A core collection of common bean from the Iberian peninsula

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## **Summary**

Characterization of crop germplasm from specific regions helps understand the patterns of genetic variation that facilitates further germplasm collection, characterization, management and their more efficient utilization in genetics, breeding and other studies. Common bean (Phaseolus vulgaris L.) is a traditional crop in the Iberian Peninsula (Spain and Portugal) where subsistence farmers have been growing and maintaining their own cultivars since their introductions from the Americas in the sixteenth century. Our objectives were to: (i) characterize diversity in the landraces collected from the Iberian Peninsula and (ii) form a core collection. Of 388 landraces from the major production regions characterized for 34 quantitative and 13 qualitative characters, including morphological, agronomic and biochemical traits, 74.7% had an Andean origin, 16.8% a Mesoamerican origin and 8.4% had seed mixtures or were recombinants between the two gene pools. Landraces of indeterminate climbing growth habit Type IV (47.2%) and bush determinate Type I (26.4%) with large (52.9%) and medium (27.4%) seeds of white (38.8%) and cream (25.9%) colour were predominant. Similarly, the "T" phaseolin pattern and common bean race Nueva Granada were the most frequent (51%). Some exceptionally large-seeded landraces of Andean (e.g., PHA-0917 with 119 g 100-seed weight<sup>-1</sup>) and Mesoamerican (e.g., PHA-0399 with 66 g 100seed weight<sup>-1</sup>) were found. These and other possible recombinants between the two gene pools merit further investigation. Fifty two landraces (13 %) were chosen to form a core collection representing the genetic diversity in the Iberian Peninsula.

### Introduction

Common bean (*Phaseolus vulgaris* L.), domesticated in Mesoamerica and Andean South America (Gepts et al., 1986), was introduced into Spain after the colonization of the Americas. The two major Andean and Mesoamerican gene pools have been identified based on morphological and agronomical traits (Singh et al., 1991b), phaseolin seed protein (Koenig et al., 1990), isozymes (Koenig & Gepts, 1989; Singh et al., 1991c), molecular markers (Becerra-Velásquez & Gepts, 1989; Freyre et al., 1996; Haley et al., 1994) and adaptation traits (Singh, 1989; Voysest & Dessert, 1991). Generally, Mesoamerican germplasm are either small- (<25 g 100 seed-weight<sup>-1</sup>) or medium-seeded (25 to 40 g 100-seed weight<sup>-1</sup>) with phaseolin patterns "S" and "B" types, different from those of their large-seeded (>40 g 100 seed-weight<sup>-1</sup>) Andean counterparts with "T", "C", "H" and "A" phaseolin types (Gepts et al., 1986; Singh et al., 1991a). Cultivars within the two gene pools have been divided into six races: Mesoamerica, Durango and Jalisco of Mesoamerican origin; and Chile, Peru and Nueva Granada of Andean origin (Singh et al., 1991a).

Although there is a lack of knowledge about bean germplasm dissemination, Gepts & Bliss (1988) and Gepts & Debouck (1991) suggested that the European germplasm was primarily from the Andes. However, it is probable that the initial common bean accessions introduced in Europe were from Mesoamerica, because Columbus arrived in Central America in 1492 and Cortés reached Mexico in 1518. In 1528, Pizarro explored Perú, opening the possibility

to introduce common bean accessions from the Andes. Thus, subsequently new cultivars may have evolved within and between the two gene pools in Southern Europe (mainly Spain and Portugal) making this region a secondary center of diversity for the common bean (Santalla et al., 2002). Moreover, seed exchange occurred mainly in the Northwest Spain (Galicia) where bean still is intercropped with maize, similar to the cropping system popular in Latin America (Santalla et al., 1994). Within Europe there was likely a quick distribution of seeds as curiosities (Zeven, 1997). Europeans still collect common beans from neighbouring and faraway regions. No records of common bean earlier than 1543 have been found in NW European herbariums, suggesting that the common bean was distributed in NW Europe after 1540, and in 1669 it was already cultivated on a large scale (Zeven, 1997).

Currently common bean is one the most important legumes for human consumption in Europe and some large germplasm collections are maintained in different countries (Amurrio et al., 2001). Lioi (1989) studied accessions from Italy and Cyprus, as well as from other non-European countries. Limongelli et al. (1996) and Piergiovanni et al. (2000) analysed seed quality of Italian beans, while Casañas et al. (1997) and Mihálikova & Benková (1995) characterized other European germplasm.

Common bean is one of the most widely cultivated grain legumes in the Iberian Peninsula. New cultivars are displacing landraces but farmers still grow landraces for personal-consumption and sale in farmer's markets. Genetic variability found for seed colour, size (although, in general, large seeded cultivars are preferred), shape and other characters is very high among genotypes grown by farmers in the Northwestern Spain and Northern Portugal (Gil & Ron, 1992; Rodiño et al., 2001). It is therefore still possible to collect valuable germplasm from this region (Ron et al., 1997). The Legumes Breeding Group at the Mision Biologica de Galicia - Spanish Council for Scientific Research (MBG-CSIC) has an Iberian common bean germplasm collection and oversees the task of further collecting, evaluating and maintaining the landraces (Ron et al., 1997).

Knowledge of patterns of genetic diversity among landraces and their relationships with new cultivars helps broaden the genetic base and maximizes use of available germplasm. In spite of increasing use of the DNA-based markers for studying genetic diversity, study of phenotypic variation in a germplasm collection is crucial for determining adaptation, agronomic potential and breeding value of landraces. Gil & Ron (1992) and Escribano et al. (1998) reported variation for morphological, agronomical and biochemical characters in common bean germplasm from NW Iberian Peninsula. Alvarez et al. (1998) estimated variation within and between common bean landraces for isozymes, random amplified polymorphic DNA (RAPD) markers, storage protein and amino acid content. These studies revealed proportionally higher variation in Spanish common bean landraces than in other germplasm previously studied.

Brown (1989) proposed formation and use of a core collection for studying and utilization of genetic diversity in large germplasm collections. Ideally, a core collection should represent most if not all diversity in the entire collection with a minimum or no redundancy (Tohme et al., 1995). With this aim in Europe core collections have been developed in different crops (Bennett et al., 2000; Liu et al., 2000; Polignano et al., 1999; Swiecicki et al., 2000). Our objectives were to: (i) determine patterns of diversity among the common bean landraces from the Iberian Peninsula and (ii) form a core collection.

#### Materials and methods

Of 388 Iberian Peninsula common bean accessions evaluated (maintained at the MBG-CSIC, Pontevedra, Spain), 286 were landraces or garden forms from Spain and 102 from Portugal. These represented different market classes (Ron et al., 1997; Santalla et al., 2001). The Portuguese landraces have been described by Rodiño et al. (2001).

Landraces were grown in field at Pontevedra (Northwestern Spain, 42 °N, 8 °W, altitude 40 m, 14 °C average temperature and 1600 mm annual rainfall) in 1998 and 1999. The experimental design was a "hill-plot", with 8 plants-plot, spaced 1 x 1 m between plots, thus giving a planting density of 80,000 plants/ha.

The following 34 quantitative characters and indices were determined (IBPGR, 1982; Santalla et al., 1994; Singh et al., 1991b): days to first flower, 50% flowering, end of flowering, flowering duration, fresh pod maturity, and first dry pod; length and width of central trifoliate leaflet (cm) and bracteole (mm); leaflet and bracteole length/width; length, width, thickness and curvature of fresh pod (mm); length/width and width/thickness of fresh pod; weight of five fresh pods (g); number of seeds per pod; number of pods per plant; pod yield (g/plant); seed dimensions (mm) (length, width, thickness, length/width and width/thickness); 100-seed weight (g); crude protein content (%); moisture content (%); crude fat content (%); total sugar content (%) (by Near Infrared Transmission spectroscopy using an Infratec 1255 Food and Feed Analyser) and seed yield (g/plant). Thirteen qualitative characters were also determined: growth

habit (from I to IV) (Hidalgo, 1991); shape of leaflet (cordate, lanceolate or ovate); leaf hairiness (sparse or dense); flower colour (white, pink or purple); shape of bracteole (cordate, lanceolate or ovate); stripes on the base of flower standard (stripped or smooth); pod fibre (scored on a 1 to 5 scale, where 1 = absence of fibre and 5 = highly fibrous ); pod colour (green, yellow or green with purple or yellow strip); pod beak position (placental or central) and curvature (curved or straight); shape of pod (flat or round) and seed shape (kidney, round or oval) and colour (white, cream, yellow, pink, red, brown, purple, black, bicolour) (Schachl & De La Rosa, 2001). Phaseolin protein pattern was analysed for five seeds for each landrace, using one-dimensional sodium dodecyl sulphate polyacrylamide gel electrophoresis (SDS-PAGE), according to the method described by Ma & Bliss (1978), Brown et al. (1981) and Gepts et al. (1986) using the reference genotypes for identification of phaseolin patterns.

The core collection was formed according to the hierarchial or branching method described by van Hintum (1995) and van Hintum et al. (2000). In that, the seed phaseolin type, an evolutionary marker (Gepts & Bliss, 1985; Gepts et al., 1986), was initially used to separate accessions into Andean (e.g., 'T', 'C', 'H') and Mesoamerican (e.g., 'S', 'B') gene pools. Within each gene pool, accessions with the same phaseolin type formed sub-groups. Morphological traits, such as, size and shape of the terminal trifoliolate leaflet, flower bracteoles, and dry seed, the presence or absence of stripes at the outer base of the flower standard, and pod beak position were used to corroborate the Andean versus Mesoamerican gene pool, determine the race affiliations, and identify recombinants or new forms

between the two gene pools. The number of accessions selected from each group to form the core collection was proportional to the size of the group and their geographical distribution. Moreover, accessions with unique characteristics (e.g., extremely large seed weight or rare recombinants between gene pools, i.e., small seed with Andean phaseolin or large seed with Mesoamerican phaseolin), despite their extremely low frequency, were included in the core collection. Subsequently, variation for each trait within the core and base collections was compared to assure consistency of genetic variability.

### **Results and discussion**

The mean, standard deviation and range for each of the 34 quantitative characters recorded for 388 common bean accessions (Table 1) indicated greater phenotypic diversity for most characters than that found by Escribano et al. (1998) in the same region. Earliness of pod maturity is an important characteristic because it helps escape or reduce incidence of insect pests and diseases. It permits farmers to bring beans to market when they can obtain high prices. It also may extend planting periods and permit farmers growing a catch crop or double cropping in temperate environments. PHA-0389, PHA-0428, PHA-0561 and PHA-0589 maturated early (< 40 days to beginning of flowering ). Because early maturity is often negatively correlated with seed yield in common bean (White & Singh, 1991), these landraces would require further evaluation in replicated trials across contrasting environments for identification of superior genotypes. PHA-0215, PHA-0252, PHA-0296 and PHA-0343 had large pods (> 170 mm long and > 14 mm width), and PHA-0917, PHA-0923, PHA-0926, PHA-0927 and PHA-0928 produced extra-large seeds (> 80 g 100-seed weight<sup>-1</sup>). PHA-0155, PHA-0262, PHA-0412, PHA-0502 and PHA-0582 produced high yields (>40 g/plant).

Regarding the nutritional quality of seed two traits could be emphasized: protein content and sugar content, the last one as component of flavour. Landraces PHA-0160, PHA-0228, PHA-0247, PHA-0280, PHA-0388, PHA-0493 and PHA-0556 scored over 30,0 % in protein content while PHA-0309, PHA-0314, PHA-0317 and PHA-0566 had more than 6,0 % sugar content that displays the value of those landraces for breeding of seed quality.

There was a predominance of landraces with indeterminate strong climbing growth habit Type IV (Table 2). This could be largely because these relatively late maturing genotypes permit farmers to take full advantage of long summer growing seasons in the traditional maize-common bean cropping systems as well as in monoculture systems grown with support, thus maximizing yield per unit of cropped land. In the Iberian Peninsula, Andean genotypes with large, white or mottled seeds of different colours and indeterminate growth habit usually are grown in association with maize. These types are preferred by growers and consumers and hence their comparatively high frequency among the 388 accessions studied. Accessions with ovate shape central leaflets and short and sparse leaf hairs were predominant. Similarly, accessions with lanceolate flower bracteoles, white colour petals and pods with placental and curved beak were more frequent. Gil & Ron (1992) also recorded very high variability among Spanish common bean landraces for seed colour, size and shape and other qualitative characters.

Because historical and linguistic information provide little evidence regarding the origin and dissemination of common bean in Europe, phaseolin protein pattern, an evolutionary marker (Gepts et al., 1986), was used to complement morphological and agronomic data. This phaseolin protein analysis was useful to identify gene pools and the origin of accessions as to the Middle American or Andean domestication centers. Also, it facilitated identification of subgroups and duplicates. The Andean "T" type phaseolin pattern was present in 36% of accessions, Andean "C" type in 22% and Andean "H" type in 21%. The Mesoamerican "S" and "B" patterns were present in 19% and 2% accessions, respectively. Among 112 accessions from the Iberian Peninsula, Gepts & Bliss (1988) found a high frequency of "C" phaseolin pattern. Lioi (1989) evaluated phaseolin protein patterns of 372 accessions from the Mediterranean (Italy and Cyprus) and other areas. More recently, Ocampo et al. (2002) compared 54 Spanish accessions with the representative wild common bean populations from its domestication range in Latin America, using phaseolin seed protein patterns and RAPD markers. Their results agreed with those of Gepts & Bliss (1988) indicating the highest frequency (57%) of accessions with the Andean "C" phaseolin, followed by "T" type (26%). The Mesoamerican "S" phaseolin was found in 15% accessions. Furthermore, the Andean accessions were domesticated from the wild populations from central and southern Peru, Bolivia and Argentina. In our study and among 66 accessions from the NW Iberian Peninsula studied by Escribano et al.(1998)., the "T" and "H" phaseolin patterns were more frequent than the type "C". Thus, the current collection of landraces were more diverse than those studied earlier probably due to the fact that bean growers and gardeners in Iberian Peninsula were regularly experimenting and introducing new bean types during the past 500 years. Such new bean types may have derived from periodic introductions from the Americas and hybridisation and mutations (Zeven, 1997).

Comparatively, high frequency (79%) of accessions with "T", "C" and "H" phaseolin patterns observed in Iberian Peninsula may indicate that genotypes originating in the Andes were better adapted to the cool and short summer of Western Europe compared to the Middle American cultivars. Their high frequency may also be due to generally greater preference for larger beans in the Iberian Peninsula and Europe because people preferred eating large-seeded fabas. Alternatively, high frequency of the Andean common beans may be due to higher proportion of cultivars grown for harvests of their green pods (Brown et al., 1982; Gepts & Bliss, 1988), usage as green shelled and dry seeds, and more attractive seed colors and shapes. However, the former is less likely because a majority of accessions were dry beans.

Approximately 8 % of the accessions were heterogeneous and exhibited two, three or four phaseolin protein patterns mainly in the most frequent Nueva Granada race. This heterogeneity may be attributed to higher level of crosspollination among beans (Ibarra-Pérez et al.,1997; Vanderborght, 1982) in the Iberian Peninsula than in other regions or deliberate maintenance of mixture of genotypes within landraces by home gardeners and subsistence farmers.

Among morphological traits, size and shape of the terminal trifoliolate leaflet, flower bracteoles, and dry seed; the presence or absence of stripes at the outer base of the flower standard and pod beak position were the most useful traits to identify the Andean versus Mesoamerican origin and the race affiliations (Table 3). As noted earlier, these characters also indicated that most landraces originated in South America, and conform to races Nueva Granada and Peru (Table 3). A noteworthy finding of this study was the identification of Mesoamerican (e.g., PHA-0399, 66 g 100-seed weight<sup>-1</sup>) and Andean (e.g., PHA-0917, 119 g 100-seed weight<sup>-1</sup>) accessions with exceptionally large seeds not found among the Latin

American collections. Moreover, there were accessions that did not have characteristics typical of either Andean or Mesoamerican gene pools. These forms had morphological traits and phaseolin and isoenzymes profiles (Santalla et al., 2002) that did not correspond with the characterization of the Andean and Mesoamerican gene pools as described by Singh et al. (1991a). Instead, they were intermediate forms or recombinants between the two gene pools (Table 3). These inter-gene pool recombinants may be of interest to breeders and geneticists because they could constitute bridging-germplasm that may aid in the transfer of useful genes between the two gene pools. Thus, these accessions may merit further studies.

The need for preserving important common bean germplasm has led to the creation of several large collections such as those maintained at CIAT, Cali, Colombia (Debouck, 1999; Hidalgo, 1991) and USDA-ARS, Pullman, Washington, USA (Silbernagel & Hannan, 1988). However, in order to effectively manage and utilize such large germplasm collections, a smaller, more manageable subsamples or core collections are being developed. For example, recently Skroch et al. (1998), Tohme et al. (1995) and Zeven et al. (1999) reported formation of common bean core collections. Thus, we also selected 52 (13 %) accessions to form a core collection (Table 4) that represent the phenotypic variation among 388 common bean accessions from the Iberian Peninsula. The range for most traits among 52 accessions of the core collection and among 388 base accessions was similar (Tables 1 & 2). Additionally, all relevant market classes identified in the base collection were represented in the core despite its reduced number of

accessions (Table 4). Thus, the core collection is representative of the common bean collection from the Iberian Peninsula. The initial screening of these core accessions for resistance to abiotic and biotic stresses and other agronomic traits could provide a clue as to in which group of landraces and garden forms to explore for additional favourable alleles and quantitative trait loci (Tohme et al., 1995; Zeven, 1999). However, the effectiveness of a core collection as a tool for understanding and utilizing crop germplasm will depend on its extensive characterization and representation of the genetic diversity it contains (Skroch et al., 1998). As an example, Miklas et al. (1999) used a subsample of the USDA common bean core collection to identify new sources of resistance to white mold [caused by *Sclerotinia sclerotiorum* (Lib) de Bary].

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#### References

Alvarez, M. T., L. E. Sáenz de Miera & M. Pérez de la Vega, 1998. Genetic variation in common bean and runner bean of the Northern Meseta in Spain. Genet Res Crop Evol 45: 243-251.

Amurrio, M., M. Santalla & A. M. De Ron (Eds.), 2001. Catalogue of Bean Genetic Resources. Fundación Pedro Barrié de la Maza / PHASELIEU - FAIR -3463 / MBG-CSIC. Pontevedra, Spain.

Becerra-Velásquez, V. L. & P. Gepts, 1994. RFLP diversity in common bean (*Phaseolus vulgaris* L.). Genome 37: 256-263.

Bennett, S.J., M. D. Hayward & D. F. Marshall, 2000. Morphological differentiation in four species of the genus *Lolium*. Genet Res Crop Evol 47: 247-255.

Brown, A. H. D., 1989. The case for core collection. In: A. H. D. Brown, O. H. Frankel, R. D. Marshall & J. T. Williams (Eds.), The Use of Plant Genetic Resources, pp. 136-156. Cambridge University Press. Cambridge, U K.

Brown, J. W. S., Y. Ma & F. A. Bliss, 1981. Genetic variation in the subunits of globulin-1 storage protein of French bean. Theor Appl Genet 59: 83-88.

Brown, J. W. S., J. R. McFerson, F. A. Bliss & T. C. Hall, 1982. Genetic divergence among commercial classes of *Phaseolus vulgaris* in relation to phaseolin patterns. HortScience 17:752-754.

Casañas, F., L. Bosch, E. Sánchez, R. Romero del Castillo, J. Valero, M. Baldi, J. Mestres & F. Nuez, 1997. Collecting, conservation and variability of the Ganxet common bean (*Phaseolus vulgaris* L.). Plant Genet Res News112: 105-106.

Debouck, D. G., 1999. Diversity in *Phaseolus* species in relation to the common bean. In: S. Singh (Ed.), Common Bean Improvement in the Twenty-Frst Century, pp. 25-52. Kluwer Academic Publ, Netherlands.

Escribano, M. R., M. Santalla, P. A. Casquero & A. M. De Ron, 1998. Patterns of genetic diversity in landraces of common bean (*Phaseolus vulgaris* L.) from Galicia. Plant Breed 117: 49-56.

Freyre, R., R. Rios, L. Guzman, D. G. Debouck & P. Gepts, 1996. Ecogeographic distribution of *Phaseolus* spp. (Fabaceae) in Bolivia. Econ Bot 50: 195-215.

Gepts, P. & F.A. Bliss. 1985. F1 hybrid weakness in the common bean: Differential geographic origin suggests two gene pools in cultivated bean germplasm. J Hered 76:447-450. Gepts, P. & F. A. Bliss, 1988. Dissemination paths of common bean (*Phaseolus vulgaris* Fabaceae) deduced from phaseolin electrophoretic variability. II. Europe and Africa. Econ Bot 42: 86-104.

Gepts, P. & D. G. Debouck, 1991. Origin, domestication and evolution of the common bean, *Phaseolus vulgaris*. In: A. van Schoonhoven & O. Voysest (Eds.), Common Beans: Research for Crop Improvement, pp. 7-53. C. A. B. Intl, Wallingford, UK and CIAT, Cali, Colombia.

Gepts, P., T. C. Osborn, K. Rashka & F. A Bliss, 1986. Phaseolin protein variability in wild forms and landraces of the common bean (*Phaseolus vulgaris*): evidence for multiple centers of domestication. Econ Bot 40: 451-468.

Gil, J. & A. De Ron, 1992. Variation in *Phaseolus vulgaris* in the Northwest of the Iberian Peninsula. Plant Breed 109: 313-319.

Haley, S. D., P. N. Miklas, L. Afanador & J. D. Kelly, 1994. Random amplified polymorphic DNA (RAPD) marker variability between and within gene pools of common bean. J Am Soc Hortic Sci 119: 122-125.

Hidalgo, R., 1991. CIAT'S world *Phaseolus* collection. In: A. Schoonhoven & O.Voysest (Eds.), Common Bean: Research for Crop Improvement, pp. 163-199.C.A.B. Intl., Wallingford, UK and CIAT, Cali, Colombia.

Ibarra-Pérez, F. J., B. Ehdaie & J. G. Waines, 1997. Estimation of outcrossing rate in common bean. Crop Sci 37: 60-65.

IBPGR (International Board for Plant Genetic Resources), 1982. Descriptors for *Phaseolus vulgaris*. IBPGR, Rome, Italy.

Koenig, R & P. Gepts, 1989. Allozyme diversity in wild *Phaseolus vulgaris*: further evidence for two major centers of genetic diversity. Theor Appl Genet 78: 809-817.

Koenig, R., S. P. Singh & P. Gepts, 1990. Novel phaseolin types in wild and cultivated common bean (*Phaseolus vulgaris* Fabaceae). Econ Bot 44: 50-60.

Limongelli, G., G. Laghetti, P. Perrino & A. R. Piergiovanni,1996. Variation of seed storage protein in landraces of common bean (*Phaseolus vulgaris* L.) from Basilicata, Southern Italy. Plant Breed 119: 513-516.

Lioi, L., 1989. Geographical variation of phaseolin patterns in an old world collection of *Phaseolus vulgaris*. Seed Sci & Technol 17: 317-324.

Liu, F. R. von Bothmer & B. Salomon, 2000. Genetic diversity in European accessions of the barley core collection as detected by isozyme electrophoresis.

Genet Res Crop Evol 47: 571-581.

Ma, Y. & F. A. Bliss, 1978. Seed proteins of common bean. Crop Sci 18: 431-437.

Miháliková, J. & M. Benková, 1995. Evaluation, maintenance and utilization of genetic resources of lentil (*Lens culinaris* Medik.) and common bean (*Phaseolus vulgaris* L.) in Slovakia. Plant Genet Res News 103: 41-44

Miklas, P. N., R. Delorme, R. Hannan & M. H. Dickson, 1999. Using of subsample of the core collection to identify new sources of resistance to white mold in common bean. Crop Sci 39: 569-573.

Ocampo, C. H., J. P. Martin, J. M. Ortiz, M. D. Sánchez-Yélamo, O. Toro & D. Debouck, 2002. Possible origins of common bean (*Phaseolus vulgaris* L.) cultivated in Spain in relation to the wild genetic pools of the Americas. Annu Rpt Bean Improv Coop 45: 236-237

Piergiovanni, A. R., D. Cerbino & C. Della Gatta, 2000. Diversity in seed quality traits of common bean populations from Basilicata (Southern Italy). Plant Breed 119: 513-516.

Polignano, G. B., E. Alba, P. Uggenti & G. Scippa, 1999. Geographical patterns

of variation in Bari faba bean germplasm collection. Genet Res Crop Evol 46: 183-192.

Rodiño, A. P., M. Santalla, I. Montero, P. A. Casquero & A. M. De Ron, 2001.Diversity in common bean (*Phaseolus vulgaris* L.) germplasm from Portugal.Genet Res Crop Evol 48: 409-417.

Ron, A. M. De, M. Santalla, N. Barcala, A. P. Rodiño, P. A. Casquero & M. C.Menéndez, 1997. *Phaseolus spp.* at the Misión Biológica de Galicia, Spain. PlantGenet Res News 112: 100.

Santalla, M., A. P. Rodiño & A. M. De Ron, 2002. Allozyme evidence supporting southwestern Europe as a secondary center of genetic diversity for common bean. Theor Appl Genet 104: 934-944.

Santalla, M., A. M. De Ron & M. R. Escribano, 1994. Effect of intercropping bush bean populations with maize on agronomic traits and their implications for selection. Field Crop Res 36: 185-189.

Santalla, M., A. M. De Ron & O. Voysest, 2001. European bean market classes. In: M. Amurrio, M. Santalla, & A. M. De Ron (Eds.), Catalogue of Bean Genetic Resources pp. 79-94. Fundación Pedro Barrié de la Maza / PHASELIEU FAIR 3463 / MBG-CSIC, Pontevedra, Spain. Schachl, R. & L. De la Rosa, 2001. Characterization of *Phaseolus* accessions. In:
C. De la Cuadra, A. M. De Ron & R. Schachl (Eds.), Handbook on Evaluation of *Phaseolus* Gemplasm pp. 29-44. PHASELIEU FAIR 3463 / MBG-CSIC, Pontevedra, Spain.

Silbernagel, M. J. & R. M. Hannan, 1988. Utilization of genetic resources in the development of commercial bean cultivars in the USA. In: P. Gepts (Ed.), Genetic Resources of *Phaseolus* Beans, pp 561-596. Kluwer Academic Publishers, Dordrecht, Netherlands.

Singh, S. P., 1989. Patterns of variation in cultivated common bean (*Phaseolus vulgaris*, Fabaceae). Econ Bot 43: 39-57.

Singh, S. P., P. Gepts & D. G. Debouck, 1991a. Races of common bean (*Phaseolus vulgaris* Fabaceae). Econ Bot 45: 379-396.

Singh, S. P., J. A. Gutiérrez, A. Molina, C. Urrea & P. Gepts, 1991b. Genetic diversity in cultivated common bean. II. Marker-based analysis of morphological and agronomic traits. Crop Sci 31: 23-29.

Singh, S. P., R. Nodari & P. Gepts, 1991c. Genetic diversity in cultivated common bean. I. Allozymes. Crop Sci 31: 19-23.

Skroch, P. W., J. Nienhuis, S. Beebe, J. Tohme & F. Pedraza, 1998. Comparison of mexican common bean (*Phaseolus vulgaris* L.) core and reserve germplasm collections. Crop Sci 38: 488-496.

Swiecicki,W.K., B. Wolko, S. Apisitwanich & P. Krajewski, 2000. An analysis of isozymic loci polymorphism in the core collection of the Polish *Pisum* genebank. Genet Res Crop Evol 47: 583-589.

Tohme, J., P. Jones, S. Beebe & M. Iwanaga, 1995. The combined use of agroecological and characteristics data to establish the CIAT *Phaseolus vulgaris* core collection. In: T. Hodgkin, A. D. H. Brown, T. J. L. van Hintum & E. A. V. Morales (Eds.), Core Collections of Plant Genetic Resources, pp. 95-107. IPGRI, Rome, Italy and John Wiley & Sons. Chinchester, UK.

Van Hintum, T. J. L., 1995. Hierarchical approaches to the analysis of genetic diversity in crop plants. In: T. Hodgkin, A. D. H. Brown, T. J. L. van Hintum & E. A. V. Morales (Eds.), Core Collections of Plant Genetic Resources, pp. 23-34. IPGRI, Rome, Italy and John Wiley & Sons, Chinchester, UK.

Van Hintum, T. J. L., A. H. D. Brown, C. Spillane & T. Hodgkin, 2000. Core collection of plant genetic resources. IPGRI Technical Bulletin No. 3. IPGRI, Rome, Italy.

Vanderborght, T., 1982. Seed increase and evaluation of the wild *Phaseolus vulgaris* L. germplasm. Mimeographed, CIAT, Cali, Colombia.

Voysest, O. & M. Dessert, 1991. Bean cultivars: classes and commercial seed types. In: A. van Schoonhoven & O. Voysest (Eds.), Common Beans: Research for Crop Improvement, pp. 119-162. C. A. B. Intl, Wallingford, UK and CIAT, Cali, Colombia.

White, J. W. & S. P. Singh, 1991. Sources and inheritance of earliness in tropically adapted indeterminate common bean. Euphytica 55: 15-19.

Zeven, A. C., 1997. The introduction of the common bean (*Phaseolus vulgaris* L.) into western Europe and the phenotypic variation of dry bean collected in the Netherlands in 1946. Euphytica 94: 319-328.

Zeven, A. C., 1999. The traditional inexplicable replacement of seed and seed ware of landraces and cultivars: A review. Euphytica 110: 181-191.

Zeven, A. C., J. Waninge, Th. Van Hintum & S. P. Singh, 1999. Phenotypic variation in a core collection of common bean (*Phaseolus vulgaris* L.) in the Netherlands. Euphytica 109: 93-106.

Table 1. Mean, standard deviation and range for 34 quantitative traits in the base and core collection of common bean from the Iberian Peninsula evaluated at

Pontevedra, Spain in 1998 and 1999

Character	Base collectio	n	Core collecti	on
	Mean - standard	Range	Mean - standard	Range
	deviation		deviation	
Davs to first flower	$48.8 \pm 7.4$	37.0 - 76.0	48.0 ± 7.5	37.0 - 67.0
Days to 50% flowering	$52.2 \pm 8.2$	39.0 - 78.0	$50.2 \pm 7.5$	39.0 - 70.0
Days to end flowering	77.2 ±13.5	56.0 - 129.0	$78.8 \pm 15.2$	56.0 - 129.0
Duration of flowering (days)	$25.0 \pm 9.7$	3.0 - 63.0	$27.8 \pm 10.6$	13.0 - 62.0
Days to pod maturity	$67.8 \pm 11.7$	51.0 - 122.0	$67.0 \pm 13.2$	55.0 - 121.0
Days to first dry pod	$90.1 \pm 18.0$	71.0 - 168.0	$90.2 \pm 19.5$	71.0 - 168.0
Length of leaflet (cm)	$9.64 \pm 1.19$	5.9 - 12.5	$9.83\pm0.97$	7.9 - 12.0
Width of leaflet (cm)	$7.33 \pm 0.99$	5.0 - 10.6	$7.53 \pm 0.76$	6.0 - 9.5
Length/width of leaflet	$1.32 \pm 0.12$	1.0 - 1.7	$1.31 \pm 0.12$	1.0 - 1.6
Length of bracteole (mm)	$5.08 \pm 0.92$	3.5 - 9.0	$5.00 \pm 0.86$	3.7 - 7.4
Width of bracteole (mm)	$3.11 \pm 0.69$	1.7 - 5.8	$3.06 \pm 0.67$	2.0 - 5.2
Length/width of bracteole	$1.66 \pm 0.23$	1.1 - 2.5	$1.66 \pm 0.22$	1.2 - 2.0
Pod length (mm)	$118.7 \pm 29.4$	51.8 - 202.6	$119.1 \pm 27.1$	75.2 - 202.6
Pod width (mm)	$12.0 \pm 1.6$	8.4 - 17.4	$12.3 \pm 1.7$	9.1 - 15.7

Character	Base collectio	n	Core collecti	on
	Mean - standard	Range	Mean - standard	Range
	deviation		deviation	
Pod thickness (mm)	5.86 ± 80	3.9 - 9.0	$5.84 \pm 0.75$	4.3 - 7.2
od curvature	$1.07 \pm 0.10$	0.7 - 1.9	$1.06\pm0.09$	1.0 - 1.5
length /width of pod	$10.0 \pm 2.6$	4.9 - 17.5	$9.60 \pm 2.15$	5.7 - 13.7
Width/thickness of pod	$2.08\pm0.33$	0.9 - 3.0	$2.13\pm0.33$	1.5 - 2.9
Weight of five pods (g)	$26.4 \pm 11.8$	2.1 - 66.6	$26.0 \pm 11.2$	7.8 - 65.3
Number of seeds per pod	$4.42 \pm 1.26$	1.0 - 8.5	$4.45 \pm 1.33$	1.3 - 7.0
Number of pods per plant	$11.4 \pm 7.2$	1.0 - 40.5	$13.7 \pm 9.6$	1.5 - 41.4
od vield (g/plant)	$60.6\pm40.8$	3.3 - 196.8	$56.6 \pm 37.2$	6.7 - 147,2
Seed length (mm)	$13.0 \pm 3.0$	7.9 - 29.0	$12.9 \pm 2.9$	7.9 - 22.0
Seed width (mm)	$8.04 \pm 1.03$	5.8 - 11.9	$7.97 \pm 1.13$	5.8 - 10.8
Seed thickness (mm)	$5.83 \pm 0.86$	3.9 - 8.1	$5.93 \pm 0.91$	4.2 - 8.1
length/width of seed	$1.62 \pm 0.26$	1.2 - 2.9	$1.58 \pm 0.25$	1.2 - 2.1
Vidth/thickness of seed	$1.39 \pm 0.17$	1.1 - 1.9	$1.37 \pm 0.17$	1.2 - 1.8
Veight of 100 seeds (g)	$40.9 \pm 14.4$	11.2 - 119.0	$42.2 \pm 16.8$	15.7 - 119.0
Seed yield (g/plant)	$11.8 \pm 8.4$	0.6 - 53.7	$12.0 \pm 8.6$	1.0 - 45.3
Protein content (%)	$27.1 \pm 1.7$	22.1 - 30.7	$27.2 \pm 1.5$	25.0 - 30.1
Fat content (%)	$1.69 \pm 0.16$	1.0 - 2.0	$1.61 \pm 0.23$	1.0 - 1.9

Character	Base collection	n	Core collection			
	Mean - standard	Range	Mean - standard	Range		
	deviation		deviation			
Starch content (%)	44.6 ± 1.5	40.1 - 49.5	44.7 ± 1.7	41.3 - 48.9		
Sugar content (%)	$4.90\pm0.47$	3.8 - 6.5	$4.88\pm0.59$	4.0 - 6.1		
Moisture content (%)	$12.2 \pm 0.5$	10.0 - 13.6	$12.2 \pm 0.5$	11.0 - 13.3		

Table 2. Frequency (%) of genotypes for 13 morphological characters in the base and core (in parenthesis) collection of common bean from the Iberian Peninsula evaluated at Pontevedra, Spain in 1998 and 1999

Growth habit	Туре І	Туре II	Type III	Type IV
	23.1 (26.9)	17.3 (17.3)	1.4 (7.7)	58.2 (48.1)
Leaflet shape	Cordate	Lanceolate	Ovate	
	14.6 (21.3)	16.8 (12.5)	68.6 (66.2)	
Bracteole shape	Cordate	Lanceolate	Ovate	
	29.5 (27.5)	53.4 (57.5)	17.1 (15.0)	
Flower wing colour	White	Pink	Purple	
	56.8 (51.3)	37.5 (43.9)	5.7 (4.8)	
Flower standard colour	White	Pink	Purple	
	55.9 (51.3)	27.4 (26.8)	16.7 (21.9)	
Stripes on flower standard	Present	Absent		
	10.9 (17.1)	89.1 (82.9)		
Leaf hairiness	Sparse	Dense		
	61.6 (65.9)	38.4 (34.1)		

Table 2.	Continued.

Pod fibre <sup>1</sup>	1	2	3	4	5
	25.8 (29.3)	30.9 (19.5)	25.4 (29.3)	15.5 (21.9)	2.40 (0.0)
Pod colour	Green	Yellow	Green/purple	Yellow/green	
	87.0 (85.3)	6.5 (2.5)	5.8 (9.8)	0.7 (2.4)	
Pod beak position	Placental	Central			
	86.9 (90.2)	13.1 (9.8)			
Pod beak curvature	Curved	Straight			
	73.5 (78.1)	26.5 (21.9)			
Pod shape	Flat	Round			
	96.6 (100.0)	3.4 (0.0)			

Primary	White	Cream	Yellow	Pink	Red	Brown	Purple	Black	Bicolour
seed colour	42.6 (44.1)	25.0 (21.2)	5.0 (0.0)	2.5 (0.0)	9.8 (5.8)	5.8 (17.3)	0.7 (0.0)	6.1 (5.8)	2.5 (5.8)
Seed	Kidney	Round	Oval	Rhombohedric					
shape	40.1 (38.5)	21.6 (28.8)	30.9 (25.0)	7.4 (7.7)					
Seed size	Large	Medium	Small						
	47.9 (51.9)	41.2 (34.6)	10.9 (13.5)						

<sup>1</sup> Pod fibre scored on a 1 to 5 scale, where 1 = absence of fibre and 5 = highly fibrous.

*Table 3.* Number of accessions of each common bean race and market classes in the base collection (the number of landraces that could be intermediate forms or recombinants between the Andean and Mesoamerican gene pools is given in italics)

Origin	Race	Market class						
Mesoamerican	Mesoamerica	Small white (29), navy (3), mulatinho (5), carioca (1), negr brillante (2), chumbinho (1), black turtle (3), small red (1 small yellow (1)						
		1 small white, 1 small yellow						
	Durango	Large great northern (10), great northern (26), pinto (6)						
		6 large great northern, 1 great northern						
	Jalisco	Small white kidney (4), rosada (1), brown mottled (1) marrow (2)						
Andean	Nueva Granada	Azufrado (12), rosada (9), bayo gordo (4), viscado (4) mottled canellini (7), black canellini (3), black mottled (2) brown garbanzo (2), brown mottled (6), canela (22), canellin (42), dark red kidney (13), cranberry (2), dark garbanzo (5) favada (15), favada pinto (1), small red pinto (14), large cranberry (18), large red mottled (5), light red kidney (1) manteca (6), sargaço (5), white kidney (29), red pinto (3)						
		5 canellini, 5 favada, 7 small red pinto, 2 white kidney						
	Peru	Ojo de cabra (8), bayo gordo (8), brown garbanzo (2), canario bola (5), cranberry (13), hen eye (1), marrow (29), morado (1), purple caparron (1), red caparron (6), rounded caparron (2)						
		1 bayo gordo, 6 marrow						
	Chile	Bayo gordo (2), cranberry (3), garbancillo (4), brown garbanzo (1), morado (1), negro brillante (16)						

Identification	Collection region / local name	Growth habit	50 % flowering (days)	Leaflet shape <sup>1</sup>	Bracteole shape <sup>1</sup>	100- seed weight	Seed color / shape	Phaseolin	Market class
Gene pool: Mes	soamerican								
Race: Mesoame	erica								
PHA-0107	Macedo de Cavaleiros, Bragança, Portugal	IV	49	С	C	28	white / oval	В	Small white
PHA-0188	Vimianzo, A Coruña, Spain	IV	61	0	С	28	white / oval	В	Small white
РНА-0523	El Barco de Ávila, Ávila, Spain / Arrocina	II	48	0	L/O	16	white / round	В	Small white
PHA-0471	Pravia, Asturias, Spain / Chichos blancos	I	54	L/O	L/O	17	white / oval	S	Navy
PHA-0663	Vilapouca de Aguiar, Vila Real, Portugal / Feijao roxo	III	49	C / O	С	25	brown / kidney	В	Mulatinho
PHA-0632	Ricla, Zaragoza, Spain	IV	55	0	C / L	29	brown/ oval	S	Chumbinho
PHA-0414	Pravia, Asturias, Spain / Chichos amarillos	Ι	44	С	L	28	cream / round	S	Small yellow
PHA-0678	Lindoso, Vila Real, Portugal / Moleirinho	II	52	L/C	0	19	red / kidney	В	Small red

Table 4. Race, market class and key characters of the 52 landraces that form the Iberian Peninsula common bean core collection.

Identification	Collection region / local name	Growth habit	50 % flowering (days)	Leaflet shape <sup>1</sup>	Bracteole shape <sup>1</sup>	100- seed weight	Seed color / shape	Phaseolin	Market class
PHA-0247	Asturias, Spain	IV	59	С	L	19	black / oval	В	Black turtle
Race: Durango				1					
РНА-0399	Pravia, Asturias, Spain / Faba pancha	IV	55	0	С	66	white / rhombohedri c	S	Large great northern
РНА-0037	Macedo de Cavaleiros, Bragança, Portugal	I	51	L	0	52	white / rhombohedri c	S	Great northern
РНА-0273	Melón, Ourense, Spain	IV	52	С	С	36	white / rhombohedri c	S	Great northern
РНА-0587	Fuentesaúco, Zamora, Spain	IV	42	0	C / O	33	white / rhombohedri c	S	Great northern
РНА-0573	Valdelacasa, Salamanca, Spain / Pinta	IV	40	С	L	28	cream / oval	S	Pinto
Race: Jalisco						•			
РНА-0065	Mirandela, Bragança, Portugal	II	47	С	0	41	white / round	S	Samll white

Identification	Collection region / local name	Growth habit	50 % flowering (days)	Leaflet shape <sup>1</sup>	Bracteole shape <sup>1</sup>	100- seed weight	Seed color / shape	Phaseolin	Market class
Gene pool: And	lean	·				·	·		
Race: Nueva G	ranada								
PHA-0582	Guernica, Vizcaya, Spain	IV	59	0	C / L	52	brown / kidney	Н	Brown mottled
PHA-0390	A Cañiza, Pontevedra, Spain	IV	51	0	L	49	brown / kidney	Н	Brown mottled
PHA-0105	Mirandela, Bragança, Portugal	II	46	L	L	72	cream / kidney	Т	Canela
PHA-0345	A Golada, Pontevedra, Spain	Ι	44	0	C / L	47	cream / kidney	С	Canela
PHA-0086	Bragança, Portugal	Ι	43	L	L	50	white / oval	С	Canellini
PHA-0252	Vila de Cruces, Pontevedra, Spain	IV	53	L	L	37	white / kidney	Н	Canellini
PHA-0441	Fustiñana, Navarra, Spain	Ι	41	0	L	35	white / oval	Н	Canellini
РНА-0563	La Bañeza, León, Spain	Ι	50	С	L	43	white / kidney	Н	Canellini

Identification	Collection region / local name	Growth habit	50 % flowering (days)	Leaflet shape <sup>1</sup>	Bracteole shape <sup>1</sup>	100- seed weight	Seed color / shape	Phaseolin	Market class
PHA-0521	Horcajo de la Ribera, Ávila, Spain / Alubia morada	I	43	0	L/O	44	red/ kidney	Т	Dark red kidney
PHA-0470	Pravia, Asturias, Spain	I	41	0	L	22	brown / kidney	Т	Dark garbanzo
PHA-0706	Villaviciosa, Asturias, Spain / Faba	IV	67	0	С	76	white / kidney	Т	Favada
PHA-0917	Lourenzá, Lugo, Spain / Faba	IV	70	0	L	119	white / kidney	Т	Favada
РНА-0654	Moimenta da Veira, Viseu, Portugal / Tourinho	I	54	L/C	0	65	white, red/ kidney	Т	Favada pinto
РНА-0486	Obécuri, Álava, Spain / Guernikesa	III	48	С	C / L	24	red, cream / round	Т	Small rec pinto
РНА-0303	Sober, Lugo, Spain	IV	50	0	L	45	cream, red / oval	С	Large cranberry
РНА-0552	La Robla, León, Spain	II	50	0	L	44	cream, red / kidney	Т	Large cranberry
РНА-0655	Moimenta da Veira, Viseu, Portugal	Ι	47	O / C	L	40	cream / kidney	С	Sargaço

Identification	Collection region / local name	Growth habit	50 % flowering (days)	Leaflet shape <sup>1</sup>	Bracteole shape <sup>1</sup>	100- seed weight	Seed color / shape	Phaseolin	Market class
PHA-0062	Mirandela, Bragança, Portugal	III	42	L	С	50	white / kidney	Т	White kidney
PHA-0178	Mañón, A Coruña, Spain / Riñón	IV	58	0	L	44	white / kidney	Т	White kidney
PHA-0225	Vilaboa, Pontevedra, Spain	IV	48	С	L	58	white / round	Т	Marrow
PHA-0590	Santa Croya de Tera, Zamora, Spain / Pinta	II	41	0	C / O	37	red, cream / kidney	T / H	Red pinto
Race: Peru					•	•		•	
PHA-0448	Peralta, Navarra, Spain	IV	50	C /O	L	37	black / oval	Н	Negro brillante
РНА-0215	Carballeda de Avia, Ourense, Spain	IV	54	0	L	43	brown / kidney	Н	Bayo gordo
РНА-0279	Vilar de Barrio, Ourense, Spain	IV	48	0	С	50	cream, red / oval	Н	Cranberry
РНА-0577	Requena, Valencia, Spain	Ι	48	0	C / L	34	cream, red / kidney	T / H	Cranberry
PHA-0476	Pola de Siero, Asturias, Spain / Ollo de pita	IV	69	L/O	L	37	white / rounded	С	Hen eye

Identification	Collection region / local name	Growth habit	50 % flowering (days)	Leaflet shape <sup>1</sup>	Bracteole shape <sup>1</sup>	100- seed weight	Seed color / shape	Phaseolin	Market class
РНА-0036	Freixo Espada, A Cinta, Bragança, Portugal	II	50	C / L	С	36	white / round	Т	Marrow
PHA-0210	Sarria, Lugo, Spain	II	48	L	0	42	white / round	Н	Marrow
PHA-0589	Muelas de los Caballeros, Zamora, Spain	Ι	39	0	C / L	43	white / rounded	Н	Marrow
PHA-0309	Sober, Lugo, Spain	II	48	L	L	48	white, purple / round	Т	Purple caparron
PHA-0565	Santo Domingo de la Calzada, La Rioja, Spain	IV	48	0	L	36	white, red / round	Н	Red caparron
PHA-0433	Peralta, Navarra, Spain	IV	69	L	0	41	brown, white / round	С	Rounded caparron
PHA-0407	Pravia, Asturias, Spain / Faba de vino tinto	IV	57	0	0	39	red / round	Н	Sangretoro
РНА-0274	Melón, Ourense, Spain	IV	50	0	С	36	brown / round	С	Dark garbanzo
Race: Chile			,					1	1
PHA-0081	Bragança, Portugal	III	41	L	L	47	cream, red/ oval	Н	Cranberry

Identification	Collection region / local name	Growth habit	50 % flowering (days)	Leaflet shape <sup>1</sup>	Bracteole shape <sup>1</sup>	100- seed weight	Seed color / shape	Phaseolin	Market class
РНА-0572	Valverde de la Casa, Salamanca, Spain / Chícharo	Ι	39	С	С	38	cream / round	С	Garbancillo
PHA-0501	Aia, Guipúzcoa, Spain / Tolosana	IV	48	0	L/O	49	black / oval	Н	Negro brillante

(1) O: ovate, L: lanceolate, C: chordate.