In protein families, amino acid sequences reflect the evolutionary history that led to biological diversity. This available information remains to be unveiled from multiple sequence alignments. Clustering sequences into groups of similar features can provide clues to the evolutionary relationship between them. A solution to cluster objects is to generate a space by dimensional reduction. Objects correspond to points whose mutual distances depend on the metric.

To carry out our project, we used the programming language Perl and the statistical environment of R. We analyzed class A G-protein-coupled receptors from five species. Distance matrices were generated from multiple sequence alignments by different similarity measures. They were analyzed by two statistic techniques: metric multidimensional scaling (cmdscale in R package stats) and principal component analysis (dudi.pca in R package ade4). Biological objects were grouped by k-means (Kmeans in R package amap). Groups were validated with the clValid functions from the R package. The clustering analysis was bootstrapped in order to assess the grouping robustness. Best parameters were determined in relation with biological meaningfulness. Our approach indicates that receptors from different genomes share a similar clustering pattern that might relate to major evolutionary determinants of class A G-protein-coupled receptors.

References

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