In accordance with proposed 21 CFR §170.36 [Notice of a claim for exemption based on a Generally Recognized As Safe (GRAS) determination] published in the Federal Register [62 FR 18938 (17 April 1997)], I am submitting in triplicate, as the notifier [George Weston Foods Limited, 1 Bradwood Street, Enfield, NSW, 2136, Australia], a Notice of the determination, on the basis of scientific procedures, that sweet lupin flour derived from sweet varieties of Lupinus spp. (lupin), produced by George Weston Foods Limited (GWF), as defined in the enclosed documents, is GRAS under specific conditions of use as a food ingredient, and therefore, is exempt from the premarket approval requirements of the Federal, Food, Drug and Cosmetic Act. Information setting forth the basis for the GRAS determination, which includes a comprehensive summary of the data available and reviewed by an independent panel of experts in support of the safety of GWF's sweet lupin flour ingredient under the intended conditions of use, as well as curricula vitae evidencing the qualifications of the members of the panel of experts for evaluating the safety of food ingredients, also are enclosed.

I trust that the enclosed Notice is acceptable. Should you have any questions or concerns regarding this GRAS Notice, please do not hesitate to contact me at any point during the review process so that we may provide a response in a timely manner.

Sincerely,

Peter Schutz
Chief Executive

George Weston Technologies
A Division of George Weston Foods Limited
peter.schultz@gwf.com.au peter.schutz@gwf.com.au

Encl.

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SWEET LUPIN FLOUR GRAS NOTICE

Prepared for: Robert L. Martin, Ph.D.
Office of Food Additive Safety (HFS-200)
Center for Food Safety and Applied Nutrition
Food and Drug Administration
5100 Paint Branch Parkway
College Park, MD 20740-3835

Prepared by: George Weston Foods Limited
1 Braidwood Street
Enfield, NSW, 2136
Australia

September 26, 2008
SWEET LUPIN FLOUR GRAS NOTICE

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George Weston Foods Limited
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I. GRAS EXEMPTION CLAIM

A. Claim of Exemption From the Requirement for Premarket Approval

Pursuant to Proposed 21 CFR §170.36(c)(1) [62 FR 18938 (17 April 1997)]
(U.S. FDA, 1997)

As defined herein, flour derived from sweet lupin has been determined by George Weston Foods Limited (GWF) to be Generally Recognized as Safe (GRAS) for use in a variety of traditional food products. This determination is based on scientific procedures, as described in the following sections, under the conditions of its intended use in food. Therefore, consistent with Section 201(s) of the Federal Food, Drug, and Cosmetic Act, the use of flour derived from sweet lupin in food as described below is exempt from the requirement of premarket approval.

Signed,

______________________________
Peter Schutz
Chief Executive
Date: 26 September 2008

George Weston Technologies
A Division of George Weston Foods Limited
peter.schutz@qwf.com.au

B. Name and Address of Notifier

George Weston Foods Limited
1 Braidwood Street
Enfield, NSW, 2136
Australia

C. Common Name of the Notified Substance

Sweet lupin flour or lupin flour

D. Conditions of Intended Use in Food

GWF intends to market flour derived from sweet varieties of Lupinus spp. (lupin) as a food ingredient in various traditional food products intended for sale on the U.S. market. The intended food uses include baked goods and baking mixes and grain products and pastas, and the lupin flour will be added to food products at a maximum use level of 25%. Sweet lupin flour is not intended for use in meat or meat-containing products.
E. Basis for the GRAS Determination

Pursuant to 21 CFR § 170.30, flour derived from sweet lupin has been determined by GWF to be GRAS on the basis of scientific procedures (U.S. FDA, 2008a). This GRAS determination is based on data generally available in the public domain pertaining to the safety of lupin and sweet lupin-derived ingredients, including lupin flour, as discussed herein, and on a consensus among a panel of experts¹ who are qualified by scientific training and experience to evaluate the safety of sweet lupin flour as a component of food [see Appendix A, "EXPERT PANEL CONSENSUS STATEMENT REGARDING THE GENERALLY RECOGNIZED AS SAFE (GRAS) STATUS OF SWEET LUPIN-DERIVED INGREDIENTS FOR USE IN FOODS"].

F. Availability of Information

The data and information that serve as the basis for this GRAS Notice will be sent to the U.S. Food and Drug Administration (FDA) upon request, or will be available for review and copying at reasonable times at the offices of:

George Weston Foods Limited
1 Braidwood Street
Enfield, NSW, 2136
Australia

Should the FDA have any questions or additional information requests regarding this notification, GWF will supply these data and information.

¹ The panel of experts consisted of Prof. Joseph F. Borzelleca, Ph.D. (Virginia Commonwealth University School of Medicine), Ashley S. Roberts, Ph.D. (Cantox Health Sciences International), and Prof. Stephen L. Taylor, Ph.D. (University of Nebraska).
II. DETAILED INFORMATION ABOUT THE IDENTITY OF THE SUBSTANCE

A. Identity

Sweet lupin flour is obtained by dehulling and milling/grinding the whole seed of sweet varieties of *Lupinus* spp. (lupin). Four (4) species of lupin have been cultivated to include both a bitter variety and a 'sweet' variety, which is so-named due to its low alkaloid content (~0.001 to 0.002%), making the sweet varieties suitable for consumption by humans and livestock (Petterson, 1998)\(^2\) (see Appendix B-1). The sweet varieties of lupin are of the following species: *L. angustifolia, L. albus, L. luteus,* and *L. mutabilis.*

B. Method of Manufacture

Sweet lupin flour is produced from the whole seed of sweet lupin. As mentioned, the species with sweet varieties of lupin used to manufacture the sweet lupin flour include *L. angustifolia, L. albus, L. luteus,* and *L. mutabilis.* The seeds are received from growers and cleaned. The flour is prepared using only physical/mechanical processing of the lupin seeds. The hull is removed from the seeds and the dehulled seeds (cotyledons or kernels) are dry-milled to produce sweet lupin flour. A schematic overview of the manufacturing process for sweet lupin flour is presented in Figure 1.

Figure 1  Schematic Overview of the Manufacturing of Sweet Lupin Flour

```
| Sweet lupin seeds* | De-hulling | Dry Milling | Sweet Lupin Flour |
```

* From sweet varieties of *L. angustifolia, L. albus, L. luteus,* and *L. mutabilis*

C. Specifications for Food-Grade Material

Sweet lupin flour is produced in accordance with current Good Manufacturing Practices (cGMP) and in order to ensure a consistent, safe product, GWF has established numerous food-grade specification parameters for the final preparation. These parameters comprise physical, chemical, and microbiological specifications, including a maximal alkaloid level of <200 ppm, as set forth by the Advisory Committee on Novel Foods and Processes (ACNFP).

---

\(^2\) Following a review of data on the safety of ingredients derived from sweet lupin, the Advisory Committee on Novel Foods and Processes (ACNFP) of the United Kingdom concluded that lupin seeds were safe for the production of human foods provided that the level of lupin alkaloids in the derived products did not exceed 200 mg/kg. A summary of the alkaloids identified in lupin is presented in Appendix B-1.
SWEET LUPIN FLOUR GRAS NOTICE

1996) of the United Kingdom (UK), and a maximum phomopsins\(^3\) level of 5 ppb, which is consistent with the maximum permitted value for human consumption of 5 μg phomopsins/kg seed established by Food Standards Australia New Zealand (FSANZ) and the Department of Health of the UK. The product specifications for sweet lupin flour are presented in Table 1.

<table>
<thead>
<tr>
<th>Specification Parameter</th>
<th>Specification</th>
<th>Method of Analysis*</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Physical and Chemical</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Description</td>
<td>Yellow-colored flour, free from foreign material and objectionable odors and flavors</td>
<td>Visual and olfactory inspection</td>
</tr>
<tr>
<td>Particle size (μm)</td>
<td>≤180</td>
<td>Passes through 180 μm sieve</td>
</tr>
<tr>
<td>Protein (TN x 6.25, % DSB m/m)</td>
<td>Minimum of 35</td>
<td>Australian Standard AS300.1.2.1-1991</td>
</tr>
<tr>
<td>Fat (% DSB m/m)</td>
<td>7 to 10</td>
<td>Australian Standard AS 2300.1.3-1988</td>
</tr>
<tr>
<td>Total Carbohydrate (% DSB m/m)</td>
<td>7 to 9</td>
<td>By Difference(^5)</td>
</tr>
<tr>
<td>Insoluble Dietary Fiber (% DSB m/m)</td>
<td>32 to 36</td>
<td>AOAC Official Method AOAC 985.29</td>
</tr>
<tr>
<td>Soluble Dietary Fiber (% DSB m/m)</td>
<td>3 to 6</td>
<td>AOAC Official Method AOAC 985.29</td>
</tr>
<tr>
<td>Moisture (%/m/m)</td>
<td>7 to 11</td>
<td>Australian Standard AS 2300.1.1-1996</td>
</tr>
<tr>
<td>Ash (% DSB m/m)</td>
<td>2 to 4</td>
<td>Australian Standard AS 2300.1.5-1998</td>
</tr>
<tr>
<td>Alkaloids (ppm)</td>
<td>&lt;200</td>
<td>GC-MS(^7)</td>
</tr>
<tr>
<td>Phomopsins (ppb)</td>
<td>&lt;5</td>
<td>Agrifood Technology Method TP/043</td>
</tr>
<tr>
<td>Cadmium (Cd) (ppm)</td>
<td>&lt;0.1</td>
<td>ICP-MS Method IEL1STIM</td>
</tr>
<tr>
<td>Lead (Pb) (ppm)</td>
<td>&lt;0.2</td>
<td>ICP-MS Method IEL1STIM</td>
</tr>
<tr>
<td><strong>Microbiological</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total plate count (CFU/g)</td>
<td>&lt;40,000</td>
<td>Australian Standard AS 1766.2.1</td>
</tr>
<tr>
<td>Coliforms (CFU/g)</td>
<td>&lt;100</td>
<td>Australian Standard AS 1766.2.3</td>
</tr>
<tr>
<td><em>Escherichia coli</em> (CFU/g)</td>
<td>&lt;10</td>
<td>Australian Standard AS 1766.2.3</td>
</tr>
<tr>
<td>Salmonella spp. (per 25 g)</td>
<td>Absent</td>
<td>AOAC Official Method 966.08, AOAC Official Method 2004.03</td>
</tr>
<tr>
<td>Yeasts and moulds (CFU/g)</td>
<td>&lt;1,000</td>
<td>Australian Standard AS 1766.2.2</td>
</tr>
<tr>
<td>Staphylococcus spp. (CFU/g)</td>
<td>&lt;100</td>
<td>Australian Standard AS 1766.2.4</td>
</tr>
<tr>
<td><em>Bacillus cereus</em> (CFU/g)</td>
<td>&lt;100</td>
<td>Australian Standard AS 1766.2.6</td>
</tr>
</tbody>
</table>

\(^3\) Phomopsins are toxins produced by fungi such as *Diaporthe toxica* or *Phomopsis leptostromiformis*, which grow on lupin plants, and phomopsin toxicity caused by phomopsin ingestion is called lupinosis (Allen, 1986; Morcombe et al., 1992; ANZFA, 2001a).

II. DETAILED INFORMATION ABOUT THE IDENTITY OF THE SUBSTANCE
George Weston Foods Limited
September 26, 2008
Table 1  Physical, Chemical, and Microbiological Specifications for Sweet Lupin Flour*

<table>
<thead>
<tr>
<th>Specification Parameter</th>
<th>Specification</th>
<th>Method of Analysis*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Listeria (per 25 g)</td>
<td>Absent</td>
<td>AOAC Official Method 999.06, AOAC Official Method 2004.06</td>
</tr>
</tbody>
</table>

* Milled from clean, non-genetically modified de-hulled sweet lupin cotyledon.
AOAC = Association of Analytical Communities CFU = Colony-forming units; DSB = dry solid basis; GC-MS = Gas chromatography/Mass spectrometry; ICP-MS = Inductively coupled plasma mass spectrometry; TN = total nitrogen
a For details of the methods of analyses please see http://www.aoac.org/ or http://www.standards.com.au/.
b By calculation: 100 - (Moisture + Fat + Protein + Ash + Dietary Fiber) = Total Carbohydrate
c Conducted at the Chemistry Centre of Western Australia

Product Analysis

Several lots of the manufactured product were analyzed to confirm that the manufacturing process produced a consistent product within the physical, chemical, and microbiological parameters of the specifications. A summary of the complete analyses of these batches is presented in Appendix B-2, along with corresponding certificates of analysis. The levels of alkaloids and phomopsins present in the sweet lupin flour produced by GWF comply with the maximal established levels of <200 ppm and 5 ppb, respectively, and therefore, are expected not to produce any adverse effects on human health.

Pesticide Residues

Sweet lupin flour is derived from a raw agricultural product, and therefore, pesticide residue analysis was conducted on the final product. Maximum Residue Limits (MRL) for grain products in the U.S. were identified in Title 40 of the Code of Federal Regulations (CFR), and where data were available, the pesticide levels in the sweet lupin flour were determined to be below the levels established by the U.S. Environmental Protection Agency (U.S. EPA, 2007). Pesticide residues in sweet lupin flour also were compared with MRL established by the Australia New Zealand Food Standards Code (ANZ Food Standards Code) (FSANZ, 2005) and/or by the Australian Pesticides and Veterinary Medicines Authority (APVMA, 2005) for grain and nut products. With the exception of some of the organochlorine compounds, the results of the analyses indicated that all tested residue components were below the MRL. With respect to the organochlorine compounds for which the sweet lupin flour did not meet the established MRL, the methods of analyses utilized for determining the levels of these compounds present in this ingredient involved limits of detection that were less sensitive than the established MRL, and hence the levels of these compounds may in fact comply with the regulatory standards. Moreover, many of the residues are sparingly soluble in water, lending to inefficient concentrating of residues in the final material. It is therefore expected that residues of pesticides that are present in the final product will not be of any concern. The analytical data of the pesticide residues in the sweet lupin flour in relation to identified MRL are presented in Appendix B-3.
Stability of Sweet Lupin Flour

The sweet lupin flour should be stored at room temperature in a dry environment, and under proper storage conditions, the ingredient has a shelf life of 6 months.
III. SELF-LIMITING LEVELS OF USE

The use of sweet lupin flour is self-limiting due to the effect of lupin flour on the baking characteristics and the impact on the sensory characteristics of the product. Sweet lupin flour is intended to replace a portion of other sources of flour. The level of substitution of sweet lupin flour for other sources of flour will be 10 to 25%.
IV. BASIS FOR GRAS DETERMINATION

A. Documentation to Support the Safety of Sweet Lupin Flour

The determination that sweet lupin flour is GRAS is on the basis of scientific procedures, and the information supporting the general recognition of the safe use of sweet lupin flour includes:

- published scientific data on the background consumption of lupin and lupin-derived ingredients;
- the entirety of pre-clinical and human studies assessing the safety and nutritional value of lupin and lupin-derived ingredients;
- the compositional similarity of sweet lupin flour to wheat flour; and
- data pertaining to the identity, intended use, and estimated intake of sweet lupin flour.

Moreover, these data were reviewed by a panel of experts, qualified by scientific training and experience to evaluate the safety of ingredients as components of food, who concluded that the proposed uses of sweet lupin flour are safe and suitable and would be GRAS based on scientific procedures [see Appendix A, "EXPERT PANEL CONSENSUS STATEMENT REGARDING THE GENERALLY RECOGNIZED AS SAFE (GRAS) STATUS OF SWEET LUPIN-DERIVED INGREDIENTS FOR USE IN FOODS]. A summary of these data is presented herein.

B. Estimated Intake of Sweet Lupin Flour

As mentioned, sweet lupin flour is intended for use in a variety of food products, including baked goods and baking mixes and grain products and pastas. The individual proposed food uses and use levels are summarized in Table 2.
Table 2 Summary of the Individual Proposed Food Uses and Use Levels for Sweet Lupin Flour in the United States

<table>
<thead>
<tr>
<th>Food Category*</th>
<th>Proposed Food Use</th>
<th>Serving Size (g or mL)</th>
<th>Maximum Use Level (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Baked Goods and Baking Mixes</td>
<td>Bagels</td>
<td>55^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Biscuits</td>
<td>55^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Cakes</td>
<td>50</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Cookies</td>
<td>30 to 40^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Combread, Corn Muffins, and Tortillas</td>
<td>55^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Crackers</td>
<td>15 to 30^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Croissants</td>
<td>55^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>English Muffins</td>
<td>50^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>French Toast, Pancakes, Waffles, and Crepes</td>
<td>85 to 110^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Muffins and Popovers</td>
<td>55^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Pastries</td>
<td>55 to 125^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Pies</td>
<td>125^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Quick Breads and Sweet Rolls</td>
<td>55^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Soft Bread Sticks</td>
<td>55^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Soft Pretzels</td>
<td>55^b</td>
<td>25</td>
</tr>
<tr>
<td></td>
<td>Yeast Breads and Rolls</td>
<td>60</td>
<td>25</td>
</tr>
<tr>
<td>Grains Products and Pastas</td>
<td>Macaroni and Noodle Products</td>
<td>240^b</td>
<td>25</td>
</tr>
</tbody>
</table>

* The food categories used to estimate the mean and 90th percentile daily intakes were obtained from National Center for Health Statistics' (NCHS) 2003-2004 National Health and Nutrition Examination Surveys (NHANES) (CDC, 2006; USDA, 2008).

^b Serving size reported was based on Reference Amounts Customarily Consumed Per Eating Occasion (RACC) (21 CFR §101.12) (U.S. FDA, 2008b). When a range of values is reported for a proposed food use, particular foods within that food use may differ with respect to their RACCs.

The consumption of sweet lupin flour from all proposed food-uses was estimated using the National Center for Health Statistics’ (NCHS) 2003-2004 National Health and Nutrition Examination Surveys (NHANES) (CDC, 2006; USDA, 2008) to estimate mean and 90th percentile daily intakes. When a range of values is reported for a proposed food use, particular foods within that food use may differ with respect to their Reference Amounts Customarily Consumed Per Eating Occasion (RACC) (21 CFR §101.12) (U.S. FDA, 2008b).

Humans and livestock have consumed lupin and lupin-derived ingredients for over 2,000 years, and various species and varieties of lupin have been historically cultivated in the Mediterranean region, northern Europe, South Africa, Australia, and New Zealand, and more recently in the southeastern United States. Sweet lupin has a crude protein level similar to that of soybeans, but contains much lower levels of potential anti-nutritional factors (ANFs), and hence has been recognized as a valuable protein source (Ballester et al., 1980; Petterson and Crosbie, 1990; Petterson, 1995). Lupin-derived ingredients are permitted for use in food for human consumption in the European Union (EU) and Australia/New Zealand (Allen, 1992; Weston Technologies, personal communication, 2005), and lupin flour is used in Europe in bread (up to 10%), pastas, cakes, and biscuits (up to 50%) (Belteky and Kovacs, 1984). In addition, lupins are enjoying application in Asia for modified traditional cultural dishes such as miso, tempeh, and tofu, as the yield of fermented products from lupin protein fractions has been reported to be greater than that for soybean (Petterson and Crosbie, 1990). Despite the documented historical consumption of lupin, quantitative consumption data were not identified.

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The consumption of sweet lupin flour from all proposed food-uses was estimated using the National Center for Health Statistics’ (NCHS) 2003-2004 National Health and Nutrition Examination Surveys (NHANES) (CDC, 2006; USDA, 2008).
Examination Surveys (NHANES) (CDC, 2006; USDA, 2008). Sweet lupin flour is proposed for use as a food ingredient at a maximum level of 25%, and the estimated daily consumption of the sweet lupin flour ingredient from all proposed food uses at the proposed use levels per serving was calculated on a g and g per kilogram body weight basis by population group.

On an all-user basis, the total population mean and 90th percentile intakes of sweet lupin flour were estimated to be 39.8 g/person/day (0.7 g/kg body weight/day) and 75.3 g/person/day (1.5 g/kg body weight/day), respectively. Of the individual population groups, the greatest mean all-user intake of sweet lupin flour on an absolute basis was estimated to be in male teenagers at 47.3 g/person/day. Infants had the lowest estimated intake of sweet lupin flour on an absolute basis, with a mean all-user intake of 21.8 g/person/day (1.8 g/kg body weight/day). On a body weight basis, estimated mean all-user intakes of sweet lupin flour were highest in infants (1.8 g/kg body weight/day) and children (1.5 g/kg body weight/day), and lowest in female and male adults (each at 0.5 g/kg body weight/day).

When heavy consumers (90th percentile) were assessed, all-user intakes of sweet lupin flour from all proposed food-uses also were estimated to be greatest in male teenagers (36.5 g/ person/day) and male adults (33.3 g/person/day), and lowest in infants (43.4 g/person/day) on an absolute basis. On a body weight basis, the highest estimated all-user 90th percentile intakes of sweet lupin flour were observed in infants (3.6 g/kg body weight/day) and children (2.8 g/kg body weight/day). The lowest all-user 90th percentile intakes of sweet lupin flour on a body weight basis were estimated to occur in male (1.1 g/kg body weight/day) and female (0.9 g/kg body weight/day) adults. A summary of the estimated all-person and all-user mean and 90th percentile intakes of sweet lupin flour by individual population groups and for the total population is presented in Tables 3 and 4.

<table>
<thead>
<tr>
<th>Population Group</th>
<th>Age Group (Years)</th>
<th>% Users</th>
<th>Actual # of Total Users</th>
<th>All-Person Consumption (g) Mean 90th Percentile</th>
<th>All-Users Consumption (g) Mean 90th Percentile</th>
</tr>
</thead>
<tbody>
<tr>
<td>Infant</td>
<td>0 to 2</td>
<td>74.6</td>
<td>694</td>
<td>17.9</td>
<td>40.9</td>
</tr>
<tr>
<td>Children</td>
<td>3 to 11</td>
<td>99.1</td>
<td>1,275</td>
<td>39.1</td>
<td>70.4</td>
</tr>
<tr>
<td>Female Teenager</td>
<td>12 to 19</td>
<td>96.2</td>
<td>974</td>
<td>38.1</td>
<td>76.5</td>
</tr>
<tr>
<td>Male Teenager</td>
<td>12 to 19</td>
<td>96.6</td>
<td>985</td>
<td>46.5</td>
<td>85.6</td>
</tr>
<tr>
<td>Female Adult</td>
<td>20 and Up</td>
<td>98.3</td>
<td>2,092</td>
<td>34.7</td>
<td>65.2</td>
</tr>
<tr>
<td>Male Adult</td>
<td>20 and Up</td>
<td>99.1</td>
<td>1,912</td>
<td>45.0</td>
<td>82.6</td>
</tr>
<tr>
<td>Total Population</td>
<td>All Ages</td>
<td>96.0</td>
<td>7,932</td>
<td>39.0</td>
<td>74.6</td>
</tr>
</tbody>
</table>

IV. BASIS FOR GRAS DETERMINATION
George Weston Foods Limited
September 26, 2008
<table>
<thead>
<tr>
<th>Population Group</th>
<th>Age Group (Years)</th>
<th>% Users</th>
<th>Actual # of Total Users</th>
<th>All-Person Consumption (g/kg body weight)</th>
<th>All-Users Consumption (g/kg body weight)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Mean</td>
<td>90th Percentile</td>
</tr>
<tr>
<td>Infant</td>
<td>0 to 2</td>
<td>74.6</td>
<td>694</td>
<td>1.5</td>
<td>3.2</td>
</tr>
<tr>
<td>Children</td>
<td>3 to 11</td>
<td>99.1</td>
<td>1,275</td>
<td>1.5</td>
<td>2.8</td>
</tr>
<tr>
<td>Female Teenager</td>
<td>12 to 19</td>
<td>98.2</td>
<td>974</td>
<td>0.7</td>
<td>1.4</td>
</tr>
<tr>
<td>Male Teenager</td>
<td>12 to 19</td>
<td>98.6</td>
<td>985</td>
<td>0.7</td>
<td>1.4</td>
</tr>
<tr>
<td>Female Adult</td>
<td>20 and Up</td>
<td>98.3</td>
<td>2,092</td>
<td>0.5</td>
<td>0.9</td>
</tr>
<tr>
<td>Male Adult</td>
<td>20 and Up</td>
<td>99.1</td>
<td>1,912</td>
<td>0.5</td>
<td>1.1</td>
</tr>
<tr>
<td>Total Population</td>
<td>All Ages</td>
<td>96.0</td>
<td>7,932</td>
<td>0.7</td>
<td>1.4</td>
</tr>
</tbody>
</table>

Sweet lupin flour consists mainly of protein (minimum of 35%) and fiber (35 to 42% soluble and insoluble fiber combined), with lesser amounts of other carbohydrates (7 to 9%) and fat (7 to 10%), and all major components of the sweet lupin flour ingredient (i.e., protein, fiber, etc.) are macronutrients that are part of a normal human diet. The source excluded, the composition of sweet lupin flour is similar to that of standardized (wheat) flour (21CFR§137.105) (U.S. FDA, 2008c). It is therefore useful to compare the estimated intake of sweet lupin flour to the background consumption of wheat flour in the United States, which was reported to be 165 g/person/day (Wheat Foods Council, 2005). The estimated 90th percentile all-user consumption for sweet lupin flour was greatest in the male teenager population group, with an intake of 86.5 g/day, which is almost 2-fold lower than the reported average wheat flour consumption. Therefore, when considering the background intake of standardized flour and that the intended uses of sweet lupin flour will replace a portion other sources of flour (including wheat) in the diet, the estimated intake of lupin flour is not expected to pose any concern.

C. Metabolic Fate of Sweet Lupin Flour

With respect to the metabolic fate of sweet lupin flour, the digestion and subsequent absorption, distribution, metabolism, and excretion of the ingredient is relevant to the metabolic fate of its macronutrient constituents. As mentioned, sweet lupin flour consists mainly of protein (minimum of 35%) and fiber (35 to 42%), with lesser amounts of other carbohydrates and additional fat. The major macronutrients present in the sweet lupin flour ingredient are expected to undergo normal metabolism. Following consumption, the protein components of the sweet lupin flour ingredient are expected to be denatured in the stomach by acid and/or cleaved by enzymes, and the dietary fibers are expected to pass relatively intact into the large intestine following consumption, where they will be subjected to fermentation.
D. Pre-Clinical Studies Pertaining to the Safe Consumption of Sweet Lupin Flour

Some traditional toxicological studies of oral exposure to lupin seed and/or ingredients derived thereof were identified in the published literature, including a subchronic oral toxicity study of lupin flour in rats, and these data support the safety of sweet lupin flour under the intended conditions of use. In addition, several studies designed to assess the nutritional equivalence of lupin seed and lupin-derived ingredients to other traditional seed crops and its acceptability for use as an alternate feed source for food-producing animals were identified, and these provide additional support for the safe consumption of GWF’s sweet lupin flour under the intended conditions of use. Summaries of these studies are presented below.

Acute Toxicity Studies

Sweet lupin and lupin fractions were reported to have low acute oral toxicity in rats, with reported oral LD₉₀ values ranging from 750 to >4,000 mg/kg body weight for *L. angustifolia* and *L. albus* whole seed and seed fractions (Stobiecki et al., 1993). No adverse effects were reported in rats following administration of a single gavage dose of conglutinin γ, a lupin seed protein isolated from *L. albus*, at levels of up to 120 mg/kg body weight, the highest dose tested (Magni et al., 2004). Moreover, lupin protein extracted from the seeds of *L. albus* and administered by gavage to male Sprague-Dawley rats for a period of 2 weeks at a dose of 250 mg lupin protein/kg body weight/day did not result in any adverse effects (Sirtori et al., 2004).

Short-term Toxicity Studies

In a number of short-term toxicity studies conducted by one research group and using the same experimental design, male Hooded-Listar rats (4 to 20/group) were provided diets containing *L. angustifolia* seed or protein fractions for 10 days (Rahman et al., 1996a,b, 1997a). The diets included: one with whole lupin seeds (supplemented or unsupplemented with essential amino acids) providing 31.5 g lupin/kg body weight/day; 1 with a soluble (LPAD) and 1 with an insoluble (LPADI) aqueous dialyzed protein fraction, providing 13.0 and 10.9 g lupin/kg body weight/day, respectively; 1 with a non-dialyzed (LPAND) aqueous protein fraction providing 17.1 g lupin/kg body weight/day; 1 with a soluble (BUSOL) and one with an insoluble (BUDI) buffer-extracted fraction providing 11.9 and 10.4 g lupin/kg body weight/day; and 1 diet containing a dialyzed residue fraction (LMR) (i.e., the fibrous material that is insoluble in both water and buffer) providing 14.9 g lupin/kg body weight/day. A diet containing lactalbumin was provided to a separate group of rats (control group) in each study. An additional 10-day study included rats supplemented or unsupplemented with whole lupin seed or LPADI in the diet at levels of 28 or 9.7 g whole seed and LPADI/kg body weight/day, respectively (Rahman, 2000). Observed effects from these studies, such as decreased body weight gain, increased urea, and decreased albumin, were suggested by the study authors to be due to a disturbance of normal protein utilization, which could have resulted from the amino acid deficiency of the lupin-containing diets, as lupin is known to
contain low levels of essential amino acids, despite supplementation of the diets with amino acids. In a clinical study by Egaña et al. (1992), lupin protein digestibility was reported to be good, and therefore a disturbance of protein utilization is not likely to occur in humans consuming an average diet. Moreover, the reported increases in plasma urea remained within reported historical control values for rats (Sharp and LaRegina, 1998), the reported increases in serum alkaline phosphatase (AP) values in lupin-treated rats were not accompanied by significant differences in the alanine aminotransferase (ALAT) or aspartate aminotransferase (ASAT) levels compared to the control group, and liver lesions were not reported in any of the other reviewed dietary studies. Rahman (2000) evaluated spleen and thymus weights, and spleen weights were reported to be significantly reduced in the LPADI group compared to the all other groups. The authors suggested that uremia might have been the cause of the immunesuppression, characterized by a significant decrease in spleen weight. The results of the study by Rahman et al. (1996b) indicated that the stomachs of rats provided unsupplemented and supplemented whole lupin seed were distended due to undigested food material and their colons were reportedly enlarged compared to lactalbumin controls. Rats in the LMR group also were reported to have enlarged spleens and colons, although no changes in abdominal organs were reported in any of the other lupin groups. Rahman et al. (1996a,b, 1997a) and Rahman (2000) did not discuss the significance of these effects, nor was the frequency of any of these effects reported. Furthermore, this study involved a number of different experimental parts, and it is unclear if organ weights and gross and microscopic examinations were conducted in only one of the experimental study parts (resulting in examination of only 4 rats/group), or in 5 different experimental parts (resulting in examination of 4 to 20 rats/group). Nonetheless, as previously mentioned, the frequency of the observed effects was not reported, and effects in the lactalbumin control group were not always reported as a means for comparison. Gross and microscopic changes were reported in the livers of all lupin-treated groups; however, liver weights were not measured. Overall, the results of these studies are poorly reported and the significance of the observed effects is unclear.

Subchronic Toxicity Study

A 90-day toxicity study in rats was identified in which the animals were fed diets containing *L. angustifolia* lupin flour spiked with lupin alkaloids providing 0 (control), 250, 1,050, or 5,050 ppm of supplemental alkaloids (Butler et al., 1996). The source of lupin used in this study was from the same agronomic area (i.e., Australia) as GWF's lupin source. The control group of this study was provided a diet containing 13.2 g lupin flour (up to 33 g/kg body weight/day), which contained a background level of ~50 ppm alkaloids (6.6 mg alkaloids/kg body weight/day), a level similar to that present in the GWF sweet lupin flour ingredient, and therefore the results of this group are considered relevant to the safety of sweet lupin flour. The group of interest is the control group in this study; however, an 'untreated' group (i.e., not provided lupin) was not available for which to compare results, and hence historical values in the rat were utilized to assess any potentially adverse effects resulting from 90-day dietary supplementation with lupin flour (Butler et al., 1996). The historical values for control rats from the laboratory in which the study was conducted were...
sought without success, and hence the data of Sharp and LaRegina (1998) were used for the purpose of this assessment. No deaths or clinical signs of toxicity were reported in the lupin flour group, and there were no significant differences in biochemical or hematological parameters or organ weights in lupin-treated rats when compared to historical values in the rat (Brown et al., 1997; Sharp and LaRegina, 1998). Furthermore, there were no histological findings in any of the organs evaluated. For the purpose of this assessment, a no-observed-effect level (NOEL) of 33 g/kg body weight/day was derived for lupin flour, which was the only dose of low-alkaloid lupin flour tested.

**Chronic Feeding Studies**

Chronic studies in rats ranging from 700 to 800 days in duration and using *L. angustifolia* seed and seed fractions were identified in the available literature (Ballester et al., 1980, 1984; Grant et al., 1993, 1995). These studies were nutritional studies and not traditional toxicity studies; however, the results of these studies support the safety of dietary lupin and lupin ingredients. Consumption of 5.6 g of whole *L. angustifolia* seed/kg body weight/day in the diet, increasing to a maximum intake of approximately 13.6 g/kg body weight/day after 15 weeks, was reported not to cause any adverse effects in rats when administered for up to 800 days (Grant et al., 1993). Body weight gain was significantly reduced in lupin-treated rats compared to controls for the first 200 days, however, was not significantly different from controls for the remainder of the study. Furthermore, lupin seed was reported to have no significant effect on pancreatic weight or composition (Grant et al., 1993). These same doses of whole lupin seed in the diet were reported to decrease body weight gain in rats dosed for up to 700 days (Grant et al., 1995). Cecum and colon weights were significantly increased in lupin-fed rats compared to control rats after 700 days of feeding, which the authors stated was not mediated by either lectin or protease inhibitors, but rather may have been the result of volatile fatty acid production due to dietary fiber digestion in these organs.

**Mutagenicity/Genotoxicity and Carcinogenicity Studies**

Studies of the mutagenic/genotoxic potential of lupin or its fractions were not identified in the literature, nor were traditional carcinogenicity studies; however, as previously mentioned, chronic/life-time studies (*i.e.*, 700 and 800 days) in rats did not reveal any evidence of carcinogenicity in lupin-treated animals, and no signs of toxicity or decreases in body weight occurred (Grant et al., 1993, 1995).

**Nutritional Studies**

Conflicting results were reported upon investigation of the potential effect of *L. angustifolia* seed and seed fractions on mineral absorption in rats, chickens, and pigs (Rubio et al., 1994; Rahman et al., 1997b; Olkowski et al., 2005; Zraly et al., 2006, 2007). Rubio et al. (1994) reported that dialyzed soluble and insoluble lupin protein fractions had no significant effect on absorption of calcium, phosphorus, or zinc in male Hooded-Listar rats, while in the same species, whole lupin seed and the LMR fraction significantly reduced phosphorus and zinc absorption due to the presence of phytate and insoluble non-starch polysaccharide fractions.
that are not present in protein fractions. Conversely, Rahman et al. (1997b) reported that equivalent doses of various lupin protein fractions, including dialyzed soluble and insoluble fractions, significantly reduced absorption of phosphorus, zinc, magnesium, and sodium (with no effects on calcium or potassium) in male Hooded-Listar rats; however, these effects were not discussed by the authors. Zraly et al. (2006) reported that plasma phosphorus concentrations in pigs were not significantly affected by the consumption of dehulled *L. angustifolia* seed meal for a period of 90 days, and the same authors also reported no significant changes in plasma calcium or phosphorus levels in pigs administered diets containing *L. albus* seed for 90 days (Zraly et al., 2007). Additionally, no significant differences in plasma zinc concentrations were reported to occur in broiler chicks from the consumption of raw or dehulled *L. albus, L. luteus,* or *L. angustifolius* seed meal for a period of 21 days (Olkowski et al., 2005). Plasma zinc concentrations in broiler chicks provided diets containing raw or dehulled lupin seed meal from *L. albus, L. luteus,* or *L. angustifolius* at doses of approximately 500 g/kg body weight/day (raw) or ranging from 252 to 650 g/kg body weight/day (dehulled) for 21 days were not significantly different from the values of control birds fed a soybean meal diet; however, plasma riboflavin concentrations were significantly increased in all lupin-fed chicks combined compared to the controls (Olkowski et al., 2005).

Sweet lupin seeds are widely used in Australia as a source of protein and energy in livestock feeds, and hence, their nutritive value has been evaluated in various feeding studies in pigs and poultry. Overall, nutritional studies in pigs and chickens indicate that lupin feeds are generally well tolerated (Dunshea et al., 2001; Rubio et al., 2003; Steenfeldt et al., 2003; Hammershoj and Steenfeldt, 2005; Martins et al., 2005; Olkowski et al., 2005; Zraly et al., 2006, 2007); however, due to the generally low levels of both methionine and lysine in lupin (Pettersson, 1998), feeds for pigs and poultry are more beneficial when they include multiple sources of protein, or supplemental amino acids (Edwards and van Barneveld, 1998). Transgenic lupin seeds have been reported to significantly improve the nutritive value of lupin as they have been modified to encode a protein which contains 16% methionine and 8% cysteine residues (Molvig et al., 1997, 2003). Furthermore, digestible energy from lupin may be compromised by ANFs (e.g., trypsin inhibitors) by interfering with digestive enzymes in monogastrics (Edwards and van Barneveld, 1998). A summary of the results of the identified nutritional studies in pigs and poultry is provided below.

**Administration of whole *L. angustifolia* seed or kernel to pigs via the diet during a 14-day feeding study, which provided a dose of 16 g lupin seed or kernel/kg body weight/day, was reported not to produce any adverse effects (Dunshea et al., 2001). Similarly, doses of 8.8 g of whole *L. angustifolia* or *L. albus* seed or kernel/kg body weight/day provided for 14 days did not result in any adverse effects (Dunshea et al., 2001). Feed intake was increased in both intact and ileorectal anastomosed pigs provided doses of 10.64 and 10.30 g**

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4 See Section F for further discussion of the phytonutrient components identified in lupin and their relevance to the safety of sweet lupin flour.
SWEET LUPIN FLOUR GRAS NOTICE

L. angustifolia seed/kg body weight/day, respectively, in the diet for a period of 3 weeks compared to pigs provided nutritionally-equivalent cholesterol-enriched casein control diets; however, there was no significant difference in body weight gains between groups (Martins et al., 2005). The difference in feed intake was likely due to unpalatability of the cholesterol-enriched control diet. Significantly decreased relative liver weights were reported in pigs provided lupin protein compared to control pigs; however, there was no significant difference in gallbladder weight between groups (Martins et al., 2005). Body weight gain was not affected following administration of 13.2 and 14.3 g of L. angustifolia seed and kernel/kg body weight/day, respectively, in the diet to pigs for 28 days, but was significantly decreased with doses of 10.2 and 10.9 g/day of L. albus seed and kernel, respectively, in the diet for the same period of time (Dunshea et al., 2001). Neither lupin diet had any significant effect on liver weight. In another feeding study, histological examination of the livers and kidneys of pigs revealed no gross lesions following administration of 1 of 3 different diets containing a combination of 2 different varieties of L. angustifolia seed levels providing doses of up to 26.9 g lupin seed/kg body weight/day and up to 23.7 mg alkaloids/kg body weight/day for a period of 7 weeks (Godfrey et al., 1985). Body weight gain and feed conversion in Large White x Landrace pigs were not affected by the administration of diets containing approximately 2.3 g dehulled lupin (L. angustifolius) seed meal/kg body weight/day for a period of 90 days (Zraly et al., 2006), although significant decreases in plasma glucose and calcium, and a significant increase in total protein were reported. Body weight gain, feed intake, and plasma and calcium levels were not affected in hybrid P x (Du x LW x L) pigs administered a diet containing approximately 4.1 g lupin (L. albus) seed/kg body weight/day for 90 days compared to animal or soy protein-fed controls (Zraly et al., 2007) or in (LW x L) x D pigs fed 5.4 and 5.1 g/kg body weight/day raw and extruded lupin (L. albus), respectively, for 42 days (Prandini et al., 2005).

Performance and some biochemical measures of toxicity were evaluated in (LW x L) x D piglets (16 males and 12 females/group, average initial body weight of 10.4 kg) weaned at 28 days of age and administered basal diets supplemented with 170 g/kg raw or extruded lupin (L. albus) seeds (providing approximately 5.4 and 5.1 g/kg body weight/day raw and extruded lupin, respectively) ad libitum for a period of 42 days (Prandini et al., 2005). No significant differences in total bilirubin or ALAT, ASAT, or AP activity were reported to occur in blood samples taken from the lupin-fed pigs at Day 42 compared to the controls; however, significant decreases in total protein and urea were reported to occur in the lupin-fed pigs compared to the controls. In 2 studies of longer duration (90 days), total protein, albumin, AP, ASAT, and ALAT levels were not significantly different between control pigs and pigs administered a diet containing approximately 2.3 g dehulled lupin (L. angustifolius)/kg body weight/day or a diet containing 4.1 g lupin (L. albus) seed/kg body weight/day (Zraly et al., 2006, 2007).

L. albus seed (unsupplemented) provided to pigs at levels of 20.7 or 31% in the diet significantly reduced growth rates, although this effect was not observed in animals provided a lower level (10.3% lupin in the diet) or in pigs provided 31% lupin in the diet supplemented with 0.2% lysine (duration not specified) (King, 1981). The authors reported that 20.7 and
31% lupin diets were deficient in lysine and therefore growth rates were significantly reduced compared to control pigs (King, 1981). \textit{L. albus} is therefore not recommended for use in pig feeds due to recognized reductions in feed intake and depressed growth rates (Edwards and van Barneveld, 1998).

Feed intake was reduced in hens provided up to 25% lupin seed in the diet for 11 weeks (providing 15 g lupin seed/kg body weight/day), although there were no significant effects on body weight (Hammershoj and Steenfeldt, 2005). Broiler chickens dosed with 1,006 g lupin seed/kg body weight/day in the diet for 14 days were reported to have decreased body weight gain, although there was no effect on feed consumption. The authors attributed the effects on body weight to a lack of appropriate digestive enzymes in chickens (Steenfeldt et al., 2003). Rubio \textit{et al.} (2003) reported that whole lupin seed provided in chicken feed at levels of up to 1,540 g/kg body weight/day for 21 days decreased both feed intake and body weight gain; however, these effects were not observed with dehulled lupin seeds consumed at a level of 1,347 g/kg body weight/day for 21 days. Olkowski \textit{et al.} (2001) examined the effects of raw, autoclaved, and dehulled lupin (\textit{L. angustifolius}) seed meal (approximately 546, 445, and 584 g/kg body weight/day, respectively) in the diet of broiler chicks for a period of 21 days and reported significantly decreased feed intake and body weight gain in all lupin-fed chicks. Raw lupin seed meal from \textit{L. albus}, \textit{L. luteus}, or \textit{L. angustifolius} at doses of approximately 500 g/kg body weight/day (raw) or ranging from 252 to 650 g/kg body weight/day (dehulled) for 21 days also resulted in significantly decreased feed intakes and growth rates compared to the controls. Dehulling was reported to significantly increase body weight gain, but the level remained significantly lower than the control group (Olkowski \textit{et al.}, 2005). Conversely, Ross 308 broiler chicks provided dehulled lupin seeds of the variety JUNO (\textit{L. luteus}) at an average level of 16.4 g/kg body weight/day for 40 days reached body weights similar to those of the control chicks fed a diet containing soy extract; chicks provided dehulled lupin seeds of the variety SONET (\textit{L. angustifolius}), however, had significantly decreased final body weights (Suchy \textit{et al.}, 2006).

Significant dose-dependent increases in relative gizzard weights were reported in Leghorn chicks receiving diets containing whole lupin (\textit{L. albus}) seeds at levels of up to 70% for a period of 14 days, and Leghorn chicks that received diets containing dehulled lupin seeds supplemented with lupin hulls had significant increases in relative intestinal organ weight and length compared to chicks that received the lupin diet without the addition of hulls (Brenes \textit{et al.}, 2002). Similarly, it was reported by the same authors that broiler chicks fed diets containing 35 and 45% whole lupin (\textit{L. albus}) (approximately 599 and 762 g lupin/kg body weight/day) for 6 weeks had significantly increased relative weights of the crop, proventriculus, gizzard, and duodenum compared to the control group receiving a wheat-soy diet (Brenes \textit{et al.}, 2002), and significant increases in the size of the duodenum, jejunum, and ileum were reported in broiler chicks fed diets containing either 40% raw (approximately 500 g/kg body weight/day) or 35% dehulled (approximately 425, 252, and 650 g/kg body weight/day for \textit{L. albus}, \textit{L. luteus}, and \textit{L. angustifolius}, respectively) lupin seed meal for 21 days when compared to chicks fed a soybean meal control diet (Olkowski \textit{et al.}, 2005); however, the relative weights of the liver, pancreas, gizzard, and heart were not significantly...
different from control values in Ross 308 broiler chicks fed wheat- and barley-based diets containing up to 20% lupin (L. luteus) for 6 weeks (Orda et al., 2006). Enlargement of some gastrointestinal organs may be interpreted as a physiological adaptation to overcome ANFs present in the lupin-based diets (Olkowski et al., 2005).

E. Studies in Humans

Data relating specifically to the safety of lupin flour consumed by human volunteers were identified in the studies conducted by Gattás Zaror et al. (1990) and Egāña et al. (1992) in which healthy volunteers consumed L. albus flour-enriched products. In a crossover study, Gattás Zaror et al. (1990) provided one 150 g cookie/day, with or without lupin flour (providing 35 and 0 g lupin flour/day and containing 13.3 and 0 g lupin protein, respectively) for a treatment period of 60 days. No compound-related changes were reported in any of the biochemical or hematological parameters tested [i.e., hematocrit, hemoglobin, prothrombin, uric acid, urea nitrogen, bilirubin, glutamic-pyruvic transaminase (GPT), ASAT, blood lipids, and creatinine], although body weight was significantly increased in both groups. The authors reported that lupin flour was well tolerated by the subjects. Egāña et al. (1992) supplemented the diet of young men (n=9) with lupin flour derived from L. albus, providing a dose of 0.4, 0.6, or 0.8 g lupin protein/kg body weight/day for a period of 10 days, which would correspond to 28, 42, and 56 g/day for the average 70 kg person. Nitrogen digestibility, complete blood count, serum total protein, albumin, urea nitrogen, globulin, ASAT, ALAT, cholesterol, and triglycerides were evaluated at the end of the study period, although hematological parameters were only measured in the low- and high-dose groups. Nitrogen digestibility was reported to range between 78.8 and 70.2%. The only significant hematological change reported was a significant increase in urea nitrogen in the high-dose group (0.8 g lupin protein/kg body weight/day) compared to the low-dose group (0.4 g lupin protein/kg body weight/day). The authors reported that lupin-containing diets were well tolerated by the subjects and were without adverse effects.

Additional studies designed to investigate parameters such as glycemic index, insulin response, plasma ghrelin response, as well as satiety and palatability indicated that ingredients derived from L. angustifolia were well tolerated in healthy volunteers and without adverse effects (Hall and Johnson, 2004; Hall et al., 2005; Lee et al., 2006). The identified studies supplied either a single dose or 2 doses in one day that provided 7.7 to 264 g lupin flour derived from L. angustifolia/serving.

F. Other Data Pertaining to the Safety of Sweet Lupin Flour

Various phytonutrients (i.e., oligosaccharides, phenolics and condensed tannins, trypsin inhibitors, phytic acid, saponins, and lectins) occur naturally in L. angustifolia, L. albus, and L. luteus at very low levels and are comparable to levels present in other grain legume species (Petterson, 1998). The possible effects of exposure to these compounds under the intended conditions of use of the sweet lupin flour ingredient of GWF are discussed below.
Moreover, lupin has recently been recognized as a potential food allergen, and therefore, the possible allergenicity of sweet lupin flour also has been considered and is discussed below.

**Other Phytonutrient Components**

As a result of their natural presence in lupin, possible additional components occurring in the final sweet lupin flour ingredient are oligosaccharides, phenolics and condensed tannins, trypsin inhibitors, phytic acid, saponins, and lectins, may occur in the final sweet lupin flour ingredient (see Appendix B-4 for results of analysis). These compounds are reported to occur naturally in sweet lupin varieties at very low levels and are comparable to levels found in other grain legume species (Pettersen, 1998).

The oligosaccharides present in lupin belong to the raffinose family and are considered to be ANFs because they cannot be metabolized by monogastrics (Pettersen, 1998). Oligosaccharides occur naturally in sweet lupin at levels of 5.2 to 11.87% (Pettersen, 1998). Following batch analysis, the level of oligosaccharides in the sweet lupin flour ingredient was determined to be 5.6% (dry solid basis). Based on the estimated total population all-user 90th percentile intake of sweet lupin flour (75.3 g/person/day), a maximum intake of 4.22 g oligosaccharide/person/day was calculated. Considering that the method of calculating the estimated intakes of the sweet lupin flour ingredient under the recommended conditions of use is 'worst-case', the actual intake of oligosaccharides will likely be much lower, and hence is not expected to produce adverse effects on human health.

Phenolic compounds are reported to have the potential to bind iron and decrease iron absorption (Disler et al., 1975; Brune et al., 1989; Hurrell et al., 1999), and condensed tannins have an affinity for binding proteins (Ricardo da Silva et al., 1991; Vallet et al., 1994; Santos-Buelga and Scalbert, 2000). The background dietary intake of phenolics (as flavonoids) from various sources, such as coffee, cocoa, red wine, and many fruits, was reported to be 1,000 mg/day, with condensed tannin intakes of 250 to 460 mg/day (Kühnau, 1976; Santos-Buelga and Scalbert, 2000). Following batch analysis, the sweet lupin flour ingredient was determined to contain phenolics at a level of 0.324%, corresponding to a maximum exposure of ~244 mg/person/day, which is a small increase over the background daily intake, and is expected to have no effect upon iron absorption. The level of condensed tannins as a component of the overall total phenolic(s) levels identified in the sweet lupin flour ingredient is negligible (<0.05%), and hence is expected not to produce any adverse effects on human health.

Trypsin inhibitors are present in whole lupin seed (<0.01 to 0.29 mg/g protein) (Pettersen, 1998) at levels which are several-fold lower than the amount occurring naturally in soybeans (34.30 to 56.14 mg/g protein), soy protein isolates (1.11 to 4.49 mg/g protein), and commercial infant soy formulas (2.2 to 15.5 mg/g protein) (Peace et al., 1992). The authorized health claim on the association between soy protein and reduced risk of coronary heart disease (CHD) includes a qualifying level of a total daily intake of 25 g soy protein for CHD risk reduction claim (U.S. FDA, 1999). Using a reported level of up to 4.49 mg trypsin inhibitors/g soy protein isolate, individuals consuming 25 g soy protein/day could be exposed
to levels of trypsin inhibitors of up to 1,400 mg/day. Analysis of sample batches of sweet lupin flour indicated that trypsin inhibitors occur at a level of 1.45 mg/g, which would provide an exposure of ~109 mg trypsin inhibitors/person/day. This level is almost 13 times less than the estimated exposure to trypsin inhibitors from consumption of 25 g soy protein/day. Therefore, the levels of trypsin inhibitors present in the sweet lupin flour ingredient are anticipated not to produce any adverse effects on human health.

Phytate is an ANF that can form insoluble complexes with cations, such as calcium and zinc, making them less available for absorption and utilization (Petterson, 1998). Phytate occurs naturally in sweet lupin at levels of 0.58 to 0.96% (Petterson, 1998), and is present in soybeans in the range of 1 to 2% (Wang and Wixon, 1999). In a review of studies investigating phytate isolated from soybeans and other studies of the effects of soy protein on iron and zinc status, the FDA concluded that evidence of potential adverse effects of phytic acid is equivocal, and noted that many other factors affect the absorption of these minerals (U.S. FDA, 1999). Analysis of sample batches of sweet lupin flour indicated levels of phytate of 0.50%, which would provide an exposure of ~377 mg phytate/person/day. This level is at least half of the estimated exposure to phytate from soybean, and therefore the level of phytate present in the sweet lupin flour ingredient is anticipated not to produce any adverse effects on human health.

Saponins are considered to be ANFs because they can lyse red blood cells (RBCs). Saponins occur naturally in L. albus at negligible levels and in L. angustifolia at levels of 480 to 730 ppm (Petterson, 1998). These compounds also are present in soybeans at levels in the range of 1 to 5 mg/g dry weight (Anderson and Wolf, 1995; Wang and Wixon, 1999). Saponins have been consumed for many years as part of the human diet without reports of ill effects. Additionally, saponins are poorly absorbed and are considered to be of low oral toxicity (Price et al., 1987; Wang and Wixon, 1999). Moreover, no adverse effects were reported in chicks, rats, or mice fed concentrations of saponins from soy that were 3- to 5-fold greater than a typical soybean meal diet (Ishaaya et al., 1969). Following batch analysis, the level of saponins in the sweet lupin flour ingredient was determined to be <0.003%, which would provide an exposure of <2.2 mg saponins/person/day. This level is at least 150-fold less than the background levels of saponin reported in L. angustifolia.

Although lectin has been reported to be present naturally in lupin, lectin activity was not detected in either L. angustifolia or L. albus following conventional agglutination assay procedures using a wide variety of red blood cell types (Petterson, 1998). Similarly, batch analysis by GWF indicated that there was no lectin activity following agglutination assays with both sheep and horse red blood cells, and therefore, the potential presence of lectin in the sweet lupin flour ingredient is anticipated not to present any concerns on human health.

Potential Allergenicity

Lupin allergy has recently been recognized with documented cases of anaphylaxis and other allergic reactions following lupin consumption (Hefle et al., 1994; Matheu et al., 1999; Novembre et al., 1999; Moneret-Vautrin et al., 2004; Smith et al., 2004; Radcliffe et al.,...
SWEET LUPIN FLOUR GRAS NOTICE

2005; Rotiroti et al., 2007; Wassenberg and Hofer, 2007). Lupin has recently been added to the list of commonly allergenic foods in the European Union but does not have such status in other parts of the world. Additionally, cross-reactivity between lupin protein allergens and allergens in peanuts has been reported (Hefle et al., 1994; Moneret-Vautrin et al., 1999; Kanny et al., 2000; Faeste et al., 2004; Wüthrich et al., 2004; Costa et al., 2005; Magni et al., 2005a,b; Peeters et al., 2007), although such cross-reactions only occur in a fraction (perhaps as high as 20%) of peanut-allergic individuals. Furthermore, lupin allergy can occur independent of peanut allergy (Peeters et al., 2007). As expected, the results of in vitro studies using GWF’s ingredients indicate that not all peanut-allergic individuals would be sensitive to these lupin-based ingredients (Nordlee, unpublished, 2004). While the levels of protein in the different lupin-based ingredients is variable, sweet lupin flour contains a minimum of 35% protein and is potentially allergenic, although thermal processing of lupin has been reported to decrease the allergic potential of lupin-derived ingredients (Álvarez-Alvarez et al., 2005).

Under the Food Allergen Labeling and Consumer Protection Act of 2004 (FALCPA), if a packaged food product contains, or contains any ingredients derived from, 1 of the 8 major allergenic foods, namely milk, eggs, fish, crustacean shellfish, tree nuts, wheat, peanuts, and soybeans, the presence of the allergenic ingredient must be identified in plain English in the list of ingredients or it should be stated adjacent to the list of ingredients that the product contains the allergenic ingredient. Although lupin is not one of the eight major allergenic foods, there have been documented cases of allergic responses to the consumption of lupin. Therefore, GWF will take steps to ensure that the presence of lupin is identified on all products that contain sweet lupin flour in order to notify consumers and to attempt to prevent exposure in sensitive populations. In order to ensure that lupin is identified on all end product labels for products containing sweet lupin flour either as a direct ingredient or as an incidental additive, GWF will indicate on the specification sheet for sweet lupin flour that the presence of lupin should be disclosed either to the food additive manufacturers’ customers and/or to the end product manufacturer so that lupin will be included on the end product label. Under FALCPA there are no labeling requirements for food ingredients that may elicit responses in individuals who are allergic to one of the eight major food allergens (i.e., there is no requirement for the labeling of the cross-reactive ingredients). As there are currently no requirements for the labeling of ingredients that may be cross-reactive, GWF intends to ensure that lupin is included on the end product label for all products containing the sweet lupin flour ingredient either as a direct ingredient or as an incidental additive rather than include a statement regarding cross-reactivity, as such a statement could be confusing to the consumer. It is therefore expected that the labeling of foods to which the sweet lupin flour ingredient is added should alert the lupin-allergic consumer to the presence of lupin.

G. Summary and Basis for GRAS Conclusion

The results of the identified animal and human studies of whole sweet lupin seed and lupin fractions have been determined by GWF not to indicate any potential for adverse effects in humans following consumption of the sweet lupin flour ingredient under the intended....
conditions of use. Sweet lupin flour is similar in composition to wheat flour, and is composed mainly of protein and fiber, with lower levels of fat and carbohydrate, and all macronutrient components of the sweet lupin flour have a history of consumption as part of a normal diet, with estimated intakes from the intended conditions of use that are within range of their background consumption from various dietary sources. Therefore, following a critical evaluation of scientific data generally available in the public domain that pertain to the safety of lupin and sweet lupin-derived ingredients, including lupin flour, under the intended conditions of use, and derivation of a consensus among a panel of experts who are qualified by scientific training and experience to evaluate the safety of ingredients as components of food that sweet lupin flour would be safe and suitable for use under the proposed conditions and also would be generally recognized as such by other experts, GWF has concluded that sweet lupin flour is GRAS under the intended conditions of use on the basis of scientific procedures.

IV. BASIS FOR GRAS DETERMINATION
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REFERENCES


Grant, G.; Dorward, P.M.; Pusztai, A. 1993. Pancreatic enlargement is evident in rats fed diets containing raw soybeans (Glycine max) or cowpeas (Vigna unguiculata) for 800 days but not in those fed diets based on kidney beans (Phaseolus vulgaris) or lupinseed (Lupinus angustifolius). J Nutr 123(12):2207-2215.


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APPENDIX A

EXPERT PANEL CONSENSUS STATEMENT REGARDING THE
GENERALLY RECOGNIZED AS SAFE (GRAS) STATUS OF SWEET
LUPIN-DERIVED INGREDIENTS FOR USE IN FOODS
EXPERT PANEL CONSENSUS STATEMENT REGARDING THE GENERALLY RECOGNIZED AS SAFE (GRAS) STATUS OF SWEET LUPIN-DERIVED INGREDIENTS FOR USE IN FOODS

September 16, 2008

INTRODUCTION

At the request of George Weston Foods Limited (GWF), an Expert Panel (the “Panel”) of independent scientists, qualified by their relevant national and international experience and scientific training to evaluate the safety of food ingredients, was specially convened to conduct a critical and comprehensive evaluation of the available pertinent data and information relevant to the safety of lupin and sweet lupin-derived ingredients, and determine whether the intended use as food ingredients of 6 ingredients derived from sweet varieties of Lupinus spp. (lupin), including lupin flour, 2 lupin protein fractions, and 3 lupin fiber products, are safe and suitable and would be Generally Recognized as Safe (GRAS), based on scientific procedures. The Panel consisted of the below-signed qualified scientific experts: Prof. Joseph F. Borzelleca, Ph.D. (Virginia Commonwealth University School of Medicine), Ashley S. Roberts, Ph.D. (Cantox Health Sciences International), and Prof. Stephen L. Taylor, Ph.D. (University of Nebraska). Curricula vitae evidencing the Panel members’ qualifications for evaluating the safety of food ingredients are provided in Attachment 1.

The Panel, independently and collectively, critically examined a comprehensive package of scientific information and data pertaining to the safety of lupin and sweet lupin-derived ingredients compiled from the literature and other published sources through July 2007 by Cantox Health Sciences International. In addition, the Panel evaluated other information deemed appropriate or necessary, including data and information provided by Weston Technologies, a division of GWF. The information evaluated by the Panel included details pertaining to the method of manufacture and product specifications, supporting analytical data, intended use-levels in specified food products, consumption estimates for all intended uses, and a comprehensive assessment of the available scientific literature pertaining to the safety of sweet lupin and sweet lupin fractions.

Following independent, critical evaluation of such data and information, the Panel convened on 15 September 2005 and unanimously concluded that the intended uses in traditional foods described herein of ingredients derived from sweet lupin, meeting appropriate food-grade specifications and manufactured consistent with current Good Manufacturing Practice (cGMP), are safe and suitable and GRAS based on scientific procedures. In August of 2007, the Panel evaluated additional data made publicly available since their initial meeting, and in March of 2008 they reviewed an amendment to the initially proposed food uses and use-levels. Subsequently, the Panel reaffirmed their consensus of the safety and suitability and the GRAS status of the intended uses of GWF’s sweet lupin-derived ingredients. A
summary of the basis for the Panel's conclusion, excluding confidential data and information, is provided below.

**SUMMARY AND BASIS FOR GRAS**

GWF intends to market ingredients derived from sweet lupin as food ingredients in various traditional food products such as bakery products, breakfast cereals, and beverages, in the United States. There are 4 species with sweet lupin varieties, namely *L. angustifolia*, *L. albus*, *L. luteus* and *L. mutabilis*, which are 'sweet' due to their low alkaloid content (Petterson, 1998). Lupin and ingredients derived thereof have been consumed by humans and livestock for over 2,000 years, and various species and varieties of lupin have been historically cultivated in the Mediterranean region, northern Europe, South Africa, Australia, and New Zealand, and more recently in the south-eastern United States (Gladstones, 1970; IPK Gatersleben, 2002). Moreover, lupin-derived ingredients are permitted for use in food for human consumption in the European Union and Australia/New Zealand (Allen, 1992; Weston Technologies, personal communication, 2005). Despite the documented historical consumption of lupin, quantitative consumption data has not been identified.

The sweet lupin ingredients are manufactured in accordance with cGMP, and include Sweet Lupin Flour, which is produced by Weston Milling, and Sweet Lupin Protein Fractions 1 and 2, Sweet Lupin Kernel Fibers 1 and 2, and Sweet Lupin Hull Fiber, which are produced by GWF. Essentially, the 6 lupin ingredients are derived from the whole seed of sweet lupin. Sweet Lupin Flour and Sweet Lupin Hull Fiber are obtained by dehulling and milling/grinding the whole lupin seeds. The protein and kernel fiber fractions require further processing to yield the final ingredients. In order to ensure consistent products, GWF has established numerous chemical and microbiological specification parameters for the final preparations, and batch samples are routinely assayed to verify that the specifications are met, ensuring a safe and consistent product. The sweet lupin-derived flour, protein, and fiber ingredients produced by GWF are intended to replace a portion of other sources of flour, protein, and fiber, and due to the self-limiting properties of the ingredients, such as viscosity and baking properties and/or sensory characteristics, the levels of substitution of the flour will be in the range of 10 to 25%, and the levels of use of the protein and fiber ingredients will be up to 20% (see Attachment 2). The ingredients are stable when stored at room temperature (approximately 25°C) in a dry environment, with a shelf life of 6 months.

The consumption of each sweet lupin-derived ingredient from all proposed food uses was estimated using the National Center for Health Statistics' (NCHS) 2003-2004 National Health and Nutrition Examination Surveys (NHANES) (CDC, 2006; USDA, 2008), which provide the most appropriate data for evaluating food-use and food-consumption patterns in the United States. Under the conditions of intended use, the total population all-user mean and 90th percentile intake of Sweet Lupin Flour was estimated to be 39.8 g/person/day (0.7 g/kg body weight/day) and 75.3 g/person/day (1.5 g/kg body weight/day), respectively. The protein fractions contain the same amount of protein (70 to 95%), and of the 2 lupin protein ingredients, Sweet Lupin Protein Fraction 1 has the highest estimated intake, with mean and 90th percentile total population all-user intakes of 18.3 g/person/day (0.3 g/kg body weight/day).
weight/day) and 35.3 g/person/day (0.7 g/kg body weight/day), respectively. Sweet Lupin Kernel Fibers 1 and 2 are intended to be used in the same food categories and at the same levels and were estimated to have a total population all-user mean intake of 35.3 g/person/day (0.6 g/kg body weight/day), and an estimated 90th percentile all-user intake of 64.9 g/person/day (1.2 g/kg body weight/day), from all proposed food-uses. The total population all-user mean and 90th percentile intakes of Sweet Lupin Hull Fiber were estimated to be 14.9 g/person/day (0.3 g/kg body weight/day) and 28.1 g/person/day (0.5 g/kg body weight/day), respectively, under the intended conditions of use.

The sweet lupin-derived ingredients are composed mainly of varying levels of protein and fiber, with lower levels of fat and carbohydrate, all of which have a long history of consumption as part of a normal diet (Harwood, 1991; IOM, 2002a,b; USDA, 2005a,b,c), and hence, are expected to undergo normal metabolism. Consumption of wheat flour in the United States was reported to be 165 g/person/day (Wheat Foods Council, 2005), and the total population U.S. mean and 90th percentile intakes of protein and fiber were reported to be 75.2 and 114.0 g protein, respectively, and 15.1 and 24.7 g fiber, respectively (IOM, 2002a,b). Background consumption of the major macronutrients of the sweet lupin ingredients from various dietary sources are within range of those estimated from the intended conditions of use of each of Sweet Lupin Flour, Sweet Lupin Protein Fractions 1 and 2, Sweet Lupin Kernel Fibers 1 and 2, and Sweet Lupin Hull Fiber.

Some of the ingredients may be utilized in the same food categories; therefore, for completeness of the data, an all-user intake of sweet lupin based on all intended food-uses of all of GWF’s sweet lupin-derived ingredients also was estimated, providing total population all-user intake mean and 90th percentile level estimates of 92.6 and 158.5 g/day, respectively. The method used to calculate the daily dietary intakes under the intended conditions of use is considered to be ‘worst case’, as it incorporates several conservative assumptions, such as the assumption that all of the ingredients will be used in all of the food use categories at the highest level of use at the same time, which is highly unlikely, and therefore, the total population all-user estimated daily intakes are considered to be gross-overestimates and it is expected that the actual exposure to lupin from all of the sweet lupin-derived ingredients will be much less.

The safety assessment of lupin and sweet lupin-derived ingredients is based on the known metabolism of the macro-components of lupin, several short- and long-term preclinical toxicity studies and nutritional studies supporting the tolerability of these ingredients, as well as several human studies investigating the effect of sweet lupin-derived ingredients on parameters such as safety, glycemic and insulinemic response, bowel function, and palatability, which demonstrated that lupin was well tolerated.

Subchronic toxicity studies in rats using lupin seed and seed fractions ranged from 10 days to 13 weeks in duration and dietary administration of lupin-derived ingredients at doses of 9.7 to 57 g/kg body weight/day resulted in few significant effects on physical, biochemical, and hematological parameters (Fudiyansyah et al., 1995; Butler et al., 1996; Rahman et al., 1996a,b, 1997a,b; Rahman, 2000; Caligari et al., 2006; Pilvi et al., 2006). Observed effects,
such as decreased body weight gains, increased urea, and decreased albumin, were suggested by study authors to be due to a disturbance of normal protein utilization, which could have resulted from the amino acid deficiency of diets containing lupin seed or lupin fractions, as lupin is known to contain low levels of the essential amino acids, lysine and methionine. The reported increases in plasma urea remained within reported historical control values for rats (Sharp and LaRegina, 1998) and similar effects on body weight gains were not observed in studies in which rats were provided adequate amino acid-supplemented diets. No significant differences in biochemical or hematological parameters, organ weights, or histopathology were reported in rats provided up to 33 g lupin flour/kg body weight/day for a period of 90 days (Butler et al., 1996). Reported increases in serum alkaline phosphatase values in lupin-treated rats were suggested by the study authors to be a result of liver necrosis; however, there were no significant differences in the alanine aminotransferase or aspartate aminotransferase levels in the lupin-treated animals compared to the control group (Rahman et al., 1996a), and liver lesions were not reported in any of the other reviewed dietary studies. A toxicity study in which lupin protein extracted from the seeds of *L. albus* was administered by gavage to male Sprague-Dawley rats for a period of 2 weeks at a dose of 250 mg/kg body weight/day did not result in any adverse effects (Sirtori et al., 2004).

Chronic studies in rats using *L. angustifolia* seed and seed fractions ranging from 700 to 800 days in duration were identified in the available literature. Consumption of up to 13.6 g whole *L. angustifolia* seed/kg body weight/day was reported not to cause any adverse effects in rats when administered in the diet for up to 800 days (Grant et al., 1993), although levels of whole lupin seed in the diet were reported to decrease body weight in rats dosed for up to 700 days (Grant et al., 1995). Additionally, cecum and colon weights were significantly increased in rats after 700 days of feeding; however, the authors attributed this effect to volatile fatty acid production due to dietary fiber digestion in these organs (Grant et al., 1995). Dietary administration of lupin protein from *L. albus* and *L. luteus* to rats at levels of 6.3 and 6.88 g/kg body weight (20% of the diet), respectively, for a period of 112 days did not result in any changes in body weights, organ weights, or gross pathology (Ballester et al., 1980), and 20% lupin protein (isolated from *L. albus*) administered in the diet to 3 generations of rats for 270 days each did not result in adverse effects on either fertility or reproductive parameters in any of the three generations (Ballester et al., 1982, 1984).

Sweet lupin seeds (*L. angustifolia* and *L. albus*) are widely used in Australia as a source of protein and energy in livestock feeds, as they are cost-competitive with a number of other protein sources (Edwards and van Barneveld, 1998). These 2 species may be used with equal success in all livestock with the exception of pigs, where *L. albus* is not recommended for use in feed due to reduced feed intake and depressed growth rates (Edwards and van Barneveld, 1998). Pigs and poultry (monogastrics) require specific levels of individual amino acids in their diets (Edwards and van Barneveld, 1998), and due to low levels of both methionine and lysine in sweet lupins, feeds for pigs and poultry are more beneficial when they include multiple sources of protein or supplemental amino acids (Edwards and van Barneveld, 1998). Additionally, non-starch polysaccharides (NSP) have been suggested to
interfere with digestive enzymes in these monogastric species, causing a large difference between net energy content and digestible energy content (Edwards and van Barneveld, 1998). Equivocal results on feed consumption and body weights have been reported in nutritional studies ranging from 5 to 90 days in pigs and chickens consuming lupin seed or kernel, or lupin protein fractions at dietary levels providing doses of 2.3 to 1,540 g/kg body weight/day (King, 1981; Godfrey et al., 1985; Dunshea et al., 2001; Olkowski et al., 2001, 2005; Brenes et al., 2002; Rubio et al., 2003; Steenfeldt et al., 2003; Hammershoj and Steenfeldt, 2005; Martins et al., 2005; Mieczkowska et al., 2005; Prandini et al., 2005; Bielecka et al., 2006; Orda et al., 2006; Suchy et al., 2006; Zraly et al., 2006, 2007), and up to 210 days in cows (dose not provided) (Krapivina and Vashchekin, 2006); however, overall, lupin-containing feeds were generally well-tolerated and without adverse effects. Dehulled lupin seeds are now used in pig feeds because dehulling is reported to greatly improve gross energy digestibility (Wigan et al., 1994), and dehulling also was reported to be beneficial for poultry (Brenes et al., 1993). Studies evaluating the potential effect of 9.4 to 10.25 g lupin/kg body weight/day on the reproductive performance of cows (Axelsen, 1980) indicated no adverse effects, and a lack of effect on ovulation in ewes was reported by Pearse et al. (1991).

Conflicting results were reported upon investigation of the potential effect of lupin seed and seed fractions on mineral absorption (Rubio et al., 1994; Rahman et al., 1997b; Olkowski et al., 2005; Zraly et al., 2006, 2007). Rubio et al. (1994) reported that dialyzed soluble and insoluble aqueous lupin protein fractions had no significant effect on absorption of calcium, phosphorus, or zinc in rats, while whole lupin seed and the dialyzed residue fraction (i.e., the fibrous material that is insoluble in both water and buffer) significantly reduced phosphorus and zinc absorption due to the presence of phytate and insoluble NSP fractions which typically are not present in protein fractions. Conversely, Rahman et al. (1997b) reported that slightly higher doses of various aqueous lupin protein fractions, including dialyzed soluble and insoluble fractions, significantly reduced absorption of phosphorus, zinc, magnesium, and sodium in rats (with no effects on calcium or potassium); however, these effects were not discussed by the authors. Olkowski et al. (2005) reported that diets containing raw or dehulled lupin (L. albus, L. luteus, or L. angustifolius) seed meal did not affect plasma zinc levels in broiler chicks but significantly reduced plasma calcium levels, and Zraly et al. (2006) reported that dehulled lupin (L. angustifolius) seed meal in the diets of pigs resulted in significant decreases in plasma calcium but no changes in plasma phosphorus, whereas L. albus seed in the diet of pigs did not have any significant effect on plasma calcium or phosphorus levels (Zraly et al., 2007).

The results of nutritional studies in humans using ingredients derived from L. angustifolia indicated that the lupin ingredients were well tolerated in healthy volunteers (Petterson et al., 1994; Johnson et al., 2003; Archer et al., 2004; Hall and Johnson, 2004; Hall et al., 2005a,b; Johnson et al., 2006; Lee et al., 2006; Naruszewicz et al., 2006; Smith et al., 2006; Joray et al., 2007). The identified studies ranged in length from a single dose to 28 days and included daily doses of 9 to 37.4 g kernel fiber and 7.7 to 264 g lupin flour derived from L. angustifolia. A dose of 35 g lupin flour from L. albus provided to young adults for a period

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of 60 days was well tolerated and was without compound-related changes in biochemical or hematological parameters (Gattas Zaror et al., 1990). A daily intake of 28 to 56 g of lupin protein from lupin flour also was reported to be well tolerated in 9 healthy men for a period of 10 days, with the only significant change being an increase in urea nitrogen in the high-dose group compared to the low-dose group (Egañ a et al., 1992). Consumption of 16.75 g of L. albus-derived lupin protein/day by otherwise healthy, chronically-smoking, volunteers with moderate hypercholesterolemia for a period of 90 days was well tolerated and resulted in significant reductions in total cholesterol, low-density lipoprotein cholesterol, blood glucose, homocysteine, high sensitivity C-reactive protein, urinary F2-isoprostane/creatinine excretion, systolic blood pressure, and diastolic blood pressure (Naruszewicz et al., 2006). Some of the doses used in these studies exceed the estimated intakes for Lupin Flour, Lupin Protein Fractions 1 and 2, Lupin Kernel Fibers 1 and 2, and Lupin Hull Fiber. There are no indications from the published literature that the intended uses of sweet lupin-derived ingredients would result in any adverse health effects in humans.

High levels of dietary alkaloids, natural toxicants in plants, may be toxic (Bradbury et al., 2004). Although some species of lupin can contain 'high' levels (2 to 3%) of alkaloids (bitter lupins) (Reinhard et al., 2006), the sweet varieties have low alkaloid levels (typically <0.02%), making them suitable for consumption by humans and livestock (Petterson, 1998). The Advisory Committee on Novel Foods and Processes (ACNFP) of the United Kingdom (ACNFP, 1996) established a limit for alkaloid levels in lupin seed fit for human consumption of 200 mg alkaloids/kg seed. Additionally, a tolerable level of exposure to lupin alkaloids for humans of 35 μg/kg body weight/day was tentatively established by Australia New Zealand Food Authority (ANZFA) (2001a). The lupin ingredients produced by GWF comply with the maximal alkaloid level set forth by ACNFP, as evidenced by the product specifications and batch data analyses, and hence the levels of alkaloids present in the sweet lupin-derived ingredients are expected not to produce any adverse effects on human health. The presence of other phyto-components in the ingredients of GWF, specifically oligosaccharides, phenols and condensed tannins, trypsin inhibitors, saponins, phytic acid, and lectins, are negligible and are not anticipated to impact the safety of the sweet lupin-derived ingredients. Lupinosis, which can occur as a result of the ingestion of lupin plants contaminated with phomopsins, which are toxins produced by fungi, has been reported in livestock that grazed on infected plants (Allen, 1986; Morcombe et al., 1992; ANZFA, 2001b), although cases of lupinosis in humans have not been identified (Lowen et al., 1995). Analysis of GWF’s sweet lupin-derived ingredients indicated compliance with maximum tolerable levels of 5 μg/kg final product, as set forth by Food Standards Australia New Zealand (FSANZ) (ANZFA, 2001b), and therefore, there is no concern for the potential occurrence of lupinosis in consumers of products containing these ingredients.

Lupin allergy has recently been recognized with documented cases of anaphylaxis and other allergic reactions following lupin consumption (Hefle et al., 1994; Matheu et al., 1999; Novembre et al., 1999; Moneret-Vautrin et al., 2004; Smith et al., 2004; Radcliffe et al., 2005; Rotiroti et al., 2007; Wassenberg and Hofer, 2007). Lupin has recently been added to the list of commonly allergenic foods in the European Union but does not have such status in
other parts of the world. Additionally, cross-reactivity between lupin protein allergens and allergens in peanuts has been reported (Hefle et al., 1994; Moneret-Vautrin et al., 1999; Kanny et al., 2000; Fæste et al., 2004; Wüthrich et al., 2004; Costa et al., 2005; Peeters et al., 2007), although such cross-reactions only occur in a fraction (perhaps as high as 20%) of peanut-allergic individuals. Furthermore, lupin allergy can occur independent of peanut allergy (Peeters et al., 2007). As expected, the results of in vitro studies using GWF's ingredients indicate that not all peanut-allergic individuals would be sensitive to these lupin-based ingredients (Nordlee, unpublished, 2004). While the levels of protein in these different lupin-based ingredients is variable, each of the ingredients is potentially allergenic, and therefore, GWF will take steps to ensure that the presence of lupin is identified on all products that contain their sweet lupin-derived ingredients in order to notify consumers and to attempt to prevent exposure in sensitive populations. As there are currently no requirements for the labeling of ingredients that may be cross-reactive, GWF intends to ensure that lupin is included on the end product label for all products containing the sweet lupin-derived ingredients either as direct ingredients or as incidental additives rather than include a statement regarding cross-reactivity. It should therefore be expected that the labeling of foods to which the sweet lupin-derived ingredients are added should alert the lupin-allergic consumer to the presence of lupin.

The results of pre-clinical and human studies of whole sweet lupin seed and lupin fractions do not indicate any potential for adverse effects in humans consuming these ingredients under the intended conditions of use. A critical evaluation of the data and information summarized in this report supports the safety and suitability and the GRAS status based on scientific procedures of the intended uses of Sweet Lupin Flour, Sweet Lupin Protein Fractions 1 and 2, Sweet Lupin Kernel Fibers 1 and 2, and Sweet Lupin Hull Fiber meeting appropriate food-grade specifications and manufactured consistent with cGMP.
CONCLUSION

We, the Expert Panel, have, independently and collectively, critically evaluated the data and information summarized above and conclude that the intended uses in traditional foods of ingredients derived from sweet lupin, meeting appropriate food-grade specifications presented herein and produced consistent with current Good Manufacturing Practices (GMP), are safe and suitable.

We further conclude that the intended uses in traditional foods of ingredients derived from sweet lupin, meeting appropriate food-grade specifications presented herein and produced consistent with current GMP, are Generally Recognized as Safe (GRAS) based on scientific procedures.

It is our opinion that other qualified experts would concur with these conclusions.

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ATTACHMENT 2

INTENDED FOOD USES AND USE LEVELS OF SWEET LUPIN-DERIVED INGREDIENTS
<table>
<thead>
<tr>
<th>Food Category</th>
<th>Proposed Food-Use</th>
<th>Sweet Lupin Ingredient</th>
<th>Maximum Use Level</th>
</tr>
</thead>
<tbody>
<tr>
<td>Baked Goods and Baking Mixes</td>
<td>Bagels; Biscuits; Cakes; Cookies; Cornbread, Corn Muffins, and Tortillas; Crackers; Croissants; English Muffins; French Toast, Pancakes, Waffles, and Crepes; Muffins and Popovers; Pastries; Pie; Quick Breads and Sweet Rolls; Soft Bread Sticks; Soft Pretzels; and Yeast Breads and Rolls</td>
<td>Flour</td>
<td>25%</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Protein Fraction 1 &amp; 2, Kernel Fibers 1 &amp; 2, Hull Fiber</td>
<td>10%</td>
</tr>
<tr>
<td>Grains Products and Pastas</td>
<td>Macaroni and Noodle Products</td>
<td>Flour</td>
<td>25%</td>
</tr>
<tr>
<td>Dairy Product Analogs</td>
<td>Condensed Milk Analogs²; Milk Powder Analogs²</td>
<td>Protein Fraction 1</td>
<td>15%</td>
</tr>
<tr>
<td></td>
<td>High Fat Powder; Imitation Milk; Soy Milk Alternatives</td>
<td>Protein Fraction 1</td>
<td>4%</td>
</tr>
<tr>
<td></td>
<td>Imitation Cheese</td>
<td>Protein Fraction 1</td>
<td>20%</td>
</tr>
<tr>
<td></td>
<td>Non-dairy Cream Substitutes and Coffee Whiteners</td>
<td>Protein Fraction 2</td>
<td>3%</td>
</tr>
<tr>
<td>Frozen Dairy Desserts and Mixes</td>
<td>Ice Cream</td>
<td>Protein Fraction 1</td>
<td>20%</td>
</tr>
<tr>
<td>Gelatins, Puddings, and Fillings</td>
<td>Mousses and Meringues</td>
<td>Protein Fraction 2</td>
<td>3%</td>
</tr>
<tr>
<td>Jams and Jellies</td>
<td>Spreadable Jelly</td>
<td>Protein Fraction 1 &amp; 2</td>
<td>3%</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Kernel Fibers 1 &amp; 2</td>
<td>2%</td>
</tr>
<tr>
<td>Meat Products</td>
<td>Commercially Processed Meats and Sandwich Ingredients</td>
<td>Protein Fraction 1</td>
<td>3%</td>
</tr>
<tr>
<td>Milk Products</td>
<td>Fermented Milk Beverages; Flavored Milk and Milk Drinks; Milk-Based Meal Replacements</td>
<td>Protein Fraction 1</td>
<td>5%</td>
</tr>
<tr>
<td>Soft Candy</td>
<td>Boiled Sweets</td>
<td>Protein Fraction 1</td>
<td>2%</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Protein Fraction 2, Kernel Fibers 1 &amp; 2</td>
<td>1%</td>
</tr>
<tr>
<td></td>
<td>Chocolate, Compound Chocolate</td>
<td>Protein Fraction 1 &amp; 2, Kernel Fibers 1 &amp; 2</td>
<td>3%</td>
</tr>
<tr>
<td></td>
<td>Soft and Firm Jellies</td>
<td>Protein Fraction 1 &amp; 2, Kernel Fibers 1 &amp; 2</td>
<td>3%</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Kernel Fibers 1 &amp; 2</td>
<td>2%</td>
</tr>
<tr>
<td>Beverages and Beverage Bases</td>
<td>Carbonated Beverages</td>
<td>Kernel Fibers 1 &amp; 2</td>
<td>3%</td>
</tr>
<tr>
<td></td>
<td>Energy, Sports and Isotonic Drinks</td>
<td>Kernel Fibers 1 &amp; 2</td>
<td>10%</td>
</tr>
<tr>
<td>Breakfast Cereals</td>
<td>Instant and Regular Hot Cereals</td>
<td>Kernel Fibers 1 &amp; 2</td>
<td>20%</td>
</tr>
<tr>
<td></td>
<td>Ready-to-Eat Breakfast Cereals</td>
<td>Hull Fiber</td>
<td>10%</td>
</tr>
<tr>
<td>Processed Fruits and Fruit Juices</td>
<td>Fruit-Flavored Drinks</td>
<td>Kernel Fibers 1 &amp; 2</td>
<td>3%</td>
</tr>
</tbody>
</table>

² No food codes were found for these categories, thus surrogate codes were chosen to represent the category. Analogues of these types were not found thus milk codes were used.
APPENDIX B

SUMMARY OF ANALYTICAL DATA, CERTIFICATES OF ANALYSIS, AND OTHER COMPOSITIONAL INFORMATION
APPENDIX B-1

SUMMARY OF THE ALKALOIDS IDENTIFIED IN *LUPINUS* SPP.
SUMMARY OF THE ALKALOIDS IDENTIFIED IN *LUPINUS* SPP.

High levels of dietary alkaloids, natural constituents in plants, may be toxic. Although some species of lupin can contain 'high' levels (2 to 3%) of alkaloids (bitter lupins), the sweet varieties have low alkaloid levels (~0.001 to 0.002%), making them suitable for consumption by humans and livestock. Alkaloids present in *L. angustifolia* comprise mainly lupanine (42 to 59%), 13-hydroxylupanine (24 to 45%), and angustifoline (7 to 15%) (Wink et al., 1995), which are derivatives of quinolizidine (Petterson, 1998). *L. albus* contains <100 mg alkaloids/kg whole seed, the majority of which is lupanine (~70%), followed by albine (~15%) and lesser amounts of 13-hydroxylupanine, sparteine, and multiflorine (Petterson and Mackintosh, 1994; Zdunczyk et al., 1994; Wink et al., 1995). The alkaloid profile of *L. luteus* is composed almost entirely of lupanine (60%) and sparteine (~30%) (Wink et al., 1995), while *L. mutabilis* contains several different alkaloids, the main ones being lupanine, sparteine, and 13-hydroxylupanine (Petterson, 1998).

Following a review of the literature, a limit for alkaloid levels in lupin seed fit for human consumption of 200 ppm was set by the Advisory Committee on Novel Foods and Processes (ACNFP) of the United Kingdom (ACNFP, 1996). Additionally, a tolerable level of exposure to lupin alkaloids for humans of 35 μg/kg body weight/day has tentatively been established by the Australia/New Zealand Food Authority (ANZFA) (ANZFA, 2001). The sweet lupin flour ingredient produced by George Weston Foods Limited complies with the maximal alkaloid level of <200 ppm set forth by the ACNFP, as evidenced by the product specifications and batch data analyses, and hence the levels of alkaloids present in the sweet lupin flour ingredient are expected not to produce any adverse effects on human health.

REFERENCES


APPENDIX B-2

BATCH ANALYSES OF SWEET LUPIN FLOUR

000112
# Table B-2-1

Summary of the Physical, Chemical and Microbiological Analyses of Batches of Sweet Lupin Flour

<table>
<thead>
<tr>
<th>Specification Parameter</th>
<th>Specification</th>
<th>Laboratory Reference Number and/or Date Received</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>E2003/014985 (Fine)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>E2003/025993 (Fine)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>25MAR04/37721 (Fine)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>E2003/014986 (Coarse)</td>
</tr>
<tr>
<td><strong>Physical and Chemical</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Protein (% DSB)</td>
<td>Minimum 35</td>
<td>42.1</td>
</tr>
<tr>
<td>Fat (% DSB)</td>
<td>7 to 10</td>
<td>8.5</td>
</tr>
<tr>
<td>Total Carbohydrate (%)</td>
<td>7 to 9</td>
<td>N/A</td>
</tr>
<tr>
<td>Insoluble Dietary Fiber (% DSB)</td>
<td>32 to 36</td>
<td>34.5&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>Soluble Dietary Fiber (% DSB)</td>
<td>3 to 6</td>
<td>32.4&lt;sup&gt;2&lt;/sup&gt;</td>
</tr>
<tr>
<td>Moisture (%)</td>
<td>7 to 11</td>
<td>7.6</td>
</tr>
<tr>
<td>Ash (% DSB)</td>
<td>2 to 4</td>
<td>N/A</td>
</tr>
<tr>
<td>Alkaloids (ppm)&lt;sup&gt;d&lt;/sup&gt;</td>
<td>&lt;200</td>
<td>120&lt;sup&gt;*&lt;/sup&gt;</td>
</tr>
<tr>
<td>Cadmium (ppm)&lt;sup&gt;e&lt;/sup&gt;</td>
<td>&lt;0.1</td>
<td>0.04&lt;sup&gt;*&lt;/sup&gt;</td>
</tr>
<tr>
<td>Lead (ppm)&lt;sup&gt;f&lt;/sup&gt;</td>
<td>&lt;0.2</td>
<td>0.13&lt;sup&gt;*&lt;/sup&gt;</td>
</tr>
<tr>
<td>Phomopsins (ppb)</td>
<td>&lt;5</td>
<td>&lt;5&lt;sup&gt;*&lt;/sup&gt;</td>
</tr>
<tr>
<td><strong>Microbiological</strong></td>
<td></td>
<td>Lab No. L65995 (31/10/03)</td>
</tr>
<tr>
<td>Total plate count (CFU/g)</td>
<td>&lt;40,000</td>
<td>250</td>
</tr>
<tr>
<td>Coliforms (CFU/g)</td>
<td>&lt;100</td>
<td>23</td>
</tr>
<tr>
<td><em>Escherichia coli</em> (CFU/g)</td>
<td>&lt;10</td>
<td>&lt;3</td>
</tr>
<tr>
<td><em>Salmonella</em> spp. (per 25 g)</td>
<td>Absent</td>
<td>ND</td>
</tr>
<tr>
<td>Yeasts and moulds (CFU/g)</td>
<td>&lt;1,000</td>
<td>300 (moulds), 3,600 (yeasts)</td>
</tr>
<tr>
<td><em>Staphylococcus</em> spp. (CFU/g)</td>
<td>&lt;100</td>
<td>NT</td>
</tr>
<tr>
<td><em>Bacillus cereus</em> (CFU/g)</td>
<td>&lt;100</td>
<td>&lt;100</td>
</tr>
<tr>
<td>Listeria (per 25 g)</td>
<td>Absent</td>
<td>ND</td>
</tr>
</tbody>
</table>

<sup>a</sup> Measured as received; <sup>b</sup> Measured on a dry solid basis
<sup>c</sup> DSB = Dry solid basis; N/A = Not available; ND = not detected
<sup>d</sup> Batch values (other than moisture) were calculated on a dry basis for comparison to product specifications
<sup>e</sup> Fiber and carbohydrate by difference
<sup>f</sup> Represents total dietary fiber
<sup>g</sup> Total alkaloids were measured by the Department of Industry and Resources Chemistry Centre [Western Australia (WA)] using Gas Chromatography/Mass Spectrometry
<sup>h</sup> Measured by the Department of Industry and Resources Chemistry Centre (WA) using inductively coupled plasma mass spectrometry (ICP-MS) Method IEL1STIM

See Attachment B-2-1 for certificates of analysis.
# REPORT OF ANALYSIS

**CLIENT (D183)**: CI 70  
**Attn**: Sarab  
**DATE RECEIVED**: 22.07.2003  
**ORDER NUMBER**: 

<table>
<thead>
<tr>
<th>LAB REPORT #</th>
<th>CLIENT REF #</th>
<th>DESCRIPTION</th>
</tr>
</thead>
<tbody>
<tr>
<td>E2003/014985</td>
<td>AR 471</td>
<td>LUPIN FLOUR COARSE</td>
</tr>
<tr>
<td>E2003/014986</td>
<td>AR 471</td>
<td>LUPIN FLOUR FINE</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Protein (Nx6.25) (0100)</th>
<th>38.8%</th>
<th>38.9%</th>
</tr>
</thead>
<tbody>
<tr>
<td>Moisture (0103)</td>
<td>9.0%</td>
<td>7.6%</td>
</tr>
<tr>
<td>Fat (Acid Hyd.) (0106)</td>
<td>7.3%</td>
<td>7.9%</td>
</tr>
<tr>
<td>Tot. Diet. Fibre (0109)</td>
<td>29.3%</td>
<td>32.2%</td>
</tr>
</tbody>
</table>

**Note**: Results on samples as received

**Signatory**  
(for Laboratory Manager)  
**Date**: 30/7/03

---

**INTERNATIONAL**  
**TELEPHONE**: +61 2 9764 8222  
**FASCIMILE**: +61 2 9742 6351

---

**.getClient**

---

**Report of Analysis**

---

**Lab Report No**  
E2003/014985  
E2003/014986

---

000115

---

**A Unit of George Weston Foods Limited**  
A.C.N. 008 420 852
REPORT OF ANALYSIS

CLIENT (D290) : FIG0048
Attn : Sarab

DATE RECEIVED : 17.12.2003
ORDER NUMBER :

LAB REPORT #  CLIENT REF #  DESCRIPTION
E2003/025993  LF1612  LUPIN FLOUR

Lab Report No  E2003/025993

Moisture (0103)  7.0%
Fat (Acid Hyd.) (0106)  8.6%
Tot.Diet.Fibre (0109)  30.1%
Protein (Dumas) (0284) (Nx6.25)  40.1%

Signatory : (for Laboratory Manager)
Date : 14/1/04

Note: Results on samples as received
LABORATORY REPORT
ON
MISCELLANEOUS POWDER

FOR: MAX SCOTT CONSULTING P/L

56A RAVENSWAY ROAD
BERWICK 3805

MAX SCOTT

ACCOUNT: M50

Date: 05/05/04

Our ref: 25MAR04/37721/MISP

SAMPLE: Data Received - 25/03/04
DETAILS: Origin -
Code/Ref. - 27 FLOUR
Order number -
No. of samples - 1
Package Type -

<table>
<thead>
<tr>
<th>TEST</th>
<th>RESULTS</th>
<th>DTS METHOD</th>
</tr>
</thead>
<tbody>
<tr>
<td>MOISTURE X%w</td>
<td>8.0</td>
<td>MOIS 16 11.96</td>
</tr>
<tr>
<td>FAT X%w</td>
<td>7.9</td>
<td>FATS 07 12.29</td>
</tr>
<tr>
<td>PROTEIN (TN x 6.25) X%i</td>
<td>39.1</td>
<td>PROT 01 02.81</td>
</tr>
<tr>
<td>ASH X%w @ 550°C</td>
<td>2.6</td>
<td>ASHS 04 04.33</td>
</tr>
<tr>
<td>DIETARY FIBRE X (INSOLUBLE)</td>
<td>32.6</td>
<td>DIET 02 03.93</td>
</tr>
<tr>
<td>DIETARY FIBRE X (SOLUBLE)</td>
<td>3.6</td>
<td></td>
</tr>
</tbody>
</table>

S. Nolan
Technical Manager
REPORT ON THE ANALYSIS OF SIX LUPIN PRODUCT SAMPLES

Sample History

Six lupin product samples were received for analysis of total alkaloids, total phenolic, oligosaccharides, trypsin inhibitor, lead, cadmium, phytic acid, lectin and total saponins.

<table>
<thead>
<tr>
<th>Lab Code</th>
<th>Sample No</th>
</tr>
</thead>
<tbody>
<tr>
<td>04D505_001</td>
<td>103</td>
</tr>
<tr>
<td>04D505_002</td>
<td>1017</td>
</tr>
<tr>
<td>04D505_003</td>
<td>1037</td>
</tr>
<tr>
<td>04D505_004</td>
<td>1046</td>
</tr>
<tr>
<td>04D505_005</td>
<td>1071</td>
</tr>
<tr>
<td>04D505_006</td>
<td>1090</td>
</tr>
</tbody>
</table>

Test Methods

Total alkaloids by in-house GC-MS method.
Oligosaccharides by method SP7.
Total phenolics by folin-ciocalton.
Trypsin inhibitor by method SP 9.
Cd (ICP-MS) = Cadmium, Cd by ICP-MS method ILE1STIM.
Pb (ICP-MS) = Lead, Pb by ICP-MS method ILE1STTIM.
Phytic acid by in house GC-MS method
Lectin: Haemagglutinin activity by in house method
Saponins by in house GC-MS method

mg/kg as = milligrams per kilogram as received.
%ar = per cent as received.
Results

<table>
<thead>
<tr>
<th>Lab No</th>
<th>Total</th>
<th>Oligosac</th>
<th>Total</th>
<th>Trypsin</th>
<th>Cd</th>
</tr>
</thead>
<tbody>
<tr>
<td>94D</td>
<td>alcohols</td>
<td>%ar</td>
<td>%ar</td>
<td>Phenolics</td>
<td>inhibitor</td>
</tr>
<tr>
<td>---------</td>
<td>---------</td>
<td>------</td>
<td>------</td>
<td>----------</td>
<td>-----------</td>
</tr>
<tr>
<td>505-001</td>
<td>0.012</td>
<td>5.6</td>
<td>0.324</td>
<td>1.45</td>
<td>0.04</td>
</tr>
<tr>
<td>505-002</td>
<td>0.008</td>
<td>1.5</td>
<td>0.168</td>
<td>2.48</td>
<td>0.03</td>
</tr>
<tr>
<td>505-003</td>
<td>0.002</td>
<td>0.4</td>
<td>0.125</td>
<td>0.099</td>
<td>0.06</td>
</tr>
<tr>
<td>505-004</td>
<td>0.003</td>
<td>0.4</td>
<td>0.307</td>
<td>0.94</td>
<td>0.06</td>
</tr>
<tr>
<td>505-005</td>
<td>0.00005</td>
<td>0.2</td>
<td>0.066</td>
<td>0.17</td>
<td>0.03</td>
</tr>
<tr>
<td>505-006</td>
<td>0.001</td>
<td>0.6</td>
<td>0.126</td>
<td>0.15</td>
<td>0.02</td>
</tr>
</tbody>
</table>

Lectin figures presented indicate the highest dilution that agglutination is still observed. A 0 result indicates no haemagglutinin activity.

Commercial standards of lupin saponins are not commercially available. The alternative standard purchased from a private company is not sufficiently pure for definitive analysis, affecting total saponin accuracy. Definitive analysis may be achieved with the provision of additional time and the isolation expense that is required to prepare specific and pure standards.

If you have any inquiries regarding these results, please contact Shao Fang Wang. These results apply specifically to the sample as received.

NEIL ROTHNIE
CHIEF
FOOD & BIOLOGICAL CHEMISTRY
LABORATORY

S.F. WANG
Chemist and Research Officer
FOOD & BIOLOGICAL CHEMISTRY
LABORATORY

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ANALYSIS REPORT

Weston Food Laboratories
PO Box 1
ENFIELD NSW 2136

ATTENTION
VIA FAX
JOB NUMBER
Sherry Duckworth
02 9742 6351
J0412-0638

DATE RECEIVED 30/12/04
OUR SAMPLE NUMBER S2004-20559
REFERENCE NUMBER Legume Flour
SAMPLE TYPE Legume Flour
Phomopsin (TP/043) <5
Phomopsin(ppb)

Note All samples are analysed on an as received basis.
This report is not to be reproduced except in full.
TP refers to the technical procedure used to conduct the analysis.

Final Report
Report Number: 5434

Lakshmi Iyer
Technical Manager
17-Jan-2005
Attention: Sherry Duckworth

Weston Food Laboratories
1 Braidwood Street, Enfield N.S.W. 2136, Phone (02) 9784 8222 Facsimile (02) 9758 8300
NATA Accredited Laboratory
A unit of George Weston Foods Limited, A.C.A. 009 490 932

Report No: 85.6.8

Sample: Lupin Protein Samples
Sender: Weston Technologies - R&D
Address: 1 Braidwood Street, Enfield NSW 2136
Received Date: 1.6.05

Condition of samples on arrival: Satisfactory

Results reported apply to sample(s) listed below. Laboratory is not responsible for sampling.

<table>
<thead>
<tr>
<th>Lab No.</th>
<th>Description</th>
<th>Plate Count</th>
<th>YEAST &amp; MOLD</th>
<th>REEDEMICROBIA</th>
<th>THERMOPHILIC COCCIDIA</th>
<th>THERMOPHILIC SPIROCHETES</th>
<th>SPOROPHILIC SPIROCHETES</th>
<th>COLIFORMS</th>
<th>E. coli</th>
<th>SALMONELLA</th>
<th>LISTERIA spp.</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>N 3317</td>
<td>924 Discharge</td>
<td>Est. 105g</td>
<td>&lt; 100g</td>
<td>Est. 30g</td>
<td>Est. 50g</td>
<td>Est. 20g</td>
<td>8 500g</td>
<td>4g</td>
<td>&lt; 3g</td>
<td>ND/ND</td>
<td>ND/ND</td>
<td>Approved</td>
</tr>
<tr>
<td>N 3318</td>
<td>1047 Coarse Flour</td>
<td>250g</td>
<td>300G, 3 600G</td>
<td>&lt; 100g</td>
<td>Est. 40g</td>
<td>Est. 40g</td>
<td>Est. 500g</td>
<td>30g</td>
<td>&lt; 3g</td>
<td>ND/ND</td>
<td>ND/ND</td>
<td></td>
</tr>
<tr>
<td>N 3320</td>
<td>1070 Soluble Fibre</td>
<td>3 000g</td>
<td>&lt; 100g</td>
<td>Est. 60g</td>
<td>Est. 10g</td>
<td>Est. 2 300g</td>
<td>&lt; 3g</td>
<td>ND/ND</td>
<td>ND/ND</td>
<td>ND/ND</td>
<td>ND/ND</td>
<td></td>
</tr>
</tbody>
</table>

Form No. WP:12027/Issue 4/7/20 May 2008

Approved by:

[Signature]
Attention: Sherry Duckworth

Weston Food Laboratories
1 Braidwood Street, Enfield N.S.W. 2136, Phone (02) 9764 8222 Facsimile (02) 9758 6300
A unit of George Weston Foods Limited, A.C.N. 006 429 632

Report No.: 03.10.334

Sample: Lupin Flour
Sender: Weston Technologies- R&D
Address: 1 Braidwood Street Enfield NSW 2136
Receive Date: 31.10.03
Condition of samples on arrival: Satisfactory

Results reported apply to sample(s) listed below. Laboratory is not responsible for sampling:

<table>
<thead>
<tr>
<th>Lab No.</th>
<th>Description</th>
<th>Plate Count</th>
<th>Yeast &amp; Mould</th>
<th>Staph. (coag.)</th>
<th>Bac. (census)</th>
<th>Thermophils</th>
<th>Meso.</th>
<th>Coliforms</th>
<th>K. oral</th>
<th>Salmonella</th>
<th>Listeria sp.</th>
</tr>
</thead>
<tbody>
<tr>
<td>L 6599</td>
<td>Lupin Flour 31.10.03</td>
<td>4.500 / g</td>
<td>900M</td>
<td>&lt; 100 / g</td>
<td>&lt; 100 / g</td>
<td>Envl. 20 / g</td>
<td>400 / g</td>
<td>40 / g</td>
<td>&lt; 3 / g</td>
<td>ND / 25g</td>
<td>ND / 25g</td>
</tr>
<tr>
<td></td>
<td>For FSA Werribee</td>
<td>700Y / g</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Key: N= Not Tested, D= Detected at, ND= Not Detected, S= Small, M=Mixed, Y= Yeast, Sp= Spore. OD1= organism counts << less than << Greater than

Comments:

Note: 
- This is an interim report, final report to be issued. Interim Report faxed on:
- This is a complete report, final copies to be issued. Complete report faxed on:
- This is a complete report, no further copies of the report will be issued. Complete report emailed on: 15.11.03 DT
- This is a final report, no further copies of the report will be issued. Final Report issued on:

Report Checked By: DT
Report Written By: MP

Certificate of Microbiological Analyses
Analysed On: 06.11.03
Operator: MP
Copies To: C.Fryirs, S.Duckworth, S.Kaur, File

NATASA Accredited Laboratory, A.C.N. 006 429 632
APPENDIX B-3

SUMMARY OF THE RESULTS OF ANALYSIS OF PESTICIDE RESIDUES IN SWEET LUPIN FLOUR AND CORRESPONDING IDENTIFIED MAXIMUM RESIDUE LIMITS (MRL) FOR VARIOUS FOOD PRODUCTS
# Table B-3-1 Pesticide Residues Identified In Sweet Lupin Flour and Corresponding Identified Maximum Residue Limits (MRL) For Various Food Products

<table>
<thead>
<tr>
<th>Food Product</th>
<th>U.S. FDA MRL (ppm)*</th>
<th>FSANZ MRL (ppm)bc</th>
<th>Sweet Lupin Flour (ppm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cereal grain</td>
<td>Not available</td>
<td>0.1</td>
<td>Complies</td>
</tr>
<tr>
<td>Dichlorodiphenyldichloroethylene (DDT)</td>
<td>Not available</td>
<td>0.020</td>
<td>MRL below LOD</td>
</tr>
<tr>
<td>Chlordane (cis and trans)</td>
<td>Not available</td>
<td>0.020</td>
<td>MRL below LOD</td>
</tr>
<tr>
<td>Oxychlordane</td>
<td>Not available</td>
<td>0.020</td>
<td>MRL below LOD</td>
</tr>
<tr>
<td>Aldrin</td>
<td>Not available</td>
<td>0.020</td>
<td>MRL below LOD</td>
</tr>
<tr>
<td>Dieldrin</td>
<td>Not available</td>
<td>0.020</td>
<td>MRL below LOD</td>
</tr>
<tr>
<td>Endrin</td>
<td>Not available</td>
<td>No limit set</td>
<td>NA</td>
</tr>
<tr>
<td>Endosulfan</td>
<td>0.1</td>
<td>0.20</td>
<td>Complies</td>
</tr>
<tr>
<td>Lindane</td>
<td>Not available</td>
<td>0.50</td>
<td>Complies</td>
</tr>
<tr>
<td>Hexachlorocyclohexanes (HCH)</td>
<td>Not available</td>
<td>0.10</td>
<td>Complies</td>
</tr>
<tr>
<td>Heptachlor</td>
<td>Not available</td>
<td>0.020</td>
<td>MRL below LOD</td>
</tr>
<tr>
<td>Heptachlor epoxide</td>
<td>Not available</td>
<td>0.020</td>
<td>MRL below LOD</td>
</tr>
<tr>
<td>Hexachlorobenzene (HCB)</td>
<td>Not available</td>
<td>0.050</td>
<td>Complies</td>
</tr>
<tr>
<td>Dichlorvos</td>
<td>0.02</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cereal grain</td>
<td>0.5</td>
<td>5</td>
<td>Complies</td>
</tr>
<tr>
<td>Peanut</td>
<td>Not available</td>
<td>2</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat bran, unprocessed</td>
<td>Not available</td>
<td>10</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat germ</td>
<td>Not available</td>
<td>10</td>
<td>Complies</td>
</tr>
<tr>
<td>Chlorpyrifos Methyl</td>
<td>0.02</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cereal grains (barley, oat, wheat, not rice)</td>
<td>6</td>
<td>10</td>
<td>Complies</td>
</tr>
<tr>
<td>Lupin (dry)</td>
<td>NA</td>
<td>10</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat bran, unprocessed</td>
<td>Not available</td>
<td>20</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat germ</td>
<td>Not available</td>
<td>30</td>
<td>Complies</td>
</tr>
<tr>
<td>Fenitrothion</td>
<td>0.02</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cereal grains</td>
<td>30</td>
<td>10</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat bran, unprocessed</td>
<td>Not available</td>
<td>20</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat germ</td>
<td>Not available</td>
<td>20</td>
<td>Complies</td>
</tr>
<tr>
<td>Pirimiphos Methyl</td>
<td>0.02</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Barley</td>
<td>Not available</td>
<td>7</td>
<td>Complies</td>
</tr>
<tr>
<td>Bran of cereal grain, unprocessed</td>
<td>Not available</td>
<td>20</td>
<td>Complies</td>
</tr>
<tr>
<td>Maize</td>
<td>8</td>
<td>7</td>
<td>Complies</td>
</tr>
<tr>
<td>Oats</td>
<td>Not available</td>
<td>7</td>
<td>Complies</td>
</tr>
<tr>
<td>Peanut</td>
<td>Not available</td>
<td>5</td>
<td>Complies</td>
</tr>
<tr>
<td>Rye</td>
<td>Not available</td>
<td>10</td>
<td>Complies</td>
</tr>
<tr>
<td>Sorghum</td>
<td>8</td>
<td>10</td>
<td>Complies</td>
</tr>
<tr>
<td>Triticale</td>
<td>Not available</td>
<td>10</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat</td>
<td>Not available</td>
<td>10</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat germ</td>
<td>Not available</td>
<td>30</td>
<td>Complies</td>
</tr>
</tbody>
</table>
Table B-3-1  Pesticide Residues Identified In Sweet Lupin Flour and Corresponding Identified Maximum Residue Limits (MRL) For Various Food Products

<table>
<thead>
<tr>
<th>Food Product</th>
<th>U.S. FDA MRL (ppm)</th>
<th>FSANZ MRL (ppm)</th>
<th>Sweet Lupin Flour (ppm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Malathion</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cereal grains</td>
<td>8</td>
<td>8</td>
<td>Complies</td>
</tr>
<tr>
<td>Peanut</td>
<td>8</td>
<td>8</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat bran unprocessed</td>
<td>Not available</td>
<td>20</td>
<td>Complies</td>
</tr>
<tr>
<td><strong>Piperonyl Butoxide</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cereal Bran unprocessed</td>
<td>Not available</td>
<td>40</td>
<td>Complies</td>
</tr>
<tr>
<td>Wheat germ</td>
<td>Not available</td>
<td>50</td>
<td>Complies</td>
</tr>
<tr>
<td>Cereal grains</td>
<td>20&lt;sup&gt;a&lt;/sup&gt;</td>
<td>20</td>
<td>Complies</td>
</tr>
<tr>
<td>Oleic acid</td>
<td>Not available</td>
<td>8</td>
<td>Complies</td>
</tr>
</tbody>
</table>

FSANZ = Food Standards Australia New Zealand; LOD = Limit of detection; MS = Mass spectrometry; NA = not applicable; U.S. FDA = Food and Drug Administration;

<sup>a</sup> U.S. EPA, 2007
<sup>b</sup> Australia New Zealand Food Standards Code, Standard 1.4.2 (FSANZ, 2005)
<sup>c</sup> APVMA, 2005
<sup>d</sup> MRL established for post-harvest barley, buckwheat, rice, rye and wheat
<sup>e</sup> MRL established for post-harvest oat and sorghum
APPENDIX B-4

SUMMARY OF THE ANALYSES OF OLIGOSACCHARIDES, PHENOLICS AND CONDENSED TANNINS, TRYSPIN INHIBITORS, PHYTIC ACID, SAPONINS, AND LECTINS PRESENT IN ONE BATCH OF SWEET LUPIN FLOUR
### Table B-4-1  Summary of the Analyses of Phytonutrients Present in One Batch of Sweet Lupin Flour

<table>
<thead>
<tr>
<th>Parameter</th>
<th>%, As Received</th>
<th>%, Dry Solids Basis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oligosaccharides</td>
<td>5.6</td>
<td>6.2</td>
</tr>
<tr>
<td>Phenolics</td>
<td>0.324</td>
<td>0.345</td>
</tr>
<tr>
<td>Trypsin Inhibitors</td>
<td>1.45&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.47&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Tannins</td>
<td>&lt;0.05</td>
<td>NR</td>
</tr>
<tr>
<td>Phylic Acid</td>
<td>0.50</td>
<td>NR</td>
</tr>
<tr>
<td>Lectin Activity (Sheep)</td>
<td>0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>NA</td>
</tr>
<tr>
<td>Lectin Activity (Horse)</td>
<td>0&lt;sup&gt;b&lt;/sup&gt;</td>
<td>NA</td>
</tr>
<tr>
<td>Saponin</td>
<td>&lt;0.003</td>
<td>NR</td>
</tr>
</tbody>
</table>

Abbreviations: NA, not applicable; NR, not reported  
<sup>a</sup> Trypsin inhibitor reported in mg/g.  
<sup>b</sup> Lectin measured by hemagglutinin activity. Figures indicate the highest dilution where agglutination is still observed. A zero (0) result indicates no activity. Cells from both sheep and horse were tested with all results equal to zero.  
See Attachment B-4-1 for summary of analyses.
### Test Methods

<table>
<thead>
<tr>
<th>Description</th>
<th>Alkaloids (%ar)</th>
<th>Oligosaccharides (%ar)</th>
<th>Phenolics (%ar)</th>
<th>Trypsin Inhibitors (mg/g ar)</th>
<th>Cadmium (ICP-MS) (mg/kg ar)</th>
<th>Lead (ICP-MS) (mg/kg ar)</th>
<th>Tannins (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Flour</td>
<td>0.012</td>
<td>0.013</td>
<td>5.6</td>
<td>0.324</td>
<td>1.45</td>
<td>1.47</td>
<td>0.04</td>
</tr>
<tr>
<td>Protein 1</td>
<td>0.002</td>
<td>0.002</td>
<td>0.4</td>
<td>0.125</td>
<td>0.099</td>
<td>0.10</td>
<td>0.06</td>
</tr>
<tr>
<td>Protein 2</td>
<td>0.003</td>
<td>0.003</td>
<td>0.4</td>
<td>0.307</td>
<td>0.94</td>
<td>0.95</td>
<td>0.06</td>
</tr>
<tr>
<td>Kernel fibre 1</td>
<td>0.008</td>
<td>0.009</td>
<td>1.5</td>
<td>0.168</td>
<td>2.48</td>
<td>2.54</td>
<td>0.03</td>
</tr>
<tr>
<td>Kernel fibre 2</td>
<td>0.0005</td>
<td>0.0005</td>
<td>0.2</td>
<td>0.066</td>
<td>0.17</td>
<td>0.17</td>
<td>0.14</td>
</tr>
<tr>
<td>Hull fibre</td>
<td>0.001</td>
<td>0.001</td>
<td>0.6</td>
<td>0.126</td>
<td>0.15</td>
<td>0.15</td>
<td>0.12</td>
</tr>
</tbody>
</table>

Analyses conducted by Dept of Industry and Resources Chemistry Centre (WA)

### Agrifood Technology have determined the following result:

- Lupin flour (#103, 250g sample) phomopsins < 5 ppb
- ANZFA and Department of Health UK have a maximum permitted value for human consumption of 5ug/kg seed (5ppb)
December 10, 2008

Robert L. Martin, Ph.D.
Office of Food Additive Safety (HFS-200)
Center for Food Safety and Applied Nutrition
Food And Drug Administration
5100 Paint Branch Parkway
College Park, MD 20740-3835

Re: Withdrawal of Generally Recognized As Safe (GRAS) Exemption Notices for Sweet Lupin Fiber, Sweet Lupin Flour, and Sweet Lupin Protein

Dear Dr. Martin:

This letter is to inform you that we would like to withdraw our GRAS Exemption Notices for Sweet Lupin Fiber, Sweet Lupin Flour, and Sweet Lupin Protein, which were forwarded to your office on September 26, 2008.

Thank you for your kind attention to this matter. Please contact me should you have any questions regarding the withdrawal of these Notices.

Sincerely,

Peter Schutz
Chief Executive
George Weston Technologies
A Division of George Weston Foods Limited
peter.schutz@gwf.com.au