

The role of liposome charge on immune response generated in BALB/c mice immunized with recombinant major surface glycoprotein of *Leishmania* (rgp12)

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[View references \(4\)](#)

Abstract

Liposomes as a lipid-based system have been shown to be an effective adjuvant formulation. In this study, the role of liposome charge in induction of a Th₁ type of immune response and protection against leishmaniasis in BALB/c mice was studied. Liposomes containing rgp12 were prepared by Dehydration-Rehydration Vesicle (DRV) method. Neutral liposomes consisted of dipalmitoylphosphatidylcholine and cholesterol. Positively and negatively charged liposomes were prepared by adding dimethyldioctadecylammonium bromide (DDAB) or dicetyl phosphate (DCP) to the neutral liposome formulation, respectively. Female BALB/c mice were immunized subcutaneously with negatively, positively charged or neutral liposomes encapsulated with rgp12, rgp12 in soluble form or PBS, three times in 2 week intervals. The extent of protection and type of immune response generated were studied in different groups of mice. The group of mice immunized with rgp12 encapsulated in neutral liposomes showed a significantly ($P < 0.01$) smaller footpad swelling upon challenge with *Leishmania* major compared with positively or negatively charged liposomes. The mice immunized with neutral liposomes also showed a significantly ($P < 0.01$) the lowest splenic parasite burden, the highest IgG₁/IgG₂ ratio and IFN- γ production and the lowest IL- ϵ level compared to the other groups. The results indicated that a Th₁ type of immune response was induced in mice immunized with neutral liposomes more efficiently than positively charged liposomes and conversely negatively charged liposomes induced a Th₂ type of immune response. © 2009 Elsevier Inc. All rights reserved.

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Indexed Keywords

EMTREE drug terms: cholesterol; dicetyl phosphate; dimethyldioctadecylammonium bromide; dipalmitoylphosphatidylcholine; gamma interferon; glycoprotein; immunoglobulin G₁ antibody; immunoglobulin G₂ antibody; interleukin ϵ ; *Leishmania* vaccine; liposome; recombinant glycoprotein gp 12; unclassified drug

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MeSH: Adjuvants, Immunologic; Animals; Antibodies, Protozoan; Antigens, Protozoan; Drug Carriers; Electrophoresis, Polyacrylamide Gel; Female; Immunization; Immunoglobulin G; Interferon-gamma; Interleukin- ϵ ; *Leishmania* major; Leishmaniasis, Cutaneous; Liposomes; Membrane Glycoproteins; Mice; Protozoan Proteins;