

Allelic composition of puroindolinium genes and confectionery properties of flour of soft winter wheat samples

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Purpose. Identification of soft winter wheat varieties and lines from the Plant Production Institute nd. a. V. Ya. Yuryev, NAAS by allelic state of *Pina-D1* and *Pinb-D1* genes for targeted use in the breeding for high confectionery properties of flour. **Methods.** Identification of the *Pina-D1* and *Pinb-D1* genes allelic state was performed by polymerase chain reaction (PCR) using allele-specific primer pairs. Confectionery properties of flour were evaluated by determining the quality indicators: the water absorption capacity (WAC) of the flour, trial baking of cookies and evaluation of its quality. **Results.** According to the results of PCR analysis, 9 samples had an allelic composition of puroindoline genes (*Pina-D1a* and *Pinb-D1a*) characteristic for soft-grained varieties. Flour of the lines 'L137-26-0-2', 'L137-26-0-3' had the best confectionery properties, it had a WAC value less than 55%, cookies diameter 85 mm, cookies height 10 mm, estimation of a surface of cookies 7–9 points, what meets the requirements for soft-grained wheat. 76% of the samples belonged to hard-grained varieties and had the corresponding alleles of the *Pina-D1* or *Pinb-D1* genes. In the studied sample, *Pina-D1* gene is represented by 2 alleles: *Pina-D1a* and *Pina-D1b*. 27 samples had *Pina-D1a* allele, which also allows them to be used in breeding programs for grain quality when crossed with soft samples, 4 ones had *Pina-D1b* allele. As to *Pinb-D1* gene, all hard grain samples had *Pinb-D1b* allele, and the 'Erythrospermum S 424-1/14' line was heterogeneous for *Pinb-D1a/Pinb-D1b*. The flour of these samples had typical for hard wheat quality indicators: WAC 68% and more, cookie diameter of 60–72 mm, cookie height of 13–15 mm, the surface evaluation of 1–4 points. **Conclusions.** The studies allowed to differentiate the breeding material and transfer a soft winter wheat cultivar of a confectionery use 'L137-26-0-3' ('Mazurok') which has an allelic structure of puroindolins genes (*Pina-D1a* and *Pinb-D1a*) characteristic for soft-grained varieties and high confectionery flour properties for qualification examination.

Keywords: common winter wheat; variety; line; *Pina* and *Pinb* genes; water absorption capacity; cookies.

Introduction

Winter wheat (*Triticum aestivum* L.) is the leading grain crop in Ukraine; it surpasses other grain crops in terms of sown areas (5.9–6.5 million hectares) and forms the basis for the formation of the country's grain balance [1]. In the context of a market economy, increased competition in the market and in connection with Ukraine's entry into the World Trade Organization, increasing the competitiveness of agricultural products becomes especially relevant.

Special attention should be paid to improving the competitiveness of grain products, including increasing its technical and quality level [2, 3]. At the same time, consumer re-

quirements for food and baking properties are growing and expanding, in particular, the creation of confectionery varieties necessary for manufacture of *cakes* and *pastries* is urgent [4].

Commodity classifications of grain in some EU countries are based on the peculiarities of the endosperm structure, in Ukraine this indicator is not taken into account [5–8]. The world is successfully breeding wheat varieties for special purposes. Soft wheat varieties 'Ami', 'MV Irma', 'Webster', 'Wisdom', 'FS 401' and others have been created among them. However, due to the low level of adaptability to growing conditions, they are not introduced on the territory of Ukraine [9]. Only two varieties of soft-grain wheat – 'Biliava' (Plant Breeding and Genetics Institute – National Center for Seed Production and Variety Research, UA) and 'Arkeoks' (Limagrains Europe, FR), which have the indices necessary for confectionery wheat have been entered into the State Register of Plant Varieties of Ukraine by 2020 [10].

The grain hardness of soft wheat is one of the most important characteristics of grain quality and directly related to grain grinding, dough kneading and making bakery products. Depending on the manifestation of this trait, wheat

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grain can be classified as a bakery or confectionery type [11]. Compared with soft endosperm caryopses, in varieties with a hard endosperm, grain grinding is a long and energy-intensive process. As a result, flour with large particles is formed. It contains a large amount of damaged starch grains, due to which it has a high water absorption capacity (WAC, %). Due to the availability of a large amount of carbohydrates as a substrate for yeast, such flour is better for yeast bread baking [12]. High-quality confectionery products are made from special types of flour, obtained when grinding grain of specific soft-grain wheat varieties. Confectionery wheat flour differs significantly in its technological properties from baking wheat flour. This flour has a characteristic consistency; it is looser than baker's and has a low specific gravity, significantly lower WAC compared to baking flour. Such technological characteristics of the flour provide a high thermoplasticity of the pastry kneadind in the first minutes of baking, due to which obtained biscuits are more loose and of better quality both in terms of organoleptic, taste, and biological characteristics of the nutritional value of the product [4]. Studies of the nature of endosperm structure inheritance revealed that the differences in cultivars in this trait are determined by several linked genes located on the short arm of chromosome 5D at the B locus (Hardness) [13, 14].

The genes encode three polypeptides that form the freeabilin protein: puroindoline a (*Pina-D1* gene), puroindoline b (*Pinb-D1* gene), and Grain Softness Protein (*Gsp-1* gene). Changes in the amino acid composition of these polypeptides are closely related to changes in grain texture. Soft wheat varieties contain both "wild" alleles of *Pina-D1* gene (*Pina-D1a* allele) and the *Pinb* gene (*Pinb-D1a* allele), while hard-grain wheat varieties contain either a deletion of the *Pina-D1* gene, or one of the "mutant" forms of *Pinb-D1* gene (alleles *Pinb-D1b-g* or *Pinb-D1l*) [15–17].

Considering the prospect of breeding directions expanding in order to increase the competitiveness of soft winter wheat varieties, the corresponding studies (the search and attraction of new source material for soft wheat breeding) are carried out at Plant Production Institute nd. a. V. Ya. Yuriev, NAAS of Ukraine. Since 2001, varieties and lines with high adaptive potential have been crossed with soft-grain varieties and lines of foreign breeding. At the National Center for Plant Genetic Resources of Ukraine, soft-grain lines [18–20] were registered, as well as an indicative collection of confectionery properties [21], a number of constant

selection lines of confectionery use were created, which are now being pre-tested, and the variety of soft-grain wheat 'Mazurok' was transferred to qualification examination at the Ukrainian Institute for Plant Varieties Examination.

The effective use of foreign material during hybridization with high-yielding local varieties adapted to the conditions of Ukraine is possible when the latter are identified by the allelic state of puroindolinium genes. Today, the allelic composition of these genes is being actively studied in varieties created at the Plant Breeding and Genetics Institute – National Center for Seed Production and Variety Research, the V. M. Remeslo Myronivka Institute of Weat, and Poltava State Agrarian Academy [23, 24].

The aim of the research is to identify *Pina-D1* and *Pinb-D1* genes of soft winter wheat samples of the Plant Production Institute nd. a. V. Ya. Yuriev, NAAS of Ukraine for further use in breeding for high confectionery properties of flour.

Materials and methods

Grains of 25 varieties of winter wheat, 12 selection lines of confectionery direction of use, which are pre-tested, 3 lines of *Triticum spelta* L were used in the study. As the standards of technological parameters of flour, we used the soft-grain variety 'Biliava' and the hard-grain variety 'Pryvablyva'.

Preliminary selection of soft-grain forms was carried out by determining the indicators: WAC of flour, test baking and evaluation of the quality of cookies [22].

To identify the allelic states of *Pina-D1* and *Pinb-D1* genes in the soft winter wheat samples, the polymerase chain reaction (PCR) method was used with the use of target gene markers.

DNA isolation was carried out from 5 grains of ten individual plants of each sample using the DiatomDNAPrep100 reagent kit (Neogen). DNA amplification was performed in tubes with a lyophilized PCR reagent kit (GenePak PCR core) in a Tertsik amplifier (Russia).

The reaction mixture for PCR analysis, which contained 20 ng of isolated DNA, 1 μM of forward and reverse primers, was brought to the final volume (20 μl) with the solvent from the PCR kit.

The identification of the allelic state of *Pina-D1* and *Pinb-D1* genes was carried out using primers proposed by Gautier et al. [25] and Klčová et al. [26]. Differentiation of *Pinb-D1* alleles (*pinb-D1a* and *pinb-D1b*) was performed with allele-specific primer pairs [27] and restriction analysis proposed by Klčová et al. [26] (Table 1).

Table 1

Primers for identification of *Pina-D1* and *Pinb-D1* gene alleles

Gene	Fragment length	Primer name	Sequence
Pina-D1	330	Pina-D1-F	CCC TGT AGA GAC AAA GCT AA
		Pina-D1-R	TCA CCA GTA ATA GCC AAT AGT G
Pinb-D1	447	Pinb-D1-F	ATG AAG ACC TTA TTC CTC CTA
		Pinb-D1-R	TCA CCA GTA ATA GCC ACT AGG GAA
	250	Pinb-glyR	CTC ATG CTC ACA GCC GCC
		Pinb-serR	CTC ATG CTC ACA GCC GCT

For identification, we used winter wheat varieties with known alleles of *Pina-D1* and *Pinb-D1* genes: 'Rheia' – *Pin a D1b*, 'Vasylyna' – *Pin a D1a*, *Pin b D1 b*, *serine*, 'Mirleben' – *Pin a D1a*, *Pin b D1a*, *glycine* [23, 25].

Amplification for all pairs of primers was performed under the following conditions: denaturation 95 °C (5 min), then 35 cycles: 94 °C (30 sec), 60 °C (30 sec), 72 °C (1 min 15 sec), final elongation 72 °C (7 min).

Amplification products were visualized by electrophoresis in 2.0% agarose gel in borate buffer; ethidium bromide was used for DNA monitoring in ultraviolet light according to

the standard procedure [28]. Electrophoresis was performed in a horizontal Hoefer Super-Sub100 device. M 50 GENPAK® was used as a molecular weight marker. The resulting gels were documented using Nikon D50 camera.

To determine the size of the amplification products, a demo version of the TotalLab 120 program (<http://www.totallab.com>) was used.

Results and discussion

Varieties and new breeding lines of winter wheat were preliminarily differentiated by confectionery properties, having determined technological indicators of confectionery quali-

Table 2

Confectionery properties of soft winter wheat samples (2016–2019)

Sample name	Evaluation of cookies				WAC, %
	diameter (D), mm	thickness (T), mm	D/T	evaluation of the cookie surface, score	
'Pryvablyva' – St	79.17	12.27	6.48	4	62.7
'Metelytsia kharkivska'	80.48	11.13	7.27	6	62.2
'Erytrospermum 533-16'	82.17	10.82	7.64	7	65.3
'Liutestsens 652-16'	73.67	12.13	6.09	7	68.4
'L 139-03 KH'	81.57	11.00	7.50	7	62.7
'Erytrospermum S 424-1/14'	80.00	11.14	7.28	6	59.1
'Erytrospermum 1002-16'	77.67	11.57	6.73	7	66.1
'Erytrospermum 1003-16'	80.33	11.07	7.26	7	61.0
'L137-26-0-2'	85.10	10.07	8.61	8	52.7
'L137-26-0-3'	86.00	9.95	8.68	9	53.1
'L202-20'	80.04	11.32	7.20	6	56.8
'S 492-3/14'	77.00	11.27	6.83	9	62.3
'VS 2019-1/15'	76.67	12.03	6.37	7	65.8
'VS 497-2/14'	77.33	11.50	6.73	7	68.4
'T. spelta 1139-16'	77.33	12.77	6.06	4	71.8
'T. spelta 1140-16'	74.67	12.30	6.07	3	68.8
'T. spelta 1145-16'	76.67	11.97	6.41	1	69.0
'Doridna'	77.33	11.73	6.59	4	69.9
'Pryvitna'	78.67	12.07	6.52	4	72.4
'Prynada'	83.33	12.03	6.93	3	68.5
'Vyhadka'	76.67	12.43	6.17	4	66.3
'Fermerka'	74.67	12.83	5.82	4	68.3
'Pronia'	77.67	12.23	6.35	4	65.9
'Doskonala'	77.33	12.37	6.25	4	66.6
'Rozkishna'	77.33	11.93	6.48	3	71.0
'Alians'	76.00	11.63	6.53	4	64.3
'Zdobna'	76.67	11.53	6.65	3	71.9
'Harmonika'	77.33	12.20	6.34	4	72.7
'Krasa laniv'	79.67	12.20	6.53	4	73.3
'Patriotka'	77.00	12.43	6.19	4	74.1
'Zapashna'	77.45	11.58	6.70	4	67.6
'Biliava' – St	86.70	10.10	8.60	6	54.6

ty: flour WAC, test baking and evaluation of the quality of cookies [22] (Table 2). The research results indicate that the quality of confectionery products varied significantly depending on the sample.

It is known that the larger the diameter of the biscuit and the smaller the thickness, the higher its quality [22]. The greatest ratio of the diameter of the biscuit to its thickness was obtained for wheat flour of the lines 'L137-26-0-3' – 8.68, 'L137-26-0-2' – 8.61, at the standard level – soft-grain varieties 'Biliava' and significantly higher in comparison with the standard hard grain variety 'Pryvablyva' – 6.48. These samples were characterized by low SPO rates of 53.1% and 52.7%, respectively. Hard-grain

winter wheat varieties 'Fermerka', 'Doskonala', 'Rozkishna', 'Alians', 'Zdobna', 'Harmonika', 'Krasa laniv', 'Patriotka' and a number of lines had a low ratio of cookie diameter to its thickness (5, 82-6.82) and, accordingly, high WAC rates of 65.9–74.1%. Lines *T. spelta* by confectionery parameters of flour are attributed to hard grain samples. Allelic state of genes *Pina-D1* and *Pinb-D1* of 40 varieties and winter wheat lines was determined (Table 3).

As a result of the study using allele-specific primers, the presence of a 330 bp fragment, that corresponds to the *Pina-D1a* allele of the puroindoline *a* gene [26], was revealed in 36 wheat samples. The absence of the amplification product on the electrophoregram corre-

Table 3

Allelic state of *Pin a* and *Pin b* genes of soft winter wheat samples

№	Sample name	<i>Pina-D1</i> allele	<i>Pinb-D1</i> allele	Amino acid
1	'Pryvablyva'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
2	'Metelytsia'	<i>Pina-D1b</i>	<i>Pinb-D1b</i>	serine
3	'Erytrospermum 533-16'	<i>Pina-D1b</i>	<i>Pinb-D1b</i>	serine
4	'Lyutestsens 652-16'	<i>Pina-D1b</i>	<i>Pinb-D1b</i>	serine
5	'L 139-03 KH'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
6	'Erytrospermum S 424-1/14'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
7	'Erytrospermum 1002-16'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
8	'Erytrospermum 1003-16'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
9	'L137-26-0-2'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
10	'L137-26-0-3'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
11	'L202-20'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
12	'S 492-3/14'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
13	'VS 2019-1/15'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
14	'VS 497-2/14'	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine
15	'T. spelta 1139-16'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
16	'T. spelta 1140-16'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
17	'T. spelta 1145-16'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
18	'Doridna'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
19	'Pryvitna'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
20	'Prynada'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
21	'Vyhadka'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
22	'Fermerka'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
23	'Pronia'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
24	'Doskonala'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
25	'Rozkishna'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
26	'Korovayna'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
27	'Aliyans'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
28	'Vasylyna' – St	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
29	'Astet'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
30	'Zdobna'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
31	'Harmonika'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
32	'Krasa laniv'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
33	'Patriotka'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
34	'Hayok'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
35	'Statna'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
36	'Zapashna'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
37	'Dyvo'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
38	'Bona Dea'	<i>Pina-D1a</i>	<i>Pinb-D1b</i>	serine
39	'Rheia' – St	<i>Pina-D1b</i>	–	–
40	'Myrleben' – St	<i>Pina-D1a</i>	<i>Pinb-D1a</i>	glycine

sponds to *Pina-D1b* allele, which was identified 'Erythrosperrum 533-16', 'Liutescens 652-16' in the samples of 'Metelytsia kharkivska', (Fig. 1).

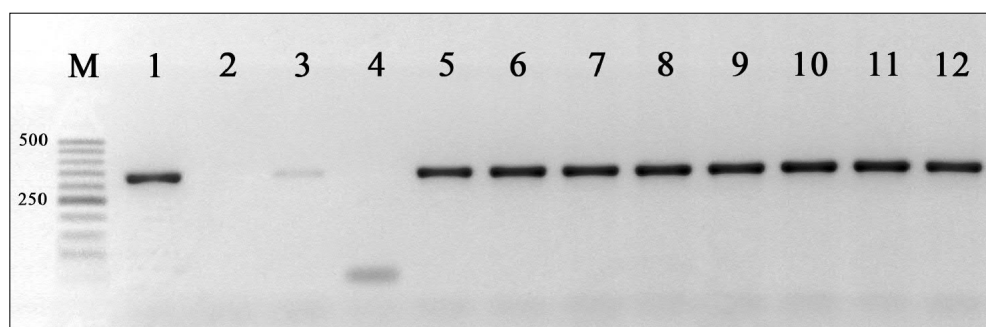


Fig. 1. Electropherogram of PCR products with specific primers for *Pina D1* gene:

M – molecular weight marker M 50 GENPAK®; 1 – 'Pryvablyva', 2 – 'Metelytsia Kharkivska', 3 – 'Erythrosperrum 533-16', 4 – 'Liutescens 652-16', 5 – 'L 139-03 KN', 6 – 'Erythrosperrum S 424- 1/14', 7 – 'Erythrosperrum 1002-16', 8 – 'Erythrosperrum 1003-16', 9 – 'L137-26-0-2', 10 – 'L137-26-0-3', 11 – 'L202-20', 12 – 'S 492-3/14'

As a result of the allelic state of the puroidoline *b* gene in the lines 'L139-03 KH', 'L137-26-0-2', 'L137-26-0-3' and others (Fig. 2), the presence

of glycine at position 46 of the polypeptide was determined, as evidenced by the amplification products of 250 bp.

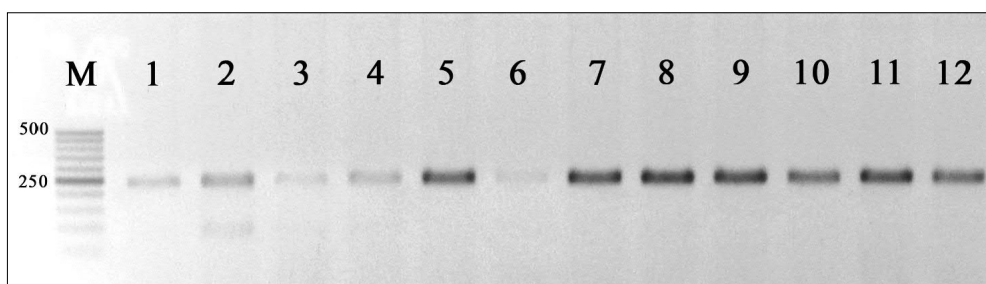


Fig. 2. Electropherogram of PCR products with primers to *Pinb-D1* alleles (see description to Fig. 1)

It is known that, despite the difference in allele-specific primer pairs by only one nucleotide, there is a possibility of errors in the determination of *Pinb-D1* alleles using primers to the glycine nucleotide sequence [27]. So, when carrying out PCR with primers that determine serine at position 46 of the *pin-b* protein, the amplification product of 250 bp in size was obtained in all studied varieties. Therefore, to obtain reliable information about the allelic state of *Pinb-D1* in the study, an

analysis was performed using the restriction endonuclease MbiI (BsrBI). Amplification was carried out using primers to *Pinb-D1* gene designed by Giroux et al. [25]; an amplification product of 447 bp was obtained, which was further cleaved with endonuclease. The enzyme recognizes the CCGICTC nucleotide sequence, which determines the change from glycine to serine. After incubation with restriction endonuclease amplification products 447 bp in the absence of mutation on the electrophoretogram,

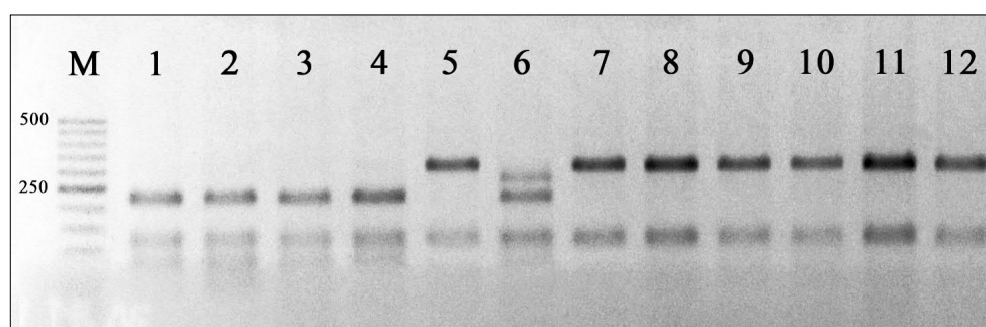


Fig. 3. Electropherogram of PCR products with primers to the nucleotide sequence of serine in determination of *Pinb-D1* alleles (see description to Fig. 1)

amplicons of 320 bp were observed. In genotypes where, as a result of the mutation, glycine was replaced by serine, the amplicons were 200 bp (Fig. 3).

The analysis revealed the presence of serine at position 46 of the puroindoline b protein in 30 out of 40 samples.

According to the results of PCR analysis of the allelic composition of puroindolinium genes out of a total of 37 samples, 9 (24%) – had the allelic composition of the puroindolinium genes characteristic for soft-grain varieties, *Pina-D1a* and *Pinb-D1a* ('L139-03KH', 'L137-26-0-2', 'L137-26-0-3', 'L202-20', 'ErythrospERMum 1002-16', 'ErythrospERMum 1003-16', 'S 492-3/14', 'VS 2019-1/15', 'VS 497-2/14'). soft winter wheat varieties 'MV Homber' (HUN), 'Warwik', 'Webster' (CAN), 'Eva' (SVK), 'FS 401' (USA), 'Oksana', 'Blond' (UKR) were used for these lines creating. The best confectionery indicators among soft-grain samples were in the lines 'L137-26-0-2', 'L137-26-0-3', which had WAC of less than 55%, biscuit diameter – 85 mm, height – 10 mm, surface assessment cookies – 8–9 points. The remaining 7 lines were characterized as satisfactory by the confectionery properties of flour.

76% of the samples had the alleles of *Pina-D1* and *Pinb-D1* genes, which are typical for hard-grain varieties. In the studied selection of hard-grain samples, the *Pina-D1* gene is represented by two alleles: *Pina-D1a* and *Pina-D1b*. 27 samples of winter wheat had *Pina-D1a* allele, which allows to use them in breeding for soft-grain quality when crossed with soft samples, 3 samples had *Pina-D1b* allele ('Metelytsia Kharkivska', 'ErythrospERMum 533-16', 'Liutescens 652-16'). All hard-grain samples had *Pinb-D1b* allele in *Pinb-D1* gene, and the 'ErythrospERMum S 424-1/14' line was heterogeneous. These samples had quality indicators characteristic of hard-grain wheat: WAC – 68% and above, diameter – 60–72 mm, biscuit height – 13–15 mm, surface score – 1–4 points.

T. spelta lines, involved in the analysis, both for alleles of puroindolinium genes and for the technological confectionery properties of flour, were included in the group of hard-grain samples.

Conclusions

The allelic state of *PINA-D1* and *Pinb-D1* genes was revealed for 37 varieties and lines of soft winter wheat breeding of Plant Production Institute nd. a. V. Ya. Yuriev NAAS.

Lines 'L139-03KH', 'L137-26-0-2', 'L137-26-0-3', 'L202-20', 'ErythrospERMum 1002-16', 'ErythrospERMum 1003-16', 'S' 492-3/14', 'VS 2019-1/15', 'VS 497-2/14' had an allelic state of the puroin-

dolinium genes characteristic for soft-grain varieties (*Pina-D1a*, *Pinb-D1a*), good and satisfactory confectionery indicators of flour quality.

76% of the studied cultivars had the allelic composition of puroindoliniums *Pina-D1a/Pina-D1b*; *Pinb-D1b*, i.e. were characterized as hard-grain samples of *Triticum aestivum* L.

The studies allowed to differentiate the created linear material and transfer for the qualification examination the soft winter wheat variety of confectionery use 'L137-26-0-3' ('Mazurok') with a genetically confirmed soft-grain structure of endosperm and high confectionery properties of flour; transfer the lines of winter wheat with high confectionery properties for registration to the NCPGRU.

'Mazurok' ('L 137-26-0-3') is a soft-grain variety of winter wheat for confectionery use. The variety is high-yielding (7.93 t/ha), highly resistant to damage by leaf septoria (7 points), powdery mildew (7 points); its frost resistance is 7 points. According to technological indicators, it is referred to soft red winter wheat, and has increased confectionery properties: low protein content in grain – 11%, vitreousness – 25%, flour strength – 73 w. i., WAC – 53%, high biscuit linear dimensions, D/T ratio, high biscuit surface score – 9 points.

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Леонов О. Ю., Шарипіна Я. Ю., Усова З. В., Суворова К. Ю., Сахно Т. В. Алельний склад генів пуриноидолінів та кондитерські властивості борошна зразків пшениці м'якої озимої. *Plant Varieties Studying and Protection*. 2020. Т. 16, № 2. С. 217–225. <https://doi.org/10.21498/2518-1017.16.2.2020.209258>

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Мета. Ідентифікувати за алельним станом гени *Pina-D1* і *Pinb-D1* сортів та ліній пшениці м'якої озимої селекції Інституту рослинництва ім. В. Я. Юр'єва НААН для цільового використання в селекції на високі кондитерські показники борошна. **Методи.** Алельний стан генів *Pina-D1* і *Pinb-D1* ідентифікували методом полімеразної ланцюгової реакції (ПЛР) з використанням алель-специфічних пар праймерів. Кондитерські властивості борошна оцінювали, визначивши показники якості: водопоглинальну здатність борошна (ВПЗ), пробне випікання печива та оцінювання його якості. **Результати.** За результатами ПЛР-аналізу 9 зразків мали алельний склад генів пуриноидолінів (*Pina-D1a* і *Pinb-D1a*), характерний для м'якозерних сортів. Кращим за кондитерськими властивостями було борошно ліній 'L137-26-0-2', 'L137-26-0-3', воно мало показник ВПЗ менший 55%, діаметр печива 85 мм, висоту – 10 мм, оцінку поверхні – 7–9 балів, що відповідало вимогам до м'якозерних пшениць. 76% зразків належали до твердозерних сортів та мали відповідні алелі генів *Pina-D1* або *Pinb-D1*. У дослі-

дженій вибірці ген *Pina-D1* був представлений 2 алелями: *Pina-D1a* та *Pina-D1b*. 27 зразків мали алель *Pina-D1a*, що також дозволило використовувати їх в селекційних програмах на якість зерна при схрещуванні зі зразками типу *soft*, 4 – алель *Pina-D1b*. За геном *Pinb-D1* всі твердозерні зразки мали алель *Pinb-D1b*, а лінія 'Еритроспермум S 424-1/14' була гетерогенною *Pinb-D1a/Pinb-D1b*. Борошно цих зразків мало характерні для твердозерної пшениці показники якості: ВПЗ 68% і більше, діаметр печива 60–72 мм, висота – 13–15 мм, оцінка поверхні – 1–4 бали. **Висновки.** Виконані дослідження дозволили ефективно диференціювати селекційний матеріал і передати на кваліфікаційну експертизу сорт пшениці м'якої озимої кондитерського напряму використання 'L137-26-0-3' ('Мазурок'), який має алельний склад генів пуриноидолінів (*Pina-D1a* і *Pinb-D1a*), характерний для м'якозерних сортів, та високі кондитерські властивості борошна.

Ключові слова: пшениця м'яка озима; сорт; лінія; гени *Pina-D1* і *Pinb-D1*; водопоглинальна здатність; печиво.

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Леонов О. Ю., Шарыпина Я. Ю., Усова З. В., Суворова Е. Ю., Сахно Т. В. Аллельный состав генов пуриноидолинов и кондитерские свойства муки образцов пшеницы мягкой озимой // *Plant Varieties Studying and Protection*. 2020. Т. 16, № 2. С. 217–225. <https://doi.org/10.21498/2518-1017.16.2.2020.209258>

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Цель. Идентифицировать аллельное состояние генов *Pina-D1* и *Pinb-D1* сортов и линий пшеницы мягкой озимой селекции Института растениеводства им. В. Я. Юрьева НААН для целевого использования в селекции на высокие кондитерские показатели муки. **Методы.** Аллельное состояние генов *Pina-D1* и *Pinb-D1* идентифицировали методом полимеразной цепной реакции (ПЦР) с использованием аллель-специфических пар праймеров. Кондитерские свойства муки оценивали, определив показатели качества: водопоглотительную способность муки (ВПС), пробную выпечку печенья и оценку его качества.

Результаты. По результатам ПЦР-анализа 9 образцов имели аллельное состояние генов пуриноидолинов, характерное для мягкозерных сортов – *Pina-D1a* и *Pinb-D1a*. Лучшей по кондитерским свойствам была мука линий пшеницы 'L137-26-0-2', 'L137-26-0-3', она имела показатель ВПС меньше 55%, диаметр печенья 85 мм, высоту – 10 мм, оценка поверхности печенья составляла 7–9 баллов, что соответствовало требованиям к мягкозерным пшеницам. 76% изученных образцов относились к твердозерным сортам и имели соответствующие аллели генов *Pina-D1* или *Pinb-D1*. В опытной выборке образцов ген

Pina был представлен 2 аллелями: *Pina D1a* и *Pina D1b*. 27 образцов имели аллель *Pina D1a*, это также позволило использовать их в селекции на качество зерна при скрещивании с сортами типа *soft, 4* – аллель *Pina D1b*. По гену *Pinb* все твёрдозёрные образцы имели аллель *Pinb D1b*, а линия 'Эритроспермум S 424-1/14' была гетерогенной *Pinb D1a/Pinb D1b*. Эти образцы имели соответствующие твёрдозёрным пшеницам показатели качества муки: ВПС 68% и выше, диаметр печенья 60–72 мм, высота – 13–15 мм, оценка поверхности – 1–4 балла. **Выводы.** Проведен-

ные исследования позволили эффективно дифференцировать селекционный материал и передать на квалификационную экспертизу сорт пшеницы мягкой озимой кондитерского направления использования 'L137-26-0-3' ('Мазурок'), который имеет аллельный состав генов puro-индолинов (*Pina-D1a* и *Pinb-D1a*), характерный для мягкозёрных сортов, и высокие кондитерские свойства муки.

Ключевые слова: пшеница мягкая озимая; сорт; линия; гены *Pina* и *Pinb*; водопоглотительная способность; печенье.

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