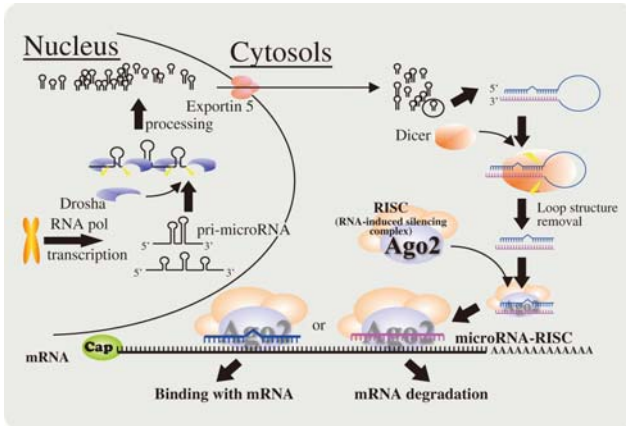
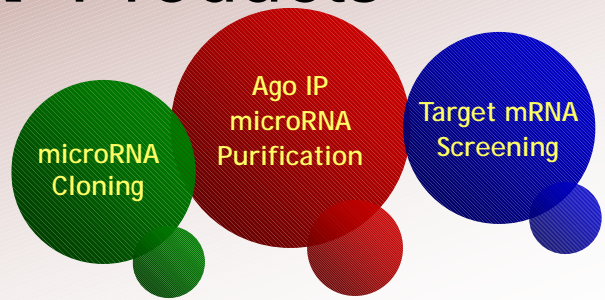




# microRNA Products

Wako Pure Chemical Industries, Ltd.



microRNAs (miRNAs) are endogenous RNAs, approximately 22 nucleotides in length, that can play important regulatory roles in animals and plants by targeting mRNAs for cleavage or translational repression. Since more than 1,000 kinds of miRNAs are suggested to exist in humans and mice, various researches such as identification of novel miRNAs with unknown functions and elucidation of the miRNA functions associated with specific diseases have been actively progressed around the world. Recently, the discovery of miRNAs in serum has opened up the possibility of using miRNAs as biomarkers of disease including cancer. microRNAs are considered to degrade the mRNA chains or inhibit the translation by the following manner: they are matured through multiple steps in cells and incorporated into a protein complex, RNA-induced silencing complex (RISC), they then

bind to argonaute subfamily of proteins which are the main components of RISC, followed by binding to a target mRNA. Four kinds of Ago subfamilies (hAgo1 to hAgo4) exist in humans and are ubiquitously expressed even though the expression levels vary with types. Among a group of Ago subfamilies, Ago2 is the most expressed protein which is considered to play a central role in miRNA pathway because it only has a slicer activity for cleavage of the target RNAs. As RISC is immunoprecipitated by antibodies against these Ago proteins, the miRNAs can be recovered from the precipitated RISC, and furthermore, because the target mRNAs are found to be coprecipitated, immunoprecipitation is becoming an essential methodology to elucidate various functions of the miRNAs.

In these circumstances, Wako offers some analytical tools for the functional analysis of miRNAs, mainly the miRNA research tools using Anti Ago Antibodies. The combination use of these tools enables you to make a comprehensive analysis of the miRNAs and mRNAs contained in the RISC that exist in cells and tissues, which can be applied not only to miRNA analyses but to target mRNA analyses.

## Product List

Description	Wako Catalog No.	Page #
<b>microRNA Purification Kits based on Immunoprecipitation method using antibody against Argonaute protein</b>		
microRNA Isolation Kit, Human/Mouse Ago1	291-70201 (10 reactions)	2
microRNA Isolation Kit, Human Ago2	292-66701 (10 reactions)	2
microRNA Isolation Kit, Mouse Ago2	292-67301 (10 reactions)	2
microRNA Isolation Kit, Human Ago3	297-70301 (10 reactions)	2
<b>Preparation Kit of cDNA encoding microRNA</b>		
microRNA Cloning Kit <i>Wako</i>	290-66501 (8 reactions)	5
Single Strand DNA Ligase, thermostable, recombinant, Solution	298-65103 (200 units); 292-65101 (500 units)	5
PCR Purification Kit <i>Wako</i>	298-67901 (30 reactions)	7
Target mRNA Cloning Kit <i>Wako</i>	298-68001 (10 reactions)	7
<b>Antibodies</b>		
Anti Ago1, Monoclonal Antibody (1F2) <for WB>	018-22401 (50 µL)	8
Anti Ago1, Monoclonal Antibody (2A7) <for IP>	015-22411 (50 µL)	8
Anti Human Ago2, Monoclonal Antibody (4G8)	011-22033 (50 µL); 015-22031 (100 µL)	8
Anti Mouse Ago2, Monoclonal Antibody (2D4)	014-22023 (50 µL); 018-22021 (100 µL)	8
Anti Human Ago3, Monoclonal Antibody	available soon!	-

microRNA Isolation Kit, Human Ago2; Mouse Ago2: patent pending (11, 30, 2007); microRNA Cloning Kit *Wako*: patent pending (1, 10, 2007)  
All listed products are for research use only. Do not administer each to human..

## microRNA Isolation Kit Series

microRNA (miRNA) Isolation Kit series can prepare high purity fractions of microRNA, which are bound with human or mouse Argonaute (Ago) protein, based on immunoprecipitation method by using a high affinity monoclonal antibody. The purified miRNA fraction contains very little contaminated degradation fragments of rRNA and tRNA. These kits will highly improve the miRNA cloning efficiency compared with conventional miRNA purification method. Furthermore, the purified miRNA are applicable to Analysis of target mRNA of miRNA.

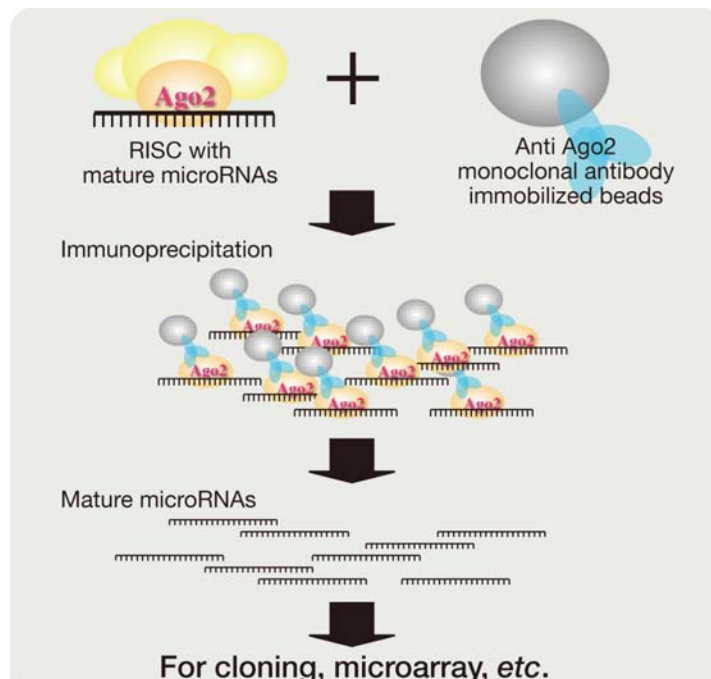
Sample		Platform			
Cultured Cells <sup>※</sup>	Tissues <sup>※</sup>	Microarray	Deep sequencer	Cloning	Quantitative RT-PCR
○	○	○	○	○	○

※Each isolation efficiency is depend on cell lines or tissues.

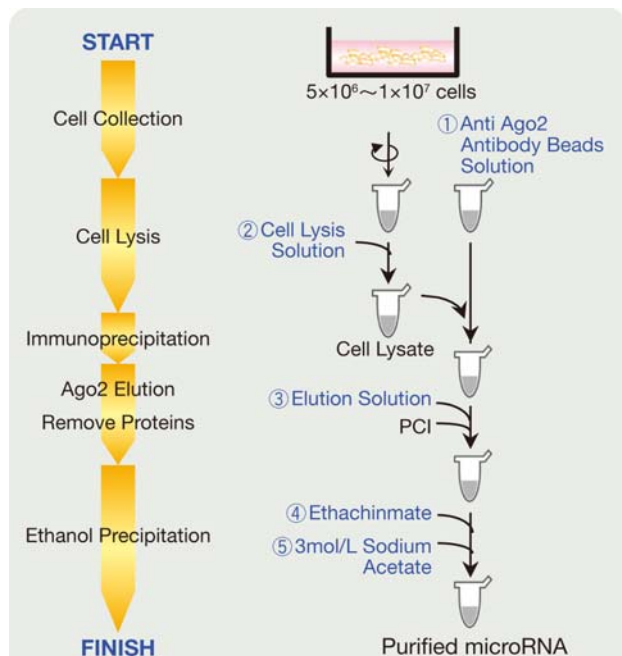
### ■ Features

- 1) Immunoprecipitation of endogenous Ago protein
- 2) High purification performance of miRNA bound with Ago protein
- 3) Little contamination of other RNAs such as degradation fragments of rRNA & tRNA
- 4) Applicable to high efficiency of microRNA cloning and microarray with purified microRNA fraction

### ■ Principle



### ■ Procedure Outline



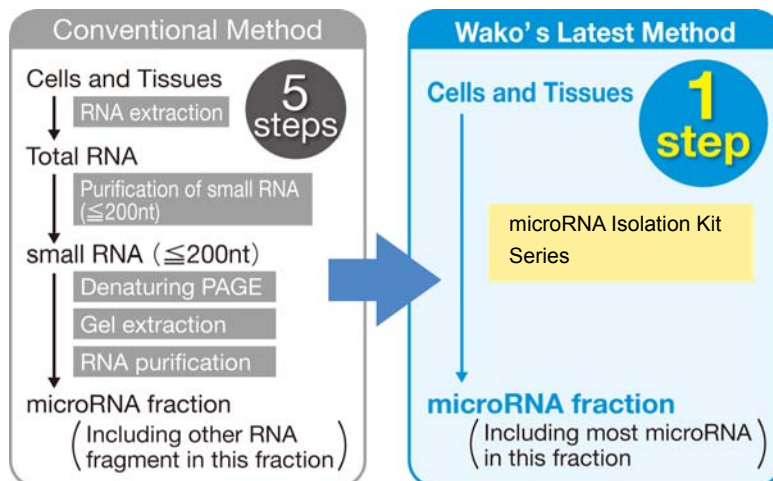
※: 2 x 10<sup>7</sup> cells are necessary for the procedure with miRNA Isolation

### ■ Kit Contents (for 10 reactions)

- |                                      |                      |
|--------------------------------------|----------------------|
| (1) Anti Ago Antibody Beads Solution | 1 vial x 500 $\mu$ L |
| (2) Cell Lysis Solution              | 1 bottle x 50 mL     |
| (3) Elution Solution                 | 1 vial x 500 $\mu$ L |
| (4) Ethachinmate                     | 1 vial x 30 $\mu$ L  |
| (5) 3 mol/L Sodium Acetate Solution  | 1 vial x 400 $\mu$ L |

### ■ Comparison with conventional method

	conventional	Wako's microRNA Isolation Kits
Total RNA Extraction	necessary	unnecessary
small RNA ( $\leq 200$ nt) Purification		
Denaturing PAGE		
Gel Extraction from denaturing PAGE	high	very low
contaminated degradation fragments of rRNA & tRNA in miRNA fraction	$\geq 1$ day	$\geq 0.5$ days
Test duration		



■ Purification of microRNA fraction from various cell lines

microRNA Isolation Kit, **Human Ago2** (Wako Cat. #292-66701)

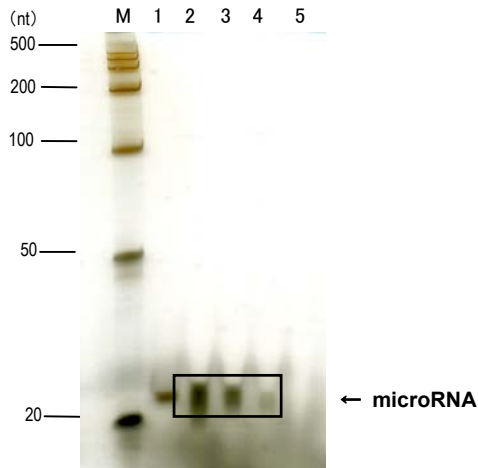


Figure: Purification of microRNA fraction by using microRNA Isolation Kit, **Human Ago2**. The purified microRNA fraction from human cultured cell lines (HeLa, HepG2, HEK293) were specifically detected by Urea-PAGE and silver staining (Wako Cat. #311-03961; CLEAR STAIN Ag). Cell number of each cell line is approximately  $5 \times 10^6$ . The applied volume per lane is half of  $10 \mu\text{L}$  of final solution prepared with an IP by this kit.

microRNA Isolation Kit, **Mouse Ago2** (Wako Cat. #292-67301)

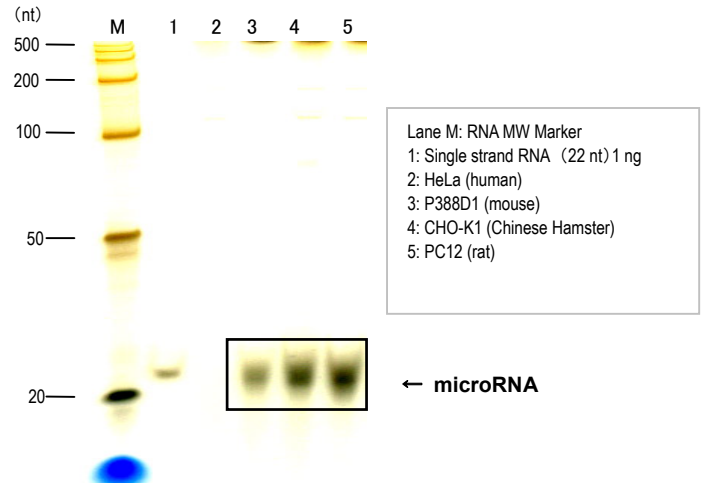


Figure: Purification of microRNA fraction by using microRNA Isolation Kit, **Mouse Ago2**. The purified microRNA fractions from rodent cultured cell line (P388D1, CHO, PC-12) were specifically detected by Urea-PAGE and silver staining (#311-03961; CLEAR STAIN Ag). Cell number of each cell line is approximately  $5 \times 10^6$ .

microRNA Isolation Kit, **Human/Mouse Ago1** (Wako Cat. #291-70201)

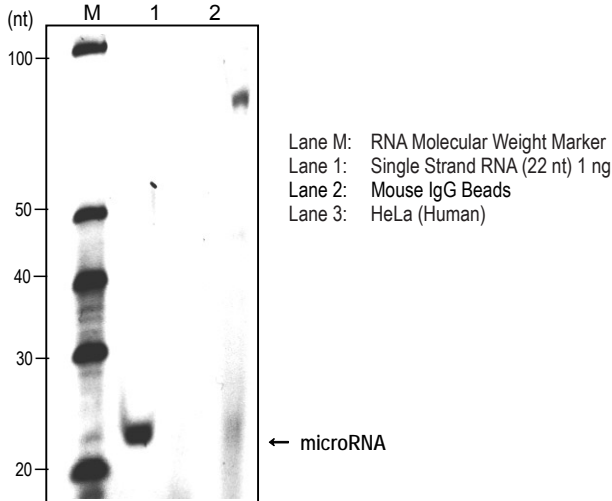


Figure: Purification of microRNA fraction by using microRNA Isolation Kit, **Human/Mouse Ago1**. The purified microRNA fraction from human cultured cell line (HeLa) was specifically detected by Urea-PAGE and silver staining (Wako Cat #311-03961; CLEAR STAIN Ag). Cell number is  $2 \times 10^7$ .

microRNA Isolation Kit, **Human Ago3** (Wako Cat. #297-70301)

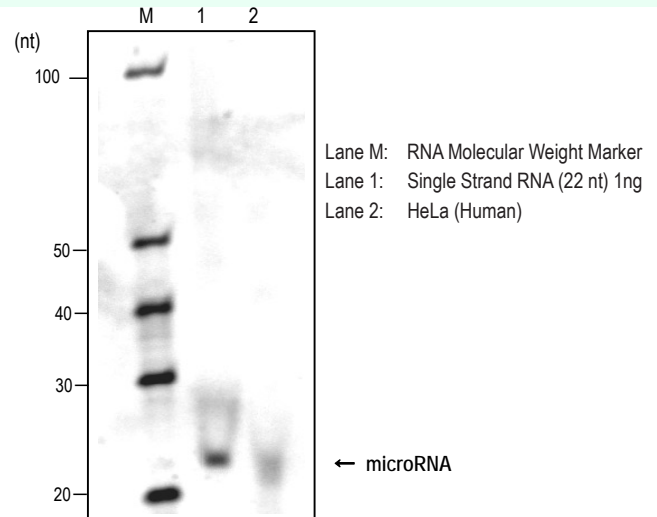


Figure: The purified microRNA fraction by using microRNA Isolation Kit, **Human Ago3**. The purified microRNA fraction from human cultured cell line (HeLa) was specifically detected by Urea-PAGE and silver staining (Wako Cat. #311-03961, CLEAR STAIN Ag). Cell number is  $5 \times 10^6$ .

■ Purification of microRNA fraction from tissues

microRNA Isolation Kit, **Human Ago2** (Wako Cat. #292-66701)

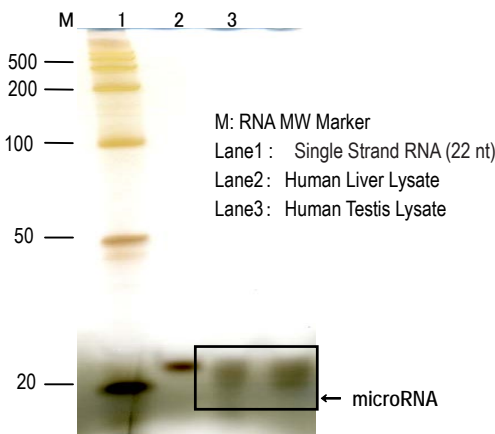


Figure: microRNA fractions, which were isolated by microRNA Isolation Kit, **Human Ago2** from human liver and testis Lysate, were specifically detected by Urea-PAGE and silver stain (Wako Cat #311-03961; CLEAR STAIN Ag) (50 mg Tissues→2 mL Total tissue lysate→Take the 1 mL→Immunoprecipitation→10  $\mu\text{L}$  Total RNA soln.→Take the 5  $\mu\text{L}$ →Urea-PAGE)

microRNA Isolation Kit, **Mouse Ago2** (Wako Cat. #292-67301)

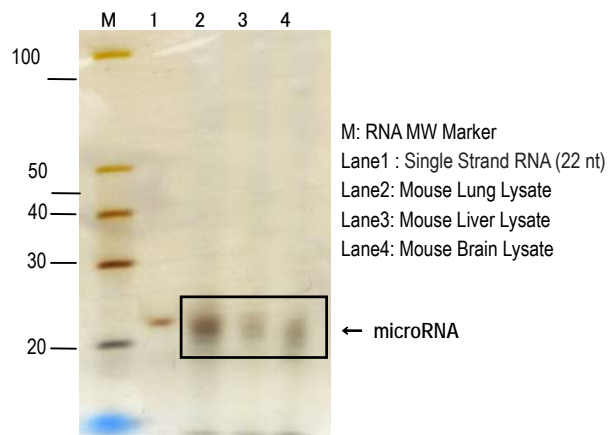
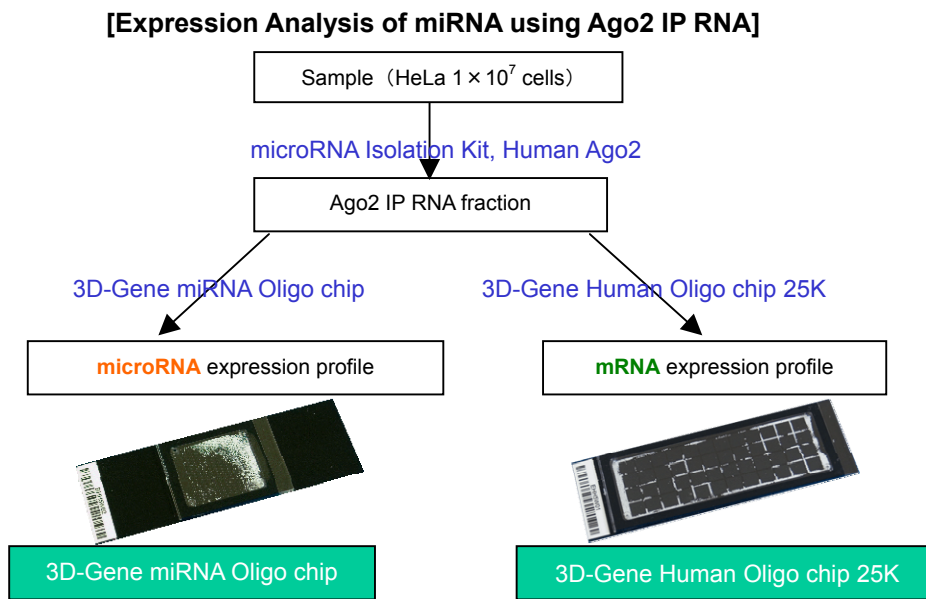


Figure: microRNA fractions, which were isolated by microRNA Isolation Kit, **Mouse Ago2** from mouse lung, liver and brain Lysate, were specifically detected by Urea-PAGE and silver stain (Wako Cat #311-03961; CLEAR STAIN Ag) (50 mg Tissues→2 mL Total tissue lysate→Take the 1 mL→Immunoprecipitation→10  $\mu\text{L}$  Total RNA soln.→Take the 5  $\mu\text{L}$ →Urea-PAGE)

■ Microarray analysis of Ago2 IP RNA fraction

Expression analysis of the miRNAs and mRNAs contained in the Ago2 IP RNA fractions can be carried out by using concomitantly the microRNA Isolation Kit, Human Ago2, and 3D-Gene miRNA Oligo chip and Human Oligo chip 25K (both available from Toray Industries, Inc.).



**[Expression analysis of microRNA using Ago2 IP RNA]**

■ Features

- 1) High-sensitivity and low-background expression analysis is possible due to highly-concentrated microRNA
- 2) Capable of exclusively profiling the microRNAs bound to RISC
- 3) Microarray analysis is possible from  $1 \times 10^6$  cells

Sample		Ago2 IP ( $1 \times 10^5$ cells)	Ago2 IP ( $1 \times 10^6$ cells)	Ago2 IP ( $1 \times 10^7$ cells)	Total RNA
Amount of RNA, average		Total IP RNA	Total IP RNA	Total IP RNA	0.5 $\mu$ g
Back ground	Average	88.41	89.18	91.75	99.18
	SD	0.73	0.70	0.80	1.35
	Baseline value (BG+2SD)	89.87	90.58	90.58	101.87
CV		0.008	0.008	0.009	0.014
Number of active spots (over BG baseline value)		312	466	522	420

↑  
*Number of Active Spot was increased.*

**[Expression analysis of mRNA using Ago2 IP RNA]**

■ Features

- 1) The signal of mRNAs in RISC is detectable from Ago2 IP RNA of  $1 \times 10^7$  cells.
- 2) Usable for target mRNA analyses, since most of the mRNAs highly concentrated in Ago2 IP RNA have the seed sequences of the miRNA detected by the miRNA array.

*(Please see the figure on the page # 7.)*

The microRNA Cloning Kit *Wako* can prepare the cDNA encoding microRNA. The cloning procedure will be completed within 1.5 days after preparation of microRNA fraction. This kit is supported by shrimp alkaline phosphatase (SAP), thermostable single strand DNA ligase (selling separately), and original modified adaptors. The cloning efficiency using this kit is improved higher than that of the conventional methods, which used bacterial alkaline phosphatase and T4 RNA ligase.

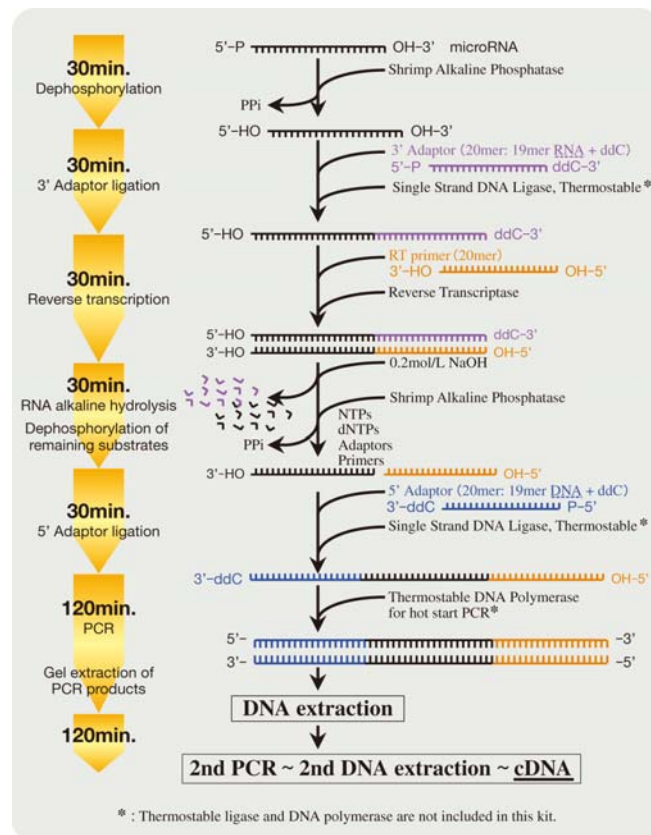
## ■ Features

- 1) High cloning efficient and accurate adaptor ligation by thermostable ligase
- 2) Suitable for cloning of microRNA forming secondary structure
- 3) Simple operation achieves forming of cDNA coding microRNA in 1.5 days

## ■ Kit Contents (for 8 reactions)

(1) SAP	16 $\mu$ L	(10) 1 mol/L Tris-HCl (pH 7.5)	160 $\mu$ L
(2) 5 $\times$ SAP Buffer	64 $\mu$ L	(11) Ethachinmate	24 $\mu$ L
(3) 40 $\times$ Ligation Buffer	16 $\mu$ L	(12) 10 mol/L Ammonium Acetate	960 $\mu$ L
(4) RNase Inhibitor	16 $\mu$ L	(13) 3' Adaptor (50 pmol/ $\mu$ L)	8 $\mu$ L
(5) 10mmol/L MnCl <sub>2</sub>	16 $\mu$ L	(14) 5' Adaptor (50 pmol/ $\mu$ L)	8 $\mu$ L
(6) Reverse Transcriptase	8 $\mu$ L	(15) RT Primer (50 pmol/ $\mu$ L)	8 $\mu$ L
(7) 10 $\times$ RT Buffer	16 $\mu$ L	(16) 5' PCR Primer (50 pmol/ $\mu$ L)	16 $\mu$ L
(8) dNTP Mixture	112 $\mu$ L	(17) 3' PCR Primer (50 pmol/ $\mu$ L)	16 $\mu$ L
(9) 0.5 mol/L EDTA (pH 8.0)	16 $\mu$ L	(18) Control RNA (30 ng/ $\mu$ L)	8 $\mu$ L

## ■ Outline of Procedure



## ■ Single Strand DNA Ligase, thermostable, recombinant, solution (Wako Cat. #298-65103; 292-65101): ※



### Features:

1. High thermal stability
2. Optimum temperature: 55 ~ 65°C
3. High ligation efficiency
4. Ligation of ssRNA, ssDNA, and ssRNA-ssDNA is applicable.

Source: *E. coli* expressed thermophilic phage TS2126 single strand DNA ligase

Appearance: 10 mmol/L Tris-HCl (pH 8.0), 50 mmol/L KCl, 0.1 mmol/L EDTA, 1 mmol/L DTT and 50 % Glycerol

Activity: Shown on each label. (approx. 10units/ $\mu$ L)

## ■ Comparison with conventional method

	Conventional method	microRNA Cloning kit <i>Wako</i>
Number of adaptor ligation steps	8	<b>3</b>
Phosphorylation of adaptors	Necessary	<b>Unnecessary</b>
Complete inactivation of Alkaline Phosphatase	Impossible	<b>Possible</b>
Apparatus for magnet beads	Necessary	<b>Unnecessary</b>
RI	Necessary	<b>Unnecessary</b>
BioAnalyzer (Agilent)	Necessary	<b>Unnecessary</b>
Adaptor ligation time	$\geq$ 8 hr	<b>2.5 hr</b>
Handling time	2.5 days	<b>1.5 days</b>
Reproducibility of adaptor ligation time	Low	<b>High</b>
Efficiency of microRNA cloning	< 10 %	<b>&gt; 70 %</b>
Detection by EtBr	Impossible	<b>Possible</b>
Cloning efficiency of secondary structured microRNA	Low (by T4 RNA Ligase)	<b>High (by Single Strand DNA Ligase, thermostable)</b>

■ High efficiency of microRNA cloning

microRNA Isolation Kit, **Human Ago2** (Wako Cat. #292-66701)

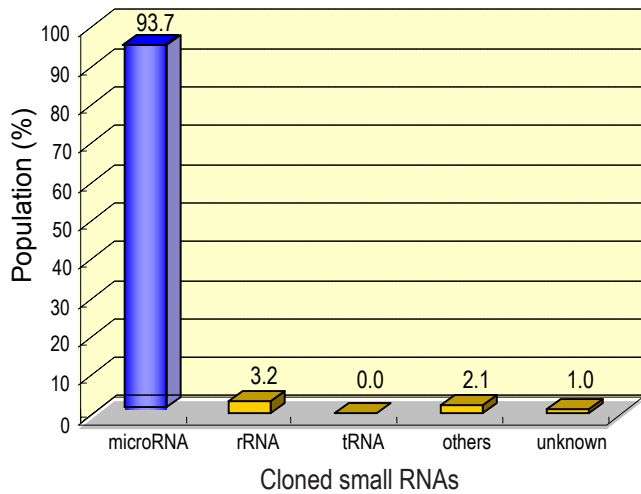


Figure: Cloning efficiency of microRNA from HeLa cell lysate. The microRNA fraction was prepared by microRNA Isolation Kit, Human Ago2 (#292-66701) from  $5 \times 10^6$  HeLa cells. The cDNA encoding microRNA was synthesized by microRNA Cloning Kit Wako (#290-66501) and was inserted into T-vector. The presence ratio of microRNA was more than 90%.

microRNA Isolation Kit, **Mouse Ago2** (Wako Cat. #292-67301)

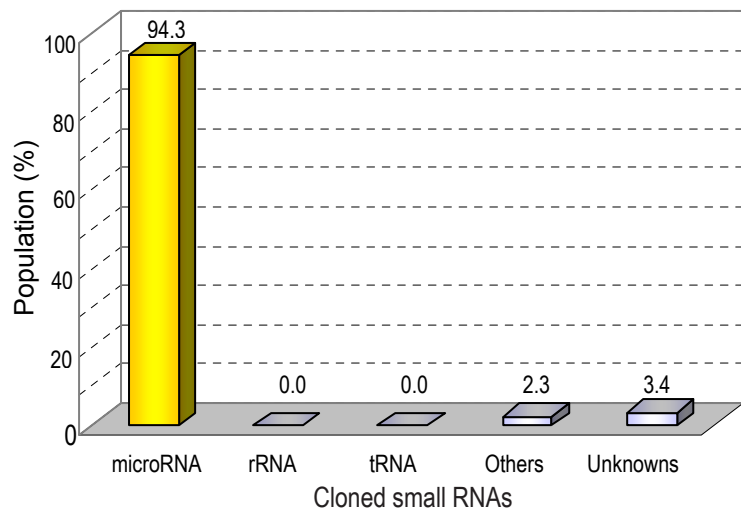


Figure: Cloning efficiency of microRNA from P388D1 cell lysate. The microRNA fraction was prepared by microRNA Isolation Kit, Mouse Ago2 (#292-67301) from  $5 \times 10^6$  P388D1 cells. The cDNA encoding microRNA was synthesized by microRNA Cloning Kit Wako (#290-66501) and was inserted into T-vector. The presence ratio of microRNA was more than 90%.

microRNA Isolation Kit, **Human/Mouse Ago1** (Wako Cat. #291-70201)

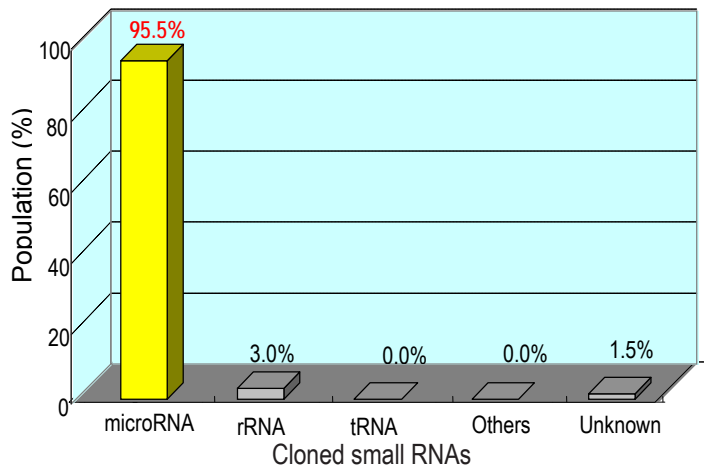


Figure: Cloning efficiency of microRNA from HeLa cell lysate. The presence ratio of microRNA was more than 90%. Others indicated cDNAs which were listed in miRBase of other organism species. Unknowns indicated cDNAs which were found in genome sequence, but not listed in miRBase.

microRNA Isolation Kit, **Human Ago3** (Wako Cat. #297-70301)

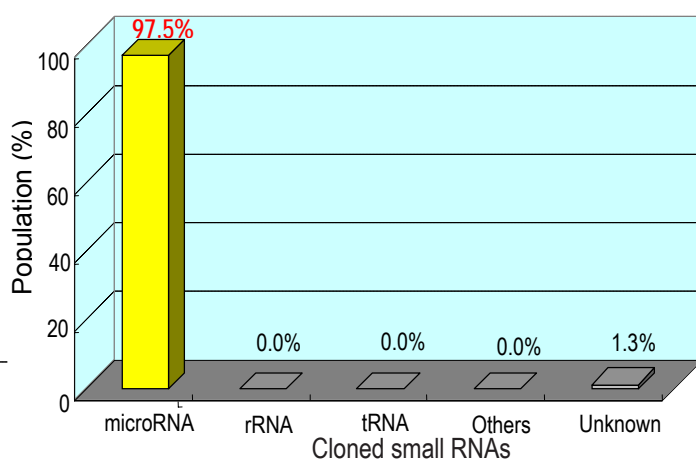


Figure: Cloning efficiency of microRNA from HeLa cell lysate. The presence ratio of microRNA was more than 90%. Others indicated cDNAs which were listed in miRBase of other organism species. Unknowns indicated cDNAs which were found in genome sequence, but not listed in miRBase.

rRNA: rRNA fragment; tRNA: tRNA fragment; Others: cDNAs which were listed in miRBase of other organism species; Unknowns: cDNAs which were found in genome sequence, but not listed in miRBase

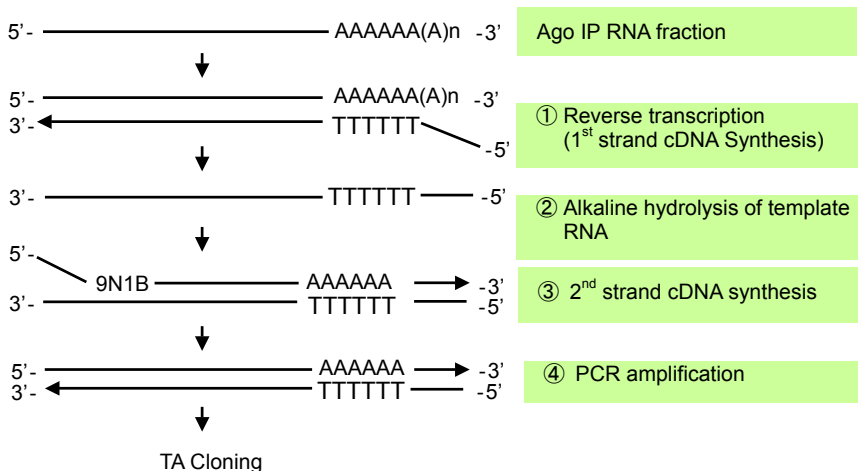
Target mRNA Cloning Kit *Wako* is used for the cDNA amplification of low population mRNA. Especially, this product will be used for the cloning of microRNA-targeted mRNA from Ago IP RNA fraction. The amplified cDNA will be the target mRNA of microRNA.



## ■ Features

- 1) Simple Protocol
- 2) Amplification of low population mRNA
- 3) Independent cDNA amplification to the length of mRNAs
- 4) Investigation of microRNA targeted mRNA

## ■ Principle



## ■ Kit Contents (10 reactions)

- (1) dT (20)-RT Primer (20 pmol/μL) 1 tube x 10 μL
- (2) 10x RT Buffer 1 tube x 20 μL
- (3) dNTP Mixture Solution (2.5 mmol/L each) 1 tube x 20 μL
- (4) RNase Inhibitor(20 U/μL) 1 tube x 10 μL
- (5) Reverse Transcriptase(200 U/μL) 1 tube x 10 μL
- (6) 0.5 mol/L EDTA Solution 1 tube x 20 μL
- (7) 1 mol/L Tris-HCl Solution 1 tube x 200 μL
- (8) Ethachinmate 1 tube x 30 μL
- (9) 10 mol/L Ammonium Acetate Solution 1 tube x 480 μL
- (10) 2nd Strand Synthesis Primer (20 pmol/μL) 1 tube x 10 μL
- (11) 5' PCR Primer(20 pmol/μL) 1 tube x 10 μL
- (12) 3' PCR Primer(20 pmol/μL) 1 tube x 10 μL
- (13) 5' Colony PCR Primer (5 pmol/μL) 1 tube x 960 μL
- (14) 3' Colony PCR Primer (5 pmol/μL) 1 tube x 960 μL

## ■ Identification of amplified cDNA and categories of cDNAs isolated from HeLa cells

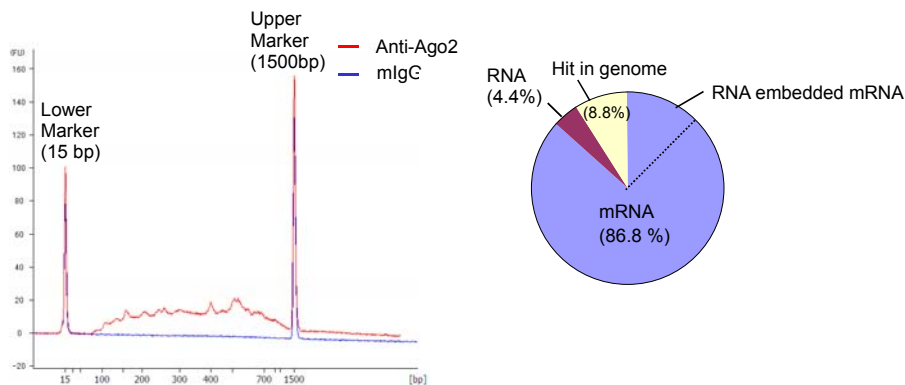


Figure: Size distribution of synthesized cDNA and categories of cDNAs isolated from HeLa cells.

### [Advantages of Target mRNA Cloning Kit *Wako*]

There is a possibility that RNA molecules other than mRNAs, which can not be identified by the RISC-trapped microarray, can be identified by cloning of the cDNAs synthesized using this kit. According to *BMC Research Notes*, **2**, 169 (2009), free Alu RNAs besides mRNAs are contained in the RISC.

### [Additional materials required in addition to Target mRNA Cloning Kit *Wako*]

(1) **THUNDER Taq Gold DNA Polymerase (Mg<sup>2+</sup> free buffer)** (Wako Cat. #317-07081; 250 U): Thermostable DNA polymerase for PCR. We recommend THUNDER Taq Gold DNA Polymerase (Mg<sup>2+</sup> free buffer) for suppression of non-specific amplification. It is used for hot start PCR and activated by 10 minute incubation at 95°C before PCR cycle. DNA fragments, which were amplified by THUNDER Taq Gold DNA Polymerase (Mg<sup>2+</sup> free buffer), are 3' protruding end of adenine base. These fragments therefore are able to directly insert into T-vectors.

(2) **PCR Purification Kit *Wako*** (Wako Cat. No. 298-67901)

PCR Purification Kit *Wako* is used for removal of primers, primer dimers, dNTPs, enzymes such as DNA polymerase and restriction enzyme from reaction buffer of various molecular biology experiments in only 10 minutes after DNA synthesis by RT-PCR or PCR.

# Anti Ago2, Monoclonal Antibodies

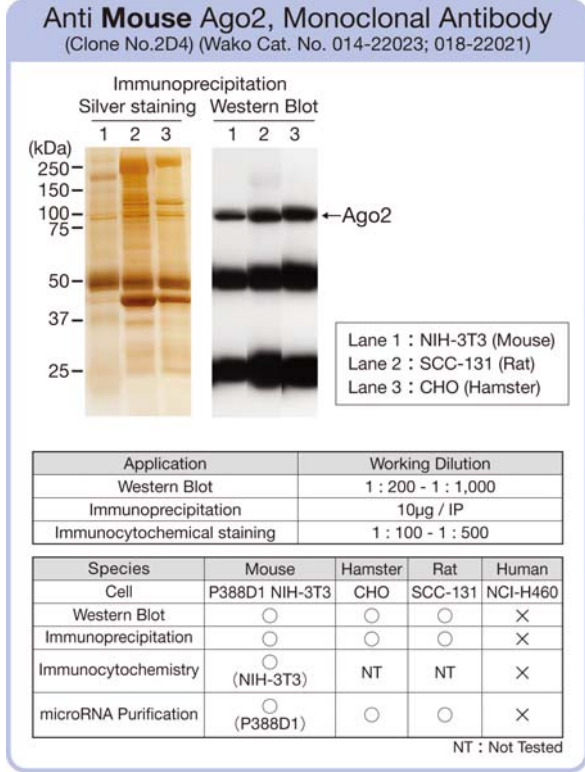
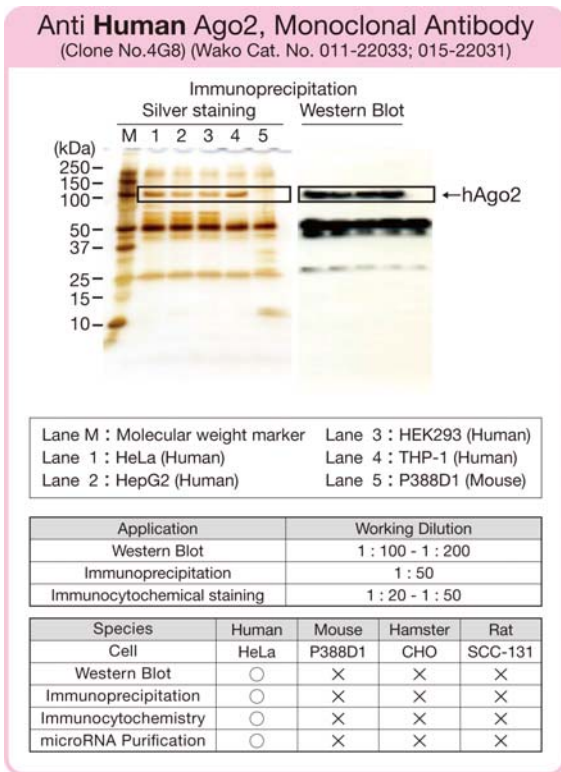
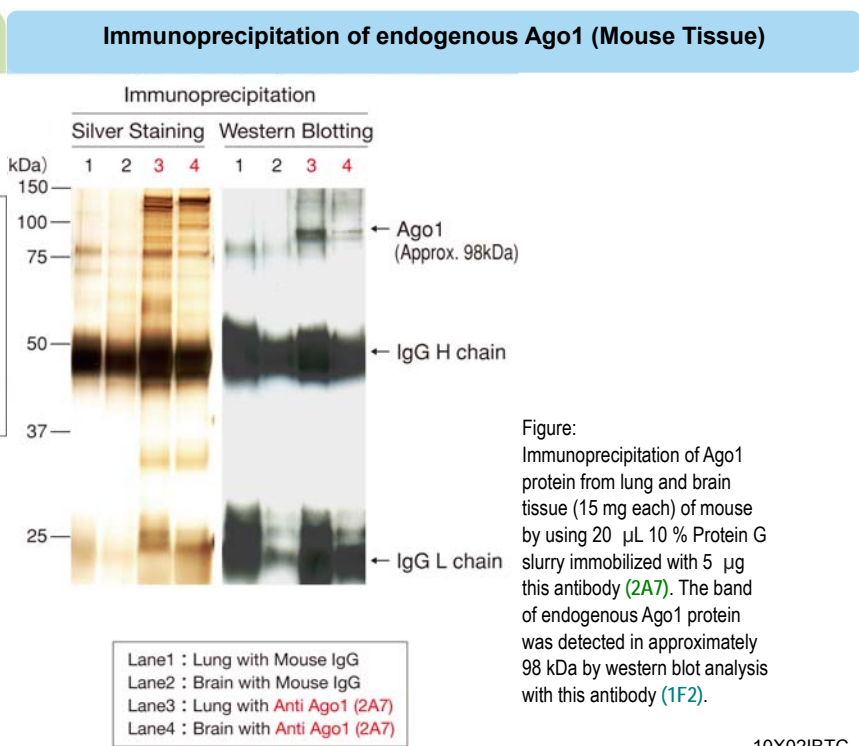
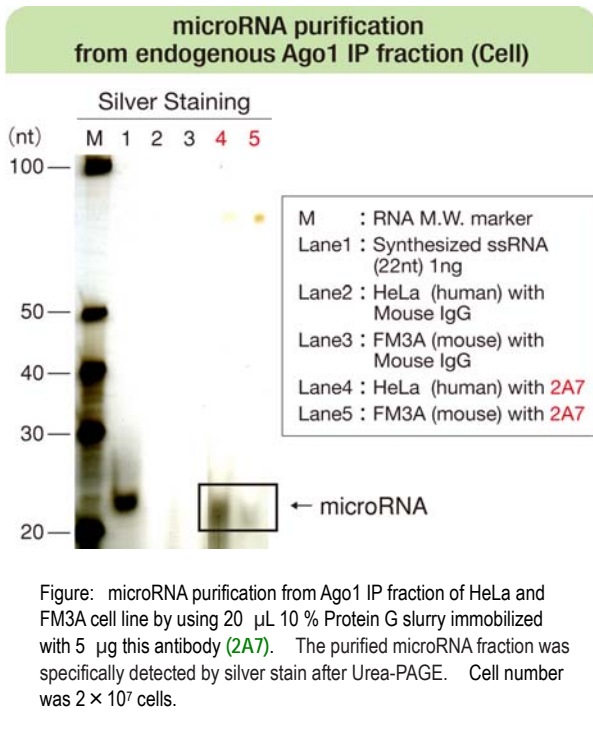


Figure: Immunoprecipitation of human Ago2 protein from human cultured cell lines (HeLa, HepG2, HEK293, THP-1) and mouse cultured cell line (P388D1) by using 20 µL 10% Protein G slurry immobilized with 10 µg of this antibody (4G8). The bands of hAgo2 protein were detected in approximately 100 kDa by using silver staining and western blot. Cell number was 5×10<sup>6</sup> cells.

Figure: Immunoprecipitation of Ago2 protein from NIH-3T3 (Mouse), SCC-131(Rat) and CHO (Hamster) cell line by using 20 µL of 10 % Protein G slurry immobilized with 5 µg of this antibody (2D4). The bands of endogenous Ago2 protein were detected in approximately 100 kDa by using silver staining and western blot. The 1/1,000 diluted antibody was used as the 1st antibody for western blot. Cell number was 5×10<sup>6</sup> cells.

# Anti Ago1, Monoclonal Antibodies

(Clone #1F2: Wako Cat. #018-22401; 2A7: #015-22411)



10X02IBTC

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Online Catalog: [www.e-reagent.com](http://www.e-reagent.com)

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[www.wakousa.com](http://www.wakousa.com)

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Head Office (Richmond, VA):

TEL: 1-804-714-1920; FAX: 1-804-271-7791

Los Angeles Sales Office (Irvine, CA):

TEL: 1-949-679-1700; FAX: 1-949-679-1701

Boston Sales Office (Cambridge, MA):

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