Mapping of major net-type net-blotch resistance loci in Ethiopian barley line c-23874 and the Russian barley cultivar Zernogradskij 813

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Materials and methods





Net blotch caused by *Pyrenophora teres* Drechs f. *teres* is an important barley disease in Russia and elsewhere. Under favorable conditions net blotch can cause significant reductions in both the yield and quality of the crop. Yield losses from this disease on susceptible cultivars in Russia can reach 40% under epidemic conditions.

Table. 1. Two barley mapping populations comprise anther culturederived doubled-haploid lines (DHLs), were developed.







Mapping population B, 114 DHLs						
Rannij 1	Russia (Siberia)	S	R			
Zernogradskij 813	Russia (South Region)	R	S			

Resistance evaluations were conducted in 2010 in greenhouse of MTT (Finland) and at laboratory conditions (VIZR, Russia) using detached leaf technique. DHLs were screened with one NTNB isolate from Finland and two isolates from the Northern part of Russia collected in Leningrad (PL 2) and Novgorod (PN 2) regions in 2010.

Results and discussion

Fig. 1. Frequency of disease reaction incited on the DHLs of two barley populations ("A" and "B") by *Pyrenophora teres* isolates.



In the greenhouse experiments two parental genotypes c-23874 and Pirkka were inoculated in four replications by isolate v-278 and



showed averaged infection types 1.5 and 8 correspondingly





The infection response was recorded on the second leaf, 10 days after inoculation, using the 10-point scale (1)



Table. 2. Number of polymorphic SNP markers, used for genotyping two DHLs populations

	Chromosom	es	Quant	ity of SNP marl	kers	Polymor	phic	polym	orphi
	Population "A"						assem	bled u	
	1H			43		12		replica	ations
1	2H			64		14			
	3H			55		20		chroin	osom
	4H			49		12			≜ ⊥c
	5H			74		19			3,0 T
	6H			55		17			
	7 H			44		14			2,4
	Bcero:			384		108			18
	Population "B"							1,0	
	1H			43		18			1,2 +
	2H			64		27			
	3H			55		31			0,6 🗕 /
	4H			49		20			A
	5H			74		30			0,0 +
	6H			55		22			0
	/H			44		16			0,53 ⁴ a0
	Bcero:			384		104			0,27+
la	tes of <i>P. teres</i> popula	f. <i>ter</i> tion o	es origin derived f	nated from from Ranni	the Nor j/ Zerno	rth of Rus ogradskij	ssia in DH ba cross	arley	-0,53↓I *
e	Origin	Exp	Infectio	Infection response		Effect of QTL in DH prog			
			Pa	Parents		OTL interval (cM)	Resistance source	e LOD/ R ²	To
			Rannij1	Zernogradsk ij 813					2. re
	Leningrad region	1	8.2	2.5	5H	4,6-19,9	Zernogradskij	2,4/0,0 8	va Ze
		2	7.1	2.2	5H	4,6-19,9	Zernogradskij	3,7/0,1 2	Ŧ
	Novgorod	1	7	2,2	5H	4,6-19,9	Zernogradskij	3,0/0,1	
	region							1	JC
		2	7,2	2,2	5H	4,6-19,9	Zernogradskij	2,7/0,1	ge T1

Segregation of resistance to PL 2 and PN 2 isolates in DHLs of "B" population





The 384 SNP set employed for genotyping of the both DH populations was selected at the JHI and covered all seven barley chromosomes. All 384 SNP markers provided good quality genotype calls for the DHLs, 108 and 164 SNP markers showed ism in population "A" and "B" respectively and were used for mapping. For both populations linkage groups were using the computer program MapManager QTX (2) with an LOD > 3.0 and the Kosambi mapping function (3). In all four of the inoculation experiment the highest LOD score (2,9) was associated with the SNP marker 11_11067 located on ne 6H (58 cM).



10	28	an	-+11
0.0 21.0 25.0 36.0 41.0 4	0.0 10.0 18.0 18.0 18.0 18.0 18.0 10.1 10.2 10.1 10.2 10.1 10.2 10.1 10.2 10.1 10.2 1	0.0 27.0 52.0 70.0 73.0 75.0 85.0 85.0 90.0 93.0 9	9,0 13,0 39,0 49,0 52,0 5
130,0 - 11_20908	118.0 123.0 125.0 125.0 11.0005	105.0 116.0 121.0 105.0 116.0 11_20628 11_20628 11_20628 11_20628 11_20628	96,0
5H		144.0	

Tabl iso

PL2

PN 2

- additive effect of c-23874 allele

Location of major net blotch resistance genes on barley chromosome

101,0 104,0 131,0 134,0

'o determine the critical LOD threshold, we executed a permutation test with 1000 permutations. A LOD threshold of about .7 in this DH population yields an experiment-wise significance level of 0.05. An identical marker interval associated with esistance to NTNB was identified for the both isolates on short arm of chromosome 5H explaining 8-12% of the total ariation. The interval was flanking by SNP markers 11_20533 and 11_20899 located in 4,6 and 19,9 cM on the Rannij 1/ Zernogradskij 813 genetic map.

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