CYTOGENETIC EFFECTS INDUCED BY DEPOSIT MYCOFLORA IN VICIA FABA BEANS FROM THE COLLECTION OF SUCEAVA GENEBANK

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Keywords: *Vicia faba* beans, deposit mycoflora, cytogenetics

Abstract: The purpose of our work was to study the cytogenetic effects induced by deposit mycoflora in *Vicia faba* (2n = 12) beans from the collection of Suceava Genebank. Cytogenetic effects were studied by means of classical plant chromosomes methods of study. We have observed that the values of the mitotic indexes are decreasing in accordance with the increasing of the storage age and also with the number of mycoflora species that are infesting the beans. Infestation with specific mycoflora produced a relatively large number, statistically significant by comparison to the controls, of interphasic aberrations and also of chromosomal aberrations in mitosis ana-telophase in all studied probes. Comparing our results with similar published data, we can strongly state that the cytogenetic effects induced by deposit mycoflora in *Vicia faba* seeds are similar with those produced by the action of a weak mutagenic agent

INTRODUCTION

The purpose of our work is to study the cytogenetic effects induced by deposit mycoflora in various species seeds from the collection of Suceava Genebank. This paper shows the cytogenetic effects induced by deposit mycoflora in $Vicia\ faba\ (2n=12)$ beans from the above mentioned collection. Cytogenetic effects were studied by means of: calculation of the mitotic index and frequency of mitosis phases, registration of the frequency and types of abnormal interphases and identification of chromosomal aberrations in mitotic ana-telophases, in accordance with classical plant cytogenetics methods.

MATERIALS AND METHODS

The biological material consisted of five probes of beans selected from the collection of Suceava Genebank:

- The control: SVGB-273, 23 years old, germination ratio 100%, not infested by mycoflora,
- SVGB 272, 18 years old, germination ratio 92%, infested by *Penicillium* sp.
- SVGB 212, 20 years old, germination ratio 92%, infested by *Penicillium* sp.
- SVGB 300, 20 years old, germination ratio 83%, infested by *Rhizopus nigricans* and *Colletotrichum lindemuthianum*
- SVGB 271, 23 years old, germination ratio 87%, infested by *Penicillium* sp. and *Aspergillus flavus*

The beans germinated in Petri dishes, on filter paper, wetted with distilled water for all variants till the roots reached 10-15 mm in length. We calculated the germination percent and the duration of germination for all experimental variants. When the roots reached the length of 10-15 mm they were fixed in Battaglia for 30 minutes. We have performed fast cytological slides according to Feulgen method (Cimpeanu at al., 2002). We have studied:

- 1. The mitotic index and frequency of mitosis phases
- 2. The frequency and types of aberrant interphases
- 3. The frequency and types of chromosomal aberrations in mitotic ana-telophases (A-T).

We have analysed this 3 steps on 5 fresh preparations. For each slide we have studied: 10 microscopic fields (objective 40x) on which we have counted all the cells in interphase, prophase, metaphase, anaphase and telophase; 10 microscopic fields on which were counted all the cells in normal and abnormal interphases and the type of interphasic aberrations and 50 ana-telophases were analysed, counting the normal, aberrant ana-telophases and the type of chromosomal aberrations. All data were statistically processed (Fowler and Cohen, 1990). Microphotographs were performed with the digital camera of Nikon research microscope with the 100X objective.

RESULTS AND DISCUSSIONS

The infestation with fungi generally determined a significant decrease of the mitotic index (Table 1). Only for the SVGB-272 probe the frequency of dividing cells is 6,45%, a very close value to

the mitotic index registered for the control (6,47%) (fig. 1). The lowest value of the mitotic index (1,69%) was observed for the SVGB-271 probe.

The values of the mitotic index are proportionally decreasing with the increase of the number of infesting species of fungi and with the storage age of beans.

For the majority of probes, and also for the controls, we have recorded a predominance of prophases, followed by metaphases and telophases. Anaphases were recorded only for controls, but in a very small percentage (Table 1 and Figure 2). An exception was noted for the probe SVGB-271 where the frequency of prophases was low and metaphases were predominant. The frequency of prophases is decreasing proportionally with the increase of infestation and of the storage age of the beans (Fig. 2). The frequency of telophases is decreased for all probes in comparison to the controls. An exception was noted for the SVGB-272 probes in which we have registered a high frequency of telophases in comparison to the controls.

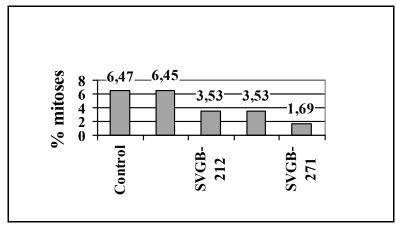


Fig. 1: Mitotic index in Vicia faba studied probes

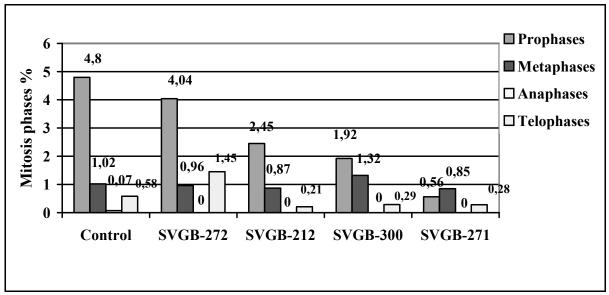


Fig. 2: The frequency of mitosis phases in *Vicia faba* studied probes

Infestation with fungi induced the increase of interphasic aberrations in all the studied probes (Table 2 and Fig. 3).

The highest frequency of aberrant interphases (1,71%) was registered for SVGB-272 probe. For the other probes the registered values are similar and quite low, but, in comparison to the very small value of aberrant interphases counted in the control probes (0,03%), we can state that the infestation with mycoflora induced a statistically significant increase of anaphase aberrations, without regard to the type of fungi or to the age of probes.

The most frequent interphasic aberration is represented by the binucleated cells, which frequency is increasing proportionally with the age of probes and with the number of infesting mycoflora species (Fig. 4).

In SVGB-272 probe the infesting mycoflora induced some interphases with one micronucleus (Fig. 5 and 6) and, in a very low percentage, interphases with two micronuclei. Analysing all the mentioned data in comparison with data from literature (Maniu et al. 2002, 2005) we can strongly state that the cytogenetic effects induced by deposit mycoflora in *Vicia faba* beans are similar with those produced by the action of a weak mutagenic agent.

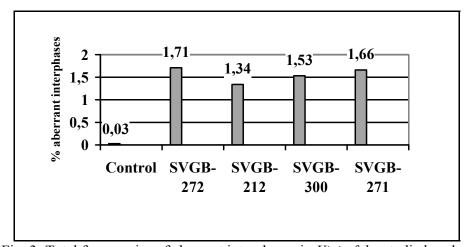


Fig. 3: Total frequencies of aberrant interphases in Vicia faba studied probes

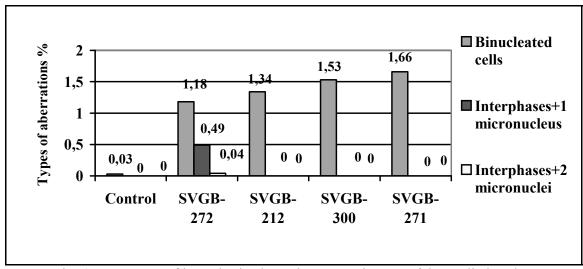


Fig. 4: Frequency of interphasic aberrations types in *Vicia faba* studied probes

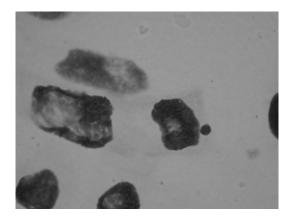


Fig. 5: Interphase with one micronucleus (SGVB-300 probe)

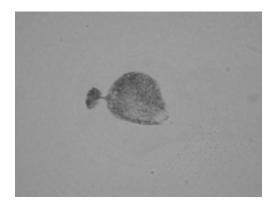


Fig.6: Chromatic continuity between nucleus and micronucleus in interphase (SVGB-271 probe)

Analysing the data from table 3 one can observe that both for the control, SVGB-272 and SVGB-271 probes were registered chromosomal aberrations in mitosis ana-telophase. In the rest of two probes the number of observed ana-telophases was too small to perform statistic analysis. The highest percentage of aberrant ana-telophases (5,67%) was observed for the SVGB-271probe (Fig. 7). The chromosomal aberrations types show a great variety, being distributed randomly (Fig.8).

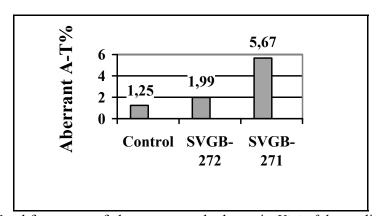


Fig. 7: Total frequency of aberrant ana-telophases in Vicia faba studied probes

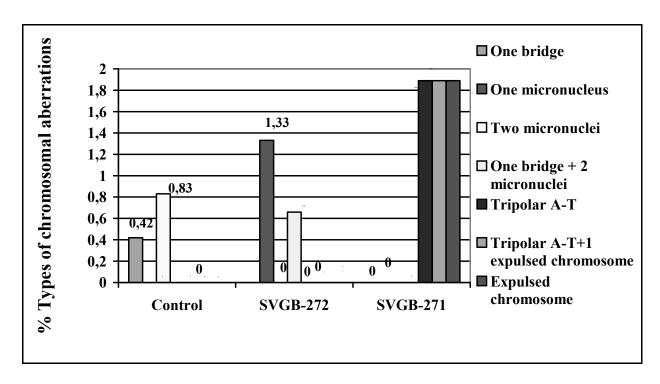
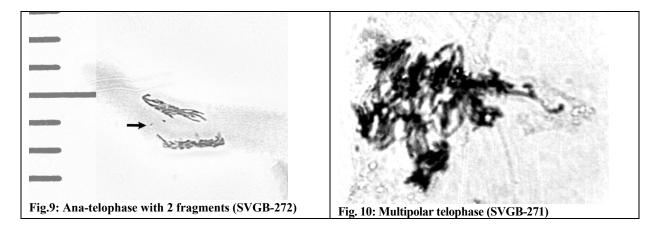


Fig. 8: Frequency of chromosomal A-T aberrations in Vicia faba studied probes

We have observed and counted many simple chromosomal aberrations such as: one bridge, one micronucleus, two micronuclei, tripolar ana-telophases, expulsed chromosomes and some complex chromosomal aberrations such as: one bridge with two micronuclei, tripolar A-T with expulsed chromosomes, etc. (fig. 9-14). The highest frequency was registered for tripolar ana-telophases and expulsed chromosomes.



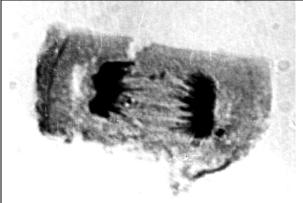


Fig.11:Ana-telophase with fragments (SVGB-300)



Fig.12: Ana-telophase with a bridge (SVGB-271)



Fig. 13: Hexapolar ana-telophase (SVGB-272)

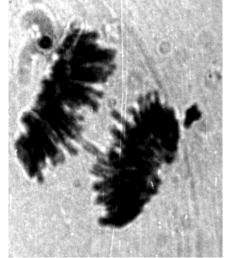


Fig. 14: Pentapolar ana-telophase with a bridge and expulsed chromosomes (SVGB-300)

The increase of the number of aberrations in ana-telophase is statistically significant reported to the controls; comparing our results with similar published data (Maniu et al. 2002, 2005) we can strongly state that the cytogenetic effects induced by deposit mycoflora in *Vicia faba* beans are similar with those produced by the action of a weak mutagenic agent.

CONCLUSIONS

The values of the mitotic indexes are decreasing in accordance with the increasing of the storage age and also with the number of mycoflora species that are infesting the beans.

Infestation with specific mycoflora produced a relatively large number, statistically significant by comparison to the controls, of interphasic aberrations and also of chromosomal aberrations in mitosis ana-telophase in all studied probes.

Comparing our results with similar published data, we can strongly state that the cytogenetic effects induced by deposit mycoflora in *Vicia faba* seeds are similar with those produced by the action of a weak mutagenic agent.

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Total anaphases Table 1: Frequency of mitoses and of mitosis phases in Vicia faba studied probes Total metaphases 96,0 0,87 0,85 28 Total prophases 4,04 1.69 6,45 Total mitoses 6.47 101 48 Total interphases 93,55 96,47 96,47 93.53 98.31 3104 Total studied cells 2753 3318 2859 3029 SVGB-212 SVGB-300 **SVGB-272** SVGB-271 Control

0,21

Table 2: Frequency of interphasic aberrations in Vicia faba studied probes

	Total								Aberration types	n types		
Probes	studied cells	Total interp	Fotal normal interphases	Total	Fotal aberrant cells	cells	Binuclea	Binucleated cells	Interphases with one micronucleus	nases one releus	Interphases with two micronuclei	ises with ronuclei
	Nr.	Nr.	%	Nr.	%	$\pm s\bar{x}$	Nr.	%	Nr.	%	Nr.	%
Control	2941	2940	76,99	1	0,03	9000,0	_	0,03				
SVGB-272	3045	2993	98,29	52	1,71	0,032	36	1,18	15	0,49	1	0,04
SVGB-212	3511	3464	99,86	47	1,34	0,003	47	1,34				
SVGB-300	3270	3220	98,47	50	1,53	0,004	50	1,53				
SVGB-271	3133	3081	98,34	52	1,66	0,004	52	1,66				

Table 3: Frequency of chromosomal aberrations in mitosis ana-telophase (A-T) in Vicia faba studied probes

×			_	_		
		Expulsed	%	-	-	1,89
		Expu	Nr	-	-	1
		Tripolar A-T with expulsed chromosome	%			1,89
	nitosis A-T		Z.	1	-	1
	n mitosis	Tri-polar A-T	%		-	1,89 1 1,89
	Types of chromosomal aberrations in m		Ż		ı	-
		One bridge + 2 micro- nuclei	%	-	1 0,66	
			Ż		1	,
		Two micro- nuclei	%	0,83	-	ı
			Nr	2	-	-
		One micro- nucleus	%	-	1,33	-
			Nr	-	2	-
		One bridge	%	0,42	-	ı
		One		1	'	,
	Total Total abnormal A-T		$\pm S\overline{X}$ Nr % Nr % Nr % Nr % Nr %	0,046	0,092	0,44
			%	1,25	1,99	5,67 0,44
			Nr	3	3	3
			%	98,75	98,01	94,33
			Nr	236	148	50
		Studied A-T	Nr	239	151	53
		Probes		Control	SVGB- 272	SVGB-