity against pathogenic *Staphylococcus aureus*. Several reports have since appeared indicating the importance of this fraction in man. It was named leukokinin, and its action was well documented (2-4).

Najjar and Nishioka (3) showed that the phagocytosis-stimulating effect of leukokinin is due to a single peptide fragment, liberated from the globulin by leuko-

kinase, a specific enzyme of the outer membrane of the neutrophil. This peptide was isolated, and its amino acid sequence was shown to be L-threonyl-L-lysyl-L-prolyl-L-arginine. It was named Tuftsin (3). Recently it was synthesized (5), and the biological action of this tetrapeptide was ascertained. Since the sera of splenectomized men (6) and dogs (7, 8) lack the stimulating effect of Tuftsin, it has been suggested that the spleen is the site of its production. Tuftsin acts in hormone-like quantities. In addition to enhancing particle ingestion, such as bacteria and latex, it also stimulates pinocytosis of radiolabeled albumin and enhances the motility and migration of polymorphonuclear leukocytes (PMN)¹ (5).

Published reports on Tuftsin have dealt with its effect on the phagocytic activity of the leukocyte. No reference whatever has been made to its possible influence on the bactericidal activity of the leukocyte. As reduction of nitrous blue tetrazolium (NBT) (9-11) is a well-known method of study of at least one important aspect of this bactericidal activity, we have investigated the effect of Tuftsin upon this process in human PMN.

METHODS

Preparation of Tuftsin and analog. Tuftsin (L-threonyl-L-lysyl-L-prolyl-L-arginine) was synthesized by two different methods²: in steps with dicyclohexylcarbodiimide as a coupling agent (12), and in steps with the polymeric-reagents approach (13). Both preparations were found to be identical and pure when analyzed by techniques such as elemental analysis, amino acid analysis, electrophoresis, and chromatography with various solvent systems. Both preparations showed identical biological activity.

Des⁴-threonyl-Tuftsin, alanyl⁴-Tuftsin, and W-nitro-Tuftsin were prepared by the polymeric reagents approach (13).

¹ Abbreviations used in this paper: analog 1, DES⁴ threonyl-Tuftsin; analog 2, alanyl⁴ Tuftsin; NBT, nitrous blue tetrazolium; PMN, polymorphonuclear leukocytes.

² Detailed synthetic procedures will be published elsewhere.

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Oligolysines (kindly donated by Dr. A. Yaron) were prepared by the solid-phase peptide approach (14).

The quantitative NBT reduction test. 25-ml samples of venous blood, donated by healthy young medical students, were collected in 30-ml disposable plastic tubes containing 50 U heparin/ml (Organon Inc., West Orange, N. J.). All samples were processed immediately after the venipuncture. 5 ml of 6% dextran (Fluka A. G., Basel, Switzerland) solution were added to each tube, and the mixture was allowed to settle for 1 h at room temperature. The next steps were strictly as described by Bachner and Nathan (9) with the following modifications:

Experiment 1. To test tubes with 0.1 ml of a leukocyte suspension (approximately 2.5×10^6 leukocytes, with a viability index of at least 95%) we added instead of latex particles one of the following materials: 20 μg of endotoxin (Difco Laboratories, Detroit, Mich.) (B_4 lipopolysaccharide) (0.2 ml); 0.1 μg of Tuftsin (0.1 ml); 0.1 μg of Tuftsin NO_2 (0.1 ml); 0.1 μg of analog 1 (0.1 ml); 0.1 μg of analog 2 (0.1 ml); 0.1 μg lysine₄ (0.1 ml); 0.22 mmol of methylene blue (BDH Chemicals Ltd., Poole, England); 20 μg of endotoxin (0.2 ml) + 0.1 μg Tuftsin (0.1 ml). All the above substances were dissolved in sterile phosphate-buffered saline, pH 7.4. After addition of 0.1 ml of 0.1% NBT (Sigma Chemical Co., Inc., St. Louis, Mo.) to each tube, the mixtures were incubated for 30 min at 37°C. The reaction was stopped by addition of 10 ml 0.5 N HCl and centrifuged. The reduced NBT (formazan) was extracted in 3 ml pyridine (BDH Chemicals Ltd.) under a boiling water bath. The color intensity of the pyridine was determined in a Unicam 800 spectrophotometer (Unicam Instruments Ltd., Cambridge, England). Results were expressed as ΔOD of reduced NBT in 2.5×10^6 cells/30 min (calculated as the difference in OD between stimulated tubes and blanks). All reagents were prepared under strictly sterile conditions with Millipore filters (Millipore Corp., Bedford, Mass.) to avoid any particulate matter that might stimulate phagocytosis.

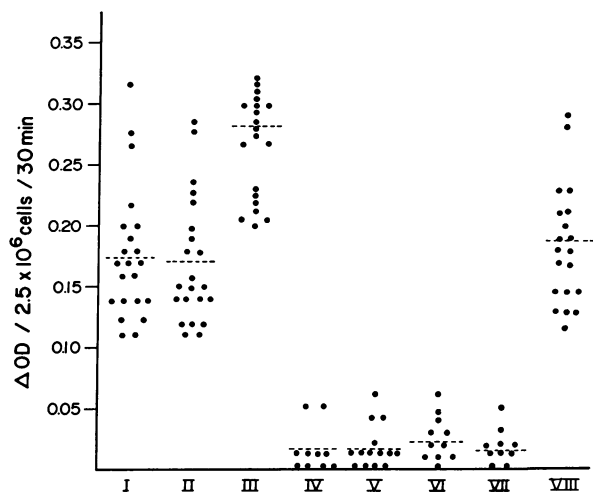


FIGURE 1 Differences in optical density (ΔOD) between stimulating and resting NBT reduction tests with the following stimulators: I, Tuftsin; II, endotoxin; III, methylene blue; IV, oligolysin; V, Tuftsin NO_2 ; VI, analog 1; VII, analog 2; VIII, endotoxin plus Tuftsin. The dotted lines indicate the mean values for each experimental protocol.

TABLE I

Results of Experiment 2

Pre-incubated	Stimulus	No. of tests	ΔOD		
			Mean	SD	Range
Analog 1	Tuftsin	11	0.030	0.021	0.00 - 0.050
	Endotoxin	10	0.027	0.024	0.00 - 0.060
	Methylene blue	8	0.25	0.06	0.15 - 0.37
Analog 2	Tuftsin	11	0.028	0.020	0.00 - 0.05
	Endotoxin	10	0.031	0.022	0.00 - 0.05
	Methylene blue	8	0.25	0.05	0.15 - 0.37

ΔOD , difference in optical density between NBT reduced by resting and stimulated cells preincubated with analog 1 and analog 2.

Experiment 2. The same cell suspensions as in experiment 1 were preincubated for 5 min at room temperature with 0.1 μg of Tuftsin analogs 1 and 2, respectively. To the test tubes we added separately: 0.1 μg Tuftsin; 20 μg endotoxin; 0.22 mmol methylene blue and 0.1 ml of 0.1% NBT. The ΔOD in experiment 2 was calculated as in experiment 1.

Experiment 3. Cell suspensions, incubated as above with Tuftsin analogs 1 and 2, were washed three times with a Krebs-Ringer solution. Tuftsin (0.1 μg) and endotoxin (20 μg), respectively, and NBT were then added to the suspension, and ΔOD was calculated.

RESULTS

Fig. 1 demonstrates the distribution of ΔOD calculated from experiment 1. It shows clearly that Tuftsin acts in all suspensions of live leukocytes to the same degree as endotoxin, 0.177 ± 0.055 and 0.171 ± 0.053 , respectively, and less than methylene blue (0.28 ± 0.062). Tuftsin NO_2 , Lysine, analog 1, and analog 2 act weakly (low ΔOD) on the stimulation of the NBT reduction and have a similar range of values: 0.014, 0.016, 0.023, and 0.016, respectively, without statistical difference between them. Simultaneous addition of endotoxin and Tuftsin gives results as for Tuftsin alone, 0.180 ± 0.050 .

Table I summarizes the results of experiment 2. It shows that Tuftsin analogs 1 and 2, when added to the cell suspension and incubated, prevent the increase of the NBT reduction by both Tuftsin and endotoxin. They do not prevent the activation by methylene blue. There is no difference between these analogs in this inhibitory effect on the NBT reduction.

Experiment 3 revealed that when washed of the analogs, the cells regain the capacity to be activated by both Tuftsin and endotoxin (values similar to in experiment 1, ΔOD 0.175 and 0.170, respectively).

DISCUSSION

The increase in the percentage of NBT-reducing leukocytes and of the total reduced dye is due to the activation of the hexosemonophosphate shunt (15). This can be induced theoretically either by increased phagocytosis, as in latex-stimulated cells (9), or by such direct activators as methylene blue, which acts without enhancing the phagocytic action (16).

We have shown that Tuftsin acts like endotoxin. The effects of both can be prevented by blocking agents such as the Tuftsin analogs. When these are removed, the cells recover their reactivity to both Tuftsin and endotoxin. The effect of methylene blue, on the other hand, is not blocked by the analogs. The effect of blocking was previously demonstrated by Najjar and Constantopoulos (6), using a synthetic Tuftsin analog and a natural pathological inactive Tuftsin molecule found in a familial disorder of Tuftsin production.

Recent observations indicate that NBT is phagocytized by the cell as a complex (with fibrinogen and/or heparin) (17). Whether Tuftsin acts as a stimulator of this phagocytosis or in some unknown way as a direct activator of the hexosemonophosphate shunt is not indicated by our experiments.

The influence of Tuftsin on CO₂ production by the hexosemonophosphate shunt activation is now under our investigation. This study will, we hope, resolve these contradictions. But whatever is the mode of action, our present observations confirm Najjar's conclusion that the simple tetrapeptide is an active substance and binds, probably by specific receptors, to the membrane of humoral PMNs.

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