




Hymenaea parvifolia Huber: dormancy breaking, morphology of fruit, seed and seedling¹

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ABSTRACT

The successful use of native species in conservation actions depends on basic knowledge of their biology, beginning with the identification of regenerating individuals in the field, up to the behavior and development of the species from the early stages of germination. Moreover, seed propagation and germination may be limited by the occurrence of dormancy. The objective of this work was to identify the best method to overcome seed dormancy of *Hymenaea parvifolia* Huber, besides characterizing the fruits, seeds, post-seminal development and seedlings of naturally occurring specimens in Amazonia. For this, (i) tests for the breakdown of dormancy were performed through thermal, chemical and mechanical treatments; (ii) analyses of their effects on seed germination were made; and (iii) morphology of germinative development of *Hymenaea parvifolia* was described. The treatments based on sulfuric acid for 15, 25 and 35 minutes were the most efficient in overcoming the integumentary dormancy. However, scarification by grinding is recommended because it had been less impact on the environment. External characteristics and shape of the fruit, seed and epicotyl can aid taxonomy of the genus and identification of its species in the field. In addition, morphological characters of the fruit, seed, germination and seedling were illustrated.

Keywords: Amazonia; post-seminal development; scarification; sulfuric acid.

INTRODUCTION

Hymenaea parvifolia Huber (jutaí-mirim) belongs to the family Fabaceae, subfamily Detarioideae and tribe Detarieae (LPWG, 2017). Is a tree native to the Amazon basin, which occurs mainly in drier areas of the central part of Brazil and the eastern Amazon region (Silva *et al.*, 2007; Pinto *et al.*, 2018). It also occurs in coastal areas of the Brazilian states of Piauí and Maranhão and in the northeast of Pará state. It is commonly an emergent tree found in “terra firme” forest on sandy soils, but also occurs to a lesser extent in secondary forests (*capoeira*), or in grasslands (Langenheim & Lee, 1974). The importance of this species for local riverine human

populations can be found in its diverse array of uses, such as human and animal food, medicines and timber (Shanley & Rosa, 2004).

Alongside direct extractivist uses, *Hymenaea* L. species are commonly recommended to support of the production of goods and services in degraded areas, such as live fences, shading of perennial and animal crops, forage production, green manure, among others (Miranda & Valentin, 2000). However, the lack of scientific knowledge about the behavior and development of native species for reforestation and recovery of degraded areas underscores the need for fundamental studies on the biology of such species, so that they can be successfully

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used in biodiversity management and conservation plans (Tonini *et al.*, 2008; Maranhão *et al.*, 2013). In addition, the phenotypic characteristics of seeds, of post-seminal development and young plant are attributes that can facilitate a more precise identification of the species (Oliveira, 2001), and alongside morphological knowledge of the vegetative and reproductive structures classically used in taxonomy, can help in the taxonomic problems and phylogenetic impasses that the traditional analysis of vegetative and floral organs alone cannot resolve (Gurgel *et al.*, 2012). For example, Almeida *et al.* (2011) concluded that the morphological characteristics of the jatobá (*Hymenaea*) fruits and seeds can be used to support taxonomic studies of the genus, to help interpret germination tests carried out in the laboratory and to contribute to clarify their propagation methods. However, little is known about the morphological aspects of the fruit, seed and post-seminal development of the *Hymenaea* species, which featured in the studies of Botelho *et al.* (2000); Cruz *et al.* (2001); Melo *et al.* (2004) and Bentes-Gama *et al.* (2008).

Additionally, the propagation of *Hymenaea* species may be limited by the occurrence of seed dormancy, delaying their germination. Cruz *et al.* (2001) detected germination of less than 20% in non-scarified seeds of *Hymenaea intermedia* Ducke. Andrade *et al.* (2010) also verified the impermeability of the tegument in *Hymenaea courbaril* L., with only 3.8% of germination when seeds were not submitted to any treatment to overcome dormancy. According to Fowler & Bianchetti (2000), the seeds forehead of some Fabaceae genera presents layers of osteosclereides, a tissue layer which prevents the entry of water and can delay germination for several years. The literature suggests methods for overcoming this integumentary (or exogenous) dormancy, such as acid scarification, immersion in hot or cold water, and mechanical scarification (Carpanezzi & Marques, 1981; Fowler & Bianchetti, 2000; Azeredo *et al.*, 2003; Bentes-Gama *et al.*, 2008; Mendes *et al.*, 2016). However, no systematic and comparative study has been performed so far to safely identify effective methods for the dormancy breakdown of *Hymenaea* seeds that provides high vigor and germination.

Thus, the aim of the present study was to answer the following questions: i) What is the best method to break seed dormancy of *H. parvifolia*? ii) Can fruit, seed and seedling morphology of *H. parvifolia* help in the taxonomy of *Hymenaea*? For this, seed dormancy tests were carried out with *H. parvifolia* seeds scarified and the characterization of morphological aspects of the fruit, seeds and seedlings to help the identification of regenerating individuals in the field and conservation strategies.

MATERIAL AND METHODS

The study was conducted at Embrapa Amazônia Oriental, Belém, Pará, Brazil. The seeds were obtained from fruits fallen naturally from the treetops and collected directly from the ground, at the Campo Experimental da Embrapa, in Mojú, Pará, Brazil (02°09'38,8 "S, 48°47'56,2" W).

The efficiency in the dormancy breaking of *H. parvifolia* seeds was evaluated for the following treatments: scarification in PA sulfuric acid for 15 min, 25 min and 35 min; scarification with an abrasive surface (electric grinder, 3450 rpm) on the side (near the hilum) and distal region (opposite the hilum) of the seed; scarification in hot water (70 °C) for 30 min; and immersion in distilled water for 48 hours. These treatments were compared to a control treatment (non-scarified seeds). For the treatment with sulfuric acid, 250 mL of acid was used for 110 seeds and after scarification the seeds were washed under running water for 10 minutes to remove the acid from the seed coat. After application of the treatments, the seeds were dried with paper towel and later sown at 1.0 cm depth. The sowing containers were made of polypropylene, with 12.5 cm (diameter) x 45.5 cm (depth), containing sand substrate and cured sawdust (1:1) boiled in water for two hours. Irrigation was done every two days during the experiment (20 days).

The number of days to initial seedling emergence (ISE) and number of emerged seedlings (ES) were counted each day. The final percentage of emerged seedlings was determined 20 days after the seeds were sown. Seedlings were considered emerged when all of their cotyledons were above the substrate surface. At the end of the experiment, the substrate was washed and the percentages of emergence (E), hard seeds (HS - seeds that did not absorb water and, at the end of experiment, looked the same as when planted), dormant seeds (DS - viable seeds that absorbed water and were swollen, but did not germinate during the experiment), dead seeds (DS) and abnormal seedlings (AS) were determined. The seedling emergence speed index (SEI) was also determined based on the procedure used by Maguire (1962). Emergence was represented by the normal seedlings, according to Brasil (2009). For the vigor test, the biometric data of hypocotyl diameter (HD), hypocotyl length (HL), primary root length (PRL), hypocotyl dry mass (HDM), eophyll dry mass (EDM) and root dry mass (RDM) were determined for the normal seedlings. The experimental design was completely randomized with four replicates of 25 seeds per treatment. The data were submitted to an analysis of variance and the averages of the treatments were compared using Tukey's test at 5% probability, it was used the software *Statistica 7.0* (Statsoft, 2007).

For the morphological analysis, 30 mature, healthy, entire and non-deformed fruits were randomly selected and described in relation to fruit type, color at maturity, dimensions, texture, pericarp consistency, dehiscence, indument and number and position of the seeds inside the fruit. The observations were made visually and with the aid of a stereomicroscope. The external morphological seed parameters evaluated were the consistency and surface color of the testa, and the shape and position of the hilum, lens, micropyle and raphe. For internal observations, the tip of the seed was cut (taking care not to damage the embryo) and left in water, changed daily, for 20 days. Transversal and longitudinal sections were made with a razor blade to observe the embryo (cotyledons, hypocotyl-radicle axis and plumule) and presence, type, shape, color and location of the reserve tissue. The plumule was described based on the classification proposed by Oliveira (1999). The base of the seed was considered the part near the apex of the radicle. The biometric characteristics of the fruits and seeds were obtained by measuring the length, width, thickness and weight of 50 fruits and 50 seeds using a digital caliper (0.01 mm accuracy) and precision scale (0.0001 g).

For the morphological analyses of the post-seminal development, seeds were sown 0.5 cm deep in 17.5 cm x 9 cm x 6 cm Tetra Pak containers containing a substrate of sifted sand and cured sawdust (1:1), which were both sterilized. Posteriorly, ten of the most vigorous normal seedlings were used, which were obtained by direct sowing and considered in the seedling phase when they were between the emission of the primary root and the expansion of the first eophyll. The vegetative elements described and illustrated were the same as those described by Gurgel *et al.* (2002). The terminology used is based on works by Duke (1965, 1969), Duke & Polhill (1981), Miquel (1987), Gunn (1991), Oliveira (1993), Spjut (1994), Barroso *et al.* (1999) and Ellis *et al.* (2009). The morphological characters of the fruits and seeds, the germination process and the seedlings were illustrated with schematic drawings and photographs.

RESULTS AND DISCUSSION

The study found significant differences among the treatments for most variables. The treatments (Table 1) with sulfuric acid, for 25 and 35 min, and hot water promoted the initial emergence of seedlings. All the treatments were efficient in overcoming seed dormancy, except where seeds were immersed in water for 48 hours. However, for the seeds submitted to water at 70°C, seedling emergence was markedly lower than all other treatments (16%). In relation to emergence speed, seeds scarified with sulfuric acid for 20 and 35 min and those scarified in the lateral region were the best treatments. These results are

similar to those encountered by Mendes *et al.* (2016), who analysed seeds of the same species and noted that treatments with sulfuric acid for 20 min and scarification with abrasion had the highest emergence speed.

The treatments with sulfuric acid and abrasion had the highest germination percentages (Table 2). These results were greater than those found by the following: Mendes *et al.* (2016), who obtained 57% and 45% seed emergence for *H. parvifolia* when treated with sulfuric acid (20 min) and mechanical scarification, respectively; and Busatto *et al.* (2013), with 80% and 60% emergence, Andrade *et al.* (2010), with 66.25% and 77.50% emergence, and Souza & Segato (2016), with 26% and 5% emergence, for *H. courbaril* treated with sulfuric acid and abrasion, respectively.

The effectiveness of sulfuric acid and abrasion in overcoming the impermeability of the seed coat was also observed by several authors in other genera of Detarieae. Silva *et al.* (2011) observed an increase in the speed of emergence of *Tamarindus indica* L. seeds when they were submitted to sulfuric acid treatment for 15 minutes and obtained 65% of germinated seeds at 15 days. Oliveira *et al.* (2017) confirmed the efficiency of chemical and mechanical scarification for *T. indica*, reaching 100% and 98% of the germinated seeds at 15 days after sowing, respectively. For *Copaifera langsdorffii* Desf., Silva *et al.* (2016) reached 50% germination after scarifying the seeds for 10 minutes in sulfuric acid.

For seeds of species of *Hymenaea*, the efficiency of scarification with an abrasive surface for is reported by Cruz *et al.* (2001), who found 100% germination (26 days after the seeds were sown) for scarified seeds of *H. intermedia*. Carpanezzi & Marques (1981) evaluated the effect of scarification on the seeds of *H. parvifolia* exposed to sulfuric acid for 35 min and observed around 95% germination, which corroborates with the results obtained in the present work.

The inefficiency of hot water and immersion in water for 48h can be seen by the percentages of hard seeds, which were over 50% (Table 2). The seeds scarified with hot water had a low emergence rate (18%), a high percentage of dormant seeds (53%) and a higher percentage of abnormal seedlings (18%). Seeds exposed to high temperature during the scarification process may have sustained injury to the embryos. Busatto *et al.* (2013) submitted seeds of *H. courbaril* to water at 90 °C, observed a reduction in seed germination and noted that this was probably due to the deterioration of the seeds at this temperature. Similarly, Lopes *et al.* (1998) found that thermal scarification of seeds of *Caesalpineia ferrea* Mart., *Cassia grandis* L. and *Samanea saman* Merrill. had a lethal effect on the embryos when the seeds were immersed in water at 100 °C for 30 and 60 seconds. Heat treatment

was also not effective in breaking dormancy of seeds of *C. langsdorffii* (Silva *et al.* 2016) and *T. indica* (Oliveira *et al.* 2017), which had a low percentage of emergence or inefficiency of treatment when their seeds were subjected to water at 80°C.

In relation to the seedling evaluation, there was a significant difference among the treatments for all variables (Table 3). The seeds scarified with sulfuric acid for 25 min had higher averages for all variables, except hypocotyl diameter. The superiority of this treatment can

Table 1: Days to initial seedling emergence, number of emerged seedlings and seedling emergence speed index for *Hymenaea parvifolia* Huber

Treatments	ISE	ES (%)	SEI
Scar. in H ₂ SO ₄ for 15 min	15.5 bc*	99 a*	1.35 b*
Scar. in H ₂ SO ₄ for 25 min	13.2 a	100 a	1.60 a
Scar. in H ₂ SO ₄ for 35 min	14.8 abc	95 a	1.40 ab
Scar. in the lateral region	16.2 cd	99 a	1.40 ab
Scar. in the distal region	17.5 d	96 a	1.03 c
Scar. in H ₂ O at 70°C for 30 min	14.2 ab	16 b	0.32 d
Immersion in water for 48h	-	x	-
Control	-	x	-

On what: ISE = Days to initial seedling emergence, ES = number of emerged seedlings and SEI = seedling emergence speed index.

(*) = Means followed by the same letter in the column do not differ by Tukey's test at 5% probability.

(-) = Missing values.

(x) = Values omitted from the analysis of variance are zero.

Table 2: Emergence, hard seeds, dormant seeds and abnormal seedlings for *Hymenaea parvifolia* Huber

Treatments	E (%)	HS	SDM	AS
Scar. in H ₂ SO ₄ for 15 min	98 a*	1 a*	x	1 a*
Scar. in H ₂ SO ₄ for 25 min	100 a	x	x	x
Scar. in H ₂ SO ₄ for 35 min	95 a	3 a	x	2 a
Scar. in the lateral region	97 a	x	1 a	2 a
Scar. in the distal region	96 a	4 a	x	x
Scar. in H ₂ O at 70°C for 30 min	18 b	53 b	11 b	18 b
Immersion in water for 48h	x	100 c	x	x
Control	x	100 c	x	x

On what: E = Emergence, HS = hard seeds, SDM = dormant seeds, AS = abnormal seedlings.

(*) = Means followed by the same letter in the column do not differ by Tukey's test at 5% probability.

(x) = Values omitted from the analysis of variance are zero.

Table 3: Hypocotyl diameter, hypocotyl length, primary root length, hypocotyl dry mass, eophyll dry mass and root dry mass for *Hymenaea parvifolia* Huber

Treatments	HD (mm)	HL (cm)	PRL (cm)	HDM g/seedling	EDM g/seedling	RDM g/seedling
Scar. in H ₂ SO ₄ for 15 min	4.6 a*	12.1 a*	16.2 a*	5.67 b*	5.18 ab*	2.67 ab*
Scar. in H ₂ SO ₄ for 25 min	3.6 b	13.7 a	18.2 a	6.48 a	7.01 a	3.13 a
Scar. in H ₂ SO ₄ for 35 min	3.8 b	12.6 a	15.8 a	6.30 a	5.87 a	2.50 b
Scar. in the proximal region	4.9 a	12.1 a	15.2 a	6.45 a	3.54 bc	2.74 ab
Scar. in the distal region	4.8 a	8.8 b	16.1 a	5.07 c	2.16 cd	2.71 ab
Scar. in H ₂ O at 70°C for 30 min	1.02 c	1.2 c	3.5 b	0.82 d	0.35 d	0.54 c
Immersion in water for 48h	-	-	-	-	-	-
Control	-	-	-	-	-	-

On what: HD = Hypocotyl diameter, HL = hypocotyl length, PRL = primary root length, HDM = hypocotyl dry mass, EDM = eophyll dry mass and RDM = root dry mass.

(*) = Means followed by the same letter in the column do not differ by Tukey's test at 5% probability.

(-) = Value nonexistent.

be explained because it had a higher emergence percentage. Souza & Sagato (2016) analysed the vigor of *H. courbaril* seeds scarified with sulfuric acid (20 min) and with abrasion opposite the hilum. Although there was no significant difference between the treatments, they observed that the chemically scarified seeds were less vigorous than those mechanically scarified. They also noted that absorption of the acid by seeds during the chemical treatment could have caused the lower vigor. In the present study, this was not observed since the seeds scarified with acid for 25 min were the most vigorous.

Fruit is a camara (Figure 1A, B), indehiscent, dry, ovoid; stipe inserted into the remnants of the uniseriate calyx (Figure 1D), margins not constricted, pedicel terete, woody, dark greyish brown, opaque, rugose, striate, glabrous (Figure 1C). Exocarp is a lighter brown, homochromous, glabrous, chartaceous, resinous; endocarp is fibrous-spongy, yellowish brown, opaque, glabrous, Funicle is chartaceous. A globose camara, with an external woody

layer and yellowish brown, velvety, farinaceous and fibrous pulp, was observed by Camargo *et al.* (2008) for the same species. These characteristics are similar to those encountered by Duarte *et al.* (2016) for *H. courbaril*. These authors observed a dry, indehiscent, polyspermous fruit with a thick mesocarp and thin, soft, spongy endocarp. Sousa *et al.* (2010) observed in *T. indica*, dry, indehiscent and polyspermic fruit, differed from the species under study, the crustacean epicarp, pulpy-gelatinous mesocarp and the epicarp detached from the seed.

Barroso *et al.* (1999) noted that the seeds of *Hymenaea* have a fleshy funiculus that is surrounded by a white aril. However, Gunn (1991) points out that the seeds of the genus do not have an aril and described this structure as the endocarp, which was corroborated by Paiva & Oliveira (2004) and adopted in the present work. The presence of aryl is a variable character among species belonging to the genera of Detarieae. *Guibourtia hymenifolia* (Moric.) J. Léonard presented spongy and whitish aryl (Battilani

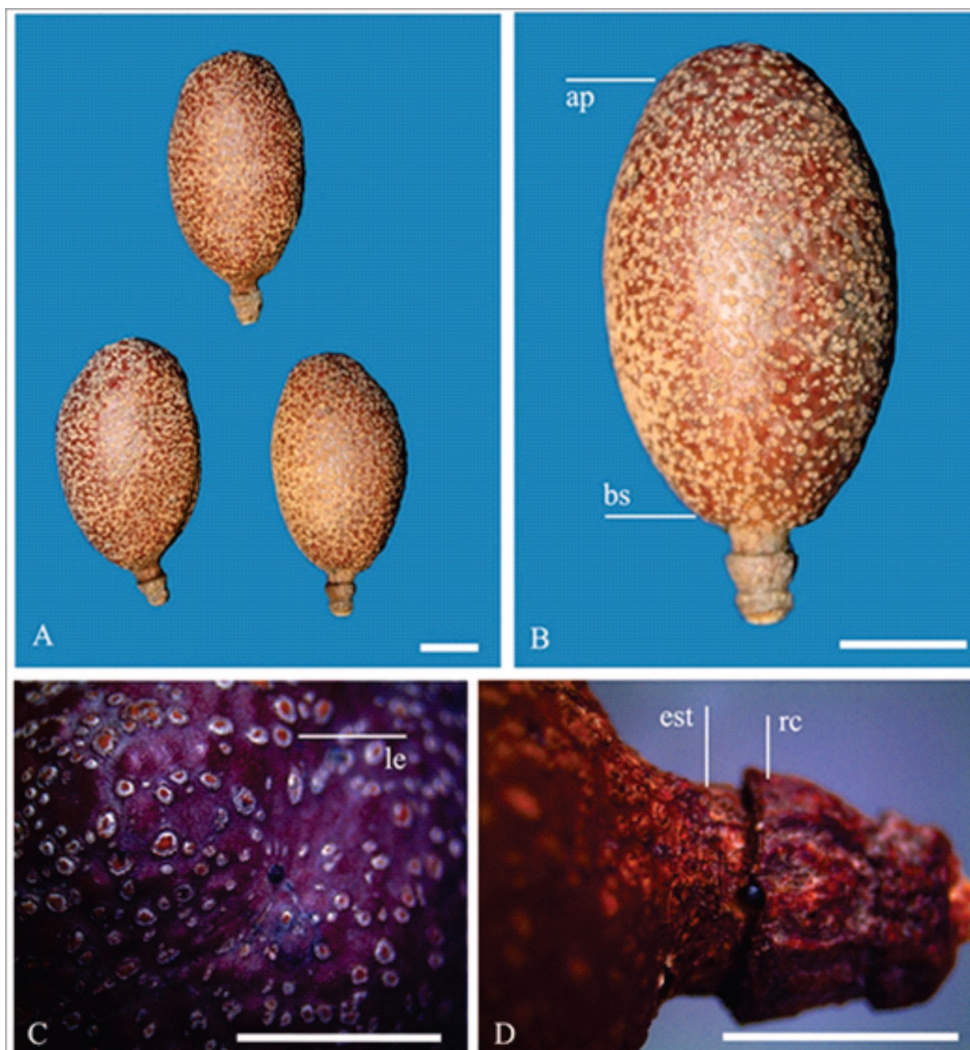


Figure 1: Fruit of *Hymenaea parvifolia* Huber A) Camara; B) entire fruit; C) lenticels in detail; D) base of camara. Legend: ap - apex, bs - base, est - stipe, le - lenticels, rc - remnants of calyx. Scale: A and B) 1 cm; C and D) 0.5 cm.

et al., 2011), *Guibourtia chodatiana* (Hassl.) J.Léonard the aryl is white and covers the entire seed (Pinto *et al.*, 2017), *Copaifera arenicola* [(DUCKE) J. COSTA AND LP QUEIROZ] the aryl is yellow (Gama & Nascimento Jr, 2019) and *C. langsdorffii* the aryl is orange (Souza *et al.*, 2017). For *T. indica*, the aryl is present in a small amount around the seeds (Sousa *et al.*, 2010).

The seeds of *H. parvifolia* are oblong and highly uniform in both shape and size (Figure 2A). The testa is homochromous, rocklike, dark brown, opaque, smooth (Figure 2B), completely surrounded by a farinose, greyish-white endocarp, pleurogram absent. The hilar region is basal (Figure 2C), near the apex of the radicle, hilum is punctiform and lens is more prominently rounded, opaque, reticulate, glabrous, imperceptible to the naked eye. Micropyle and raphe are imperceptible. Embryo is cotyledonar and axial, plumule is moderately developed or poorly differentiated (*mitaforme* type), in the form of a glove (Figure 2D, E). Camargo *et al.* (2008) described the seeds of *H. parvifolia* as ellipsoid with a smooth and glabrous testa, punctate hilum and pleurogram absent. In *H. courbaril* the seeds presents the following characteristics: tegument stony, reddish to dark brown; hilum oblong, micropyle discreet, embryo axial, cotyledonar

differing from *H. parvifolia* by the ovoid and globular shape (Duarte *et al.*, 2016). These characteristics corroborate the circumscription of the genus. However, some characteristics such as: dark brown forehead, hilum punctiform, crass cotyledons and axial embryo are common to *T. indica*, which differ from the species under study, due to the presence of a pleurogram on both sides, a plumule with bifoliated rudiments and the hypocotyl axis, radicle of the asciform type (Sousa *et al.*, 2010).

Germination is phanerocotyledonar, epigeous, and emergence is curved and then straight. Similar features were described by Guerra *et al.* (2006) para *C. Langsdorffii*, Kadoma & Sartori (2007) for seedlings of *Hymenaea stilocarpa* var. *stilocarpa* Mart., Camargo *et al.* (2008) for *H. parvifolia*, Sousa *et al.* (2010) para *T. indica*, Battilani *et al.* (2011) para *G. hymenifolia* and Duarte *et al.* (2016) for *H. courbaril*. In the present study, the hydrated seeds increased in volume and on the ninth day after sowing the main root broke the seed coat in the basal region (Figure 3A). The main root is yellowish, glabrous, root cap is not evident, root collar is asymmetrically differentiated, pinkish, with hairs numerous, simple, hyaline, long, straight, with curved, adpressed apex (Figure 3B), lenticels are reddish brown. Cotyledons are oblong, thick, reddish,

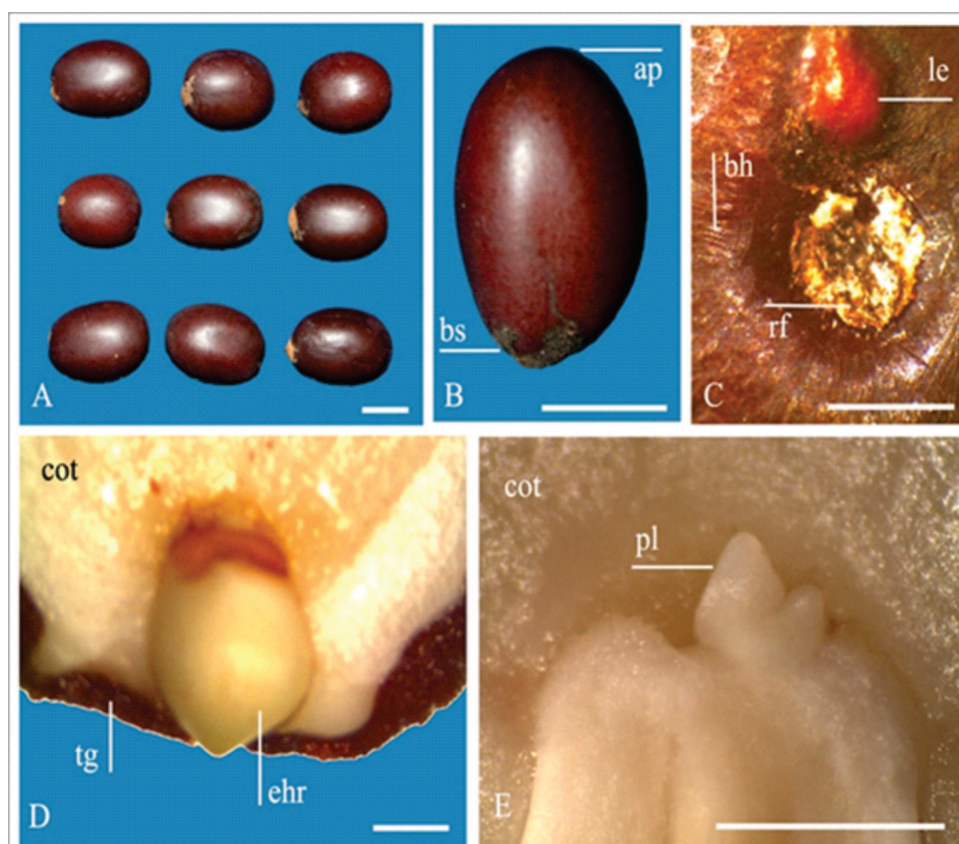


Figure 2: Seeds of *Hymenaea parvifolia* Huber A) oblong seeds; B) apex and base; C) hilar region in detail; D) embryonic axis; E) mitaformplumule. Legend: ap - apex, bh - hilar border, bs - base, cot - cotyledons, ehr - hypocotyl-radicle axis, le - lens, pl - plumule, tg - tegument, rf - funicular remnants. Scale: A) 1 cm; B) 0.5 cm; C and E) 1 mm.

parallel and unilateral, opposite when completely open, sessile, trichomes are rare, simple, hyaline, reduced on the margins and adaxial surface. *H. courbaril* emits the main root on the third day after sowing, with similar characteristics to those described in this study, they differ by the little differentiated neck (Duarte *et al.*, 2016).

Roots are pivotal, the main root is axial, cylindrical, sinuous, subwoody, rarely with hairs that are simple, hyaline, reduced and straight, streaks and lenticels are dark brown, and side roots are numerous (Figure 3D). Hypocotyl and root collar are also elongate, which is similar to what was observed by Melo *et al.* (2004) in a study of *H. intermedia* seedlings. Root collar is evident, asymmetrical, reddish, hairs are numerous, simple, hyaline and short, posteriorly unilateral (Figures 3E; 4A). Hypocotyl is vigorous, epigeous, straight, rectangular with rounded vertices, reddish, subsequently turning ferruginous, with many simple trichomes that are hyaline, short and straight, lenticels are present, reddish and intact (Figures 3F; 4B, C). These characteristics confirm the descriptions made by Camargo *et al.* (2008) for the same species, observed the elongated hypocotyl, initially pink, becoming brown, initially smooth, becoming striated and covered by short whitish hairs. The morphology of the *H. parvifolia* hypocotyl differs of *H. intermedia* var. *adenotricha* that has a cylindrical shape and a light green color (Melo *et al.*, 2004), from *T. indica*

with a cylindrical, light green and glabrous shape (Sousa *et al.*, 2010).

Cotyledons are fleshy, pinkish, elliptical, initially covered by the seed coat and rarely with trichomes around them (Figure 3E, F). This differs from what was encountered by Kadoma & Sartori (2007) for seedlings of *H. courbaril*, which lacked the trichomes and elliptical shape of the cotyledons. Epicotyl was straight, cylindrical, greenish, glabrous, with light brown lenticels, a bud in development and a stipule (Figures 3G; 4D, E, F, I). These characteristics corroborate with Duarte *et al.* (2016) who described the *H. courbaril* epicotyl as long, green with the presence of stipules in the first two metaphyles. Eophylls were simple, unifoliolate, reddish brown, opposite, ovoid, apex obtuse, margin ciliate, rarely with simple, hyaline, reduced, straight trichomes, base reniform, venation penninerved, secondary veins eucaptodromous, tertiary veins reticulate, immersed on the adaxial surface, impressed on the abaxial surface, coriaceous, translucent glands throughout the laminar surface; pulvinus and pulvinule green, striated, rugose, short, both with simple, hyaline trichomes (Figure 4D), the pulvinule more frequently lateral (Figures 3G; 4G, H, D). The characteristics of pulvino and pulvino were similar to those described by Melo *et al.* (2004) for the eofilos of *H. intermedia* that differ from *H. parvifolia* by their light green color and for being composed.

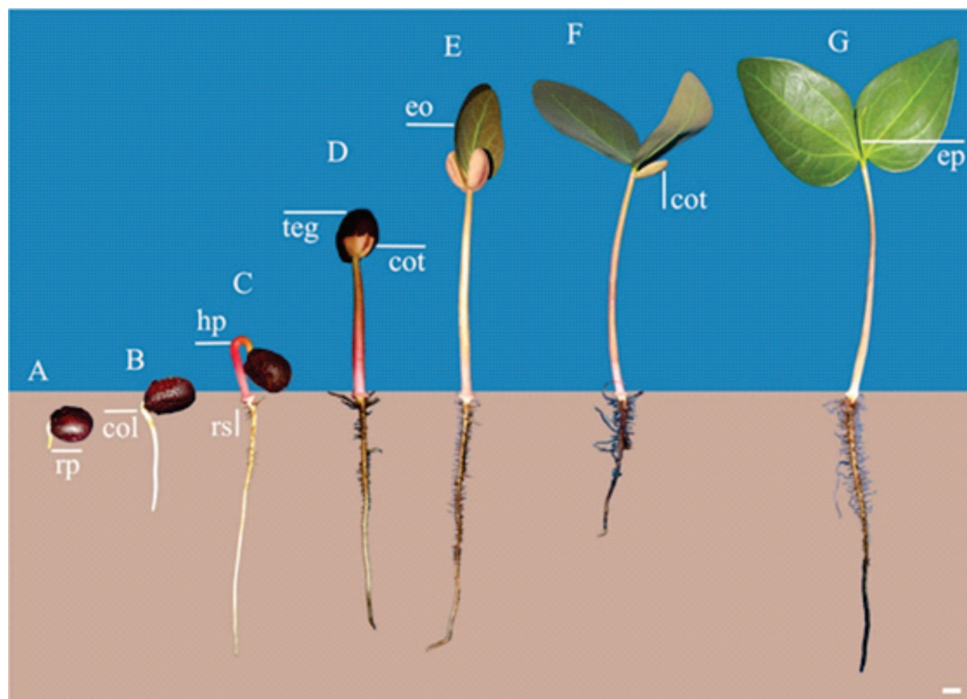


Figure 3: Stages of post-seminal development and seedlings of *Hymenaea parvifolia* Huber A) primary root emission; B) elongated primary root; C) hypocotyl handles; D) straight hypocotyl; E) emergence of eophylls; F) free and fully expanded eophylls from cotyledons; G) elongated epicotyl. Legend: col - differentiated root collar, hp - hypocotyl, rs - secondary roots, teg - tegument; cot - cotyledons, eo - eophylls, ep - epicotyl, rp - primary root. Scale: 1 cm.

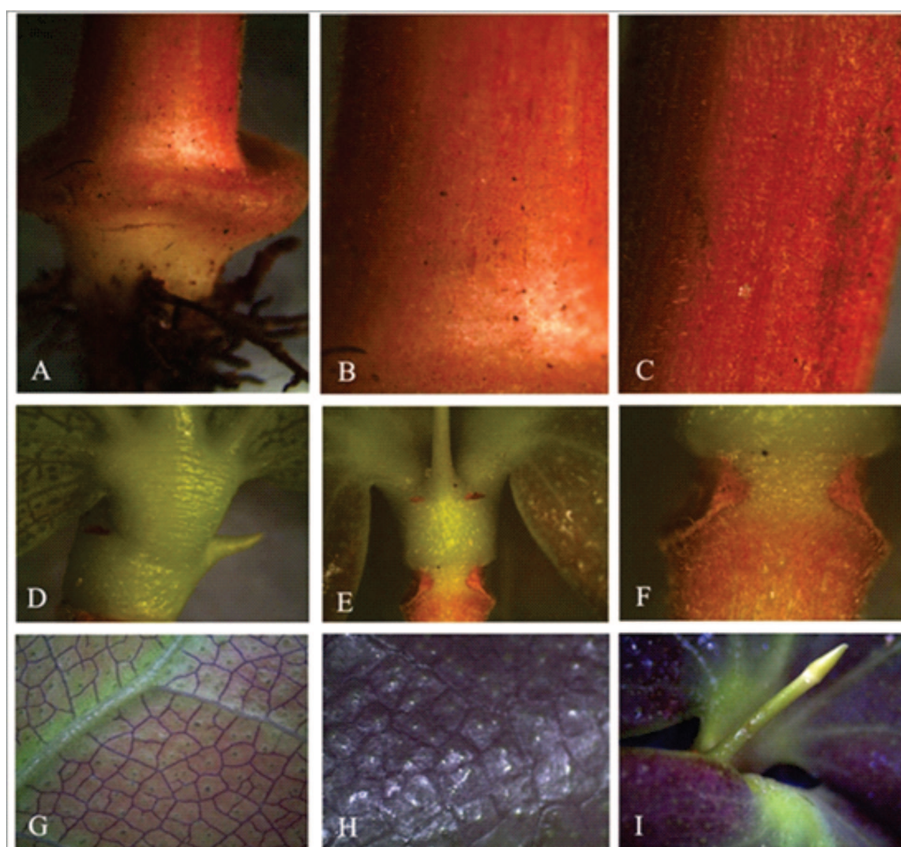


Figure 4: Details of the seedling of *Hymenaea parvifolia* Huber A) Root collar; B) Hypocotyl marks; C) hyaline trichomes on the hypocotyl; D) striate pulvinus and pulvinule; E) sessile eophylls and stipule scar; F) Cotyledon scars; G) Glands on the abaxialeophyll lamina; H) Glands on the adaxialeophyll lamina; I) Epicotyl. Magnification: 25x.

CONCLUSIONS

The treatments used in this study to break the dormancy of *H. parvifolia* Huber can make a difference in seed germination time and seedling vigor. Thus, treatments based on sulfuric acid for 15, 25 and 35 minutes were the most efficient in overcoming the integumentary dormancy and better in all aspects analysed. However, scarification by grinding the side near the hilum is recommended because it has less impact on the environment.

Morphological characteristics of fruits, seeds, post-seminal development and seedlings highlighted here have been shown to be a useful resource for the differentiation between this and other species of *Hymenaea*. Among them, those characteristics with higher taxonomic value are the fruit shape of *Hymenaea parvifolia*, size and shape of the seed and characteristics of the epicotyl.

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