The Effects of Packaging Gas, Temperature and Storage Time on Germination, Loaf Volume and Protein Solubility of Wheat

A Thesis

Presented to the

Department of Food Science and Nutrition

Brigham Young University

In Partial Fulfillment

of the Requirements for the Degree

Master of Science

by

Scott W. Myers

August 1991

This thesis, by Scott W. Myers is accepted in its present form by the Department of Food Science and Nutrition of Brigham Young University as satisfying the thesis requirement for the degree of Master of Science.

John Hal Johnson, Committee Chairman

H. Gill Hilton, Committee Member

Clayton S. Huber, Committee Member

7/9/9/ Date

Mark J. Rowe, Department Chairman

Table of Contents

	Page
List of Tables	iv
List of Figures	v
Introduction and Literature Review	1
Materials and Methods	8
Results and Discussion	12
References	30
Appendix	33

Table of Contents

	Page
List of Tables	iv
List of Figures	v
Introduction and Literature Review	1
Materials and Methods	8
Results and Discussion	12
References	30
Appendix	33

List of Tables

Table		Page
1	The effect of temperature and storage time on the percent oxygen of wheat sealed in cans under air or nitrogen	13

List of Figures

I	igure		Page
	1	The effect of storage temperature and time on the germination of wheat	17
	2	The effect of storage temperature and time on loaf volume of wheat	20
	3	The effect of storage temperature and time on loaf volume of wheat packaged with air	22
	4	The effect of storage temperature and time on loaf volume of wheat packaged with carbon dioxide	23
	5	The effect of storage temperature and time on loaf volume of wheat packaged with nitrogen	24
	6	The effect of storage temperature and time on protein solubility of wheat	26

Introduction and Literature Review

Barton (1961) stated that the three major factors which influence seed viability during storage are temperature, moisture content and oxygen pressure. Wheat functionality may be damaged in the field before or during harvest, during artificial drying operations or during subsequent storage and handling (Halverson and Zeleny, 1988). The extent and nature of this damage will govern the viability and quality of wheat. Temperature

Heat damage occurs in wheat exposed to relatively high temperatures, causing decreases in germination and breadmaking quality. Glass et al. (1959) stored wheat in flasks sealed under air or nitrogen at 20°C and 30°C for up to 48 weeks. Storage at 30°C showed accelerated decreases in germination and bread-making quality when compared to storage at 20°C. Storage in nitrogen delayed deterioration somewhat at 30°C and significantly at 20°C. Burges et al. (1963) stored barley at 13.2% moisture content in sealed jars at 18°C, 32°C, 35°C and 40°C for 26 weeks. They found that at 18°C, germination remained close to 95% for the entire 26 weeks. Within 10 weeks, germination levels at 40°C dropped to essentially 0%, while germination levels at 32°C and 35°C were similar to those at 18°C. After 26 weeks, however,

germination levels at 32°C and 35°C had dropped to less than 20%.

One of the signs of exposure to high temperatures and aging in seeds is a decrease in α -amylase activity. This decrease in \alpha-amylase activity is partially due to thermal inactivation of α -amylase which can lead to decreased germination due to the lack of utilizable substrate [Bhattacharyya and Sen-Mandi, 1985]. The release of substrate to the germinating embryo is controlled by the rate at which α -amylase is synthesized and reaches the starch granules (Wrigley and Bietz, 1988). Bhattacharyya and Sen-Mandi (1985) demonstrated that embryos of aged non-germinating wheat germinated well when placed on agar containing 3% sucrose or 2% glucose. The same aged wheat completely failed to germinate when placed on agar alone. An amylase activity assay showed that the aged wheat had no α -amylase activity. Moisture Content

Physical and chemical changes which take place in stored grain increase greatly with moisture content of the grain (Hunt and Pixton, 1974). Seeds with high moisture show increased susceptibility to germination and development of germ damage. Germ damage is apparent when the kernels take on a dull appearance and when the germ becomes darkened (Milner and Geddes, 1954). Petruzzelli (1986) stored wheat, having moisture contents ranging from 15 to 33 percent, under either hermetic and aerobic conditions at 25°C. Under

hermetic storage, wheat lost viability more rapidly with increased moisture. Wheat stored aerobically showed enhanced viability as moisture increased from 24 to 31 percent but had a drastic drop in viability at 33 percent. At 24 to 31 percent moisture, the wheat maintained viability longer when stored aerobically than stored hermetically. Glass et al. (1959) reported that for wheat stored in air at 30°C, germ damage occurred much earlier at 16 to 18% moisture than at 13 to 15% moisture. After eighteen weeks, half of the wheat at 16% and all of the wheat at 18% moisture had developed germ damage. Germ damage was much slower at 15% moisture; 36 weeks were required before 75% of the wheat developed germ damage. Germ damage was virtually nonexistent in wheat with moisture contents below 14% at 30°C. Hunt and Pixton (1974) stated that for most cereal grains, critical values for moisture content are about 14%. Most species of seeds remain viable longer during storage when moisture is reduced to five percent (Roberts, 1972).

Oxygen Level

Long-term storage experiments were conducted with seeds stored in sealed ampoules filled with air, oxygen, carbon dioxide or nitrogen. Temperature, moisture content and gaseous composition were maintained at constant levels. Roberts (1961) stored rice at temperatures of 27°C to 47°C for 1.5 years sealed with air, oxygen, carbon dioxide or nitrogen. Germination results were complex; the influence on

viability by each gas was not the same under all temperatures and moisture contents. Harrison (1966) stored 10 varieties of lettuce seeds for three years at 18°C and 6% moisture content under various gases. At the end of three years, he found that seeds stored in oxygen had a mean viability of 8%; in air it was 57%; whereas in both nitrogen and carbon dioxide it was 78%. Similar results were obtained using onion seed at 18°C and 8% moisture.

Roberts and Abdalla (1968) found the viability of barley, broad beans and peas decreased when stored under various levels of oxygen. Increasing the oxygen from 0 to 21% significantly accelerated the loss of viability. Peterson et al. (1956) reported that wheat stored 16 days at 25°C gradually decreased in mold growth, germ damage and respiration as oxygen was lowered from 21 to 0.2%.

Glass et al. (1959) stored wheat at 30°C for 42 weeks and at 20°C for 48 weeks in nitrogen or air. At 30°C, storage in nitrogen delayed the onset of deterioration, measured by germ damage and loaf volume, as compared with air. At 20°C, deterioration was much slower in wheat stored in nitrogen. Roberts (1961) reported that for seeds that remain viable for a year when stored in air, storage in nitrogen increased the period of viability.

Storing lettuce and onion seeds sealed under carbon dioxide or air for 18 years, Harrison (1966) showed that carbon dioxide extended viability. Lettuce seeds in air

remained viable for about eight years while seeds stored in carbon dioxide remained viable for more than nine years. Similar results were obtained with the onion seed. et al. (1984) reported early storage in carbon dioxide at 22°C prolonged seed longevity. Wheat was stored in carbon dioxide for 18 months and then in air for 16 years under ambient conditions. Every two years, the viability of the wheat was measured by germination tests. The wheat treated with early storage in carbon dioxide exhibited 58% germination after 10 years while seeds stored in air for the same time period completely failed to germinate. Peterson et al. (1956) stored wheat at 30°C and 18% moisture for 16 days under a carbon dioxide-oxygen mixture with 21% oxygen and up to 79% carbon dioxide. The carbon dioxide-oxygen mixture had little effect on respiration and mold growth until the carbon dioxide concentration exceeded 13.8% where very sharp inhibition occurred. Seeds stored in 50 and 79% carbon dioxide exhibited high viability.

Respiration

Under aerobic conditions, respiration occurs as molecular oxygen reacts with carbohydrates or lipids to produce carbon dioxide, water and energy. Respiration is accelerated by increases in temperature until it is limited by such factors as thermal inactivation of key enzymes, exhaustion of substrate, limitation in oxygen supply or accumulation of inhibitory concentrations of carbon dioxide (Pomeranz, 1974).

Roberts and Abdalla (1968) showed that when pea seeds were stored in sealed ampoules at 25°C, there was a linear increase in carbon dioxide content and a corresponding decrease in oxygen content. After 11.3 weeks, the oxygen content of the ampoules had dropped from 21 to 1.4 percent and the carbon dioxide content rose from 0.03 to about 12 percent. Roberts (1961) stated that if seeds are stored in nitrogen or carbon dioxide instead of air, changes in gaseous environment would be less and stated that storage in oxygen would result in greater changes.

Methods of Measurement

The ultimate measurement of wheat flour performance is established by a bake test (Mailhot and Patton, 1988). Germination is the most sensitive test for detecting heat-damage, however, it does not correlate well with predicting bread baking quality because it is an all-or-nothing measurement. Every (1987) developed a protein solubility test (PST) that more accurately predicts baking quality by detecting the extent of heat damage to soluble wheat proteins. His test was based on Pence et al. (1953) who reported that thermal denaturation of wheat proteins, especially glutenin, drastically reduced germination and baking quality of wheat. Every (1987) showed that heat damaged wheat produced significantly reduced loaf volumes in addition to decreased solubility in a dilute sodium chloride solution. Reductions in solubility are detected by decreases in absorption due to

decreased dye binding. Carpenter (1973) demonstrated that heat damaged proteins bind to dyes less than raw proteins. Kratzer et al., (1990) confirmed that heat damage to proteins could be measured using Coomassie Blue G dye-binding. This relationship between dye binding and heat damage establishes the significance of the PST as an indicator of baking quality. The PST, like baking tests, shows gradual decreases in wheat quality and thus is a better indicator of bread-making quality than germination. To better understand the changes in wheat viability and quality under different storage conditions, measurements of percent germination, loaf volume and protein solubility are desirable.

Objective |

Most of the early experiments were designed to predict the effects of open air storage. There is little information on the effects of storage conditions on wheat for home use. A five year study has been undertaken to measure the effects of packaging gas, storage temperature and time on the viability and quality of wheat stored in sealed containers. This is a report on the changes occurring during the first 18 months of storage of wheat sampled at 0, 6, 12 and 18 months.

Materials and Methods

Materials

Two varieties of wheat, Freemont and Ute, were obtained from the Brigham Young University Spanish Fork farm in Utah. The Freemont variety is a hard spring wheat harvested in August, 1988. The Ute variety is a hard winter wheat harvested in June, 1988. The wheat was stored in an open air shed in 100 lb. polypropylene sacks until September, 1988.

Packaging

The wheat was shipped to the Vacu-Dry Company in Sebastopol, California, where it was packaged and sealed in No. 2 1/2 cans. Each variety was randomly divided and packaged under one of three packaging gases: air, carbon dioxide or nitrogen.

Storage

Upon return to Provo, Utah, the canned wheat was randomly assigned to one of four temperatures: 4°C, 21°C, 37°C and 54°C. Cans were taken from storage at six month intervals for analysis. While waiting to undergo analysis, the canned wheat was stored at -17°C for two to three months.

Percent Oxygen Test

The oxygen content of each can was measured with a Beckman Altex Oxygen Analyzer (model 0260). A 4 inch, 7 gauge stainless steel hollow needle inserted through a rubber stopper was used to puncture the can so that a percent oxygen

reading could be taken. The rubber stopper served as a temporary gasket to prevent air intrusion. The oxygen content recorded was the lowest oxygen value that registered before it increased due to the intrusion of outside air.

Moisture Test

The percent moisture of the wheat was determined by weighing out 5 grams of whole wheat into a petri dish, drying it in an oven at $95^{\circ}C \pm 5^{\circ}C$ for 36 hours and then reweighing the wheat. The loss in weight was used to calculate the percent moisture of the wheat.

Germination Test

One hundred undamaged wheat kernels were counted out and soaked in a solution of 0.05% sodium benzoate (w/v) for 2 hours. The sodium benzoate solution served to inhibit mold growth. After soaking, the kernels were placed on moist filter paper in a covered petri dish and stored at 20°C in a dark place for four days. The kernels that germinated were then counted and recorded as a percentage.

Grinding and Milling

Whole wheat flour for the baking and protein solubility tests was obtained by grinding the wheat in a high speed rotor-stator pin-mill, Magic Mill III Plus (Magic Mill Co., Salt Lake City, UT 84104). To obtain a bread-type flour, the wheat was ground at the most coarse setting on the mill. The flour was stored in the freezer at -28°C for one to two weeks until used.

Baking Test le Volume was brough

Bread was made in an automatic bread maker: Panasonic Bread Bakery, (Model SD-BT2P, Panasonic Inc., 1 Panasonic Way, Secaucus, NJ, 07094). The formulation of the bread was 270 g whole wheat flour, 11 g margarine, 11.9 g sucrose, 8.4 g nonfat dried milk, 4.7 g NaCl, 3.2 g dry active yeast and 210 ml water. After baking and cooling, loaf volume was measured by rapeseed displacement.

Protein Solubility Test

The protein solubility test (PST) (Every, 1987) was used to assess the heat damage of the whole wheat flour. Fifty ml of 2% (w/v) NaCl were measured in a 100 ml graduated cylinder. 1.000 gram of whole wheat flour was added to the salt solution and mixed by inverting the cylinder by hand once every second for 30 seconds. After allowing the flour to settle for 30 seconds, a 100 μ l extract was removed from the top layer of the liquid and immediately mixed with 3 ml of Coomassie Brilliant Blue G (Sigma Chemical Company, St. Louis, Mo, No. B-0770) protein stain reagent. After 2 minutes, the absorbance was read at 595 nm by a Perkin-Elmer Coleman 124 spectrophotometer (Hitachi Ltd., Tokyo, Japan) against a blank of the protein stain reagent.

Preparation of the Protein Stain Reagent

The preparation of the reagent was according to Bradford (1976). Ten mg of Coomassie Brilliant Blue G were dissolved in 5 ml of 95% ethanol. Ten ml 85% (w/v) phosphoric acid were

added and the volume was brought up to 100 ml with distilled water. Consistent results were obtained by shaking the reagent before use and periodically during use to prevent settling. The reagent was refrigerated when not in use to prolong its lifetime from one month at room temperature to three or more months. No reagent, however, was more than 2 weeks old when used.

Statistical Analysis not assessed apparatoly due to

A multivariate analysis was performed on the data from the percent oxygen test, germination test and baking test. An analysis of variance was conducted on the data from the protein solubility test. Fisher's LSD multiple comparison procedure was conducted on the main effects or interactions where significance occurred. All hypotheses were tested at a 0.05 level of significance.

these readings and those from the atudy were probably the nethods of measurement and the number of dama person

aboved a three-way interaction between package temperature and storage time. Table 1 shows oxygen for cans seeled with air or nitrogen. Canadian showed the most significant phanges in oxygen tout. It sealed with air or nitrogen. It is also the most significant phanges in oxygen the canadian change in oxygen the canadian change in oxygen content over 18 souths of accress. The pages

Results and Discussion

With multivariate analysis, the tests of percent oxygen, germination and baking showed a significant three-way interaction between packaging gas, temperature and storage time. The two-way interactions between variety and packaging gas and between variety and time were also significant. The main effects were not examined separately due to the significance of these interactions.

Percent Oxygen Test

The Vacu-Dry Company that packaged the wheat into cans reported that the cans sealed under air had 16 to 18% oxygen, those sealed under carbon dioxide 9.5 to 11.5% oxygen, and those sealed under nitrogen had less than 2% oxygen. The oxygen content from cans with carbon dioxide and nitrogen was determined using a flushing procedure. The differences in these readings and those from the study were probably due to the methods of measurement and the number of cans measured.

Percent oxygen results using the Altex oxygen analyzer showed a three-way interaction between packaging gas, temperature and storage time. Table 1 shows the percent oxygen for cans sealed with air or nitrogen. Cans with air showed the most significant changes in oxygen content. Cans sealed with air stored at 4°C showed no significant changes in oxygen content over 18 months of storage. Cans stored at 21°C for 6 months and 12 months had significantly greater

Table 1: The effect of temperature and storage time on the percent oxygen of wheat sealed in cans under air or nitrogen.

Percent Oxygen Present

months air <u>6</u> <u>12</u> <u>18</u> 4°C 17.1% 17.3% 17.6% 21°C 14.4% 13.5% 11.4% 37°C 12.3% 6.8% 5.2% 54°C 8.9% 6.3% 3.7% nitrogen <u>6</u> <u>12</u> <u> 18</u> 4°C 4.9% 4.0% 4.4% 21°C 4.1% 4.5% 3.4% 37°C 4.7% 3.1% 2.8% 54°C 5.6% 4.4% 2.8%

oxygen contents than those cans stored at 21°C for 18 months. Cans stored at 37°C and 54°C had significant decreases in oxygen content as time progressed during the study. As respiration rates and other processes such as oxidation increase at higher temperatures, oxygen uptake occurs resulting in decreases in percent oxygen as shown. Cans packaged under air and stored at high temperatures would probably reach a point where carbon dioxide exerts a greater influence over the wheat than does oxygen.

The cans sealed with carbon dioxide did not show any significant changes in oxygen content with temperature or storage time according to our method of analysis. This did not agree with the expected results, therefore, the oxygen data from the carbon dioxide cans is considered to be inaccurate for the following reasons. First, the analyses of the Vacu-Dry Company measured 50% less oxygen than our analyses. Second, the data for carbon dioxide is not consistent with that of the other gas treatments. For both air and nitrogen, oxygen was diminished with storage at the warmer temperatures as respiration occurred. The carbon dioxide results should have reflected similar changes. Third, the oxygen levels would be expected to be lower as a result of flushing as seen for the nitrogen cans. The flushing should have also lowered the oxygen content of the carbon dioxide cans. The reason for the inaccuracy of the carbon dioxide oxygen data is unknown. An alternative method of

measuring oxygen should show that oxygen levels for carbon dioxide are more consistent with the expected results.

Cans sealed with nitrogen had more complex interactions with time and temperature for oxygen content. There were no significant differences in oxygen content at 4°C and 21°C. At 37°C, the oxygen content at 6 months was significantly greater than that at 12 and 18 months. At 54°C, the oxygen content at 6 and 12 months was greater than that at 18 months. This data is starting to show that the oxygen content decreases at higher temperatures as it did in the cans packaged under air. The smaller changes in oxygen content for nitrogen are probably due to the low initial amount of oxygen and possible inhibition of respiration by nitrogen.

Multivariate analysis showed that percent oxygen in cans was affected by a three-way interaction between packaging gas, temperature and storage time. It was not influenced by the wheat variety. This was expected since both varieties were packaged under the same conditions and nothing occurred that should cause differences.

Moisture Test

The moisture content of the wheat was determined for each variety. The Freemont variety averaged $8.1\%\pm0.3\%$ moisture and the Ute variety averaged $8.7\%\pm0.6\%$ moisture. These values were considerably lower than the 14% moisture levels of commercially stored grain (Hunt and Pixton, 1974) but well within the levels needed for the wheat seeds to remain viable

(Roberts, 1972). The lower moisture levels in the wheat were encouraged by drought conditions in Utah during the past few years.

Germination Test

The percentage of wheat that germinated was affected by an interaction between variety and packaging gas. Germination results for the Freemont variety for air and nitrogen were not significantly different. Both gases produced significantly better germination results than carbon dioxide. The percentage of wheat that germinated for carbon dioxide averaged about 6% lower than those of the other gases. the Ute variety, germination with nitrogen was significantly greater than with air, there being a 6.5% difference. The results for carbon dioxide fell between these two gases and were not significantly different from either one. months of storage, no trends were found that would establish which packaging gas was best to ensure optimal germination because of the variation between the varieties.

The interaction of temperature and time on germination is depicted in Figure 1. The 4°C and 21°C treatments did not significantly affect germination over 18 months. Some variation in the wheat itself or in the testing may account for small temperature differences. The 6 month germination levels at 37°C did not differ significantly from those at 4°C and 21°C. Germination levels at 37°C for 6 months were significantly higher than those at 12 and 18 months, dropping

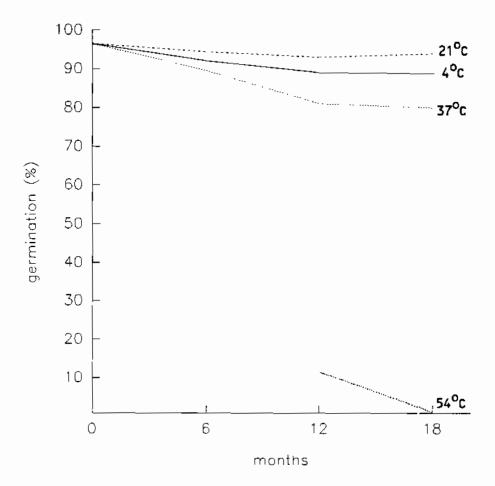


Figure 1: The effect of storage temperature and time on the germination of wheat (SEM = 2.79).

from 90% \pm 3% to 81% \pm 8% and 80% \pm 6%, respectively. This trend was even more significant at 54°C. After only 6 months, 71% \pm 9% of the wheat germinated. Germination further dropped to 11% \pm 9% after 12 months and 0% \pm 0% after 18 months of storage at 54°C. From these results it is clear that storage at elevated temperatures becomes increasingly detrimental to wheat germination over time.

These results correspond with those of other researchers. Every (1987) showed that heat-treating wheat, which had an initial moisture content of 18%, for 3 hours at 60°C and 80°C resulted in 63% and 0% germination respectively (SEM = 6). Lesser heat treatments had germination values ranging from 82 to 97%. While the heat treatment in his study was more harsh than those of this study, his results also indicate that heat seriously affects germination. The prolonged 54°C heat treatment of the current study could have caused a loss of α -amylase activity after 18 months of storage. Bhattacharyya and Sen-Mandi (1985) showed that aged wheat, that lacked α -amylase activity, germinated when placed on agar containing sucrose.

Baking Test

The baking quality of the stored wheat was measured by loaf volume. An automatic bread maker was used to bake the loaves to avoid the variation associated with making the bread by hand. Multivariate analysis showed interactions between variety and time and between temperature and time.

Significant differences in loaf volume were also found among the three packaging gases.

The relationship between variety and time showed that at 6 months, the Ute variety produced significantly larger loaves than the Freemont variety. However, the Freemont variety gave significantly larger loaf volumes at 12 months. There were no significant differences between loaf volumes of Freemont and Ute varieties at 18 months. The reason for this variation between the varieties is not known. It might be due to the varieties undergoing changes in proteins at different times during storage.

As with germination, there is a significant interaction between temperature and time on loaf volume as shown in Figure After 6 months, there were no significant differences between loaf volumes of wheat stored at 4°C, 21°C and 37°C. Loaf volumes at all three temperatures were significantly greater than those at 54°C. Twelve months of storage showed no significant differences between loaf volumes of wheat stored at 4°C and 21°C and between those of wheat stored at 37°C and 54°. Loaf volumes of wheat stored at 4°C and 21°C were significantly greater than those of wheat stored at 37°C 54°C. After 18 months, loaf volume decreased and significantly as the storage temperature increased. At 4°C and 18 months, the average of loaf volumes was 1328 ml \pm 34 The average loaf volumes at 18 months dropped to 1242 ml \pm 46 ml for 21°C, 1204 ml \pm 32 ml for 37°C and 939 ml \pm 51 ml

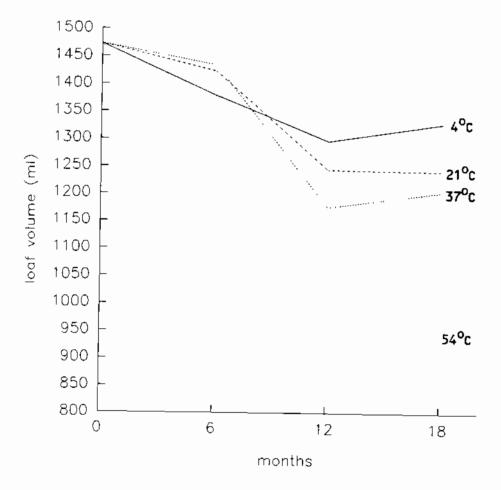


Figure 2: The effect of storage temperature and time on loaf volume of wheat (SEM = 13.00).

for 54°C. While no measurements were made, the average loaf volume for 18 months at 54°C appeared only slightly larger than the volume of the ingredients.

As the stored wheat received increasing amounts of temperature and time, the loaves showed signs of protein damage. The loaves rose but could not hold their structure and the bread became increasingly more compact. Storage at 37°C and 54°C adversely affected bread quality in a relatively short time period. Every (1987) also conducted baking tests on heat treated wheat and found that temperatures of 60°C and 80°C for 3 hours produced loaves of less than 50% volume of the controls.

The effect of packaging gas on loaf volume was slight but significant. The average loaf volume from wheat packaged under the three gases differed by only 41 ml. The loaf volume of the wheat packaged under air averaged 1271 ml ± 40 ml, which was significantly greater than the average loaf volume of carbon dioxide, 1230 ml ± 49 ml. The nitrogen packaged wheat averaged 1250 ml ± 43 ml, which was not significantly different from the results of either of the other two gases. The effects of each packaging gas versus temperature and time are shown in Figures 3, 4 and 5 for comparison. From these graphs it can be seen that for carbon dioxide at 18 months and 21°C, wheat gave loaves lower in volume than loaves at 37°C for air or nitrogen. While the above results are significant, only slight overall differences between loaf volumes of the

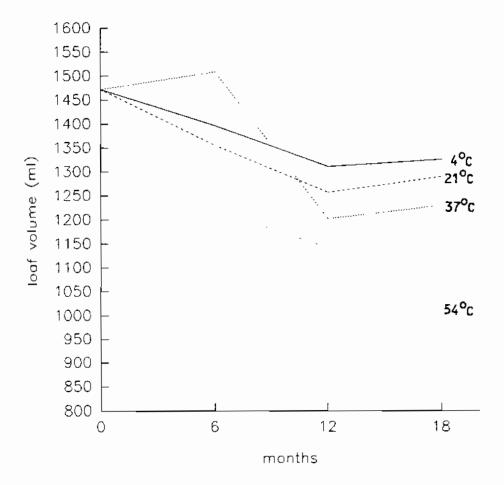


Figure 3: The effect of storage temperature and time on loaf volume of wheat packaged with air (SEM = 20.65).

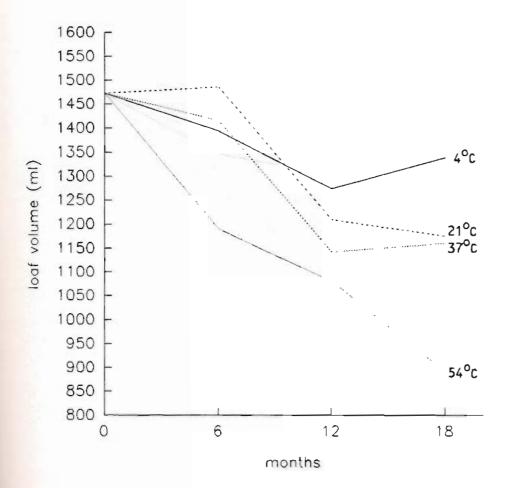


Figure 4: The effect of storage temperature and time on loaf volume of wheat packaged with carbon dioxide (SEM = 24.85).

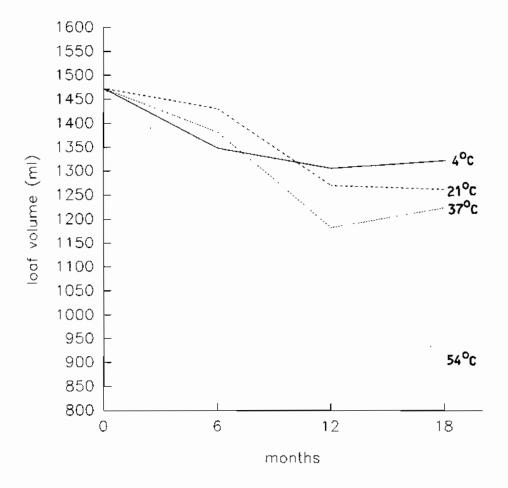


Figure 5: The effect of storage temperature and time on loaf volume of wheat packaged with nitrogen (SEM = 21.91).

packaging gases exist as shown by overlapping confidence intervals for air and carbon dioxide. With additional storage, the effects of packaging gases may become more pronounced.

Protein Solubility Test

An analysis of variance on the results from the protein solubility test (PST) showed that, like the other tests, there was a significant interaction between temperature and time. Additionally, significant differences were shown between varieties and between packaging gases.

Temperature and time effects on protein solubility are shown in Figure 6. The PST for the 4°C wheat showed that absorbance at 6 months was significantly greater than at 12 and 18 months. The increase in absorbance at 6 months over the control absorbance at 0 months cannot be explained except for by experimental error. The PST requires exact methodology to get good results and this probably accounts for the variation in results. There were no significant differences in absorbance during 18 months of storage at 21°C. month absorbance at 21°C was significantly lower than that at At 37°C, there were significant differences in 37°C. absorbance at each time period. The average absorbance at 6 months, 0.532 + 0.014, was significantly greater than that at 12 months, 0.482 ± 0.018. Similarly, the average absorbance at 12 months was significantly greater than that at 18 months, 0.445 ± 0.012 . The 6 and 12 month absorbances at 54° C,

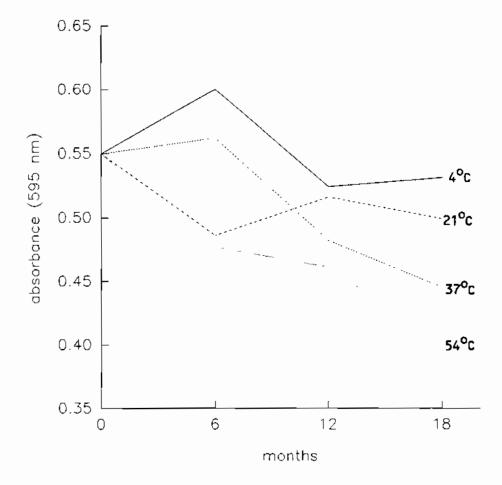


Figure 6: The effect of storage temperature and time on protein solubility of wheat (SEM = 0.0069).

 0.447 ± 0.011 and 0.461 ± 0.032 , were not significantly different from each other. They both were significantly greater than the 18 month absorbance, 0.400 ± 0.014 . At 12 and 18 months of storage, significant decreases in absorbance were seen with each sequential increase in storage temperature. From this, it is evident that there is a significant trend for decreased absorbance at higher temperatures. This indicates protein degradation is occurring and concurs with decreasing germination and lower loaf volumes under the same conditions.

These results concur with the findings of Every (1987) who showed heat damaged wheat exhibited decreased absorbance with the PST. Three hours at 60°C and 80°C showed decreased absorbances of 0.286 and 0.145, respectively, from the control absorbance of 0.349. These values are lower than those of the current study but Every used more severe heat treatments and different varieties of wheat.

Freemont and Ute varieties of wheat had initial average absorbances of 0.481 \pm 0.010 and 0.513 \pm 0.011 respectively. The Ute variety had significant and consistent higher absorbances than the Freemont variety due to initially higher levels of soluble protein in the Ute variety.

There were also significant differences in protein absorbance among the packaging gases. Wheat packaged in nitrogen had a significantly higher average absorbance, 0.516 \pm 0.012, than wheat packaged in carbon dioxide, 0.494 \pm 0.014,

or air, 0.480 ± 0.014. There was no significant difference in average absorbance between the carbon dioxide and air packaged wheat. Nitrogen, the most inert gas, had the mildest effect on protein solubility. The absorbance values of the three packaging gases, however, are within 0.036 of each other so further study is needed before decisive conclusions are made about these effects.

Conclusion

In summary, the percent oxygen test showed that time, temperature and packaging gas all played a role in the oxygen content of the wheat cans. This interaction was most pronounced for the wheat sealed under air. The germination, baking and protein solubility tests showed that temperature significantly affects wheat quality as a function of time. At higher storage temperatures, wheat suffers increasing degradation over shorter periods of time. This was especially pronounced at 54°C. Eventually, wheat is neither able to germinate nor to produce bread that is acceptable. The germination test showed that the wheat varieties responded differently to each packaging gas. The Freemont variety responded best to air and nitrogen packaging while the Ute variety responded best to nitrogen and carbon dioxide packaging. The baking test showed that varieties responded differently over time and that wheat packaged under air produced significantly larger loaf volumes than wheat packaged under carbon dioxide. The PST showed the Ute variety and

packaging under nitrogen had the highest average absorbances. In all three tests, the differences between packaging gases were small but significant. Within 18 months of storage, a single packaging gas did not stand out as the best so no conclusions on packaging gas could be made. Differences due to packaging gases may become more significant during the three and a half years left in this study.

References

- Barton, L. V., 1961. Seed Preservation and Longevity, Leonard Hill, London.
- Bennici, A., Bitonti, M. B., Floris, C., Gennai, D., Innocenti, A. M. 1984. Aging in triticum durum wheat seeds: early storage in carbon dioxide prolongs longevity. Env. Exp. Bot. 24:159-165.
- Bhattacharyya, S. and Sen-Mandi, S. 1985. Studies into the causes of non-germination of aged wheat seeds. Annals Bot. 56:475-479.
- Bradford, M. M. 1976. A rapid and sensitive method for the quantitation of microgram quantities of protein utilizing the principle of dye binding. Anal. Biochem. 72:248-254.
- Burges, H. D., Edwards, D. M., Burrell, N. J., and Cammell, M. E. 1963. Effect of storage temperature and moisture content on the germinative energy of malting barley, with particular reference to high temperature. J. Sci. Food Agric. 14:580-584.
- Carpenter, K. J. 1973. Damage to lysine in food processing: its measurement and significance. Nutr. Abs. & Rev. 43:423.
- Every, D. 1987. A simple, four-minute, protein-solubility test for heat damage in wheat. J. Cereal Sci. 6:225-236.
- Glass, R. L., Ponte, J. G., Christensen, C. M., and Geddes, W. F. 1959. Grain storage studies XXVIII. The influence of temperature and moisture level on the behavior of wheat stored in air or nitrogen. Cereal Chem. 36:341-356.
- Halverson, J. and Zeleny, L. 1988. Criteria of wheat quality, pp. 15-45. In Wheat: Chemistry and Technology, vol. 1, ed. Y. Pomeranz. American Association of Cereal Chemists, Inc., St. Paul, Minnesota.
- Harrison, B. J. 1966. Seed deterioration in relation to storage conditions and its influence upon germination, chromosomal damage and plant performance. J. Nat. Inst. Agric. Bot. 10:644-663.

- Hunt, W. H. and Pixton, S. W. 1974. Moisture Its significance, behavior, and measurement, pp. 1-55. In Storage of Cereal Grains and their Products, ed. C. M. Christensen. Monograph series, Vol. V. American Association of Cereal Chemists, Inc., St. Paul, Minnesota.
- Kratzer, F. H., Bersch, S. and Vohra, P. 1990. Evaluation of heat-damage to protein by Coomassie Blue G dyebinding. J. Food Sci. 55:805-807.
- Mailhot, W. C. and Patton, J. C. 1988. Criteria of flour quality, pp. 69-90. In Wheat: Chemistry and Technology, Vol. 2, ed. Y. Pomeranz. American Association of Cereal Chemists, Inc., St. Paul, Minnesota.
- Milner, M. and Geddes, W. F. 1954. Respiration and heating, pp. 152-220. In Storage of Cereal Grains and their Products, eds. J. A. Anderson and A. W. Alcock. Monograph series, Vol. II. American Association of Cereal Chemists, Inc., St. Paul, Minnesota.
- Pence, J. W., Mohammad, A. and Mecham, D. K. 1953. Thermal denaturation of gluten. Cereal Chem. 30:115-126.
- Peterson, A., Schlegel, V., Hummel, B., Cuendet, L. S., Geddes, W. F., and Christensen, C. M. 1956. Grain storage studies XXII. Influence of oxygen and carbon dioxide concentrations on mold growth and grain deterioration. Cereal Chem. 33:53-66.
- Petruzzelli, L. 1986. Wheat viability at high moisture content under hermetic and aerobic conditions. Annals Bot. 58:259-265.
- Pomeranz, Y. 1974. Biochemical, functional, and nutritive changes during storage, pp. 56-114. In Storage of Cereal Grains and their Products, ed. C. M. Christensen. Monograph series, Vol. V. American Association of Cereal Chemists, Inc., St. Paul, Minnesota.
- Roberts, E. H. 1961. The viability of rice seed in relation to temperature, moisture content, and gaseous environment. Annals Bot. 25:381-390.
- Roberts, E. H. 1972. Storage environment and the control of viability, pp. 14-58. In Viability of Seeds, ed. E. H. Roberts. Chapman & Hall, London.

- Roberts, E. H. and Abdalla, F. H. 1968. The influence of temperature, moisture and oxygen on period of seed viability in barley, broad beans and peas. Ann. Bot. 32:97-117.
- Wrigley, C. W. and Bietz, J. A. 1988. Proteins and amino acids, pp. 159-276. In Wheat: Chemistry and Technology, Vol. 1. ed. Y. Pomeranz. American Association of Cereal Chemists, Inc., St. Paul, Minnesota.

Appendix

Wheat Data Summary Sheet - Percent Oxygen - 6 months

c	code	%O ₂	ave		code	%O ₂	ave
FA	4-6	17.0% 	17.0%	עט	4-6	17.1% *	17.1%
FN	4-6	5.9% ª	5.9%	Uì	1 4-6	4.6% 4.1%	4.4%
FA	21-6	15.9% 12.8%	14.4%	UZ	21-6	15.2% 13.6%	14.4%
FN	21-6	5.1% 3.3%	4.2%	U	1 21-6	4.3% 3.5%	3.9%
FA	37-6	11.8% 16.5%	14.2%	עט	37-6	10.6% 10.2%	10.4%
FN	37-6	5.7% 4.6%	5.2%	Uì	I 37 − 6	4.1% 4.4%	4.3%
FA	54-6	8.5% 9.7%	9.1%	U	A 54-6	9.1% 8.4%	8.8%
FN	54-6	5.6% 5.9%	5.8%	UI	54-6		5.4%

a - these cans were opened improperly so the percent oxygen could not be measured.

note: the percent oxygen readings from the carbon dioxide cans were inaccurate and are not reported. See text for details.

code:

F - Freemont

U - Ute

A - air

C - carbon dioxide

N - nitrogen

the numbers signify temperature and time i.e. $4-6 = 4 \,^{\circ}\text{C} - 6$ months

Wheat Data Summary Sheet - Percent Oxygen - 12 months

C	code	%O ₂	ave		code	%O ₂	ave
FA	4-12	16.2% 18.6%	17.4%	U	A 4-12	16.4% 17.8%	17.1%
FN	4-12	2.9% 5.2%	4.1%	UI	N 4-12	3.5% 4.4%	4.0%
FA	21-12	14.0% 14.0%	14.0%	U	A 21-12	13.1% 12.8%	13.0%
FN	21-12	4.1% 3.9%	4.0%	וט	V 21-12	5.4% 4.7%	5.1%
FA	37-12	7.5% 5.5%	6.5%	U	A 37-12	7.8% 6.3%	7.1%
FN	37-12	4.3% 2.6%	3.5%	UI	7 37-12	2.8% 2.8%	2.8%
FA	54-12	7.9% 6.6%	7.3%	U	A 54-12	6.9% 3.7%	5.3%
FN	54-12	4.3% 5.1%	4.7%	Uì	7 54-12	4.1% 4.0%	4.1%

note: the percent oxygen readings from the carbon dioxide cans were inaccurate and are not reported. See text for details.

code:

F - Freemont

U - Ute

A - air

C - carbon dioxide

N - nitrogen

the numbers signify temperature and time i.e. $4-12 = 4 \,^{\circ}\text{C} - 12 \, \text{months}$

Wheat Data Summary Sheet - Percent Oxygen - 18 months

(code	%O ₂	ave	c	ode	%O ₂	ave
FA	4-18	18.0% 17.2%	17.6%	UA	4-18	18.0% 17.3%	17.7%
FN	4-18	4.6%	4.0%	UN	4-18	4.6%	4.7%
FA	21-18	12.5% 11.2%	11.9%	UA	21-18	11.5% 10.3%	10.9%
FN	21-18	4.0% 2.7%	3.3%	UN	21-18	3.1% 3.7%	3.4%
FA	37-18	5.7% 5.0%	5.4%	UA	37-18	5.4% 4.7%	5.0%
FN	37-18	2.9% 2.7%	2.8%	UN	37-18	2.9% 2.6%	2.8%
FA	54-18	4.0% 3.2%	3.6%	UA	54-18	3.9% 3.5%	3.7%
FN	54-18	3.2% 2.8%	3.0%	UN	54-18	2.8% 2.4%	2.6%

note: the percent oxygen readings from the carbon dioxide cans were inaccurate and are not reported. See text for details.

code:

F - Freemont

A - air

N - nitrogen

U - Ute

C - carbon dioxide

the numbers signify temperature and time i.e. $4-18 = 4 \,^{\circ}\text{C} - 18 \, \text{months}$

Wheat Data Summary Sheet - Germination Test - control

code	%germ	ave	code	%germ	ave
Fre ctrl	98% 94%	96%	Ute ctrl	95% 93%	94%

Wheat Data Summary Sheet - Germination Test - 6 months

354

(code	%germ	ave	(code	%germ	ave
FA	4-6		93.5%	UA	4-6	95%	91.5%
		93%				888	
FC	4-6		90%	UC	4-6	93%	93.5%
		91%				948	
FN	4-6	928	91%	UN	4-6	90%	92.5%
		90%				95%	
FA	21-6	96%	94.5%	UA	21-6	97%	94.5%
		93%				92%	
FC	21-6	948	92.5%	UC	21-6	95%	94.5%
		91%				948	
FN	21-6		94%	UN	21-6	968	96%
		948				96%	
FA	37-6	86%	88%	UA	37-6	81%	83%
		90%				85%	
FC	37-6	888	88.5%	UC	37-6	92%	94.5%
		898				97%	
FN	37-6	97%	91%	UN	37-6	87%	92%
		85%				97%	
FA	54-6	83%	78.5%	UA	54-6	50%	55%
		74%				60%	
FC	54-6	41%	56.5%	UC	54-6		77%
		72%				75%	
FN	54-6		77.5%	UN	54-6		81.5%
1	11		HO!			76%	
		, , ,				,	

code:

F - Freemont U - Ute

A - air

C - carbon dioxide

N - nitrogen

the numbers signify temperature and time i.e. $4-6 = 4 \,^{\circ}\text{C} - 6$ months

1.8 1-12 = 4°C - 12 months

Wheat Data Summary Sheet - Germination Test - 12 months

c	code	%germ	ave	(code	%germ	ave
FA	4-12	92% 95%	93.5%	UA	4-12	91% 88%	89.5%
FC	4-12	91% 87%	89%	UC	4-12	85% 89%	87%
FN	4-12	92% 87%	89.5%	UN	4-12	89% 82%	85.5%
FA	21-12	89% 98%	93.5%	UA	21-12	95% 94%	94.5%
FC	21-12	92% 90%	91%	UC	21-12	93% 92%	92.5%
FN	21-12	91% 93%	92%	UN	21-12	93% 95%	94%
FA	37-12	84% 94%	89%	UA	37-12	78 % 85%	81.5%
FC	37-12	498 878	68%	UC	37-12	70% 91%	80.5%
FN	37-12	79 % 91 %	85%	UN	37-12	81% 83%	82%
FA	54-12	13% 0%	6.5%	UA	54-12	8% 0%	4 %
FC	54-12	0% 0%	0%	υc	54-12	23% 3%	13%
FN	54-12	9% 10%	9.5%	UN	54-12	45% 24%	34.5%

code:

F - Freemont

A - air

N - nitrogen

U - Ute

C - carbon dioxide

the numbers signify temperature and time i.e. 4-12 = 4°C - 12 months

Wheat Data Summary Sheet - Germination Test - 18 months

c	code	%germ	ave	(code	%germ	ave
FA	4-18	91% 90%	90.5%	UA	4-18	91% 88%	89.5%
FC	4-18	86% 83%	84.5%	UC	4-18	92% 87%	89.5%
FN	4-18	84% 88%	86%	UN	4-18	938 928	92.5%
FA	21-18	978 908	93.5%	UA	21-18	98% 94%	96%
FC	21-18	90% 95%	92.5%	UC	21-18	94% 88%	91%
FN	21-18	95% 96%	95.5%	UN	21-18	95% 93%	94%
FA	37-18	87 % 73 %	80%	UA	37-18	74 % 83%	78.5%
FC	37-18	63% 68%	65.5%	UC	37-18	82% 90%	86%
FN	37-18	75ዩ 82ዩ	78.5%	UN	37-18	87% 95%	91%
FA	54-18	0% 0%	0%	UA	54-18	0% 0%	0%
FC	54-18	0% 0%	0%	UC	54-18	0% 0%	0%
FN	54-18	0% 0%	0%	UN	54-18	0% 0%	0%

code:

F - Freemont A - air

U - Ute

C - carbon dioxide

N - nitrogen

the numbers signify temperature and time i.e. $4-18 = 4 \,^{\circ}\text{C} - 18 \,^{\circ}\text{months}$

Wheat	Data	Summary	Sheet	_	Baking	Test	_	control
-------	------	---------	-------	---	--------	------	---	---------

code	volume	ave	code	volume	ave
Fre ctrl	1450 ml 1480 ml	1465 ml	Ute ctrl	1490 ml 1470 ml	1480 ml

Wheat Data Summary Sheet - Baking Test - 6 months

cod	le t	volum	ne	ave		c	code	volu	ne	ave	
FA 4	-6 <u>1</u>	1285 1360		1323	ml	UA	4-6	1540 1400		1470	m1
FC 4	-6		ml	1313	ml	ÜC	4-6		ml	1475	ml
FN 4		1250 1390		1320	ml	UN	4-6		ml	1375	ml
FA 21		1500 1320		1410	ml			1300	ml	1300	ml
FC 21		1560 1360		1460	ml	υc	21-6	1560 1460		1510	ml
FN 21		1590 1370		1480	ml	UN	21-6		ml	1380	ml
FA 37	-	1460 1440		1450	ml	UA	37-6	1495 1635		1565	ml
FC 37		1500 1230		1365	ml	UC	37-6	1450 1480		1465	ml
FN 37		1460 1340	_	1400	ml	UN	37-6		ml	1363	ml
FA 54		1130		1105	ml	UA	54-6	1315 1360		1338	ml
FC 54	-6		ml	1198	ml	ŪC		1135 1230	ml	1183	ml
FN 54	-6		ml	1175	ml	UN			ml	1370	ml

code:

F - Freemont

U - Ute

A - air

C - carbon dioxide

N - nitrogen

the numbers signify temperature and time i.e. $4-6 = 4 \,^{\circ}\text{C} - 6$ months

Wheat Data Summary Sheet - Baking Test - 12 months

c	code	volume	ave	code	volume	ave
FA	4-12	1355 ml 1310 ml	1333 ml	UA 4-12	1340 ml 1235 ml	1286 ml
FC	4-12		1290 ml	UC 4-12	1305 ml 1210 ml	1258 ml
FN	4-12		1373 ml	UN 4-12		1238 ml
FA	21-12	1270 ml 1245 ml	1258 ml	UA 21-12	1315 ml 1195 ml	1255 ml
FC	21-12	1260 ml 1275 ml	1268 ml	UC 21-12	1125 ml 1175 ml	1150 ml
FN	21-12	1280 ml 1260 ml	1270 ml	UN 21-12	1240 ml 1295 ml	1268 ml
FA	37-12	1300 ml 1190 ml	1245 ml	UA 37-12	1115 ml 1200 ml	1158 ml
FC	37-12	1265 ml 1175 ml	1220 ml	UC 37-12	1000 ml 1130 ml	1065 ml
FN	37-12	1125 ml 1225 ml	1175 ml	UN 37-12	1150 ml 1230 ml	1190 ml
FA	54-12	1310 ml 1130 ml	1220 ml	UA 54-12	1025 ml 1100 ml	1063 ml
FC	54-12	1075 ml 1140 ml	1108 ml	UC 54-12		1053 ml
FN	54-12	1105 ml 1175 ml	1140 ml	UN 54-12		1060 ml

code:

F - Freemont A - air N - nitrogen

U - Ute

C - carbon dioxide

the numbers signify temperature and time i.e. $4-12 = 4 \, ^{\circ} \text{C} - 12 \text{ months}$

Wheat Data Summary Sheet - Baking Test - 18 months

c	code	volume	ave	code	volume	ave
FA	4-18	1285 ml 1395 ml	1340 ml	UA 4-12	1395 ml 1225 ml	1310 ml
FC	4-18	1325 ml 1360 ml	1343 ml	UC 4-18	1315 ml 1355 ml	1335 ml
FN	4-18	1350 ml 1360 ml	1355 ml	UN 4-18	1245 ml 1330 ml	1288 ml
FA	21-18	1315 ml 1290 ml	1303 ml	UA 21-18	1240 ml 1310 ml	1275 ml
FC	21-18	1185 ml 1195 ml	1190 ml	UC 21-18	1100 ml 1220 ml	1160 ml
FN	21-18	1250 ml 1260 ml	1255 ml	UN 21-18	1175 ml 1360 ml	1277 ml
FA	37-18	1200 ml 1265 ml	1233 ml	UA 37-18	1230 ml 1220 ml	1225 ml
FC	37-18	1170 ml 1230 ml	1200 ml	UC 37-18	1140 ml 1100 ml	1120 ml
FN	37-18	1235 ml 1240 ml	1238 ml	UN 37-18	1260 ml 1160 ml	1210 ml
FA	54-18	960 ml 1090 ml	1025 ml	UA 54-18	950 ml 1060 ml	1005 ml
FC	54-18		875 ml	UC 54-18		910 ml
FN	54-18	900 ml 920 ml	910 ml	UN 54-18	920 ml 900 ml	910 ml

code:

F - Freemont

A - air

N - nitrogen

U - Ute

C - carbon dioxide

the numbers signify temperature and time i.e. $4-18 = 4 \,^{\circ}\text{C} - 18 \,^{\circ}\text{months}$

Wheat	Data	Silmmariz	Shoot	_	Drotein	Solubility	_	control
wneat	Data	Summary	Sueer	_	Protein	SOLUDILLEA	_	COULLOT

code	abs.	ave		code	abs.	ave
Fre ctrl	0.469 0.521 0.518	0.503		Ute ctrl	0.593 0.617 0.582	0.597
Wheat Data	a Summa	ry Sheet -	Protein S	olubility	- 6 mont	ths

code	abs.	ave	c	code	abs.	ave
FA 4-6	0.521 0.558 0.554	0.544	UA	4-6	0.631 0.624 0.628	0.628
FC 4-6	0.551 0.512 0.569	0.544	UC	4-6	0.648 0.653 0.642	0.648
FN 4-6	0.602 0.625 0.632	0.620	UN	4-6	0.612 0.613 0.618	0.614
FA 21-6	0.432 0.486 0.493	0.470	UA 2	21-6	0.537 0.521 0.520	0.526
FC 21-6	0.509 0.515 0.491	0.505	UC 2	21-6	0.479 0.445 0.453	0.459
FN 21-6	0.473 0.488 0.469	0.477	UN 2	21-6	0.468 0.479 0.489	0.479
FA 37-6	0.488 0.505 0.503	0.499	UA 3	37-6	0.493 0.511 0.507	0.504
FC 37-6	0.561 0.559 0.590	0.570	UC 3	37-6	0.521 0.541 0.545	0.536
FN 37-6	0.491 0.539 0.538	0.523	UN 3	37-6	0.539 0.575 0.561	0.558
FA 54-6	0.455 0.461 0.423	0.446	UA 5	54-6	0.462 0.483 0.497	0.481
FC 54-6	0.451 0.478 0.450	0.460	UC 5	54-6	0.455 0.508 0.485	0.483
FN 54-6	0.448 0.501 0.511	0.487	UN 5	54-6	0.484 0.513 0.522	0.506

Wheat Data Summary Sheet - Protein Solubility - 12 months

c	ode	abs.	ave	code	abs.	ave
FA	4-12	0.483 0.504 0.489	0.492	UA 4-12	0.555 0.509 0.518	0.527
FC	4-12	0.531 0.515 0.525	0.524	UC 4-12	0.518 0.539 0.532	0.530
FN	4-12	0.521 0.518 0.541	0.527	UN 4-12	0.531 0.550 0.556	0.546
FA	21-12	0.522 0.480 0.522	0.508	UA 21-12	0.535 0.507 0.511	0.518
FC	21-12	0.453 0.482 0.492	0.476	UC 21-12	0.495 0.489 0.507	0.497
FN	21-12	0.501 0.515 0.519	0.512	UN 21-12	0.570 0.583 0.610	0.588
FA	37-12	0.440 0.472 0.451	0.454	UA 37-12	0.430 0.449 0.439	0.439
FC	37-12	0.455 0.449 0.469	0.458	UC 37-12		0.509
FN	37-12	0.479 0.522 0.493	0.498	UN 37-12	0.513 0.544 0.546	0.534
FA	54-12	0.359 0.359 0.362	0.360	UA 54-12	0.502 0.519 0.507	0.509
FC	54-12	0.401 0.399 0.429	0.410	UC 54-12	0.514 0.519 0.489	0.507
FN	54-12	0.459 0.468 0.482	0.470	UN 54-12	0.499 0.521 0.522	0.514

Wheat Data Summary Sheet - Protein Solubility - 18 months

code	abs.	ave	·	code	abs.	ave
FA 4-1	8 0.481 0.498 0.480	0.486	UA	4-18	0.502 0.512 0.550	0.521
FC 4-1	8 0.488 0.493 0.530	0.504	UC	4-18		0.563
FN 4-1		0.538	UN	4-18		0.582
FA 21-1	0.452 0.450 0.493	0.465	UA	21-18	0.481 0.542 0.524	0.516
FC 21-1	8 0.481 0.495 0.502	0.493	υc	21-18	0.520 0.519 0.508	0.516
FN 21-1	0.472 0.468 0.395	0.445	UN	21-18	0.562 0.578 0.539	0.560
FA 37-1	0.421 0.435 0.462	0.439	UA	37-18	0.420 0.411 0.440	0.424
FC 37-1	0.432 0.456 0.441	0.443	UC	37-18	0.425 0.441 0.444	0.437
FN 37-1	0.463 0.460 0.489	0.471	UN	37-18	0.480 0.478 0.498	0.485
FA 54-1	.8 0.372 0.412 0.398	0.394	UA	54-18	0.391 0.332 0.375	0.366
FC 54-1		0.384	υc	54-18	0.392 0.409 0.383	0.395
FN 54-1	0.391 0.439 0.389	0.406	UN	54-18	0.428 0.441 0.487	0.452

The Effects of Packaging Gas, Temperature and Storage Time on Germination, Loaf Volume and Protein Solubility of Wheat

Scott W. Myers

Department of Food Science and Nutrition

M. S. Degree, August 1991

ABSTRACT

Freemont and Ute wheat cultivars were stored for 18 months in cans under air, carbon dioxide or nitrogen at temperatures of 4°C, 21°C, 37°C or 54°C. Replicate cans were removed from storage at 6 month intervals and analyzed for percent oxygen, percent germination, loaf volume, and protein solubility.

After 18 months of storage, temperature had the greatest effect on wheat viability and quality. Storage at 37°C and 54°C, resulted in significant decreases in oxygen, germination, loaf volume and protein solubility. Effects at 21°C were not as severe, however, loaf volume and protein solubility were significantly reduced. Effects at 54°C differed significantly from those at 4°C after 6 months. After 12 months at 37°C and 18 months at 21°C significant differences were also found. The three packaging gases performed differently depending on the test involved. No packaging gas was the best among all of the tests.

COMMITTEE APPROVAL:

hn Hal Johnson, Committee Chairman

H. Gill Wilton, Committee Member

Clayton/S. Huber, Committee Member

Mark J. Rowe Department Chairman