



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<b>(21) International Application Number:</b> PCT/US98/09575  <b>(22) International Filing Date:</b> 12 May 1998 (12.05.98)  <b>(30) Priority Data:</b> 60/046,200                      12 May 1997 (12.05.97)                      US 09/023,809                      13 February 1998 (13.02.98)                      US  <b>(71) Applicant:</b> PHOENIX PHARMACOLOGICS, INC. [US/US]; 115 John Robert Thomas Drive, Exton, PA 19341 (US).  <b>(72) Inventor:</b> CLARK, Mike, A.; 3940 Hall Road, Big Pine, FL 33043 (US).  <b>(74) Agent:</b> ELDERKIN, Dianne, B.; Woodcock Washburn Kurtz Mackiewicz & Norris LLP, 46th floor, One Liberty Place, Philadelphia, PA 19103 (US).		<b>(81) Designated States:</b> AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
<b>(54) Title:</b> MODIFIED ARGININE DEIMINASE		
<b>(57) Abstract</b>  <p>The present invention is directed to arginine deiminase modified with polyethylene glycol, to methods of treating cancer, and to methods of treating and/or inhibiting metastasis.</p>		

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## MODIFIED ARGININE DEIMINASE

### Related Applications

This application claims priority benefit to U.S. Provisional Patent Application Serial No. 60/046,200 filed on May 12, 1997 and U.S. Patent Application  
5 Serial No. 09/023,809 filed February 13, 1998.

### Field of the Invention

The present invention is directed to arginine deiminase modified with polyethylene glycol, to methods for treating cancer, and to methods for treating and/or inhibiting metastasis.

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### Background of the Invention

Malignant melanoma (stage 3) and hepatoma are fatal diseases which kill most patients within one year of diagnosis. In the United States, approximately 16,000 people die from these diseases annually. The incidence of melanoma is rapidly increasing in the United States and is even higher in other countries, such as Australia. The incidence  
15 of hepatoma, in parts of the world where hepatitis is endemic, is even greater. For example, hepatoma is one of the leading forms of cancer in Japan and Taiwan. Effective treatments for these diseases are urgently needed.

Selective deprivation of essential amino acids has been used to treat some forms of cancer. The best known example is the use of L-asparaginase to lower levels of  
20 asparagine as a treatment for acute lymphoblastic leukemia. The L-asparaginase most frequently used is isolated from *E. coli*. However, clinical use of this enzyme is compromised by its inherent antigenicity and short circulating half-life, as described by Y.K. Park, et al, *Anticancer Res.*, 1:373-376 (1981). Covalent modification of *E. coli* L-asparaginase with polyethylene glycol reduces its antigenicity and prolongs its circulating  
25 Phalf-life, as described, for example, by Park, *Anticancer Res.*, *supra*; Y. Kamisaki et al, *J. Pharmacol. Exp. Ther.*, 216:410-414 (1981); and Y. Kamisaki et al, *Gann.*, 73:47-474

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(1982). Although there has been a great deal of effort to identify other essential amino acid degrading enzymes for the treatment of cancer, none have been approved, primarily because deprivation of essential amino acids, by definition, results in numerous, and severe, side effects.

5               It has been reported that enzymes which degrade non-essential amino acids, such as arginine, may be an effective means of controlling some forms of cancer. For example, arginine deiminase (ADI) isolated from *Pseudomonas putida* was described by J.B. Jones, "The Effect of Arginine Deiminase on Murine Leukemic Lymphoblasts," Ph.D. Dissertation, The University of Oklahoma, pages 1-165 (1981). Although effective in  
10   killing tumor cells *in vitro*, ADI isolated from *P. putida* failed to exhibit efficacy *in vivo* because it had little enzyme activity at a neutral pH and was rapidly cleared from the circulation of experimental animals. Arginine deiminase derived from *Mycoplasma arginini* is described, for example, by Takaku et al, *Int. J. Cancer*, 51:244-249 (1992), and U.S. Patent No. 5,474,928, the disclosures of which are hereby incorporated by reference  
15   herein in their entirety. However, a problem associated with the therapeutic use of such a heterologous protein is its antigenicity. The chemical modification of arginine deiminase from *Mycoplasma arginini*, via a cyanuric chloride linking group, with polyethylene glycol was described by Takaku et al., *Jpn. J. Cancer Res.*, 84:1195-1200 (1993). However, the modified protein was toxic when metabolized due to the release of cyanide  
20   from the cyanuric chloride linking group.

              There is a need for compositions which degrade non-essential amino acids and which do not have the problems associated with the prior art. The present invention is directed to these, as well as other, important ends.

### **Summary of the Invention**

25               The present invention is directed to arginine deiminase modified with polyethylene glycol. In a preferred embodiment, the arginine deiminase is modified with polyethylene glycol, having a total weight average molecular weight of about 1,000 to about 50,000, directly or through a biocompatible linking group.

              Another embodiment of the invention is directed to methods of treating  
30   cancer, including, for example, sarcomas, hepatomas and melanomas. The invention is also directed to methods of treating and/or inhibiting the metastasis of tumor cells.

              These and other aspects of the present invention will be elucidated in the

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following detailed description of the invention.

### **Brief Description of the Drawings**

**Figure 1** depicts the amino acid sequences of arginine deiminase cloned from *Mycoplasma arginini* (the top amino acid sequence identified as ADIPROT),

5 *Mycoplasma arthritides* (the middle amino acid sequence identified as ARTADIPRO), and *Mycoplasma hominus* (the bottom amino acid sequence identified as HOMADIPRO).

**Figures 2A and 2B** are graphs showing the effect of a single dose of native arginine deiminase and arginine deiminase modified with polyethylene glycol (*e.g.*, molecular weight 5,000) on serum arginine levels and serum citrulline levels in mice.

10 **Figure 3** is a graph showing the effects on serum arginine levels when PEG10,000 is covalently bonded to ADI via various linking groups.

**Figure 4** is a graph showing the effect that the linking group and the molecular weight of the polyethylene glycol have on citrulline production in mice injected with a single dose of PEG-ADI.

15 **Figures 5A and 5B** are graphs showing the dose response that ADI-SS-PEG5,000 had on serum arginine and citrulline levels. **Figures 5C and 5D** are graphs showing the dose response that ADI-SS-PEG20,000 had on serum arginine and citrulline levels.

**Figure 6** is a graph showing the antigenicity of native ADI, ADI-SS-PEG5,000, and ADI-SS-PEG20,000.

**Figure 7** is a graph showing the effect that treatments with ADI-SS-PEG5,000, ADI-SS-PEG12,000 or ADI-SS-PEG20,000 had on tumor size in mice which were injected with SK-mel 2 human melanoma cells.

25 **Figure 8** is a graph showing the effect that treatments with ADI-PEG20,000 had on tumor size in mice which were injected with SK-mel 28, SK-mel 2 or M24-met human melanoma cells.

**Figure 9** is a graph showing the effect that treatments with ADI-PEG5,000, ADI-PEG12,000 or ADI-PEG20,000 had on the survival of mice which were injected with human hepatoma SK-Hep1 cells.

### 30 **Detailed Description of the Invention**

Normal cells do not require arginine for growth, since they can synthesize arginine from citrulline in a two step process catalyzed by arginosuccinate synthase and

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arginosuccinate lyase. In contrast, melanomas, hepatomas and some sarcomas do not express arginosuccinate synthase; therefore, they are auxotrophic for arginine. This metabolic difference may be capitalized upon to develop a safe and effective therapy to treat these forms of cancer. Arginine deiminase catalyzes the conversion of arginine to  
5 citrulline, and may be used to eliminate arginine. Thus, arginine deiminase may be utilized as a treatment for melanomas, hepatomas and some sarcomas.

Native arginine deiminase may be found in microorganisms and is antigenic and rapidly cleared from circulation in a patient. These problems may be overcome by covalently modifying arginine deiminase with polyethylene glycol (PEG).  
10 Arginine deiminase covalently modified with polyethylene glycol (with or without a linking group) may be hereinafter referred to as "ADI-PEG." When compared to native arginine deiminase, ADI-PEG retains most of its enzymatic activity, is far less antigenic, has a greatly extended circulating half-life, and is much more efficacious in the treatment of tumors.

15 "Polyethylene glycol" or "PEG" refers to mixtures of condensation polymers of ethylene oxide and water, in a branched or straight chain, represented by the general formula  $H(OCH_2CH_2)_nOH$ , wherein  $n$  is at least 4. "Polyethylene glycol" or "PEG" is used in combination with a numeric suffix to indicate the approximate weight average molecular weight thereof. For example, PEG5,000 refers to polyethylene glycol  
20 having a total weight average molecular weight of about 5,000; PEG12,000 refers to polyethylene glycol having a total weight average molecular weight of about 12,000; and PEG20,000 refers to polyethylene glycol having a total weight average molecular weight of about 20,000.

"Melanoma" may be a malignant or benign tumor arising from the  
25 melanocytic system of the skin and other organs, including the oral cavity, esophagus, anal canal, vagina, leptomeninges, and/or the conjunctivae or eye. The term "melanoma" includes, for example, acral-lentiginous melanoma, amelanotic melanoma, benign juvenile melanoma, lentigo maligna melanoma, malignant melanoma, nodular melanoma, subungual melanoma and superficial spreading melanoma.

30 "Hepatoma" may be a malignant or benign tumor of the liver, including, for example, hepatocellular carcinoma.

"Patient" refers to an animal, preferably a mammal, more preferably a

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human.

“Biocompatible” refers to materials or compounds which are generally not injurious to biological functions and which will not result in any degree of unacceptable toxicity, including allergenic and disease states.

5           Throughout the present disclosure, the following abbreviations may be used: PEG, polyethylene glycol; ADI, arginine deiminase; SS, succinimidyl succinate; SSA, succinimidyl succinamide; SPA, succinimidyl propionate; and NHS, N-hydroxy-succinimide.

10           The present invention is based on the unexpected discovery that ADI modified with polyethylene glycol provides excellent results in treating certain types of cancer and inhibiting the metastasis of cancer. ADI may be covalently bonded to polyethylene glycol with or without a linking group, although a preferred embodiment utilizes a linking group.

15           In the present invention, the arginine deiminase gene may be derived, cloned or produced from any source, including, for example, microorganisms, recombinant biotechnology or any combination thereof. Preferably, arginine deiminase is cloned from microorganisms of the genus *Mycoplasma*. More preferably, the arginine deiminase is cloned from *Mycoplasma arginini*, *Mycoplasma hominus*, *Mycoplasma arthritides*, or any combination thereof. In particular, the arginine deiminase used in the  
20           present invention may have one or more of the amino acid sequences depicted in Figure 1.

25           In one embodiment of the present invention, the polyethylene glycol (PEG) has a total weight average molecular weight of about 1,000 to about 50,000; more preferably from about 3,000 to about 40,000, more preferably from about 5,000 to about 30,000; more preferably from about 8,000 to about 30,000; more preferably from about 11,000 to about 30,000; even more preferably from about 12,000 to about 28,000; still more preferably from about 16,000 to about 24,000; even more preferably from about 18,000 to about 22,000; even more preferably from about 19,000 to about 21,000, and most preferably about 20,000. Generally, polyethylene glycol with a molecular weight of 30,000 or more is difficult to dissolve, and yields of the formulated product are greatly  
30           reduced. The polyethylene glycol may be a branched or straight chain, preferably a straight chain. Generally, increasing the molecular weight of the polyethylene glycol decreases the immunogenicity of the ADI. The polyethylene glycol having a molecular

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weight described in this embodiment may be used in conjunction with ADI, and, optionally, a biocompatible linking group, to treat cancer, including, for example, melanomas, hepatomas and sarcomas, preferably melanomas.

In another embodiment of the present invention, the polyethylene glycol  
5 has a total weight average molecular weight of about 1,000 to about 50,000; preferably about 3,000 to about 30,000; more preferably from about 3,000 to about 20,000; more preferably from about 4,000 to about 12,000; still more preferably from about 4,000 to about 10,000; even more preferably from about 4,000 to about 8,000; still more preferably from about 4,000 to about 6,000; with about 5,000 being most preferred. The  
10 polyethylene glycol may be a branched or straight chain, preferably a straight chain. The polyethylene glycol having a molecular weight described in this embodiment may be used in conjunction with ADI, and optionally, a biocompatible linking group, to treat cancer, including, for example, melanomas, hepatomas and sarcomas, preferably hepatomas.

The linking group used to covalently attach ADI to PEG may be any  
15 biocompatible linking group. As discussed above, "biocompatible" indicates that the compound or group is non-toxic and may be utilized *in vitro* or *in vivo* without causing injury, sickness, disease or death. PEG can be bonded to the linking group, for example, via an ether bond, an ester bond, a thiol bond or an amide bond. Suitable biocompatible linking groups include, for example, an ester group, an amide group, an imide group, a  
20 carbamate group, a carboxyl group, a hydroxyl group, a carbohydrate, a maleimide group (including, for example, succinimidyl succinate (SS), succinimidyl propionate (SPA), succinimidyl carboxymethylate (SCM), succinimidyl succinamide (SSA) or N-hydroxy succinimide (NHS)), an epoxide group, an oxycarbonylimidazole group (including, for example, carbonyldimidazole (CDI)), a nitro phenyl group (including, for example,  
25 nitrophenyl carbonate (NPC) or trichlorophenyl carbonate (TPC)), a trysylate group, an aldehyde group, an isocyanate group, a vinylsulfone group, a tyrosine group, a cysteine group, a histidine group or a primary amine. Preferably, the biocompatible linking group is an ester group and/or a maleimide group. More preferably, the linking group is SS, SPA, SCM, SSA or NHS; with SS, SPA or NHS being more preferred, and with SS or  
30 SPA being most preferred.

Alternatively, ADI may be coupled directly to PEG (i.e., without a linking group) through an amino group, a sulfhydryl group, a hydroxyl group or a carboxyl group.



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ADI may be covalently bonded to PEG, via a biocompatible linking group, using methods known in the art, as described, for example, by Park et al., *Anticancer Res.*, 1:373-376 (1981); and Zaplinsky and Lee, *Polyethylene Glycol Chemistry: Biotechnical and Biomedical Applications*, J.M. Harris, ed., Plenum Press, NY, Chapter 21 (1992), the  
5 disclosures of which are hereby incorporated by reference herein in their entirety.

The attachment of PEG to ADI increases the circulating half-life of ADI. Generally, PEG is attached to a primary amine of ADI. Selection of the attachment site of polyethylene glycol on the arginine deiminase is determined by the role of each of the sites within the active domain of the protein, as would be known to the skilled artisan. PEG  
10 may be attached to the primary amines of arginine deiminase without substantial loss of enzymatic activity. For example, ADI cloned from *Mycoplasma arginini*, *Mycoplasma arthritides* and *Mycoplasma hominus* has about 17 lysines that may be modified by this procedure. In other words, the 17 lysines are all possible points at which ADI can be attached to PEG via a biocompatible linking group, such as SS, SPA, SCM, SSA and/or  
15 NHS. PEG may also be attached to other sites on ADI, as would be apparent to one skilled in the art in view of the present disclosure.

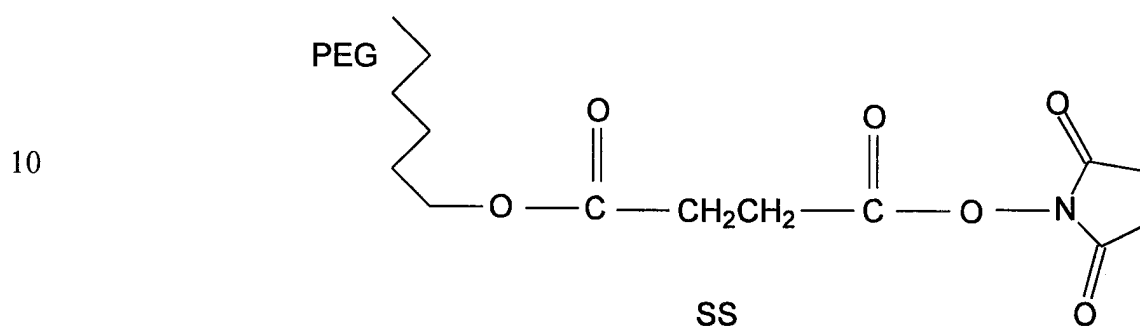
From 1 to about 30 PEG molecules may be covalently bonded to ADI. Preferably, ADI is modified with about 7 to about 15 PEG molecules, more preferably from about 9 to about 12 PEG molecules. In other words, about 30% to about 70% of the  
20 primary amino groups in arginine deiminase are modified with PEG, preferably about 40% to about 60%, more preferably about 45% to about 55%, and most preferably about 50% of the primary amino groups in arginine deiminase are modified with PEG. When PEG is covalently bonded to the end terminus of ADI, preferably only 1 PEG molecule is utilized. Increasing the number of PEG units on ADI increases the circulating half life of the  
25 enzyme. However, increasing the number of PEG units on ADI decreases the specific activity of the enzyme. Thus, a balance needs to be achieved between the two, as would be apparent to one skilled in the art in view of the present disclosure.

In the present invention, a common feature of the most preferred biocompatible linking groups is that they attach to a primary amine of arginine deiminase  
30 via a maleimide group. Once coupled with arginine deiminase, SS-PEG has an ester linkage next to the PEG, which may render this site sensitive to serum esterase, which may release PEG from ADI in the body. SPA-PEG and PEG2-NHS do not have an ester

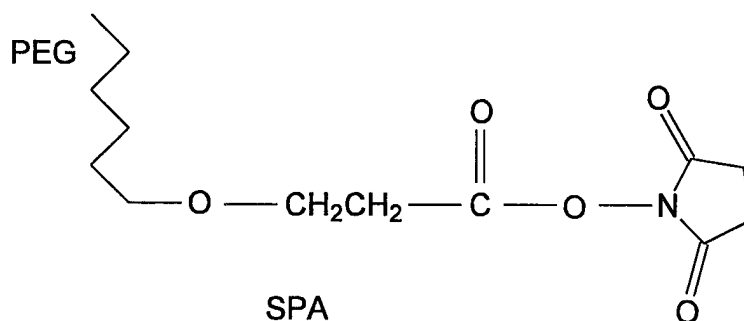
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linkage, so they are not sensitive to serum esterase.

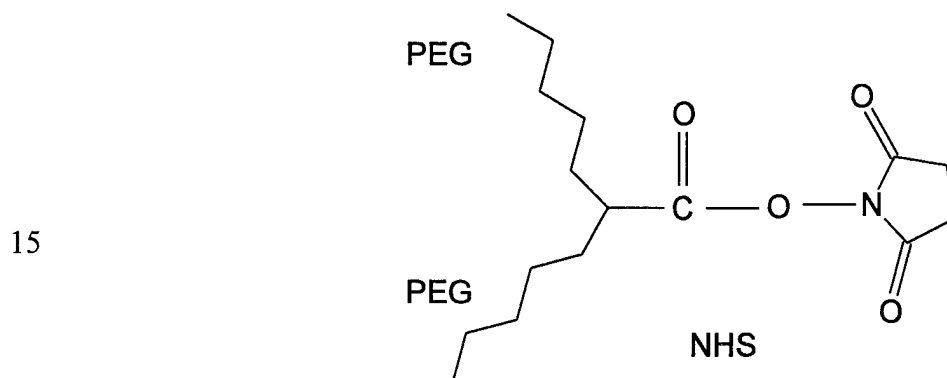
In the present invention, the particular linking groups do not appear to influence the circulating half-life of PEG-ADI or its specific enzyme activity. However, it is critical to use a biocompatible linking group in the present invention. PEG which is  
5 attached to the protein may be either a single chain, as with SS-PEG, SPA-PEG and SC-PEG, or a branched chain of PEG may be used, as with PEG2-NHS. The structural formulas of the preferred linking groups in the present invention are set forth below.  
SS-PEG:



SPA-PEG:



PEG2-NHS:



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A therapeutically effective amount of one of the compounds of the present invention is an amount that is effective to inhibit tumor growth. Generally, treatment is initiated with small dosages which can be increased by small increments until the optimum effect under the circumstances is achieved. Generally, a therapeutic dosage of compounds of the present invention may be from about 1 to about 200 mg/kg twice a week to about once every two weeks. For example, the dosage may be about 1 mg/kg once a week as a 2 ml intravenous injection to about 20 mg/kg once every 3 days. The optimum dosage with ADI-SS-PEG5,000 may be about twice a week, while the optimum dosage with ADI-SS-PEG20,000 may be from about once a week to about once every two weeks. PEG-ADI may be mixed with a phosphate buffered saline solution, or any other appropriate solution known to those skilled in the art, prior to injection. The PEG-ADI formulation may be administered as a solid (lyophalate) or as a liquid formulation, as desired.

The methods of the present invention can involve either *in vitro* or *in vivo* applications. In the case of *in vitro* applications, including cell culture applications, the compounds described herein can be added to the cells in cultures and then incubated. The compounds of the present invention may also be used to facilitate the production of monoclonal and/or polyclonal antibodies, using antibody production techniques well known in the art. The monoclonal and/or polyclonal antibodies can then be used in a wide variety of diagnostic applications, as would be apparent to one skilled in the art.

The *in vivo* means of administration of the compounds of the present invention will vary depending upon the intended application. As one skilled in the art will recognize, administration of the PEG-ADI composition of the present invention can be carried out, for example, orally, intranasally, intraperitoneally, parenterally, intravenously, intralymphatically, intratumorally, intramuscularly, interstitially, intra-arterially, subcutaneously, intraocularly, intrasynovial, transepithelial, and transdermally.

### Examples

The invention is further demonstrated in the following examples, which are for purposes of illustration, and are not intended to limit the scope of the present invention.

#### **Example 1: Production of Recombinant ADI**

Cultures of *Mycoplasma arginini* (ATCC 23243), *Mycoplasma hominus* (ATCC 23114) and *Mycoplasma arthritides* (ATCC 23192) were obtained from the

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American Type Culture Collection, Rockville, Maryland.

Arginine deiminase was cloned from *Mycoplasma arginini*, *Mycoplasma hominus* and *Mycoplasma arthritides* and expressed in *E. coli* as previously described by S. Misawa et al, *J. Biotechnology*, 36:145-155 (1994), the disclosure of which is hereby  
5 incorporated herein by reference in its entirety. The amino acid sequences of arginine deiminase from each of the above species is set forth in Figure 1. The top amino acid sequence, identified as ADIPROT, is from *Mycoplasma arginini*; the middle amino acid sequence, identified as ARTADIPRO, is from *Mycoplasma arthritides*; and the bottom amino acid sequence, identified as HOMADIPRO, is from *Mycoplasma hominus*. Each of  
10 the amino acid sequences are more than 96% conserved. Characterization, by methods known to those skilled in the art, of each of the proteins with respect to specific enzyme activity,  $K_m$ ,  $V_{max}$  and pH optima revealed that they were biochemically indistinguishable from each other. The pH optima was determined using a citrate buffer (pH 5-6.5), a phosphate buffer (pH 6.5-7.5) and a borate buffer (pH 7.5-8.5). The  $K_m$  and  $V_{max}$  were  
15 determined by incubating the enzyme with various concentrations of arginine and quantifying citrulline production. The  $K_m$  for the various enzymes was about 0.02 to 0.06  $\mu$ M and the  $V_{max}$  was about 15-20  $\mu$ mol/min/mg, the values of which are within standard error of each other.

The arginine deiminase genes were amplified by polymerase chain reaction  
20 using the following primer pair derived from the published sequence of *M. arginini*, as described, for example, by T. Ohno et al, *Infect. Immun.*, 58:3788-3795 (1990), the disclosure of which is hereby incorporated by reference herein in its entirety:

5'-GCAATCGATGTGTATTTGACAGT-3'

5'-TGAGGATCCTTACTACCACTTAACATCTTTACG-3'

25 The polymerase chain reaction product was cloned as a Bam HI-Hind III fragment into expression plasmid pQE16. DNA sequence analysis indicated this fragment had the same sequence for the arginine deiminase gene as described by Ohno et al, *Infect. Immun.*, *supra*. The five TGA codons in the ADI gene which encode tryptophan in *Mycoplasma* were changed to TGG codons by oligonucleotide-directed mutagenesis prior to gene  
30 expression in *E. coli*, as taught, for example, by J.R. Sayers et al, *Biotechniques*, 13:592-596 (1992). Recombinant ADI was expressed in inclusion bodies at levels of 10% of total cell protein.

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The proteins from each of the above three species of *Mycoplasma* have approximately 95% homology and are readily purified by column chromatography. Approximately 1.2 g of pure protein may be isolated from 1 liter of fermentation broth. Recombinant ADI is stable for about 2 weeks at 37°C and for at least 8 months when  
5 stored at 4°C. As determined by methods known to those skilled in the art, the proteins had a high affinity for arginine (0.04 µM), and a physiological pH optima of about 7.2 to about 7.4.

**Example 2: Renaturation and Purification of Recombinant ADI**

ADI protein was renatured, with minor modifications, as described by  
10 Misawa et al, *J. Biotechnology*, 36:145-155 (1994), the disclosure of which is hereby incorporated herein by reference in its entirety. 100 g of cell paste was resuspended in 800 ml of 10 mM K<sub>2</sub>PO<sub>4</sub> pH 7.0, 1 mM EDTA (buffer 1) and the cells were disrupted by two passes in a Microfluidizer (Microfluidics Corporation, Newton, MA). Triton X-100 was added to achieve a final concentration of 4% (v/v). The homogenate was stirred for 30  
15 min at 4°C, then centrifuged for 30 min at 13,000 g. The pellet was collected and resuspended in one liter of buffer 1 containing 0.5% Triton X-100. The solution was diafiltered against 5 volumes of denaturation buffer (50 mM Tris HCl, pH 8.5, 10 mM DTT) using hollow-fiber cartridges with 100 kD retention rating (Microgon Inc., Laguna Hills, CA). Guanidine HCl was added to achieve a final concentration of 6 M and the  
20 solution was stirred for 15 min at 4°C. The solution was diluted 100-fold into refolding buffer 1, 10 mM K<sub>2</sub>PO<sub>4</sub>, pH 7.0 and stirred for 48 hours at 15°C, particulates were removed by centrifugation at 15,000 x g.

The resulting supernatant was concentrated on a Q Sepharose Past Flow (Pharmacia Inc., Piscataway, NJ) column preequilibrated in refolding buffer. ADI was  
25 eluted using refolding buffer containing 0.2 M NaCl. The purification procedure yielded ADI protein, which was >95% pure as estimated by SDS-PAGE analysis. 8 g of pure renatured ADI protein was produced from 1 kg of cell paste which corresponds to 200 mg purified ADI per liter of fermentation.

ADI activity was determined by micro-modification of the method  
30 described by Oginsky et al, *Meth. Enzymol.*, (1957) 3:639-642. 10 µl samples in 0.1 M Na<sub>2</sub>PO<sub>4</sub>, pH 7.0 (BUN assay buffer) were placed in a 96 well microliter plate, 40 µl of 0.5 mM arginine in BUN assay buffer was added, and the plate was covered and incubated at

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37°C for 15 minutes. 20 µl of complete BUN reagent (Sigma Diagnostics) was added and the plate was incubated for 10 minutes at 100°C. The plate was then cooled to 22°C and analyzed at 490 nm by a microliter plate reader (Molecular Devices, Inc). 1.0 IU is the amount of enzyme which converts 1 µmole of L-arginine to L-citrulline per minute.

- 5 Protein concentrations were determined using Pierce Coomassie Blue Protein Assay Reagent (Pierce Co., Rockford, IL) with bovine serum albumin as a standard.

The enzyme activity of the purified ADI preparations was 17-25 IU/mg.

**Example 3: Attachment of PEG to ADI**

PEG was covalently bonded to ADI in a 100 mM phosphate buffer, pH 7.4.

- 10 Briefly, ADI in phosphate buffer was mixed with a 100 molar excess of PEG. The reaction was stirred at room temperature for 1 hour, then the mixture was extensively dialyzed to remove unincorporated PEG.

- A first experiment was performed where the effect of the linking group used in the PEG-ADI compositions was evaluated. PEG10,000 and ADI were covalently  
15 bonded via four different linking groups: an ester group or maleimide group, including SS, SSA, SPA and SSPA, where the PEG had a total weight average molecular weight of 5,000, 10,000, 12,000, 20,000, 30,000 and 40,000; an epoxy group, PEG-epoxy, where the PEG had a total weight average molecular weight of 5,000; and a branched PEG group, PEG2-NHS, where the PEG had a total weight average molecular weight of 10,000,  
20 20,000 and 40,000.

- 5.0 IU of the resulting compositions were injected into mice (5 mice in each group). To determine the serum levels of arginine, the mice were bled from the retro orbital plexus (100 µl). Immediately following collection an equal volume of 50% (w/v) of trichloroacetic acid was added. The precipitate was removed by centrifugation (13,000 x g  
25 for 30 minutes) and the supernatant removed and stored frozen at -70°C. The samples were then analyzed using an automated amino acid analyzer and reagents from Beckman Instruments using protocols supplied by the manufacturer. The limits of sensitivity for citrulline by this method was approximately 2-6 µM and the reproducibility of measurements within about 8%. The amount of serum arginine was determined by amino  
30 acid analysis. As can be seen from the results in Figure 3, the linking group covalently bonding the PEG and ADI did not have an appreciable effect on the ability of ADI to reduce serum arginine *in vivo*. In other words, the linking group may not be critical to the

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results of the experiment, except that a non-toxic linking group must be used for *in vivo* applications.

A second experiment was performed wherein the effect of the linking group and molecular weight of PEG on serum citrulline levels *in vivo* was evaluated. Mice (5 in each group) were given various compositions of ADI and PEG-ADI in an amount of 5.0 IU. To determine the serum levels of citrulline, the mice were bled from the retro orbital plexus (100 ul). Immediately following collection an equal volume of 50% (w/v) of trichloroacetic acid was added. The precipitate was removed by centrifugation (13,000 x g for 30 minutes) and the supernatant removed and stored frozen at -70°C. The samples were then analyzed using an automated amino acid analyzer and reagents from Beckman Instruments using protocols supplied by the manufacturer. The limits of sensitivity for citrulline by this method was approximately 2-6 uM and the reproducibility of measurements within about 8%. The amount of citrulline was determined, and the area under the curve approximated and expressed as  $\mu\text{mol}$  days.

In Figure 4, the open circles indicate the amount of citrulline produced by native ADI, the filled circles are ADI-SC-PEG, the open squares are ADI-SS-PEG, the open triangles are ADI-SPA-PEG, and the filled triangles are branched chain PEG-NHS-PEG<sub>2</sub>. The results in Figure 4 demonstrate that the molecular weight of the PEG determines the effectiveness of the PEG-ADI composition. The effectiveness of the PEG-ADI compositions is not necessarily based on the method or means of attachment of the PEG to ADI, except that a biocompatible linking group must be used for *in vivo* applications.

The results in Figure 4 also demonstrate that the optimal molecular weight of PEG is 20,000. Although PEG30,000 appears to be superior to PEG20,000 in terms of its pharmacodynamics, PEG30,000 is less soluble, which makes it more difficult to work with. The yields, which were based on the recovery of enzyme activity, were about 90% for PEG5,000 and PEG12,000; about 85% for PEG20,000 and about 40% for PEG30,000. Therefore, PEG20,000 is the best compromise between yield and circulating half life, as determined by citrulline production.

In a third experiment, the dose response of serum arginine depletion and the production of citrulline with ADI-SS-PEG5,000 and ADI-SS-PEG20,000 was determined. Mice (5 in each group) were given a single injection of 0.05 IU, 0.5 IU or 5.0 IU of either

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ADI-SS-PEG5,000 or ADI-SS-PEG20,000. At indicated times, serum was collected, as described above, and an amino acid analysis was performed to quantify serum arginine (Figures 5A and 5C) and serum citrulline (Figures 5B and 5D). Both formulations induced a dose dependent decrease in serum arginine and an increase in serum citrulline.

- 5 However, the effects induced by ADI-SS-PEG20,000 were more pronounced and of longer duration than the effects induced by ADI-SS-PEG5,000.

**Example 4: Selectivity of ADI Mediated Cytotoxicity**

The selectivity of arginine deiminase mediated cytotoxicity was demonstrated using a number of human tumors. Specifically, human tumors were tested *in*  
 10 *vitro* for sensitivity to ADI-SS-PEG5,000 (50 ng/ml). Viability of cultures was determined after 7 days. For a culture to be defined as "inhibited," greater than 95% of the cells must take up Trypan blue dye. A host of normal cells were also tested, including endothelial cells, smooth muscle cells, epithelial cells and fibroblasts, and none were inhibited by ADI-SS-PEG5,000. Although arginine deiminase has no appreciable toxicity  
 15 towards normal, and most tumor cells, ADI-SS-PEG5,000 greatly inhibited all human melanomas and hepatomas that were commercially available from the ATCC, MSKCC and Europe.

**Table 1: Specificity of Arginine Deiminase Cytotoxicity**

20	Tumor Type	Number of Tumors Tested	Tumors inhibited (%)
	Brain	16	0
	Colon	34	0
	Bladder	3	0
	Breast	12	0
	Kidney	5	0
25	Sarcoma	11	64
	Hepatoma	17	100
	Melanoma	37	100

In a parallel set of experiments, mRNA was isolated from the tumors. Northern blot analyses, using the human arginosuccinate synthase cDNA probe, indicated



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complete concordance between the sensitivity to arginine deiminase treatment and an inability to express arginosuccinate synthase. This data suggests that ADI toxicity results from an inability to induce arginosuccinate synthase. Therefore, these cells cannot synthesize arginine from citrulline, and are unable to synthesize the proteins necessary for growth.

**Example 5: Circulating Half-Life**

Balb C mice (5 in each group) were injected intravenously with a single 5.0 IU dose of either native arginine deiminase or various formulations of arginine deiminase modified with polyethylene glycol, as indicated in Figures 3A and 3B. To determine the serum levels of arginine and citrulline, the mice were bled from the retro orbital plexus (100 ul). Immediately following collection an equal volume of 50% (w/v) of trichloroacetic acid was added. The precipitate was removed by centrifugation (13,000 x g for 30 minutes) and the supernatant removed and stored frozen at -70°C. The samples were then analyzed using an automated amino acid analyzer and reagents from Beckman Instruments using protocols supplied by the manufacturer. The limits of sensitivity for arginine by this method was approximately 6 pM and the reproducibility of measurements within about 8%.

A dose dependent decrease in serum arginine levels, as shown by the solid circles in Figure 2A, and a rise in serum citrulline, as shown by the open triangles in Figure 2B, were detected from the single dose administration of native ADI (filled circles) or ADI-SS-PEG (open triangles). However, the decrease in serum arginine and rise in serum citrulline was short lived, and soon returned to normal. The half life of arginine depletion is summarized in the Table below.

**Table 2: Half-Life of Serum Arginine Depletion**

Compound	Half-Life in Days
Native ADI	1
ADI-SS-PEG5,000	5
ADI-SS-PEG12,000	15
ADI-SS-PEG20,000	20
ADI-SS-PEG30,000	22

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This experiment demonstrates that normal cells and tissues are able to convert the citrulline back into arginine intracellularly while melanomas and hepatomas cannot because they lack arginosuccinate synthetase.

**Example 6: Antigenicity of PEG modified ADI**

5 To determine the antigenicity of native ADI, ADI-SS-PEG5,000, and ADI-SS-PEG20,000, the procedures described in, for example, Park, *Anticancer Res.*, *supra*, and Kamisaki, *J. Pharmacol. Exp. Ther.*, *supra*, were followed.. Briefly, Balb C mice (5 in each group) were intravenously injected weekly for 12 weeks with approximately 0.5 IU (100 ug of protein) of native ADI, ADI-SS-PEG5,000 or ADI-SS-PEG20,000. The  
10 animals were bled (0.05 ml) from the retro orbital plexus at the beginning of the experiment and at weeks 4, 8 and 12. The serum was isolated and stored at -70°C. The titers of anti-ADI IgG were determined by ELISA. 50 ug of ADI was added to each well of a 96 well micro-titer plate and was incubated at room temperature for 4 hours. The plates were rinsed with PBS and then coated with bovine serum albumin (1 mg/ml) to  
15 block nonspecific protein binding sites, and stored over night at 4°C. The next day serum from the mice was diluted and added to the wells. After 1 hour the plates were rinsed with PBS and rabbit anti-mouse IgG coupled to peroxidase was added to the wells. The plates were incubated for 30 min and then the resulting UV absorbance was measured using a micro-titer plate reader. The titer was defined as the highest dilution of the serum which  
20 resulted in a two-fold increase from background absorbance (approximately 0.50 OD).

The results are shown in Figure 6. The open circles represent the data obtained from animals injected with native ADI, which was very antigenic. The filled circles represent the data obtained from the animals injected with ADI-SS-PEG5,000, while the open triangles represent the data obtained from the animals injected with ADI-SS-PEG20,000. As can be seen from Figure 6, ADI-SS-PEG5,000 and ADI-SS-  
25 PEG20,000 are significantly less antigenic than native ADI. For example, as few as 4 injections of native ADI resulted in a titer of about  $10^6$ , while 4 injections of any of the PEG-ADI formulations failed to produce any measurable antibody. However, after 8 injections, the ADI-PEG5,000 had a titer of about  $10^2$ , while ADI-PEG20,000 did not  
30 induce this much of an immune response until after 12 injections. The results demonstrate that attaching PEG to ADI blunts the immune response to the protein.

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**Example 7: Tumor Inhibition of Human Melanomas**

The effect of PEG-ADI on the growth of human melanoma (SK-Mel 28) in nude mice was determined. Nude mice (5 in each group) were injected with  $10^6$  SK-mel 2 human melanoma cells which were allowed to grow until the tumors reached a diameter of about 3-5 mm. The mice were left untreated (open circles) or were treated once a week for 12 weeks with 5.0 IU of ADI-SS-PEG5,000 (filled triangles), ADI-SS-PEG12,000 (open triangles) or ADI-SS-PEG20,000 (filled circles). The tumor size was measured weekly, and the mean diameter of the tumors is presented in Figure 7.

Figure 8 shows the effectiveness of ADI-SS-PEG20,000 on three human melanomas (SK-mel 2, SK-mel 28, M24-met) grown *in vivo* in nude mice. Nude mice (5 in each group) were injected with  $10^6$  SK-mel 2, SK-mel 28 or M24-met human melanoma cells. The tumors were allowed to grow until they were approximately 3-5 mm in diameter. Thereafter, the animals were injected once a week with 5.0 IU of ADI-SS-PEG20,000. The results are shown in Figure 8, and show that PEG-ADI inhibited tumor growth and that eventually the tumors began to regress and disappear. Because the tumors did not have argininosuccinate synthetase, they were unable to synthesize proteins (because ADI eliminated arginine and the tumors could not make it) so that the cells "starved to death."

Since M24-met human melanoma is highly metastatic, the animals injected with M24-met human melanoma cells were sacrificed after 4 weeks of treatment and the number of metastases in the lungs of the animals was determined. The control animals had an average of 32 metastases, while the animals treated with ADI-SS-PEG20,000 did not have any metastases. The results appear to indicate that ADI-SS-PEG20,000 not only inhibited the growth of the primary melanoma tumor, but also inhibited the formation of metastases.

It is of interest to note that in over 200 animals tested, the average number of metastases in the control group was  $49 \pm 18$ , while only a single metastasis was observed in 1 treated animal.

**Example 8: Tumor Inhibition of Human Hepatomas**

The ability of PEG5,000-ADI to inhibit the growth of a human hepatoma *in vivo* was tested. Nude mice (5 in each group) were injected with  $10^6$  human hepatoma SK-Hep1 cells. The tumors were allowed to grow for two weeks and then the animals were

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treated once a week with 5.0 IU of SS-PEG5,000-ADI (solid circles), SS-PEG12,000-ADI (solid triangles) or SS-PEG20,000-ADI (open triangles). The results are set forth in Figure 9. The untreated animals (open circles) all died within 3 weeks. In contrast, animals treated with ADI had a far longer life expectancy, as can be seen from Figure 9.

- 5 All the surviving mice were euthanized after 6 months, and necropsy indicated that they were free of tumors.

Surprisingly, PEG5,000-ADI is most effective in inhibiting hepatoma growth *in vivo*. The exact mechanism by which this occurs is unknown. Without being bound to any theory of the invention works, it appears that proteins formulated with SS-PEG5,000-ADI become sequestered in the liver. Larger molecular weights of PEG do not, which may be due to the uniqueness of the hepatic endothelium and the spaces (fenestrae) being of such a size that larger molecular weights of PEG-ADI conjugates are excluded.

**Example 9: Application to Humans**

PEG5,000-ADI and PEG20,000-ADI were incubated *ex vivo* with normal human serum and the effects on arginine concentration was determined by amino acid analysis, where the enzyme was found to be fully active and capable of degrading all the detectable arginine with the same kinetics as in the experiments involving mice. The reaction was conducted at a volume of 0.1 ml in a time of 1 hour at 37°C.

Additionally, the levels of arginine and citrulline in human serum are identical with that found in mice. PEG-proteins circulate longer in humans than they do in mice. For example, the circulating half life of PEG conjugated adenosine deiminase, asparaginase, glucocerbroidase, uricase, hemoglobin and superoxide dismutase all have a circulating half life that is 5 to 10 times longer than the same formulations in mice. What this has meant in the past is that the human dose is most often 1/5 to 1/10 of that used in mice.

- 25 Accordingly, PEG-ADI should circulate even longer in humans than it does in mice.

Each of the patents, patent applications and publications described herein are hereby incorporated by reference herein in their entirety.

Various modifications of the invention, in addition to those described herein, will be apparent to one skilled in the art in view of the foregoing description. Such modifications are also intended to fall within the scope of the appended claims.

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What is claimed is:

1. A compound comprising arginine deiminase covalently bonded via a linking group to polyethylene glycol, wherein the polyethylene glycol has a total weight average molecular weight in the range of about 12,000 to about 40,000.
2. The compound of claim 1, wherein said linking group is a maleimide group.
3. The compound of claim 2, wherein said maleimide group is succinimidyl succinate, succinimidyl propionate, succinimidyl carboxymethylate, succinimidyl succinamide, N-hydroxy succinimide or combinations thereof.
4. The compound of claim 3, wherein said maleimide group is succinimidyl succinate, succinimidyl propionate or combinations thereof.
5. The compound of claim 1, wherein said arginine deiminase is derived from a microorganism of the genus *Mycoplasma*.
6. The compound of claim 5, wherein said microorganism is selected from the group consisting of *Mycoplasma arginini*, *Mycoplasma hominus*, *Mycoplasma arthritides* and combinations thereof.
7. The compound of claim 1, wherein said arginine deiminase is covalently bonded to about 7 to about 15 polyethylene glycol molecules.
8. The compound of claim 7, wherein said arginine deiminase is covalently bonded to about 9 to about 12 polyethylene glycol molecules.
9. The compound of claim 1, wherein said polyethylene glycol has a total weight average molecular weight of from about 12,000 to about 30,000.
10. A method of enhancing the circulating half life of arginine deiminase comprising modifying said arginine deiminase by covalently bonding said arginine deiminase via a linking group to polyethylene glycol, wherein the polyethylene glycol has a total weight average molecular weight in the range from about 12,000 to about 40,000.

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11. A method of treating a tumor in a patient comprising administering to said patient a therapeutically effective amount of a compound of Claim 1.

12. The method of claim 11, wherein said tumor is a melanoma.

13. The method of claim 12, wherein said polyethylene glycol has a total weight average molecular weight of about 20,000

14. The method of claim 12, wherein said linking group is a maleimide group.

15. The method of claim 14, wherein said maleimide group is succinimidyl succinate, succinimidyl propionate, succinimidyl carboxymethylate, succinimidyl succinamide, N-hydroxy succinimide or combinations thereof.

16. A method of treating hepatoma in a patient comprising administering to said patient a therapeutically effective amount of a compound comprising arginine deiminase covalently bonded via a linking group to polyethylene glycol, wherein the polyethylene glycol has a total weight average molecular weight in the range of about 5,000.

17. The method of claim 16, wherein said linking group is a maleimide group.

18. The method of claim 17, wherein said maleimide group is succinimidyl succinate, succinimidyl propionate, succinimidyl carboxymethylate, succinimidyl succinamide, N-hydroxy succinimide or combinations thereof.

19. The method of claim 11, wherein said tumor is a sarcoma.

20. A method of treating and inhibiting metastases in a patient comprising administering to said patient a therapeutically effective amount of a compound of claim 1.

21. A compound comprising arginine deiminase covalently bonded via a linking group to polyethylene glycol, wherein the polyethylene glycol has a total weight average molecular weight of from about 1,000 to about 40,000, and wherein the linking group is selected from the group consisting of a maleimide group, an amide group, an

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imide group, a carbamate group, an ester group, an epoxy group, a carboxyl group, a hydroxyl group, a carbohydrate, a tyrosine group, a cysteine group, a histidine group and combinations thereof.

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The alignment was done on 3 Protein sequences.

Character to show that a position in the alignment is perfectly conserved: '\*'

Character to show that a position is well conserved: '.'

# Alignment

```

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ARTADIPRO MSVFDSKFKGIHVYSEIGELESVLVHEPGREIDYITPARLDELLFSAILE 50
HOMADIPRO MSVFDSKFNHGIHVYSEIGELETVLVHEPGREIDYITPARLDELLFSAILE 50
          *****.*****.*****.*****.*****.*****.*****.*****.*****.*****

ADIPROT   SHDARKEHKQFVAELKANDINVVELIDLVAETYDLASQEAKDKLIEEFLE 100
ARTADIPRO SHDARKEQSQFVAILKANDINVVETIDLVAETYDLASQEAKDRLIEEFLE 100
HOMADIPRO SHDARKEHQSFVKIMKDRGINVVELTDLVAETYDLASKAAKEEFIETFLE 100
          *****.  **  .*.*****.*****.***.  **.***

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ARTADIPRO DSEPVLSEAHKKVVVRNFLKAKKTSRKLVELMMAGITKYDLGVEADHELIV 150
HOMADIPRO ETVPVLTEANKKAVRAFLLSKPT-HEMVEFMMSGITKYELGVESENELIV 149
          ..  ***.***.***.***.***.***.***.***.***.***.***.***.***.***.***.***

ADIPROT   DMPNLYFTRDPFASVGNVGTIHYMRYKVRQRETLFSRFVFSNHPKLINT 200
ARTADIPRO DMPNLYFTRDPFASVGNVGTIHFMRKVRRRRETLFSRFVFRNHPKLINT 200
HOMADIPRO DMPNLYFTRDPFASVGNVGTIHFMRKVRRRRETLFARFVFRNHPKLINT 199
          *****.*****.***.***.***.***.***.***.***.***.***.***.***.***.***.***

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ARTADIPRO PWYYDPAMKLSIEGGDVFIYNNDTLVVGVSERTDLQTVTLAKNIVANKE 250
HOMADIPRO PWYYDPAMKMPIEGGDVFIYNNDTLVVGVSERTDLQTVTLAKNIVANKE 249
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ADIPROT   CEFKRIVAINVPKWTNLMHLDTWLTMLDKDKFLYSPIANDVFKFWDYDLV 300
ARTADIPRO CEFKRIVAINVPKWTNLMHLDTWLTMLDKDKFLYSPIANDVFKFWDYDLV 300
HOMADIPRO VEFKRIVAINVPKWTNLMHLDTWLTMLDKDKFLYSPIANDVFKFWDYDLV 299
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ARTADIPRO NGGAEPQPVENGLPLEGLLQSIINKKPVLPIAGEGASQMEIERETHFDG 350
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          *****.*****.***.***.***.***.***.***.***.***.***.***.***.***.***.***

ADIPROT   TNYLAIRPGVVIGYSRNEKTNAALEAAGIKVLPFHGNQLSLGMGNARCMS 400
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HOMADIPRO TNYLAIRPGVVIGYSRNEKTNAALKAAGIKVLPFHGNQLSLGMGNARCMS 399
          ***.***.***.***.***.***.***.***.***.***.***.***.***.***.***.***

ADIPROT   MPLSRKDVK 409
ARTADIPRO MPLSRKDVK 409
HOMADIPRO MPLSRKDVK 408
          *****

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FIGURE 1

ADIPROT = *Mycoplasma arginini*  
ARTADIPRO = *Mycoplasma arthritides*  
HOMADIPRO = *Mycoplasma hominus*



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FIG. 2A

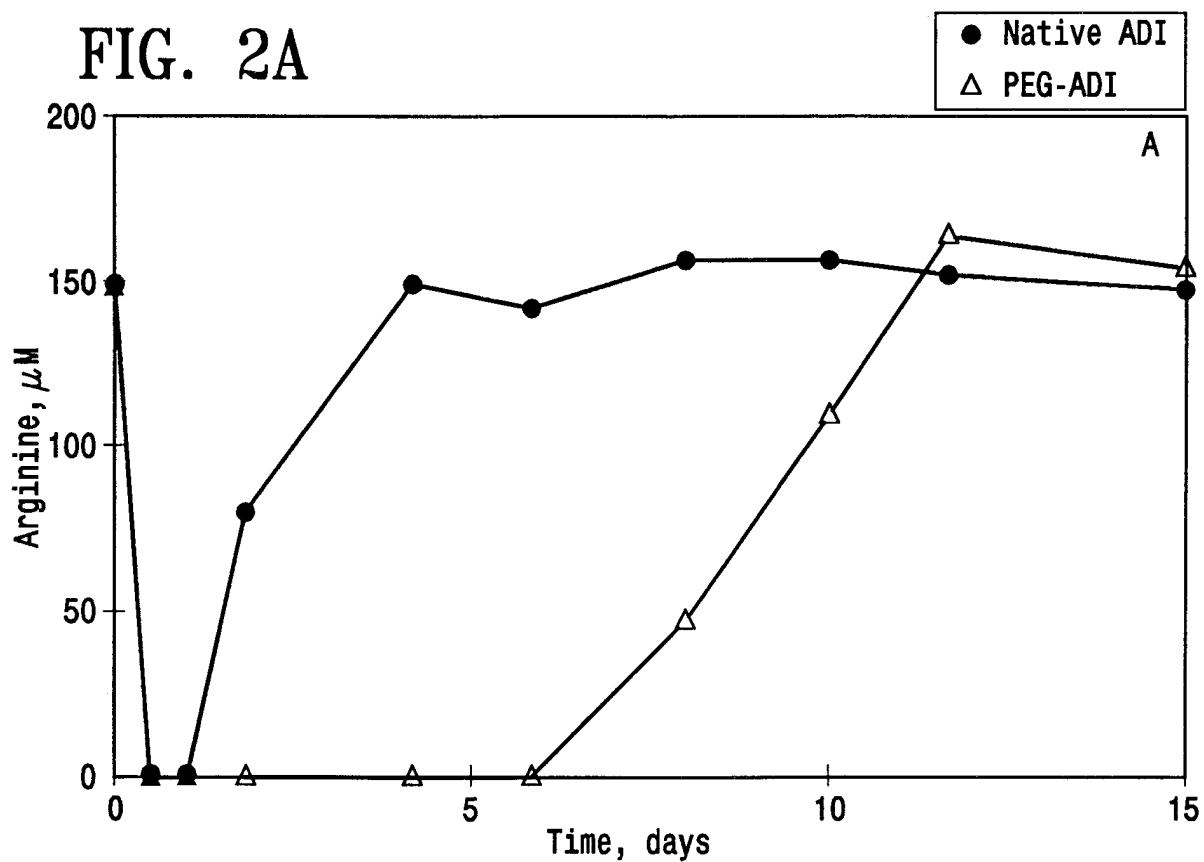
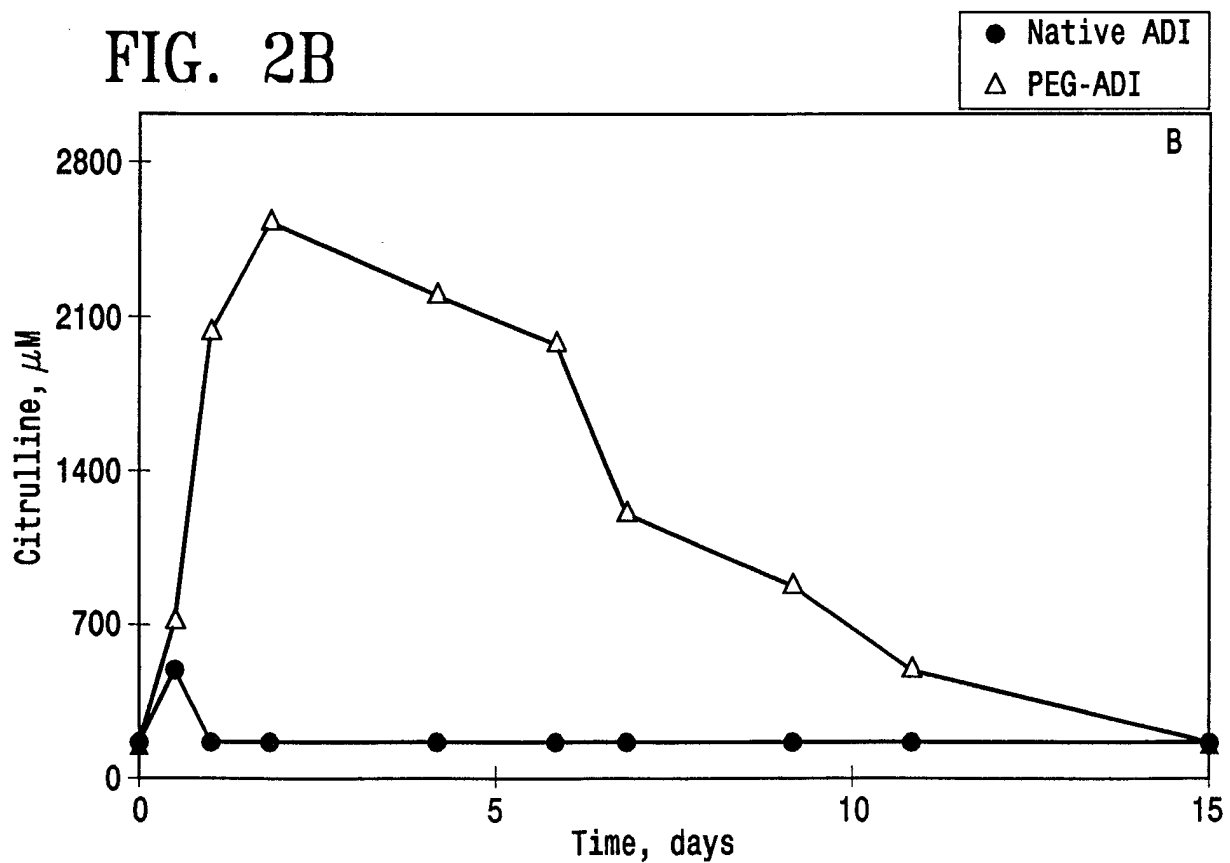
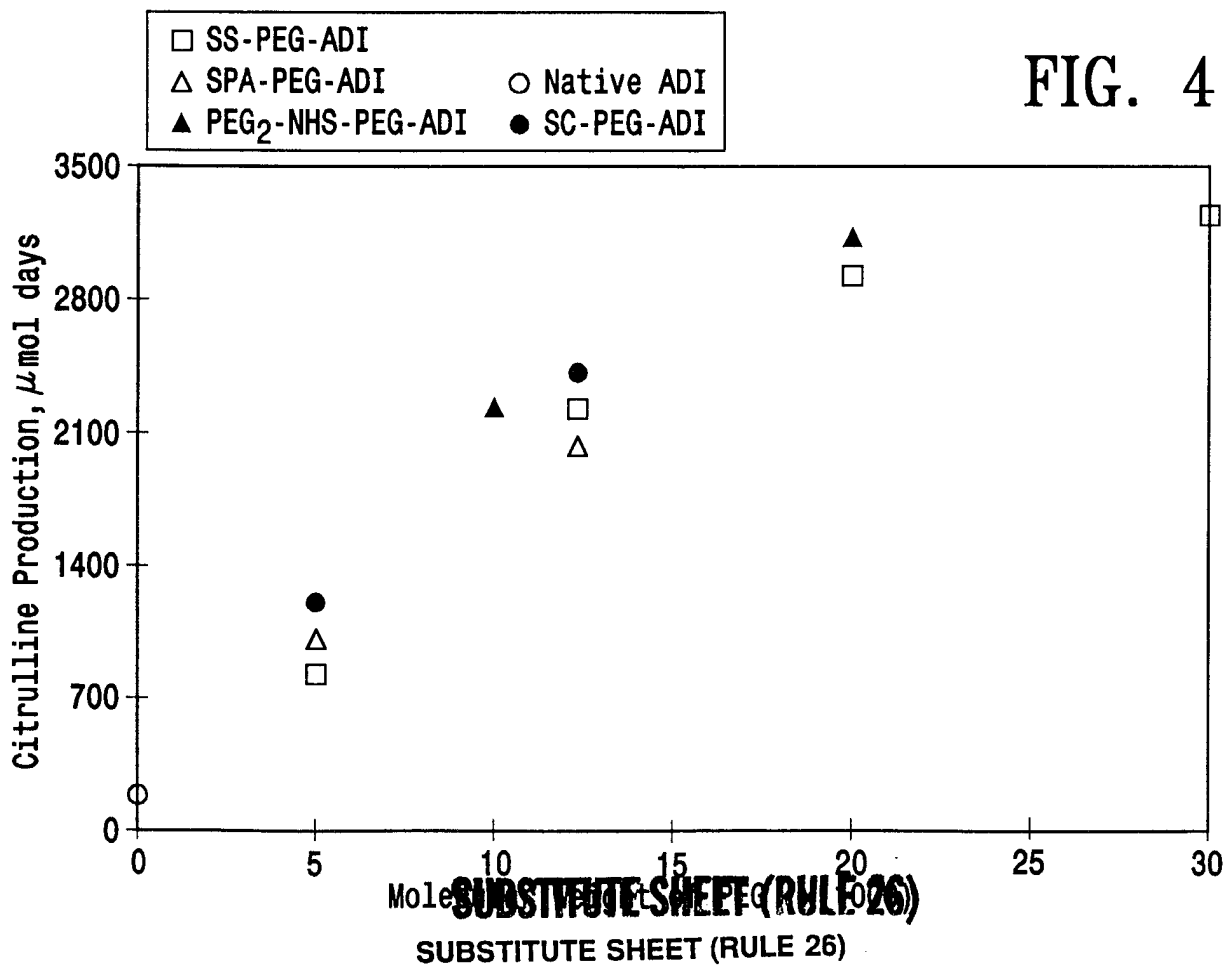
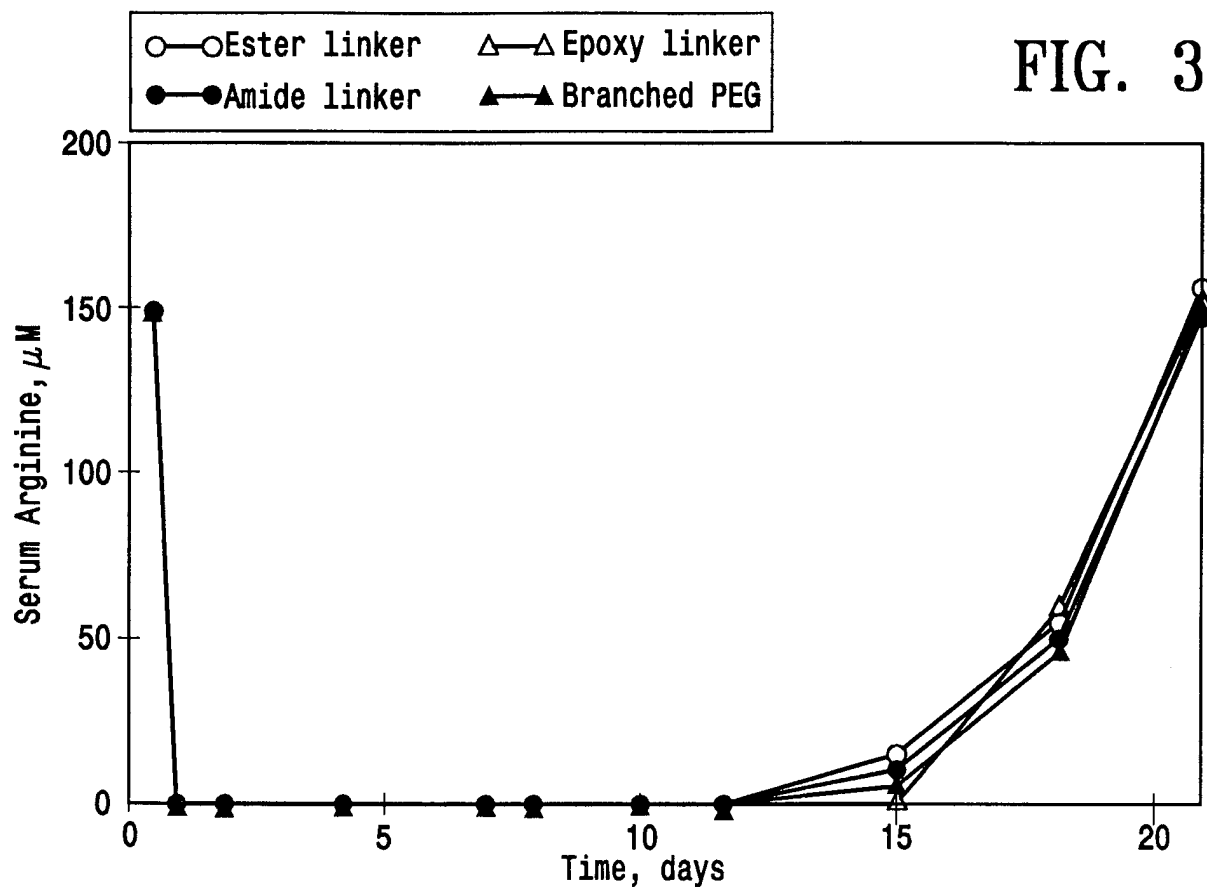
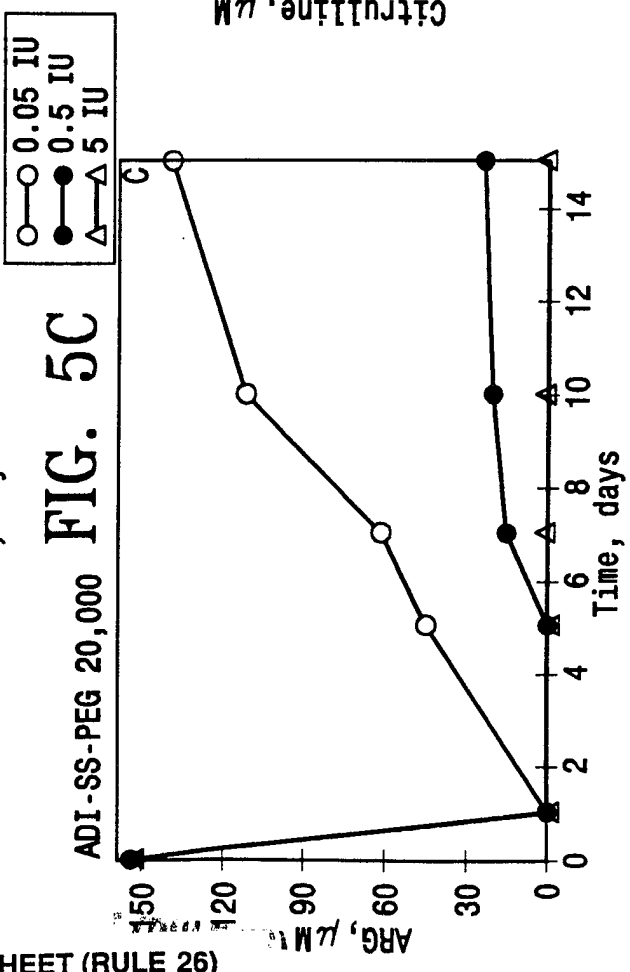
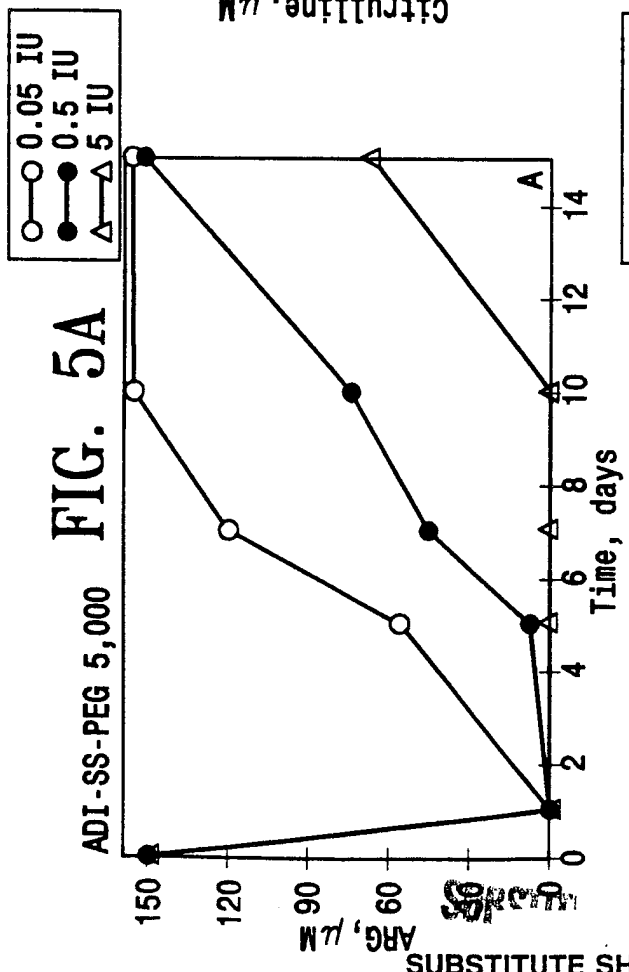
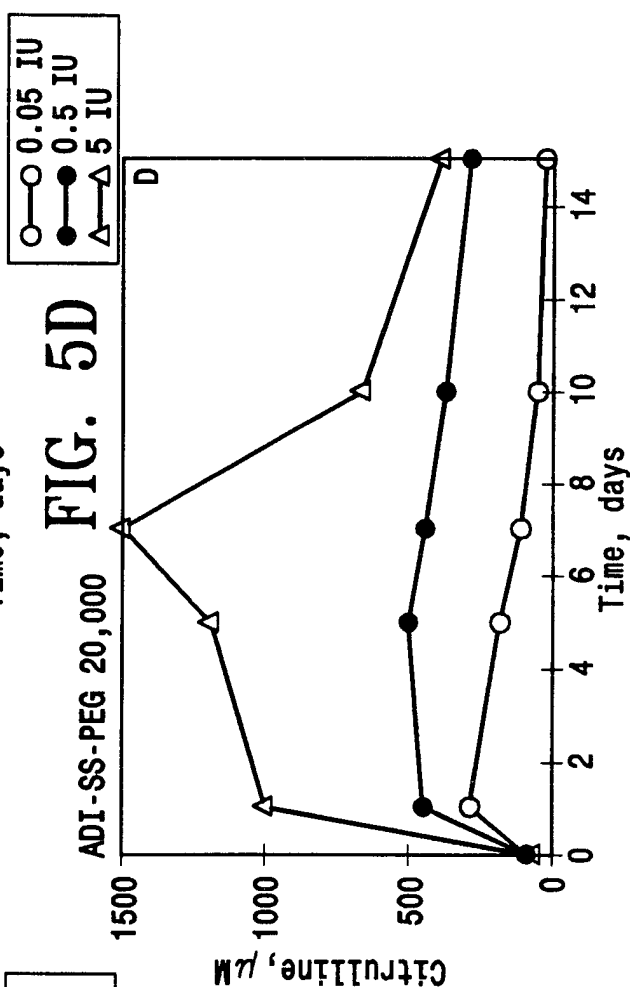
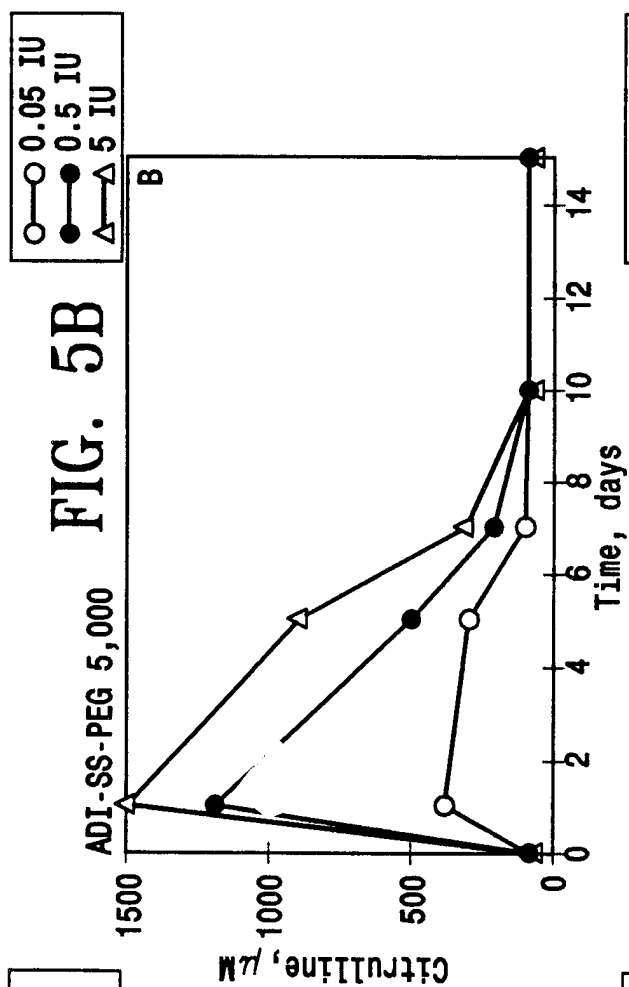


FIG. 2B



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FIG. 6

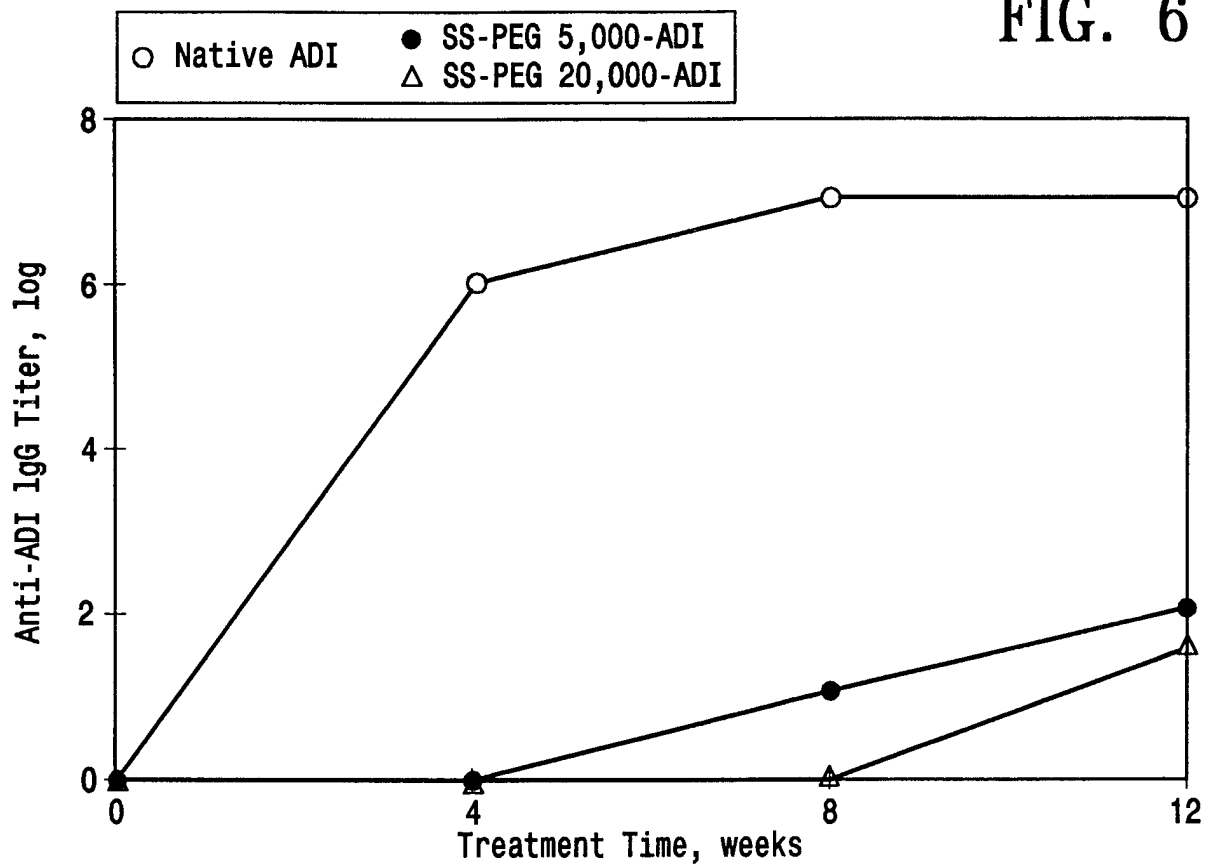
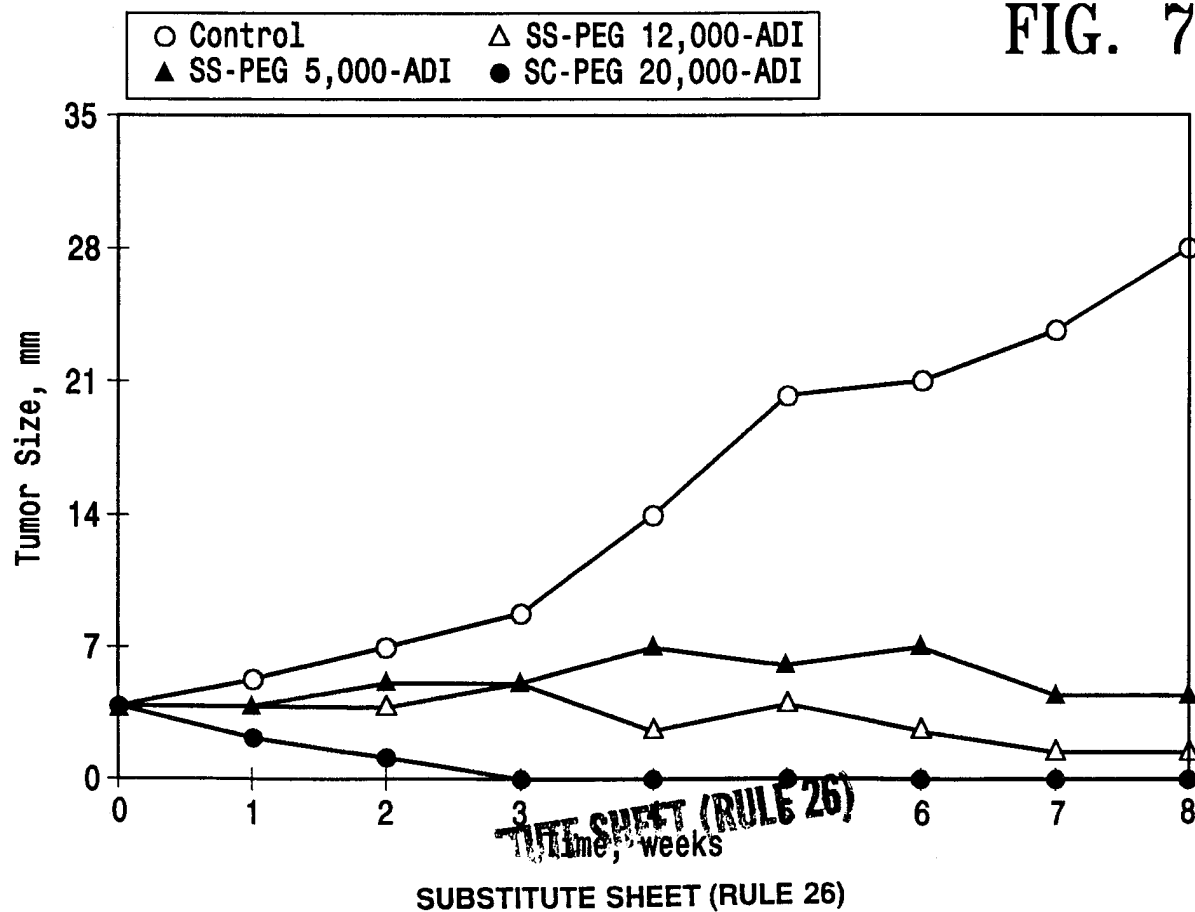
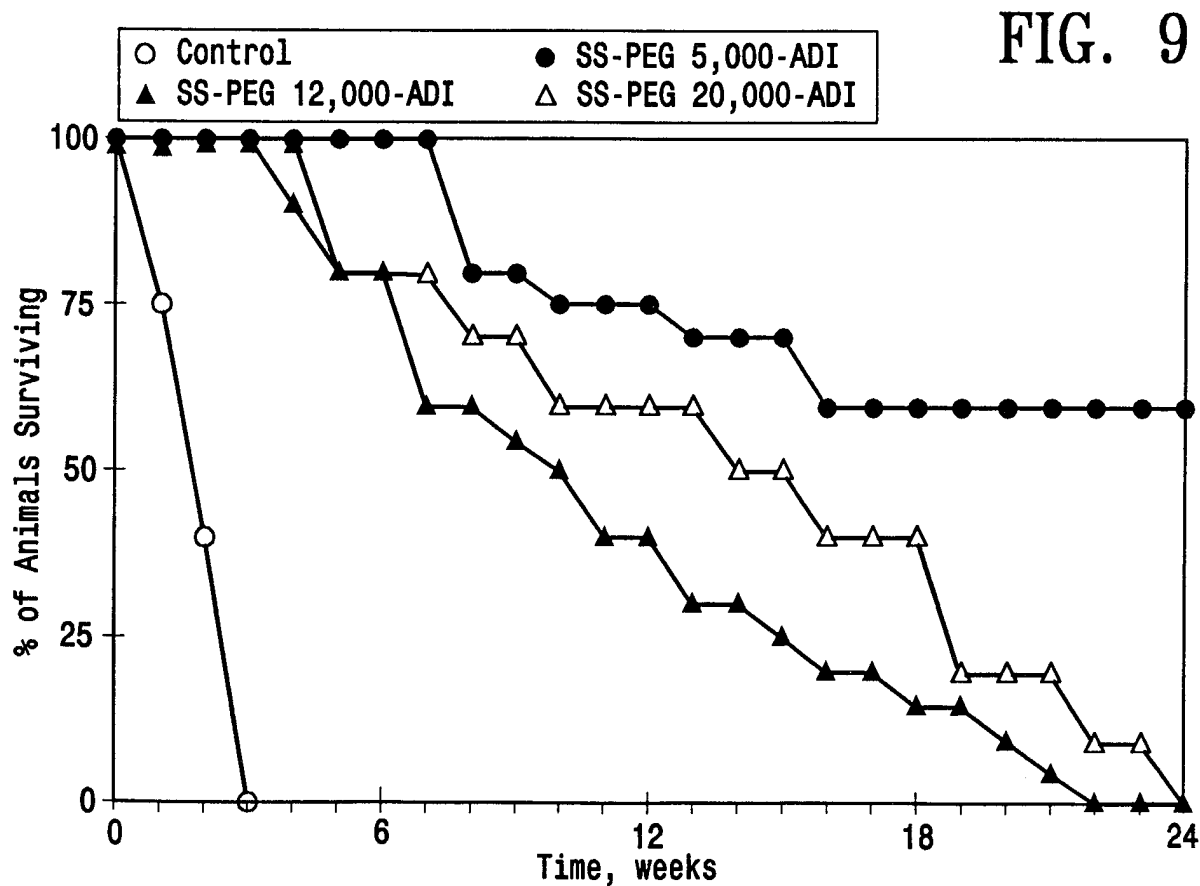
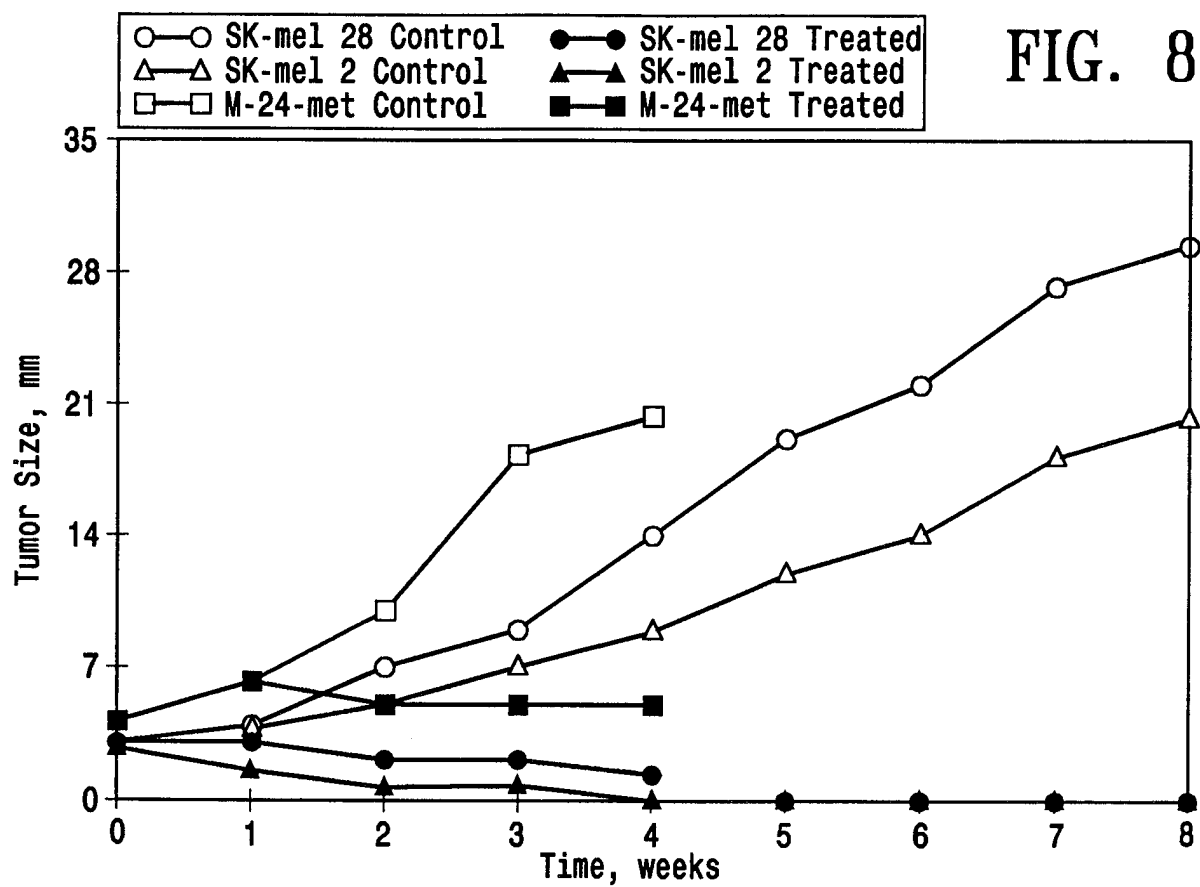


FIG. 7



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# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US98/09575

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :C12N 9/80

US CL :435/228

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 435/228

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS, CAS Online

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5,474,928 A (TAKAKU et al) 12 December 1995, see entire document.	1-21



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A* document defining the general state of the art which is not considered to be of particular relevance	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*B* earlier document published on or after the international filing date	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*&* document member of the same patent family
*O* document referring to an oral disclosure, use, exhibition or other means	
*P* document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

29 JULY 1998

Date of mailing of the international search report

08 SEP 1998

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