# Certificate of Analysis



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

 Catalog No.
 Amount
 Lot Number

 630480
 5 x 1 ml
 2312163A

# **Description**

This yeast two-hybrid library was constructed from human cDNA that had been previously normalized to preferentially remove abundant cDNAs derived from high-copy-number mRNAs. The normalization process incorporates a Duplex-Specific Nuclease (DSN) treatment and SMART<sup>TM</sup> (Switching Mechanism at 5' end of RNA Template) technology and increases the representation of low-copy-number transcripts in the library. This reduces the number of clones that must be screened to identify positive interactions and facilitates the identification and characterization of novel protein-protein interactions. A Universal human cDNA library transformed into yeast strain Y187. The library can be readily mated to a MATa GAL4 reporter strain, such as AH109 or Y2HGold.

### **Package Contents**

- 5 x 1.0 ml Mate & Plate Library Universal Human (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.

# **Expiration Date**

• JAN. 02, 2027

#### **mRNA Source**

 Mixture of Poly A<sup>+</sup> RNAs from a collection of adult human tissues chosen to represent a broad range of expressed genes. Both male and female donors are represented. Modeled after our Human Universal Reference Total RNA (Cat. No. 636538).

#### **Cloning Vector**

pGADT7-RecAB

## **Cloning Site**

• Sfi I A/Sfi I B

### **Priming Method**

• Sfi I (dT)<sub>30</sub> primed

#### **Yeast Genotype**

• (Y187): MATα, 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

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## **Shipping Conditions**

• Dry ice

#### **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 User Manual
- pGADT7-RecAB Vector Information
- pGADT7-RecAB Vector Sequence in GenBank Format

#### References

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

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)	$>5 \times 10^{7} \text{ cfu/ml}$
Number of independent clones	$5.91 \times 10^6$
Average cDNA size	<u>1.25 kb</u>
cDNA size range*	0.8 - 2.0  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).

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Lane M. 1 kb DNA ladder

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15 of 15 colonies contained inserts as determined by PCR.

1. 1.10 kb 2. 2.02 kb 10 11 12 13 14 15 3. 1.83 kb 4. 1.20 kb 5. 1.07 kb 6. 1.20 kb 3000 7. 0.97 kb 2500 8. 0.80 kb 2000 1500 9. 1.15 kb 1000 10. 1.18 kb 750 11. 1.11 kb 12. 1.40 kb

# 3. cDNA Normalization

13. 1.37 kb 14. 1.24 kb 15. 1.13 kb

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

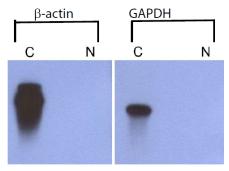


Figure 1. DSN-Normalization reduces the amount of highly abundant transcripts. Normalized (Lanes N) and non-normalized (Lanes C) Human Universal cDNA samples (PCR products) were electrophoresed on a 1.5% agarose gel and transferred to Hybond-N membrane. PCR-amplified probes of GAPDH and β-actin were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

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

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# CATALOG NO.

630480

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