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Mutational Analysis of the Human 2B4 (CD244)/CD48 Interaction: Lys$^{68}$ and Glu$^{70}$ in the V Domain of 2B4 Are Critical for CD48 Binding and Functional Activation of NK Cells

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Interaction between receptors and ligands plays a critical role in the generation of immune responses. The 2B4 (CD244), a member of the CD2 subset of the Ig superfamily, is the high affinity ligand for CD48. It is expressed on NK cells, T cells, monocytes, and basophils. Recent data indicate that 2B4/CD48 interactions regulate NK and T lymphocyte functions. In human NK cells, 2B4/CD48 interaction induces activation signals, whereas in murine NK cells it sends inhibitory signals. To determine the structural basis for 2B4/CD48 interaction, selected amino acid residues in the V domain of the human 2B4 (h2B4) were mutated to alanine by site-directed mutagenesis. Following transient expression of these mutants in B16F10 melanoma cells, their interaction with soluble CD48-Fc fusion protein was assessed by flow cytometry. We identified amino acid residues in the extracellular domain of h2B4 that are involved in interacting with CD48. Binding of CD48-Fc fusion protein to RNK-16 cells stably transfected with wild-type and a double-mutant Lys$^{68}$Ala-Glu$^{70}$Ala h2B4 further demonstrated that Lys$^{68}$ and Glu$^{70}$ in the V domain of h2B4 are essential for 2B4/CD48 interaction. Functional analysis indicated that Lys$^{68}$ and Glu$^{70}$ in the extracellular domain of h2B4 play a key role in the activation of human NK cells through 2B4/CD48 interaction.


N atural killer cells are bone marrow-derived lymphocytes that function as key players in innate immunity by recognizing viral, bacterial, and parasitic infections and neoplastic target cells (1–3). NK cell recognition is regulated by specific receptors that, upon interaction with their respective ligands, may send stimulatory or inhibitory signals (4–7). These receptors belong to two structurally distinct superfamilies: the Ig superfamily and the lectin superfamily. In the Ig superfamily, the CD2 family plays a major role in lymphocyte function and regulation (8, 9). The CD2 family includes the CD2, CD58 (LFA-3), CS1 (CD2-like receptor activating cytotoxic cells), CD84, B lymphocyte activator macrophage expressed, CD229 (Ly-9), SF-2001, NTB-A, CD150 (signaling lymphocytic activation molecule), 2B4 (CD244), and CD48 (10). Members of the CD2 family have been shown to interact with other members of the same family or with themselves, and these interactions play an important role regulating a variety of immune responses. The 2B4 is expressed on all NK cells, a subset of CD8+ T cells, TCRγδ+ T cells, monocytes, and basophils (11–13). Ligation of surface 2B4 with a specific mAb or its ligand CD48 sends activation signals to NK cells, as evident from increased cytolytic function, induction of IFN-γ production, and NK cell invasiveness (13–17). Recently, we have generated and characterized 2B4-deficient mice that revealed an in vivo role for 2B4 in immune response against tumor (18). Further studies on these 2B4-deficient mice revealed that murine 2B4 acts as a non-MHC-binding inhibitory receptor both in vitro and in vivo, and this inhibition is independent of signaling lymphocytic activation molecule-associated protein expression (19, 20). The 2B4 is expressed early in NK cell development, before the expression of other NK cell-specific markers such as NK1.1 and Ly-49 family members, suggesting a vital role in NK cell development and function (21, 22). Defective signaling via 2B4 due to a genetic defect in signaling lymphocytic activation molecule-associated protein/Src homology 2 domain protein 1A has been implicated in the pathogenesis of X-linked lymphoproliferative disease (23–26).

Unlike 2B4, its ligand CD48 is widely expressed on cells of hemopoietic origin. Anti-CD48 mAb can inhibit the proliferation of T cells in vitro and suppress cell-mediated immunity in vivo, suggesting an important role for T cell activation (27). CD48-deficient mice have severe defect in CD4+ T cell activation (28). Earlier studies have shown that CD48 also binds to CD2, strongly in rodents and weakly in humans (27, 29–31). However, the affinity of the interaction between human CD2 (hCD2)$^4$ and hCD48 is ~100-fold lower than hCD2 and hCD58 (31). The 2B4/CD48 interaction has ~10-fold higher affinity than CD2/CD48 interaction (32). Moreover, the strength of the 2B4/CD48 interactions is conserved across the species, with similar affinities being observed for mouse and human 2B4 (h2B4)/CD48 interactions (32). Because 2B4/CD48 interaction has a much higher affinity than CD2/CD48 interaction, 2B4 is the physiological ligand for CD48. The

4 Abbreviations used in this paper: hCD, human CD; [Ca$^{2+}$], intracellular Ca$^{2+}$ concentration; 3-D, three-dimensional; h2B4, human 2B4.
2B4 and CD48 are structurally related, and the genes that encode 2B4 and CD48 are located in chromosome 1 at band 1q21–24 (33, 34).

The 2B4/CD48 interaction not only regulates NK cell functions, but also induces B cell and T cell proliferation (14, 35). Recently, it has been shown that 2B4 molecule acts as a ligand for enhancing activation and proliferation of neighboring T cells through CD48 (36). The 2B4/CD48 interaction among T cells themselves augments CTL lysis of specific targets (37). In the CD2 family, CD2 (36). The 2B4/CD48 interaction among T cells themselves augments CTL lysis of specific targets (37). In the CD2 family, CD2

Materials and Methods
Cell lines and reagents
B16F10 (mouse melanoma cell line), COS-7, and YT (human NK cell line) cells were cultured in medium RPMI 1640 supplemented with 10% FBS (HyClone), 2 mM glutamine, 100 U/ml penicillin, 100 U/ml streptomycin,
10 mM HEPES, and 10 mM nonessential amino acids. Cells were maintained at 37°C in a humidified 5% CO2/95% air incubator. All of the cell culture reagents were obtained from Invitrogen Life Technologies, unless otherwise noted. The mAb that specifically recognizes h2B4 (mAb Cl.17), and all the secondary Abs were purchased from Corixa. All enzymes were purchased from New England Biolabs, unless otherwise stated. All custom synthesized oligonucleotides used in this study were supplied by Integrated DNA Technologies.

Construction and expression of soluble CD48-Fc fusion protein

Soluble CD48-Fc fusion protein was produced by fusing the extracellular domain of the CD48 with Fc portion of the human IgG. The extracellular domain of the hCD48 (GenBank Accession M59904) was amplified by PCR using forward primer, CD48FP, 5'-TGG TGG CTA GCG ATT CAA GGT CAC TTG-3' and reverse primer, CD48RP, 5'-AAG AAT GGA TCC ACC GTG ACC ACT AGC C-3'. The amplified product was subcloned in front of human Fc gene at Nhel and BamHI cloning sites in pCD5lneg1 vector (kindly provided by B. Seed, Harvard Medical School, Boston, MA), which contain the C2 and C3 regions of the human IgG1. To verify the orientation and folding of the fusion protein, 2 μg of pure pDNA was transfected into 50% confluent COS-7 cells using Fugene-6 transfection reagent (Roche Diagnostic Systems). After 48 h, the cells were washed three times with PBS and changed to cysteine- and methionine-depleted RPMI 1640 medium (Mediatech). A total of 150 μCi/ml 35S-labeled cysteine and methionine (ICN Radiochemicals) was added to medium, and after 48 h supernatant was collected. A total of 5 μg of protein A-agarose beads (Bio-Rad) was added in the supernatant to precipitate the CD48-Fc fusion protein. The beads were washed well with PBS to remove the nonspecific binding proteins. Then beads were split into two halves and eluted with SDS-loading buffer with and without DTT. The proteins were separated by 10% SDS-PAGE under reducing and nonreducing conditions, and the gel was dried to expose for autoradiography. For checking the CD48 portion of fusion protein, immunoprecipitated proteins were electrophoretically transferred to polyvinylidene difluoride (Pierce) membrane. Membranes were incubated with anti-CD48 mAb (1/100 dilution) in blocking buffer overnight. Peroxidase-conjugated secondary Abs were used at 1/2000 dilution, and immunoreactive bands were visualized using ECL (Kirkgegaard & Perry Laboratories), as per manufacturer’s description. For isolation of large quantities of the CD48-Fc fusion protein, COS-7 cells were transfected in large quantities; supernatants were collected at days 3, 6, and 9, and protein was isolated by protein A affinity chromatography.

Production of polyclonal Ab against h2B4

The 2B4-Fc fusion protein was produced, as described above, for CD48-Fc fusion protein. The extracellular domain of the h2B4 was PCR amplified using forward primer, h2B4FP, 5’-AAG AAT GGA TCC ACC GTG ACC ACT AGC C-3’ and h2B4RP, 5’-AAG AAT GGA TCC ACC GTG ACC ACT AGC C-3’. The amplified product was cloned in front of human Fc gene at Nhel and BamHI cloning sites in pcDNA5 neo (kindly provided by Lingzhi Zhang, Texas Southwestern Medical Center, Dallas, TX). Constructs were confirmed by sequencing the mutated pcDNA5 neo clones. For the double mutation, a single PCR primer contained both mutations because the two mutations were close to each other.

Transfection and flow cytometry analysis for h2B4-mutated clones

B16F10 cells were transiently transfected with h2B4-mutated pDNA using Fugene-6 transfection reagent (Roche Diagnostic Systems). After 48 h, cells were incubated with specific mAbs (Cl.17), polyclonal Ab against 2B4, and CD48-Fc fusion protein to detect the expression of the molecules. They were then washed with cold PBS at low speed and incubated with FITC-conjugated goat anti-mouse IgG for 1 h, FITC-conjugated goat anti-rabbit IgG for 40 min at 4°C. Samples were washed twice and were analyzed by flow cytometry.

Cytotoxicity assay

Target cells were labeled by incubating 1 × 10^6 cells with 2 MBq of Na235CrO4 (NEN Research Products) for 90 min at 37°C under 5% CO2 in air. The target cells (100 μl) were incubated with effector cell suspension (100 μl) under various conditions. After incubation for 4 h at 37°C under 5% CO2 in air, the cells were pelleted at 250 × g for 5 min, 100 μl of the supernatants were removed, and their radioactivity was measured. The percentage of specific lysis was calculated by the following equation: (a - b/c - d) × 100, where a is the radioactivity of the supernatant of target cells mixed with effector cells, b is that in the supernatant of target cells incubated alone, and c is that in the supernatant after lysis of target cells with 1% Nonidet P-40.
Calcium flux assay

We used a protocol similar to that used by Prasanna et al. (41). Briefly, to measure intracellular calcium mobilization, RNK-16 cells stably expressing wild-type and double-mutant K68AE70A h2B4 were seeded on coverslips and loaded in 3 μM fura 2 dye (Molecular Probes) in a modified Krebs-Ringer buffer solution (in millimoles: 115 NaCl, 2.5 CaCl₂, 1.2 MgCl₂, 24 NaHCO₃, 5 KCl, 5 glucose, and 25 HEPES, pH 7.4) for 30 min at 37°C. Then, sCD48-IgG (Fc) fusion protein or anti-NKR-P1A (10/78) mAb was used to induce receptor cross-linking, and cross-linking goat anti-human or goat anti-mouse IgG (20 μg/ml) (BD Pharmingen) was added. The fura 2 fluorescence from these cells was monitored at 37°C by the ratio technique (excitation at 340 and 380 nm, emission at 500 nm) under a Nikon Diaphot microscope using Metafluor software (Universal Imaging). Calibrations were performed in vivo, and conditions of high intracellular Ca²⁺ concentration ([Ca²⁺]ᵢ) were achieved by adding the Ca²⁺ ionophore 4-Bromo-A23187 (1–3 μM; Calbiochem), whereas conditions of low [Ca²⁺]ᵢ were obtained by adding EGTA (4–5 mM). Statistical significance of [Ca²⁺]ᵢ between control and treatments was determined parametrically by Student’s t test at p < 0.05.

Results

Recombinant CD48-Fc fusion protein is dimeric and binds h2B4

To study 2B4/CD48 interactions, we generated soluble CD48-Fc fusion protein by ligating the extracellular domain of CD48 with Fc portion of human IgG at the C terminus. ³⁵S-labeled CD48-Fc protein was immunoprecipitated by protein A-agarose and separated on SDS-PAGE to check purity and biochemical structure. Autoradiogram of the gel showed bands of 135 and 65 kDa under nonreducing and reducing conditions, respectively, indicating CD48-Fc fusion protein as a dimer and in proper folding (Fig. 1A). Affinity-purified fusion protein was used to detect the binding of CD48 to h2B4 by flow cytometry (Fig. 1B). The CD48-Fc fusion protein binds specifically to B16F10 cells transfected with h2B4 and not to untransfected cells, suggesting that it is functionally active. The binding was inhibited after blocking with h2B4 Abs, suggesting that the interaction is specific for h2B4 (Fig. 1B).

Amino acid residues in the V domain of 2B4 contribute to 2B4/CD48 interaction

The CD2/CD58 pair was selected to compare 2B4/CD48 interaction studies because both pairs are structurally and functionally related. CD2 also binds to CD48 weakly as compared with 2B4 (29). Early studies suggested involvement of the charged residues in protein recognition and cell adhesion (42, 43). Recently, CD2-CD58 receptor-ligand interaction is well studied using x-ray crystallography, nuclear magnetic resonance, and site-directed mutation analysis (44–46). To identify the charged amino acids that may be involved in 2B4/CD48 interaction, the extracellular domains of 2B4, CD48, CD2, and CD58 were aligned using the GCG software (Fig. 2). Based on the earlier point mutational and the crystallographic studies, the key amino acids involved in the CD2/CD58 interaction were selected. In CD2, mutations K34, K43, K41, K51, D31, D32, R48, and K91 and in CD58, mutations E25, K32, E39, E42, R44, D33, K34, E37, E78, E84, and E95 showed to interrupt the CD2-CD58 interaction modest to completely (42, 43, 45). The amino acid residues in 2B4 selected for mutagenesis are given in bold. Table I gives the list of amino acids selected for point mutation along with PCR primer sequences used for site-directed mutagenesis.

Preliminary experiments were conducted by expressing all of the 11 single mutants and wild-type 2B4 in B16F10 melanoma cells by transient transfection. The expression of the h2B4 mutants was verified by FACS analysis with C1.7 (mAb against h2B4) as well as polyclonal antisera against h2B4. Figs. 3 and 4 show the expression pattern of wild-type h2B4 and mutants in B16F10 cells. All of the mutants are expressed on the cell surface, like wild-type h2B4. Binding of C1.7 indicated that some of the point mutations affect the mAb binding; in particular, K68A and E70A severely disrupted the C1.7 binding, indicating that these two residues may be involved in recognition of h2B4 by mAb C1.7. The role of these two residues in C1.7 binding is further evident, as seen in Fig. 5C. This indicates that some residues involved in ligand binding and mAb C1.7 binding may overlap. This is not surprising because mAb C1.7 activates human NK cells. The expression of the mutated clones was also checked by Western blot (data not shown). Both the FACS and Western blot analysis depict the same results that the anti-h2B4 Abs bind to all h2B4 mutants like wild-type h2B4. This suggests that by mutating to alanine, the three-dimensional structure of h2B4 is not altered and can be used to identify
the key amino acids involved in binding with its ligand CD48. The expression pattern of the mutants compared with the wild type is shown in Table II and Figs. 3 and 4. FACS analysis was performed with CD48-Fc fusion protein with wild-type h2B4 and h2B4 mutants. Mean fluorescence intensity of binding of CD48-Fc to h2B4 mutants in comparison with its binding to wild-type h2B4 is also shown in Table II. Statistical analysis using \( t \) test (\( p < 0.05 \)) showed that K47A, D49A, S50A, K55A, N61A, and T110A did not significantly alter the CD48 binding, whereas K54A, G114A, and K115A significantly altered the binding. In particular, mutants K68A and E70A were shown to highly significantly alter the binding to CD48. To determine the relative locations of these residues on the V domain and to verify whether these modifications alter the 3-dimensional (3-D) structure of the h2B4, a predicted 3-D structure of h2B4 and its mutants was determined using the 3D-PSSM software (www.sbg.bio.ic.ac.uk/~3dpssm) (47). The predicted residues based on computer simulation that affect CD48 binding to h2B4 lie closely together on the surface of the V domain (Fig. 6). Corresponding to CD2/CD58 interaction, E70 lies in the C' strand and K54 lies on the F strand of the predicted \( \beta \) strand, whereas the other amino acids alter the binding map within the predicted loops: two residues in FG loop (G114 and K115) and one residue in C'C' loop (K68). Thus, several charged amino acid residues make relative contributions to h2B4/CD48 interaction.

To further confirm the results obtained by transient transfection, we generated stable expression of wild-type and double-mutant K68A,E70A h2B4 in RNK-16 and K562 cells. Both wild-type and double-mutant K68A,E70A h2B4 showed similar levels of expression on RNK-16 as well as K562 cells (Fig. 5, A and B). The significance of the double mutant was further confirmed when it

**FIGURE 5.** Stable expression of wild-type and double-mutant K68AE70A h2B4 in RNK-16 and K562 cells. Wild-type and double-mutant K68AE70A h2B4 were stably expressed in RNK-16 and K562 cells. A, Expression of stably transfected wild-type and double-mutant K68AE70A h2B4 in RNK-16 cells determined by flow cytometry using preimmune sera (open) and anti-h2B4 polyclonal Ab (filled). FITC-conjugated anti-rabbit IgG was used as the secondary Ab. B, Expression of stably transfected wild-type and double-mutant K68AE70A h2B4 in K562 cells determined by flow cytometry using preimmune sera (open) and anti-h2B4 polyclonal Ab (filled). FITC-conjugated anti-rabbit IgG was used as the secondary Ab. C, Binding of mAb C1.7 to RNK-16 cells stably transfected with wild-type and double-mutant K68AE70A h2B4. Open histogram control Ab and filled histogram mAb C1.7. FITC-conjugated anti-mouse IgG was used to detect C1.7 by flow cytometry. Results are representative of three independent experiments.

**FIGURE 4.** The double mutation K68AE70A abolishes CD48 binding to h2B4. The double mutation K68AE70A abolishes CD48 binding to h2B4. B16F10 cells transfected with pCI-neo empty vector (open) and B16F10 cells transfected with mutants K68A, E70A, and double-mutant K68AE70A (filled) were incubated with mAb (C1.7) denoted by mAb, h2B4 polyclonal Ab denoted by pAb, and soluble CD48-Fc fusion protein denoted by CD48. FITC-conjugated anti-mouse IgG, anti-rabbit IgG, and anti-human IgG were used to detect C1.7, polyclonal anti-h2B4, and CD48-Fc fusion protein, respectively, by flow cytometry. Results are representative of three independent experiments.

Mutations Lys\(^{68}\)Ala and Glu\(^{70}\)Ala abolish h2B4/CD48 interaction

The h2B4 mutants K68A and E70A showed mean fluorescence intensity values of 3.59 and 1.60 with CD48-Fc fusion protein, respectively. This suggests that these amino acids play a significant role in CD48 binding to h2B4. When both the amino acid residues were mutated, it completely abolished the binding of CD48 to h2B4 (Fig. 4 and Table II). The predicted 3-D structure of the double mutant was similar to the wild-type 3-D structure (Fig. 6). More importantly, Glu\(^{70}\) lies in the \( \beta \) sheet of the Ig domain, implicating that h2B4/CD48 interaction is similar to the face-to-face interaction of CD2 and CD58, which involves \( \beta \) sheets of both counterreceptors. This indicates that amino acid residues Lys\(^{68}\) and Glu\(^{70}\) are essential for h2B4/CD48 interaction.
almost completely abolished the binding of CD48 to 2B4 (Fig. 5).

Mutations Lys\textsuperscript{68}Ala and Glu\textsuperscript{70}Ala inhibit NK cell function

h2B4 has previously been shown to activate NK cell-mediated cytotoxicity (12, 15, 16, 48, 49). To determine whether interaction through Lys\textsuperscript{68} and Glu\textsuperscript{70} of h2B4 with CD48 plays any functional role in NK cell activation, we performed cytotoxicity assays as well as calcium flux experiments. For cytolytic function, we used human NK cell line NK92 as effectors against 51Cr-labeled K562 target cells stably transfected with h2B4 and double-mutant K68AE70A. The expression of wild-type and double-mutant K68AE70A h2B4 in K562 cells was determined by flow cytometry using polyclonal anti-h2B4 (Fig. 5B). Because K562 is a very susceptible target for NK killing, the baseline killing is very high in the absence of any other stimulation, as expected. Expression of wild-type h2B4 in K562 cells increased the cytolytic function of NK92 cells. This may be due to enhanced adhesion of these transfected cells with NK92 as well as activation signals resulting from 2B4/CD48 interactions. The cytotoxic activity of NK92 against K562 transfected with the double-mutant K68AE70A reduced the killing to the basal level (Fig. 5B). Because K562 is a very susceptible target for NK killing, the baseline killing is very high in the absence of any other stimulation, as expected. Expression of wild-type h2B4 in K562 cells increased the cytolytic function of NK92 cells. This may be due to enhanced adhesion of these transfected cells with NK92 as well as activation signals resulting from 2B4/CD48 interactions. The cytotoxic activity of NK92 against K562 transfected with the double-mutant K68AE70A reduced the killing to the basal level (Fig. 7A). Similarly, wild-type and double-mutant K68AE70A h2B4 expressed in the rat NK cell line RNK-16 was able to mediate cellular cytotoxicity of 51Cr-labeled P815 target cells stably transfected with hCD48. As compared with the wild-type h2B4-RNK-16 transfectants, the double-mutant K68AE70A-transfected RNK-16 cells showed significantly reduced cytolytic activity against P815 cells transfected with hCD48 (Fig. 7B). This suggests that 2B4/CD48 interaction involving the amino acid residues Lys\textsuperscript{68} and Glu\textsuperscript{70} does play a role in the cytolytic function of NK cells.

We also performed intracellular calcium flux experiments to confirm the functional data obtained in the cytotoxicity assay. Wild-type and the double-mutant K68AE70A h2B4 were expressed in RNK-16 cells, and induction of calcium release was determined upon interaction with soluble CD48-Fc fusion protein. An Ab against NKR-P1, mAb 10/78, was used as a positive control, as it has been shown previously that it stimulates phosphoinositide turnover and induces calcium flux within RNK-16 cells (50–52). Cross-linking of h2B4 using soluble CD48-Fc fusion protein on h2B4 RNK-16 transfectants resulted in a significant calcium flux, as measured by fura 2 over time, whereas cross-linking of mutated h2B4 on double-mutant K68AE70A RNK-16 transfectants did not show any discernible calcium mobilization (Fig. 8), which convincingly corroborates with the cytotoxicity assays.

**Discussion**

Direct cell-cell interactions are important for immune cell function to fight against foreign pathogens and tumor cells. Multiple receptor-ligand interactions present on the cell membrane determine the cell functions in both adaptive and innate immunity. In humans, CD58 is the high affinity receptor for CD2 and is the well-studied
heterophilic adhesive interaction model. Wang et al. (45) crystal-
lized this receptor-ligand complex and studied the interface of the
structure in detail. The CD2/CD58 interactions are predominantly
determined by charged amino acids present in the extracellular
domain. The nature of the interface is derived from charge comple-
mentarity rather than shape complementarity. In support of this
view, CD2 has more positively charged residues, whereas its coun-
terreceptor CD58 has more negatively charged residues at the ex-
tracellular domain. Like CD2/CD58, 2B4 has seven positively
charged and two negatively charged amino acid residues, whereas
CD48 has five positively charged and seven negatively charged
amino acid residues in the V domain. This strongly suggests that
2B4/CD48 interactions may be charge driven.

In the CD2/CD58 model, 10 salt bridges and 5 hydrogen bonds
form an interdigitated salt bridge network. This array not only
ensures high coligand specificity, but also contributes binding en-
ergy, because the unfavorable like-charge residues clustering in
each of the binding surfaces will be neutralized upon complex
formation (44). In our study, mutation at K54, G114, and K115
reduced the CD48 binding, suggesting that these amino acids are
involved in initial binding by forming salt bridge as well as hy-
drophobic bond with CD48. This initial recognition and V domain
elasticity help for corecognition of other charged amino acids to
form a strong salt bridge between 2B4 and CD48. Changing K68
and E70 to alanine significantly reduces the binding and totally
abolishes it in the K68AE70A double-mutated 2B4 protein, sug-
gestting that these two are primarily involved in the CD48 binding
(Figs. 4 and 5D). Interestingly, our data also identified a critical
role for these two amino acid residues in the recognition of h2B4
by mAb C1.7 (Fig. 5C). CD48 has a Ser and Lys residue corre-
sponding to positions 68 and 70 in 2B4 (Fig. 2). Therefore, it is
tempting to speculate that Glu70 of 2B4 may form salt bridges
electrostatic interactions) with Lys 70 of CD48. Moreover, E70 is
reduced and two negatively charged amino acid residues, whereas
CD48 has five positively charged and seven negatively charged

amino acid residues in the V domain. This strongly suggests that
2B4/CD48 interactions may be charge driven.

In conclusion, we have shown that charged amino acids in the V
domain of 2B4 are involved in 2B4/CD48 interaction. We further
demonstrated that the amino acid residues Lys68 and Glu70 in the
V domain of h2B4 play a critical role in physical interaction with
CD48, and disruption of this interaction impairs functional activa-
tion of NK cells through both receptors. The identification of
h2B4/CD48 binding site has implication in designing small mol-
ecules that could block/modulate NK and T cell functions resulting
from 2B4/CD48 interaction.
**FIGURE 8.** The double mutation, K68AE70A, inhibits calcium mobilization in RNK-16 transfectants. Calcium mobilization determined by the ratio technique fura 2 over time. A, RNK-16 cells stably expressing wild-type h2B4; B, double-mutant K68AE70A (Fig. 5A) were seeded on coverslips and loaded in 3 μM fura 2 dye in a modified Krebs-Ringer buffer solution for 30 min at 37°C. They were treated with 10 μg/ml soluble CD48-Fc fusion protein (a), followed by 20 μg/ml goat anti-human IgG (b). C, Untransfected RNK-16 were treated either with 5 μg/ml anti-NKR-P1A (10/78) mAb (positive control) (thick) or 10 μg/ml soluble CD48-Fc fusion protein (thin) (a) and subsequently exposed to 20 μg/ml goat anti-mouse IgG or goat anti-human IgG (b), respectively. The data are representative of three independent experiments.

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**Disclosures**

The authors have no financial conflict of interest.

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