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# **ACCESS BUSINESS GROUP RAW MATERIAL QUESTIONNAIRE**

# **SUPPLIER DETAILS**

SUPPLIER / MANUFACTURER:					
Product Code/Name:					
Company Name:					
Address:					
City, State, ZIP Code:					
Country:					
Contact Name:					
Email Address:					
Phone Number:					
Fax Number:					
<b>Type of supplier</b> : (Broker, Grower, Manufacturer, Other):					
Name of Manufacturer:					
Country of Manufacture:					
Can this material be ordered directly from the manufacturer (if different)?					
Can this material be ordered directly from					
RAW	MATERIAL STATUS				
	MATERIAL STATUS				
RAW		☐ Yes ☐ No			
RAW  REGULATORY STATUS:  Has this product or raw material been sold on Regulatory status (GRAS, etc):  Pharmacopoeial reference (USP, Commission	the USA retail market prior to 1994?  n E, etc.)				
RAW  REGULATORY STATUS:  Has this product or raw material been sold on Regulatory status (GRAS, etc):  Pharmacopoeial reference (USP, Commission Total Lead ppm (Required for CA Prop 65 Commission CA)	the USA retail market prior to 1994?  n E, etc.)  mpliance):				
RAW  REGULATORY STATUS:  Has this product or raw material been sold on Regulatory status (GRAS, etc):  Pharmacopoeial reference (USP, Commission Total Lead ppm (Required for CA Prop 65 Con Conforms to established European purity criteria)	the USA retail market prior to 1994?  n E, etc.)  mpliance):				
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RAW  REGULATORY STATUS:  Has this product or raw material been sold on Regulatory status (GRAS, etc):  Pharmacopoeial reference (USP, Commission Total Lead ppm (Required for CA Prop 65 Con Conforms to established European purity crite Approved in the following markets:	the USA retail market prior to 1994?  n E, etc.)  mpliance):  ria for this material :				
RAW  REGULATORY STATUS:  Has this product or raw material been sold on Regulatory status (GRAS, etc):  Pharmacopoeial reference (USP, Commission Total Lead ppm (Required for CA Prop 65 Con Conforms to established European purity criteria)	the USA retail market prior to 1994?  n E, etc.)  mpliance):  ria for this material :				
RAW  REGULATORY STATUS:  Has this product or raw material been sold on Regulatory status (GRAS, etc):  Pharmacopoeial reference (USP, Commission Total Lead ppm (Required for CA Prop 65 Con Conforms to established European purity crite Approved in the following markets:	the USA retail market prior to 1994?  n E, etc.)  mpliance):  ria for this material :				
RAW  REGULATORY STATUS:  Has this product or raw material been sold on Regulatory status (GRAS, etc):  Pharmacopoeial reference (USP, Commission Total Lead ppm (Required for CA Prop 65 Con Conforms to established European purity crite Approved in the following markets:  KOSHER / HALAL STATUS (provide copy of the copy	the USA retail market prior to 1994?  n E, etc.)  mpliance):  ria for this material :	/			

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### **BREAKDOWN & SOURCE OF RAW MATERIAL**

INGREDIENT /				European	(√) source				
SUB-INGREDIENT  (name as it should appear on product labeling)	Percent (%) (Range is acceptable)	(e.g. sour carri	., Nutrient ce, binder, er, solvent, etener, etc.)	E-number  (Conforms to European purity criteria)	Natural	Synthetic	Animal	Other? (bio- fermentatio	Please List Source.  If plant or animal source, also fill out sections below
TOTAL =	100%								
Genus & Species (variety / cultivar if available)	Pla Part			of reference s o identify spe				try Of (Crop)	Endangered species (Y/N)
* Source of reference standard ex	kamples: botani	cal vouch	ner specimen,	chemical refere	ence sta	ndard,	chain of	custody	r, etc.:
ANIMAL SOURCE (example	e: fish, cattl	e, swir	ne, birds, n	nollusks, et	c.)				
Complete this section if Mate	erial or Sub-o	compon	ent <u>is deri</u>	ved in whole	or pa	rt from	anima	al sour	ces
Animal Common Name:									_
Genus and species:									<u> </u>
Country of Origin of Animal(s	s):								<u> </u>
Is this ingredient a milk deriv	ative?		Yes	No					
If <b>not a milk derivative</b> , plea 1. Country of Animal		g:							<u> </u>
2. Animal Body Part	(s) Utilized:								<u> </u>
3. Animal Sub-ingred	dient(s):								

\* If your product is bovine-derived (other than from milk) and sourced from a country other than the United States, please attach a **BSE certificate** from the **exporting country** and **an import certificate from the USDA**.

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## **PROCESSING INFORMATION**

MANUFACTURING PROCESS:	
Please provide a Manufacturing Process Flow Chart to satisfy foreign market registration requirements	<b>S</b> .
Brief Process description (ex. "single step alcohol extraction"):	
Are solvents used in the manufacturing process?   Yes   No	
Solvent(s) used & percentage (strengths) of each solvent:	
Solvent residual level (ppm):	
Is this a plant/botanical material?	
Plant Materials only: Native extraction ratio (Crude botanical: finished product, less excipients):	
Final extraction ratio (Crude botanical: finished product):	

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### **GMO STATUS OF RAW MATERIAL**

Please  $\underline{\textbf{READ ALL 5 OPTIONS}}$  below and fill out the  $\underline{\textbf{ONE}}$  which most accurately describes the status of your raw material (including subcomponents/excipients).

1. IDENTITY PRESERVED ( IP );				
Must be able to confirm all the following and provide documentation:				
DOES NOT contain raw materials originating from a GM-crop or GM-microorganism, either main ingredient or any sub component/excipient (such as MALTODEXTRIN, ETHYL ALCOHOL etc.)				
DO HAVE a QA system in place to verify the non-GM origin of all raw materials used in products not declared GMO and to ensure proper separation of GM derived raw material from non-GM derived raw material throughout your total production process.				
HAVE PROVIDED with this questionnaire a detailed description of your internal quality system (including procedures for cleaning, segregation, seed origin etc. and copies of any certificates or validations mentioned in the quality system) and/or a copy of a 3 <sup>rd</sup> Party Certification.				
3 <sup>rd</sup> Party Certifying Body: Expiration date:				
2. PCR NEGATIVE:				
Must be able to confirm all the following and provide documentation:				
CONTAINS raw materials originating from a crop or microorganism in which there exists GM versions (for instance: from corn, soy, potato, tomato, cotton, sugar beet, rapeseed, alfalfa, etc.), for either main ingredient or any sub component/excipient (such as MALTODEXTRIN, ETHYL ALCOHOL etc.) but unable to provide traceability to support not from a GM-crop or GM-microorganism.				
Which component(s) in this material may be derived from GM?				
If known, what commercial GM crop was used ? (example: Roundup Ready Corn GA21)				
DOES test PCR Negative for presence of GM-protein/DNA.				
☐ HAVE PROVIDED with this questionnaire a Certificate of Analysis with PCR Negative test results.				
3. GMO THRESHOLD:				
Must be able to confirm all the following and provide documentation:				
CONTAINS raw materials originating from a crop or microorganism in which there exists GM versions (for instance: from corn, soy, potato, tomato, cotton, sugar beet, rapeseed, alfalfa, etc.), for either main ingredient or any sub component/excipient (such as MALTODEXTRIN, ETHYL ALCOHOL etc.) but unable to provide traceability to support not from a GM-crop or GM-microorganism.				
Which component(s) in this material is suspect as GM?				
DOES contain less than 0.9% GM-protein/DNA				
☐ HAVE PROVIDED with this questionnaire a Certificate of Analysis with PCR test results.				

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# **GMO STATUS OF RAW MATERIAL (continued)**

4. CONSIDERED GMO:
Must be able to confirm all the following:
□ <b>CONTAINS</b> raw materials originating from a crop or microorganism in which there exists GM versions (for instance: from corn, soy, potato, tomato, cotton, sugar beet, rapeseed, alfalfa, etc.), for either main ingredient or any sub component/excipient (such as MALTODEXTRIN, ETHYL ALCOHOL etc.) but unable to provide traceability to support not from a GM-crop or GM-microorganism.
□ <b>DO NOT TEST</b> material in order to guarantee either PCR negative or less than 0.9% GM-protein/DNA.
5. GMO FREE:
Must be able to confirm all the following and provide documentation:
NO GM-crop or GM-microorganism exists for the raw material sources of any ingredient or any sub component/excipient (such as MALTODEXTRIN, ETHYL ALCOHOL etc.) in this material. For instance: NOT from corn, soy, potato, tomato, cotton, sugar beet, rapeseed, alfalfa, etc.
NO GM-labeling within the EU of the product(s) containing this raw material(s) will be required per the legal provisions of EC Directives 1829/2003 and 1830/2003 concerning the traceability & labeling of GM organisms & the traceability of food and feed products produced from GM organisms.
☐ HAVE PROVIDED with this questionnaire a Non-GMO Statement on your company letterhead.
OTHER GMO QUESTIONS – Animal Feed:
If this ingredient <u>or</u> any sub component/excipient is derived from animals (such as stearic acid, etc.), please check the ONE that applies below:
☐ We do <b>not</b> process any raw material derived from animals fed with GM-crops. *See Note.
- or -
☐ We do process raw material derived from animals fed with GM-crops. *See Note
*Note: A negative answer is not compulsory to avoid future GM-labeling in EU, but would be favorable

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## **ALLERGEN STATUS**

Does this ingredient contain, or is it derived from any of the following? (check appropriate box)
Are any of the following processed on the same equipment as the material you provide to us? (check box below)

	Conta Deri fro	ved	Processed on same Equipment		Contai Deriv	ved	Processed on same Equipment
Allergen	YES	NO	YES	Allergen	YES	NO	YES
Artificial Colors				Oats			
Artificial Flavors				Peach			
Artificial Preservatives				Peanut			
Barley				Rye			
Buckwheat				Sesame			
Caffeine				Shellfish			
Celery				Soy			
Corn				Spelt			
Crustaceans				Sugar			
Egg				Sulfites			
Fish				> Naturally occurring	g? 🗌		
Gluten				> Sulfite level (ppm)			<del></del>
Kamut				Tomato			
Lactose				Tree nuts			
Lupin				Wheat			
Milk				Yeast			
Molluscs				Other Nuts			
Mustard				> Specify			

Comments:

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# PLANT/BOTANICAL MATERIAL INFORMATION

CONTAMINANTS:				
Aflatoxin test results:				
Aflatoxin testing is performed on:	Crude Botanical	Finished Pro	duct	Both
Radioactivity testing is performed on:	Crude Botanical	Finished Pro	duct	Both
Microbiological testing is performed on:	Crude Botanical	Finished Pro	duct	Both
Heavy metals testing is performed on:	Crude Botanical	Finished Pro	duct	□Both
Heavy metals test method used (e.g. ICP-	MS vs. AA):			
ORGANIC STATUS:				
Is the crude botanical certified <b>Organic</b> ?			∐Yes	□No
If <b>NO</b> , are you willing to use a certified organic crude botanical or to move toward certified organic farming practices?			∐Yes	□No
If YES, please provide the name of the	e certifying agency AND cop	y of certificate:		
			_	
CULTIVATION DETAILS:				
Name of Manufacturer and grower/wild-co	llector (plant materials):			
Manufacturing location(s):				
State of crude botanical prior to processing	g:	Other_		
Can this material be ordered directly from	the manufacturer (if differen	t)?	∐Yes	□No
Do you (the supplier) directly grow the sou	rce crop on a farm controlle	d by you?	∐Yes	□No
If <b>NO</b> , do you (the supplier) have any of cultivation of the crude botanical us			□Yes	□No
Would you be willing to partner with Acces for the purpose of alignment of farming me			∐Yes	□No
For processing operations, would you be v	villing to use feedstock grow	n by ABG?	□Yes	□No

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### **BASIC MATERIAL INFORMATION**

Individual answers are optional if information is included on the product specification that  $\underline{\text{must}}$  accompany this document.

Item	Specification
Appearance (powder, liquid, etc)	
Bulk density (Tap and loose):	
Powder flow characteristics (e.g., Flowdex):	
Particle size distribution (USS sieve):	
Odor (If applicable):	
Taste (If applicable):	
Color (If applicable, provide expected L* a* and b* values):	
Moisture (Loss On Drying or similar):	
Coliform:	
E. coli:	
Aerobic Plate Count:	
Mold:	
Yeast:	
Salmonella:	
Shelf life:	
Storage Conditions:	
Packaging options (type of package, size options, etc.):	
Is bar-coding available on your packaging?	☐ Yes ☐ No
Marker(s) level(s) and analytical method or related activity such as ORAC:	
Pesticides (third party test results, including method):	
Radioactivity testing (third party test results):	
Heavy metals (Pb, Hg, Cd, As - third party test results):	
Heavy metals test method used (e.g. ICP-MS vs. AA):	
Sanitizing treatments used in raw material processing, including concentration:	
Clinical testing performed on this product	
(protocol, results, publications, attach if necessary):	
Describe any intellectual property or exclusivity agreements that would affect this product:	

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### **CERTIFICATION**

DOCUMENTATION CHECKLIST
Please attach the following documents to this completed questionnaire:
<ul> <li>Statement of Non-Irradiation, non-ETO and non-chemical sterilization of raw materials (signed)</li> <li>Certified Organic</li> <li>Nutrient Profile or Proximate Analysis</li> <li>Certificate of Analysis (C of A)</li> <li>MSDS</li> </ul>
I certify that the information provided in this document is true and correct.
Signature
Print Name

NOTE: Any modifications or revisions to this product must meet raw material specifications and should be forwarded to the Product Development Department:

NUTRILITE, A Division of Access Business Group 5600 Beach Blvd P.O. Box 5940 Buena Park, CA 90622-5940 (714) 562-6200 (phone)

Title