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Organ/tissue expression patterns based on UniGene EST frequency calculations and Cytomer® Alexander Kel, Ellen Fricke, TRANSPATH Team

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We made an estimation of the gene expression levels in different tissues based on the UniGene database (release of March, 2003). The following steps were performed:

- 1. For all UniGene clusters (**G** is the total number of clusters for human genes), we collected all of the information about libraries that was used for sequencing the ESTs included in the clusters.
- 2. From the list of libraries we excluded all cancer-related libraries. Only the libraries of normal tissues and organs were considered.
- 3. All tissue and organ names used in the selected list of libraries were linked to corresponding terms in the Cytomer[®] database [1]. All libraries linked to the same term were grouped into "organ library groups" (e.g. liver library group). *P* is the total number of formed organ library groups.
- 4. Compute n_{ij} which is the number of ESTs of a gene i (Unigene cluster) linked to a organ library group j.

5. Compute
$$a_{ij} = \frac{n_{ij}}{\sum_{i=1,G} n_{ij}}$$
 - an "abundancy score" of each gene in each group.
$$\overline{a_{i\bullet}} = \frac{\sum_{j=1,P} a_{ij}}{P}$$
 - an average "abundancy score" of gene *i*. And
$$\frac{\sum_{i=1,P} a_{ij}}{P}$$

$$sigma(\overline{a}_{i\bullet}) = \sqrt{\frac{\sum\limits_{j=1,P} (a_{ij} - \overline{a}_{i\bullet})^2}{P(P-1)}} - standard error.$$

7. Compute
$$S_{ij} = \frac{a_{ij}}{\sum_{j=1,p} a_{ij}}$$
 - a "specificity score" of each gene in each group.

8. Compute
$$\overline{S}_{\bullet j} = \frac{\sum\limits_{i=1,G}^{S_{ij}}}{G} \quad \text{- an average "specificity score" of group } \textbf{\textit{j}}. \text{ And }$$

$$sigma(\overline{S}_{\bullet j}) = \sqrt{\frac{\sum\limits_{i=1,G}(S_{ij} - \overline{S}_{\bullet j})^2}{G(G-1)}} \quad \text{- standard error.}$$

$$da_{ij} = \frac{a_{ij} - \overline{a}_{i\bullet}}{sigma(\overline{a}_{i\bullet})}$$
 - an "abundancy parameter" which shows a difference of the abundancy score from an average abundancy score. In the case of
$$da_{ij} = \frac{a_{ij} - \overline{a}_{i\bullet}}{sigma(\overline{a}_{i\bullet})}$$
 - an "abundancy parameter" which shows a difference of the abundancy score from an average abundancy score. In the case of
$$da_{ij} = \frac{a_{ij} - \overline{a}_{i\bullet}}{sigma(\overline{a}_{i\bullet})}$$

$$da_{ij} \ge 2.0, \text{ we can consider this difference as statistically significant.}$$

$$ds_{ij} = \frac{s_{ij} - \bar{s}_{\bullet j}}{sigma(\bar{s}_{\bullet j})}$$

$$ds_{ij} \ge 2.0, \text{ we can consider this difference as statistically significant.}$$

$$- a \text{ "specificity parameter" which shows a difference of the specificity score from an average specificity score. In the case of $ds_{ij} \ge 2.0$, we can consider this difference as statistically significant.$$

11. We used the following default cut-offs to establish links between genes i and tissue/organs j and to form groups of genes expressed in a given tissue/organ: $da_{ij} \ge 2.0$ AND $ds_{ij} \ge 2.0$

[1] PMID: 15089753

Wingender, E.

9. Compute

TRANSFAC®, TRANSPATH® and CYTOMER® as starting points for an ontology of regulatory networks In Silico Biol. 4, 55-61 (2004).