

Food Chemistry 79 (2002) 251–254

www.elsevier.com/locate/foodchem

Food Chemistry

Analytical, Nutritional and Clinical Methods Section

Analysis of β -glucan content in barley cultivars from different locations of China

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Received 4 September 2001; received in revised form 9 January 2002; accepted 9 January 2002

Abstract

b-glucan content of barley cultivars collected from various areas of China, as well as from Canada and Australia, were assayed by an enzymatic method. Meanwhile a multi-location trial was conducted to determine b-glucan content of eight barley cultivars in each location. For 164 cultivars, originating from China, β -glucan content ranged from 2.98% for Sumei 21 to 8.62% for OB25, with a mean of 4.58%. Tibet barleys, all of the hull-less type, had the highest β -glucan content. Winter barley cultivars from various areas of China had basically the same β -glucan content as those from Canada and Australia. The seeds produced in Hangzhou had lower b-glucan contents than those produced locally. The effects of cultivars, environments (locations) and their interactions on bglucan content variation was all highly significant, with the cultivars having the largest effect. The highly significant interaction between cultivars and environment suggests that it is important to plant the appropriate cultivar in a particular area in order to obtain barley seeds with reasonable β -glucan contents. \odot 2002 Elsevier Science Ltd. All rights reserved.

Keywords: Barley; b-glucan; Cultivar; Environment

1. Introduction

The mixed-linked $(1\rightarrow 3)$, $(1\rightarrow 14)$ - β -D-glucan (hereafter referred to as simple β -glucan) is a cell wall polysaccharide found only in grass and cereals (Carpita, 1996). Compared with other cereals, barley and oats have relatively high β -glucan contents. In barley, β -glucan has become a growing concern at multi-discipline levels due to its positive role in human health when used as food and its potential deterioration on malting. In the brewing industry, a high content of β -glucan in barley may lead to problems such as reduced rate of wort filtration and haze formation in beer, and adversely affect the recovery of malt extract (Barrett, Clapperton, Divers, & Rennie, 1973; McCleary & Glennie-Homes, 1985). The high viscosity of aqueous solutions of barley β -glucan has also been implicated in the problems encountered in the stock-feed industry; therefore it is considered as a anti-nutritional factor, particularly in chicken and poultry diets where it may give sticky droppings and affect nutritional intake,

growth and feed conversion efficiency (Hasselman & Thomke, 1982; Wang, Newman, Newman, & Hofer, 1992). On the other hand, positive roles of β -glucan in human nutrition and health, such as lowering of blood cholesterol and blood glucose, increasing mineral and vitamin bio-availability and controlling colon cancer, have been reported (Klopfenstein, 1988; Martinez, Newman, & Newman, 1991).

Variation in barley β -glucan content is associated with genetically controlled starch and hull type. High β glucan content is reported in hull-less cultivars (Bengtsson, Aman, Graham, Newman, & Newman 1990; Bhatty, MacGergor, & Rossnagel, 1991) and in cultivars with 100% amylopectin (waxy) starch (Ullrich, Clancy, Eslick, & Lance, 1986; Xue, Newman, Newman, & Maguire, 1991). Environment also influences β glucan content (Fastnaught, Berglund, Holm, & Fox, 1996; Narasimhalu, Kong, Choo, Ferguson, Therrien, Ho, May, & Jui, 1995; Stuart, Loi, & Fincher, 1998). Barley b-glucan contents, reported so far, have ranged from less than 2% to more than 10%, and variation is not only caused by genetics and environment, but is also due to the analysis methodology. The earlier analytical method, by measuring viscosity gave high values, while the enzymatic method, developed by McCleary and

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Glennie-Homes (1985), by use of a specific assay kit, has proved to be a simple procedure given good repeatability.

Barley is one of the most important cereal crops in China; it grows in widely ecological locations and finds manifold uses. Commonly, it is used as a material for malting and feed processing while, in Tibet, barley is a stable food crop. However, there has been little research on b-glucan content in China. The objective of this study was to determine β -glucan content of barley cultivars from different locations and to examine its variation as affected by genotypes and environments.

2. Materials and methods

One-hundred and eighty-five barley cultivars or breeding lines were collected from the main productive areas of China. Among them, 75 cultivars (lines) came from Tibet (kindly provided by Professor Qiang X-L at the Tibet Institute of Agricultural Research, Lasa, 91.1° E, 29.6° N), 27 from Xingjian (kindly provided by Professor Cao L-P at the Agricultural college, Shihezi University; Shihezi, 87.6° E, 43.7° N), and others from Jiangsu (Yanchen, 118.2° E, 31.5° N), Hunan (Zhangsai, 1130° E, 28.2° N), Hubei (Wuhan, 114.2° E, 30.6° N), Sichuan (Chengdu, 104.1° E, 30.6° N), Shandong (Taian, 117.2° E, 36.2° N), Gansu (Lanzhou, 103.8° E, 36.05° N) and Zhejiang (Hangzhou, 120.2° E, 30.5° N). All of the Tibet barleys were of the hull-less type while other Chinese barleys were the hulled type. Except for Xinjiang, where barley is sown in spring, in all other regions it is sown in winter. In addition, 21 commercial barley cultivars were collected from Canada and Australia for comparative study.

All collected cultivars or lines were planted at the experimental farm of Zhejiang University (Huajia Chi Campus) in early November 1999. The materials were grown in two replicates of two-row plots, 2 m long, 30 cm between rows and 100 seeds per row. The trial was

managed according to locally recommended cultural practice. At maturity, thirty spikes of main shoots were harvested from the cultivars with similar mature dates.

According to a previous experiment (Wang, Zhang, Chen, & Ding, 1998), eight two-rowed barley cuitivars with similar growth durations and substantial differences in β -glucan contents were used and grown at six diverse environments of Zhejiang in the 1999–2000 growing season. All cultivars were grown adjacently in the same field and each cultivar consisted of three 2-mlength rows with three replicates. At maturity, 30 spikes of main shoots were harvested from each replicate.

Grain samples were ground through a 0.5-mm sieve in a Cyclotec mill after drying in an oven for 24 h. Total bglucan was analyzed according to the McCleary and Glennie-Holmes (1985) methods using a commercial kit (Megazyme Ltd. Ireland).

3. Results and discussion

There were great differences in β -glucan contents among the samples, both from the different areas and within the same location (Table 1). For 164 cultivars (lines) from China, β -glucan content ranged from 2.98% for Sumei 21, a commercial cultivar in Jiangsu province, to 8.62% for QB25, a breeding line in Tibet. On average, for all cultivars, the value was 4.58% and approximately two third of samples fell into the range 3.0–5.0%. In terms of location distribution, the cultivars or lines with β -glucan contents higher than 8% were found only in the samples from Tibet, in which the mean of all 75 cultivars was 5.25%, ranked the highest among various collections, and 20% of samples had β glucan contents higher than 6%. The mean content of 27 Xinjiang cultivars was 5.09%, this being the second highest among collections. Although no cultivar containing more than 8% of β -glucan was found in Xinjiang barley samples, its proportion of cultivars with

Table 1

Frequency distribution of barley cultivars originating from the different areas on the basis of grain b-glucan content

| β -Glucan content | Total | Tibet | Xinjiang | Other areas in China | | Canada and Australia |
|-------------------------|-------|------------------|--------------|---------------------------|----------------------------|----------------------|
| | | | | Seeds from original areas | Seeds produced at Hangzhou | |
| | | | | $\frac{0}{0}$ | | |
| <3.0 | 5.34 | $\left(\right)$ | θ | 3.23 | 12.9 | 14.3 |
| $>$ 3.0 $-$ < 4.0 | 28.6 | 8.0 | 7.41 | 45.2 | 67.7 | 38.10 |
| $>4.0-5.0$ | 37.4 | 46.7 | 51.9 | 35.5 | 16.1 | 28.6 |
| $> 5.0 - 6.0$ | 18.0 | 24.0 | 18.5 | 16.13 | 3.23 | 19.1 |
| $>6.0 - < 7.0$ | 6.8 | 12.0 | 18.5 | θ | | θ |
| $>7.0 - 8.0$ | 2.91 | 6.67 | 3.7 | θ | | θ |
| > 8.0 | 0.97 | 2.67 | $\mathbf{0}$ | 0 | | 0 |
| Sample number | 164 | 75 | 27 | 62 | 62 | 21 |
| Mean | 4.58 | 5.25 | 5.09 | 4.24 | 3.61 | 4.02 |
| Standard deviation | 1.08 | 1.06 | 0.92 | 0.67 | 0.55 | 0.81 |

more than 6% of β -glucan was nearly as high as that in Tibet samples. Sixty-two cultivars from other areas of China had a 4.24% mean β -glucan content, obviously lower than those both from Tibet and Xinjiang. Compared to original seeds, the Hangzhou-productive seeds were significantly lower in β -glucan content, suggesting the modification of β -glucan content by environment. Twenty-one beer cultivars from Canada and Austria were within the range 2.76% for AB168–11 to 5.14% for Gateways 3, with a mean content of 4.02%.

The wide diversity in β -glucan content could be significant in the commercial utilization of barley. Since the enzymatic method of measuring β -glucan content was developed by McCleary and Gennie-Holmes (1985), grain b-glucan contents among barley cultivars have been reported to range from 3.0 to 8.2% (Aman & Graham, 1987; Carr, Glatter, Jeraci, & Lewis, 1989; Fastnaught et al., 1996; Henry, 1987; Lehtonen & Aikasalo, 1987). The wider range of β -glucan content for the barley collections in this study may be attributed to more samples and the greater differences in the environments of a particular cultivar origin. The higher b-glucan content in Tibet barley cultivars may be at least partly related to their hulless type and is desirable in light of their use as feed.

In China, more and more malt barley is being imported with the increase of beer production, as local barley is considered to be inferior, in malting quality, to imported barley. One of the reasons for inferior quality is attributed to high β -glucan content. In this experiment, we found that most of the cultivars, except for those from Xinjiang, had β -glucan contents not higher than those from Canada and Australia, indicating that b-glucan content is not a predominant factor leading to inferior malting quality in Chinese-productive barley. Moreover, the barley grown in Zhejiang had a much lower β -glucan content. Therefore, the higher β -glucan content in malt for Chinese-productive beer barleys should be attributed to lower β -glucanase activity in grains.

Variance analysis for β -glucan content of eight barley cultivars, grown at six locations in Zhejiang, showed that the effects of cultivars, locations and their interaction were all highly significant (Table 2). Comparatively,

Table 2 Variance analysis on β -glucan contents of eight barley cultivars grown at six locations

** Significant at 0.01 probability level.

cultivars shared a larger contribution to the variation in b-glucan content than locations or interaction.

On average in six locations, β -glucan content of eight cultivars in this study ranged from 3.24% for Zhepi 4 to 4.62% for Xue 92–47, with a mean of 3.95% (Table 3). The cultivars differed widely in the coefficient of variation (CV) when planted at different locations. Thus, Xuemei 3 and ZAU 7 had the greatest and least CV, respectively, suggesting a great difference among cultivars in their response to environment in terms of β -glucan content.

On average among eight cultivars, b-glucan content of six locations ranged from 3.86% for Yuyao to 4.14% for Teizhou (Table 4). Although the locations also differed significantly, the absolute difference between the highest and lowest was much smaller than that between cultivars. Coefficient of variation in β -glucan content for eight cultivars grown within a location was smaller than for a cultivar grown in the different locations.

Genetic and environmental variations in the B-glucan content of barley have been reported in several studies (Aastrup, 1979; Henry, 1986; Molina-Cano, Francesch, Perez-Vendrell, Ramo, Voltas, & Brufau, 1997). Genetic factors have been found to be the more important (Henry, 1986; Molina-Cano et al., 1997). In this study, significant differences among cultivars are also found, irrespective of planting locations, and cultivars differ in the variation of β -glucan content over locations. The

Table 3

Means and coefficients of variation of β -glucan contents corresponding to barley cultivars grown at six locations

| Cultivar | β -Glucan% | $CV\%$ |
|------------|-------------------|--------|
| Xu92-47 | 4.62 ^a | 12.3 |
| ZAU 7 | 3.90 _b | 9.81 |
| Xumai 3 | 4.54a | 21.9 |
| ZAU6 | 4.04 _b | 12.2 |
| $92 - 11$ | 3.81b | 11.7 |
| ZAU 3 | 3.49c | 10.4 |
| Zhepi 4 | 3.92 _b | 18.8 |
| Zheyuan 18 | 3.24d | 11.9 |

^a Means within a column followed by a different letter are significantly different $(P<0.05)$.

Table 4

Means and coefficients of variation of β -glucan contents of eight barley cultivars grown at each location

| Location | $Mean\%$ | $CV\%$ | |
|----------|--------------------|--------|--|
| Hangzhou | 4.07a ^a | 7.45 | |
| Jiaxing | 4.06a | 6.03 | |
| Yuvao | 3.68 _b | 7.84 | |
| Ouzhou | 3.99a | 8.94 | |
| Taizhou | 4.14a | 7.82 | |
| Lishui | 3.75 _b | 7.68 | |

^a Means within a column followed by a different letter are significantly different $(P<0.05)$.

interaction of cultivar and location is also highly significant, suggesting the importance of cultivar selection in a particular area.

Acknowledgements

The authors are deeply indebted to both the National Natural Science Foundation of China (No. 39870506) and Zhejiang Natural Science Foundation (No. 397259) for their support to this project.

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