Cloning and expression analysis of hemoglobin genes from maize (*Zea mays* ssp. *mays*) and teosinte (*Zea mays* ssp. *parviglumis*)

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1. Introduction

Hemoglobins (Hbs) are heme proteins that have been detected in all phyla, and function by binding and transporting oxygen and other gaseous ligands [1–3]. In plants, non-symbiotic Hbs have been detected in organs of vascular and non-vascular species; however, the function of this group of proteins is still not known [4, 5]. Among flowering plants, genes coding for non-symbiotic Hbs were cloned from dicot and monocot species. In dicots, non-symbiotic Hbs were first detected in root nodules and roots of Parasponia [6], and roots of Trena tomentosa [7], and more recently non-symbiotic hb genes were cloned from a number of dicot plants, including soybean [8] and Arabidopsis [9]. Apparently, the expression of non-symbiotic hb genes varies significantly, and the highest levels of expression were detected in metabolically active or stressed organs. For example, highest levels of non-symbiotic Hb transcripts were detected in soybean stems [8], which are metabolically active organs. In Arabidopsis, two hb genes were cloned and analyzed: ahb1 gene is expressed in roots and rosette leaves but overexpressed under hypoxic conditions, whereas ahb2 gene is expressed at low levels in rosette leaves but overexpressed at low temperatures [9].

In monocots, genes coding for non-symbiotic Hbs were first cloned from barley [10] and subsequently from rice [11], but Southern blot analysis using a heterologous hb probe from barley showed that hb genes also exist in a number of cereals, such as maize, rye and wheat [10]. In barley, a single copy of the non-symbiotic hb gene exists, which is expressed in plant roots, and overexpressed in roots from plants growing in microaerobiosis. In rice four copies of the hb gene exist, and different promoters were identified suggesting that rice hb genes are regulated by different trans-acting factors.
2. Materials and methods

2.1. Seed germination and plant growing

Maize (Zea mays ssp. mays) seeds were obtained commercially, and teosinte (Zea mays ssp. parviglumis) seeds were either provided by Dr. Mark J. Millard (Iowa State University, USA) or collected at Teloloapan (Guerrero State) in the Balsas Region, México. Maize and teosinte seeds were germinated for 5–7 days in paper towels imbibed in water. Embryonic organs, such as coleoptiles, seminal roots and embryos, were collected from germinated seeds. To avoid any contamination from the endosperm, embryos were carefully detached from the cotyledon using a microspatula and, if necessary, endosperm remnants were removed using a razor blade. Organs were immediately frozen in liquid nitrogen and stored at −80°C until used. Germinated seeds were also planted in pots containing vermiculite and then grown in a greenhouse at 22°C with light/dark periods of 12 h/12 h. Plants were watered with tap water every third day and with nutrient solution [17] every sixth day. Young and mature plants were grown for 2 or 16 weeks after germination, respectively, and roots and leaves were collected as above and stored at −80°C until used.

2.2. Total DNA and RNA isolation

Total DNA was isolated from 1–3 g of roots or leaves using a modification of the hexadecyltrimethylammonium bromide method [18]. Total RNA was isolated from 100 mg of frozen tissues using the Trizol reagent (Gibco BRL), and poly(A)^+ RNA was isolated using a Quick-Prep mRNA purification kit (Pharmacia-Amersham) following the instructions of the manufacturers. DNA and total RNA and poly(A)^+ RNA were quantitated by spectrophotometry assuming $A_{260} = 50$ or 40 μg/ml for DNA and RNA, respectively [19].

2.3. Primers and PCR amplification

PCR primers were designed to amplify the maize and teosinte hb genes (hbm and hbt, respectively) using sequences from the Start and Stop codons of the maize Hb cDNA (GenBank accession No. AF236080). Oligonucleotide sequences were 5’-ATGGCAGCTCCGGAG-GCGAC-3’ and 5’-CTAAGGATCCGGTTCAT-3′ for the sense (Maize Hb/ATG) and antisense (Maize Hb/ TAG) primers, respectively. Total plant DNA (~0.5 μg) was used as template for PCR amplification. PCR components and concentrations were: 0.5 μM of each sense and antisense primer, 200 μM of each dNTP and 0.5 U of Taq DNA polymerase (Gibco BRL) in 1×PCR buffer containing 1.5 mM MgCl$_2$. PCR was done in a final volume of 25 μl using a thermal cycler (Progene). Amplification was carried out for 35 cycles at 60°C for annealing, and PCR products were detected in a 1.4% (w/v) agarose gel after staining with ethidium bromide.

2.4. Cloning of hbm and hbt genes

PCR products were isolated from the agarose gel using the Genelclean kit (Bio 101) and then cloned into the vector pCR2.1 (Invitrogen) following standard protocols [19]. Cloned fragments were fully sequenced in both directions at the DNA Sequencing Facility of the University of Nebraska-Lincoln, USA, and DNA sequences were compared with sequences deposited in the GenBank database using the BLAST program [20]. Additional computer analyses were performed using the GCG package (Genetics Computer Group).

2.5. Detection of Hbm and Hbt transcripts in plant organs

Detection of Hb transcripts in plant organs was performed by RT-PCR [21] using a RNA-PCR kit (Cetus or Gibco BRL). Reverse transcription was done using 1 μg of total RNA or 45 ng of poly(A)^+ RNA as the template for embryonic and vegetative organs, respectively, and oligo d(T)$_{16}$ as primer following the manufacturer’s protocol. PCR amplification was performed with the primers Maize Hb/ATG and Maize Hb/TAG that had been used for the amplification of hbm and hbt and the same conditions described above, but for 40 cycles. For a positive control, ubiquitin transcripts were amplified using the primers Ubi sense (5’-GTTCCTCGCGCT-CAGGCGAGC-3’) and Ubi antisense (5’-GCTTA-AATGAACCATGGCTTCAT-3’) from a maize ubiquitin cDNA (GenBank accession No. U29158). PCR products were detected in a 1.4% (w/v) agarose gel after staining with ethidium bromide.
2.6. Purification of recombinant rice Hb1 and generation of anti-rice Hb1 antibodies

Purification of recombinant rice Hb1 from transgenic Escherichia coli pEMBL18/rHb1 [11] was performed essentially as described by Arredondo-Peter et al. [22]. Pure rice Hb1 was used to generate polyclonal anti-rice Hb1 antibodies in rabbits by the Monoclonal Antibody Core Facility of the University of Nebraska–Lincoln, USA.

2.7. Western blot analysis of maize and teosinte organs

Plant organs (0.5–1 g) were homogenized with liquid nitrogen, and the homogenate was resuspended in 2 vols. of extraction buffer (20 mM Tris–HCl, pH 8, containing 1 mM phenylmethylsulfoxide fluoride). The resulting solution was centrifuged at 13 000×g at 4°C, and the supernatant was used for determination of total proteins by a dye-binding assay (Sigma-Aldrich) using bovine serum albumin as standard. To adjust to a protein concentration of 1–2 μg/μl, samples were concentrated by centrifugation using centrifugal concentrators (Centricon). Samples consisting of fresh extracts (40 μg of total soluble proteins/lane) were subjected to electrophoresis in a 15% (w/v) sodium dodecyl sulfate–polyacrylamide gel, and then transferred in transfer buffer (5 mM Tris–HCl, pH 8.0, 192 mM glycine, 20% (v/v) methanol) to a Hybond C-extra nitrocellulose membrane (Pharmacia–Amersham) at 4°C overnight.

Western blotting was performed at least in triplicate following standard protocols [23]. Briefly, membranes were blocked by incubation with 3% (w/v) non-fat dry milk in TBS buffer (100 mM Tris–HCl, pH 8.0, 150 mM NaCl) containing 0.05% (v/v) Tween 20. Protein blots were incubated with a 1/5000 dilution of the anti-rice Hb1 antibodies for 1 h at room temperature. Blots were washed with TBS-Tween and then incubated with a 1/3000 dilution of the anti-rabbit IgG conjugated with alkaline phosphatase (Roche) for 1 h at room temperature. After washing with TBS-Tween, blots were developed using the chromogenic mix nitroblue tetrazolium/5-bromo-4-chloroindolyl phosphate (Roche).

3. Results and discussion

3.1. Cloning and characterization of hbm and hbt genes

When total DNA from maize and teosinte was used as the template with primers designed for the maize Hb cDNA, two PCR products of approx. 650 and 880 bp were obtained (Figure 1). These PCR products were purified, cloned and sequenced, and the sequences were compared with sequences deposited in the GenBank database. The sequence of the smaller fragment (650 bp) showed no similarity with Hbs and was a non-specific amplification. The sequence of the larger fragment (880 bp) from maize (Figure 1A) and teosinte (Figure 1B) was identical or very similar to the maize Hb cDNA sequence, respectively, indicating that these fragments corresponded to the hbm and hbt genes.

The hbm and hbt sequences were compared to the maize Hb cDNA to identify exon and intron (IVS) sequences. Both hbm and hbt genes have four exons and three introns (Figure 2) located at identical positions as all of the known plant hb genes [11, 24, 25]. However, the nucleotide sequence at the 3′ splice site of the first intron and 5′ splice site of the third intron was unusual in hbm and hbt (agGG and AGgc, respectively) (Figure 2) compared to those sequences from Parasponia [25] and rice [11] hb genes (agGA and AGgt, respectively). Sequence comparison between hbm and hbt showed that exons were highly conserved (~99% similar), but intron sequences were rather variable (similarity values were 68, 95 and 83% for IVS-I, IVS-II and IVS-III, respectively).

Maize and teosinte hb genes code for predicted proteins of 164 amino acid residues in length (Figure 3), with a calculated molecular mass of 18.3 kDa. Sequence comparison showed that the predicted Hbm and Hbt proteins are identical to each other. Hbm and Hbt contain distal (H76) and proximal (H111) His residues, as well as the P51, F57, F81, and F11 that are highly conserved in plant Hbs [24]. Also, Hbm and Hbt contain a single Cys residue, C85, that is conserved in non-symbiotic Hbs (Figure 3) [11].

![Figure 1. Amplification of maize (A) and teosinte (B) hb genes by PCR using the primers Maize Hb/ATG and Maize Hb/TAG. Samples were subjected to electrophoresis in a 1.4% (w/v) agarose gel and stained with ethidium bromide. Lane 1, 1 kb ladder; lane 2, total PCR reaction. Arrows show the 880 bp fragment corresponding to maize and teosinte hb genes. Molecular size markers are shown in bp.](image-url)
Figure 2. Sequence alignment of maize and teosinte *hb* genes. Coding (exon) and non-coding (intron) sequences are shown in upper and lowercase letters, respectively. Vertical bars show identical bases in both sequences. Dots correspond to indels (insertion/deletion). Sequences flanking the exon/intron boundaries are underlined. N at position 99 of maize *hb* gene corresponds to C or A. Sequences were aligned using the PileUp routine of the GCG program. Maize and teosinte *hb* gene sequences were deposited in the GenBank database under accession Nos. AY005818 and AF291052, respectively.
Computer analysis showed that the hydropathy profile of Hbm and Hbt is highly similar to that of recombinant rice Hb1 (Figure 4). The tertiary structure of rice Hb1 was elucidated recently [26], and it was shown that rice Hb1 is a dimer when the concentration is approx. 0.8 mM [27]. Thus, the above observation suggests that the tertiary structure of Hbm and Hbt is very similar to that of rice Hb1. Moreover, it has been proposed that residues I/V49, S52, E122, V123 and F126 form the dimer interface of rice Hb1 and other non-symbiotic Hbs [6]. These residues are also conserved in Hbm and Hbt (Figure 3), which strongly suggests that Hbm and Hbt can also form dimers. Thus, we predict that the biochemical properties of Hbm and Hbt are the same to those reported for the rice Hb1 [11, 26, 27], i.e. Hbm and Hbt have a very high affinity for oxygen.

Figure 3. Sequence alignment of cereal and selected non-symbiotic Hbs. Distal (H76) and proximal (H111) His residues are underlined, and Cys residues are underlined. Asterisks show the most conserved residues. Helix regions are calculated from the structure of rice Hb1 [26]. Amino acid sequences were obtained from the GenBank database using the following accession Nos.: U76030 (rice Hb1), U76031 (rice Hb2), U01229 (barley Hb), M36509 (Parasponia andersonii Hb), X00296 (Trema tomentosa Hb), X53950 (Casuarina glauca Hb2), U47143 (soybean Hb) and U94998 (Arabidopsis thaliana Hb1). Alignment of sequences was done using the PileUp routine of the GCG program.
3.2. Expression analysis of hbm and hbt genes in plant organs

Non-symbiotic Hbs have been detected in plant organs by Northern blot [8], RT-PCR [11] and Western blot [28, 29]. In this work we analyzed the expression of hbm and hbt genes in plant organs by evaluating the levels of Hb transcripts and proteins by RT-PCR and Western blot, respectively.

To determine the levels of Hb transcripts in maize and teosinte organs, total and poly(A)* RNA were isolated from embryonic (coleoptiles, seminal roots and embryos) and vegetative (young and mature leaves and roots) organs, and then subjected to PCR using primers for Hbm cDNA (above). Low levels of amplification products of the expected size (498 bp) were detected in maize coleoptiles, seminal roots and embryos (Figure 5A), indicating that hbm expresses in maize embryonic organs. However, no amplification products were detected in teosinte embryonic (Figure 5B) and maize and teosinte vegetative (Figure 6) organs. Thus, to explore the possibility that the amount of Hb transcripts was below the detection limit in the above organs an (10 μl) aliquot of the RT-PCR reaction was subjected to a re-PCR for 40 additional cycles; however, no Hb transcripts were detected after re-PCR.

Maize and teosinte hb genes code for proteins with a predicted molecular mass of 18.3 kDa, which is nearly identical to the molecular mass of recombinant rice Hb1 (18.4 kDa) [11]. Our previous analysis by Western blot showed that soluble extracts of maize and teosinte organs contain proteins that comigrate with rice Hb1, and that specifically cross-react with anti-rice Hb1 antibodies (not shown), suggesting that these proteins were maize and teosinte Hbs. To determine the level of Hb proteins in maize and teosinte organs the same amount of fresh plant soluble extracts were Western blotted with anti-rice Hb1 antibodies. Results showed that high levels of Hbm protein exist in maize embryonic organs (Figure 5A), and that the highest levels of Hbm were detected in seminal roots and embryos, and the lowest in coleoptiles. Hbm proteins were also detected in maize vegetative organs (young and mature leaves and roots) (Figure 6A), but the Hb levels were considerably lower than those detected in embryonic organs. No differences were observed between Hbm levels from leaves and roots, or young and mature organs. In contrast to maize, Hbt proteins were not detected in teosinte embryonic organs (Figure 5B), which is consistent with the
Figure 6. Expression of hb genes in vegetative organs from maize (A) and teosinte (B). Arrows show the Hb proteins (18.3 kDa) detected by Western blot. Maize ubiquitin (Ubi) (500 bp) and rice Hb1 (Hb1) (18.4 kDa) were used as controls for RT-PCR and Western blot, respectively. These results are representative of at least three replicates.

inability to detect Hbt transcripts by RT-PCR and re-PCR (above). However, low levels of Hbt proteins were detected in teosinte young and mature leaves and roots (Figure 6B); apparently, Hbt levels were higher in leaves than in roots from young and mature plants.

The above observations suggest that hb genes are differently expressed in maize and teosinte organs. In maize embryonic organs hb genes express and transcripts are translated at considerable levels of Hbm protein; in contrast, in teosinte embryonic organs apparently hb genes are shut down as no Hbt transcripts and proteins were detected. These results were surprising given the relatedness of teosinte to maize, and the existence of Hbs in seeds of other cereals [10, 29]. However, it is still possible that Hbt could be below detection limits, or that hbt gene has a different type of regulation. In vegetative organs from maize and teosinte no Hb transcripts were detected, but low levels of Hb proteins were detected. This observation suggests that in vegetative organs hbm and hbt express at very low levels (below the detection limit) resulting in low abundance of Hbm and Hbt transcripts, which are translated in detectable levels of proteins. Therefore, the results obtained in this work suggest that the expression of hb genes is down- or up-regulated in maize and teosinte, respectively, from germination to vegetative growing.

4. Conclusions

The results presented above show that hb genes from maize and teosinte are highly conserved and code for identical Hb proteins. This suggests that the domestication of teosinte to maize did not substantially affect the evolution of Hbs in these plants. The function of non-symbiotic Hbs in plants is not known [4, 30], but the existence of Hbs in specific tissues of plant organs suggests that non-symbiotic Hbs play specialized rather than housekeeping functions (Ross et al., submitted). In this work we showed that the hb genes are expressed and Hb proteins are synthesized in maize and teosinte organs. However, apparently the regulation of hb genes is different in maize and teosinte seeds suggesting that Hbm and Hbt play different roles in the plant embryonic organs. Taylor et al. [10] and Lira-Ruan et al. [29] detected Hbs in germinating seeds from barley and rice, respectively, and it was postulated that Hbs play a role in germination, i.e. by providing O₂ to sustain an active metabolism. However, no Hbs were detected in germinating seeds from teosinte suggesting that these proteins are not essential for germination in this plant. Also, as judged by the level of sensitivity of the procedures used in this work, the expression of hb genes in maize and teosinte vegetative organs is extremely low but high enough for the synthesis of detectable Hb proteins. Level of Hb proteins in maize and teosinte vegetative organs were comparable, thus it is likely that Hbs have a similar function in leaves and roots from maize and teosinte.

Acknowledgments

The authors wish to express their gratitude to Drs. Mark J. Millard (Iowa State University, USA) and Cesáreo Catalán Everad (University of Guerrero, México) for providing the teosinte seeds. To Dr. Robert Hill (University of Manitoba, Canada) for providing information about the maize Hb cDNA clone. To three anonymous reviewers for helpful suggestions. This work was partially funded by Consejo Nacional de Ciencia y Tecnología (25229N) and DGAPA/PAPIIT-UNAM (IN202399), Mexico.

References


