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MHC Class II Molecules Traffic into Lipid Rafts during Intracellular Transport

Neil J. Poloso, Aura Muntasell, and Paul A. Roche

There have been many studies demonstrating that a portion of MHC class II molecules reside in detergent-insoluble membrane domains (commonly referred to as lipid rafts). We have proposed that the function of raft association is to concentrate specific MHC class II-peptide complexes in plasma membrane microdomains that can facilitate efficient T cell activation. We now show that MHC class II becomes lipid raft associated before binding antigenic peptides. Using pulse-chase radiolabeling techniques, we find that newly synthesized MHC class II and MHC class II-invariant chain complexes initially reside in a detergent-soluble membrane fraction and acquire detergent insolubility as they traffic to lysosomal Ag processing compartments. Monensin, an inhibitor of protein transport through the Golgi apparatus, blocks association of newly synthesized MHC class II with lipid rafts. Treatment of cells with leupeptin, which inhibits invariant chain degradation, leads to the accumulation of MHC class II in lipid rafts within the lysosome-like Ag-processing compartments. Raft fractionation of lysosomal membranes confirmed the presence of MHC class II in detergent-insoluble microdomains in Ag-processing compartments. These findings indicate that newly synthesized MHC class II complexes are directed to detergent-insoluble lipid raft microdomains before peptide loading, a process that may facilitate the loading of similar peptides on MHC class II complexes in these microdomains. *The Journal of Immunology*, 2004, 173: 4539–4546.

Activation of CD4+ T cells requires engagement of the TCR with specific MHC class II-peptide complexes on the surface of an APC. Less than 500 MHC class II-peptide complexes are required for T cell activation, a number that can represent <0.1% of the total amount of class II on the APC surface (1–3). We are interested in understanding the mechanism that would allow a T cell to engage such a small number of specific class II-peptide complexes. One possible mechanism could be that these relevant class II-peptide complexes are not randomly distributed on the surface of APCs but are instead locally concentrated in small plasma membrane microdomains.

Lipid rafts are membrane microdomains that have received considerable attention in recent years. These microdomains are characterized as having a high concentration of cholesterol and sphingolipids and are often enriched in GPI-anchored and other lipid-modified proteins. Although individual rafts are too small to visualize by light microscopy, rafts and raft components are often characterized by their biophysical property of exhibiting relative insolubility in nonionic detergents and can be isolated in discontinuous sucrose density gradients (4). As assessed by both detergent insolubility and immunofluorescence microscopy, there is considerable evidence that a significant proportion of MHC class II resides in these detergent-insoluble lipid raft membrane microdomains in a variety of APC subtypes, including B cells, macrophages, and activated dendritic cells (5–12). Disruption of APC lipid rafts profoundly inhibits T cell activation by APCs only under conditions in which the number of Ag-specific complexes is limiting (13). This evidence has led us to propose that association with lipid rafts serves to increase the local surface density of MHC class II peptide complexes to facilitate TCR cross-linking and activation (11, 13).

Because perturbing MHC class II association with lipid rafts can inhibit Ag presentation to T cells, it follows that under these conditions each “raft” must contain more than one class II molecule with any given peptide (i.e., each raft must contain at least two copies of identical class II-peptide complexes). However, how MHC class II containing identical peptides could come to reside in the same lipid raft microdomain is unclear. Our hypothesis is that it occurs by a mechanism that is intimately related to the cell biology of class II trafficking and antigenic peptide loading in APCs. Newly synthesized class II binds to a chaperone protein, termed the invariant chain (II)2 in the endoplasmic reticulum (ER), which escorts nascent MHC class II αβ complexes out of the ER, through the Golgi apparatus, and ultimately into lysosome-like Ag processing compartments (14). In these compartments, Ii is degraded by proteases and dissociates from MHC class II (15). Only a small CLIP remains associated with the peptide-binding groove of the MHC class II αβ dimer in this compartment (16, 17). The peptide editor HLA-DM, which also resides in these Ag-processing compartments, catalyzes the exchange of the class II-associated CLIP fragment with high affinity antigenic peptides that are generated by proteolysis of internalized foreign Ag (16, 17). Upon antigenic peptide loading the MHC class II-peptide complexes travel via vesicles and/or tubules from these peptide-loading compartments to the plasma membrane (14, 18, 19).

In this study, we have examined at what point in the intracellular MHC class II trafficking pathway MHC class II first associates with lipid raft microdomains. We considered two distinct, testable possibilities: 1) that newly arriving MHC class II-peptide complexes are already raft-associated before delivery to the plasma

1 Intracellular Transport

2 Abbreviations used in this paper: Ii, invariant chain; ER, endoplasmic reticulum; MCD, methyl-β-cyclohextrin; endo H, endoglycosidase H; NEM, N-ethylmaleimide; pH, pH gradient electrophoresis; LIP, leupeptin-induced polypeptide.
membrane or 2) that MHC class II-peptide complexes associate with pre-existing lipid rafts only after arrival at the plasma membrane. Using pulse-chase biosynthetic labeling techniques, we find that the former of the two possibilities is correct and that newly synthesized MHC class II associates intracellularly with lipid rafts in a post-Golgi Ag-loading compartment. Additionally, we show that association of MHC class II with lipid rafts is not dependent upon class II binding to Ii, demonstrating that raft association of MHC class II is an intrinsic property of the MHC class II molecule itself and does not require Ii-dependent sorting of MHC class II molecules to intracellular Ag-processing compartments.

Materials and Methods

Cell lines

Human JY B cells were maintained in RPMI 1640 containing 10% FBS, 10 mM glutamine, 10 mM HEPES, and 50 µg/ml gentamicin (complete RPMI) at 37°C. ALLa cells were maintained in DMEM containing 10% FBS, 10 mM HEPES, 10 mM glutamine, and 50 µg/ml gentamicin at 37°C.

Transfection studies

cDNA constructs used in this study were pcDNA-3 (Invitrogen Life Technologies, Carlsbad, CA), pcDNA3-human I-µ33, and CD8α-HLA-DR3α, CD8α-HLA-DR3β (20). Subconfluent HeLa cells were transiently transfected using LipofectAmine Reagent (Invitrogen Life Technologies) according to manufacturer’s instructions and harvested for biochemical studies 24 h later.

Abs and reagents

Abs to ICAM-1 and CD59 were purchased from Serotec (Raleigh, NC). mAb LA243 and isotype control IgG Abs were purchased from BD Pharmingen (San Diego, CA). Anti-calnexin mAb was purchased from BD Transduction Laboratories (San Diego, CA). The mAbs to the HLA-DR-α-chain (DA6.147), HLA-DR-β-chain (XDS.A11), and human Ii (Pin1.1) have been previously described (20). A rabbit antiserum recognizing the cytosolic domain of the HLA-DR-β-chain was the generous gift of Dr. E. Long (National Institute of Allergy and Infectious Diseases, National Institutes of Health, Bethesda, MD). The control Ab, MOPC21, and protein A-Sepharose beads was purchased from Sigma-Aldrich (St. Louis, MO). The anti-I-α Ab, LL1, has been previously described (21). The anti-HLA-DM mAb K-76 was a kind gift of Dr. S. Pierce (National Institute of Allergy and Infectious Diseases, National Institutes of Health). [35S]Methionine was purchased from ICN Biosciences (Irvine, CA). Methyl-β-cyclodextrin (MCD) was purchased from Sigma-Aldrich. Endoglycosidase H (endo H) was purchased from New England Biolabs (Beverly, MA).

Radiolabeling (pulse-chase) experiments

Human JY B cells were harvested from culture (in log phase growth) and washed twice with methionine-free DMEM containing 3% dialyzed FBS, 10 mM HEPES, 50 µg/ml gentamicin, and 50 µM 2-ME. Cells (usually 75 × 10^6 cells per time point/condition) were resuspended at 10 × 10^6/ml and incubated in methionine-free medium at 37°C for 9 min. [35S]Methionine was added to a final concentration 1.5 mCi/75 × 10^6 cells and incubated with the cells at 37°C in a humidified CO2 atmosphere for various time points. The pulse radiolabeling was stopped by the addition of ice-cold media containing a 20-fold excess of unlabeled methionine and washed. Cells were then either immediately used for subsequent raft fractionations or chased in complete RPMI medium containing a 5-fold excess of unlabeled methionine at a concentration of 2 × 10^6/ml for various time points. In some experiments, monensin (100 nM; Sigma-Aldrich) or leupeptin (0.5 mg/ml; Sigma-Aldrich) were added during the chase. In some studies, either unlabeled or radiolabeled cells were treated with 10 mM MCD for 12 min before raft isolation as described previously (11).

Isolation of lipid rafts

Isolation of membrane rafts was performed as described in Ref. 13. Briefly, cells were lysed at a concentration of 10^7 cells/ml in MES-saline buffer (20 mM MES, 150 mM NaCl, pH 6.5) containing 1% Brij-58 and protease inhibitors (500 µM PMSF, 100 µM Na-p-tosyl-l-lysine chloromethyl ketone (TLCK), 5 mM iodoacetamide, 10 µg/ml aprotinin, and 5 µg/ml leupeptin) for 1 h on ice. The lysate was mixed 1:1 with 90% sucrose in MES saline and plated at the bottom of an ultracentrifuge tube. The sample was overlaid with 6 ml of 30% sucrose and 4 ml of 5% sucrose in MES buffer and centrifuged at 100,000 × g in a SW41 Ti swinging bucket rotor at 4°C for 18 h. Fractions (1 ml) were gently removed from the top of the gradient and each sample was incubated with N-octylglucoside (1% final) at 4°C for 1 h to completely solubilize Brij-58 insoluble membrane fractions.

Isolation of lysosomal Ag-processing compartments

Lysosomes from JY B cells were isolated by standard protocol as described (22). JY B cells (10^7) were washed twice in HBSS (Mg and Ca-free) and once in TEA/sucrose buffer (10 mM tris(hydroxymethyl)aminomethane (Tris), 10 mM HEPES, 10 mM acetic acid, 250 mM sucrose, pH 7.4). Cells were then resuspended in ice-cold TEA/sucrose (~35 × 10^6/ml) and passed through a ball-bearing homogenizer (0.01 cm clearance) on ice for eight passes (resulting in ~80% cracked cells). The homogenate was subjected to centrifugation at 1300 × g for 10 min twice to remove intact cells, nuclei, and cell debris. Two milliliters of postnuclear supernatant was mixed with 9 ml of ice-cold TEA-buffered 27% Percoll (Amersham Biosciences, Piscataway, NJ) and placed into a 12-ml ultracenfuge tube. A 0.6-ml cushion of TEA-buffered 27% Histodenz (Sigma-Aldrich) was underlaid beneath the Percoll and the sample was subjected to centrifugation at 17,000 rpm in a SW41Ti rotor for 1 h at 4°C. The low density membranes (2 ml) were visualized and collected from the top of the gradient and the high-density lysosomal membranes (0.6 ml) were visualized just above the Histodenz cushion and collected by needle puncture. Aliquots of each pool were saved for immunoblot analysis. The membranes from each pool were pelleted by centrifugation at 100,000 × g for 4°C, washed once in PBS, and the membrane pellets lysed in an equal volume of MES buffer containing 2% Brij-58 and protease inhibitors. Lipid raft isolation was conducted as described in the previous section.

Immunoprecipitations

To detect radiolabeled proteins or protein complexes in membrane raft fractions, pools of raft and nonraft (soluble) fractions were subjected to immunoprecipitation. BSA (1 mg/ml final) was added to each pool to inhibit nonspecific protein adsorption to Sepharose beads. Protein A-Sepharose beads (Sigma-Aldrich) were precoated to rabbit anti-mouse Ig serum (Sigma-Aldrich) and specific mouse mAbs were incubated with the beads for 1 h at 4°C. Beads were washed in Tris-saline buffer (10 mM Tris, 150 mM NaCl, pH 7.4) and once in immunoprecipitation buffer (1% Triton X-100, 1 mg/ml BSA in Tris saline containing protease inhibitors as described in raft isolation). Pooled fractions were precleared with beads coupled to the control mAb MOPC21 for 1 h at 4°C. Aliquots of precleared lysates were added to specific immunoprecipitation beads and the mixtures were incubated 2–4 h at 4°C on a rotator. Immunoprecipitates were extensively washed in immunoprecipitation buffer and the pellets were frozen at −80°C until analysis. Immunoprecipitates were analyzed by either one-dimensional SDS-PAGE or two-dimensional gel with nonequilibrium pH gradient electrophoresis (NEPHGE) in the first dimension followed SDS-PAGE in the second dimension (23). All radiolabeled gels were fixed in 20% methanol/10% acetic acid, and signals enhanced by treatment with Enlightening (ICN Biosciences) before drying and exposure. Aliquots of each immunoprecipitate were also analyzed by SDS-PAGE and immunoblotting. In some experiments, immunoprecipitates were split and treated with or without endo H for 1 h at 37°C as per the manufacturer’s instructions.

Immunoblotting

To visualize proteins residing in membrane rafts, equal portions of each fraction or pooled fractions were analyzed by separation on 10.5% SDS-PAGE, transferring to Sequi-gel PVDF membranes (Bio-Rad, Hercules, CA) and immunoblotting with specific mAbs. Blots were analyzed by using rabbit, mouse, or rat Abs (either cytoplasmic, purified Abs, or ascite fluids) as described in the figure legends. Blots were developed with HRP-conjugated anti-mouse IgG (1:2500) or anti-rabbit IgG (1:5000) (Southern Biotechnologies Associates, Birmingham, AL) followed by ECL (PerkinElmer, Boston, MA). Quantitative analysis of protein expression in various samples was performed by laser densitometry of either immunoblots or autoradiograms using a Molecular Dynamics Densitometer and ImageQuant software (Sunnyvale, CA). Scans of multiple exposures were obtained to ensure that the results fell into the linear range of the instrument.

Results

MHC class II is insoluble in the detergent Brij-58

Previously, we have reported that ~15–20% of MHC class II molecules reside in lipid rafts as shown by TX-100 detergent insolubility (11). Subsequent studies have shown that anywhere from 2 to 50% of MHC class II is detergent insoluble as analyzed by

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sucrose density gradient centrifugation depending on the cell type examined, the ratio of cell protein-to-detergent, and detergent used for membrane protein solubilization (6, 8–10, 12, 24). Therefore, before investigating the kinetics of MHC class II association with detergent insoluble membrane microdomains, we sought to fully characterize the distribution of MHC class II and other integral membrane proteins in these raft microdomains isolated from EBV-transformed B cell lines lysed in the nonionic detergent Brij-58. Sucrose density gradient centrifugation revealed that both calnexin (an ER-resident protein) and ICAM-1 (a plasma membrane protein) were excluded from the low buoyant density raft fractions and were found in the soluble fractions of the gradient (98 and 100%, respectively) (Fig. 1A). By contrast, CD59, a GPI-anchored protein, exclusively partitions with the detergent-insoluble raft fractions (Fig. 1A). Approximately 65% of the total pool of membrane-associated MHC class II partitioned with the lipid raft fractions in these gradients (Fig. 1, A and C). This behavior was cholesterol dependent, because perturbation of plasma membrane cholesterol using MCD before raft isolation reduced MHC class II association with lipid rafts by 85% (Fig. 1, B and C). From these results, we conclude that the low buoyant density of MHC class II is not an artifact of incomplete membrane solubilization and is a cholesterol-dependent phenomenon.

MHC class II becomes detergent-insoluble during transport to the cell surface

In our attempts to understand more completely the biological role of MHC class II association with detergent-insoluble lipid raft membrane microdomains, we set out to determine whether MHC class II molecules that have not yet bound antigenic peptides were raft-associated or if class II association with lipid rafts only occurred after peptide loading. To address this question, we “pulse” radiolabeled B cells with [35 S]methionine for only 10 min and followed the fate of these labeled MHC class II molecules during different times of “chase” to determine whether raft association of newly synthesized MHC class II changed as the class II molecules trafficked through the cell. Very little newly synthesized MHC class II molecules were found in lipid rafts following the 10 min pulse (<2%, Fig. 2A). After a 3-h chase, a time when some (but not all) MHC class II would have reached the cell surface (25), 45% of newly synthesized MHC class II had now associated with rafts (Fig. 2, A and C). The absence of the high-mannose, unprocessed form of Ii-p33 in the MHC class II immunoprecipitate suggests that either raft-associated MHC class II is devoid of Ii or the immunoprecipitate contains MHC class II αβI complexes possessing mature, processed forms of Ii-p33 (indicated as Ii-p33). By 16 h (a time point in which almost all labeled MHC class II is on the cell surface (25)) ~60% of newly synthesized MHC class II is associated with lipid rafts (Fig. 2A), a value that is essentially identical to that of the total pool of membrane-associated class II (as determined by Western blotting) (Fig. 2B). The loss of [35 S]methionine signal at the 16 h chase point is a consequence of the dissociation of the methionine-rich Ii from the MHC class II αβI complex. These studies demonstrate that MHC class II was initially present in a detergent soluble membrane domain and acquired detergent insolubility during intracellular transport.

![FIGURE 1](https://example.com/figure1.png)

**FIGURE 1.** Detergent insolubility of MHC class II in Brij-58. JY B cells (5 x 10⁶) were either mock-treated (A) or treated with 10 mM MCD (B) for 12 min. Following treatment, cells were washed in HBSS, lysed in 1% Brij-58, and analyzed by sucrose density gradient centrifugation as described in Materials and Methods. Equal portions of each fraction were analyzed by immunoblotting using Abs recognizing MHC class II α-chains (mAb DA6.147), CD59, ICAM-1, or calnexin. C, The percentage of the total fraction of each protein partitioning with the detergent-insoluble lipid raft fractions was determined by quantitative immunoblot analysis. The graph represents combined data from the total number of experiments indicated, where n is equal to the number of experiments for the particular data point.

![FIGURE 2](https://example.com/figure2.png)

**FIGURE 2.** MHC class II traffics into rafts during biosynthesis. For each time point indicated, 75 x 10⁶ JY B cells were pulsed with [35 S]methionine for 10 min at 37°C. Aliquots of the incorporation were then harvested immediately (Pulse) or chased in complete media containing excess non-radioactive methionine for either 3 h or for 16 h. Cells were harvested, lysed, and rafts were isolated as described in Materials and Methods. Following raft fractionation, raft (R) and nonraft soluble (S) fractions were pooled and subjected to immunoprecipitation using a mixture of MHC class II-specific mAbs (DA6.147 and XD5.411). Immunoprecipitates were boiled and reduced in sample buffer containing 2-ME before loading. A, Immunoprecipitates were analyzed by SDS-PAGE and autoradiography to visualize newly synthesized [35 S]methionine-labeled MHC class II. Aliquots of these same immunoprecipitates were analyzed by immunoblotting using a MHC class II α-chain specific mAb (DA6.147) to determine the total amount MHC class II present in each sample under steady state conditions. C, The percentage of class II molecules partitioning with lipid raft microdomains (as a fraction of the total amount of class II present in all gradient fractions) was determined by quantitative densitometry of either immunoblots (for total MHC class II) or autoradiograms for [35 S]methionine-labeled class II. The graph represents the results of four independent experiments.
Mature MHC class II-Ii complexes are enriched in lipid rafts

During transit from the ER to the plasma membrane, MHC class II αβ complexes traverse the Golgi apparatus and ultimately enter lysosome-like Ag processing compartments (14). In these compartments, Ii is proteolytically degraded, dissociates from the MHC class II αβ complex, and antigenic peptides load onto nascent MHC class II (14). We next examined whether newly synthesized raft-associated MHC class II remains associated with Ii as αβI complexes, or if MHC class II only associates with lipid rafts after Ii dissociation. To determine the subunit composition of newly synthesized raft and nonraft MHC class II, we analyzed MHC class II immunoprecipitates from cells metabolically labeled and chased for either 3 h or 16 h by two-dimensional PAGE (NEPHGE followed by SDS-PAGE). After 3 h of chase, [35S]methionine-labeled, raft-associated MHC class II molecules remained associated with Ii, revealing the presence of MHC class II αβI complexes in lipid rafts (Fig. 3A). Two-dimensional PAGE demonstrated that the carbohydrate moieties present on raft-associated class II α-, β-, p33Ii-, and p35Ii chains were sialylated and fully mature, revealing that these protein complexes had entered the trans-cisternae of the Golgi apparatus (Fig. 3A). Notice also the relatively low abundance of high-mannose (unprocessed) p33-Ii and p35-Ii in class II immunoprecipitates from the lipid raft fraction. By contrast, carbohydrate moieties present on MHC class II molecules in the detergent-soluble (nonraft) fraction were not fully sialylated and existed predominantly in their high-mannose glycoform, consistent with the location of these complexes in the ER and early Golgi compartments. After 16 h of chase essentially all Ii had dissociated from the newly synthesized MHC class II (25) and the subunits of the class II αβ dimer exist in their completely sialylated form (Fig. 3A).

To further confirm that MHC class II αβ complexes in the soluble fraction were primarily present in the ER, we treated class II immunoprecipitates from pooled raft and soluble fractions with endo H. Endo H removes immature high-mannose carbohydrates from newly formed proteins that have not traversed the medial-cisternae of the Golgi apparatus, where the carbohydrates are processed and become resistant to endo H digestion. Based on the two-dimensional PAGE analyses, we would expect Ii complexed with class II to be endo H-resistant in the raft fraction, but mostly endo H-sensitive in the soluble (nonraft) fraction. As expected, Ii bound to class II in the raft fraction is predominantly endo H-resistant, while in the soluble fraction it is predominantly endo H-sensitive (Fig. 3B). Therefore, we conclude that MHC class II αβI complexes are detergent soluble early in the biosynthetic (secretory) pathway and begin to acquire detergent insolubility after transport through the Golgi apparatus.

MHC class II is raft-associated in lysosomal Ag-processing compartments

To further define the stage in which MHC class II begins to accumulate in lipid rafts we treated cells with either monensin or leupeptin, well-characterized inhibitors of MHC class II trafficking (15, 23, 26, 27). Monensin is an ionophore that inhibits protein transport from the medial- to trans-cisternae of the Golgi apparatus, prevents terminal glycosylation of MHC class II-associated Ii, and results in an accumulation of MHC class II αβI complexes in the medial-Golgi apparatus (23). By contrast, leupeptin is a protease inhibitor that prevents complete Ii proteolysis, and as a consequence class II molecules are retained in lysosome-like Ag-processing compartments (15, 26, 27). Furthermore, incomplete Ii proteolysis generates a 21-kDa leupeptin-induced polypeptide (LIP) fragment of Ii that remains associated with class II and contains a lysosomal-retaining motif, and for this reason class II αβ-LIP complexes reveal peptide-free MHC class II that resides in internal Ag-processing compartments (15). Using these inhibitors we can therefore examine the detergent insolubility of MHC class II “trapped” at various points in the biosynthetic pathway. To restrict our analysis to newly synthesized MHC class II, we once again used a pulse-chase approach. B cells were metabolically labeled with [35S]methionine and chased in the absence or presence of either monensin or leupeptin for 3 h before raft isolation. Once again, very little newly synthesized MHC class II was present in lipid rafts after the initial 10 min pulse labeling, but after 3 h ~50% of newly synthesized MHC class II had acquired detergent insolubility (Fig. 4A). Treatment with monensin almost completely

![FIGURE 3. Mature MHC class II-Ii complexes are present in rafts. A. Pulse-chase radiolabeling of JY B cells, cell lysis, and raft isolation were performed as described in Fig. 2. MHC class II was immunoprecipitated from pools of the raft and the soluble (nonraft) fractions using MHC class II mAb-coupled beads. The beads were washed and then eluted in 8 M urea for 30 min at 37°C. The immunoprecipitates were subjected to two-dimensional PAGE and visualized by autoradiography. The arrows indicate the positions of the MHC class II α-chain, MHC class II β-chain, the high mannose forms of p33-Ii or p35-Ii, and the sialylated/processed form of p33-Ii (p33Ii) or p35-Ii (p35Ii). B, MHC class II was immunoprecipitated from pooled raft and soluble (nonraft) fractions using an anti-MHC class II β-chain rabbit serum and treated with or without endo H as described in Materials and Methods. The samples were analyzed by immunoblotting using an Ii-specific mAb (LL1). The mobility of endo H-resistant and endo H-sensitive Ii is indicated by arrows. The results are representative of three independent experiments.](http://www.jimmunol.org/content/151/11/4542/F3.large.jpg)
prevented newly synthesized MHC class II from acquiring detergent insolubility (Fig. 4A). It should be noted that this short 3-h treatment had very little effect on the distribution of the entire pool of cell-associated class II (Fig. 4, B and C). Unlike monensin, treatment with leupeptin did not inhibit MHC class II association with lipid rafts (Fig. 4A). The inclusion of leupeptin during the 3-h chase resulted in the appearance of the \( \text{Ii} \) breakdown product \( \text{LIP} \) associated with MHC class II, confirming the presence of these complexes in lysosome-like Ag-processing compartments (Fig. 4A). Because class II-LIP complexes are not capable of binding antigenic peptides (28), these data demonstrate that MHC class II acquires detergent insolubility before binding antigenic peptides.

To further determine whether MHC class II was associated with lipid rafts in intracellular Ag-loading compartments, we isolated these compartments by subcellular fractionation on 27% Percoll gradients. The purity of the low-density membranes (which includes plasma membrane) and the high-density lysosome fraction was confirmed by immunoblotting. Analysis of each fraction for MHC class II and HLA-DM (a lysosomal resident protein) revealed that our lysosomal fraction contained >90% of HLA-DM but <2% of the total class II (Fig. 5A). The pooled membrane fractions were then solubilized in Brij-58 and subjected to raft fractionation on sucrose density gradients. Approximately 42% of MHC class II was associated with lipid rafts when purified lysosomal membranes were analyzed on these sucrose gradients (Fig. 5, B and C). This is virtually identical to the amount of MHC class II associated with lipid rafts in the low density membrane fraction (Fig. 5, B and C). (The difference between this value and that obtained using intact cells is likely due to different membrane-detergent ratios used in each procedure.) Therefore, MHC class II

![Figure 4](http://www.jimmunol.org/)

**FIGURE 4.** MHC class II associates with lipid rafts in late Golgi/lysosomal compartments. Following a 10-min pulse radiolabeling with \([\text{35}^{S}]\)methionine, JY B cells were washed and chased for 3 h in complete medium alone (no treatment, NT), medium containing monensin (Mon), or medium containing leupeptin (Leu). As a control, a sample of the 10-min pulse radiolabeled sample (P) was included. Following lipid raft fractionation, raft (R) and soluble/nonraft (S) fractions were pooled and were subjected to immunoprecipitation by MHC class II mAb beads (DA6.147 and XD5.All). A, The immunoprecipitates were analyzed by SDS-PAGE and autoradiography to visualize radiolabeled MHC class II molecules. B, Aliquots of the same immunoprecipitates were analyzed by immunoblotting using the anti-MHC class II \( \alpha \)-chain-specific mAb (DA6.147) to determine the total amount of MHC class II present in each sample under steady state conditions. C, The percentage of class II molecules partitioning with lipid raft microdomains under each condition (as a fraction of the total amount of class II present in all gradient fractions) was determined by quantitative densitometry of either immunoblots (for total MHC class II) or autoradiograms (for \([\text{35}^{S}]\)methionine-labeled class II). The graph represents the results of three independent experiments.

![Figure 5](http://www.jimmunol.org/)

**FIGURE 5.** MHC class II is raft associated in isolated Ag-processing compartments. JY B cells were broken, internal compartments isolated using 27% Percoll gradients, and low density (LD) and high density lysosomal membranes were isolated as described in the Materials and Methods. A, Equal volumes of each pooled fraction were analyzed by immunoblotting using Abs recognizing MHC class II \( \alpha \)-chain (DA6.147), HLA-DM (K-76), and CD59. B, Purified low density membrane and the high density lysosomal membrane pools were solubilized in Brij-58 and lipid rafts were isolated by fractionation on sucrose density gradients. Equivalent portions of each fraction were subjected to immunoblot analysis for MHC class II (DA6.147). C, The percentage of class II molecules partitioning with lipid raft microdomains under each condition (as a fraction of the total amount of class II present in all gradient fractions) was determined by quantitative densitometry of immunoblots. The graph represents the results of three independent experiments.
Association with Ii is not required for MHC class II raft association

Our earlier radiolabeling experiments revealed the presence of MHC class II αβI complexes in lipid rafts (Fig. 3). Therefore, we examined whether expression of Ii was required for initial MHC class II raft association. To address this question, we isolated lipid rafts from HeLa cells expressing MHC class II α- and β-chains in the presence or absence of Ii and analyzed the raft association of MHC class II from each cell type. As expected, the control proteins CD59 and ICAM-1 resided in raft and nonraft (soluble) fractions, respectively (Fig. 7A). The expression of Ii (Fig. 7B) did not significantly change the distribution of MHC class II with detergent-insoluble raft microdomains (Fig. 7, A and C). This data demonstrates that the Ii-dependent sorting of MHC class II to Ag-processing compartments is not strictly required for the association of MHC class II with detergent-insoluble membrane microdomains.

Discussion

In this study, we investigated at which point during the biosynthetic pathway MHC class II enters detergent-insoluble lipid raft microdomains, and whether this event preceded loading of class II with antigenic peptides. We found that upon synthesis, MHC class II αβI complexes were present in detergent-soluble membranes and over time chased into detergent-insoluble lipid raft membranes, while the distribution of the total unlabeled pool of class II in rafts remained stable. This demonstrates that newly synthesized MHC class II is not initially present in lipid rafts. This pattern of movement into detergent-insoluble membranes or lipid rafts following initial synthesis is reminiscent of findings described for another raft resident protein, placental-like alkaline phosphatase, which acquires detergent insolubility and moves into lipid rafts during transport through the Golgi apparatus (31). This is consistent with our findings by two-dimensional PAGE analysis and endo H treatment that MHC class II αβI complexes containing Golgi-processed carbohydrates are enriched in detergent-insoluble lipid rafts, while the carbohydrates on the majority of detergent-soluble (nonraft) αβI complexes have not been processed. Furthermore, addition of monensin, which blocks transport of MHC class II from the medial to trans-Golgi (23), almost completely prevented the acquisition of MHC class II detergent insolubility. Taken together, this data supports the idea that MHC class II must traffic through the medial-Golgi to become raft associated.

Although biochemical isolation of rafts using the criteria of detergent insolubility may not completely reflect the true nature of lipid rafts on living cells, our pulse-chase data clearly demonstrate
that the observed insolubility of MHC class II is not a detergent-induced artifact. Detergent insolubility of newly synthesized MHC class II molecules was observed only after these proteins traversed the medial-Golgi even in cells in which the partitioning of the total pool of class II molecules was unchanged, highlighting this critical point. Therefore, it is important to emphasize that this phenomenon of detergent insolubility requires the trafficking of MHC class II molecules into distinct compartments in the APC.

Three pieces of evidence support the idea that detergent insolubility or raft association of MHC class II occurs before peptide loading. First, we found MHC class II αβ complexes in lipid rafts, although the majority of these αβ complexes did possess late Golgi-processed carbohydrates. This points to raft association before II dissociation in Ag-loading compartments. Second, MHC class II-CLIP complexes were present in lipid rafts, demonstrating that even MHC class II that had not yet been “edited” by HLA-DM was sensitive to treatment of DCs with raft-disrupting MHC class II-peptide complexes formed under these different conditions and the role of endocytosis in regulating DC function, they strongly suggest that the mechanism of antigenic peptide loading (and therefore the effective concentration of specific MHC class II-peptide complexes is rafts) may affect the quality of the signal delivered to the T cell by the APC (affecting T cell polarization) as well as the quality of a signal delivered to the DC from the T cell (affecting DC lysosome tubulation). We propose that internal loading of MHC class II enriches specific MHC class II-peptide complexes in small numbers of distinct detergent-insoluble membrane microdomains, while surface loading by preprocessed peptides does not concentrate specific class II-peptide complexes in the same microdomains (i.e., they are more dispersed).

This leads us to the simple question: what makes MHC class II associate with lipid rafts? In the data presented here, we find that Ii expression does not influence MHC class II raft association (Fig. 6). Additionally, we have previously shown that >50% of surface MHC class II is associated with lipid rafts (11). This is in marked contrast to a recent study showing that expression of Ii in tumor cell lines inhibits the association of surface MHC class II with lipid rafts (35). Other than citing cell type differences or experimental protocol, we have no explanation for the discrepancy between our data and that of Dolan et al. (35).

Biophysical studies have led investigators to propose that individual rafts could contain as many as 60 protein molecules (36) and a recent study suggests that each raft could contain as few as three protein molecules (37). Given such data, it is difficult to envision how a significant concentration of ligands, such as specific MHC class II-peptide complexes, could occur in large enough density to activate a T cell. Nevertheless, disruption of these lipid rafts does indeed abrogate activation of T cells by APCs, even if these APCs are formaldehyde fixed after raft disruption (11). One could argue that while only a very small percentage of surface MHC class II molecules contain the correct peptide for T cell recognition, given enough time the T cell could “scan” the APC for enough MHC class II-peptide complexes that are passively diffusing on the APC plasma membrane to become fully activated. However, it is unlikely that large numbers of class II molecules could diffuse to the T cell:APC contact site given the small amount of time an APC and T cell are engaged in vivo before TCR cross-linking (38). Additionally, the requirement for TCR cross-linking to generate the initial “stop” signal for the T cell (39) would be more likely to occur if the specific MHC class II complexes were preclustered. The data we present here shows the presence of MHC class II preclustered in lipid raft microdomains before peptide loading, thereby facilitating the loading of specific peptides on the MHC class II within an individual raft. It is interesting to speculate that the importance of membrane microdomains in clustering a few specific MHC class II-peptide complexes may be critical at this early time, when it would facilitate the initial cross-linking of the TCR. This engagement could then trigger a signal for the T cell to stop and allow itself to rearrange its cytoskeleton (and membrane organization), thereby allowing time for the T cell to “scan” for enough specific MHC class II-peptide complexes to become fully activated.

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References


