Peg-urate oxidase conjugates and use thereof
Peg-urate oxidase conjugates and use thereof
Conjugés peg-urate oxydase et utilisation associée

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• SAVOCA K V ET AL: "Induction of tolerance in mice by uricase and monomethoxypolyethylene glycol-modified uricase." INTERNATIONAL ARCHIVES OF ALLERGY AND APPLIED IMMUNOLOGY, 75 (1) 58-67., 1984, XP002125212
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The present invention relates to chemical modification of proteins to prolong their circulating lifetimes and reduce their immunogenicity. More specifically, the invention relates to conjugation of poly(ethylene glycols) or poly(ethylene oxides) to urate oxidases, which substantially eliminates urate oxidase immunogenicity without compromising its uricolytic activity.

Urate oxidases (uricases; E.C. 1.7.3.3) are enzymes that catalyze the oxidation of uric acid to a more soluble product, allantoin, a purine metabolite that is more readily excreted. Humans do not produce enzymatically active uricase, as a result of several mutations in the gene for uricase acquired during the evolution of higher primates. Wu, X, et al., (1992) J Mol Evol 34:78-84. As a consequence, in susceptible individuals, excessive concentrations of uric acid in the blood (hyperuricemia) and in the urine (hyperuricosuria) can lead to painful arthritis (gout), disfiguring urate deposits (tophi) and renal failure. In some affected individuals, available drugs such as allopurinol (an inhibitor of uric acid synthesis) produce treatment-limiting adverse effects or do not relieve these conditions adequately. Hande, KR, et al., (1984) Am J Med 76:47-56; Fam, AG, (1990) Baillière's Clin Rheumatol 4:177-192. Injections of uricase can decrease hyperuricemia and hyperuricosuria, at least transiently. Since uricase is a foreign protein in humans, however, even the first injection of the unmodified protein from Aspergillus flavus has induced anaphylactic reactions in several percent of treated patients (Pui, C-H, et al., (1997) Leukemia 11:1813-1816), and immunologic responses limit its utility for chronic or intermittent treatment. Donadio, D, et al., (1981) Nouv Presse Med 10:711-712; Leaustic, M, et al., (1983) Rev Rhum Mal Osteoartic 50:553-554.


In nearly all of the reported attempts to PEGylate uricase (i.e. to covalently couple PEG to uricase), the PEG was attached primarily to amino groups, including the amino-terminal residue and the available lysine residues. In the uricases commonly used, the total number of lysines in each of the four identical subunits is between 25 (Aspergillus flavus (U.S. Patent 5,382,518)) and 29 (pig (Wu, X, et al., (1989) Proc Natl Acad Sci USA 86:9412-9416)). Some of the lysines are unavailable for PEGylation in the native conformation of the enzyme. The most common approach to reducing
the immunogenicity of uricase has been to couple large numbers of strands of low molecular weight PEG. This has invariably resulted in large decreases in the enzymatic activity of the resultant conjugates.

Previous investigators have used injected uricase to catalyze the conversion of uric acid to allantoin in vivo. See Pui, et al., (1997). This is the basis for the use in France and Italy of uricase from the fungus Aspergillus flavus (Uricozyme®) to prevent or temporarily correct the hyperuricemia associated with cytotoxic therapy for hematologic malignancies and to transiently reduce severe hyperuricemia in patients with gout. Potaux, L, et al., (1975) Nouv Presse Méd 4:1109-1112; Legoux, R, et al., (1992) J Biol Chem 267:8565-8570; U.S. Patents 5,382,518 and 5,541,098. Because of its short circulating lifetime, Uricozyme® requires daily injections. Furthermore, it is not well suited for long-term therapy because of its immunogenicity.

A single intravenous injection of a preparation of Candida utilis uricase coupled to 5 kDa PEG reduced serum urate to undetectable levels in five human subjects whose average pre-injection serum urate concentration was 6.2 mg/dL, which is within the normal range. Davis, et al., (1981). The subjects were given an additional injection four weeks later, but their responses were not reported. No antibodies to uricase were detected following the second (and last) injection, using a relatively insensitive gel diffusion assay. This reference reported no results from chronic or subchronic treatments of human patients or experimental animals.

A preparation of uricase from Arthrobacter protoformiae coupled to 5 kDa PEG was used to temporarily control hyperuricemia in a single patient with lymphoma whose pre-injection serum urate concentration was 15 mg/dL. Chua, et al., (1988). Because of the critical condition of the patient and the short duration of treatment (four injections during 14 days), it was not possible to evaluate the long-term efficacy or safety of the conjugate.

In this application, the term "immunogenicity" refers to the induction of an immune response by an injected preparation of PEG-modified or unmodified uricase (the antigen), while "antigenicity" refers to the reaction of an antigen with preexisting antibodies. Collectively, antigenicity and immunogenicity are referred to as "immunoreactivity." In previous studies of PEG-uricase, immunoreactivity was assessed by a variety of methods, including: 1) the reaction in vitro of PEG-uricase with preformed antibodies; 2) measurements of induced antibody synthesis; and 3) accelerated clearance rates after repeated injections.

Previous attempts to eliminate the immunogenicity of uricases from several sources by coupling various numbers of strands of PEG through various linkers have met with limited success. PEG-uricases were first disclosed by FF Davis and by Y Inada and their colleagues. Davis, et al., (1978); U.S. Patent 4,179,337; Nishimura, et al., (1979); Japanese Patents 55-99189 and 56-55079. The conjugate disclosed in the ’337 patent was synthesized by reacting uricase of unspecified origin with 2,000-fold molar excess of 750 dalton PEG, indicating that a large number of polymer molecules was likely to have been attached to each uricase subunit. The ’337 patent discloses the coupling of either PEG or poly(propylene glycol) with molecular weights of 500 to 20,000 daltons, preferably about 500 to 5,000 daltons, to provide active, water-soluble, non-immunogenic conjugates of various polypeptide hormones and enzymes including oxidoreductases, of which uricase is one of three examples. In addition, the ’337 patent emphasizes the coupling of 10 to 100 polymer strands per molecule of enzyme, and the retention of at least 40% of enzymatic activity. No test results were reported for the extent of coupling of PEG to the available amino groups of uricase, the residual specific uricolytic activity, or the immunoreactivity of the conjugate.

Data from 13 citations relating to PEGylation of uricase are summarized in Table 1. Some of these results are also presented graphically in Figures 1A-2B. Seven of these publications describe significant decreases in uricolytic activity measured in vitro caused by coupling various numbers of strands of PEG to uricase from Candida utilis. Coupling a large number of strands of 5 kDa PEG to porcine liver uricase gave similar results, as described in both the Chen publication and a symposium report by the same group. Chen, et al., (1981); Davis, et al., (1978).

Among the studies summarized in Table 1, the immunoreactivity of uricase was reported to be decreased by PEGylation in seven of them and eliminated in five of them. In three of the latter five studies, the elimination of immunoreactivity was associated with profound decreases in uricolytic activity - to at most 15%, 28%, or 45% of the initial activity. Nishimura, et al., (1979) (15% activity); Chen, et al., (1981) (28% activity); Nishimura, et al., (1981) (45% activity). In the fourth report, PEG was reported to be coupled to 61% of the available lysine residues, but the residual specific activity was not stated. Abuchowski, et al., (1981). However, a research team that included two of the same scientists and used the same methods reported elsewhere that this extent of coupling left residual activity of only 23-28%. Chen, et al., (1981). The 1981 publications of Abuchowski et al., and Chen et al., indicate that to reduce the immunogenicity of uricase substantially, PEG must be coupled to approximately 60% of the available lysine residues (Table 1). The fifth publication in which the immunoreactivity of uricase was reported to have been eliminated does not disclose the extent of PEG coupling, the residual uricolytic activity, or the nature of the PEG-protein linkage. Veronese, FM, et al., (1997) in JM Harris, et al., (Eds.), Poly(ethylene glycol) Chemistry and Biological Applications. ACS Symposium Series 680 (pp. 182-192) Washington, DC: American Chemical Society.

Conjugation of PEG to a smaller fraction of the lysine residues in uricase reduced but did not eliminate its immunoreactivity in experimental animals. Tsuji, J, et al., (1985) Int J Immunopharmacol 7:725-730 (28-45% of the amino groups coupled); Yasuda, Y. et al., (1990) Chem Pharm Bull 38:2053-2056 (38% of the amino groups coupled). The
residual uricolytic activities of the corresponding adducts ranged from <33% (Tsuji, et al.) to 60% (Yasuda, et al.) of their initial values. Tsuji, et al., synthesized PEG-uricase conjugates with 7.5 kDa and 10 kDa PEGs, in addition to 5 kDa PEG. All of the resultant conjugates were somewhat immunogenic and antigenic, while displaying markedly reduced enzymatic activities (Table I; Figures 1A-1B).

[0015] A PEGylated preparation of uricase from Candida utilis that was safely administered twice to each of five humans was reported to have retained only 11% of its initial activity. Davis, et al., (1981). Several years later, PEG-modified uricase from Arthrobacter protoformiae was administered four times to one patient with advanced lymphoma and severe hyperuricemia. Chua, et al., (1988). While the residual activity of that enzyme preparation was not measured. Chua, et al., demonstrated the absence of anti-uricase antibodies in the patient’s serum 26 days after the first PEG-uricase injection, using an enzyme-linked immunosorbent assay (ELISA).

[0016] As summarized in Table 1, previous studies of PEGylated uricase show that catalytic activity is markedly depressed by coupling a sufficient number of strands of PEG to decrease its immunoreactivity substantially. Furthermore, most previous preparations of PEG-uricase were synthesized using PEG activated with cyanuric chloride, a triazine derivative (2,4,6-trichloro-1,3,5-triazine) that has been shown to introduce new antigenic determinants and to induce the formation of antibodies in rabbits. Tsuji, et al., (1985).

**TABLE I: Characteristics of PEG-uricases from Previous Studies**

<table>
<thead>
<tr>
<th>Source of Uricase</th>
<th>Coupling Linkage</th>
<th>Molecular Weight of PEG (kDa)</th>
<th>Percent of Lysines with PEG Attached</th>
<th>Residual Uricolytic Activity (%)</th>
<th>Antigenicity or Immunogenicity Comments</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Not reported</td>
<td>Azide</td>
<td>0.7 (diol)</td>
<td>Not reported</td>
<td>Not reported</td>
<td>Antigenicity with rabbit serum (% of that of the unmodified enzyme)</td>
<td></td>
</tr>
<tr>
<td>Candida utilis</td>
<td>Triazine (Cyanuric chloride)</td>
<td>5</td>
<td>% of -98-:</td>
<td></td>
<td>70%</td>
<td>Nishimura, et al., 1979</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>20</td>
<td>31</td>
<td>6%</td>
<td></td>
</tr>
<tr>
<td>Candida utilis</td>
<td>PEG₂₅ triazine</td>
<td>2 x 5</td>
<td>22</td>
<td>87</td>
<td>86%</td>
<td>Nishimura, et al., 1981</td>
</tr>
<tr>
<td>Candida utilis</td>
<td>Triazine</td>
<td>5</td>
<td>71</td>
<td>11</td>
<td>Five men tolerated two injections in 30 days.</td>
<td>Davis, et al., 1981</td>
</tr>
<tr>
<td>Candida utilis</td>
<td>Triazine according to Chen, et al., 1981</td>
<td>5</td>
<td>49</td>
<td>Not reported</td>
<td>Similar immunogenicity in birds to native uricase immunogenicity negative</td>
<td>Abuchowski, et al., 1981</td>
</tr>
<tr>
<td>Porcine liver</td>
<td></td>
<td>5</td>
<td>37</td>
<td>60</td>
<td>Accelerated clearance in mice Constant Clearance (half-life ca. 8 hours)</td>
<td>Chen. et al.: 1981</td>
</tr>
<tr>
<td>Source of Uricase</td>
<td>Coupling Linkage</td>
<td>Molecular Weight of PEG (kDa)</td>
<td>Percent of Lysines with PEG Attached</td>
<td>Residual Uricolytic Activity (%)</td>
<td>Antigenicity or Immunogenicity Comments</td>
<td>Reference</td>
</tr>
<tr>
<td>------------------</td>
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<td>-------------------------------------</td>
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<td>----------------------------------------</td>
<td>-----------</td>
</tr>
<tr>
<td>Candida utilis</td>
<td>Triazine</td>
<td>5</td>
<td>Not reported</td>
<td>Not reported</td>
<td>PEG-uricase was given orally to chickens in liposomes (ance).</td>
<td>Tsuji, et al., 1985</td>
</tr>
<tr>
<td>Candida utilis</td>
<td>Triazine</td>
<td>5</td>
<td>57</td>
<td>23</td>
<td>Imnenogenicity reduced. but positive in rabbits. (Antibodies are not to uricase: they cross react with PEG-superoxide dismutase.) Antigenicity tested with guinea pig antibodies was reduced.</td>
<td>Chua, et al., 1988</td>
</tr>
<tr>
<td>Arthrobacter protoformiae</td>
<td>Not reported</td>
<td>5</td>
<td>Not reported</td>
<td>Not reported</td>
<td>No antibodies were detected by EUSA 26 days after the first of four PEG-uricase injections.</td>
<td></td>
</tr>
<tr>
<td>Candida utilis</td>
<td>PEG₂ triazine</td>
<td>2 x 5</td>
<td>57</td>
<td>23</td>
<td>Not reported Not reported Not reported Antigenicity tested with rabbit serum was reduced by 75%.</td>
<td>Yasuda, et al., 1990</td>
</tr>
</tbody>
</table>
Japanese Patent 3-148298 to A Sano, et al., discloses modified proteins, including uricase, derivatized with PEG having a molecular weight of 1-12 kDa that show reduced antigenicity and "improved prolonged" action, and methods of making such derivatized peptides. However, there are no disclosures regarding strand counts, enzyme assays, biological tests or the meaning of "improved prolonged." Japanese Patents 55-99189 and 62-55079, both to Y Inada, disclose uricase conjugates prepared with PEG-triazine or bis-PEG-triazine (denoted as PEG2 in Table 1), respectively. See Nishimura, et al., (1979 and 1981). In the first type of conjugate, the molecular weights of the PEGs were 2 kDa and 5 kDa, while in the second, only 5 kDa PEG was used. Nishimura, et al., (1979) reported the recovery of 15% of the uricolytic activity after modification of 43% of the available lysines with linear 5 kDa PEG, while Nishimura, et al., (1981) reported the recovery of 31% or 45% of the uricolytic activity after modification of 46% or 36% of the lysines, respectively, with PEG2.

<table>
<thead>
<tr>
<th>Source of Uricase</th>
<th>Coupling Linkage</th>
<th>Molecular Weight of PEG (kDa)</th>
<th>Percent of Lysines with PEG Attached</th>
<th>Residual Uricolytic Activity (%)</th>
<th>Antigenicity or Immunogenicity Comments</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Candida utilis</td>
<td>PEG2 triazine</td>
<td>2 x 5</td>
<td>22</td>
<td>68</td>
<td>Single injection. PEG increased the half-life from ca. 1 h to ca. 8 h in mice. PEG blocked clearance by liver, spleen and kidney (24-h study duration).</td>
<td>Fuiita, et al., 1991</td>
</tr>
<tr>
<td>Not reported</td>
<td>PEG2</td>
<td>Not reported</td>
<td>Not reported</td>
<td>Not reported</td>
<td>Immunogenicity in mice was decreased by 98% (PEG) or 100% (PEG2).</td>
<td>Veronase, et al., 1997</td>
</tr>
</tbody>
</table>

Summary of the Invention

Previous studies teach that when a significant reduction in the immunogenicity and/or antigenicity of uricase is achieved by PEGylation, it is invariably associated with a substantial loss of uricolytic activity. The safety, convenience and cost-effectiveness of biopharmaceuticals are all adversely impacted by decreases in their potencies and the resultant need to increase the administered dose. Thus, there is a need for a safe and effective alternative means for lowering elevated levels of uric acid in body fluids. Including blood and urine. The present disclosure provides a substantially non-immunogenic PEG-uricase that retains all or nearly all of the uricolytic activity of the unmodified enzyme.

One embodiment of the present invention is a conjugate of urate oxidase (uricase) as defined in claim 1. Orber embodiments are defined in claim 2 to 6. The uricase of this aspect of the invention may be recombinant. Whether recombinant or not, the uricase may be of mammalian origin. In one aspect of this embodiment, the uricase may be porcine, bovine or ovine liver uricase. In another aspect of this embodiment, the uricase may be chimeric. The chimeric uricase may contain portions of porcine liver and/or baboon liver uricase. For example, the chimeric uricase may be pig-baboon chimeric uricase (PBC uricase) or porcine uricase containing the mutations R291K and T301S (PKS uricase) (see sequences in Figure 6 and results of physiological and immunological studies in Figures 7-12). Alternatively, the uricase may be baboon liver uricase in which tyrosine 97 has been replaced by histidine, whereby the specific activity of the uricase may be increased by at least about 60%. The uricase of the invention, whatever the origin, may also be in a form that is truncated, either at the amino terminal, or at the carboxyl terminal, or at both terminals. Likewise, the uricase may be fungal or microbial uricase. In one aspect of this embodiment, the fungal or microbial uricase may be a naturally occurring or recombinant form of uricase from Aspergillus flavus, Arthrobacter globiformis or Candida utilis. Alternatively, the uricase may be an invertebrate uricase, such as, for example, a naturally occurring or recombinant...
form of uricase from *Drosophila melanogaster* or *Drosophila pseudoobscura*. The uricase of the invention may also be a plant uricase, for example, a naturally occurring or recombinant form of uricase from soybean root nodule (*Glycine max*). The PEG may have an average molecular weight between about 5 kDa and 100 kDa; preferably the PEG may have an average molecular weight between about 10 kDa and 60 kDa; more preferably, the PEG may have an average molecular weight between about 20 kDa and about 40 kDa, such as, for example, 30 kDa. The average number of covalently coupled strands of PEG may be 2 to 10 strands per uricase subunit; preferably, the average number of covalently coupled strands may be 3 to 8 per subunit: more preferably, the average number of strands of PEG may be 4 to 6 per subunit. In one aspect of this embodiment the uricase may be tetrameric. The strands of PEG may be covalently linked to uricase via urethane (carbamate) linkages, secondary amine linkages, and/or amide linkages. When the uricase is a recombinant form of any of the uricases mentioned herein, the recombinant form may have substantially the sequence of the naturally occurring form.

Another embodiment of the present invention is a pharmaceutical composition as defined in claim 6. The composition may be stabilized by lyophilization and also may dissolve promptly upon reconstitution to provide solutions suitable for parenteral administration.

The present disclosure also provides the use of PEG uricase for lowering uric acid levels in body fluids and tissues of a mammal. The use includes administering to a mammal an effective uric acid-lowering amount of PEG-uricase. The PEG-uricase may be a purified uricase of two or more subunits in which each subunit may be covalently linked to an average of 2 to 10 strands of linear or branched PEG, wherein each molecule of PEG may have a molecular weight between about 5 kDa and 100 kDa, in a pharmaceutically acceptable carrier. The mammal may be a human.

A method may include the steps of: applying the solution to at least one separation column at a pH between about 9 and 10.5, such as, for example, 10.2; recovering fractions of the eluate and identifying those that may contain isolated tetrameric uricase, wherein the fractions are substantially free of uricase aggregates; and pooling the fractions of the isolated tetrameric uricase. The separation column may be based on ion exchange, size exclusion, or any other effective separation property. The method may also include analysis of the fractions to determine the presence of tetrameric uricase and/or the absence of uricase aggregates. For example, such analysis may include high performance liquid chromatography (HPLC), other chromatographic methods, light scattering, centrifugation and/or electrophoresis. In one aspect of this embodiment, the purified tetrameric uricase may contain less than about 10% uricase aggregates.

**Brief Description of the Drawings**

Figure 1A shows the retention of activity by PEGylated uricase from *Candida utilis* as a function of the number of strands of PEG coupled per subunit.

Figure 1B shows the retention of activity by PEGylated uricase from *Candida utilis* as a function of the total mass of PEG coupled per subunit.

Figure 2A shows the retention of activity by PEGylated uricase from porcine liver as a function of the number of strands of PEG coupled per subunit.

Figure 2B shows the retention of activity by PEGylated uricase from porcine liver as a function of the total mass of PEG coupled per subunit.

Figure 3A shows the retention of activity by PEGylated pig-baboon chimeric (PBC) uricase as a function of the number of strands coupled per subunit.

Figure 3B shows the retention of activity by PEGylated PBC uricase as a function of the total mass of PEG coupled per subunit.

Figure 4A shows the retention of activity by PEGylated uricase from *Aspergillus flavus* as a function of the number of strands of PEG coupled per subunit.

Figure 4B shows the retention of activity by PEGylated uricase from *Aspergillus flavus* as a function of the total mass of PEG coupled per subunit.

Figure 5A shows the retention of activity by PEGylated recombinant soybean root nodule uricase as a function of the number of strands of PEG coupled per subunit.

Figure 5B shows the retention of activity by PEGylated recombinant soybean root nodule uricase as a function of the total mass of PEG coupled per subunit.

Figure 6 shows the deduced amino acid sequences of pig-baboon chimeric uricase (PBC uricase), PBC uricase that is truncated at both the amino and carboxyl terminals (PBC-NT-CT) and porcine uricase containing the mutations
The present disclosure provides improved conjugates of water-soluble polymers, preferably poly(ethylene glycols) or poly(ethylene oxides), with uricases. The invention also provides pharmaceutical compositions of the improved conjugates. These conjugates are substantially non-immunogenic and retain at least 75%, preferably 85%, and more preferably 95% or more of the uricolytic activity of the unmodified enzyme. Uricases suitable for conjugation to water-soluble polymers include naturally occurring urate oxidases isolated from bacteria, fungi and the tissues of plants and animals, both vertebrates and invertebrates, as well as recombinant form of uricase, including mutated, hybrid, and/or truncated enzymatically active variants of uricase. Water-soluble polymers suitable for use in the present disclosure include linear and branched poly(ethylene glycols) or poly(ethylene oxides), all commonly known as PEGs. Examples of branched PEG are the subject of U.S. Patent 5,643,575. One preferred example of linear PEG is monomethoxyPEG of the general structure CH$_3$O-(CH$_2$CH$_2$O)$_n$H, where n varies from about 100 to about 2,300.

One preferred mammalian uricase is recombinant pig-baboon chimeric uricase, composed of portions of the sequences of pig liver and baboon liver uricase, both of which were first determined by Wu, et al., (1989). One example of such a chimeric uricase contains the first 225 amino acids from the porcine uricase sequence (SEQ ID NO: 1) and the last 79 amino acids from the baboon uricase sequence (SEQ ID NO: 2) (pig-baboon uricase, or PBC uricase; see Figure 6). Another example of such a chimeric uricase contains residues 7-225 of the porcine sequence (SEQ ID NO: 1) and residues 226-301 of the baboon sequence (SEQ ID NO: 1) and residues 226-301 of the baboon sequence (SEQ ID NO: 2); this is equivalent to PBC uricase that is truncated at the first 288 amino acids from the porcine sequence (SEQ ID NO: 1) and the last 16 amino acids from the baboon sequence (SEQ ID NO: 2). Since the latter sequence differs from the porcine sequence at only two positions, having a lysine (K) in place of arginine at residue 291 and a serine (S) in place of threonine at residue 301, this mutant is referred to as pig-K-S or PKS uricase. PKS, PBC and PBC-NT-CT uricases each have one more lysine residue and, hence, one more potential site of PEGylation than either the porcine or baboon sequence.


In one embodiment of the invention, uricase may be conjugated via a biologically stable, nontoxic, covalent linkage to a relatively small number of strands of PEG. Such linkages may include urethane (carbamate) linkages, secondary amine linkages, and amide linkages. Various activated PEGs suitable for such conjugation are available commercially from Shearwater Polymers, Huntsville, AL.

For example, urethane linkages to uricase may be formed by incubating uricase in the presence of the succinimidyl carbonate (SC) or 4-nitrophenyl carbonate (NPC) derivative of PEG. SC-PEG may be synthesized using the procedure described in U.S. Patent 5,612,460, which is hereby incorporated by reference. NPC-PEG may be synthesized by reacting PEG with 4-nitrophenyl chloroformate according to methods described in Veronese. FM, et al., (1985) Appl Biochem Biotechnol 11:141-152, and in U.S. Patent 5,286,637, which is hereby incorporated by reference. The methods described in the ’637 patent are adapted to PEGs of higher molecular weight by adjusting the concentrations of the reactants to maintain similar stoichiometry. An alternative method of synthesis of NPC-PEG is described by Boner, W, et al., East German Patent Specification DD 279 486 A1.

Amide linkages to uricase may be obtained using an N-hydroxysuccinimide ester of a carboxylic acid derivative
of PEG (Shearwater Polymers). Secondary amine linkages may be formed using 2,2,2-trifluoroethanesulfonyl PEG (tresyl PEG; Shearwater Polymers) or by reductive alkylation using PEG aldehyde (Shearwater Polymers) and sodium cyanoborohydride.

[0030] In conjugates containing PEGs with molecular weights between 5 kDa and 30 kDa, the maximum number of strands of PEG that were coupled per subunit, while retaining at least 75% of the uricolytic activity of the unmodified enzyme, ranged from an average of 2 strands for soybean uricase to more than 10 strands for PBC uricase (see assay conditions in Example 1 and results in Figures 1A-5B). The latter extent of PEGylation corresponds to approximately one third of the total amino groups. In one embodiment of the invention, the average number of strands of PEG coupled per uricase subunit is between 2 and 10. In a preferred embodiment, the average number of strands of PEG coupled per uricase subunit is between 3 and 8. In a more preferred embodiment, the average number of covalently linked strands of PEG per uricase subunit is between 4 and 6. In another embodiment, the molecular weight of PEG used for the coupling reaction is between 5 kDa and 100 kDa, preferably between 10 kDa and 60 kDa, and more preferably between 20 kDa and 40 kDa, such as, for example 30 kDa.

[0031] There are several factors that may affect the choice of the optimal molecular weight and number of strands of PEG for coupling to a given form of uricase. In general, the reduction or elimination of immunogenicity without substantial loss of uricolytic activity may require the coupling of relatively more strands of PEG of lower molecular weight, compared to relatively fewer strands of PEG of higher molecular weight. For example, either 6 strands of 20 kDa PEG per subunit or 4 strands of 30 kDa PEG per subunit might be optimally effective. Likewise, each different form of uricase may have a different optimum with respect to both the size and number of strands. See Figures 1A-5B.

[0032] PEG conjugation rendered all of the tested uricases soluble and stable in buffers at physiological pH, without the addition of a substrate analog or inhibitor, such as 8-azaxanthine that is used as a stabilizer in the fungal uricase (Uricozyme®) sold by Sanofi Winthrop in France and Italy. Two different conjugates of PBC uricase, one containing approximately 6 strands of 10 kDa PEG per subunit and the other containing approximately 2 strands of 19 kDa PEG per subunit, retained significant activity after incubation in mouse serum for more than one month at 37°C. In addition, several of the conjugates of this invention had circulating half-lives in mice that were greater than two days, in contrast to the approximately 8-hour or 24-hour half-lives previously reported for PEG-modified mammalian and microbial uricases. Chen, et al., (1981); Fuertges, F, et al., (1990) J Conrr Release 11: 139-148; Fujita, T, et al., (1991) J Pharmacobiodyn 14:623-629. Longer half-lives of injected protein drugs make them more cost-effective and can lead to improved patient compliance. Prolonged half-life is also indicative of products that are better tolerated by the body.

[0033] When PEG conjugates of PBC uricase were prepared from the purified tetrameric form of the enzyme (four 35 kDa subunits), they displayed profoundly reduced immunogenicity in mice (Figure 7), in contrast to the moderate immunogenicity of PEG conjugates of larger forms of the enzyme (e.g. octamers of the 35 kDa subunit; see Figure 12), and the very high immunogenicity of the unmodified enzyme. Repeated injections of uricase-deficient mice with PEG-uricase of the present invention eliminated their hyperuricemia for more than 2 months and protected the structure and function of their kidneys against uric acid-related damage (Figures 8-11).

[0034] Injections of fully active conjugates of PBC uricase with 10 kDa PEG (Figures 3A-3B) reduced dramatically the hyperuricemia of homozygous, uricase-deficient mice (Figure 8). Uric acid levels in the urine were also reduced dramatically in all uricase-deficient mice treated with PEG-modified PBC uricase. Uricase-deficient mice received a series of injections with a preparation of PEG-uricase similar to that used to obtain the data in Figure 8. This treatment reduced the severity of a urine-concentrating defect, as demonstrated by measurements of urine osmolality under normal conditions and after a 12 - hour period of water deprivation (Figure 9) and by their water consumption and urine output (Figure 10), compared to the corresponding measurements in untreated, genetically similar mice. It was also demonstrated that ten weeks of treatment, starting within the first ten days of life, of homozygous uricase-deficient (uox -/-) "knockout" mice with a PEG-uricase of this invention decreased the severity of urate-induced disruption of the renal architecture, as visualized by magnetic resonance microscopy (Figure 11). For microscopy methods, see Hedlund, LW, et al., (1991) Fund Appl Toxicol 16:787-797; Johnson, GA, et al., (1992) in JC Gore, (Ed.), Reviews of Magnetic Resonance in Medicine. Vol. 4 (pp187-219) New York: Pergamon Press.

[0035] Purified preparations of naturally occurring and recombinant uricases usually contain a mixture of aggregates of the enzyme, in addition to the tetrameric (140 kDa) form. The percentage of each uricase preparation that is in the tetrameric form generally varies from approximately 20% to 90%. Despite evidence that unPEGylated aggregates of several other proteins are highly immunogenic (see, e.g., Moore, WV, et al., (1980) J Clin Endocrinol Metab 51:691-697), previous studies of PEG-uricase do not describe any efforts to limit the content of aggregates, suggesting that the potential immunogenicity of the PEG-modified aggregates was not considered. On the basis of the observations of the present inventors, it appears likely that such aggregates were present in the enzyme preparations used for previous syntheses of PEG-uricase. Their presence may have rendered the task of preparing non-immunogenic conjugates more difficult. It also appears that the large losses of uricolytic activity observed in previous efforts to PEGylate uricase were related to the large number of strands of low molecular weight PEG that were coupled. On the other hand, the methods of uricase purification and PEGylation described herein permit the covalent attachment of as many as 10 strands of
The results presented herein indicate that, even when extensively PEGylated, forms of PBC uricase larger than constituting as little as about 10%, 3%, 2%, or less, of the total isolated uricase. Of the uricase thus pooled, at least 90% may be in the tetrameric form; the undesirable aggregates may thus constitute as little as about 10%, 3%, 2%, or less, of the total isolated uricase.

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In another preferred embodiment, substantially all aggregates of the tetrameric form of the enzyme may be removed by ion-exchange or size-exclusion chromatography at a pH between about 9 and 10.5, preferably 10.2, prior to PEG conjugation of the resulting substantially tetrameric preparation of uricase. The molecular weight of the uricase in each fraction from the preparative column may be monitored by any size-dependent analytical technique, including, for example, HPLC, conventional size-exclusion chromatography, centrifugation, light scattering, capillary electrophoresis or gel electrophoresis in a non-denaturing buffer. For tetrameric uricase isolated using size-exclusion chromatography, fractions containing only the 140 kDa form of the enzyme may be pooled and used for conjugation to PEG. For tetrameric uricase isolated using ion-exchange chromatography, fractions from the ion-exchange column may be analyzed with respect to size to determine which fractions contain substantial amounts of the tetrameric form without detectable aggregate. Of the uricase thus pooled, at least 90% may be in the tetrameric form; the undesirable aggregates may thus constitute as little as about 10%, 3%, 2%, or less, of the total isolated uricase.

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The tetrameric form of uricase (molecular weight ca. 140 kDa) was purified from a solution of porcine liver uricase by preparative size-exclusion or ion-exchange chromatography, followed by analytical size-exclusion chromatography. Porcine liver uricase was obtained from Sigma-Aldrich, St. Louis, MO, catalog No. U2350 or U3377; or Boehringer Mannheim, Indianapolis, IN.

Preparative and analytical size-exclusion chromatography were performed at pH 10-10.5, preferably 10.2, in 10 mM sodium carbonate buffer containing 0.1 M NaCl on Superdex 200 columns that had been previously calibrated with proteins of known molecular weight. Superdex was obtained from Amersham Pharmacia, Piscataway, NJ. Any buffer may be used that is capable of maintaining the desired pH and that is compatible with the chemistry to be used for subsequent PEG coupling. Such buffers are well known in the art. The ultraviolet absorbance of the eluate from the preparative column was monitored at 280 nm, and uricase-containing portions of the eluate corresponding to the molecular weight of the desired tetrameric form, but free of higher molecular weight species, were collected for use in synthesizing substantially non-immunogenic PEG-uricase as described in Example 2. Alternatively, tetrameric forms of uricase can be isolated using other size-exclusion media such as, for example, Superose 12 (Amersham Pharmacia) or any other medium that is compatible with mildly alkaline solutions and that has an appropriate size fractionation range. Such media are readily available and are well known in the art.

Ion-exchange chromatography was performed at pH 10-10.5, preferably 10.2, on Mono Q columns (Amersham Pharmacia, Piscataway, NJ) that had been equilibrated with 0.1 M sodium carbonate buffer. Any buffer that is compatible with the chemistry of PEG coupling and that is capable of maintaining the desired pH may be used at sufficiently low ionic strength to permit the adsorption of uricase to the column. Such buffers are well known in the art. The ultraviolet absorbance of the eluate was monitored at 280 nm during elution of the uricase from the ion-exchange resin by increasing the ionic strength to permit the adsorption of uricase to the column. Such buffers are well known in the art. The ultraviolet absorbance of the eluate was monitored at 280 nm during elution of the uricase from the ion-exchange resin by increasing the ionic strength to permit the adsorption of uricase to the column. Such buffers are well known in the art.
concentrations in the assay of 6-150 μM. Uricase preparations were diluted in this borate buffer containing bovine serum albumin (Sigma-Aldrich, St. Louis, MO, catalog No. A-7030), so that the final concentration of albumin in the assay was 0.1 mg/mL. After mixing various dilutions of the enzyme with the substrate in the wells of a microtiter plate in a microplate reader, the rate of disappearance of uric acid at 25°C was monitored at 292 nm every 4 seconds for 3 minutes. From samples in which between 10% and 40% of the substrate was consumed within 3 minutes, at least 20 data points were used to calculate the maximal rate of decrease in the absorbance per minute. One international unit (IU) of uricase activity is defined as the amount of enzyme that consumes one micromole of uric acid per minute; specific activities are expressed as IU/mg protein. Some of the data for relative uricase activities in Figures 1A-5B were obtained using 100 μM uric acid in the assay. Other results for the velocity at 100 μM uric acid (V100) were calculated from the values of the Michaelis constant (K_M) and the maximal velocity (V_max) for the respective enzyme preparations, using the formula:

\[ V_{100} = 100 \times \frac{V_{\text{max}}}{(K_M + 100)} \]

where K_M is expressed in micromolar units.

EXAMPLE 2

PEG coupling to tetrameric porcine uricase

[0049] To a solution of tetrameric uricase in 0.1 M sodium carbonate buffer, pH 10.2, 10 - 200 moles of an activated derivative of monomethoxyPEG, e.g., the 4-nitrophenyl carbonate (NPC-PEG), of various sizes (5 kDa to 30 kDa) were added for each mole of uricase subunit (molecular weight 35 kDa). These and other suitable activated PEGs are available from Shearwater Polymers. Instructions for coupling these PEGs to proteins are given in the catalog of Shearwater Polymers, on the internet at www.swpolymers.com, and in JM Harris, et al., (Eds.) (1997) Poly(ethylene glycol) Chemistry and Biological Applications. ACS Symposium Series 680, Washington, DC: American Chemical Society. The coupling reaction was allowed to proceed at 0-8°C until the extent of PEG coupling no longer changed significantly with time. Unreacted PEG was then removed from the reaction product by chromatography and/or ultrafiltration.

[0050] The number of strands of PEG coupled per subunit of uricase was determined by an adaptation of the methods described by Kunitani, M, et al., (1991) J Chromatogr 588:125-137; Saifer, et al., (1997) and Sherman, et al., (1997). Briefly, aliquots of the PEGylation reaction mixtures or fractions from the preparative ion-exchange or size-exclusion columns were characterized by analytical size-exclusion HPLC on a TSK 5,000 RWXL column at room temperature in 10 mM sodium carbonate buffer, pH 10.2, containing 0.1 M NaCl. The HPLC column was obtained from Tosohaaas, Montgomeryville, PA. Proteins and PEGs were monitored by ultraviolet absorbance and refractive index detectors. The amount of protein in the conjugate was calculated from the ultraviolet absorbance relative to that of the appropriate unmodified uricase standard. The amount of PEG in the conjugate was then calculated from the area of the refractive index peak, corrected for the contribution of the protein to refractive index, relative to the area of the refractive index peak of the appropriate PEG standard.

[0051] Figure 2A shows the retention of activity by PEGylated porcine liver uricase as a function of the number of strands of PEG coupled per subunit. Data of the present inventors (▲, □) are compared with those of Chen, et al., (1981). The data point within a large circle denotes a conjugate reported to be non-immunoreactive by Chen, et al., (1981). As shown in Figure 2A, conjugates of tetrameric porcine uricase with up to 6 strands of 30 kDa PEG per subunit or up to 7 strands of 5 kDa PEG per subunit retained at least 75% of the activity of the unmodified enzyme. The apparent increase in specific activity with an increasing number of strands of 5 kDa or 30 kDa PEG (up to about 4 strands per subunit) may reflect the relative insolubility or instability of the unmodified enzyme compared to the conjugates. As shown in Figure 2B, conjugates of porcine uricase with an average of more than 3 strands of 30 kDa PEG per subunit contain a greater mass of PEG than was found sufficient to preclude immunoreactivity by Chen, et al., (1981).

EXAMPLE 3

Properties of PEG conjugates of tetrameric recombinant PBC uricase

[0052] Recombinant pig-baboon chimeric (PBC) uricase cDNA was subcloned into the pET3d expression vector (Novagen, Madison, WI) and the resultant plasmid construct was transformed into and expressed in a strain of Escherichia coli BL21(DE3)pLysS (Novagen). These procedures were carried out using methods well known in the art of molecular biology. See Erlich (1989); Sambrook, et al., (1999); Ausubel, F, et al., (Eds.), (1997) Short Protocols in Molecular Biology. New York: John Wiley & Sons.
Figure 6 shows the deduced amino acid sequence of PBC uricase (amino acids 1-225 of SEQ ID NO: 1 and amino acids 226-304 of SEQ ID NO: 2), compared with the porcine (SEQ ID NO: 1) and baboon (SEQ ID NO: 2) sequences. Residues in the baboon sequence that differ from those in the porcine sequence are shown in bold type. The porcine and baboon sequences were first determined by Wu, et al., (1989) and were confirmed by the present inventors. SEQ ID NO. 1 is identical to Accession Number p16164 of GenBank, except for the absence of the initial methionyl residue in the GenBank sequence. SEQ ID NO. 2 is identical to Accession Number p25689 of GenBank, except for the absence of the initial methionyl residue and a change from histidine to threonine at residue 153 in the GenBank sequence (residue 154 in Figure 6).

The tetrmeric form of PBC uricase was isolated and coupled to PEGs of various molecular weights as described in Examples 1 and 2. Conjugates prepared with 5 kDa, 10 kDa, 19 kDa or 30 kDa PEG contained up to 10 strands of PEG per subunit. Those prepared with PEGs of at least 10 kDa retained more than 95% of the initial specific activity of the recombinant uricase (Figures 3A-3B).

The following properties of a conjugate of tetrameric PBC uricase with approximately 6 strands of 10 kDa PEG per subunit are illustrated in the indicated figures: the lack of immunogenicity (Figure 7) and the efficacy in uricase-deficient mice in 1) correcting hyperuricemia and hyperuricosuria (Figure 8); 2) decreasing the severity of a urine-concentrating defect (Figure 9), and 3) decreasing the severity of nephrogenic diabetes insipidus (Figure 10). In addition, this PEG-uricase decreased the severity of uric acid-related renal damage, as visualized by magnetic resonance microscope (Figure 11).

Figure 7 shows the activity of PBC uricase in mouse serum 24 h after each of four or five intraperitoneal injections of PEG-uricase, relative to the value 24 h after the first injection. PEG conjugates were prepared from three different preparations of PBC uricase using two different techniques for PEG activation. One preparation (∗) was tested in uricase-deficient (uox -/-) mice; the other two (∆, ■) were tested in normal BALB/c mice. The most immunoreactive preparation (Δ) was prepared from purified PBC uricase containing an unknown quantity of uricase aggregates coupled to an average of 7 strands of 5 kDa PEG per subunit, using the succinimidyl carbonate derivative of PEG (SC-PEG). Zalipsky, U.S. Patent 5,612,460, hereby incorporated by reference. The moderately immunoreactive preparation (∗) was prepared by coupling a PBC uricase preparation containing 11% aggregates to an average of 2 strands of 19 kDa PEG per subunit, using a 4-nitrophenyl carbonate derivative of PEG (NPC-PEG). Sherman, et al., (1997). The least immunoreactive conjugate (∗) was prepared by coupling an average of 6 strands of 10 kDa NPC-PEG per subunit to a preparation of PBC uricase containing <5% aggregated uricase.

Figure 8 shows the inverse relationship between the concentrations of uric acid in the serum and urine and the activity of injected PEG-uricase in the serum of a uricase-deficient (uox -/-) mouse. Injections at zero time and after 72 h contained 0.43 IU of PBC uricase conjugated to an average of 6 strands of 10 kDa PEG per enzyme subunit.

Figure 9 shows that treatment of uricase-deficient mice with PEG-modified PBC uricase decreased the severity of a urine-concentrating defect. The mean and standard deviation of data for urine osmolality are shown for two mice containing one copy of the normal murine uricase gene (uox +/-), six untreated homozygous uricase-deficient mice (uox -/-) and six homozygous uricase-deficient mice that were injected ten times between the third and 72nd day of life with either 95 or 190 mIU of PEG-uricase. Mice of each genetic background either had received water ad libitum (solid bars) or had been deprived of water for 12 h (hatched bars) prior to collection of their urine.

Figure 10 shows that treatment of uricase-deficient mice with PEG-modified PBC uricase decreased the severity of nephrogenic diabetes insipidus, characterized by abnormally high consumption of water and abnormally high urine output. The genetic backgrounds of the mice and treatment protocol were the same as in Figure 9. The mean and standard deviation of the daily water consumption (solid bars) and urine output (hatched bars) are shown for three groups of six mice.

Figure 11 shows that treatment of uricase-deficient mice with PEG-modified PBC uricase decreased the severity of uric acid-induced nephropathy, as visualized by magnetic resonance microscopy. The genetic backgrounds of the three groups of mice and the treatment protocol were the same as in Figures 9 and 10. Magnetic resonance microscopy was performed at the Center for in vivo Microscopy, Duke University Medical Center, Durham, North Carolina.

In addition to the results summarized in Figures 8-11, it was demonstrated that the uric acid levels in the urine of all uricase-deficient mice decreased dramatically after treatment with PEG-modified PBC uricase. Finally, Figure 12 shows that, unlike the PEG-modified tetrmeric form of PBC uricase, the octameric form (molecular weight = 280 kDa), even when extensively PEGylated, is immunogenic in mice. This property is reflected in the accelerated clearance of the PEG-modified octamer within 5 days after a single intraperitoneal injection. The same mice were re-injected with the same dose of the same PEG-uricase preparations on days 8 and 15. Twenty-four hours after the second and third injections, uricolytic activity was undetectable in the sera of mice injected with the PEGylated octamer, but was readily detected in the sera of those injected with the PEGylated tetramer. These findings, in combination with the accelerated clearance of the PEGylated octamer observed after the first injection (Figure 12), support the utility of removing all forms of uricase larger than the tetramer prior to PEGylation of the enzyme.
EXAMPLE 4

PEG conjugation of uricase from Candida utilis

Uricase from Candida utilis was obtained from either Sigma-Aldrich (St. Louis, MO; catalog No. U1878) or Worthington Biochemical Corporation (Freehold, NJ; catalog No. URYW). Proceeding as described in Examples 1 and 2, the tetrameric form was isolated and PEG conjugates were synthesized with 5 kDa, 10 kDa or 30 kDa PEG (Figures 1A-1B). Figure 1A shows the retention of activity by PEGylated uricase from Candida utilis as a function of the number of strands of PEG coupled per subunit. Data of the present inventors (▲, ●, □) are compared with those of Nishimura, et al. (1979; Nishimura, et al., 1981); Chen, et al. (1981); Davis, et al., (1981); Tsuji, et al., (1985); Yasuda, et al., (1990), and Fujita, et al., (1991). Data points within large circles denote conjugates reported to be non-antigenic by Nishimura, et al., (1979 or 1981) or non-immunoreactive by Chen, et al., (1981).

Figure 1B shows the retention of activity by PEGylated uricase from Candida utilis as a function of the total mass of PEG coupled per subunit. Data of the present inventors (▲, ●, □) are compared with those of the same reports as in Figure 1A. Data points within large circles have the same meaning as in Figure 1A.

As shown in Figures 1A and 1B, conjugates with an average of up to 6 strands of 5 kDa or 30 kDa PEG or 9 strands of 10 kDa PEG per subunit retained at least 75% of the activity of the unmodified enzyme. The apparent increase in specific activity as an increasing number of strands of 30 kDa PEG is attached (up to 5 or 6 strands per subunit) may reflect the relative insolubility or instability of the unmodified enzyme compared to the conjugates.

EXAMPLE 5

PEG conjugation of uricase from Aspergillus flavus

Uricase from Aspergillus flavus was obtained from Sanofi Winthrop (Gentilly Cédex, France). Proceeding as described in Example 2, conjugates with PEGs of various molecular weights were synthesized (Figures 4A-4B). Conjugates prepared by coupling the enzyme from A. flavus with an average of up to 12 strands of 5 kDa PEG or up to 7 strands of 30 kDa PEG per subunit retained at least 75% of the initial specific activity of this fungal uricase.

EXAMPLE 6

PEG conjugation of soybean uricase

Recombinant uricase from soybean root nodule (also called nodulin 35) was prepared and purified as described by Kahn and Tipton (Kahn, K, et al., (1997) Biochemistry 36:4731-4738), and was provided by Dr. Tipton (University of Missouri, Columbia, MO). Proceeding as described in Examples 1 and 2, the tetrameric form was isolated and conjugates were prepared with PEGs of various molecular weights (Figures 5A-5B). In contrast to uricase from Candida utilis (Figure 1A), porcine uricase (Figure 2A), pig-baboon chimeric uricase (Figure 3A) and uricase from Aspergillus flavus (Figure 4A), the soybean enzyme tolerated coupling of only approximately 2 strands of 5 kDa or 30 kDa PEG per subunit with retention of at least 75% of the initial uricolytic activity.

EXAMPLE 7

PEG conjugation of uricase from Arthrobacter globiformis

Uricase from Arthrobacter globiformis was obtained from Sigma-Aldrich (catalog No. U7128). See Japanese Patent 9-154581. Proceeding as described in Examples 1 and 2, the tetrameric form was isolated and conjugates with 5 kDa and 30 kDa PEG were prepared. While conjugates with an average of more than 3 strands of 5 kDa PEG per subunit retained less than 60% of the initial specific activity, conjugates with an average of approximately 2 strands of 30 kDa PEG per subunit retained at least 85% of the initial specific activity.

EXAMPLE 8

PEG conjugation of amino-truncated porcine and PBC uricases

Recombinant porcine and PBC uricases from which the first six amino acids at the amino terminal are deleted are expressed in and purified from E. coli by standard techniques, as described in Example 3. Proceeding as described in Examples 1 and 2, PEG conjugates of the amino-truncated uricases are synthesized to produce substantially non-
immunogenic conjugates that retain at least 75% of the initial specific activity.

EXAMPLE 9

PEG conjugation of porcine and PBC uricases truncated at the carboxyl terminal or both the amino and carboxyl terminals

[0069] Recombinant porcine and PBC uricases from which the last three amino acids at the carboxyl terminal are deleted are expressed in and purified from E. coli by standard techniques, as described in Example 3. This carboxyl-terminal deletion may enhance the solubility of the unmodified enzymes, since it removes the peroxisomal-targeting signal. See Miura, et al., (1994). Proceeding as described in Examples 1 and 2, PEG conjugates of the carboxyl-truncated uricases are synthesized to produce substantially non-immunogenic conjugates that retain at least 75% of the initial specific activity. The sequence of recombinant PBC uricase truncated by six residues at the amino terminal and by three residues at the carboxyl terminal (PBC-NT-CT) is shown in Figure 6. This uricase is expressed, purified and PEGylated as described in Examples 1, 2 and 3 to produce substantially non-immunogenic conjugates that retain at least 75% of the initial specific activity.

EXAMPLE 10

PEG conjugation of porcine uricase mutants containing an increased number of PEG attachment sites

[0070] Recombinant porcine uricases are prepared as described in Example 3, in which the potential number of sites of PEG attachment is increased by replacing one or more arginine residues with lysine. See Hershfield, MS, et al., (1991) Proc Natl Acad Sci USA 88:7185-7189. The amino acid sequence of one example of such a mutant (PKS uricase), in which the arginine at residue 291 is replaced by lysine and the threonine at residue 301 is replaced by serine, is shown in Figure 6. Proceeding as described in Examples 1 and 2, PEG is conjugated to this uricase to produce substantially non-immunogenic conjugates that retain at least 75% of the initial specific activity of the recombinant uricase.

EXAMPLE 11

PEG conjugation of a recombinant baboon uricase mutant

[0071] Using standard methods of molecular biology, as in Example 3, recombinant baboon uricase is constructed having an amino acid substitution (histidine for tyrosine) at position 97 (see baboon sequence in Figure 6). Proceeding as described in Examples 1 and 2, PEG conjugates of the tetrameric form of the recombinant baboon uricase mutant are synthesized to produce conjugates of substantially reduced immunogenicity that retain at least 75% of the initial specific activity of the recombinant uricase.

EXAMPLE 12

Immunogenicity of PEG conjugates from Candida utilis, Aspergillus flavus, and Arthrobacter globiformis

[0072] Uricase from Candida utilis, Aspergillus flavus, and Arthrobacter globiformis are obtained as described in Examples 4, 5, and 7, respectively. Proceeding as described in Examples 1 and 2, PEG conjugates are synthesized with 5 kDa, 10 kDa, 20 kDa or 30 kDa PEG. The immunogenicity of these conjugates is substantially reduced or eliminated.

[0073] Preferred embodiments of the present disclosure are disclosed below and are designated as embodiment E1 to embodiment E8.

E1. A conjugate comprising uricase conjugated to PEG, wherein the PEG has an average molecular weight of about 10,000 to about 100,000 Daltons, and wherein the uricase retains at least about 75% of the uricolytic activity of unconjugated uricase, and wherein the immunogenicity of the uricase is substantially reduced.

E2. The conjugate of E1, wherein said PEG is poly(ethylene glycol).

E3. The conjugate of E2, wherein said poly(ethylene glycol) has an average molecular weight of about 20,000 Daltons.

E4. The conjugate of E2, wherein said poly(ethylene glycol) has an average molecular weight of about 10,000 Daltons.

E5. The conjugate of E2, wherein said poly(ethylene glycol) has an average molecular weight of about 30,000 Daltons.
E6. The conjugate of E1, wherein the uricase is tetrameric.

E7. A pharmaceutical composition for lowering uric acid levels in a body fluid or tissue, comprising the conjugate of E1 and a pharmaceutically acceptable carrier.

E8. A method for isolating a tetrameric form of uricase from a solution of uricase, said solution comprising tetrameric uricase and uricase aggregates, comprising the steps of:

applying said solution to at least one separation column at a pH between about 9.0 and 10.5; and

recovering from said column one or more fractions that contain isolated tetrameric uricase, wherein said one or more fractions are substantially free of uricase aggregates.

SEQUENCE LISTING

[0074]

<110> Williams, L. David Hershfield, Michael S. Kelly, Susan J. Saifer, Mark G.P. Sherman, Merry R.

<120> PEG-URATE OXIDASE CONJUGATES AND USE THEREOF

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 Claims

1. A conjugate comprising a purified uricase conjugated to poly(ethylene glycol) (PEG), wherein at least 90% of said uricase is in a tetrameric form and wherein said PEG has an average molecular weight of about 10,000 to about 60,000 Daltons.

2. The conjugate of claim 1, wherein said PEG has an average molecular weight of about 30,000 Daltons.

3. The conjugate of claim 1, wherein said PEG has an average molecular weight of about 20,000 Daltons.

4. The conjugate of claim 1, wherein said PEG has an average molecular weight of about 10,000 Daltons.

5. The conjugate of claim 1, wherein an average of 2 to 10 strands of PEG are covalently linked to each subunit of said uricase.

6. A pharmaceutical composition suitable for lowering uric acid levels in a body fluid or tissue, comprising the conjugate of any one of claims 1-5.
Patentansprüche

1. Konjugat, umfassend aufgereinigte Uricase, konjugiert an Poly(ethylenglycol) (PEG), wobei zumindest 90 % der Uricase in tetramerer Form vorliegen und wobei das PEG ein mittleres Molekulargewicht von etwa 10.000 bis etwa 60.000 Dalton besitzt.

2. Konjugat nach Anspruch 1, wobei das PEG ein mittleres Molekulargewicht von etwa 30.000 Dalton besitzt.


4. Konjugat nach Anspruch 1, wobei das PEG ein mittleres Molekulargewicht von etwa 10.000 Dalton besitzt.

5. Konjugat nach Anspruch 1, wobei ein Mittelwert von 2 bis 10 PEG-Strängen an jede Untereinheit der Uricase verknüpft sind.

6. Pharmazeutische Zusammensetzung, die zur Verminderung der Harnsäurekonzentrationen in einer Körperflüssigkeit oder einem Gewebe geeignet ist, umfassend das Konjugat nach einem der Ansprüchen 1 bis 5.

Revendications

1. Conjugué comprenant une uricase purifiée conjuguée à du polyéthylène glycol (PEG), au moins 90% de l'uricase étant sous une forme tétramère et le PEG ayant une masse moléculaire moyenne d’environ 10,000 à environ 60,000 Daltons.

2. Conjugué suivant la revendication 1, dans lequel le PEG a une masse moléculaire moyenne d’environ 30,000 Daltons.

3. Conjugué suivant la revendication 1, dans lequel le PEG a une masse moléculaire moyenne d’environ 20,000 Daltons.

4. Conjugué suivant la revendication 1, dans lequel le PEG a une masse moléculaire moyenne d’environ 10,000 Daltons.

5. Conjugué suivant la revendication 1, dans lequel une moyenne de 2 à 10 brins de PEG sont reliés par covalence à chaque sous-unité de l’uricase.

6. Composition pharmaceutique propre à baisser les taux d’acide urique dans un corps fluide ou un tissu corporel, comprenant le conjugué suivant l’une quelconque des revendications 1 à 5.
Figure 1A: Retention of Activity by PEGylated Candida Uricase

- ▲ 30-kDa PEG by MVP
- ● 10-kDa PEG by MVP
- □ 5-kDa PEG by MVP
- X 2 x 5-kDa PEG by Yasuda et al. 1990
- ● 2 x 5-kDa PEG by Nishimura et al. 1981
- ○ 2 x 5-kDa PEG by Fujita et al. 1991
- + 10-kDa PEG by Tsuji et al. 1985
- ■ 5-kDa PEG by Nishimura et al. 1979
- ▲ 5-kDa PEG by Chen et al. 1981
- ● 5-kDa PEG by Davis et al. 1981
- ○ 5-kDa PEG by Tsuji et al. 1985
- X 7.5 kDa PEG by Tsuji et al. 1985
- ○ Non-immunoreactive

Percent of Specific Activity without PEG

PEG Strands Coupled / Uricase Subunit
Figure 2A: Retention of Activity by PEGylated Porcine Uricase

- ▲ 30-kDa PEG by MVP
- □ 5-kDa PEG by MVP
- △ 5-kDa PEG by Chen et al. 1981
- ○ Non-immunoreactive

Percent of Specific Activity without PEG

PEG Strands Coupled / Uricase Subunit
Figure 2B: Retention of Activity by PEGylated Porcine Uricase
Figure 4A: Retention of Activity by PEGylated Uricase® (A. flavus Uricase)

- ▲ 30-kDa PEG by MVP
- □ 5-kDa PEG by MVP

PEG Strands Coupled / Uricase Subunit

Percent of Specific Activity without PEG
Figure 4B: Retention of Activity by PEGylated Uriczyme® (A. flavus Uricase)

Percent of Specific Activity without PEG

Mass of PEG Coupled (kDa) / Uricase Subunit

- ▲ 30-kDa PEG by MVP
- □ 5-kDa PEG by MVP
Figure 5B: Retention of Activity by PEGylated Soybean Uricase

- △ 30-kDa PEG by MVP
- □ 5-kDa PEG by MVP

Percent of Specific Activity Without PEG

Mass of PEG Coupled (kDa)/Uricase Subunit
**Figure 6:** Deduced amino acid sequences of Pig-Baboon Chimeric (PBC) uricase, PBC uricase that is truncated at the amino and carboxyl terminals (PBC-N-T-CT) and Porcine uricase containing the mutations R291K and T301S (PKS uricase), compared with the porcine sequence (SEQ ID NO: 1) and the baboon sequence (SEQ ID NO: 2).

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Figure 7: Serum Uricase Activity 24 Hours after Each PEG-Uricase Injection, Relative to the First Injection

PEG-Uricase (Tetramer)
6 x 10-kDa PEGs/Subunit

uox-/- mice

PEG-Uricase (Tetramer + Aggregates)
2 x 19-kDa PEGs/Subunit

PEG-Uricase
7 x 5-kDa PEGs/Subunit

Relative Serum Uricase Activity

Days after First Injection
Figure 8: Inverse Relationship between Serum PEG-Uricase Activity and Uric Acid Levels in the Serum and Urine of a Uricase-Deficient Mouse.
Figure 9: Decreased Severity of Urine-Concentrating Defect
in Uricase-Deficient Mice Treated with PEG-Uricase

- Water ad libitum
- Water deprived

Urine Osmolality, mOsm/kg

\[ uox^{+/-}, uox^{-/-}, uox^{-/-} \text{ treated} \]
Figure 10: Decreased Severity of Nephrogenic Diabetes Insipidus

In Uricase-Deficient Mice Treated with PEG-Uricase

Water Consumed or Urine Output, mL/day

\[ uox^{+/−} \quad uox^{−/−} \quad uox^{−/−} \text{ treated} \]
*Figure 11:*

Decreased Severity of Uric Acid-Induced Nephropathy after Treatment with PEG-Uricase, as Visualized by Magnetic Resonance Microscopy

Kidney of normal mouse

Kidney of untreated uricase knockout mouse

Kidney of PEG-uricase treated uricase knockout mouse
**Figure 12:** Clearance from the Circulation of BALB/c Mice of PBC Uricase Tetramer and Octamer Coupled to 5-6 Strands of 10-kDa PEG/Subunit
REFERENCES CITED IN THE DESCRIPTION

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