CIPRO 2.5: *Ciona intestinalis* Protein Database - a unique integrated repository of large-scale omics data, bioinformatic analyses, and curated annotation, with ability for user rating and comments

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From Beyond the Genome: The true gene count, human evolution and disease genomics
Boston, MA, USA. 11-13 October 2010

CIPRO database (http://cipro.ibio.jp/2.5) is an integrated protein database for a tunicate species *Ciona intestinalis* that is part of the Urochordata. Although CIPRO provides proteomic and transcriptomic data on a single species, the animal is considered unique in the evolutionary tree, representing a possible origin of the vertebrates and is a good model for understanding chordate evolution, including human evolution. Furthermore, *C. intestinalis* has been one of the favorites of developmental biologists; therefore, a lot of amount of accumulated knowledge on its development, morphology, in addition to the recent genome sequence and gene expression data exists. The CIPRO database aims to collect published data and to present unique information, including the unpublished transcriptomic and proteomic data and human curated annotation, for the use of researchers in biology and bioinformatics.

(i) Original experimental data Unpublished experimental data, including 2D-PAGE with the identified protein spots by protein mass fingerprint (PMF) MS analysis, expressions or localizations of protein and RNA across developmental stages and tissues, altogether summarized in a single chart for the comparison among status and methods. RNA expressions are observed by microarray and EST. Each protein is linked to an independent Ascidian Proteome Database summarizing large-scale MS-based proteomic analyses.  
(ii) Whole *Ciona intestinalis* proteome database Proteins across gene models are presented: all protein models derived from published gene models are incorporated, including Kyoto model (KG), KH (successor of KG model), PROCITS, JGI's versions 1 and 2, and Ensembl (version 58.2) are incorporated. Identical sequences across gene models are shown.  
(iii) Original comprehensive user-friendly interfaces Bioinformatic analyses and prediction results are summarized in pictures for grasp at a glance: homology search, cytolocalization, secondary structure prediction combined with modification sites, such include phosphorylation and three-dimensional structures.
(iv) **Comparative analysis data for disease association**
Comparison with human genome: map location of human homologues is graphically shown with associated disease information. Comparative data for other model organisms are also included.

(v) **Community-wide curation capability opened to users**
To facilitate progressive improvement of annotation by visited users, users can place additional annotation for the protein name and/or comments, which will be subjected to rating by the followed data viewers. To aid curation by wide community, information for literature and essence of matched motif patterns and other related protein information are shown with the links.

(vi) **Useful search facilities**
Various search methods are provided including blast homology, free text, partial sequence, protein mass fragment, and cross item searches.

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Published: 11 October 2010

doi:10.1186/gb-2010-11-S1-P11
Cite this article as: Endo et al. CIPRO 2.5: Ciona intestinalis Protein Database - a unique integrated repository of large-scale omics data, bioinformatic analyses, and curated annotation, with ability for user rating and comments. Genome Biology 2010 11(Suppl 1):P11.