Female gametogenesis and early seed development in *Amaranthus hypochondriacus* L.

**Abstract**

**Background.** Attention to amaranth grains has increased in recent years due to the nutritional value of their seed proteins, which have high levels of the amino acid lysine. However, there is no detailed study describing the stages of seed development in *Amaranthus hypochondriacus*.

**Question.** How are the developmental patterns of the female gametophyte and young seed in *Amaranthus hypochondriacus*?

**Species studied.** *Amaranthus hypochondriacus* L ‘Revancha’ (Amaranthaceae).

**Study site and years of study.** Plants were grown and collected from 2014 to 2016, in a greenhouse at Instituto Tecnológico de Celaya, Guanajuato, Mexico.

**Methods.** Glomerules were collected before pollination and two weeks after anthesis. The ovules at different development stages were fixed and cleared and were analyzed by light microscopy. A clearing protocol was used to observe the developmental stages during female gametogenesis and embryogenesis.

**Results.** We observed that the *Amaranthus hypochondriacus* ovule has a campylotropous form. The female gametophyte showed a *Polygonum*-type pattern of development. We were also able to identify all the stages from the megaspore mother cell to the cotyledon embryo stage. After meiosis, the micropylar megaspore differentiates into the functional megaspore. The embryo did not show symmetric divisions, although the final pattern is similar to that of in eudicotyledons. The suspensor showed additional longitudinal divisions, giving rise to a 2-rowed suspensor, while the endosperm showed a helobial development.

**Conclusions.** These results will be used as baseline to identify morphological changes during seed development and to develop new strategies to improve seed quality or increase the yield.

**Key words.** Amaranth, embryogenesis, megagametogenesis, megasporogenesis.

**Resumen**

**Antecedentes:** La atención al grano de amaranto ha aumentado en los últimos años debido al valor nutricional de las proteínas de su semilla con altos niveles de lisina. Sin embargo, no existe un estudio detallado que describa las etapas tempranas del desarrollo de su semilla.

**Pregunta:** ¿Cómo es el patrón de desarrollo del gametófito femenino y la semilla en *Amaranthus hypochondriacus*?

**Especie de estudio:** *Amaranthus hypochondriacus* L ‘Revancha’ (Amaranthaceae).

**Sitio de estudio y fechas:** Las plantas fueron cultivadas y se colectaron entre los años 2014 y 2016, en invernadero, en el Instituto Tecnológico de Celaya, Guanajuato, Mexico.

**Métodos:** Los glomérulos se colectaron antes de fecundación y hasta dos semanas después de la antesis. Los óvulos se fijaron y aclararon en diferentes estados de desarrollo, se analizaron bajo microscopía de luz en campo claro. Mediante un protocolo de clareo se logró observar todas las etapas de desarrollo de la gametogénesis femenina y la embriogénesis.

**Resultados:** Los óvulos de *Amaranthus hypochondriacus* son del tipo camplótrofo. El gametófito femenino muestra un patrón de desarrollo del tipo *Polygonum*. Se identificaron todas las etapas de desarrollo desde la célula madre de la megaspora al estado embionario cotiledonar. Después de la meiosis la megaspora micropilar se diferencia en la megaspora funcional. El embrión no mostró divisiones simétricas aunque el patrón final es similar al observado en eudicotiledóneas El suspensor mostró divisiones longitud-

This is an open access article distributed under the terms of the Creative Commons Attribution License CC BY-NC (4.0) international.
los óvulos adicionales, dando lugar a un suspensor con dos filas de células. El endospermo mostró un patrón helobial.

**Conclusiones:** Estos resultados servirán de base para identificar cambios morfológicos durante el desarrollo de la semilla y generar nuevas estrategias que mejoren la calidad y la producción de las mismas.

**Palabras clave.** Amaranto, embriogénesis, megagametogénesis, megasporogénesis.


Entre los pseudocereales, los amarantos son considerados de tener gran potencial para reducir problemas de malnutrición, especialmente en países en desarrollo, y son prometedores cultivos para los próximos decenios con el objetivo de seguridad alimentaria mundial. Son capaces de adaptarse a suelos pobres, toleran sequía y salinidad, y tienen alto valor nutricional, propiedades funcionales y una amplia variedad de usos.

Durante las últimas décadas, las posibilidades nutricionales y otras cualidades únicas de amarantos han sido resaltadas a nivel mundial, no obstante, no se han regresado a su lugar como cultivo básico y estratégico (Das 2016, Espitia-Rangel 2016). La grano es el producto comercial principal de *A. hypochondriacus*, por lo tanto es importante desarrollar información sobre el ontogénesis de la semilla amaranto y su biología reproductiva.


El desarrollo de las semillas no es sincronizado en todas la inflorescencia, debido a la indeterminada crecimiento del inflorescencia de amaranto. Se realizó una análisis para determinar cómo la posición influye el tamaño del óvulo y el ovario. El objetivo del presente estudio fue analizar female gametophyte and early seed development of amaranth, para llenar los vacíos en los informes y generar información que pudiera ser usada como base para identificar cambios morfológicos o para investigación adicional.

**Materials and methods**

El ciclo de vida de amaranto plantas crecidas en condiciones de invernadero se completó en 70-90 días. Después de la anversión, la etapa reproductiva (desde la anversión hasta la madurez fisiológica) se completó en 30 días (en condiciones de campo el ciclo de vida se completó en 130 días).

**Plant material.** Nosotros estudiaron el ciclo reproductivo de *Amaranthus hypochondriacus* L. ‘Re-vancha’. Las plantas fueron cultivadas en macetas con 1 kg de substrato (tierra de turba:perlita:vermiculita 2:1:1 v/v) en un invernadero, donde el mínimo y máximo temperatura fueron 11 ± 2 y 27 ± 2 °C. Los ovarios femeninos (OF) fueron recogidos antes de que el estaménos (SF) libera polen y dos semanas después de la anversión. Los ovarios femeninos (OF) fueron recogidos durante dos años y aproximadamente 25 óvulos de cada etapa fueron observados para estudios morfológicos y estadísticos.

**Variations of the ovary and seed, by branch and inflorescence position.** En orden a correlacionar el tamaño de los óvulos con su desarrollo, estas fueron medidas en diferentes etapas. Inflorescencias fueron divididas en tres secciones: basal (ST1), media (ST2) y apical (ST3) según el procedimiento descrito por Olufolaji & Odeleye (2011), también se dividieron en tres ramas para experimentar...
basal (BR1) middle (BR2) and apical (BR3) sections (Figure 2S). Ovary length, ovule length and ovule width were measured in each of the inflorescence positions. The data were analyzed using a randomized complete block design (three inflorescence and three branch sections) using three replicates for each treatment, using the GLM procedure of the SAS; the comparisons of means used Tukey’s honestly significant difference (HSD) test ($P < 0.05$). Twenty-five ovules of each stage were observed for female gametophyte development with their respective ovule dimensions.

**Histological analysis.** Glomerules were fixed in FAA solution (18:1:1 ethanol: formaldehyde: acetic acid), placed under vacuum three times for 5 minutes and left in that solution for 24 h at 4 °C. After fixing, the samples were dehydrated through an ethanol series (35, 50, 75, 80, 90 and 96 % ethanol). The ethanol was gradually replaced with xylene (3:1, 1:1, 1:3 v/v) up to 100 % xylene, then the samples were embedded in Paraplast wax (Leica); samples remained in 100 % Paraplast for 24 h (Johansen 1940). Sections (8-12 µm) were cut with a microtome (Leica), mounted on glass slides and stained with hematoxylin-eosin.

**Clarification.** Ovules and immature seeds were fixed for 24 h in FAA solution. After fixing, the ovules and immature seeds were dehydrated in an ethanol series up to 96 % ethanol. Ovules were cleared through a series of methyl salicylate-ethanol solutions in proportions of 3:1 (for 24 h), 1:1 (for 2 h), 1:3 (for 2 h) up to 100 % methyl salicylate (Salinas-Gamboa et al. 2016). They were incubated at 4 °C from 5 to 25 days and observed with a Leica microscope (Wetzlar, Germany), under Nomarski optics.

**Results**

**Effect of the position on the branch and inflorescence on ovule and ovary size.** PF contains an ovary with one ovule, short style and three long stigmas (Figures 1C, D, 2S). After 71 days of planting, most gametes were fertilized and the seed stages ranged from early embryogenesis to grain filling (Figure 1S, Table 1). The ovules from the basal and middle sections were larger than the ovules from apical sections; only ovary length was not affected by the position on inflorescence (Figures 1S, 2S Table 2).

**Inflorescences.** The glomerules are dichasial cymes; the first flower is a single terminal SF on the branch on either side; the second and third PF are terminal on the secondary branches (Figures 1A, B, 2S). Well-developed SF with five mature anthers containing pollen grains was observed. These flowers finished development before the PF on the same branch (Figure 1B). The development of the central PF was not synchronized with others PF, we observed PF with ovules that are developing while others already have developed ovules (Figure 1C, D).

**Type of ovule.** Each *A. hypochondriacus* ovary contains only one ovule; before pollination, when the stigmas were closed (Figures 1C, D, 2S), the nucellar tissue and the integuments were still developing and growing. The amaranth ovule is of the campylotropous type, in which the micropyle is directed toward the chalaza. The chalaza is situated at a right angle to the funiculus. The young ovule was transiently observed as anatropous (Figure 2). During its development the micropyle of this ovule is oriented towards the funicle by the curvature of the nucellus (Figure 2A, B).

**Megasporogenesis and megagametogenesis.** Megasporogenesis began when in the apical region of the nucellus a subepidermal cell developed as an archesporial cell (AR, Figure 3A, B), which will give rise to the megaspore mother cell (MMC) or megasporeocyte (Figure 3C, D). After the MMC undergoes meiosis, it gives rise to four haploid megaspore cells (Figure 3E). At the end of the meiosis (Figure 3F), the megaspores located in the chalazal end die, while the micropylar becomes the functional megaspore (FM).

Megagametogenesis begins when the FM becomes enlarged, starting the female gametophyte development. We identified the initial coenocytic phase, in which three rounds of nuclear
Figure 1. Longitudinal sections of glomerules and reproductive structures of amaranth. A-B: Glomerules of amaranth showing staminate (SF) and pistillate flowers (PF), SF showed anthers (ANT) with pollen (PO). C-D: Early stages of ovule (O) development inside the ovary (OV) with inner (II), outer integuments (OI), ovule primordium (P), nucellus (N), chalaza (CL), micropyle (M) and funicle (F). Scale bars = 50 µm.

Figure 2. Mature ovules before fertilization in *Amaranthus hypochondriacus*. A: Mature campylotropous ovule type before fertilization under light microscopy showing the polar nuclei (PN) and the synergid cells (Sy). B: Ovule at mature female gametophyte (FG) stage showing the egg cell (EC) and central cell (CC). Outer integument (OI), inner integument (II); micropyle (M), chalaza (CL) and funicle (F). Scale bars = 50 µm.
Female gametogenesis and early seed development of Amaranth

**Embryogenesis.** After the double fertilization, the initial stage is characterized by rapid cell division and the segregation of developmental programs in the descendants of the zygote (Figure 5) that form the embryo and the suspensor (Figure 6). The fertilized egg cell began to elongate and differentiate into the zygote. We observed that one of the synergids persists after fertilization (Figure 5A). The first division of the zygote was transverse (Figure 5B) and gave rise to a basal cell and an apical cell (Figures 5C, 1S), which became the suspensor and embryo, respectively. The suspensor underwent cell divisions to the four cell stage before the linear proembryo underwent its first division (Figures 5D, E). The endosperm and the perisperm were differentiated at this stage (Figures 5F, 1S, Table 1S), forming the structure that allows the growth of the embryo. The first embryo division was longitudinal, (Figures 5E, 6A, 6B, 3S), while the second division was transverse; the hypophysis was defined when the suspensor was at the seven-cell stage (Figures 6A, B, 1SA, 1SB, Table 1S).

Cell division and differentiation continued in the embryo proper, which developed into an early to late globular embryo (Figures 7A-E). Following the four-cell stage the embryo underwent another division to form the octant stage, and with this the early globular stage began (Figure 7A). Periclinal cell divisions were observed at the mid-globular stage, and the first divi-

<table>
<thead>
<tr>
<th>Development stage</th>
<th>Ovule length (mm)</th>
<th>SD</th>
<th>Ovule width (mm)</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Megasporegenesis and megagametogenesis</td>
<td>&lt; 0.123</td>
<td>0.042</td>
<td>&lt; 0.191</td>
<td>0.059</td>
</tr>
<tr>
<td>Mature ovule</td>
<td>0.263</td>
<td>0.044</td>
<td>0.299</td>
<td>0.055</td>
</tr>
<tr>
<td>From early globular embryo to early cotyledon stage</td>
<td>0.890</td>
<td>0.169</td>
<td>0.851</td>
<td>0.166</td>
</tr>
<tr>
<td>Seed in plant</td>
<td>1.628</td>
<td>0.089</td>
<td>1.515</td>
<td>0.058</td>
</tr>
<tr>
<td>Mature seed</td>
<td>1.392</td>
<td>0.053</td>
<td>1.352</td>
<td>0.050</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Inflorescence position</th>
<th>Branch position</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>ST1</td>
<td>ST2</td>
</tr>
<tr>
<td>Ovary length (mm) (OL)</td>
<td>0.947 a</td>
<td>1.029 a</td>
</tr>
<tr>
<td>Ovule length (mm) (OvL)</td>
<td>0.642 ab</td>
<td>0.693 a</td>
</tr>
<tr>
<td>Ovule width (mm) (OvW)</td>
<td>0.641 a</td>
<td>0.691 a</td>
</tr>
</tbody>
</table>

Values with different letters in a row are significantly different. Measurements at basal (ST1 or BR1), middle (ST2 or BR2) and apical (ST3 or BR3) position.
sion of the hypophysis occurred (Figure 7B). Then the hypophysis gave rise to two small, lens-shaped cells during the late globular stage of the embryo (Figure 7C). At the end of the globular stage the embryo has undergone periclinal and anticlinal cell divisions, and continued to the transition of the heart embryo stage (Figures 7D, E). The endosperm showed a helobial type, as we observed cellularized chalazal endosperm and the presence of a structure that perhaps functions as a scaffold to the positioning of the free nuclei (Figures 7F, 3S). There are regional specification processes that organize two main domains in the globular embryo; these domains or regions contribute to specific parts of the mature embryo.

**Figure 3.** Megasporogenesis-megagametogenesis in *Amaranthus hypochondriacus*. A-B: Early stages of megasporogenesis showing archespore cell (AR), C-D: meagaspore mother cell (MMC), (E) tetrad of megaspores (Me), (F) functional megaspore (FM) and degenerated megaspore cells (DM). G-H: Female gametophyte at two nuclei stage, I-J: four nuclei stage; nuclei are shown with arrows. Panels B, H and J are close-up of the rectangle in the panels A, G and I, respectively. Scale bars = 50 µm in A, C, E, G, I, 25 µm in B, H, J and 20 µm in D, F.
At the late globular stage there was a transition from a globular to a heart-shaped polarized embryo, which is caused by the differential growth of the apical domain to form the cotyledon primordia and the shoot meristem (Figures 8A, 1S, Table 1S). At the heart stage (Figure 8A) the embryo has already differentiated most of the tissues and cell types. During the torpedo stage the endosperm has been almost consumed (Figures 8B, C, D). At the early cotyledon stage (Figure 8E) the embryo showed well-differentiated shoot and root meristems. Although in most of the suspensors the cell divisions were ordered, in some cases they were disordered (Figures 7A-E, 8B, C).

At the maturation stage (Figures 8F, 1S, Table 1S) the embryo increased dramatically in size due to cell expansion and the accumulation of storage products, primarily in the cotyledons. The endosperm is almost consumed; however, the perisperm is maintained, perhaps maintaining nutrition and providing appropriate space for the growth of the mature embryo.

Discussion

Ovule and ovary. Despite apparent overall simplicity in architecture, ovules are composed of a number of clearly differentiated cell types (Figure 3S). Although the size of the ovary seems

Figure 4. Megagametogenesis in *Amaranthus hypochondriacus*. A: Cellularization at the chalaza pole (CL). B: Detail of mature female gametophyte with antipodals (AN), egg cell (EC), synergid (SY) polar nuclei (PN) and micropyle (M). C: Cellularization at the micropyle pole. D: Female gametophyte after degeneration of one synergid (DSY) showing the central cell (CC); arrowhead shows the structure that allows free nucleus migration. Scale bars = 50 µm in A, C, D and 25 µm in B.
not to be affected by the position on the branch or inflorescence, the ovule size is affected. Our analysis showed that ovule size is larger when it is formed at the basal or middle position of the inflorescence, suggesting a higher yield in these regions and perhaps the presence of more nutrients. The clearing analysis also showed that at first the *A. hypochondriacus* ovule is tenuinucellate. The MMC is originated from the L2 dermal layer and was localized at the L3 layer (Figure 3S).

**Figure 5.** Early-stage embryos in *Amaranthus hypochondriacus*. A: Detail of elongated zygote (Zy), endosperm (EN), synergid (Sy) and micropyle (M). B-C: One cell stage and differentiation of the basal cell (bc) and apical cell (ac) of embryo (E). D: One cell stage and suspensor (SUS) at the two cell stage of embryo. E: Embryo starts the first division and the suspensor at the three-cell stage. F: Linear embryo in the seed which shows the perisperm (PE). Scale bars = 50 µm in A, B, F, and 25 µm in C, D, E.

**Figure 6.** Early embryogenesis in *Amaranthus hypochondriacus*. A: Embryo (E) at two-cell stage. B: Embryo at 2-4 cell transition stage showing the suspensor (SUS) and hypophysis (hy). Scale bars = 50 µm.
Figure 7. Mid embryogenesis in *Amaranthus hypochondriacus*. A-B: Early globular stage embryo (E) and suspensor (SUS). C: Mid-globular stage embryo. D: Late globular stage embryo. E: Globular-heart transition stage embryo. F: Cellularized chalazal-endosperm (CE) and free nuclear-endosperm is shown by arrowheads. Scale bars = 25 µm in A, B, and 50 µm in C-F.

Figure 8. Late embryogenesis in *Amaranthus hypochondriacus*. A: Heart transition stage embryo. B-C: Heart-torpedo transition stage embryo. D: Torpedo stage embryo and general aspect of shoot apical meristem (SAM) and cotyledons (C). E: Cotyledon stage embryo with radicle (R). F: Mature embryo stage showing the main components: cotyledons, shoot apical meristem, radicle and perisperm (PE). Scale bars = 50 µm in A-F.
The development and form of the ovule in Amaranthaceae shows variations, as reported in *Celosia argentea*, *Amaranthus viridis*, *Cyathulatomentosa*, *Pupalia lappacea* and *Digeria arvensis* (Kajale 1935, 1940). During the early stages of ovule development their form is very similar, however, when fertilization occurs it assumes an ana-campylotropous form. In *Allmania nodiflora* the final form of the ovule is campylotropous, but it is derived from an amphitropous form. In *Alternanthera sessilis* the ovule begins to curve when the integuments are differentiated, and once the megaspores are formed it shows its typical campylotropous form (Bouman 1984). The ovule type in the Amaranthaceae quinoa (*Chenopodium quinoa* W) has been reported as amphitropous and campylotropous (Bouman 1984, López-Fernández & Maldonado 2013a, b).

The *A. hypochondriacus* ovule was reported as anatropous and crassinucellate during the early stages (Coimbra & Salema 1999). The nucellus is enveloped by two integuments, leaving only a straight opening at the apex; however, our results based on histological and clearing analysis showed that at the mature female gametophyte stage the ovule changes to campylotropous; this type of ovule has a curved body but its curvature is less than the anatropous ovule.

*Megasporogenesis and megagametogenesis in the Amaranthaceae.* The differentiation of the MMC is the first step in the transition from the sporophyte to the gametophyte phase of the plant life cycle (Guo & Zheng 2013). In *Digeria arvensis*, although the MMC usually produces a tetrad of haploid megaspores, in some cases it gives rise to 3 megaspores due to the suppression of division in the upper dyad cell, which is normal in *Alternanthera sessilis*; nevertheless, the chalazal megaspore is always the functional one and the development of the female gametophyte is the normal 8-nucleate type (Kajale 1935). The antipodals are always three in number and are persistent, very often up to the early stages of embryo development. Once fertilization is completed, the female gametophyte begins to elongate along one side of the antipodals and they are left in a lateral position (Kajale 1935, Joshi 1936). We observed that the MMC divides by meiosis to produce four haploid megaspore cells. Then, as happens in quinoa (Chehregani et al. 2009), the micropylar megaspore gives rise to the functional megaspore.

On the other hand, an increase in the number of antipodal cells has been observed in several families of the Centrospermales, including the Nyctaginaceae, Chenopodiaceae and Phytolaccaceae. The Amaranthaceae do not follow a rigid sequence of development in having only three antipodals (Kajale 1954). In *Pupalia lappacea* several antipodals are formed as a small mass of cells. During the early stages of embryo development the antipodals degenerate. Previous reports (Kajale 1954, Coimbra & Salema 1999) observed that in *A. hypochondriacus* the antipodal cells degenerate before the egg apparatus is differentiated. Nevertheless, in this study we were able to observe antipodal cells in mature ovules even after the degeneration of one synergid or fertilization, suggesting that amaranth female gametophytes in some conditions require the antipodal cells for longer periods.

*Embryology in the family Amaranthaceae.* Embryo development stages in many plants correspond approximately to the proembryo/suspensor stage, globular stage, heart stage, maturation and dormancy, and the root apical meristem is fully constituted when the cotyledons begin to be curved (García-Campayo et al. 2011, Bartoli et al. 2016, Zhao et al. 2017). In *Atriplex alimux* only some stages of embryo development were described (Talamali et al. 2007). In *Alternanthera sessilis* the first division in the fertilized egg is transverse. The next division is again transverse and takes place in the micropyle cell, and then more transverse divisions follow. It forms a horseshoe-shaped structure in the outer part of the seed (Joshi & Kajale 1937). Similarly, in *A. hypochondriacus* we observed that the basal cell divides transversally, giving rise to a row of seven to nine cells (suspensor), while the apical cell (the embryo proper) undergoes a longitudinal division first. The hypophysis was differentiated when the suspensor was at the five to seven-cell stage. As the embryo develops, it becomes greatly elongated and curved. Longitudinal and asymmetric divisions in the suspensor were also observed.

The nucellus of Amaranthaceae persists even after fertilization, becoming a main source of nutrients called perisperm (Prego et al. 1998, López-Fernández & Maldonado 2013a, b, Burrieza et al. 2014).

Joshi & Kajale (1937) observed that the structure of the endosperm of *Alternanthera sessilis*
differs from that of *Digera arvensis*. In *D. arvensis* the growth of the endosperm and the embryo destroys a part of the nucellus, but its central region remains intact. The endosperm becomes cellular in *Alternanthera*, but remains free nuclear at the chalazal end. In the mature seed most of the endosperm is absorbed by the growing embryo. Only a layer or two is left surrounding the radicle, as we observed in *A. hypochondriacus*. In *A. hypochondriacus* the endosperm showed a helobial development, as in the central region single nuclei were observed, but the chalazal endosperm was cellularized and at the torpedo stage the endosperm was almost consumed. Nevertheless, the perisperm is maintained, and thus acquires great importance in the nutritive value of the amaranth seed.

Finally, studies on micro and megagametogenesis in different species are an important prerequisite for developing robust hybrid seed production systems or attempting to alter reproduction (Moco & Marianth 2004, Salinas-Gamboa et al. 2016, Bartoli et al. 2017). The generation of information about reproductive development in *Amaranthus* species has biological and agronomic importance such as establishment of reproductive schedules, monitoring fertility problems and growth kinetics, which in turn may be the base for breeding programs and crop management. Our results show that the female reproductive development of amaranth is unique, since it can be defined as a variant of the Polygonum type considering that the micropylar megaspore is the functional one. These will also be useful to identify morphological changes during seed development and to develop new strategies to improve seed quality.

**Acknowledgments**

We are grateful for discussions and helpful comments on the manuscript from Jean-Philippe Vielle-Calzada and Humberto R. Medina. We acknowledge the Consejo Nacional de Ciencia y Tecnología (CONACYT, México) for the scholarship 175327 to ABL. This study was mostly performed using the facilities of Instituto Tecnológico de Celaya. We thank the Instituto Nacional de Investigaciones Forestales Agrícolas y Pecuarias (INIFAP) for providing facilities and the Convenio de Desempeño en Educación Superior Regional UTA-1401.

**Literature cited**


Huerta OJA, Barba de la Rosa AP. 2012. Caracterización bioquímica y estructural de las proteínas de


López-Fernández MP, Maldonado S. 2013b. Ricinosomes provide an early indicator of suspensor and endosperm cells destined to die during late seed development in quinoa (*Chenopodium quinoa*). *Annals of Botany* 112: 1253-1262. DOI:10.1093/aob/mct184


