

Chemical Vendor/Supplier Application Form (STEP 1)

Company Name _____
Address _____
City/Province/Postal Code _____
Phone _____ Fax _____
CEO Name _____ CEO Title _____
CEO E-mail _____ CEO Phone _____
Business Info E-mail _____
State Business License Number _____ State Tax ID Number _____

COMPANY INFORMATION

Organization Type: _____
Province of Incorporation _____
Domestic/Foreign Owned _____
Is your company owned by a parent company? Yes No
Parent Company Name _____
Parent Company Address _____
Parent Company Tax ID _____
Are you chemical business only? Yes No
Other business 1: _____
Other business 2: _____
Does your company accept credit cards? Yes No
Primary Standard Industrial Code _____
Additional SICs _____
Products/Services (short narrative): _____

Company's Web Site(s): _____
Did your company have a name change in the past 12 months? Yes No
Name _____
Company Emergency Contact: Name _____ Phone/E-mail _____
Quality Assurance Contact: Name _____ Phone/E-mail _____

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GENERAL INFORMATION

Area in Sq. Meter.: Manufacturing ____ Office ____ Total ____

Number of Personnel: Manufacturing ____ Quality Assurance ____ Engineering ____

Are clean room facilities used for manufacturing product? ____ Yes ____ No

What percentage of present work is: Government ____ Commercial ____ Other ____

Describe any special processes that you perform (synthesis, chromatography, distillation, high/low temperature reactions, reactions under high pressure, hydrogenation reactions, micronization, etc.)

Are you ISO-9000 certified? ____ Yes ____ No ISO Certificate Type _____

Registrar _____ Certificate Number _____

Expiration Date: ISO READY/Not Certified _____ Date of Certification _____

Registered or certified to any other Quality Management System or model?

Model _____

QUALITY MANAGEMENT SYSTEM

Do you maintain operation policies and procedures for your quality management system? ____ Yes
____ No

Is an internal audit program maintained that reviews compliance with all aspects of the quality program?
____ Yes ____ No

Does the organizational structure define quality responsibility and authority? ____ Yes ____ No

Does the organizational structure provide access to top management? ____ Yes ____ No

Is the health and status of your quality management system periodically reviewed with management?
____ Yes ____ No

Do you have a documented employee training program? ____ Yes ____ No

Is the quality organization responsible for acceptance of product and services? ____ Yes ____ No

Are records of inspections and tests maintained? ____ Yes ____ No

Are quality data used in reporting results and trends to management? ____ Yes ____ No

Are quality records available to support customer certifications? ____ Yes ____ No

RAW MATERIALS SOURCING CONTROL

Are procurement sources evaluated and monitored? ____ Yes ____ No

Are quality requirements and inspection procedures specified? ____ Yes ____ No

Is a documented system maintained for the evaluation of purchased materials? ____ Yes ____ No

Are incoming materials identified and segregated until acceptance? ____ Yes ____ No

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MATERIAL CONTROL

Do procedures exist for storage, release, and movement of material? Yes No
Are materials in storage identified and controlled? Yes No
Are in-process materials identified and controlled? Yes No
Are materials inspections identified and controlled? Yes No
Do storage areas and facilities provide control to protect material from degradation? Yes
 No
Are nonconforming items identified, segregated, and controlled? Yes No
If required, do you have the ability to provide tractability? Yes No
Does your company have a return policy? Yes No;
If Yes,
describe _____

REGULATORY COMPLIANCE

Are you a US FDA approved facility? Yes No.
Please
explain _____

If you are a US FDA approved facility for manufacturing pharmaceutical products, can you describe your most recent two inspectional results?
Please
describe _____

I certify that the information provided above is true.

Name _____ Signature _____ Business Seal _____

Date _____
