

1 60 120 180 240 300

Descriptions

Legend for links to other resources: [U](#) UniGene [E](#) GEO [G](#) Gene [S](#) Structure [M](#) Map Viewer [P](#) PubChem BioAssay

Sequences producing significant alignments:

Accession	Description	Max score	Total score	Query coverage	E value	Max ident	Li
NT_167190.1	Homo sapiens chromosome 11 genomic contig, GRCh37.p2 r	446	446	71%	4e-123	100%	
NW_001838028.2	Homo sapiens chromosome 11 genomic contig, alternate as	446	446	71%	4e-123	100%	

Alignments

Select All [Get selected sequences](#) [Distance tree of results](#)

```
> ref|NT\_167190.1 D Homo sapiens chromosome 11 genomic contig, GRCh37.p2 reference
primary assembly
Length=4159379

Features in this part of subject sequence:
  tensurin-4

Score = 446 bits (241), Expect = 4e-123
Identities = 241/241 (100%), Gaps = 0/241 (0%)
Strand=Plus/Minus

Query  96      TCTCCCTCACCATAGTCGATGGGCTCAGTTCCTCGAAGCATAGTATGGTGGACAGTCA 155
      |||
Sbjct  23763477 TCTCCCTCACCATAGTCGATGGGCTCAGTTCCTCGAAGCATAGTATGGTGGACAGTCA 23763418

Query  156     AGGGCTCTGGTCTTGTGTACACTTTTATCAGCCAGCAAAAATATCCAGAAGATTCCTA 215
      |||
Sbjct  23763417 AGGGCTCTGGTCTTGTGTACACTTTTATCAGCCAGCAAAAATATCCAGAAGATTCCTA 23763358

Query  216     TGGGAGGAGACTCAGCCTGAAGCATCAGCATTGAGCAGCCAGGCTGCCACAGCCTTGAC 275
      |||
Sbjct  23763357 TGGGAGGAGACTCAGCCTGAAGCATCAGCATTGAGCAGCCAGGCTGCCACAGCCTTGAC 23763298

Query  276     AGATTGATCTGCAGCCTGGAGCTCAGCCTCCCTGGTGTCTGAAGGAGCTCAACAGCC 335
      |||
Sbjct  23763297 AGATTGATCTGCAGCCTGGAGCTCAGCCTCCCTGGTGTCTGAAGGAGCTCAACAGCC 23763238

Query  336      A 336
      |
Sbjct  23763237 A 23763237

> ref|NW\_001838028.2 D Homo sapiens chromosome 11 genomic contig, alternate assembly
HuRef SCAF_1103279187758, whole genome shotgun sequence
Length=16912879

Features in this part of subject sequence:
  tensurin-4

Score = 446 bits (241), Expect = 4e-123
Identities = 241/241 (100%), Gaps = 0/241 (0%)
Strand=Plus/Plus
```