The Poplar Genome Project



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Introduction

Poplar trees have been used all over the world to produce a large variety of wood-based products such as timber, pulp, and paper. Besides their great economical value, poplars are also rapidly becoming the model organism for forest biology and tree biotechnology. It is therefore not surprising that, in 2001, poplar was selected as the first woody plant to have its genome sequenced.

Now that the poplar genome is publicly available, genomewide bioinformatics analyses of the genome sequence can take a start.

Genome annotation

The annotation of the genome was performed by 3 different labs: 2 in the USA (using FgenesH and Grail_EXP) and 1 in Belgium (using Eugene). From these 3 different annotations one consensus annotation was constructed using a rule-based system that for a given locus selected the most plausible genemodel. Before the post-processing steps (like removing putative transposons) we ended up with +- 58000 genes.

method	gene count	
Eugene	25018	43.11%
FgenesH	22546	38.85%
Grail_exp	7352	12.66%
genewise	3120	5 38%

Table 1 gives an overview of the selection of genes from the different gene finders. It is clear that our tool performs better than all others used in this annotation effort



Figure 1. Screenshot of the Poplar Genome Browser

The Genome Browser is available at: http://genome.igi-psf.org/Poptr1/Poptr1.home.html

Exploring the duplication past of poplar

For all paralogous genes in the poplar genome a Ks-estimation was performed. After corrections and filtering a histogram was constructed by plotting the number of paralogs against their age (Ks). As can be seen on the histogram there are many paralogs with the similar Ks-values (0.20-0.30) which indicate the fact that a large number of genes were created in a short period of time. The most plausible explanation for this observation is that a genome duplication has occurred in the poplar genome.

We performed the same procedure on ESTs of 7 different Poplar species and came to the same conclusion for all of them. This suggests that the genome duplication has occurred before the divergence of all poplar species.



Figure 1. Ks age distribution of the paralogous genes in the poplar genome

The most striking result of this analysis is not the fact that there was a genome duplication, but rather when it has happened. When we infer the time in My from the Ks distribution we dated the duplication event some 8 My ago, which clearly contradicts with the fossil records on poplar.

This indicates that the average Ks-substitution rate used for plants is probably not applicable to poplar. Due to its much longer generation time it is very likely that poplar has a much smaller Ks-substitution rate than all other plants studied so far.

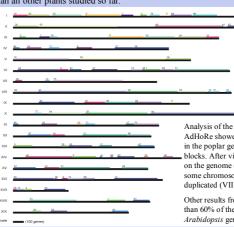


Figure 2. Map with duplicated blocks in the poplar genome

Analysis of the poplar genome with the software i-AdHoRe showed that more then 80% of all genes in the poplar genome are present in duplicated blocks. After visualization of the duplicated blocks on the genome (see figure 2) it became clear that some chromosomes have almost been completely duplicated (VIII and X).

Other results from this analysis showed that more than 60% of the poplar genome is colinear with the Arabidopsis genome.

Gene family comparison



between Poplar and Arabidopsis of the number of transcription factors. For most of the TFs the numbers are comparable although there are some differences, eg. the Myb TFs, which seem to have undergone lineage specific expansion in poplar. Also the pathogen-related TFs are more abundant in Poplar, which is perhaps correlated with the perennial nature of Poplar.

Table 1 shows a comparison

Repetitive sequences and transposons

- Repeats occupy 28% of the assembled genome and more than 50% of the unassembled scaffolds.
- Small scaffolds and unassembled reads contained substantially more repetitive sequence than large scaffolds.

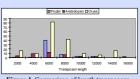


Figure 1. Comparison of length transposons



When we compare the length and frequency of LTRtransposons in Arabidopsis, poplar and rice it is clear that in rice there are more and longer transposons, although the longest ones are identified in poplar.

This comparison included 37 Ath, 58 poplar and 181 rice transposons.

Conclusions

- · Outstanding international collaboration has placed poplar on equivalent footing with other model species
- Assembly was heavily complicated by repeats, but facilitated by genetic maps
- Unprecedented opportunities to explore molecular basis of commercial and environmental traits
- · Most of the hard work and many exciting discoveries are

References:

- ov A., Solovyev V. (2000) FgenesH: Ab initio gene finding in Drosophila genomic DNA. Genome Res., 10,516-522
- ex.T., Moisan , A. and Rouzé, P. (2001). EuGène : An Eucaryotic Gene Finder that combines several sources of evidence. Computational Biology, Eds. O. Gascuel and M-F. Sagot, LNCS 2066, pp. 111-125
- Simillion, C., Vandepoele, K., Saeys, Y. & Van de Peer, Y. (2004) Building genomic profiles for uncovering segmental homology in the twilight zone. Genome Res. 14, 1095-1106
- Sterck, L., Rombauts, S., Jansson, S., Sterky, F., Rouzé, P. & Van de Peer, Y. (2005) EST data suggest that popular is an ancient polyploid. New Phytol. (In Press)
- Ying Xu, Manesh Shah, Doug Hyatt, Richard Mural, and Edward C. Uberbacher, GRAIL-EXP. Multiple Gene Modeling Using Pattern Recognition and Homology, The Seventh Department of Energy Contractor and Graniee Workshop, January 1999