# All about Bioneer's

# **Custom Services**

Molecular Biology & Protein Expression



# **Contents**

1.	. Oligonucleotide Synthesis	
	DNA/RNA Synthesis  CRISPR Cas9/Cpf1-gRNA  siRNA/miRNA Synthesis  In vivo siRNA delivery using SAMiRNA™  MALDI-TOF MS Analysis	9 12
2.	. Molecular Biology	
	DNA/RNA Amplification  Gene Expression Analysis  Gene Synthesis  Sanger Sequencing  mRNA Profiling	26
3.	. Protein Synthesis	
	Cell-free Protein Synthesis	45

With the long years of expertise, Bioneer have developed unique, well-established patented technologies and production systems.

We provide customized services through 1:1 consultation with our experts who have accumulated experiences in the field of biotechnology for a long time.

All our services are conducted under ISO 9001 quality management system for accurate, reproducible results.

#### Oligonucleotide Synthesis Molecular Gene **RNAi Expression Synthesis Analysis** DNA/RNA **Target PCR Synthesis** Gene DNA/RNA siRNA/miRNA DNA/RNA Gene Expression **Synthesis Amplification Synthesis Analysis** Customized DNA/RNA Standard Oligo Predesigned siRNA qPCR Array Service Amplification Kit HT-Oligo™ Customized qPCR Panel Kit Premade siRNA Control siRNA Primer & Probe Design Service Modification Oligo **Dual-labeled Probes** Custom siRNA Single Gene qPCR Primer Set Extendamers™ miRNA Mimic / Inhibitor Large Scale Oligo Control miRNA Custom RNA Oligo *In vivo* siRNA delivery using SAMiRNA™ CRISPR Cas9/Cpf1-gRNA MALDI-TOF MS Analysis

# **Service Overview**

Biology **Protein Synthesis** 

Gene Cloning

Sequencing

Protein Synthesis

#### Gene Synthesis

Gene Synthesis Rapid Gene Synthesis AccuGeneBlock Gene Cloning Mutagenesis Codon Library Mutagenesis

#### Sanger Sequencing

Standard Sequencing Full-length Sequencing Difficult Sequencing Additional Service

#### mRNA Profiling

#### Cell-free Protein Synthesis

Standard Protein Synthesis One-day Protein Synthesis Standard Gene to Protein Cloning-free Gene to Protein

# **DNA/RNA Synthesis**

# Oligonucleotide Synthesis

► DNA/RNA Synthesis ·····	· 4 - 8 Page
CRISPR Cas9/Cpf1-gRNA ·····	· 9 - 11 Page
siRNA/miRNA Synthesis · · · · · · · · · · · · · · · · · ·	12 - 18 Page
<i>In vivo</i> siRNA delivery using SAMiRNA™ ······	19 - 21 Page
MALDI-TOF MS Analysis ·····	22 - 23 Page



# **DNA/RNA Synthesis**

#### **List of Services**

Standard Oligo **Dual-labeled Probes** Custom RNA Oligo

HT-Oligo™ Extendamers™ Modification Oligo Large Scale Oligo

#### **Key Features**

- © Provide exact concentration of oligos by utilizing our patented Accuoligo® technology
- O Supply DNase, RNase, DNA-free oligos by synthesizing in a clean room
- Undergo strict quality control using a total of 7 MALDI-TOF spectrometers to perform full QC inspection.
- Utilizes the world's first automated production line by processing all protocols from order to synthesis, QC, and packaging automatically using "Computer Aided Manufacturing (CAM) System", allowing consistent quality of oligos

#### What is AccuOligo® Technology?

AccuOligo® is our patented technology which prevents dried oligos from falling or slipping from the tubes during their transit. While the conventional synthesis service had a danger of dried oligos falling from the top of the tube due to impacts and shakes, leading to loss of yield after opening it, AccuOligo® technology not only allows to deliver the oligos with precise concentration and yield by allowing them to stick at the bottom even at intense vibrations, but also to prevent cross-contamination between the wells when packaged in plates. Furthermore, all the components used to prevent the falling of oligos do not affect experimental results such PCR, sequencing, or enzyme cutting.

#### **Purification System**

- Bio-RP: Bio-RP is a proprietary, high-quality purification method developed by Bioneer which allows to supply nuclease-free oligos by synthesizing in a clean room and eliminating truncated failures including n-1 mer.
- HPLC: HPLC allows to extract oligos being about 30 mer long with high purity over 90%. This method is recommended for fluorescent dye modification or protocols requiring high-quality oligos such as cloning, site directed mutagenesis, and quantitative gene detection.
- PAGE: PAGE method is for isolating up to 200 mer of long oligos having high purity over 95% by using electrophoresis on. polyacrylamide gel.

#### Contact Us

E-mail: oligo-support@bioneer.co.kr, Tel: +82-42-930-8777

## **Standard Oligo**

#### O Description

Standard Oligonucleotide synthesis service is for ordering oligos of 130 mer or less. Various synthetic scales and purification methods can be selected depending on the applications. Selectable purification services include Bio-RP, HPLC, PAGE, etc.

#### Ordering Information

Synthesis Scale (nmol)	Base Limitation (mer)	Guaranteed Yield(O.D.) Based on 20 bases			Service Period (Working Day)		
(HIIIOI)	(iiiei)	Bio-RP	PAGE	HPLC	Bio-RP	PAGE	HPLC
25	15 - 60	2	1	1.5	2	3	3
50	10 - 75	4	2	2.5	2	3	3
200	5 - 110	8	6	7	2	3	3
1,000	5 - 130	30	18	25	2	3	3
10,000	5 - 50	300	150	200	2	3	3
15,000	5 - 50	Inquire		3	4	4	

<sup>\*</sup> The synthesis scale (nmol) represents the start volume of synthesis. Therefore, all the final yield after synthesis and purification depends on the oligo length, synthesis efficiency, and nucleotide sequences.

### HT-Oligo™

#### O Description

HT-Oligo™ (High Throughput Oligonucleotide) synthesis service can synthesize more than 96 different types of oligos simultaneously at a reasonable price. Furthermore, for this synthesis, we provide concentration normalization service for free.

Synthesis Scale (nmol)	Base Limitation	Base Limitation Guaranteed Yi (mer) Based on 20					
(IIIIIOI)	(IIICI)	Bio-RP	PAGE	HPLC	Bio-RP	(Working Day) D-RP PAGE HPLC 2 3 3 2 3 3 2 3 3 2 3 3 Inquire	
25	15 - 60	2	1	1.5	2	3	3
50	10 - 75	4	2	2.5	2	3	3
200	5 - 110	8	6	7	2	3	3
1,000	5 - 130	30	18	25	2	3	3
10,000	5 - 50	300	150	200	Inquire		
15,000	5 - 50	Inquire			Inquire		

## **Modification Oligo**

#### O Description

Modification Oligo service provides various 3', 5', or internal modification oligonucleotide synthesis service ranging from biotin-labeled primers to fluorescence-labeled primers to be used for real-time PCR analysis.

#### Ordering Information

Synthesis Scale (nmol)	Base Limitation (mer)	Guaranteed Yield(O.D.)	Service Period (Working Day)
50	10 - 75		
200	5 - 110	Varies depending on the types of	
1,000	5 - 50		4
10,000	5 - 50	modifications done	
15,000	5 - 50		

<sup>\*</sup> Refer to our Bioneer website (www.bioneer.com) for more information on the type of modification that can be done.

### **Dual-labeled Probes**

#### Operation

Duel-labeled Probe service synthesizes oligos with a fluorescent dye (reporter) positioned at the 5' end while a quencher is on the 3' end. Those are mostly used for Real-time PCR analysis, but can also be used for multiplex analysis, done through searching several genes at once, by using various fluorescent materials. Our Dual-labeled Probes guarantees high quality products by performing MALDI-TOF Mass spectrometer QC method and fluorescence test using fluoroskan.

#### Ordering Information

Synthesis Scale (nmol)	Base Limitation (mer)	Guaranteed Yield(O.D.)	Service Period (Working Day)
50	10 - 75		
200	5 - 110	Varies depending on the types of	
1,000	5 - 50		4-5
10,000	5 - 50	modifications done	
15,000	5 - 50		

### Extendamers<sup>™</sup>

#### O Description

Extendamers™ (Long Oligonucleotides) synthesis service is for ordering long oligos with 130~200 bp, which is mainly applied in cloning, siRNA, and gene construction.

Base Limitation	Guaranteed	l Yield(O.D.)	Service Period (Working Day)		
(mer)	Bio-RP	PAGE	Bio-RP	PAGE	
130 - 200	3-4	0.25 - 0.3	5-6	7 - 8	

### Large Scale Oligo

#### O Description

Large Scale Oligonucleotide (Bulk Oligo) service is for ordering oligos with 15,000 nmol scales. By using our proprietary instrument developed solely by us, Bulk Oligo Synthesizer, we can provide comprehensive range of oligo scales ranging from mg to kg units, along with advanced oligo synthesis and purification technologies for antisense researches and large-scale protocols such as decoy and RNAi oligo.

#### Ordering Information

Guaranteed Yield	Base Limitation	Service Period (Working Day)		
(mg)	(mer)	Bio-RP	HPLC	
50				
100				
250	5 - 50	10	10	
500				
1,000				

### **Custom RNA Oligo**

#### O Description

Custom RNA Oligo service provide synthesis of RNA sequences in single strand or double strand forms as our customers wish. Purification methods in this service can be selected between Bio-RP and HPLC while various modification of up to 32 different 3' overhang selection can be made. Moreover, customers can select annealing, drying or ready-to-use options to determine how their RNA is finalized and provided.

Guaranteed Yield	Base Limitation	Service Period (Working Day)		
(mg)	(mer)	Bio-RP	HPLC	
10				
20	5 - 85	2 2	2 /	
50		2 - 3	3 - 4	
100				

<sup>\*</sup> Both Sense and anti-sense RNAs in single-stranded form are provided and dried at the same concentration.

<sup>\*</sup> Annealing buffer is provided by default.

<sup>\*</sup> Additional costs will be charged for ordering siRNA in an annealing form.

<sup>\*</sup> All synthetic RNAs are shipped with MALDI-TOF QC Data.

# CRISPR Cas9/Cpf1-gRNA

# Oligonucleotide Synthesis

DNA/RNA Synthesis · · · · · · · · · · · · · · · · · ·	· · 4 - 8 Page
► CRISPR Cas9/Cpf1-gRNA ······	· 9 - 11 Page
siRNA/miRNA Synthesis ·····	12 - 18 Page
<i>In vivo</i> siRNA delivery using SAMiRNA™ ·····	19 - 21 Page
MALDI-TOF MS Analysis ·····	22 - 23 Page



# CRISPR Cas9/Cpf1-gRNA

#### **List of Services/Product**

CRISPR Cas9-gRNA CRISPR Cpf1-gRNA AccuCRISPR™-Cas9 tracrRNA

#### **Key Features**

- Ready-to-use guide RNAs to be immediately used for experimental purposes with nucleases (without the need of cloning process)
- © Guaranteed high-quality products through synthesis in a clean room
- Advanced RNA oligo synthesis technology
- © Comprehensive gRNA design compatible with various CRISPR nuclease

#### What is CRISPR Cas9/Cpf1-gRNA?

The CRISPR Cas9/Cpf-gRNA system is genome editing technology that can modify genes using nuclease (Cas9 or Cpf1) and gRNA (guide RNA) complex. Our CRISPR Cas9/Cpf1-gRNA service can synthesize gRNA with the sequence that the customers desire. The ordered gRNA can be used in the form of ribonucleoprotein (RNP) with Nuclease (Cas9 or Cpf1). Since the RNP form can be rapidly degraded by endogenous protease or RNase, It has the advantage of lower off-target. With our rich experiences on oligo synthesis, we provide CRISPR gRNA service with the best quality that have passed our QC tests by synthesizing in our advanced clean-room facility and purifying with MALDI-TOF mass spectrometry.

		CRISPR-Ca	as9 System	CRISPR-Cpf1 system
	Nuclease	Ca	as9	Cpf1
Construction	Construction Guide RNA	crRNA	or Chimeric RNA (sgRNA)	GrDNA
		tracrRNA		crRNA
PAM sequence		NGG		TTTN
Cutting method		Blunt	Blunt ends 5'overhang	

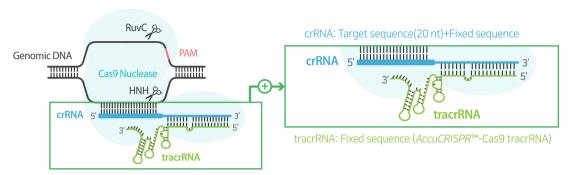
#### **Contact Us**

E-mail: CRISPR@bioneer.co.kr, Tel: +82-42-930-8752

### **CRISPR Cas9-gRNA**

#### O Description

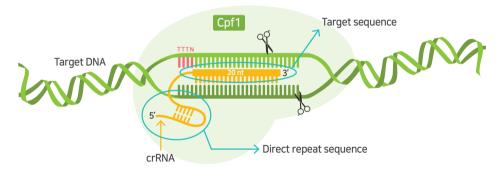
The gRNA in the CRISPR-Cas9 system consists of crRNA and tracrRNA containing Cas9 nuclease binding sequences, We sell crRNA and tracrRNA together so that our customer can easily construct a crRNA-tracrRNA complex in a ribonucleoprotein form to be directly used in the CRISPR system.



### **CRISPR Cpf1-gRNA**

#### Operation

Bioneer provides gRNA that can be used on two types of Cpf1: AsCpf1 and LbCpf1. By inputting 20 nt target-specific sequences in our online order page, the fixed direct repeated sequences are automatically added in the order. After receiving the Cpf1 gRNA(crRNA), it can be used in the form of RNP (Ribonucleoprotein).



### AccuCRISPR™-Cas9 tracrRNA

#### O Description

AccuCRISPR™-Cas9 tracrRNA works in the form of duplex with crRNA. As it has a fixed sequence, simply select its scale and quantity during your orders to be used with crRNA for your CRISPR system.

Cat. No.	Product Description
S-5120-5	AccuCRISPR™-Cas9 tracrRNA (5 nmole)
S-5120-10	AccuCRISPR™-Cas9 tracrRNA (10 nmole)

# siRNA/miRNA Synthesis

# Oligonucleotide Synthesis

DNA/RNA Synthesis · · · · · · · · · · · · · · · · · ·	· · 4 - 8 Page
CRISPR Cas9/Cpf1-gRNA ·····	· 9 - 11 Page
► siRNA/miRNA Synthesis ·····	12 - 18 Page
<i>In vivo</i> siRNA delivery using SAMiRNA™ ·····	19 - 21 Page
MALDI-TOF MS Analysis ·····	22 - 23 Page



# siRNA / miRNA Synthesis

#### **List of Services/Products**

#### Genome-wide Predesigned siRNA & Primer

*AccuTarget*™ Genome-wide Predesigned siRNA Library AccuTarget™ Real-Time PCR Primer Library

#### Premade siRNA Set & Primer

*AccuTarget*™ Human Premade siRNA Sets *AccuTarget*™ Real-Time PCR Primers for Premade siRNA Sets

#### Control siRNA

*AccuTarget*™ Control siRNAs (Positive / Negative)

#### **Custom siRNA**

*AccuTarget*™ Custom Designed siRNA Synthesis

#### miRNA Mimics & Inhibitors

AccuTarget™ Human miRNA Mimic / Inhibitor

#### **Control miRNA**

AccuTarget™ Control miRNAs (Positive / Negative)

#### **Key Features**

- © Provide 164,643 presigned siRNA for 54,881 target genes of Human, mouse, rat and 1,786 human mature miRNA present in the miRBase Sequence Database (version 22)
- © Guarantee more than 80% with the siRNA designed with validated Turbo si-Designer
- O Undergo strict quality control by synthesizing in a clean room

#### What is Turbo si-Designer?

Turbo si-Designer is a siRNA design program developed by collaboration between BIONEER corporation and National Genome Information Center (NGIC). It uses algorithms to find the best siRNA target sites to construct predesigned siRNA with more than 80% guaranteed knockdown efficiency.

#### **Contact Us**

E-mail:siRNA-support@bioneer.co.kr, Tel: +82-42-930-8777

## AccuTarget™ Genome-wide Predesigned siRNA Library

#### O Description

AccuTarget™ Genome-wide Predesigned siRNA Library uses Turbo si-Designer to design siRNA for 18,462 human genes, 19,859 mouse genes, and 16,560 rat genes, allowing us to immediately synthesize and ship the desired products to our customers. Furthermore, we have minimized the off-target effect to maximize the siRNA knockdown efficiencies and guaranteed to inhibit more than 80% of the target expression levels for three out of two siRNA for the same genes.

#### Ordering Information

Product Description	Purification	Guaranteed Yield (nmol)	Service Period (Working Day)	
		10		
Accustoractin Conomo mido Prodocianos ciDNA Library	Bio-RP/HPLC	20	2-3	
AccuTarget™ Genome-wide Predesigned siRNA Library		50	2-3	
		100		

### AccuTarget™ Real-Time PCR Primer Library

#### O Description

AccuTarget™ Real-Time PCR Primer Library is a Real-time PCR primer for 11,154 human predesigned siRNA. All primers of this product have passed the amplification efficiency tests and QC tests using MALDI-TOF mass spectrometer.

Cat. No.	Product Description
PHS-P01	AccuTarget™ Real-Time PCR Primer for Individual Gene, 100 reactions
PHS-P02	AccuTarget™ Real-Time PCR Primer for Individual Gene, 200 reactions

# *AccuTarget*™ Human Premade siRNA Sets

#### O Description

AccuTarget™ Human Premade siRNA set consists of 54,144 types of siRNAs that can be immediately used in transfection. This siRNA library set is constructed with 25 biological pathways (or gene family) specialized in research purposes such as specific cellular processes, cancers, and diseases. Furthermore, it can be either provided separately with 10, 20, 50, or 100 nmol or a whole set with 0.1, 0.25, 0.5, or 1 nmol each.

Product Description	Purification	Guaranteed Yield
		1 siRNA (0.1, 0.25, 0.5, 2 nmol)
AccuTarget™ Human Premade siRNA Set		2 siRNAs (0.1, 0.25, 0.5, 2 nmol)
	Bio-RP/HPLC	3 siRNAs (0.1, 0.25, 0.5, 2 nmol)
AccuTarget ™ Human Premade siRNA Subset	DIO-RE/TIELC	0.1, 0.25, 0.5, 2 nmol
AccuTarget ™ Human Druggable siRNA Library Set		0.1, 0.25, 0.5, 2 nmol (minimum order 10 siRNAs)

Cono Family	No of No of		Cana Family	Humar	Human Genes	
Gene Family Functional Class				No. of Genes	No. of siRNA	
Antioxidant	38	114	Lyase	123	369	
Apoptosis	290	870	Motor	122	366	
Cancer	1,158	3,471	NF-kB pathway	37	111	
Caspase	37	111	Nucleic acid binding	2,573	7,719	
Cell cycle	112	336	Oxidoreductase	551	1,653	
Cyclase	21	63	Peptidase	491	1,473	
Cytochrome P450	52	156	Phosphatase	188	564	
Deaminase	22	66	Receptor	1,516	4,548	
GPCR signaling pathway	727	2,181	Transferase	1,428	4,284	
Helicase	114	342	Transporter	1,021	3,063	
Isomerase	104	312	Tubulin	20	60	
Kinase	699	2,097	Ubiquitin	77	231	
Ligase	272	816				

# AccuTarget™ Real-Time PCR Primers for Premade siRNA Sets

#### Operation

AccuTarget™ Real-Time PCR Primers for Premade siRNA Sets are provided in 11,154 primer sets according to the signaling classification by gene function.

#### Ordering Information

Product Description	The Number of Genes	Rxn/Gene
AccuTarget™ Human Real-Time PCR Primer Set	11,154 genes	50 reactions

<sup>\*</sup> For more information, visit our Bioneer website (www.bioneer.com).

## AccuTarget™ Control siRNAs (Positive / Negative)

#### O Description

 $AccuTarget^{\text{TM}}$  Positive Control siRNA has a high knockdown efficiency being more than 90% for its target genes. Moreover,  $AccuTarget^{\text{TM}}$  Negative Control siRNA is a non-targeting siRNA with low homology for all genes of humans, mice, and rats to be used as a negative control for knockdown experiments.

Cat. No. Product Description		Purification	Guaranteed Yield		
Positive Control siRNA					
SP-1001/1002/1003	AccuTarget™ GAPDH Positive Control siRNA	Bio-RP			
SP-1011/1012/1013	AccuTarget™ GAPDH Positive Control siRNA	HPLC			
SP-2001/2002/2003	AccuTarget™ GFP Positive Control siRNA	Bio-RP			
SP-2011/2012/2013	AccuTarget™ GFP Positive Control siRNA	HPLC			
SP-3001/3002/3003	AccuTarget™ Luciferase Positive Control siRNA	Bio-RP	F/10/20 pmal		
SP-3011/3012/3013	AccuTarget™ Luciferase Positive Control siRNA	HPLC	5/10/20 nmol		
SP-4001/4002/4003	AccuTarget™ Mouse Lamin A/C Positive Control siRNA	Bio-RP			
SP-4011/4012/4013	AccuTarget™ Mouse Lamin A/C Positive Control siRNA	HPLC			
SP-5001/5002/5003	AccuTarget™ Mouse cyclophilin B c Positive Control siRNA	Bio-RP			
SP-5011/5012/5013	AccuTarget™ Mouse cyclophilin B c Positive Control siRNA	HPLC			
Negative Contro	ol siRNA				
SN-1001/1002/1003	AccuTarget™ Negative Control siRNA	Bio-RP			
SN-1011/1012/1013	AccuTarget™ Negative Control siRNA	HPLC	5/10/20 nmol		
SN-1021/1022/1023	AccuTarget™ Fluorescein-labeled Negative Control siRNA	HPLC			
Control siRNA Sets					
SS-1001/1002/1003	AccuTarget™ GAPDH / GFP / Luciferase Control siRNA Set	Bio-RP	5 nmol PC + 2 nmol NC		
SS-1011/1012/1013	AccuTarget™ GAPDH / GFP / Luciferase Control siRNA Set	HPLC	STITIOLEC + ZTITIOLING		

### AccuTarget™ Custom Designed siRNA Synthesis

#### O Description

AccuTarget™ Custom Designed siRNA Synthesis service synthesizes siRNA with the requested sequences by customers, Not only we use Turbo si-designer for the siRNA design, but also can apply various modifications to your RNA sequences as they want.

#### Ordering Information

Cat. No.	Product Description	Purification	Guaranteed Yield	
S-1017-1/2/3/4	AccuTarget™ Custom Designed siRNA Synthesis	Bio-RP	10/20/F0/100 pmol	
S-1018-1/2/3/4	AccuTarget™ Custom Designed siRNA Synthesis	HPLC	10/20/50/100 nmol	

<sup>\*</sup> For more information on the available modification services, visit our Bioneer website (www.bioneer.com).

### AccuTarget™ Human miRNA Mimic & Inhibitor

#### O Description

AccuTarget™ miRNA mimic is a double-stranded RNA oligonucleotide that is chemically synthesized and manufactured from 1,786 human mature miRNA present in the miRbase Sequence Database (version 22). AccuTarget™ miRNA Inhibitor is a single-stranded synthetic RNA designed to target each of human miRNA and inhibits its functions, Ready-to-transfect miRNA shows the same activity as endogenous miRNA in the cells after its transfection, while miRNA inhibitor suppresses target miRNA activity to be used in the loss-of-function studies of miRNA.

Cat. No.	Product Description	Purification	Guaranteed Yield
Custom miRNA			
SMM-001/002/003	<i>AccuTarget</i> ™ miRNA Mimic	Bio-RP	F/10/20 pmol
SMI-001/002/003	AccuTarget™ miRNA Inhibitor	Bio-RP	5/10/20 nmol
Library miRNAs			
SML-1001/1002/1003/1004	<i>AccuTarget</i> ™ miRNA Mimic	Bio-RP	0.25/0.5/1/2.550
SML-2001/2002/2003/2004	AccuTarget™ miRNA Inhibitor	Bio-RP	0.25/0.5/1/2 nmol

# **AccuTarget** ™ Control miRNAs

#### O Description

AccuTarget™ miRNA Housekeeping Positive Control targets 3'UTR(untranslated region) of a housekeeping gene GAPDH with over 90% of knockdown efficiency. Also, miRNA Negative Control has a minimum sequence identity for all miRNA in humans, mice, and rats.

Cat. No.	Product Description	Purification	Guaranteed Yield	
Positive Control siF	RNA			
SMC-1001/1002/1003	AccuTarget™ miRNA Housekeeping Positive Control (GAPDH)	Bio-RP	5/10/20 nmol	
Negative Control m	niRNA			
SMC-2001/2002/2003	AccuTarget™ miRNA Mimic Negative Control #1	Bio-RP		
SMC-3001/3002/3003	AccuTarget™ miRNA Mimic Negative Control #2	Bio-RP		
SMC-4001/4002/4003	AccuTarget™ Fluorescein-labeled miRNA Mimic Negative Control #1	HPLC		
SMC-5001/5002/5003	AccuTarget™ Fluorescein-labeled miRNA Mimic Negative Control #2	HPLC	E/10/20 pmol	
SMC-2101/2102/2103	AccuTarget™ miRNA Inhibitor Negative Control #1	Bio-RP	5/10/20 nmol	
SMC-3101/3102/3103	AccuTarget™ miRNA Inhibitor Negative Control #2	Bio-RP		
SMC-4101/4102/4103	AccuTarget™ Fluorescein-labeled miRNA Inhibitor Negative Control #1	HPLC		
SMC-5101/5102/5103	AccuTarget™ Fluorescein-labeled miRNA Inhibitor Negative Control #2	HPLC		

# *In vivo* siRNA delivery using SAMiRNA™

# Oligonucleotide Synthesis

	DNA/RNA Synthesis · · · · · · · · · · · · · · · · · ·	• 4 - 8 Page
	CRISPR Cas9/Cpf1-gRNA ······	9 - 11 Page
	siRNA/miRNA Synthesis ·····	12 - 18 Page
>	· <i>In vivo</i> siRNA delivery using SAMiRNA™ ······	19 - 21 Page
	MALDI-TOF MS Analysis ·····	22 - 23 Page



# In vivo siRNA delivery using SAMiRNA™

#### **List of Service**

SAMiRNA™ Custom Service

#### **Key Features**

- © The world's only technology delivering the native siRNA/miRNA through the single molecule nanoparticle transporter substance
- No side effects: a safe drug delivery system with no innate immune and inflammatory responses
- O Low toxicity: Only ~ 100 mpk of NOAEL in mouse
- OPERFORM Preferential delivery to tumors and inflammatory tissues: suitable for solid tumors, fibrotic diseases
- Advanced mechanism: single step automated synthesis process without the need of additional formulation or complicated quality control tests.

#### **Contact Us**

E-mail:siRNA@bioneer.co.kr, Tel: +82-42-930-8777

# SAMiRNA™ Custom Service (Self-Assembled-Micelle-inhibitory-RNA)

#### O Description

SAMiRNA $^{\text{TM}}$  is a RNAi drug substance in the form of nanoparticle that uses an innovative new technology developed by our company which has overcome the limitation of traditional siRNA-based therapeutic technology to safely deliver the siRNA to the *in vivo* target organ and cells efficiently. As the world's only siRNA prodrug, SAMiRNA $^{\text{TM}}$  is a revolutionary RNAi substance composed of a single molecule structure of a nanoparticle which maintains its stability even when it is injected to the blood-stream. It can be selectively delivered to the cancerous and diseased cells. As it enters the target cell, it is converted to siRNA which in turn degrade the cancer-inducing RNA.

Most of the siRNA therapeutic technologies currently undergoing clinical trial require liposome-based formulation or encapsulation step of the synthesized siRNA as the delivery vehicle to transport into the target tissue. However, at this time, not only it is difficult to control the loading efficiency of synthesized siRNA into the liposome, but as siRNA normally has low loading efficiency, the drug has to undergo complicated QC process for isolation and purification of the packaged siRNA for mass production. Furthermore, some reports state that some lipid components consisting the liposome have toxicity including innate immune response.

Unlike this liposome-based formulation, SAMiRNA $^{\text{m}}$  has overcome the liposome siRNA drugs by being prepared as a single entity in a fully automated synthesis system by self-assembling in an aqueous solution to be made into a stable nanoparticle without formulation. The therapeutic effect of this product for cancer or fibrosis has been already tested by mouse model experiments. Moreover, SAMiRNA $^{\text{m}}$  has less toxicity than the conventional siRNA carriers, allowing our product to be more advantageous than the others as the new generation drug candidate.

SAMIRNA	Order Scale of SAMiRNA (nmol)	Number of Mouse (intravenous injections)
	100	5 mg/ kg $\rightarrow$ 5 mouse / 1 mg/ kg $\rightarrow$ 25 mouse
Custom Designed SAMiRNA	500	5 mg/ kg → 25 mouse / 1 mg/ kg → 125 mouse
	1000	5 mg/ kg → 50 mouse / 1 mg/ kg → 250 mouse
	100	5 mg/ kg → 5 mouse / 1 mg/ kg → 25 mouse
Control SAMiRNA	500	5 mg/ kg → 25 mouse / 1 mg/ kg → 125 mouse
	1000	5 mg/ kg → 50 mouse / 1 mg/ kg → 250 mouse
El	100	5 mg/ kg → 5 mouse / 1 mg/ kg → 25 mouse
Fluorescence conjugated SAMiRNA (FITC, Cyanine 5.5 etc)	500	5 mg/ kg → 25 mouse / 1 mg/ kg → 125 mouse
(FITC, Cyallille 5.5 etc)	1000	5 mg/ kg → 50 mouse / 1 mg/ kg → 250 mouse

# MALDI-TOF MS Analysis

# Oligonucleotide Synthesis

DNA/RNA Synthesis · · · · · · · · · · · · · · · · · ·	· · 4 - 8 Page
CRISPR Cas9/Cpf1-gRNA ······	· 9 - 11 Page
siRNA/miRNA Synthesis ······	12 - 18 Page
<i>In vivo</i> siRNA delivery using SAMiRNA™ ·····	19 - 21 Page
MALDI-TOF MS Analysis ·····	 22 - 23 Page



# **MALDI-TOF MS Analysis**

#### **List of Service**

MALDI-TOF MS Analysis Service

#### **Key Features**

- © Trustworthy methology: Provide reliable data through periodically qualified MALDI-TOF Mass spectrometer
- O Accurate data: Purify samples by desalting pre-treatment to remove salt, impurities, etc. in the samples
- Rich experience: Undergo accurate and fast analysis results based on 20 years of experience in oligo nucleotide synthesis
  and quality control

#### **Contact Us**

E-mail:maldims@bioneer.co.kr, Tel: +82-42-930-8554

### **MALDI-TOF MS Analysis Service**

#### Operation

MALDI-TOF MS analysis involves ionization of biomolecules and large organic molecules with lasers to measure their molecular weight using matrix. This allows to quickly analyze polymeric materials without sample degradation. Our analysis service uses MALDI-TOF Mass spectrometers to provide reliable molecular weight analysis results only within two days with our rich experiences and expertise after receiving the single strand oligonucleotide from our customers.

#### Workflow



- \* The Service Order Form can be downloaded from our Bioneer website (www.bioneer.com).
- \* Please print out the completed form and sent it together with the sample to be analyzed to the following address: 8-11, Munpyeongseo-ro, Daedeok-gu, Daejeon 34302, Republic of Korea, Bioneer Quality Management Team.

Cat. No.	S-2400
Sample target	Single strand oligo nucleotide (DNA, RNA)
Required amount of measurement	500 pmol~1 nmol
Maximum measuring range	1000 - 15,000 Da (within about 50 mer)
Using matrix	3-hydroxy-2-pyridinecarboxylic acid
Analysis period	Within 2 days after receipt of sample
Result of analysis	Send report (attachment) by e-mail

# DNA/RNA Amplification

# **Molecular Biology**

► DNA/RNA Amplification ······	24 - 25 Page
Gene Expression Analysis	26 - 30 Page
Gene Synthesis · · · · · · · · · · · · · · · · · ·	31 - 35 Page
Sanger Sequencing	36 - 38 Page
mRNA Profiling · · · · · · · · · · · · · · · · · · ·	39 - 44 Page



# **DNA / RNA Amplification**

#### **List of Service**

AccuPower® Customized DNA/RNA Amplification Kit

#### **Key Features**

- O Draw reliable results from repetitive DNA/RNA amplification tests
- Manufactures DNA/RNA amplification kits specially made for each customer
- Provide simplified procedures containing all the components necessary for PCR packaged and mixed in tubes/plates for reactions to be performed immediately just by adding the template DNA/RNA and D.W.
- Maintain enzyme activity for a month at room temperature, a year in a freezer by using our patented stabilizer

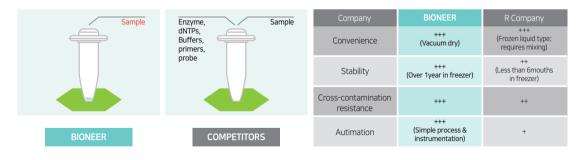
#### **Contact Us**

E-mail:customized\_service@bioneer.co.kr, Tel: +82-42-930-8577

### AccuPower® Customized DNA/RNA Amplification Kit

#### Operation

AccuPower® Customized DNA/RNA Amplification Kit Service provides tailor-made DNA/RNA amplification PreMix products optimized to draw out highly reproducible results under customer's experiential conditions by undergoing multiple PCR tests. This has a great advantage of high reaction specificity and PCR amplification efficiency, along with minimized amplified products from non-specific reactions and primer-dimer formation. Comprehensive types of premixes for various types of PCR reactions can be ordered, such as complex genomics, cDNA templates, low-copy targets, multiplex PCR, and etc. Moreover, each tube contains mixtures, consisting of all the necessary reactants like DNA polymerase, dNTPs, reaction buffers, and thermostable PPase, mixed together and vacuum dried with the ordered primer sets, packaged with amounts sufficient for one PCR run for it to be easily started just be adding the template and the D.W.



Production Period	Single	~ 10 working days	
	2-3 Multiplex	~ 12 working days	
	4 Multiplex	~ 16 working days	
	5 Multiplex Inquire		
Minimum Quantity	1,920 tests (480 x 4 plates)		

<sup>\*</sup> The time taken for the manufacture will be longer if no template is provided.

<sup>\*</sup> The duration for the QC tests is not included in the production period.

<sup>\*</sup> The fee for the oligo synthesis is billed separately from the quoted price.

# Gene Expression Analysis

# **Molecular Biology**

DNA/RNA Amplification ······	24 - 25 Page
► Gene Expression Analysis ·····	26 - 30 Page
Gene Synthesis · · · · · · · · · · · · · · · · · ·	31 - 35 Page
Sanger Sequencing	36 - 38 Page
mRNA Profiling · · · · · · · · · · · · · · · · · · ·	39 - 44 Page



# **Gene Expression Analysis**

#### **List of Services**

AccuPower® qPCR Array ServicePrimer & Probe Design ServiceAccuPower® Customized qPCR Panel KitSingle Gene qPCR Primer Set

#### **Key Features**

- O Directly use the results in SCI papers as all the procedures are conducted strictly to the MIQE guideline
- © Perform service according to the customer's needs and quickly provide the customized publication grade analysis results
- Use 12,000 pairs of human primer libraries with proven amplification efficiency to provide accurate results

#### What is MIQE Guideline?

MIQE (Minimum Information for Publication of Quantitative Real-time PCR Experiments) Guideline\* is an evaluation standard agreed upon by qPCR experts around the world for quality control of the "qPCR Gene Expression Analysis" papers. A step-by-step checklist is presented throughout the entire experiment to determine and evaluate the reliability, accuracy, and reproducibility of the qPCR experiments. This service provides all the information of the conducted experiences to the customers for them to evaluate the results by themselves.

\*Bustin, S.A., et al. 2009. The MIQE Guidelines: Minimum Information for Publication of Quantitative Real-Time PCR Experiments, *Clinical Chemistry* 55:4, 611-622.

#### **qPCR Array Library**

Primer Library is designed using primer blast and its bioinformatics tool. It is classified into 10 categories, including an unknown set of genes. See qPCR Array Library when selecting the target gene.

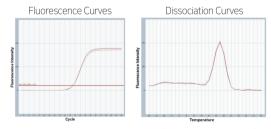


Figure 1. Primer validation result (Example. ACTB gene)

qPCR Array Library Categories	
1. Cancer	6. Signal transduction
2. Cellular differentiation & Regulation	7. Stem cell & Cell development
3. Disease	8. Toxicology & Drug metabolism
4. Immunology	9. Biological process
5. Metabolism	10. Unknown genes

#### **Contact Us**

E-mail:qPCRarray@bioneer.com, Tel: +82-42-930-8673

### AccuPower® qPCR Array Service

#### O Description

qPCR Array Service undergoes quantitative analysis on various gene expression. We provide this service for gene analysis requested by customers and pathways for numerous diseases and metabolism.

The qPCR Array combines real-time PCR, being the most sensitive and reliable analysis for gene expression, and microarray, allowing to profile multiple gene expression patterns once. This can enable accurate quantitative analysis of multiple genes at a low cost.

#### Workflow

#### 1. Consultation and Sample shipping

The starting stage differs by the types of the requested sample. We accept cells, tissues, blood, total RNA, cDNA, etc. Please refer to our homepage or contact us to discuss more information on sample preparation.

#### 2. Total RNA extraction and gDNA removal

We choose the suitable RNA extraction kit for maximized quality and purity of total RNAs depending on the types of sample. Furthermore, we treat the sample with RNase free DNase I to eliminate gDNAs that may affect the mRNA quantification analysis.

#### 3. Quality Check

As impurities in the total extracted RNA can inhibit the PCR, thus affecting the amplification efficiency, we use spectrophotometers to measure both the quantity and purity of nucleic acids. Not only that, as it is difficult to accurately quantify the degraded DNA, capillary gel electrophoresis will be also used to measure the RNA quality score (RQS). All the measurements will be recorded in the reports and sent to the customers. We proceed to the next step only when the purity and degree of degradation of the RNA satisfy the recommended value (RQS > 8.0).

#### 4. cDNA synthesis

We synthesize the full-length cDNA, even from RNAs with complex secondary structures, to get accurate expression analysis results using our two patented technologies: Thermostable RTase ( $RocketScript^{TM}$ ) and Cyclic Temperature Reverse Transcription (CTRT). We use different RT primers, Oligo dT or Random primers (dN6, dN12), depending on the customer's research purposes.

#### 5. qPCR

We use Exicycler™ 96 Real-Time Quantitative Thermal Cycler having our patented highly-sensitive optical detection system to draw exceptionally reliable qPCR data. We also have all our equipment regularly inspected and managed under ISO regulations While the qPCR primer sets, the standard specimen mixed with various cDNA, are used with stratagene reference total RNA(Human, Mouse, Rat) or the specimen provided by the customers to check the target specific amplification.

#### 6. Analyzing and providing results

We carry out relative quantitative analysis of qPCR by using  $2^{-\Delta\Delta Ct}$  method. Customers will receive Excel files having the following data: raw Ct values, melting curve analysis, p-values, fold changes, scatter plots, volcano plots, and heat map images. If the customer has installed  $Exicycler^{TM}$  analysis software, we can also provide qPCR raw data. Additionally, information on primers such as genetic information, amplicon size, and PCR conditions following the MIQE can be also provided upon the customer's request.

Customized Service		≤ 16 genes	≤ 32 genes	≤96 genes
cDNA Sample	cDNA Sample Minimum Volume 200 µl		400 µl	900 µl
DNA Cample	Minimum Volume /Minimum Concentration	15 µl / 100 ng/µl		
RNA Sample	Total RNA Amount	> 5 µg	> 10 µg	> 20 µg
	Service Period	2 weeks	2~3 weeks	3~5 weeks

<sup>\*</sup> Service period may be prolonged if the request is made by using the primer not owned by Bioneer.

<sup>\*</sup> Please inquire Single gene qPCR separately.

<sup>\*</sup> For details about the request, visit the Bioneer website(www.bioneer.com)

### AccuPower® Customized qPCR Panel Kit

#### O Description

AccuPower® Customized qPCR Panel Kit has a batch of proven primer pairs that can be easily used for analysis just by adding cDNA samples and 2X Master Mix. Additionally, we can provide customized plate layout designed and made according to the customer's needs. We have our specialized experts to manufacture the plates according to the MIQE (Minimum Information for Publication of Quantitative Real-time PCR Experiments) Guidelines\* for accurate, reliable, and quick analysis results to be used in SCI papers.

\*Bustin, S.A., et al. 2009. The MIQE Guidelines: Minimum Information for Publication of Quantitative Real-Time PCR Experiments, Clinical Chemistry 55:4, 611-622

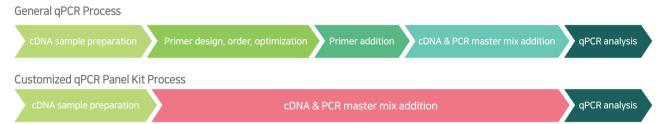


Figure 1. Comparison with the general qPCR and customized qPCR panel kit.

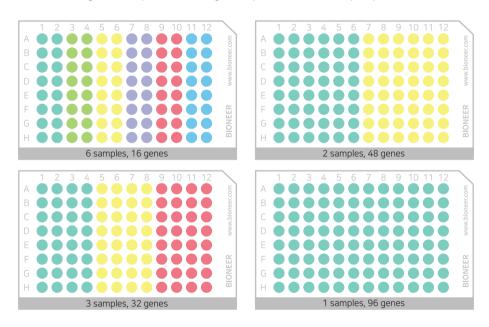


Figure 2. Examples of customized qPCR panel kit by number of genes.

The Number of Genes	1 - 16 genes	17 - 32 genes	33 - 96 genes	
Minimum Order Quantity	3 plates	4 plates	9 plates	
Service Period	Inquire			

<sup>\*</sup> The service can be only ordered through email. Please download the order form in our website (www.bioneer.com) and send it to us.

<sup>\*</sup> Services requested with the primers not owned by Bioneer will take longer time than usual.

<sup>\*</sup> Refer to our qPCR Array Library for the target gene selection (genes not found in the library is also available.).

<sup>\*</sup> All the qPCR primer sets have been validated by *Exicycler™* 96 and *AccuPower®2X GreenStar™* Master Mix.

### **Primer & Probe Design Service**

#### O Description

Accurate primer and probe designs are essential for obtaining a reliable real-time PCR data. Nevertheless, those processes require years of experience and knowledge in gene tests and construction.

Through our continuous researches and development, our gene expression analysis team has gained about 12,000 validated primer pairs to obtain accurate expression analysis results. With this expertise, our experts can quickly provide the best optimized primer designs & probes fitting customers' research purposes the most.

#### Ordering Information

You can find the service information in our Bioneer website (www.bioneer.com). Please send the application form through our email (qPCRarray@bioneer.com).

Number of Genes	Two primary genes	For two additional genes	
Service Period	Reports are sent on the very day	Adds one more day	
Cat. No.	S-6000		

### Single Gene qPCR Primer Set

#### Operation

Our Single Gene qPCR Primer Sets not only validate primers for gene expression analysis, but also synthesize and delivers them to the customers. Qualification processes are the most important step in obtaining reliable real-time PCR analysis data. To make precise and accurate qPCR primers for expression analysis, it not only requires constructive designs, but also needs numerous validation tests. Thus, even the most experienced experts require considerable time and money.

Our experts in Bioneer meticulously test the target specificity and amplification efficiency of the primers strictly based on the sequences provided by the customers. Then, they will be synthesized and delivered. All the tests will comply with MIQE Guidelines, the guide for reliable qPCR test results. Primers will be synthesized and sent with the amount sufficient for 200 qPCR tests. Our products can be immediately used in research papers without requiring the customers to do any validation tests prior to their experiments.

If the customers do not know, or cannot provide their primer sequences, we can still synthesize, verify, and ship the sets by sending us the target genomic information for expression analysis with "Primer & Probe Design Service," which will require a separate charge.

#### Ordering Information

You can find the service information in our Bioneer website (www.bioneer.com). Please send the application form through our email (qPCRarray@bioneer.com).

Cat. No.	Product Description	
S-6042-S200	Single Gene qPCR Primer Set	
Related product	Related products	
K-3140	AccuPrep® Universal RNA Extraction Kit (100 rxn)	
K-2201	AccuPower® RocketScript™ Cycle RT PreMix (dT₂₀) (96 T, 20 μl)	
K-6254	AccuPower® 2X GreenStar™ qPCR Master Mix (-ROX Dye) (5 ml)	

<sup>\*</sup>Single Gene qPCR Primer Set is optimized for the above products.

# **Gene Synthesis**

# **Molecular Biology**

	DNA/RNA Amplification ······	24 - 25 Page
	Gene Expression Analysis	26 - 30 Page
<b>&gt;</b>	Gene Synthesis · · · · · · · · · · · · · · · · · ·	31 - 35 Page
	Sanger Sequencing	36 - 38 Page
	mRNA Profiling	39 - 44 Page



# **Gene Synthesis**

#### **List of Services**

Gene Synthesis Service

Rapid Gene Synthesis Service

AccuGeneBlock Service

Gene Cloning Service

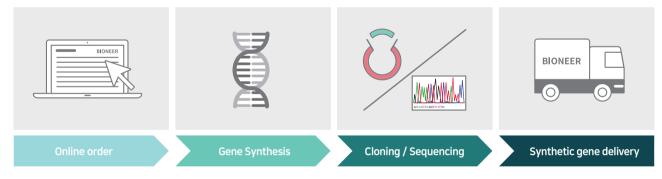
Mutagenesis Service

Codon Library Mutagenesis Service

### **Key Features**

- © Use integrated production system utilizing high-speed oligo synthesizer allowing to quickly synthesize uniformed quality of oligos from raw materials
- © Guarantee 100% sequence accuracy using Automatic DNA sequencer (ABI 3730)
- © Provides codon optimization services with our accumulated experience and techniques
- © Select from wide range of choices for compatible vectors according to the desired applications at a reasonable price

#### Workflow



#### **Contact Us**

E-mail:geneorder@bioneer.com, Tel: +82-42-930-8793, 8515

### **Gene Synthesis Service**

#### O Description

Gene Synthesis Service synthesizes and provides genes with 100% accuracy only by the gene or protein sequence information.

#### Ordering Information

Service Period	1 – 1,200 bp	(Guarantee) 12 working days
	1,201 – 2,000 bp	(Guarantee) 17 working days
	2,001 – 3,000 bp	(Guarantee) 22 working days
	3,001 bp -	Inquire
Delivery Form	2 ~ 5 ug of lyophilized plasmid (*Low copy plasmid: 1~2 μg)	
Cloning Vector	Basic vector -pBHA	
	Other - Inquire	
Subcloning	Combined with Gene Cloning Service if the order contains commercial or custom vectors	
Additional Service	Plasmid Amplification Service	

<sup>\*</sup> The service period will be extended along with the price increase if the sequences contain complicated structures such as high or low GC, repeats, homo-polymeric runs, etc.

### Rapid Gene Synthesis Service

#### O Description

Rapid Gene Synthesis Service provides synthetic genes just within 5 to 8 days after the order date.

Service Period	1 - 500 bp	5 working days
	501 – 1,500 bp	8 working days
Delivery Form	2 ~ 5 ug of lyophilized plasmid	
Cloning Vector	рВНА	
Additional Service	Plasmid Amplification Service	

<sup>\*</sup> Synthesis of genes having complex sequences such as high or low GC, repeats, homo-polymeric runs, etc. may be limited. In this case, Please use our General Gene Synthesis service.

<sup>\*</sup> Genes inhibiting *E.coli* growth or producing substances toxic to its cellular system will be delivered either in the mutated form of plasmid DNA or in the form of PCR products.

<sup>\*</sup> Please use the "Gene to Protein Service" to undergo protein synthesis and purification after our gene synthesis service to obtain the proteins directly.

<sup>\*</sup> Visit our Bioneer website (www.bioneer.com) for more information on vectors.

<sup>\*</sup> Synthesis of genes inhibiting *E.coli* growth or producing toxic substances to its cellular system may be limited. In this case, Please use our General Gene Synthesis service. We cannot guarantee the service time if we find those characteristics during the synthesis process.

<sup>\* 40%</sup> of discount will be applied if the synthesis is completed after our guarantee period.

### AccuGeneBlock Service

#### O Description

AccuGeneBlock Service provides synthetic genes less than 1 kb in the form of double stranded DNA within at least 3 days.

#### Ordering Information

Service Period	500 bp or less	Average 3 - 5 working days
	501 – 1,000 bp	Average 4 - 6 working days
Delivery Form	500 ng ~ 1 ug of lyophilized PCR product (3' A-tailing or blunt end form)	
Additional Service	Amplification Service	

<sup>\*</sup> Synthesis of genes having complex sequences such as high or low GC, repeats, homo-polymeric runs, etc. may be limited. In this case, Please use our General Gene Synthesis service.

## **Gene Cloning Service**

#### O Description

Gene Cloning service provides a simple, straightforward cloning and sequence check using our long periods of expertise in the field of molecular biology.

Available Materials	Plasmid DNA	≥ 10 µl, 150-200 ng/µl
	PCR Product	≥ 10 µl, 50 ng/µl
Service Period	1 - 8 kb (vector+insert)	Average 5 - 10 working days
	8 - 11 kb (vector+insert)	Average 10 - 15 working days
	11 kb - (vector+insert)	Inquire
Additional Service	Plasmid Amplification Service *Excluded for low copy plasmid service	
	Discount when combined with Gene Synthesis Service	

<sup>\*</sup> The price and period can increase depending on the structures and characteristics of the genes.

<sup>\*</sup> Only 1 Kb or less genes can be synthesized.

<sup>\*</sup> All the products are shipped after a meticulous sequence check.

<sup>\*</sup> Vector costs will be charged separately if the cloning service is requested with a commercial vector.

<sup>\*</sup> Sending the wrong sample will charge a fee of 50% of the total cost, so please check before the packaging.

<sup>\*</sup> A fee of 50% of the total cost will be charged for the cancelation.

<sup>\*</sup> If you want us to pause the service due to your personal reasons, we can hold for 1 month at maximum. Afterwards, we will automatically cancel your order and charge 50% of the total cost.

## **Mutagenesis Service**

#### O Description

Mutagenesis service synthesizes mutant genes essential for studies on structures of proteins and improvements on enzyme functions. This can be used with Gene Synthesis Service to prepare your researches quickly and easily.

#### Ordering Information

Available Materials	Plasmid DNA	≥ 10 µl, 150-200 ng/µl
Service Period	- 1 kb	Average 5 -10 working days
	1 kb - 3 kb	Average 10 - 15 working days
	3 kb - 5 kb	Average 15 - 25 working days
	5 kb -	Inquire
Plasmid Increase Service		
Additional Service	Discount when combined with gene synthesis	

<sup>\*</sup> The price and period can increase depending on the structure and characteristics of the gene.

# **Codon Library Mutagenesis Service**

#### O Description

Codon Library Mutagenesis Service can provide gene libraries by codon-based mutagenesis based on the codon bias of a species of the customer's interest. Simply select the species to express the genes and specify the location of the codons, along with the name of the amino acids for us to undergo the codon mutation process in the DNA form according to the codon usage of the selected species. A maximum of 7 consecutive codons can be selected for the mutation sites (maximum of 21 bp for DNA).

#### Ordering Information

Available Materials	Plasmid DNA	≥ 10 µl, 150-200 ng/µl
	- 1 kb	Average 5 - 10 working days
Service Period	1 kb - 3 kb	Average 10 - 15 working days
Service Period	3 kb - 5 kb	Average 15 - 25 working days
	5 kb -	Inquire
Plasmid Amplification Service		
Additional Service	Discount when combined with gene synthesis	

<sup>\*</sup> The price and period may increase depending on the ordered structure and characteristics of the gene.

<sup>\*</sup> Sending the wrong sample will charge a fee of 50% of the total cost, so please check before the packaging.

<sup>\*</sup> Sending the wrong sample will charge a fee of 50% of the total cost, so please check before the packaging.

# Sanger Sequencing

# **Molecular Biology**

	DNA/RNA Amplification ······	24 - 25 Page
	Gene Expression Analysis	26 - 30 Page
	Gene Synthesis · · · · · · · · · · · · · · · · · ·	31 - 35 Page
<b>&gt;</b>	Sanger Sequencing · · · · · · · · · · · · · · · · · · ·	36 - 38 Page
	mRNA Profiling · · · · · · · · · · · · · · · · · · ·	39 - 44 Page



# Sanger Sequencing

#### **List of Services**

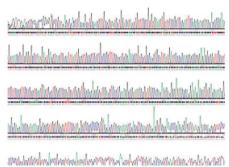
Standard Sequencing Service
Difficult Sequencing Service

Full-length Sequencing Service (primer walking) Additional Service

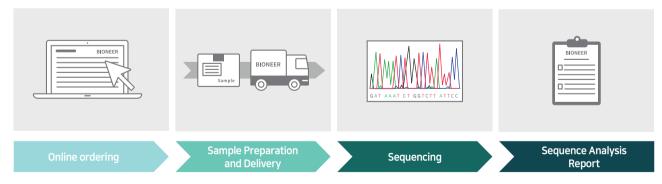
## **Key Features**

- High quality data with accurate results through automated process (Phred Score (QV): ≥ 20, Guaranteed read lengths: ≥ 700bp)
- © Supply of commonly used universal primer capable of being selected upon order





#### Workflow



#### **Contact Us**

E-mail:sequencing@bioneer.co.kr, Tel: +82-42-930-8777

## Sanger Sequencing

#### O Description

#### **Standard Sequencing**

Standard Sequencing Service undergo analysis of desired plasmid or PCR product regions using a primer optimized for most of the normal samples.

#### Full-length Sequencing (Primer walking)

Full Length Sequencing Service utilizes primer walking to analyze plasmids, long inserts, or PCR products. The service time depends on the size, but it takes about a week for a plasmid being about 3 kb long. The data will be provided in the form of contig file for each single data and assembly while the primer and its design will be sent in a dried form after the analysis.

#### **Difficult Sequencing**

This product is for analyzing templates difficult to be done using the Standard Sequencing service such as those having GC/AT rich regions, repeated sequences, secondary structures, etc. This service is optimized using our long time experiences and techniques by utilizing different conditions and reagents from the Standard Sequencing Service.

#### **Additional Service**

Optional choices of services can be provided depending on the requested type of sample, such as PCR purification, Gelextraction, and PCR clean-up.

#### Ordering Information

#### Sample and primer preparation

Sample Type	Concentration	Volume/rxn	Purity
Plasmid DNA	≥ 100 ng/µl	≥ 10 µl	
PCR product	≥ 50 ng/µl	≥ 10 µl	A <sub>260</sub> /A <sub>280</sub> : ~1.8
Non-purified PCR product	≥ 50 ng/µl	≥ 20 µl	A <sub>260</sub> /A <sub>230</sub> : 2.0-2.2
Genomic DNA	≥ 30 ng/µl	≥ 30 µl	

Primer Type	Concentration	Volume/rxn
Specific primer	5 pmol/μl (5 μm)	5 µl
Universal primer	Simply write the primer name if owned by our company; free primers are available for use.	
Ordered primer	Can order with oligo and sequencing service at the same time	

<sup>\*</sup> Please prepare the sample in a 1.5 ml tube or plate based on 1 rxn basis also with an amount sufficient for a re-reaction.

<sup>\*</sup> Samples and primers will be discarded after a month. Please ask us separately if you want us to store them longer.

<sup>\*</sup> Although you can keep the primers, we are not responsible for their degradation caused by long-term storage. We recommend to change the primers every month and freeze them in several tubes.

# mRNA Profiling

# **Molecular Biology**

DNA/RNA Amplification	24 - 25 Page
Gene Expression Analysis	26 - 30 Page
Gene Synthesis · · · · · · · · · · · · · · · · · ·	31 - 35 Page
Sanger Sequencing	36 - 38 Page
► mRNA Profiling ······	39 - 44 Page



# mRNA Profiling

#### **List of Services**

Next-generation RNA Sequencing (RNA-seq) + Gene Expression Analysis Service (qPCR Array Service)

#### **Key Features**

© Undergo RNA-seq and qPCR array to provide both single-step gene discover and verification services at once

#### **RNA Sequuncing**

- Total RNA Sequencing
- mRNA Sequencing
- Small RNA Sequencing



#### **Gene Expression Analysis Service**

<Service>
qPCR Array Service

<Primer Optimization Service>
qPCR Array System: Customized qPCR panel kit
Single Gene qPCR Primer Set

#### Workflow

E-mail order  $\rightarrow$  Sample shipment  $\rightarrow$  RNA extraction and QC  $\rightarrow$  Library Prep.  $\rightarrow$  NGS  $\rightarrow$  Data report  $\rightarrow$  qPCR Array Service sample shipment  $\rightarrow$  RNA extraction and QC  $\rightarrow$  cDNA synthesis  $\rightarrow$  qPCR and data analysis  $\rightarrow$  Data report

#### **Contact Us**

RNA-seq E-mail: NGS\_support@bioneer.co.kr, Tel: +82-42-930-8553 qPCR Array Service E-mail:qPCRarray@bioneer.com, Tel: +82-42-930-8673

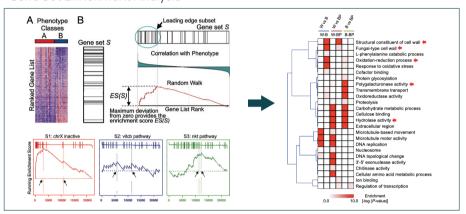
#### O Description

RNA-seq and qPCR Array Systems are high quality services performed by our experienced experts. Not only we can provide RNA Sequencing Service to check the differences between the sample expression values, but also offer differentiated analysis packages (such as Gene Expression, DEG, GO, etc.) from rich bioinformatics infrastructure. Furthermore, our qPCR Array Service complies with the MIQE Guidelines\* made for real-time PCR experts to quickly deliver accurate real-time PCR data which can be directly published in research papers.

\* Bustin, S.A., et al. 2009. The MIQE Guidelines: Minimum Information for Publication of Quantitative Real-Time PCR Experiments, *Clinical Chemistry* 55:4, 611-622.

## RNA-Seq (NGS)

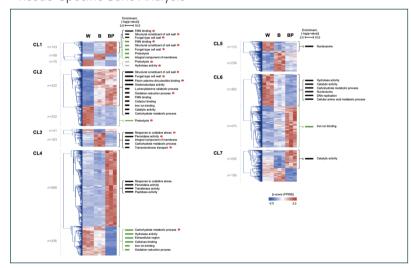
• Gene Set Enrichment Analysis



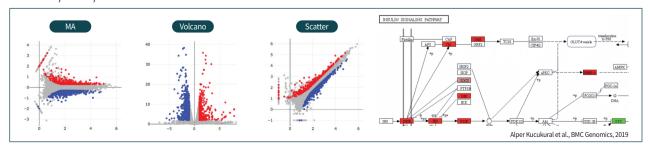
#### Functional Annotation

# GO, Pathway, PPI, Disease, etc. Kind of Sequences, Gene IDs, etc. Statistics Calculate enrichment paulus with methods like Fieher exact, typergement; Binomial distribution, etc.] Pigure 1. The infrastructure of typical enrichment tools. Even though the enrichment analysis tools have distinct features, they can be generally described as three major layers: backend annotation database; data mining, and result presentation. Each of the layers, rather than statistical methods alone, greatly influences the analytic results. Da Wei Huang et al., Nucleic Acids Research, 2009

#### • Tissue-Specific Gene Analysis

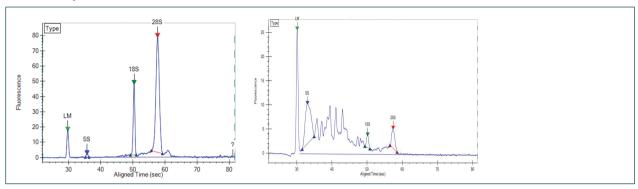


#### Pathway Analysis

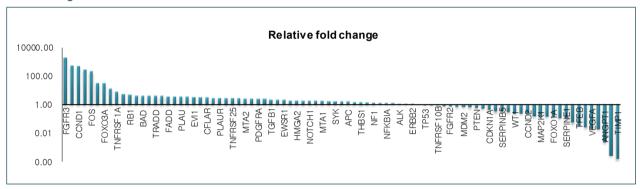


## **qPCR Array Service**

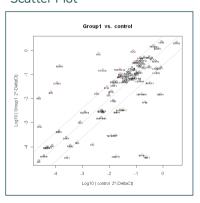
#### • RNA Quality Check



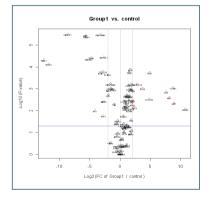
#### • Fold change



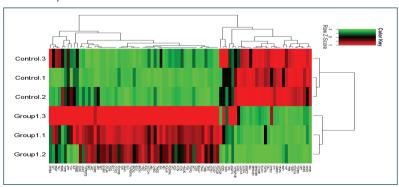
#### Scatter Plot



#### Volcano Plot



#### Heat map



#### Ordering Information

#### Sample Preparation

#### 1. RNA sample

Quickly proceed in an RNase-free environment during the RNA extraction. You must remove the genomic DNA for us to go through accurate analysis. If both RNA-seq and qPCR array services have been requested for the same type of samples, please provide each of them separately with the sufficient amounts and store them at  $-80^{\circ}$ C.

#### Sample Requirements

- RNA-seg

Minimum Quantity	Minimum Concentration	Total RNA Quantity
200 ng	65 ng/µl	≥1 µg

#### - qPCR Array Service

Number of Analysis Genes	Minimum Volume	Minimum Concentration	Total RNAQuantity
≤ 16 genes			> 5 µg
≤ 32 genes	15 µl	100 ng/μl	> 10 µg
≤ 96 genes			> 20 µg

#### Sample Purity & Integrity Standard

	Method	Standard
Purity	Spectrophotometer	260/280nm > 1.8
loto arity (	Capillary electrophoresis	RIN or RQS or RIS > 8.0 (NGS: > 7)
Integrity	Normal electrophoresis	28S/18S ratio 2:1

#### 2. Tissue and Cell sample

For harvested cell pellets or extracted tissues, prepare the samples by using one of the following methods to minimize RNA degradation.

- Method 1 Immediately freeze the samples with liquid nitrogen and store them at 80°C.
- Method 2 Process the samples with the RNA stabilization solution by following its manual and store them at -80°C.
- **Method 3** Process with 1% Beta-mercaptoethanol in the Lysis buffer (Guanidinium thiocyanate buffer\*). Add 400 μl of lysis buffer per 10<sup>4</sup> ~ 10<sup>6</sup> of cells, crush using a homogenizer, and store them at 80°C (\*RB buffer in *AccuPrep*® Universal RNA Extraction Kit).

#### 3. Requesting blood or other samples

- Those will require consultation before processing the orders.
- We do not provide the service for cDNA as we cannot guarantee the accuracy of its RNA Seq analysis as it is difficult to undergo QC tests.

\*\* qPCR Array System: For Customized qPCR panel kits and Single Gene qPCR Primer Sets, download the order form from our Bioneer websites (www. bioneer.com) and send it to our gene expression team email (qPCRarray@bioneer.com)

#### Address to send the samples

- RNA-seq: 8-11, Munpyeongseo-ro, Daedeok-gu, Daejeon 34302, Republic of Korea, Bioneer Sequencing Team.
- qPCR Array Service: 8-11, Munpyeongseo-ro, Daedeok-gu, Daejeon 34302, Republic of Korea, Bioneer gene expression analysis Team.

#### **Precautions**

- Please refrain shipping on Friday or holiday eves to avoid any deterioration in sample qualities due to delays in deliveries.
- Please include enough dry ice to prevent RNA degradation during the shipment period.

#### Service period

- RNA-seq: about 4~6 weeks
- qPCR Array Service:

Number of Analysis Genes	Service Period
≤ 16 genes	about 2 weeks
≤ 32 genes	about 2~3 weeks
≤ 96 genes	about 3~5 weeks

<sup>\*</sup> Service time can vary according to the methods used for sample extraction and data analysis.

<sup>\*</sup> Samples will be discarded after 1 month of service termination.

# Cell-free Protein Synthesis

# **Protein Synthesis**

► Cell-free Protein Synthesis · · · · · 45 - 48 Page



# **Cell-free Protein Synthesis**

#### **List of Services**

Standard Protein Synthesis Service One-day Protein Synthesis Service

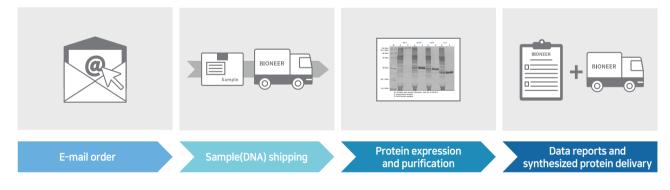
Standard Gene to Protein Service Cloning-free Gene to Protein Service

#### **Key Features**

- © Quickly provide purified proteins for customer's experimental purposes using E. coli cell-free protein expression
- O Synthesize various kinds of proteins, even toxic ones that are unable to be done through cell culturing
- © Can allow customers to synthesize proteins directly by themselves using our protein synthesis kit if necessary

## What is cell-free protein expression system?

Cell-free protein expression system allows to express proteins from nucleic acids without culturing  $E.\ coli$ . The recombinant proteins are expressed and purified through  $in\-vitro$  transcription, translation and magnetic separation using the protein synthesis kit with  $E.\ coli$  cell extract and  $ExiProgen^{TM}$  instrument. This allows to synthesize not only enzymes, growth factors, hormones, antigen proteins and antibodies, but also toxic proteins that could not be produced through cell culture.



#### **Contact Us**

E-mail: sales@bioneer.co.kr or proteinorder@bioneer.com, Tel: +82-42-930-8777

## **Standard Protein Synthesis Service**

#### O Description

Our Standard Protein Synthesis Service can provide recombinant proteins within at least a week by using our automatic protein synthesis system ( $ExiProgen^{TM}$  Protein Synthesis System) to synthesize various proteins such as enzymes, growth factors, hormones, antigen proteins, and antibodies.

#### Workflow

According to the type of the requested template DNA, Protein Synthesis Service is divided into two different ways.

E. coli expression vectors (ex. T7-based expression vector)

**Quotation and Order** 

Collection of sample DNA

Protein expression and purification with  $ExiProgen^{\text{TM}}$ 

Non-expression vectors (ex. Cloning vector/ target gene)

**Quotation and Order** 

Collection of sample DNA and design of primers

Preparation of template DNA (PCR product)

Protein expression and purification with *ExiProgen*™

#### Ordering Information

Amount of Samples	E. coli expression vector	0.3kb -4.0kb by ORF(open reading frame)
		> 10 µg (A <sub>260/280</sub> : 1.7 - 2.0, A <sub>260/230</sub> : >1.5)
	Non-expression vector	0.3kb -1.5kb by ORF(open reading frame)
		> 2 µg
Service Period	Average 7 – 14 working days	
Delivery Form	Data report	E-mail
	Protein	100 μg (0.1 - 1.0 mg/ml) of solution

<sup>\*</sup> Providing plasmid DNA samples not reaching the standard for amount and purity conditions will charge additional costs as DNA amplification services must be done beforehand.

# **One-day Protein Synthesis Service**

#### Operation

Our One-day Protein Synthesis Service provides synthesis result of recombinant proteins within a single day. We use our automatic protein synthesis systems ( $ExiProgen^{TM}$  Protein Synthesis System) to synthesize various proteins such as enzymes, growth factors, hormones, and antiqen proteins.

#### Ordering Information

Amount of Sample	E. coli expression vector1)	> 10 µg (A <sub>260/280</sub> : 1.7 - 2.0, A <sub>260/230</sub> : > 1.5)
Service Period <sup>2)</sup>	1 working days	
Dalinama Farma?)	Data report	E-mail
Delivery Form <sup>3)</sup>	Protein	0.1 - 1.0 mg/ml of solution

<sup>1)</sup> Providing plasmid DNA samples not reaching the standard for amount and purity conditions will charge additional costs as DNA amplification services must be done beforehand.

<sup>\*</sup> Requesting non-expression vectors such as cloning vector of PCR products will charge additional costs as primary primer sets and template DNA preparation services must be done beforehand.

<sup>\*</sup> The service price is for synthesis of 100 μg which is the total protein amount regardless of the target protein purity.

<sup>2)</sup> Service period does not include the time taken during the shipment.

<sup>3)</sup> Synthesized proteins are provided after one reaction. Ordering more proteins will shift the service to the "Standard Protein synthesis Service".

## Standard Gene to Protein Service

#### O Description

Standard Service synthesizes genes based on our Gene Synthesis Service by cloning into an expression vector for *E. coli*, followed by the protein synthesis service.

#### Workflow

Online Order E. coli Codon Optimized Gene synthesis Protein synthesis Gene and Protein delivery

#### Ordering Information

Service Period	1 - 1,500 bp	Average 20 - 35 working days
	1,501 - 3,000 bp	Average 30 - 40 working days
Delivery Form	Gene	2 - 5 μg of lyophilized plasmid
	Protein <sup>1)</sup>	100 μg (0.1 - 1.0 mg/ml) of solution

<sup>1)</sup> We will only charge the set up fee after failing the protein synthesis while the gene synthesis fee will be charged normally, and send the synthesized genes along with the data report.

## Cloning-free Gene to Protein Service

#### Operation

Cloning-free Service provides proteins based on the PCR products after synthesizing genes using AccuGeneBlock Service. As cloning is not required, this service will take relatively shorter time compared to other services.

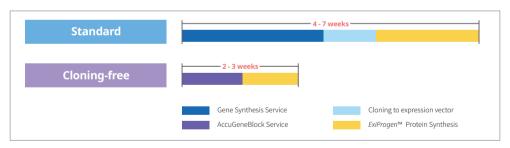


Figure 1. Comparsion of service time

#### Workflow

Online Order E. coli Codon Optimized Gene synthesis Protein synthesis Gene and Protein delivery

#### Ordering Information

Service Period	Average 10 - 15 working days	
Delivery form	Gene	500 ng - 1 μg of lyophilized plasmid PCR product
	Protein <sup>1)</sup>	100 μg (0.1 - 1.0 mg/ml) of solution

<sup>1)</sup> We will only charge the set up fee after failing the protein synthesis while the gene synthesis fee will be charged normally, and send the synthesized genes along with the data report.

<sup>\*</sup> Canceling your order within 5 days after receiving the confirmation email will charge 50% of the original price. while 80% of the fee will be charged after that time.

<sup>\*</sup> Ordering complicated sequences such as those containing high or low GC content, repeats, or homopolymeric runs will charge additional costs.

All about Bioneer's

# **Custom** Services





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