

# Product datasheet for TR500505

## Dbt Mouse shRNA Plasmid (Locus ID 13171)

## **Product data:**

#### OriGene Technologies, Inc.

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| Product Type:                | shRNA Plasmids   |
|------------------------------|--|
| Product Name:                | Dbt Mouse shRNA Plasmid (Locus ID 13171)   |
| Locus ID:                    | 13171  |
| Synonyms:                    | BCKAD-E2; D3Wsu60e   |
| Vector:                      | pRS (TR20003)  |
| E. coli Selection:           | Ampicillin   |
| Mammalian Cell<br>Selection: | Puromycin  |
| Format:                      | Retroviral plasmids  |
| Components:                  | Dbt - Mouse, 4 unique 29mer shRNA constructs in retroviral untagged vector(Gene ID =<br>13171). 5µg purified plasmid DNA per construct<br>29-mer scrambled shRNA cassette in pRS Vector, TR30012, included for free.   |
| RefSeq:                      | <u>BC055890, NM 010022, NM 001357344, NM 010022.1, NM 010022.2, NM 010022.3</u>  |
| UniProt ID:                  | <u>P53395</u>  |
| Summary:                     | The branched-chain alpha-keto dehydrogenase complex catalyzes the overall conversion of<br>alpha-keto acids to acyl-CoA and CO(2). It contains multiple copies of three enzymatic<br>components: branched-chain alpha-keto acid decarboxylase (E1), lipoamide acyltransferase<br>(E2) and lipoamide dehydrogenase (E3). Within this complex, the catalytic function of this<br>enzyme is to accept, and to transfer to coenzyme A, acyl groups that are generated by the<br>branched-chain alpha-keto acid decarboxylase component.[UniProtKB/Swiss-Prot Function] |
| shRNA Design:                | These shRNA constructs were designed against multiple splice variants at this gene locus. To be certain that your variant of interest is targeted, please contact <u>techsupport@origene.com</u> . If you need a special design or shRNA sequence, please utilize our <u>custom shRNA service</u> .  |



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### **CRIGENE** Dbt Mouse shRNA Plasmid (Locus ID 13171) – TR500505

Performance Guaranteed: OriGene guarantees that the sequences in the shRNA expression cassettes are verified to correspond to the target gene with 100% identity. One of the four constructs at minimum are guaranteed to produce 70% or more gene expression knock-down provided a minimum transfection efficiency of 80% is achieved. Western Blot data is recommended over qPCR to evaluate the silencing effect of the shRNA constructs 72 hrs post transfection. To properly assess knockdown, the gene expression level from the included scramble control vector must be used in comparison with the target-specific shRNA transfected samples.

For non-conforming shRNA, requests for replacement product must be made within ninety (90) days from the date of delivery of the shRNA kit. To arrange for a free replacement with newly designed constructs, please contact Technical Services at techsupport@origene.com. Please provide your data indicating the transfection efficiency and measurement of gene expression knockdown compared to the scrambled shRNA control (Western Blot data preferred).

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