

## **Quotation Request Form**

## **MOLECULAR BIOLOGY CORE**

Dept of Surgery/Cardiac division MBClab@osumc.edu

1. Contact Information				
Name*:				
Title (Position):				
Institution/Company:				
Address:				
E-mail* / Telephone:				

\* Please describe your mutagenesis (Section 2-1) or subcloning (Section 2-2) project.

2-1. Request of mutagenesis					
Vect (\	Target site  Mutagensis  Wt construct  Wector + target gene)  Mutant construct  (vector + mutant gene)				
Mutagenesis	<ul> <li>□ Substitution</li> <li>□ Deletion</li> <li>□ Insertion</li> <li>□ Chimeragenesis</li> <li>□ Multiple-site mutagenesis</li> <li>□ Random mutagenesis</li> </ul>				
Quantity of mutants	I want to generate total ( ) mutant constructs.				
Sequencing analysis	<ul> <li>□ Basic sequencing to confirm target mutation (free up to 1.0 kb sequencing)</li> <li>□ Full-length sequencing of gene insert</li> <li>: Free 1.0 kb, \$25/extra 800 bp sequencing for analysis of at least 3 colonies</li> </ul>				
Information of wt construct	☐ Construct name: (Size: kb) ☐ Bacterial selection marker: Ampicilin, kanamycin, or other ☐ Gene name or GenBank Accession No:				

Vector for mutant gene	☐ Original con	struct [	☐ New customer's vector:		
Target mutations	Examples of description				
	1. Substitution:	AAATTT <u>GCG</u> AAATTT	T → AAATTTCgCAAATTT		
	2. Deletion:	AAATTT <u>GCG</u> AAATTT	T → AAATTTAAATTT		
	3. Insertion:	AAATTT <u>G</u> CGAAATTT	TT → AAATTT <u>Gaaa</u> CGAAATTT		
	Please describe your mutagenesis plan or target mutations.				
	1.				
	2.				
	3.				
	4.				
	5.				

→ Or please send us *wt* construct and mutagenesis information by e-mail. We will provide a quote soon.

2-2. Request of subcloning							
	Subcloning Subclone Subclone	n sites ning vector					
Subcloning method	□ PCR-subcloning □ Gene synthesis and cloning in a	vector					
Quantity of subcloning	I want to generate total ( ) subclone constructs.						
Sequencing analysis	☐ Basic service: Free sequencing of the whole gene insert up to 3.0 kb☐ Whole sequence confirmation for gene insert larger than 3.0 kb☐ Free up to 3.0 kb, \$25/extra 800 bp sequencing for analysis of at least 3 colonies						
→ For PCR subcloning	Information of template construct						
	□ Construct name:	(Size:	kb)				
	☐ Gene name or GenBank Accession No:	(Size:	kb)				

Form ver. 2019, 07-23 ☐ Bacterial selection marker: Ampicilin, kanamycin, or other **Subcloning destination vector** (Size: ☐ Vector name: kb) ☐ If commercially available, please indicate a commercial name and provider: Commercial name: Provider: ☐ Bacterial selection marker: Ampicilin, kanamycin, or other \_\_\_ → For gene synthesis The size of the gene inserts to be synthesized 3. \_\_\_\_\_bp 4. \_\_\_\_\_ bp 5. \_\_\_\_\_ bp Gene property, if applicable: ☐ GC rich ☐ Tandem repeat ☐ Inverted repeat (shRNA, LoxP, ...) Codon optimization, if necessary: □ No □ Yes (□ E. coli □ Yeast □ Mammalian □ other\_\_\_\_\_) Subcloning destination vector

□ pUC19 or pBluescript (\$90 of subcloning charge per construct)
 □ Other customer's vector (\$180 of subcloning charge per construct)

☐ Bacterial selection marker: Ampicilin, kanamycin, or other

☐ If commercially available, please indicate a commercial name and provider:

Provider:

(Size:

kb)

→ Or please send us your information of gene of interest and subcloning plan by e-mail. We will provide a quote soon.

Commercial name:

☐ Vector name:

For customer's vector, please describe

## **Comments:**