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Poly- $\beta$ -1 $\rightarrow$ 4-N-Acetylglucosamine1. INTRODUCTION

The present invention relates, first, to a  
5 purified, easily produced poly- $\beta$ -1 $\rightarrow$ 4-N-  
acetylglucosamine (p-GlcNAc) polysaccharide species.  
The p-GlcNAc of the invention is a polymer of high  
molecular weight whose constituent monosaccharide  
sugars are attached in a  $\beta$ -1 $\rightarrow$ 4 conformation, and which  
10 is free of proteins, and substantially free of single  
amino acids, and other organic and inorganic  
contaminants. In addition, derivatives and  
reformulations of p-GlcNAc are described. The present  
invention further relates to methods for the  
15 purification of the p-GlcNAc of the invention from  
microalgae, preferably diatom, starting sources.  
Still further, the invention relates to methods for  
the derivatization and reformulation of the p-GlcNAc.  
Additionally, the present invention relates to the  
20 uses of pure p-GlcNAc, its derivatives, and/or its  
reformulations.

2. BACKGROUND OF THE INVENTION

There exists today an extensive literature on the  
25 properties, activities, and uses of polysaccharides  
that consist, in part, of p-GlcNAc. A class of such  
materials has been generically referred to as  
"chitin", while deacetylated chitin derivatives have  
been referred to as "chitosan". When these terms were  
30 first used, around 1823, it was believed that chitin  
and chitosan always occurred in nature as distinct,  
well-defined, unique, and invariant chemical species,  
with chitin being fully acetylated and chitosan being  
fully deacetylated compositions. It was approximately  
35 a century later, however, before it was discovered

that the terms "chitin" and "chitosan" are, in fact, very ambiguous. Rather than referring to well-defined compounds, these terms actually refer to a family of compounds that exhibit widely differing physical and chemical properties. These differences are due to the products' varying molecular weights, varying degrees of acetylation, and the presence of contaminants such as covalently bound, species-specific proteins, single amino acid and inorganic contaminants. Even today, the terms "chitin" and "chitosan" are used ambiguously, and actually refer to poorly defined mixtures of many different compounds.

For example, the properties of "chitins" isolated from conventional sources such as crustacean outer shells and fungal mycelial mats are unpredictably variable. Such variations are due not only to species differences but are also due to varying environmental and seasonal effects that determine some of the biochemical characteristics of the "chitin"-producing species. In fact, the unpredictable variability of raw material is largely responsible for the slow growth of chitin-based industries.

No reports exist today in the scientific literature describing the isolation and production, from material sources, of pure, fully acetylated p-GlcNAc, *i.e.*, a product or products uncontaminated by organic or inorganic impurities. While McLachlan *et al.* (McLachlan, A.G. *et al.*, 1965, Can. J. Botany 43:707-713) reported the isolation of chitin, subsequent studies have shown that the "pure" substance obtained, in fact contained proteins and other contaminants.

Deacetylated and partially deacetylated chitin preparations exhibit potentially beneficial chemical properties, such as high reactivity, dense cationic

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charges, powerful metal chelating capacity, the ability to covalently attach proteins, and solubility in many aqueous solvents. The unpredictable variability of these preparations, as described above, however, severely limits the utility of these heterogenous compounds. For example, the currently available "chitins" and "chitosans" give rise to irreproducible data and to unacceptably wide variations in experimental results. Additionally, the available preparations are not sufficiently homogenous or pure, and the preparation constituents are not sufficiently reproducible for these preparations to be acceptable for use in applications, especially in medical ones. Thus, although extremely desirable, a true, purified preparations of chitin and chitosan, whose properties are highly reproducible and which are easily manufactured, do not currently exist.

### 3. SUMMARY OF THE INVENTION

The present invention relates, first, to an isolated, easily produced, pure p-GlcNAc species. The p-GlcNAc of the invention is a polymer of high molecular weight whose constituent monosaccharides are attached in a  $\beta$ -1 $\rightarrow$ 4 conformation, and which is free of proteins, substantially free of other organic contaminants, and substantially free of inorganic contaminants.

The importance of the present invention resides in the fact that the problem of unpredictable raw material variability has been overcome. It is, for the first time, possible to produce, by simple means, and on a commercial scale, biomedically pure, p-GlcNAc of high molecular weight and consistent properties. The material produced in the present invention is highly crystalline and is produced from carefully

controlled, aseptic cultures of one of a number of marine microalgae, preferably diatoms, which have been grown in a defined medium.

The present invention further describes  
5 derivatives and reformulations of p-GlcNAc as well as methods for the production of such derivatives and reformulations. Such derivatizations may include, but are not limited to polyglucosamine and its  
10 derivatives, and such reformulations may include, but are not limited to membranes, filaments, non-woven textiles, sponges, and three dimensional matrices. Still further, the present invention relates to methods for the purification of the p-GlcNAc of the invention from microalgae, preferably diatom, sources.  
15 Additionally, the present invention relates to the uses of the purified p-GlcNAc, its derivatives, and/or its reformulations. Among these uses are novel commercial applications relating to such industries as the biomedical, pharmaceutical, and cosmetic  
20 industries, all of which require starting materials of the highest degree of purity. For example, the p-GlcNAc materials of the invention may be formulated to exhibit controllable biodegradation properties, and, further, may be used as part of slow drug delivery  
25 systems, as cell encapsulation systems, and as treatments for the prevention of post-surgical adhesions.

#### 4. BRIEF DESCRIPTION OF THE FIGURES

30 FIG. 1. Chemical structure of 100% p-GlcNAc.  
"n" refers to an integer ranging from about 4,000 to about 150,000, with about 4,000 to about 15,000 being preferred.

FIG. 2. Carbohydrate analysis of p-GlcNAc, Gas Chromatography-Mass Spectroscopy data. Solid squares represent p-GlcNAc purified using the acid treatment/neutralization variation of the  
5 Chemical/Biological method, as described in Section 5.3.2, below.

FIG. 3A. Circular dichroism spectra of solid membranes of pure p-GlcNAc.  
10

FIG. 3B. Circular dichroism spectra of solid membranes of Deacetylated p-GlcNAc. The disappearance of the 211 nm minimum and 195 nm maximum observed in pure p-GlcNAc (FIG. 3A) indicates complete  
15 Deacetylation under the conditions used, as described in Section 5.4 below.

FIG. 4A. Infra-red spectra analyses of thin membranes of pure diatom p-GlcNAc prepared by the  
20 mechanical force purification method, top, and the chemical/biological purification method, bottom.

FIG. 4B. Infra-red spectra analyses of two preparations of commercial "chitin" cast into  
25 membranes according to the methods detailed in Section 5.5, below.

FIG. 4C. Infra-red spectra analyses of pure p-GlcNAc which was modified by heat denaturation (top)  
30 and by chemical deacetylation (bottom), according to the methods detailed in Section 5.4, below.

FIG. 4D. Infra-red spectrum analysis of a p-GlcNAc membrane derived from the diatom Thalassiosira  
35 fluviatilis, using the chemical/biological

purification method, as detailed in Section 5.3.2, below.

5 FIG. 4E. Infra-red spectrum analysis of a p-GlcNac membrane prepared by the mechanical force purification method, as described in Section 5.3.1, below, following autoclaving.

10 FIG. 5A. NMR analysis of p-GlcNac purified using the chemical/biological purification method as described in Section 5.3.2, below. Chart depicting peak amplitudes, areas, and ratios relative to reference controls. Ratio of total areas of peaks.

15 FIG. 5B. NMR analysis of p-GlcNac purified using the chemical/biological purification method as described in Section 5.3.2. The graph depicts the ratios of total areas of peaks.

20 FIG. 6A-6B. Transmission electron micrographs (TEM) of a p-GlcNac membrane prepared by the mechanical force purification method as described in Section 5.3.1, below. Magnification: 6A: 4190x; 6B: 16,250x.

25 FIG. 7A-7B. Transmission electron micrographs (TEM) of a p-GlcNac membrane by HF treatment as described in the discussion of the chemical/biological purification method in Section 5.3.2, below.  
30 Magnification: 7A: 5270x; 7B: 8150x.

35 FIG. 8A-8B. Transmission electron micrographs (TEM) of a p-GlcNac membrane prepared by the acid treatment/neutralization variation of the chemical/biological purification method, as described

in Section 5.3.2, below. Magnification: 8A: 5270x;  
8B: 16,700x.

FIG. 9A. Scanning electron micrograph depicting  
5 a p-GlcNAc membrane prepared by the acid  
treatment/neutralization variation of the  
chemical/biological purification method as described  
in Section 5.3.2, below.  
Magnification: 200x.

10

FIG. 9B. Scanning electron micrograph depicting  
a p-GlcNAc membrane prepared by the acid  
treatment/neutralization variation of the  
chemical/biological purification method as described  
15 in Section 5.3.2, below.  
Magnification: 1000x.

FIG. 9C. Scanning electron micrograph depicting  
a p-GlcNAc membrane prepared by the acid  
20 treatment/neutralization variation of the  
chemical/biological purification method as described  
in Section 5.3.2, below.  
Magnification: 5000x.

25 FIG. 9D. Scanning electron micrograph depicting  
a p-GlcNAc membrane prepared by the acid  
treatment/neutralization variation of the  
chemical/biological purification method as described  
in Section 5.3.2, below.  
30 Magnification: 10,000x.

FIG. 9E. Scanning electron micrograph depicting  
a p-GlcNAc membrane prepared by the acid  
treatment/neutralization variation of the  
35

chemical/biological purification method as described in Section 5.3.2, below. Magnification: 20,000x.

FIG. 10A-10B. Scanning electron micrographs of a  
5 pure p-GlcNAC membrane made from material which was initially produced using the cell dissolution/neutralization purification method described in Section 5.3, below, dissolved in dimethylacetamide/lithium chloride, and  
10 reprecipitated in H<sub>2</sub>O into a mat, as described below in Section 5.5. Magnification: 10A: 1000x; 10B: 10,000x.

FIG. 11A-11B. Scanning electron micrographs of a deacetylated p-GlcNAC mat. Magnification: 11A:  
15 1000x; 11B: 10,000x.

FIG. 12A-12B. Photographs of diatoms. Note the p-GlcNAC fibers extending from the diatom cell bodies.

20 FIG. 13. Diagram depicting some of the possible p-GlcNAC and deacetylated p-GlcNAC derivatives of the invention. (Adapted from S. Hirano, "Production and Application of Chitin and Chitosan in Japan", in "Chitin and Chitosan", 1989, Skjak-Braek, Anthonsen,  
25 and Sanford, eds. Elsevier Science Publishing Co., pp. 37-43.)

FIG. 14. Cell viability study of cells grown in the presence or absence of p-GlcNAC membranes. Closed  
30 circle (●): cells grown on p-GlcNAC matrix; open circles (○): cells grown in absence of matrix.

FIG. 15A-15B. SEM micrographs of transformed mouse fibroblast cells grown on p-GlcNAC membranes.  
35 Magnification: 15A: 1000x; 15B: 3000x.

FIG. 16A. Scanning electron micrograph (SEM) of a collagen-only control material prepared according to the method described, below, in Section 13.1. Magnification 100x.

5

FIG. 16B. Scanning electron micrograph (SEM) of a collagen/p-GlcNAc hybrid material prepared according to the method described, below, in Section 13.1.

Ratio collagen suspension:p-GlcNAc suspension equals  
10 3:1, with final concentrations of 7.5 mg/ml collagen and 0.07 mg/ml p-GlcNAc. Magnification 100x.

FIG. 16C. Scanning electron micrograph (SEM) of a collagen/p-GlcNAc hybrid material prepared according  
15 to the method described, below, in Section 13.1.

Ratio collagen suspension:p-GlcNAc suspension equals  
1:1, with final concentrations of 5.0 mg/ml collagen  
and 0.12 mg/ml p-GlcNAc. Magnification 100x.

20 FIG. 16D. Scanning electron micrograph (SEM) of a collagen/p-GlcNAc hybrid material prepared according to the method described, below, in Section 13.1.

Ratio collagen suspension:p-GlcNAc suspension equals  
2:2, with final concentrations of 10.0 mg/ml collagen  
25 and 0.25 mg/ml p-GlcNAc. Magnification 100x.

FIG. 16E. Scanning electron micrograph (SEM) of a collagen/p-GlcNAc hybrid material prepared according to the method described, below, in Section 13.1.

30 Ratio collagen suspension:p-GlcNAc suspension equals

35

1:3, with final concentrations of 2.5 mg/ml collagen and 0.25 mg/ml p-GlcNAc. Magnification 100x.

FIG. 17A. SEM of mouse 3T3 fibroblast cells  
5 cultured on the collagen-only control material of FIG. 16A, above. Magnification 100x.

FIG. 17B. SEM of mouse 3T3 fibroblast cells  
cultured on the collagen/p-GlcNAc material of FIG.  
10 16B, above. Magnification 100x.

FIG. 17C. SEM of mouse 3T3 fibroblast cells  
cultured on the collagen/p-GlcNAc material of FIG.  
16C, above. Magnification 100x.

15

FIG. 17D. SEM of mouse 3T3 fibroblast cells  
cultured on the collagen/p-GlcNAc material of FIG.  
16D, above. Magnification 100x.

20 FIG. 18. Transformed NMR data curves, used to obtain areas for each carbon atom and to then calculate the CH<sub>3</sub>(area) to c-atom(area) ratios.

FIG. 19. Typical p-GlcNAc C<sup>13</sup>-NMR spectrum. The  
25 individual peaks represent the contribution to the spectrum of each unique carbon atom in the molecule.

FIG. 20. Transformed NMR spectrum data representing values calculated for CH<sub>3</sub>(area) to C-atom(area) ratios. Top: Graphic depiction of data;  
30 bottom: numerical depiction of data.

FIG. 21A-G. Three dimensional p-GlcNAc matrices produced in various solvents. Specifically, the p-  
35 GlcNAc matrices were produced in distilled water (FIG.



21A, FIG. 21D), 10% methanol in distilled water (FIG. 21B), 25% methanol in distilled water (FIG. 21C), 10% ethanol in distilled water (FIG. 21E), 25% ethanol in distilled water (FIG. 21F) and 40% ethanol in  
5 distilled water (FIG. 21G). Magnification: 200x. A scale marking of 200 microns is indicated on each of these figures.

FIG. 22A-G. Fibroblast cells grown on three  
10 dimensional p-GlcNAc matrices prepared by lyophilizing p-GlcNAc in distilled water. Magnification: 100x (FIGS. 22A, 22E), 500x (FIG. 22B), 1000x (FIGS. 22C, 22F), 5000x (FIGS. 22D, 22G). Scales marking 5, 20, 50, or 200 microns, as indicated, are included in each  
15 of the figures.

FIG. 23. A typical standard curve obtained using the procedure described, below, in Section 18.1. A standard curve such as this one was used in the  
20 lysozyme-chitinase assay also described, below, in Section 18.1.

FIG. 24. p-GlcNAc lysozyme digestion data. The graph presented here depicts the accumulation of N-  
25 acetylglucosamine over time, as p-GlcNAc membranes are digested with lysozyme. The graph compares the degradation rate of fully acetylated p-GlcNAc to partially (50%) deacetylated p-GlcNAc, and demonstrates that the degradation rate for the  
30 partially deacetylated p-GlcNAc was substantially higher than that of the fully acetylated p-GlcNAc material.

FIG. 25. p-GlcNAc lysozyme digestion data. The  
35 graph presented here depicts the accumulation of N-

acetylglucosamine over time, as p-GlcNAc membranes are digested with lysozyme. The graph compares the degradation rate of two partially deacetylated p-GlcNAc membranes (specifically a 25% and a 50%  
5 deacetylated p-GlcNAc membrane). The data demonstrate that the degradation rate increases as the percent of deacetylation increases, with the degradation rate for the 50% deacetylated p-GlcNAc membrane being  
10 substantially higher than that of the 25% deacetylated p-GlcNAc membrane.

FIG. 26A-26E. p-GlcNAc in vivo biodegradability data. FIG. 26A-26C depict rats which have had prototype 1 (fully acetylated p-GlcNAc) membrane  
15 abdominally implanted, as described, below, in Section 18.1. FIG. 26A shows a rat at day 0 of the implantation; FIG. 26B shows a rat at day 14 post-implantation; FIG. 26C shows a rat at day 21 post-implantation. FIG. 26D-26E depict rats which have had  
20 prototype 3A (lyophilized and partially deacetylated p-GlcNAc membrane) abdominally implanted, as described, below, in Section 18.1. FIG. 26D shows a rat at day 0 of the implantation; FIG. 26E shows a rat at day 14 post-implantation.

25

FIG. 27. The graph depicted here illustrates data concerning the percent increase in tumor size of animals which either received no treatment (●) or received p-GlcNAc-lactate/5'Flurouracil (FU) (O), as  
30 described, below, in Section 20.1.

FIG. 28. The graph depicted here illustrates data concerning the percent increase in tumor size of animals which either received p-GlcNAc-lactate alone  
35

(●) or received p-GlcNAc-lactate/5'Flurouracil (FU) (O), as described, below, in Section 20.1.

5 FIG. 29. The graph depicted here illustrates data concerning the percent increase in tumor size of animals which either received no treatment (●) or received p-GlcNAc-lactate/mitomycin (mito) (O), as described, below, in Section 20.1.

10 FIG. 30. The graph depicted here illustrates data concerning the percent increase in tumor size of animals which either received p-GlcNAc-lactate alone (●) or received p-GlcNAc-lactate/5' mitomycin (mito) (O), as described, below, in Section 20.1.

15 FIG. 31. The bar graph depicted here illustrates the average percent change in tumor size per animal of animals treated with p-GlcNAc/5'FU high dose (bar 1), p-GlcNAc/5'FU low dose (bar 2), p-GlcNAc membrane  
20 alone (bar 3), and untreated (bar 4). N=4 for bars 1 and 2, n=2 for bars 3 and 4.

##### 5. DETAILED DESCRIPTION OF THE INVENTION

Presented below, is, first, a description of  
25 physical characteristics of the purified p-GlcNAc species of the invention, of the p-GlcNAc derivatives, and of their reformulations. Next, methods are described for the purification of the p-GlcNAc species of the invention from microalgae, preferably diatom,  
30 starting sources. Third, derivatives and reformulations of the p-GlcNAc, and methods for the production of such derivatives and reformulations are presented. Finally, uses are presented for the p-GlcNAc, p-GlcNAc derivatives and/or p-GlcNAc  
35 reformulations of the invention.

### 5.1 p-GlcNAc

The p-GlcNAc polysaccharide species of the invention is a polymer of high molecular weight ranging from a weight average of about 800,000 daltons to about 30 million daltons, based upon gel permeation chromatography measurements. Such a molecular weight range represents a p-GlcNAc species having about 4,000 to about 150,000 N-acetylglucosamine monosaccharides attached in a  $\beta$ -1 $\rightarrow$ 4 configuration, with about 4,000 to about 15,000 N-acetylglucosamine monosaccharides being preferred (FIG. 1).

The variability of the p-GlcNAc of the invention is very low, and its purity is very high, both of which are evidenced by chemical and physical criteria. Among these are chemical composition and non-polysaccharide contaminants. First, chemical composition data for the p-GlcNAc produced using two different purification methods, both of which are described in Section 5.3, below, is shown in Table I below. As can be seen, the chemical composition of the p-GlcNAc produced by both methods is, within the bounds of experimental error, the same as the formula compositions of p-GlcNAc. Second, as is also shown in Table I, the p-GlcNAc produced is free of detectable protein contaminants, is substantially free of other organic contaminants such as free amino acids, and is substantially free of inorganic contaminants such as ash and metal ions (the p-GlcNAc of the invention may contain up to about 0.05% trace metals). Further, the p-GlcNAc of the invention exhibits a very low percentage of bound water.

TABLE I

CHEMICAL ANALYSIS DATA (% by weight)

5           Theoretical Values for Pure p-GlcNAc:

	Carbon	-	47.29
	Hydrogen	-	6.40
	Nitrogen	-	6.89
	Oxygen	-	39.41
10	Protein	-	0.00

Experimental Data on p-GlcNAc Mats:  
 (Number of experimental batches for each membrane type being greater than 30 for each membrane type)

15	<u>MECHANICAL FORCE</u>		<u>CHEMICAL/BIOLOGICAL</u>		
	<u>METHOD</u>		<u>METHOD</u>		
	Normalized <sup>1</sup>	% Dev.	Normalized <sup>1</sup>	% Dev.	
	Carbon	47.21 ± 0.08	-0.17	47.31 ± 0.11	+0.04
	Hydrogen	6.45 ± 0.08	+0.78	6.34 ± 0.08	-0.94
	Nitrogen	6.97 ± 0.18	+0.87	6.94 ± 0.16	+0.73
20	Oxygen	39.55 ± 0.36	+0.36	39.41 ± 0.10	0.00
		<u>Average Values</u>		<u>Average Values</u>	
	Protein	0.00		0.00	
	Ash	1.30		0.98	
	Moisture	2.0		1.2	

25   -----  
<sup>1</sup> Raw analytical data have been normalized to account for ash and moisture content of the samples.

The pure p-GlcNAc of the invention exhibits a carbohydrate analysis profile substantially similar to that shown in FIG. 2. The primary monosaccharide of the pure p-GlcNAc of the invention is N-acetylglucosamine. Further, the pure p-GlcNAc of the invention does not contain the monosaccharide glucosamine.

The circular dichroism (CD) and sharp infra-red spectra (IR) of the p-GlcNAc of the invention are shown in FIGS. 3A, and FIGS. 4A and 4D, respectively, which present analyses of material produced using the methods described in Section 5.3, below. Such physical data corroborates that the p-GlcNAc of the invention is of high purity and crystallinity. The methods used to obtain the CD and IR data are described, below, in the Working Example in Section 6.

10 NMR analysis of the pure p-GlcNAc of the invention exhibits a pattern substantially similar to that seen in FIGS. 5A, 5B, 18A and 18B. Such an NMR pattern indicates not only data which is consistent with the p-GlcNAc of the invention being a fully acetylated polymer, but also demonstrates the lack of contaminating organic matter within the p-GlcNAc species.

The electron micrographic structure of the p-GlcNAc of the invention, as produced using the methods described in Section 5.3, below and demonstrated in the Working Examples presented, below, in Section 8 and 9, is depicted in FIGS. 6A through FIG. 9E.

The p-GlcNAc of the invention exhibits a high degree of biocompatibility. Biocompatibility may be determined by a variety of techniques, including, but not limited to such procedures as the elution test, intramuscular implantation, or intracutaneous or systemic injection into animal subjects. Briefly, an elution test (U.S. Pharmacopeia XXII, 1990, pp. 1415-1497; U.S. Pharmacopeia XXII, 1991, Supplement 5, pp. 2702-2703) is designed to evaluate the biocompatibility of test article extracts, and assays the biological reactivity of a mammalian cell culture line which is sensitive to extractable cytotoxic articles (such as, for example, the L929 cell line) in

response to the test article. The Working Example presented in Section 10, below, demonstrates the high biocompatibility of the p-GlcNAc of the invention.

5                   5.2 METHODS OF PRODUCING MICROALGAL SOURCES OF p-GlcNAc

                  5.2.1 MICROALGAL SOURCES OF p-GlcNAc

                  The p-GlcNAc of the invention is produced by, and  
10               may be purified from, microalgae, preferably diatoms.  
                  The diatoms of several genera and numerous species  
                  within such genera may be utilized as p-GlcNAc  
                  starting sources. Each of these diatoms produce  
                  fibers composed of p-GlcNAc which extend from their  
15               cell bodies. See FIG. 12A-12B for photographs of such  
                  diatoms. The diatoms which may be used as starting  
                  sources for the production of the p-GlcNAc of the  
                  invention include, but are not limited to members of  
                  the *Coscinodiscus* genus, the *Cyclotella* genus, and the  
20               *Thalassiosira* genus, with the *Thalassiosira* genus  
                  being preferred.

                  Among the *Coscinodiscus* genus, the species of  
                  diatom that may be used to produce the p-GlcNAc of the  
                  invention include, but are not limited to the  
25               *concinnus* and *radiatus* species. The diatoms among the  
                  *Cyclotella* genus which may be used include, but are  
                  not limited to the *caspia*, *cryptica*, and *meneghiniana*  
                  species. The *Thalassiosira* diatoms that may be  
                  utilized to produce the starting material for the p-  
30               GlcNAc of the invention include, but are not limited  
                  to the *nitzschoides*, *aestivalis*, *antarctica*,  
                  *deciphens*, *eccentrica*, *floridana*, *fluviatilis*,  
                  *gravida*, *guillardii*, *hyalina*, *minima*, *nordenskioldii*,  
                  *oceanica*, *polychorda*, *pseudonana*; *rotula*, *tubifera*,  
35               *tumida*, and *weissflogii* species, with the *fluviatilis*  
                  and *weissflogii* species being preferred.

Diatoms such as those described above may be obtained, for example, from the culture collection of the Bigelow Laboratory for Ocean Sciences, Center for Collection of Marine Phytoplankton (McKown Point, West Boothbay Harbor, Maine, 04575).

#### 5.2.2 METHODS FOR GROWING DIATOMS

Any of the diatoms described in Section 5.2.1, above, may be grown by utilizing, for example, the methods described in this section. New diatom cultures are initiated by inoculating, under sterile conditions, Nutrient Medium with an aliquot of a mature diatom culture. The Nutrient Medium must be free of all other microorganisms, therefore all materials, including water, organic components, and inorganic components used in the preparation of the Nutrient Medium must be sterile. In addition, it is mandatory that all procedures involved in this operation be conducted under strictly sterile conditions, *i.e.*, all containers, all transfers of substances from one vessel to another, etc. must be performed in a sterile environment. The quantity of Nutrient Medium to be prepared at one time should not exceed what is necessary to start a new culture. For example, Fernbach flasks which occupy approximately one square foot of surface may be used as vessels for the diatom cultures, and such vessels require one liter of Nutrient Medium for optimum growth of the diatom organism.

Preparation of the nutrient medium involves the following operations:

- a) Acquisition and processing of seawater
- b) Preparation of distilled and deionized water.
- c) Preparation of primary nutrient stocks



- d) Preparation of nutrient working stocks
- e) Preparation of the final nutrient medium

Filtered seawater may be obtained, for example, from the Marine Biology Laboratory (Woods Hole, Massachusetts). Seawater containers should be stored at 5° C. When required, the necessary volume of water may be filtered through a Buchner filtration unit, using a nitrocellulose filter membrane with 0.45 micron pore size (Millipore, Inc.). The seawater is then sterilized by autoclaving at, for example, 121° C. for 15 minutes per liter. On completion of the sterilization process, the capped are immediately cooled, preferably by transfer to a cold room capable of allowing the solutions to reach a temperature of approximately 5° C. When it is to be used, solutions are allowed to reach room temperature.

Tap water is distilled and deionized using standard equipment and procedures, and collected and stored in sterile, securely capped, preferably glass, containers.

Listed below are formulas which may be followed in preparing the stock solutions necessary for the preparation of the Nutrient Medium. It is to be understood that while such formulas are to be used as guides, it is intended that routine variations of such formulas which contribute to the preparation of a Nutrient Medium capable of sustaining microalgal diatom growth sufficient for the p-GlcNAc preparative processes described here also be within the scope of the present invention.

I. Trace Metal Primary Stocks (TMPS)

- a. 39 mM  $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$  (copper [II] sulfate pentahydrate) (9.8g copper [II] sulfate/L)

- 20 -

- b. 7.5 mM  $\text{ZnSO}_4 \cdot 7\text{H}_2\text{O}$  (Zinc sulfate heptahydrate) (22g zinc sulfate/L)
- c. 42 mM  $\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$  (Cobalt [II] chloride hexahydrate) (10g cobalt [II] chloride/L)
- 5 d. 91 mM  $\text{MnCl}_2 \cdot 4\text{H}_2\text{O}$  (Manganese [II] chloride tetrahydrate) 18g manganese [II] chloride/L)
- e. 26 mM  $\text{NaMoO}_4 \cdot 2\text{H}_2\text{O}$  (Sodium molybdate dihydrate) 6.3g sodium molybdate/L)
- f. 153.5 mM  $\text{H}_2\text{SeO}_3$  (Selenious acid) (12.9g
- 10 selenious acid/L).
- Sterile filter each nutrient with a filter of no greater than 0.2mm pore size.

II. Vitamin Primary Stocks (VPS)

- 15 a. 1 mg/ml Vitamin B12
- b. 0.1 mg/ml Biotin

Sterile filter both stocks with a filter of no greater than 0.2mm pore size.

20 III. Sodium Salts Working Stocks (SSWS)

- a. Sodium nitrate working stock: 0.88 M (75 g  $\text{NaNO}_3$ /L)
- b. Sodium phosphate monobasic monohydrate working stock: 36.2 mM  $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$  (5 g  $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$ /L)
- 25 c. Sodium metasilicate nonahydrate working stock: 0.11 M  $\text{Na}_2\text{SiO}_3 \cdot 9\text{H}_2\text{O}$  (30 g  $\text{Na}_2\text{SiO}_3 \cdot 9\text{H}_2\text{O}$ /L)

Sterile filter each of the SSWS with a filter of no greater than 0.2mm pore size.

30 IV. Trace Metal Working Stocks (TMWS)

- 11.7 mM  $\text{Na}_2\text{EDTA}$  (Ethylenediamine Tetraacetic acid, disodium salt dihydrate) (4.36 g/L)
- 11.7 mM  $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$  (Iron [III] chloride hexahydrate) (3.15 g/L)

35

1 ml/L of each of the six primary trace metal stocks listed above.

Sterile filter with a filter of no greater than 0.2mm pore size. Note that the trace metal working stock  
5 must be prepared fresh each time a new Nutrient Medium is assembled.

V. Vitamin Working Stock (VWS)

1.0  $\mu\text{g/ml}$  Biotin (1.0 ml primary Biotin  
10 Stock/100 ml)

1.0  $\mu\text{g/ml}$  Vitamin B12 (0.1 ml Vitamin B12 primary stock/100 ml)

20 mg of Thiamine HCl (Thiamine hydrochloride/100 ml).

15 Sterile filter with a filter of no greater than 0.2mm pore size. Note that a new Vitamin Working Stock should be prepared fresh every time a new nutrient medium is being assembled.

20 Described below are techniques which may be followed for the preparation of Nutrient Medium and for diatom culturing. It is to be understood that, in addition to these techniques, any routine variation in the formulas and/or procedures described herein which  
25 result in a Nutrient Medium and in procedures capable of sustaining diatom growth sufficient for the preparative processes described herein is intended to be within the scope of the present invention.

Nutrient Medium may be prepared, for example, as  
30 follows: To each liter of filtered and sterilized seawater may be added 1 ml of the  $\text{NaNO}_3$  working stock, 1 ml of the  $\text{NaH}_2\text{PO}_4 \cdot \text{H}_2\text{O}$  working stock, 1 ml of the Trace Metal working stock, and 1 ml of the  $\text{Na}_2\text{SiO}_3 \cdot 9\text{H}_2\text{O}$  working stock. Simultaneously with the addition of  
35  $\text{Na}_2\text{SiO}_3 \cdot 9\text{H}_2\text{O}$ , 2 mls of 1 N HCl may be added and the

solution may be shaken to mix. Next, 1.5 mls 1 N NaOH may be added and the solution may again be shaken to mix. Finally, 0.5 ml of the Vitamin working stock may be added.

5           In order to grow a new diatom culture, 7 ml of a mature culture, (having a cell density of approximately  $1 \times 10^5$  cells/ml), may be transferred to a sterile container containing 100 ml of sterile Nutrient Medium, which may be prepared according to  
10 the methods described above. The inoculated culture may then be incubated for 8 days under the following conditions:

          Temperature: 20 degrees Centigrade

          Constant illumination.

15           Agitation: Gentle swirling of flasks once for two or three seconds every morning and every evening.

          After 8 days of incubation, 80 ml of this incubated culture may be transferred, under sterile  
20 conditions, to 1000 ml of Nutrient Medium, which may, for example, be contained in a 2.8 L Fernbach flask, protected by a cotton wool plug covered by cheesecloth. Such a culture may be allowed to incubate and grow to the desired cell density, or  
25 alternatively, may be used to inoculate new diatom cultures. Once a culture reaches a desired cell density, the culture's p-GlcNAc fibers may be harvested, and the p-GlcNAc of the invention may be purified, using methods such as those described below  
30 in Section 5.3, below.

          CO<sub>2</sub> may be dissolved in the culture solution in order to maintain a culture pH of approximately 7 to 8, with approximately 7.4 being preferred. The maintenance of such a neutral pH environment, greatly

increases the p-GlcNAc yield that may be obtained from each diatom culture.

5.3 METHODS FOR ISOLATION, PURIFICATION, AND CONCENTRATION OF p-GlcNAc FIBERS

5 Presented in this Section are methods which may be utilized for the preparation of p-GlcNAc fibers from diatom cultures such as those described, above, in Section 5.2.

10 While each of the methods described below for the purification of p-GlcNAc from microalgae, preferably diatom, starting sources produces very pure, unadulterated, crystalline p-GlcNAc, each of the methods yields p-GlcNAc having specific  
15 characteristics and advantageous features. For example, the p-GlcNAc of the invention purified via the Mechanical Force method presented in Section 5.3.1, below, produces a p-GlcNAc membrane that provides a superior substrate for the attachment of  
20 cells to the p-GlcNAc. The second method, described below in Section 5.3.2, the Chemical/Biological method, produces a much higher average yield than the average p-GlcNAc yield produced by the Mechanical Force method. Additionally, the acid treatment/  
25 neutralization variation described as part of the Chemical/Biological method of Section 5.3.2, below, produces extremely long p-GlcNAc fibers, with some fibers being in excess of 100  $\mu\text{m}$ , and of very high molecular weight, as high as 20-30 million daltons.

30 5.3.1 MECHANICAL FORCE METHOD FOR PREPARATION OF PURE p-GlcNAc

The p-GlcNAc fibers may be separated from diatom cell bodies by subjecting the contents of the culture to an appropriate mechanical force. Such a mechanical  
35 force may include, but is not limited to, a shear

force generated by, for example, a colloid mill, an ultrasound device, or a bubble generator, or a cutting force generated by, for example, a Waring blender.

The resulting suspension of diatom cell bodies and p-GlcNAc fibers are then segregated. For example, the suspension may be subjected to a series of centrifugation steps which segregate the p-GlcNAc fibers from the cell bodies, yielding a clear supernatant exhibiting little, if any, visible flocculent material. A fixed angle rotor, and a temperature of about 10° C. are preferred for the centrifugation steps. The speed, duration, and total number of centrifugation steps required may vary depending on, for example, the specific centrifugation rotor being used, but the determination of the values for such parameters will be apparent to one of ordinary skill in the art.

The p-GlcNAc fibers in the supernatant may then be concentrated using techniques well known to those of skill in the art. Such techniques may include, but are not limited to suction and filtration devices.

Finally, the concentrated p-GlcNAc fibers are washed with, for example, distilled-deionized water, HCl and ethanol, or other appropriate solvents, preferably solvents, such as alcohols, in which both organic and inorganic materials dissolve.

The Working Example presented in Section 7, below, demonstrates the use of this method for the purification of p-GlcNAc.

30

#### 5.3.2. CHEMICAL/BIOLOGICAL METHOD FOR PURIFICATION OF p-GlcNAc

In this method, p-GlcNAc fibers are separated from diatom cell bodies by subjecting them to chemical and/or biological agents as described in more detail below.

35

Diatom cultures may be treated with a chemical capable of weakening diatom cell walls, which leads to a release of the p-GlcNAc fibers without altering their structure. Such a chemical may include, but is  
5 not limited to, hydrofluoric acid (HF).

Alternatively, a mature diatom culture may be treated with a biological agent capable of altering a biological process may be used to inhibit p-GlcNAc fiber synthesis, thus releasing the fibers already  
10 present. For example, such an agent may include, but is not limited to, polyoxin-D, an inhibitor of the enzyme N-acetylglucosaminyl-P-transferase.

The cell bodies and p-GlcNAc-containing fibers of diatom cultures treated with a member of the above  
15 described chemical or biological agents are then segregated. For example, the contents of treated diatom cultures may be allowed to settle such that the contents of the cultures are allowed to form two distinct layers. The upper layer will contain  
20 primarily the p-GlcNAc fibers, while the bottom layer will contain the cell bodies. The upper p-GlcNAc fiber-containing layer may be siphoned off, leaving behind the settled cellular material of the bottom layer.

25 The siphoned off p-GlcNAc fiber-containing layer may then be further purified to remove protein and other unwanted matter by treatment with a detergent that will not damage the p-GlcNAc fibers. Such a detergent may include, but is not limited to, sodium  
30 dodecyl sulfate (SDS).

When acid treatment, such as HF treatment, is used to separate p-GlcNAc fibers from diatom cell bodies, a step may be included for the dispersal of the fibers. Such a step may include, but is not  
35 limited to, the use of mechanical force for fiber

dispersal, such as a step in which the fibers are subjected to a Waring blender dispersal.

Alternatively, the acid-treated suspension may, in an optional step, be neutralized prior to further  
5 purification by detergent treatment. Such neutralization will, in general, change the pH of the suspension from approximately 1.8 to approximately 7.0, and may be accomplished by, for example, the  
10 addition of an appropriate volume of 1M Tris (pH 8.0) or the addition of an appropriate volume of sodium hydroxide (NaOH). Neutralization, in general, yields pure p-GlcNAc fibers of a substantially greater length than the other purification methods discussed herein.

The purified p-GlcNAc fibers may then be  
15 concentrated using techniques well known to those of skill in the art, such as by utilizing a suction and filtration device. Finally, the p-GlcNAc fibers are washed, in a series of steps with distilled-deionized water, HCl and ethanol, or other appropriate solvents,  
20 preferably solvents, such as alcohols, in which both organic and inorganic materials dissolve.

The Working Example presented, below, in Section 8 demonstrates the successful utilization of such a purification method.

25

#### 5.4 DERIVATIZATION OF p-GlcNAc

The pure, fully acetylated p-GlcNAc of the invention may be derivatized, by utilizing a variety of controlled conditions and procedures, into a large  
30 range of different compounds. See FIG. 13 for a diagram depicting some of these compounds. Such derivatized compounds may include, but are not limited to, partially or completely deacetylated p-GlcNAc, which has been modified via chemical and/or enzymatic  
35 means, as described in further detail, below.



Additionally, p-GlcNAc, or its deacetylated derivative, may be derivatized by being sulfated, phosphorylated, and/or nitrated. Further, as detailed below, O-sulfonyl, N-acyl, O-alkyl, N-alkyl, 5 deoxyhalogen, and N-alkylidene and N-arylidene and other derivatives may be prepared from the p-GlcNAc or deacetylated p-GlcNAc of the invention. The deacetylated p-GlcNAc of the invention may also be used to prepare a variety of organic salts and/or 10 metal chelates. Further, the p-GlcNAc, or a derivative thereof, of the invention may have attached to it, either covalently or non-covalently, any of a variety of molecules. Still further, the p-GlcNAc of the invention, or a derivative thereof, may be 15 subjected to controlled hydrolysis conditions which yield groups of molecules having uniform and discrete molecular weight characteristics.

One or more of the monosaccharide units of the p-GlcNAc of the invention may be deacetylated to form a 20 poly- $\beta$ -1 $\rightarrow$ 4-N-glucosamine species. A poly- $\beta$ -1 $\rightarrow$ 4-N-glucosamine species of the invention in which each of the monosaccharide units of the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of the invention has been deacetylated will have a molecular weight of about 25 640,000 daltons to about 24 million daltons, with about 640,000 daltons to about 2.4 million daltons being preferred. A species with such a molecular weight range represents a species having about 4000 to about 150,000 glucosamine monosaccharides covalently 30 attached in a  $\beta$ -1 $\rightarrow$ 4 configuration, with about 4,000 to about 15,000 glucosamine monosaccharides being preferred. At least one of the monosaccharide units of the poly- $\beta$ -1 $\rightarrow$ 4-N-glucosamine species may remain acetylated, with about 25% to about 75% acetylation

being preferred, and about 30% acetylation being most preferred.

The p-GlcNAc of the invention may be deacetylated by treatment with a base to yield glucosamines with  
5 free amino groups. This hydrolysis process may be carried out with solutions of concentrated sodium hydroxide or potassium hydroxide at elevated temperatures. To precisely control the extent of deacetylation and to avoid degradation of the main  
10 carbohydrate chain of the polysaccharide molecule, however, it is preferable that an enzymatic procedure utilizing a chitin deacetylase enzyme be used for p-GlcNAc deacylation. Such a deacetylase enzymatic procedure is well known to those of skill in the art  
15 and may be performed as in (U.S. Patent No. 5,219,749), which is incorporated herein, by reference, in its entirety.

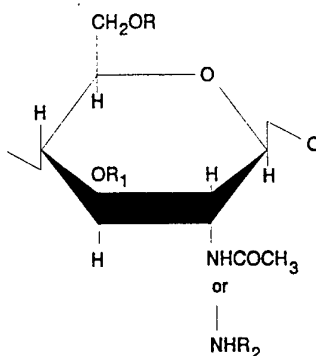
One or more of the monosaccharide units of the p-GlcNAc of the invention may be derivatized to contain  
20 at least one sulfate group, or, alternatively, may be phosphorylated or nitrated, as depicted below:

25

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5



10 where, R and/or R<sub>1</sub>, in place of a hydrogen, and/or R<sub>2</sub>, in place of -COCH<sub>3</sub>, may be a sulfate (-SHO<sub>3</sub>), a phosphate (-P(OH)<sub>2</sub>), or a nitrate (-NO<sub>2</sub>) group.

15 Described below are methods by which such p-GlcNAc derivatives may be prepared. Before performing methods such as those described in this Section, it may be advantageous to first lyophilize, freeze in liquid nitrogen, and pulverize the p-GlcNAc starting material.

20 Sulphated p-GlcNAc derivatives may be generated, by, for example, a two step process. In the first step, O-carboxymethyl p-GlcNAc may be prepared from the p-GlcNAc and/or p-GlcNAc derivatives of the invention by, for example, utilizing techniques such as those described by Tokura *et al.* (Tokura, S. *et al.*, 1983, *Polym. J.* 15:485). Second, the sulfation step may be carried out with, for example, N, N-dimethyl-formamide-sulfur trioxide, according to techniques well known to those of skill in the art, such as are described by Schweiger (Schweiger, R.G., 1972, *Carbohydrate Res.* 21:219). The resulting product may be isolated as a sodium salt.

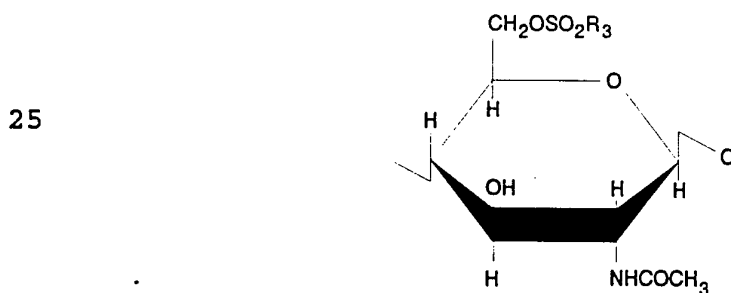
30 Phosphorylated p-GlcNAc derivatives of the invention may be prepared, for example, by utilizing techniques well known to those of skill in the art, such as those described by Nishi *et al.* (Nishi, N. *et al.*, 1986, in "Chitin in Nature and Technology,

35

Muzzarelli et al., eds. Plenum Press, New York, pp. 297-299). Briefly, p-GlcNAc/methanesulfonic acid mixture may be treated with phosphorus pentoxide (in an approximately 0.5 to 4.0 molar equivalent) with stirring, at a temperature of about 0° C. to about 5° C. Treatment may be for about 2 hours. The resulting product may then be precipitated and washed using standard techniques well known to those of skill in the art. For example, the sample may be precipitated with a solvent such as ether, centrifuged, washed with a solvent such as ether, acetone, or methanol, and dried.

Nitrated p-GlcNAc derivatives may be prepared by utilizing techniques well known to those of skill in the art, such as those described by Schorigin and Halt (Schorigin, R. and Halt, E., 1934, Chem. Ber. 67:1712). Briefly, p-GlcNAc and/or a p-GlcNAc derivative may be treated with concentrated nitric acid to form a stable nitrated product.

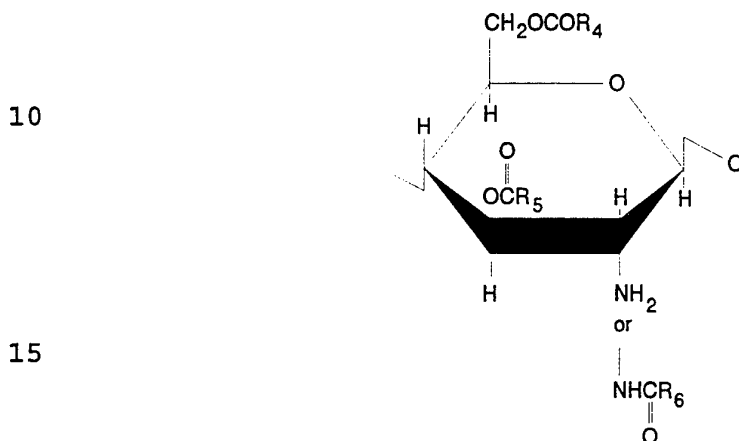
One or more of the monosaccharide units of the p-GlcNAc of the invention may contain a sulfonyl group, as depicted below:



30 where  $\text{R}_3$  may be an alkyl, an aryl, an alkenyl, or an alkynyl moiety. Such a derivative may be generated by well known methods such as the method described in Kurita et al. (Kurita, K. et al., 1990, Polym. Prep [Am. Chem. Soc., Div. Polym. Chem.] 31:624-625).  
35 Briefly, an aqueous alkali p-GlcNAc solution may be

reacted with a chloroform solution of tosyl chloride, and the reaction may then be allowed to proceed smoothly at low temperatures.

One or more of the monosaccharides of the p-GlcNAc of the invention or its deacetylated derivative may contain one or more O-acyl groups, as depicted below:

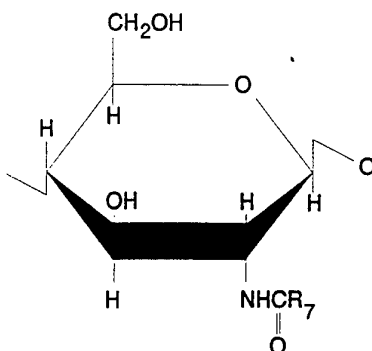


where  $R_4$  and/or  $R_5$ , in place of hydrogen, may be an alkyl, an alkenyl, or an alkynyl moiety, and  $R_6$  may be an alkyl, an alkenyl, or an alkynyl moiety. An example of such a derivative may be generated by well known methods such as those described by Komai (Komai, T. et al., 1986, in "Chitin in Nature and Technology", Muzzarelli et al., eds., Plenum Press, New York, pp. 497-506). Briefly, p-GlcNAc may be reacted with any of a number of suitable acyl chlorides in methanesulfonic acid to yield p-GlcNAc derivatives which include, but are not limited to, caproyl, capryl, lanroyl, or benzoyl derivatives.

30 One or more of the monosaccharides of the deacetylated p-GlcNAc of the invention may contain an N-acyl group, as depicted below:

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5



10 where R<sub>7</sub> may be an alkyl, an alkenyl, or an alkynyl moiety. Such a derivatization may be obtained by utilizing techniques well known to those of skill in the art, such as the technique described in Hirano *et al.* (Hirano, S. *et al.*, 1976, Carbohydrate Research

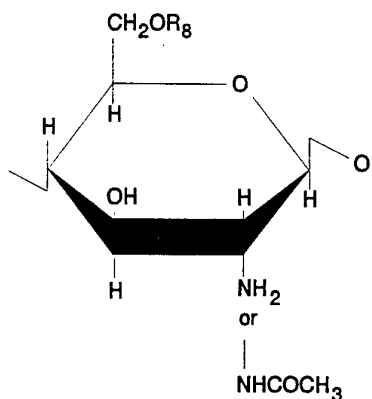
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Deacetylated p-GlcNAc is soluble in a number of aqueous solutions of organic acids. The addition of selected carboxylic anhydrides to such p-GlcNAc-containing solutions, in aqueous methanolic acetic acid, results in the formation of N-acyl p-GlcNAc derivatives.

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One or more of the monosaccharides of the deacetylated p-GlcNAc of the invention or of its deacetylated derivative, may contain an O-alkyl group, as depicted below:

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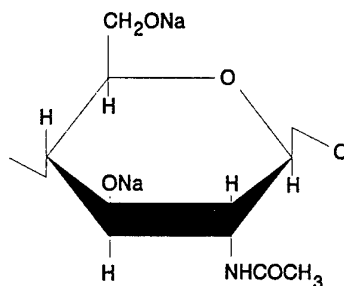


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where  $R_6$  may be an alkyl, and alkenyl, or a alkynyl moiety. Such a derivatization may be obtained by using techniques well known to those of skill in the art. For example, the procedure described by Maresh et al. (Maresh, G. et al., in "Chitin and Chitosan," Skjak-Braek, G. et al., eds., 1989, Elsevier Publishing Co., pp. 389-395). Briefly, deacetylated p-GlcNAc may be dispersed in dimethoxyethane (DME) and reacted with an excess of propylene oxide. The period of the reaction may be 24 hours, and the reaction takes place in an autoclave at 40 to 90° C. The mixture may then be diluted with water and filtered. The DME may be removed by distillation. Finally, the end-product may be isolated via lyophilization.

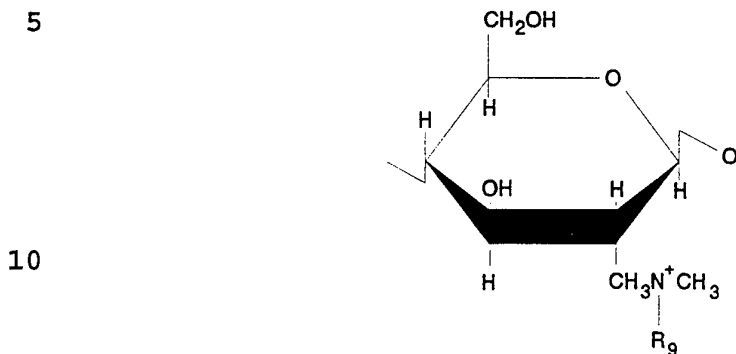
One or more of the monosaccharide units of the p-GlcNAc of the invention may be an alkali derivative, as depicted below:



Such a derivative may be obtained by using techniques well known to those of skill in the art. For example, a method such as that described by Noguchi et al. (Noguchi, J. et al., 1969, Kogyo Kagaku Zasshi 72:796-799) may be utilized. Briefly, p-GlcNAc may be steeped, under vacuo, in NaOH (43%, preferably) for a period of approximately two hours at about 0°C. Excess NaOH may then be removed by, for example, centrifugation in a basket centrifuge and by mechanical pressing.

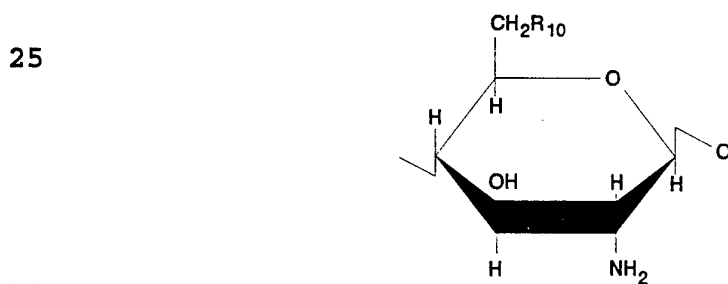
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One or more of the monosaccharide units of the deacetylated derivative of the p-GlcNAc of the invention may contain an N-alkyl group, as depicted below:



where  $R_9$  may be an alkyl, an alkenyl, or an alkynyl moiety. Such a derivatization may be obtained by utilizing, for example, a procedure such as that of Maresh et al. (Maresh, G. et al., in "Chitin and Chitosan," Skjak-Brack, G. et al., eds. 1989, Elsevier Publishing Co., pp. 389-395), as described, above, for the production of O-alkyl p-GlcNAc derivatives.

20 One or more of the monosaccharide units of the deacetylated derivative of the p-GlcNAc of the invention may contain at least one deoxyhalogen derivative, as depicted below:



where  $R_{10}$  may be F, Cl, Br, or I, with I being preferred. Such a derivative may be obtained by using techniques well known to those of skill in the art. For example, a procedure such as that described by Kurita et al. (Kurita, K. et al., 1990, Polym. Prep.

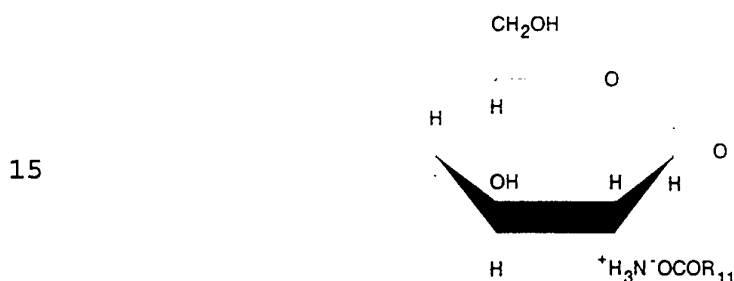
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[Am. Chem. Soc. Div. Polym. Chem.] 31:624-625) may be utilized. Briefly, a tosylated p-GlcNAC is made to react with a sodium halide in dimethylsulfoxide, yielding a deoxyhalogen derivative. p-GlcNAC

5 tosylation may be performed by reacting an aqueous alkali p-GlcNAC solution with a chloroform solution of tosyl chloride. Such a reaction may proceed smoothly at low temperatures.

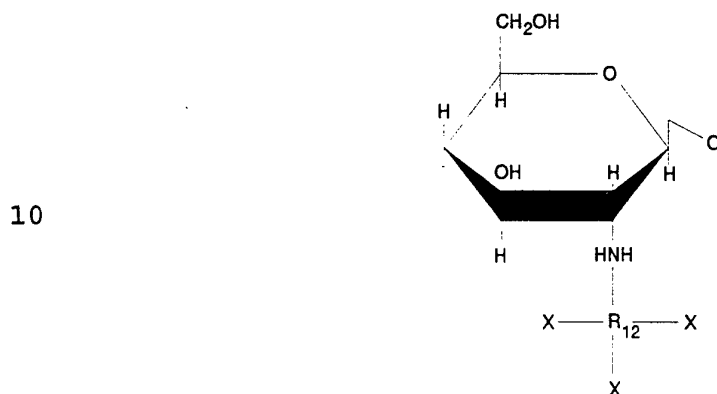
One or more of the monosaccharide units of the  
10 deacetylated derivative of the p-GlcNAC of the invention may form a salt, as depicted below:



where  $R_{11}$  may be an alkyl, an alkenyl, or an alkynyl  
20 moiety. Such a derivatization may be obtained by using techniques well known to those of skill in the art. For example, a procedure such as that described by Austin and Sennett (Austin, P.R. and Sennett, S., in "Chitin in Nature and Technology," 1986,  
25 Muzzarelli, R.A.A. *et al.*, eds. Plenum Press, pp. 279-286) may be utilized. Briefly, deacetylated p-GlcNAC may be suspended in an organic medium such as, for example, ethyl acetate or isopropanol, to which may be added an appropriate organic acid such as, for  
30 example, formic, acetic, glycolic, or lactic acid. The mixture may be allowed to stand for a period of time (1 to 3 hours, for example). The temperature of reaction and drying may vary from about 12° to about 35° C., with 20° to 25°C being preferred. The salts

may then be separated by filtration, washed with fresh medium, and the residual medium evaporated.

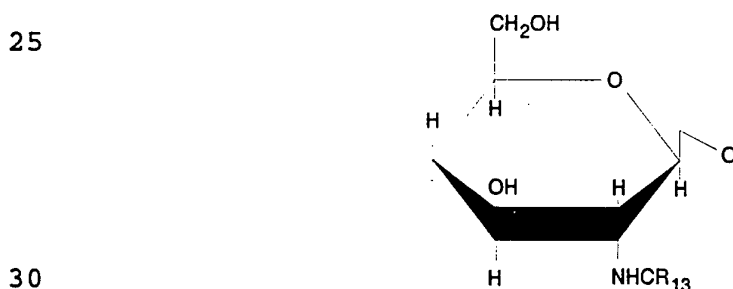
One or more of the monosaccharide units of the deacetylated derivative of the p-GlcNAc of the invention may form a metal chelate, as depicted below:



15 where  $R_{12}$  may be a metal ion, particularly one of the transition metals, and X is the dative bond established by the nitrogen electrons present in the amino and substituted amino groups present in the deacetylated p-GlcNAc.

20

One or more of the monosaccharide units of the deacetylated derivative of the p-GlcNAc of the invention may contain an N-alkylidene or an N-arylidene group, as depicted below:



where  $R_{13}$  may be an alkyl, an alkenyl, an alkynyl, or an aryl moiety. Such a derivatization may be obtained by using techniques well known to those of skill in the art. For example, a procedure such as that described by Hirano *et al.* (Hirano, S. *et al.*, 1981,

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J. Biomed. Mat. Res. 15:903-911) may be utilized. Briefly, an N-substitution reaction of deacetylated p-GlcNAc may be performed with carboxylic anhydrides and/or arylaldehydes to yield acyl- and/or arylidene  
5 derivatives.

Further, the p-GlcNAc of the invention, or its deacetylated derivative, may be subjected to controlled hydrolysis conditions, which yield groups of molecules having uniform, discrete molecular weight  
10 and other physical characteristics. Such hydrolysis conditions may include, for example, treatment with the enzyme, lysozyme. p-GlcNAc may be exposed to lysozyme for varying periods of time, in order to control the extent of hydrolysis. In addition, the  
15 rate of hydrolysis may be controlled as a function of the extent to which the p-GlcNAc that is being lysozyme treated has been deacetylated. Deacetylation conditions may be as described earlier in this Section. The more fully a p-GlcNAc molecule has been  
20 deacetylated, the more fully the molecule will be hydrolyzed. Changes in physical characteristics, in addition to the lowering of molecular weight, may be elicited by hydrolysis and/or deacetylation treatments. Extensive hydrolysis causes liquefaction  
25 of the p-GlcNAc. The results of a hydrolysis/deacetylation procedure are presented below in the Working Example of Section 9, below.

Further, heat denaturation may function to modify the crystalline structure of the p-GlcNAc. Such a  
30 modification of the p-GlcNAc product crystalline structure may advantageously affect, for example, the reactivity of the p-GlcNAc.

Further, a variety of molecules may be covalently or non-covalently functionally attached to the  
35 deacetylated derivatives of the p-GlcNAc of the

invention. Such molecules may include, but are not limited to such polypeptides as growth factors, such as nerve growth factor, proteases, such as pepsin, hormones, or peptide recognition sequences such as RGD sequences, fibronectin recognition sequences, laminin, integrins, cell adhesion molecules, and the like. Covalent attachment of molecules to the exposed primary amines of deacetylated p-GlcNAc may be accomplished by, for example, chemical attachment utilizing bi-functional cross-linking reagents that act as specific length chemical spacers. Such techniques are well known to those of skill in the art, and may resemble, for example, the methods of Davis and Preston (Davis, M. and Preston, J.F. 1981, Anal. Biochem. 116:404-407) and Staros et al. (Staros, J. V. et al., 1986, Anal. Biochem. 156:220-222). Briefly, carboxylic residues on the peptide to be attached to the deacetylated or partially deacetylated p-GlcNAc of the invention may be activated and then crosslinked to the p-GlcNAc. Activation may be accomplished, for example, by the addition of a solution such as carbodiimide EDC (1-ethyl-3-(3-dimethylaminopropyl)carbodiimide) to a peptide solution in a phosphate buffer. Preferably, this solution would additionally contain a reagent such as sulpho-NHS (N-hydroxysulphosuccinimide) to enhance coupling. The activated peptide may be crosslinked to the deacetylated p-GlcNAc by mixing in a high pH buffer, such as carbonate buffer (pH 9.0-9.2).

Alternatively, such molecules such as those described above may be non-covalently attached to deacetylated p-GlcNAc using techniques well known to those of skill in the art. For example, a molecule or molecules of choice may be mixed with a deacetylated p-GlcNAc solution prior to lyophilization.

Alternatively, hybrids comprising p-GlcNAc and/or p-GlcNAc derivatives may be formed. Such hybrids may contain any of a number of natural and/or synthetic materials, in addition to p-GlcNAc and/or p-GlcNAc derivatives. For example, hybrids may be formed of p-GlcNAc and/or p-GlcNAc derivatives plus one or more extracellular matrix (ECM) components. Such ECM components may include, but are not limited to, collagen, fibronectin, glycosaminoglycans, and/or peptidoglycans. Hybrids may also be formed of p-GlcNAc and/or p-GlcNAc derivatives plus one or more synthetic materials such as, for example, polyethylene. Such a p-GlcNAc/polyethylene or p-GlcNAc derivative/polyethylene hybrid may be made by thermally linking the hybrid components via, for example, autoclaving.

Additionally, an iodo-p-GlcNAc derivative may be copolymerized with, for example, styrene, for the manufacture of novel plastic materials. Iodo-p-GlcNAc can be prepared by a process similar to that described by Kurita and Inoue (Kurita, K. and Inoue, S., 1989, in "Chitin and Chitosan", Skjak-Braek et al., eds., Elsevier Science Publishing Co., Inc., p. 365), via tosylation and iodination of p-GlcNAc. The iodo derivative of p-GlcNAc can then be dispersed in nitrobenzene and reacted with styrene, with tin (IV) chloride being used as a catalyst.

In the case of a collagen/p-GlcNAc hybrid, briefly, a p-GlcNAc suspension and a collagen suspension may be mixed and lyophilized, and crosslinked, preferably dehydrothermally crosslinked. The collagen species of such hybrids may be native or synthetic, and may be of human or non-human, such as bovine, for example, origin. p-GlcNAc/collagen and/or p-GlcNAc derivative/collagen hybrid materials exhibit

uniform properties, and form a porous matrix that may act, for example, as an efficient three-dimensional matrix for the attachment and growth of cells. The Working Example presented in Section 13, below

5 demonstrates the formation, properties and usefulness of such a p-GlcNAc/collagen hybrid.

Hybrids comprising combinations of deacetylated p-GlcNAc and such compounds as, for example, heparin, sodium alginate, and carboxymethyl p-GlcNAc may be  
10 formulated using techniques such as those described herein. Such combinations may be formed or reformed into, for example, membranes and fibers.

Complexes of deacetylated p-GlcNAc with polyanions such as, for example, polyacrylic acid or  
15 pectin, possessing both positive and negative charges, may be formulated. The formation of such complexes may be accomplished according to a method similar to that described by Mireles et al. (Mireles, C. et al., 1992, in "Advances in Chitin and Chitosan", Brine, C.J. et al., eds., Elsevier Publishers, Ltd.).  
20 Deacetylated p-GlcNAc and polyacrylic acid, carrageenan or pectin, for example, are dissolved in HCl and NaCl, respectively, and the reactant solutions, with equal pH, are mixed. This operation  
25 produces effective flocculating molecules possessing both positive and negative characteristics, useful, for example, in the processing of waste waters.

#### 5.5 REFORMULATIONS

30 The p-GlcNAc of the invention, as well as its deacetylated derivatives and/or their derivatizations, such as those described, above, in Section 5.4, may be dissolved and subsequently reformulated into a variety of shapes and configurations.

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Solution of the p-GlcNAC of the invention can be achieved by treatment with dimethyl acetamide (DMA)/lithium chloride. p-GlcNAC may be readily dissolved by stirring in a DMA solution containing 5% LiCl (by weight of the DMA). Water soluble p-GlcNAC derivatives, such as p-GlcNAC salts, may be dissolved in water. P-GlcNAC which has been at least about 75% deacetylated may be put into solution in, for example, a mild acidic solution, such as 1% acetic acid.

10 p-GlcNAC derivatives that are water-insoluble may be put into solution in organic solvents.

Derivatization of p-GlcNAC in DMA:LiCl with phenyl isocyanates may be used to produce carbanilates. Further, derivatization of p-GlcNAC in DMA:LiCl with toluene-p-sulphonylchloride may be used to produce toluene-p-sulfonate.

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The p-GlcNAC of the invention, its deacetylated derivatives, and/or their derivatizations in solution may then be precipitated and reformulated into shapes which include, but are not limited to, mats, strings, ropes, microspheres, microbeads, membranes, fibers, powders, and sponges. Further, ultrathin (i.e., less than about 1 micron thick) uniform membranes may be formulated.

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Such reformulations may be achieved, by, for example, taking advantage of the fact that pure p-GlcNAC is insoluble in solutions such as water and alcohol, preferably ethanol. Introduction, by conventional means, such as by injection, for example, of the p-GlcNAC-containing DMA/LiCl mixture into such a water or alcohol, preferably ethanol, solution will bring about the reprecipitation, and therefore reformulation, of the dissolved p-GlcNAC. Such a pure p-GlcNAC reformulation is demonstrated in the Working Example presented, below, in Section 11. In the case

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of water soluble p-GlcNAc derivatives, reformulations may be achieved by reprecipitating in such organic solvents as, for example, ethyl acetate or isopropanol. Reformulations of p-GlcNAc which has  
5 been at least about 75% deacetylated may be achieved by reprecipitating in an alkaline solution. Water-insoluble p-GlcNAc derivatives may be reformulated by reprecipitation in aqueous solutions, such as, for example, water.

10 Deacetylated p-GlcNAc, in conjunction with oxidized cellulose, may be formulated to produce p-GlcNAc/cellulose hybrid materials improving the wet-strength of paper products. An oxidized cotton substrate can be approached closely by the  
15 deacetylated p-GlcNAc chain which has a flat ribbon-like shape, similar to that of cotton. Such proximity maximizes the contribution of the van der Waals forces to the forces promoting adsorption, thus enhancing the wet-strength properties of the hybrid p-GlcNAc-  
20 cellulose materials.

p-GlcNAc membranes and three dimensional p-GlcNAc matrices may be produced via methods which provide for the formation of controlled average pore sizes within either the membranes or the matrices. Pore size can  
25 be controlled in membranes and matrices by varying the amount of p-GlcNAc material used, and by the addition of certain solvents such as methanol or ethanol, with ethanol being preferred, in specific amounts, ranging from about 5% to about 40%, prior to the formation of  
30 membranes and/or matrices. In general, the greater the percentage of solvent, the smaller the average pore size formed will be. The Example presented, below, in Section 15, demonstrates the synthesis and characterization of such porous p-GlcNAc structures.



## 5.6 USES

The p-GlcNAc of the invention, as well as its deacetylated derivatives and their derivatizations, such as those described, above, in Section 5.4, and reformulations, such as those described above, in Section 5.5, have a variety of uses. For example, the non-toxic, non-pyrogenic, biodegradable, and biocompatible properties of the molecules of the invention, in addition to the advantageous properties of the p-GlcNAc and its derivatives, as described herein, lend themselves to applications in such diverse fields as agriculture, cosmetics, the biomedical industry, animal nutrition and health, and the food, chemical, photographic, and pharmaceutical industries.

### 5.6.1 BIOMEDICAL USES OF p-GlcNAc MATERIALS

#### 5.6.1.1 DRUG IMMOBILIZATION/DELIVERY USES

Biomedical uses of p-GlcNAc material may include, for example, enzyme and/or drug immobilization/delivery methods. For example, the p-GlcNAc of the invention or its derivatives, may have peptides of interest (growth factors, for example) covalently attached to them, as described, above, in Section 5.4. Peptide-containing p-GlcNAc may be administered to a patient using standard procedures well known to those of skill in the art, which include, but are not limited to injection, implantation, arthroscopic, laparoscopic or similar means. Upon introduction of the peptide-containing p-GlcNAc into a patient, the p-GlcNAc of the invention biodegrades, such that the attached peptides are gradually released into the bloodstream of the patient, thus providing a method for controlled drug delivery.

Deacetylated or partially deacetylated p-GlcNAc species may be produced having a predictable rate of biodegradability. For example, the percentage of deacetylation affects the rate at which the p-GlcNAc species degrades. Generally, the higher the percentage of deacetylation, the faster the rate of biodegradability and resorption will be. Thus, the degree of p-GlcNAc biodegradability and the in vivo rate of resorption may be controlled during the p-GlcNAc's production. Examples of the production and characterization of such p-GlcNAc materials are presented in Section 18, below. p-GlcNAc materials having such controllable biodegradability rates may be formulated into membranes, gels, sponges, microspheres, fibers, and the like. These p-GlcNAc products adhere and mold to tissues, both soft and hard tissues, in the human body with no need for suturing. The p-GlcNAc materials may, for example, be applied during general or minimally invasive surgery, such as laparoscopic surgery.

p-GlcNAc materials having a controllable rate of biodegradation may be useful, for example, to promote hemostasis in bleeding tissues, organs and blood vessels, to provide periodontal barriers for the separation of soft and hard tissue during the repair process following periodontal surgery, to provide surgical space fillers, to promote soft tissue augmentation, particularly in the skin for the purpose of reducing skin wrinkles, and as urinary sphincter augmentation, for the purpose of controlling incontinence. The Example presented in Section 19, below, demonstrates the use of such p-GlcNAc materials in one such application, namely, to promote hemostasis.

In addition, the molecules of the invention may serve as slow release drug delivery vehicles wherein the drug of interest has been encapsulated by the p-GlcNAc, or a derivative thereof. A drug/p-GlcNAc  
5 encapsulation may be produced, for example, by following a modification of the acid treatment/neutralization variation of the chemical/biological purification method presented, above, in Section 5.3.2. Rather than raising the pH  
10 of the p-GlcNAc solution to approximately neutral pH range (i.e., approximately 7.4), one may create a basic pH environment, by raising the pH to approximately 9.0 after the purification of the p-GlcNAc is completed. At a more basic pH, the  
15 structure of the p-GlcNAc of the invention, or a derivative thereof, assumes a more three dimensional or "open" configuration. As the pH is lowered, the molecule's configuration reverts to a more compact, "closed" configuration. Thus, a drug of interest may  
20 be added to a p-GlcNAc at a high pH, then the pH of the p-GlcNAc/drug suspension may be lowered, thereby "trapping" or encapsulating the drug of interest within a p-GlcNAc matrix.

Such p-GlcNAc encapsulations may be administered  
25 to a patient using standard techniques well known to those of skill in the art, so that, upon administration, the encapsulated drug is slowly released into the system of the patient as the p-GlcNAc of the encapsulation degrades.

30 p-GlcNAc-based gels and membranes have a variety of applications as therapeutic drug delivery systems. Such applications include, for example, anti-tumor drug delivery systems. The drug delivery systems described herein are feasible for use with any anti-  
35 tumor drug. Such drugs are well known to those of

skill in the art, and may be formulated into p-GlcNAc gels or membranes, for example, so as to provide site-specific slow-release delivery directly to the tumor or to the region vacated by the tumor following  
5 surgery. Such an immobilized slow-release p-GlcNAc drug product can act as an important initial defensive procedure after surgery. Such p-GlcNAc anti-tumor drug delivery systems are particularly useful in treating tumors which are totally or partially  
10 inaccessible through surgery, such as, for example, is the case with certain brain tumors.

Additional targets for p-GlcNAc anti-tumor systems include, but are not limited to, skin, GI tract, pancreatic, lung, breast, urinary tract and  
15 uterine tumors, and HIV-related Kaposi's sarcomas.

Antitumor drugs that are radiation enhancers are preferred for instances in which radiation therapy treatment is to be prescribed, either in lieu of, or following surgery. Examples of such drugs include,  
20 for example, 5'-fluorouracil, mitomycin, cis-platin and its derivatives, taxol, adriamycin, actinomycin, bleomycins, daunomycins, and methamycins.

Dose ranges for anti-tumor drugs may be lower than, equal to or greater than the typical daily doses  
25 prescribed for systemic treatment of patients. Higher doses may be tolerated in that the drugs are delivered locally at the site of a tumor. Other tissues, therefore, including blood cells, are not as readily exposed to the drugs. Doses of such drugs are well  
30 known to those of skill in the art, and may, alternatively, routinely be determined using standard techniques well known to those of skill in the art, such as, for example, are described, below, at the end of this Section.

The p-GlcNAc/drug delivery systems of the invention may, additionally, be used for the treatment of infections. For such an application, antibiotics, either water soluble or water insoluble, may be

5 immobilized/formulated in p-GlcNAc based materials, such as, for example, gels and membranes. Antibiotics are well known to those of skill in the art, and include, for example, penicillins, cephalosporins, tetracyclines, ampicillin, aureothicin, bacitracin,

10 chloramphenicol, cycloserine, erythromycin, gentamicin, gramacidins, kanamycins, neomycins, streptomycins, tobramycin, and vancomycin. Doses of such drugs are well known to those of skill in the art, and may, alternatively, routinely be determined

15 using standard techniques well known to those of skill in the art, such as, for example, are described, below, at the end of this Section.

Such p-GlcNAc antibiotic products may be used to treat bacterial infections that occur either

20 externally, e.g., on skin, scalp, dermal ulcers or eyes, or internally, e.g., localized infections of the brain, muscles, abdomen. A prominent application is for treatment of HIV-related opportunistic infections.

The p-GlcNAc/drug delivery systems of the invention may be formulated with anti-inflammatory drugs to control dysfunctional activity of the inflammatory and immune processes. For example, p-GlcNAc may be formulated with non-steroidal anti-inflammatory drugs (NSAIDs) and used to the reduction

25 of local pain and inflammation induced by diseases such as Rheumatoid arthritis, osteoarthritis and systemic lupus, to name a few. The localized delivery of such NSAIDs using the p-GlcNAc gel or membrane/drug delivery systems of the invention may serve to reduce

30 NSAID side effects, which may include gastric

irritation, azotemia, platelet disfunction and liver function abnormalities. NSAIDs are well known to those of skill in the art and include inhibitors of cyclooxygenase, such as aspirin, etodolac, fenoprofen and naproxen. Other anti-inflammatory drugs may be utilized as part of the p-GlcNAc/drug delivery systems of the invention, such as, for example, inhibitors of lipid inflammatory mediators, such as leucotrienes. Doses for such drugs are well known to those of skill in the art, and may, alternatively, routinely be determined using standard techniques well known to those of skill in the art, such as, for example, are described, below, at the end of this Section.

The p-GlcNAc/drug delivery systems of the invention may additionally be formulated with antifungal agents, using techniques described above, for the treatment of specific fungal diseases. Antifungal agents are well known to those of skill in the art, and may include, for example, amphotericin, anisomycin, antifungone, blastomycin, griseofulvins, and nystatin. Doses of such drugs are well known to those of skill in the art, and may, alternatively, routinely be determined using standard techniques well known to those of skill in the art, such as, for example, are described, below, at the end of this Section.

The p-GlcNAc/drug delivery systems of the invention may also be formulated with antiprotozoal agents, using techniques described above, for the treatment of specific protozoal infections. Antiprotozoal agents are well known to those of skill in the art, and may include, for example, antiamoebin, antiprotozin, monomycin, paromomycin and trichomycin. Doses of such drugs are well known to those of skill in the art, and may, alternatively, routinely be

determined using standard techniques well known to those of skill in the art, such as, for example, are described, below, at the end of this Section.

The p-GlcNAc drug delivery systems of the invention may be formulated with spermicidal compounds, using techniques such as those described, above, to produce effective contraceptives. Appropriate spermicides are well known to those of skill in the art. Doses of such spermicides are well known to those of skill in the art, and may, alternatively, routinely be determined using standard techniques well known to those of skill in the art, such as, for example, are described, below, at the end of this Section.

The p-GlcNAc drug delivery systems of the invention may, still further, be formulated using therapeutic protein agents. Such formulations may be produced using, for example, techniques such as those described above. By utilizing such p-GlcNAc therapeutic protein systems, it is possible to deliver specific proteins directly to desired target sites and to effect slow release of the proteins at such sites. Examples of possible proteins include, but are not limited to insulin, monoclonal antibodies, breast cancer immunotoxin, tumor necrosis factor, interferons, human growth hormone, lymphokines, colony stimulating factor, interleukins and human serum albumin. Doses of such therapeutic protein agents are well known to those of skill in the art, and may, alternatively, routinely be determined using standard techniques well known to those of skill in the art, such as, for example, are described, below, at the end of this Section.

Because the p-GlcNAc materials of the invention are themselves immunoneutral, in that they do not

elicit an immune response in humans, such p-GlcNAc devices, as described above, comprising p-GlcNAc membranes, 3D porous matrices and/or gels that harbor immobilized drugs, may deliver such drugs in a manner  
5 that there is no immune response. Certain additional materials, such as natural alginates and synthetic polymers, may be used in some cases to construct such devices in combination with the p-GlcNAc material.

The therapeutically effective doses of any of the  
10 drugs or agents described above, in conjunction with the p-GlcNAc-based systems described herein, may routinely be determined using techniques well known to those of skill in the art. A "therapeutically effective" dose refers to that amount of the compound  
15 sufficient to result in amelioration of symptoms of the processes and/or diseases described herein.

Toxicity and therapeutic efficacy of the drugs can be determined by standard pharmaceutical procedures in cell cultures or experimental animals,  
20 e.g., for determining the LD<sub>50</sub> (the dose lethal to 50% of the population) and the ED<sub>50</sub> (the dose therapeutically effective in 50% of the population). The dose ratio between toxic and therapeutic effects is the therapeutic index and it can be expressed as  
25 the ratio LD<sub>50</sub>/ED<sub>50</sub>. Compounds which exhibit large therapeutic indices are preferred. While compounds that exhibit toxic side effects may be used, care should be taken to design a delivery system that targets such compounds to the site of affected tissue  
30 in order to minimize potential damage to uninfected cells and, thereby, reduce side effects.

The data obtained from the cell culture assays and animal studies can be used in formulating a range of dosage for use in humans. The dosage of such  
35 compounds lies preferably within a range of



circulating concentrations that include the ED50 with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilized. For any  
5 compound used in the method of the invention, the therapeutically effective dose can be estimated initially from cell culture assays. A dose may be formulated in animal models to achieve a circulating plasma concentration range that includes the IC50  
10 (*i.e.*, the concentration of the test compound which achieves a half-maximal inhibition of symptoms) as determined in cell culture. Such information can be used to more accurately determine useful doses in humans. Levels in plasma may be measured, for  
15 example, by high performance liquid chromatography.

#### 5.6.1.2. p-GlcNAc CELL ENCAPSULATION USES

p-GlcNAc encapsulated cells may be formulated, and such p-GlcNAc encapsulated cells may be  
20 administered to a patient, via standard techniques well known to those of skill in the art. See, for example, the administration techniques described, above, in Section 5.6.1.1. Alternatively, see, for example, Aebisher et al. (Aebisher, P. et al., in  
25 "Fundamentals of Animal Cell Encapsulation and Immobilization", 1993, CRC Press, pp. 197-224), which is incorporated herein by reference in its entirety. Cells may be encapsulated by, on, or within p-GlcNAc or partially deacetylated p-GlcNAc membranes, three  
30 dimensional p-GlcNAc porous matrices, or p-GlcNAc gels.

Three dimensional matrices can be seeded with cells and used in certain applications without further encapsulation. Alternatively, cells can be  
35 encapsulated into microspheres or droplets of p-

GlcNAc-based polymer gels such as, for example, a p-GlcNAc-lactate polyelectrolyte polymer (a polycationic polymer). Gels, droplets or microspheres into which cells have been encapsulated may then be coated with a  
5 second polyelectrolyte of opposite charge (e.g., with a polyanion, such as an alginate) to form an outer capsule which provides immuno-isolation for the encapsulated cells, thus reducing the risk of immune rejection by the host organism.

10        Additionally, cells entrapped in p-GlcNAc gels, three dimensional p-GlcNAc matrices, or both, can be loaded into thermoplastic capsules in yet another method of formulation. Thermoplastic-based capsules can also be utilized to provide immuno-protection for  
15 implanted cells in a host organism. Such thermoplastic capsules are made of materials such as hydroxyethyl methylacrylate-methylmethacrylate copolymer (HEMA-MMA). Thermoplastic-derived microcapsules are formed, for example, by the  
20 coextrusion of a solution of HEMA-MMA in polyethylene glycol and the cell-containing p-GlcNAc matrix and/or gel medium, into an appropriate organic solvent such as hexadecane. See, for example, the method described by Aebisher et al. (Aebisher, P. et al., in  
25 "Fundamentals of Animal Cell Encapsulation and Immobilization", 1993, CRC Press, pp. 197-224).

The p-GlcNAc cell encapsulations have a variety of applications. First, they may be utilized for the delivery of therapeutic compounds, synthesized and  
30 secreted by cells attached to and encapsulated in the membranes, matrices or gels. For example and not by way of limitation, the p-GlcNAc/cell encapsulations may be used for delivery of insulin in the treatment of diabetes, nerve growth factor for the treatment of  
35 Alzheimer's disease, factor VIII and other clotting

factors for the treatment of hemophilia, dopamine for the treatment of Parkinson's disease, enkephalins via adrenal chromaffin cells for the treatment of chronic pain, dystrophin for the treatment of muscular  
5 dystrophy, and human growth hormone for the treatment of abnormal growth.

Further, because the p-GlcNAc materials of the invention are themselves immunoneutral, as they do not elicit an immune response in humans, it is possible to  
10 engineer and construct devices consisting of p-GlcNAc membranes, three-dimensional porous p-GlcNAc matrices and/or p-GlcNAc gels that harbor attached cells which can deliver cell-based therapeutics in a manner such that the cells are immuno-isolated, i.e., no anti-cell  
15 host immune response is elicited. Certain additional materials, such as, for example, natural alginates and synthetic polymers, may be used to construct such devices in addition to the p-GlcNAc material itself.

p-GlcNAc/cell encapsulation compositions may  
20 additionally be utilized for the delivery of cells to seed tissue regeneration. Applications of specific cell types encapsulated for the seeding of cell growth leading to tissue regeneration at the site of an injury may include, but are not limited to  
25 regeneration of skin, cartilage, nerves, bone, liver, and blood vessels. The tissue regeneration applications of cells encapsulated in p-GlcNAc materials are advantageous, in part, because of the ability of the p-GlcNAc material to adhere to injured  
30 tissue, to provide a substrate for mammalian cell growth, and to undergo bioresorption coincident with the growth of new healthy tissue during the tissue regeneration process at the site of injury. Examples include, but are not limited to the regeneration of

skin, bone, cartilage, liver, tendon, and ligament tissues.

5.6.1.3 UTILIZING p-GlcNAc MATERIALS FOR THE  
PREVENTION OF POST SURGICAL ADHESIONS

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Additionally, p-GlcNAc membranes may be used to provide a biodegradable, biocompatible mechanical barrier to prevent post-surgical adhesions. The Example presented in Section 17, below, demonstrate such a p-GlcNAc application. Solid p-GlcNAc or p-GlcNAc derivatives formulated into membranes or sponges may be utilized for such an application. Preferred membranes are thin, generally less than about 1 mm in thickness. Preferable p-GlcNAc derivatives are p-GlcNAc derivatives which have been about 50-80% deacetylated. Such p-GlcNAc derivatives will generally be resorbed approximately 7-21 days post implantation.

Liquid p-GlcNAc derivatives are also suitable for use in the prevention of post surgical adhesions. Preferable liquid p-GlcNAc derivatives for such an application are deacetylated p-GlcNAc salt derivatives and carboxymethyl p-GlcNAc derivatives. A p-GlcNAc derivative which is particularly preferred for the prevention of post surgical adhesions is a p-GlcNAc-lactate derivative, especially a p-GlcNAc-lactate gel derivative. Such p-GlcNAc-lactate derivatives may be formulated using propylene glycol and water, as, for example, described in Section 17.1. p-GlcNAc-lactate derivatives may be produced having high and low viscosities, which allows for the ability to tailor the p-GlcNAc used to the specific indication of interest. For example, it may be useful to use a p-GlcNAc product having a lower viscosity for delivery through a syringe or via a spray, while it may be desirable to use a p-GlcNAc product having a higher

viscosity, and therefore greater lubrication properties, when the indication is an orthopedic one.

For the prevention of post surgical adhesions, solid p-GlcNAc formulations are suitable for clearly circumscribed wound sites. Such p-GlcNAc formulations should be applied following the surgical procedure and the material should completely cover the traumatized tissue. It can be applied either in conjunction with either general or minimally invasive (e.g., laparoscopic) surgical procedures. The solid p-GlcNAc formulations can be cut and applied using standard surgical procedures and instrumentation well known to those of skill in the art.

The liquid p-GlcNAc formulations can be applied, for the prevention of post surgical adhesions, in larger areas prone to form such postoperative adhesions. The p-GlcNAc-lactate gel, for example, can be applied before the surgical procedure to provide additional lubrication and thus reduce the amount of traumatized tissue. Alternatively, the liquid p-GlcNAc formulation, such as p-GlcNAc-lactate, can be applied following the surgical procedure to form a physical barrier to prevent postoperative adhesion formation.

The p-GlcNAc material can be painted, sprayed or dropped from a syringe device onto the wounded site. In laparoscopic procedures, low viscosity materials can, for example, be delivered with standard suction irrigation devices. Higher viscosity materials will require pressure to reach its target. The pressure can be provided by a compressed gas powered piston or a syringe type device.

The amount of liquid p-GlcNAc formulation, such as the p-GlcNAc-lactate gel formulation, required for prevention of post surgical adhesions is proportional

to the extent of the traumatized tissue. The p-GlcNAc material administered should be applied in the range of 0.1 ml to 1.5 ml per sq. cm of surface area.

5     5.6.1.4 OTHER BIOMEDICAL USES OF p-GlcNAc MATERIALS

Other biomedical uses of p-GlcNAc materials include, for example, the use of such materials as cell culture substrates. For example, as shown in the Working Example presented in Section 12, below, the p-  
10 GlcNAc of the invention acts as a very efficient substrate for mammalian cells grown in culture. Further, three dimensional configurations of p-GlcNAc may be used as a medium components which will allow three dimensional cell culture growth.

15     The cell substrate capabilities of the p-GlcNAc of the invention may also be utilized in vivo. Here, the p-GlcNAc of the invention, or a derivative thereof, as described herein, may act to facilitate tissue regeneration (e.g., regeneration of connective  
20 tissue covering teeth near the gum line, vascular grafts, ligament, tendon, cartilage, bone, skin, nerve tissues). The p-GlcNAc molecules of the invention may, therefore, for example, have extensive plastic surgery applications.

25     Deacetylated p-GlcNAc is preferred for use as a sealant of vascular grafts. Deacetylated p-GlcNAc derivatives such as N-carboxymethyl and N-carboxybutyl deacetylated p-GlcNAc are preferred as tissue regeneration reagents. N-carboxymethyl deacetylated  
30 p-GlcNAc may, for example, be inoculated into the cornea to induce neovascularization.

Further biomedical applications of the p-GlcNAc of the invention or of its derivatives, as described herein, may involve the molecules' use in wound

dressings, wound healing ointments, and surgical sutures, sponges, and the like.

Still further, such molecules may be used, for example, in the treatment of osteoarthritis, in the  
5 reduction of blood serum cholesterol levels, as anti-viral agents, as anti-bacterial agents, as immunomodulators, as anticoagulants, as dialysis and ultrafiltration membranes, as anti-tumor agents, as contact lens material, and as oral adsorbents for  
10 iremic toxins when administered to kidney failure patients. Microcrystalline p-GlcNAC suspensions or water soluble p-GlcNAC derivatives are preferred for the treatment of arthritis, by, for example, injection directly into arthritic joints.

15 p-GlcNAC has additional applications as a component of artificial or donor skin. For example, p-GlcNAC, preferably as non-woven p-GlcNAC films, may be applied to split thickness skin donor sites, over, for example, donor dermis.

20 Deacetylated p-GlcNAC to which a protease, such as pepsin, has been attached may be used for the controlled digestion of proteins in contact with such p-GlcNAC/protease compounds.

Certain derivatizations of the p-GlcNAC of the  
25 invention, or of its derivatives, may be preferred for specific applications. (Derivatizations are described in Section 5.4, above.) For example, sulfated, phosphorylated, and/or nitrated p-GlcNAC derivatives may be preferred as anticoagulants or as lipoprotein  
30 lipase activators. N-acyl p-GlcNAC derivatives may also be preferred for anticoagulants, in addition to being preferred for, for example, use in production of artificial blood vessels, anti-viral compounds, anti-tumor (specifically, cancer cell aggregating  
35 compounds), dialysis and ultrafiltration membranes,

and in the production of controlled release drug delivery systems. O-alkyl p-GlcNAc and its deacetylated derivatives may also be preferred in the production of controlled release drug delivery systems. N-alkyl p-GlcNAc derivatives may be preferred as anti-bacterial agents. Oxido deaminated derivatives may be preferred as anti-cancer agents, specifically their use in conjunction with immunotherapy for cancer cells. Deacetylated p-GlcNAc derivatives may be preferred as wound healing agents. N-alkylidene and N-arylidene p-GlcNAc derivatives may be preferred for the enzyme immobilization applications.

15           5.6.2 AGRICULTURAL USES OF p-GlcNAc MATERIALS

The p-GlcNAc of the invention or its derivatives may be used in various agricultural applications, as well. Such applications include, but are not limited to insecticide, fungicide, bactericide, and nematocide applications. N-carboxymethyl deacetylated p-GlcNAc derivatives are preferred for use as effective bacteriostatic reagents. N-alkyl p-GlcNAc derivatives may be preferred for fungicide applications. Additionally, the molecules of the invention may be used in various soil treatment applications, including, but not limited to, fertilizer compositions. Further, controlled release of agrochemicals may be achieved by entrapping such chemicals via the immobilization, encapsulation, and other methods described, above, in this Section. Additionally, analogs of, for example, Rhizobium nodulation factors and/or nitrogen fixation inducers may be immobilized onto, and administered via, the p-GlcNAc and/or p-GlcNAc derivatives of the invention.

35



### 5.6.3 NUTRITION/FOOD INDUSTRY USES OF p-GlcNAc MATERIALS

The p-GlcNAc of the invention and its derivatives as described herein additionally have applications in the fields of animal and human nutrition. For  
5 example, the molecules of the invention may be used as feed ingredients. Techniques such as those described, above, in this Section, may be used in the production of controlled release products in animal systems.  
10 Additionally, the biomedical applications described above may be utilized in animal systems by incorporating routine modifications well known to those of ordinary skill in the art.

Food industry applications of the p-GlcNAc of the invention and of its derivatives, as described herein,  
15 may include, but are not limited to anticholesterol (i.e., hypocholesterolemic compounds), fat-binding compounds, emulsifiers, carriers, preservatives, seasonings, and food texturizers, in addition to fruit  
20 coatings, and food packaging products.

### 5.6.4 COSMETIC USES OF p-GlcNAc MATERIALS

Cosmetic applications of the p-GlcNAc of the invention may include, but are not limited to, the  
25 production of products for hair and skin care. Skin care products may include, for example, cosmetics utilizing deacetylated p-GlcNAc salts, carboxymethyl p-GlcNAc-containing products, and cosmetic packs containing deacetylated p-GlcNAc and such derivatives  
30 as hydroxypropyl-, N-succinyl-, and quaternary p-GlcNAc derivatives. Hair products may include, for example, carboxymethyl p-GlcNAc-containing products, and film-forming p-GlcNAc derivatives.

#### 5.6.5 CHEMICAL ENGINEERING APPLICATIONS OF p-GlcNAC MATERIALS

5 The p-GlcNAC of the invention and its derivatives have a variety of applications that are useful in the chemical engineering industry. For example, p-GlcNAC may be used as a coupling agent for adhesion of metals to polymers, membranes formed by glycol p-GlcNAC may be used in desalination applications, and membranes formed by other p-GlcNAC derivatives may be used for  
10 transport of halogen ions. Other applications may include the production of flame retardants, and the manufacture of metal chelating compounds and compounds capable of removing trace and heavy metals from liquids as well as water-soluble industrial  
15 pollutants, such as PCBs, for example. p-GlcNAC and/or p-GlcNAC derivatives may be used in photographic applications. For example, the ability of p-GlcNAC and/or p-GlcNAC derivatives to chelate metals, such as silver halides, may be utilized by  
20 contacting photographic solutions to recast mats, such as thin membranes, of p-GlcNAC and/or p-GlcNAC derivatives.

#### 25 6. EXAMPLE: PHYSICAL CHARACTERIZATION OF PURE PREPARATIONS OF p-GlcNAC

Presented in this Example, are circular dichroism (CD) and infra-red spectra (IR) analyses of p-GlcNAC and deacetylated p-GlcNAC membranes.

#### 30 6.1 MATERIALS AND METHODS

##### p-GlcNAC and commercial "chitin" preparations:

The p-GlcNAC used in the CD studies was prepared using the Mechanical Force purification method described, above, in Section 5.3.1.

35

Commercial "chitin" was purchased from NovaChem, Ltd., PO Box 1030 Armdale, Halifax, Nova Scotia, Canada, B3L 4K9.

The p-GlcNAC membranes used in the IR studies were prepared by either the Mechanical Force purification method as described, above, in Section 5.3.1, or by the Chemical/Biological purification method, as described, above, in Section 5.3.2, as indicated.

The commercial "p-GlcNAC" preparations were cast into membranes by dissolving in a dimethylacetamide solution containing 5% lithium chloride, and layering onto distilled, deionized water until membranes precipitated.

p-GlcNAC derivatives and treatments: The Deacetylated p-GlcNAC used in both the CD and IR studies was prepared by treatment of the p-GlcNAC with 50% NaOH at 60° C. for 2 hours. The heat-denatured p-GlcNAC membranes used in the IR studies were modified by boiling in 0.2mM EDTA for 3 minutes. Autoclaved p-GlcNAC was autoclaved or 30 minutes at 122° C.

CD techniques: Solid state CD techniques were carried out essentially according to Domard (Domard, A., 1986, Int. J. Macromol. 8:243-246).

25

## 6.2 RESULTS

### 6.2.1 CD ANALYSIS

In the CD spectra obtained from untreated p-GlcNAC (FIG. 3A), the expected  $n-\pi^*$  and  $\pi-\pi^{**}$  optically active electronic transitions (220-185nm) were observed due to the presence of the carbonyl group in the acetyl moiety of p-GlcNAC are present. Such peaks are completely absent in the CD spectrum obtained from the deacetylated p-GlcNAC product, as shown in FIG.

35 3B.

### 6.2.2 IR SPECTRA ANALYSIS

The IR spectra obtained in this study are consistent with the chemical structure of p-GlcNAc. Additionally, the sharp definition of each IR peak is indicative of the presence of an ordered and regular (i.e., pseudocrystalline) structure in the p-GlcNAc fibers. See FIG. 4A for the IR spectrum of p-GlcNAc purified via the Mechanical Force purification method, and FIG. 4D for the IR spectrum of p-GlcNAc purified via the Chemical/Biological method. For comparison, see FIG. 4B, which demonstrates the IR spectrum of a commercial "chitin" preparation.

The IR spectrum obtained from the autoclaved p-GlcNAc material (FIG. 4E) does not differ visibly from the IR spectrum observed in FIG. 4A. This data indicates that the p-GlcNAc material may be sterilized by autoclaving with no loss of polymer structure.

## 7. EXAMPLE: PURIFICATION OF p-GlcNAc USING THE MECHANICAL FORCE PURIFICATION METHOD

In this section, p-GlcNAc was purified using the Mechanical Force technique described above, in Section 5.3.1.

### 7.1 MATERIALS AND METHODS/RESULTS

Diatom culture conditions: The diatom species Thalassiosira fluviatilis was grown in culture according the procedures described, above, in Sections 5.1 and 5.2.

SEM procedures: The SEM techniques used here are as those described, below, in Section 12.1.

p-GlcNAc Purification procedure: p-GlcNAc was purified from the diatom culture by utilizing the Mechanical Force technique described above, in Section 5.3.1. Specifically, the p-GlcNAc fibers were separated from the diatom cell bodies by subjecting

the contents of the culture to three short bursts of top speed mixing motion in a Waring blender. Total time of the three bursts was about one second. The resulting suspension was centrifuged at 3500 rpm in a Sorvall GS-4 fixed angle rotor, for 20 minutes at about 10°C. The supernatant was decanted, and centrifuged again, this time at, 4000 rpm in a Sorvall GS-4 fixed angle rotor for 20 minutes at about 10°C. Once again, the supernatant was decanted and centrifuged at 4000 rpm at 10° C. The final supernatant of the third centrifugation was clear, with little, if any, visible flocs floating in the liquid. The clear supernatant was decanted into a Buchner filtration unit equipped with nitrocellulose with 0.8µm pore size, suction was then applied and the liquid was filtered from the fiber suspension, allowing the fibers to be collected onto the membrane. The collected fibers were washed with 1 liter of distilled, deionized H<sub>2</sub>O at 70° C. When almost all of the water had been drained, fibers were washed, with suction, with 1 liter of 1 N HCl at 70°C. When most of the acid solution had been drained, the fibers were washed with 1 liter of distilled, deionized H<sub>2</sub>O at 70°C, using suction. When most of the wash water had been drained, the fibers were washed with 1 liter of 95% ethanol at room temperature, and vacuum was applied. The filter membrane on which the white fiber membrane had been collected was then removed from the filtration unit and the membrane and its membrane support was dried in a drying oven at 58°C for 20 minutes, after which the membrane and its support was placed in a desiccator for 16 hours.

Following this purification procedure, the yield of p-GlcNAc from a 1000 ml culture was 6.85 milligrams per liter of diatom culture. SEM photographs of the

membrane formed by the collection of the p-GlcNAC fibers via this technique is shown in FIG. 6A-6B.

5 8. EXAMPLE: PURIFICATION OF p-GlcNAC USING THE  
BIOLOGICAL/CHEMICAL PURIFICATION  
METHOD

10 In this section, p-GlcNAC was purified using two of the Chemical/Biological techniques described above, in Section 5.3.2. Briefly, p-GlcNAC was purified via HF treatment, in one case, and via acid treatment/neutralization in the second case.

8.1 MATERIALS AND METHODS/RESULTS

15 Diatom culture conditions: The diatom species Thalassiosira fluviatilis was grown in culture according to the procedures described, above, in Sections 5.1 and 5.2.

SEM procedures: The techniques utilized in this study were as described, below, in Section 12.1.

20 Purification procedure: First, p-GlcNAC was purified by HF treatment, the results of which are shown in FIG. 7A-7B. Specifically, under a fume hood, 2.42 ml of a 49% (29N) HF solution was added to the diatom contents of the culture, at room temperature, 25 for each 1000 ml of the volume of the original cell culture, resulting in a 0.07 M HF solution. The mixture was then shaken vigorously for about 30 seconds, causing persistent foam to appear over the liquid. The container was allowed to stand 30 undisturbed for 5-6 hours to allow heavy particulates to settle. At the end of this time, a layer of foam had formed, while the liquid itself was divided into two strata: first, a narrow, very dark green layer resting on the bottom of the container below a second, 35 much lighter colored grayish-green and murky phase which represented perhaps 85-90% of the total volume

of liquid. The foam layer was carefully siphoned off, using a capillary glass tube and vacuum suction. The grayish cloudy supernatant was then siphoned off, with care being taken to not disturb the dark bottom layer, which consisted mainly of settled cell bodies, and was transferred to a separate plastic container. The grayish cloudy supernatant was allowed to stand undisturbed for an additional 16 hours. The liquid was initially almost colorless, light grey, but not transparent. After 16 hours settling time, a small amount of foam remained on top of the main body of liquid and a small amount of green matter had settled on the bottom of the container. The liquid was lighter in color, but still not transparent. The foam on top of the liquid was siphoned off as before. The main body of liquid was then carefully siphoned off, leaving behind the small amount of settled green material at the bottom of the container. The liquid which had thus been isolated, contained the majority of the p-GlcNAc fibers and some impurities.

To remove proteins and other unwanted matter liberated by the diatoms during the preceding steps in the procedure from the fiber-containing liquid, the suspension of fibers and cell remnants was washed with sodium dodecyl sulfate (SDS). Specifically, the necessary volume of a 20% SDS solution was added to make the final concentration of the liquid 0.5% SDS by volume. The container holding the liquid was sealed, secured in a horizontal position on a shaking machine, and agitated for 24 hours at about 100 shakes a minute. Soon after shaking began, large flocs of white p-GlcNAc fibers appeared in the suspension, and a considerable amount of foam accumulated in the head space of the containers. At the end of the SDS washing, the contents of the containers were

transferred to Buchner filtration equipment equipped with a 0.8  $\mu\text{m}$  (Supor Filter, Gelman) filter membrane. The liquid was filtered with suction, and the p-GlcNAc fibers in the liquid were collected on the filter  
5 membrane.

The p-GlcNAc fibers collected on the filter membrane were then washed further. First, the fibers were washed with hot (70° C.) distilled, deionized H<sub>2</sub>O, using three times the volume of the original  
10 suspension. With a water jet using distilled, deionized H<sub>2</sub>O, the white fiber clumps collected on the filter membrane of the Buchner filter were transferred to a Waring blender, and the fiber clumps were disintegrated with about 10 short mixing bursts. The  
15 suspension of disintegrated fibers was transferred to a Buchner filter funnel equipped with a nitrocellulose filter membrane as described above, and the liquid was removed under suction. The collected fibers were washed with 1000 ml of hot (70°C) 1N HCl solution, and  
20 subsequently further washed with 1000 ml hot (70°C) distilled, deionized H<sub>2</sub>O. Finally, the fibers were washed with 1000 ml 95% ethanol at room temperature, and filtered to dryness. The fiber membrane and the filter membrane supporting the fiber membrane were  
25 then dried in a drying oven at 58°C for 20 minutes. The membrane and membrane support was then placed in a desiccator for 16 hours. The membrane was then carefully detached from the filter membrane.

Second, p-GlcNAc was purified by using the acid  
30 treatment/neutralization method described, above, in Section 5.3.2. Specifically, the p-GlcNAc was processed as described earlier in this Section, until prior to the SDS wash step, at which point the solution was neutralized to a pH of approximately 7.0  
35 by the addition of a 2.9M Tris solution. The p-GlcNAc



yield from this purification procedure was 20.20 milligrams per liter of diatom culture. On average, approximately 60 milligrams per liter diatom culture are obtained. SEM micrographs of membranes formed during the purification procedure are shown in FIGS. 8A-8B and 9A-9E.

9. EXAMPLE: p-GlcNAc DEACETYLATION

A p-GlcNAc membrane was suspended in a solution containing 50% NaOH. The suspension was heated at 80°C for 2 hours. The resulting deacetylated membrane was dried and studied by scanning electron microscopy, as shown in FIG. 11A-11B.

10. EXAMPLE: p-GlcNAc BIOCOMPATIBILITY

In this Example, it is demonstrated that the p-GlcNAc of the invention exhibits no detectable biological reactivity, as assayed by elution tests, intramuscular implantation in rabbits, intracutaneous injection in rabbits, and systemic injections in mice.

10.1. MATERIALS AND METHODS

10.1.1. ELUTION TEST

Conditions for the elution test conformed to the specifications set forth in the U.S. Pharmacopeia XXII, 1990, pp. 1415-1497 and to U.S. Pharmacopeia XXII, Supplement 5, 1991, pp. 2702-2703.

Cell culture: Mouse fibroblast L929 cell line (American Type Culture Collection Rockville, MD; ATCC No. CCL1; NCTC clone 929) was utilized. A 24 hour confluent monolayer of L929 cells was propagated in complete Minimum Essential Medium (MEM).

p-GlcNAc: a solid membrane of p-GlcNAc which had been prepared according to the Mechanical Force method of purification described, above, in Section 5.3.1,

was extracted in 20 ml serum-supplemented MEM as per U.S. Pharmacopeia XXII (1990) requirements.

Controls: Natural rubber was used as a positive control, and silicone was used as a negative control.

5 Controls were tested in the same manner as the test article, p-GlcNAc.

Extracts: Extracts were prepared at 37°C, in a humidified atmosphere containing 5% carbon dioxide, for 24 hours. Extracts were evaluated for a change in  
 10 pH, and adjustments were made to bring the pH to within +/- 0.2 pH units of the original medium. Adjustments were made with HCl lower extract pH on with NaHCO<sub>3</sub> to raise the extract pH. Extracts were sterile filtered by passage through a 0.22 micron  
 15 filter, prior to being applied to the cell monolayer.

Dosing: 3 mls of p-GlcNAc or control extracts were used to replace the maintenance medium of cell cultures. All extracts were tested in duplicate.

Evaluation Criteria: Response of the cell  
 20 monolayer was evaluated either visually or under a microscope. The biological reactivity, i.e., cellular degeneration and/or malformation, was rated on a scale of 0 to 4, as shown below. The test system is suitable if no signs of cellular reactivity (Grade 0)  
 25 are noted for the negative control article, and the positive control article shows a greater than mild reactivity (Grade 2). The test article (i.e., p-GlcNAc) meets the biocompatibility test if none of the cultures treated with the test article show a greater  
 30 than mild reactivity.

Grade	Reactivity	Description of Reactivity Zone
0	None	Discrete intracytoplasmic granules; No cell Lysis

35

1	Slightly	Not more than 20% of the cells are round, loosely attached, and without intra-cytoplasmic granules; occasional lysed cells are present
5	2 Mild	Not more than 50% of the cells are round and devoid of intracytoplasmic granules; extensive cell lysis and empty areas between cells
10	3 Moderate	Not more than 70% of the cell layers contain rounded cells and/or are lysed
	4 Severe	Nearly complete destruction of the cell layers

15                    10.1.2. INTRAMUSCULAR IMPLANTATIONS

20                    Animals: Healthy, New Zealand White Rabbits, male and female, (Eastern Rabbit Breeding Laboratory, Taunton, MA) were used. Rabbits were individually housed using suspended stainless steel cages. Upon receipt, animals were placed in quarantine for 8 days, under the same conditions, as for the actual test. Hardwood chips (Sani-chips™, J.P. Murphy Forest Products, Montvale, NJ) were used as non-contact bedding under cages. The animal facility was maintained at a temperature of 68° +/- 3°F, with a relative humidity at 30-70%, a minimum of 10-13 complete air exchanges per hour, and a 12-hour light/dark cycle using full spectrum fluorescent lights. Animals were supplied with commercial feed (Agway ProLab, Waverly, NY) under controlled conditions and municipal tap water ad libitum. No known contaminants were present in the feed, bedding, or water which would be expected to interfere with the test results. Animals selected for the study were chosen from a larger pool of animals. Rabbits were

35

- 70 -

weighted to nearest 10g and individually identified by ear tattoo.

p-GlcNAc: The p-GlcNAc used was as described, above, in Section 10.1.1.

5        Implantation Test: Two rabbits were used for each implantation test. On the day of the test, the animal skin on both sides of the spinal column was clipped free of fur. Each animal was anesthetized to prevent muscular movement. Using sterile hypodermic needles  
10 and stylets, four strips of the test p-GlcNAc (1mm x 1mm x 10mm) were implanted into the paravertebral muscle on one side of the spine of each of two rabbits (2.5 to 5cm from the midline, parallel to the spinal column, and about 2.5 cm from each other). In a  
15 similar fashion, two strips of the USP negative control plastic RS (1mm x 1mm x 10mm) were implanted in the opposite muscle of each animal. Animals were maintained for a period of 7 days. At the end of the observation period, the animals were weighed and  
20 euthanized by an injectable barbituate, Euthanasia-5 (Veterinary Laboratories, Inc., Lenexa, KS). Sufficient time was allowed to elapse for the tissue to be cut without bleeding. The area of the tissue surrounding the center portion of each implant strip  
25 was examined macroscopically using a magnifying lens. Hemorrhaging, necrosis, discolorations and infections were scored using the following scale: 0=Normal, 1=Mild, 2=Moderate, and 3=Severe. Encapsulation, if present, was scored by first measuring the width of  
30 the capsule (i.e., the distance from the periphery of the implant to the periphery of the capsule) rounded to the nearest 0.1mm. The encapsulation was scored as follows:

35

	Capsule Width	Score
	None	0
	up to 0.5 mm	1
5	0.6 - 1.0 mm	2
	1.1 - 2.0 mm	3
	Greater than 2.0 mm	4

10 The differences between the average scores for  
the p-GlcNAC and the positive control article were  
calculated. The test is considered negative if, in  
each rabbit, the difference between the average scores  
for each category of biological reaction for the p-  
15 GlcNAC and the positive control plastic implant sites  
does not exceed 1.0; or, if the difference between the  
mean scores for all categories of biological reaction  
for each p-GlcNAC article and the average score for  
all categories for all the positive control plastic  
20 implant sites does not exceed 1.0, for not more than  
one of four p-GlcNAC strips.

#### 10.1.3. INTRACUTANEOUS INJECTIONS

Animals: New Zealand white rabbits were used and  
maintained as described, above, in Section 10.1.2.

25 p-GlcNAC: A solid membrane of p-GlcNAC which had  
been prepared according to the mechanical force method  
of purification described, above, in Section 5.3.1,  
was placed in an extraction flask, to which 20 ml of  
the appropriate medium were added. Extractions were  
30 performed by heating to 70° for 24 hours. Following  
this procedure, extracts were cooled to room  
temperature. Each extraction bottle was shaken  
vigorously prior to administration.

Intracutaneous Test: On the day of the test,  
35 animals were clipped free of fur on the dorsal side.  
A volume of 0.2 ml of each p-GlcNAC extract was

injected intracutaneously at five sites on one side of each of two rabbits. More than one p-GlcNAc extract was used per rabbit. At five sites on the other side of each rabbit, 0.2 ml of the corresponding control  
 5 was injected. Injection sites were observed for signs of erythema, edema, and necrosis at 24, 48, and 72 hours after injection. Observations were scored according to the Draize Scale for the Scoring Skin Reaction (USP Pharmacopeia XXII, 1990, 1497-1500; USP  
 10 Pharmacopeia XXII, Supplement 5, 1991, 2703-2705) as shown in Table II, below:

**TABLE II**  
 Draize Scale for Scoring Skin Reactions

15	<u>Value</u>
<u>Erythema and Eschar Formation</u>	
No erythema . . . . .	0
Very slight erythema (barely perceptible) . . . . .	1
20 Well defined erythema . . . . .	2
Moderate to severe erythema . . . . .	3
Severe erythema (beet redness) to slight eschar formation (injuries in depth) . . . . .	4
Total possible erythema score = 4	
25	
<u>Edema Formation</u>	
No edema . . . . .	0
Very slight edema (barely perceptible) . . . . .	1
30 Slight edema (edges are well defined by definite raising) . . . . .	2
Moderate edema (raised approximately 1mm and extending beyond area of exposure) . . . . .	3
Severe edema (raised more than 1mm and extending beyond area of exposure) . . . . .	4
Total possible edema score = 4	
35	

All erythema and edema scores at 24, 48, and 72 hours were totaled separately and divided by 12 (i.e., 2 animals x 3 scoring periods x 2 scoring categories) to determine the overall mean score for the p-GlcNAc  
5 versus the corresponding control. Animals were weighed at the end of the observation period and euthanized by injection of a barbituate, Euthanasia-5 (Veterinary Laboratories, Inc., Lenexa, KS). The results of the test are met if the difference between  
10 the p-GlcNAc and the control means reaction scores (erythema/edema) is 1.0 or less).

#### 10.1.4. SYSTEMIC INJECTIONS

Animals: Albino Swiss mice (Mus musculus),  
15 female, (Charles River Breeding Laboratories, Wilmington, MA) were used. Groups of 5 mice were housed in polypropylene cages fitted with stainless steel lids. Hardwood chips (Sani-chips™, J.P. Murphy Forest Products, Montvale, NJ) were used as contact  
20 bedding in the cages. The animal facility was maintained as a limited access area. The animal rooms were kept at a temperature of 68 +/- 3°F, with a relative humidity of 30-70%, a minimum of 10-13 complete air exchanges per hour, and a 12 hour  
25 light/dark cycle using full spectrum fluorescent lights. Mice were supplied with commercial feed and municipal tap water ad libitum. There were no known contaminants present in the feed, bedding, or water which would be expected to interfere with the test  
30 results. Animals selected for the study were chosen from a larger pool of animals. Mice were weighed to the nearest 0.1g and individually identified by ear punch.

p-GlcNAc: The samples used were as described,  
35 above, in Section 10.1.1. Extracts were prepared

according to the procedures described in Section 10.1.3, above.

Systemic Injection Test: Groups of 5 mice were injected with either p-GlcNAc extract or a corresponding control article, in the same amounts and by the same routes as set forth below:

	Test Article or Control Article Extracts	Dosing Route	Dose/Kg	Injection Rate
10	0.9% Sodium Chloride Injection, USP (0.9% NaCl)	Intravenous	50 ml	0.1 ml/sec
15	1 in 20 Alcohol in 0.9% Sodium Chloride Injection USP (EtOH:NaCl)	Intravenous	50 ml	0.1 ml/sec
20	Polyethylene Glycol 400 (PEG 400)	Intraperitoneal	10 g	— —
	Cottonseed Oil (CSO)	Intraperitoneal	50 ml	— —

Extracts of the p-GlcNAc prepared with PEG 400, and the corresponding control, were diluted with 0.9% NaCl, to obtain 200 mg of PEG 400 per ml. For the Intracutaneous Test, PEG 400 was diluted with 0.9% NaCl to obtain 120 mg of PEG 400 per ml.

The animals were observed immediately after injection, at 24 hours, 48 hours, and 72 hours after injection. Animals were weighed at the end of the observation period and euthanized by exposure to carbon dioxide gas. The requirements of the test are met if none of the animals treated with the p-GlcNAc shows a significantly greater biological reactivity than the animals treated with the control article.



10.2 RESULTS10.2.1. ELUTION TEST

The response of the cell monolayer to the p-GlcNAc test article was evaluated visually and under a microscope. No cytochemical stains were used in the evaluation. No signs of cellular biological reactivity (Grade 0) were observed by 48 hours post-exposure to the negative control article or to the p-GlcNAc. Severe reactivity (Grade 4) was noted for the positive control article, as shown below in Table III:

TABLE III

Time	REACTIVITY GRADES					
	p-GlcNAc		Control Articles			
	A	B	A	B	A	B
0 Hours	0	0	0	0	0	0
24 Hours	0	0	0	0	4	4
48 Hours	0	0	0	0	4	4

The p-GlcNAc of the invention, therefore, passes requirements of the elution test for biocompatibility, and, thus, is non-cytotoxic.

10.2.2 INTRAMUSCULAR IMPLANTATIONS

Both rabbits (A and B) tested increased in body weight and exhibited no signs of toxicity. See Table IV for data. In addition, there were no overt signs of toxicity noted in either animal. Macroscopic evaluation of the test and control article implant sites showed no inflammation, encapsulation, hemorrhage, necrosis, or discoloration. See Table IV for results. The test, therefore, demonstrates that

the p-GlcNAc assayed exhibits no biological reactivities, in that, in each rabbit, the difference between the average scores for all of the categories of biological reaction for all of the p-GlcNAc implant sites and the average score for all categories for all the control implant sites did not exceed 1.0.

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TABLE IV

IMPLANTATION TEST  
 (Macroscopic Observations)  
 Test Article: p-GlcNAC  
 Animal Specie: Rabbit  
 Animal #: A

TISSUE SITE:	TEST					CONTROL AVERAGE
	T1	T2	T3	T4	AVERAGE	
Inflammation	0	0	0	0	0	0
Encapsulation	0	0	0	0	0	0
Hemorrhage	0	0	0	0	0	0
Necrosis	0	0	0	0	0	0
Discoloration	0	0	0	0	0	0
Total	0	0	0	0	0	0
MEAN SCORE: (total/5)	0	0	0	0	0	0

AVERAGE CONTROL VALUE: 0

TABLE IV (cont.)

Animal #: B

TISSUE SITE:	TEST AVERAGE					CONTROL AVERAGE	
	T1	T2	T3	T4	C1	C2	
Inflammation	0	0	0	0	0	0	0
Encapsulation	0	0	0	0	0	0	0
Hemorrhage	0	0	0	0	0	0	0
Necrosis	0	0	0	0	0	0	0
Discoloration	0	0	0	0	0	0	0

Total

0

0

0

0

0

0

0

MEAN SCORE:  
(total/5)

0

0

0

0

0

0

0

AVERAGE CONTROL VALUE: 0

10.2.3. INTRACUTANEOUS TEST

All of the animals increased in weight. See Table V for data. There were no signs of erythema or edema observed at any of the p-GlcNAc or control  
5 article sites. Overt signs of toxicity were not observed in any animal. Because the difference between the p-GlcNAc and control article mean reaction scores (erythema/edema) was less than 1.0, the p-GlcNAc meets the requirements of the intracutaneous  
10 test. See Table VI for results. Therefore, as assayed by this test, the p-GlcNAc demonstrates no biological reactivity.

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TABLE V

Intracutaneous and Implant Tests  
 Body Weights and Clinical Observations  
 Test Article: p-GlcNAc Animal Species: Rabbit

Group	Animal #	Sex	Body Weight (Kg)		Weight Change	Signs of Toxicity*
			Day 0	Day 3		
0.9% NaCl & CSO	23113	Male	2.51	2.55	0.04	None
0.9% NaCl & CSO	23114	Female	2.43	2.46	0.03	None
EtoH: NaCl & PEG 400	23115	Male	2.47	2.50	0.03	None
EtoH: NaCl & PEG 400	23116	Female	2.59	2.63	0.04	None
			Day 0	Day 7		
Implant A		Male	2.74	2.80	0.06	None
Implant B		Female	2.66	2.74	0.08	None

\*Summary of Observations Day 0 Through Day 7 (Implant) and Day 0 through Day 3 (Intracutaneous).







TABLE VI (Cont.)

NaCl/EtOH Extract

Animal ID #	Vehicle	SITE NUMBERS					SCORING (ER/ED)					Averages						
		T-1	C-1	T-2	C-2	T-3	C-3	T-4	C-4	T-5	C-5	Time:	T	C				
23115	NaCl	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	24hr.	0/0	0/0
	EtOH	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	48hr.	0/0	0/0
	Total	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	72hr.	0/0	0/0
23116	NaCl	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	24hr.	0/0	0/0
	EtOH	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	48hr.	0/0	0/0
	Total	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	72hr.	0/0	0/0

TABLE VI (Cont.)

PEG Extract

Animal ID #	Vehicle	SITE NUMBERS						SCORING (ER/ED)					Averages					
		T-1	C-1	T-2	C-2	T-3	C-3	T-4	C-4	T-5	C-5	Time:	T	C				
23115	PEG	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	24hr.	0/0	0/0
		0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	48hr.	0/0	0/0
		0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	72hr.	0/0	0/0
	Total	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0			0/0	0/0
23116	PEG	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	24hr.	0/0	0/0
		0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	48hr.	0/0	0/0
		0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	72hr.	0/0	0/0
	Total	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0	0/0			0/0	0/0

10.2.4. SYSTEMIC TEST

All of the mice treated with the p-GlcNAc extract or the control article increased in weight. See Table VII for data. In addition, there were no overt signs of toxicity observed in any p-GlcNAc or control animal. See Table VI for results. It is concluded, therefore, that none of the p-GlcNAc test animals showed a significantly greater biological reactivity than the animals treated with the control article.

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TABLE VII

## ANIMAL WEIGHTS AND CLINICAL OBSERVATIONS

5	Group	Sex	Dose (ml)	Animal #	Body Weight (g)		Weight Change	Signs of Toxicity *
					Day 0	Day 3		
10	NaCl:	Female	1.03	I.	20.6	22.8	2.2	None
	EtOH	Female	1.06	II.	21.1	23.4	2.3	None
	Test	Female	1.02	III.	20.4	22.6	2.2	None
	50ml/kg	Female	1.11	IV.	22.2	24.5	2.3	None
		Female	1.05	V.	21.0	23.2	2.2	None
				Mean		21.1	23.3	
			SD +/-		0.7	0.7		
15	NaCl:	Female	1.04	VI.	20.7	23.2	2.5	None
	EtOH	Female	1.04	VII.	20.8	23.5	2.7	None
	Control	Female	1.02	VIII.	20.3	22.3	2.0	None
	50ml/kg	Female	0.91	IX.	18.2	20.6	2.4	None
		Female	0.94	X.	18.7	20.9	2.2	None
				Mean		19.7	22.1	
			SD +/-		1.2	1.3		
20	PEG	Female	1.02	XI.	20.3	22.7	2.4	None
	Test	Female	0.96	XII.	19.2	21.4	2.2	None
	10ml/kg	Female	0.95	XIII.	18.9	21.6	2.7	None
		Female	1.05	XIV.	20.9	22.7	1.8	None
		Female	0.94	XV.	18.7	21.2	2.5	None
				Mean		19.6	21.9	
			SD +/-		1.0	0.7		
25	PEG	Female	1.01	XVI.	20.1	22.3	2.2	None
	Control	Female	0.99	XVII.	19.8	22.0	2.2	None
	10g/kg	Female	1.10	XVIII.	22.0	24.3	2.3	None
		Female	1.07	XIX.	21.4	23.6	2.2	None
		Female	1.03	XX.	20.6	22.4	1.8	None
				Mean		20.8	22.9	
			SD +/-		0.9	1.0		
30								

\*Summary of observations - 0, 4, 24, 48, and 72 h after injection

11. EXAMPLE: p-GlcNAC REFORMULATION

In the Working Example presented in this Section, a p-GlcNAC membrane (16.2 mg) was dissolved in 1 ml of a dimethylacetamide solution containing 5% LiCl. The p-GlcNAC-containing solution was placed in a syringe and extruded into 50 ml of pure water to precipitate a fiber. The resulting fiber was studied with scanning electron microscopy, as shown in FIG. 10A-10B.

12. EXAMPLE: CELL ATTACHMENT TO p-GlcNAC

In this working example, it is demonstrated that p-GlcNAC represents an efficient substrate for cell attachment and growth in culture.

12.1 MATERIALS AND METHODS

Cells: The transformed mouse 3T3 fibroblast cell line was used, and was grown in DMEM supplemented with 10 fetal bovine serum (FBS).

p-GlcNAC membranes: p-GlcNAC was prepared according to the methods described, above, in Sections 5.3.1 and 8.

p-GlcNAC membranes were initially stuck to a #1 (18mm) Corning cover glass using one drop of water, and were attached by autoclaving at 121° C. for 30 minutes. Membranes prepared in this manner were then placed in culture wells of 6 well culture plates.

Cell counts: Cell numbers were determined in media by direct counting with a hemocytometer, and on matrix by first rinsing membranes with fresh medium DMEM + 10% FBS) followed by treatment with trypsin (10%, at 37° C. for 5 minutes) prior to counting.

SEM operating conditions: A Zeiss 962 instrument was utilized with an accelerating voltage of 10kv, and a working distance of 15mm. Polaroid type 55 p/n (u4)

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was utilized at various magnifications, as indicated.  
Sample coat: carbon coat (100Å) & 100Å aupd.

Specimen preparation: For primary fixation, the culture growth medium was replaced with 2%  
5 glutaraldehyde in Eagle's DMEM without serum. Several changes were performed to ensure a complete transition from growth media to Fixative. Fixation proceeded for 0.5 hours at room temperature. Cover slips were transferred to fresh vials containing 2%  
10 Glutaraldehyde in 0.1M Na Cacodylate pH 7.2 with 0.1M Sucrose and fixed for a further 1.5 hours at room temperature.

Dehydration, CPD, Mount and Sputter Coating:

Samples were rinsed in 0.1M Na Cacodylate pH 7.2,  
15 and cover slips were transferred to a CPD holder. Dehydration was performed in ethanol series (30%, 50%, 75%, 85%, 95% and 3 x 100%, 5 mins each), and samples were critical point dried. Cover slips were then mounted on Al stubs, carbon coated, using vacuum  
20 Evaporator (@ 100Å) and Sputter Coated with 100 Å AuPd.

## 12.2 RESULTS

p-GlcNAc membranes were tested for an ability to  
25 form a substrate on which cells may be grown in culture. Mouse fibroblast cells were grown in wells in the presence or absence of p-GlcNAc membranes and cell counts were taken daily to assay the viability of cultures. The results of one such series of cell  
30 counts in shown in FIG. 14. As indicated, by day 5 after plating, only the wells containing p-GlcNAc membranes were able to continue to sustain viable cells, demonstrating that p-GlcNAc membranes are capable of acting as efficient substrates for the  
35 continued growth of cells in culture.

Further, the SEM micrographs depicted in FIG. 15A-15B show healthy cells strongly attached to p-GlcNAc membranes.

5           13.   EXAMPLE: p-GlcNAc/COLLAGEN HYBRIDS

Presented in this Working Example is the formation and characterization of a p-GlcNAc/collagen hybrid material.

10                   13.1   MATERIALS AND METHODS

Materials: Bovine Type I collagen was used in preparation of the hybrids described in this study. p-GlcNAc was prepared according to the mechanical force method described, above, in Section 5.3.2.

15           Hybrid preparation: Collagen (10 milligrams/ml) and p-GlcNAc (0.25 milligrams/ml) suspensions were mixed, in different ratios, frozen in liquid N<sub>2</sub> (-80°C.), thermal soaked at -9° C. for 4 hours, and lyophilized. Material was dehydrothermally cross-  
20 linked under vacuum (approximately 0.030 Torr) at 60°C. for 3 days.

Cell Culture: Mouse 3T3 fibroblast cells were grown on the collagen/p-GlcNAc hybrids produced. Standard culturing procedures were followed, and SEM  
25 micrographs were taken after 8 days in culture.

13.2   RESULTS

Collagen and p-GlcNAc suspensions were mixed in differing ratios (namely, 3:1, 1:1, 2:2, and 1:3  
30 collagen:p-GlcNAc suspension ratios), frozen, lyophilized, and crosslinked. Such a procedure yielded collagen/p-GlcNAc slabs. SEM micrographs of the resulting materials are shown in FIGS. 16 B-E. Fig. 16A represents a collagen-only control material.  
35 Note the porous structure of the hybrid material.

The collagen/p-GlcNAc hybrids of the invention provide an efficient three-dimensional structure for the attachment and growth of cells, as shown in the SEM micrographs in FIGS. 17A-D.

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14.EXAMPLE: NMR CHARACTERIZATION OF PURE PREPARATIONS OF p-GlcNAc

Presented in this Example is an NMR (nuclear magnetic resonance) analysis of pure p-GlcNAc preparations.

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14.1 MATERIALS AND METHODS

p-GlcNAc preparations: The p-GlcNAc used in the NMR studies described here was prepared using the chemical purification method described, above, in Section 5.3.2, with hydrofluoric acid utilized as the chemical reagent.

15

NMR techniques: Solid state NMR data was obtained using a Bruker 500MH NMR spectrometer. Computer image analysis was used to transform the raw NMR spectrum data so as to eliminate background and to normalize baselines. An example of such transformed data are shown in FIG. 18. Transformed NMR curves such as that in Figure 18 were used to obtain areas for every carbon atom type, and to then calculate the CH<sub>3</sub>(area) to C-atom(area) ratios. Such values, obtained as described are provided in FIG. 20.

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14.2 RESULTS

Solid state NMR data was obtained by measuring the C<sup>13</sup>-NMR spectrum of a 500mg sample of p-GlcNAc. A typical NMR spectrum is shown in FIG. 19. The individual peaks represent the contribution to the spectrum of each unique carbon atom in the molecule. The relative percentage of each type of carbon atom in the molecule was determined dividing the area of the

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peak generated by that carbon species by the total sum of the areas under all of the NMR peaks obtained in the spectrum. Thus, it was possible to calculate the ratio of each of the atoms of the molecule measured by a reference atom. All p-GlcNAC molecules consist of N-acetylated glucosamine residues having C1, C2, C3, C4, C5 and C6 atoms, by definition. The ratio, then, of the area of the N-acetyl CH3 carbon atom peak to the areas of any of the glucosamine residue carbon atom peaks, above, should be 1.0 if all of the glucosamine residues in the polymer are N-acetylated. Data such as those in FIG. 20 were used to obtain values for the CH3(area) ratios.

The calculated ratios in Fig. 20 are in many cases equal to or nearly equal to 1.0, within experimental error, e.g. CH3/C2=1.097, CH3/C6=0.984, CH3/C5=1.007, CH3/C1=0.886. These results are consistent with the conclusion that the p-GlcNAC material of the invention is free of contaminants and is fully acetylated (i.e. that essentially 100% of the glucosamine residues are N-acetylated).

15. EXAMPLE: SYNTHESIS AND BIOLOGICAL CHARACTERIZATION OF CONTROLLED PORE SIZE THREE-DIMENSIONAL p-GlcNAC MATRICES

Described below, are methods for the production of three-dimensional p-GlcNAC based porous matrices having controlled average pore sizes. Such matrices have a variety of important applications, particularly, for example, as means for the encapsulation of cells. Such cell encapsulation compositions are useful as transplantable cell-based therapeutics, and in other cell & tissue engineering applications such as in cartilage regeneration. The capability to manipulate the morphology and

dimensionality of p-GlcNAc materials, as demonstrated here, provides a powerful tool in expanding the potential applications of the p-GlcNAc material of the invention.

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#### 15.1 MATERIALS AND METHODS

p-GlcNAc starting material: p-GlcNAc was prepared using the chemical purification method described, above, in Section 5.3.2, with hydrofluoric  
10 utilized as the chemical reagent.

Matrix formation: Suspensions (5mls) containing 20 mg p-GlcNAc samples were made in the solvents listed below in Section 15.2, prior to lyophilization. Samples were then poured into wells of tissue culture  
15 dishes and frozen at -20°C. The frozen samples were then lyophilized to dryness, and the resulting three dimensional matrices were removed.

Scanning electron microscopy techniques: The procedures utilized here were performed as described,  
20 above, in Section 12.1. The images shown in FIGS. 21A-G. are 200X magnifications of the matrix material, and a scale marking of 200 microns is indicated on each of these figures.

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#### 15.2 RESULTS

p-GlcNAc samples were obtained from each of the following solvents, as described, above, in Section  
15.1:

- A. Distilled water
- 30 B. 10% methanol in distilled water
- C. 25% methanol in distilled water
- D. Distilled water only
- E. 10% ethanol in distilled water
- F. 25% ethanol in distilled water
- G. 40% ethanol in distilled water

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Samples of matrix formed using each of the solvents were subjected to scanning, electron microscopic (SEM) analysis, as shown in FIGS. 21A-G. These figures reveal that the average matrix pore size  
5 decreases as the percentage of either methanol or ethanol increases in each suspension.

Specifically, pore diameter in the two water suspensions (FIGS. 21A and 21D) approach 200 microns on average. Pore size in the samples depicted in  
10 FIGS. 21C and 21F (25% methanol and ethanol, respectively) are between 30 and 50 microns on average.

The results shown here suggest that while both ethanol and methanol may successfully used to control  
15 p-GlcNAc pore size, ethanol may be more efficient than methanol in enabling the control of the p-GlcNAc matrix pore size.

16. EXAMPLE: CELL GROWTH ON THREE DIMENSIONAL  
20 POROUS p-GlcNAc MATRICES

Described in this Section are results demonstrating the successful use of three dimensional p-GlcNAc porous matrices as substrates for the culturing of cells.

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16.1 MATERIALS AND METHODS

p-GlcNAc starting material: p-GlcNAc was prepared using the chemical purification method described, above, in Section 5.3.2, with hydrofluoric  
30 acid utilized as the chemical reagent.

Matrix formation: Three-dimensional p-GlcNAc matrices were prepared by the lyophilization of suspensions of p-GlcNAc in water, water-ethanol, or water-methanol mixtures.

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Suspensions (5 mls) containing 20 mgs p-GlcNAc were prepared in the following solvents prior to lyophilization:

1. Distilled water only
- 5 2. 10% methanol in distilled water
3. 25% methanol in distilled water
4. Distilled water only
5. 10% ethanol in distilled water
6. 25% ethanol in distilled water
7. 40% ethanol in distilled water

Samples were poured into circular wells of plastic tissue culture dishes and were frozen at -20°C. The frozen samples were then lyophilized to dryness, and the resulting three dimensional matrices were removed. Samples of each matrix were subjected to scanning electron microscopic (SEM) analysis.

15 Cells: Mouse embryo BALBC/3T3 fibroblast cell line (clone A31), obtained from the ATCC, were used for culturing on the three dimensional porous p-GlcNAc matrices.

Culturing conditions: One cm<sup>2</sup> samples of porous 20 matrices were placed in tissue culture wells and were covered with a standard tissue-culture growth medium. Each well was seeded and cells were cultured for 6 days at 37°C in a CO<sub>2</sub> incubator (5% CO<sub>2</sub>).

SEM procedures: Matrix samples were fixed and 25 subjected to SEM analysis as described, above, in Section 12.1. The matrices were prepared by lyophilizing p-GlcNAc in distilled water. Images vary in magnification from 100X to 5000X, as indicated in figure legends (FIGS. 22A-G).

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## 16.2 RESULTS

SEM photographs of p-GlcNAc matrices containing attached mouse fibroblast cells attached are shown in FIGS 22A-G. These photographs show that the three 35 dimensional p-GlcNAc matrices contain attached mouse

fibroblast cells. Further, the photographs reveal that there is a close interaction and connection between the cells and the p-GlcNAc matrix material. It is also notable that the cells have a rounded three-dimensional morphology which is different from the flat, spread shape of the cells when cultured directly onto plastic culture dishes. Cell viabilities were determined to be greater than 95%.

10 17. EXAMPLE: p-GlcNAc SUCCESSFULLY PREVENTS POST SURGICAL ADHESIONS

The Example presented herein demonstrates the successful use of p-GlcNAc materials, specifically a p-GlcNAc membrane and gel formulation, to prevent the formation of post surgical adhesions in a series of animal models for such adhesions.

17.1 MATERIALS AND METHODS

Synthesis p-GlcNAc-lactate: p-GlcNAc membrane starting material was produced by the chemical method, as described, above, in Section 5.3.2, with hydrofluoric acid utilized as the chemical reagent.

The p-GlcNAc was converted to deacetylated p-GlcNAc by the following method. (It should be noted that approximately 1.4 g of p-GlcNAc are needed to produce each 1 g of p-GlcNAc lactate, given the expected loss in mass of approximately 15% which occurs upon deacetylation.) Approximately 200mg of p-GlcNAc membrane material were mixed vigorously with approximately 200 ml 60% NaOH. The vigorous shaking served to separate the p-GlcNAc membrane material to the extent possible. The NaOH solution used was made at least 12 hours before using. Samples were placed in an 80°C water bath for 6 hrs, with periodic shaking to separate and wet p-GlcNAc material. After 6 hrs, the samples were taken from water bath and the NaOH

solution was immediately removed. The membrane materials were washed with ddH<sub>2</sub>O, at room temperature, until a pH of 7 was reached. The membranes were removed from the water and dried on a Teflon sheet.

5 At this point a 2 mg sample was collected for C, H, N analysis in order to determine extent of deacetylation. Further, solubility in 1% acetic acid was checked, with a solubility of 1 mg/ml indicating that the p-GlcNAc material was appropriately  
10 deacetylated.

The partially deacetylated pGlcNAc was then converted to pGlcNAc-lactate using the following method: Sufficient 2-propanol (containing 10% water) to wet all of the partially deacetylated pGlcNAc  
15 material and to allow for stirring was added to 1g of the partially deacetylated p-GlcNAc in a 250 ml Erlenmeyer flask. (Approximately 30 mls 2-propanol necessary.) 2-propanol must be reagent grade, and fresh prior to each synthesis. With stirring, 1.1 mL  
20 of a 50% aqueous lactic acid solution. Lactic acid should be reagent grade, and must be analyzed to determine exact concentration of available (i.e., non-esterified) lactic acid present. This was generally accomplished by titration with 0.1N NaOH to the  
25 phenolphthalein end-point (pH 7.0). The concentration of lactic acid used must be constant, i.e., must be +/- 1 percent, for each p-GlcNAc synthesis. The mixture was allowed to stir for at least two hours. It is possible to add low heat in order to elevate the  
30 reaction rate. Reaction time may be extended, or the amount of 50% aqueous lactic acid may be increased so that the reaction goes to completion. After stirring, the contents of the flask were poured through a Buchner funnel using quantitative ashless filter  
35 paper. The material was washed with 15 ml of

anhydrous 2-propanol. The material was allowed to air dry in a fume hood for 2 hours and then placed in an oven at 40°C overnight. For every gram of partially deacetylated p-GlcNAc starting material, a final p-GlcNAc-lactate weight of approximately 1.4 g, (i.e., an increase of 40% in mass) should be obtained.

Formulation of p-GlcNAc-lactate as a gel: The p-GlcNAc-lactate material was formulated into a gel as follows: p-GlcNAc-lactate starting was dissolved in dd-deionized water to a concentration of between 0.1-4.0% p-GlcNAc-lactate, by weight. Reagent grade propylene glycol (2-propandiol) was then added to a final propylene glycol concentration of between 1-10%. In some cases, a preservative was added to prevent bacterial and/or fungal contamination of the product. Typically, concentrations of p-GlcNAc-lactate of between 0.1%-4.0% were prepared. The viscosity of these preparations increases dramatically as the p-GlcNAc-lactate percentage increases, such that formulations having 0.5% or more of the p-GlcNAc-lactate behave as gels.

Animal models:

Sprague-Dawley rats: Adhesions are produced in this model by abrading or irritating the serosal surface of the cecum and apposing it to an area of parietal peritoneum. The success rate for inducing adhesions in control animals with this method is reported at an average 80%.

Specifically, the surgical procedure for inducing post surgical adhesions in these rats involved the following. Animals were placed in dorsal recumbency and prepared and draped accordingly for aseptic surgery. Abdominal cavities were exposed through a midline incision. An area, approximately 0.5 cm x 1.0 cm, of parietal peritoneum on the left abdominal wall

was carefully excised, removing a thin layer of muscle, along with the peritoneum.

The cecum was then elevated and isolated. An area, approximately 0.5 cm x 1.0 cm, on the lateral surface of the proximal end of the cecum was abraded by rubbing ten times with a dry gauze. The cecum was then scraped with a scalpel blade to cause minute petechial hemorrhages. The cecal abrasion and the peritoneal incision were left exposed for 15 minutes.

After 15 minutes, the test article (i.e., the p-GlcNAc material) or control article was applied to the cecal wound. The cecal abrasion and the peritoneal wound were then opposed and held in contact with Allis tissue forceps for an additional 15 minutes.

The cecum was then replaced into the abdomen such that the abraded area of the cecum was adjacent to the peritoneal site. The abdominal incision was closed and the animal was allowed to recover from the anesthesia.

Fourteen days after surgery, animals were euthanized and the abraded area was examined for the formation of post surgical adhesions. If adhesions were present, the entire area involved in the adhesion (i.e., body wall, test or control article, and internal organs) were dissected free of the animal.

The extent of involvement and tenacity of adhesions was evaluated according to the following scales:

Extent of involvement scores:

- 0 no adhesion
- 1 adhesion  $\leq$  25% of the area
- 2 adhesion  $\leq$  50% of the area
- 3 adhesion  $\leq$  75% of the area
- 4 adhesion  $>$  25% of the area



Tenacity Scores:

- 0 no adhesion  
1 adhesion freed with blunt dissection  
2 adhesion freed with aggressive dissection  
3 adhesion requiring sharp dissection

5

Additional animal models: Pig and horse large animal bowel models were used to assess the prevention of peritoneal adhesions.

10 Surgical procedure: The animals were placed in dorsal recumbency and prepared and draped accordingly for aseptic surgery. The abdominal cavity was exposed through a midline incision. The small intestine was elevated and a 2 cm X 2 cm section was identified, extensively abraded (approximately 200 strikes using a  
15 scalpel), and allowed to dry for 10 minutes. The test article (i.e., p-GlcNAc material) or control article was then applied to the abraded wound, and the wounded section of the small intestine was replaced into the abdomen. In such a large bowel type of animal model,  
20 six wounds, each separated by 4 inches of bowel from the adjacent wound provides an environment highly prone to form adhesions. Following the last of the wounds, the abdominal incision is closed and the animal is allowed to recover from the anesthesia.

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Analysis of peritoneal adhesions: Twenty one days after surgery, animals were euthanized and the abraded area was examined, with adhesion formation being evaluated following a procedure similar to that  
30 of the Sprague-Dawley rat cecum model.

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17.2 RESULTS

When injury occurs, the body sets in motion a complex set of responses designed to restore the injured area. In the final stages of healing,  
35 connective tissue forms at the wound site to

regenerate the body structure and protect the affected area from further damage. In some instances this cascade of events does not work properly and can lead to life threatening conditions.

5 For example, as a visceral organ heals following surgery, a fibrin clot generated during the surgical procedure may invade the surface of adjoining organs forming a link which allows for fibroblast migration. This migration leads to collagen deposition and tissue  
10 growth, which in turn causes the organs involved to adhere to one another.

Such adhesions, referred to as post surgical adhesions, may produce pain, obstruction and malfunction by distorting the organ or organs  
15 involved. Immobilized joints, intestinal obstruction and infertility are often linked to the formation of post-surgical adhesions. Furthermore, post surgical adhesion will complicate and extend the length of future surgical procedures in the surrounding region.  
20 This last issue is of particular relevance to open heart surgeries and cesarean section obstetrical procedures where additional surgeries may be required. The formation of adhesions is very common following abdominal, cardiovascular and orthopedic surgical  
25 procedures.

When adhesions become pathological and seriously interfere with organ function, surgical adhesiolysis (sharp or blunt dissection of the adhesion in conjunction with meticulous surgical techniques) is  
30 the treatment that is currently used to eliminate adhesions. In 1991, approximately 500,000 adhesiolysis procedures were performed. This procedure is, however, notoriously ineffective, with the frequency of recurrence of adhesion formation  
35 reported to be as high as 90%. Further, no other

technique or composition has proven effective in the prevention of such post surgical adhesions.

The results presented herein, therefore, are significant in that they demonstrate the effectiveness of the p-GlcNAc materials of the invention for the prevention of post surgical adhesions. Specifically, the results presented here demonstrate the efficacy of p-GlcNAc based solid and liquid formulations as barriers to the formation of abdominal post surgical adhesions in accepted rat and pig animal model systems.

One of the accepted animal models used to study adhesion formation employs visceral-parietal peritoneal adhesions in Sprague-Dawley rats. Both partially deacetylated p-GlcNAc membranes and p-GlcNAc-lactate gel formulations prevented and/or considerably reduced the incidence of adhesion formation as compared with either non-treated controls treated with InterCEED™ (Johnson & Johnson), the only commercially available product for this indication.

Specifically, a total of 18 rats were used to test p-GlcNAc-lactate gel formulations. 12 animals were used as controls, with 6 receiving no treatment and 6 receiving InterCeed™. 6 animals received 0.25% p-GlcNAc-lactate gel, 10% propylene glycol, water. Animals receiving the p-GlcNAc-lactate gel treatment showed a significantly reduced incidence of postoperative adhesion formation, compared to either of the controls, as shown, below, in Table VIII.

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TABLE VIII

	Extent of Involvement	Tenacity
Control (No treatment)	1 +/- 2.1	1 +/- 1.5
InterCEED™	1 +/- 1.8	1 +/- 1.5
p-GlcNAc-lactate gel	0 +/- 0.8	1 +/- 1.2

Partially deacetylated p-GlcNAc membranes were also tested for their ability to prevent to occurrence of post surgical adhesions in the rat animal model. A total of 22 rats were used in the study. 12 animals were used as controls, with 6 receiving no treatment and 6 receiving InterCEED™. Ten animals each received a 1cm x 1cm membrane of an approximately 60% deacetylated p-glcNAc formulation. The animals which received the partially deacetylated p-GlcNAc membrane showed a significant reduction in the incidence of formation of postoperative adhesions, as compared with the non-treated controls and InterCEED™, as shown, below, in Table IX.

TABLE IX

	Extent of Involvement	Tenacity
Control (No treatment)	3 +/- 1.8	1 +/- 0.6
InterCEED™	3 +/- 1.6	1 +/- 0.4
p-GlcNAc-membrane	1 +/- 0.8	1 +/- 0.3

Large animal bowel models for the prevention of peritoneal adhesions were also used to test p-GlcNAc compositions. Specifically, six pigs and one horse were used to study both the partially deacetylated p-GlcNAc membrane and the p-GlcNAc-lactate gel. The partially deacetylated p-GlcNAc membrane consisted of a 2 cm X 2 cm piece of 60% deacetylated p-GlcNAc membrane, while the p-GlcNAc-lactate gel consisted of 0.25% p-GlcNAc lactate formulated with 10% propylene glycol and water. Control animals received no treatment to the wounded site.

The results of these large animal studies revealed that, while the control sites formed multiple adhesions and scare tissue in the surrounding site, both the p-GlcNAc membrane and gel formulations effectively prevented the formation of adhesions.

Samples from control and treated sites were additionally examined using SEM, which showed an increased amount of fibrosis in the control sites as compared to the treated tissues.

#### 18. EXAMPLE: BIODEGRADABILITY OF p-GlcNAc MATERIALS

The Example presented in this Section demonstrates that p-GlcNAc materials of the invention may be prepared which exhibit controllable in vitro and in vivo biodegradability and rates of resorption.

##### 18.1 MATERIALS AND METHODS

p-GlcNAc materials: Prototype I was made by the method described, above, in Section 5.3.2, via the chemical method, with hydrofluoric acid being utilized as the chemical reagent. Prototype I represented 100% acetylated p-GlcNAc.

The p-GlcNAc starting material of prototype 3A was made by the method described, above, in Section

5.3.2, via the chemical method, with hydrofluoric acid being utilized as the chemical reagent. The p-GlcNAc material was then deacetylated by the method described, above, in Section 5.4. Specifically, the  
5 p-GlcNAc material was treated with a 40% NaOH solution at 60°C. for 30 minutes. The resulting prototype 3A was determined to be 30% deacetylated.

The p-GlcNAc starting material of prototype 4 was made by the method described, above, in Section 5.3.2,  
10 via the chemical method, with hydrofluoric acid being utilized as the chemical reagent. The p-GlcNAc material was then deacetylated by treatment with a 40% NaOH solution at 60°C. for 30 minutes. Next, the fibers were suspended in distilled water, frozen at -  
15 20°C., and lyophilized to dryness. Prototype 4 was also determined to be 30% deacetylated.

Abdominal implantation model: Sprague Dawley albino rats were utilized for the abdominal implantation model studies. Animals were anesthetized  
20 and prepared for surgery, and an incision was made in the skin and abdominal muscles. The cecum was located and lifted out. A 1 cm x 1 cm membrane of p-GlcNAc material was placed onto the cecum, and the incision was closed with nylon. Control animals were those in  
25 which no material was placed onto the cecum.

Animals were opened at 14 and 21 days post implantation. Photographs were taken during the implant and explant procedures (FIGS. 23A-E). Samples of cecum were prepared for histopathology after the  
30 explant procedure.

p-GlcNAc in vitro degradation lysozyme-chitinase assay: The assay is a colorimetric assay for N-acetyl glucosamine, and was performed as follows: 150 $\mu$ l of a reaction sample was pipetted into 13x100mm glass  
35 disposable test tubes, in duplicate. 25 $\mu$ l of 0.25M

potassium phosphate buffer (pH 7.1) was added to each test tube, followed by the addition of 35 $\mu$ l of 0.8M potassium borate solution (pH 9.8). Tubes were immediately immersed into an ice-bath for a minimum of 5 2 minutes. Samples were then removed from the ice-bath, 1ml of freshly prepared DMAB reagent was added, and the samples were vortexed. DMAB (Dimethyl aminobenzaldehyde) reagent was made by adding 70mls of glacial acetic acid and 10mls of 11.6N (concentrated) 10 HCl to 8 grams of p-dimethyl aminobenzaldehyde. Samples were then incubated at 37°C for 20 minutes.

To prepare a standard curve, the following procedure was utilized. A GlcNAc stock solution was diluted to 0.1mg/ml with 0.010M sodium acetate buffer 15 (pH 4.5), and 0 $\mu$ l, 20 $\mu$ l, 30 $\mu$ l, 90 $\mu$ l or 120 $\mu$ l of the diluted GlcNAc solution was added to a set of test tubes. This was followed by the addition of 150 $\mu$ l, 130 $\mu$ l, 60 $\mu$ l or 30 $\mu$ l, respectively, of 0.010M sodium acetate buffer (pH 4.5) to the test tubes. Next, 25 $\mu$ l 20 of 0.25M potassium phosphate buffer (pH 7.1) and 35 $\mu$ l of 0.8M potassium borate buffer (pH 9.8) were added to each test tube. A duplicate set of test tubes is prepared by the same procedure.

The test tubes are capped and boiled at 100°C. for 25 for exactly 3 minutes. The tubes are then immersed in an ice bath. The tubes are removed from the ice bath and 1ml of DMAB reagent, freshly prepared according to the method described above in the Section, is added to each tube. The tubes are incubated at 37°C for 20 30 minutes. The absorbance of the contents of each tube is read at 585nm. Absorbance should be read as quickly as possible. The standard curve is plotted on graph paper and used to determine the concentration of N-acetyl glucosamine in the reaction samples. A 35 typical standard curve is shown in FIG. 23.

18.2 RESULTS

The in-vitro biodegradability of p-GlcNAc materials was studied in experiments which assayed the relative susceptibility of p-GlcNAc membrane materials to degradation by lysozyme. p-GlcNAc membranes were exposed to an excess of lysozyme in a 10mM acetate buffer, and the subsequent release of N-acetyl glucosamine was determined using the assay described, above, in Section 18.1.

10 The results of these experiments indicated that partially deacetylated membranes are more susceptible to digestion by lysozyme (see FIG. 24) and, further, that the rate of lysozyme degradation is directly related to the extent of deacetylation (see FIG. 25, 15 which compares the degradation rates of a 50% to a 25% deacetylated p-GlcNAc membrane).

Additionally, experiments were performed which addressed the in-vivo biodegradability of p-GlcNAc materials. Such experiments utilized an abdominal 20 implantation model. Three p-GlcNAc materials, as listed below, were tested.

p-GlcNAc materials tested:

- 25 1) p-GlcNAc, fully acetylated (designated prototype 1);
- 2) partially deacetylated p-GlcNAc membrane (designated prototype 3A); and
- 3) lyophilized and partially deacetylated p-GlcNAc membrane (designated prototype 4).

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The fully acetylated p-GlcNAc (prototype 1) was resorbed within 21 days, as shown in FIGS. 26A-26C. The partially deacetylated p-GlcNAc membrane (prototype 3A) was completely resorbed within 14 days, 35 as shown in FIGS. 26D-26E. The lyophilized and



partially deacetylated p-GlcNAc membrane (prototype 4) had not yet been completely resorbed after 21 days post-implantation.

Histopathology analyses showed that once the p-GlcNAc material has been resorbed there were no histological differences detectable between tissue samples obtained from the treated and from the control animals.

10 19. EXAMPLE: p-GlcNAc HEMOSTASIS STUDIES

The experiments described herein study the efficacy of the p-GlcNAc materials of the invention for the control of bleeding. The success of the p-GlcNAc materials in controlling bleeding is, further, compared against commercially available hemostatic products.

19.1 MATERIALS AND METHODS

p-GlcNAc and control materials: partially deacetylated (approximately 70%) p-GlcNAc membranes were made using the chemical separation technique described, above, in Section 5.3.2, with hydrofluoric acid being utilized as the chemical reagent, and the techniques described, above, in Section 5.4. 2 cm x 1 cm pieces were used. p-GlcNAc-lactate gel (4% p-GlcNAc-lactate, formulated in propylene glycol and water) was produced following the methods described, above, in Section 17.1. The control material utilized for the study of bleeding in the spleen and liver was Gelfoam™ (Upjohn Company). Gelfoam™ and Avitene™ (Medchem Products, Inc.) were the control materials used in the study of small blood vessel bleeding.

Test animals: New Zealand White rabbits were used. 3 animals received two wounds in the spleen and one wound in the liver. 4 animals received five

surgical wounds to blood vessels of similar size in the caudal mesenteric artery system.

Surgical preparation: The animals were anesthetized with ketamine HCl and Xylazine. The animals were placed in dorsal recumbency, and all the hair from the abdomen was removed. The abdomen was then scrubbed with povidone-iodine and 70% isopropyl alcohol and draped for aseptic surgery.

Liver/spleen surgical procedures: A midline incision was made and either the spleen or liver was exteriorized and packed with moist lap sponges. A 3-4 mm diameter cork bore was used to make a circular wound of about 2 mm depth at one end of the organ. Once the splenic tissue was removed, a pre-weighed 4" X 4" gauze sponge was used to absorb all the blood lost from the splenic wound for a period of one minute. The sponge was re-weighed to quantify the amount of blood lost from that particular wound. The test animal was then treated by application to the wound of one of the treatment materials. The time until hemostasis and the amount of blood lost prior to hemostasis was recorded.

After hemostasis in the first wound was achieved, a second wound in the spleen and one wound in the liver were made following the same procedure.

Small blood vessel surgical procedure: A midline incision was made and the small bowel was exteriorized exposing the caudal mesenteric artery system. The bowel was packed with moist lap sponges and five blood vessels of about the same size were identified. A scalpel was used to make a wound of about 1 mm depth at one of the vessels. A pre-weighed 4" x 4" gauze sponge was used to absorb all the blood lost from the vessel wound for a period of one minute. The sponge was re-weighed to quantify the amount of blood lost

from that particular wound. The animal was then treated by application to the wound of one of the treatment materials. The time until hemostasis and the amount of blood lost prior to hemostasis was  
5 recorded.

After hemostasis in the first wound was achieved, four more wounds were made following the same procedure.

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### 19.2 RESULTS

p-GlcNAc materials were tested for their ability to control bleeding in the spleen and liver of rat animal models. The p-GlcNAc materials tested were:  
1) partially deacetylated (approximately 70%) p-  
15 GlcNAc; and 2) p-GlcNAc-lactate gel (4% p-GlcNAc-lactate, formulated in propylene glycol and water). The effectiveness of these p-GlcNAc materials was compared to Gelfoam™ (Upjohn Company).

Each material was tested three times (twice in  
20 the spleen and once in the liver). Both of the p-GlcNAc based materials exhibited an effectiveness in controlling bleeding within the first minute after application which was comparable to that of Gelfoam™.

The p-GlcNAc based materials have additional  
25 advantages. Specifically, the p-GlcNAc materials do not need to be held in place during the procedure, may be left in the body, where they will be resorbed within two to three weeks (Gelfoam™ is not indicated for this purpose), are compatible with both general  
30 and minimally invasive surgical procedures.

Next, the efficacy of p-GlcNAc based materials in the control of bleeding in small blood vessels was studied, and compared against commercially available hemostatic products.

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Each material was tested five times (twice in one of the animals and once in the other three animals). The p-GlcNAc membrane and gel formulations were easily applied to the site and controlled the bleeding within 5 2 minutes. Gelfoam™, which had to be held in place in order to perform its function achieved hemostasis within the same 2 minute range as the p-GlcNAc materials. Avitene™, a fibrous material made of collagen, was difficult to handle and required more 10 than five minutes to control the bleeding.

Thus, the results described herein demonstrate that the p-GlcNAc materials tested here represent effective, convenient hemostatic agents.

15 20. EXAMPLE: p-GlcNAc DRUG DELIVERY SYSTEMS

Described herein are studies demonstrating the successful use of p-GlcNAc materials to deliver anti-tumor drugs to the site of malignant skin cancer and colon cancer tumors such that the delivered anti tumor 20 drugs exhibit a therapeutic impact upon the tumors.

20.1 MATERIALS AND METHODS

p-GlcNAc-lactate drug delivery compositions:

Mixtures of 5'-fluorouracil (5'-FU) and p-GlcNAc-lactate were formulated as follows; 0.5mL of 5'-FU 25 (50mg/mL) was mixed with 0.5mL of propylene glycol, and 2.0mL of 4% p-GlcNAc-lactate was added and mixed. The p-GlcNAc-lactate was produced using the techniques described, above, in Section ---. Even after 30 extensive mixing, the 5'FU did not completely dissolve into the p-GlcNAc-lactate gel. Assuming complete mixing, the final concentration of 5'-FU would be 6.25mg/mL.

Mixtures of mitomycin (Mito) and p-GlcNAc-lactate 35 were formulated as follows; 0.5mg of Mito (lyophilized

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powder) were dissolved in 5ml of propylene glycol, and 0.5ml of the Mito solution was mixed with 0.5mL of MPT's 4% p-GlcN-lactate preparation to give a final Mito concentration of 0.2mg/ml and a final p-GlcNac-lactate concentration of 2%. The materials were compatible, with the Mito dissolving easily into the p-GlcNac-lactate gel.

p-GlcNac membrane 5'FU delivery compositions:

Samples of 5'-fluorouracil (5'-FU) were immobilized into discs of pure p-GlcNac membrane material produced using the chemical separation method described, above, in Section 5.3.2, with hydrofluoric acid being utilized as the chemical reagent. Each disc described here had a diameter of 1.5cm, as described here.

For the preparation of high dose (HD) discs, 0.64mL of a 50mg/mL solution of 5'-FU was mixed with suspensions containing approximately 8 mg of pure p-GlcNac. The mixtures were allowed to stand for several hours to promote the absorption of 5'-FU into the p-GlcNac, and were then dried at 55°C for 3.5 hours. The resulting HD discs contained a total of 32mg 5'-FU, which is equivalent to approximately twice the normal total 14 day dose of 5'-FU typically given to a cancer patient,

Low dose (LD) 5'-FU containing p-GlcNac discs were prepared in the same manner, except that the LD discs contained 17mg of 5'-FU, an amount equivalent to equal the normal total human dose for a 14 day period, normalized to the weight of the experimental mice based on the typical dose of 5'-FU per Kg body weight for humans.

Test Animals: For the 5'FU study, SCID (severe combined immunodeficiency) mice were inoculated with subcutaneous flank injections of HT-29 colon cancer cells (ATCC;  $1 \times 10^5$  cells per inoculum) obtained by

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standard tissue culture methods, in order to produce HT-29 colon cancer tumors. These injections led to palpable tumors which were harvested in 14-21 days. Tumors were dissected and necrotic tissue was cut  
5 away. The HT-29 colon cancer tumors were sliced into 3x3x3mm pieces.

The experimental SCID mice were anesthetized via intra-peritoneal injections with a standard dose of avetin, and a slice of HT-29 colon cancer tumor was  
10 implanted onto the cecum of each mouse. Specifically, each mouse was surgically opened to expose its abdomen and the cecum was located, which was nicked with a scalpel to make a small incision. A 3x3x3mm tumor slice was sutured over the incision onto the cecum  
15 using 5.0 silk sutures. The abdomen was then closed using Clay Adams staples.

All mice were caged singly and fed for two weeks. All mice were healthy and none had obstructed colons at the end of the two week period.

20 On day 14, each mouse was anesthetized, and its abdomen was reopened. The growing tumors were measured (length and horizontal dimensions). Tumors were then treated with the p-GlcNAc/anti-tumor drug or were used as controls.

25 Six mice were used for the p-GlcNAc-lactate 5'FU study, and 15 mice were used for the p-GlcNAc membrane 5'FU study.

For the mitomycin study, nine SCID mice were inoculated with sub-cutaneous injections of A431  
30 squamous cell skin cancer cells (ATCC;  $1 \times 10^5$  cells per inoculum). Tumors resulted in all mice within 14 days.

Treatments: For the p-GlcNAc-lactate 5'FU study, animals were treated once daily by "painting" the 5'-  
35 fluorouracil (5'-FU)-containing p-GlcNAc gel mixture

onto the skin area over the tumor mass. Measurements of the tumor size were obtained daily. Control animals included animals treated with p-GlcNAc alone, without 5'-FU, and animals which received no  
5 treatment.

For the p-GlcNAc membrane 5'FU study, the HT29 colon tumors in the SCID mice were treated by surgically implanting discs of the drug-containing p-GlcNAc membrane material directly onto their surface,  
10 after having allowed the tumor to grow on the colon for 14 days. Mice were sacrificed 14 days following the implant procedure. Measurements of tumor volumes were made immediately prior to implanting the drug-containing p-GlcNAc membranes on day 0 and at the  
15 termination of the experiment on day 14. Control animals included ones treated with the p-GlcNAc membrane without 5'-FU, and controls which received no treatment. Additionally, two animals received daily systemic injections of 5'-FU in doses equivalent to  
20 the HD and LD regimen.

For the p-GlcNAc-lactate Mito study, animals were treated daily as in the p-GlcNAc-lactate 5'-FU study, with 3 animals being treated with the Mito containing mixture. In addition, 3 animals were treated with p-  
25 GlcNAc minus Mito, 2 animals received no treatment, and 1 animal received propylene glycol.

## 20.2 RESULTS

### 20.2.1 p-GlcNAc-LACTATE 5'FU

30 Experiments designed to study the effect of p-GlcNAc-lactate 5'FU drug delivery systems on tumor size were conducted, as described, above, in Section 20.1.

The largest length and width dimension were  
35 measured for each tumor and the cross-sectional area

using these dimensions was calculated. The cross-sectional area values are shown in Table X, below.

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Table X

	Animal #	Treatment	Tumor Size (cm <sup>2</sup> )			
			Day 0	Day 4	Day 11	Day 15
5	1	CL + 5FU	63	90	168	156
	2	CL + 5FU	48	56	70	88
10	3	CL Control	21	36	88	108
	4	CL Control	58	110	150	195,30
15	5	Nothing	40	64	132	234
	6	Nothing	28	42	100	132
			<u>% Increase in Size</u>			
			Day 0	Day 4	Day 11	Day 15
20	1	CL + 5FU	0	43	167	147
	2	CL + 5FU	0	17	47	84
	3	CL Control	0	71	319	414
25	4	CL Control	0	90	160	289
	5	Nothing	0	61	232	488
30	6	Nothing	0	48	253	366

The data comparing p-GlcNAc-lactate 5'FU treated animals with controls are shown in FIGS. 27-28. The data summarized in Table X and FIGS. 27-28 clearly suggest that the HT-29 subcutaneous tumors in the rats treated with the 5'-FU containing p-GlcNAc-

lactate gels have a significantly retarded rate of growth compared to controls. Their growth has been slowed 2.5-fold in comparison to the p-GlcNAc-lactate gel controls and 4-fold compared to the no treatment controls.

#### 20.2.2 p-GlcNAc-LACTATE MITO

Experiments designed to study the effect of p-GlcNAc-lactate 5'FU drug delivery systems on tumor size were also conducted, as described, above, in Section 20.1.

The largest length and width dimensions were measured for each tumor and the cross sectional area using these dimensions was calculated. The cross-sectional area values were as shown in Table XI, below.

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Table XI

	Animal #	Treatment	Tumor Size (cm <sup>2</sup> )			
			Day 0	Day 3	Day 5	Day 8
5	1	pGlcN-L + Mito	23	23	42	49
	2	pGlcN-L + Mito	23	16	54	63
10	3	pGlcN-L + Mito	72	99	Term	Term
	4	pGlcN-L control	27	54	140	203
15	5	pGlcN-L control	30	54	96	140
	6	pGlcN-L control	30	58	200	221
20	7	Nothing	48	75	126	300
	8	Nothing	44	80	207	Dead
	9	Propylene glycol	49	86	180	216
25			<u>% Increase in Size</u>			
			Day 0	Day 3	Day 5	Day 8
	1	pGlcN-L + Mito	0	0	83	135
30	2	pGlcN-L + Mito	0	-30	135	174
	3	pGlcN-L + Mito	0	38	Term	Term
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	4	pGlcN-L control	0	100	419	652
5	5	pGlcN-L control	0	80	220	367
	6	pGlcN-L control	0	93	567	637
	7	Nothing	0	56	163	525
10	8	Nothing	0	82	370	Dead
	9	Propylene glycol	0	76	267	341

15 The data comparing p-GlcNAc-lactate Mito treated  
animals with controls are shown in FIGS. 29-30. The  
data summarized in Table XI and FIGS. 29-30 clearly  
suggest that the tumors growing in the rats treated  
with the Mitomycin-containing p-GlcNAc-lactate gels  
20 animals have a significantly retarded rate of growth.  
Their growth was been slowed by 4-fold in comparison  
to the p-GlcNAc-lactate gel controls and 4-fold  
compared to the no treatment controls.

### 25 20.2.3 p-GlcNAc MEMBRANE 5'FU

Next, experiments designed to study the effect of  
p-GlcNAc membrane 5'FU drug delivery systems on skin  
cancer tumor size were conducted, as described, above,  
in Section 20.1.

30 The tumor volume data obtained during the study,  
including percent change in volume caused by the  
different treatments, is summarized in Table XII,  
below. Tumors were assumed to be cylindrical in  
shape. Their volumes were determined by measuring  
their width and length, and using the following  
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equation:  $V = \pi r^2 l$ , where the radius  $r$  is 0.5 times the width and  $l$  is the length.

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TABLE XII  
p-GlcNAc Membrane + 5'-Fluorouracil Animal Data

Animal #	Treatment	Tumor Volume Pre-Treatment (mm <sup>3</sup> )	Tumor Volume Post-treatment (mm <sup>3</sup> )	% Change	Comments
<u>A. 5'-FU High Dose:</u>					
1.	5'-FU HD	393	283	-28.0	
2.	5'-FU HD	785	308	-60.8	
3.	5'-FU HD	98.1	62.8	-36.0	
4.	5'-FU HD	785	550	-30.0	
Average per Animal				-38.7	
<u>B. 5'-FU Low Dose:</u>					
5.	5'-FU LD	603	170	-71.9	
6.	5'-FU LD	603	615	2.0	
7.	5'-FU LD	269	198	-26.0	
8.	5'-FU LD	169	226	33.3	
Average per animal				-15.7	
<u>C. p-GlcNAc Control:</u>					
9.	p-GlcNAc Control	170	550	320.0	
10.	p-GlcNAc Control				Died day 12
Average per Animal				320.0	

TABLE XII (Cont'd)

<u>D. No Treatment Control:</u>					
11. No Treatment	402	864	215.0	Sat. tumors	
12. No Treatment	21.2	572	2700.0	Sat. tumors	
Average per Animal			1457.0		
<u>E. 5'-FU via Intravenous Injection - Control:</u>					
13. Low Dose	402	703	175.0		
14. Low Dose	402	402	0.0	Died day 13	
15. High Dose	402	132	-67.0	Died day 13	

FIG. 31 summarizes a portion of the data presented, above, in Table XII. as shown in FIG. 31, the data strongly suggest that tumors treated with the high dose (HD) 5'-FU-containing p-GlcNAc membranes have stopped growing and have, in all cases, actually gotten significantly smaller. The low dose (LD) polymer materials resulted in disease stability and slight decrease in tumor size. In contrast, the tumors in the control animals continued to rapidly increase in size. It is interesting to note that two of the three control animals which were treated via IV died during the study, indicating that systemic delivery of the equivalent amount of 5'-FU is lethal, whereas site-specific delivery via the p-GlcNAc polymer is efficacious in ridding the animal of the disease.

### 20.3 CONCLUSION

The data presented in this Section strongly suggest that the site-specific delivery of anti-tumor drugs has a positive effect in retarding and reversing tumor growth. Successful results were obtained using p-GlcNAc drug delivery compositions produced having two different formulations, namely p-GlcNAc-lactate and p-GlcNAc membrane formulations. Further, successful results were obtained using two different anti-tumor drugs, 5'-FU and Mito. Thus, the p-GlcNAc drug delivery systems of the invention exhibit anti-tumor activity, useful, for example, in the delivery of drugs specifically to the site of the tumor cells of interest.

It is apparent that many modifications and variations of this invention as set forth here may be made without departing from the spirit and scope



thereof. The specific embodiments described above are given by way of example only, and the invention is limited only by the terms of the appended claims.

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WHAT IS CLAIMED IS:

1. An isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species comprising about 4,000 to about  
5 150,000 N-acetylglucosamine monosaccharides covalently attached in a  $\beta$ -1 $\rightarrow$ 4 conformation, free of protein, substantially free of other organic contaminants, substantially free of inorganic contaminants, and having a molecular weight of about 800,000 daltons to  
10 about 30 million daltons.

2. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 wherein the species comprises about 4,000 to about 15,000 N-  
15 acetylglucosamine monosaccharides covalently attached in a  $\beta$ -1 $\rightarrow$ 4 conformation, and having a molecular weight of about 800,000 daltons to about 3 million daltons.

3. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 or 2 wherein the species  
20 is a cell culture substrate.

4. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 or 2 wherein the species  
25 is a mat, string, rope, microsphere, microbead, membrane, fiber, powder, or sponge.

5. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 or 2 wherein the species  
30 is a three dimensional matrix formulation.

6. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 or 2 wherein at least one N-acetylglucosamine monosaccharide has been  
35 deacetylated.

7. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 6 wherein at least about 25% to about 75% of the N-acetylglucosamine monosaccharides have been deacetylated.

5

8. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 7 wherein at least about 70% of the N-acetylglucosamine monosaccharides have been deacetylated.

10

9. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 6 wherein the species is a mat, string, rope, microsphere, microbead, membrane, fiber, powder, or sponge.

15

10. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 6 wherein the species is a three dimensional matrix formulation.

20

11. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 or 2 wherein at least one monosaccharide contains a sulfate group, a sulfonyl group, an O-acyl group, an N-acyl group, an O-alkyl group, an N-alkyl group, an N-alkylidene

25

group, or an N-arylidene group.

12. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 or 2 wherein at least one monosaccharide is a phosphorylated derivative, a

30

nitrate derivative, an alkali derivative, or a deoxyhalogen derivative.

13. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 or 2 wherein at least

35

one monosaccharide forms a salt or a metal chelate.

14. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 or 2 wherein the species is isolated from a microalgae source.
- 5 15. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 14 wherein the microalgae source is a diatom source.
- 10 16. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 15 wherein the diatom is of the genus Thalassiosira.
- 15 17. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 16 wherein the diatom of the genus Thalassiosira is Thalassiosira fluviatilis or Thalassiosira weissflogii.
- 20 18. An isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species comprising about 4,000 to about 150,000 glucosamine monosaccharides covalently attached in a  $\beta$ -1 $\rightarrow$ 4 conformation, free of protein, substantially free of other organic contaminants, substantially free of inorganic contaminants, and having a molecular weight of about 640,000 daltons to about 24 million daltons.
- 25 19. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 18 wherein the species comprises about 4,000 to about 15,000 glucosamine monosaccharides covalently attached in a  $\beta$ -1 $\rightarrow$ 4 conformation, having a molecular weight of about 640,000 daltons to about 2.4 million daltons.
- 30 20. An isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species comprising about 4,000 to about 150,000 glucosamine monosaccharides covalently attached in a  $\beta$ -1 $\rightarrow$ 4
- 35

conformation, free of protein, substantially free of other organic contaminants, substantially free of inorganic contaminants, wherein at least one glucosamine monosaccharide has been acetylated.

5

21. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 20 wherein at least about 25% to about 75% of the glucosamine monosaccharides have been acetylated.

10

22. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 21 wherein at least about 30% of the glucosamine monosaccharides have been acetylated.

15

23. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 18 or 20 wherein the species is a mat, string, rope, microsphere, microbead, membrane, fiber, powder, or sponge.

20

24. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 18 or 20 wherein the species is a three dimensional matrix formulation.

25

25. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 18 or 20 wherein at least one monosaccharide contains a sulfate group, a sulfonyl group, an O-acyl group, an N-acyl group, an O-alkyl group, an N-alkyl group, an N-alkylidene group, or an N-arylidene group.

30

26. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 18 or 20 wherein at least one monosaccharide is a phosphorylated derivative, a nitrated derivative, an alkali derivative, or a deoxyhalogen derivative.

35

27. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 18 or 20 wherein at least one monosaccharide forms a salt or a metal chelate.

5 28. A method for isolating a poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species comprising about 4,000 to about 150,000 N-acetylglucosamine monosaccharides covalently attached in a  $\beta$ -1 $\rightarrow$ 4 conformation, free of protein, substantially free of  
10 other organic contaminants, substantially free of inorganic contaminants, and having a molecular weight of about 800,000 daltons to about 30 million daltons comprising:

- 15 (a) subjecting a microalgae comprising a cell body and a poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber to a mechanical force for a time sufficient to separate the cell body from the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber;
- 20 (b) segregating the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber from the cell body; and
- (c) removing all protein, substantially all other organic contaminants, and substantially all inorganic contaminants from the segregated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber,
- 25 so that the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species is isolated.

29. The method of Claim 28 wherein the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species isolated  
30 comprises about 4,000 to about 15,000 N-acetylglucosamine monosaccharides and has a molecular weight of about 800,000 daltons to about 3 million daltons.

30. The method of Claim 28 wherein the mechanical force is a shear force or a cutting force.

31. The method of Claim 28 wherein the  
5 microalgae is a diatom.

32. The method of Claim 31 wherein the diatom is of the genus Thalassiosira fluviatilis.

10 33. The method of Claim 32 wherein the diatom of the genus Thalassiosira is Thalassiosira fluviatilis or Thalassiosira weissflogii.

34. A method for isolating a poly- $\beta$ -1 $\rightarrow$ 4-N-  
15 acetylglucosamine (p-GlcNAc) species comprising about 4,000 to about 150,000 N-acetylglucosamine monosaccharides covalently attached in a  $\beta$ -1 $\rightarrow$ 4 conformation, free of protein, substantially free of other organic contaminants, substantially free of  
20 inorganic contaminants, and having a molecular weight of about 800,000 daltons to about 30 million daltons comprising:

(a) treating a microalgae comprising a cell body and a poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber with a  
25 chemical capable of weakening a diatom cell wall for a sufficient time so that the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber is released from the cell body;

(b) segregating the poly- $\beta$ -1 $\rightarrow$ 4-N-  
30 acetylglucosamine fiber from the cell body; and

(c) removing all protein, substantially all other organic contaminants, and substantially all inorganic contaminants from the segregated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber,

35

so that the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species is isolated.

35. The method of Claim 34 wherein the poly- $\beta$ -  
5 1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species isolated comprises about 4,000 to about 15,000 N-acetylglucosamine monosaccharides and has a molecular weight of about 800,000 daltons to about 3 million daltons.

10

36. The method of Claim 34 wherein the chemical capable of weakening a microalgae cell wall is hydrofluoric acid.

15 37. The method of Claim 34 further comprising neutralizing the segregated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber prior to step (c).

20 38. The method of Claim 37 further comprising, after step (c):

(d) placing the isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species in a basic pH environment.

25 39. The method of Claim 38 further comprising:  
(e) adding a drug to the isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species so that the drug and the species are in the basic pH environment; and  
(f) lowering the pH of the basic pH environment,  
30 so that the drug is encapsulated within the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species.

40. The method of Claim 34 wherein the microalgae is a diatom.

35



41. The method of Claim 40 wherein the diatom is of the genus Thallasiosira.

42. The method of Claim 41 wherein the diatom of the genus Thallasiosira is Thallasiosira fluviatilis or Thallasiosira weissflogii.

43. A method for isolating a poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species comprising about 4,000 to about 150,000 N-acetylglucosamine monosaccharides covalently attached in a  $\beta$ -1 $\rightarrow$ 4 conformation, free of protein, substantially free of other organic contaminants, substantially free of inorganic contaminants, and having a molecular weight of about 800,000 daltons to about 30 million daltons comprising:

- (a) treating a microalgae comprising a cell body and a poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber with a biological agent capable of poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber synthesis for a sufficient time so that the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber is released from the cell body;
  - (b) segregating the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber from the cell body; and
  - (c) removing all protein, substantially all other organic contaminants, and substantially all inorganic contaminants from the segregated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine fiber,
- so that the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species is isolated.

44. The method of Claim 43 wherein the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species isolated comprises about 4,000 to about 15,000 N-

acetylglucosamine monosaccharides and has a molecular weight of about 800,000 daltons to about 3 million daltons.

5           45. The method of Claim 43 wherein the biological agent is polyoxin-D.

          46. The method of Claim 43 wherein the microalgae is a diatom.

10

          47. The method of Claim 46 wherein the diatom is of the genus Thalassiosira.

          48. The method of Claim 47 wherein the diatom of  
15 the genus Thalassiosira is Thalassiosira fluviatilis or Thalassiosira weissflogii.

          49. A poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species isolated by the method of Claim 28, 29, 34,  
20 35, 43, or 44.

          50. A drug/poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine encapsulation made by the method of Claim 39.

25           51. An isolated drug/poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine encapsulation comprising a drug encapsulated by the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 1.

30           52. The isolated drug/poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine encapsulation of Claim 51 wherein the drug encapsulated is an anti-tumor drug.

          53. The isolated drug/poly- $\beta$ -1 $\rightarrow$ 4-N-  
35 acetylglucosamine encapsulation of Claim 52 wherein

the anti-tumor drug is 5'-fluorouracil, mitomycin, cis-platin, taxol, adriamycin, actinomycin, a bleomycin, a daunomycin or a methamycin anti-tumor drug.

5

54. The isolated drug/poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine encapsulation of Claim 51 wherein the drug encapsulated is an antibiotic, an anti-inflammatory, an antifungal, an antiprotozoal or a spermicidal drug.

10

55. An isolated drug/poly- $\beta$ -1 $\rightarrow$ 4-glucosamine encapsulation comprising a drug encapsulated by the poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 20.

15

56. The isolated drug/poly- $\beta$ -1 $\rightarrow$ 4-N-glucosamine encapsulation of Claim 55 wherein the drug encapsulated is an anti-tumor drug.

20

57. The isolated drug/poly- $\beta$ -1 $\rightarrow$ 4-N-glucosamine encapsulation of Claim 56 wherein the anti-tumor drug is 5'-fluorouracil, mitomycin, cis-platin, taxol, adriamycin, actinomycin, a bleomycin, a daunomycin or a methamycin anti-tumor drug.

25

58. The isolated drug/poly- $\beta$ -1 $\rightarrow$ 4-N-glucosamine encapsulation of Claim 55 wherein the drug encapsulated is an antibiotic, an anti-inflammatory, an antifungal, an antiprotozoal or a spermicidal drug.

30

59. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 1 or 2 crosslinked to a collagen species.

35

60. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 6 crosslinked to a collagen species.

5 61. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 11 crosslinked to a collagen species.

10 62. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 12 crosslinked to a collagen species.

15 63. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 13 crosslinked to a collagen species.

20 64. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 18 or 19 crosslinked to a collagen species.

25 65. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 20 crosslinked to a collagen species.

30 66. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 25 crosslinked to a collagen species.

35 67. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 26 crosslinked to a collagen species.

68. An isolated hybrid composition comprising the poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 27 crosslinked to a collagen species.

69. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 6 wherein at least one peptide is functionally attached to a deacetylated monosaccharide.

5

70. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 69 wherein the peptide is covalently attached to a deacetylated monosaccharide.

10 71. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 69 wherein the peptide is non-covalently attached to a deacetylated monosaccharide.

15 72. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 69 wherein the peptide is a growth factor, hormone, peptide recognition sequence, laminin, integrin, or cell adhesion molecule.

20 73. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 18 or 19 wherein at least one peptide is functionally attached to a glucosamine monosaccharide.

25 74. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 73 wherein the peptide is covalently attached to a glucosamine monosaccharide.

30 75. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 73 wherein the peptide is non-covalently attached to a glucosamine monosaccharide.

35 76. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 73 wherein the peptide is a growth factor, hormone, peptide recognition sequence, laminin, integrin, or cell adhesion molecule.

77. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 20 wherein at least one peptide is functionally attached to a glucosamine monosaccharide.

5 78. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 77 wherein the peptide is covalently attached to a glucosamine monosaccharide.

10 79. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 77 wherein the peptide is non-covalently attached to a glucosamine monosaccharide.

15 80. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of claim 77 wherein the peptide is a growth factor, hormone, peptide recognition sequence, laminin, integrin, or cell adhesion molecule.

20 81. An isolated cell/poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine encapsulation species comprising a cell encapsulated by the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine species of Claim 1.

25 82. The isolated cell/poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine encapsulation species of Claim 81 wherein the species is contained within a thermoplastic capsule.

30 83. An isolated cell/poly- $\beta$ -1 $\rightarrow$ 4-glucosamine encapsulation species comprising a cell encapsulated by the poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 20.

84. The isolated cell/poly- $\beta$ -1 $\rightarrow$ 4-N-glucosamine encapsulation of Claim 83 wherein the species is contained within a thermoplastic capsule.

85. A method for controlled drug delivery comprising: administering to a patient the poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 65 to which a multiplicity of identical peptides are functionally attached so that, upon administration, the attached peptides are released as the poly- $\beta$ -1 $\rightarrow$ 4-acetylglucosamine species degrades.

86. The method of Claim 81 wherein the multiplicity of identical peptides is covalently attached to the poly- $\beta$ -1 $\rightarrow$ 4-acetylglucosamine species.

87. The method of Claim 81 wherein the multiplicity of identical peptides is non-covalently attached to the poly- $\beta$ -1 $\rightarrow$ 4-acetylglucosamine species.

88. A method for slow molecular drug delivery comprising: administering to a patient the drug/poly- $\beta$ -1 $\rightarrow$ 4-acetylglucosamine encapsulation of Claim 51 or 55 so that, upon administration, the drug is slowly released into the system of the patient as the poly- $\beta$ -1 $\rightarrow$ 4-acetylglucosamine species degrades.

89. The isolated poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine (p-GlcNAc) species of Claim 1 wherein the species is a poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine-lactate species.

90. The isolated poly- $\beta$ -1 $\rightarrow$ 4-glucosamine species of Claim 18 or 20 wherein the species is a poly- $\beta$ -1 $\rightarrow$ 4-glucosamine-lactate species.

91. A method for reduction of post surgical adhesions comprising: administering, prior to a surgical procedure, to a patient the poly- $\beta$ -1 $\rightarrow$ 4-N-acetylglucosamine-lactate species of Claim 85 so that,

after the surgical procedure, the patient exhibits a reduction in the amount of traumatized tissue relative to a patient not having received the species, thus reducing the occurrence of post surgical adhesions.

5

92. A method for reduction of post surgical adhesions comprising: administering, prior to a surgical procedure, to a patient the poly- $\beta$ -1 $\rightarrow$ 4-glucosamine-lactate species of Claim 86 so that, after  
10 the surgical procedure, the patient exhibits a reduction in the amount of traumatized tissue relative to a patient not having received the species, thus reducing the occurrence of post surgical adhesions relative to a patient not having been administered the  
15 species.

93. A method for reduction of post surgical adhesions comprising: administering, following a surgical procedure, to a patient the poly- $\beta$ -1 $\rightarrow$ 4-N-  
20 acetylglucosamine-lactate species of Claim 85 so that a physical barrier between traumatized tissue and non-traumatized tissue is produced, thus reducing the occurrence of post surgical adhesions relative to a patient not having been administered the species.  
25

94. A method for reduction of post surgical adhesions comprising: administering, following a surgical procedure, to a patient the poly- $\beta$ -1 $\rightarrow$ 4-N-  
30 acetylglucosamine-lactate species of Claim 86 so that a physical barrier between traumatized tissue and non-traumatized tissue is produced, thus reducing the occurrence of post surgical adhesions relative to a patient not having been administered the species.

35



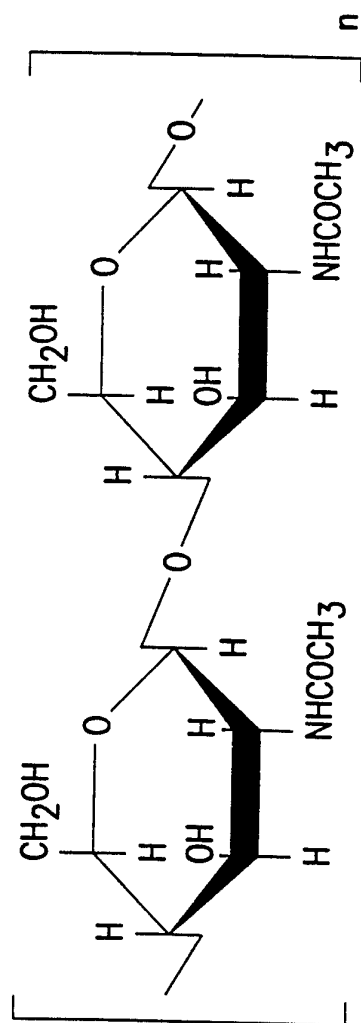


FIG.1

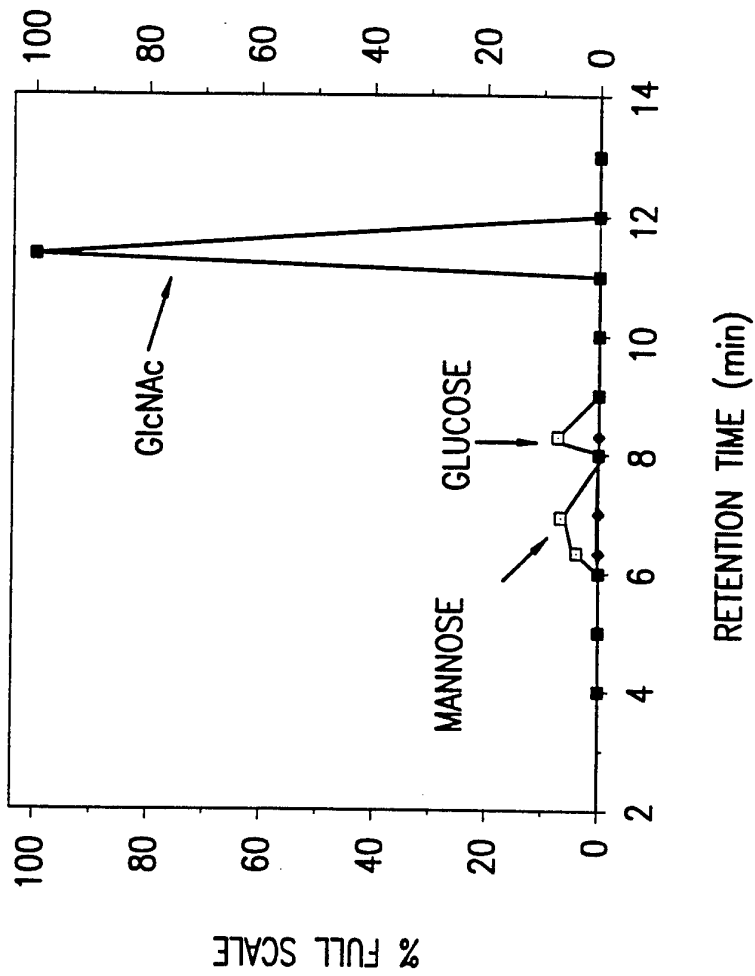


FIG.2

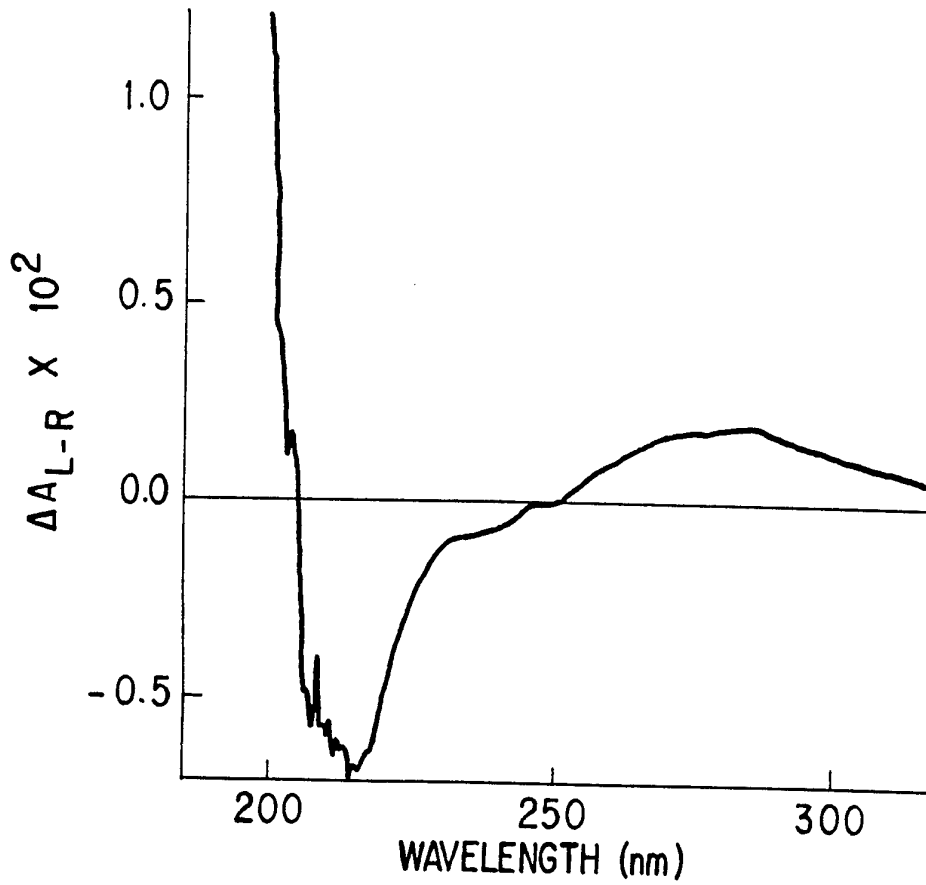


FIG. 3A

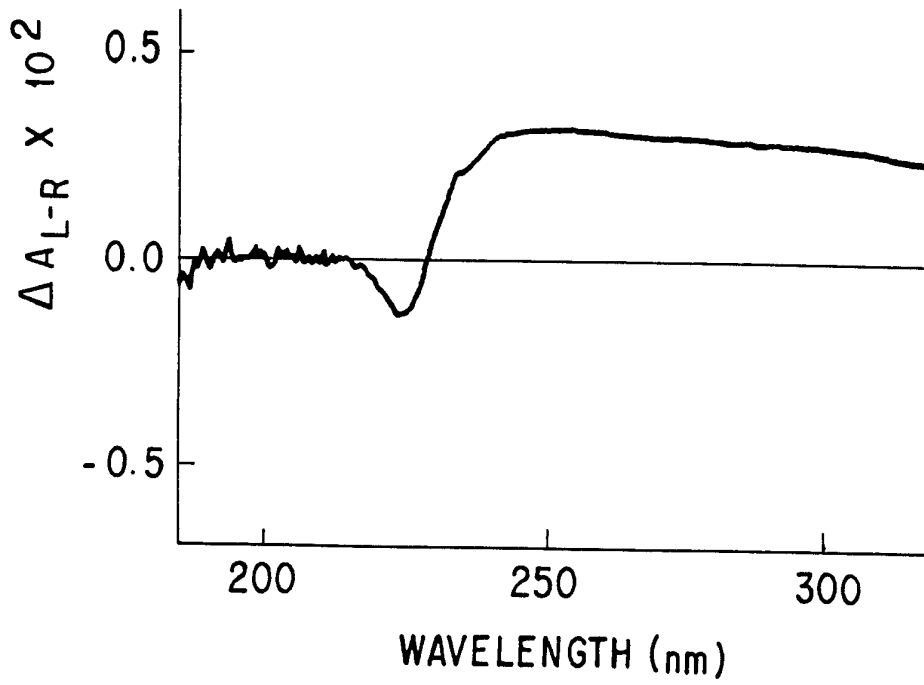


FIG. 3B

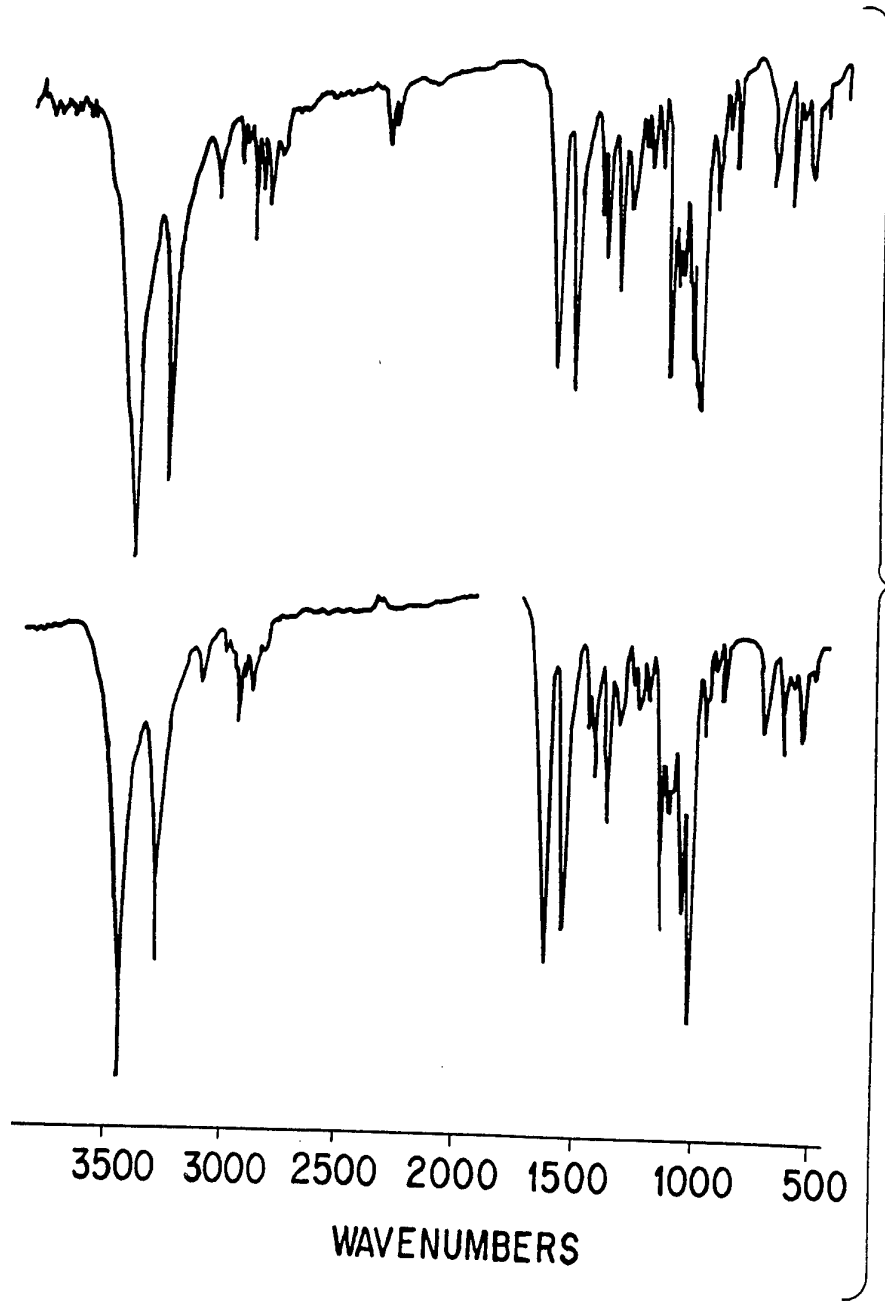


FIG. 4A

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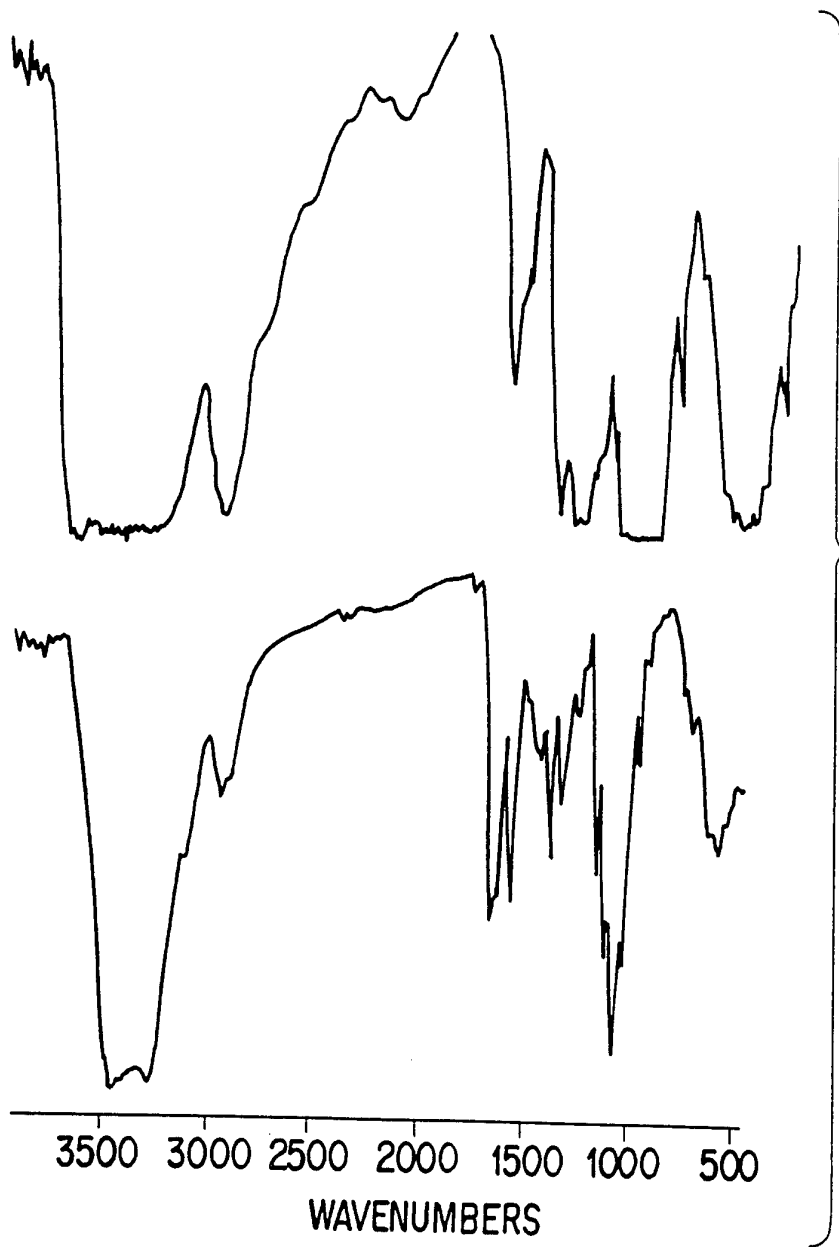


FIG. 4B

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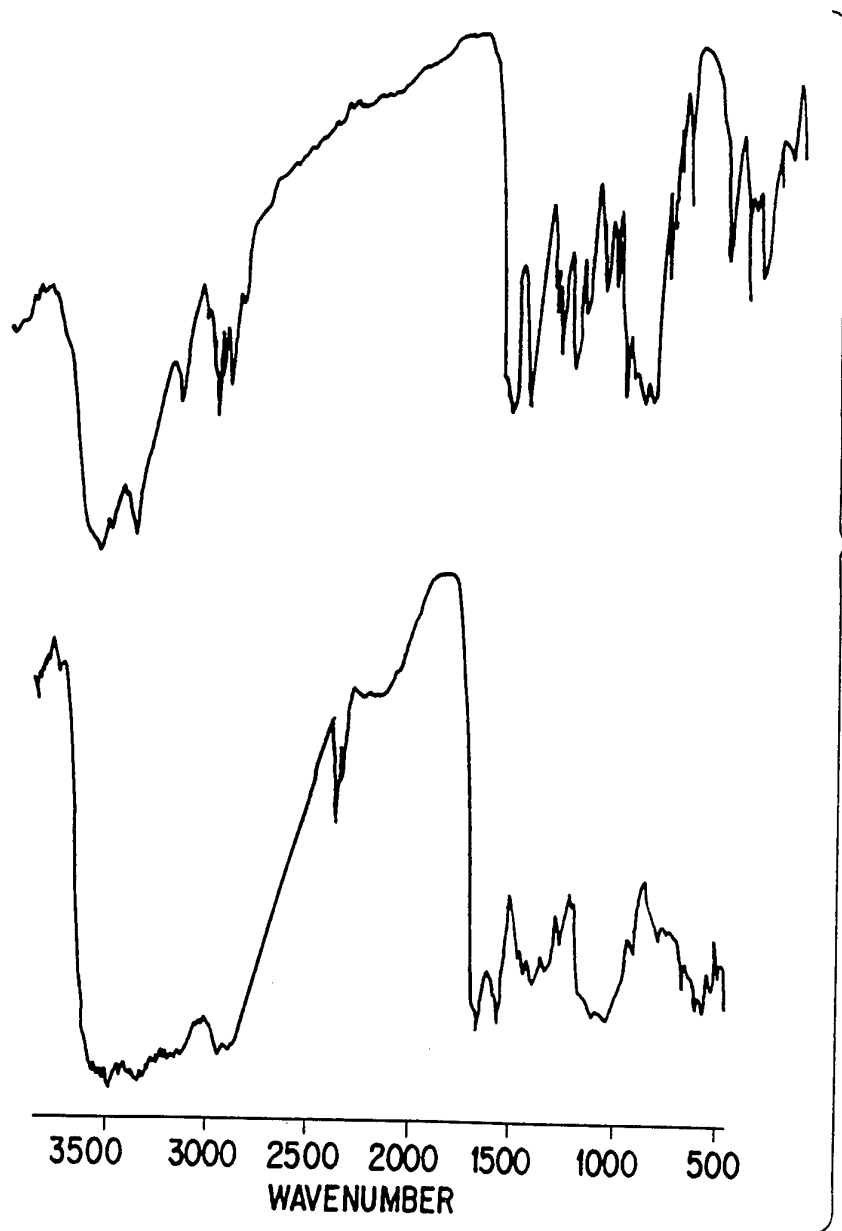


FIG. 4C

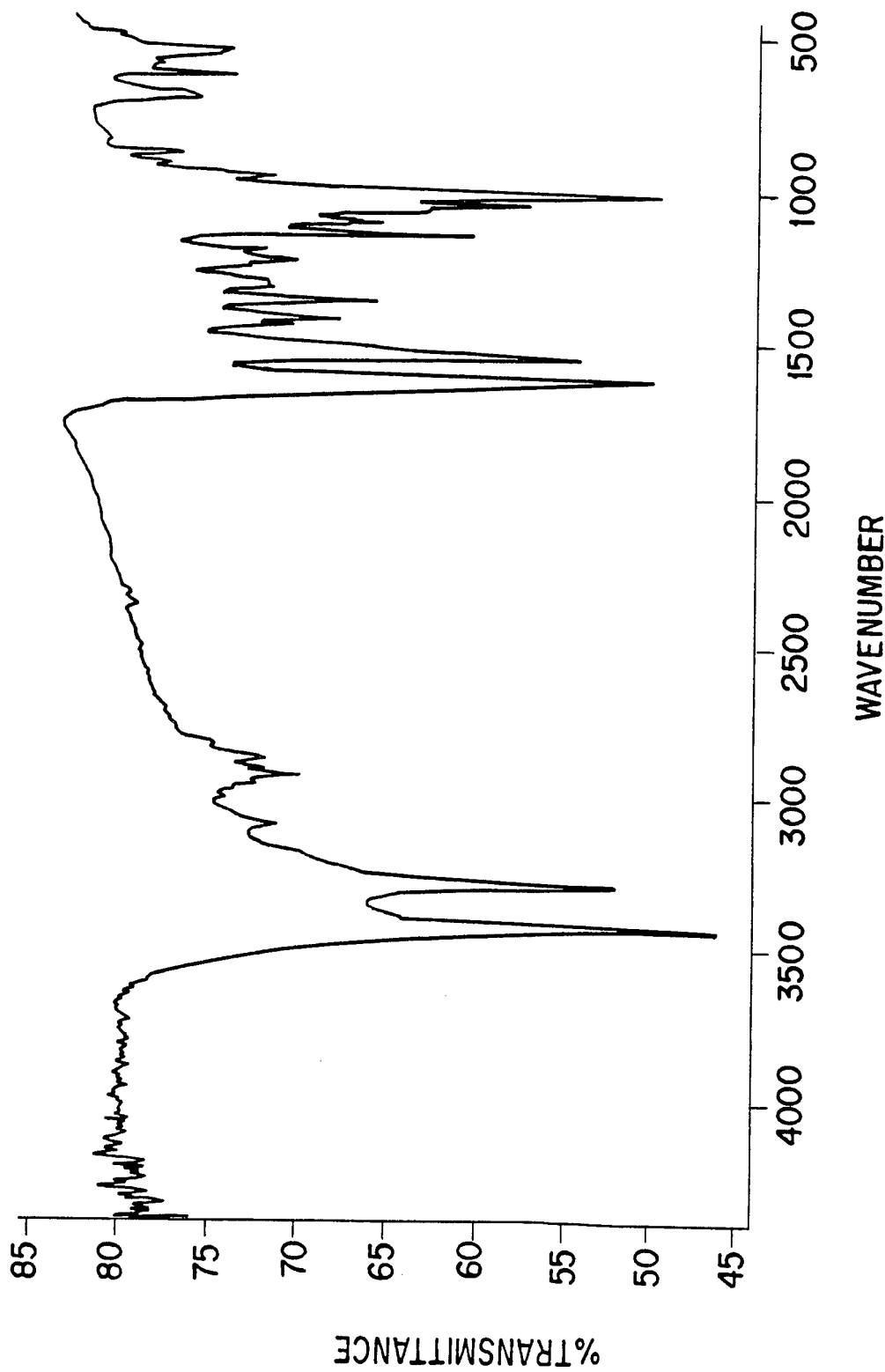


FIG. 4D



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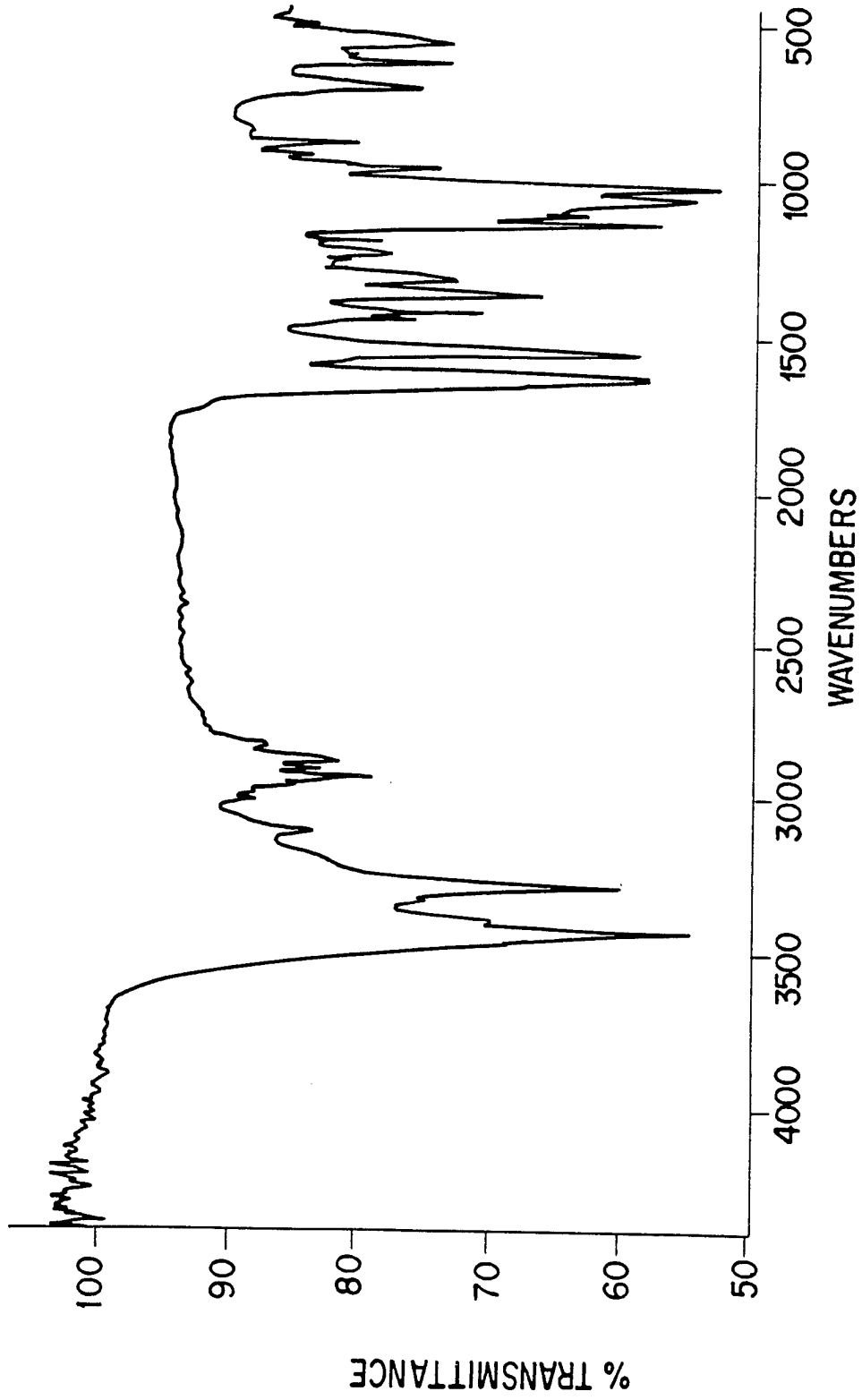


FIG. 4E

PEAK	%RF		AMPLITUDE	%SIGMA	AREA	%TOTAL		RATIO	RATIO
1	5	C=0	0.087	0.29	1.304	1.674			5.609
2	6	C=0	0.146	0.38	2.855	3.664	4.159	1.759	2.562
3	47	C1	0.563	0.38	11.153	14.314	11.153	0.656	0.656
4	59	C4	0.452	0.32	7.41	9.51	7.410	0.987	0.987
5	63	C5	0.311	0.49	7.906	10.147	7.906	0.925	0.925
6	64	C3	1.195	0.16	9.816	12.598			0.745
7	65	C3	0.533	0.4	11.11	14.259	20.926	0.350	0.658
8	72	C6	0.148	1.1	8.419	10.805			0.869
9	73	C6	0.21	0.18	1.98	2.541	10.399	0.703	3.694
10	74	C2	0.026	0.2	0.27	0.346			27.089
11	75	C2	0.227	0.72	8.38	10.755	8.650	0.846	0.873
12	94	CH3	0.377	0.38	7.314	9.387			1.000

FIG.5A

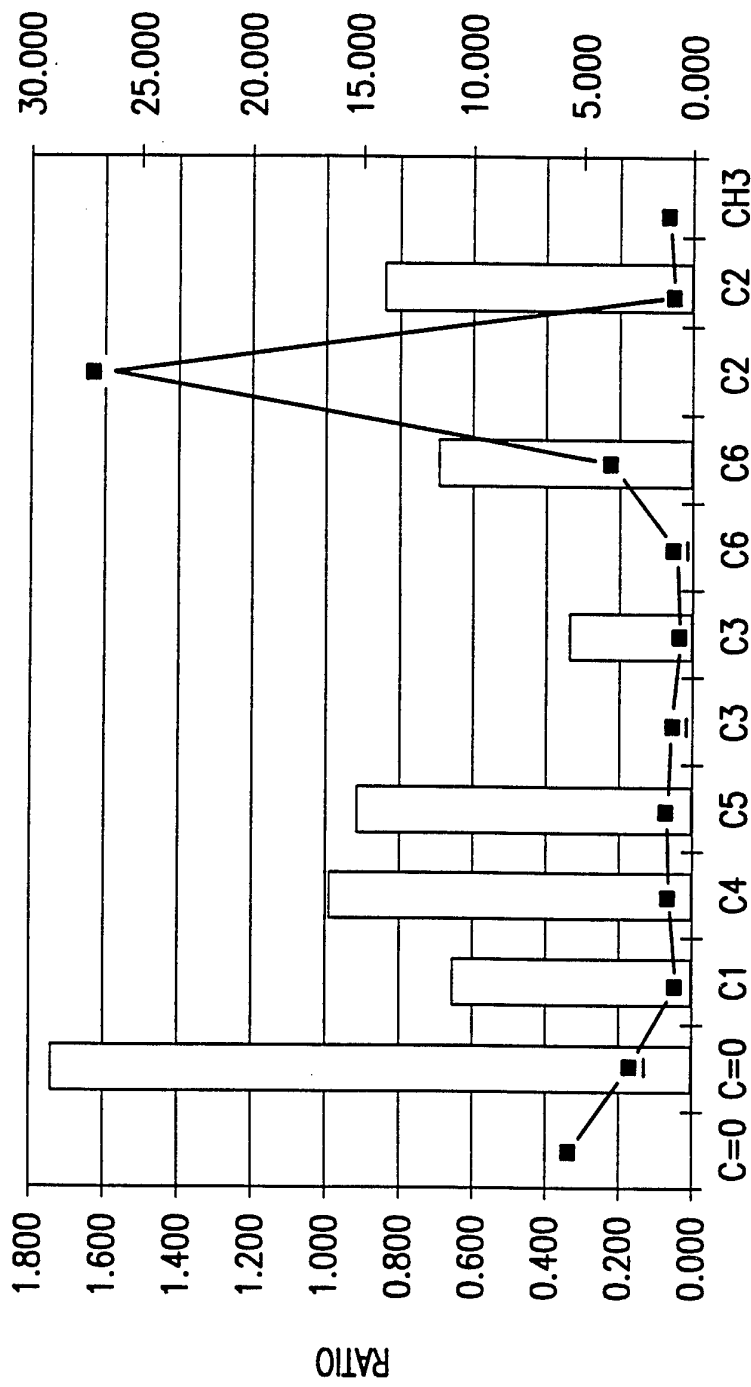


FIG.5B

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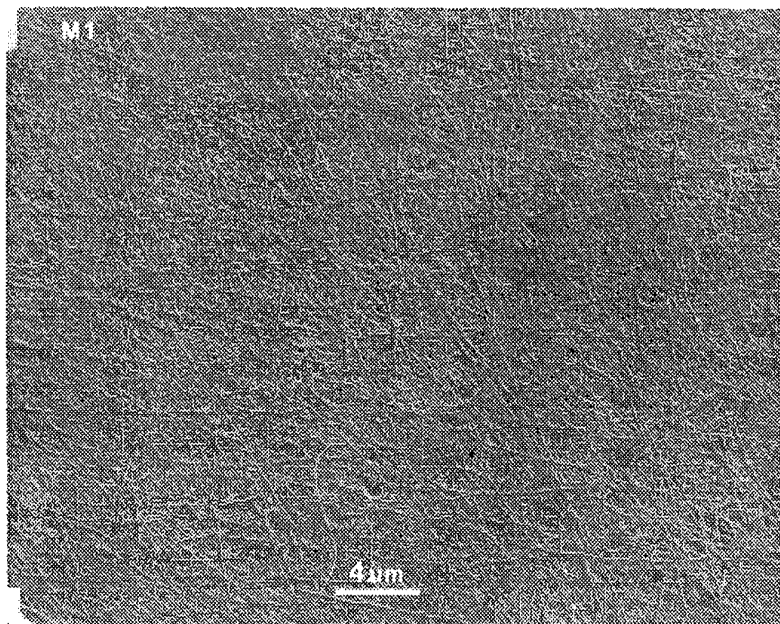


FIG.6A

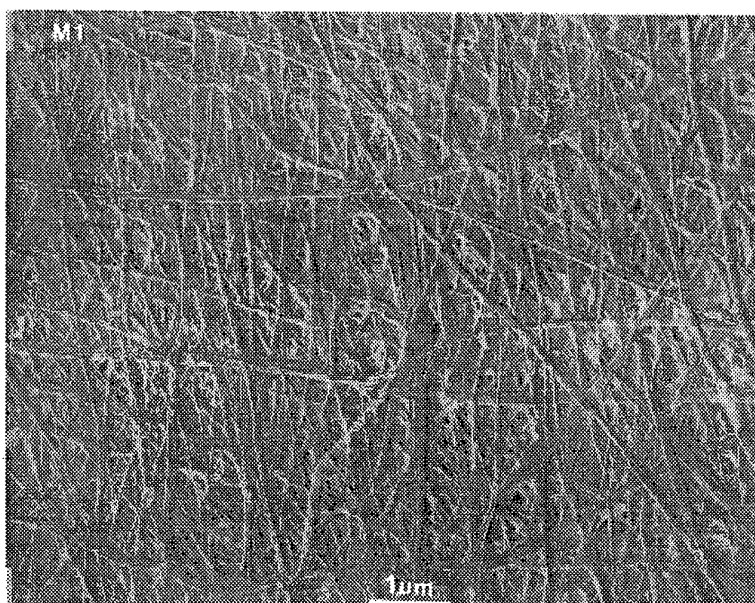


FIG.6B

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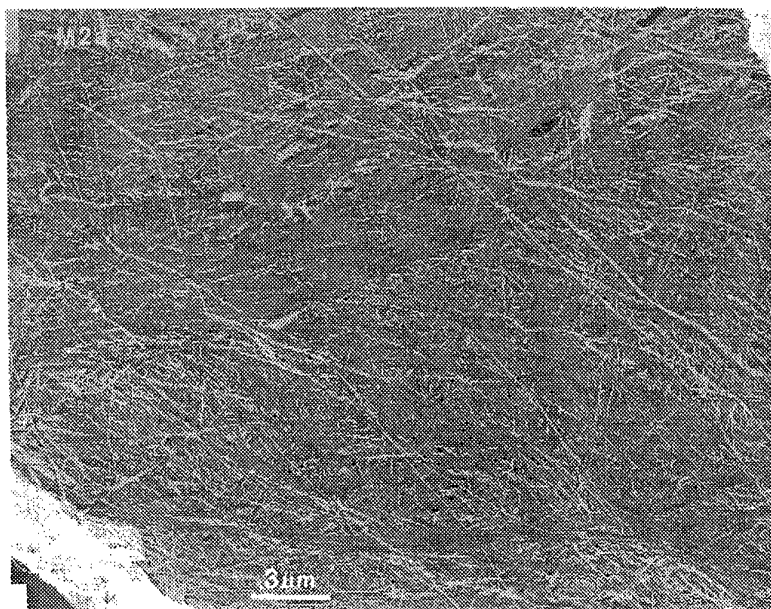


FIG.7A

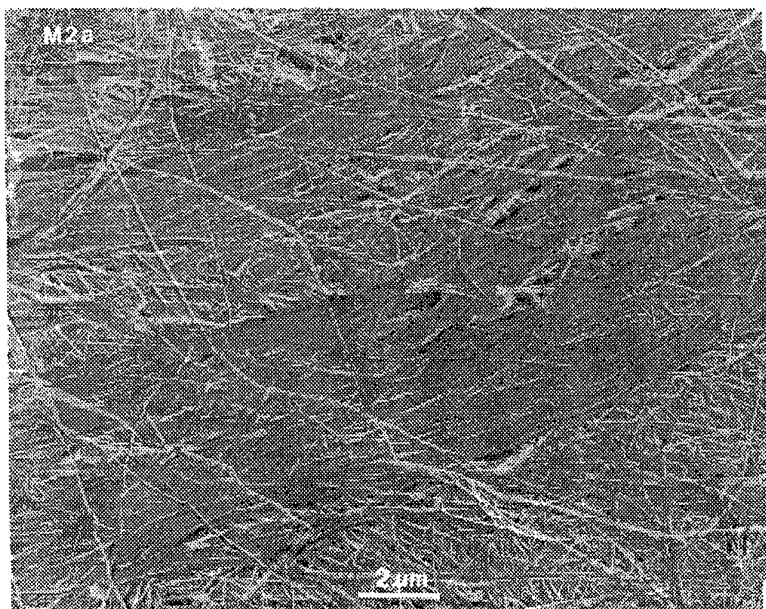


FIG.7B

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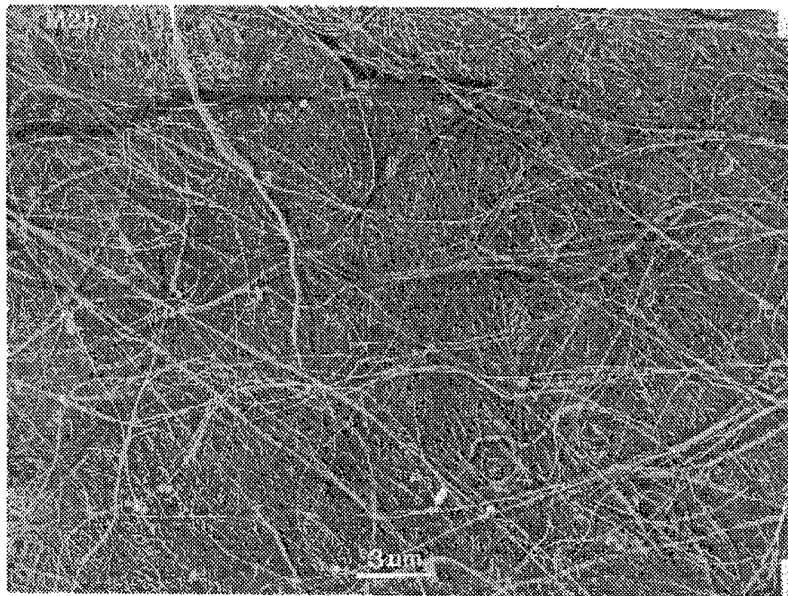


FIG.8A

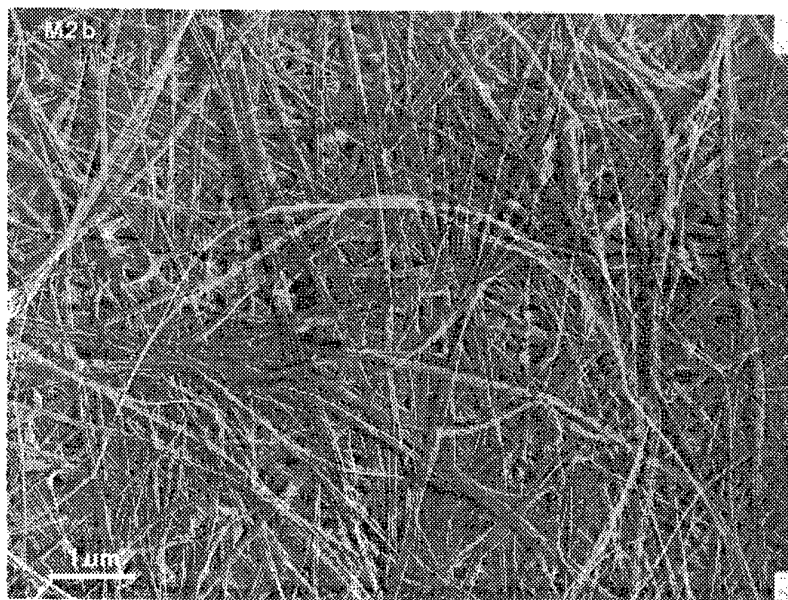


FIG.8B

RECTIFIED SHEET (RULE 91)

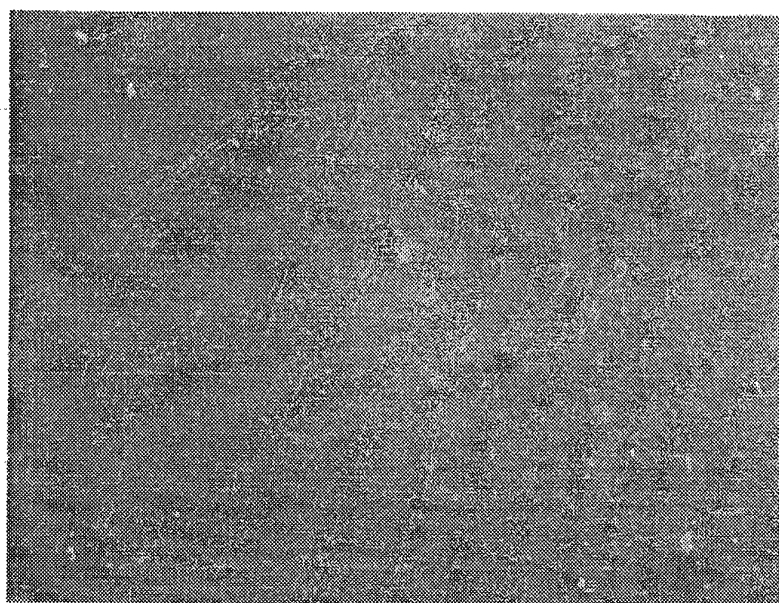
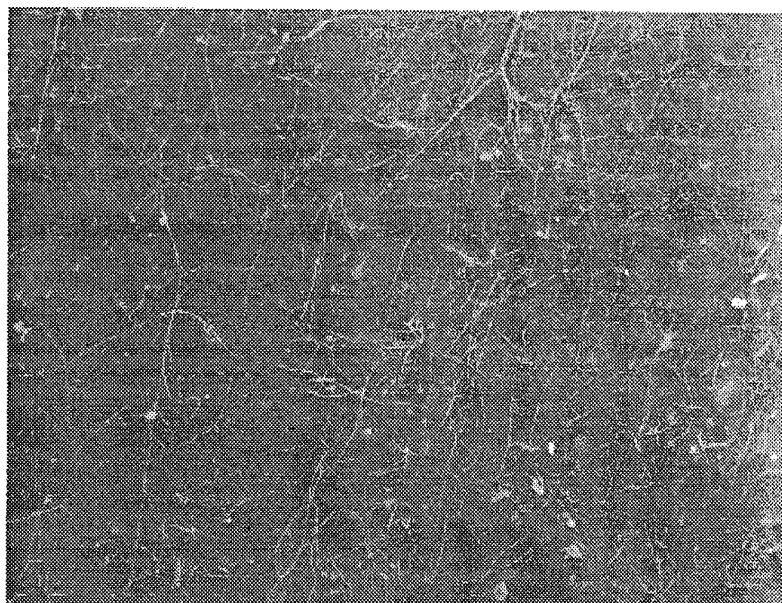


FIG.9A



**FIG.9B**



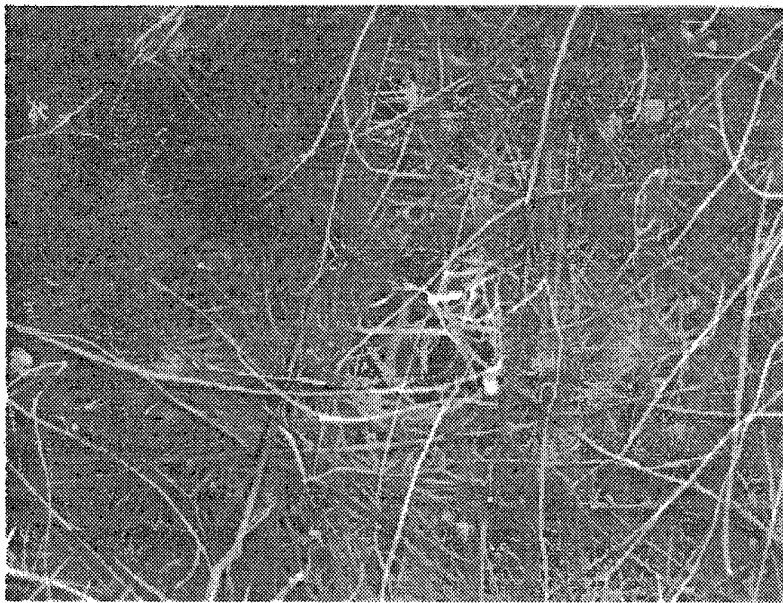


FIG.9C

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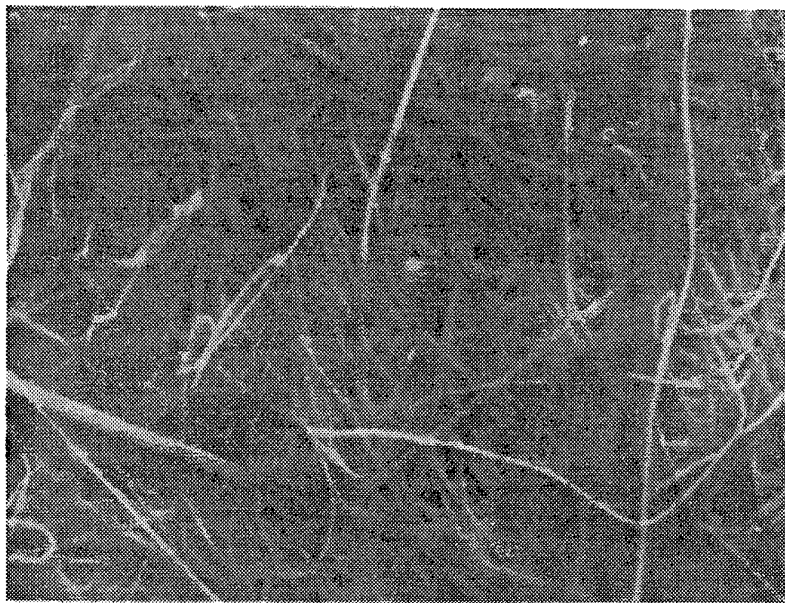


FIG.9D

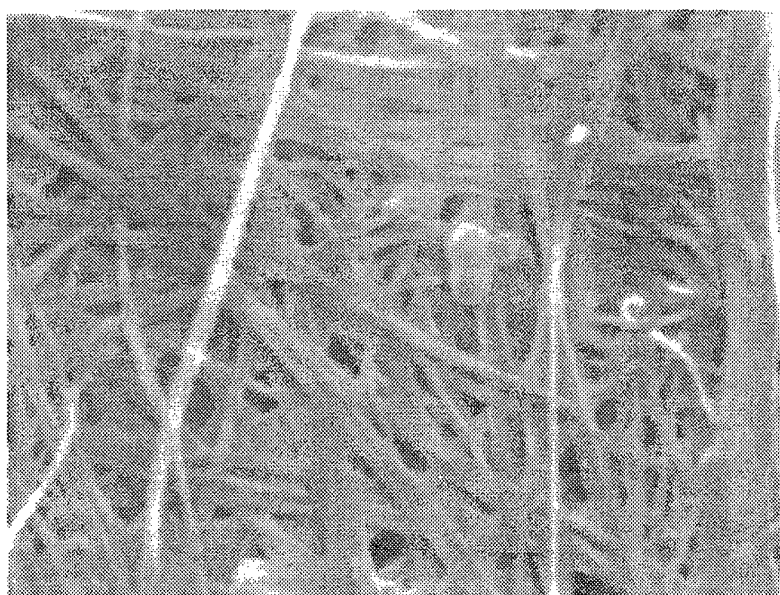


FIG.9E

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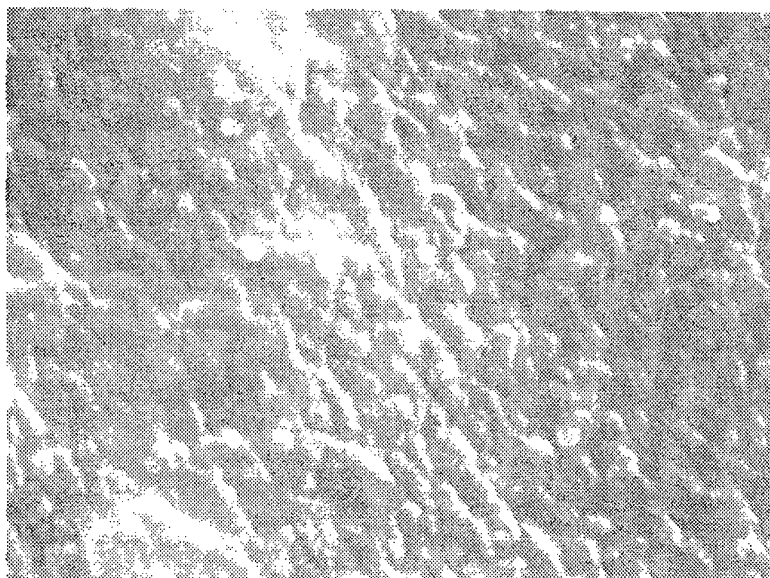


FIG. 10A

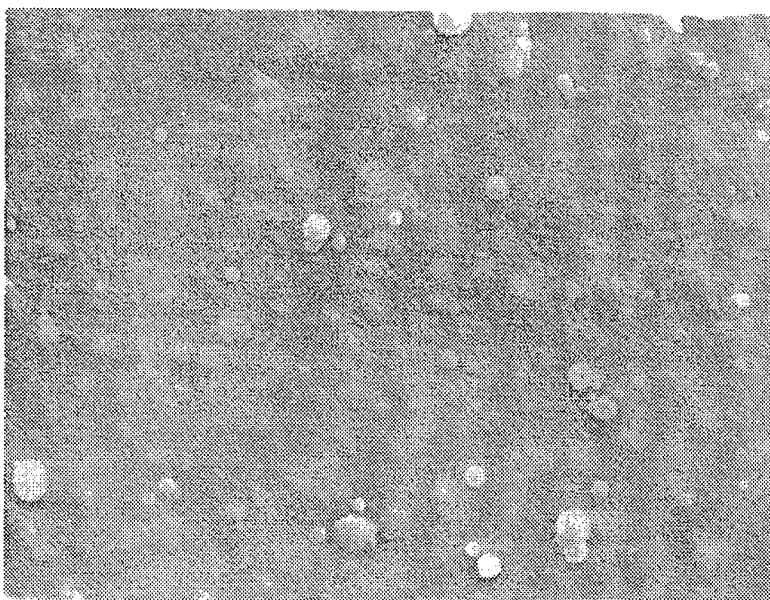


FIG. 10B

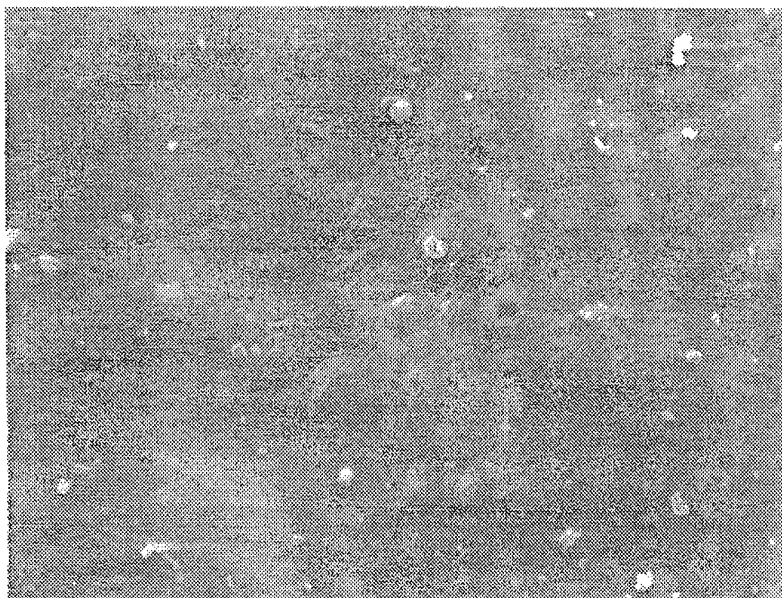


FIG. 11A

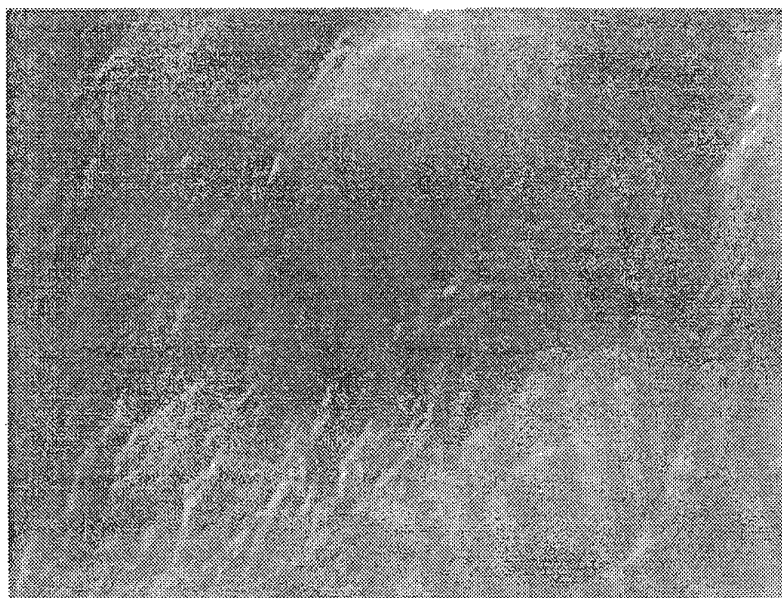


FIG. 11B

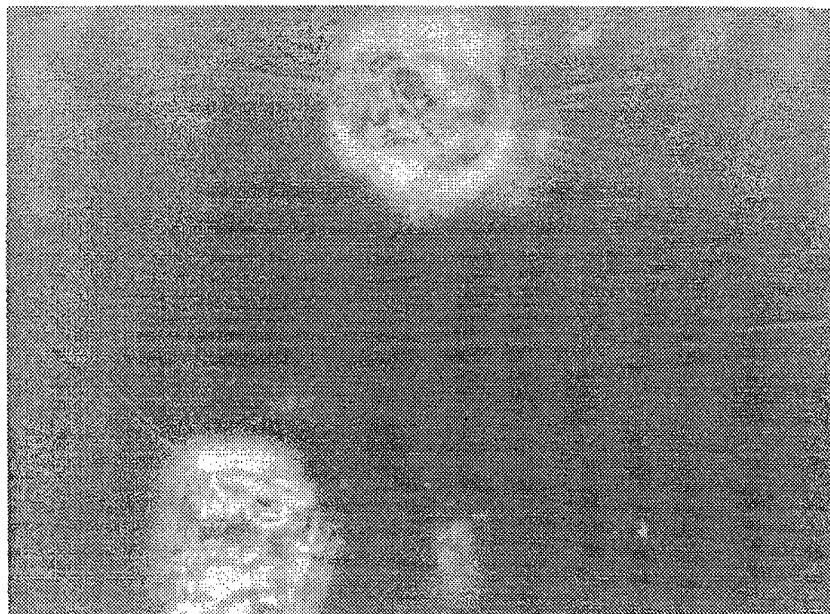


FIG. 12A

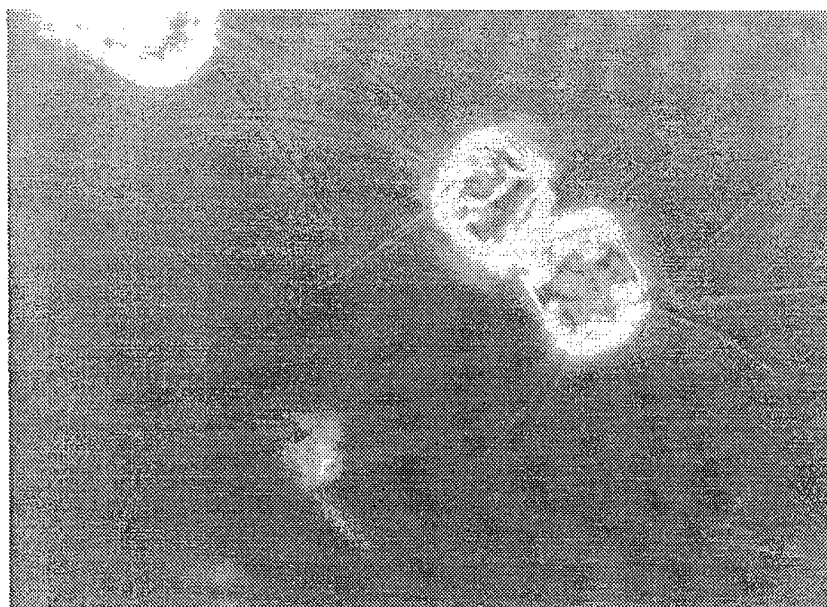


FIG. 12B

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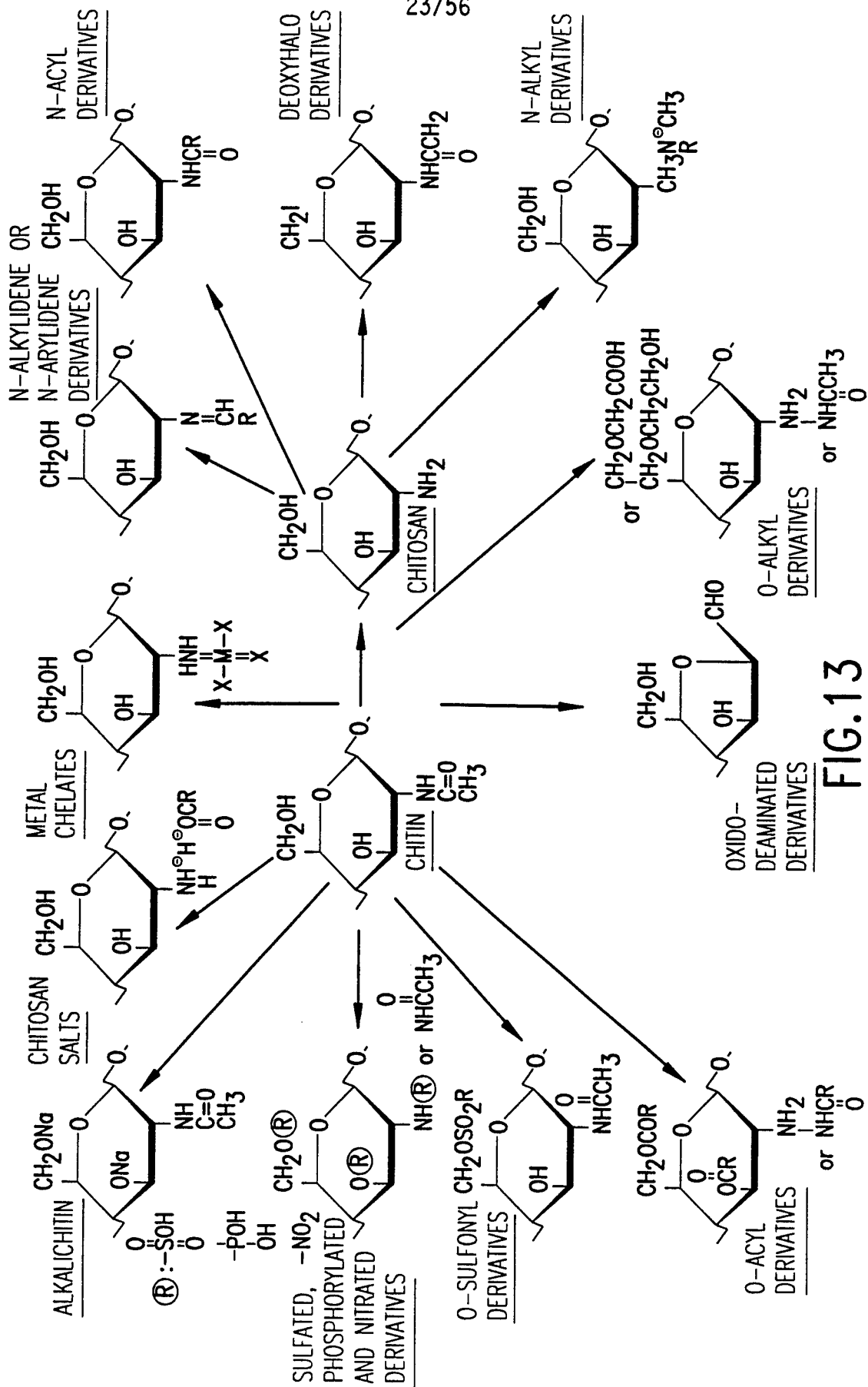


FIG.13

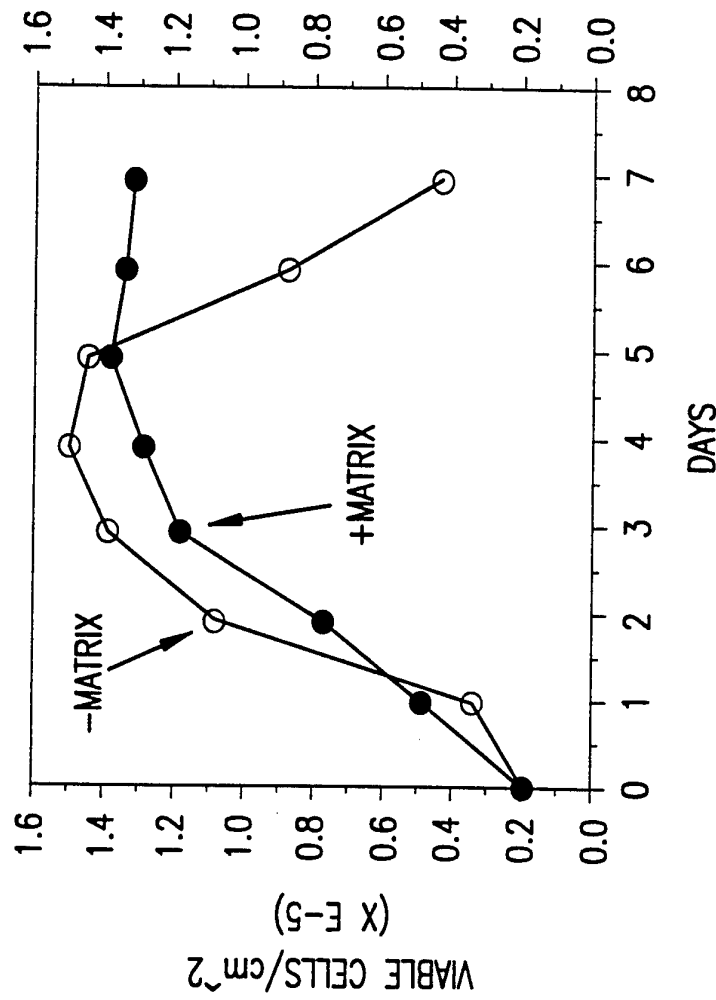


FIG.14



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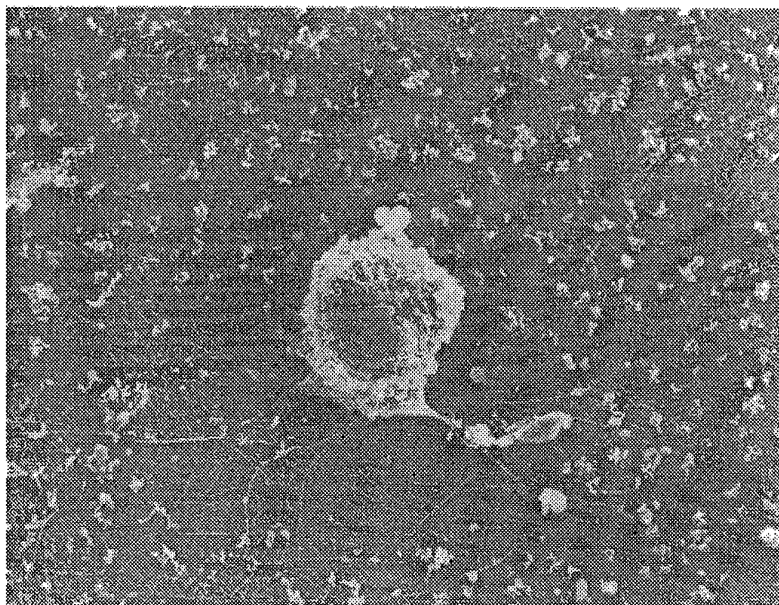


FIG. 15A

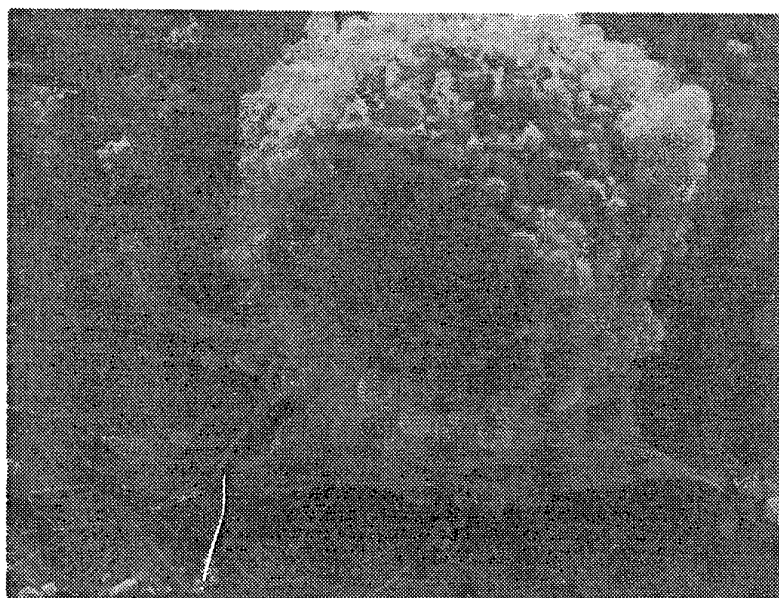


FIG. 15B

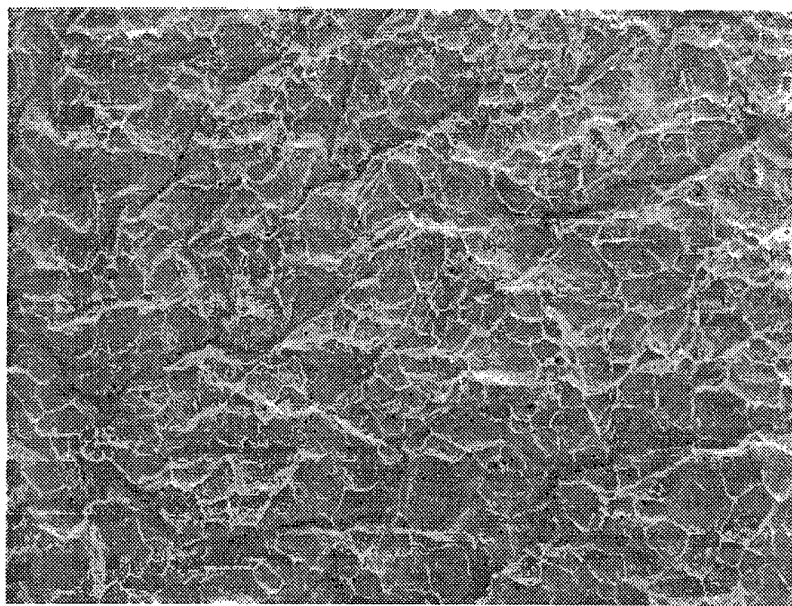


FIG.16A

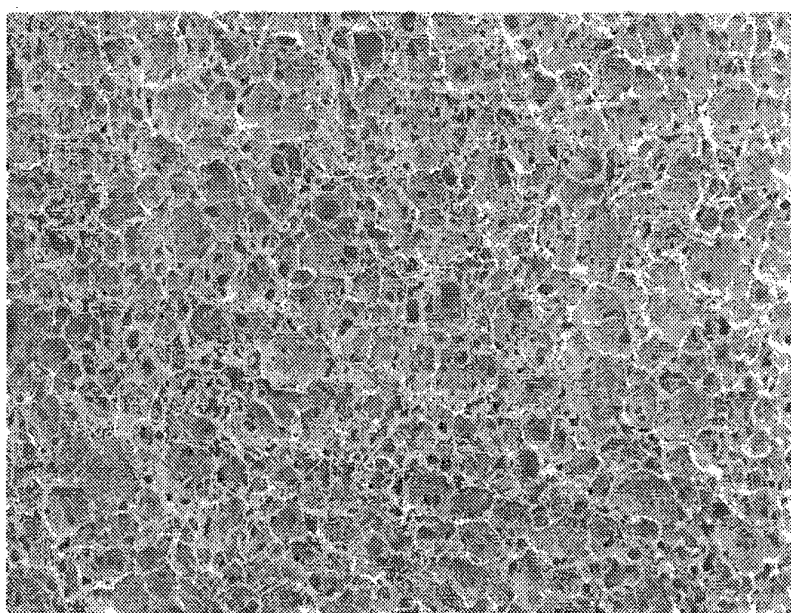


FIG. 16B

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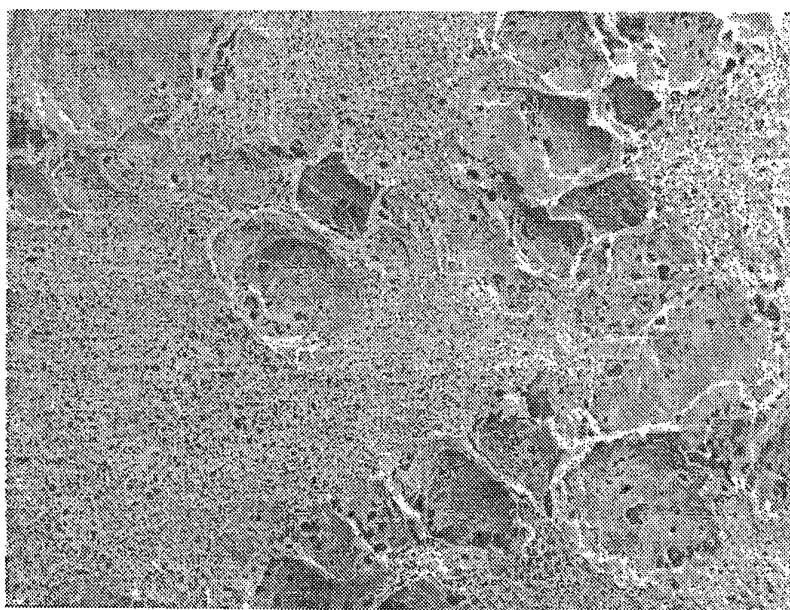


FIG.16C

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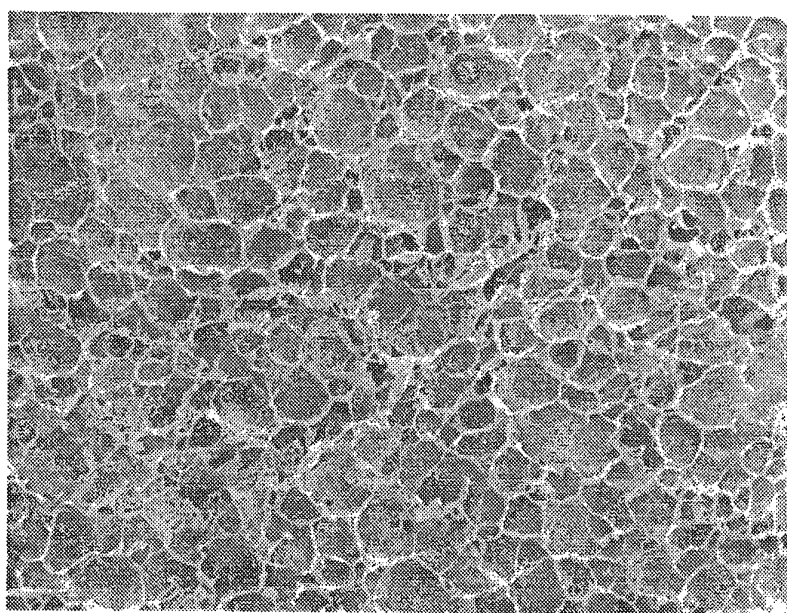


FIG. 16D

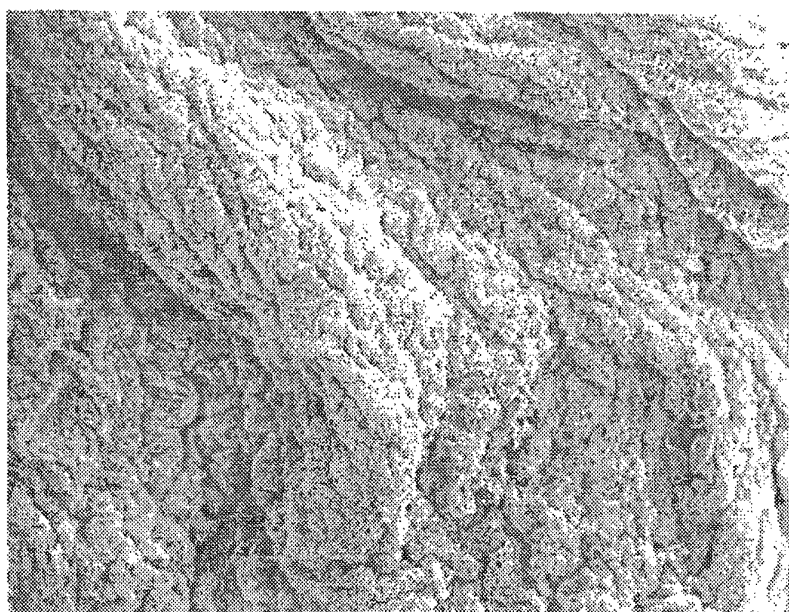


FIG. 16E

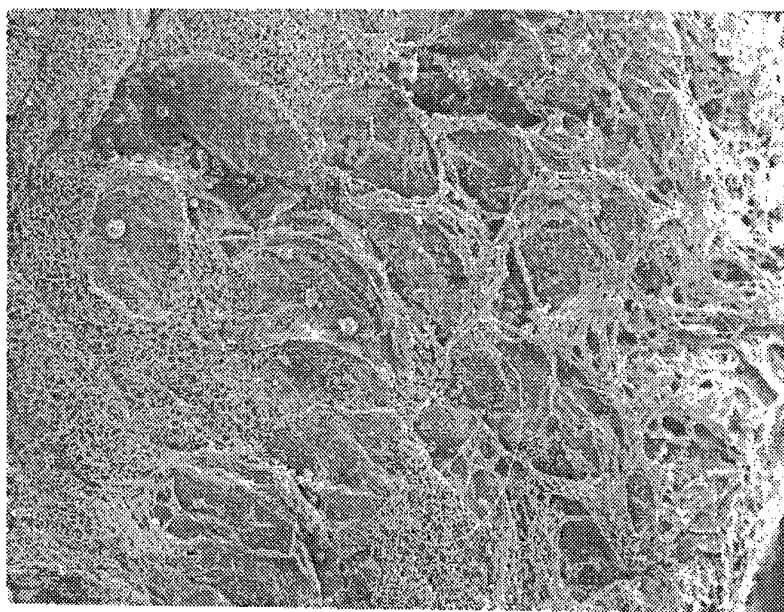


FIG.17A

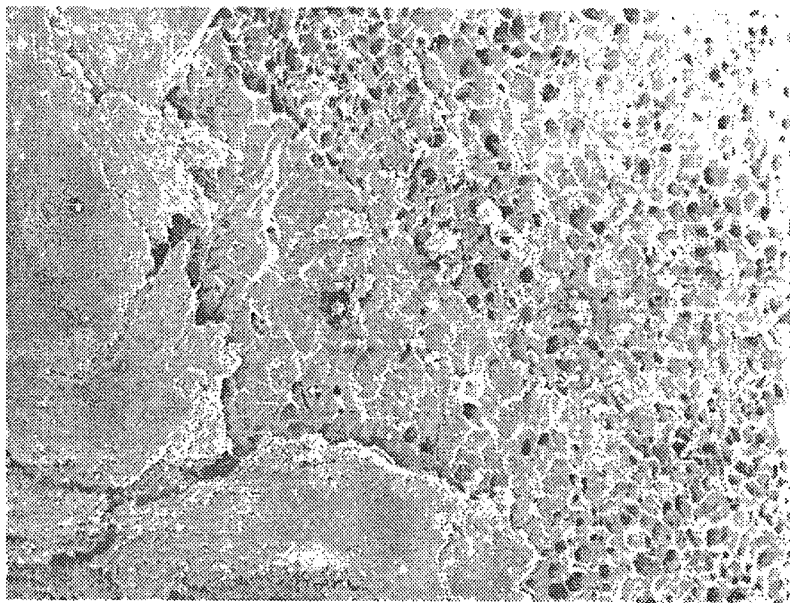


FIG.17B



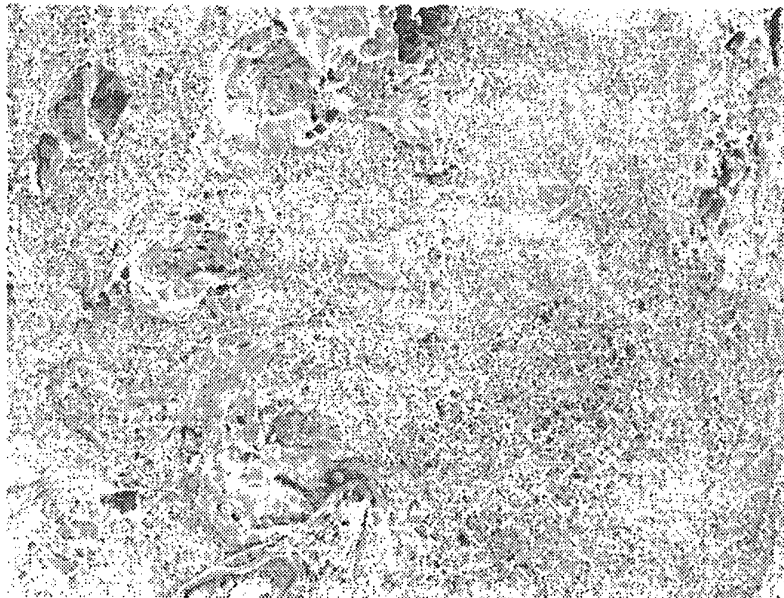


FIG. 17C



FIG.17D

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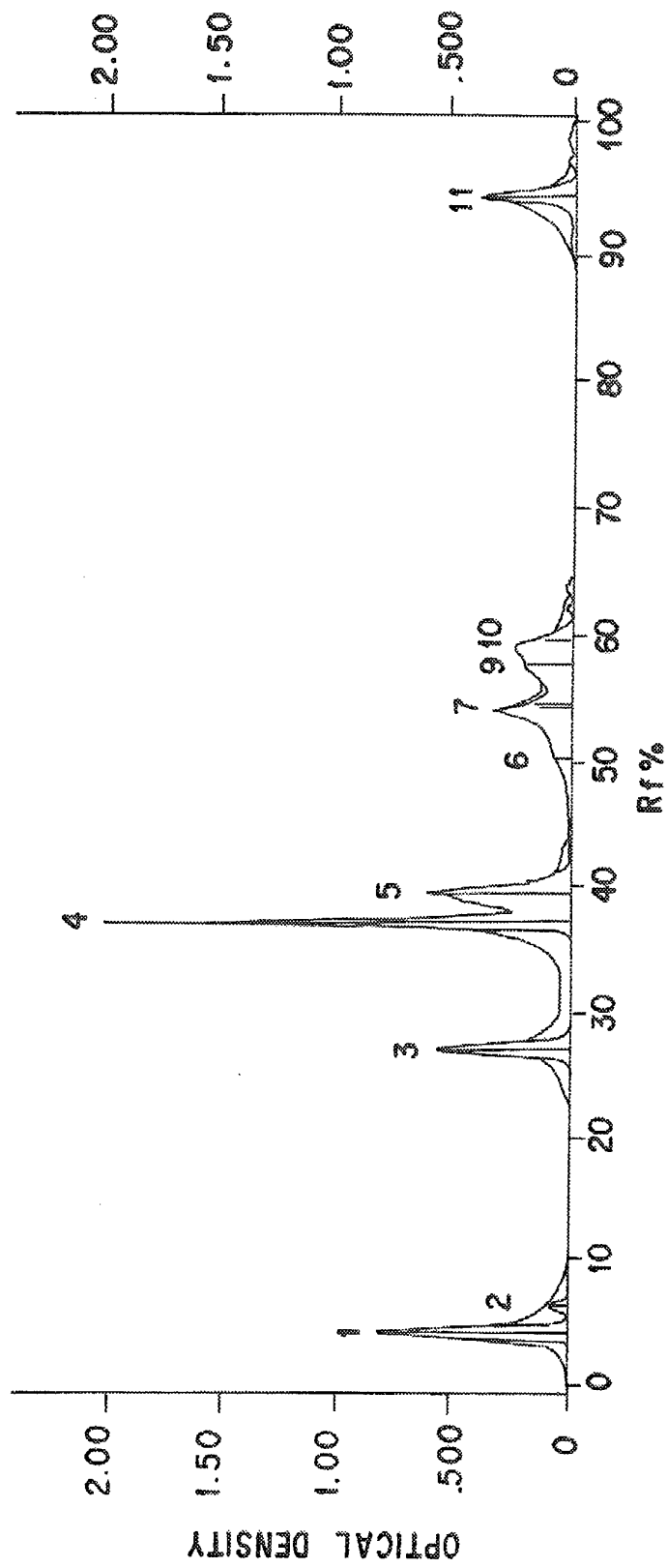


FIG. 18

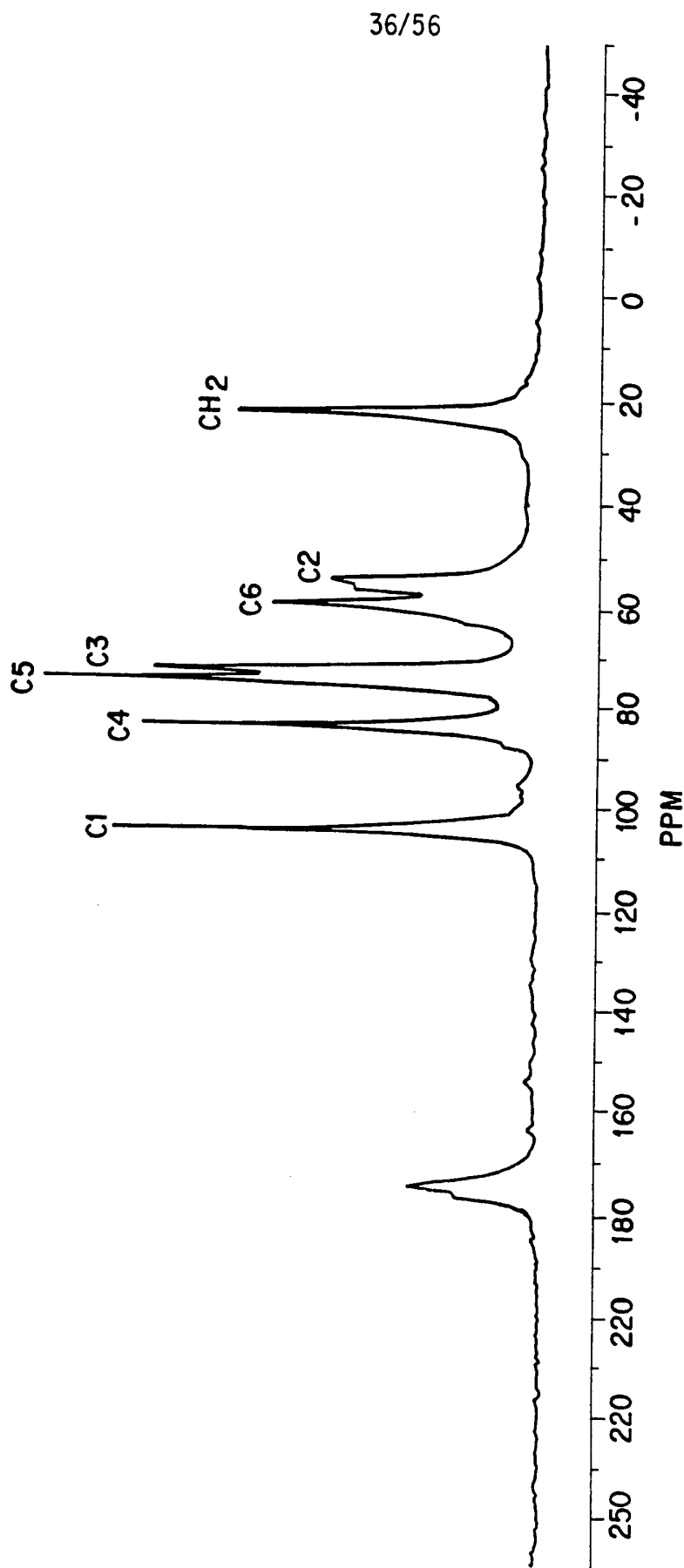
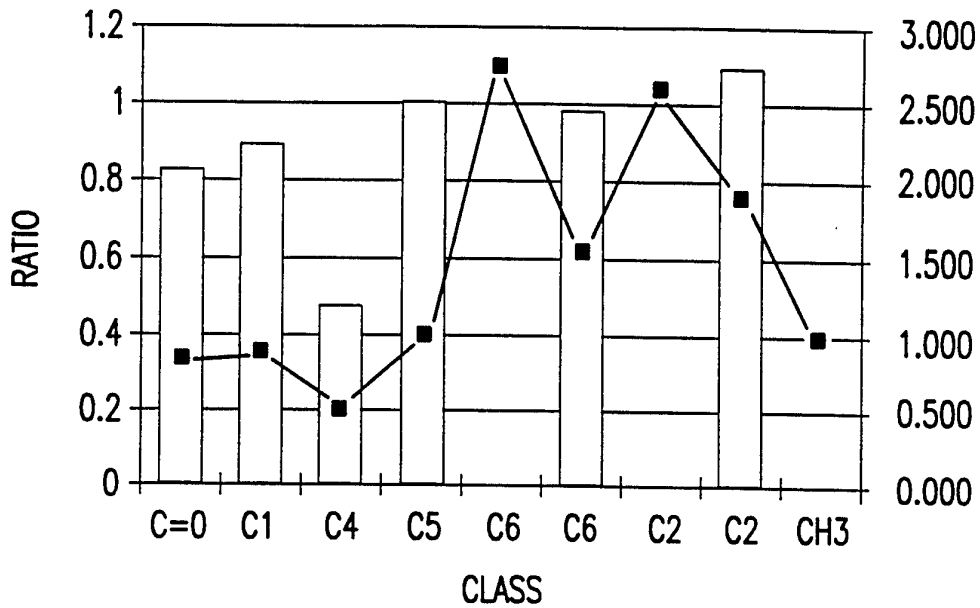


FIG. 19

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PEAK	%RF		AMPLITUDE	%SIGMA	AREA	%TOTAL		RATIO	RATIO
1	4	C=0	0.803	0.42	18.08	14.5	18.080	0.8308	0.831
2	27	C1	0.594	0.53	16.959	13.6	16.959	0.8857	0.886
3	37	C4	2.073	0.28	30.787	24.68	30.787	0.4879	0.488
4	39	C5	0.581	0.48	14.915	11.96	14.915	1.007	1.007
5	51	C6	0.096	1.06	5.504	4.413			2.729
6	54	C6	0.324	0.56	9.767	7.831	15.271	0.9836	1.538
7	57	C2	0.197	0.55	5.848	4.689			2.568
8	59	C2	0.226	0.64	7.843	6.289	13.691	1.0971	1.915
9	95	CH3	0.363	0.77	15.02	12.04	15.020		1.000

FIG.20

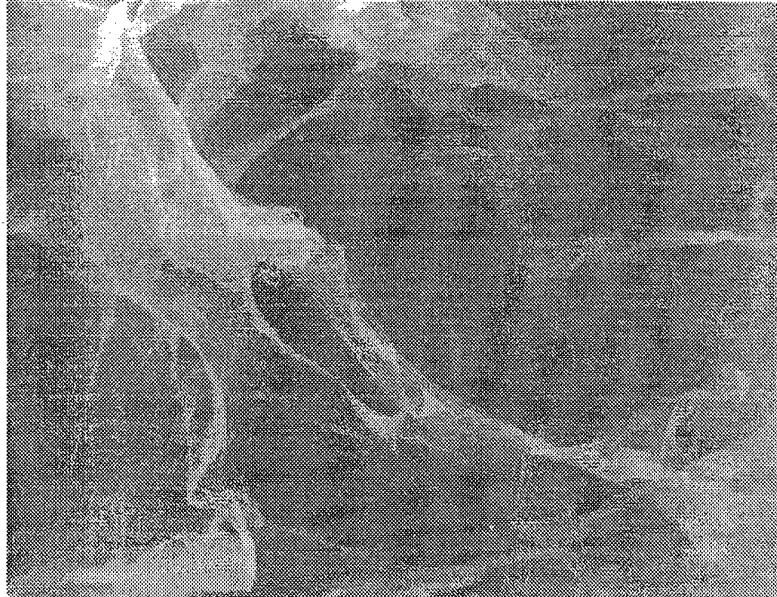


FIG.21A 200 μm

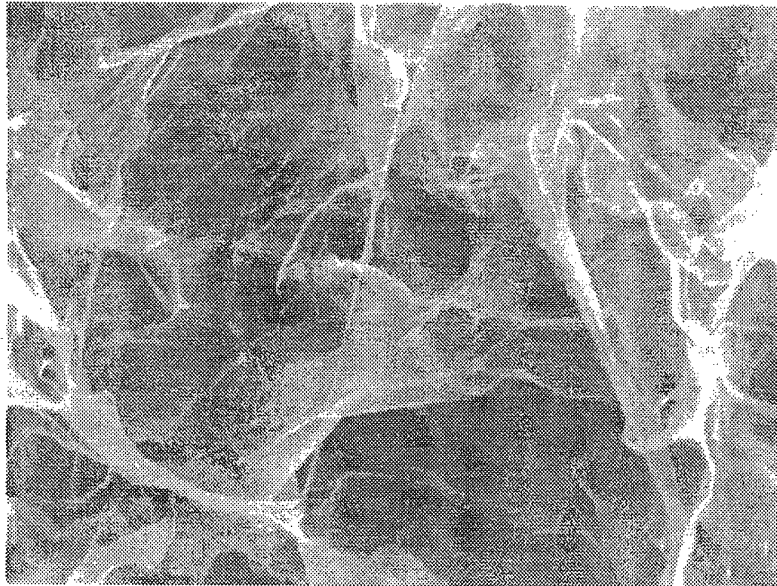
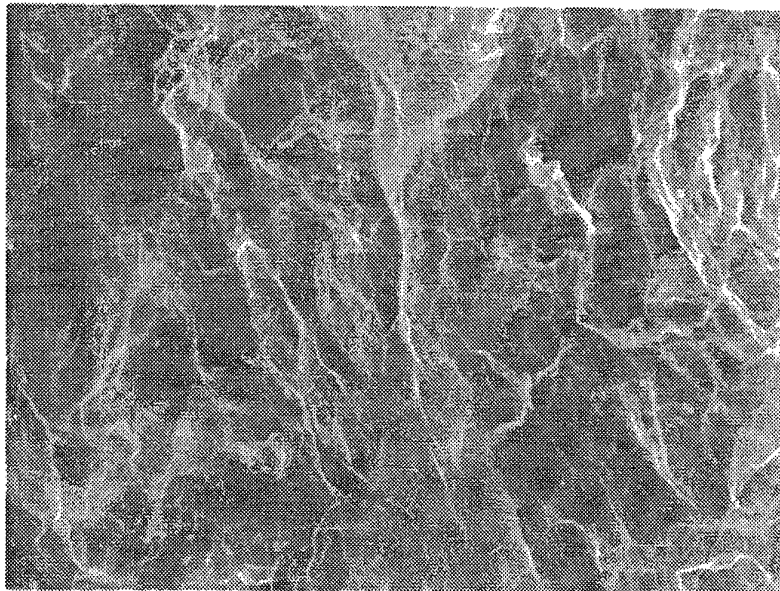


FIG.21B 200 μm

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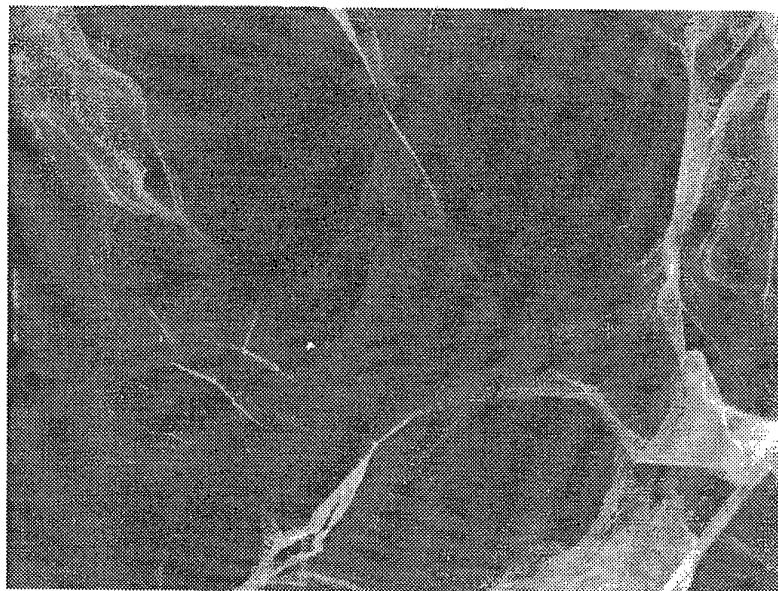
200  $\mu\text{m}$  

FIG.21C




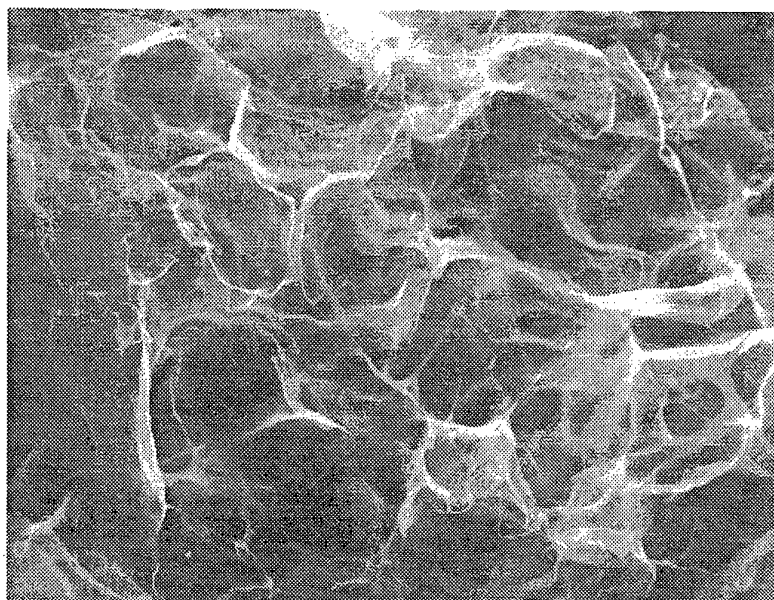
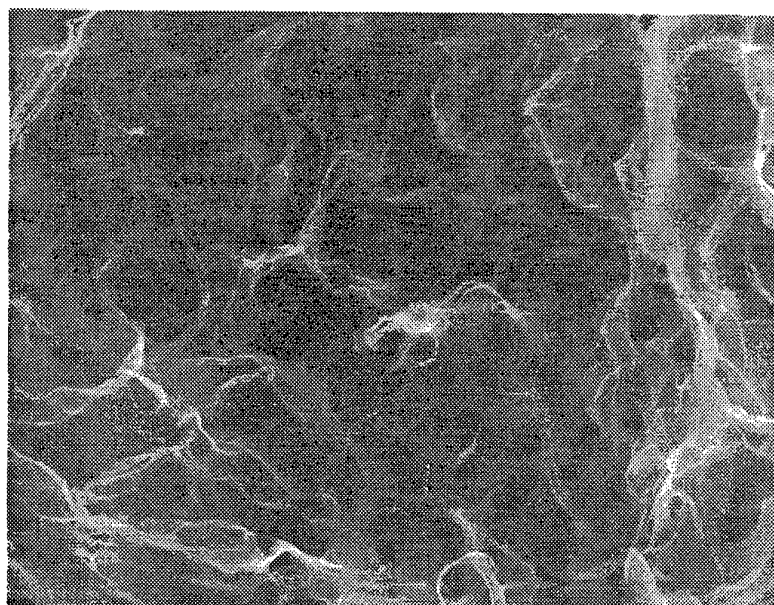
200  $\mu\text{m}$  

FIG.21D



200  $\mu$ m

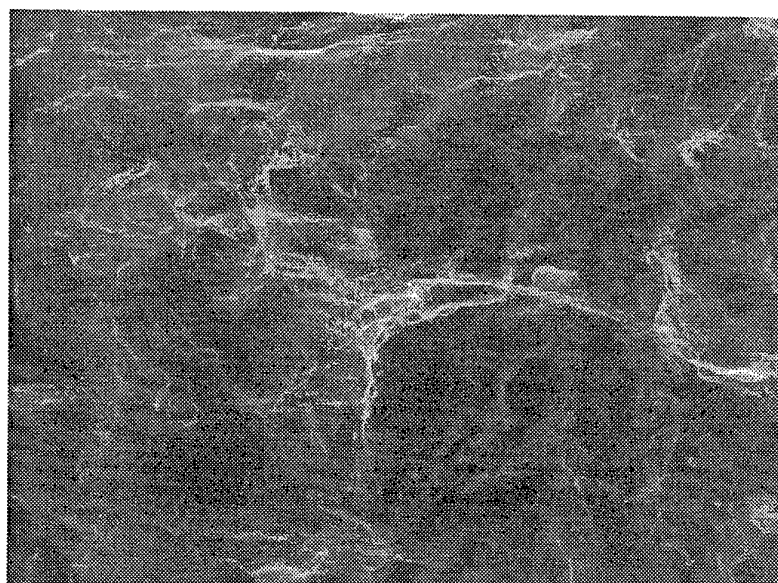
FIG.21E



200  $\mu$ m

FIG.21F

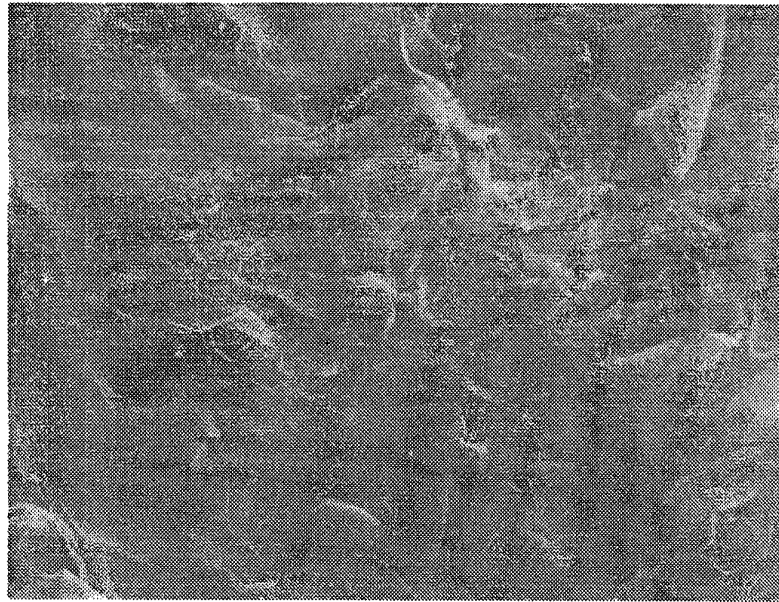




200 μm

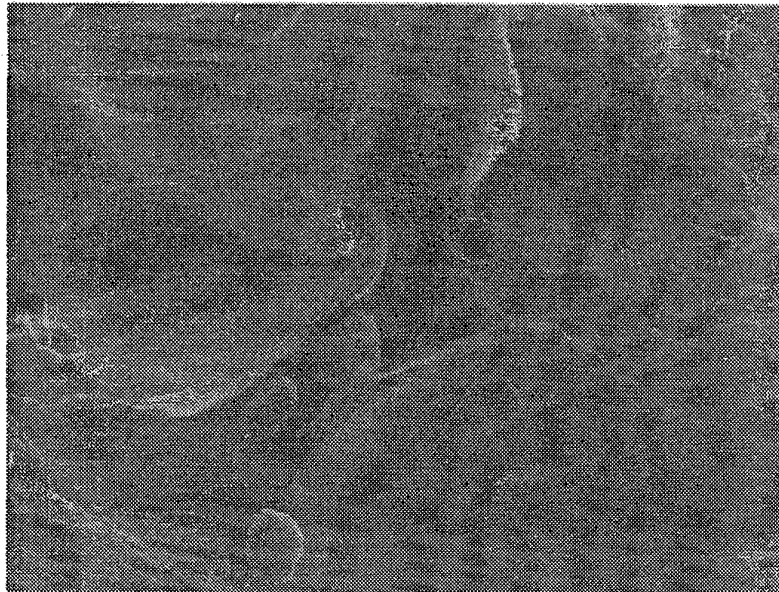
FIG.21G

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200 μm

FIG.22A



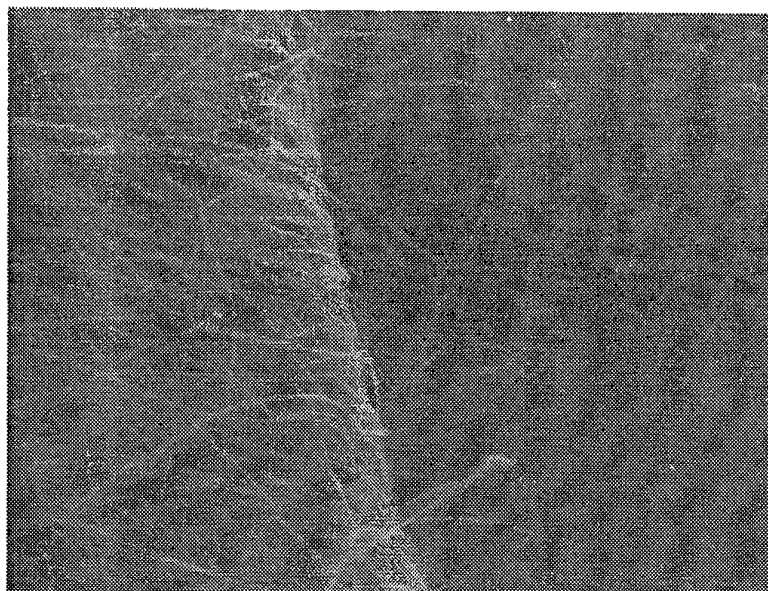
50 μm

FIG.22B



20µm

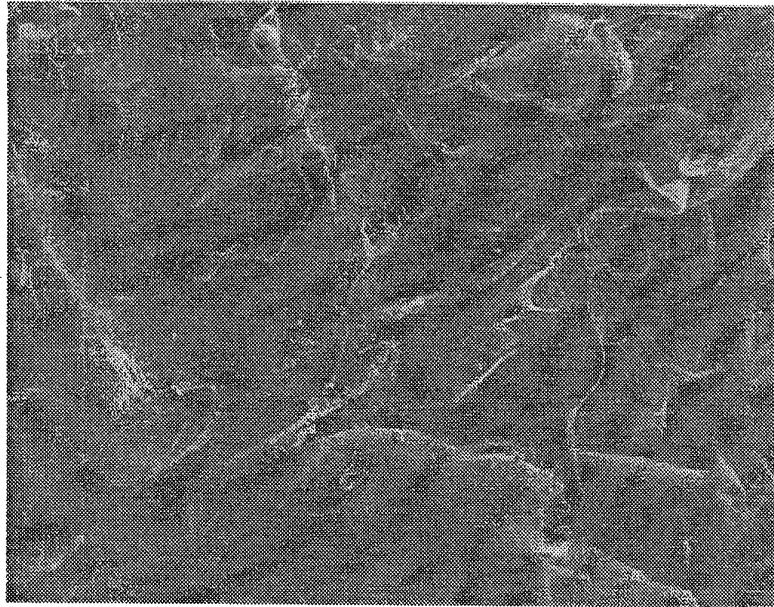
FIG.22C



5µm

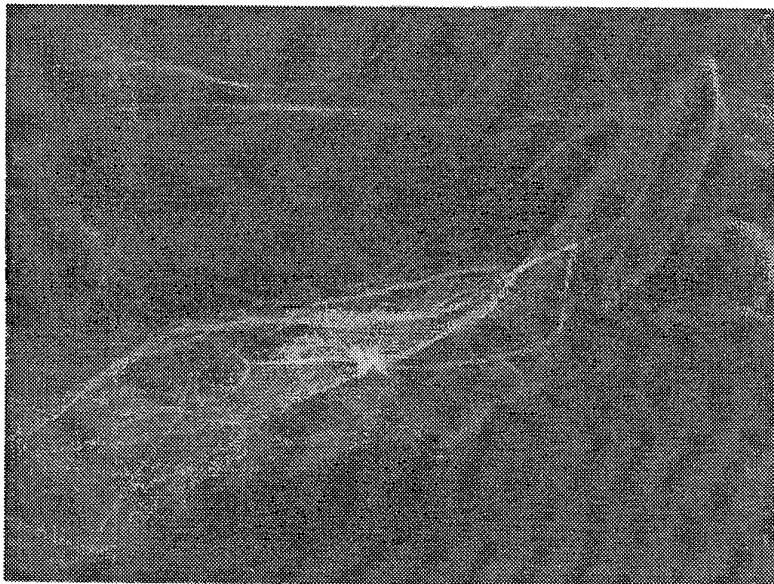
FIG.22D

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200µm

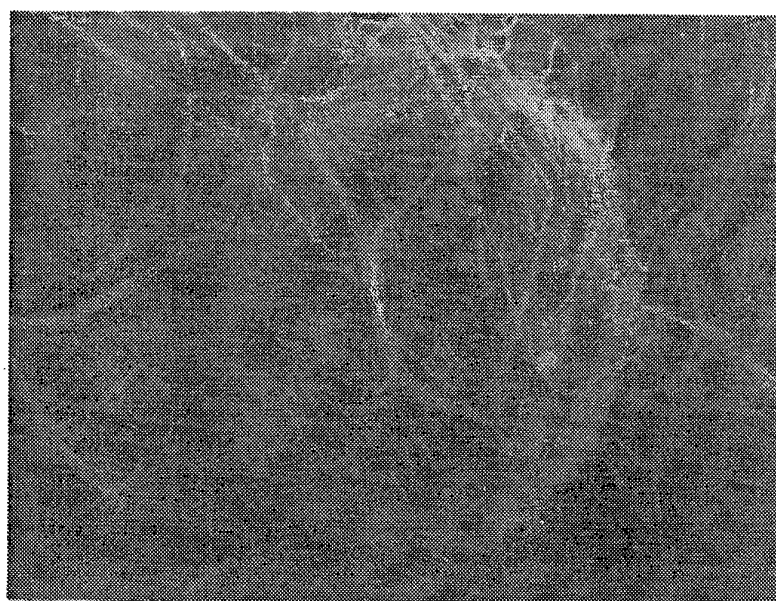
FIG.22E



20µm

FIG.22F

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5μm

FIG.22G

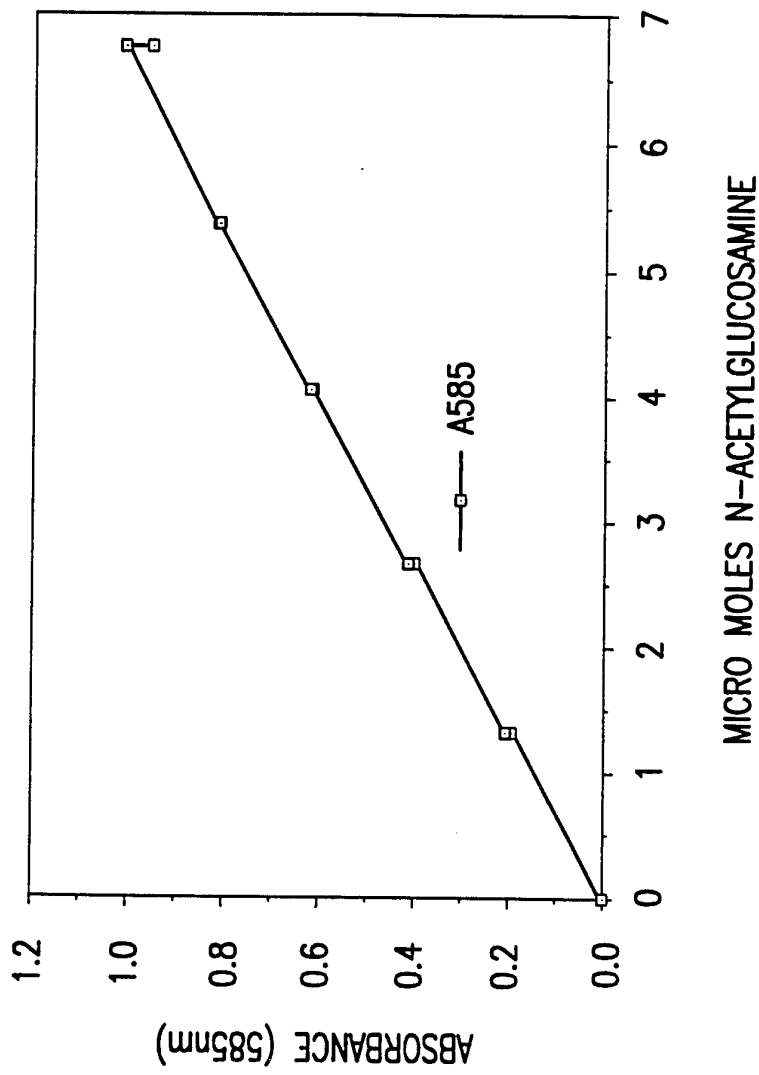


FIG.23

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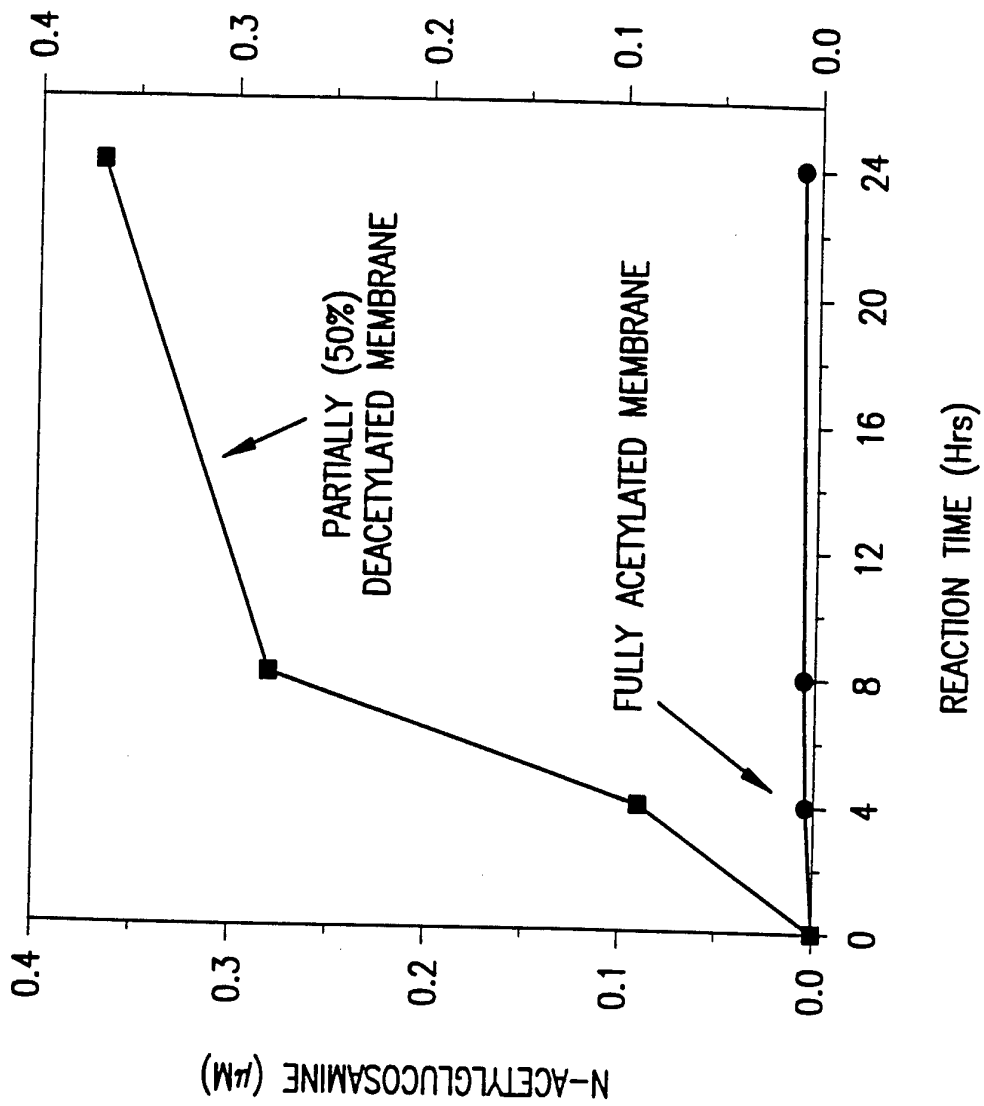


FIG.24

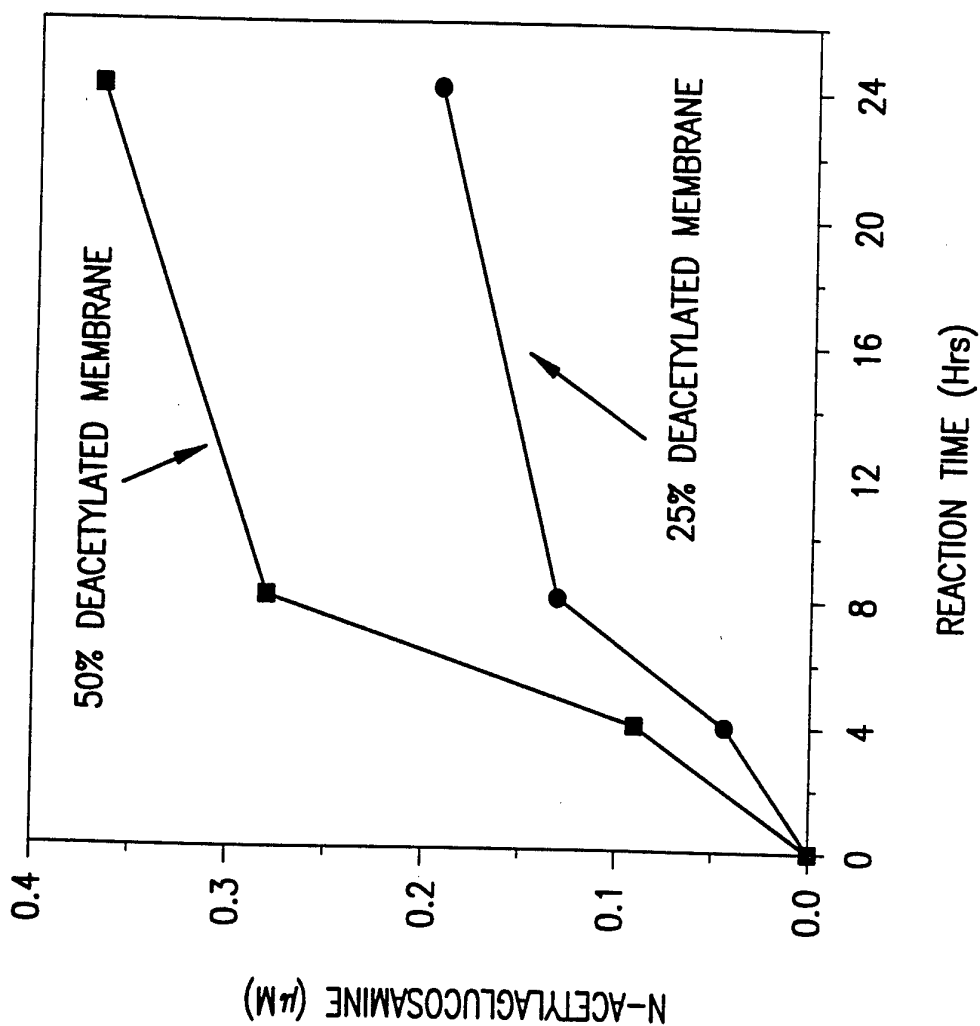


FIG.25



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PROTOTYPE 1: DAY 0

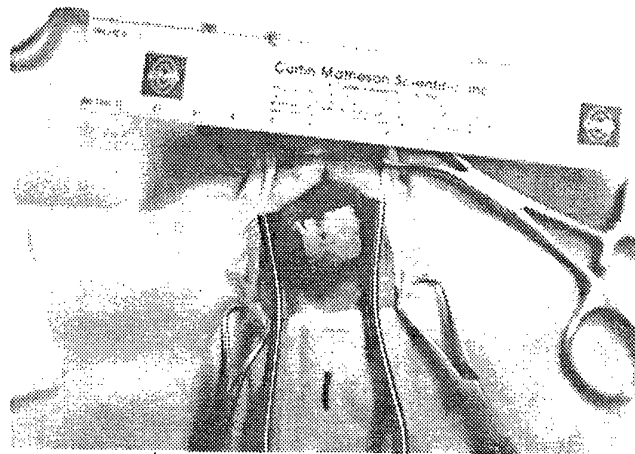


FIG.26A

PROTOTYPE 1: DAY 14

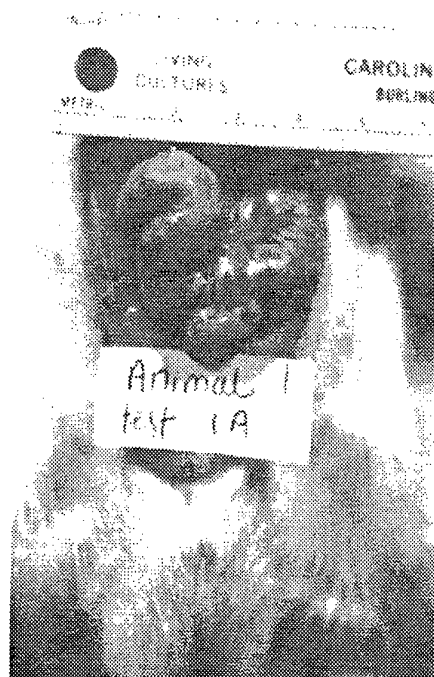


FIG.26B

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PROTOTYPE 1: DAY 21

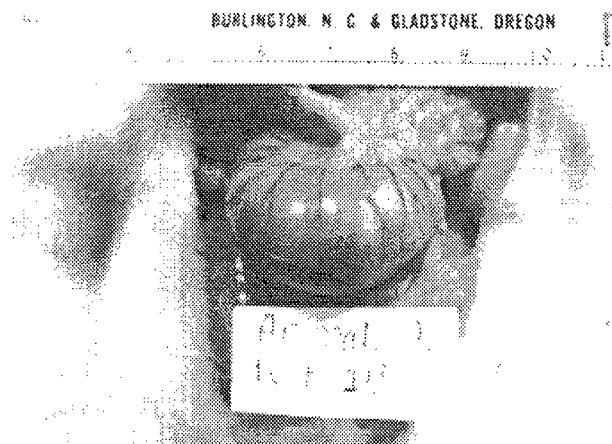


FIG.26C

PROTOTYPE 3A: DAY 0

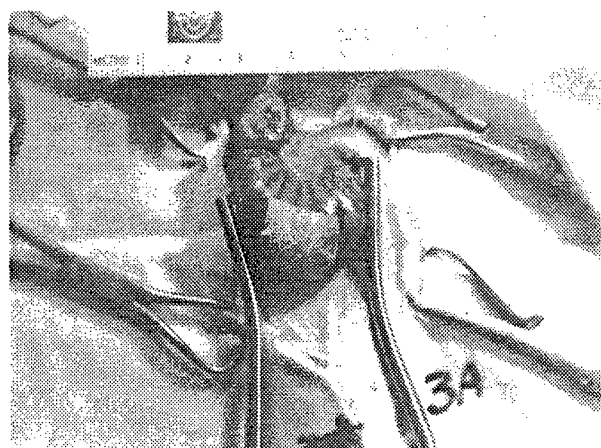


FIG.26D

PROTOTYPE 3A: DAY 14

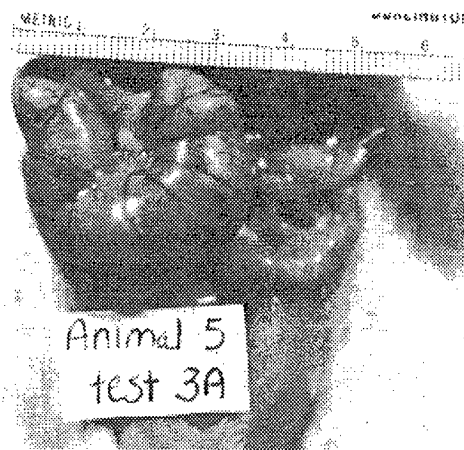


FIG.26E

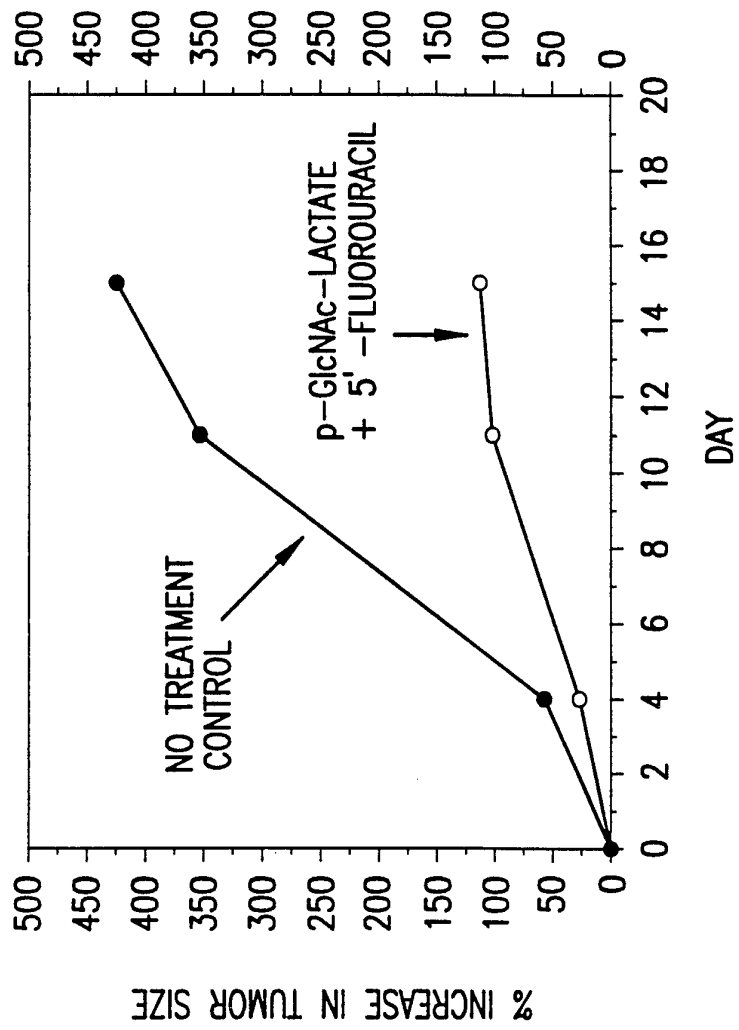


FIG.27

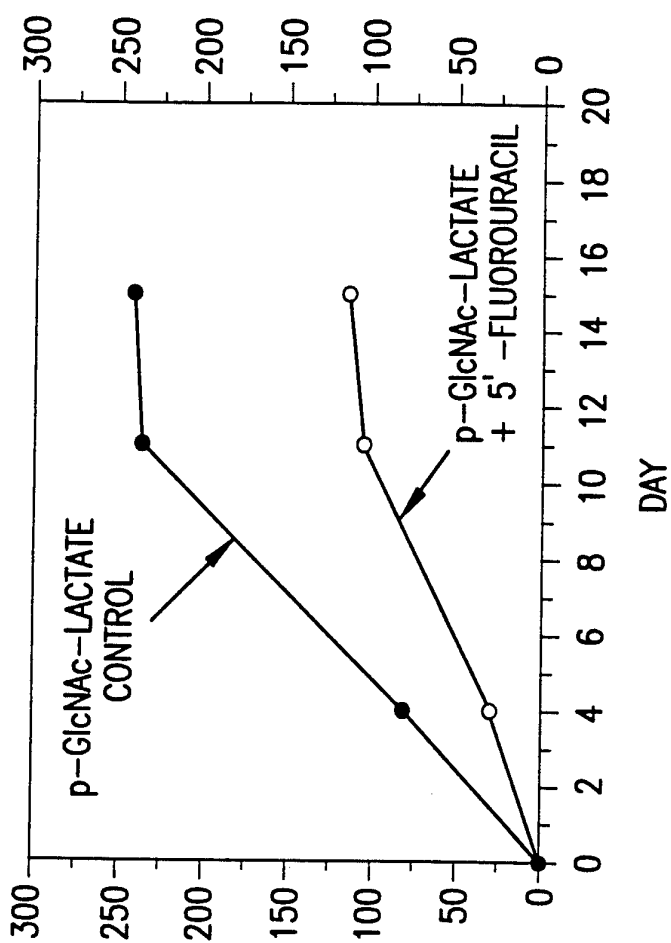


FIG.28

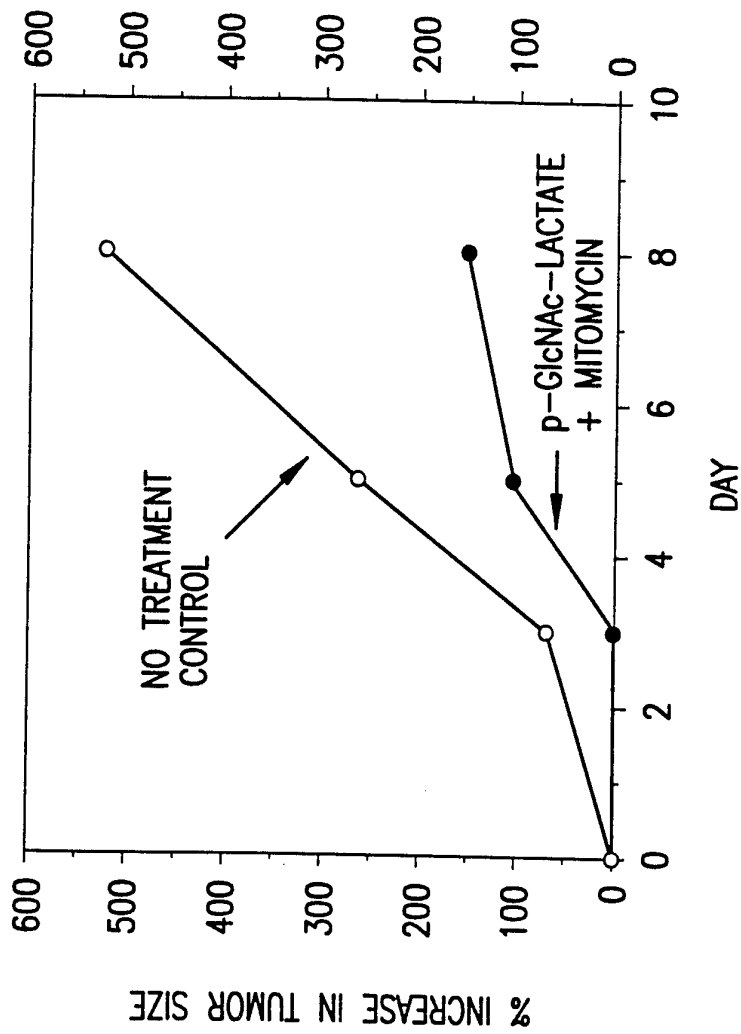


FIG.29

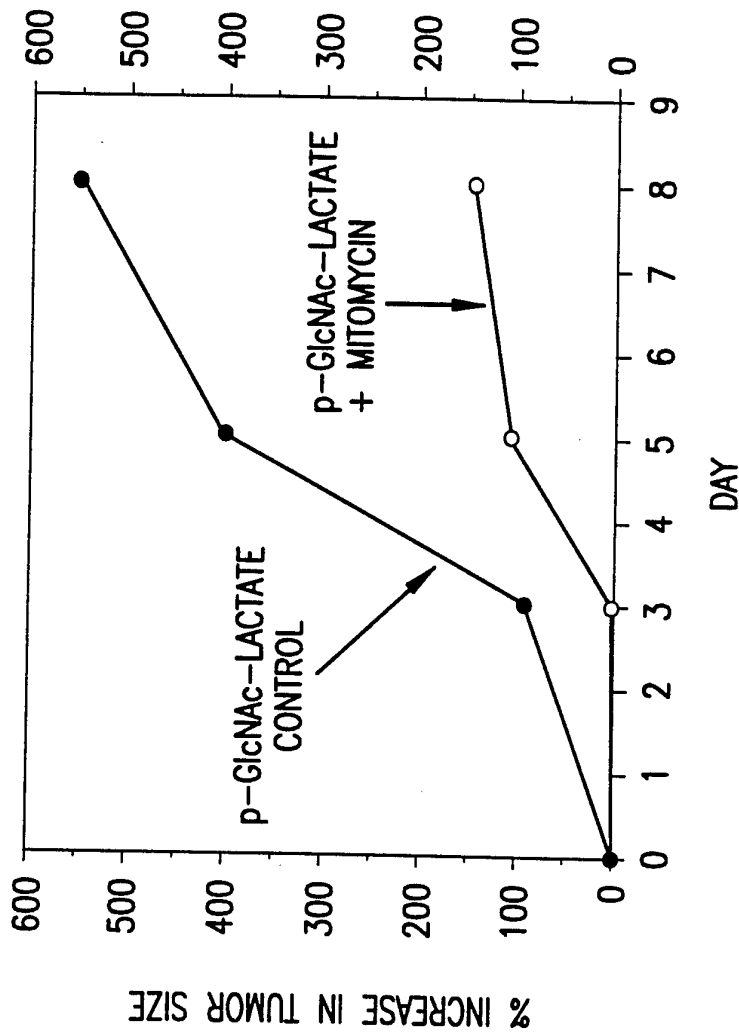


FIG.30

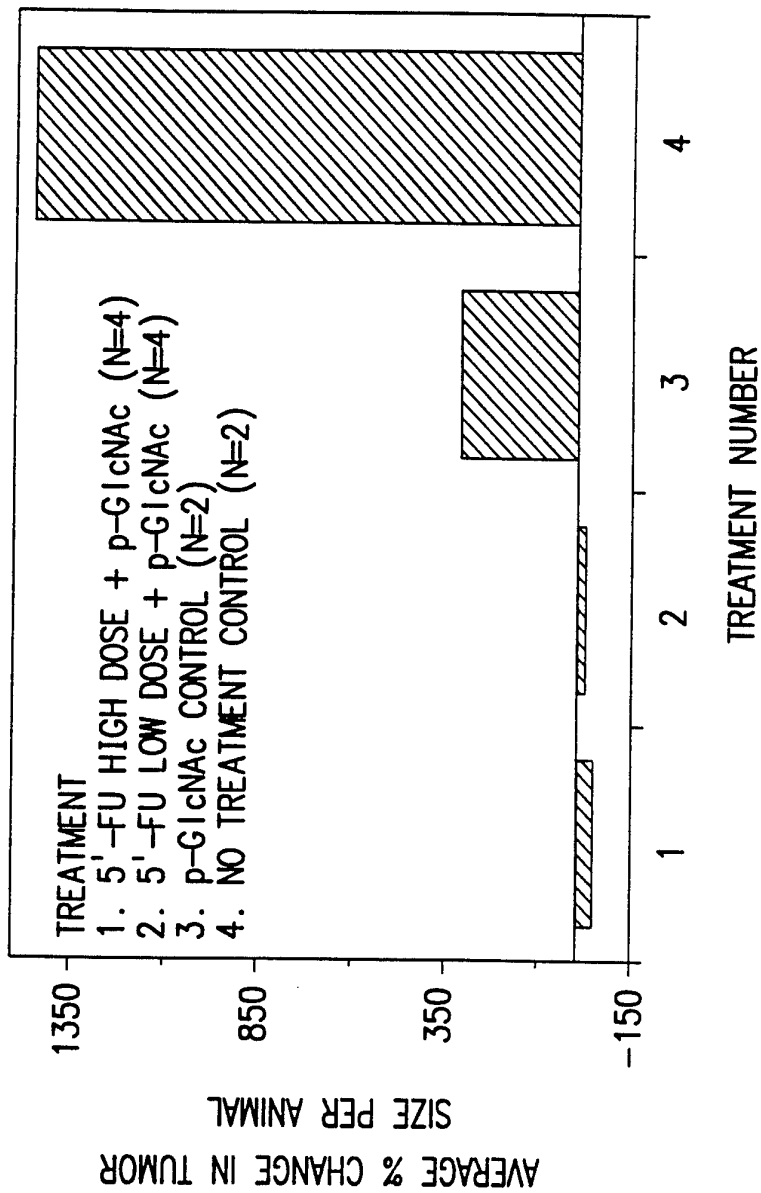


FIG. 31



# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US94/13706

**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(6) : C08B 37/08; C12P 19/26  
US CL : 514/55; 536/20; 435/84

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. : 514/55; 536/20; 435/84

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)

Please See Extra Sheet.

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y, P	BIOMATERIALS, Volume 15, Number 13, issued 1994, B. A. Zielinski et al., "Chitosan as a Matrix for Mammalian Cell Encapsulation", pages 1049-1056, see the abstract.	39, 50-58, 81-84
X, P	KEYSTONE SYMPOSIUM ON TISSUE ENGINEERING, 20-26 February 1994, J. N. Vournakis et al., "Isolation & Characterization of Pure Poly-N-Acetylglucosamine: Controlled Enzymatic Deacetylation and Formulation for Tissue Engineering Applications", column 2, abstract no. PZ313, Journal of Cellular Biochemistry, Supplement 0, Number 18C, page 283, see the entire abstract.	1-27
-----		-----
Y, P		28-94

Further documents are listed in the continuation of Box C.     See patent family annex.

<p>* Special categories of cited documents:</p> <p>*A* document defining the general state of the art which is not considered to be of particular relevance</p> <p>*E* earlier document published on or after the international filing date</p> <p>*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)</p> <p>*O* document referring to an oral disclosure, use, exhibition or other means</p> <p>*P* document published prior to the international filing date but later than the priority date claimed</p>	<p>*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</p> <p>*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</p> <p>*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</p> <p>*&amp;* document member of the same patent family</p>
--	---

Date of the actual completion of the international search <b>03 FEBRUARY 1995</b>	Date of mailing of the international search report <b>23 FEB 1995</b>
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231	Authorized officer <i>K. Kahler Fonda</i> <b>KATHLEEN KAHLER FONDA</b>
Facsimile No. (703) 305-3230	Telephone No. (703) 308-0196

## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US94/13706

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	BIOTECHNOLOGY PROGRESS, Volume 9, Number 5, issued 1993, H. W. Matthew et al., "Complex Coacervate Microcapsules for Mammalian Cell Culture and Artificial Organ Development", pages 510-519, see the abstract and pages 514-515.	39, 50-58, 81-84
Y	BIOTECHNOLOGY AND BIOENGINEERING, Volume 42, Number 10, issued 1993, A. F. Groboillot et al., "Membrane Formation by Interfacial Cross-Linking of Chitosan for Microencapsulation of Lactococcus lactis", pages 1157-1163, see the abstract.	39, 50-58, 81-84
Y	WO, A, 93/12875 (WANG) 08 July 1993, see pages 19-21.	4, 9, 23
Y	EP, A, 0,543,572 (FREEMAN) 26 MAY 1993, see columns 3-4.	39, 50-58, 81-84
Y	JOURNAL OF PHARMACY AND PHARMACOLOGY, Volume 44, Number 4, issued 1992, B. C. Thanoo et al., "Cross-Linked Chitosan Microspheres: Preparation and Evaluation as a Matrix for the Controlled Release of Pharmaceuticals", pages 283-286, see the abstract.	4, 9, 23, 39, 50-58, 81-84
Y	US, A, 3,989,535 (CAPOZZA) 02 NOVEMBER 1976, see columns 6-12.	1-13, 18-27, 39, 50-58, 81-84
Y	US, A, 3,988,411 (CAPOZZA) 26 OCTOBER 1976, see columns 6-12.	1-13, 18-27, 39, 50-58, 81-84
Y	CANADIAN JOURNAL OF BOTANY, Volume 43, issued 1965, J. McLachlan et al., "Studies on the Chitan (Chitin: Poly-N-Acetylglucosamine) Fibers of the Diatom Thalassiosira Fluviatilis Hustedt", pages 707-713, see the abstract and pages 708-709.	1-37, 40-44, 46-49
Y	CANADIAN JOURNAL OF CHEMISTRY, Volume 44, issued 1966, M. Falk et al., "Studies on Chitan ( $\beta$ -(1 $\rightarrow$ 4)-Linked 2-Acetamido-2-deoxy-D-Glucan) Fibers of the Diatom Thalassiosira Fluviatilis Hustedt", pages 2269-2281, see the abstract and pages 2271-2277.	1-37, 40-44, 46-49

# INTERNATIONAL SEARCH REPORT

International application No.

PCT/US94/13706

## B. FIELDS SEARCHED

Electronic data bases consulted (Name of data base and where practicable terms used):

databases: APS, Biosis, HCA, Registry

search terms: chitin#, chitosan#, microalga#, micro(w)alga#, thalas?, thallas?, diatom#, chrysophyt?, bacteria#, acetylglucosamine#, acetyl(w)glucosamine#, poly(w)glnac, encapsulat?, microencapsulat?, manif, purifn, prepn. Also inventor name search.