Effects of Long Term Storage Factors on Nutritional Value of Conserved Barley Accessions in Ethiopia

Asnakech Senbeta*, Dejene Dida

Crop & Horticulture Biodiversity Directorate, Ethiopian Biodiversity Institute, Addis Ababa, Ethiopia

Email address:

asni.senbeta@gmail.com (A. Senbeta)

*Corresponding author

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Abstract: Ethiopia is one of the world centers of genetic diversity and origin of barley crop. Now a day the country is conserve many cereal crops under cold room storage, gene bank. One of the conserved crops is barley. Barley is the cereal crop, that mainly grown in the central highland of Ethiopia. The samples of 45 barley accessions were used for the current analysis. The aim of this study is to evaluate the effects of long term storage factors on nutritional value of barley that tested in 1993 and compare the results with nutritional value of barley that tested after 25 years (2018). Sample preparation and nutritional values were determined by using appropriate standard official procedures, AOAC (Association of Official Analytical Chemists). Descriptive graphs and paired t-test were used to analyze the data obtained on nutritional values of barley crop from two test years. The nutritional value of barley crop after 25 years of storage time was decreased as compared with its nutritional value tested in 1993, except total carbohydrate. As a result, the study found statistically significant difference of average percentage content of crude protein, total mineral ash, moisture and total carbohydrate between 1993 and after 25 years of storage time. The current study concluded that, when the storage time increases, the nutritional values of barley crop were significantly decreased for all nutritional values except crude fat content and total carbohydrate.

Keywords: Barley, Ethiopia, Long Term Storage, Nutritional Value, Paired T-test

1. Introduction

Ethiopia is known as a center of genetic diversity and origin for a number of cultivated and wild plants [1, 2]. The richness and range of genetic diversity in Ethiopia, particularly of landraces, is currently subject to serious genetic erosion and irreversible losses due to the changing nature of agricultural production. Widespread adoption of modern varieties, technological change (such as use of fertilizer and irrigation), land use change, habitat destruction and drought, among other important factors have lowered the demand for landraces adapted to marginal growing conditions in Ethiopia [3-5]. With the objective of addressing conservation of this diminishing plant genetic diversity in the country, the Plant Genetic Resources Centre of Ethiopia (PGRC/E), now the Ethiopian Biodiversity Institute (EBI) was established in 1976. The primary mandates of Ethiopian Biodiversity Institute include the preservation of genetic diversity of crop plants, their wild relatives, and native

species important to Ethiopian agriculture and biodiversity.

The conservation of plant genetic materials depended on collection. The collection priorities have been expeditions for agriculturally important, rare and threatened genotypes, to minimize genetic erosion and loss of varieties. Over 84 000 accessions from more than 120 plant species have been collected from across the country and preserved ex-situ at Ethiopian Biodiversity Institute. The ex-situ collection is managed in accordance with international standards for gene banks, with optimum treatment developed for seeds of most food crops [6]. The gene bank serves as a reservoir of genes potentially useful for many purposes, including resistance to diseases, pests and other environmental stresses, as well as for traits that increase yield or food quality. The storing seeds requirements are adequately dried (moisture contents as low as 5% for starchy seeds such as barley), careful production of high quality seed for maximum longevity, and carefully avoiding any sub-optimal effect of very low moisture content [7].

Hermetically sealed aluminum foil bags are used for long-term (-10°C) and short-term (+4°C) seed storage. From the base and active collections, the seeds are regularly regenerated and multiplied, based on thresholds for viability or seed quantity. In addition to this, for some of conserved materials the nutritional values were analyzed. One of the most conserved materials in Ethiopian Biodiversity Institute is Barley. Barley (*Hordeum vulgare* L.) is the fourth most important cereal crop worldwide, after wheat, corn and rice, belonging to family Poaceae [8]. Ethiopia is a center of diversity for barley, because of the presence of great diversity in ecology [9]. It has a long history of cultivation in the highlands [10]. The principal uses of barley are as feed for animals, in the form of barley meal, and as grain for malting and brewing in the manufacture of beer and whisky [11, 12]. But barley in Ethiopia is mainly used for making local recipes and drinks, such as: Bread, Injera, kolo, Genfo, Animal feed, Beso, Tela and Borde. Over the last forty years Ethiopian Biodiversity Institute was conserved 16,224 accessions of barley, by ten years interval viability tested were done and the status of the material is evaluate, when it failed below standard it must be regenerate and multiply it. Accessions are tested and monitored both at the conserved time and during conservation in order to ensure the quality of the material in the gene bank. In the case of nutritional values analysis some of barley accessions were analyzed at the conserved time (25 years ago) but the conserved material was not analyzed again for the second time or during conservation. Due to this reason do not know the statuses of the nutritional values of the conserved materials were distorted or not. Now the current study is fill this research gaps. 45 accessions of conserved barley materials were used for analysis and observe the difference of nutritional values of the same barley accessions that analyzed 25 years ago. And compare the nutritional values performance of the results of these accessions. Therefore the objective of this study was to evaluate and determine the significant difference of the effect of long-term storage conditions on nutritional value of conserved barley grain at gene bank. This help to profile the overall barley nutritional information and long term storage effect for both consumption and research purpose.

2. Materials and Methods

2.1. Sample Size and Sample Preparations

45 barley accessions were used for the current research experiment. 30 g of each samples accession were received from cold room storage of Ethiopian biodiversity institute. The Seeds were hand-sorted to clean and remove foreign materials. The raw barely seeds prepared without removing the seed coat and made to flour with electric grinder or Retch sample mill (Retsch, ISO9001) until to pass through 0.5 mm sieve mesh size, and finally packed into airtight polyethylene plastic bags. The nutritional analysis was performed in 2 replicates. All laboratory experiment was conducted in

Ethiopian biodiversity institute

2.2. Nutritional Content Analysis

The nutritional value of the samples was carried out according to Association of Official Analytical Chemists (AOAC, 2000) method for nutrient analysis [13].

2.2.1. Determination of Moisture Content

The moisture content of the barley samples was determined by oven drying method. A clean, empty aluminum dishes, and its lids were dried in drying oven (Memmert, Germany) at 100°C for 1hr, and cooled in a desiccator (with granular silica gel) for about 30 min, and weighed. The samples prepared for each accession were mixed thoroughly, and about 3.0 g of flour was weighed in duplicate. The dishes and the samples were placed in the drying oven, and dried for 3hr at 105°C. After drying, the samples were cooled in a desiccator for 30 min, and weighed it. Then percentage of moisture calculates as follow:

$$\% \text{ Moisture} = \frac{((\text{wt of empty dishes} + \text{sample}) - (\text{wt of dishes} + \text{dry sample})) \times 100}{\text{wt of sample}} \quad (1)$$

2.2.2. Determination of Crude Protein

The Protein content of the barley flour samples was determined by the Kjeldahl method. 1g of barely sample was used for this process, the experiment was passed three steps, such as digestion, Distillation and titration was involved. All nitrogen is converted to ammonia by digestion with a mixture of concentrated sulfuric acid and Kjeldahl tablet containing copper sulfate and potassium sulfate as a catalyst by 420°C temperature. The ammonia released after alkalization with sodium hydroxide is steam distilled into boric acid and titrated with 0.1N hydrochloric acid. Then the result is calculated as based on conversion factor form Nitrogen to Protein. Protein content (% N × 5.83) was determined by the Kjeldahl method described by [13].

$$\% \text{ total Protein} = (\% \text{ N} \times 5.83) \quad (2)$$

Where,

% N = Percentage of Nitrogen.

5.83 = barley conversion factor.

2.2.3. Determination of Crude Fat Content

The crude fat content of barley flour samples was determined according to AOAC (2000) by Soxtec using Petroleum ether (b. pt 40-60°C) solvent. About 3.00 g of barley flour sample was added into the extraction thimbles and covered it with fat free cotton. The thimbles with the sample content were placed into soxtec extraction chamber and then switched on the heater. The cooling water was switched on, and a 25 ml of Petroleum ether was added to the extraction can and attached it into corresponding thimbles in soxtec extraction chamber. Over all extraction processes was conducted for about 1hr including solvent recover and evaporate. Then the extraction can with their content were removed from the extraction chamber and placed in the

drying oven at 100°C for about 30 min, cooled to room temperature in the desiccator for about 30 min and weighed it. The amount of fat was quantified gravimetrically and calculated from the difference in weight of the extraction can before and after extraction as percentage.

Calculate the fat content as follows:

$$\% \text{Crude fat} = \frac{((\text{weight of can} + \text{fat}) - (\text{weight of can})) \times 100}{\text{weight of sample}} \quad (3)$$

2.2.4. Determination of Total Ash Content

Total ash content of barely flour samples was determined by the dry ash method in a muffle furnace according to AOAC (2000). The crucible were placed in the muffle furnace (Carbolite CSF 1200) at 550°C overnight to ensure that impurities on the surface of crucible are burned off and the crucible cooled in the desiccator for 30 min. Then the crucible was weighed. About 3.0 g of barely flour of each sample in duplicate was added into each crucible, then put it in to the muffle furnace and heated at 550°C overnight. Finally, taken out of the crucible from the Muffle Furnace and immediately place it in a desiccator till cooled to room temperature, and each crucible with ash was weighed. Weight of total ash was calculated by difference, and expressed as percentage of samples.

Calculate the percentage ash content (wet weight basis) as follows:

$$= \frac{\frac{\% \text{ ash (wet)}}{((\text{weight of crucible} + \text{ash}) - (\text{weight of crucible}))} \times 100}{\text{weight of sample}}$$

Calculation of ash content on dry basis (when moisture content is known) as follows:

$$\% \text{ASH (dry)} = \frac{\% \text{ ash (wet)} \times 100}{(100 - \% \text{ moisture})} \quad (4)$$

2.2.5. Total Carbohydrate Content

Total Carbohydrate content was obtained by difference from the combined percent of moisture, protein, ash and fat from 100. The result was calculated as follows;

$$\text{Total Carbohydrate content} = 100 - \{\text{moisture (g \%)} + \text{crude protein (g \%)} + \text{total fat (g \%)} + \text{total ash (g \%)}\} \quad (5)$$

Where (g %) = grams per 100 grams.

3. Statistical Analysis

All laboratory experiments were carried out duplicate for each extract (n=2). The mean, tables, graphs, charts and standard deviation (SD) of means were used to describe the research data. The statistical analyses were performed by paired t-test of mean comparisons at 5% level of significance.

4. Results and Discussion

The nutritional values of 45 conserved barley accessions

were tested in 1993 were evaluated for the second time on the same barley accession after 25 years in 2018. The result of all accessions analysis for nutritional values was presented in the table 1. The long term storage time was influenced the nutritional composition results of the barley samples. The graph illustrate below shows different result of the same accession by different years of test; crude protein, crude fat, total mineral ash, moisture content and total Carbohydrate content. As indicated in the figure 1, the percentage content of crude protein was slightly decreases and fluctuates as storage time increase. The highest percentage of crude protein (%15.43) was obtained in accession number 4963 that test in 1993, and the lowest (%6.11) was observed in accession number 4988 that test in 2018. Therefore a storage time is affecting the percentage content of crude protein in barley crop. On contrary, the figure 2 shows the percentage content of crude fat was fluctuate randomly when the storage time increases. But the highest percentage of crude fat (%2.90) was determined in accession number 4907 in test year 1993 whereas, the lowest (%1.61) was determined in accession number 4944 in 1993 and accession number 4995 in the test year of 2018. More or less a storage time is affecting the percentage content of crude fat in barley.

Moreover, figure 3 shows, the percentage content of total mineral ash was slightly decreased and sometimes fluctuated as storage time increase. The highest percentage of total mineral ash (%3.04) was obtained in accession number 4927 that test in 2018, and the lowest amount of total mineral ash content (%2.11) was observed in accession number 4961 that test in 2018. But percentage of total mineral ash decreases as storage time increases. It indicates that, a storage time is affecting the percentage of total mineral ash in barley. Similarly, in figure 4, it indicates percentage of moisture content was decreased and sometime fluctuated as storage time increase. Thus, it indicate that when storage time increase, the conserved material is loss its moisture. The highest percentage of moisture content (%11.59) was determined in accession number 4907 that test in 2018, and the lowest amount of moisture content (%6.30) was obtained in accession number 4923 that test in 2018. Logically, moisture content decrease as storage time increase. So that storage time is affecting the moisture content of barley accession. Furthermore, figure 5 indicates that percentage of total carbohydrate content was increase as storage time increase. The highest percentage of total carbohydrate (%80.02) was obtained in accession number 4988 that test in 2018, and the lowest percentage of total carbohydrate content (%69.28) was obtained in accession number 4963 that test in 1993. But percentage of total carbohydrate content increase as storage time increase. This is due to the material loss its moisture content and the substance concentrates. Therefore storage time is affecting total carbohydrate content of barley.

Table 1. Nutritional values of Barley accession result and year of test.

No	Accession No.	% Crude Protein		% Crude Fat		% Mineral ash	
		1995 GC	2018 GC	1995 GC	2018 GC	1995 GC	2018 GC
1	4907	11.89	12.48	2.90	1.77	2.30	2.99
2	4908	10.60	11.85	1.82	1.84	2.42	2.57
3	4919	12.31	10.77	1.75	2.36	2.33	2.49
4	4921	13.14	11.68	1.82	1.70	2.23	2.35
5	4923	12.97	12.17	1.85	1.70	2.22	2.69
6	4927	13.40	11.77	1.84	1.63	2.75	3.04
7	4929	12.60	12.34	1.89	1.76	2.32	2.56
8	4939	10.00	11.97	1.97	1.86	2.61	2.58
9	4942	13.11	11.77	1.78	1.73	2.68	2.41
10	4943	12.56	11.11	1.98	1.79	2.62	2.66
11	4944	12.85	11.09	1.61	2.01	2.69	2.50
12	4945	11.99	10.77	2.04	1.86	2.46	2.34
13	4951	12.38	10.80	1.70	1.71	2.31	2.65
14	4952	11.26	11.14	1.70	1.86	2.53	2.41
15	4955	11.42	12.09	1.78	1.87	2.52	2.24
16	4958	11.53	12.34	1.87	1.90	2.66	2.31
17	4960	12.22	13.13	1.68	2.24	2.80	2.20
18	4961	12.46	12.18	1.71	2.26	2.53	2.11
19	4963	15.43	11.04	1.82	1.85	2.74	2.35
20	4966	12.29	11.12	1.75	1.83	2.71	2.41
21	4967	13.66	11.61	1.79	1.97	2.70	2.28
22	4969	12.19	10.59	1.75	1.94	2.70	2.36
23	4970	12.45	10.86	1.80	1.95	2.65	2.39
24	4972	11.42	10.82	1.72	1.77	2.59	2.47
25	4973	12.14	11.12	1.69	2.38	2.57	2.34
26	4974	12.93	12.67	1.72	1.90	2.45	2.34
27	4976	12.89	12.68	1.93	1.81	2.63	2.36
28	4977	12.40	12.53	1.83	2.09	2.68	2.46
29	4978	12.66	12.73	1.77	2.18	2.71	2.34
30	4980	12.98	14.15	1.79	1.83	2.40	2.40
31	4981	13.23	13.22	1.75	1.76	2.53	2.37
32	4982	13.01	12.63	1.78	1.80	2.43	2.22
33	4983	12.57	12.06	1.68	1.77	2.53	2.39
34	4984	13.22	13.05	1.75	1.85	2.54	2.18
35	4985	13.58	14.12	1.87	1.68	2.32	2.23
36	4987	13.93	12.84	1.99	2.23	2.55	2.26
37	4988	14.10	6.11	1.82	2.11	2.47	2.48
38	4989	13.23	12.76	1.80	1.95	2.34	2.41
39	4990	13.57	13.69	1.86	1.89	2.41	2.34
40	4991	14.14	11.70	1.65	1.75	2.60	2.76
41	4992	14.63	12.61	1.77	1.71	2.46	2.23
42	4993	13.92	11.96	1.70	1.78	2.66	2.44
43	4994	14.13	13.58	1.82	2.02	2.31	2.27
44	4995	14.80	12.35	2.11	1.61	2.57	2.34
45	4996	13.97	12.06	1.96	2.03	2.52	2.33

Table 1. Continued.

No	Accession No.	% Moisture content		% Total Carbohydrate content	
		1995 GC	2018 GC	1995 GC	2018 GC
1	4907	10.42	11.59	72.49	71.17
2	4908	10.63	11.23	74.53	72.50
3	4919	10.35	11.32	73.26	73.06
4	4921	10.88	11.07	71.93	73.20
5	4923	9.73	6.30	73.23	77.15
6	4927	10.45	8.03	71.56	75.53
7	4929	10.64	6.68	72.55	76.65

No	Accession No.	% Moisture content		% Total Carbohydrate content	
		1995 GC	2018 GC	1995 GC	2018 GC
8	4939	10.76	9.08	74.66	74.51
9	4942	9.81	9.28	72.62	74.81
10	4943	10.72	9.55	72.12	74.89
11	4944	10.96	9.23	71.89	75.17
12	4945	10.75	9.40	72.76	75.64
13	4951	10.73	9.54	72.88	75.30
14	4952	10.73	9.43	73.78	75.17
15	4955	10.84	9.42	73.44	74.38
16	4958	10.45	9.38	73.49	74.07
17	4960	9.75	9.17	73.55	73.27
18	4961	10.13	9.37	73.17	74.07
19	4963	10.73	9.23	69.28	75.53
20	4966	10.82	9.51	72.43	75.14
21	4967	10.23	9.57	71.62	74.57
22	4969	10.27	9.42	73.09	75.70
23	4970	11.10	9.33	72.00	75.47
24	4972	10.18	9.28	74.09	75.67
25	4973	10.01	9.30	73.59	74.87
26	4974	10.01	9.92	72.89	73.17
27	4976	10.15	8.28	72.40	74.87
28	4977	10.56	7.40	72.53	75.53
29	4978	10.69	7.63	72.17	75.12
30	4980	10.06	9.32	72.77	72.29
31	4981	10.78	9.71	71.71	72.95
32	4982	10.85	9.70	71.93	73.65
33	4983	10.88	9.22	72.34	74.56
34	4984	11.38	9.37	71.11	73.55
35	4985	11.38	9.74	70.85	72.23
36	4987	10.96	9.03	70.57	73.64
37	4988	11.36	9.28	70.25	80.02
38	4989	10.74	9.44	71.89	73.44
39	4990	10.72	9.40	71.44	72.68
40	4991	10.85	9.50	70.76	74.30
41	4992	11.02	9.52	70.12	73.93
42	4993	11.15	9.42	70.57	74.40
43	4994	11.13	9.39	70.61	72.74
44	4995	11.05	9.30	69.47	74.40
45	4996	10.93	9.26	70.62	74.32

Descriptive analysis for the percentage content of crude protein in barley crop by year of test for each sampled accession

Figure 1. Percentage of crude protein content by year of test.

Descriptive analysis for the distribution of crude fat content in barley crop by year of test for each sampled accession.

Figure 2. Percentage of crude fat content by year of test.

Descriptive analysis for the distribution of mineral ash content in barley crop by year of test for each sampled accession

Figure 3. Percentage of mineral ash content by year of test.

Descriptive analysis for the distribution of moisture content in barley crop by year of test for each sampled accession.

Figure 4. Percentage of moisture content by year of test.

Descriptive analysis of the distribution of total carbohydrate content in barley crop by year of test for each sampled accession.

Figure 5. Percentage of total carbohydrate content by year of test.

Figure 6. Average percentage contents of Barley crop by ingredients and year of test.

Test of average percentage contents of nutritional values in the barley crop in 1993 and after 25 years of storage time (in 2018) were decreased. The data was obtained from the test of percentage crude protein, crude fat, Mineral ash, moisture and total carbohydrate contents conducted on the same

sampled barley accession in 1993 and 2018. The paired t-test of mean difference of percentage content ingredients in barley crop in 1993 and after 25 years of storage (in 2018), Significant at 5% level

Table 2. Test of difference in average percentage contents of nutritional values of barley accession in 1993 and after 25 years (2018).

Paired Samples Statistics for Barley Crop between percentage content in 1993 and after 25 years of storage time (2018)		Percentage Content Mean	Paired Differences 95% Conf. Interval		t
			Lower	Upper	
Pair 1	Crude Protein Contents in 1993 test	12.848±0.16	0.407	1.373	3.714*
	Crude Protein Contents in 2018 test	11.958±0.19			
Pair 2	Crude Fat Contents in 1993 test	1.83±0.029	-0.154	0.024	-1.477
	Crude Fat Contents in 2018 test	1.895±0.028			
Pair 3	Mineral Ash Contents in 1993 test	2.53±0.023	0.033	0.185	2.876*
	Mineral Ash Contents in 2018 test	2.42±0.028			
Pair 4	Moisture Contents in 1993 test	10.638±0.063	1.036	1.639	8.936*
	Moisture Contents in 2018 test	9.3±0.147			
Pair 5	Total Carbohydrate Contents in 1993 test	72.156±0.191	-2.875	-1.670	-7.601*
	Total Carbohydrate content in 2018 test	74.428±0.223			

*Significant at 5% level.

As it can be depicted in the Table 2, the average percentage difference of crude protein contents in barley crop was found statistically significant. It indicates that, there is significant difference in average percentage contents of crude protein of barley after 25 years of storage time. The high average percentage of crude protein shown in 1993 test year (12.848 ± 0.16) and the lowest was shown in 2018 test year (11.958 ± 0.19). The percentage of crude protein in barley were in general agreement with the values (7.50–12.12%) [14]. But this study shows little different for that agreement value, but still in the range. Such a difference could be due to variety and environmental conditions [15]. Other studies have also found that protein content of barley is highly dependent on the cultivar [16] and differs with growth conditions, particularly with the rate and timing of nitrogen fertilization [17]. According to [18] the crude protein concentration in barley varieties ranged from 9.44% to 13.57% it is similar to these studies. The average percentage difference of average crude fat contents in barley crop was found statistically insignificant. The average crude fat concentration in 1993 test year was (1.83 ± 0.029) and 2018 test year was (1.895 ± 0.028). A crude fat value is similar in barley varieties that determined in earlier studies [19].

As indicated in the Table 2, average percentage difference of mineral ash contents in barley crop was found statistically significant. It indicates that, there is significant difference in average percentage contents of mineral ash of barley after 25 years of storage time and before. The results shows that the high mineral ash content was in test year 1993 (2.53 ± 0.023) and the lowest result obtained after 25 years of storage time (2.42 ± 0.028). The obtained results, of the chemical composition in barley grains are comparable with those reported by [20].

As it can be depicted in the Table 2, the average percentage difference of moisture content in barley crop was found statistically significant. It indicates that, there is significant difference in average percentage contents of moisture of barley after 25 years of storage time and before. The result showed that the percentage content of moisture was obtained (10.638 ± 0.063) and (9.3 ± 0.147) in 1993 and 2018 respectively. Notably, the value of moisture content was decreases as storage time increases. This was because of the moisture content depends on the storage condition and hygroscopic capacity of the seed [21].

Moreover, the average percentage difference of total carbohydrate contents in barley crop was found statistically significant. It indicates that, there is significant difference in average percentage contents of total carbohydrate of barley after 25 years of storage time. The result showed that the percentage content of total carbohydrate was found (72.156 ± 0.191) and (74.428 ± 0.223) respectively in 1993 and 2018. As storage time increases percentage value of total carbohydrate also increase, this is due to all nutritional value of the sample decrease. Barley was higher in total carbohydrate than other crops as many reviewers indicated in their studies.

5. Conclusion

The study revealed that long term storage was affect nutritional value of conserved barley crop. As a result, the study found; statistically significant difference of average percentage content of crude protein, mineral ash, moisture and total carbohydrate between 1993 and after 25 years of storage time. The current study concluded that, when the storage time increases, the nutritional values of barley crop were significantly decreased for all nutritional values except crude fat content and total carbohydrate. Thus, it helps to profile the overall barley nutritional information and long term storage effect for both consumption and research purpose. Therefore, this study recommends that, the conserved material in Gene Bank should regenerate at possible short period of time.

Conflict of Interest

The author declares that there is no conflict of interests regarding the publication of this paper.

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