

# Pigeonpea Nutrition and Its Improvement

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**SUMMARY.** Pigeonpea (*Cajanus cajan* [L.] Millsp.), known by several vernacular and trade names such as red gram, tuar, Angola pea, Congo pea, yellow dhal and oil dhal, is one of the major grain legume crops of the tropics and sub-tropics. It is a favorite crop of small holder dryland farmers because it can grow well under subsistence level of agriculture and provides nutritive food, fodder, and fuel wood. It also improves soil by fixing atmospheric nitrogen. India by far is the largest pigeonpea producer where it is consumed as decorticated split peas, popularly called as 'dhal.' In other countries, its consumption as whole dry seed and green vegetable is popular. Its foliage is used as fodder and milling by-products form an excellent feed for domestic animals. Pigeonpea seeds contain about 20-22% protein and appreciable amounts of essential amino acids and minerals. Dehulling and boiling treatments of seeds get rid of the most antinutritional factors such as tannins and enzyme inhibitors. Seed storage causes considerable losses in the quality of this legume. The seed protein of pigeonpea has been successfully enhanced by breeding from 20-22% to 28-30%. Such lines also agronomically performed well and have acceptable seed size and color. The high-protein lines were found

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nutritionally superior to the cultivars because they would provide more quantities of utilizable protein and sulfur-containing amino acids. [Article copies available for a fee from The Haworth Document Delivery Service: 1-800-HAWORTH. E-mail address: <getinfo@haworthpressinc.com> Website: <<http://www.HaworthPress.com>> © 2002 by The Haworth Press, Inc. All rights reserved.]

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## INTRODUCTION

In Asia and Africa population growth is the prime development constraint. Recently, India has crossed the alarming 'one billion' population mark and it is catching up fast the most populous country China. Providing significant quantity of quality food to the growing population with limited resources is a big challenge. Due to increase in rural family size, where the population growth is over 2% per year, the land holdings are dividing by each passing generation leading to increased pressure on unit land and reduction in the capacity of farmers to produce sufficient quantity of nutritive food for their families. In producing food crops, such farmers try to keep a balance between cereals and legumes, besides aiming to obtain some fodder for cattle. However, their small holdings and lack of basic resources like water, fertilizer, and pesticides restrict the food production. In general, the food production from small holdings is short both in quantity and quality and under such circumstances, the expectant and nursing mothers and young children are the most vulnerable lots. Recently, a number of weaning and supplementary foods have come in market but due to high prices they remain a luxury of urban middle and upper classes. Since the animal protein is also out of their reach, the problem of malnutrition among poverty-ridden masses is achieving a serious dimension in the country. To meet this challenge, a concerted effort is needed to increase the production of protein-rich legume crops which can be grown under subsistence level of farming and no crop other than pigeonpea suits most because it is drought tolerant, need minimum inputs, and can produce reasonable amounts of food, fodder, and fuel wood. Pigeonpea seeds contain about 20-22% protein and reasonable amounts of essential amino acids. India is the largest producer of pigeonpea with annual production of three million tons harvested from about four million hectares. In the past 10

years the area under pigeonpea is consistently increasing at the rate of 2% each per year (Ryan, 1997) but still its demand is out-scoring the supply and serious efforts are needed at every level to boost the production of this important legume. In India pigeonpea is predominantly consumed as *dhal* (decorticated dry split peas) but its whole seeds are also consumed in Africa and the Caribbean islands. In this chapter various aspects of pigeonpea nutrition are reviewed.

## GRAIN QUALITY OF PIGEONPEA

Pal (1939) published perhaps the first review on the nutrition value of pulses in India. He compared different pulse crops for their digestibility coefficient, biological value, net protein value, and four essential amino acids. For biological value he judged pigeonpea as the best pulse crop and concluded that it makes the most nutritive food when eaten with rice. However, using an arbitrary scale for the overall nutrition value of the pulses, chickpea (*Cicer arietinum*) and black gram (*Phaseolus mungo*) were considered superior to pigeonpea. The nutritional value of food is determined by its chemical constituents and in pigeonpea a wide range is reported for these vital elements (Tripathi et al., 1975; Sharma et al., 1977; Narsimha and Desikachar, 1978; Manimekalai et al., 1979; Singh et al., 1984 a & b). Besides inherent genotypic differences, such variation can be attributed to environment where the crop was grown, methods of sampling and analyses, and method and length of seed storage periods.

### *Chemical Composition of Dry Seeds*

The distribution of some dietary nutrients in different parts of dry pigeonpea seed as reported by Faris and Singh (1990) is given in Table 1. Broadly, pigeonpea seed contains 85% cotyledons, 14% seed coat, and less than 1% embryo. Carbohydrates and proteins are major constituents of cotyledons, embryo, and seed coat. Quantitatively, the cotyledons (66.7%) and seed coat (58.7%) are rich in carbohydrates while protein (49.6%) constitutes a major portion of embryo. Carbohydrates and fat are also present in significant quantities in embryo. About one-third of seed coat is made of fibers. The seed also contains amino acids, calcium, fiber, and iron. The contents of methionine and cystine, the sulfur-containing amino acids, range around 1% and they predominantly reside in cotyledons and embryo. Calcium is predominantly

TABLE 1. Distribution of nutrients in mature pigeonpea seed.

Constituent	Whole seed	Cotyledons	Embryo	Seed coat
Carbohydrates (%)	64.2	66.7	31.0	58.7
Protein (%)	20.5	22.2	49.6	4.9
Fat (%)	3.8	4.4	13.5	0.3
Fiber (%)	5.0	0.4	1.4	31.9
Ash (%)	4.2	4.2	6.0	3.5
Lysine <sup>1</sup>	6.8	7.1	7.0	3.9
Threonine <sup>1</sup>	3.8	4.3	4.7	2.5
Methionine <sup>1</sup>	1.0	1.2	1.4	0.7
Cystine <sup>1</sup>	1.2	1.3	1.7	-
Calcium <sup>2</sup>	296	176	400	917
Iron <sup>2</sup>	6.7	6.1	13.0	9.5
Thiamine <sup>2</sup>	0.63	0.40	-	-
Riboflavin <sup>2</sup>	0.16	0.25	-	-
Niacin <sup>2</sup>	3.1	2.2	-	-

Adapted from Faris and Singh (1990)

1: g 100<sup>-1</sup> g protein

2: mg 100<sup>-1</sup> g dry matter

found in seed coat and embryo. Singh and Jambunathan (1982) studied distribution of major protein fractions in different components of pigeonpea seed and found that globulins constitute about 65% of total protein (Table 2). In comparison to other protein fractions globulin is inferior in sulfur-containing amino acids. Albumin, though in relatively small quantity, is rich in sulfur amino acids. The portion of prolamin in seed is low. According to Eggum and Beames (1983) pigeonpea is rated inferior to most other legumes as far as sulfur-containing amino acids is concerned but, unlike other legumes, the high protein content of pigeonpea seed is not tightly linked to its low methionine content (Singh and Eggum, 1984). Nigam and Giri (1961) observed that stachyose and verbascose constitute major component among sugars of pigeonpea. The pigeonpea starch has been found to be stable to heat up to 90°C (Modi and Kulkarni, 1976).

TABLE 2. Major protein fractions (%) in dry pigeonpea seed.

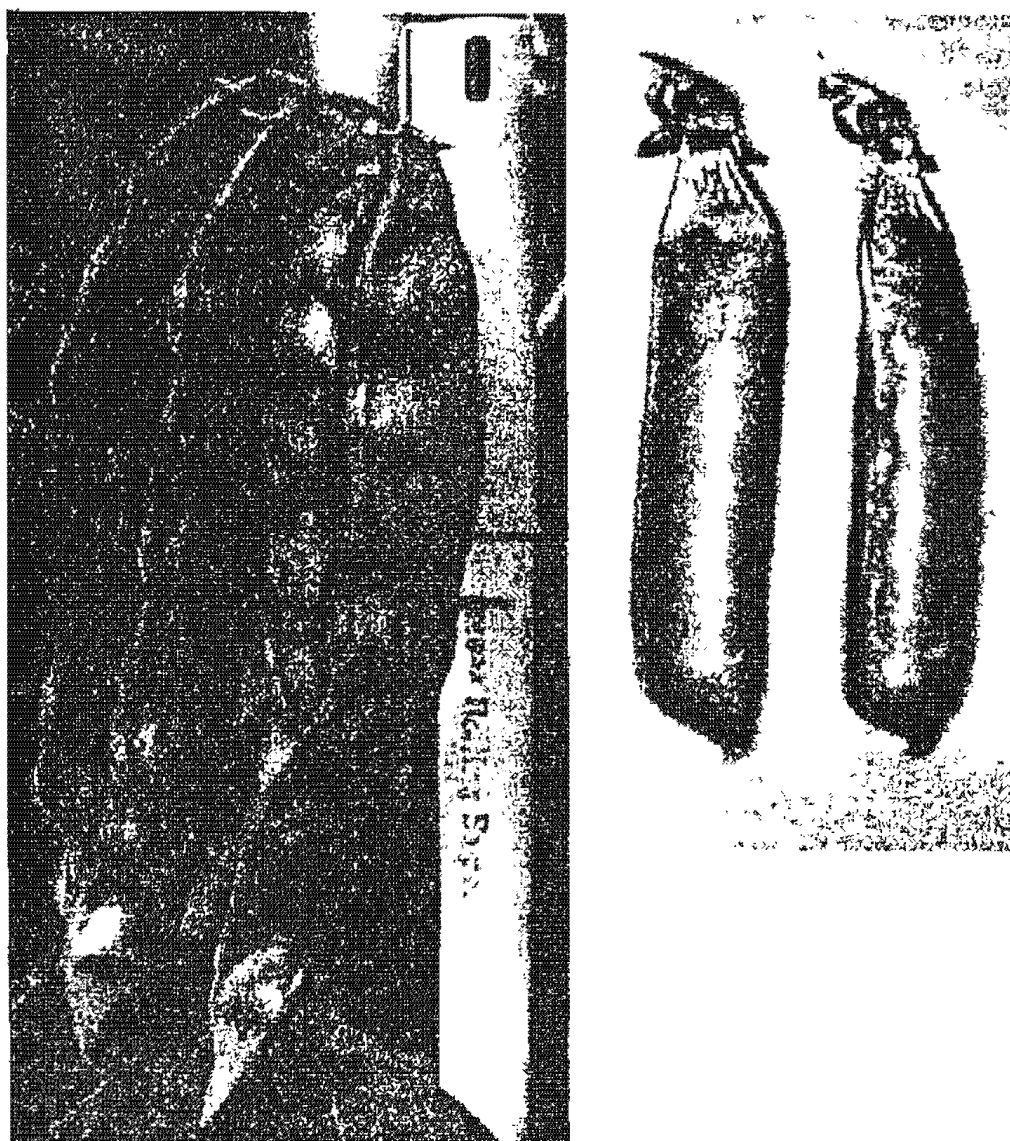
Component	Albumin	Globulin	Glutelin	Prolamin
Whole seed	10.2	59.9	17.4	3.0
Embryo	17.0	52.7	21.3	2.7
Cotyledons	11.4	64.5	18.2	3.5
Seed coat	2.6	26.3	32.8	4.2

Adapted from Singh and Jambunathan (1982)

### *Chemical Composition of Immature Seeds*

Physiologically mature green seeds of pigeonpea are consumed as vegetable in Dominican Republic, the Caribbean islands and some parts of India. In Dominican Republic 80% of the crop is exported as canned or frozen vegetable. Singh et al. (1977) compared vegetable pigeonpea (Figure 1) with that of pea (*Pisum sativum*) and found that pea seed had higher protein than pigeonpea but it was similar in crude fibre content. The trypsin inhibitors were more in pigeonpea when compared to pea but far less when compared with soybean. Nutritionally, green pigeonpea seed is considered superior to the *dhal* (Table 3). According to Faris et al. (1987) green pigeonpea seed is a rich source of iron, calcium, and magnesium when compared with its *dhal*. The green seed contains lower quantities of trypsin and amylase inhibitors and flatulence-causing sugars. The green seed cooks quickly and is also a better source of vitamin A. The protein and starch digestibility of green seed are higher than mature seed (Singh et al., 1984a). Singh et al. (1991) while studying chemical changes in the developing pigeonpea seeds, found that dry matter accumulation increased up to 28-32 days after flowering in different cultivars and recommended that for the best green pea yield, the crop should be harvested at nearly 30 days after flowering. The protein content and soluble sugars showed a gradual decrease with advancing maturation of seeds while its starch continued to increase. ICP 7035, a known vegetable type Indian pigeonpea landrace, was found to be more biochemically active in accumulating soluble sugars. This landrace was marginally low in calcium and magnesium at all the stages of seed development. The iron content of seeds also decreased as they approached maturity. In pigeonpea a range of pod color is found but it is not related to any quality parameter (Saxena et al., 1983).

FIGURE 1. Vegetable pigeonpea (left) and green pea pods (right).



### *Antinutritional Factors*

Like most legumes, pigeonpea also contains certain amounts of antinutritional factors. These include oligosaccharides (raffinose, stachyose, verbascose), polyphenols (phenols, tannins), phytolectins, and enzyme (trypsin, chymotrypsin, amylase) inhibitors. Singh (1988) studied a number of genotypes for quantifying important antinutritional factors and toxic substances present in pigeonpea seeds and found a large variation among genotypes for these traits (Table 4). Amylase and trypsin inhibitors and phenols were found in significant quantities. In addition, the flatulence causing sugars were also present in appreciable quantities. According to Kamath and Belavady (1980) pigeonpea seeds also contain certain unavailable carbohydrates which characteristically reduce the bioavailability of important nutrients.

TABLE 3. Comparison of pigeonpea green seed and *dhal* for various nutritional constituents.

Constituent	<i>Dhal</i>	Green seed
Starch content (%)	57.6	48.4
Protein (%)	24.6	21.0
Protein digestibility (%)	60.5	66.8
Trypsin inhibitor (unit mg <sup>-1</sup> )	13.5	2.8
Soluble sugars (%)	5.2	5.1
Crude fiber (%)	1.2	8.2
Fat (%)	1.6	2.3
Calcium <sup>1</sup>	16.3	94.6
Magnesium <sup>1</sup>	78.9	113.7
Copper <sup>1</sup>	1.3	1.4
Iron <sup>1</sup>	2.9	4.6
Zinc <sup>1</sup>	3.0	2.5

Adapted from Paris et al. (1987)

1 = mg 100 g<sup>-1</sup> sample

TABLE 4. Genotypic variation for major antinutritional factors in pigeonpea.

Factor	Genotypes	Range
Total phenols (mg g <sup>-1</sup> )	14	3.0-18.3
Tannins (mg g <sup>-1</sup> )	10	0.0-0.2
Trypsin inhibitor (units mg <sup>-1</sup> )	9	8.1-12.1
Chymotrypsin inhibitor (units mg <sup>-1</sup> )	9	2.1-3.6
Amylase inhibitor (units g <sup>-1</sup> )	9	22.5-34.2
Raffinose (g 100 g <sup>-1</sup> )	10	0.24-1.05
Stachyose (g 100 g <sup>-1</sup> )	9	0.35-0.86
Stachyose + verbascose (g 100 g <sup>-1</sup> )	4	1.60-2.30

Adapted from Singh (1988)

Godbole et al. (1994) reported the presence of protease inhibitor in seven-day old seed of variety TAT-10 while Ambekar et al. (1996) found that the inhibitors are either not synthesized or inactive up to 28 days of the seed development. No other plant part except seed recorded the presence of trypsin and chymotrypsin inhibitors (Mutimani and Paramjyothi, 1995). Since seed coat is rich in antinutritional factor it assumes greater importance where whole pigeonpea seeds are consumed. In many African, South American and Caribbean countries where dehulling facilities are not available and whole seed is consumed, predominantly white seeded cultivars are grown which contain relatively less quantity of polyphenols. Singh (1984) compared white, brown, and light brown seeded pigeonpea types for antinutritional factors (Table 5) and established strong relationship between seed coat color and antinutritional factors. He found three times greater quantity of polyphenols in the red seeded lines in comparison to the white seeded types. Similarly, the enzyme inhibition activity was much larger in the colored pigeonpea. In India almost entire pigeonpea production is converted into *dhal*. In this process the seed coat is removed and therefore the large amounts of tannins present in the colored pigeonpea pose no problem in its consumption. The amounts of polyphenols in *dhal* made

TABLE 5. Polyphenol contents and varietal differences in the enzyme-inhibitory property of pigeonpea polyphenols.

Cultivar	Testa color	Polyphenols (mg g <sup>-1</sup> sample)	Enzyme inhibition <sup>1</sup> (%)			
			Trypsin	Chymo- trypsin	Human saliva	Hog pancreas
Hy 3C	White	3.7	37.9	36.0	34.5	21.8
NP (WR) 15	White	6.0	40.5	38.6	32.7	19.7
C 11	Light brown	14.2	91.5	90.3	86.0	80.9
BDN 1	Brown	15.2	90.3	91.6	79.4	69.3
No. 148	Brown	14.9	88.0	85.9	75.8	68.5
Mean		10.8	69.7	68.5	61.7	52.0
SE		±0.2	±2.1	±1.7	±1.4	±1.3

<sup>1</sup> Based on assay using 200 mg polyphenols for trypsin and chymotrypsin, and 250 µg polyphenols for amylase inhibitions.

Adapted from Singh (1984)



either from red or white grain were found to be similar and ranged between 1.4 to 1.9 mg g<sup>-1</sup> samples. Pichare and Kachole (1994) found no association between trypsin inhibitor activity and pod borer resistance in pigeonpea.

## HUMAN NUTRITION

### *Role of Pigeonpea in Rural Diets*

Methionine and cystine followed by tryptophan and threonine are the limiting essential amino acids in pigeonpea whereas lysine is the limiting amino acid in rice and wheat (Faris and Singh, 1990). A food combining cereals and pulses provide a balanced diet because they complement the amino acid profiles of each other. According to Hulse (1977), the mutual compensation is closest to the ideal value when the ratio by weight of cereals to legume is roughly 70:30. In southern and eastern Africa this ratio is 90:10 reflecting shortage of legume protein in the diet. Daniel et al. (1970) studied supplementation of cereal diets with various proportions of pigeonpea in rats and reported that supplementation of ration with pigeonpea significantly enhanced the nutritive value of the diet. Supplementation of rice diet with 8.5% and 16.7% pigeonpea *dhal* markedly improved the quality of diet (Table 6). Similarly, Kurien et al. (1971) demonstrated that a supplement of pigeonpea in maize diet significantly improved the quality of food.

TABLE 6. Effect of supplementary rice diets with varying levels of pigeonpea on the growth of young rats.<sup>1</sup>

Diet	Protein (%)	Gain in mass (%)	Protein intake (%)	Protein efficiency ratio
Rice	7.2	25.5	11.8	1.78
Rice + 8.5% pigeonpea	8.7	32.8	15.5	2.13
Rice + 16.7% pigeonpea	10.0	45.2	19.6	2.32
Rice + 25.0% pigeonpea	11.4	48.9	21.8	2.25

<sup>1</sup> Based on an experimental period of 4 weeks.  
Adapted from Daniel et al. (1970)

Bidinger and Nag (1981) conducted a village level study including 240 families of different resource groups, representing six villages located in three agro-climatic zones of India. They observed that pigeonpea was by far the most preferred pulse crop and its consumption patterns differed widely by age group, farm size, and the village. The consumption rate was found linear with small farmers consuming the least amount and the large farmers the most. The consumption of pigeonpea was also found to be related to the production. National Institute of Nutrition in India recommends cereal:pulse ratio of 3:1 for very young children, 5:1 for women, and 6:1 for men. In most cases in villages, these standards could not be met (Table 7). Bidinger and Nag (1981) also reported that 10% of the protein and 5% of energy in the village diets came from pigeonpea. The maximum lysine provided from the diet was 21.7%. These values are low and reflect the low consumption of legumes. Prema and Kurup (1973) reported that pigeonpea contains cholesterol and phospholipid lowering effect. Globulin fraction of pigeonpea protein was found to have a significant hypolipidaemic action in rats fed with a high-fat and high-cholesterol diet. They reported marked reduction in the total and free cholesterol, phospholipids, and triglycerides contents in serum, liver, and aorta tissues of rat.

### *Nutrition Losses in Dehulling*

In the Indian subcontinent pigeonpea is predominantly consumed in the form of *dhal* and conversion whole seed into *dhal* is a big industry in

TABLE 7. Relative cereal:pulse consumption by different age groups in six villages of central India.

Village	Age group		
	1 to 6	7 to 18	Adults
Aurepalle	31:1	35:1	37:1
Dokur	23:1	31:1	42:1
Shirapur	15:1	14:1	17:1
Kalman	14:1	18:1	20:1
Kanzara	7:1	9:1	10:1
Kinkheda	9:1	10:1	10:1

Adapted from Bidinger and Nag (1981)

the country. For commercial purposes, big machines are used for dehulling while in rural areas, dehulling is done by using traditional grinding stones called *chakki* or quern. Since the cotyledons of pigeonpea are attached tightly with seed coat by gums, the processing primarily involves loosening of husk followed by dehusking and splitting of the two cotyledons. Therefore, pigeonpea dehulling is not only difficult but also a specialized function when compared with other legumes. Losses of seed mass during the process of dehulling is a common event. Excluding the husk which accounts for about 15%, the *dhal* recovery in pigeonpea is around 60% by *chakki* and around 70% by machines (Singh and Jambunathan, 1981). This means even by using advanced technology about 15-17% of grain mass is lost. By using *chakki* such losses shoot up to 20-25%.

Reddy et al. (1979) studied the protein deposition pattern in pigeonpea seed and reported that the outer layers of the cotyledons are richer in protein in comparison to inner layers of seed. From nutrition point of view, this is a matter of concern since dehulling not only removes protein-rich germ but also the outer layers of the cotyledons where relatively more protein constituents are housed. Fortunately, the protein quality in terms of amino acids is not adversely affected by dehulling. Singh and Jambunathan (1990) further reported that dehulling also removes about 20% calcium and 30% iron. To preserve the nutritive value of pigeonpea seed and minimizing the nutrient losses during dehulling it is essential that more efficient dehulling technology is developed and transferred to rural areas where by and large milling is still carried out by inefficient old-age techniques. According to Kurien (1981) under controlled conditions the *dhal* yield achieves the maximum efficiency of 80-84% but at commercial level the recovery remains around 70%. He also reported large varietal differences (72 to 82%) for *dhal* yield. Therefore, it can be assumed that with a combination of a superior variety and an efficient pigeonpea processing technology, the nutrient losses can be minimized.

### *Cooking Quality*

Pigeonpea seeds dry, green, or processed in the form of *dhal* and other products are consumed after cooking. Therefore, besides various nutritional aspects the cooking time and other related parameters assume significant importance. Consumers always prefer a *dhal* that cooks fast and produces more volume upon cooking with high consistency and flavor. Cooking time recorded between 22 and 44 minutes for

*dhal* and between 45 and 67 minutes for whole seed by Sharma et al. (1977) indicate the extent of genotypic variation present for this trait.

Cooking time of *dhal* was found to be independent of taste and flavor (Maninekalai et al., 1979). Jambunathan and Singh (1981) studied various physico-chemical characters in 25 pigeonpea cultivars and reported a considerable range (Table 8) for various quality parameters. The cooking time ranged between 24-68 minutes. They also found that quick cooking trait was associated with large seed size, high solid dispersal, water absorption, nitrogen solubility indices, and nitrogen content of the solids dispersed. Lines with high protein and small seeds in general take more time to cook. Narsimha and Desikachar (1978), Singh et al. (1984c), and Sharma et al. (1977) reported positive association of cooking time with its calcium and magnesium contents. The issue of pre-cooking, soaking, and cooking time needs to be resolved. In

TABLE 8. Variation for various physico-chemical characteristics in 25 pigeonpea cultivars.

Constituent	Range	Mean
Solids dispersed (%)	20.8-54.7	37.4
Water absorption (g g <sup>-1</sup> sample)		
<i>dhal</i>	1.69-2.65	2.25
whole grain	0.63-1.34	1.02
Increase in volume (v/v)		
<i>dhal</i>	1.18-1.86	1.51
whole grain	0.91-1.54	1.13
Starch (%)	51.5-63.4	58.6
Soluble sugars (%)	3.6-5.3	4.8
Nitrogen solubility index (%)	28.7-42.5	36.4
Nitrogen content in solids dispersed (%)	19.6-31.8	27.3
Protein (%)	19.7-25.2	22.1
100-seed mass (g)	6.2-20.7	9.6
Cooking time (min)	24-68	38

Adapted from Jambunathan and Singh (1981)

some experiments pre-soaking in water reduced cooking time (ICRISAT, 1987) while in others (Saxena et al., 1992) this treatment increased the cooking time significantly. Soaking in sodium bicarbonate helped in reducing cooking time in pigeonpea but it increased pH and thereby adversely affected the organoleptic quality of *dhal*.

According to Salunkhe (1982) cooking of pigeonpea improved the bioavailability of nutrients and also destroyed some antinutritional factors. Heat treatment also enhanced the starch digestibility. Lines, which take long time to cook also face the danger of losing vital vitamins from the food. Cooking of seed after germination not only enhances the digestibility of starch (Jyothi and Reddy, 1981) but also reduces the levels of oligosaccharides (Iyenger and Kulkarni, 1977). The fermentation of seeds also helps in reducing inhibitory activity of the digestive enzymes (Rajalakshmi and Vanaja, 1967). It appears that a little research has been conducted in the past in this critical aspect of pigeonpea quality. Studies in understanding the role of various chemical constituents on cooking time in diverse genetic materials will help in resolving this issue.

Geervani (1981) studied the effect of boiling, pressure-cooking, and roasting on the quality of pigeonpea and reported that thiamine and riboflavine were destroyed by heat but niacin content was unaltered during all the treatments. Availability of lysine and methionine decreased more on roasting but the available methionine increased on boiling and pressure-cooking.

### ***Quality Losses in Storage***

Throughout the world and particularly in India and Africa, pigeonpea is predominantly cultivated by small holder resource poor farmers for meeting their domestic protein needs and to generate some income. These farmers generally store the whole seeds for about 8-12 months for sowing and round-the-year consumption. They process small quantities of grain through hand-operated mills as and when needed. In rural areas, the seeds are generally stored in gunny bags or bins made of mud and husk and during the storage period a considerable damage is caused by storage pests. Among these, bruchid (*Callosabruchus* spp.) is the major pest. In most cases the ripening pigeonpea pods are infested on the plants in the field and this infestation is carried to the storage bins through seeds. Since the bruchids complete their life cycle in about four weeks, they multiply fast inside the bin and cause considerable seed damage. This damage not only reduces *dhal* recovery and deteriorates

germination but also adversely affects the hygiene and nutritive value of seeds (Parpia, 1973). In pigeonpea, although some genotypic variation for bruchid resistance is reported (Uma Reddy and Pushpamma, 1981) but these differences are inconsistent and are not large enough to ignore the issue of storage pests.

According to the standard set by Prevention of Food Adulteration Act of 1967, food grains containing more than 10 mg uric acid per 100 g of food, arising as a result of insect infestation, are unfit for human consumption. The pigeonpea seeds when stored for five months turned unfit for consumption as their total uric acid content crossed the prescribed limit of 200 mg per 100 g of sample (Daniel et al., 1977). Cooking time of food legumes in general increases with storage time and pigeonpea is no exception. Vimla and Pushpamma (1983) reported that by storing pigeonpea for eight months the safety level of uric acid was crossed. They also found that cooking time of both undamaged and damaged pigeonpea seeds increased significantly after storing them for about 12 months, indicating that even improved storage methods failed to retain the quality traits of stored seeds (Vimla and Pushpamma, 1985). Srivastava et al. (1988) reported that with the increase in insect infestation and the advancement of storage period the parameters such as seed moisture, total ash, crude fibre, protein, and reducing sugar contents increased while fat, carbohydrates, and non-reducing sugars decreased. Daniel et al. (1977) found that lysine, threonine, and protein efficiency ratios were significantly and adversely affected in pigeonpea when the seeds were stored in jute bags. Uma Reddy and Pushpamma (1981) reported significant reduction in the amino acid contents in the infested seed samples and the decline in lysine was greater than those of methionine and tryptophan. Daniel et al. (1977) observed significant decrease in the protein efficiency ratio due to storage. The storage of pigeonpea seeds also resulted in the loss of vitamins. Such losses were less (10-26%) in the protected seed and high (32-49%) in the unprotected infested seeds (Uma Reddy and Pushpamma, 1981). Thiamine and niacin contents also registered decline during storage. A number of factors have been identified which determine the extent of quality loss during storage. These include moisture, temperature, relative humidity, corneous thickness, hardness of grain, and ovipositional differences of storage pests (Squire, 1933; Singh et al., 1977). The storage losses can be reduced to some extent by improving the storage conditions but the decline in some quality parameters is inevitable. A well directed research is needed to provide pigeonpea farmers a cost-effective and efficient seed storing technology.

## ANIMAL NUTRITION

Pigeonpea is a wonderful plant because besides providing nutritious food for human beings it is a preferred animal fodder and feed also. Its fodder is relished by cattle, goats, and sheep while its harvest-trash, grain, and milling by-products form an excellent feed for various domestic animals.

### *Fodder*

The perennial nature of pigeonpea plant allows it to produce tender leaves shortly after cutting the plants during its vegetative growth period and also after the harvest of seed crop. The fodder yields of pure stand cuts depend on both genotype and management practices, which include height and frequency of cuttings, availability of soil moisture and nutrition. The genotypic differences for vegetative growth have also been observed at ICRISAT. The long-duration pigeonpea cultivars that are photo-thermal sensitive produce large biomass when planted around the longest day of the year. The same genotype if planted later in the reducing daylengths produces less quantity of biomass. The selection of a suitable cultivar and appropriate agronomic management practice can produce plenty of quality forage from this crop.

Singh and Kush (1981) in India, Herrera et al. (1966) in Columbia, Parbery (1967) in Australia, and Shiyong et al. (1999) in China have reported around  $50 \text{ t ha}^{-1}$  fodder yields in multiple cuttings from pigeonpea. The actual edible forage, however, is about 50% of the total yield because of the woody stem of the plant (Whiteman and Norton, 1981). Pigeonpea stands are also used for grazing purposes. In Hawaii the live-weight gain of over  $1,120 \text{ kg ha}^{-1} \text{ year}^{-1}$  have been reported by Krauss (1932). Whiteman and Norton (1981) concluded that pigeonpea forage was superior to grass in gain head<sup>-1</sup> indicating that the crop had a higher nutritive value and could carry a higher stocking rate than those of the grasses.

Generally, pigeonpea is used as forage for supplementing protein when the pasture quality is sub-standard. The young tender leaves and fresh flowers and pods form nutritive fodder for all grades of livestock. Leaves are the major forage component during the vegetative growth. As the plant approaches reproductive stage the fodder quality is enhanced due to the development of high-protein seeds. Therefore, the forage quality at a particular time will depend on the proportion of different plant parts. A proximate analysis of different pigeonpea plant

parts as reported by various workers is summarized in Table 9. According to Krauss (1921) the fresh green foliate contain 23.7% crude protein and 35.7% fibre and seed meal has 25.3% protein and 7.3% fiber. Henke et al. (1940) compared pigeonpea fodder with other better known forage crops and concluded that pigeonpea produced the highest economic value of digestible nutrients per unit area when compared with leucern (*Medicago sativa*) and other species.

Whiteman and Norton (1981) conducted sheep feeding trials using pigeonpea pods and pangola grass (*Digitaria decumbens*) in Australia. They concluded that pigeonpea pods with 7.5% crude protein fed as a sole diet were of low nutritive value and sheep lost 2% body weight. However, the inclusion of 33% of high quality forage such pangola grasses in the diet considerably improved the nutritive value of feed. They also pointed out that the harvest trash, which contains a significant proportion of leaves, would be a more nutritive feed than pods alone.

In China, pigeonpea is being promoted to meet the growing need of fresh quality fodder in the country because it can be grown well in the eroded soils of southern hilly regions for providing quality fodder under dry conditions. The ability of pigeonpea to allow 3-5 fodder cuttings within a year also makes it a useful stall-feeding crop. Pigeonpea being a perennial drought tolerant crop has shown high adaptation in a range of soil types of mountain regions of Du Au, Dahua, Huan Jiang, and Feng Shan counties of Guangxi province of China. According to Fuji

TABLE 9. Major nutrition constituents in different pigeonpea plant parts.

Component	Crude protein (%)	Crude fiber (%)	N-free extract (%)	Fat (%)	Ash (%)	Reference
Fresh green forage	23.7	35.7	26.3	5.3	8.7	Krauss (1921)
Whole tops, mature	18.8	29.4	40.0	5.2	5.6	Work (1946)
Whole tops, young	15.8	31.2	37.7	4.6	5.6	Work (1946)
Seed meal	25.3	7.3	61.2	1.7	4.1	Krauss (1921)
Mature dry seed	21.3	-	63.7	1.7	4.2	Morton (1976)
Pod meal	10.1	40.7	45.0	1.6	3.1	Krauss (1921)
Pods intact	7.0	-	42.8	0.4	5.7	Morton (1976)



and Zhanghong (1995) the foliage of pigeonpea is a quality fodder and goats, buffalo, cattle, and pig relish it.

A preliminary evaluation of ICRISAT pigeonpea varieties at Guangxi Academy of Agricultural Sciences in China showed that with multiple cuttings within a year variety ICPL 93047 produced 54 t ha<sup>-1</sup> of fresh and 29 t ha<sup>-1</sup> of dry fodder (Shiying et al., 1999). This experiment also showed that pigeonpea could grow well during winter and can meet the fodder needs when normal fodder supply is limited. It was observed that the goats and cattle liked the dry forage of pigeonpea better than green matter. S.C. Rao (Personal communication) compared tall and dwarf pigeonpea lines for forage production and their nutrition value in the southern plains of USA and reported that the dwarf genotype (PBNA) produced tender branches resulting in relatively less stem dry matter. In comparison to tall types (23 g kg<sup>-1</sup>) the dwarf line produced greater (28.6 g kg<sup>-1</sup>) nitrogen. The digestibility of the forage harvested from the dwarf line was also greater than the tall genotypes.

### *Feed*

Whole grain, threshing trash and milling by-products are used as feed for cattle, poultry, and pigs. These pigeonpea by-products provide protein-rich substitute for domestic animals at cheaper rates. In countries where climate is hot and dry and other legume crops are difficult to grow, pigeonpea is an attractive alternative. In the first quarter of 20th century pigeonpea was extensively used as poultry meal in Hawaii. According to Krauss (1932) an equal mixture of cracked pigeonpea and cracked maize seed was considered the best poultry ration. Draper (1944), Springhall et al. (1974), and Wallis et al. (1986) considered pigeonpea as an ideal protein substitute for all types of poultry rations. Since whole pigeonpea seeds contain some amount of antinutritional factors heat treatment of grains was introduced in the animal ration preparations. This resulted in a significant increase in the apparent metabolizable energy content of pigeonpea meal (Nowkolo and Oji, 1985). Wallis et al. (1986) reported little effect of heating on growth rates, feed intake, and in feed conversion efficiency. Falvey and Visitpanich (1980) conducted pig-feeding trials using 30% ground pigeonpea seed in Thailand. They reported live-weight gain from 25 g day<sup>-1</sup> to 159 g day<sup>-1</sup>. By using boiled pigeonpea the live-weight gain was further increased to 205 g day<sup>-1</sup>. This increase was attributed to the reduction of trypsin inhibitor activity that in turn improved the feed conversion ratio. The re-

cently bred high-protein pigeonpea lines at ICRISAT are likely to enhance the utility of pigeonpea in animal ration.

Use of pigeonpea seeds as feed is a common practice in rural China and it is primarily fed to pigs and chickens and some times to cattle and goats also. For pigs, the boiled seeds of pigeonpea are used to prepare feed mixtures with other ingredients while raw seeds are fed to chickens. In 1992, Research Institute of Resource Insects in China studied the nutritional value of pigeonpea feed. In this experiment pigs were fed with feed mixtures prepared with different concentrations of pigeonpea (Fuji et al., 1995). They found that a mixture with 6-12% pigeonpea in the meal mixture, the gain in the meat-mass production was  $78 \text{ g day}^{-1}$  with a ratio of meat-mass to feed input of 3.54:1 and this efficiency-mark meets the Chinese National Standards. Based on this information, Fuji et al. (1995) developed various feed mixtures using pigeonpea seed (22% protein) and dry leaf powder (19% protein) as major source of protein.

As mentioned earlier that in India over 3 million tons of pigeonpea is converted into *dhal* annually by processing either at household level by *chakki* or at commercial mills. This conversion of whole dry seeds into *dhal* yields significant quantity (about 30%) of the by-products. These include approximately 3-8% brokens, 15% powder, and 10% husk. The powder and brokens are important source of protein for cattle feed (Pathak, 1970). Whiteman and Norton (1981) evaluated non-seed material collected from machine harvester and reported that it contains 13.9% crude protein, 0.35% phosphorus, 0.06% sulfur, and 7.3% ash. This ration when fed sole was inadequate for live-weight maintenance and they attributed it to its low sulfur content which is associated with nitrogen requirement of cattle and suggested that sulfur supplement is essential for utilization of this forage.

### **GENETIC ENHANCEMENT OF PROTEIN**

Increasing yielding capacity of the food crops is the primary task of plant breeders and production of varieties resistant to diseases and pests is their perennial target. As far as demand for quality is concerned, it assumes importance after a certain quantitative level of food production has been achieved. In most third world countries food supplies have not kept pace with the rising population and therefore quality breeding never reached a priority level in any institution. Considering the state of malnutrition in most developing and under-developed countries and the

role pigeonpea can play in subsistence farming, the genetic enhancement of protein content in pigeonpea is a logical approach for addressing this issue. A small increase in protein content of the adapted cultivars will lead to significant protein yield on a sustainable basis. For increased adoption of the enhanced-protein cultivars it is essential that they perform as good as normal cultivars in most agronomic traits such as seed size, disease resistance, and yield. This will ensure adequate returns to farmers. Since ICRISAT has global responsibility for pigeonpea improvement, it took this challenge and a project on breeding high-protein lines was implemented. To have an effective breeding program studies on genetic control were also conducted to develop efficient breeding methodology and selection and testing procedures. The results are discussed herein.

### ***Genetic Control of Protein Content***

Information on the genetic control of protein content in pigeonpea is limited. Saxena and Sharma (1990), while reviewing the subject, reported the presence of both additive and non-additive genetic variation in determining protein content in pigeonpea and this variation was found to be controlled by 3-4 genes (Dahiya et al., 1977). Reddy et al. (1979) reported that the magnitude of heterosis for protein was in the negative direction. Dahiya and Brar (1977) reported strong maternal influence in determining protein content of seed.

For better understanding of the nature of genetic parameters the parents should have a large variation for the traits. Since the genetic material used in earlier genetic studies had limited variability for protein, Durga (1989) conducted genetic analysis for protein in two high (30-31% protein), two medium (26-27% protein), and two low (22-23% protein) lines of pigeonpea to develop basic information on various genetic parameters. She concluded that (i) reciprocal differences in  $F_1$  generation for protein were large, (ii) protein content was under additive and complementary gene effects, (iii) low-protein content was dominant or partially dominant over high-protein content, and (iv) protein content had moderately high (65.2%) narrow-sense heritability.

### ***Breeding Methodology***

As the first step in breeding for high protein, a search for high-protein trait was made in literature and in the pigeonpea germplasm available in ICRISAT gene bank. Swaminathan (1973) analyzed about 2000 pigeon-

pea germplasm and reported little variation for protein content. Although significant genotypic differences were reported for protein content in some studies (Yadav, 1984; Esh et al., 1959), the variation was not found large enough to allow selection of lines within the germplasm. Further, it was also noticed that the reported observations were based on single year data and perhaps had significant confounded influence of the sampling and environment. Considering all the factors together, it was decided to search for high-protein trait in the secondary gene pool and use them in breeding program. Since all the wild relatives of pigeonpea cannot be crossed with cultivated types, only the crossable species were examined. The results indicated that *Cajanus sericeus*, *C. lineatus*, and *C. scarabaeoides* had high-protein (Table 10). The *dhal* protein levels in this group ranged up to 31%. Therefore, these were selected as donor parents for hybridization. Breeding for high protein was carried out using pedigree method. Since the wild relatives of pigeonpea differ grossly from the cultivated types with respect to all agronomic traits and are unfit for cultivation and consumption, a breeding strategy was developed to select simultaneously for improved agronomic traits and high protein content.

TABLE 10. Protein content and seed size of high-protein lines and their parents.

Species/genotype	Protein (%)	100-seed mass (g)	Protein seed <sup>-1</sup> (mg)
Cultivated species			
Baigani	23.7	11.2	26.5
Pant A-2	22.7	7.5	17.0
T.21	24.4	7.5	18.3
Wild species			
<i>C. scarabaeoides</i>	28.4	2.3	6.5
<i>C. sericea</i>	29.4	1.9	5.6
<i>C. albicans</i>	30.5	2.8	8.5
High-protein lines			
HPL 2	29.0	12.1	35.1
HPL 7	28.0	10.0	28.0
HPL 40	27.0	10.4	28.1
HPL 51	27.9	10.6	29.6

Adapted from Saxena et al. (1987a)

### Hybridization and Selection

Crosses were made using the wild relative as female parent. In the subsequent segregating generation ( $F_2$ ), as expected, a large variation was observed for plant type and seed characters. In each cross over 200 plants were examined for protein content and the segregants with desirable protein content were selected. In each subsequent generation the individuals with improved plant type were selected in field and the final selection of the single plants for generation advance was done after determining their protein content in the laboratory. Each plant sample was evaluated in duplicate and compared with a control cultivar grown in the same field. The selected plants were ratooned and selfed using muslin cloth bags to harvest genetically pure seed. After 10 generations of pedigree selection simultaneously for agronomic traits in field and seed characters in laboratory, several breeding lines were identified (Tables 10 and 11). In these selections the high-protein trait of the wild relative of pigeonpea and seed characters of cultivated type were recovered.

TABLE 11. Performance of some high-protein pigeonpea selections at Patancheru.

Year/line	Days to mature	100-seed mass (g)	Grain yield (t ha <sup>-1</sup> )	Protein	
				(%)	(kg ha <sup>-1</sup> )
1985					
HPL 40-5	169	9.6	2.10	26.9	452
HPL 40-17	169	8.5	2.07	26.5	440
BDN 1 (control)	168	9.6	2.02	23.2	373
SE	±0.9	±0.18	±0.18	±0.46	±37.3
CV %	0.9	3.4	17.3	3.0	17.0
1986					
HPL 8-10	163	10.5	1.66	26.5	353
HPL 8-16	162	10.5	1.57	27.4	344
ICPL 211 (control)	162	14.3	1.46	21.6	251
SE	±1.1	±0.15	±0.19	±0.21	±38.5
CV %	1.3	2.5	27.0	1.7	25.8

### ***Relationship Between Seed Size and Protein***

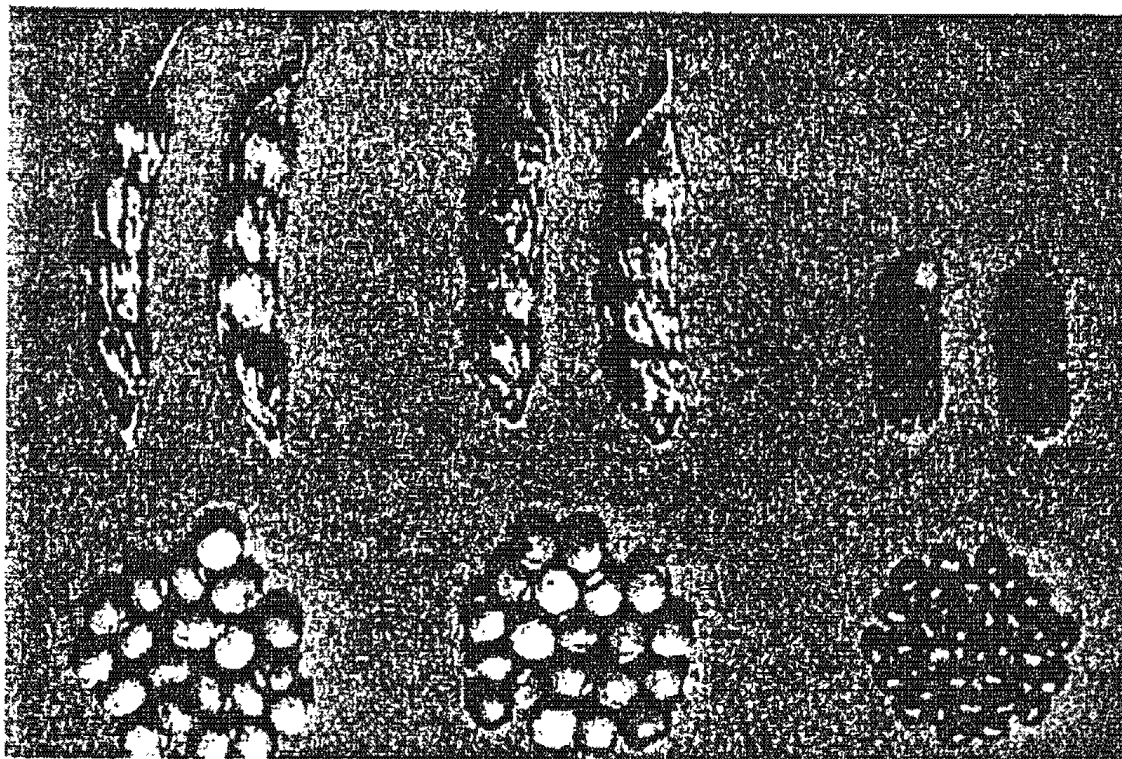
In pigeonpea, seed size is an important parameter from consumption and dehulling points of view. Negative correlations between seed size and protein have been reported in cereals such as pearl millet (Kumar et al., 1983) and sorghum (Wayne and Casady, 1974) and legumes (Blixt, 1979; Imam, 1979). In pigeonpea, Dahiya and Brar (1976) found no association between seed size and protein in 220 germplasm accessions, while Reddy et al. (1979) reported negative association between these two variables in inter-specific derivatives. Saxena et al. (1987b) while studying this relationship in 192 pigeonpea cultivars, found that the association between seed size and protein content was negative and partly controlled by genetic factors. Therefore, at the commencement of the project, aimed to develop high-protein lines in pigeonpea, there was some concern that the high-protein level of the wild species donors would remain associated with small seed size and antinutritional factors in the derived lines. According to Bahl et al. (1979) a negative relationship between seed size and protein was a result of the deposition of an increased amount of starch in seed which alters the starch:protein ratio. However, it has been found in *Vicia faba* (Abo-Hegazi, 1979), *Phaseolus vulgaris* (Gridley and Evans, 1979), and soybean (Hartwig and Hinson, 1972) that the negative correlation could be changed through breeding and selection of exceptional genotypes in segregating generations that appeared more efficient than expected in their protein synthesis and combine superior agronomic traits is possible.

The estimates of correlation between seed size and protein obtained in the breeding materials developed at ICRISAT indicated that in pigeonpea improved seed size and protein can be selected simultaneously (Saxena et al., 1987c). From the inter-specific crosses, some promising high-protein lines identified are HPL 2, HPL 7, HPL 40, and HPL 51 (Table 10). These lines combined high-protein and large seed size (Figure 2).

### ***Agronomic Evaluation of High-Protein Lines***

In pigeonpea, seed size is also associated with yield. Sharma and Saxena (1977) showed that these two variables are independent of each other in the 100-seed mass range of 9-12 g. This relationship, however, was positive in small seeded materials and negative in the materials larger than the seed size range indicated above. These observations in-

FIGURE 2. Seeds and pods of wild relative of pigeonpea (right), high-protein line (middle), and control cultivar (left)



dicated that within the medium seed size range simultaneous improvement for seed protein, seed size as well as yield could be made.

The results of agronomic evaluation of the promising high-protein selections were very encouraging (Table 11). The high-protein selections were found similar to the control cultivars in important agronomic traits such as days to maturity, seed size, and dry seed yield. For protein content the selections were significantly superior to the controls and their protein content ranged between 26-27%. An estimate of total protein harvest in this trial revealed that by growing these high-protein lines in one hectare about 350-450 kg crude protein could be harvested. These values reflect the additional advantage of 80-100 kg protein ha<sup>-1</sup>. Cultivation of these lines will markedly improve availability of the nutritive protein to the farmers without sacrificing the seed yield.

### *Nutritional Quality of High-Protein Lines*

Since the high-protein trait in pigeonpea was transferred using traditional breeding methods from its wild relatives, known to possess various antinutritional factors, it was necessary to compare the derived lines with traditional cultivars for various nutritional quality parameters be-

fore releasing them for human or animal consumption. For this purpose, Singh et al. (1990) compared two high-protein lines (HPL 8 and HPL 24) with two control cultivars (C 11 and ICPL 211). The main findings are discussed below.

*Chemical composition:* There were large differences between the protein levels of high-protein lines (28.7 to 31.1%) and control cultivars (23.1 to 24.8%). As expected the starch component (54.3 to 55.6%) of the high-protein lines was relatively less than that of controls (58.7 to 59.3%). Also the high-protein lines (2.5 to 2.6%) were marginally lower in fat content when compared with control cultivars (2.9 to 3.1%). The differences in the major protein fractions of the high and normal-protein lines were large. In comparison to controls (60.3 to 60.5%), the globulin fraction was higher (63.5 to 66.2%) in the high-protein lines and the reverse was true for glutelin (Table 12). This variation in the storage proteins, however, was not large enough to influence the amino acid profiles of high and normal-protein lines (Singh et al., 1990). Also the activities of trypsin and chymotrypsin inhibitors were found more or less similar in the high-protein lines and the controls.

*Biological evaluation:* A series of rat feeding trials and laboratory

TABLE 12. Comparison of high-protein lines and control cultivars for starch, major protein fractions, and sulphur containing amino acids.

Constituent	High-protein lines		Controls		SE
	HPL 8	HPL 40	C 11	ICPL 211	
Starch <sup>1</sup>	54.3	55.6	58.7	59.3	±0.30
Protein <sup>1</sup>	28.7	31.1	24.8	23.1	±0.09
Albumin <sup>2</sup>	9.1	8.0	7.7	8.6	±0.34
Globulin <sup>2</sup>	63.5	66.2	60.5	60.3	±1.08
Prolami <sup>2</sup>	2.9	3.2	3.6	2.1	±0.06
Glutelin <sup>2</sup>	20.2	19.7	23.3	22.8	±0.75
Methionine <sup>2</sup>	1.0	1.0	1.1	1.1	±0.02
Cystine <sup>2</sup>	0.8	0.8	0.7	0.7	±0.01

Adapted from Singh et al. (1990)

1: (g 100<sup>-1</sup> g *dha*)

2: (g 100<sup>-1</sup> g protein)



evaluations were conducted to assess various nutritional parameters for the high-protein lines. The raw seeds and cooked *dhal* samples from the high and normal-protein lines were found more or less similar in true protein digestibility, biological value, and net protein utilization. However the high-protein lines were found significantly superior in utilizable protein (Table 13). Singh et al. (1990) concluded that the high-protein lines are nutritionally superior to normal-protein cultivars as the former contain quantitatively more utilizable protein and sulfur containing amino acids. The whole seeds of the high-protein lines for animals and their *dhal* for human beings is nutritionally beneficial and its promotion will help in addressing the nutritional issues in rural areas.

TABLE 13. Biological evaluation (g 100<sup>-1</sup> g) of raw whole pigeonpea seed and cooked *dhal* samples of high-protein selections and normal-protein control cultivars.

Parameter	High-protein lines		Control cultivars		SE
	HPL 8	HPL 40	C 11	ICPL 211	
<i>Raw whole seed</i>					
Protein	25.6	27.3	21.9	21.0	±0.48
TD	58.5	58.0	59.5	60.6	±1.08
BV	68.7	70.5	64.3	64.0	±1.13
NPU	40.2	40.9	38.3	38.8	±0.64
UP	10.3	11.2	8.4	8.1	±0.23
<i>Cooked dhal</i>					
Protein	27.6	30.8	23.9	22.8	±0.26
TD	83.7	82.9	84.3	85.7	±2.14
BV	67.0	65.3	66.7	62.9	±1.68
NPU	56.1	54.1	56.2	53.9	±1.06
UP	15.5	16.7	13.5	12.3	±0.25

TD : True protein digestibility

BV : Biological value

NPU : Net protein utilization

UP : Utilization protein

Adapted from Singh et al. (1990)

### **GENOTYPE-ENVIRONMENT INTERACTION FOR SEED PROTEIN**

Environment plays a significant role in the expression of both morphological and biochemical traits in crop plants. Some characters such as seed size show less environmentally induced variability while others exhibit relatively large variation and the seed protein belongs to the later group. The observed variation for protein over locations could be influenced by environment where it is grown including the soil type and its moisture and nutrient level. Some sites favor higher nitrogen accumulation in seed as compared to others and such marked differences have been demonstrated in almost all the cereals and legumes. Hamilton et al. (1951) observed a linear relationship between protein accumulation and the increase in the alcohol-soluble protein fraction of the total protein in maize. This resulted in reduced biological value of protein as neither lysine nor tryptophan is alcohol-soluble protein.

In crops like pigeonpea where flowering is determined by photoperiod and temperature, the rate of development and duration of vegetative and reproductive periods vary widely. The meteorological conditions of certain months may exert diverse effects on the nutritional metabolism in the varieties, thus influencing the crude protein content of seeds. Pietri et al. (1971) found no response of fertilization on protein content of pigeonpea while Sham (1976), Singh et al. (1974), and Esh et al. (1959) reported significant effects of location and fertilizer application on pigeonpea seed protein. Oke (1969) reported that incorporation of 20 ppm sulfur alone or in combination with phosphorus increased methionine content of pigeonpea. Jain et al. (1986) observed significant effect of location in the advanced breeding lines of short and medium maturity duration. Singh et al. (1984c) reported significant effects of growing season (rainy and post-rainy) on various quality parameters.

Saxena et al. (1984) reported the results of an extensive study conducted by ICRISAT in 1975 to characterize environmental variation for protein content in six pigeonpea cultivars at Hyderabad (17°N), Sehore (24°N), Mandasore (25°N), Pantnagar (29°), and Hisar (29°). At each location the plantings were done in different (4-12) months. They observed large and significant differences among locations and among months within a location (Table 14). In general, the protein levels were high at higher latitudes. Within a variety grown at a particular location in different months, the variation for protein was also large. For example, in cv. 'Prabhat', planted at Hyderabad over 12 months, the seed protein content ranged from 21.6 to 25.2%; similarly at Pantnagar in 10

TABLE 14. Variation for *dhal* protein (%) in six pigeonpea cultivars planted in various months at different locations during 1975.

Cultivar (%)	Location	No. of months	Mean	Range	Variance	C.V.
Prabhat	Hyderabad	12	23.4	21.6-25.2	1.93	5.94
	Pantnagar	10	25.9	24.5-27.9	1.41	4.59
	Sehore	6	23.0	20.9-25.2	2.24	6.52
	Hisar	6	25.3	24.3-27.0	0.95	3.85
	Mandsore	7	24.5	23.2-26.2	0.90	3.88
Pusa Ageti	Hyderabad	12	23.8	21.0-26.4	2.34	6.45
	Pantnagar	7	26.7	24.7-29.3	2.73	6.19
	Sehore	8	23.9	22.1-25.4	1.29	4.76
	Hisar	6	24.9	24.2-25.4	0.18	1.72
	Mandsore	6	25.1	23.7-26.6	1.31	4.56
T. 21	Hyderabad	12	24.3	22.3-26.4	2.43	6.41
	Pantnagar	8	26.8	25.6-28.6	0.81	3.36
	Sehore	10	22.8	20.2-24.4	1.52	5.40
	Hisar	4	25.1	24.3-25.7	0.33	2.29
	Mandsore	8	26.2	24.5-28.2	1.79	5.10
No. 148	Hyderabad	12	24.0	20.7-26.8	2.79	6.96
	Pantnagar	4	26.4	25.3-28.1	1.43	4.53
	Sehore	9	23.9	22.1-26.2	1.56	5.25
	Hisar	6	25.2	24.0-26.3	0.79	3.52
	Mandsore	8	25.0	23.5-25.8	0.72	3.40
ST 1	Hyderabad	12	23.6	22.3-24.6	0.56	3.18
	Pantnagar	7	26.5	24.5-27.5	0.95	3.68
	Sehore	9	22.8	19.8-25.6	3.29	7.97
	Hisar	6	25.3	24.4-26.2	0.56	2.96
	Mandsore	7	24.8	21.4-27.4	4.45	8.50
PDM 1	Hyderabad	12	23.6	20.1-26.9	4.15	8.63
	Pantnagar	6	26.4	25.4-28.1	1.01	3.82
	Sehore	8	24.1	20.7-26.4	4.31	8.62
	Hisar	5	26.5	24.9-27.6	1.13	4.01
	Mandsore	7	23.9	22.1-26.1	3.44	7.74

plantings within a year the protein content of cv. Prabhat ranged from 24.5 to 27.9%. Besides environments, the pigeonpea lines used in this study differed considerably in their response to photoperiod and temperature and this caused a large variation in flowering and maturity. Therefore no attempt was made to identify the effect of any particular location, date of planting or any other specific factor on the protein values. The differences observed in maximum and minimum protein values within varieties amply demonstrated that the environment could

have a significant role in determining the seed protein content in pigeonpea.

Genotype-environment interaction in the high-protein lines developed at ICRISAT was also studied (Saxena et al., 1987a). The replicated trials were conducted at six locations. Although statistically significant genotype-environment interactions were recorded, the high-protein lines recorded significantly superior protein content to controls. For example in HPL 24 the protein ranged between 30.9 to 32.3% while in control it ranged between 21.4 to 24.5 (Table 15). The data also showed that the extent of variation for protein-content was more or less similar in high-protein lines and control cultivar, but the high-protein trait was maintained at each location. A summary of performance of the

TABLE 15. Protein content of high-protein selections at different locations during 1985 and in different years at Patancheru.

Locations/year	HPL 24	HPL 25	HPL 26	HPL 28	Control	SE
Locations in 1985						
Patancheru	31.3	28.6	29.7	27.8	23.3	±0.26
Jalna	32.2	28.9	29.7	30.4	23.1	±0.69
SK Nagar	30.9	28.4	29.0	27.3	21.4	±0.36
Gulbarga	32.1	29.9	-	27.6	23.0	±0.49
Gwalior	32.3	30.4	28.2	27.3	22.0	±0.71
Hisar	31.1	29.6	31.7	29.2	24.5	-
Years at Patancheru						
1981	28.3	28.3	27.6	27.6	20.9	-
1982	33.1	31.8	30.9	31.8	22.6	-
1983	29.3	29.8	29.7	29.5	23.6	-
1984	33.8	31.6	30.7	31.4	22.7	-
1985	31.4	29.1	29.0	27.9	22.9	-
1986	32.6	31.4	30.9	29.2	23.3	-
Mean	31.4	30.3	29.8	29.6	22.7	-

high-protein lines grown at Patancheru (Table 15) for six years indicated significant year-to-year variation. For example in 1982, 1984, and 1986 relatively high-protein estimates were recorded and in 1981 the estimates were relatively low. In spite of such variation over the years the superiority of high protein line was maintained.

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