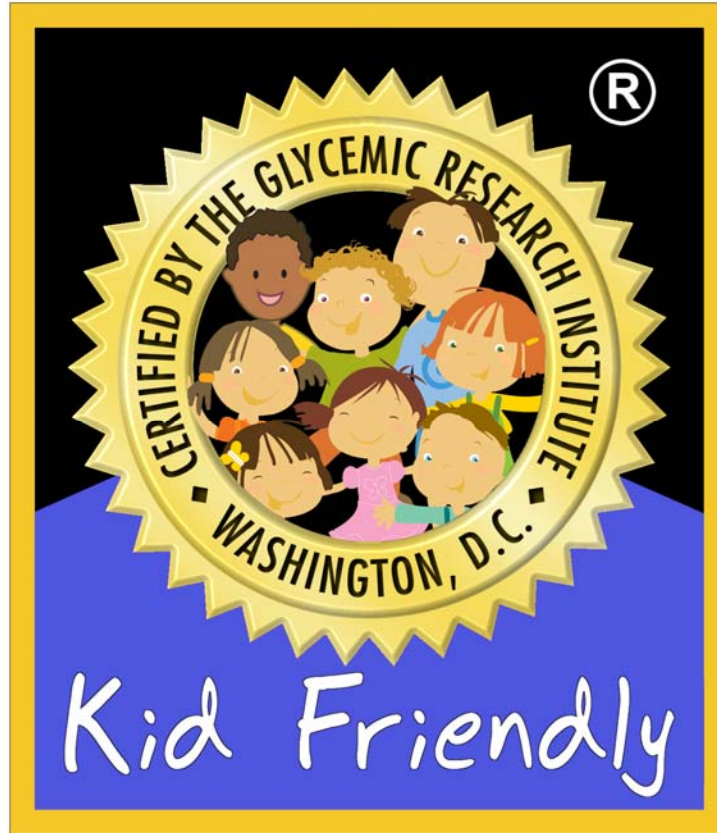


FINALLY-AN ENDORSEMENT THAT MATTERS  
GLYCEMIC RESEARCH INSTITUTE "KID FRIENDLY" CERTIFICATION



- A Professional Clinical Research Organization Specializing in FDA Claim Substantiation
- The only organization providing Kid Friendly Certification & Clinical Trials
- Serving the Largest Food & Research Companies in the World
- Official Clinical Testing Facility for the Glycemic Research Institute
- Board Approved Human In Vivo Clinical Trials
- Member of Association of Clinical Research Professionals
- 25 + Years of Glycemic Expertise

**Glycemic Research Laboratories**  
**Glycemic Solutions**  
111 Second Avenue N.E. Suite 512  
St. Petersburg, Florida 33701

THE OFFICIAL CLINICAL TESTING FACILITY FOR  
GLYCEMIC RESEARCH INSTITUTE

**GLYCEMIC RESEARCH**  
**LABORATORIES**  
WWW.GLYCEMICRESEARCHLABORATORIES.COM

**Glycemic Solutions Corporation**

111 Second Avenue N.E., Suite 512  
St. Petersburg, Florida 33701  
www.GlycemicIndexTesting.com

**INSTRUCTIONS FOR SUBMISSION OF TEST FOODS**

**REVIEW PROCESS**

To submit a product (herein the “**Test Food**”) for “**Kid Friendly**” clinical studies:

- 1) Fill out and sign Documents **A, B, & C** and submit via fax to:  
**Glycemic Solutions Corporation: FAX (727) 894-6969**

Document “ <b>A</b> ”	Product Submission Form	1 Page
Document “ <b>B</b> ”	Label and Ingredient Information	1 Page
Document “ <b>C</b> ”	Standard Terms & Conditions Agreement for Services	2 Pages

- 2) The Glycemic Solutions/Glycemic Research Laboratories (GS/GRL) Medical Advisory Board will review the application and respond via email or phone. Clients are assigned a Clinical Studies Coordinator who will work one-on-one to answer any questions, and to coordinate the project.
- 3) If the product is accepted by GS/GRL for Board Approved Human In Vivo Clinical Trials, directions on submitting the Test Food (s) will be forthcoming.

### **CLINICAL TRIAL COSTS**

The cost of clinical testing will depend on the Protocol utilized during the trials. The GS/GRL Medical Advisory Board will review the Application form and product label, as well as the client's desired legal label claims. Per this review, the client will be contacted by GS/GRL with specific pricing.

### **SUBMISSION FEES**

Submission fees are due prior to beginning the clinical trials. All fees are based on U.S. dollars. Payment may be in the form of a check, money order, or bank wire. Contact the Clinical Studies Coordinator (CSC) for bank wiring instructions.

### **SUBMISSION OF TEST FOODS**

Approved Test Foods (products submitted for trials) will be sent to GS/GRL per *Test Food Submission Guidelines*. The Clinical Studies Coordinator will direct the client in all phases of the Application process.

### **TRIAL RESULTS**

Following GS/GRL Human In Vivo Clinical Trials, the client will be advised of the results in a comprehensive written report. The final Official report is sent directly to the client. Follow-up conference calls are available to clients who wish to discuss the results of the final report.

### **GOVERNMENT REGULATION: GLYCEMIC CLAIMS**

Per government regulations regarding Claims and label statements, Glycemic Solutions/Glycemic Research Laboratories does *not* quantify the Glycemic Index of any product without Human In Vivo Clinical Trials.

*For more information on Legal/Illegal Glycemic Claims, see*

***AVOID FDA ACTION  
AGAINST ILLEGAL GLYCEMIC LABEL CLAIMS  
WWW.GLYCEMIC.COM***



## PRODUCT SUBMISSION OVERVIEW

Fill out the appropriate documents provided herein (Documents A, B, & C). Submit Documents **A, B, & C** to Glycemic Solutions/Glycemic Research Laboratories (GS/GRL) via **FAX** to **(727) 894-6969**.

Products submitted cannot be returned to the client.

Any product sample suspected of, or containing a pathogen or substance considered hazardous will be rejected. Glycemic Solutions/Glycemic Research Laboratories reserves the right to refuse any product deemed unfit for human consumption.

Products accepted for submission must arrive in a sanitary condition in appropriate packaging. The product must be packed with a label and ingredient panel (see Doc C).

Products should *not* be physically submitted until the client has been contacted by Glycemic Solutions/Glycemic Research Laboratories. If the GS/GRL Medical Advisory Board has determined that the product(s) are acceptable for clinical trials, the client will be notified via phone by a Clinical Studies Coordinator.



## INSTRUCTIONS FOR SHIPPING TEST FOODS FOR CLINICAL TRIALS

### ACCEPTANCE: HUMAN IN VIVO CLINICAL TRIALS

Once the Test Food (s) has been approved by the Glycemic Solutions/Glycemic Research Laboratories Medical Advisory Board, a Clinical Studies Coordinator will contact the client to arrange shipping of the Test Food (s).

### SHIPPING THE PRODUCT FOR CLINICAL TRIALS

The Test Food (s) are to be shipped to Glycemic Solutions/Glycemic Research Laboratories (see address below). The product may be sent via Fed-Ex, UPS or Postal Service (depending on shelf-life and stability of the Test Food). Overnight shipping is preferred. If the product is heat-sensitive, pack with a dry-ice unit, and ship via Fed-Ex-Overnight.

### SHIPPING PERISHABLE FOODS

When shipping fresh foods, frozen foods, perishable foods (foods that need to be refrigerated or frozen), pack the food in dry ice and send via Fed-Ex overnight to be delivered Monday-Friday, 9 AM-5 PM. If the food does not arrive in an acceptable state, human testing cannot proceed. Notify your assigned Glycemic Solutions Clinical Studies Coordinator as to when the product will arrive.

### **TEST FOODS SHIPPING ADDRESS:**

#### **GLYCEMIC SOLUTIONS CORPORATION**

111 Second Avenue N.E. Suite 512  
St. Petersburg, Florida 33701  
(727) 894-6900 Fax (727) 894-6969

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**Glycemic Research Laboratories**  
**Glycemic Solutions**  
111 Second Avenue N.E. Suite 512  
St. Petersburg, Florida 33701



### **CALCULATING THE AMOUNT OF TEST FOOD TO BE SHIPPED FOR CLINICAL STUDIES**

The Medical Advisory Board will determine the amount (gram weight and serving size) of the Test Food required for clinical testing. The guidelines for selecting the amount of the Test Food to be fed to human subjects is determined by FDA guidelines related to appropriate food servings in foods, Nutraceuticals, and Pharmaceuticals and Glycemic Solutions Board Approved Clinical Trials Protocols. The client will be advised as to the amount of servings and total product required.

### **RIGHT TO REFUSE TEST FOODS**

Glycemic Solutions/Glycemic Research Laboratories (GS/GRL) reserves the right to refuse any product (Test Food) submitted to GS/GRL, per the Rules & Regulations as set forth by the GS/GRL Medical Advisory Board.

### **QUESTIONS**

Questions regarding the status of products may be addressed to the Clinical Studies Coordinator at (727) 894-6900. Prior to beginning clinical studies, specific protocols will be discussed with the client.



**PRODUCT  
SUBMISSION  
FORM  
DOCUMENT "A"**

**Glycemic Solutions Corporation**

111 Second Avenue N.E., Suite 512  
St. Petersburg, Florida 33701  
www.GlycemicIndexTesting.com

Company Name & Address

\_\_\_\_\_

\_\_\_\_\_

\_\_\_\_\_

Contact Person \_\_\_\_\_

Title \_\_\_\_\_

Date \_\_\_\_\_

PO # \_\_\_\_\_

Email \_\_\_\_\_

Web \_\_\_\_\_

Phone \_\_\_\_\_

Fax \_\_\_\_\_

Name of Product Submitted \_\_\_\_\_

Description of Product \_\_\_\_\_



**Kid Friendly Clinical Study**

**Total Number of Products Submitted** \_\_\_\_\_  
(For each product submitted, fill out an additional form.)

THE OFFICIAL CLINICAL TESTING FACILITY FOR  
GLYCEMIC RESEARCH INSTITUTE

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**LABEL AND INGREDIENT INFORMATION:  
DOCUMENT “B”**

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**LABEL AND INGREDIENT INFORMATION:**

Place the product label and ingredients panel on this page



**STANDARD TERMS & CONDITIONS AGREEMENT FOR SERVICES  
DOCUMENT “C”**

All services provided by Glycemic Solutions/Glycemic Research Laboratories (GS/GRL) and/or its affiliates are subject to the terms and conditions stated herein.

Client signature on this Document constitutes acceptance of the Terms and Conditions as stated herein.

**A. USE OF NAME**

The client agrees not to use Glycemic Solutions/Glycemic Research Laboratories and/or its affiliates name or data in any manner which might cause harm to said companies reputation and/or business. Under no circumstances is the name of Glycemic Research Institute (GRI), Glycemic Solutions (GS), Glycemic Research Laboratories, and/or its affiliates to be published, either alone or in association with that of any other party, without prior written approval from the corporate offices of GRI/GS.

**B. PAYMENT TERMS**

Total payment is due prior to the onset of the clinical studies. This fee is non-refundable.

**C. TRIAL RESULTS**

Following completion of the Human In Vivo Clinical Trials, the client will be notified of the results. An Official Report will be generated by GS and sent to the client.

**INITIAL**\_\_\_\_\_

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**Glycemic Research Laboratories  
Glycemic Solutions**  
111 Second Avenue N.E. Suite 512  
St. Petersburg, Florida 33701

**D. REFORMULATION**

If the Test Food (s) submitted does not qualify as “LOW GLYCEMIC” and/or “LOW GLYCEMIC FOR DIABETICS,” the client may elect to reformulate and resubmit the product. Prior to the trial, or post-trial, Low Glycemic product development is available upon request.

**F. FORMULA CHANGES**

Any change or modifications in the product formula (Test Food) as tested by GS invalidates prior clinical testing results and/or use of any GRI Mark. No claims or statements as related to GS Glycemic Testing may be made in relation to altered Test Foods as previously tested by GS. Any formula or ingredient changes to Test Foods previously approved as “Low Glycemic” by GS requires updated Clinical Trials for validation by GS and/or GRI.

**SIGNATORY PAGE**

The client agrees to the Terms & Conditions as stated herein.

\_\_\_\_\_  
Client Signature Date \_\_\_\_\_

\_\_\_\_\_  
Client Printed Name

\_\_\_\_\_  
Corporation

**LEGAL COPYRIGHT © NOTICE:** *This report is the sole property of Glycemic Research Laboratories/Glycemic Solutions (GS) Corporation, and may not be copied in any format or portion without express prior written permission from GS.*