The low level of variability is depicted in plastomes of early soybean varieties

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Motivation and aim

The chloroplast genome of land plants is usually conservative in structure, but variable in sequence. Intraspecific diversity of organelle possibly plays an important role in successful breeding and creation of the new varieties in plants.

The aim of our study was to evaluate the level of chloroplast genomes diversity in specific group of *Glycine max* varieties. The study sample contained 24 early-maturing *G. max* cultivars from our collection. All studied soybean varieties have different geographic origin.
Methods

1) Isolation of the chloroplast fraction by differential centrifugation from the 7-day-old soybean seedlings*

2) Phenol-chlorophorm DNA extraction

3) DNA sequencing on Illumina MiSeq using Illumina DNA Prep library preparation kit and MiSeq Reagent Kit v3 (600-cycle)

4) The NGS data processing algorithm included:

   - FASTQ
   - SAM
   - BAM
   - VCF
   - FASTA

Quality of the isolated chloroplast DNA was checked by RFLP-analysis

Ethidium bromide stained agarose gel of electrophoretically-separated soybean chloroplast DNA restriction fragments

1, 2 – undigested chloroplast DNA; 3 – Lambda DNA/EcoRI Marker; 4 – Lambda DNA/HindIII Marker; 5-12 – chloroplast DNA of different soybean varieties digested by EcoRI

Variability was detected in **8 positions** along the whole cpDNA molecule.
Comparative analysis of 24 complete soybean chloroplast genome sequences revealed 3 SSRs, 1 INDEL and 5 SNP between samples. 4 SNPs were located in coding sequences of \textit{atpB}, \textit{rps4}, \textit{accD} and \textit{rps3} genes. All of them were synonymic.

### Discovered polymorphic sites of the soybean chloroplast genome

<table>
<thead>
<tr>
<th>Location in the reference sequence (MW357264.1)</th>
<th>Reference allele</th>
<th>Alternative allele</th>
<th>Cultivars carrying the alternative allele</th>
</tr>
</thead>
<tbody>
<tr>
<td>6967 rbcL – atpB intergenic region</td>
<td>(T)$_{11}$</td>
<td>(T)$_{10}$</td>
<td>Viliya</td>
</tr>
<tr>
<td>8408 atpB coding sequence</td>
<td>G</td>
<td>A</td>
<td>Lyubasha, Kitrossa, Oressa</td>
</tr>
<tr>
<td>15507 rps4 coding sequence</td>
<td>T</td>
<td>A</td>
<td>Lyubasha, Kitrossa, Oressa</td>
</tr>
<tr>
<td>38504 rpoC1 intron</td>
<td>(A)$_{4}$</td>
<td>(A)$_{5}$</td>
<td>Lyubasha, Kitrossa, Oressa</td>
</tr>
<tr>
<td>51525 atpA - trnR-UCU intergenic region</td>
<td>(A)$_{19}$</td>
<td>(A)$_{13}$</td>
<td>Lyubasha, Kitrossa, Oressa</td>
</tr>
<tr>
<td>57873 accD coding sequence</td>
<td>C</td>
<td>A</td>
<td>Lyubasha, Kitrossa, Oressa</td>
</tr>
<tr>
<td>75657 petD intron</td>
<td>(T)$_{14}$</td>
<td>(T)$_{15}$</td>
<td>Lyubasha, Kitrossa, Oressa</td>
</tr>
<tr>
<td>82035 rps3 coding sequence</td>
<td>G</td>
<td>T</td>
<td>Lyubasha, Kitrossa, Oressa</td>
</tr>
<tr>
<td>116598 ndhA intron</td>
<td>T</td>
<td>A</td>
<td>Lyubasha, Kitrossa, Oressa</td>
</tr>
</tbody>
</table>

On the basis of revealed diversity 3 haplotypes can be defined:

- 6967 (T)$_{11}$/ (T)$_{10}$
- 8408 G/A
- 15507 T/A
- 38504 (A)$_{4}$/ (A)$_{5}$
- 51525 (A)$_{19}$/ (A)$_{13}$
- 57873 C/A
- 75657 (T)$_{14}$/ (T)$_{15}$
- 82035 G/T
- 116598 T/A
Conclusions

Analysis of 24 complete soybean chloroplast genome sequences demonstrated the drastically low level of genetic diversity in studied varieties of *Glycine max*. Possibly it is the result of the founder effect – evidently a narrow range of ancestors was used while this group of varieties was created.

Our results are consistent with earlier works, that revealed the low level of intraspecific diversity in soybean.

Quite another data was obtained earlier in our laboratory for Poaceae (barley): 9 INDELs, 19 SSRs and 79 SNPs. *

Three varieties (Lyubasha, Kitrosa and Oresa) were found to be extremely different from the other samples studied: 2 SSRs, 1 INDEL and 5 SNP. It is of special interest that maternal parents of these three varieties originated from China.

The future plans:

To verificate of NGS data by Sanger

To obtain and compare full mitochondrial genomes of 24 *G. max* varieties listed above

To enlarge the range of *G. max* varieties for further study of their organelle genome diversity

To submit the obtained complete organelle sequences into NCBI GenBank database

To realize: Is there a link between successful hybrid performance and type of maternal cytoplasm?

Acknowledgments

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