# Certificate of Analysis



# Mate & Plate™ Library - Human Brain (Normalized)

Catalog No. Lot Number 630486 1707659A

# **Description**

This yeast two-hybrid library was constructed from mRNA isolated from human brain tissue and transformed into yeast strain Y187. The cDNA was normalized prior to library construction to reduce the copy number of abundant cDNAs derived from highly represented mRNAs, thereby increasing the representation of low copy number transcripts. The normalization process combines a Duplex-Specific Nuclease (DSN) treatment and SMART® technology, reduces the number of clones that must be screened in your yeast two-hybrid assay, and facilitates the identification and characterization of novel protein-protein interactions.

The library was transformed into yeast strain Y187 and can be readily mated to a *MATa* GAL4 reporter strain, such as AH109 or Y2HGold (Clontechniques, 2008), for screening.

# **Package Contents**

- 5 x 1.0 ml Mate & Plate Library Human Brain (Normalized)
- 1 x 1.0 ml Mate & Plate Library Control (pGADT7-T in Y187)

## **Storage Conditions**

- Store all components at -70°C
- Do not refreeze

#### **Shelf Life**

• 1 year from date of receipt under proper storage conditions.

#### mRNA Source

 Mixture of poly A+ RNA isolated from normal, whole brains from 8 male Caucasians, ages: 43–66; cause of death: sudden death.

## **Cloning Vector**

• pGADT7-RecAB

#### **Cloning Site**

• Sfi I A/Sfi I B

#### **Priming Method**

• Sfi I (dT)<sub>30</sub> primed

## **Yeast Genotype**

Y187: MATa, ura3-52, his3-200, ade2-101, trp1-901, leu2-3, 112, gal4Δ, gal80Δ, met—, URA3 :: GAL1<sub>UAS</sub>-GAL1<sub>TATA</sub>-LacZ, MEL1

Mate & Plate<sup>TM</sup> Library - Human Brain (Normalized)

## **Shipping Conditions**

• Dry ice  $(-70^{\circ}\text{C})$ 

#### **Product Documents**

Documents for our products are available for download at <u>takarabio.com/manuals</u> The following documents apply to this product:

- Matchmaker® Gold Yeast Two-Hybrid System User Manual (PT4084-1)
- pGADT7-RecAB Vector Information (PT3718-5)

#### References

Franz, O., Bruchhaus, I. & Roeder, T. Verification of differential gene transcription using virtual northern blotting. *Nucleic Acids Res.* **27**, e3 (1999).

Pretransformed Mate & Plate Libraries. Clontechniques 24, 26–27 (2008).

Shagin, D. A. *et al.* A novel method for SNP detection using a new duplex-specific nuclease from crab hepatopancreas. *Genome Res.* **12**, 1935–1942 (2002).

Zhulidov, P. A. *et al.* Simple cDNA normalization using kamchatka crab duplex-specific nuclease. *Nucleic Acids Res.* **32**, e37 (2004).

# **Quality Control Data**

## 1. Quality Control Data

Test	Result
Titer (yeast colonies)	$>5 \text{ x } 10^7 \text{ cfu/ml}$
Number of independent clones	$3.2 \times 10^6$
Average cDNA size	<u>1.40 kb</u>
cDNA size range*	1.0-1.8 kb

<sup>\*</sup>the cDNA was size-selected by excision from an agarose gel prior to cloning

# 2. Quality Control Data for the Pretransformed Library in Yeast

Library Insert Size Screening

15 yeast colonies were randomly picked and screened by PCR using the Matchmaker AD LD-Insert Screening Amplimer Set (Cat. No. 630433).

(082917) Page 2 of 3

15 of 15 colonies contained inserts as determined by PCR.

LANE M. 1 kb DNA ladder

10.

11.

12.

13.

14.

15.

1. 1.16 kb 2. 1.47 kb 3. 1.03 kb 4. 1.31 kb 5. 1.86 kb 6. 1.31 kb 7. 1.23 kb 8. 1.75 kb 9. 1.13 kb

1.27 kb

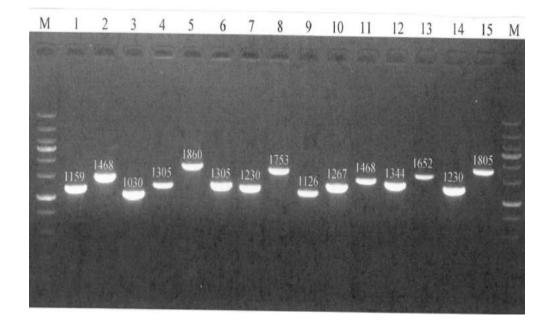
1.47 kb

1.34 kb

1.65 kb

1.23 kb

1.81 kb



#### 3. cDNA Normalization

cDNA generated using SMART technology was normalized using Duplex-Specific Nuclease (DSN) normalization (Zhulidov et al., 2004; Shagin et al., 2002). Prior to cloning, the efficiency of normalization was assessed by virtual Northern blot analysis (Franz et al., 1999) comparing the abundance of  $\beta$ -actin and GAPDH in normalized and non-normalized human brain cDNA.

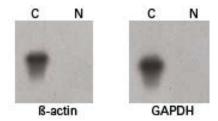


Figure 1. DSN-Normalization reduces the amount of highly abundant transcripts. Normalized (Lanes N) and non-normalized (Lanes C) human brain cDNA samples (PCR products) were electrophoresed on a 1.5% agarose gel and transferred to Hybond-N membrane. PCR-amplified probes of β-actin and GAPDH were labeled with  $^{32}$ P-dATP and hybridized to the membrane. GenBank Accession numbers: β-actin, NM\_001101 and GAPDH, NM\_002046.

It is certified that this product meets the above specifications, as reviewed and approved by the Quality Department.

(082917) Page 3 of 3



# Mate & Plate<sup>TM</sup> Library - Human Brain (Normalized)

# CATALOG NO.

630486

#### NOTICE TO PURCHASER:

Our products are to be used for research purposes only. They may not be used for any other purpose, including, but not limited to, use in drugs, in vitro diagnostic purposes, therapeutics, or in humans. Our products may not be transferred to third parties, resold, modified for resale, or used to manufacture commercial products or to provide a service to third parties without prior written approval of Takara Bio USA, Inc.

Your use of this product is also subject to compliance with the licensing requirements listed below and described on the product's web page at <a href="http://www.takarabio.com">http://www.takarabio.com</a>. It is your responsibility to review, understand and adhere to any restrictions imposed by these statements.

#### **TRADEMARKS:**

#### ©2017 Takara Bio Inc. All Rights Reserved.

All trademarks are the property of Takara Bio Inc. or its affiliate(s) in the U.S. and/or other countries or their respective owners. Certain trademarks may not be registered in all jurisdictions.

Takara Bio USA, Inc.

1290 Terra Bella Avenue, Mountain View, CA 94043, USA U.S. Technical Support: techUS@takarabio.com

United States/Canada Asia Pacific **Europe** Japan 800.662.2566 +1.650.919.7300 +81.(0)77.565.6999

+33.(0)1.3904.6880

8/31/2017