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# Improving the Mutritive Value of Cereal Based Foods

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Progress Report No. 1

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### Improving the Nutritive Value of Cereal Based Foods

Progress Report No. 1

Jan. - Dec., 1968

Final authorization for initiation of this research project was received in January, 1968. This report constitutes essentially the progress made during the first year under the contract.

The general objective of this research project is designed to improve the nutritional value of wheat-based food by supplementation and process modification without decreasing the food acceptability to the consuming people in North Africa and Pakistan. The studies were to be conducted in two parts. One deals with wheat product supplementation with lysine or protein concentrates and vitamins and minerals to determine the best nutritionally economical blends which will result in highly acceptable finished foods. The other part deals with process modifications to maximize the recovery of the nutritionally valuable constituents of wheat.

Both specific objectives are to be carried out in three phases; i.e., a survey phase, an experimental phase, on a field testing phase. The survey phase has been essentially completed although a follow-up trip may be needed as more information is developed as the experimental phase progresses.

This report will cover the progress made in the survey phase and experimental phase of the project.

#### PHASE I - THE SURVEY PHASE

The purposes of the survey phase were as follows:

- 1. To gain first-hand knowledge about North Africa and Pakistan including:
  - a. geography
  - b. agriculture
  - c. economy
  - d. characteristics of the population
  - e. history
- 2. To learn as much as possible about couscous and chapatis
  - a. composition
  - b. preparation
  - c. consumption
  - d. commercial processing
- 3. To determine dietary patterns and food consumption data
  - a. average
  - b. typical daily diets and intakes
- 4. To obtain information about the nutritional status of the population segments
- 5. To evaluate grain production, marketing and processing
  - a. production, current and potential
  - b. marketing and prices
  - c. milling
  - d. commercial processing of milled food products
- 6. To determine potential indigenous sources of nutrients
- 7. To obtain samples of grains, nutrient sources, and sample foods for analysis and evaluation
- 8. To determine pertinent research groups and research in each country
- 9. To establish contacts for cooperative in-country testing and development

To carry out these objectives, a trip was made by Dr. Charles

Dayce, Dr. William Hoover, and Prof. Arlin Ward during March, 1968, to Morocco, to FAO headquarters in Rome, Italy, and to Pakistan. It would be impractical to detail the mass of knowledge and information accumulated on this trip. It was felt that we had accomplished to varying degrees the nine purposes previously listed.

One finding is worthy of special mention. From information about typical daily food intekes, it became obvious that couse as, while important in the Moroccan diet, no longer represents the basic food position of bread. For this reason, we decided to include Arab bread along with couseous in the supplementation studies to be carried out in the experimental phase.

However, to give an indication of the extent of the indormation gathered on the pages immediately following will be found:

- 1. Informational materials acquired
  - a. Morocco
  - b. Pakistan
  - c. Food and Agricultural Organization
- 2. Samples and process information obtained. (Actual samples not attached to sheets in this report. These sheets are duplicates of those appearing in our project master files which do have the samples mounted in the small squares on the sheets.) Enough of each sample was acquired for chemical and physical analysis.
  - a. Morocco
  - b. Pakistan
- 3. Pertiment contacts established

A number of photos were taken to tell the story to the team working on this project. Slides and movies were also made to show the techniques used in Morocco and West Pakistan in producing and processing their grain and grain products.

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- 2. Simplified Handbook on Mutrition for Personnel Conducting Feeding Programs Overseas, Melvin B. Myers (2 copies)
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- Morocco Role of Fertilizer in Agricultural Development with special emphasis on wheat, Thurman M. Kelso, Ralph E. McKnight, John L. Nevins, and Darrell A. Russell
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- 12. Final Report to The Agency for International Development U. S. Dept. of State on Phase 1 of Feasibility Study for Locally produced Wheat Based Protein Food Products in Tunisia by the International Milling Co., Inc.
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- 19. Alimentation at Travail Manuel Agricale (French)
- 20. Rapport d'une enquate par sondage sur l'etat de nutrition d'enfants Marocains de 1 jour a 14 ans inclus, Professeur Andre Recult (French)

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- 22. A Resume of population and health statistics for Morocco (3 copies)
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- 18. Pakistan Council of Scientific and Industrial Research (2 copies)
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- 21. Data on Oil Seeds: West Pakistan
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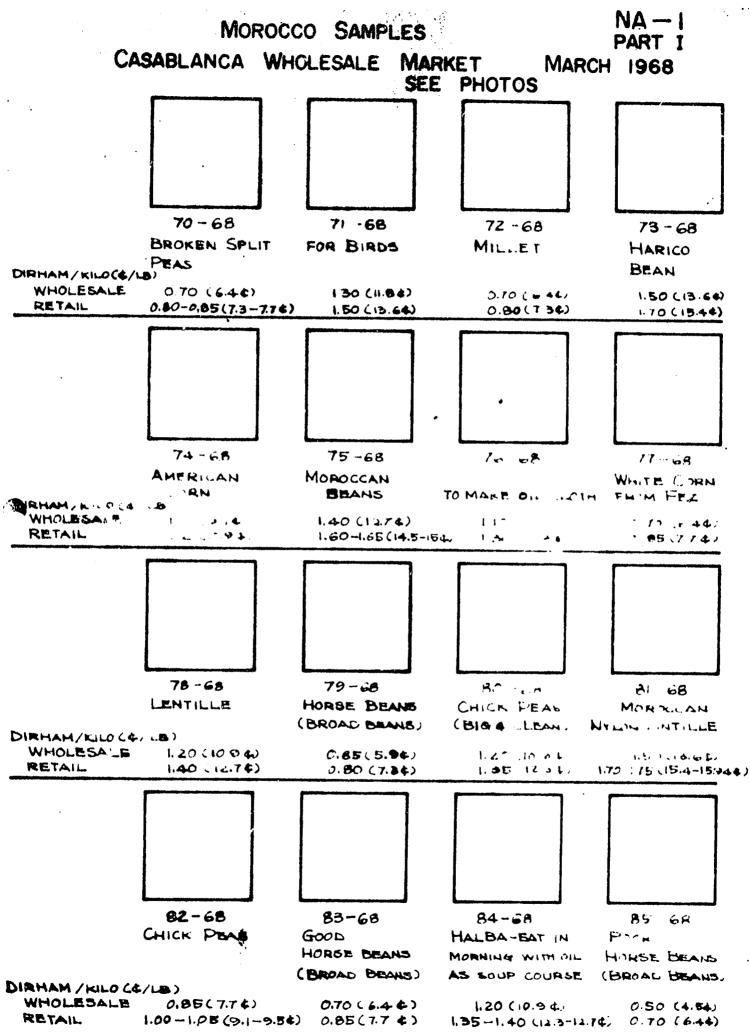
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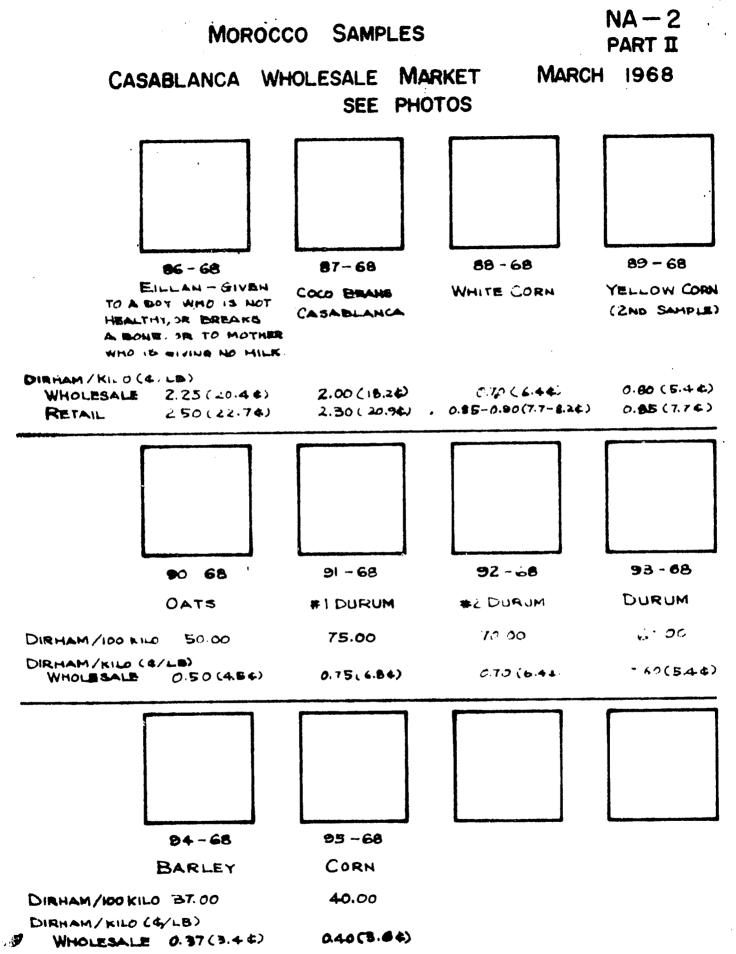
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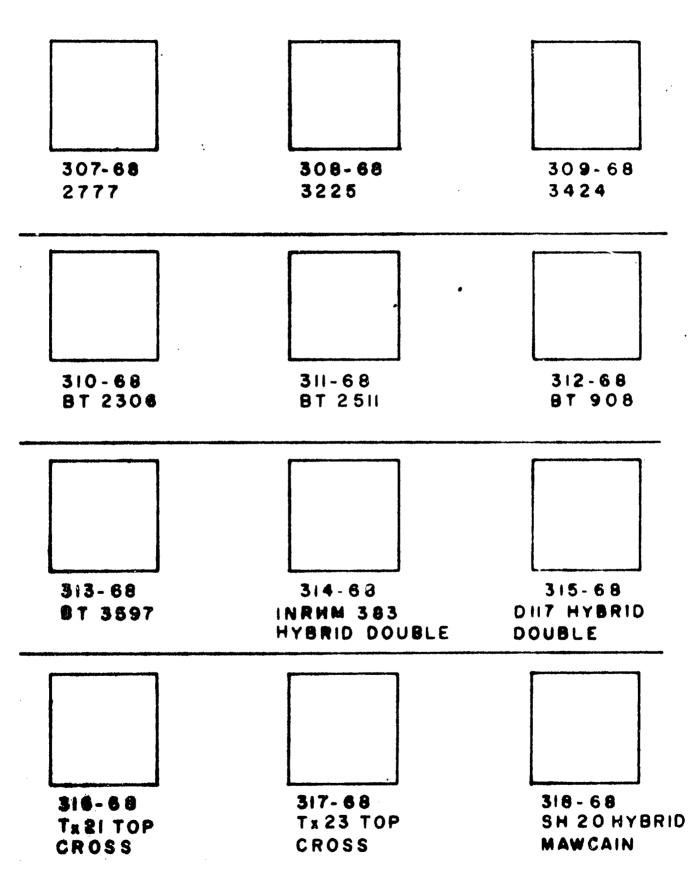
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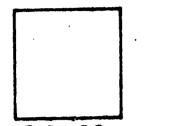


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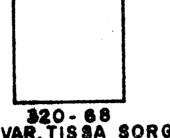
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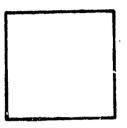




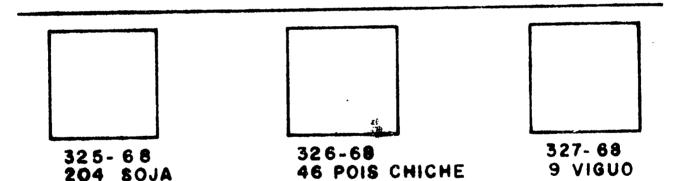
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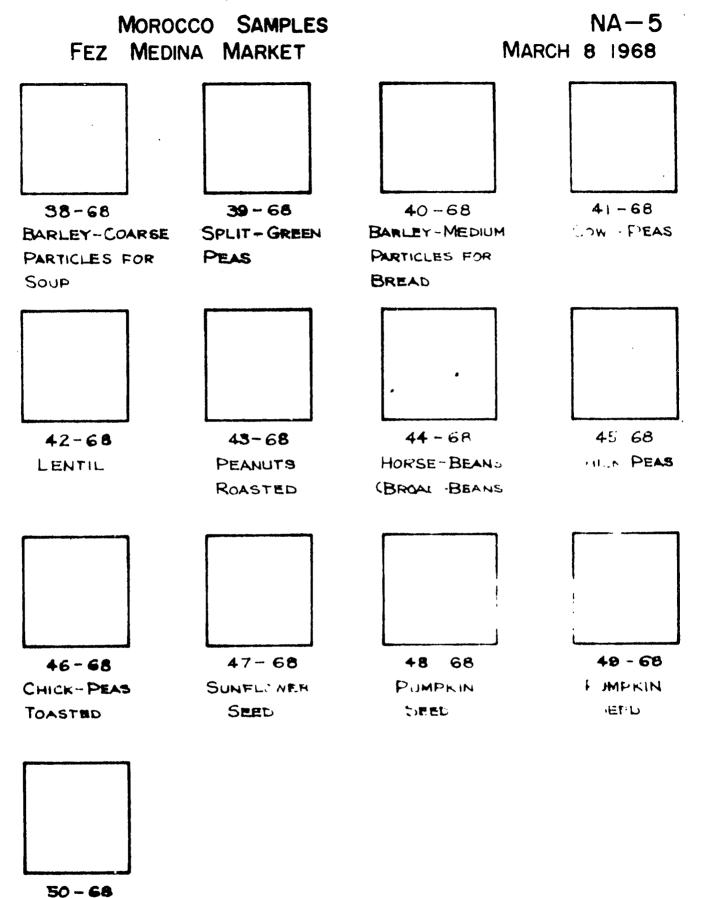


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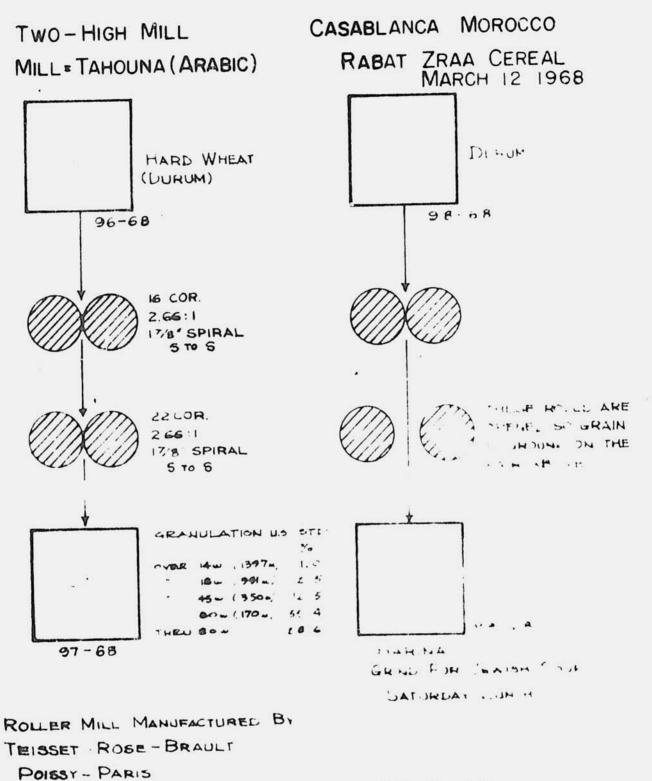


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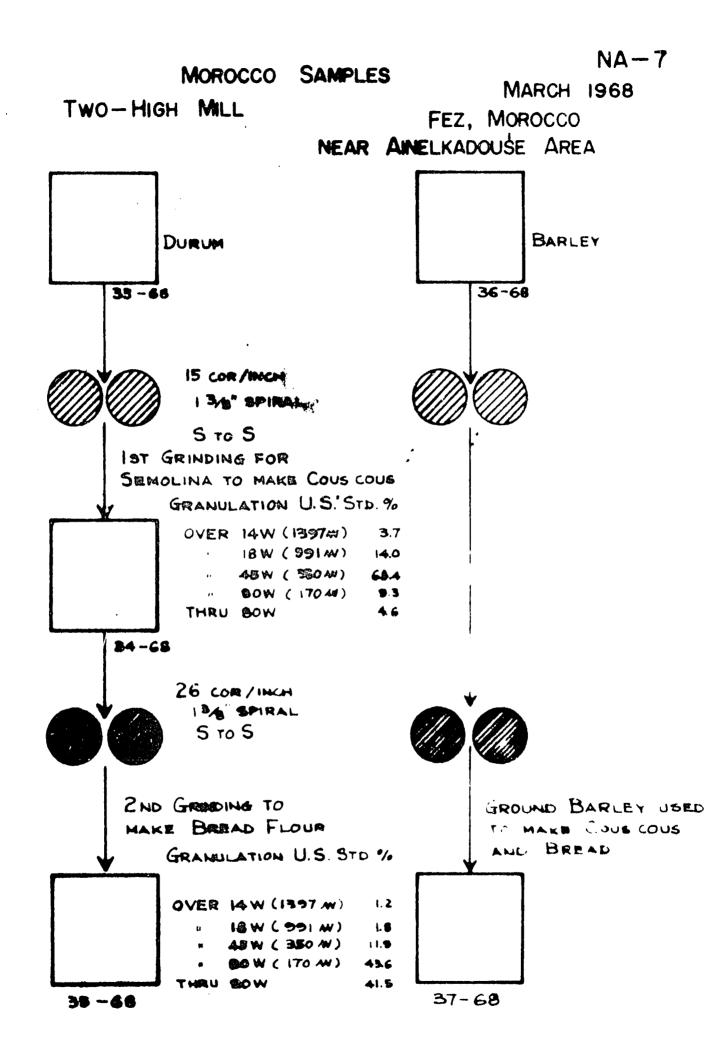
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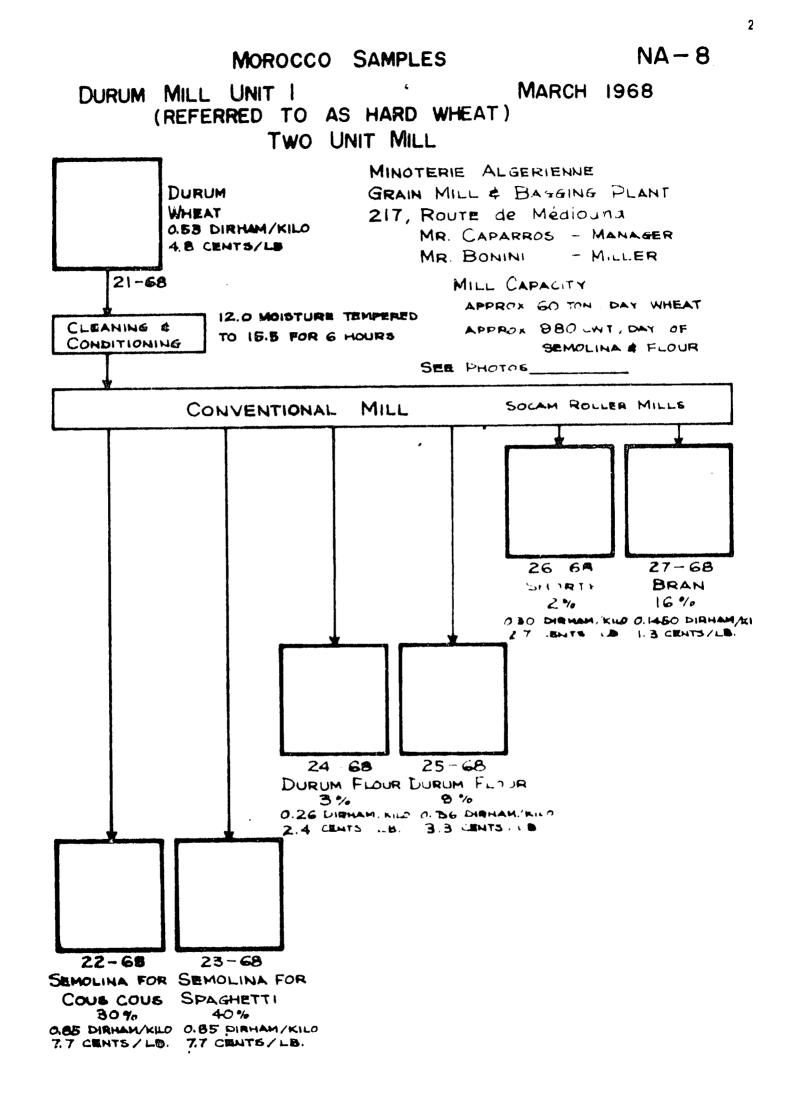


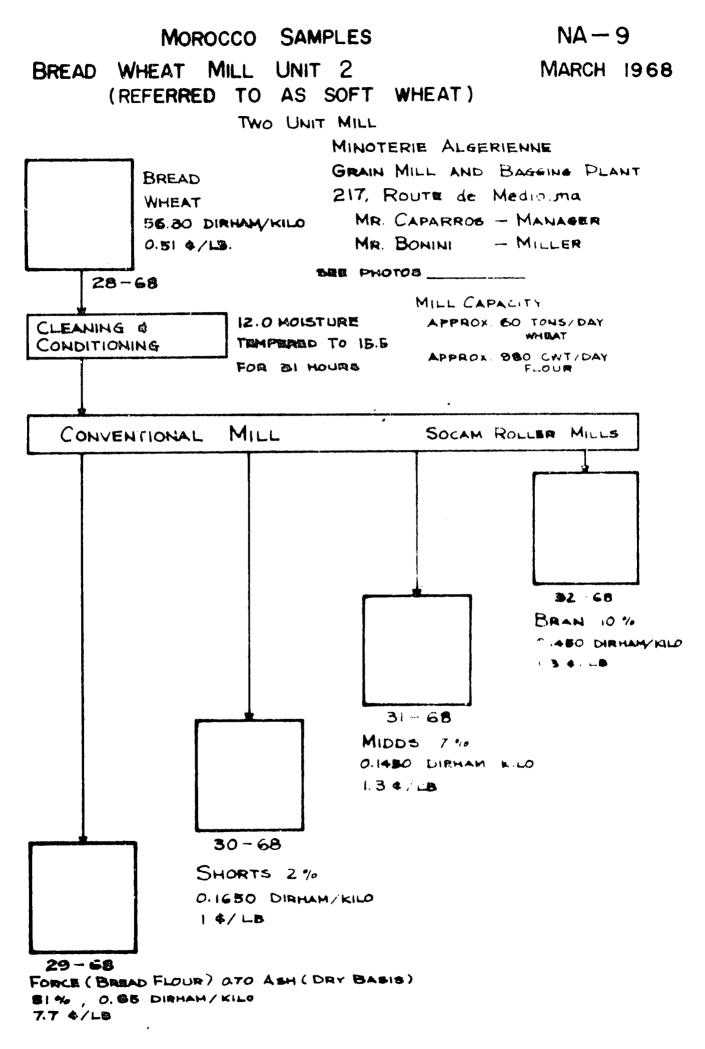
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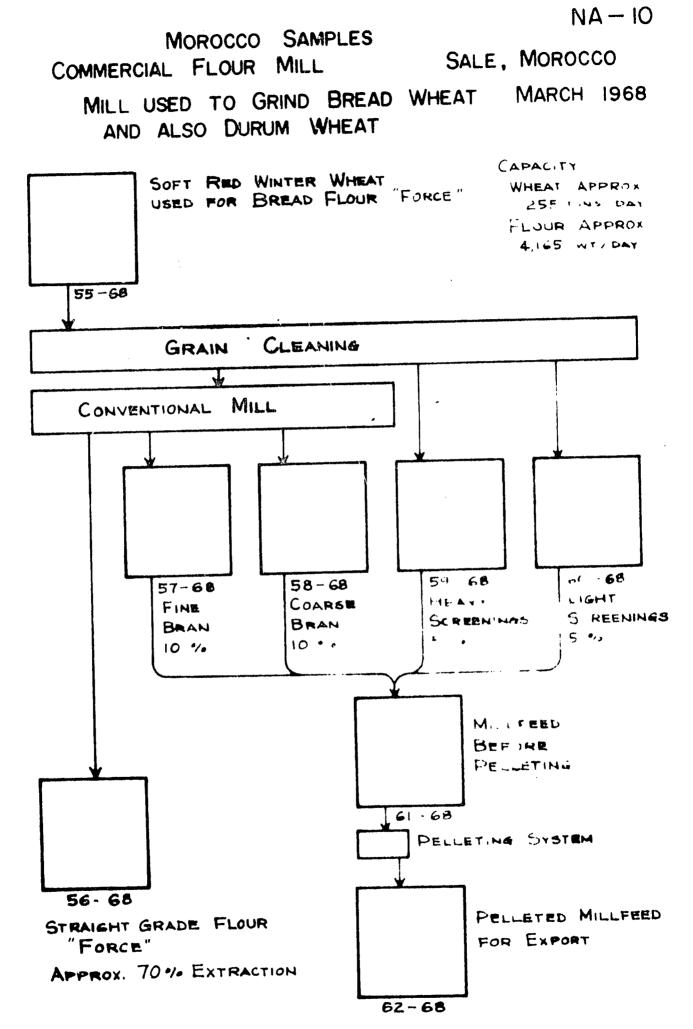
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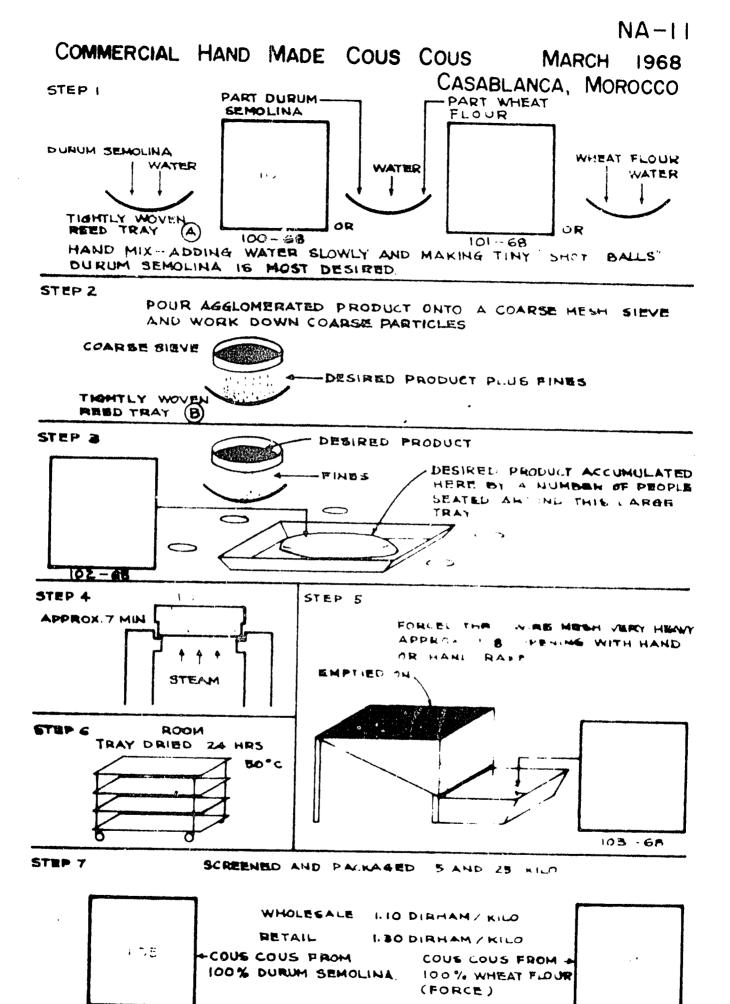
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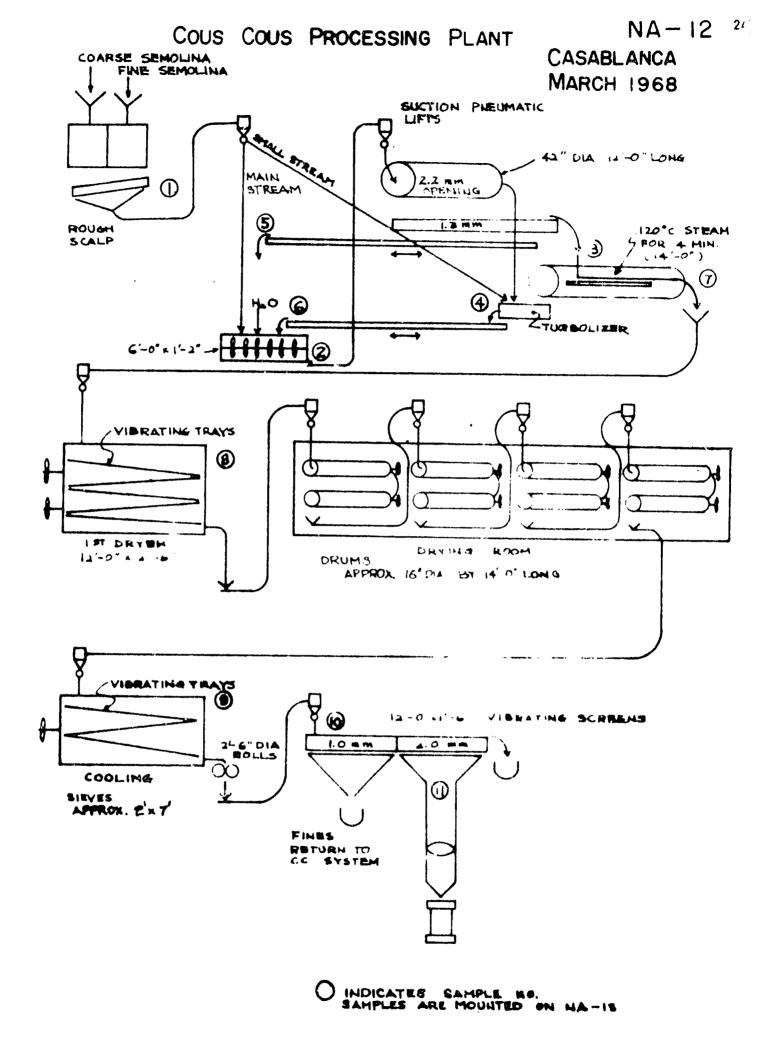


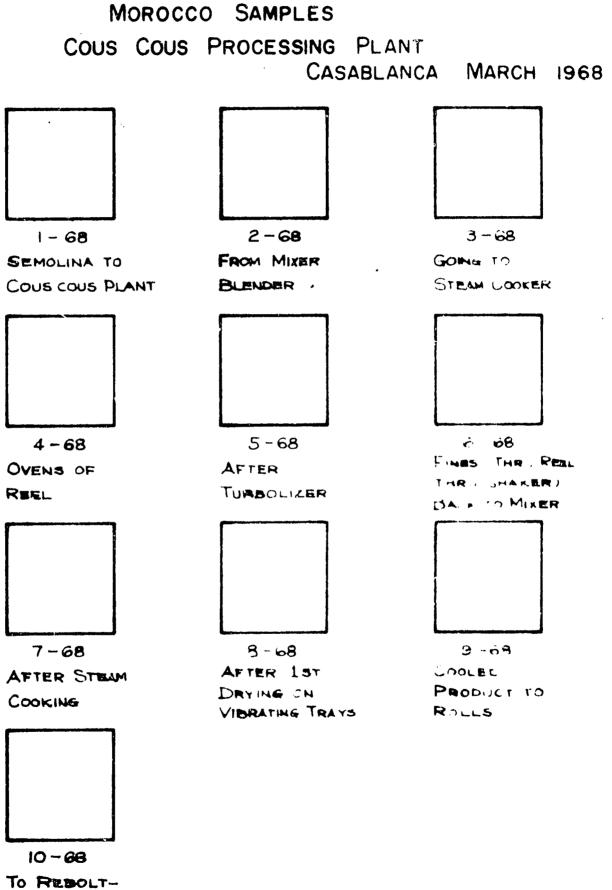


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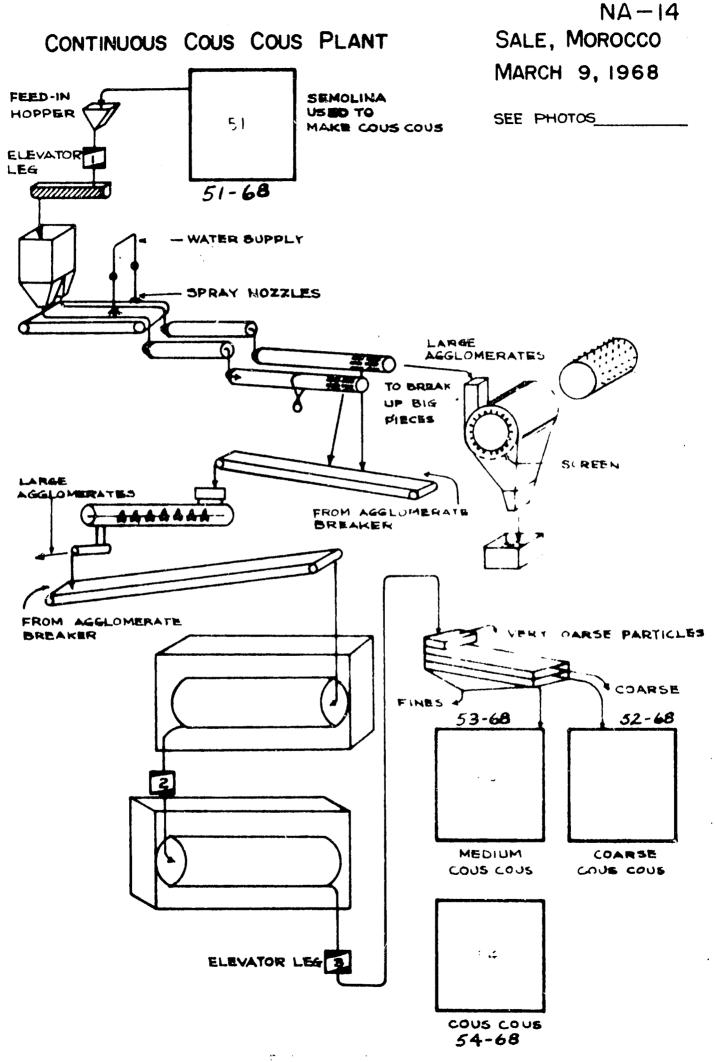
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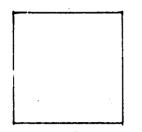


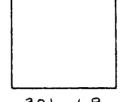
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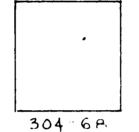
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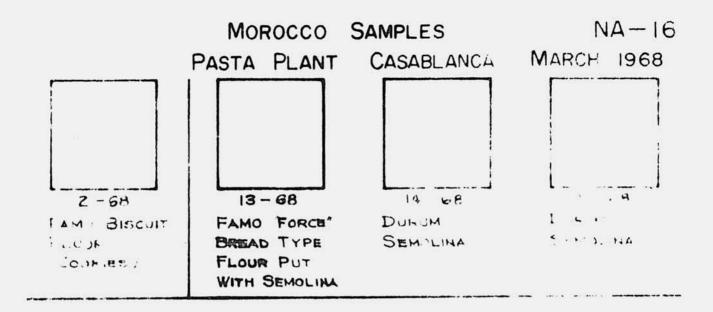


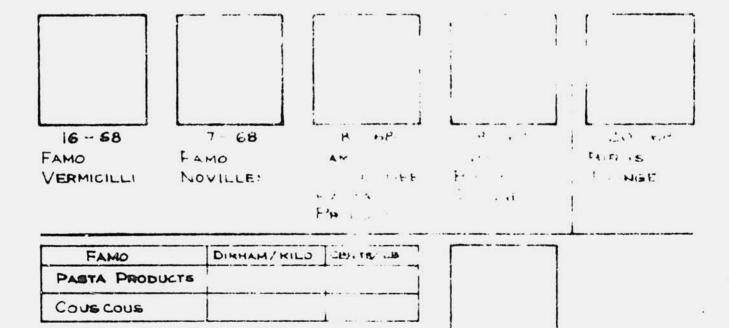
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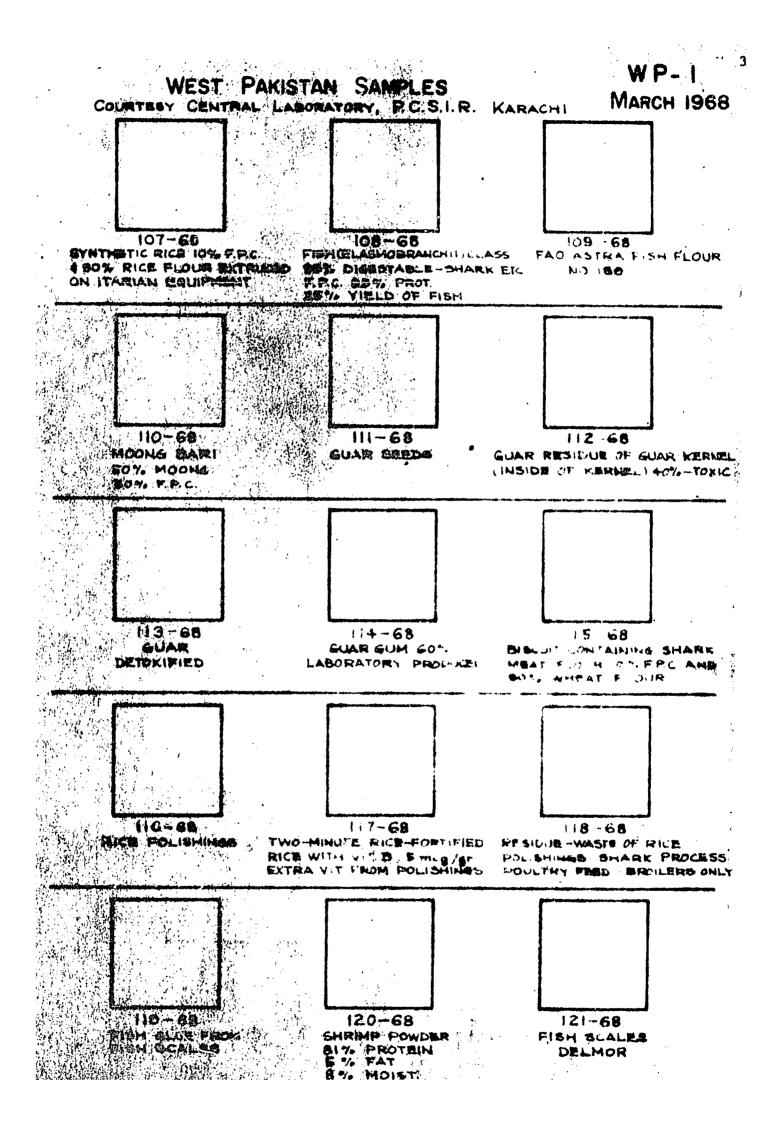


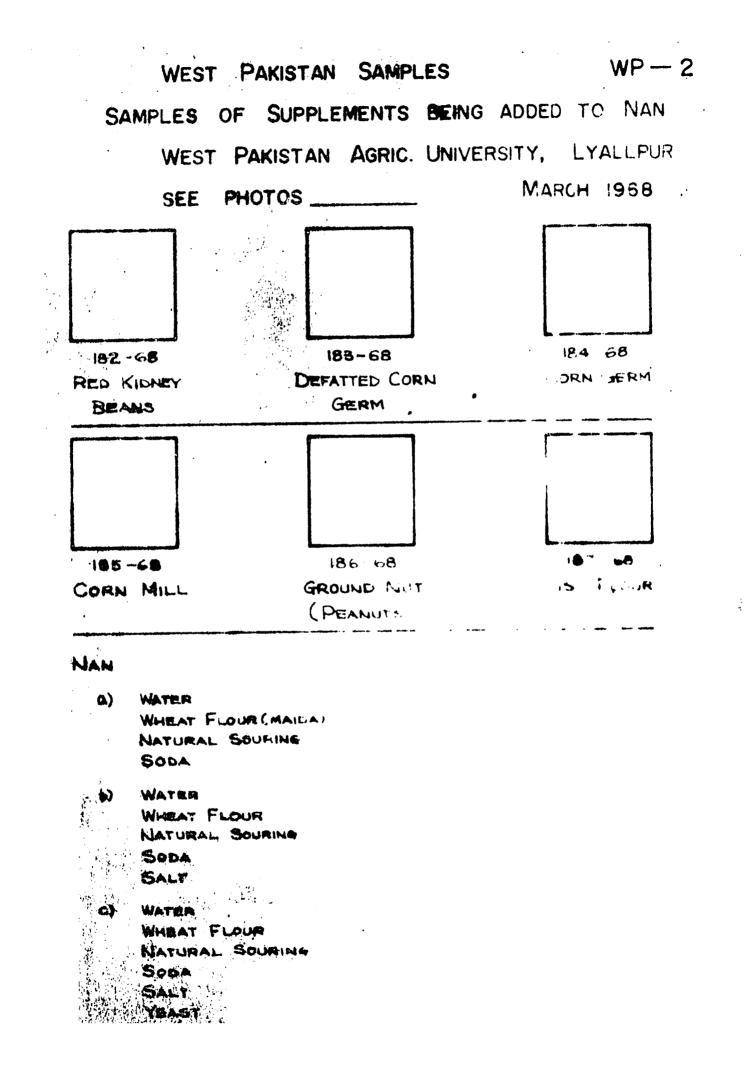


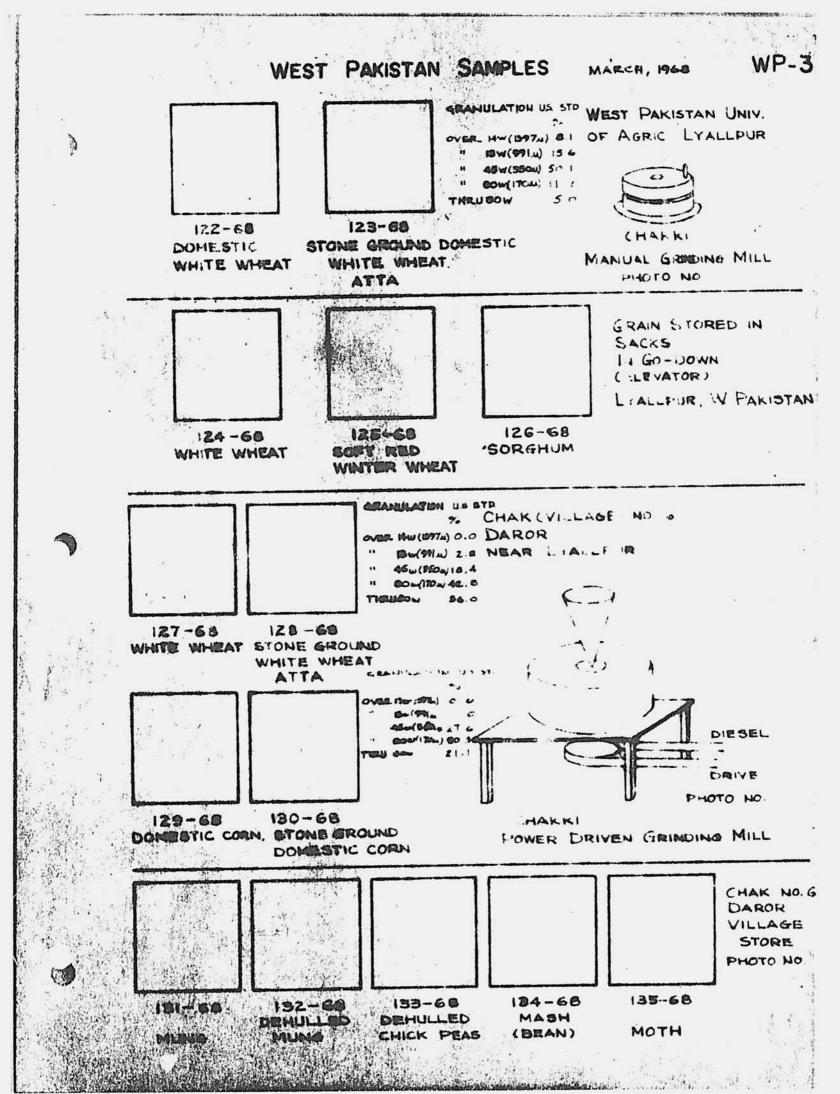


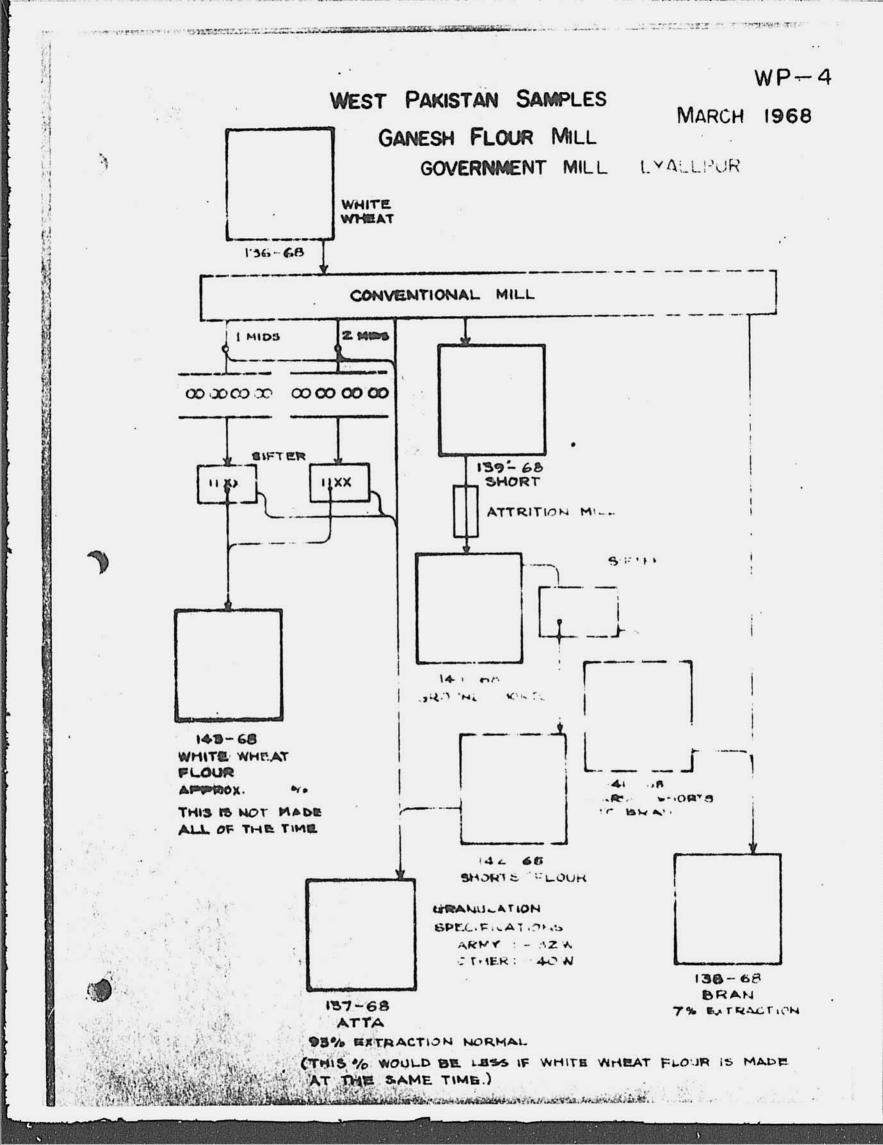
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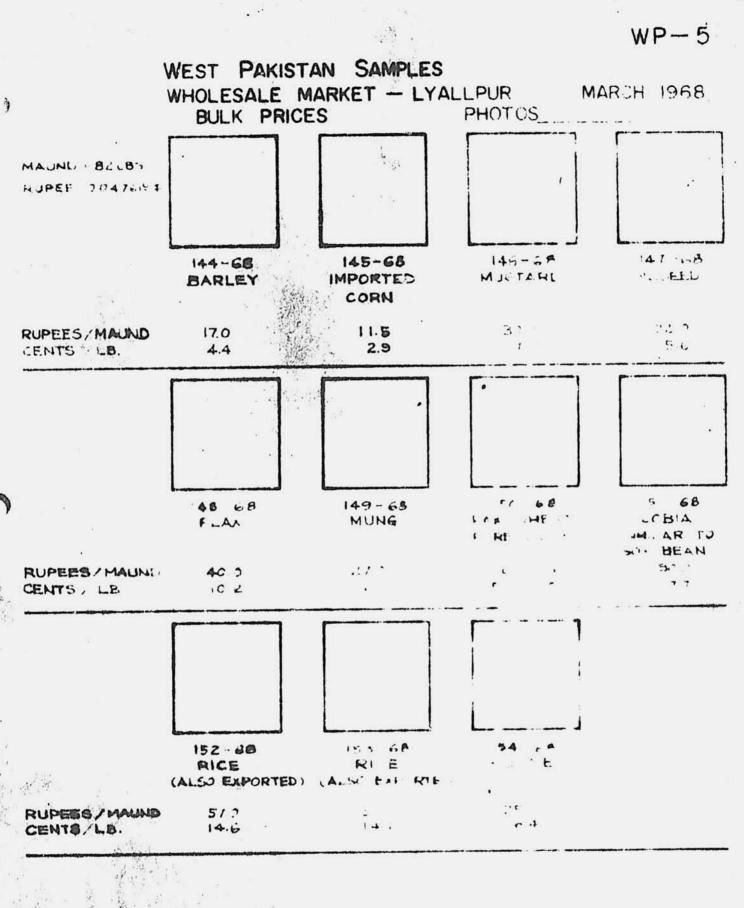
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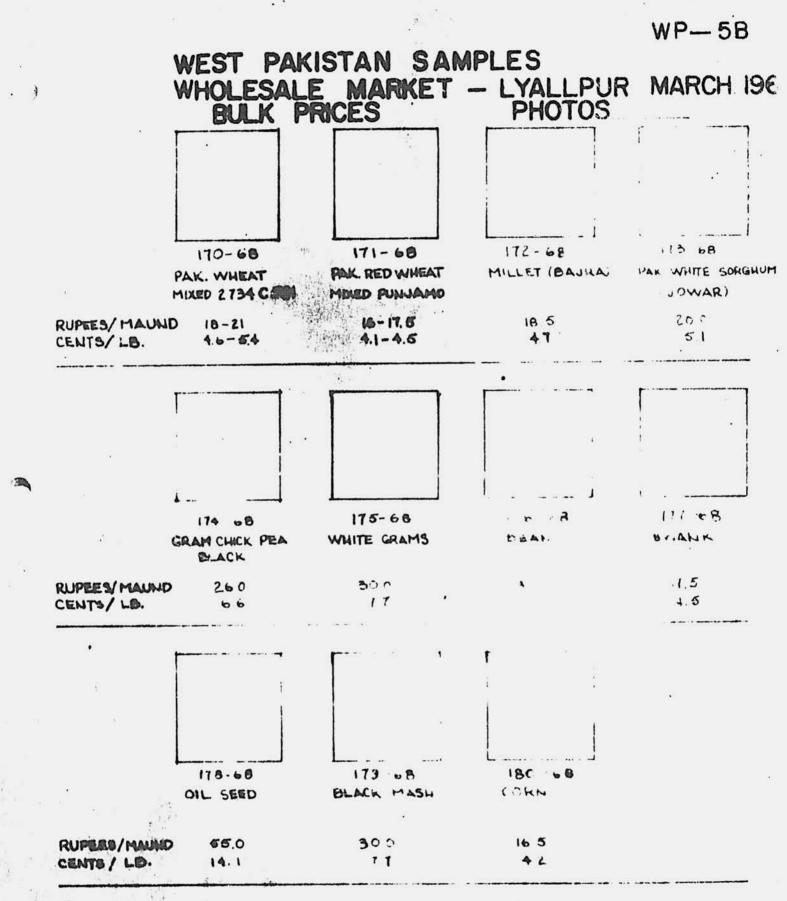






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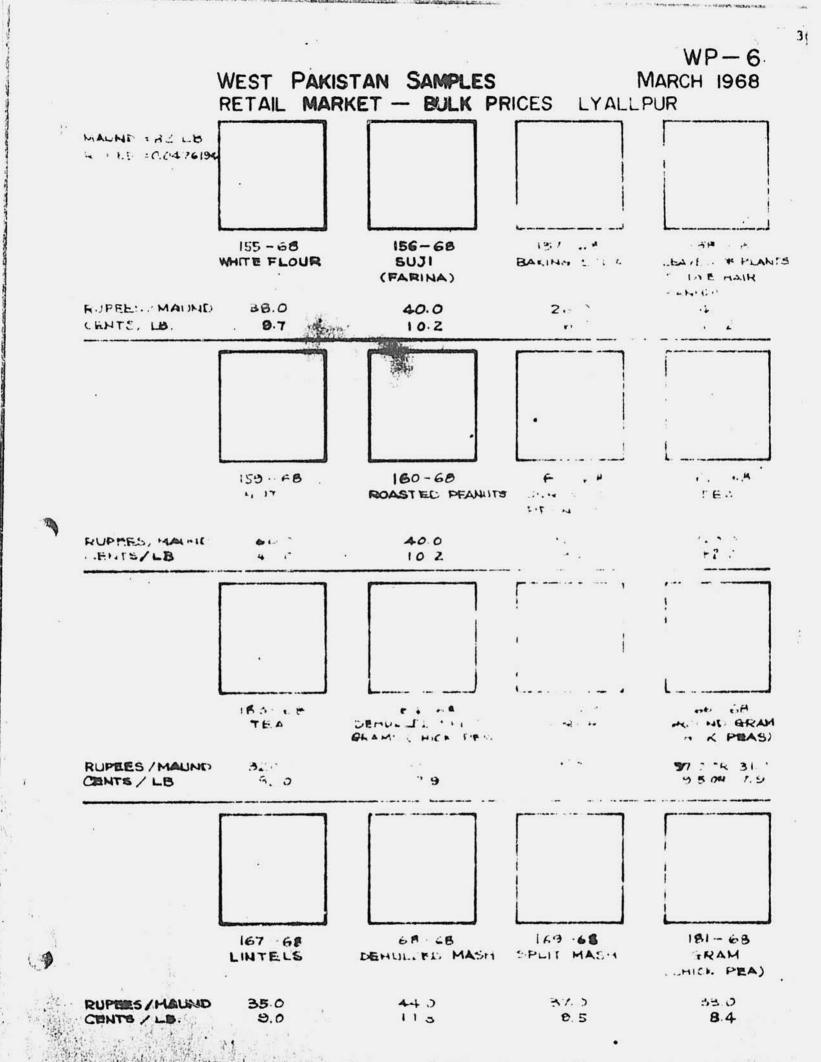
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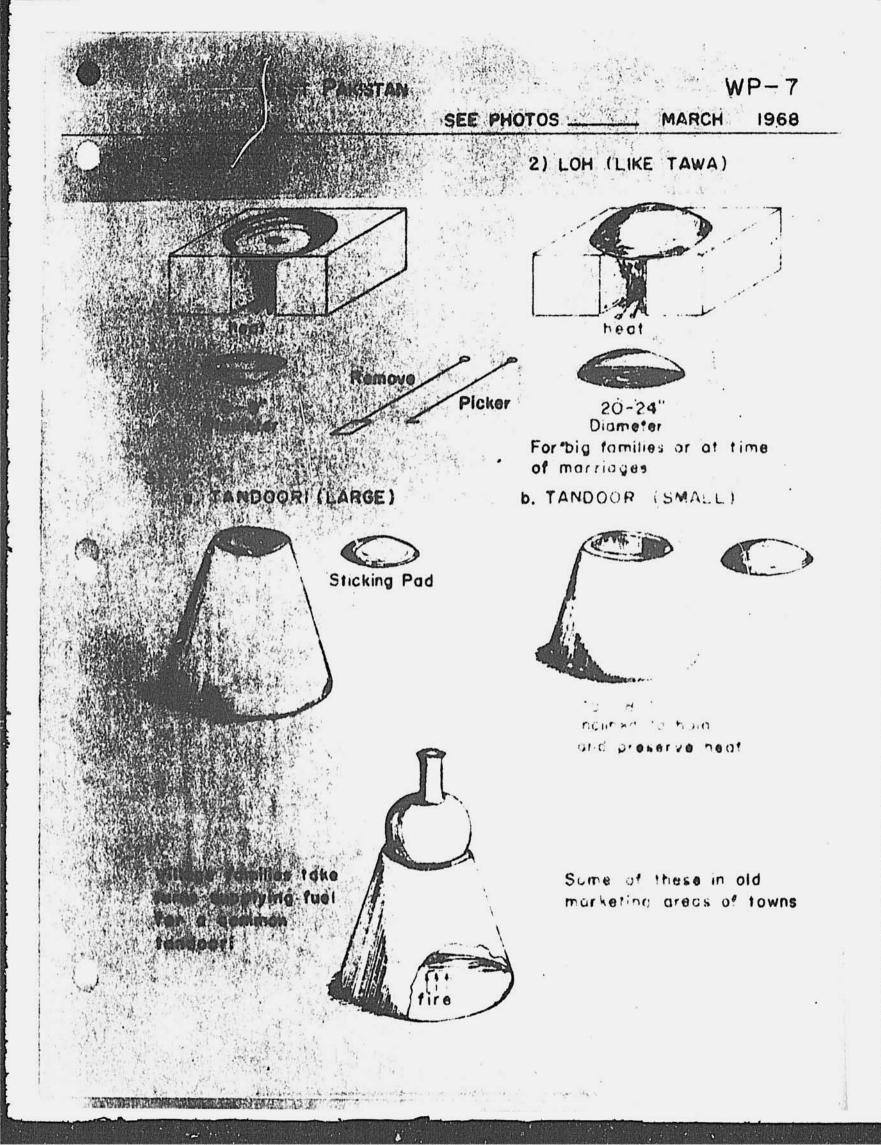
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WEST PAKISTAN INDEX

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#### PHASE II - THE EXPERIMENTAL PHASE

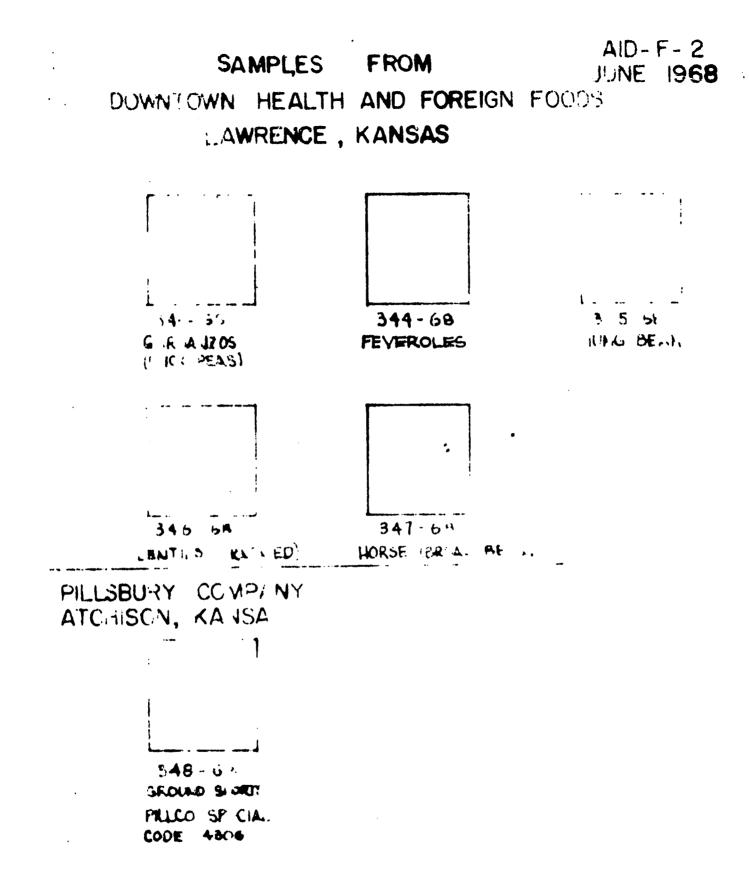
#### MATERIALS

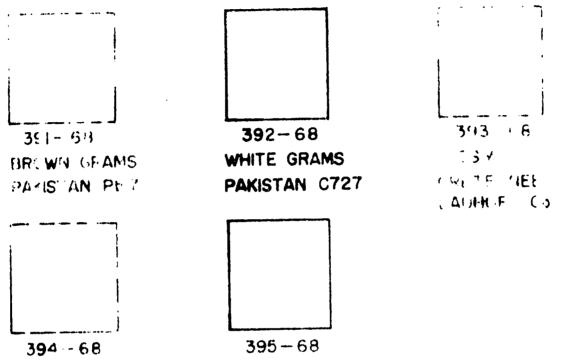
The processing of grain and other food products is done in many different ways. A study of the present methods used in Morocco and West Pakistan was made. Samples of the grain and grain products were procured from the processing plants.

Grain and other available products that would be potential supplements for cereal based foods were procured. When possible simplified flow diagrams were made showing the procedures used in processing. The local market place--both wholesale and retail--were visited and samples were obtained wherever possible. The section of this report beginning on page 12 indicates the type of fmaterials and process products obtained.

After identifying the samples obtained in Morocco and West Pakistan, we have been procuring similar materials in the United States with which to work. Some of these are depicted on the next two pages.

The potential nutrient supplements can be derived from fractions of the native materials, from protein concentrates, or from synthetic nutrients. The following supplemental materials have been obtained and are currently under study: Peanut flour; fish protein concentrate; soy isolate; soy flour; high lysine corn grits; corn-soy-milk (CSM); grain sorghum protein concentrate; wheat protein concentrate; glandless cottonseed flour; horsebean flour; chick pea flour; yeast concentrate; and L. Lysine-HC1. 41





PEANUT FLOUR

FISH FLOUR

## MILLING AND PROCESSING OF PRODUCTS AND SUPPLEMENTS

Techniques have been worked out to produce materials similar to the native products in analytical properties as well as physical properties such as particle size. A considerable amount of material for couscous manufacture and also atta for making chapati and roti products has been produced. We are developing a better understanding of the effect of processing on the end use of the products. <u>COUSCOUS</u>

An outline of work underway and the experimental milling flours developed are given on the pages 46 through 48. Preliminary milling studies have been completed on a 97% extraction flour and semolina from durum wheat. The granulation and protein content of the 97% extraction flour are very nearly the same as those obtained in Morocco. The couscous preparations have been compared with the commercial Moroccan products and have met the approval of a taste panel of international students familiar with couscous. Improved cooking utensils have been obtained through the assistance of the Moroccan Embassy. 45

#### An Outline of a Work Schedule for a Couscous Study

#### Allen Kirleis

- I. Milling procedure
  - A. 97% extraction flour
  - B. Semolina
  - C. 97% extraction flour and "horsebean flour" of "chick pea flour"
- II. Develop a standard method for preparation of couscous
- JTI. Keeping quality
  - A. Of 97% extraction flour and semolina and 97% extraction flour and horsebeans
    - 1. Development of fat acidity
  - B. Of dried couscous made from 97% extraction flour and semolina and 97% extraction flour and horsebeans
    - 1. Development of fat acidity
    - 2. Possibly organoleptic test
- IV. Nutritive quality
  - A. Determine vitamins and availability of protein on 97% extraction flour and semolina and 97% flour and horsebeans
  - B. Determine vitamins and availability of protein on couscous made from 97% extraction flour and semolina and 97% flour and horsebeans
  - C. Possibly rat studies of 97% extraction flour, semelina and couscous and blend
- V. Acceptability of couscous from A, B, and C in I above

# SAMPLES PRODUCED AT MANSAS STATE UNIVERSITY

## AIL - F - 3 J INE 1968

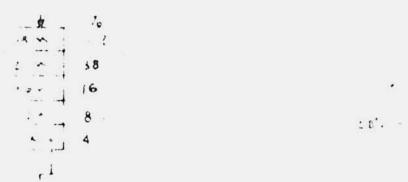


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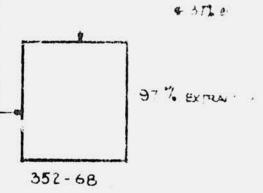
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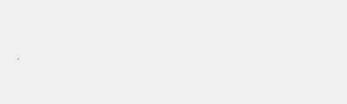


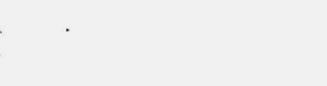
SEMOLINA FOR BREAD (FINE)





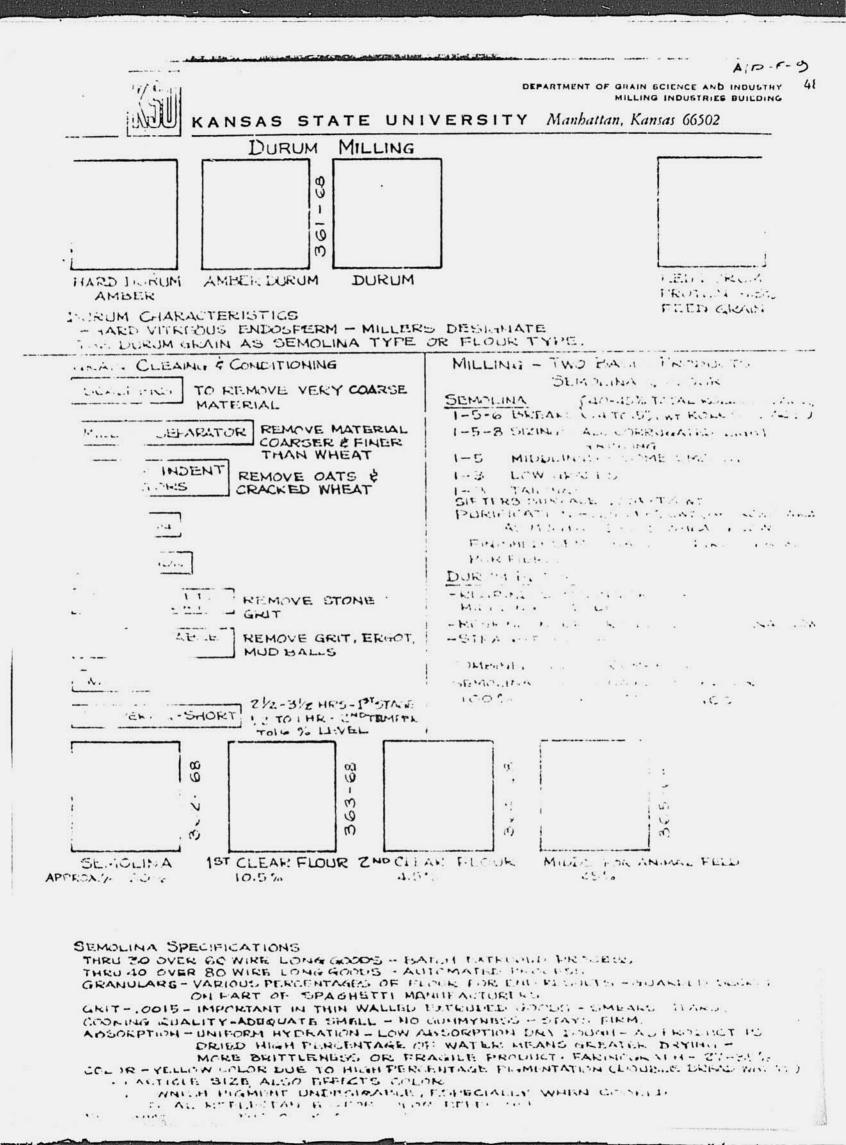












#### MOROCCAN BREAD

Two studies are underway on Moroccan Breads. An Outline of Work on Characterizing and Supplementing Moroccan bread is given on page and a milling flow developed for Moroccan bread flour is given on page

A program of study investigating new methods of milling wheat for Moroccan bread flour production and for processing chick pea and horsebean flours for supplementation is outlined on pages 52 and 53.

Chick peas and horsebean samples were obtained from Morocco. Their preliminary analyses indicated a high level of protein content. The amino acid contents for these legumes showed good levels of essential amino acids. It was proposed to use the flour of these legumes in fortifying the local bread to increase its nutritive value.

Samples similar to these legumes were obtained from a foreign food stored in Kansas City (Scimico Italian Supermarket). It was found that the chick pea was a product of Morocco, the horsebean was an Italian product.

Experimental milling was done with these legumes and experimental flows were developed (pages 54 and 55). Analyses were made for each fraction obtained. Horsebean flour was fractionated according to particle size and analyses for moisture content, protein, ash, crude fat and crude fiber were made.

#### An Outline of a Work Schedule for Study of Moroccan Breads

#### Luis Aira

This involves the following grains--rye, barley, millet and wheat (soft red winter)

1. Moroccan Bread

Formulation

Test Baking for Standard

Characterization of Standard

Evaluation

2. Grains -- Physical Properties

Wheat

Rye

Barley

Millet

3. Supplementation and Enrichment

Legumes

Lysine

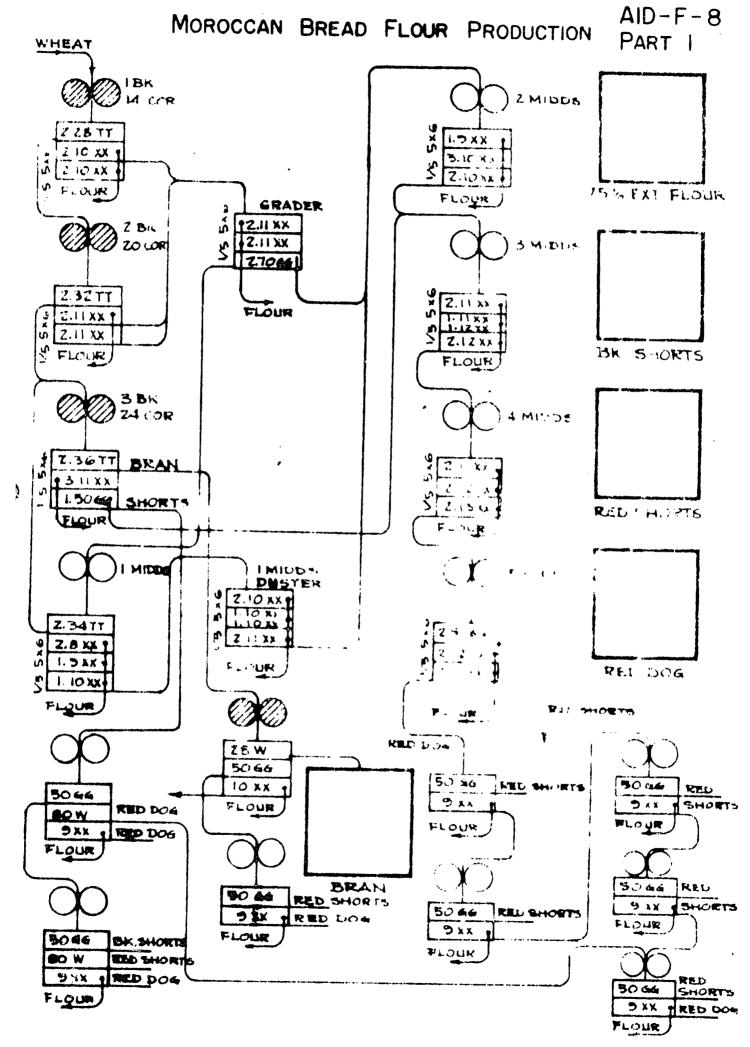
Standard Enrichment

Physical Properties and Baking

4. Characterization and Evaluation of Finished Products

Organoleptic Evaluation Panel

**Biological** 



The Proposed Outline and Schedule of Work on the Use of Chick Peas and Horsebeans In Moroccan Bread

M. Al Suaidy

I. Increasing yield and nutritive value of wheat products.

A. Grain and Treatment

.

Origin of Grain

From two Agricultural Stations in Kansas (Manhattan and Hays)

Variety:

Guide

Manhattan Protein 12.0% " 12.8% Hays Blended and tested here at KSU 13.9%

Grain: . . .

B. Treatments of Grain

Grain CONTROL ----- STANDARD Processing

Control ----- Sorghum Peeler

Cold Treatment

- 1. Temper at 4°C. water bath to 20% moisture.
- 2. Divide in half,
  - a. Freeze one half at -10°C. for 72 hours. b. Second half storage at 40°F. for 72 hours.
- 3. Remove grain from freeze and cold room and dry in air oven at 110°F. over night.
- C. Treatment on Sorghum "Peeler"

Control

Frozen Grain

Non-Frozen Grain

Non-frozen grain (cont.)

1. Short Tempering 25% moisture for 30 minutes.

- 2. Run through sorghum peeler at 1350 RPM with feeding gage at 4.5.
- 3. Sieve and Air separate--determine yield.
- 4. Dry in air oven at 110° F. over night,
- 5. Approximate analysis of fraction.
- 6. Store grain at 40°F.
- D. OTHER TREATMENTS: (Proposed)

Using

- 1. Softner
- 2. Pearling or Engelburg Huller
- II. Supplementation Studies
  - A. MILLING STUDIES

Temper to 16% moisture for 24 hours.

Mill using a flow for 85 and 97% extraction.

B. FLOUR STUDIES

Analyses: Moisture, ash, protein, fat and fiber.

Physical Dough Tests; Farinograph, Extensograph and Amylograph and Flour Colorimeter.

C. BAKING TEST

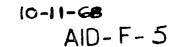
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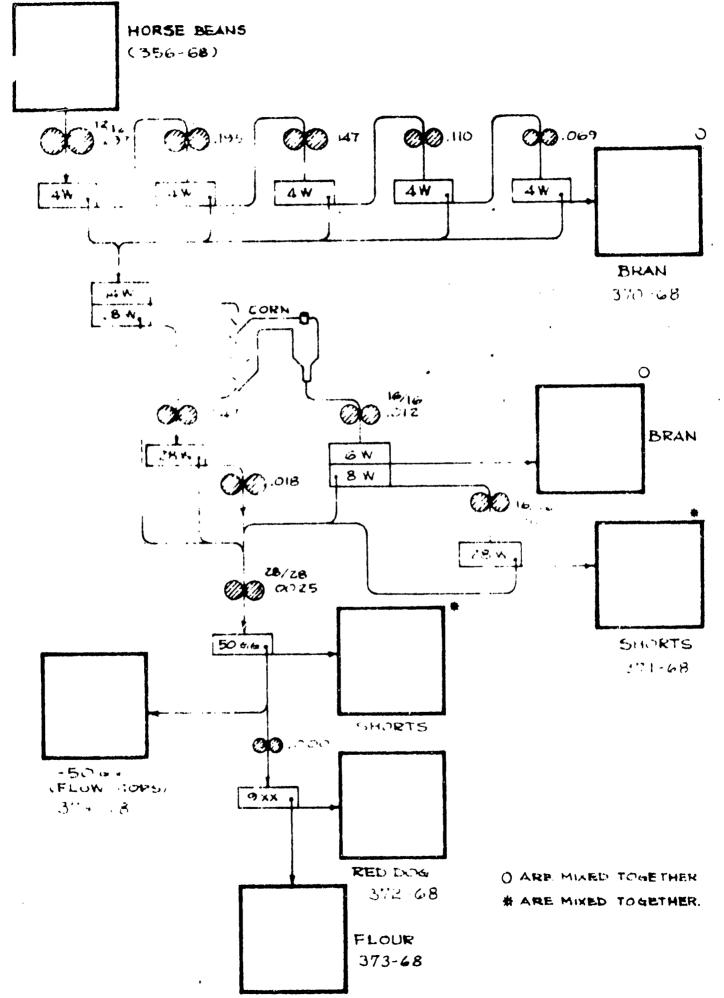
85% extraction American Standard 97% extraction Moroccan Bread

D. Supplementation on 85% and 97% Flour Extraction

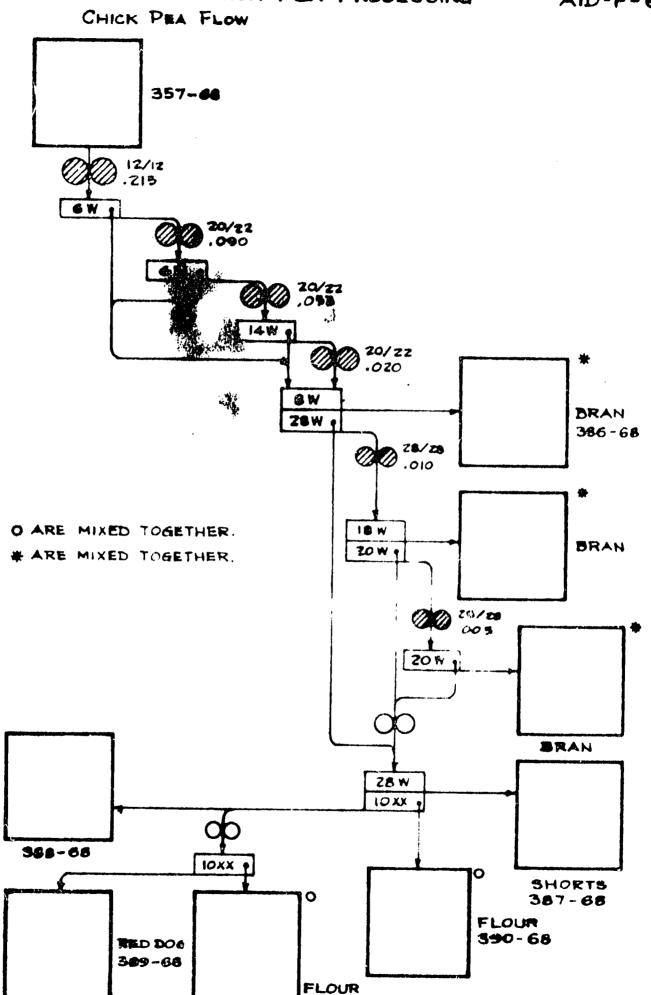
Horsebean Flour (0, 1, 3, 6, 9, 12 - 50%) Chick Pea Flour (0, 1, 3, 6, 9, 12 - 50%) 1. Flour analyses - proximate, minerals and amino acids.

- 2. Granulation and yield.
- 3. Effect on physical dough properties.
- Baking test scoring.
  Taste panel.
- 6. Bread analyses proximate minerals and amino acids.





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#### ROTI TRODUCTS

Considerable progress has been made on the milling and processing of roti products. The bulk of this progress is reported in the nutrition section as three papers submitted for publication (pages 10) through 151).

An outline of additional work is given on page 58.

Studies on the shelflife of Chapatis on Nan have been carried out as outlined on page 59. Each study was extended over a six-week period. In the packaging study, daily moisture losses were recorded during the initial week and then followed by weekly measurements.

The developing microbial population during storage was studied. Microbial evaluations were made on the initial ingredients, the mixes and the final products. It was found that the most undesirable microorganisms were <u>Aspergillus niger</u>, <u>Aspergillus flavus</u>, <u>Penicillium</u> <u>lilacium</u>, and <u>Aspergillus fumigatus</u>. A study is now underway to add mold inhibitors that are acceptable under the laws of the countries concerned.

The shelflife and preserving studies are deemed necessary before marketing of chapatis and nan can be considered. Village or neighborhood bakeries are looked upon as advantageous developments to a nutrient supplementation program.

#### An Outline of the Work Schedule on Roti Products

#### Robert Tang

#### ATTA AND CHAPATI

1. Analysis for

Amino Acid

Vitamin - 3, B, Nicotinic Acid, Pyrodovine

Minerals

Fats - Fats Acidity

2. Supplemented ( 5, 10, 20, 30% )

Fish Protein (2)

Sorghum Protein

Wheat Protein Conc.

Lysine

Corn Protein Conc.

Corn Soy Milk

Legumes

Analysis As Above

3. Atta and Chapati From Other Grains

Corn

Sorghum

Millet

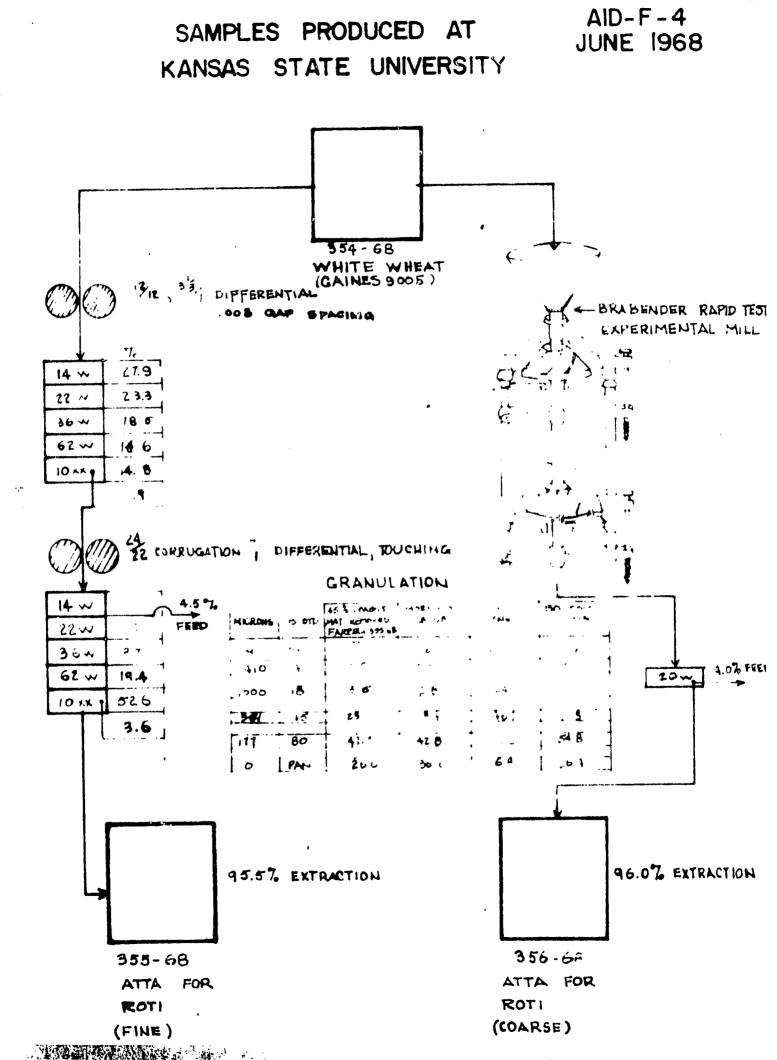
Barley

4. Characterization and Evaluation of

Finished Product

Crganoleptic Evaluation

Biological



## Microbiological Studies of Chapatis and Nan for Quality Control and Keeping or Shelf Life

#### Abdul Rashid Mann

- 1. Microbiological survey of raw materials including atta.
  - a. Bacterial
  - b. Mold
  - c. Yeast
- 2. Prepared chapatis and nans.

lst steps in preparation - Samples for microbiological testing 2nd step in preparation - Samples for microbiological testing 3. The prepared chapatis and nan

- 3. Storage
  - a. Keeping quality
    - General microbial determination
      Specific bacteria, yeast and mold
  - b. Pathogens Enterics
- 4 Packaging
  - a. open no coverage
  - b. Poly bags
  - c. Saran bag
  - d. Foil wrap
  - e. Waxed paper
  - f. Ordinary wrapping paper (Brown or white)

#### Protein Concentrate from Whast Bran and Shorts

Preliminary studies on producing protein concentrate for human food from wheat millfeeds have been undertaken. Samples of bran and shorts were sacked off the stream while milling a Kansas hard red winter wheat. Germ and red dog were sacked off separately.

Samples of bran and shorts were ground separately by a MIKRO Grinder with the rotor at 7300 r.p.m.; the separator at 2000 r.p.m.; the process air wt 400 cu. ft./min. Each ground sample was mixed by a tumbling blender and sifted by a G.W. lab sifter in 500 gram lots. Samples of the throughs were taken after each 30 seconds of sifting, weighed and analyzed separately.

Results of the analysis of fractions obtained are given on page 61. High Lysine Corn

Many 5-lb. lots of yellow corn and high lysine corn (both white and yellow) ware milled by many different flow diagrams to develop experimental milling procedures for the different kinds of corn, at low moistures (15-16%) and high moistures (22-23%) with and without impact type germination.

White high lysine corn grown in Kansar in 1967 was milled on experimental milling equipment in sufficient quantities to obtain 50 lbs. or more of flour, fine meal, and medium meal for testing by fine grinding and air classification for protein quantity and quality in the many fractions. This work is scheduled to start the week of December 11.

The analytical evaluation of the milling fractions obtained is given on page 62.

#### Sorghum Protein Concentrate

Considerable work has been done on developing high protein fractions from grain sorghum. Results of this work are found in the nutrition section on pages 158 to 172.

				Ground	500 g.		A SIEVE	*		
Jode	Sieve	Time Sec,	Cum. Time Sec.	Through g.	Cum. Through g.	Moist. %	Pro. %	Pro. 14‰ m.b.	Ash %	Ash 147. m.b.
FS-1	9XX	30	30	135	135	9.0	19.6	18.5	1.9	3.7
FS-2	11	30	< O	74	209	9.1	14.5	18.4	3.8	3.6
FS-3	11	30	40	61	270	9.3	19.0	18.0	3.7	3.5
FS-4		30	120	28	298	9.3	17.6	16.7	3.9	3.7
FS-5		30	150	16	314	9.2			4.1	3.9
FS-6	+9XX					9.2	13.5	12.8	4.4	4.6

# Ground Shorts Sifted on 9XX Sieve

-----

### Ground Shorts Sifted on 10XX Sieve 500 g. sample

	-11	10XX	30	30	75	75	9.3	20.4	19. 1	4.2	4.0
	-12	11	30	60	55	130	9.3	20.2	14.2	3.9	3.7
201	-13	"	30	90	57	187	9.3	19.9	18.9	3.8	3.6
10	14		30	120	43	230	4.I	19.2	18.2	3.6	3.4
	15		30	150	41	271	9.0	20.6	19.4	3.7	3.5
	10	+10XX					8.9	13.9	1 . 1	4.7	4.4

### Ground Bran Sifted on 9XX Sieve 500 g. sample

FB-1	9XX	30	30	71	71	9.8	15	i <sup>7</sup> .4	15.5	5.2
FB -2		30	60	36	107	4.7	1.24 1	1 . 4	5.6	5 3
FB - 3		30	40	27	134	9.7	1.5	1.4	5.7	5.4
FB -4		30	120	27	161	4.8	18 1		5.1	5.8
FB -5		30	150	19	180	9. H	10.1	12.	6.5	6.2
FB -6	+9XX					1, 8		12		6.2

### Ground Bran Sifted on JXA Steve 500 g. sample

	FB-11	10XX	30	30	-41	41	10.1	20.4	19.5	6.1	5.8
	FB -12		30	60	37	78	9.9	20.7	19.8	5.6	5.3
	FB -13		30	90	38	116	9.8	19.8	18.9	5.3	5.1
2	FB -14		30	120	31	147	9.8	18.7	17.8	5.6	
	FB -15		30	150	31	178	9.8	18.2	17.4	6.3	5.3
	FB -16	+10XX					9.3	12.9	12.2	6.4	6.0 6.1

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Lab No.		Identification	ht f	Motor TH	Protein	Ash	·. Fat
					′ <b>.</b>	۶.	<b>'•</b>
G 1168	White	Hulls	29-Jan-66	5.5	12.8	2.2	6.0
G 1169	Corn	Germ ,	••	5.2	16.	4.7	14.7
G 1170	High Lysine	+ 14W C. Grits	**	6.4	4.4	. 85	1.7
G 1171	'67	+ 20W M. Grits	••	. 4	8 1	.62	1.0
G 1172		+ 50gg F. Grits	12	12.1	6. N	. 3.4	0.3
G 1173		+ 50gg Flour	T.	10.2	5.8	. 26	0.5
G i174	Yellow	Hulls	п	5.0	12.9	1.8	6.2
G 1175	Corn	Germ	**	14.4	14.4	4.3	18.4
G 1176	High Lysine	+ 14W C. Grits	••	8.4	9.2	. 84	2.2
G 1177	'67	+ 20W M. Grits	••	1.8	8.6	.66	1.5
G 1178		+ 50gg F. Grits	۰,	8.0	7.5	. 59	0.9
G 1174		+ 55gg Flour		8.4	6.5	. 35	0.9

## High 'vs me cont. Sel mg

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#### PHYSICAL PROPERTIES

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Physical property evaluations are underway to determine the effect o % supplements to the properties of wheat flour.

The list of supplements and their suggested addition level is as follows:

Supplement	Levels of additive
Peanut flour	0, 1, 2, 5, 8 and 50%
Fish protein concentrate (100, 200 and 300 mesh)	0, 1.5, 3, 5, 8 and 10%
L. Lysine - HCl	0, 0.1, 0.2, 0.3, and 0.4%
Soybean products	
Soy Isolate (fine, medium, coarse granulation)	0, 1, 3, 6, 9, 12 and 15%
Soy flour concentrate	0, 1, 3, 6, 8 and 50%
High lysine corn meal	0, 1, 3, 6, 9 and 50%
Corn-Soy-Milk (CSM)	0, 1, 6, 9 and 50%
Sorghum protein concentrate	0, 1, 6, 9 and 50%
Wheat protein concentrate	0, 1, 3, 6, 9 and 30%
Glandless Cottonseed flour	0, 1, 3, 6, 9 and 15%
Horsebean flour	0, 1, 3, 6, 9, 12 and 50%
Chickpea flour	0, 1, 3, 6, 9, 12 and 50%
Yeast concentrate	0, 1, 3, 4 and 6%

The general dough rheology is based upon the use of the farinograph, extensograph, amylograph and gassing power tests (See Table on page 64.

PHYSICAL PRO			ED FLOURS WITH	فالمغربية بالانتجاب والجروب والمتبع والمتبع والمتعادي والتقا	JRS
	% FLOUR	%	<b>%</b>	% FLOUR ABSORP-	mm GASSING
No.	PROTEIN*	ASH	MOISTURE	TION	POWER
Peanut flour	57.7	4.5	8.2		46.6
Low protein white flour	10.4	.4	12.9	60.3	20.2
-10 peanut flour	14.6	.8	12.4	64.3	36.5
+20 peanut flour	18.8	1.2	11.9	69.0	33.4
+30 peanut flour	21.8	1.6	11.5	73.5	38,6
+40 peanut flour	27.1	2.1	11,1	79.9	41.8
+50 peanut flour	31.2	2.5	10.8	83,6	41.9
Medium protein white flour	12.5	4.3	12.8	61.0	36.1
+10 peanut flour	16.2	.83	12.1	65.4	41.3
+20 peanut flour	20.9	1.2	11.9	71.8	38.9
±30 peanut flour	24.6	1.7	11.3	76.2	41.2
+40 peanut flour	28.7	2.1	10.9	80.5	43,5
+50 peanut flour	34.4	2.7	10.0	85.5	45.0
high protein white flour	13.0	.44	12.1	63.5	34.4
+10 peanut flour	17.6	.90	14.4	70.3	36.5
+20 peanut flour	20.6	2.0	11.5	72.3	42.4
+°C peanut flour	25.0	1.6	11.1	77.0	38.6
+4C peanut flour	29.0	2.0	10.7	81.5	52.2
+50 peanut flour	32.8	2.4	10.2	85.0	40.0

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\* Protein Factor 5.75

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The peanut flour was blended with a low protein, a medium protein and a high protein wheat flour at the 5, 10, 15, 20, 30, 40, and 50 percent levels. The general increase in protein content resulting at each increment of blending is shown in the table on page 64. Results indicate that flour absorption increased as the increments of blending increased.

Farinograph data indicated a strengthening in the lower protein flour up to the 30% level, but above this level it became difficult to distinguish between the low, medium, or high protein flours with respect to the farinograph curve. The farinograph picture did indicate that there had been heat damage to the protein of the peanut flour. The extensograph data paralleled that of the farinograph.

The peanut flour blends were baked according to the standard pup loaf method. Results indicate that blends beyond the 30% level of peanut flour destroys all loaf volume. The loaf volume tends to decrease at the 30% level. The flavor beyond that point is not to be recommended.

The upper levels of the blends were tested by baking into nan, the Pakastani bread. The resultant product proved to be heavy but had good keeping quality. The flavor improved with age.

The physical dough testing with the soy isolate is divided into coarse, medium and fine granulation material. Thus for results indicate that the medium granulation is better with respect to flavor, loaf volume, grain and texture of finished bread but these studies are incomplete.

Preliminary physical property studies have been made upon the chick pea and horsebean flours produced. With the exception of the water absorption characteristics, the preliminary physical dough tests do not show the same trend as soy and peanut flour blends perhaps because these beans were not heat treated.

## IMPROVING NUTRITIONAL VALUE OF CEREAL BASED FOODS (NUTRITIONAL EVALUATION)

The enclosed attachments cover literature reviews which have been conducted on various aspects on legumes which may be important in nutritional evaluation of these products if they are to be additives in human diets.

Of specific interest has been information on toxic factors and/or inhibitors found in the chick peas and the bread beam.

Summaries covering some of the more pertinent references in these areas is given in the first attachments. Secondly, progress that has been made in analysis of samples collected in Morroco and Pakistan are summarized in the attached tables with data giving values for amino zoid composition and protein values of the samples on which analysis has been completed. Also in progress is the evaluation of the mineral content of these samples using activation analysis.

Data available on some of the samples have been examined and it would appear that some of the data look reasonable, but that other samples need to be checked further to determine the significance of the values which have been obtained. Data on some samples for nitrogen level have not agreed well with the more common Kjeldahl analysis. These values are being re-checked and the work on the analysis is being evaluated to determine the accuracy of the information

Information covering three separate growth studies using rats are included. These studies cover work which was done to evaluate the effect of lysine and vitamin and mineral additions to atta used in making chapatis. In the second study the nutritional value of protein in fractions from sorghum grain is evaluated. These studies were designed to evaluate the protein quality of endosperm. The studies evaluated protein from fractions of floury endosperm and horny endosperm. In the third study evaluations were designed to determine the effect of minimum additions of vitamin and minerals which might result in the most improvement in growth rates. Complete vitamin and mineral additions to diets containing wheat flour or the cereal flours will result in good growth. In this study only those minerals calculated or most deficient were added in test diets.

If such additions are not made poor growth can be expected since high extration wheat flours and other cereal products contain certain amounts of vitamin and minerals but not adequate levels of all nutrients. Enrichment which supplied the lacking nutrients would aid in avoiding nutritional imbalance which might occur on the addition of single nutrients. Preliminary data obtained in the third rat study indicate the need for further evaluations related to the effects of only limited supplements of certair vitamins and minerals.

#### TOXIC FACTORS IN CHICK PEAS

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The presence of toxic factors in legumes have been recognized by man since the dawn of history. Among the legume seeds of nutritional importance investigated, the trypsin inhibitor was found in the chick peas and other beams. The trypsin inhibitor was not found in non legumes.

Brochers and Ackerson (1950) have presented a table containing seeds of legumenosae and other seeds. Chick peas contain 32.3 units of inhibitor/ml. extract of sample. (Table attached).

They also studied the effect of autoclaving on the nutritive value of pulses. There was no correlation between the improvement in nutritive value after autoclaving and the presence or absence of the trypsin inhibitor.

Hirwo and Magar (1952) showed that trypsin inhibitor content of pulses was less them that of beans. They found that methionine supplementation to rew pulses increased nutritive values.

Chattepadiya and Banerjee (1953) studies the biological value of proteins of five variaties of pulses (including chick peas) by the rat growth attail, both before and after 48 hours of germination. They reported that, trypists - inhibitor activity did not change with germination indicating that the alternal biological value of the protein is not due to either increases or decreases in their trypsin inhibitor activity.

Sahanie and Shandarkar (1954) found that the trypsin inhibitor from chick peas are destroyed by heating at 100% for one hour. Whereas the fibilities from double beans are stable under these conditions. Autoclaving destroys most inhibitors with the exception of double bean inhibitor.

Stachuba in (1931) worked on other legumes. Lathyrism is the only Glabalin lower to arise from their use. The poisonous principle is an acid, 61

white and hygroscopic, extracted by macerating the ground peas in cold water or alcohol for several days, percipitating albumen, filtering and concentrating the liquid. Experiments were done with the active principle and the extract on monkeys, rabbits and frogs. It caused lesions in the brain and spinal cord and paralysis of motor nerves.

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According to Altschul (1958), diseases caused by toxic factors are referred to as lathyrism, cicerism or odoratism. A species of cicer have been presumably pointed out as causative agent. He mentioned that the common reasons for lathyrism may be due to:

- a Abnormally high levels of selenium and manganese in certain legume seeds,
- b' Or a particular amino acid deficiency.

We has also presented a table on the "effect on amino acid composition of wheat and corn flours by addition of chick pea products and soybean meal." The table is attached with the report.

Altschul pointed out the common practice in preparing dishes by soaking first and throwing away the steeping water may help in diminishing the risks derived from the ingestion material containing the toxic factor. (He means the ingestion of toxic factor).

Abramova and Chernikov (1964) investigated the proteinase inhibitor in chick peas along with other legumes. The fat free flour was extracted, precipitated, filtered and dialyzed until salt free. The inhibitory effect of this material was investigated. The highest antitryptic and antichymotryptic activity was found in <u>P. Vulgaris</u> seeds followed in <u>G. Max. C. Arictinum</u> (chick peas), <u>L. Esculenta</u> and <u>P. Sativum</u>. None of these seeds exhibited antipaptic activity. 292

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## RAYMOND BORGERRS AND C. W. ACEBRSON

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### VAPLE I

Description Description of the Legensinnoos and Others	10 ju
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Inhibitor U. da/mi facutynicus to g	
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Canapalia maifanuia jack boan 16 0	2
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Corea serentencia method to	
Chamadorida faminulata pantai i	mu
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Laganus angustifaling-blue lupine 6.9	111
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of animal officering in the second in the second se	
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	61
Phononius accurate acarlist runner beng 21 a	<b>I</b> - ·
	1 2.1
Phenolus volgaris-garden bean	
Pieren entieren -garden pen	A
Pissen sabirum var. aromes field pea	ų.
Gaja mas-soybush 0	
Sephere jaronica-Japanese pageda tros 35.0	R
Trifelium protence-manmoth red clover	9
Vicia fabe-horse bean 0	
Vicia satise common vetch	
Vigna sinensis-black synd pan	
Non-lagumes (43 ?	
Aleverico fordis-tung bean	
Avena satis-onto	
Hordeum sulgare-barley	
Linum unitaliprimum fan	
Sends coverie-rye	
Sorghum sulpars-Leoti	
Triticum velgare-wheat 0	
Zas mayo meleniaio-com 0	
Nerrenclature accust	
"Nocaemclature sorrding to GRAHAM, E. H., Legumes for Fromen Centrel a Wildhife, U.S.Dopl. Agr. Misc. Publ. No. 453 (1964)	
Wildhife, U.S.Dopi. Agr. Musc. Publ. No. 453 (1941).	

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TABLE IV ESTRET ON AMERIC ACES COL. POSTION OF WEILAT AND COMP PLOUDS ST ADDITION OF CHICK PER PROPORT AND SOTEMAN MEAL . 50

	Proteint	Amount containing 100 g-	inter Amounts of series which is see					0 g. of protein		
Source of protoin	( <del>day</del> basis)	protein (dry basis)	Locations	Methia-	Planyt-	Loto		las-	Three-	Trypto-
Floam	%	8.	e.	F	6.	G	6			
Typical wheat flour*				12.00			<b>B</b> -	G-	<b>g</b> .	8-
Wheth care four	19.5	800	70	80		1.9	1000			
Additives	9.6	1042	15.0	31	8 0		4.1	8.8	27	0.8
		2018				8.9		6 4	87	0.6
Soybas mal	65 0	222	8.0							
Whats shigh per form	13.5	4051	0.0	1.7	5.8	0.0	8.8	6 6		14
Undefatted chick pas protain	60 0			120000			÷.			• •
Contract parts president propulate	63.0	167	B.Q	1.6	6.8	6.6	0.8	8.9		1.0
Exclude Chant from markening		111)							••	
10% Baybana mani										
10% Chick pas faur	10 0	805	7.8	1.9		4.5				12012
40% China pro farm	23 6	755	78	1.9	8.7	8.7	4.7	6 1	8 8	1.1
100 Testa in the second	6 64	600	8.4	17	6 2		4 8	48	S.1	0 8
10% Undefatted chick pas protes	17 8	670		- 1 8		6.9	8.8	5 8	41	0 9
10% Chick pes protein provier	<b>60 1</b>	655	7.8			8.5	4.8	48	3 5	0.9
Lanchod cers four products				18	5 9	3 9	80	4.0	5 7	0.0
20% Soyhana meal	16 7									
10% Chick and flow		600	11 6	8.8	6 2	67	8 8		58	10
80% Chieft man flower	11 e	600	10 7	8.8	5 3	3 2	1.5	6 5	3 9	
10% Dedelatiod chieft pes protesa	10 6	600	10 6			6.5	10	6 0		07
10% Chieft pas protein perder	14 6	334	18 5	2 4	56		. 7		4 2	0 9
amatial amine and	17 6	1933	11 9	8.8		4.4			4 2	08
annalisi annino acid regenerate per de	7 (adats)							6 2	\$ 3	08
Nitrogan X 6.65						16	10	1.6	10	0 5

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R. Block and D. Belling, "The Ammo Acad Compatition of Proteins and Fonds," and ed. Charles C. Thomas, Springfield, Illinois, 1951.
 E. Louis C., of the Institute Maximum de Investigances Temeligens, espublished work performed at the University of Wisconsis, 1952.
 Based on the recommended ash daily intake which is twice the minimal amount listed in Table III, Chapter 2, and in Table VIII, Chep-ter 18.

#### PROTEIN REQUIREMENTS

Mason et al. (1964) found that the American and European women had significantly higher basal metabolic rates than the Indian womer. The differences could not be attributed to differences in age, neight, weight, or muscle mass. They suggested that the differences may be due to several generations of Indians adapting to the tropical climate. Since the metabolic rates of the Indian women are lower, their protein needs may be lower.

The amino acid requirements for adult humans and for rats have been reported by Rose and others and is given in Processed Plant Protein Foodstuffs, p.24, (Altschul, 1958):

amino ecid	adult human g/day	rat i F die
arginine	0.00	)
histidine	0.00	0.4
isoleucine	0 70	0 5
leucine	1.10	n s
lysine	0.80	10
methionine	1 10	0.5
phenylalanine	1 10	1 1
chreonine	0.50	• 1
tryptophan	0.25	•
valine	0.80	. 7

Hegsted (1957) estimated that children of the giver ages require the amounts of protein listed below:

1 month -- 2.5 g/kg body weight 18 months -- 1.0 g/kg body weight 2.5 vears -- 0.76-0.83 g/kg body weight 6.5 years -- 0.62-0.67 g/kg body weight

Howe <u>et al</u>. (1965) reported the following recommendations: infants (at tirth) -- 2.2 g/kg body weight

imanus lat cirtr	1) C.C g/kg body weight
2.months	2.0 g/kg body weight
l year	1.5 g/kg body weight
1-5 years	1.0-1.5 g/kg body weight

Plenert et al. (1965) found that increasing the protein intake of infants above 3 g/kg did not improve weight gain or utilization of food and energy.

Nitrogen balance studies (Nakagawa et al., 1952 have shown that 10-12 year old boys require approximately 1.9 g threenine (35 mg/kg), 0.9 g valine (33 mg kg), and 3.7 g : e glalanine (27 mg/kg) daily.

Fomon and Filer (1966) reported that the sole ina for infants and value requirements, were slightly lower than previous reports had indicated. Twenty infants were fet a formula which contained protein (6% of formula in the form of methionine-supplemented soy isolate. They reported that 50-85 mg isolaucine/kg/day and 53-70 mg valine/kg/day were sufficient for most children. These values they compared with 79-126 mg isolaucine/kg/day and 85-105 mg valine/kg/day reported by Snyderman <u>et al</u>. (1959, 1964).

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# PROTEIN AVAILABILITY

Autoclaving (121°C, 30 and 60 minutes) reduced the lysine content of chickpeas by 10% (González del Cueto <u>et al.</u>, 1960). In contrast to reports that autoclaving reduced the arginine content of cottonseed meal, no measurable reduction in arginine concentration was observed in chickpeas (Gonzáles del Cueto <u>et al.</u>, 1953). Kuiken and Lyman (1948) reported the true availability (ar.no acids fed minus amino acids in faces of rates of Wirest as follows:

arginine	96.4%
istidine	98.8%
soleucine	95.0%
eucine	95.4%
vsine	92.8%
methionine	94.9%
phenylalanine	96.9%
threonine	92.2%
trvptophane	93.2%
valine	93.2%
total N	95 <b>.0%</b>

Supta <u>et ai</u>. (1958) reported that on the basis of rat feeding experiments, lysine is 81-865 available in rice and 61+79% available in wheat.

The true digestibility of properly cooked legumes nas been estimated to be between 85 and 95%, with beans being digested slightly poorer than peas (Avdrovd and Doughty, 1964). The protein content of some Indian pulses has been observed to increase during germination (Chattopadhyay and Banerjee, 1953). Genetic strains of chickpeas vary as much as 38% in protein content (Esh et al., 1960).

# SUPPLEMENTATION OF DIETS

Howe et al. (1967) stated that fish protein concentrate and sovbean meal are the only potentially available protein concentrates containing a sufficiently high lysine content to be useful as a supplement. In general they advocated the addition of pure amino acids to the staple foods. On the other hand, Hegsted (1968) stated that purified amino arids are too expensive at the present to be useful as supplements for developing areas.

Growth of children 2-5 years of are has been improved chighly significantly) by supplementing their usual rice and pulse diets with one ounce of a high protein food inity commet al., 1966). The food contained role ar coinct protein isolate, 15% roasted chickpea flour, vitamins and minerals and supplied 22.2 g protein per cunce.

Guttikar <u>et al</u>. (1965) evaluated a protein food based on groundnut, chickpea, and sesame flour (4:4:2)as being quite effective as a source of protein when fed at the rate of 1.5 g protein per kg body weight.

The effect of supplementing the rice diet of Indian girls (8-9 years) with lysine, methionine, and threonine has been studied (Parthasarathy <u>et al.</u>, 1964). The rice diet contained 250 g rice, small amounts of legumes, vegetables, 45 g oil, and 5 g skim milk powder, vitamins, sugar, and a salt mixture. It furnished 1500 10

.

Barness <u>et al.</u> (1961) observed that wheat 'cream of wheat cereal) fortified with lvsime and potassium appeared to supply adequate protein for growing Latin American infants 3-17 months of age. The daily caloric intake was 75-125 kcal/kg; protein intake was 1.2-4.0 g/kg.

Chickpeas and broad beans have been used in foods in the Near East and other parts of the world to supplement the liet of small children (Autret and Van Veen, 1955; Astour et al., 1965). In India legume flour is sometimes added to wheat flour (up to 10%) to make chapatties (Avarant and Doughty, 1964).

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		81-68		324 - 68	321-68	7 <b>9-68</b>
	78-68	Moroccan	346-68	Lentile	111	Horse
	Lentils	Lentils	Lentiis	No. 53	<b>Peies</b>	Beans
Protein	24.1	23.0	17.0	25.8	24.1	26.4
Noisture	10.6	10.7	11.7	10.7	1	11.0
Lysine	2.019	1.776	1.246	1.812	1.732	1.676
Ristidine	0.661	0.613	0.651	6 648	0.650	0.613
Ammonia	0.428	0.417	0.317	0.468	0.488	0.433
Arginine	2.045	1.846	1.537	2.192	2.640	2.440
Apartic acid	2.904	2.551	2.291	3.242	2.887	2.901
Threenine	0.920	0.851	0.740	0.974	0.929	0.949
Serine	1.206	1.148	0.970	1.302	1.201	1.243
Glutamic acid	4.063	3.924	3.487	4.550	4.492	4.607
Proline	1.416	0.827	1.337	1.089	1.016	1.014
Glycine	0.964	0.902	0.775	1.053	1.067	1.046
Alanine	1.018	0.924	0.796	1.105	1.052	0.062
Half Cystine	0.026	0.000	0.0	0.0	0.365	0.0
Valine	1.266	1.114	1.016	1.182	0.690	1.232
Methionine	0.096	0.500	0.274	0.099	0.122	0.113
Isoleuc ine	1.047	0.922	0.780	0.904	1.074	1.060
Leucine	1.612	1.405	1.364	1.844	1.662	1.687
Tyreeine	0.797	0.644	0.574	0.786	0.832	0.798
Phenylalanine	1.171	1.075	1.040	0.285	1.090	1.039
Mitrogen Recovery Ouidations:	94.78	89.42	106.60	91.83	98.00	88.41
Cystine				0.150	0.272	
Methionine				0.093	0.148	

7 SAMPLE

81

	83-68 Good Horse Beans	331-68 Werthorn Beans	335-66 Lg. Kidney Baana	344-68 Peveroles	347-68 Norse Beans	182-68 Rod Kidney Boans	73-68 Nericot Beens
Protein	26.3	23.3	22 🕈	25.8	د 27	21.4	20.8
<b>Woisture</b>	10.1	13.4	11.5	10.3	9.8	10.3	10.6
Lysine	1.308	1.705	1.559	1.555	1.613	1.575	1.419
Histidine	0.501	0.723	0.677	0.613	0.612	0.653	0.507
Ammonia	0.360	0.454	0.432	0.452	0.474	0.414	0.322
Arginine	2.206	1.673	1.491	2.674	2.354	1.592	1.119
Aspartic acid	2.319	2.823	2.898	2.789	2.760	2.852	2.555
Threenine	0.779	1.032	0.991	0.848	0.952	0.932	0.924
Serine	0.983	1.452	1.377	1.207	1.175	1.374	1.236
Slutamic acid	3.649	3.694	4.075	3.965	4.452	4.059	3.465
Proline	0.789	1.215	0.850	1.007	0.874	0.914	0.894
ylcine	0.882	0.950	0.896	1.022	1.005	0.917	0.819
lenine	0.887	1.016	0.933	0.977	1.018	0.990	0.876
lelf Cystine	0.000	0.255	0.0	0.349	0.0	0.210	0.211
Valine	0.993	1.192	1.316	1.119	1.197	0.971	1.001
<b>i</b> ethionine	0.062	0.210	0.213	0.115	0.063	0.210	0.174
seleucine	2.006	1.077	1.064	0.964	1.040	1.075	
eucine	1.332	1.869	1.659	1.745	1.585	1.884	0.943
lyrosine	0.633	0.757	0.742	0.784	0.799	0.797	1.634
Phonylalanine	0.791	1.295	1.309	1.035	1.047	1.325	0.709 2.305
litrogen Recovery Dridations:	76.08	94.99	92.03	89.52	83.26	98 <b>.8</b> 4	87.50
Cystine		0.243	0.213			0.194	
Methionine		0.265	0.255			0.170	

7 SAPLE

			% SAMPLE				
	75-68 Noroccan Beans	85-68 Poor Horse Beans	322-68 319 Feveroles	348-68 G <b>rou</b> nd Short <b>s</b>	124A-68 White Wheat	128 <b>A-68</b> Stone Ground White Wheat	127-68 White Wheat
Protein	22.7	22.0	24.9	21.1	10.7	11.6	12.2
Moisture	10.6	10.7	10 9	13.1	13.2	11.7	11.9
Lysine	1.628	1.441	1.580	0.586	0.324	0.333	0.322
Histidine	0.721	0.527	0.631	U.529	0.244	0.260	0.270
Amonia	0.517	0.448	0.459	0.864	0.340	0.318	0.385
Arginiae	1.619	1.979	2.402	1.126	0.509	0.573	0.575
Aspartic acid	2.783	2.355	2.883	1.225	0.598	0.661	0.601
Threenine	0.915	0.815	0.886	0.657	0.342	0.377	0.366
Serine	1.183	1.031	1.172	1.107	0.535	0.557	0.564
Slutamic acid	4.375	3.758	4.314	8,613	3.517	3.885	3.814
roline	1.016	0.891	1.013	2.533	1.024	1.276	1.169
Glycine	0.984	0.922	1.044	0.928	0.456	0.474	0.471
Manine	1.003	0.901	1.008	0.777	0.427	0.438	0.471
alf Cystine	0.0	0.268	0.257	0.702	0.353	0.310	0.438
<b>/aline</b>	1.096	0.965	1.143	1.006	0.505	0.606	0.471
Methionine	0.331	0.093	0.075	0.159	0.123	0.181	0.478
lsoleucing	0.941	0.882	1.018	0.800	0.394	0.445	0.128
Joucine	1.704	1.432	1.628	1.390	0.764	0.720	0.423
<b>Nyrosine</b>	0.730	0.697	0.792	0.708	0.362	0.344	0.368
Phenylalanine	1.306	0.928	1.044	1.101	0.522	0.512	0.368
Ritrogen Recovery Dxidations:	95.48	90.99	91.97	108.51	96.11	94.41	91.37
Systine			U. 294		0.255	0.276	0.331
Mathionine			U. ito		0.160	0.177	0.190

	307-68 NA-3 277	313-68 BT 3597	311-68 BT 3597	58-68 Coarse Bran	91-68 #1 Burum	312-68 BT 908	308-68 3225
Protein	11.8	11.5	11.1	14.1	11.6	10.5	10.3
Noisture	9.2	11.6	10.7	11.4	11.8	11.3	10.9
Lysine	~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~ ~	0.339	0.317	G.596	0.316	0.304	0.294
Lysine	0.315	0.332	0.239	0.392	0.238	0.232	0.231
	0.260	0.254		0.322	0.394	0.339	0.308
Amonia	0.307	0.353	0.366	0.922	0.540	0.499	0.523
Arginine	0.546	0.549	0.539	1.130	0.545	0.624	0.589
Aspartic acid	0.671	0.626	0.652			0.328	0.301
Threesins	0.335	0.344	0.337	0.494	0.329	0.523	0.492
Sering	0.584	0.527	0.538	0.672	0.580		3.143
Qlutasie acid	3.929	3 " 698	3.530	2,941	3.891	3.457	
Prolize	1.252	1.218	1.210	0.930	1.266	1.186	1.114
Gircina	0.431	0.436	0.455	0.817	0.421	0.431	0.404
Alenino	0.402	0.405	0.414	9.731	0.402	0.393	0.374
Half Cystine	0.232	0.372	0.383	0.392	0.308	0.353	0.265
Valine	0.515	0.301	0.496	0.697	0.474	0.476	0.463
Methionine	0.055	0.151	0.069	0.151	0.035	0.052	0.086
Isoleucine	0.426	8.386	0.330	0.399	0.374	0.352	0.332
Leucine	0.701	0.729	0.736	0.868	0.732	0.686	0.660
	0.339	0.316	0.326	0.399	0.321	0.323	0.297
Tyrocine	0.557	0.517	0.479	9.570	0.530	0.472	0.469
Phenylalanias	0.331	U. 347	0.477	2.274			
	89.54	92.61	<b>96.</b> 47	91.78	93 <b>.90</b>	95.71	92.24
Residetions:				0.317		0.930	0.247
'Cystine	0.245	0.267	0.276	0.315		0.230	
Machicaiae	0.120	0.158	0.164	0.194		0.1 <b>29</b>	0.034

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	310-68 BT 2306	93-68 Durum	57-68 Fine Bran	50-68 fr Rec winter	3.7 <del>9</del> - 1 - 34 24	908 42 Durum	ob-68 Strat, Curade Flour force
Protein	10.9	13.0	14.9		:: I	11.6	9.0
Moisture	11.7	11.7	11.1		1 1 N	4	12.4
Lysine	0.280	0.324	0.614			0.342	0.20
Histidine	0.227	0.266	0.371	158	. 251	0.305	6.200
Ammonia	0.300	0.371	0.333	. 174	1.326	0.372	0.3/1
Arginine	0.486	0.553	0.957	0.580	0.534	0.652	0.392
Asy attic acid	0.525	0.631	0.103	0.586	0.566	0.757	0.392
Threonine	0.324	0.385	0.521	0.343	0.338	0.398	0.272
Serine	0.503	0.605	0.665	0.525	0.538	0.699	0.461
Glutenic acid	3.364	4.070	3.068	3.584	3.449	4.780	3.668
Proline	1.005	1.255	1.092	1.032	1.023	1.557	1.042
- Glycine	0.425	0.527	0.773	0.461	0.445	0.499	0.337
Alaning	0.375	0.444	0.728	0.413	0.416	0.470	0.291
Half Cystine	0.430	0.436	0.405	0.335	0.354	0.391	0.366
Valine	0.455	0.549	0.800	0.581	0.487	0.621	0.440
Methionine	0.041	0.145	0.200	0.131	0.083	0.129	0.104
Isoleucine	0.368	0.444	0.457	0.412	0.399	0.419	0.341
Leucine	0.703	0.846	0.883	0.699	0.761	0.865	0.611
Tyrosine	0.337	0.406	0.428	0.352	0.351	0.412	0.316
Fhenylalanine	0.488	0.582	0.591	0.531	0.520	0.696	0.477
Nitrogen Bacovery	87.45	88.50	88.65	91.30	91.20	93.37	97.67
Cystine	0 250				0.240		0.215
Hethionine	6.151				0.192		0.112

7. SAMPLE

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	84-68 Helba-Ext.	71-68 Bird Seed	151 <b>-68</b> Lobin	325-68 2 <b>04 Soja</b>	70-68 Broken Split Peas	332-68 Gorden Peas	333-68 Small Br. Crowder
Protein	25.5	16.3	23.7	38.4	22.4	. 27.3	20.4
Noisture	9.70	10.0	10.0	8.1	10.8	12.4	12.7
Lysine	1.637	0.357	1.746	2.800	1.753	2.350	1.495
Histidina	0.617	0.329	G.780	1.094	0.564	0.695	0.653
Amonie	0.401	0.395	0.415	U.799	0.357	0.577	0.428
Arginine	2.344	0.947	1.725	3.275	2.433	2.855	1.482
Aspartic acid	2.864	0.904	3.065	4.996	2.897	3 124	2.248
Threenine	0.950	0.403	0.959	1.654	0.877	1.128	0.858
Serine	1.248	0.680	1.286	2.150	1.118	1.393	1.121
Glutamic acid	4.266	4.842	4.725	8.301	4 660	5.126	3.821
Proline	1.030	1.198	1.388	1.999	4.203 3 1.177	1.146	0.983
Glycine	1.183	0.479	1.015	1.723	0.986	1.305	0.947
Alamine	0.997	0.634	1.046	0.757	1.006	1.314	0.994
Balf Cystine	0.143	0.491	0.0	1.101	0.153	0. <b>27</b> 1	0.107
Valine	1.167	0.791	1.300	1.925	1.121	0.883	1.058
Methionine	0.159	0.208	0.277	0.467	0.311	0.0	0.267
Isoleucine	1.245	0.669	1.061	1.942	0.974	1.159	0.903
Leucine	1.498	0.159	1.727	2.822	1.817	1.981	1.621
Tyros inc	0.783	0.421	9.7 <b>99</b>	1.519	0.763	0.898	0.728
Phenylalanine	1.042	0.902	1.418	2.124	1.069	1.272	1.175
Nitrogen Recovery Oridations:	89.62	86.62	97.08	102.93	101.2	100.51	97.36
Cystine			0.218			0.349	0.207
Methionine			0.189			0.285	0.281

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			7. SAMPL	Ľ			
	.336-68 Crowdar Cow Pee	328-68 191 Pois	36-68 Wheat é Barley	144- <b>68</b> Barley	94-68 Barley	338- <b>68</b> Hgtreated Sorghum	126A-68 Sorghum
otein	23.3	23.7	12.1	10.8	9.4	10.2	· 9.5
lsture	14.1	9.9	11.6	11.0	11.3	12.6	12.3
ine	1.799	1.849	0.302	0.354	0.322	0.220	0.182
stidine	0.836	0.555	0.231	0.219	0.185	0.225	0.193
onia	0.536	0.365	0.232	0.304	0.226	0.283	0.261
inine	2.004	1.940	0.484	0.516	0.441	0.363	0.333
rtic acid	3.061	2.697	0.578	0.696	0.540	0.818	0.664
mine	1.033	0.984	0.348	0.383	0.324	0.365	0.314
28	1.377	1.111	0.515	0.493	0.398	0.481	0.314
mmic acid	4.937	4.013	3.501	3.043	2,293	2.441	2.072
ine	1.283	0.876	1.227	1.215	0.968	0.736	0.743
ine	1.092	1.067	0.411	0.455	0.363	0.283	0.314
ine	1.117	.084	0.411	0.455	0.368	0.925	0.894
E Cystine	0.0	0.402	0.307	0.326	0.242	0.045	0.190
.DC	1.394	1.120	0.492	0.537	0.457	0.606	0.481
hionine	0.363	0.183	9.136	0.036	0.068	0.199	0.043
loucine	1.060	1.029	0.394	0.402	0.321	0.532	0.384
cine	1.978	1.519	0.750	0.762	0.638	1.410	1.254
sine	0.838	0.824	0.355	0.369	0.292	0.341	0.384
ylalanine	1.489	1.162	0.395	0.570	0.482	0.388	0.511
ogen Recovery lations:	107.51	90.78	83.95	93.10	85.75	93.50	90.87
tine	0.235		0.146	0.219			0 141
hionine	0.332		0.182	0.139			0.141 0.120

	3 <b>18-58</b> SH2O-Hybrid Hewcain	319-68 Sorghum Vulgare	77 <b>-68</b> White Corn from Fer	334-68 American Corn	129-68 Domestic Corn	315-68 D117 Hybrid Double	74-68 American Corn
Protein	16.3	10.5	10.4	11.4	9.3	10.1	16.1
Noisture	11.6	12.1	11.3	13.6	1.5	11:4	11.0
Lysine	0.280	0.238	0.231	0.259	0.283	0. <b>26</b> 7	0.298
Histidine	0.330	0.210	0.227	0.343	0.269	0.280	0.294
Amonia	0.475	0.327 •	0.255	0.307	0.186	0.233	0.234
Arginine	0.568	0.361	0.433	0.428	0.434	0.443	0.442
Aspertic acid	1.115	0.862	0.761	0.775	0.632	0.764	0.832
Threenine	ü.507	0.341	0.365	0.424	0,351	0.367	0.444
Serine	0.705	0.530	0.472	0.592	0.454	0.528	0.594
<b>Clytanic</b> acid	3.921	2.495	2.393	2.712	1.901	2.213	2.569
Proline	1.263	1.169	1.193	1.277	0.839	1.081	1.156
Glycine	0.441	0.329	0.341	0.384	0.344	0.362	0.401
Alenine	1.575	1.028	0.994	0.957	0.685	0.816	0.942
Balf Cystine	0.301	0.0	0.110	0.351	0.171	0.213	0.126
Valine	0.818	0.517	0.588	0.577	0.468	0.461	0.603
Nethionine	0.200	0.021	0.239	0.204	0.019	0.040	0.153
Isoleucine	0.664	0.414	0.456	0.443	0.334	0.324	0.439
Laucina	2.089	1.446	1.387	1.630	1.017	1.399	1.427
Tyrosine	0.696	0.399	0.468	0.521	0.371	0.412	0.485
Phenylelanine	0.550	0.558	0.568	0.641	0.427	0.537	0.563
Nitrogen Recovery Oxidations:	92.07	97.17	96.98	98.84	88.66	94.39	65.08
Cystine	0.269	0.158		0.251	0.208	0.210	
Methionine	0.267	0.099		0.227	0.171	0.176	

		345-66				341-68	
	148-68	Mang	1314-66	337-68	72 <b>A-68</b>	Cenary	340-68
	Mang	Besa	Ming	White Mellet	Millet	Seed	Mustard
Protein	23.4	21.9	22.9	10.6	10.1	16.6	26.0
Neisture	11.7	11.9	11.1	12.2	10.9	12.7	7.10
Lysine	1.696	1.653	1.757	0.189	0.186	0.535	1.210
Histidine	0.651	0.612	0.660	0.217	0.204	0.318	<b>6.576</b>
Amercia	0.419	0.376	0.266	0.319	0.242	0.479	0.532
Arginine	1.629	1.550	1.656	0.357	0.402	1.134	1.444
Apartic acid	3.026	2.731	3.060	0.692	0.677	1.357	1.477
Throwshe	0.851	0.804	0.833	0.340	0.326	0.486	0.891
Serine	1.333	1.238	. 1.281	0.733	0.615	0.700	0.936
<b>Electonic</b> acid	4.665	4.196	<b>4.598</b>	2.359	2.279	3.840	4.109
Proling	1.136	1.037	1.297	0.679	0.774	0.841	1.260
Glyuine	0.966	0.901	0.913	0.252	0.262	0.625	1.073
Alenine	1.130	0.993	1.001	1.173	1.003	0.740	0.910
Half Cystine	0.107	0.200	0.0	0.182	0.085	0.457	0.666
Valine	1.321	0.405	1.339	0.242	0.532	0.778	0.919
Methicaine	0.266	0.278	0.250	0.160	0.256	0.220	0.091
Isolaucine	1.128	1.010	1.058	0.422	0.602	0.636	·0 <b>.850</b>
Loucine	2.002	1.825	1.736	1.311	1.047	1.089	1.442
Tyresing	0.798	0.764	0.663	0.423	0.345	C. 504	0.584
Phonylelenine	1.515	1.421	1.414	0.613	0.515	0.849	0.872
Nitrogen Recovery Oxidations:	96.76	92.87	96.00	91.10	89.48	89.06	74.10
Cystine	0.186		0.163	0.157		0.355	0.546
Nethioning	0.277		0.290	0.510		0.272	0.354

## I PROTEIN

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		81-68		324-68		79-68	83-68
	78-68	Moroccan	346-68	Lontils	1.1-68	Horse	Good Horse
	Lentils	Lentils	Lentils	Lentils No. 53		Beans	Beans
Protein	24.1	23.0	17.0	. 8	•	26.4	26.3
Moisture	10.6	10.7	11.7	10 '	1.0	11 0	10.1
Lysine	8.377	7.721	7.329	7.02.	/ 185	6.347	4.974
Histidine	2.741	2.665	3.828	2.513	2.698	2.322	1.906
Ammonia	1.778	1.812	· 1.864	1.815	2.024	1.641	1.367
Arginine	8.484	8.026	9.040	8.49	10.953	9.241	8.387
Aspartic acid	12.049	11.090	13.479	12.566	11.980	10.987	8.818
Threonine	3.819	3.701	4.351	3.774	3.854	3.593	2.963
Serine	5.003	4.991	.5.705	5.048	4.984	4.707	3.736
Glutanic acid	16.859	17.062	20.509	17.634	18.638	17.451	13.675
Proline	5.877	3.596	7.862	4.222	4.215	3.839	2.998
Glycine	3.999	3.922	4.560	4.082	4.427	3.961	3.352
Alamine	4.224	4.020	4.693	4.285	4.367	4.022	3.374
Half Cystine	0.106	0.000	0.000	0.000	1.516	0.000	0.000
Valine	5.252	4.864	5.979	4.582	2.863	4.665	3.777
Methionine	0.405	0.777	1.613	0.382	0.508	0.428	0.236
Isoleucine	4.344	4.010	4.587	3.506	4.458	4.016	7.633
Leucine	6.690	6.110	8.025	7.149	6.897	6.390	5.065
Tyrosine	3.305	2.801	3.377	3.048	3.450	3.024	2.407
Phenylalanine	4.859	4.676	6.117	4.982	4.524	3.937	3.086
Oxidation:							8
Oysting				0.581	1.130		
Hathionine				0.360	0.615		

# Z PROTEIN

	331-68 Northern Beans	335-68 Lg. Kidney Beans	344-68 Fever, ies	34'-68 Hurse Beans	15: 65 Red Kidnev Brane	hi-68 Haricot Beans	75-68 Moroccan Beans
Protein	23.3	22.9	25.8				
Moisture	13.6	11.5	10.3	27.3 4.4	10. 1	20.8	22.7 10.6
Lysine	7.317	6.983	6 01 :				10.0
Histidine	3.102	2.958	6.02	2.908	7.362	6.822	7.172
Ammonia	1.950	1.884	2.378	40	3.050	2.724	3.177
Arginine	7.180	6.509	1.751	: 135	1 933	1.548	2.279
Aspartic acid	12.115		10.364	8.623	7.437	5.378	7.134
Threenine	4.430	12.656	10.811	10.111	13.329	12.283	12.258
Serine	6.232	4.327 .	5.205	3.487	4.357	4.441	4.033
Glutanic acid		6.012	4.677	4.303	6.420	5.941	5.211
Proline	15.853	17.795	15.370	16.309	18.969	16.660	19.273
Glycine	5.213	3.710	3.904	3.203	4.271	4.296	4.476
Alanine	4.076	3.911	3.963	3.682	4.287	3.936	4.337
	4.361	4.076	3.785	3.729	4.579	4.211	4.421
Half Cystine Valine	1.094	0.000	1.353	0.000	0.982	1.012	0.000
Mathionine	5.115	5.746 28	4.337	4.385	4.538	4.955	4.830
	0.900	6.929	0.445	0.231	0.982	0.838	1.460
Isoleucine	4.624		3.736	3.808	5.023	4.534	4.147
Leucine	8.020	7.246	6.762	5.807	8.803	7.855	
Tyrosine	3.248	3.238	3.037	2.926	3.723	3.364	7.503
Phenylalanine	5.558	5.716	4.010	3.836	6.240		3.217
Omidation:				5.050	0.240	6.274	5.755
Cystine	1.044	0.929			0.000		54
Methionine	1.136	1.114			0.905 0.793		
					0.795		

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# Z PROTEIN

	85-68 Poor Horse Beans	322-68 319 Feveroles	348-68 Ground Shorts	124-68 White Wheat	128-68 Stone Ground White Wheat	127-68 White Wheat	307-68 MA-3 277
Protein	22.0	24.9	21.1	10	11.6	12.2	
Moisture	10.7	10.9	13.1	13.2	.1	11.9	11.8 9.2
Lysine	6.551	6.345	2.776	3.926	2.872	2.639	2.667
Histidine	2.396	2.532	2.505	2.278	2.243	2.214	2.205
Ammonia	2.035	1.843	4.097	3.174	2 741	3.157	2.602
Arginine	8.994	9.647	5.337	4.761	4.941	4.716	4.631
Aspartic acid	10.706	11.578	5.806	5.589	5.703	4.927	5.682
Threonine	3.704	3.558	3.114	3.192	3.250	2.996	2.839
Serine	<b>4.688</b>	4.705 •	5.248	4.996	4.806	4.623	4.946
Glutamic acid	17.081	17.324	40.821	32.865	33.488	31.262	33.296
Proline	4.048	4.069	12.004	9.568	10.999	9.581	10.614
Glycine	4.189	4.194	4.396	4.260	4.090	3.862	3.653
Alanine	4.095	4.047	3.681	3.991	3.772	3.592	
Half Cystine	1.217	1.030	3.325	3.297	2.675	3.859	3.403
Valine	4.385	4.589	4.770	4.721	5.221	3.915	1.966
Methionine	0.421	0.302	0.755	1.153	1.563	1.056	4.361
Isoleucáne	4.010	4.088	3.792	3.678	3.836	3.465	0.470
Leucine	6.511	6.539	6.586	7.144	6.206	6.624	3.610
Tyrosine	3.167	3.182	3.355	3.388	2.969	3.020	5.943
Phenylalanine	4.220	4.193	5.219	4.881	4.410	.547	2.874 4.724
Oxidation:							
Cystine		1.182		2.382	2.380	2.711	2 075
Methionine		0.661		1.493	1.523	1.638	2.075 1.016

# 2 PROTEIN

	313-68 BT 3597	311-68 BT2511	56-68 Coarse Bran	61-68 #I Durum	312-68 BT-908	308-68 3225	310-68 BT2 306
Protein Moisture	11.5 11.6	11.1 10.7	14.1 11.4	1:6 11.8	10.5 3	10.3 10.9	10.9 11.7
Lysine Histidine Amonia Arginine Aspartic acid Threonine Serine Glutamic acid Proline Glycine Alanine Half Cystine Valine Methionine Isoleucine Leucine Tyrosine Phenylalanine	2.891 2.207 3.072 4.777 5.441 2.988 4.585 32.152 10.589 3.787 3.520 3.233 4.355 1.313 3.360 6.340 2.745 4.498	2.852 2.151 3.301 4.775 5.875 3.039 4.843 31.798 10.901 4.103 3.730 3.446 4.469 0.624 2.972 6.634 2.933 4.315	4.176 2.777 2.281 7.052 8.015 3.501 4.764 20.857 6.948 5.794 5.187 2.781 4.945 1.072 2.832 6.158 2.831 4.042	2.053 3.398 4.659 5.903 2.837 4.997 33.547 10.913 3.626 3.466 2.658 4.090 0.303 3.224 6.309 2.767 4.571	2.899 2.210 3.229 4.752 5.938 3.125 4.984 32.922 11.297 4.107 3.742 3.362 4.530 0.499 3.352 6.534 3.072 4.500	2.854 2.247 2.993 5.075 5.716 2.927 4.778 30.516 10.820 3.927 3.634 2.574 4.492 0.838 3.225 6.441 2.884 4.550	2.567 2.082 2.757 4.462 4.812 2.974 4.611 30.864 9.222 3.899 3.438 3.946 4.170 0.378 3.374 6.495 3.093
Oxidation: Cystine Methionine	2.317 1.371	2.488 1.473	2.236 .1.379		2.191 1.226	2.397 0.328	4.478 2.294 1.383

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## Z PROTEIN

			55-68		56 <b>-68</b>			
	93-68 Durum	57-68 Fine Bran	Soft Red Winter	309-68 3424	92-58 #2 Durum	Straight Grade Flour "Force"	84-68 Halba Ext.	
Protein	13.0	14.9	11.7	<b>. .</b>		<del>9</del> 6	25.5	
Moisture	11.7	11.1	12.6	11 5	11.4	12.4	9.70	
Lysine	2.495	4.124	2.837	2 807	2.514	2.161	6.421	
Histidine	2.043	2.492	2.209	2.257	2.240	2.084	2.421	
Ammonia	2.852	2.235	3.194	2.933	2.737	3.861	1.572	
Arginine	4.255	6.426	4.960	4.808	<b>a</b> . 796	4.083	9.193	
Aspartic acid	4.857	7.401	5.013	5.097	5.564	4.084	11.231	
Threonine	2.964	3.495	2.934	3.044	2.929	2.834	3.724	
Serina	4.676	4.465	4.485	4.850	5.142	4.801	4.896	
Glutamic acid	31.306	20.593	30.629	31.074	35.1 <b>50</b>	38.209	16.728	
Proline	9.654	7.327	8.817	9.220	11.450	11.374	4.038	
Glycine	4.051	5.188	3.939	4.011	3.668	3.505	4.460	
Alanine	3.414	4.886	3.526	3.747	3.455	3.034	3.910	
Half Cystine	3.355	2.716	2.863	3.185	2.875	3.809	0.563	
Valine	4.221	5.371	4.964	4.388	4.567	4.579	4.576	
Methiopine	1.117	1.342	1.118	0.752	0.949	1.088	0.624	
Isoleucine	3.412	3.067	3.521	3.595	3.082	3.554	4.883	
Leucine	6.509	5.927	5.975	6.856	6.359	6.367	5.876	
Tyrosine	3.125	2.876	3.012	3.160	3.029	3.291	3.069	
Phenylalanine	4.478	3.964	4.536	4.684	5.116	4.965	4.088	
Oxidation:								
Cystine			2.526	2.219		2.235		
Methionine			1.392	1.731		1.165		

# Z PROTEIN

	71-68 Bird Seed	151-68 Lobia	325-68 204 Soja	70-68 Draken Spli: Pean	332-68 Garden Peas	<b>JJJ-68</b> Smell Br. Crowder	336-68 Crower Cow Pea
Protein Moisture	16.3	23.7	38.4	· . 4	27 3	20.4	
	10.0	10.0	8.1	10.8	12.6	12.7	23.3
Lysine	2.192	7.366	7.293	7.825	8 (00	<b>.</b>	
Eistidine	2.019	3.292	2.850	2.518	8.609	7.329	7.723
Ammonia	2.421	1.751	2.081	1.595	2.544	3.201	3.587
Arginine	5.810	7.279	8.527	10.862	2.114	2.100	2.299
Aspartic acid	5.543	12.932	13.011		10.456	7.266	8-600
Threonine	2.472	4.046	4.306	12.934	11.442	11.021	13.139
Serine	4.173	5.427	5.600	3.914	4.131	4.205	4.435
Glutanic acid	29.705	19.937	21.618	4.989	5.101	5.497	5.911
Proline	7.349	5.855	5.207	18.785	18.851	18.728	21.189
Glycine	2.936	4.283	4.487	5.255	4.197	4.816	5.508
Alamine	4.198	4.413	4.576	4.402	4.781	4.640	4.688
Half Cystine	3.011	0.000		4.501	4.812	4.673	4.793
Valine	4.855	5.483	2.867	0.685	0.992	0.524	0.000
Methionine	1.274	1.167	5.013	5.005	3.236	5.184	5.984
Isoleucine	4.103	4.478	1.217	1.388	0.000	1.309	1.643
Leucine	6.498	7.286	5.056	4.348	4.247	4.426	4.637
Tyrosine	2.583	3.370	7.348	8.113	7.255	7.948	8.499
Phenylalanine	5.531	5.985	3.955	3.407	3.291	3.569	3.596
		J. 90J	5.532	4.773	4.658	5.760	6.388
Oxidation:							
Cystine		0.921	•				
Methionine		0.796			1.278	1.012	1.009
		,.			1.045	1.375	1.425

# 2 PROTEIN

	- 32 <b>8-</b> 68 191 Pois	36-68 Wheat 6 Barley	144-68 Berley	94-68 Barley	338-68 Hg-treated Sorghum	126-68 Sorghum	318-68 SH20 Hybrid Nawcein
Protein	23.7	12.1	10.8	9.4	10.2	• 9.5	16.3
Moisture	9.9	11.4	11.0	11.3	12.6	12.3	11.6
Lysine	7.901	2.492	3.277	3.424	2.153	1.915	1.716
Histidine	2.342	1.908	2.028	1.969	2.201	2.035	
Ammonia	1.542	2.668	2.815	2.402	2.776	2.745	2.023
Arginine	8.186	4.004	4.779	4.697	3.557	3.510	2.917 3.486
Aspartic acid	11.382	4.773	6.442	5.746	8.017	6.991	6.841
Threonine	4.151	2.874	3.542	3.450	3.579	3.302	3.110
Serine	4.687	4.264	4.560	4.234	4.716	4.608	4.347
Glutamic acid	16.932	28.934	28.175	24.393	23.929	21.810	24.054
Proline	3.694	10.138	11.251	10.299	7.216	7.817	7.750
Glycine	4.502	3.397	4.213	3.857	2.778	3.308	2.708
Alanine	4.573	3.398	4.233	3.915	9.073	9.411	
Half Cystine	1.694	2.538	3.015	2.574	0.439		9.660
Valine	4.725	4.067	4.972	4.860	5.942	2.003	1.846
Kethionine	0.771	1.120	0.332	0.718	1.950	5.067	5.020
Isoleucíne	4.344	3.258	3.723	3.419	5.220	0.450	1.228
Leucine	6.410	6.200	7.052	6.787	13.819	4.039	4.076
Tyrosine	3.479	2.931	3.421	3.107	3.339	13.203	12.819
Phenylalanine	4.904	4.920	5.279	5.123	3.807	4.041 5.381	<b>4.270</b> 5.399
Gxidation:	•						
Cystine		1.208	2.027			1 100	
Methionine		1.500	1.286			1.486	1.648
						1.267	1.637

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8	319-68 Corghum Vulgare	77-68 White Corn from Pez	334-68 American Corn	129-68 Domesti: Corn	315-88 Dll' Hybrid Double	74-68 American Corn	149-68 Hung
Protein	10.5	10.4	11.4	11 - A		•	
Moisture	12.1	11.3	13.6	4.5 1; 5	10 1 11 4	16.1 11.0	23.4 11.7
Lysine	2.271	2.221	2.275	\$ 114 -			
Histidine	2.000	2.180	3.013		2.643	1.849	7.247
Ammonia	3.117	2.456	2.693	2.894	2.772	1.827	2.782
Arginine	3.440	4.162		1 997	2.307	1.455	1.792
Aspartic acid	8.214	7.316	3.753	4.665	4.390	2.746	6.963
Threonine	3.244		6.797	6.791	7.566	5.169	12.931
Serine	5.052	3.512	3.723	3.771	3.635	2.760	3.636
Glutamic acid	23.764	4.555	5.189	4.885	5.227	3.692	5.698
Proline	11.131	23.012	23.789	20.441	21.913	15.959	19.936
Glycine	3.135	11.470	11.204	9.019	10.599	7.180	4.855
Alanine		3.283	3.369	3.702	3. 87	2.488	4.213
Half cystine	9.788	9.555	8.394	7.366	8.081	5.852	4.830
Valine	0.000	1.055	3.079	1.837	2.105	0.784	0.457
	4.928	5.656	5.061	5.034	4.563	3.745	5.645
Methionine	0.199	2.294	1.789	0.201	0.391	0.952	1.137
Isolaucine	3.947	4.389	3.886	3.593	3.207	2.728	4.822
Leucine	13.775	13.340	14.296	10.939	13.855	8.861	
Tyrosine	3.798	4.503	4.556	3.985	4.075		8.557
Phenylalanine	5.314	5.463	5.623	4.595	5.314	3.014 3.496	3.409 6.475
Oxidation:							
Cystine	1.509		2.205	2 2 2 2			
Methionine	6.945			2.232	2.077		0.797
			1.989	1.842	1.739		1.183

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345-68		- 10
Mung	131-68	What
Beans	Hung	×+11
21.9	22.9	:u e
11.9	11.1	12.2
	Mung Beans 21.9	Hung      131-68        Beans      Hung        21.9      22.9

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	345-68		1 50		· · · · ·		339-68
	Mung	131-68	White		at la CV	3411-08	Pumpkin
	Beans	Hung	<b>Y</b> illet	-		Mustard	Seeds
Protein	21.9	22.9	.u o		4.4	26 0	20.0
Moisture	11.9	11.1	12.2	. ÷	4	1	30.0 6.9
Lysine	7.547	7.673	1.785	a.,	1 . 2t	46.	
Histidine	2.746	2.883	2.04		1 41-	4.656 2.215	4.020
Ammonia	1.715	1.600	3.005		. 88 .		2.363
Arginine	7.079	7.230	3.369	1 954	5 833	2.044	1.316
Aspartic acid	12.469	13.451	6.530	6.701	576	5.679	15.235
Threonine	3.671	3.639	3.210	3.245	2.925	3.425	9.845
Serine	5.655	5.594	6.914	6.089	4.219		2.826
Glutamic acid	19.160	20.078	22.259	22.565	23.131	3.600	5.202
Proline	4.734	5.665	6.405	7.666		15.805	19.725
Glycine	4.113	3.985	2.376	2.591	1 766	4.847	3.674
Alanine	4.534	4.371	11.068	9.926		4.126	5.935
Half Cystine	0.912	0.000	1.718	0.840	4.460	3.502	4.159
Valine	1.851	5.848	2.286	5.265	2.756	2.556	2.321
Methionine	1.270	1.091	1.509	2.534	4.683	3.533	4.298
Isoleucine	4.610	4.621	3.982		1.323	0.351	1.579
Leucine	8.334	7.581	12.367	3.979	3.829	3.268	3.556
Tyrosine	3.489	2.897	3.988	10.366 3.418	6.559	5.547	6.589
Fhenylalanine	6.490	6.177	5.780		3.033	2.246	3.717
	0.490	0.177	3.700	5.100	5.113	3.355	4.829
Oxidation:							
Cystine		0.710	1 8-		2.140	2.099	1.357
Methionine		1.264	4.810		1.640	1.362	2.221

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		80-68	8++	8		1 <b>.</b>	392-01
	152-65	thick	(h:			hi .	
	Rice	Peas	Резн	* 46	•	Peas	Chick Peas
Protein	· .	21	14 1	<b>.</b> .			
Doisture	12.8	9.40	19 B		*	e .	9.0
Lysine	3.989	6.832	1400 F.				
Histidine	3.156		141	<b>H</b> 43	•*··)	• • · · q	7.312
Ammonia	2.236	2.534	1.628	1 4	f	2.817	2.642
Arginine	8.046	1.802	1.61.	. :	· · · · ·	1 '71	1.714
Aspartic acid	10.649	10.339	8.825	· •	* *	- 928	10.216
Threonine	3.870	12.745	12.353	1. 884	1	12.542	.2.147
Serine		3.770	3.879	4.4.3		3.998	3.685
Glutamic acid	5.223	5.877	5.520	5.479	4.393	5.606	5.247
Proline	19.659	18.404	17.740	19.444	14. 797	17.689	17.108
Glycine	4.815	3.593	4.111	3.743	1 :16	4.418	4.131
Alanine	4.551	4.163	4.183	4.263	1.251	4 378	4.177
	5.790	4.374	4.537	4.473	1.584	4.519	4.329
Half Cystine	2.554	1.557	1.859	0.861	J. 188	2.627	2.191
Valine	6.297	4.995	4.895	4.929	3.741	4.591	4.365
Methionine	1.519	0.883	0.964	1.238	0.734	0.906	1.026
Isoleucine	4.352	4.017	4.442	4.530	3.050	4.459	
leucine	7.202	6.878	7.833	7.000	5.948	8.101	4.259
fyrosine	4.706	1.940	3.126	3.226	2.332		7.646
Phenylalanine	5.191	4.169	6.171	6.216	5.037	3.099 5.837	2.868 5.623
xidation:							2.023
Cystine	2.434						
lethionine	2.575						

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Studies on Attas and Chapatis I  $\frac{1}{}$ 

Experimental Production of Atta

M. Shafiq Chaudhry, <sup>2/</sup> M. M. MacMasters, E. P. Farrell and W. J. Hoover Department of Grain Science and Industry

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### Summary

An experimental method was developed for making atta, the coarse tour from which unleavened bread (chapatis) is made in Pakistur and India. Attas of 80% and 95% extraction were prepared from Hard it winter, Haro Fed Spring, Durum, White Club, Soft Red worder and Pakistania wheats Particle size distribution of experimental atous were to the range of that of typical Pakistani attas. Proximate componention and dough characteristics of the experimental attas were tetermined. A standard method was developed and used to produce chapatis from the experimental attas. The chapatis were evaluated organoleptically by a panel using a scoring system devised for the purpose. All attas except those from Durum wheat yielded satisfactory chapatis. color apparently is a major factor in determining acceptability, the White

Contribution No. <u>649</u>, Department of Grain Science and Industry, Annsas Agricultural Experiment Station, Kansas State University, Manhattan, Kansas 66502.

 $\frac{27}{1}$  This paper is based in part on the dissertation presented by M. Shafiq Chaudhry in partial fulfillment of the requirements for the Ph.D. degree. Dr. Chaudhry's present address is Department of Food Science, West Pakistan Agricultural University, Lyallpur, Pakistan.

Section Carter Security

Club and Pakistani wheat products were found to be significantly more acceptable than those from any red wheat.

#### Introduction

Unleavened bread is a staple article of food in Pakistan and India. It is similar in appearance to a tortilla but made of course wheat flour and known as a "chapati". The flour from which chapatis are made is called "atta". The typical chapati is prepared from atta and water mixed into a dough that is cooked on a flat, ungreased, hot surface. The chapati is turned several times during cooking and finally, if of good quality, it puffs, i.e., the two surfaces separate because of considerable expansion of gases, probably mostly steam, between them. The puffing subsides as the chapati cools, so the cooked chapati resembles a light tan tortilla flecked with dark brown spots. When cold, a cooked chapati is soft and pliable.

There currently is much interest in possible improvement of average diets in both Pakistan and India. As chapatis are the staple food and sometimes practically the only food in large parts of both countries, improved nutrition must depend primarily on improvement of the nutritive value of chapatis. Particle size or conventional wheat flour of Europe and the Americas is, however, usually considered too small for good chapatis. There is very little in the literature on the production and characteristics of atta. Yet it is obviously very difficult to study nutritional improvement of a product unless the raw material is either readily available or can easily be produced.

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#### Materials and Methods

Seven samples of wheat were used for the study. Each was fumigated when received.

<u>Hard Red Winter</u>: One sample, Triumph variety, from a farm near Scott City, Kansas.

<u>Hard Red Spring</u>: One sample, Selkirk variety from Minnesota. <u>Durum</u>: One commercial blend from Peavey Mills, Minneapolis. <u>White Club</u>. Two samples, Burt and Gaines varieties, respectively, grown in the state of Washington.

Soft Red Winter: One composite sample already on hand. Pakistani: One blend of improved varieties C-273 and C-228 from the Department of Plant Breeding and Genetics, West Pakistan Agricultural University, Lyallpur.

Five hundred grams of each sample was milled and the resulting atta tested for making chapatis. (Wichita variety, (Hard Ked Winter) a commercial blend of Hard Red Spring and Omar variety, (White (inb) also were used in preliminary tests and found to be not appreciably different from other samples in their respective classes.)

A milling method was developed that yielded attas lowely approximating in properties samples obtained from Pak stan. A Hart-Carter Dockage Tester was used for cleaning, with all settings as recommended by the manufacturer. Each sample was conditioned to 15% moisture content by adding the calculated amount of water as a spray to the grain as it was tumbled in a small rotating drum. The conditioned wheat was held in polyethylene bags 24-48 hours before milling. The milling flow sheet is shown in Fig. 1. Ross experimental roller mills were used.

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All except the smooth pair of rolls were of "Getchell" type, set dull to dull. The sifting was carried out on a Smico laboratory sifter for 2 min., except in the fourth step where the time was 15-30 sec., depending on the wheat being milled, to obtain approximately 5% overs of 44 W.

Two types of atta, 95% and 80% extraction, were obtained. For 95%' extraction, 5% bran (overs of 44 W from the fourth grind) was removed. For 80% extraction, 15% fines (thrus of 10%%) was also removed. After appropriate removal the remaining millstreams were pooled and blended 25 min. in a laboratory blender.

Eleven samples of atta that represented products made (a) on steel roller mill (A-E), (b) on the manually operated stone grinder locally known as a "chakki" in Pakistan (F-H), and (c) on animal-powered stone burr mills (I-K) were employed to determine the particle size distribution of acceptable attas. (Letters refer to designations in Fig. 3.) Two hundred grams of atta were sifted for 2 min. on a Rotap sifter through sieves indicated in Fig. 1. The overs of each sieve and thrus of 150 W were used to calculate a cumulative particle size curve.

Analyses for moisture, crude protein, ash, crude fat and crude fiber were made according to methods 44-15, 46-10, 08-10, 30-20, and 32-15, respectively, in Cereal Laboratory Methods (1962).

Preliminary studies were made of doughs prepared by housewives from Pakistan and India. The amounts of water used by three housewives with 300 grams of atta of 95% extraction (from Gaines variety wheat) were 49%, 59%, and 71.5%, respectively. A small amount of vegetable oil was added when 49% water was used. Each dough was immediately

-4-

placed in a large size farinograph bowl and the consistency determined. The values were 1,000 B.U., 780 B.U. and 410 B.U., respectively. All gave acceptable chapatis.

A mechanical procedure to evaluate doughs, for chapatis was devised on the basis of knowledge obtained of machinery used for making tortillas. Two sets of sheeting rolls were adapted to determine machinability. The first set, a product of National Machinery Co., had a roll speed of 85 r.p.m., peripheral speed 84 ft. per min.; this set was adjusted to a clearance of one-eighth inch between the rolls. The second set, a product of Anetsburger Bros., Inc., had a roll speed of our.p.m., peripheral speed 52 ft. per min., adjustment was made to a clearance of 0.050 in, between the rolls. Attas were evaluated for machinability by starting at 71% absorption and decreasing absorption by 12 at each trial until a dough was obtained that made a clean pass through each set of rolls. Wrinkling after the second rolls could be reduced by (a) substituting 2% water with an equal amount of vegetable oil (Wesson oil), (b) reducing absorption by 5% from the level that gave no sticking to the rolls, (c) generous use of dusting flour (atta). Combining the three factors completely eliminated wrinkling and improved the handling properties of the chapatis. In Pakistan and India, the housewife commonly uses over 70% absorption and employs dusting flour (atta) to improve handling properties. Fat is sometimes used in making chapatis in India; the product is then not a chapati but a "paratha". Economic considerations often prohibit the use of butter fat or processed vegetable oils in India and Pakistan.

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Procedures 54-21 and 54-10 in Cereal Laboratory Methods (1962) were used to evaluate dough characteristics in the farinograph and extensiograph, respectively.

Cooked chapatis were evaluated organoleptically by a panel of judges composed of seven students from Pakistan and India to whom chapatis were a common and well-known food. A scoring system was devised by which each characteristic was evaluated on a scale from 0 to 10, a score of 0 representing the poorest quality and 10 the best (Fig. 2). Each chapati was prepared from 50 gm. of atta, sheeted, cooked on a hot plate, cooled for about 5 minutes and wrapped in wax paper until it was presented to the judges. Organoleptic data were analyzed by students "t" distribution test and analysis of variance (Alder and Roessler 1958).

## Results and Discussion

The particle size distribution of attas obtained from Pakistan varied considerably (Fig. 3); that of sample A, from a modern mill, was considered to represent what would probably be available for largescale production of chapatis, similar to production of bread in a commercial bakery.

The method developed for milling wheat to atta gave products with particle size distribution well within the limits of distribution found for Pakistani attas (Figs. 4, 5, 6), although there were large variations among both the commercial and the experimentally milled samples. Attas of 95% extraction from Pakistani and Hard Red Spring wheats were similar in particle size distribution. However, Aziz and Bhatti (1962) found a

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wide range in the granularity of attas from various sources, yet all made good chapatis. Probably color is more important than granularity, within a wide range, in determining quality of atta.

Proximate composition of the attas is shown in Table I. Protein . intent was from il.uot to 15.42%, ether extract was from 1.40% to 2.02% and as content was from 1.38% to 1.81%. Hard Red Winter wheat most nearly approached Pakistani wheat in the protein content of the atta, whereas the hard wheats gave attas of appreciably higher ash content the that of atta from Pakistani wheat.

Note that we could that permitted preparation of a machinable fought of minimum area. Attas of the other wheats were machinable when ware was added at 5% below maximum absorption, 2% of the water was replaid with vegetable oil, and generous use was made or dusting fl

Far regraph curves (titration to 500 B.U.) are some of Figs 7 and to extension raph curves are shown in Figs. 9 and 10. Neither showed differences that could be related to differences in quality of chapatis made from attas.

Rate of extraction of the atta had no significant effect on acceptability of chapatis except in the case of Gaines wheat, where the atta of 95% extraction was judged to yield a slightly better chapati than the atta of 80% extraction (Table II).

When chapatis from artas of 95% extraction were compared, those from Burt, Gaines and Pakistani wheats were significantly better than the others, chiefly on the basis of color, but with a slight influence from flavor differences (Table III).

-7-

A similar trend was noted when chapatis from attas of 80% extraction were compared. In this series, flavor played a more pronounced role in the differences. The judges may have been more influenced by color than they realized, as the tests were carried out under ordinary fluorescent lighting. The tests may not have shown actual differences in flavor, but simulated the usual conditions of consumption closely enough to give good indication of consumer preference.

The results explain why U.S. wheats are often judged inferior for making atta. If only White Club wheats were supplied to atta producers, the quality of U.S. wheats might well be judged excellent.

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The authors are grateful to the Western Wheat Quality Laboratory for supplying the White Club wheats and to those who furnished the other samples; to the Ganesh Flour Mills, Lyallpur, Pakistan, for a sample of atts; to the members of the judging panel. The senior author appreciates the support of himself by A.I.D. during the study and the facilities, equipment and supplies furnished by Kansas State University.

Atte	Extraction rate	Noisture	Crude protein	Crude fat	Ash
		(7.)	(%)	(%)	(%)
Hard Red Winte	r 80%	12.95	14.56	1.66	1.72
Hard Red Winte	r 95%	13.03	15.04	. 1.47	1.60
Hard Red Sprin	g <b>807.</b>	13.43	13.69	2.02	1.76
Hard Red Sprin	g 95%	13.21	14.11	1.85	1.61
Soft Red Winte	r 80%	13.35	12.92	1.85	1.81
Soft <b>Red</b> Winte	r 95%	13,35	12.46	1.70	1.62
lurt	80%	13.30	11.65	1.59	1.44
Burt	95%	13.02	12.02	1.40	1.38
Gaines	807.	13.28	12.45	1,76	1,64
Gainge	95%	12.79	12.04	1.42	1.55
Pakistani	95%	12.45	15,42	1.59	1.44

TAB	LE	Ĩ

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Chemical analysis of atta samples milled from indicated wheats\*

\* Crude protein, crude fat, and ash are expressed on moisture free basis

TABLE II

Effect of the rate of extraction on the organoleptic properties of chapatis prepared from attas of 80% and 95% extraction of indicated wheats.

Type of wheat	Rate of	Average score of seven panel members					
	extraction,	Color	Flavor	Texture	Acceptability		
Hard Red Winter	80	7.00		7.71	8,00		
	95	7.29	<b>n.s</b> 7.43	n.e 7.29	n.# 8.00		
Hard Bad Cantas	80	<b>.</b> .86	7.29	7,43	7.43		
Hard Red Sprim	95	n.#	. n.s 7.43	<b>D. 4</b>	n.s 7.21		
Boft <b>Red Wi</b> nter	80		7.00	7.14	7.14		
Net and winter	95		n. <b>s</b> 7,43	n.s 7.57	n.e 7.57		
urt Wheat	80		<b>8.</b> 79		9.36		
	95	n.s 9.79	n.s 8,43	n.s 8,57	n. s 9.07		
aines Wheat	80		8.86	8.64	8.93		
	95	n. <b>s</b> 9.57	n.s 9.00	n.s 8.71	* 9.36		

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significant at 0.05 level \*

nonsignificant 3,8

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## TAPTE III

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Analyses of variance for color, flavor, texture and acceptability of chapatis

Source of	Degrees of		Mean squar	re and significa	ince
variance	freedom	Coler	Flavor	Texture	Acceptability
Wheat	5	22.768 <del>**</del> *	3,820*	2.114 n.s	7.394**
Error	36	0.710	1.483	1.609	0.831
Total	41				

prepared from attas of 95% extraction of different wheats.

n.s. nonsignificant

significant at 0.05 level

\*\* significant at 0.01 level

## TABLE IV

Analyses of variance for color, flavor, texture and acceptability of chapatis prepared from attas of 80% extraction of different wheats.

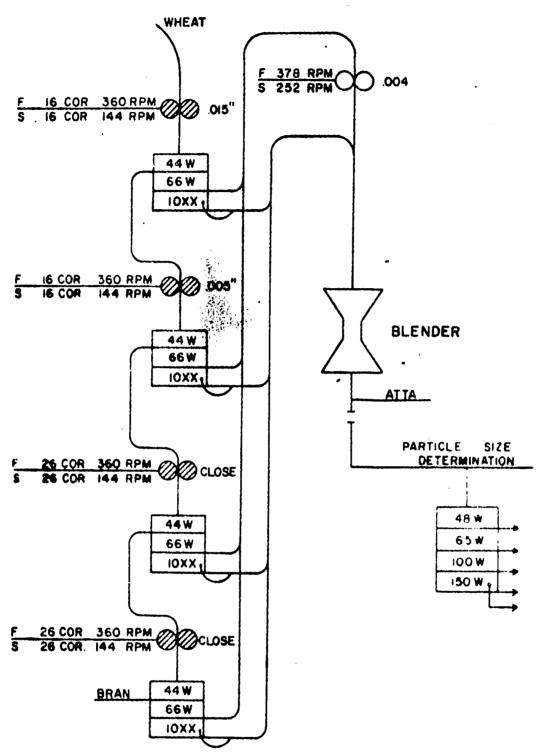
Source of	Degrees of		Mean squar	e and signification	nce
variation	freedom	Color	Flavor	Texture	Acceptability
Wheat type	. 5	23.84**	8.148**	2.024 D.S.	13.924**
Error	36	0.78	1.286	2.175	0.944
Total	41				

n.s. nonsignificant

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\*\* significant at 0.01 level



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Figure 1. A schematic flow sheet for milling of atta from wheat and set of Tyler sieves used in particle size determination.

Score Sheet for Chapatis		
Name	Late	
Elease examine the Charatis -	amples and score these wi	th respect to the
qualities in question, using	the following scale:	
	Best	10 point
	• Foorest	0 point
Luality in question	Score	_
1. Color		
2. Flavor	· · · · · · · · · · · · · · · · · · ·	
3. Jexture		
4. Acceptablisty		

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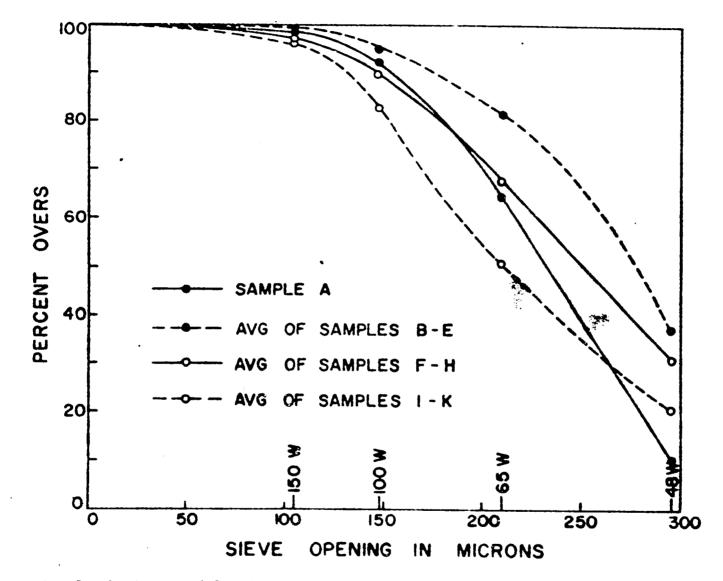


Figure 3. Cumulative particle size distribution curves of atta samples imported from Fakistan.

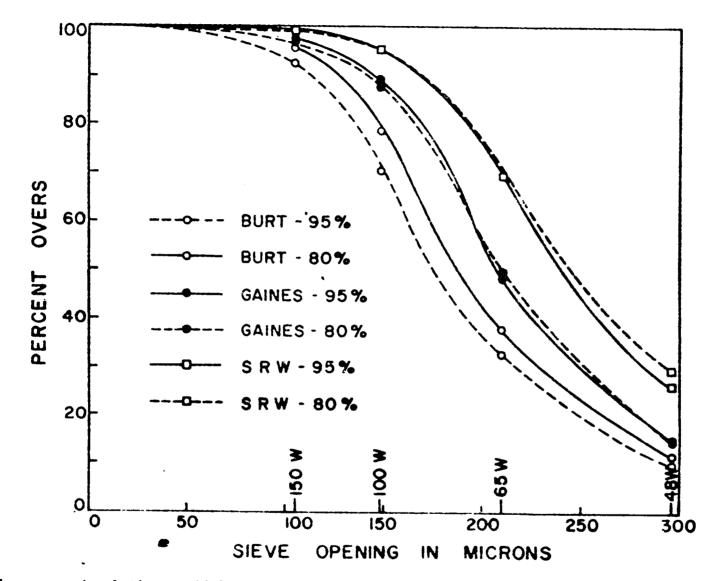


Figure 4. Annulative carticle size listre ution curves of attas milled from Burt, Gaines and loft and Winter wheats.

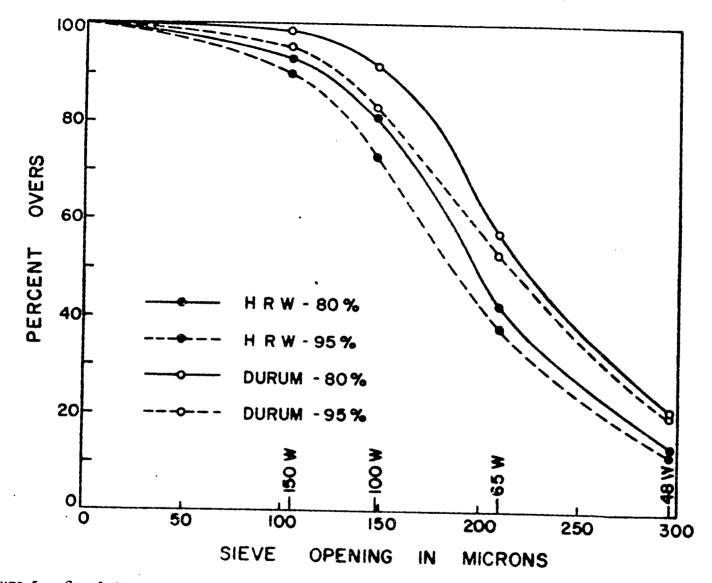


Figure 5. Cumulative marticle size distribution curves of attas milled from Hard Red Winter and Durum wheats.

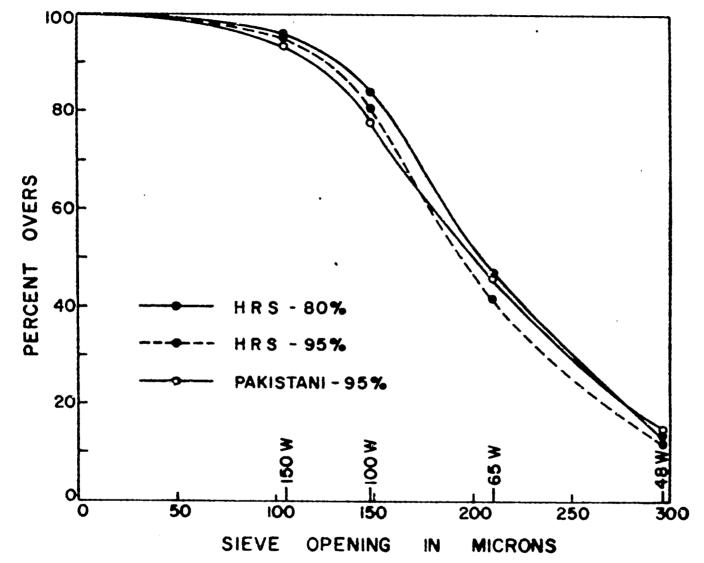


Figure 6. Cumulative particle size distribution curves of attas milled from Hard Red Spring and Pakistani wheats.

Figure 7. Farinograph curves of attas from Gaines, Burt, and HRS Wheats

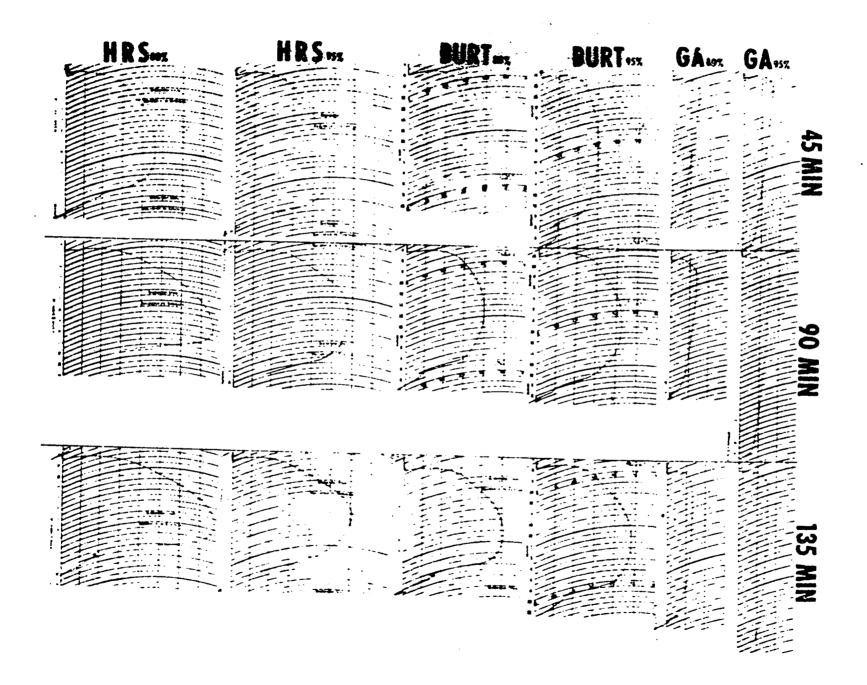


Figure 8. Farinograph curves of attas from SRW, HRW and C 273 Wheats

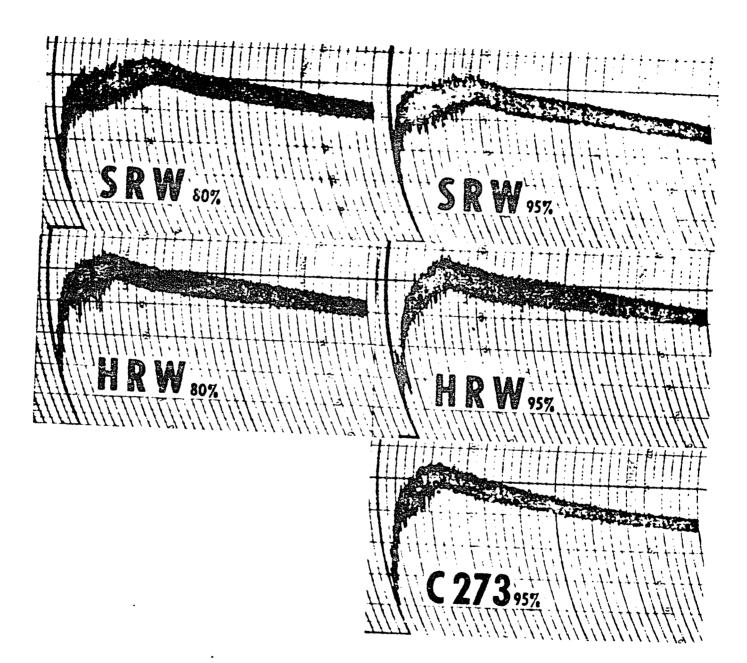


Figure 9. Extensiograph curves of attas from Gaines, Burt, and HRS Wheats .

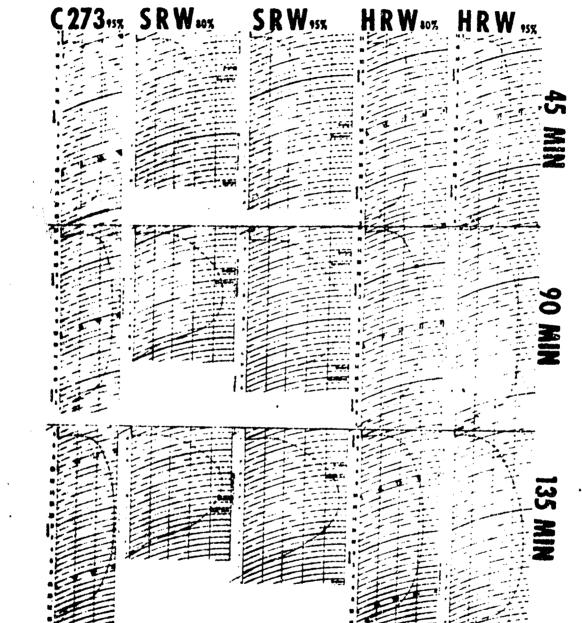
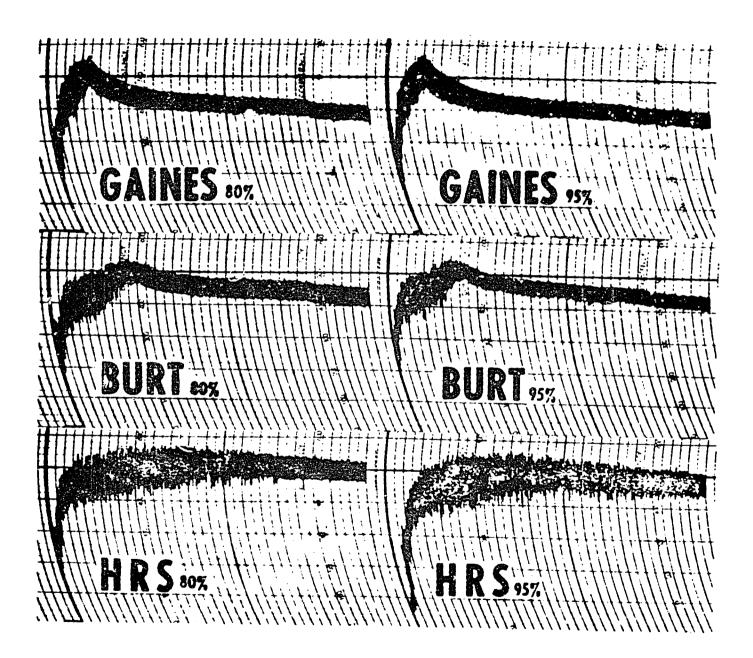


Figure 10. Extensiograph curves of attas from HRW, SHW, and C 273 Wheats

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Studies on Attas and Chapatis II

Nutritional Value of Chapatis With and Without Added Lysine

M. Shafiq Chaudhry<sup>1</sup>, M. M. MacMasters, and W. J. Hoover

### Summery

Rate were fed ground whole wheat, attas of 80% and 95% extraction and chapatim made from them with and without supplementation with vitemins and minerals and with lysins at two levels of supplementation but without supplementary vitamins and minerals. Growth rates and protein efficiency ratios (FER) were evaluated usekly over an eight week period. The PER values were higher for chapatis then for attas from which they were prepared. Supplementation with vitamins and minerals resulted in an increase in PER during the 8 week feeding trial. A similar improved PER resulted during the first four weeks on dists with added lysing but without added vitamins and minerals, but an adverse effect on PKR during the second four week period was found using this dist. Livers of rate fed lysine-supplemented dists had lower moisture content and higher protein content than those of the rate on ether diets. Fortification of cereal products with lysine in the absence of adequate concurrent fortification with vitamins and minerals may not be of volue when the products form assentially the only article of dist as is often true in developing countries.

Contribution No. \_\_\_\_\_. Department of Grain Science and Industry, Kansas Agricultural Experiment Station, Kansas State University, Manbattan, Kansas 66502.

This paper is taken from a portion of the dissertation presented by H. Shafiq Chaudhry in partial fulfillment of the requirements for the Ph.D. degree at Eanses State University. Br. Chaudhry's present address is Department of Food Science, West Pakistan Agricultural University, Lyallpur.

#### Introduction

Improving the nutritional status of developing nations, most of which produce insufficient protein for the national need, is currently of much interest. Often a cereal food constitutes the major portion of the diet of most of the people and may even be essentially their sole source of mutrition. In West Pakisten and a large portion of India, unleavened bread, known as "chapatis", holds that position. Improvement of the nutritional status of the people in those areas will be most easily and acceptably accemplished by improvement of the nutritional value of chapatis.

Flour enrichment programs in the United States, Newfoundland, the Philippines and electroner have established the beneficial nutritional effects of vitamin and mineral supplementation of basic careal foods. There have been very few studies made of the nutritional value of chapatis. The Protein Efficiency Ratio (RER) of chapatis was reported by Shayamala and Kannedy (1962) to be about 20% higher than that of unbeated flour, and replacement of 10% flour with soy flour or dry milk solids was found to further increase the FER. Initias (1962) reported that chapatis prepared from whola-wheat pastry flour with the addition of 15% medium fat soy flour and 10% dry skimmed milk supported emcellent growth of rate.

In areas where chapatis form the staple food, it has been felt that the limiting amino acid in the dist is lysina, since the lysine content of wheat protein is known to be the limiting factor in that protein for humans. It is likely that vitamins and minerals are not at optimal levels in dists based langely on chapatis. The losses of thismin that occur during milling of wheat to the coarse flour, called "atts", from which ehapatis are made, was studied by Singh <u>et al</u>. according to Asis and Bhatti (1962). Losses of 20% to over 50% of thismine were reported, the less being dependent upon the type of milling.

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The present study was undertaken to explore the possibility of improving the nutritional value of chapatis by the fortification of atta with lysine and some vitamins and minerals.

#### Materials and Methods

Gaines variety White Club wheat was milled experimentally to produce two attas of 80% and 95% extraction respectively; the wheat and the milling procedure were described by Chaudhry et al. (1968).

Chapatis were prepared in batches, each from 1000 grams of atta (d.b.), to which 70-75% distilled water was added to produce a dough of the proper consistency. The dough was kneaded by hand, divided into balls weighing 50 grams each, and each ball shaped with a rolling pin into a chapati of 6to 7-inch diameter. Each chapati was cooked on an ungreased hot plate  $(290^{\circ} - 300^{\circ}C.)$  for approximately two minutes.

Cooked chapatis were air-dried in the laboratory  $(65^{\circ} - 75^{\circ}F.)$  for 48-72 hours, then ground in a Wiley Hammer Mill No. 1 to pass through a 1-mm. sieve.

Diets containing the two attas and whole ground wheat, as well as the cooked, dried and ground chapatis, were prepared and fed to weanling male rate, as shown in Table I. Each prepared diet was analyzed for moisture content, and 2% sodium chloride and 5% refined cottonseed oil (d.b.) were added to each. The moisture content of each diet was then adjusted to 15% by the addition of distilled water. Vitamin and mineral supplementation levels were based upon multiple increments of the amounts available in the original grain, rather than upon known dietary requirements of the test animals.

Male weanling rats (Sprague-Dawley strain) were used in the studies. They were fed a stock diet for one day before being housed individually, Five randomly selected rats were maintained on each diet. Initial weight was taken after the animals had been on the diets for 5 days and weekly thereafter, for a total of 8 weeks. Food and water were provided <u>ad libitum</u>. At the end of 8 weeks, the rats were sacrificed and the liver of each was removed for analysis for moisture, crude fat and crude protein.

Whole wheat, attas and chapatis were analyzed for moisture, crude fat, nitrogen, ash, crude fiber, thiamine, niacin, riboflavin, calcium, and iron contents by methods 44-15, 30-20, 46-10, 08-10, 32-15, 86-80, 86-51, 86-70, 40-20, and 40-41 respectively, in Cereal Laboratory Methods (1962). The factor 5.7 was used to convert nitrogen to crude protein value. <u>Lactobacillus</u> <u>plantarum NRRL B-531 was the organism used in determining niacin.</u>

Each liver was wrapped in aluminum foil and frozen. The frozen liver was aliced rapidly, and appropriate amounts weighed for analysis. Moisture was determined by the vacuum oven method  $(100^{\circ}C., 5 \text{ hrs.})$ ; dried samples were extracted for 8 hours with ethyl ether (high heat, Goldfish extractor) for crude fat determination. The Kjeldahl method was used to determine nitrogen.

Data on weight gain and on protein efficiency were analyzed by two way classification analysis of variance, Fryer (1966). Duncan's New Multiple Range Test, as outlined by Fryer (1966) was used to determine the significance of differences among means of percentage gain in weight and protein efficiency ratio.

# Results and Discussion

Chemical analyses of the wheat, attas and chapatis are shown in Table II, where each value is an average of 4 to 6 replications. No significant difference in vitamin contents was found between the attas of 80% and 95% extraction. The three vitamins that were determined decreased

-4-

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in amount during cooking of the chapatis, but the difference was not significant when atta of one extraction rate was compared with chapatis made from it. With the exception of minerals, atta of 80% extraction and chapatis prepared from it showed higher contents of nutrients than atta of 95% extraction and chapatis prepared from it.

Average cumulative weight gain curves are shown in Figs. 1, 2, and 3. Atta of 95% extraction (Niet III) promoted significantly better growth than that of 80% extraction (Diet II) and than whole wheat (Diet I) during the first week, otherwise the three diets yielded no significant differences, (Figs. 1 and 2).

Feeding studies to determine the nutritive value of vitamins and minerals added in making chapatis from atta of 80% extraction showed that supplementation at levels of 50% (Diet VIII) or 100% (Diet IX) above the level in the original wheat (Diet I) gave a significant improvement (Fig. 2). Similar results were obtained with chapatis made from atta of 95% extraction in which the vitamins and minerals were added to make them equal in the atta to the amounts present in the original whole wheat (Diet VII). Comparison of the data suggested that rats fed diets based on atta of 80% extraction (Diet IV) and chapatis made from it performed better than those fed diets based on stta of 95% extraction (Diet III) and chapatis made from it. The differences were, however, not statistically significant. Hepburn et al. (1960) found less than half as high a concentration of lysine in the best patent flour than in germ. Removal of 15% fines during production of atta of 80% extraction would therefore, mean removal of 15% lysine-poor material, with the result that the amino acid balance of the atta would be improved. The chapatis made from atta of 80% extraction also contained more vitamins than those made from atta of 95% extraction.

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Supplementation with lysine led to unexpected results (Fig. 3). Addition of lysine at 0.2% (Diet X) and 0.4% (Diet XI) the weight of the atta significantly improved the performance of the test animals during the first four weeks, but caused a decline in the growth rate during the subsequent four weeks. Such an effect has not been reported in any of the numerous studies on lysine supplementations of foods. Rosenberg and Rohdenberg (1952) found significantly improved nutritional value to result from addition of lysine at 0.2% to 0.8% levels to a diet of which 90% was air dried bread; those workers considered 0.2% to 0.4% to be about the optimal level for lysine supplementation, and other workers have come to similar conclusions.

Lack of fortification with vitamins and minerals of the diets to which lysine was added might have caused the observed results. Portification of wheat products with lysine to provide better nutrition for developing nations has generally been recommended. Apparently this is a promising procedure when the diet of the people contains other sources of vitamins, minerals and even small amounts of methionine. In aconomically poor areas of Pakistan and India, chapatis often form the sole article of food consumed over long periods of time. Little fruit, vegetables, fats or oils, meats or fish are eaten with the chapatis in such areas. In view of the results of the present study, it appears that atta supplementation with lysine is of questionable value unless adequate enrichment with vitamins, minerals and other amino acids in marginal supply is also practiced.

Data showing consumption of feed and of protein, gain in weight and PER are given in Table III, and data on analysis of variance are

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shown in Table IV. Differences in PER among the diets in both four week periods were tested for significance by Duncan's NMRT. During the first four weeks, Diet XI (0.4% lysine supplemented) gave the highest PER, but that diet and Diet X (supplemented with lysine at 0.2% level) gave the lowest PER values during the second four week period. PER values were higher for chapatis than for the attas from which they were prepared. This may be due to nutrient availability rather than protein quality. Shayamala and Kennedy (1962) attributed a similar difference found in their studies to the destruction of Trypsin inhibitor during baking. Two other possible factors may be involved. P.rst, Parihar and Charterji (1956) determined by X-ray diffraction studies that starch is gelatinized during the baking of chapatis. The starch would therefore be more susceptible to the action of digestive enzymes. Second, although no information is available on the fate of phytin during the baking of chapatis, Kent (1966) stater that phytin is hydrolyzed during the baking of bread. If hydrolysis occurs as chapatis are baked, phosphorus would be freed, and there would be less probable formation of complexes of calcium and iron with phytin.

During the first four weeks, supplementation with vitamins, minerals and lysine resulted in an increase of the PER and the increased value continued during the second four weeks, except in the cases of supplementation with lysine.

Data obtained on the livers at the end of the feading experiment indicated that the supplementation of the dist with lysine increased protein content and decreased moisture content of the liver. No consistent effects were obtained as the result of rate of extraction of atta, baking or supplementation with vitamins and minerals.

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# (Acknowledgments)

The authors are grateful to the Northern Regional Research Laboratory, Peoria, Illinois, for the culture of <u>Lactobacillus plantarum</u>. The senior author appreciates the support of AID during the course of the study, and the provision by Kansas State University of all supplies and equipment used.

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## TABLE 1

# Composition of diets

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	•	
(2% sodium chloride and 5% fat not shown in this table, were added to dieta before		
and not shown in this table. Here added to doore but	<b>•</b> • •	
	tending	

Diet	Code	Description					e feeding to re	
No.		Procription	The amoun	t of vari	ous nutrients	added (expr	essed as mg./1(	0 8.)
			Thiamine HCl	Niacin	Riboflavin	Calcium Carbonate	Fe SO4 *7H20	Lysine HCl
I	WW	Whole ground wheat	-	-	-			
11	<b>AT-</b> 80	Atta, 80% extraction	-	-	-	-	-	-
III	AT-95	Atta, 95% extraction	-	-	-	•	-	-
IV	<b>CH-80</b>	Chapati from atta of			•	-	-	-
V	Сн-95	80% extraction Chapati from atta of	•	. <b>-</b>	-	-	-	-
VI	CH-80 IX	95% extraction Chapati from atta of	-	-	-	-	-	
VII	CH-95 IX	80% extraction Chapati from atta of	0.008	1.25	0.031	6.71	-	-
VIII	CH-80 1.5x	95% extraction Chapati from atta of	0.188	1.41	0.046	-	-	-
x	CH-80 2X	80% extraction Chapati from atta of	0.368	2.35	0.056	93.64	11.67	-
x	CH-80 2L	80% extraction Chapati from atta of	0.735	4.70	0.112	187.28	23.34	-
XI	CH-80 4L	80% extraction Chapati from atta of	-	-	-	. <u>-</u>	-	200
		80% extraction	-	-	-	-	-	400

## TAPLE 11

Sample	Moisture	Crude protein	Crude fat	Crude fiber	Ash	1:	Calcium	Niacin	Thiamin	Ribo- flavin
	~ 7	%	7	7.	2	mg7	mg7	mg7.	mg7	
Whole wheat	9.97	12.22	1.70	2.62	1.713	4.89	58.4	4.70	0.733	0.112
Atta of 80% extraction	12.39	12.16	2.04	1.95	1.611	4.96	56.3	4.28	0.728	0.096
Atta of 95% extraction	12.57	12.07	1.81	1.72	1.482	5.07	53.9	3.78	0.680	0.074
Chapatis from atta of 80% extraction	7.28	12.15	1.21	2.02	1.643	5.77	56.5	3.58	0.728	0.081
Chapatis from atta of 95% extraction	6.54	12.09	1.00	1.79	1.451	7.52	65.1	3.42	0.548	0.066
LSD 0.01 level			0.204)	ĵ <b></b> •	0,035	0.177	3.89		0.203	0,0199

• Analysis of when, attis all chapates (moisture free basis)

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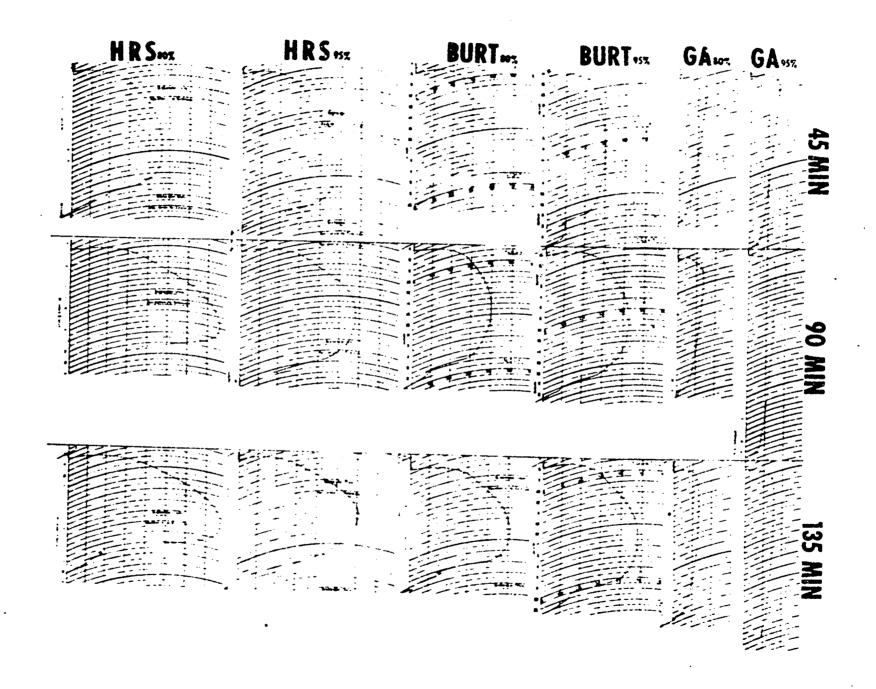
Diet No.	F	irst four we	er period		Second four week period				
	Feed consumed (8.)	Protein consumed (g.)	Gain in weight (g.)	P.E.R.	Feed consumed (g.)	Protein consumed (g.)	Gain in weight (8.)	P.E.R.	
I	225.4	21.66	22.20	1.031	201.2	19.34	13.20	0.680	
II	205.0	20.40	22.60	1.107	200.2	19.92	18.80	0.797	
111	235.2	22.44	22.20	1.011	198.6	18.95	16.40	0.873	
IÅ	201.8	19.84	26.80	1.348	191.4	18.81	22.00	1,172	
V	185.6	17.76	23.00	1.290	176.2	16.86	12.80	0.786	
VI	217.2	21.52	27.20	1.265	197.8	19.60	22.60	1.156	
VII	185.0	17.95	23.80	1.311	193.8	18.18	17.80	0.968	
111	216.2	21.31	32.00	1.4 **	201.2	19.84	22.20	1,105	
IX	261.2	25.62	39.20	1.535	272.8	26.76	35.00	1.317	
X	200.2	20.00	36.40	1.724	190.0	18.98	11.00	0.544	
XI	191.6	19.35	35.20	1.821	225.2	22.74	8.8	<b>0.</b> 410	

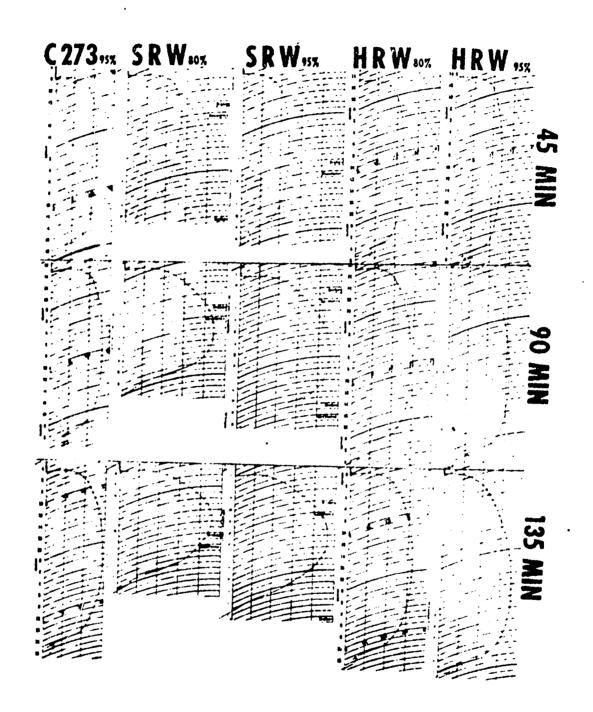
## TABLE III

Average amount of feed consumed, protein consumed, gain in weight a diprotein officiency ratio (P.E.R.

Figure 10. Extensiograph curves of attas from HRW, SRW, and C 273 Wheats

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# Studies on Attas and Chapatis II

Nutritional Value of Chapatis With and Without Added Lysing

M. Shafiq Chaudhryl, N. M. MacMasters, and W. J. Hoover

#### Summery

Rate were fed ground whole wheat, attas of 80% and 95% extraction and chapatie made from them with and without supplementation with vitamins and minerals and with lysine at two levels of supplementation but without supplementary vitamins and minerals. Growth rates and protein efficiency ratios (PER) were evaluated weekly over an eight week period. The PER values were higher for chapatis than for attas from which they were prepared. Supplementation with vitamins and minerals resulted in an increase in PER during the 8 week feeding trial. A similar improved PER resulted during the first four weeks on diets with added lysine but without added vitamins and minorals, but an adverse effect on PER during the second four week period was found using this diet. Livers of rats fed lysine-supplemented diets had lower moisture content and higher protein content than those of the rats on other diets. Fortification of cereal products with lysine in the absence of adequate concurrent fortification with vitamins and minerals may not be of value when the products form essentially the only article of diet as is often true in developing countries.

1/ Contribution No. \_\_\_\_\_ Department of Grain Science and Industry, Kansas Agricultural Experiment Station, Kansas State University, Manhattan, Kansas 66502.

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### Introduction

- 2 -

Improving the nutritional status of developing nations, most of which produce insufficient protein for the national need, is currently of much interest. Often a cereal food constitutes the major portion of the diet of most of the people and may even be assentially their sole source of nutrition. In West Pakistan end a large portion of India, unleavened bread, known as "chapatis", holds that position. Improvement of the nutritional status of the people in those areas will be most easily and acceptably accomplished by improvement of the nutritional value of chapatis.

Flour enrichment programs in the United States, Newfoundland, the Philippines and elsewhere have established the beneficial nutritional effects of vitamin and mineral supplementation of basic cereal foods. There have been very few studies made of the nutritional value of chapatis. The Protein Efficiency Ratic (RER) of chapatis was reported by Shayamala and Kannedy (1962) to be about 20% higher than that of unbeated flour, and replacement of 10% flour with soy flour or dry milk solids was found to further increase the PER. Imitian (1962) reported that chapatis prepared from whole-wheat pastry flour with the addition of 15% medium fat soy flour and 10% dry skinmed milk supported excellent growth of rate.

In areas where chapatis form the steple food, it has been felt that the limiting amino acid in the dist is lysine, since the lysine content of wheat protein is known to be the limiting factor in that protein for bumans. It is likely that vitamins and minerals are not at optimal levels in diets based largely on chapatis. The losses of thismin that occur during milling of wheat to the coarse flour, called "atta", from which chapatis are made, was studied by Singh <u>et al</u>. according to Amis and Ehatti (1962). Losses of 20% to over 50% of thismine were reported, the loss being dependent upon the type of milling. The present study was undertaken to explore the possibility of improving the nutritional value of chapatis by the fortification of atta with lysine and some vitamins and minerals.

#### Materials and Methods

Gaines variety White Club wheat was milled experimentally to produce two attas of 80% and 95% extraction respectively; the wheat and the milling procedure were described by Chaudhry <u>et al.</u> (1968).

Chapatis were prepared in batches, each from 1000 grams of atta (d.b.), to which 70-75% distilled water was added to produce a dough of the proper consistency. The dough was kneaded by hand, divided into balls weighing 50 grams each, and each ball shaped with a rolling pin into a chapati of 6to 7-inch diameter. Each chapati was cooked on an ungreased hot plate  $(290^{\circ} - 300^{\circ}C.)$  for approximately two minutes.

Cooked chapatis were air-dried in the laboratory  $(65^{\circ} - 75^{\circ}F.)$  for 48-72 hours, then ground in a Wiley Hammer Mill No. 1 to pass through a 1-mm. sieve.

Diets containing the two attas and whole ground wheat, as well as the cooked, dried and ground chapatis, were prepared and fed to weanling male rats, as shown in Table I. Each prepared diet was analyzed for moisture content, and 2% sodium chloride and 5% refined cottonseed oil (d.b.) were added to each. The moisture content of each diet was then adjusted to 15% by the addition of distilled water. Vitamin and mineral supplementation levels were based upon multiple increments of the amounts available in the original grain, rather than upon known dietary requirements of the test animals.

Male weanling rats (Sprague-Dawley strain) were used in the studies. They were fed a stock diet for one day before being housed individually. Five randomly selected rats were maintained on each diet. Initial weight was taken after the animals had been on the diets for 5 days and weekly thereafter, for a total of 8 weeks. Food and water were provided <u>ad libitum</u>. At the end of 8 weeks, the rats were sacrificed and the liver of each was removed for analysis for moisture, crude fat and crude protein.

Whole wheat, attas and chapatis were analyzed for moisture, crude fat, nitrogen, ash, crude fiber, thiamine, niacin, riboflavin, calcium, and iron contents by methods 44-15, 30-20, 46-10, 08-10, 32-15, 86-80, 86-51, 86-70, 40-20, and 40-41 respectively, in Cereal Laboratory Methods (1962). The factor 5.7 was used to convert nitrogen to crude protein value. <u>Lactobacillus</u> <u>plantarum</u> NRRL B-531 was the organism used in determining niacin.

Each liver was wrapped in aluminum foil and frozen. The frozen liver was sliced rapidly, and appropriate amounts weighed for analysis. Moisture was determined by the vacuum oven method  $(100^{\circ}C., 5 \text{ hrs.})$ ; dried samples were extracted for 8 hours with ethyl ether (high heat, Goldfish extractor) for crude fat determination. The Kjeldahl method was used to determine nitrogen.

Data on weight gain and on protein efficiency were analyzed by two way classification analysis of variance, Fryer (1966). Duncan's New Multiple Range Test, as outlined by Fryer (1966) was used to determine the significance of differences among means of percentage gain in weight and protein efficiency ratio.

# Results and Discussion

Chemical analyses of the wheat, attas and chapatis are shown in Table II, where each value is an average of 4 to 6 replications. No significant difference in vitamin contents was found between the attas of 80% and 95% extraction. The three vitamins that were determined decreased

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in amount during cooking of the chapatis, but the difference was not significant when atta of one extraction rate was compared with chapatis made from it. With the exception of minerals, atta of 80% extraction and chapatis prepared from it showed higher contents of nutrients than atta of 95% extraction and chapatis prepared from it.

Average cumulative weight gain curves are shown in Figs. 1, 2, and 3. Atta of 95% extraction (Diet III) promoted significantly better growth than that of 80% extraction (Diet II) and than whole wheat (Diet I) during the first week, otherwise the three diets yielded no significant differences, (Figs. 1 and 2).

Feeding studies to determine the nutritive value of vitamins and minerals added in making chapatis from atta of 80% extraction showed that supplementation at levels of 50% (Diet VIII) or 100% (Diet IX) above the level in the original wheat (Diet I) gave a significant improvement (Fig. 2). Similar results were obtained with chapatis made from atta of 95% extraction in which the vitamins and minerals were added to make them equal in the atta to the amounts present in the original whole wheat (Diet VII). Comparison of the data suggested that rate fed diets based on atta of 80% extraction (Diet IV) and chapatis made from it performed better than those fed diets based on atta of 95% extraction (Diet III) and chapatis made from it. The differences were, however, not statistically significant. Hepburn et al. (1960) found less than half as high a concentration of lysine in the best patent flour than in germ. Removal of 15% fines during production of atta of 80% extraction would therefore, mean removal of 15% lysine-poor material, with the result that the amino acid balance of the atta would be improved. The chapatis made from atta of 80% extraction also contained more vitamins than those made from atta of 95% extraction.

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Supplementation with lysine led to unexpected results (Fig. 3). Addition of lysine at 0.2% (Diet X) and 0.4% (Diet XI) the weight of the atta significantly improved the performance of the test animals during the first four weeks, but caused a decline in the growth rate during the subsequent four weeks. Such an effect has not been reported in any of the numerous studies on lysine supplementations of foods. Rosenberg and Rohdenberg (1952) found significantly improved nutritional value to result from addition of lysine at 0.2% to 0.8% levels to a diet of which 90% was air dried bread; those workers considered 0.2% to 0.4% to be about the optimal level for lysine supplementation, and other workers have come to similar conclusions.

Lack of fortification with vitamins and minerals of the diets to which lysine was added might have caused the observed results. Fortification of wheat products with lysine to provide better nutrition for developing nations has generally been recommended. Apparently this is a promising procedure when the diet of the people contains other sources of vitamins, minerals and even small amounts of methionine. In economically poor areas of Pakistan and India, chapatis often form the sole article of food consumed over long periods of time. Little fruit, vegetables, fats or oils, meats or fish are eaten with the chapatis in such areas. In view of the results of the present study, it appears that atta supplementation with lysine is of questionable value unless adequate envichment with vitamins, minerals and other amino acids in marginal supply is also practiced.

Data showing consumption of feed and of protein, gain in weight and PER are given in Table III, and data on analysis of variance are

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shown in Table IV. Differences in PER among the diets in both four week periods were tested for significance by Duncan's NMRT. During the first four weeks, Diet XI (0.4% lysine supplemented) gave the highest PER, but that diet and Diet X (supplemented with lysine at 0.2% level) gave the lowest PER values during the second four week period. PER values were higher for chapatis than for the attas from which they were prepared. This may be due to nutrient availability rather than protein quality. Shayamala and Kennedy (1962) attributed a similar difference found in their studies to the destruction of Trypsin inhibitor during baking. Two other possible factors may be involved. First, Parihar and Chatterji (1956) determined by X-ray diffraction studies that starch is gelatinized during the baking of chapatis. The starch would therefore be more susceptible to the action of digestive enzymes. Second, although no information is available on the fate of phytin during the baking of chapatis, Kent (1966) states that phytin is hydrolyzed during the baking of bread. If hydrolysis occurs as chapatis are baked, phosphorus would be freed, and there would be less probable formation of complexes of calcium and iron with phytin.

During the first four weeks, supplementation with vitamins, minerals and lysine resulted in an increase of the PER and the increased value continued during the second four weeks, except in the cases of supplementation with lysine.

Data obtained on the livers at the end of the feeding experiment indicated that the supplementation of the diet with lysine increased protein content and decreased moisture content of the liver. No consistent effects were obtained as the result of rate of extraction of atta, baking or supplementation with vitamins and minerals.

-7-

## (Acknowledgments)

The authors are grateful to the Northern Regional Research Laboratory, Peoria, Illinois, for the culture of <u>Lactobacillus plantarum</u>. The senior author appreciates the support of AID during the course of the study, and the provision by Kansas State University of all supplies and equipment used.

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# TABLE 1

# Composition of diets

(2% sodium chloride and 5% fat not shown in this table, were added to diets before feeding to rats)

Diet Code No.		Description The amount of various nutrients added (expressed as mg./100 g.						
			Thiamine HCl	Niacin	Riboflavin	Calcium Carbonate	Fe 504 7H20	Lysine HCl
I	w	Whole ground wheat	-	-	-	-	•	•
II	AT-80	Atta, 80% extraction	-	-	-	-	•	<b>-</b> .
III	AT-95	Atta, 95% extraction	-	-		-	-	-
IV	CH-80	Chapati from atta of						
۷	СН-95	80% extraction Chapati from atta of	-	-	• .	-	-	-
VI	CH-80 IX	95% extraction Chapati from atta of	-	-	-	-	-	•
VII	CH-95 IX	80% extraction Chapati from atta of	0.008	1.23	0.031	6.71	-	-
VIII	CH-80 1.5x	•	0.188	1.41	0.046	-	-	-
IX	CH-80 2X	80% extraction Chapati from atta of	0.368	2.35	0.056	93.64	11.67	-
X	CH-80 21	80% extraction Chapati from atta of	0,735	4.70	0.112	187.28	23.34	-
XI	CH-80 41	<b>807, extraction</b> Chapati from atta of	-	-	-	-	-	200
		80% extraction	-	-	-	-	-	400

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	(mo:	istu	:•	free	bas	4		

Sample	Moisture	Crude	Crude	Crude	Ash	1:	Calcium	Niacin	Thiarir	Ribo-
•	Z	protein 7	fat <b>7</b>	fiber 7.	7.	mg7.	mg7.	mg7.	mg <b>?</b> _	flavin
Whole wheat	9.97	12.22	1.70	2.62	1.713	4.89	58.4	4.70	0.733	0.112
Atta of 80% extraction	12.39	12.16	2.04	1.95	1.611	نه . ليو	56.3	4.28	0.728	0.096
Atta of 95% extraction	12.57	12.07	1.81	1.72	1.482	5.07	53.9	3.78	0.680	0.074
Chapatis from atta of 80% extraction	7.28	12.15	1.21	2.02	1.643	5.77	56.5	3.58	0.728	0.081
Chapatis from atta of 95% extraction	6.54	12.09	1.00	1.:-	1.4%.	7.52	65.1	3.42	C.548	0.066
LSD 0.01 level		 •	ο.	·)•••		0.177	- 3.89		0.203	0.0199

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17	r	٠	. •		1	*	

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Average amount of feed consumed, protein consumed, gain to weapon and protect officiency ratio (P.E.R.

	Feed		e- period	_		Second Iou	r week perio	0d
	consumed (g.)	Protein consumed (g.)	Gain in weight (g.)	P.E.R.	Feed consumed (g.)	Protein consumed (g.;	Gain in weight (g.)	P.E.R.
I	225.4	21.66	22.20	1.031	201.2	19.34	13.20	0.680
II	205.0	20.40	22.60	1.107	200.2	19.92	18.80	0.797
III	235.2	22.44	22,20	1.011	198.6	18,95	16.40	0.873
ĨV	201.8	19.84	26.80	1.348	191.4	18.81	22.00	1.172
V	185.6	17.76	23.00	1.290	176.2	16.86	12.80	0.786
VI	217.2	21.52	27.20	1.266	197.8	19.60	22.60	1.156
VII	185.0	17.95	<b>23.</b> 80	1.311	143.8	18.18	17.80	0.968
VIII	216.2	21.31	32.00	1.4 · ·	201.2	19.84	22.20	1,105
IX	261.2	25.62	39.20	1.5 **	272.8	26.76	35.00	1.317
X	200.2	20.00	36	1.724	190.0	18.98	11.00	0,544
XI	191.6	19.35	35.2%	1.821	225.2	22.74	8.8	0.410

of different diets i t - periods of four weeks each.

# TABLE IV

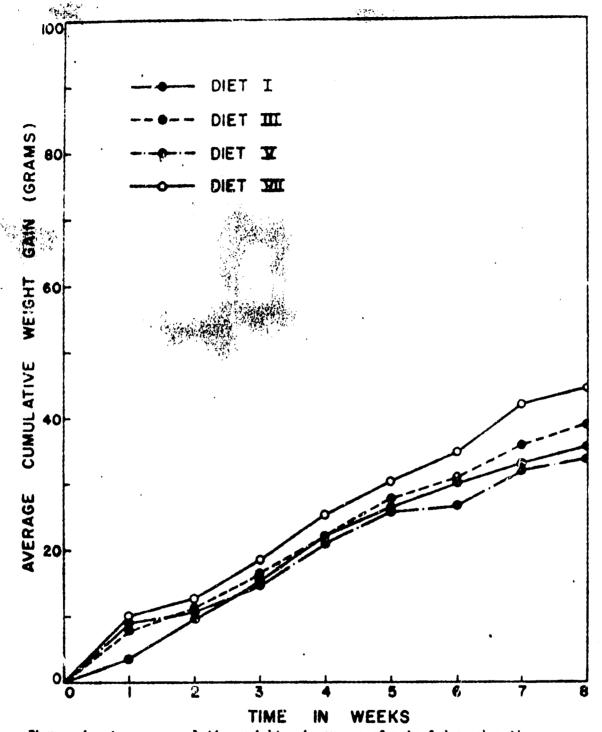
Analysis of variance for protein efficiency ratio of various diets during two growth periods of four weeks each.

Source + ;	Destee of	Sum of		
variation	freedom	squares	<b>Mea</b> n square	F value
Perioda	ì	5,54268	5.5427	109,76**
Diets	1.	2.6688	0.2664	5,285**
Periods X				
Diet•	10	4.78756	0,4788	9,481**
Error	88	4.44127	0.0505	
Total	109	17.44031		

**\*\*** significant at 0.01 level,

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TIME IN WEEKS Figure 1. Avorage cumulative weight gain curves of rate fed on chapatis prepared from 95% extraction atta and supplemented with vitening and minorals.

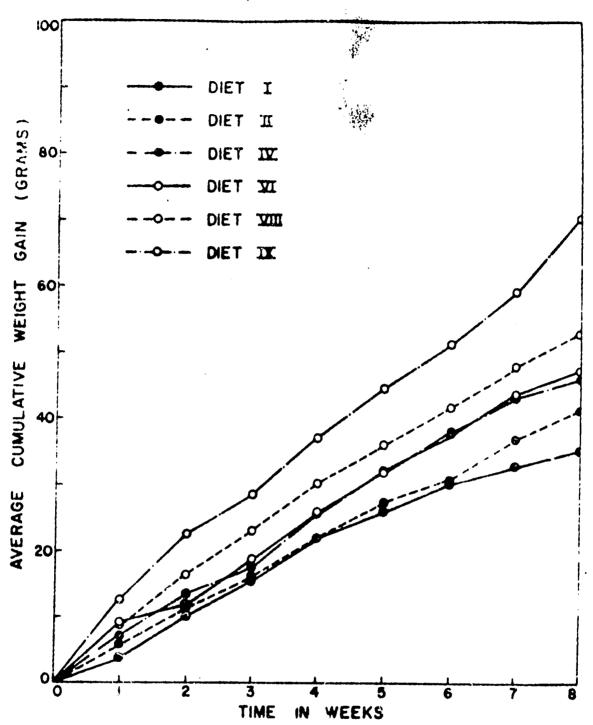


Figure 2. Average cumulative weight gain curves of rats fed on chapatis propared from 80% extraction atta and supplemented with vitaminu and minerals.

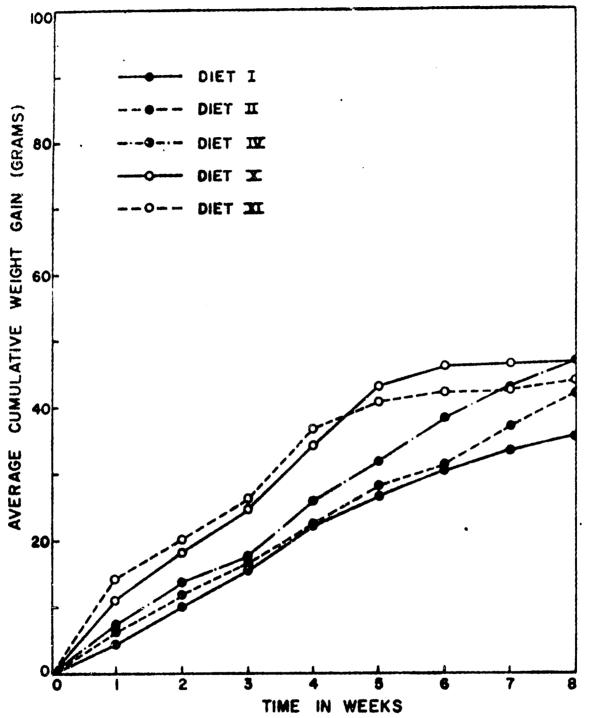


Figure 3. Average cumulative weight gain curves of rats fed en chapatis prepared from 80% extraction atta and supplemented with lysine.

## Studies on Attas and Chapatis III

# A Note on the Relative Experiance of Vitamins and Minerels as Pictury Supplements.

Hoin Y. Chung, N. M. MacMioters and W. J. Hoever

#### Seminary

Proding experiments with vecaling unle rate indicate that adequate supplementation of chapitic with vitamino and minerals is as nocessary for optimal growth as is supplementation with lysine. Supplementation of flour or atta with lysine for mutuiticant improvement in protein-poor countries may be of questionable value unless adequate vitamins and minerals are also supplied.

#### Introduction

Chaushry et al. (196-b), feeding abapatis supplemented with 0.4% lysime to wanning mile rate, found loss in weight of the rate by the eighth week. It was postulated that the natural vitemin and mineral supply perheps been inadequate because of increased metabolic activity brought about by the stimulation of growth caused by the lysime added in the feeding trials.

Exploratory studies were undertaken with the limited mount of stan grepared by Chaudhry <u>et al</u>. (195-a) that remained after the feeding experiments (Chaudhry <u>et al</u>. 195-b) were sensined.

V Contribution HD. . . Bepartment of Grain Science and Industry, Hancas Agricultural Happeriment Station, Reases State University, Manhattan, Reases 66368.

This note is board on part of the dissertation presented by Hoin Y. Chung in partial fulfilizent of the requirements for the Ph.D. degree.

#### Interiale and Mothodo

- 2 -

The amount of atta (90% extraction) available was estimated to be sufficient to make enough chapatis to feed mine rate for six vesks. Three venaling male rate (Sprague-Bawley strain), chosen at random, were therefore placed on each of three dicts after two days on a communical feed (Perime Dog Chev) and water of Libitums

- A. Ground character + 0.4% -- 100
- B. Ground chapping & C.C. Lysian & viennins and minerals
- C. Ground chapatis + 0.4% lysims + 0.04% methiculae + vitanine and minarcls.

The composition of the vitamin-winners's supplement is shown in Table I. In the study made by Chandlery <u>at al</u>. (195-b) vitamine and minerals were added to dists, not supplemented with lysing on the basis of multiple ineroments of the amounts present in the original whole wheat. In the present study, supplementation of the dists was based upon the lowels recommended by the Matiemal Research Council.

Proximate compositions of the three dists are above in Table II.

#### Rocults and Discussion

healte of the amperiment are shown in Figure 1. Supplementation with 0.42 Lysims alone gave a commistive weight gain that use only elightly above that obtained by Chemilary <u>at al</u>. (196-b). The rate used in the present sumfice were delayed in transit, so were several days older when entering the trials then the rate used in the formar studies. Weights of the elder sale were understandedly semathet higher then weights of these proviously used.

The addition of the complete vitamin and minoral supplement used in the present study greatly increased the gain in weight of the rate over that obtained when 0.4% lysine alone was added. Hereover, the gain was much greater them was obtained by Chaudhay gi al. (1968 b) using a limited vitamin supplement.

Addition of 0.64% authioning to the lysing-supplemented dist containing vitamins and minerals caused no approxisble difference in weight gained.

Purther work is unsided to determine the minimal effective amounts of vitumine and minorals movied to supplement shapstis for busine matrition. The present coplexatory copariments, hereway, give strong evidence that vitumine and minorals must be supplied in adaptite amounts if good results are to be obtained through supplementation of atta or shapstis with lysime.

# Table I

.

Vitamia and Minural Contants of Diets

Vitania or Mineral	Supplones Present stydy	Cheadley ALAL (1963)	by wheat	Recommended
			flow, 80% estra.	by IAC
Vitamin A, <u>L.W.</u>	1000	0	0	200
Vitamin D, <u>L.H.</u>	100	0	-	•
Vitamin 2, mg	10	0	-	6
Vitamin K, mg	0.5	C	•	0.01
Thiomine BC1, ag	0.5	0.73	0.26	0.25
Vitemin B <sub>6</sub> , mg	0.4	0	•	0.12
Nidein, ug	4,0	3.50	2.0	1.5
On peatotheaste, ag	25	0	•	0.8
Cheline chloride, ag	200	0	-	75
Vitamin B <sub>12</sub> , SMA-	2	0	-	0.5
Ribeflövin, og	1	0.081	0.00	•
24 <b>34 , co</b> s	10	0	•	•
Biotia, ng	0.02	0	-	•
Polis selú, ng	0.2	0	-	•
On Le tum, mg	700	56.5	24	600
Phosphorus, ng	500	0	191	500
Sodium, ag	50	79	1	<b>3</b> 0
Potassiga, ag	180	٥	95	180
Chlories, eg	241	122	•	50
Nagaoo Lan, ug	49	0	•	40
Waghases, ag	5	0	•	5
Irea, an	5	5.77	1.3	2.5
Copper, as	0.5	0	•	0.5
Ilue, eg	2.5	0	-	1.2
Iodias, mg	0.15	0	•	0.015

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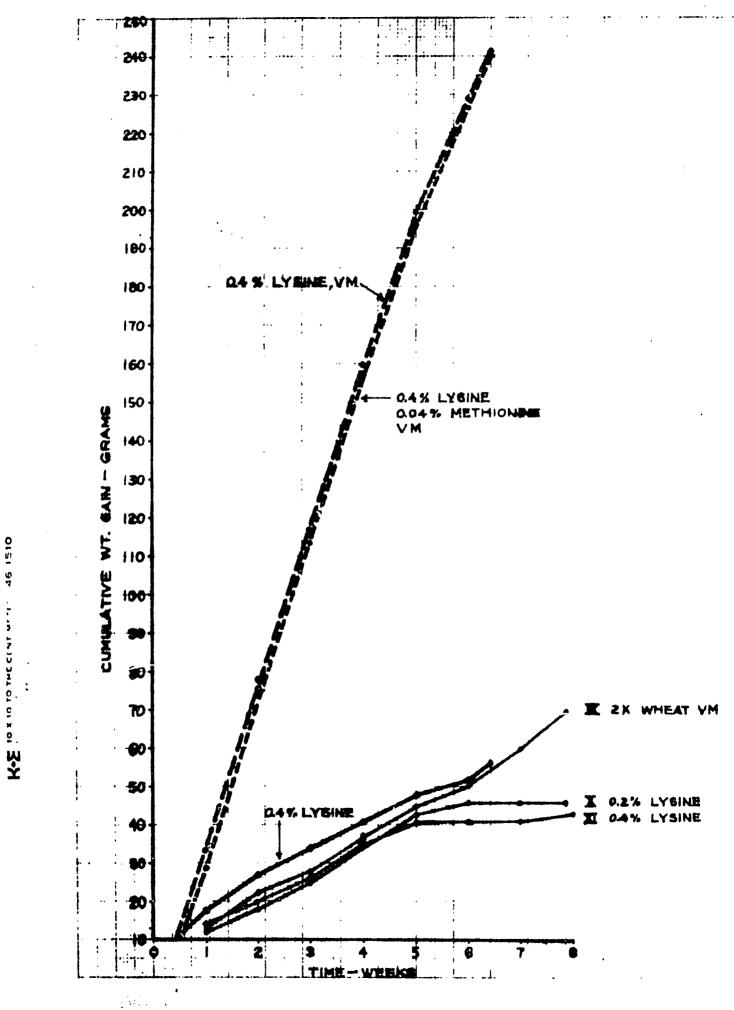
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# Table II

# Proximate Composition of Biets (dry basis)

•

12.3	12.6
5.6	4.8
2.3	2.2
	2.3



15C

#### REFERENCES

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- Chaudhry, N. S., Machasters, N. N., and Masver, W. J. 196-b. Studies on attas and emeratic IS. Matriticaal value of chapatic with and without added lysing. Food Technol.
- U.S.D.A. 1953. Composition of Spade (new, processed, prepared). Agricultural Research Service, U. S. Bepartment of Agriculture, Agricultural Mendbook No. 6.

#### Nutritional Evaluation of Wheat Based Foods:

Rat Study I. Studies on minimum supplementation of high extraction flour.

Objective: To study the effect of supplementing 97% extraction wheat flour with limiting amino acids, minerals, and vitamins, each of the rations shown in Table 1 were fed to eight weanling (21-day old) female albino rats of the Sprague-Dawley strain. The duration of the study was 28 days following an adjustment period of three days. Procedure: Diet I, the control diet, was designed to meet all MRC minimal requirements for the rat. Dist I was supplemented with four amino acide (lysine, threonine, valine, and methionine), six minerals (mangamose, iodine, calcium, phosphorus, sodium, and chlorine), and four vitamins (vitamins A, D, and B<sub>12</sub>, and riboflavin). Supplementation of these nutrients was based on analyzical values on the wheat products and a comparison with requirement values. Diets II through WII had one or more of these supplements omitted as follows: Diet II-threenine, valine, and mothionine; Diet III--calcium; Diet IV--onehalf the level of calcium supplied in the control; Diet V -- phosphorus; Diet VI--iodine; Diet VII--vitamin 8,2. Diet VIII was supplemented with lysine and with vitamins and mimerals greater than the minimal requirements. Diet VIII was included to compare results of the present experiment with those of previous experiments of Chung ( ) and Chaminty ( ).

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	Contro I		Mo Ca	SCO SUPPL	NO P SUPPOL.	NI I	no la	Pean Gapt.		
		1								╺╋╸
6 LysH+1	0.60	0,60	0.60	0.60	0,60	0.40	9.60	0,60		
d1 Inc	9.19		0.19	0.19	2.19	0.19	0,19		-	
L VA.	12.24		0.24	434	AIBY	A124				
d  ne	0.17	-	0,17	919	all	0,17	0.24 917	-		
Mn J 14 "24	6.0057	0.000	9.0087	6189 8	0.0079	0.0059		-		
KI	1	3 6.98.8002		I			0.9587		-+	-+
C.C.O.7	0.011	SAU.			1.521	1	8.60000A	-		
NIPOAR.			0.91	Q.15.7V	ILLAN	<u>8.141</u>	<u>arp</u> 4			
Dices V	1.894	1.3.84		1		1	1.001	+	+	
NGC.			7	ALPTR-		1.834	1.934	2.20		
Ms ?	<u> </u>	6411K3	8.1143	Lands_	611.19	AMAN -	0.1148	0.127	-	
								0.066		
c	+						+	1.05		
KC.		+	<u> </u>	<u> </u>	+	<b> </b>		0.344	+	
P.,	+					÷	ł	<b> </b>	+	-
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VIT E		+		•••••		<u> </u>		10-91	+	
Vπ K								0.014	+	<u> </u>
THA-HCI	<u>                                      </u>							0,0005	+	
Typione Mil	+			-		<u> </u>		20904	+	
NAC N								B	+	<b>_</b>
CA General	<u>↓</u>		-					0.075	+	<b></b>
H1	<u>↓ −</u>			* <b>~</b>				01910	+	
B10							-	0000 p		
Foure - Di								6.690 h	<b>_</b>	
CHOLINEC				-			/	9,80		<b>_</b>
									<b> </b>	<b> </b>
	9.98 3	3.940				2.9/2		s.ce	╡	<b> </b>
Corn St Pup	2.7.962	3.396 2.	1.770	B.//30 2.	3,1410	2, Roba	2.7476	P17	<b> </b>	
7A 1	10.0	10.0	10.0	10.0	10,0	10.0	10.0	10.0	Liva	
4' AT FLOUR	90,0	10.0	90.0	90.0	90.0		90.0	90.0		
GRAND TO TAL	100.0	100.0	100.0	100.D	100.0	100.0	100 0	100.0	an:	Ι
Table	1. Co	maas	<u> 11 100</u>	.1	115 . 7				I	
4		7				1				
				T						T

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The nitrogen content of the dists was determined by the Kjeldahl method (A.O.A.C. 2.036). The dists contained the following amounts of protein (N x 6.25):

Diet I	10.85%
Diet II	10.29%
Diet III	10.75%
Diet IV	10.75%
Diet V	10.7 <b>5%</b>
Diet VI	10.86%
Diet VII	10.86%
Diet VIII	10.427

Animals were weighed weekly. Protein efffciency ratios (PER) were calculated at the end of the 28-day study. The cumulative weight gains and PER's are shown in Table 2.

Diet		Cumletive Weight Goine, g.							
	7 days	14 dayo	21 days	28 dave					
I	14.4	39.6	64.4	84 . 2	1.99				
11	18.8	44.5	69.8	91.9	2.16				
111	17.0	38,0	. 61.6	73.9	1.80				
IV	17.4	42.5	69.8	89.0	2.13				
v	13.1	34.9	55.6	75.5	1.95				
VI	15.4	40.1	66.9	86.2	2.03				
• 11	12.9	92.6	56.0	77.0	1.97				
VIII	16.2	44.5	71.4	92.6	2.11				

Table 2. Cumulative weight gains and protein efficiency ration of rate fod experimental diets.

The analysis of variance (Table 3) showed that differences among PER's were significant at the 1% level.

	Table 3.	Analysis of varia	nce of PER data.	
Source	d, f.	S.S.	M.S.	P
Diets	7	0.824	0.118	4,701**
Brror	56	1.405	0.025	
- 2.10	8	LSD = 0.19 .05	584	
1.01 = 2.98	B			
Differences	among diets	at the 5% level as	e summarized be	low

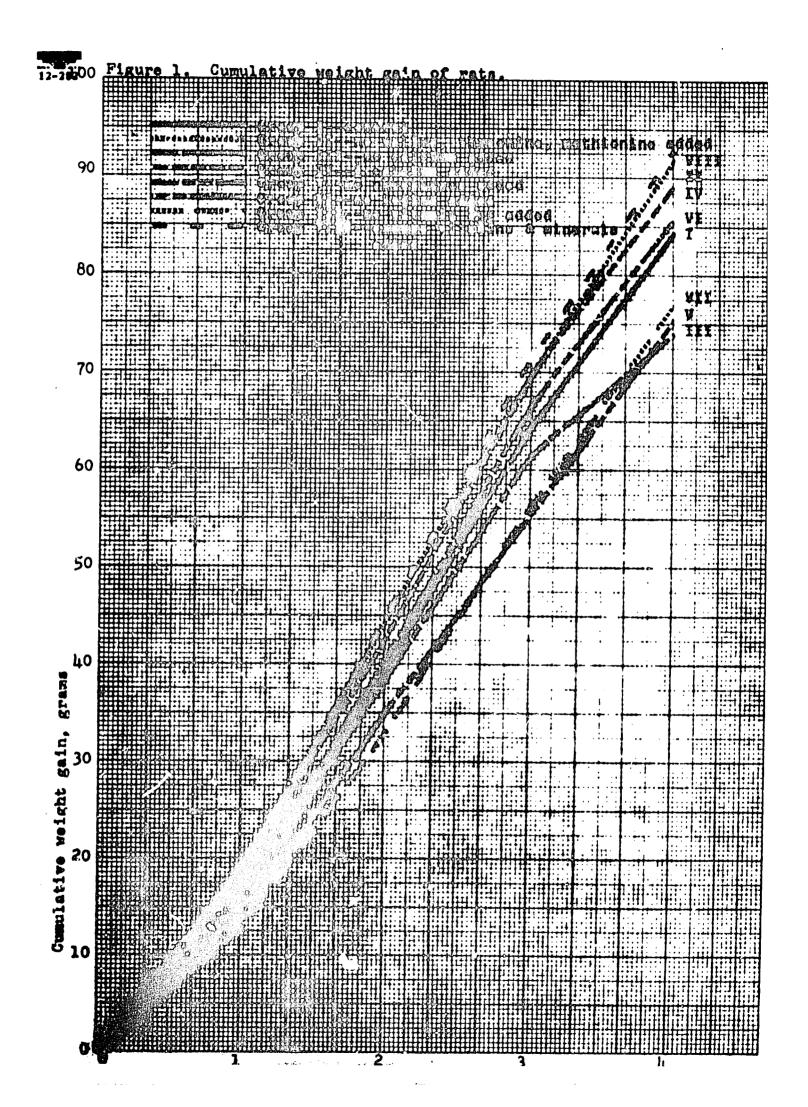
VIII) V, III VI > III Ιÿ III VII > III

In this experiment it was observed that omission of mothionine, valine, and thraonine improved the lysine-supplemented rations as measured by PER's. The supplementation of calcium was necessary, but adding half of the difference between the calcium supplied by the wheat flour and the MRC requirement produced results as favorable as adding an amount equivalent to the entire difference. The inclusion of an excess of vitemins and minerals above NRC minimal requirements did not improve the growth of rats. The addition of phosphorus and vitemin  $B_{12}$  appeared to improve the quality of the rations. The addition of iedina did not alter the quality of the dists significantly.

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Data presented in Figure 1 indicate that the effect of limited calcium was more apparent near the end of the study. These results indicate the need to carry similar experiments for longer periods to determine longer term effects.

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### STUDIES TO EVALUATE THE NUTRITIONAL VALUE OF THE PROTEIN IN MILLED SORGHUM GRAIN FRACTIONS

Preliminary studies of sorghum grain have indicated that the nutritional quality of the protein in sorghum grain varies. The highest nutritional values is observed in the sorghum grain fraction derived from the floury endosperm.

Horny endosperm has a poor nutritional balance being deficient in lysine and other basic essential amino acids. The following experiment was designed to evaluate the nutritional value of the protein of various endosperm fractions.

Two varieties were selected having equal protein content. The grains were milled and various endosperm fractions were supplemented with amine acids for biological study. The additional amino acid supplementation studies were planned to offset the protein parameters.

## EXPERIMENT OBJECTIVES

The objectives of the study were as follows:

(1) To compare the protein quality of milled endosperm fractions from two hybrids.

(2) To study the effects of supplementing endosperm protein with lysine and methionine.

(3) To study the effects of supplementing both high and low protein milled products with lysine and methionine.

(4) To compare the growth of rats fed milled endosperm printies with that of casein.

#### MATERIALS AND METHODS

Two hybrids having equal protein content (9.50% protein, 12% moisture basis) were dry milled using conventional dry milling equipment as described by the milling flow in Figure 1. Amino acid composition of all endosperm fractions are given in Table Ia. The yield and approximate analysis of the milled products are given in Table 1.

Fractions 3 and 7 from each hybrid were divided and half was supplemented with methionine and lysine and the remainder was fed without supplementation. Diets containing fraction 3 (floury endesperm) were fed at approximately a 5.6% protein level while diets containing fraction 7 (horny endosperm) was adjusted with starch to provide approximately 10.2% protein. Casein control diets were fed at both dietary protein levels. Approximate analysis of the diets are shown in Table II.

Table III and IV show the composition of the diets and the levels of lysine and mothionine addition. The amino acids were incorporated to provide those diets with 100% of the NRC requirement.

The ten diets were fed to 22-day old weanling female white rats in individual cages. There were six replications per diet with one ret per replication. The sixty rats, weights ranging from 43 to 50 grams, were randomized to the various diets.

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#### **RESULTS AND CONCLUSIONS**

-4-

Average weight gain, feed consumption, and protein efficiency ratios (PER) for rats fed the floury endosperm products in the low protein (5.6%) diets are shown in Table V. The supplementation of lysine and methionine to diets 2 and 6 resulted in a marked increase in weight gain and PER above the non-supplemented diets. The supplemented diets were similar to the casein diet which was fed at the same protein level.

The same information is provided in Table VI for rats fed the horny endosperm products. These diets were isonitrogencus at approximately 10.2% protein. Again the amino acid supplementation caused a marked increase in gain and PER's. The weight gain of the supplemented diets was superior to the casein diet. The PER's, however, were similar. Horny endosperm from Frontier 400C produced greater gain and had a higher PER than Paymenter Kiowa when supplemented with the amino acids. However, when the PER's were adjusted for the control diets (as shown in Table VII) differences were not significant.

Table VII provides a means of comparison of floury and horny endosperm. The PER's of the experimental dists were corrected by the fraction as follows:

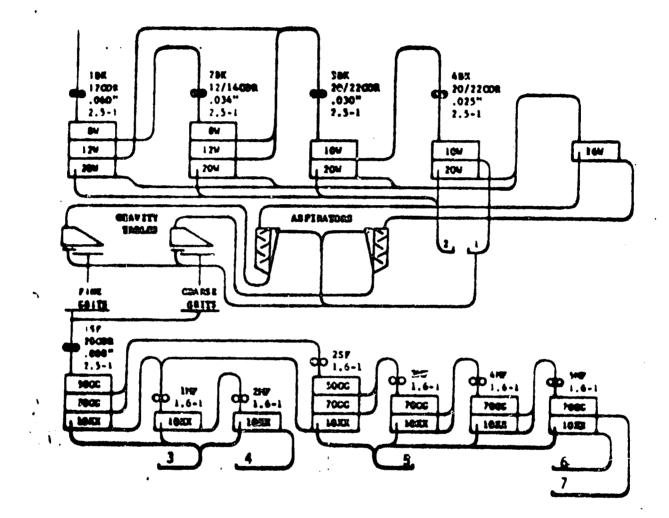
The 2.5 is the assumed PER for a reference standard casein diet. The various experimental diets were corrected by the fraction obtained from their respective control diets having similar protein content, i.e. diets 1, 2, 5 and 6 were corrected by the use of control diet 9; diets 3, 4, 7 and 8 were corrected by the use of control diet 10.

When the diets were not supplemented the floury endosperm was superior to the horny endosperm within a given hybrid. When the floury and horny fractions were supplemented with amino acids, however, differences were not present indicating the protein was utilized equally when the amino acid deficit was supplied. Table VII also shows no differences between the PER's obtained from the two hybrids irrespective of amino acid supplementation.

#### SUMMARY

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The data from this study substantiates differences between floury and horny endosperm. The protein quality of floury endosperm appears to be superior to that of horny endosperm when fed on an isonitrogenous basis. Differences between floury and horny fractions, however, were not present when both were supplemented with sufficient lysine and methionine to meet adjusted NRC requirements. When lysine and methionine is incorporated into the diets containing floury and horny endosperm, gain and PER's are high and comparable to those obtained with casein. Significant differences were not observed between fractions obtained from the two hybrids used in this study.



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	YI	eld <sup>3</sup>		ude itein		ude at		rude iber	A	sh	
	A	8	A	8	A	8	A	8		B	
										· · · ·	
	27.33	25.32	*								
	18. <b>88</b>	<b>19.6</b> 2									
	12.54	12.23	7.40	6.82	0.91	1.02	0.91	0.80	0.48	0.47	
	5.06	5.06	9.49	10.62	0.98	0.94	1.13	1.36	0.50	0.51	
	9.95	10.31	9.88	8.78	0 <b>.96</b>	1.09	0. <b>70</b>	1.13	0.50	0.47	
	8. <b>38</b>	8.19	13.98	12.58	0.87	1.12	0.72	1.12	0.47	0.46	
	14.29	16.61	15.96	15.47	1.12	1.23	1.12	1.34	0. <b>59</b>	0.55	
	9.95 8.38	10.31 8.19	9.88 13.98	8.78 12.58	0. <b>96</b> 0.87	1.09 1.12	0.70 0.72	1.13 1.12	0.50 0.47	)	0.47

TABLE I ANAL"SIS OF FRACTIONS OBTAINED FROM DRY MILLING SORGHUM GRAIN

<sup>1</sup>Moisture free basis.

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<sup>2</sup>Hybrid A is Paymaster Kiowa. Hybrid B is Frontier 400C

<sup>3</sup>Percent of whole sorghum grain

\*Blank spaces indicate no determinations made.

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Fraction		3		4		5	ł	6		7	
ibbrid <sup>1</sup>	A	8	<b>A</b>	t	A	8	A	<b>B</b> .		В	Casein
Protein <sup>2</sup>	7.40	6.82	9.49	10.62	9.88	8.78	13.98	12.58	15.96	15.47	94.31
Lysine <sup>3</sup>	1.72	1.80	- 1.33	1.35	1 30	1.54	0 00	1.10	1 05		
Histidine	1.98	2.10	2.03	2.16	1.96		0.99	1.15	1.05	1.16	8.98
Amonta	2.95	2.75	3.08	3.38	2.76	2.25	1.81	2.18	1.93	2.05	3.21
Arginine	2.85	3.01	2.61	2.71		3.23	2.94	3.16	3.15	2.90	1.86
Aspartic acid	6.13	6.15	5.93	6.23	2.30	3.07	2.15	2.54	2.27	2.63	4.09
Aspereic ecia	0.13	0.13	J. <b>JJ</b>	0.23	6.33	6.32	6.43	5.97	6.39	7.38	8.04
Threonine	3.09	3.16	3.14	3.26	3.16	3.38	3.00	3.22	3.03	3.34	4.72
Serine	4.45	4.31	4.54	4.92	4.66	4.78	4.61	4.70	4.63	4.91	6.36
Glutamic acid	21.73	20.67	23.67	24.85	24.93	25.11	25.24	25.68	25.59	27.05	
Proline	8.06	8.17	8.93	9.20	9.03	9.28	9.47	9.42			27.15
Glycine	2.61	2.79	2. <b>50</b>	2.56	2.36	2.75	2.15	2.40	<b>8.86</b> 2.18	10.87 2. <b>56</b>	12.63
Alanine	9.34	8.92	10.41	10.81	10.66	10.82	11.21	11.53	10.95	11.42	
Cystine	1.77	2.13				10.0L		11.33	1.47		3.31
Valine	4.28	4.86	4.38	4.67	4.81			4.87		1.77	0.53
Nethionine	1.62	1.72		••••/	4.01			4.0/	5.05	5.39	7.00
Isoleucine	3.98	3.83	3.99	4.12	4.27	<b>A</b> 10	A 99	A 06	1.38	1.56	2.19
	3.30	J. 04	3.33	7.12	4.27	4.18	4.22	4.26	4.30	4.45	5.80
Leucine	13.63	12.95	15.53	16.53	16.02	16.47	16.29	16.32	16.40	16.83	10.48
Tyrosine	4.08	4.02	4.15	4.41	4.43	4.45	4.58	4.44	4.50	4.53	
Phenylalanine	5.13	5.08	5.30	5.46	5.79	5.40	5.82	5.94	6.01	4.53 6.12	6.33 5.64
I-Recovery	86.26	85.31	92.28	97.51	92 46	95.94	91.68	97.04	94.42	99.76	103.44

Tuble le Amino Acid Distribution in Protein of Sorghum Grain Hilled Fractions and Casein

<sup>1</sup>Hybrid A is Paymaster Kiowa; Hybrid B is Frontier 400 C

<sup>2</sup>Protein, S (N x 6.25) moisture free basis.

Grams of amino acid per 16 grams nitrogen, duplicate determinations on samples 3A, 3B, 7A and 7B.

Hybrid	Milling Fraction	Amino Acid Supplement	Diet No.	Moisture	Protein	Fat	Ash
	3	-	1	31.2	5.6	4.2	3.5
P <b>ayma</b> ster K		+	2	11.3	5.8	4.3	3.4
	7	-	3	10.6	9.8	4.3	3.4
	*****	+	4	10.7	30.4	4.3	3.4
	3	-	5	10.7	5.4	4.3	3.4
Frontier 40		<b>*</b>	6	10.8	5.8	4.4	3.4
	7	-	7	10.5	10.0	4.2	3.4
	·	+	8	10.8	ì0.7	4.0	3.5
Casein		-	9	10.0	5.8	4.0	3.2
		-	۲,۲	9.9	10.0	4.0	3.2

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TABLE II APPROXIMATE ANALYSIS OF EXPERIMENTAL DIETS

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Diets	1	2	3	4	5	6	7	8	<u> </u>	10
Ingredient	s: X	r	x	x	1	ĩ	1	1	٤	z
Milled Fraction	85.0 <b>8</b>	85.08	67.98	67. <b>98</b>	92.26	92.26	70.49	70 <b>. 49</b>		•
Starch	7.18	7.18	23.03	23. <b>03</b>			<b>20.8</b> 0	20.80	84.75	80.26
Corn 011	3.40	3.40	3.40	3.40	3.40	3.40	3.40	3.40	3.86	3.86
Water <sup>1</sup>			1.22	1.22			0.97	0.97	0.97	0.87
¥itamin Premix <sup>2</sup>	0.63	0. <b>63</b>	0.63	0.63	0.63	0.63	0.63	0.63	0.63	0.63
Mineral Premix <sup>2</sup>	3.71	3.71	3.71	3.71	3.71	3.71	3.71	3.71	3.71	3.71
Lysine		0.33		0.64		0.33		0.64		
Methionine		0.12		0.25		0.12		0.25		
Casein									6.08	10.66

TABLE III COMPOSITION OF EXPERIMENTAL DIETS

Water was added to enjust moisture content of diet.

<sup>2</sup>Premix composition given in Table IV.

		En entre Diers	
	INGREDIENT	1000 g. Diet	
Mineral	Premix:	g.	
	Dicalcium phosphate <sup>1</sup>	30.00	
	Salt	3.00	
	Trace minerals <sup>2</sup>	0.50	
	Potassium chloride	3.05	
	Magnesium sulfate	1.77	
litamin	Premix	mg.	
	Vitamin A (30.0 IU per mg.)	66.70	
	vitamin D (15.0 IU per mg.)	133.00	8
	Alpha tocopherol (110.1 U per.g.)	544 90	
	Manadiona	) 10	
	Thiamine HC1	1.25	
	Riboflavin	2.50	
	Pyridoxine HCl	1 20	
1	Niacin	15 00	
	Calcium pantothenate	8.00	
1	Vitamin B <sub>12</sub>	5.00	
(	Choline chloride	750.00	
(	Carrier (starch)	5000.00	

TABLE IV COMPOSITION OF PREMIXES FOR EXPERIMENTAL DIETS

<sup>1</sup>Sargents Calcium Company, Das Moines, Iowa: P, 18.5%; Ca, 19.0-22.5%.

<sup>2</sup>CCC trace mineral mix contained: (ppm) Mn, 10; Fe, 10; Ca, 14; Cu, 1; Zn, 5; I, 0.3; and Co, 0.1; Calcium Carbonate Company, Quincy, Illinois.

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Hybrid <sup>1</sup>	Amino Acíd Supplement	Diet <sup>2</sup>	Gain <sup>3</sup>	Feed Consumed <sup>4</sup>	PER <sup>5</sup>
 A		. ]	5.50 <sup>ª</sup>	193.5 <sup>8</sup>	0.49 <sup>8</sup>
B	-	5	8.83 <sup>a</sup>	210.3 <sup>ab</sup>	0. <b>76</b> ª
Α	. +	. 2	35.50 <sup>b</sup>	242.0 <sup>cb</sup>	2.54 <sup>t</sup>
8	+	6	36.67 <sup>b</sup>	257.0 <sup>C</sup>	2.44 <sup>b</sup>
Casein	-	9	37.00 <sup>5</sup>	259.3 <sup>c</sup>	2.46 <sup>b</sup>

TABLE \	V	AVERA	GE V	<b>IE</b> IGHT	GAI	N, FEE	D CONSUMPTION,	AND
		PER'S	FOR	RATS	FED	FLOURY	ENDOSPERM	

<sup>1</sup>Hybrid A is Paymaster Kiowa. Hybrid B is Frontier 400C.

<sup>2</sup>Diet contains approximately 5.6% protein.

<sup>3</sup>Average weight gain per rat 0-4 weeks,  $LSD_{.01} = 8.22$  g.

 $^{4}$ Gm feed consumed 0-4 weeks, LSD<sub>.01</sub> = 46.12 g.

<sup>5</sup>Gm of gain/gm of protein consumed, LSD  $_{.01} = 0.38$ .

Hybrid <sup>1</sup>	Amino Acid Supplement	Diet <sup>2</sup>	Gain <sup>3</sup>	Feed 4 Consumed	PER <sup>5</sup>
A .	-	3	2.50 <sup>ª</sup>	168.7ª	0.15 <sup>a</sup>
8	-	7	4.50 <sup>a</sup>	176.7ª	0.25 <sup>a</sup>
A	+ •	4	75.70 <sup>C</sup>	342.5 <sup>C</sup>	2.13 <sup>b</sup>
8	+	8	91.67 <sup>d</sup>	<b>366</b> .0 <sup>C</sup>	2.34 <sup>C</sup>
Casein	-	10	62.17 <sup>b</sup>	272.0 <sup>b</sup>	2.29 <sup>bc</sup>

IARTE AI	AVERAGE WEIGHT	GAIN, FEED CONSUMPTION, AND	
	PER'S FOR RATS	FED HORNY ENDOSPERM	

Hybrid A is Paymaster Kiowa. Hybrid B is Frontier 400C.

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Diets contain approximately 10.2% protein.

<sup>3</sup>Average weight gain per rat 0-4 weeks, LSD = 11.5, a

<sup>4</sup>Gm of feed consumed 0-4 weeks, LSD = 42.08 g.

<sup>5</sup>Gm of gain/gm of protein consumed,  $LSD_{.01} = 0.20$ 

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Nilled Fraction	Amino Acia Supplement	Hybrid <sup>2</sup>	Diet	PER <sup>3</sup>
Floury	. •	A	1	0.50 <b>ab</b>
· · · • • • • • • • • • • • • • • • • •	**	B	5	0.78ª
	-	A	3	0.16 <sup>C</sup>
Horny		<b>B</b>	7	0.27 <sup>cb</sup>
	+	A	2	2.58 <sup>d</sup>
Floury	+	В	6	2.49 <sup>d</sup>
	◆	· A	4	2.32 <sup>d</sup>
lorny	<b>+</b>	В	8	2 55 <sup>d</sup>

TABLE VII	PER'S (	OF EXPE	ERIMENTA	DIETS	ADJUSTED	BY	THE
	1	CASEIN	CONTROL	DIETS			

<sup>1</sup>Correction factor for the low protein, fleury endosperm diets 1, 2, 5 and 6 is equal to:

(2.5 constant PER for casein) (2.458 PER of reference std. casein, diet 9)

Correction factor for the high protein, horny endosperm diets 3, 4, 7 and 8 is equal to:

(2.291 PER of reference std. casein, Diet 10)

2 Hybrid A is Paymaster Kiowa. Hybrid B is Frontier 400C.

 $^{3}$ Gm of gain/gm of protein consumed, corrected by the control dists; numbers having different letters are significantly different. LSD<sub>.01</sub> = 0.30.

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2. Effect of Cooking on Nutritive Quality of Common Native Foods.

A number of reports are available on the effect of heat on tie-up or destruction of amino acids in pure proteins, grains, and proteincarbohydrate mixtures. There are few studies on effect of heat on nutritive value of food products.

Studies will be made to extend the knowledge of effects of normal cooking of chapatis, cous-cous, and Arab bread on nutritive value. A preliminary study was made two years ago on the effect of toasting bread on tie-up of lysine. Toasting to a dark brown color markedly lowered growth of rats, but addition of lysine almost fully overcame the adverse effect.

Cooking can have both a favorable as well as unfavorable effect. In addition to the well-known effect on starch granules, there is some evidence that heat may make phosphorus of grains more available. Heat may destroy factors such as the anti-trypsin factor and those causing Lathyrism or Odoratism. Since most literature reports on these effects were done years ago, and often crudely, more needs to be known of the possible hazards from these factors in chick peas, a likely source of protein supplement for chapatis. Cooking also could tie up or destroy lysine and other supplements added to chapatis. This has not been investigated.

There are chemical tests for destruction or freeing of nutrients, such as solubility for phosphorus and FDNB method for lysine. In some

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cases comparisons of the animal and chemical methods need to be made, since results are not the same, and they could give clues to nutrient availability to animals.

3. The nutrition group has a major responsibility to evaluate and decide some questions relative to the project, which are not primarily based on our laboratory work directly. We are at work on these problems and will continue discussing them until conclusions or at least working assumptions can be made.

Some of these are:

- A. Can improvement of nutrition of adults and children, with the foods to be employed be considered as one or separate problems?
- B. To what extent will the nutritional plans need to take into account parasitism, other disease, and pre-existing nutritional status of subjects?
- C. What size individual in the population are we planning for (i.e. 55-60 kg man, or larger or smaller)? Should we plan for minimal or optimal nutrient intake? What level of energy, type and level of protein and levels of vitamins and minerals will become our working standard?
- D. What would be the most satisfactory supplement to grain-pure nutrients or natural foodstuffs.