

(12) STANDARD PATENT
(19) AUSTRALIAN PATENT OFFICE

(11) Application No. **AU 2003286065 B2**

(54) Title
Isolation of inositol from plant materials

(51) International Patent Classification(s)
C12P 7/02 (2006.01)

(21) Application No: **2003286065** (22) Date of Filing: **2003.11.28**

(87) WIPO No: **WO04/050887**

(30) Priority Data

| | | |
|------------------|-------------------|--------------|
| (31) Number | (32) Date | (33) Country |
| 2,413,240 | 2002.11.29 | CA |

(43) Publication Date: **2004.06.23**

(43) Publication Journal Date: **2004.07.29**

(44) Accepted Journal Date: **2009.12.24**

(71) Applicant(s)
MCN BioProducts Inc.

(72) Inventor(s)
Newkirk, Rex W.;Classen, Henry L.;Maenz, David D.

(74) Agent / Attorney
FB Rice & Co, Level 23 200 Queen Street, Melbourne, VIC, 3000

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
17 June 2004 (17.06.2004)

PCT

(10) International Publication Number
WO 2004/050887 A3

(51) International Patent Classification⁷: **C12P 7/02**

[CA/CA]; 316 Hugo Avenue, Saskatoon, Saskatchewan S7N 1J8 (CA). **CLASSEN, Henry, L.** [CA/CA]; 235 Sylvian Crescent, Saskatoon, Saskatchewan S7H 5G1 (CA).

(21) International Application Number:
PCT/CA2003/001849

(22) International Filing Date:
28 November 2003 (28.11.2003)

(74) **Agent: BLAKE, CASSELS & GRAYDON LLP**; World Exchange Plaza, 20th Floor, 45 O'Connor Street, Ottawa, Ontario K1P 1A4 (CA).

(25) Filing Language: English

(81) **Designated States (national):** AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(26) Publication Language: English

(30) Priority Data:
2,413,240 29 November 2002 (29.11.2002) CA

(71) Applicant (for all designated States except US): **MCN BIOPRODUCTS INC.** [CA/CA]; Suite 860, Saskatoon Square, 410, 22nd Street, East, Saskatoon, Saskatchewan S7K 5T6 (CA).

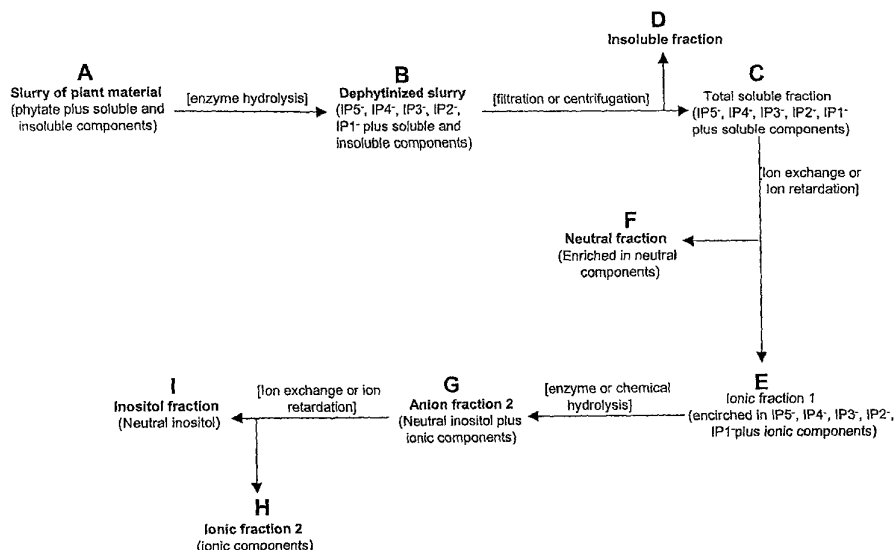
(84) **Designated States (regional):** ARIPO patent (BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

(72) Inventors; and
(75) Inventors/Applicants (for US only): **NEWKIRK, Rex, W.** [CA/CA]; 339 Haslam Crescent, Saskatoon, Saskatchewan S7S 1E6 (CA). **MAENZ, David, D.**

[Continued on next page]

(54) Title: ISOLATION OF INOSITOL FROM PLANT MATERIALS

Purification of inositol from plant materials



(57) **Abstract:** Phytate and/or phytin and/or phytic acid in an aqueous slurry of plant material is partially hydrolyzed by incubating the slurry with an enzyme product enriched in phytase. The soluble fraction of the slurry is separated into anionic and neutral fractions. The anionic fraction is then hydrolyzed further, and the hydrolyzate is separated into second ionic and neutral fractions. The second neutral fraction thus obtained is rich in inositol, and does not contain significant quantities of other sugars which would be hard to separate from it.



WO 2004/050887 A3



Published:

— with international search report

(88) Date of publication of the international search report:

23 September 2004

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

Purification of Inositol from Plant Materials

Field of the Invention

This invention relates to production of inositol from plant materials.

5 Background to the Invention

Inositol is a highly valued B-vitamin. Plants contain phytic acid {myoinositol 1,2,3,4,5,6-hexakis (dihydrogen phosphoric acid)} as the storage form of phosphorus. Phytic acid is found within plant cell structures as mineral bound complexes termed phytin. Phytin
10 is largely insoluble at neutral pH. Phytic acid can also exist in solution in the salt form termed phytate. The terms "phytin" and phytate are often used interchangeably. In this disclosure, the term "phytate" is intended to refer to phytic acid, phytate and phytin, except where a distinction between these materials is
15 made specifically.

Some of the partial hydrolysis products of phytate are inositol pentaphosphate (IP5), inositol tetraphosphate (IP4), inositol triphosphate (IP3), inositol diphosphate (IP2) and inositol monophosphate (IP1). These partial hydrolysis products of
20 phytate can be hydrolyzed further to yield inositol. The obtaining of inositol from a plant material requires conversion of the phytate to inositol and purification of the inositol from other components in the plant starting material.

Producing inositol from plant material is difficult. One approach is
25 to hydrolyze the phytate in an aqueous slurry, to yield various sugars including inositol. However, inositol is a neutral soluble sugar that is very similar in molecular size and charge

characteristics to other sugars such as glucose that are often present in high levels in plant materials. Because of this, it can be difficult to separate the inositol from the other carbohydrates in the slurry.

- 5 Another approach to production of inositol from plant materials is to purify the phytate from the starting slurry and to hydrolyze the purified phytate to inositol in a later step in the overall process. However, because phytate in plants usually exists in the form of phytin, direct phytate purification from an aqueous slurry of plant materials requires solubilization of phytin and then separation of the phytate from the remainder of the
- 10 components of the slurry. Efficient extraction, solubilization of phytin and separation from the remaining components of the slurry is difficult.

Description of the Invention

This invention describes a useful and novel process for overcoming the inherent

15 difficulties in obtaining inositol from plant materials.

In accordance with the inventive process, phytate in an aqueous slurry of plant material is partially hydrolyzed by incubating the slurry with an enzyme product enriched in phytase. The soluble fraction of the slurry is separated into anionic and neutral

20 fractions. The anionic fraction is then hydrolyzed further, and is in turn separated into ionic and neutral fractions.

The neutral fraction thus obtained is rich in inositol, and does not contain significant quantities of other sugars which would be hard to separate from it.

25

The present invention provides a process for producing inositol from plant materials comprising the steps of:

- (a) treating an aqueous slurry of plant material with a phytase enzyme to partially hydrolyse at least one of phytate, phytic acid and phytin to
- 30 inositol phosphates, under conditions which do not promote full hydrolysis in inositol;
- (b) separating said slurry into a water soluble fraction and a water-insoluble fraction;

35

- (c) separating said water soluble fraction into a first ionic fraction which contains anionic components comprising inositol phosphates and a first other fraction which contains the neutral components;
- 5 (d) hydrolysing the inositol phosphates in said first ionic fraction; and
- (e) separating the hydrolyzed first ionic fraction into a second ionic fraction and a second neutral fraction which contains inositol.

- 3 -

Description of the Drawings

Figure 1 is a process flow chart depicting processing stages in accordance with the invention.

5

Detailed Description of the Invention

According to the invention, an aqueous slurry of plant material is partially hydrolyzed using phytase enzyme. Figure 1 shows a process flow chart of the various steps in the process of the invention.

10

As shown in Figure 1, an aqueous slurry of plant material A is subjected to partial hydrolysis with phytase. This is preferably done by incubating the aqueous slurry with phytase, at suitable temperature and pH to encourage partial hydrolysis.

15 The phytase enzyme can hydrolyze phytate to inositol pentaphosphate (IP5), inositol tetraphosphate (IP4), inositol triphosphate (IP3) and inositol diphosphate (IP2). However, the phytase has little activity for hydrolysis of inositol 2-monophosphate inositol (IP1). Acid phosphatase can readily hydrolyze IP1 to free inositol, which is not desired at this point in the inventive process. Thus, the source of phytase used preferably contains little or no acid phosphatase. A source of phytase containing acid phosphatase activity can also be used, if reaction conditions are chosen to favour phytase activity and avoid substantial hydrolysis of IP1 by acid phosphatase. In using a source of phytase containing acid phosphatase the preferred

20

25

- 4 -

pH of the reaction is greater than 3.0 and less than 7 for optimum phytase activity without substantial hydrolysis of IP1 to inositol.

IP5, IP4, IP3, IP4, IP2 and IP1 are the major products of the reaction. They are highly soluble negatively charged compounds
5 that exist in solution in the partially hydrolyzed slurry (shown as "B" in Figure 1). The partially hydrolyzed slurry is separated by physical separation means, such as filtration or centrifugation, to generate an inositol phosphate-containing soluble fraction (a "total soluble" fraction called C in Figure 1,) and an insoluble
10 fraction (called "D" in Figure 1).

Unlike inositol, inositol phosphates have a negative charge. It is therefore possible to separate the total soluble fraction into an anionic fraction and a first neutral fraction, with the inositol phosphates passing into the anionic fraction. Depending on how
15 the separation is carried out, any cationic soluble materials present may remain either with the anionic fraction or the first neutral fraction.

Total soluble fraction C is therefore separated into a first ionic fraction enriched in anionic constituents – called "ionic fraction 1" or "E" in Figure1) and a first fraction enriched in neutral constituents (and possibly cationic constituents as well, called "F" in Figure 1). Ionic fraction 1 (E) contains most of the inositol phosphates, and the neutral fraction contains most of the neutral soluble constituents of the total soluble fraction. The separation is
20 done using known techniques for the separation of charged ionic species from soluble neutral compounds in a solution. Such techniques are, for example, ion exchange, ion exclusion, or ion retardation column separations. If it is desired to retain the cationic components in the neutral fraction, a cationic ion

- 5 -

exchange resin can be used, which will separate out only the anionic components into the first ionic fraction. If it is desired that the cationic components are separated out as well, then mixed anionic and cationic exchange resins can be used. The important
5 thing at this stage is to end with one fraction which contains the anionic components and a second which contains the neutral components. Cationic components are not of concern in the process of the invention, so they can remain in either fraction.

The next step in the process is to complete the hydrolysis of
10 inositol phosphates in the ionic fraction. This process can be done with enzymes such as phytase or acid phosphatase or without enzyme-based catalysis under controlled conditions of temperature, pressure and pH. Suitable conditions for inositol phosphate hydrolysis are known, and can be chosen according to
15 the particular reaction equipment available. The preferred approach is to use an enzyme source containing acid phosphatase at a pH of less than 4 for optimum activity. Complete hydrolysis of inositol phosphates will generate an anionic fraction (G in Figure 1) which contains various anionic
20 compounds from fraction E as well as neutral inositol.

Inositol can be separated from the remainder of the soluble compounds in the anionic fraction G using known techniques for separating charged from neutral compounds in solution, such as, for example, an ion exchange, ion exclusion or ion retardation
25 column. This process generates a second ionic fraction (—called herein ionic fraction 2, and indicated in Figure 1 as “H’). and a second neutral I fraction (“I” in Figure1) rich in inositol. The inositol in the second neutral fraction can then be concentrated, crystallized and dried to form a final dry purified inositol product.

The invention has been described by reference to preferred embodiments, but it will be understood that other embodiments will be evident to a person skilled in the art. It is therefore desired that the invention shall not be limited by the particular embodiments shown, but shall include such other embodiments as would occur to a skilled person.

5

Any discussion of documents, acts, materials, devices, articles or the like which has been included in the present specification is solely for the purpose of providing a context for the present invention. It is not to be taken as an admission that any or all of these matters form part of the prior art base or were common general knowledge in the
10 field relevant to the present invention as it existed before the priority date of each claim of this application.

Throughout this specification the word "comprise", or variations such as "comprises" or "comprising", will be understood to imply the inclusion of a stated element, integer
15 or step, or group of elements, integers or steps, but not the exclusion of any other element, integer or step, or group of elements, integers or steps.

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. A process for producing inositol from plant materials comprising the steps of:
 - 5 (a) treating an aqueous slurry of plant material with a phytase enzyme to partially hydrolyse at least one of phytate, phytic acid and phytin to inositol phosphates, under conditions which do not promote full hydrolysis in inositol;
 - 10 (b) separating said slurry into a water soluble fraction and a water-insoluble fraction;
 - (c) separating said water soluble fraction into a first ionic fraction which contains anionic components comprising inositol phosphates and a first other fraction which contains the neutral components;
 - 15 (d) hydrolysing the inositol phosphates in said first ionic fraction; and
 - (e) separating the hydrolyzed first ionic fraction into a second ionic fraction and a second neutral fraction which contains inositol.
 - 20
2. The process of claim 1 wherein said phytase enzyme does not include acid phosphatase.
3. The process of claim 1 or claim 2 wherein said step of treating the aqueous
25 slurry is carried out at a pH between about 3.0 and about 7.0.
4. The process of claim 3 wherein said phytase enzyme includes acid phosphatase.
5. The process of any one of claims 1 to 4 wherein said step of separating the
30 slurry into a water-soluble fraction and an insoluble fraction is carried out by centrifugation.
6. The process of any one of claims 1 to 4 wherein said step of separating the
35 slurry into a water-soluble fraction and an insoluble fraction is carried out by filtration.

7. The process of any of claims 1-6, in which the step of hydrolyzing the inositol phosphates in said first ionic fraction comprises treatment of said first ionic fraction with phytase.
- 5 8. The process of any of claims 1-7, in which the step of hydrolyzing the inositol phosphates in said first ionic fraction comprises treatment of said first ionic fraction with acid phosphatase.
9. The process of claim 8, wherein said hydrolysis is carried out at a pH of less
10 than 4.
10. The process of any of claims 1-6, in which the step of hydrolyzing the inositol phosphates in said first ionic fraction comprises subjecting of said first ionic fraction in the absence of added phytase to conditions of temperature, pressure and pH which
15 promote hydrolysis.
11. A process as claimed in any of claims 1-10, comprising the step of separating purified inositol from said second neutral fraction.
- 20 12. A process according to claim 1 substantially as hereinbefore described.

Fig1. Purification of inositol from plant materials

