

INTERACTION OF TARGET CELL-BOUND C3bi AND
C3d WITH HUMAN LYMPHOCYTE RECEPTORS
Enhancement of Antibody-mediated Cellular Cytotoxicity*

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In antibody-dependent, lymphocyte-mediated cytotoxicity (ADCC),¹ the interaction between IgG antibodies on the target cells and Fc IgG receptors (FcγR) on the effector cells affords close contact between the cells resulting in lysis of the targets. Present evidence suggests that the Fc-FcγR interaction triggers the cytolytic reaction. (1). It has been shown earlier that lymphoid effector cells with C3 receptors can bind C3-fragment-bearing erythrocytes as assessed by rosette formation. However, this interaction did not result in lysis of the target cells in absence of IgG antibody (2-4). In the presence of antibody to the target cells, target cell-bound C3 fragments strongly enhanced ADCC (1, 5). It was concluded that target cell-associated C3 amplifies ADCC by improving effector cell-target cell contact.

Cell-bound C3 may exist in several molecular forms, designated C3b, C3bi, and C3d (reviewed in [6]). C3 is deposited on particles as C3b which subsequently is cleaved by factor I to form C3bi without reduction of molecular weight. C3bi is then converted to C3d by enzymatic removal of the C3c fragment. The relative proportions of the three fragments on the surface of a target are determined by exposure to controlling enzymes.

The question of the role of these different C3 fragments in regulating cellular effector functions such as ADCC or phagocytosis is largely unresolved. Although

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¹ *Abbreviations used in this paper:* ADCC, antibody-dependent, lymphocyte-mediated cytotoxicity; complete TCM, Hepses (10 mM) -buffered RPMI-1640 containing 0.4% HSA and supplemented with 100 U penicillin, 100 μg Fungizone, and 2 mM L-glutamine; E_b, E_c, E_s, bovine, chicken, or sheep erythrocytes, respectively; E_bC3b, E_bC3bi, E_bC3d, C3b-fragment-, C3bi-fragment-, and C3d-fragment-bearing E_b, respectively; FcγR, Fc IgG receptor; FIP, Ficoll-Isopaque; GBV⁺⁺, veronal-buffered physiological saline, pH 7.4, containing 0.15 mM CaCl₂; HSA, human serum albumin; NF, nephritic factor; PFC, plaque-forming cells; PMN, polymorphonuclear leukocytes; TH, Hanks' balanced salt solution containing 15 mM Tris, pH 7.2; UV, ultraviolet.

previous studies suggested that target cell-bound C3b and C3d might enhance ADCC, no conclusions as to their relative importance could be drawn (3). Moreover, C3bi was not clearly defined at that time. This is the first description of a role for C3bi in lymphocyte function. The present study shows that cell-bound C3bi can directly interact with certain lymphocytes and that in the case of ADCC, this interaction greatly enhances lysis of target cells. The enhancing effect of C3bi is greater than that of C3d, and considerably greater than that of C3b.

Materials and Methods

Buffers. Buffers used were as follows: GVB⁺⁺: veronal-buffered physiological saline, pH 7.4, containing 0.15 mM CaCl₂, 0.5 mM MgCl₂, and 0.1% gelatin; and TH: Hanks' balanced salt solution containing 15 mM Tris, pH 7.2.

Purified Proteins. The proteins C3 (7), factor B (8), C3b inactivator (9), and β 1H (9) were prepared as described previously. Partially purified nephritic factor (NF) was prepared as described previously (10). 1 ml of NF was absorbed three times for 20 min at 0°C with 10⁹ erythrocytes and one time with 10⁹ erythrocytes bearing C3 fragments (EC3b). Bovine conglutinin was prepared by the method of Lachmann (11). Trypsin (TPCK) was purchased from Worthington Biochemical Corp. (Freehold, N. J.).

Erythrocytes. Bovine erythrocytes (E_b) and sheep erythrocytes (E_s) were purchased from Colorado Serum Co., Denver, Colo. and used within 2 wk. Chicken erythrocytes (E_c) were prepared from heparinized blood of 10–20-wk-old white Leghorns as previously described. (12).

Preparation of EC3b. EC3b were prepared by a modification of the procedure published elsewhere (13). 6 × 10⁹ washed erythrocytes were mixed with 800 μ g C3 and 1.5 μ g ¹²⁵I-C3 in a total vol of 500 μ l GVB⁺⁺. Deposition of C3b from the fluid phase onto the cell surface was effected by addition of 24 μ g trypsin and incubation for 2.5 min at 23°. Cells were washed 3 times with 10 ml GVB⁺⁺ and resuspended in 500 μ l GVB⁺⁺. Formation of cell-bound, alternative C3 convertase was accomplished by addition of 500 μ g factor B, 300 ng factor D, 30 μ g NF, and sufficient MgSO₄ to give a final concentration of 1.2 mM and incubation for 15 min at 37°C. Cells were washed two times in GVB⁺⁺ and suspended in 0.5 ml GVB⁺⁺ containing 800 μ g C3 and 1.5 μ g ¹²⁵I-C3. C3 deposition by the cell-bound C3 convertase was effected by incubation for 2.5 h at 37°C or 1 h at 37°C and 16 h at 4°C. Cells were washed three times into the appropriate buffer and standardized to 10⁹/ml. The number of C3b bound per cell was determined by assessing the radioactivity associated with a 50- μ l sample of cells. Erythrocytes bearing C3bi fragments (EC3bi) were produced by incubation for 45 min at 37°C of EC3b with purified C3b inactivator (8.0 μ g/ml) and β 1H (40.0 μ g/ml). Erythrocytes bearing C3d fragments (EC3d) were produced from EC3bi by incubation with 10 μ g/ml trypsin.

Testing of Cellular Intermediates. The functional state of C3 fragments on erythrocytes was ascertained using a series of assays. C3/C5 convertase formation was determined by the ability of erythrocytes to be lysed by incubation with factors B and D and guinea pig C3-9 as described (14). Immune adherence reactivity was determined in microtiter plates using human erythrocytes as published previously (15). Conglutinin-dependent hemagglutination was performed in microtiter plates containing 2.5 × 10⁶ erythrocytes, serial-dilutions of bovine conglutinin, and GVB⁺⁺ to a total vol of 75 μ l.

Antisera. Antisera to E_b or E_c were prepared in rabbits by six intravenous injections during a period of 10 d. IgG fractions of the antisera were prepared by gel-exclusion chromatography on Sephadex G-200 (Pharmacia Fine Chemicals, Uppsala, Sweden) (16). Anti-E_b reacted with E_b but not with E_c as determined by hemagglutination (titer: 21 μ g/ml) and ADCC-induction, whereas the reverse was true for anti-E_c. Both antibody preparations reacted with E_s.

Lymphocytes. Human peripheral blood lymphocytes were isolated and purified as previously described (12). In brief, after defibrination of the blood with glass beads, a leukocyte-enriched fraction was prepared by gelatin sedimentation. Phagocytes and adhering cells were removed by treatment with carbonyl iron powder, followed by centrifugation through Ficoll-Isopaque (FIP; Pharmacia Fine Chemicals). The lymphocytes, collected at the interface, were kept at 4°C overnight in Hepes (10 mM)-buffered RPMI-1640 (Grand Island Biological Co., Grand Island, N. Y.) containing 0.4% human serum albumin (HSA; Cutler Laboratories, Berkeley,

Calif.) and supplemented with 100 U penicillin, 100 $\mu\text{g/ml}$ Fungizone (Grand Island Biological Co.), and 2 mM L-glutamine (complete TCM). They were used the next day after centrifugation and suspension in fresh complete TCM. HSA was used as protein supplement throughout these experiments.

Fc γ R Modulation of Lymphocytes. 1.6×10^8 lymphocytes and 4×10^6 E_b or E_c were mixed in a 50-ml Falcon tube (Falcon Labware, Div. of Becton, Dickinson & Co., Oxnard, Calif.) in a total vol of 12 ml complete TCM in the presence of anti-E_b or anti-E_c (IgG fractions). The antibody concentrations were chosen to give 80–90% lysis of ⁵¹Cr-labeled erythrocytes when mixed with aliquots of the lymphocytes at the same ratios under ADCC conditions. After 20 h (or, in one experiment, 40 h) of incubation at 37°C, the cells were washed once in TH, and were then suspended in complete TCM at a concentration of 4×10^6 lymphocytes/ml. Control lymphocytes were mixed with erythrocytes and processed similarly but in the absence of antibody. When lymphocytes, modulated by exposure to antibody-coated erythrocytes, were mixed with fresh ⁵¹Cr-labeled target erythrocytes (E_a) under ADCC conditions but without addition of fresh antibody, no lysis ensued (cf. Table II), indicating that no antibody had been transferred with the lymphocytes in these experiments (17).

Fractionation of Lymphocytes by Adsorption to Monolayers of C3-fragment-bearing E_b. Tissue culture dishes (Falcon 3002; 60 \times 15 mm) were treated for 1 h with 2 ml TH containing 50 $\mu\text{g/ml}$ poly-L-lysine (70,000 mol wt; Sigma Chemical Co., St. Louis, Mo.). After washing, 2 ml TH containing 4×10^8 E_b bearing either C3b, C3bi, or C3d, were added to the dishes. The erythrocytes were allowed to settle at room temperature. After 40 min of incubation, nonadsorbed E_b were washed off with TH. 8×10^6 lymphocytes in 3 ml of complete TCM were added to the monolayers and incubated for 15 min at 37°C. The dishes were then centrifuged for 5 min at 100 g. After 15 min at 4°C, the nonadhering lymphocytes were carefully decanted with a Pasteur pipette and pooled with three additional washes (2 ml of TH wash per dish). The lymphocytes were then washed in TH, and finally suspended in complete TCM at a concentration of 4×10^6 living cells/ml. Recovery of lymphocytes from C3-free E_b control monolayers was ~100%.

Fractionation of Lymphocytes by Rosetting with C3-fragment-bearing E_b. 1.6×10^8 lymphocytes, suspended in 1 ml complete TCM, were mixed in conical 15-ml tubes (Falcon 2098) with 1 ml TH containing 4×10^8 E_b bearing either C3b, C3bi, or C3d (E_bC3b, E_bC3bi, and E_bC3d). After 5 min of incubation at 37°C, the tubes were centrifuged for 5 min at 100 g. After 1 h additional incubation at 4°C, the cell mixtures were gently suspended, diluted with 3 ml complete TCM, and layered over 1.5 ml FIP. After centrifugation for 20 min at 1,200 g, the nonrosetting lymphocytes were collected at the interface washed in TH and suspended in complete TCM at a concentration of 3×10^6 living cells/ml.

ADCC. ADCC of erythrocytes (E_b, E_s, or E_c, respectively) was assayed as described (12). 4×10^6 lymphocytes were mixed with 10^6 ⁵¹Cr-labeled erythrocytes, either untreated or bearing C3 fragments, in the presence or absence of anti-erythrocyte antibodies (IgG fractions). The total vol of the incubation mixture was 0.3 ml. All mixtures were prepared in duplicate and incubated for 18 h at 37°C. After centrifugation for 10 min at 800 g, the release of ⁵¹Cr was determined.

Rosette Formation with C3-Fragment-bearing E_b. 0.1 ml lymphocytes in complete TCM (4×10^6 lymphocytes/ml) were thoroughly mixed with 0.1 ml E_b, E_bC3b, E_bC3bi, or E_bC3d (10^9 /ml). The mixtures were incubated for 15 min at 37°C, centrifuged for 5 min at 100 g and held at 4°C overnight. Raji and Daudi cells were washed three times with TH and suspended in complete TCM at a concentration of 2×10^6 cells/ml. For rosetting, 0.1 ml of the cells was mixed with 0.1 ml C3-fragment-bearing E_b as described above. Drops of the cell mixtures were transferred to slides and covered with cover slips, coated with a 0.001% ethanol solution of acridine orange for nuclear staining of the lymphocytes (18). The percentage of rosetting cells was determined by counting 600–800 cells (two slides per sample) in the ultraviolet (UV) microscope under visible and UV light.

Results

Identification of Target Cell-bound C3 Fragments. Table I defines the characteristics of C3 fragments present on target cells. Cell-bound C3b was capable of producing

TABLE I
 Identification of C3 Fragments on Cellular Intermediate Complexes

Property	Intermediate complex		
	EC3b	EC3bi	EC3d
C3-associated $^{125}\text{I}^*$	100	100	10
C3/C5 convertase ‡	100	0	0
Immune adherence §	100	0	0
Conglutination §	0	100	0

* Data are represented as percentages of C3b radioactivity.

‡ Data are represented as percentages of lysis.

§ Data are represented as percentages of activity.

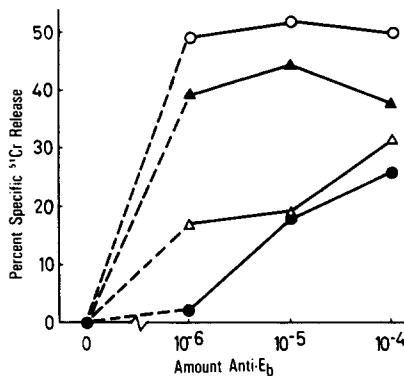


FIG. 1. Enhancement of ADCC by target cell-bound C3 fragments. ^{51}Cr -labeled E_b lacking C3 fragments (●) or bearing 45,000 C3b (△), C3bi (○), or C3d (▲) molecules per cell were incubated with lymphocytes and anti- E_b antibody (13 mg IgG/ml stock solution) for 18 h. at an E:T ratio of 4:1 as described in Materials and Methods. Spontaneous ^{51}Cr release was 8% for E_b and $\text{E}_b\text{C3b}$, and 13% for $\text{E}_b\text{C3bi}$ and $\text{E}_b\text{C3d}$. The release in the antibody-free lymphocyte controls was the same as the spontaneous release ($\pm 1\%$).

positive immune adherence reactivity with human erythrocytes and of forming the alternative pathway C3/C5 convertase with factors B and D and magnesium ions. Cells bearing C3bi were lacking the above functions, but exhibited the unique ability to be agglutinated by purified bovine conglutinin in the presence of calcium ions. Cells bearing C3d lacked all the above functions, but retained 10% of the radioactivity previously associated with C3b. A minimum of 100 C3b molecules or 500 C3bi molecules could be detected per cell by the functional tests.

Enhancement of ADCC by Target Cell-bound C3bi and C3d. To establish the effect of target cell-bound C3 fragments on cell-mediated lysis, $\text{E}_b\text{C3b}$, $\text{E}_b\text{C3bi}$, or $\text{E}_b\text{C3d}$ were incubated with highly purified lymphocytes either in the absence or in the presence of rabbit anti- E_b (IgG fraction). Fig. 1 shows specific ^{51}Cr release from the target cells after 18 h of incubation. The spontaneous ^{51}Cr release from C3-fragment-bearing erythrocytes was comparable to that observed with erythrocytes lacking C3 fragments. Target cells with C3b, C3bi, or C3d were not lysed by the lymphocytes in absence of antibody. However, all three fragments enhanced ADCC. Enhancement was between 5- and 15-fold at suboptimal antibody concentration. The number of C3 molecules per target cell varied from 15,000 to 100,000, and very similar results were obtained in replicate experiments. The enhancing effects of the C3 fragments ranked in the

order of C3bi > C3d \gg C3b. The enhancing effect of C3b was very weak. The magnitude of enhancement by the fragments varied with different lymphocyte preparations.

To study the effect of enhancement by C3 fragments on modulated lymphocytes, lymphocytes were incubated at 37°C with antibody-coated E_b or E_c before they were used as effector cells. This treatment is known to remove Fc γ R from the lymphocyte surface (19, 20) and, hence, to strongly reduce their ADCC potential. However, their natural cytotoxicity is unchanged or even increased as determined by lysis of nucleated target cells in the absence of antibody (21, 22). Table II shows the results of three experiments in which modulated lymphocytes were used with E_s with or without C3 fragments in absence or in presence of antibody. In experiments 2 and 3, modulation markedly reduced ADCC. The presence of C3 fragments on the target, however, markedly enhanced the cells' cytotoxicity, which was dependent on antibody dose. In experiment 3, lysis of C3-fragment-bearing cells could be detected even in the absence of antibody.

Demonstration of C3bi Receptors on Lymphocytes in Addition to Receptors for C3b and C3d. The interaction of lymphocytes with E_bC3b, E_bC3bi, or E_bC3d was studied using a rosetting assay. Table III shows the results obtained with lymphocytes from six different donors. Three different preparations of C3 were used. As control, the lymphoblastoid cell lines Raji and Daudi were included. Peripheral blood lymphocytes formed rosettes with all three types of indicator cells. There was no difference in the size of the rosettes, regardless of the type of fragments used. However, the percent of rosettes formed with different indicator cells varied significantly. The percent rosettes formed with E_bC3b was almost twice that with E_bC3bi and 5- to 10-fold higher than with E_bC3d. Both Raji and Daudi cells formed very strong rosettes with

TABLE II
Effect of Target Cell-bound C3 Fragments on Cytotoxicity of Fc γ R-modulated Lymphocytes

Exp.*	Target cells	Spontaneous release \ddagger	⁵¹ Cr release at different concentrations of anti-E _b §											
			Control lymphocytes (μ g IgG/ml)						Modulated lymphocytes (μ g IgG/ml)					
			0	0.13	0.4	1.3	4.0	13	0	0.13	0.4	1.3	4.0	13
1	E _s	5	9	23	40	51	5	7	25	30				
	E _s C3b	10	12	47	62	71	16	20	53	60				
2	E _s	20	12			28	11				17			
	E _s C3b	23	11			39	14			43				
	E _s C3bi	18	16			56	15			53				
3	E _s	13	12	44			14	17						
	E _s C3b	20	26	58			33	64						
	E _s C3bi	22	30	77			35	71						
	E _s C3d	24	27	72			24	57						

* Experiment 1: ~50,000 C3b-E_s; lymphocytes modulated with E_b-anti-E_b for 20 h. Experiment 2: ~20,000 C3b/E_s; lymphocytes modulated with E_c-anti-E_c for 40 h. Experiment 3: ~90,000 C3b/E_s; lymphocytes modulated with E_c-anti-E_c for 20 h.

\ddagger Percent ⁵¹Cr-release from E_s in lymphocyte-free controls.

§ Percent total ⁵¹Cr-release (uncorrected) after 18 h, lymphocyte:target cell ratios of 4:1.

TABLE III
Binding of C3-Fragment-bearing E_b by Lymphoid Cells

Exp.	Percent rosettes*								
	E_b C3b			E_b C3bi			E_b C3d		
	Lympho- cytes‡	Raji	Daudi	Lympho- cytes	Raji	Daudi	Lympho- cytes	Raji	Daudi
1	13.0	64	<1	8.8	>95	>95	1.4	>95	>95
2	10.1	20	0	5.2	>95	>95	5.2	>95	~80
3	10.5	<5	0	7.1	>95	>95	1.7	>95	~60
4	10.4	~90	ND	3.0	>95	ND	0.5	>95	ND
5	11.3	42	0	4.3	>95	ND	1.0	>95	~60
6	10.2	ND	ND	5.5	ND	ND	1.0	ND	ND
	10.9 ± 1.1§	~44	0	5.7 ± 2.1	~95	~95	1.8 ± 1.7	~95	~70

* C3-fragment-bearing E_b made up freshly for each experiment. No rosettes were formed in E_b controls included in every experiment.

‡ From six different donors.

§ Means ± SD.

|| Not done.

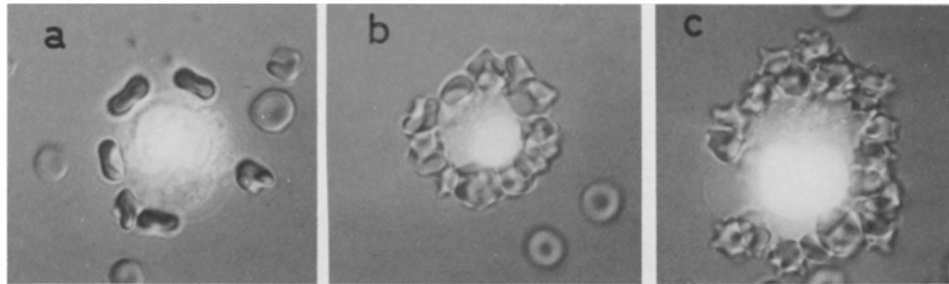


FIG. 2. Photomicrographs of Raji cells forming rosettes with E_b C3b (a), E_b C3bi (b), and E_b C3d (c).

E_b C3bi and E_b C3d. In both instances, a large number of indicator cells was bound to lymphoblastoid cells. Raji cells, but not Daudi cells, also bound E_b C3b. However, the frequency of these rosettes was low and the rosettes were weak, and contained fewer indicator cells. (Fig. 2)

Heterogeneity of Lymphocytes with Respect to Type of Complement Receptor. C3-fragment-binding lymphocytes were depleted from the effector cell preparations by two different methods: (a) adsorption of lymphocytes to monolayers of C3-fragment-bearing E_b , and (b) formation of rosettes with C3-fragment-bearing E_b , followed by separation from nonrosetting lymphocytes by centrifugation.

Table IV shows the distribution of C3 receptors among cells that did not adsorb to E_b C3d monolayers. The cells recovered in the supernate were then layered onto E_b C3bi monolayers. Each depletion step removed ~20% of the lymphocytes originally added. There was no selective loss of E_b C3b rosetting cells. The yield of E_b C3bi-binding cells after C3d depletion was slightly lower than the total yield. C3bi depletion resulted in a very significant, but not complete, reduction of E_b C3bi rosetting cells. Similarly, C3d depletion reduced the number of E_b C3d rosetting cells. The extent of

TABLE IV
Fractionation of Lymphocytes by Sequential Adsorptions to C3-fragment-bearing E_b Monolayers

Lymphocytes	Percent recovery after depletion*	Percent rosettes					
		E _b C3b		E _b C3bi		E _b C3d	
		Frequency‡	Recovery§	Frequency	Recovery	Frequency	Recovery
Control	100	10.5	100	7.6	100	1.9	100
E _b C3d depleted	83	12.3	~99	6.3	74	<0.5	<20
E _b C3bi depleted after E _b C3d depletion	63	11.0	67	1.6	15	<0.5	<20

* Recovery of lymphocytes in supernates after monolayer adsorption, given as percent of original input (= 100).

‡ Frequencies of rosette-forming lymphocytes in supernates after monolayer adsorption assayed by addition of fresh E_b-intermediates (no rosettes present in supernates before assay).

§ Recoveries of rosette-forming lymphocytes in supernates after adsorption, given as percent of original input (= 100).

TABLE V
Fractionation of Lymphocytes by Rosetting with C3-Fragment-bearing E_b

Lymphocytes	Percent recovery after depletion*	Percent rosettes					
		E _b C3b		E _b C3bi		E _b C3d	
		Frequency‡	Recovery§	Frequency	Recovery	Frequency	Recovery
Control	100	10.5	100	5.5	100	1.0	
E _b C3d depleted	75	10.0	71	3.5	48	<<0.5	NC
E _b C3b + E _b C3d depleted	66	<1	<6	2.0	24	<<0.5	NC

* Recovery of lymphocytes in supernates after rosettes depletion, given as percent of original input (= 100).

‡ Frequencies of rosette-forming lymphocytes in supernates after depletion, assayed by addition of fresh E_b-intermediates (no rosettes present in supernates before assay).

§ Recoveries of rosette forming lymphocytes in supernates after depletion, given as percent of original input (= 100).

|| Not calculated.

this depletion is not certain because of the low frequency of E_bC3d rosetting cells in this experiment.

Table V summarizes the results of an experiment in which depletion was performed by rosetting with either E_bC3d or with a mixture consisting of equal parts of E_bC3b and E_bC3d. It will again be noted that C3d depletion did not selectively diminish the number of C3b rosetting cells. Only very few remaining C3d rosettes were seen after this step. The frequency of C3bi rosetting cells was also decreased. Depletion with mixtures of E_bC3b and E_bC3d removed >90% of the E_bC3b rosetting cells and apparently all E_bC3d rosetting cells, whereas ~25% of the E_bC3bi rosetting cells remained in the supernate.

Inhibition of ADCC by C3-Fragment-bearing Bystander Cells. To explore whether interaction of lymphocytes with erythrocyte-bound C3 fragments results in the release of a soluble ADCC-enhancing factor, unlabeled E_b, bearing either C3b, C3bi or C3d, were mixed with an equal number of ⁵¹Cr-labeled erythrocytes not bearing C3

fragments. Different dilutions of antibody to E_b were added to the lymphocyte- E_b mixtures. Fig. 3 indicates that no soluble factor that would enhance lysis of C3-free cells was released under these conditions. On the contrary, the presence of C3-fragment-bearing E_b in the incubation mixtures inhibited lysis of the C3-free targets. Inhibition by C3-fragment-bearing E_b was also observed when E_c was used as target cells in the presence of anti- E_c antibodies which did not cross-react with E_b .

Discussion

The data presented in this paper show that lymphocytes that participate in ADCC may have receptors reacting with target cell-bound C3bi in addition to those reacting with C3b and C3d. All three cell-bound C3 fragments enhanced ADCC, but C3bi and C3d enhanced the reaction much more strongly than C3b. The data also show that in absence of specific antibody, cytotoxicity was not induced in spite of the presence of up to 100,000 C3-fragments/target cell, which is in agreement with our earlier results (2).

Approximately 40% of the lymphocytic effector cells of ADCC have receptors for activated C3 as assessed by rosetting of individual effector cells in the ADCC plaque assay (23, 24). At least two distinct receptors reacting with C3b or C3d have previously been found on human peripheral blood lymphocytes and other cells (25, 26). The C3b receptor (CR1 or immune adherence receptor) is present on erythrocytes, lymphocytes, polymorphonuclear leukocytes (PMN), and monocytes and reacts with the intact C3b-molecule. The C3d receptor (CR2) is present on lymphocytes, but not on PMN or monocytes and reacts with the C3d-fragment (26-28). Recently, the occurrence of an additional receptor on human PMN and monocytes with specificity for C3bi (CR3) was proposed (29). The C3bi receptor has also been found on human glomeruli (24) and on rat peritoneal mast cells (30). In this study, using human peripheral blood lymphocytes, ~11% of the cells bound EC3b, 6% bound EC3bi, and 2% bound EC3d. Raji cells bound EC3b weakly and with different frequencies when different batches of these cells were tested. In contrast, most Raji cells formed large rosettes with either C3bi- or C3d-indicator cells. The latter was also the case with Daudi cells which, however, did not bind E_b C3b. This is in accord with previous

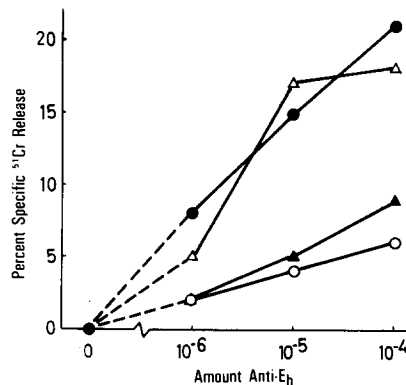


FIG. 3. Inhibition of ADCC by C3-fragment-bearing bystander cells. The target cell samples were mixtures of equal numbers of ^{51}Cr -labeled E_b and unlabeled E_b (●), E_b C3b (Δ), E_b C3bi (○), or E_b C3d (▲). Bystander cells carried 67,500 C3 fragments/cell. Experiment was performed as outlined in Materials and Methods and Fig. 1.

findings indicating that Daudi cells lack the C3b receptor, but possess the C3d receptor (31, 32).

Evidence was obtained indicating that the distribution of C3 receptors among human peripheral blood lymphocytes is nonhomogeneous. The results of the depletion experiments indicated that lymphocytes bearing C3b receptors are distinct from those bearing C3bi and/or C3d receptors. No definite conclusion could be reached regarding the differential expression of C3bi and C3d receptors on these lymphocytes. However, in unpublished studies, further insights in these questions have recently been gained by using the ADCC plaque assay (23, 24). C3bi receptor bearing plaque-forming cells (PFC) were more than twice as frequent than C3b receptor PFC. C3d receptor PFC were even fewer than C3b receptor PFC. Further, plaques were formed on monolayers of E_b that bore antibody and either C3b, C3bi, or C3d. Subsequently the PFC were incubated with E_b that bore one of the three fragments but not antibody. Under those conditions, the majority of the PFC rosetted with those cells that carried the same C3 fragments as the target cells. Most of the PFC with receptors for C3d also had receptors for C3bi. The PFC that carried both the C3bi and C3d receptors appeared to be T cells, as judged by surface-marker analysis, whereas the C3b receptor PFC were null cells. A detailed account of these results will be published elsewhere (B. Wählin, H. Perlmann, P. Perlmann, R. Schreiber, and H. Müller-Eberhard, manuscript in preparation).

It has been reported that EAC-rosetting of C3-receptor bearing lymphocytes induces the release of lymphotoxin-like factors, which enhance the natural cytotoxicity of lymphocytes to nucleated target cells (33). The present results do not support a role for soluble factors in this system. The exposure of lymphocytes to C3 fragments before the cytotoxicity assay did not increase their natural cytotoxicity to E_b, nor their ADCC potential. Admixture of C3-fragment bearing target cells to ⁵¹Cr-labeled, C3-fragment-free target cells did not enhance ADCC of the labeled cells. Enhancement should have occurred if a soluble enhancing factor had been generated in the presence of C3 fragments.

The target cell-mixing experiments gave additional information. To achieve enhancement of ADCC, antibody and C3 fragments had to reside on the same cell. When cells were introduced into the reaction mixture which carried C3 fragments but no antibody, lysis of antibody-coated cells was inhibited. The inhibitory capacity of the fragments ranked in the same order as their enhancing capacity: C3bi > C3d ≫ C3b. Presumably, this inhibition of ADCC is a result of competitive binding of the C3-fragment-bearing cells to the effector cells.

Taken together, the results described in this paper show that cell-bound C3 fragments have an important role in regulating ADCC. Because this reaction requires target cell-effector cell contact (1) C3 fragments on the surface of the target cells may enhance cytotoxicity through interaction with C3 receptors on the effector cells. That C3bi and C3d are more efficient in enhancing ADCC than C3b may be biologically significant. Although complement activation via the classical or the alternative pathway leads to deposition of C3b on the target cell surface, the structural and functional integrity of C3b is under rigid temporal constraints. Owing to the cooperative action of β1H and the C3b inactivator, C3b is converted to C3bi, which in turn is degraded to C3d. Because C3b conversion is a rapid process, the predominant forms of cell-bound C3 are C3bi and C3d. Neither of these fragments is capable of sustaining

the cytolytic reaction which would result in the formation of the membrane attack complex and cell death. However, because C3bi and C3d are more efficient in enhancing ADCC than C3b, these fragments may facilitate cellular surveillance in the presence of minute amounts of antibody.

Summary

The occurrence and distribution of distinct receptors for three C3 fragments on purified human blood lymphocytes were studied by rosette formation. Indicator cells were bovine, chicken, or sheep erythrocytes (E) bearing up to 100,000 molecules of human C3b (EC3b) without antibody. EC3b was converted to C3bi-bearing-E (EC3bi) with purified C3b inactivator (factor I) and β 1H (factor H), and to C3d-bearing E (EC3d) by treatment of EC3bi with trypsin. Using bovine E (E_b) as indicators, ~11% of the lymphocytes bound E_b C3b, 6% bound E_b C3bi and 2% bound E_b C3d. Fractionation of the lymphocytes by adsorption to monolayers of C3-fragment-bearing E_b or by rosetting indicated that most of the cells with receptors for C3b were distinct from those having receptors for C3bi and/or C3d. Cells from two lymphoblastoid cell lines (Raji and Daudi) formed strong rosettes with either EC3bi or EC3d. A fraction of the Raji cells, but not of Daudi cells formed rosettes with EC3b, which were weak. ^{51}Cr -labeled E was used as a target in antibody-dependent, lymphocyte-mediated cytotoxicity (ADCC). In the absence of antibody, C3-fragment-bearing E was not lysed by the lymphocytes. However, at suboptimal concentrations of IgG anti-E antibody, ADCC of C3-fragment-bearing E was strongly enhanced. The enhancing capacity of the fragments occurred in the order of C3bi > C3d \gg C3b. In addition, C3-fragment-bearing cells inhibited the lysis of antibody-coated cells not bearing C3 fragments. Inhibition ranked in the same order as enhancement. It is concluded that target cell bound C3 fragments enhance ADCC by improving contact between target cells and those effector cells which have C3 receptors. Cell-bound C3 fragments inhibit ADCC of C3-free targets by impeding their contact with such effector cells. It is proposed that certain lymphocytes are capable of interacting with C3bi in addition to C3b and C3d and that C3bi and C3d have a greater regulatory effect on their cytolytic function than C3b.

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