



# RFQ TEMPLATE

Please fill in as much information as possible. Complete information in all categories will help to ensure an accurate quote for your project.

Company: \_\_\_\_\_ Phone: \_\_\_\_\_

Address: \_\_\_\_\_

Key Personnel, phone numbers, and Email addresses:

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## Basic Project Information:

Type of dosage form needed: \_\_\_\_\_

Indication: \_\_\_\_\_ Strength: \_\_\_\_\_

Molecular Structure of the API: \_\_\_\_\_

Is this a new formulation or a modification of existing formulation? \_\_\_\_\_

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Please outline your timeline/deadlines: \_\_\_\_\_

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## Analytical Development:

Analytical methods available for API?      Yes      No

Transfer existing methods or develop new methods? \_\_\_\_\_

Any analytical method outside of the typical HPLC Assay, Dissolution, CU and Moisture? \_\_\_\_\_

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**Formulation Development:**

Has there been previous formulation development?      Yes    No

If yes, is tech transfer sufficient?    Yes    No

If no, please provide details on formulation development need: \_\_\_\_\_

\_\_\_\_\_

How many prototype strengths or formulations to be considered? \_\_\_\_\_

Any stability study of prototypes? \_\_\_\_\_

**Manufacturing Requirement (CTM):**

How many strengths and units? (please list): \_\_\_\_\_

\_\_\_\_\_

Matching placebos?      Yes    No      If yes, # of units: \_\_\_\_\_

Packaging and labeling requirements: \_\_\_\_\_

\_\_\_\_\_

**Stability Studies:**

Conduct per ICH guidelines?      Yes    No

If No then specify stability study details for CTM: \_\_\_\_\_

\_\_\_\_\_

Please indicate:      Bulk product stability      Packaged product

**Special Handling Requirements:**

Indicate if API is:      Potent      Cytotoxic      Controlled Substance

Indicate if API is light sensitive, hygroscopic, etc.: \_\_\_\_\_

\_\_\_\_\_

Provide MSDS if available.