



Final Report Protocol GIL-1747

October 12, 2017

Holista Biotech Corp.

Glycemic Index determination of:

Low GI Noodles

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Table of Contents

Table of Contents	2
Investigators	3
Lead Investigator:	
Co-Investigators:	
Summary	
Background	
Study Objective	
Methods	
Subjects	6
Protocol	
Informed Consent	
Test Meals	
Table 1. Nutrient content of test meals	
Palatability	
Blood Samples	
Data Analysis	
Results	
Table 2. Subject details	
Analytical Variation for Blood Glucose and Within Subject Variation of Reference Foo	
Adverse Events and Protocol Deviations	
Palatability	
Blood Glucose Response	
Glycemic Index	
Table 3. Palatability, Incremental Area Under the Curve, Glycemic Index and Glycemic	
Index Category	
References	
Appendix 1	
Macronutrients and ingredients of test foods	
Appendix 2	
Abbreviations	
Definitions	







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GIL-1747 Page 3 of 16



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Summary

The glycemic index (GI) value of one food was determined using the standard ISO method (ISO 26642:2010). Ten subjects consumed the test food (noodles) once and the control meal from white bread twice. All meals contained 25g of available carbohydrate. The GI values are expressed on the glucose scale where the GI of glucose=100 and white bread=71. The GI values (Mean±SEM) of the foods tested were:

Test Meal	Glycemic Index	GI Category
White Bread Control	71	High
Low GI Noodles	38±4	Low

^{*} The listed GI value(s) is/are only valid as long as product ingredients, formulation, processing, and/or any other material production factors remain unchanged.



GIL-1747 Page 4 of 16

^{**} Using the classification of ISO 26642:2010, products with a glycemic index (GI) less than or equal to 55 are classified as being low GI, those with a GI of 56 to 69 are classified as medium, while those with a GI equal to or greater than 70 are high GI.



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Background

The glycemic index (GI) was proposed in the 1980s as a means to classify carbohydrate foods according to their effect on postprandial blood glucose responses (Jenkins et al, 1981). Low GI foods release their carbohydrate slowly and elicit a lower glycemic response while high GI foods are rapidly digested with a corresponding higher glycemic response. The rate of glucose absorption and extent and duration of elevated blood glucose levels induce many hormonal and metabolic changes that may affect health or disease parameters. Low GI diets may help in weight maintenance (Larsen et al, 2010) and weight loss (Ebbeling et al, 2003) in addition to being protective against chronic disease such as diabetes (Salmeron et al, 1997a,b), heart disease (van Dam et al, 2000; Lui et al, 2002) and certain cancers (Augustin et al, 2001; Francheschi et al, 2001). Interest in identifying low GI foods and the food factors responsible for the low GI of foods has therefore increased. Several food factors have been identified that influence *in vivo* absorption and therefore potentially the GI of a food or meal. Some of these factors include: the gross matrix structure, cell wall and starch structure (i.e. ripening), amylose to amylopectin ratio, and viscous fibre (Brand et al, 1985).

The methodology for determining the GI is now well established (Wolever et al, 1991; Brouns et al, 2005) and has been shown to be reproducible by laboratories across the world (Wolever et al, 2003, 2008). The protocol used by Glycemic Index Laboratories to determine the GI of foods adheres to or exceeds the methods specified by the International Organization for Standardization in: ISO 26642:2010 "Food products - Determination of the GI and recommendation for food classification".

Study Objective: The objective of this study was to determine the GI of Holista Low GI Noodles.



GIL-1747 Page 5 of 16



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Methods

Subjects

Inclusion criteria: Subjects were males or non-pregnant females aged 18-75 years and in good health.

Exclusion and withdrawal criteria: Subject less than 18 years old or older than 75 years; with a known history of AIDS, hepatitis, diabetes or a heart condition. Subject taking medication or with any condition which might, in the opinion of Dr. Wolever either make participation dangerous to the subject or to others or affect the results.

Number of Subjects: A total of ten (10) subjects were studied. Using the t-distribution and assuming an average coefficient of variation (CV) of within individual variation of incremental area under the blood glucose curve (IAUC) values of 25%, n=10 subjects has 80% power to detect a 33% difference in IAUC with 2 tailed p<0.05.

Protocol

Glycemic Index Testing: The protocol used at GI labs for determining the GI of foods follows the methods described in ISO 26642:2010 - "Food products - Determination of the glycaemic index and recommendation for food classification".

The study used an open-label, randomized crossover design. Each subject underwent treatments on separate days, with each subject performing up to 3 tests per week separated by at least one day. On each test day, subjects came to Glycemic Index Laboratories (20 Victoria Street, 3rd floor) in the morning after a 10-14 hour overnight fast. After being weighed and having two fasting blood samples obtained by finger-prick five minutes apart, the subject then consumed a test meal within 15 minutes. Further blood samples were obtained at 15, 30, 45, 60, 90 and 120 minutes after the start of the test meal. Subjects remained seated quietly during the 2 hours of the test. After the completion of the test they were offered a snack and then allowed to leave.

Informed Consent

The GI Laboratories protocol has been approved by the Western Institutional Review Board® which meets all the requirements of the US Food and Drug Administration (FDA), the Department of Health and Human Services (DHHS), the Canadian Health Protection Branch (HPB), Canadian Institutes of Health Research (CIHR) and the European Community Guidelines. All subjects provided written informed consent prior to starting the study.



GIL-1747 Page 6 of 16



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Test Meals

The test meals consisted of a portion of the test food or white bread control containing 25g available carbohydrate (defined as total carbohydrate minus dietary fiber/non-digestible carbohydrates). The portion size of the test foods was calculated based on the results of nutrition analysis provided by the client (Table 1). Each subject was given a choice of a beverage (water, coffee or tea with 30ml of 2% milk and non-caloric sweetener if desired) to consume with the test meal; the beverage chosen was kept the same for all test meals. The control meal was tested 2 times by all subjects. Test meals were given in random order.

Food preparation:

White bread Control: Bread was baked in a bread maker in loaves containing 500g available carbohydrate. The ingredients for each loaf (510ml warm water, 694g all-purpose flour, 14g sugar, 8g salt and 13g yeast) were placed into the bread maker according to instructions, and the machine turned on. After the loaf had been made, it was allowed to cool for an hour, and then weighed and after discarding the crust ends, the remainder was divided into portion sizes containing 25g available carbohydrate. These portions were frozen prior to use, and reheated in the microwave prior to consumption.

Low GI Noodles: 37.3g was weighed out and added to boiling water. The noodles were boiled for 3 minutes drained and then served immediately to the subject.

Table 1. Nutrient content of test meals

Test Meal	Abbr	Weight (g)	Protein (g)	Fat (g)	T CHO	Fibre (g)	AvCHO (g)
White bread	WB25	56	4.07	0.31	27.12	1.33	25.8
Low GI Noodles	Noodl	37.3	4.82	0.88	26.32	1.32	25

^{*}Average of last 3 macronutrient analyses by Silliker Labs.

Palatability

After consuming the test meal, subjects rated the palatability of the test meal using a visual analogue scale consisting of a 100mm line anchored at the left end by "very unpalatable" and at the right end by "very palatable". Subjects made a vertical mark along the line to indicate their perceived palatability. The distance from the left end of the line to the mark made by the subject is the palatability rating; the higher the value, the higher the perceived palatability.



GIL-1747 Page 7 of 16



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Blood Samples

Blood samples (2-3 drops each) were collected into 5 mL tubes containing a small amount of anticoagulant (sodium fluoride/potassium oxalate). The samples were mixed by rotating the tube vigorously and then refrigerated during the testing session. After completion of the test session, samples were stored at -20°C prior to glucose analysis. Blood glucose analysis, using a YSI (Yellow Spring Instruments, OH) analyzer, took place within five days of collection.

Data Analysis

Data were entered into a spreadsheet by 2 different individuals and the values compared to assure accurate transcription. IAUC values were calculated using the trapezoid rule, ignoring area beneath the baseline. For the purpose of the IAUC calculation, fasting glucose was taken to be the mean of the first measurement of the glucose concentration at times -5 and 0min. Glucose was measured in the 0min fasting sample 2 times and the data used to determine the standard deviation (SD) of the analytical variation as follows:

SD=
$$\sqrt{(\sum d^2/n)}$$

The GI was calculated by expressing each subject's glucose IAUC for the test food as a percentage of the same subject's mean response after reference meal and, if required, adjusting the GI to the glucose scale where glucose = 100 and white bread =71. A second statistical analysis was done on the GI values after excluding those values >2SD above the mean in which case excluded values were replaced by the mean of the remaining values and the error degrees of freedom in the ANOVA was reduced by the number of outliers excluded. After demonstrating significant heterogeneity, the differences between individual means were assessed using Tukey's test to control for multiple comparisons, with the criterion for significance being 2-tailed p<0.05. Means which differ by more than the least significant difference (LSD) differ significantly.



GIL-1747 Page 8 of 16





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Results

Ten (10) subjects (5 males and 5 females), aged 49±14 years with a body mass index of 29.4±4.1kg/m² participated in the study. The individual details are shown in Table 2.

Table 2. Subject details

ID	Sex	Ethnicity	Age	•		Weight		BMI
		·	(yrs)	(cm)	(in)	(kg)	(lb)	(kg/m²)
103	F	Caucasian	60	165.0	64.4	84.5	185.9	31.0
143	F	Caucasian	55	164.0	64.0	91.4	201.1	34.0
191	М	Caucasian	58	192.0	74.9	111.2	244.6	30.2
299	F	Caucasian	58	165.0	64.4	92.5	203.5	34.0
333	F	South Asian	59	151.0	58.9	76.2	167.6	33.4
436	М	Chinese	55	167.4	65.3	74.9	164.8	26.7
554	М	Caucasian	40	165.0	64.4	85.8	188.8	31.5
572	F	Black	49	159.5	62.2	59.6	131.1	23.4
628	М	Caucasian	19	164.0	64.0	63.8	140.4	23.7
643	М	South Asian	32	169.5	66.1	75.2	165.4	26.2
ME	AN		48.5	166.2	64.8	81.5	179.3	29.4
S	D		13.8	10.4	4.0	15.0	33.1	4.1



GIL-1747 Page 9 of 16



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Analytical Variation for Blood Glucose and Within Subject Variation of Reference Food

Analytical variation: Duplicate analysis was performed on 28 samples taken at 0min. The mean \pm SD of blood glucose in these samples was 4.39 ± 0.035 mmol/L for a CV of 0.8%, which is <3.6% and, thus, satisfactory (ISO 26642:2010). The mean \pm SD for the 30 -5 and 0min samples was 4.40 ± 0.093 mmol/L for a CV of 2.1%, which is greater than analytical variation because it reflects both analytical variation and minute-to-minute variation in blood glucose.

Within subject variation of reference food: There was no significant effect of order on the IAUC values after the repeated white bread controls. The mean within- CV of the IAUC values after the 2 repeated white bread controls was 19.7%. The tests appeared to be technically satisfactory, as judged by the average within-subject variation of glycemic responses for the repeated white bread tests. Values less than 30% are considered to be satisfactory (ISO 26642:2010).

Adverse Events and Protocol Deviations

All test meals were well tolerated and no adverse events were reported.

Palatability

Palatability scores are given in Table 3. There were no statistically significant differences between palatability scores of the noodles and white bread.

Blood Glucose Response

The blood glucose responses to the test meals are shown in figure 1. Mean fasting blood glucose was identical before each test meal within each series. The tabulated blood glucose responses can be viewed on the analysis pages (GICalc Report GIL-1747 Holista data noodles.pdf).



GIL-1747 Page 10 of 16





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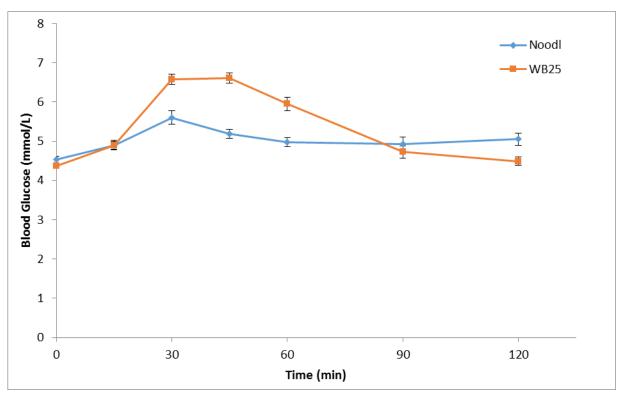


Figure 1: Postprandial glucose responses to Low GI Noodles (Noodl) and the White Bread Control (WB25) (mean of 2 meals) 10 subjects each. All meals contained 25g of available carbohydrate. Data are expressed as Mean±SEM

Glycemic Index

The data and calculations for the GI are given in the pdf sheets entitled: GICalc Report GIL-1747 Holista data noodles.pdf. There were no statistical outliers and all data were used to calculate the GI.



GIL-1747 Page 11 of 16





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Table 3. Palatability, Incremental Area Under the Curve, Glycemic Index and Glycemic Index Category

Test Meal	Abbr	Palatability (mm)	IAUC (mmol×min/L)	GI	GI Category
White bread Control	WB25	69±6ª	127.2±10.9ª	71 ^a	High
Low GI Noodles	Noodl	66±8 ^a	66.1±7.9 ^b	38±4 ^b	Low

Results are given as Mean±SEM.

Report Prepared by:

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Director of Research



GIL-1747 Page 12 of 16

^{*} GI values are given after outliers were excluded and by category (Category: high (GI ≥70), medium (56≥GI≤-69), or low,(GI≤55) (ISO 26642:2010)

ab Numbers within the same column with a different letter in the superscript are statistically significantly different (P<0.05).



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GIL-1747 Page 13 of 16





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Appendix 1

Macronutrients and ingredients of test foods

	Kipling Quality Systems	Issue Date:	Aug 02 2017
WINGS	R&D Product Spec Sheet	Revision:	Aug 23 2017
Wing's Food Products	Low GI Noodles	Pag	e 1 of 1

Low GI Noodles



Sample Overview

Product Name	Low GI Noodles
Ingredients	Enriched wheat flour, water, liquid whole egg, liquid egg white, barley, lentil, fenugreek, okra, salt.
Allergen Information	Contains: Wheat, Egg
Recommended Cooking Instructions	Add $100\mathrm{g}$ noodles to $2\mathrm{L}$ boiling water. Cook for 3 minutes, stirring occasionally. Drain and serve.

Approved By	Document Location:
	Y:\2016 Low GI Noodles\2017.08.17 - Samples for GI testing\Spec Sheet - Low GI Noodles 4% - revised cooking time -
	Aug 23 2017.docx



GIL-1747 Page 14 of 16

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Appendix 2

Abbreviations

BMI body mass index

CHO carbohydrate

CV coefficient of variation

GI glycemic index GL glycemic load

IAUC incremental area under the curve

min minutes

SEM standard error of the mean

SD standard deviation

WB white bread

ANOVA analysis of variance

Definitions

Glycemic Index

a ranking of carbohydrate-containing foods according to the extent to which they raise blood glucose levels after being eaten

Glycemic Load

a ranking of carbohydrate-containing foods according to the amount of available carbohydrate in the serving and its glycemic index

In vivo

occurring in living organisms (ie. humans)

Macronutrient

one of the nutritional components of the diet: fat, protein, or carbohydrate

Postprandial

following a meal



GIL-1747 Page 15 of 16



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GI Labs offers a full range of research services, in healthy populations or specific target populations, including:

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