
	PREMIER MAGNESIA - GILES CHEMICAL COMPANY FORM			
	Title: Vendor Assessment Survey	Page : 1 of 5	Revision : 00 Effective Date : 08/27/12	
	Author: Deborah Durbin	Form Number: QA-012-F03		

In order to better understand your business, and as a step in establishing and maintaining a good business relationship with you, we request that you please provide us with the following information. Thank You.

GENERAL INFORMATION:

Company Name: _____

Contact Name: _____ Telephone: _____

Primary Address: _____

City, State, Zip Code: _____

Web Site: _____

Division or Subsidiary of (if applicable): _____

BUSINESS INFORMATION:

When was your company established? _____

Please describe your usual payment terms: _____

Are you willing to provide us with a financial statement? ☐ Yes ☐ No

Are you willing to provide us with a change-notification agreement? ☐ Yes ☐ No

Is your company ISO/QS registered? ☐ Yes ☐ No

If yes, who is the regulating body? _____

Please list three customer references:

1. Company Name: _____

Contact Name: _____ Telephone: _____

Address: _____

2. Company Name: _____



Contact Name: _____ Telephone: _____

Address: _____

3. Company Name: _____

Contact Name: _____ Telephone: _____

Address: _____

	PREMIER MAGNESIA - GILES CHEMICAL COMPANY FORM			
	Title: Vendor Assessment Survey	Page : 2 of 5		Revision : 00 Effective Date : 08/27/12
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FACILITIES INFORMATION:

Does your facility shut down for any extended periods (more than one week)? _____

Do your plants operate in accordance with EPA and OSHA standards? ☐ Yes ☐ No

Do you have a contingency plan in case of disaster? ☐ Yes ☐ No

PRODUCT INFORMATION:

Please list your top three product lines and their percentage of your business: % of Business

1. _____
2. _____
3. _____

QUALITY SYSTEM INFORMATION:



QUALITY SYSTEM	Yes	No	N/A
Do you have a formal Quality Manual?			
Do you have formal, company-wide quality procedures?			
Do you have procedures for performing, documenting and responding to internal audits?			

MANAGEMENT RESPONSIBILITY	Yes	No	N/A
Does your Quality organization have a designated Senior Management Representative?			
Do senior management representatives routinely review the quality system for effectiveness?			

PERSONNEL TRAINING	Yes	No	N/A
Does your training system identify all relevant training needs of each employee performing all processes?			
Are training records maintained for individual employees?			

DOCUMENT CONTROLS	Yes	No	N/A
Do you have procedures for the control of production and system documents?			
Do you have procedures to ensure that only current documents are being utilized?			
Do you maintain records of changes made to documents?			

PURCHASING CONTROLS	Yes	No	N/A
Do you have procedures for qualifying/approving suppliers that result in an approved supplier list?			
Are inspections performed on incoming materials?			

	PREMIER MAGNESIA - GILES CHEMICAL COMPANY FORM			
	Title: Vendor Assessment Survey Author: Deborah Durbin	Page : 3 of 5 Form Number: QA-012-F03	Revision : 00 Effective Date : 08/27/12	

PRODUCTION AND PROCESS CONTROLS/STATISTICAL TECHNIQUES	Yes	No	N/A
Do you use statistical methods to control your processes?			
Is the sampling inspection defined by a procedure?			
Is your sampling inspection adjusted on the basis of inspection/test results?			
Do you have a procedure for the approval and release of new processes?			
Do you have a procedure for the approval and release of new equipment?			
Do you have procedures for the maintenance/replacement of production equipment?			

MANUFACTURING	Yes	No	N/A
Do you have manufacturing procedures?			
Do you maintain Device History Records?			
Do you maintain schedules for maintenance of manufacturing equipment?			



INSPECTION, MEASURING AND TEST EQUIPMENT (CALIBRATION)	Yes	No	N/A
Is the calibration of your inspection, measuring and test equipment defined by a procedure?			
Is all inspection, measuring and test equipment identified as to its calibration status?			

ACCEPTANCE ACTIVITIES	Yes	No	N/A
Are Acceptance Inspection activities performed on raw materials, intermediate products and finished goods, where applicable?			

NON-CONFORMING PRODUCT	Yes	No	N/A
Are there written procedures for controlling non-conforming materials in Receiving?			
Are there written procedures for controlling in-process non-conforming materials?			
Are there written procedures for controlling non-conforming materials in Final Inspection?			
Are procedures used for the repair, rework and disposition of non-conforming materials?			

CORRECTIVE AND PREVENTIVE ACTION (CAPA)	Yes	No	N/A
Do you have a CAPA system implemented for customer complaints?			
Do you have a CAPA system implemented for supplier defects?			
Do you have a CAPA system implemented for internal defects?			
Is your CAPA system defined by a procedure?			
Is CAPA effectiveness reviewed?			
Are CAPA reports maintained as part of quality records?			

HANDLING, STORAGE AND DELIVERY	Yes	No	N/A
Is the identification and inspection status of each article maintained from the time of receipt of the material until delivery to the customer?			
Is there adequate control to prevent co-mingling of parts, lots and batches?			
Are limited shelf life materials controlled and identified?			
Are products maintained in proper storage conditions at all times, including during delivery?			

	PREMIER MAGNESIA - GILES CHEMICAL COMPANY FORM			
	Title: Vendor Assessment Survey	Page : 4 of 5	Revision : 00 Effective Date : 08/27/12	
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RECORDS	Yes	No	N/A
Do you maintain quality records on the quality and the manufacturing processes?			
Do you define which records are included and the time of retention?			
Are quality records current, complete and accurate?			
Does Management review quality records?			
Do quality/test records show failure and cause of failure?			

SERVICE ACTIVITIES	Yes	No	N/A
Do you provide detailed service reports explaining all work completed?			
Do you inspect completed work and validate the effectiveness?			
Do you provide service quotes prior to initiating work?			

TO BE COMPLETED BY GILES CHEMICAL:

COMMENTS (Please attach additional sheets if necessary)

Approved? Yes _____ No _____ If no, state why and list corrections necessary within this section. Indicate timing of corrections if known.

SURVEY INFORMATION REVIEWED BY:

_____ : Printed Name

_____ : Signature

_____ : Date



COMPANY FORM

Author: **Deborah Durbin**

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