

PROTEIN EXPRESSION SERVICE QUOTATION REQUEST FORM

Name:

Institution:

Shipping Address:

Phone:

E-mail:

PROTEIN INFORMATION

Protein Name:

Molecular Weight:

Protein Accesion Number or Sequence:

If mutants are included in your protein sequence, please enter the protein sequence

Is your protein a monomer?:

Yes

No

Do you want Innoprot to synthesize and clone the gene for you?

Yes

No

If YES, please paste the desired DNA sequence including epitope tags here:

Do you want Innoprot to subclone your gene into our expression construct?

Yes

No

If supplying your own expression construct, please give a brief description of your vector:

Please indicate the preferred expression host system:

Bacteria

Baculovirus / Insect cells

Mammalian

PROTEIN EXPRESSION INFORMATION

Quantity of Protein Required (in mg):

Special Requirements

PROTEIN PURIFICATION INFORMATION

Is your protein: Membrane-bound Secreted A protease Other enzyme Toxic to *E.coli*

Other features that may cause difficulty in purification (specify):

Innoprot will select 1 or 2 tags for your (e.g. 6xHis, GST, or MBP) to facilitate purification:

OK No, I want to select on my own (specify):

Do you want Innoprot to remove the tag: Yes No

If YES, Which protease should be use to remove the tag:

Doesn't matter Thrombin Enterokinase rTev Other (specify):

Is your protein insoluble? Yes No I don't know

If the expressed protein is insoluble, do you want Innoprot to attempt the refolding?: Yes No

Are there known expression, purification or refolding protocols for this protein?: Yes No

Special Requirements (buffers, handling, etc):

QUOTATION REQUEST CHECKLIST

Please make sure that the following materials and data are included with the quotation request:

- 1. Gene accession number or sequence
- 2. If requesting gene synthesis, *desired gene sequence including epitope tags*

REMEMBER: THESE STARTING MATERIALS WILL BE REQUIRED WHEN YOU SUBMIT YOUR ORDER TO INNOPROT

- 1. Starting construct**, if requesting subcloning (~ 5ug of plasmid DNA)
- 2. Expression construct**, if supplying your own (~ 5ug of plasmid DNA)
- 3. Known expression, purification and refolding protocols, if available**
- 4. If requesting subcloning**, detailed map of the starting construct, including both your gene and vector
- 5. If supplying your own expression vector**, detailed map of the expression vector