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olidin-2-yl)methyl]-Nchloroacetylbenzamides with keyhole limpet hemocyanin. Conjugates of N-[(pyrrolidin-2-yl)methyl]-Nchloroacetylbenzamides of type (II) and N-[(pyrrolidin-2-yl)methyl]-Nchloroacetylmethoxybenzamides of types (IV) and (V) with keyhole limpet hemocyanin (KLH) were prepared in order to study the immunogenic character of the conjugates. Routine for the synthesis of these conjugates included the cleavage of the benzyl esters with hydrazine and the conjugation with KLH. The immunogenic character of

the conjugates was assessed by miceimmunization. Following the immunization of the conjugates to mice IgM titers were determined in the serum of the mice and complete antisera against each of the conjugates were obtained. The conjugate of N-[(pyrrolidin-2-yl)meth yl]-N-chloroacetylbenzamide of type (II) reacted with KLH more effectively than other conjugates in immunizing the mice. The highest IgM titer of these conjugates were obtained when the reaction was performed at pH 9.5. The results indicated that the hydrophobic

characteristics of the amino group of the side chains of the benzamides and the length of the spacer between the carboxylic acid and the aromatic rings in the benzamides were critical factors for the binding of the compounds to KLH and the conjugating efficiency of the compounds with KLH. The conjugate of N-[(pyrrolidin-2-yl)methyl]-N

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errors. The software costs money in order to maintain production and support the programmers. If we can't ship any longer, do not ask us for the refund. If you like our product, use it! There is no refund if you do not like the software. The present invention relates to a biocompatible polymer of monomers, such as methyl methacrylate and 2-hydroxyethyl methacrylate with reticulated pores, having a high polar index, a high ion exchange capacity and low capacity for water, the polymer being usable as a biomaterial, preferably for implantation in the body, particularly

as a bone substitute. For a long time, the development of new biomaterials for orthopedic surgery has been of considerable interest to provide the surgeon with an autologous substitute for bone, in order to avoid the disadvantages connected with the use of alloplastic materials, such as the risk of implant rejection, secondary surgery, increased costs of medical care, and the like. Consequently, a great number of materials, principally biocompatible polymers of low elasticity, such as, for example, polyand copolymers of 2-hydroxyethyl methacrylate (HEMA), which are

resorbed slowly, have been used. Such polymers are associated with the disadvantage that their in-vivo resorption can take up to about two years, so that not only is the desired substitution effect not achieved but also the implant may be resorbed to such an extent that it is no longer suitable for its intended purpose or for new purposes. Thus, for example, the use of a HEMA polymer in bone, both for substitution and for the production of implants, has the disadvantage that its low elasticity results in bone fractures at relatively low forces (e.g., less than

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