

Contents lists available at ScienceDirect

### Data in Brief





#### Data Article

# Data on four apoptosis-related genes in the colonial tunicate *Botryllus schlosseri*



Nicola Franchi, Francesca Ballin, Lucia Manni\*, Filippo Schiavon, Loriano Ballarin

Department of Biology, University of Padova, Italy

#### ARTICLE INFO

Article history:
Received 13 April 2016
Received in revised form
3 May 2016
Accepted 11 May 2016
Available online 20 May 2016

Keywords:
Sequence databases
Alignments
BLAST analysis
Botryllus schlosseri
Apoptosis-related transcripts
Bax
AIF1
PARP1

#### ABSTRACT

The data described are related to the article entitled "Recurrent phagocytosis-induced apoptosis in the cyclical generation change of the compound ascidian *Botryllus schlosseri*" (Franchi et al., 2016) [1]. Four apoptosis-related genes, showing high similarity with mammalian Bax (a member of the Bcl-2 protein family), AIF1 (apoptosis-inducing factor-1), PARP1 (poly ADP ribose polymerase-1) and IAP7 (inhibitor of apoptosis-7) were identified from the analysis of the trascriptome of *B. schlosseri*. They were named BsBax, BsAIF1, BsPARP1 and BsIAP7. Here, their deduced amino acid sequence were compared with known sequences of orthologous genes from other deuterostome species together with a study of their identity/similarity.

© 2016 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY license (http://creativecommons.org/licenses/by/4.0/).

#### Specifications Table

Subject area Biology

More specific sub- Developmental Biology

ject area

IAP7

Type of data Tables, figures

DOI of original article: http://dx.doi.org/10.1016/j.dci.2016.04.011

E-mail address: lucia.manni@unipd.it (L. Manni).

<sup>\*</sup> Corresponding author at: Department of Biology, University of Padova via U. Bassi 58/B 35131 Padova Italy. Tel.: +39 049 8276252; fax: +39 049 8276199.

How data was Bioinformatic analysis, RACE

acquired

Data format Raw and analysed data

Experimental The partial transcripts present in the transcriptome, identified by BLAST analysis,

were elongated through 5'and 3' RACE according to the 2nd generation of 5'/3'

RACE kit

Experimental Analysis with BLAST, LALIGN, SMART, Clustal Omega

features

factors

Data source Padova, Italy

location

Data accessibility Data are available in this article and at GenBank via accession numbers Gen-

Bank: KU948200 for BsBAX, GenBank: KU948201 for BsPARP1, GenBank:

KU948202 for BsAIF1, GenBank: KU948203 for BsIAP7.

#### Value of the data

- The data provide the full-length sequences of four apoptosis-related transcripts from the colonial ascidian *B. schlosseri* useful to study the phylogeny trees of the corresponding proteins in chordates.
- From the data, the protein primary structures can be deduced and, from that, three-dimensional models can be obtained, useful to compare the domain organization of the corresponding chordate proteins.
- Expression studies, exploiting the present data, can contribute to elucidate the dynamics of the
  cyclical apoptosis, which characterizes the colonial blastogenetic cycle of the ascidian B. schlosseri.

#### 1. Data

The data reported include supporting information to the phylogenetic analyses of Franchi et al. [1]. They consist of transcript sequences, sequence alignments and comparisons of four apoptosis-related genes identified in the recently-obtained transcriptome of *B. schlosseri* [2]. The sequences show high similarity with mammalian transcripts for Bax, AIF1, PARP1 and IAP7 and were named BsBax, BsAIF1, BsPARP1 and BsIAP7, respectively. The expression of these genes was studied further in the above-reported paper [1].

#### 2. Experimental design, materials and methods

Amplification and cloning of transcripts for BsBax, BsAIF1, BsPARP1 and BsIAP7 was achieved with specific primers designed on sequences found in our collection of transcriptomes [2]. In order to verify and complete the full length cDNA, PCR reactions were carried out with a denaturing step at 94 °C for 2 min, 40 cycles of 30 s at 94 °C, 40 s at 60 °C and 90 s at 72 °C, and a final extension at 72 °C for 10 min. Amplicons were separated using 1.5% agarose gel, purified, cloned and sequenced. The partial transcripts were elongated through 5′ and 3′ RACE according to the 2nd generation of 5′/3′ RACE kit (Roche). Supplementary Table 1 reports the specific primers used for amplicons production and their elongation through 5′- and 3′-RACE and for the *in situ* hybridisation experiments reported in [1].

The sequences from GenBank, reported in Supplementary Tables 2–5, were used for alignments and sequence comparisons with the sequences of BsBax, BsAIF1, BsPARP1, BsIAP7, respectively. The

## Download English Version:

# https://daneshyari.com/en/article/174653

Download Persian Version:

https://daneshyari.com/article/174653

<u>Daneshyari.com</u>