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PRODUCT: Mate & Plate[™] Library - Universal Drosophila (Normalized)

CATALOG No. 630485

LOT NUMBER: 1503559A

STORAGE CONDITIONS

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

SHELF LIFE

1 year from date of receipt under proper storage conditions.

SHIPPING CONDITIONS

Dry ice $(-70^{\circ}C)$.

mRNA SOURCE

Equal quantities of poly A+ RNA isolated from embryo (~20 hr), larval, and adult stage Drosophila melanogaster.

CLONING VECTOR: pGADT7-RecAB

CLONING SITE: SfilA/SfilB

PRIMING METHOD: Sfil (dT)₃₀ primed

YEAST GENOTYPE (Y187): MATα, ura3-

52, his3-200, ade2-101, trp1-901, leu2-3, 112, gal4D, gal80D, met–, URA3 :: GAL1_{UAS}-GAL1_{TATA}-LacZ, MEL1

DESCRIPTION

This yeast two-hybrid library was constructed from mRNA isolated from Drosophila melanogaster 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 (1), for screening.

PACKAGE CONTENTS

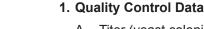
- 5 x 1.0 ml Mate & Plate Library Universal Drosophila (Normalized)
- 1 x 1.0 ml Mate & Plate Library Control (pGADT7-T in Y187)

OTHER

- Matchmaker[™] Gold Yeast Two-Hybrid User Manual (PT4084-1)
- pGADT7-RecAB Vector Information (PT3718-5)

FOR RESEARCH USE ONLY

QUALITY CONTROL DATA



- A. Titer (yeast colonies): $\geq 5 \times 10^7$ cfu/ml
- B. Number of
 - independent clones: 3.2×10^6
- C. Average cDNA size: <u>1.5 kb</u>

D. cDNA size range: 0.7–2.5 kb

(The cDNA was size-selected by excision from an agarose gel prior to cloning)

Continued on back of page.

Certificate of Analysis



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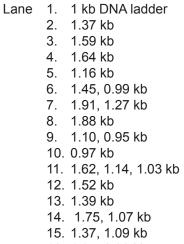
QUALITY CONTROL DATA continued

2. Quality Control Data for the Pretransformed Library in Yeast

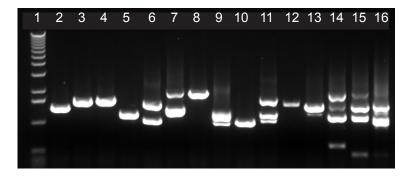
Library Insert Size Screening

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

15 of 15 colonies contained inserts as determined by PCR.



16. 1.37, 0.98 kb



3. cDNA Normalization

cDNA generated using SMART technology was normalized using Duplex-Specific Nuclease (DSN) normalization (2, 3). Prior to cloning, the efficiency of normalization was assessed by virtual Northern blot analysis (4) comparing the abundance of β -actin in normalized and non-normalized Drosophila melanogaster cDNA.

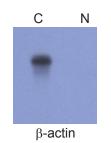


Figure 1. DSN-Normalization reduces the amount of highly abundant transcripts. Normalized (Lane N) and non-normalized (Lane C) Drosophila Universal cDNA samples (PCR products) were electrophoresed on a 1.5% agarose gel and transferred to Hybond-N membrane. A PCR-amplified probe for β -actin was labeled with 32P-dATP and hybridized to the membrane. GenBank Accession number: β -actin, NM 001101.

REFERENCES

- 1. Pretransformed Mate & Plate[™] Libraries (January 2008) Clontechniques XXIV(1):26–27.
- 2. Zhulidov, P. A., et al. (2004) Nucleic Acids Res. 32:e37.
- 3. Shagin, D. A., et al. (2002) Genom. Res. **12**:1942–1953.
- 4. Franz, O., et al. (1999) Nucleic Acids Res. **27**:e3.



Mate & PlateTM Library - Universal Drosophila (Normalized)

CATALOG NO.

630485

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