

The Role of Formylpeptide Receptors, C5a Receptors, and Cytosolic-Free Calcium in Neutrophil Priming

Werner Zimmerli, Anne-Marie Reber,
and Clemens A. Dahinden

From the Departments of Medicine and Research, University Hospital,
Basel, and the Institute for Clinical Immunology, University Hospital,
Bern, Switzerland

Polymorphonuclear leukocytes (PMNL) exposed to chemoattractants or cytokines change their functional capacity. The effect of endotoxin-activated serum as a priming agent on human PMNL was tested. Pretreatment of PMNL with endotoxin-activated serum increased their oxidative burst in response to formylpeptide (FMLP) ($P < .02$) and C5a ($P < .05$). Priming for membrane depolarization was observed in PMNL preincubated with either endotoxin-activated serum, low concentrations of purified C5a, or endotoxin but not with decomplexed plasma. Primed PMNL had an increased number of FMLP but not C5a receptors as compared with control PMNL. The "resting" cytosolic free calcium was increased in primed PMNL ($P < .02$). Intracellular calcium buffering abolished the priming effect of endotoxin-activated serum. Thus, endotoxin-activated serum can prime cellular responsiveness for membrane depolarization and superoxide production in response to FMLP and to C5a. Priming may be due to an increased resting cytosolic-free calcium.

Polymorphonuclear leukocytes (PMNL) are end-stage cells that play an important role in host defense and inflammation [1]. Upon exposure to chemoattractants, they are capable of improving their functional capacity [2]. Immature PMNL such as those from patients with myelodysplastic syndromes or leukemia have functional defects [3, 4]. In contrast, extravascular PMNL, that is, those in an advanced life stage, have an increased responsiveness to *N*-formylmethionylleucylphenylalanine (FMLP) compared to blood PMNL from the same species [2]. The exact mechanisms of this functional improvement (priming) are unknown but may at least be partially due to FMLP-receptor upregulation [2].

In this study we analyzed the effect of prestimulation of human PMNL *in vitro* on their subsequent responsiveness. The phenomenon of priming was investigated by incubating PMNL with either endotoxin-activated serum (EAS) or some of its components, namely endotoxin, C5a, and decomplexed plasma. EAS was used because it most closely simulates the clinical situation of gram-negative sepsis, during which PMNL may be primed. The importance of migration in the cellular priming was analyzed by using a chemotactic chamber as an *in vitro* model of locomotion. Cellular responsiveness of EAS-pretreated PMNL to FMLP and purified C5a was measured. The mechanisms of priming were analyzed in terms of FMLP-

and C5a-receptor studies. Because not every priming could be explained by receptor upregulation, we investigated whether an increased "resting" cytosolic-free calcium ($[Ca^{2+}]_i$) could be involved.

Materials and Methods

Reagents. Reagents used and their sources were: bovine serum albumin, cytochalasin B, cytochrome C (type IV), dimethyl sulfoxide, HEPES, FMLP, phenolphthalein glucuronic acid solution, phenolphthalein, phenolphthalein standard solution, phorbol myristate acetate (PMA), quin 2 acetoxymethylester (quin 2-AM), quin 2 free acid, and triton X-100 (Sigma Chemical, St. Louis); dextran T-500 and Percoll (Pharmacia Fine Chemicals, Uppsala, Sweden); endotoxin lipopolysaccharide-B *Escherichia coli* O26 (Difco Laboratories, Detroit); ^{57}Co -B12 and FMLP [3H]P (specific activity 60 Ci/mmol; New England Nuclear, Boston); EDTA and activated charcoal (Merck AG, Zurich); polymyxin B and ionomycin (Calbiochem, San Diego); 3,3'-dipentylloxycarbonyl iodide (di-O-C5(3); Molecular Probes, Junction City, OR), and Harleco Diff-Quik (Merz & Dade AG, Düringen, Switzerland). HEPES-Hanks' balanced salt solutions with (HBSS) or without calcium and magnesium (mHBSS) were prepared according to standard methods [5]. FMLP, PMA, and EAS were prepared as described by Metcalf et al. [5].

Isolation of PMNL. Human blood PMNL were purified from EDTA-anticoagulated (5 mM final concentration) blood, drawn by venipuncture from healthy human volunteers. Purification included dextran T-500 (3% in 0.9% saline) sedimentation and subsequent centrifugation (350 g for 30 min at 4°C) on a 53%/67% Percoll gradient, similar to that previously described [2, 6]. PMNL accumulating at the interphase of the gradient were washed twice in 0.9% saline. Hypotonic lysis was performed, if necessary, with distilled water and correction of the osmolarity with 1.8% NaCl after 15–30 s.

Pretreatment of PMNL *in vitro*. Purified PMNL (>98% pure) were suspended in HBSS supplemented with 0.1% glucose and 0.1% bovine serum albumin at a concentration of 1×10^7 PMNL/ml.

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Reprints and correspondence: Dr. Werner Zimmerli, Departments of Medicine and Research, University Hospital, CH-4031 Basel, Switzerland.

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The priming agents were added in different concentrations (see Results). Incubations were performed in Falcon plastic tubes (Becton Dickinson, Oxnard, CA) on an overhead rotator at 37°C, and terminated after 30 min by adding ice-cold mHBSS. Pretreated PMNL were washed twice with mHBSS and then resuspended at 10^7 PMNL/ml for use in the functional assays. Migrated PMNL (CTX-EAS-PMNL) were washed away from the lower part of a 10- μ m thick, 5- μ m pore diameter chemotaxis filter (Nucleopore, Pleasanton, CA), and from the lower wells of a chemotaxis microchamber (Neuro Probe, Cabin John, MD) containing 1.7% EAS. These PMNL, which migrated through the filter within 30 min, were washed twice before use in the functional assays. Control PMNL were incubated in HBSS/0.1% glucose/0.1% albumin alone.

Control experiments. Endotoxin modulates PMNL functions at low concentrations [7-11]. Therefore, any solution used for the purification and preincubation procedure was tested for its endotoxin contamination by the amoebocyte lysate assay (Pyrogel; Skan AG, Basel, Switzerland). H₂O, saline, EDTA, dextran, and C5a (10^{-9} M) contained <1 ng/ml endotoxin. HBSS-albumin and Percoll (53% and 67%) had an endotoxin contamination of <100 ng/ml. Since polymyxin B binds endotoxin, we compared PMNL preparations isolated in either the presence or absence of polymyxin B (10 μ g/ml) [12]. The priming effect of endotoxin (5 μ g/ml) in the superoxide assay was identical in the two different PMNL preparations (data not shown).

Membrane potential. Membrane potential was determined in a fluorescence spectrophotometer (LS5; Perkin-Elmer, Norwalk, CT) with the fluorescent probe di-O-C5(3) as previously described [13].

Superoxide production. Superoxide (O_2^-) production was determined at 37°C spectrophotometrically (550 nm, mM extinction coefficient of 21,100 $M^{-1} cm^{-1}$) by continuously monitoring the O_2^- -dismutase (300 units/ml)-inhibitable reduction of cytochrome-C (120 μ g/ml) in a double-beam spectrophotometer (model 35, Beckman Instruments, Fullerton, CA) with a temperature regulator set, as described previously [6].

C5a purification and labeling. C5a was purified from human yeast-activated plasma as described by Hugli et al. [14]. C5a was homogenous as shown by a single band on PAGE and on microzone electrophoresis at pH 8.6. The protein content was determined by amino acid analysis. Purified C5a was radiolabeled with ^{125}I using a solid-phase lactoperoxidase-glucose oxidase method according to Hugli and Chenoweth [15]. The initial specific activity was 160 Ci/mmol.

Receptor assays. FMLP [3H] binding to PMNL was carried out at 4°C for 20 min using a silicon-oil technique as described previously [2, 16]. For Scatchard analyses final concentrations of FMLP [3H] in binding assays varied from 3 to 200 nM. Nonspecific binding was determined with a parallel series of samples by adding a 1000-fold excess of unlabeled FMLP. Scatchard plots of the experimental data (specific bindings) were fitted by linear regression. The dissociation constant (K_d) and receptor number (B_{max}) were estimated from the fitted slopes and x intercepts [2]. ^{125}I -C5a binding to PMNL was carried out at 4°C using a silicon-oil technique, similar to that described by Huey and Hugli [17]. The incubation time was 45 min. For Scatchard analyses final concentrations of ^{125}I -C5a in binding assays were 0.08-20 nM. To determine the nonspecific binding, purified cold C5a was added to a parallel series of samples at a final concentration of 2.5 μ M. Nonspecific binding was routinely <10%.

The data were treated as described above. The specific activity of ^{125}I -C5a was determined for each separate assay.

Measurement of cytosolic-free calcium ($[Ca^{2+}]_i$). Control and EAS-PMNL were loaded with quin 2-AM as follows: Cells were incubated in HBSS supplemented with glucose (0.1%) at a concentration of 1×10^7 PMNL/ml with quin 2-AM at a final concentration of 1 μ M. Quin 2-AM was diluted 10-fold with 50% dimethyl sulfoxide/HBSS from a 2 mM stock solution in dimethyl sulfoxide before addition to PMNL. Quin 2 loading of PMNL was performed in a 30-min incubation with overhead rotation. Loading of 0.2-0.4 nmol of quin 2/ 10^6 PMNL was obtained. After loading, cells were washed twice and resuspended in glucose-supplemented HBSS. Cells were diluted to 2.5×10^6 PMNL/ml, maintained at room temperature, and protected from light until use. Control cells were incubated with an equivalent concentration of dimethyl sulfoxide and treated identically.

To determine the role of $[Ca^{2+}]_i$ in cellular priming, $[Ca^{2+}]_i$ was buffered with quin 2 in control experiments. For this purpose PMNL (1×10^7 /ml) were incubated 30 min in calcium-free HBSS/0.1% glucose with quin 2-AM at a final concentration of 10 μ M, which resulted in a loading of 2 nmol of quin 2/ 10^6 PMNL.

Fluorescent measurements were performed with a Perkin-Elmer fluorimeter (LS5; Perkin-Elmer) with two cutoff filters as described by Lew et al. [18]. The calculation of the loading and $[Ca^{2+}]_i$ was performed as described by Metcalf et al. [5].

$[Ca^{2+}]_i$ dependency of O_2^- production. To transiently increase $[Ca^{2+}]_i$, ionomycin (500 nM final concentration) was added to PMNL in calcium-free medium (mHBSS), as described by Lew et al. [19]. This increase was recorded with the quin 2 method. In concurrent experiments, FMLP-induced O_2^- production was measured in identically treated PMNL at different intervals after the addition of ionomycin.

Statistics. As an estimate of variance, standard deviation of the mean was used in the text and tables and standard error in the figures. Means were compared by Student's t test.

Results

Specific and primary granules in primed PMNL. Limited degranulation of specific granules is possibly one of the critical events in the cellular priming process. We therefore determined the content of specific and primary granules in PMNL preincubated either in HBSS/0.1% glucose/0.1% albumin or in 1.7% EAS or that had migrated through 5- μ m holes of a 10- μ m polycarbonate filter toward 1.7% EAS. Table 1 summarizes the results. Pretreatment of PMNL with 1.7% EAS and migration toward EAS induced a preferential loss of specific granules. The primary granule marker β -glucuronidase did not significantly differ in the three cell types.

Superoxide production. We next studied the effect of EAS preincubation on the subsequent O_2^- production in response to FMLP and C5a. O_2^- production was determined in the presence and absence of cytochalasin B, since pretreatment of PMNL with cytochalasin B by itself enhances and prolongs the respiratory burst induced by FMLP and C5a [20]. Figure 1 shows the O_2^- production rate in EAS-PMNL compared

Table 1. Granule content of untreated (control) and pretreated polymorphonuclear leukocytes (PMNL).

Cell type	B-12 binding protein (%)	<i>P</i> (paired <i>t</i> test)	β -glucuronidase (μ g phenolphthalein released/4 h/ 10^6 PMNL) (%)
Control PMNL	496 \pm 62 (100)	.002 } .001 }	174 \pm 21 (100)
EAS-PMNL*	429 \pm 62 (87 \pm 6.4)		170 \pm 16 (98 \pm 4.4)
EAS-CTX PMNL†	417 \pm 64 (84 \pm 6.2)		169 \pm 17 (97 \pm 4.1)

NOTE. Total granule content was determined by lysing the PMNL suspension (5×10^6 PMNL/ml) with triton X-100 (0.1%). Experiments were performed on six different days with PMNL from six different volunteers. All three PMNL types were prepared from the same cell batch. Results are mean \pm SD.

* PMNL were preincubated with endotoxin-activated serum (EAS).

† PMNL that migrated previously toward EAS through a 5- μ m-pore nitrocellulose filter.

with control cells. Interestingly, when PMNL were stimulated with 10^{-6} M FMLP, the priming effect of EAS was observed only in the absence of cytochalasin B. However, at a submaximal concentration of FMLP (10^{-7} M) priming was observed both in the absence (2.7 ± 0.2 vs. 1.7 ± 0.4 nmol of O_2^- /min/ 10^6 PMNL; $P < .05$) and in the presence (5 ± 1.0 vs. 3.3 ± 0.25 nmol of O_2^- /min/ 10^6 PMNL; $P < .05$) of cytochalasin B. The responsiveness of EAS-PMNL was also increased by stimulation with 10^{-7} M C5a (figure 1B), regardless of the presence of cytochalasin B.

By preincubating PMNL with a substimulatory concentration of C5a (10^{-9} M), their responsiveness toward FMLP

(10^{-6} M) was increased (5.0 ± 0.6 vs. 2.7 ± 0.6 nmol of O_2^- /min/ 10^6 PMNL; $P < .001$). In contrast, the responsiveness toward cytochalasin B plus C5a (10^{-7} M) decreased (1.1 ± 0.6 vs. 2.0 ± 1.2 nmol of O_2^- /min/ 10^6 PMNL; $P < .05$).

Effect of different priming factors on FMLP- and C5a-induced membrane depolarization. Using the membrane depolarization assay, previously shown to be a convenient test to detect priming [13], we analyzed which component of EAS might be responsible for PMNL priming and whether migration influenced the priming process. Figure 2A shows that the response of PMNL previously incubated with EAS, C5a, or endotoxin, but not decomplexed plasma, was similarly enhanced toward 10^{-6} M FMLP, as was the response of PMNL which had previously migrated through a chemotactic filter. Priming did not result in a modified dose-dependency response; that is, the FMLP concentration leading to half-maximal depolarization was similar between EAS- and control PMNL ($2 \times 10^{-8} \pm 1.4 \times 10^{-8}$ M vs. $2.4 \times 10^{-8} \pm 1.3 \times 10^{-8}$ M, respectively; $n = 8$ pairs).

Figure 2B shows the membrane depolarization of different PMNL preparations in response to 10^{-8} M C5a. With this stimulus, the priming effect of EAS (1.7%), C5a (10^{-9} M) or endotoxin (5 μ g/ml) was even higher, reaching between 170% (C5a) and 210% (EAS) above controls. However, the C5a concentration eliciting half maximal depolarization was somewhat higher in EAS-PMNL than in control cells ($4.3 \times 10^{-10} \pm 9.4 \times 10^{-11}$ M vs. $3.6 \times 10^{-10} \pm 9.4 \times 10^{-11}$ M, respectively; $n = 8$ pairs, $P < .05$). At a nondepolarizing concentration of C5a (10^{-10} M) in the preincubation, the priming for C5a (10^{-8} M)-induced membrane depolarization was in the same range, $159\% \pm 7\%$ of the control. Thus, in contrast to O_2^- production, pretreatment with a low concentration of a stimulus identical to that used in the restimulation led to priming and not to deactivation of the membrane depolarization.

Requirements for priming. Priming by EAS was a rapid process. Figure 3 shows that it was detectable at 5 min and reached a plateau within 20 min. The different requirements are summarized in table 2. Priming for FMLP-induced O_2^- production occurred at low concentrations of EAS; it was tem-

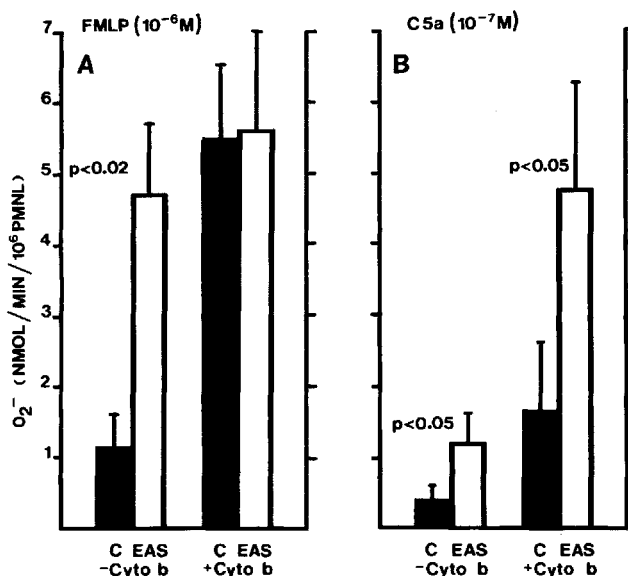


Figure 1. Superoxide production (O_2^-) of blood polymorphonuclear leukocytes (PMNL) incubated (37°C for 30 min) either in HBSS/0.1% glucose/0.1% albumin (C) or in 1.7% endotoxin-activated serum (EAS). PMNL were washed twice before testing O_2^- production. A, Maximal initial rate in response to 10^{-6} M formylpeptide (FMLP) in the presence or absence of cytochalasin B. B, Corresponding results in response to 10^{-7} M C5a. Statistical analysis was performed by paired *t* test. Results are means \pm SE of six different experiments.

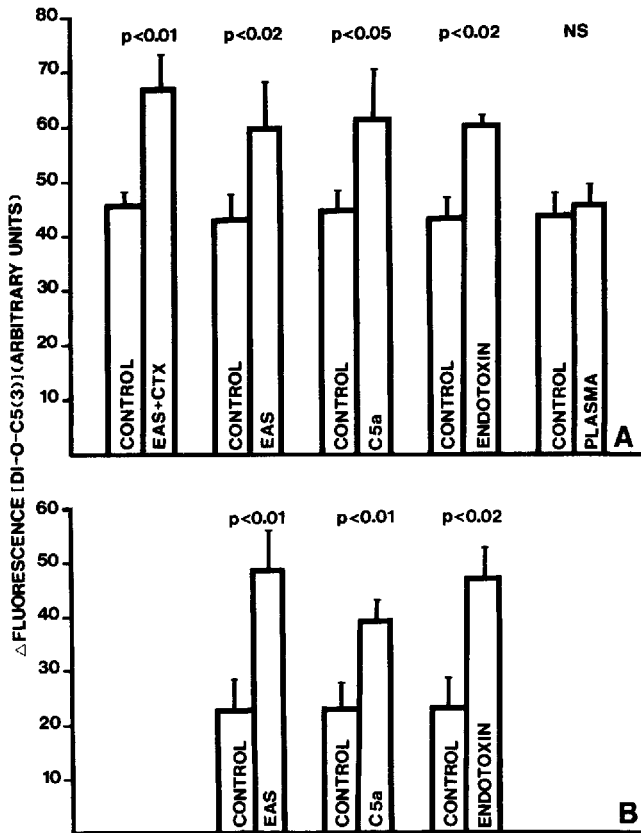


Figure 2. Effect of priming factors on 10^{-6} M formylpeptide (FMLP)- (A) and 10^{-8} M C5a- (B) induced membrane depolarization. Polymorphonuclear leukocytes (PMNL) were incubated for 30 min in either HBSS/0.1% glucose/0.1% albumin (control), 1.7% endotoxin-activated serum (EAS), purified C5a (10^{-9} M), endotoxin (5 μ g/ml), or decompemented plasma (1.7%). PMNL that had previously migrated through a chemotactic filter toward 1.7% EAS were labeled with EAS + CTX. PMNL were washed twice before testing membrane depolarization. Results are means \pm SE of 4–12 different experiments. Statistical analysis was performed by paired *t* test.

perature dependent and required neither protein synthesis nor extracellular calcium. However, no priming was observed when intracellular calcium was buffered with quin 2 before incubation of PMNL with EAS in calcium-free buffer.

Mechanisms of priming. To determine whether the observed limited degranulation of specific granules increased receptor number, FMLP and C5a receptors were determined. Figure 4 shows that the number of FMLP receptors was 1.4-fold higher in EAS-preincubated PMNL ($P < .05$) but this was not accompanied by a significant alteration in affinity. Preincubation with endotoxin alone did not significantly increase FMLP receptors: $22,000 \pm 5400$ per control PMNL and $26,400 \pm 11,100$ per endotoxin-primed PMNL ($n = 5$ pairs). Figure 5 shows that in cells, whether EAS primed or not, C5a receptors were similar in number and affinity under conditions in which the number of FMLP receptors was increased.

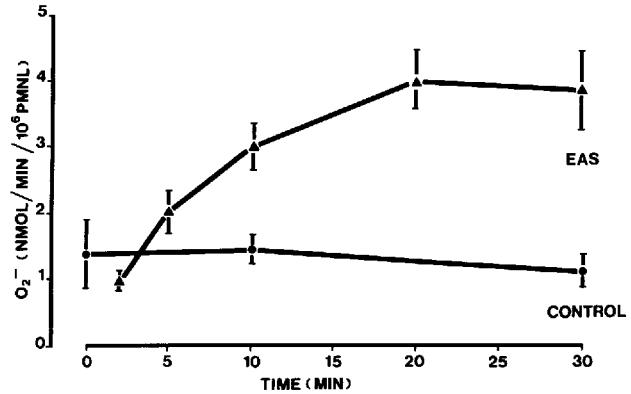


Figure 3. Influence of priming time on formylpeptide (FMLP)-induced superoxide (O_2^-) production. PMNL were incubated either in HBSS/0.1% glucose/0.1% albumin or in 1.7% endotoxin-activated serum. PMNL were diluted into ice-cold HBSS and washed twice at corresponding time points. Results are means \pm SE of triplicates from a representative experiment.

Table 2. Requirements for the priming process: O_2^- production rate in response to 10^{-6} M formyl peptide (FMLP) (nmol/min/ 10^6 PMNL).

Experimental conditions	EAS-PMNL*	Control-PMNL†	Significance‡
EAS-concentrations			
0.007%	2.3 ± 0.2	1.5 ± 0.3	<.025
0.33%	3.0 ± 0.8		
3.3%	3.2 ± 1.3		
Temperature			
0°C	1.7 ± 0.1	1.3 ± 0.2	NS
37°C	4.0 ± 0.2	1.1 ± 0.3	<.001
Protein synthesis			
- cycloheximide	2.8 ± 0.5	1.3 ± 0.3	<.001
+ cycloheximide (10 μ g/ml)	2.6 ± 0.4	ND	
Calcium			
1 mM Ca^{2+}	3.7 ± 1.7	1.3 ± 0.2	<.001
No Ca^{2+} /2 mM EGTA	3.9 ± 1.3	ND	
Quin 2-buffering (2 nmol quin 2/10^6 PMNL), no Ca^{2+}/2 mM EGTA			
	1.7 ± 0.2	1.7 ± 0.4	NS

NOTE. EAS = endotoxin-activated serum, PMNL = polymorphonuclear leukocytes. Each experiment was performed in triplicate at least three times.
 * Incubation for 30 min with 1.7% EAS if not otherwise stated.
 † Incubation for 30 min with HBSS/0.1% glucose/0.1% bovine serum albumin.
 ‡ Comparison between control and EAS-PMNL (paired *t* test); *P* value; NS = not significant.

As an alternative possible priming mechanism, we examined the intracellular calcium in primed and unprimed PMNL. Figure 6 shows that primed cells had a consistently higher “resting” $[Ca^{2+}]_i$ than control PMNL. This difference was significant ($P < .02$) with cells isolated from five different donors.

To investigate whether an increased $[Ca^{2+}]_i$ at the time of stimulation with an agonist can enhance the PMNL response, we measured the O_2^- production at different $[Ca^{2+}]_i$ concentrations. PMNL were pretreated with ionomycin to evaluate

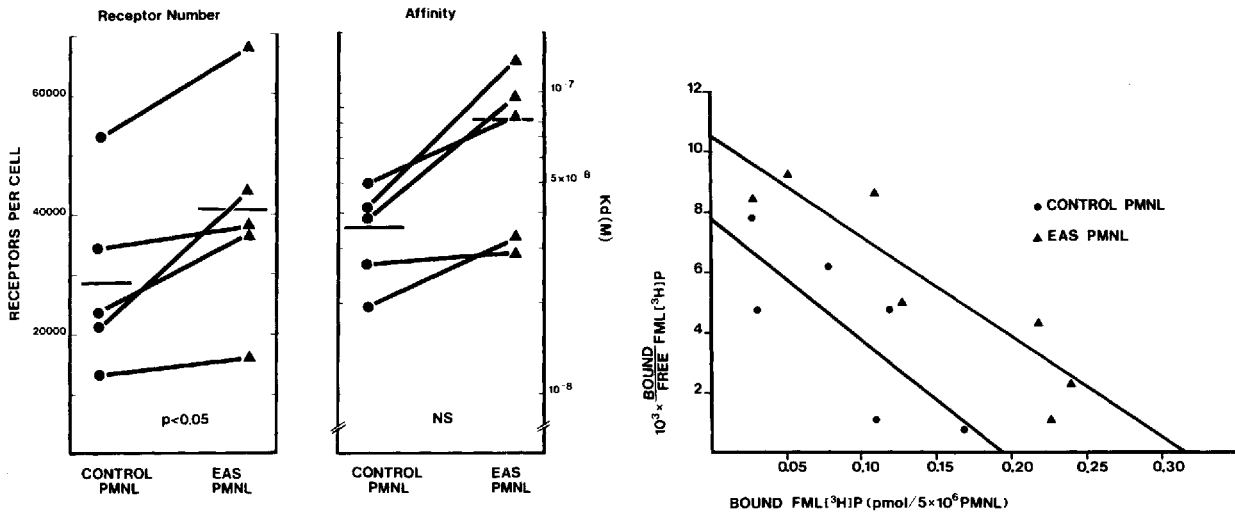


Figure 4. Total formylpeptide (FMLP) receptors per cell (left) and affinity constant (Kd) (middle). Polymorphonuclear leukocytes (PMNL) were preincubated and washed as described in Methods. Left and middle panels present results from Scatchard analyses performed on control and endotoxin-activated serum (EAS)-treated PMNL (pairs). Each pair represents an individual experiment. Significance of difference between means was determined by two-tailed Student's *t* test. Right panel shows Scatchard plot of representative experiment with control and EAS-incubated PMNL.

the [Ca²⁺]_i and thereafter stimulated with FMLP at different intervals and levels of [Ca²⁺]_i. Figure 7 shows a good relationship between the [Ca²⁺]_i at the time of stimulation and the O₂⁻ production rate that rose to values 2.2–4.2-fold above the baseline in five different experiments.

Discussion

It has been observed previously that PMNL modify their functional characteristics by sequential stimulation [16, 21–24].

These modifications are many and can be summarized as deactivation, adaptation, and priming. In our present study, we asked the following questions: Does EAS prime PMNL for FMLP- and C5a-induced functions? And, if so, which are the requirements for the EAS-induced priming? Also, what are the priming mechanisms?

By using highly purified C5a, we analyzed the effect of EAS prestimulation, even on cell functions that require high concentrations of the complement-derived chemotactic factor for stimulation. The use of C5a and FMLP allowed us to detect

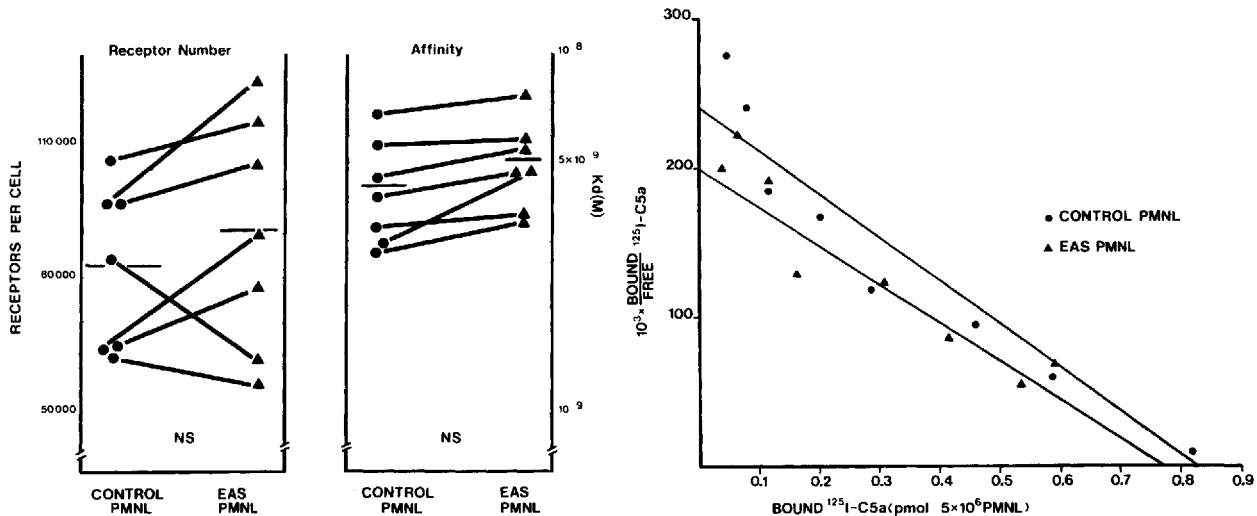


Figure 5. Total C5a receptors per cell (left) and affinity constant (Kd) (middle). Polymorphonuclear leukocytes (PMNL) were preincubated and washed as described in Methods. Left and middle panels present results from Scatchard analyses performed on control and endotoxin-activated serum (EAS)-treated PMNL (pairs). Each pair represents an individual experiment. Significance of difference between means was determined by two-tailed Student's *t* test. Right panel shows Scatchard plot of representative experiment with control and EAS-incubated PMNL.

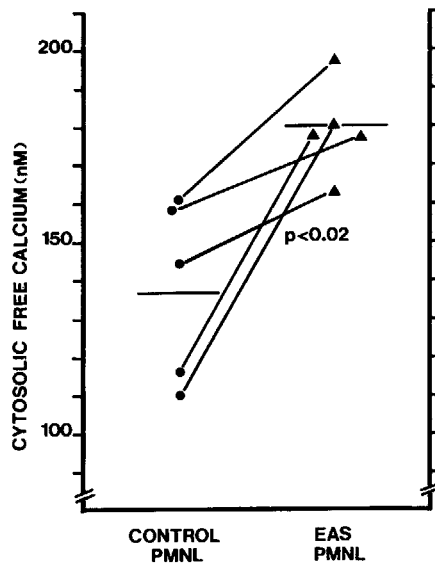


Figure 6. Resting cytosolic free calcium ($[Ca^{2+}]_i$) in control and endotoxin-activated serum (EAS)-treated polymorphonuclear leukocytes (PMNL) (pairs). PMNL were preincubated and washed as described in Methods. $[Ca^{2+}]_i$ was determined in resting cells by the quin 2 method. Significance of difference between means was determined by two-tailed *t* test.

possible discordant reactivity of EAS-PMNL, namely deactivation for one stimulus and priming for the other [2].

Control experiments confirmed that our PMNL preparations, isolated on a Percoll gradient, were not activated to a significant degree, since the priming was similar for PMNL prepared either in the presence or in the absence of polymyxin B. Further, significant priming with <100 ng/ml endotoxin occurs only after an incubation of >30 min [7].

EAS-pretreated PMNL had a lower residual content of specific granules than did control PMNL. This effect was even more pronounced in PMNL that had migrated through a filter in response to EAS. However, the enhancement of cellular responsiveness toward a second agonist was identical regardless of whether the cells had migrated, indicating that a larger degree of degranulation of secondary granules does not necessarily lead to a stronger priming.

In the presence of cytochalasin B, priming was observed only with a submaximal concentration of FMLP (10^{-7} M). This indicates that the respiratory burst was maximally stimulated by cytochalasin B and 10^{-6} M FMLP. PMNL preincubated with 10^{-9} M C5a showed a primed membrane depolarization but a deactivated O_2^- production in response to C5a. This discrepancy remains unexplained and argues against a causal relationship between changes in membrane potential and subsequent oxidative burst.

The mechanisms of priming remain unknown. Many mediators, including low concentrations of LTB_4 [25, 26], 5-HETE [27], platelet activating factor [26, 28], FMLP [21, 22, 24], and C5a (as shown here), plus cytokines such as granulocyte-

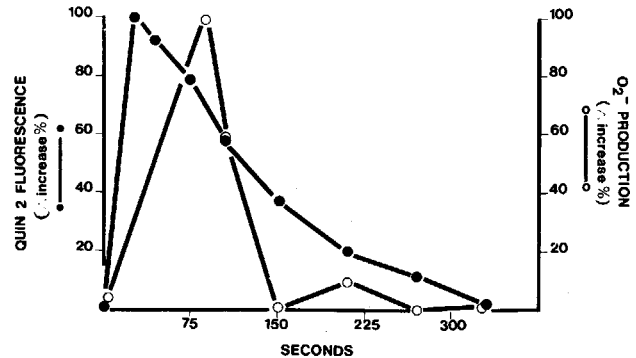


Figure 7. Representative experiment showing the dependency of superoxide (O_2^-) production on $[Ca^{2+}]_i$ in response to formylpeptide (FMLP). $[Ca^{2+}]_i$ was transiently increased by adding ionomycin ($500 \mu M$ final concentration) to polymorphonuclear leukocytes (PMNL) in calcium-free medium (mHBSS) [13]. Changes in $[Ca^{2+}]_i$ were recorded with the quin 2 method. In concurrent experiments, FMLP-induced O_2^- production was measured in identically treated PMNL at different intervals after the addition of ionomycin. Values of 100% represent the maximal increases for $[Ca^{2+}]_i$ and O_2^- production above the baseline value. Experiments with PMNL from four other donors gave similar results.

macrophage colony-stimulating factor [29–32], tumor necrosis factor- α [33], and endotoxin [10, 11, 28], have been shown to enhance the response triggered by a second agonist. Priming with EAS occurs rapidly (figure 3), whereas prolonged incubation is required for the action of lipopolysaccharide or cytokines [7, 10, 11, 28]. Thus, the mechanism of priming by EAS might differ from that by cytokines and endotoxin.

Most priming agents such as chemotactic factors induce a limited degranulation of secondary granules and mobilize FMLP and C3b_i receptors. It has been suggested that FMLP-receptor upregulation might be responsible for the enhanced responsiveness of PMNL [2, 16, 34]. We found an increase of FMLP receptors after preincubation of PMNL with EAS (figure 4) but not with endotoxin. Further, preincubation of PMNL with EAS did not result in an upregulation of C5a receptors. Since we demonstrated that priming does not result in an increased number or affinity of C5a receptors but still enhances a response triggered by C5a, mechanisms other than receptor upregulation must be operative in the priming process.

Forehand et al. [35] recently offered an explanation for the priming of PMNL with lipopolysaccharide. They found a 1.6-fold increase of the resting $[Ca^{2+}]_i$ in lipopolysaccharide-treated PMNL. We found a similar increase of the resting $[Ca^{2+}]_i$ in PMNL primed with EAS. Finkel et al. [36] showed that increasing the $[Ca^{2+}]_i$ of PMNL with ionomycin potentiated their oxidative burst in response to different stimuli. The positive association between the O_2^- production rate and the $[Ca^{2+}]_i$ at the time of stimulation with FMLP (figure 7) points out the relevance of the increased resting $[Ca^{2+}]_i$ as priming factor.

This proposed priming mechanism is compatible with the observed priming requirements (table 2). Since priming also occurred in the absence of extracellular calcium, the small increase of intracellular free calcium arises via mobilization from intracellular pools [37]. This is also supported by the lack of priming after buffering $[Ca^{2+}]_i$ with quin 2. A similar observation was made by Forehand et al. [35], who could abolish the endotoxin-induced priming by MAPTAM, another intracellular Ca^{2+} chelator.

In conclusion, we demonstrated that EAS primes PMNL for both FMLP- and C5a-induced function without affecting C5a receptor number and affinity. Therefore receptor upregulation cannot be the priming mechanism for every stimulus. EAS-treated PMNL had an increased resting $[Ca^{2+}]_i$, which may be responsible for their priming, since FMLP-induced O_2^- production was increased in PMNL with a higher $[Ca^{2+}]_i$ at time of stimulation and EAS did not prime PMNL in which $[Ca^{2+}]_i$ was buffered during preincubation.

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