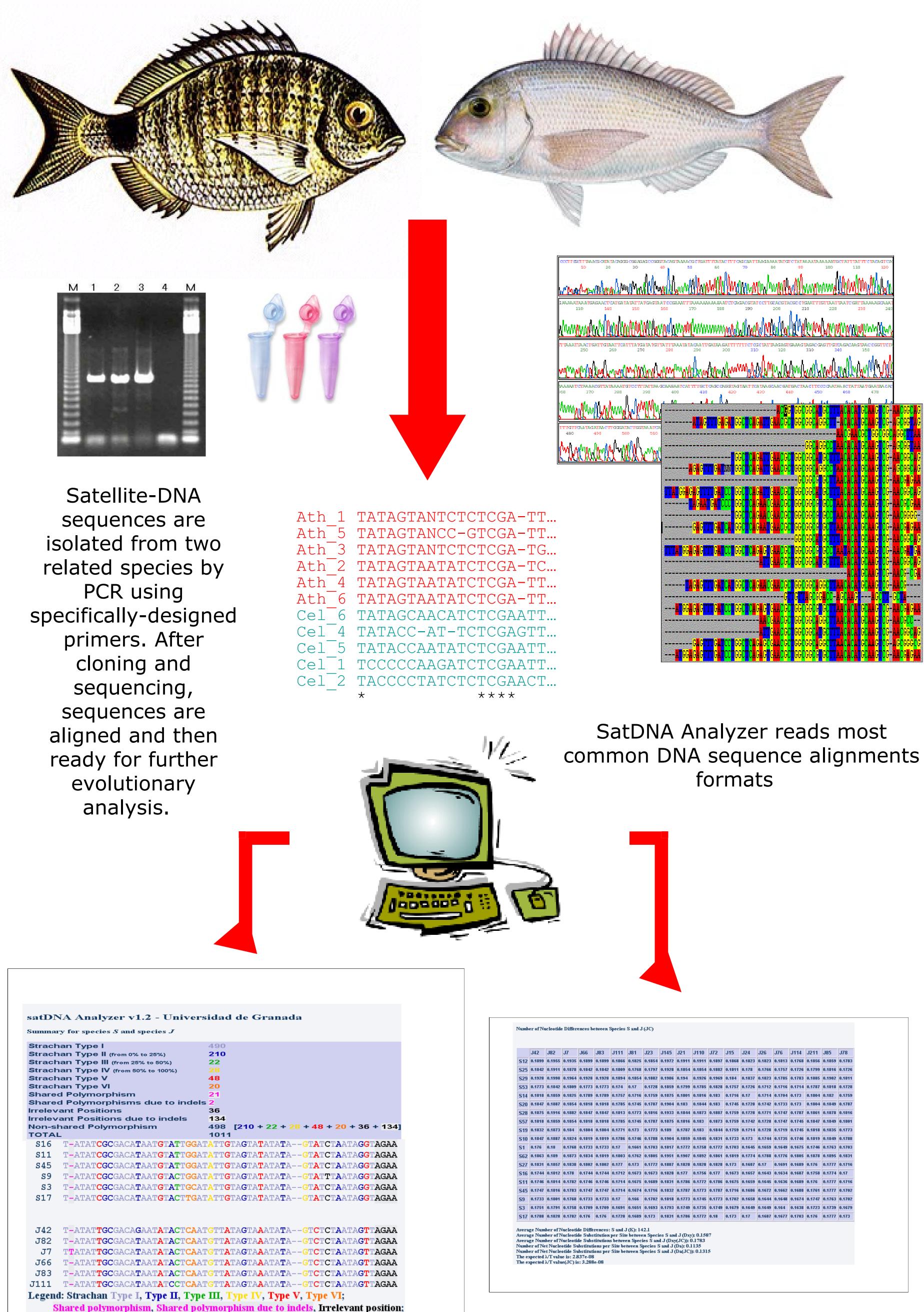
SatDNA Analyzer, the first computing solution for satellite-**DNA evolutionary analysis**

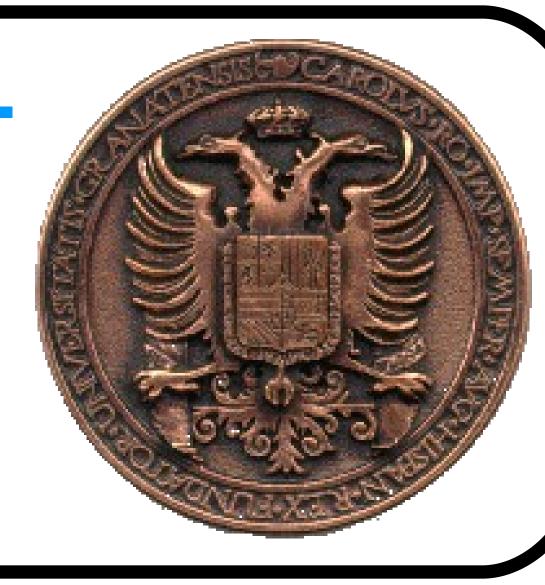
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satDNA Analyzer (satellite-DNA Analyzer), is a software package for the analysis of satellite-DNA sequences from aligned DNA sequence data. It allows for the analysis of the patterns of variation at each nucleotide position considered independently amongst all units of a given satellite-DNA family when comparing two species. The program classifies each

Make it easy, use satDNA Analyzer!!





site accordingly as monomorphic or polymorphic, discriminates shared from non-shared polymorphisms and classifies each non-shared polymorphism according to the model proposed by Strachan et al. (1985) in six different stages of transition during the spread of a variant repeat unit toward its fixation. Unfortunately, by this date, such calculations are manually performed in an arduous and timeconsuming manner. Trying to fill this gap, we have designed satDNA Analyzer 1.2. This software supposes a time saving feature since every utility is automatized and collected in a single software package, so the user does not need to use different programs. Additionally, it significantly avoids data miscalculations due to human errors, very prone to occur specially in large files.

Satellite DNAs are tandemly arrayed highly repetitive DNA sequences of the eukaryotic genomes located in the constitutive

heterochromatin. The repeats comprising a satellite-DNA family do not evolve independently of one another but rather follow concerted evolution. Basically, it states that once a mutation arises it spreads through the rest of repeats of one species toward the fixation of the new variant, being the rate of fixation higher than the rate of mutation. This leads to high levels of intraspecific homogeneity and interspecific divergence. Assuming molecular clock, Strachan et al. (1985) designed a method for concerted evolution quantification. This method compares position to position two sets of aligned sequences belonging to two different species. It allows classifying any position in terms of six stages. In brief, the Class 1 site represents complete homogeneity across all repeat units sampled from a pair of species, whereas Classes 2, 3 and 4 represent intermediate stages in which one of the species shows a polymorphism. The frequency of the new nucleotide variant at the site considered is low in Class 2 and intermediate in Class 3, while Class 4 represents sites in which a mutation has replaced the progenitor base in most members of the repetitive family in the other species. Class 5 represents diagnostic sites in which a new variant is fully homogenized and fixed in all the members of one of the species while the other species retains the progenitor nucleotide. A Class 6 site represents an additional step over stage 5 (new variants appear in some of the members of the repetitive family at a site fully divergent between the two species).

Furthermore, this program implements several other utilities for satellite-DNA analysis evolution such as the design of the average consensus sequences, the average base pair contents, the distribution of variant sites, the transition to transversion ratio, and different estimates of intra and inter-specific variation that make it a powerful tool for satellite-DNA evolutionary analysis.

Screenshots of satDNA Analyzer html output (left) Alignment showing in colors the different types of positions the software recognizes. (right) Example of a table display for statistical calculations.

References:

Strachan, T., Webb, D. and Dover, GA. (1985). Transition stages of molecular drive in multiplecopy DNA families in *Drosophila* EMBO J. 1985 July; 4(7): 1701–1708.

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