

Research progress on the bulb expansion and starch enrichment in taro (*Colocasia esculenta* (L). Schott)

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Background. Taro is an important potato crop in the world, which can be used as food, vegetable, feed and industrial raw materials. The yield and quality of taro are mainly determined by the expansion degree of taro bulb and the filling condition of starch, while the expansion of taro bulb is a complex biological process. However, few information is reviewed on research progress on the bulb expansion and starch enrichment in taro.

Methodology. PubMed, Web of Science and China National Knowledge Infrastructure databases were searched for relevant articles. After removing duplicate articles and the articles with little relevance, 73 articles were selected for the review.

Results. This article introduces the formation and development of taro bulb for the workers engaged in taro research. The content includes the process of amyloplast formation at the cytological level and the changes in bulb expansion and starch enrichment at physiological levels, which involve endogenous hormones and key enzyme genes for starch synthesis. The effects of environment and cultivation methods on taro bulb expansion were also reviewed.

Conclusions. The future research directions and research focus about the development of taro bulb were prospected. Limited research has been conducted on the physiological mechanism and hormone regulation pathway of taro growth and development, taro bulb expansion, key gene expression, and starch enrichment. Therefore, the above research will become the key research direction in the future.

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12

13 Abstract

14 **Background.** Taro is an important crop in the world, and it can be used as food, feed, and
15 industrial raw material. The yield and product of taro are mainly determined by the expansion
16 degree and the state of starch enrichment. The expansion of taro bulb is a complex biological
17 process, which involves the increase in the number and volume of bulb cells and the starch
18 enrichment in the cells.

19 **Methodology.** PubMed, Web of Science and China National Knowledge Infrastructure
20 databases were searched for relevant articles. After removing duplicate articles and the articles
21 with little relevance, 73 articles were selected for the review.

22 **Results.** This article introduces the formation and development of taro bulb for the workers
23 engaged in taro research. The content includes the process of amyloplast formation at the
24 cytological level and the changes in bulb expansion and starch enrichment at physiological
25 levels, which involve endogenous hormones and key enzyme genes for starch synthesis. The
26 effects of environment and cultivation methods on taro bulb expansion were also reviewed.
27 Finally, the future research directions and research focus about the development of taro bulb
28 were prospected.

29 **Conclusions.** The development of taro bulbs is a complex biochemical process, including the
30 accumulation of morphogenesis and assimilation products, involving gene expression, material
31 metabolism, nutrient input, and the effect of external environmental conditions. At present, the
32 research on the development of taro mainly focuses on evolutionary classification, genotype and
33 isozyme analysis, cultivation, production, processing, and utilization. Limited research has been
34 conducted on the physiological mechanism and hormone regulation pathway of taro growth and
35 development, taro bulb expansion, key gene expression, and starch enrichment. There-fore, the
36 following research will become the key research direction in the future.

37

38 Introduction

39 Taro is an underground bulbous crop planted in the tropical and subtropical regions (Figure
40 1). It originated in China, India, Malaysia, and other regions, and is widely cultivated in Asia,
41 Africa, and other regions. **It has a cultivation history of more than 2,000 years.** At present, taro
42 can be classified using three methods. According to the ecological type, it can be divided into
43 aquatic taro and dry taro, and it can be classified into petiole, flower, and bulb taro according to
44 the eating part, and it can be classified into Kui taro, multi-cormels taro, and multi-head taro
45 according to bulb tillering habit (Wu et al., 2021). The three classifications of taros in agricultural
46 production of China exhibit various characteristics (Figure 2). Kui taro has few sub taro and
47 single mother-taro which is the main edible part. In multi-cormels taro, the sub taros are
48 numerous and grow in groups which have strong tillering ability. In multi-head taro, there are few
49 sub taros and the tillers of the mother taros grow in groups. The fundamental difference of taro

50 bulb tiller is the difference of expansion between mother and sub taro. This phenomenon is
51 caused by the difference in the number of chromosome and differential expression of genes (Zhu
52 et al., 2018). Studies on taro chromosomes has proved that Kui taro is diploid ($2n=2x=28$), while
53 the multi-head is triploid ($2n=3x=42$) (Huang et al., 2014). The chromosome multiples are related
54 to the geographical distribution. Diploid is more common in hot and humid areas with low
55 altitudes, and triploid is more common in dry and cold areas with high altitudes [4]. Taro is
56 essential for the body, and it can be used as food, vegetable, feed, and health product. The taro
57 bulb, which is rich in starch, carbohydrates, and energy, is one of the staple food in Asia, Africa,
58 and India (Njintang et al., 2008). In comparison with potato, sweet potato, and cassava, taro
59 starch granules are small and has a diameter of approximately $1.5\ \mu\text{m}$, making it easy to digest
60 and providing therapeutic and health care functions. It has been used for the preparation of infant
61 food in Haweei and some other islands in the pacific. Taro starch particles are small and uniform
62 and have good cold and hot stability, good whiteness, and good adhesion. Therefore, it can be
63 used as a brightener in cosmetics (Qi et al., 2015). Taro bulbs are rich in many nutrients, such as
64 starch, and the starch content of different varieties of taro could reach 10%–36% (Falade et al.,
65 2013). Taro contains a large amount of dietary fiber and various essential amino acids, but its fat
66 content is low (Sefa-Dedeh et al., 2002). It is also rich in vitamins and amino acid, making it is a
67 food for people of all ages (Han et al., 2018). Taro is a food and medicine homologous crop,
68 which is often used for the treatment of diarrhea, internal bleeding, asthma, and skin diseases
69 (Prajapati et al., 2011). It could also lower blood sugar and cholesterol (Sebnem et al., 2012).
70 Therefore, taro has good prospects for medicinal development. Taro starch can be used as an
71 industrial raw material. It could improve the stability of Pickering emulsion (Zhang et al., 2020),
72 and it could be used for the treatment of wastewater generated in the printing and dyeing process
73 (Zhou et al., 2018) and as an enhancer of starch film (Dai et al., 2015).

74 According to the statistics of the Food and Agriculture Organization of United Nations, taro
75 is the 14th largest vegetable crop, with a global planting area of 1.6 million hectares and an
76 annual output of 11.7 million tons. The cultivation area in China ranks first in the world and has a
77 large number of wild resources and local varieties, which are mainly distributed in southern
78 regions such as Yunnan, Taiwan, and the Yangtze River Basin. In recent years, taro has become
79 an important input and export trade industry. The total export trade volume and total value were
80 higher than the imports (Chang et al., 2019). With the rapid development of the taro industry, the
81 demand for taro has increased. It has also introduced new and high requirements for taro
82 production and basic theoretical research.

83 **Survey methodology**

84 PubMed, Web of Science and China National Knowledge Infrastructure databases were
85 searched for relevant articles. A total of 2525 articles appeared Pubmed database using “taro” as
86 the search term and the date of publication from 1975/1/1 to 2022/5/1. After narrowing the search
87 with keywords including “bulb of taro”, “bulb expansion in taro”, “bulb starch enrichment in
88 taro”, “amyloplast enrichment process of taro bulbs”, “development of taro bulbs”,
89 “regulation of hormones on bulb development”, “genes in starch synthesis”, and “regulation of
90 hormones on bulb swelling”, 1532 studies were obtained. With “taro” as the search term, 2525
91 articles published between 1975–2022 appeared in the Web of Science database, of which 914
92 articles were selected as above. After removing duplicate articles and the articles with little
93 relevance, 73 articles were selected for the review.

94

95 **Taro expansion and development process**

96 Taro is a perennial herbaceous plant, which belongs to the Araceae family, but it is generally
97 cultivated as an annual crop in agricultural production. According to the growth and development

98 characteristic of taro, its life circle is divided into five periods, namely, the germination, seedling,
99 growth, taro expansion, and bulb dormancy period (Sun et al., 2014). Taro bulbs undergo
100 metamorphosis during evolution. Take multi-cormels taro and multi-head taro for example. The
101 bulb of taro is composed of mother and sub taro, and some varieties include grandson and great
102 grandson taro (Figure 3). The mother taro of taro bulb is developed by the top bud of the sub taro.
103 The sub taro is developed from the lateral bud on the mother taro, and the grandson taro is
104 developed from the lateral bud of the sub taro. An axillary bud is present on each node of mother
105 taro, and it can develop into sub taro. The axillary bud grown in the leaf axil and the leaf have a
106 co-growth relationship. The growth site grows clockwise, and the angle between adjacent axillary
107 buds is 144° . The positions of each axillary bud are roughly in a straight line, showing typical 2/5
108 phyllodes features (Xu et al., 2020). The sub taro of Kui taro grows first and then thickens, and
109 the top swells to form a smaller sub-taro; the sub taro of taro with multi-cormels elongates and
110 thickens, and the whole bulb also swells to form a conical shape; three small-molecular-weight
111 proteins, namely, CSP2, CSP3 and CSP4, are present in the sub taro, which have spatiotemporal
112 specificity and are related to the occurrence and expansion of the sub taro. the relative differences
113 in the contents of CSP1, CSP2, and CSP4 in the mother taro are related to the apical dominance
114 of the terminal buds of the mother taro. The greater the difference is, the more obvious the apical
115 dominance and the lower the developmental degree of the sub taro is (Zhu et al., 2018). However,
116 Luiz found that the expression of TC1 gene during bulb development is spatiotemporal and acts
117 by encoding globulin Gd1, but TC1 gene is not a regulator gene of bulb expansion (Castro et al.,
118 1992). The gene that determines bulb expansion is present at the beginning of bulb
119 differentiation. The role of TC1 gene in the process of bulb expansion remains to be further
120 studied. The proportion of fresh weight of taro first increases and then decreases in the growth
121 process. Similarly, the water content of the bulbs first in-creases and then decreases, while the
122 dry matter content gradually increases, showing an S-shaped growth trend. The dry matter
123 accumulation of the sub taro in the later stage is greater than that of mother taro (Xu et al., 2020).

124

125 During the growth and development process of mother taro, the weight gradually increased,
126 the water content gradually decreased, the total starch content gradually increased, and the
127 soluble sugar content first increased and then decreased. The volume of parenchyma cells
128 continued to increase, and the internal starch content gradually increased. The number and
129 diameter of starch bodies showed an upward trend, and the number of starch granules in a single
130 starch body also increases (Zhu et al., 2017). During taro bulb expansion, the number of links on
131 the surface of the bulb gradually increased, the fresh and dry weight of a single bulb continued to
132 increase; the volume of the parenchyma cells in the bulb continued to increase, the number and
133 size of amyloplasts gradually increased, and the entire cell was enriched; with the expansion of
134 the bulb, the vascular tissue gradually developed and perfected, and the area of the sieve tube
135 increased in number and became irregularly arranged; the taro bulb was densely covered with
136 mucous cavities, which spread out from the center of the bulb, but no mu-cus cavity was
137 observed in the epidermal cells (Sheng et al., 2021).

138 **Amyloplast enrichment process of taro bulbs**

139 At present, the research on the development and proliferation of amyloplasts mainly focuses
140 on the endosperm cells of grains. In the development of endosperm amyloplasts, the number of
141 plastids continued to divide, and the number increased; then the number and volume of starch
142 granules increased in the plastid; individual starch granules increased in size and then began to
143 form small amyloplasts. As the development progressed, the envelope of the small amyloplasts
144 was further extended and expanded. Finally, the newly expanded envelope is filled with starch
145 granules, thereby forming the large amyloplasts. The amyloplasts of the large starch granules in

146 the wheat endosperm formed small starch granules by budding, while the amyloplasts of the
147 small starch granules proliferated by constriction or budding (Wei et al., 2002; Wei et al., 2008).

148 A few reports have focused on the proliferation mode of amyloplasts in under-ground
149 rhizome crops. The starch bodies of the complex starch contained in sweet potato tubers exist in
150 the form of “diplex”, “triad”, and “multiple”; for the growth mode of the amyloplasts, single grain
151 starch directly expands and grows; the amyloplasts of multigranular starch first split into
152 monomers, and then grow in the form of mono-mers; the amyloplasts of complex starch do not
153 divide during the growth process and form large complex amyloplasts (Jing et al., 2013).
154 According to the morphology of amyloids observed by scanning electron microscope, the cassava
155 root amyloid membrane is either "constricted", "wrinkled", or "sprouted", forming multiple
156 differentiation centers of amyloplasts; with the continuous expansion of the starch body, the
157 membrane structure was degraded and disappeared, and the irregular starch granules were
158 released (Min et al., 2010).

159 Zhu et al. (2018) divided the accumulation process of taro bulb starch into three stages,
160 namely, starch formation, rapid starch accumulation, and starch body enrichment (Zhu et al.,
161 2017). The specific performance is the formation of amyloplasts and the continuous increase in
162 diameter, followed by the continuous increase in the number of starch-es, and finally the increase
163 in the number of starch granules in the amyloplasts. The surface of taro starch body is mostly
164 round and oval, belonging to complex starch, containing multiple polyhedral starch granules.
165 Take Kui taro for example. in the early stage of development, amyloid was mostly distributed at
166 the edge of the cell, and then gathered at the center of the cell. Later in development, the number
167 and size of amyloplasts continued to increase (Figure 4). Transmission electron microscopy
168 showed that the amyloplasts of taro bulbs were large, and the large amyloplasts split into several
169 small starch granules. Some small starch granules (Figure 5A) were observed close to the free
170 state at the edge of amyloplasts, and these granules were loosely arranged and smaller in size.
171 With the development of the bulb, the starch granules with large diameters in the amyloplasts
172 were mostly concentrated in one area, and their arrangement was relatively compact. The small
173 starch granules were mostly distributed on the other side or around, and their arrangement was
174 loose (Figure 5B).

175 Taro bulb starch originates from the precursor plastid, and starch granules are formed in the
176 plastid and free in the cytoplasm of the membrane (Sheng, 2021). As the starch granules increase,
177 the volume of the plastids increases, and the starch granules develop from the original irregular
178 shape to a round shape. In the later stage, they gradually extruded each other into polygons. The
179 starch bodies of taro bulbs were mostly com-plex starches. Amyloid proliferation is divided into
180 two stages. In the first stage, the number of starch granules in the starch body changes; the starch
181 granules gradually increase, fill up, squeeze, and deform each other. The large starch granules
182 split into many small starch granules. The second stage involves the proliferation of amyloplasts.
183 Taro amyloplasts can be split via membrane constriction proliferation and capsular vesicle
184 proliferation. Capsule constriction and proliferation squeeze the starch gran-ules in the starch
185 body to both sides through the inward depression of the amyloplasts envelope. Further, it is
186 constricted into multiple amyloplasts, and the encapsulated vesicles proliferate in which the
187 original amyloplasts spit out vesicles from the envelope, and new amyloplasts are generated and
188 proliferated in the vesicles.

189 **Effects of environment and cultivation methods on the development of** 190 **taro bulbs**

191 Taro adapts to high-temperature and -humidity environment and is not resistant to low
192 temperature and frost. The optimum temperature for germination is 12–15 °C. The optimum
193 temperature for growth is generally 25–30 °C. If the temperature is very low, the growth of taro

194 will slow down or stop. If the temperature is very high, the condition will not be conducive to the
195 development and expansion of taro bulb. Different varieties of taro have different requirements
196 for temperature. Taro with Multi-cormels can be planted at low temperatures, making it widely
197 distributed in temperate zones. Kui taro has strict requirements for high temperature. Therefore,
198 Kui taro is mostly produced in tropical and subtropical regions with high temperature and
199 humidity (Chang et al., 2019). Taro requires sunlight, and the light saturation point is
200 approximately 50,000 lux. Taro is shade-tolerant and can grow under scattered light. The light
201 intensity, composition, and light time remarkably affect the growth of taro, but strong light is
202 conducive to the growth of taro and improves yield and quality. Under blue-violet light, the
203 leaves of taro are large and thick, and the petioles are thick and short, and this condition is
204 conducive to the growth and development of bulbs. In red and yellow light, the leaves are small,
205 and the petioles are slender, and this condition is not conducive to the growth and development of
206 bulbs. In the early stage of taro development, a longer light time is required to promote the
207 increase of leaf area and the accumulation of photosynthetic products. The later stage of taro
208 development requires a shorter light time to facilitate the formation and expansion of bulbs
209 (Chang et al., 2019). Taro requires dampness. Calla taro is grown in paddy fields, but dry taro
210 cannot be flooded for a long time. Taro has different requirements for humidity before different
211 growth stages. The field should be kept moist during the germination stage to induce the
212 germination of taro. During the taro-forming stage, the water demand is large, and the water
213 supply needs to be guaranteed; the soil should be kept dry before harvesting to maintain good
214 condition for the harvesting and storage of bulbs (Huang et al., 2016). Taro does not require very
215 strict soil texture. Loose and fertile soil with deep soil layer and convenient irrigation and
216 drainage is conducive to the growth of taro and the expansion of bulbs. Taro can grow normally
217 in soil with pH of 4.1–9.1, but the optimum pH is 5.5–7.0. A highly acidic or highly alkaline soil
218 is not conducive to the growth and development of taro (Huang et al., 2016).

219 Different cultivation methods remarkably affect the yield of taro. Film mulching can provide
220 soil temperature in the early stage of taro growth, which is beneficial to the growth of taro and the
221 expansion of bulbs. The growth and yield of taro bulbs in perforated film-covering cultivation
222 was better than that in ridge film-covering cultivation. However, considering the inconvenience
223 of cultivating a large amount of soil during the growth period of taro, ridge and perforation film-
224 covering cultivation easily form green taro, thus affecting the quality of bulbs and the taste of
225 eating (Wang et al., 2001). If no freezing damage is observed after emergence, early sowing is
226 conducive to the development of taro root. The plant height of taro increases, the number of taro
227 and taro increases, and the yield increases, but these changes only slightly affect the number of
228 taro and the shape index of taro. If the planting is late, the life cycle of the taro will be shortened,
229 which is not conducive to the growth and development of the bulb. This condition will lead to
230 insufficient bulb expansion, thereby reducing the yield (Zheng, 2008). Nitrogen and potassium
231 fertilizers have obvious effects on the yield and quality of taro, and potassium fertilizer has a
232 greater effect than nitrogen fertilizer. A significant interaction effect was observed among
233 nitrogen, potassium, and phosphorus fertilizer. Within the reasonable range of potassium and
234 nitrogen fertilizer application, the yield gradually increases with the increase of the amount of
235 fertilizer. Excessive fertilizer application will reduce the yield. Phosphate fertilizer alone only
236 slightly affects the development of taro bulbs, and no obvious rule has been established.
237 Reasonable fertilization is beneficial to the growth, development, and yield increase of taro bulbs
238 (Song et al., 2004).

239 **Regulation of hormones on bulb development**

240 **Regulation of endogenous hormones on bulb expansion and starch enrichment**

241 Hormones is an important endogenous substance that regulates plant growth and is a key
242 factor in bulb formation (Durbak et al., 2012). Some genes and proteins related to bulb for-

243 mation are also closely related to plant hormone signaling pathways (Aksenova et al., 2012).
244 Different plant hormones have different functions in bulb expansion, and gibberellin (GA) can
245 inhibit or delay tuber formation (Vreugdenhil et al., 1989). Abscisic acid (ABA) does not
246 participate in the induced metamorphosis process of tubers, but it counteracts the antagonism of
247 other hormones (Shu et al., 2017). Auxin (IAA) can promote the metamorphic development of
248 tubers and promote plant root development, and its concentration affects tuberous root thickening
249 (Wang et al., 2006). Although GA and ABA are not directly related to tuber formation, they are
250 related to the ratio of GA3/ABA. The balance of “inducing substances” and “inhibiting
251 substances” is a key factor for tuber formation (Liu, 2001). Cytokinins (CTK) are mainly in-
252 volved in the formation of tubers. Matsuo and Mitsuzono (1988) reported that the content of
253 zeatin riboside (ZR) is significantly positively correlated with the formation and thickening of
254 sweet potato tubers (Matsuo et al., 1988). The overexpression of CTK synthesis gene *ipt* in potato
255 could form more tubers (Tao et al., 2010). IAA-related genes such as ARFs and Aux/IAAs are
256 specifically expressed in early tuber development (Kloosterman et al., 2008). IAA and GA3 are
257 necessary for potato stolon elongation. ABA and jasmonic acid (JA) are positive regulators for
258 inducing tuber formation. GA3 is a negative regulator (Liu et al., 2019). JA and methyl jasmonate
259 (MeJA), as classes of plant growth regulators, play an important role in tuber and bulb formation
260 (Sarkar et al., 2006).

261 Plant endogenous hormone regulation is closely related to starch anabolism (Kim et al.,
262 2005). The enlargement of plant bulbs mainly depends on starch accumulation and cell divi-sion
263 enlargement, and starch accumulation mainly depends on sucrose synthesis and transportation.
264 Plant hormone signal transduction affects starch accumulation. ABA can induce the expression of
265 starch synthesis genes and enhance the transduction of sugar signals to promote starch
266 accumulation (Akihiro et al., 2005). The level of GA at the grain filling stage of wheat is
267 positively correlated with the final grain yield and starch yield. GA plays an important role in
268 starch accumulation in wheat grains. Changes in endogenous hormone levels may indirectly
269 affect starch accumulation in grains by affecting regulatory enzymes and regulatory processes
270 (Xie et al., 2003). Scientists added IAA to MS medium, and the potato tuber starch content and
271 starch granule size increased by 15%–30% (Gukasyan et al., 2005). In the study of tulip bulbs,
272 IAA and ZR indirectly promoted starch accumulation by increasing the activity of ADP-glucose
273 pyrophosphorylase (AGPase), thus catalyzing the production of a large number of products.
274 Endogenous hormones may promote starch accumulation by participating in the starch synthesis
275 pathway (Miao et al., 2016). Hormones have multiple roles and interact to form a regulatory
276 network, thereby regulating tuber development (Jung et al., 2013). In the early stage of
277 development, the content of endogenous hormones ABA, Z, and ZR showed an upward trend,
278 while the content of IAA, GA3, and JA showed a downward trend. In the later stage of
279 development, the content of endogenous hormones ABA, IAA, Z, and GA3 showed an upward
280 trend. The content of ZR and JA showed a downward trend, but the contents of IAA, GA3, and
281 JA hormones were generally high during the whole development process (Sheng, 2021). Other
282 related studies on endogenous hormones on the growth and development of taro bulbs and starch
283 enrichment have not been conducted.

284 **Regulation of growth, development, and starch enrichment of bulbs by exogenous** 285 **hormones**

286 The effect of exogenous hormones on the rhizome expansion of potato, sweet potato, and
287 other potato crops has been studied. Yang (2005) used four auxins to spray potatoes (Yang,
288 2005). The results showed that the four auxins increased plant height and stem diameter and
289 prolonged the photosynthetic accumulation in the later stage. This condition allowed the tubers to
290 accumulate more organic matter during the expansion stage, thereby substantially increasing the
291 yield. The exogenous application of IAA can promote the formation of potato stolons and the

292 development of tubers. It is achieved by accelerating starch accumulation and starch granule
293 enlargement, which are beneficial to the formation and development of tubers (Gukasyan et al.,
294 2005; Roumeliotis et al., 2012). GA also promotes the occurrence of stolons. In the present
295 experiment, stolons appeared on the second day after the medium containing GA3 and IAA was
296 added, and the occurrence continued throughout the tuber setting period (Lian et al., 2002).
297 However, the addition of GA alone could inhibit or delay the formation of potato tubers, and
298 inactivation of the active GA gene could promote potato tuber formation (Xu et al., 1998;
299 Roumeliotis et al., 2012). Treatment with exogenous GAs inhibited SS and SSS activity, thus
300 decreasing the sucrose and starch contents in tubers (Vreugdenhil et al., 1999). ABA is a
301 promoting factor for the formation of potato tubers, and timely spraying is beneficial to potato
302 formation (Krauss et al., 1982; Garcia et al., 2014). Varying results have been obtained about the
303 role of ABA in the development of tuber plants. GA3 inhibits the formation of potatoes in vitro,
304 while ABA promotes its tuber formation (Hu et al., 2017). Exogenously applied ABA can
305 promote tuber expansion (Xu et al., 2022). However, in vitro, ABA cannot make stolons
306 metamorphose into tubers smoothly (Yang, 2005). ABA does not participate in the induced
307 metamorphosis process of tubers, but its presence counteracts the respective physiological ac-
308 tivities of other hormones (Xu et al., 2022). CTK can promote potato tuber development, regulate
309 the balance between source and sink, and participate in the transport of nutrients to storage
310 organs (Roitsch et al., 2000). When a certain concentration of CTK is applied exogenously, the
311 biomass of tubers remarkably increases, and the transformation of stolons to tubers is accelerated
312 (Romanov, 2009). In vitro, CTK inhibits sucrose invertase activity but activates phosphorylase
313 and AGPase, thereby promoting starch accumulation (Zhu et al., 2016). Therefore, CTK is an
314 important factor in inducing tuber formation (Quan et al., 2002). The exogenous application of
315 JA and its derivatives can induce the swelling of the stolon top, and the content of endogenous JA
316 increases during this process (Abdala et al., 2002). After exogenous JA treatment, the in-
317 tracellular sucrose accumulates, thus increasing the osmotic pressure of the cell wall, changing
318 the structure of the cell wall, and increasing the cell ductility. More polysaccharides such as
319 cellulose, hemicellulose, and pectin accumulate, indicating that JA controls the expansion of the
320 cell by regulating the synthesis of intracellular sugar (Takahashi et al., 1995). This phenomenon
321 induces the formation of the apical meristem of potato stolon and promotes tuber development
322 (Cenzano et al., 2003).

323 The development of taro bulbs is remarkably affected by exogenous hormones. The
324 diameter of taro bulbs that were irrigated with auxin increased significantly, the weight increased,
325 and the filling degree of amyloplasts in parenchyma cells increased. Low concentration of 6-BA
326 can promote the development of bulbs, but it is not conducive to the enrichment of amyloplasts,
327 and high concentrations of 6-BA have a certain inhibitory effect on the development of bulbs.
328 GA3 promotes the elongation of the petioles of taro plants, but it does not promote the expansion
329 of the bulbs. High concentrations of GA3 (100–200 mg/L) have an inhibitory effect on the
330 development of taro bulbs, but it promotes the development of taro and increases the number of
331 taro (Sheng, 2021). Limited studies have focused on the effects of exogenous hormones on the
332 development of taro bulbs and their enrichment of amyloplasts, and further research is need-ed.

333 **Role of key enzyme genes in starch synthesis in starch enrichment**

334 In crops mainly harvesting underground storage organs, the synthesis and accumulation of
335 starch is a complex physiological and biochemical process, which is the result of the synergistic
336 interaction of multiple enzymes. The key enzymes of starch synthesis in root crops, such as
337 potato and lotus root, have been widely studied. The changes in AGPase and soluble starch
338 synthase (SSS) activities have important effects on starch synthesis in potato tubers (Tang, 2015).
339 However, SS and AGPase can remarkably promote the synthesis of starch in the process of lotus
340 root rhizome expansion, and their activities affect the starch content of lotus root rhizomes at the

341 mature stage (Li et al., 2006). Based on the study of substance accumulation and changes in
342 related enzyme activities during the development of yam, sucrose phosphate synthase activity
343 plays a key regulatory role in the development of yam tubers and is closely related to the main
344 functional substances (Liang et al., 2011). Based on the study of taro bulbs, AGPase activity is
345 positively correlated with total starch content (Zang et al., 2016). With the gradual deepening of
346 the re-research on starch metabolism pathways, people have new understanding of the key enzyme
347 gene sequences and related expression regulators in the pathway.

348 In sweet potato, the key enzyme genes of starch synthesis such as AGPase and SS have been
349 cloned into the gene sequence. The expression and regulation of these genes have been studied,
350 and these key enzyme genes play a key role in the sweet potato starch metabolism pathway (Tang
351 et al., 2011). The genes controlling sweet potato starch synthesis include granule bound starch
352 synthase (GBSS) gene I, SSS genes I and II, starch branching enzyme (SBE) genes I and II,
353 starch de-branching enzyme (DBE) gene, AGPase gene A/B/C, sucrose synthase (SS) genes I and
354 II, and isoamylase (ISA) gene (Kim et al., 2009).

355 GBSS I is a key enzyme that controls starch synthesis, and it catalyzes the synthe-sis of
356 amylose. Otani et al. (2007) interfered with the expression of GBSS I by RNAi technology to
357 make sweet potato taste more glutinous. SSSII can affect the structure of amylopectin and reduce
358 the gelatinization temperature of starch (Otani et al., 2007). The reduction of starch gelatinization
359 temperature is conducive to simplifying the starch hydrolysis process and reducing the
360 production cost of starch fermentation (Takahata et al., 2010). AGPase im-proves the starch
361 content of potato tubers (Song et al., 2005). However, the synergistic expression of starch
362 synthesis-related genes under exogenous sucrose treatment promotes the con-version of sucrose
363 to starch (Ahn et al., 2010). Peak synthase has been widely studied, but no direct research has
364 been conducted on taro starch synthase.

365 **Conclusions and future direction**

366 The development of taro bulbs is a complex biochemical process, including the
367 accumulation of morphogenesis and assimilation products, involving gene expression, material
368 metabolism, nutrient input, and the effect of external environmental conditions (Figure 6).
369 Limited studies have been conducted locally and abroad. Under-standing the development
370 process, expansion mechanism, and regulation mechanism of taro bulbs has a guiding role in the
371 production of taro and is important to ensuring food security and responding to food crises. At
372 present, the research on the development of taro mainly focuses on evolutionary classification,
373 genotype and isozyme analysis, cultivation, production, processing, and utilization. Limited
374 research has been conducted on the physiological mechanism and hormone regulation pathway of
375 taro growth and development, taro bulb expansion, key gene expression, and starch enrichment.
376 Therefore, the following research will become the key research direction in the future.

377 **Expansion of taro bulbs and the regularity of the development and spatial 378 distribution of starch bodies**

379 Starch is the main storage material of taro bulbs, and amyloplasts are the orga-nelles that
380 synthesize and accumulate starch. The development of amyloplasts deter-mines the yield and
381 quality of taro. Limited studies locally and abroad have focused on the development of taro corm
382 and amyloplast, and they remain in the preliminary stage. The fine structure observation of
383 amyloplast development, its proliferation mode, and the spatial distribution characteristics of taro
384 corm amyloplast are not clear. In the future, the occurrence, division, proliferation, and
385 enrichment of amyloplasts in parenchyma cells of different types of taro bulbs, the differences of
386 physical and chemical properties of taro starch at different development stages, and the develop-
387 ment and enrichment characteristics of amyloplasts in different spatial parts should be focused
388 on.

389 **Role of key enzyme genes in starch synthesis in starch enrichment**

390 Starch is the main storage material of taro bulbs. The expansion process of taro is closely
 391 related to the synthesis of starch, and the genes related to starch synthesis are closely related to
 392 starch synthesis, which directly determine the starch content of taro. The research on starch
 393 synthase gene has remarkably progressed in wheat, rice, potato and other crops, but limited
 394 research has been conducted on taro starch synthase gene. Therefore, the differences in the
 395 expression of key enzyme genes (e.g., AGPase, GBSS, SSS, and SBE) for taro starch synthesis,
 396 the roles of these genes in regulating the starch enrichment process, and the exploration of
 397 individual gene functions will become the focus of research.

398 **Regulation of hormones on bulb swelling and starch enrichment**

399 The expansion of taro bulbs mainly depends on the increase in the number and volume of
 400 parenchyma cells, and this process results from the synergistic action of various hormones,
 401 especially IAA, GA, CTK, and other hormones. The changes of hormones during bulb
 402 development, the relationship between hormones and bulb expansion, and the relationship
 403 between the expression of hormone synthesis genes and signal transduction-related genes and
 404 taro starch enrichment need to be investigated.

405 **Hormone-regulated pathways promoting bulb expansion and starch enrichment**

406 Exogenous hormones and plant growth regulators have important regulatory effects on taro
 407 bulb swelling, starch enrichment, and yield increase. The effects of exogenous substances such as
 408 6-BA, 2,4-D, GA₃, PP333, and 5-aminolevulinic acid (5-ALA) on the development, yield, and
 409 quality of taro corm, as well as the type and concentration of the best exogenous hormone to
 410 promote corm expansion and starch enrichment need to be studied, and plant growth regulators
 411 for increased yield and improved quality of taro should be developed to provide an important
 412 theoretical basis for taro production.

413 In a word, the development of taro bulbs still requires a lot of research. With the deepening
 414 of research and the solution of key problems, the production of taro will continue to improve, and
 415 the development and utilization of taro will be more efficient.

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421 The authors declare there are no competing interests.

422 **Author Contributions**

423 E.J.Z., F.X. and W.Y.S. conceived the outline of the manuscript.

424 E.J.Z. and F.X. wrote the manuscript.

425 W.Y.S., W.J.J., W.L.L., X.R.Y., X.P.W. provided revisions.

426 All authors read and approved the final manuscript.

427 **Data Avail**

428 The following information was supplied regarding data availability:

429 There is no raw data or code in this literature review.

430 **Ability Statement**

431 All data were collected from the published research papers.

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436 **References**

437 **Abdala G, Castro G, Miersch O, Pearce, D. 2002.** Changes in jasmonate and gibberellin levels
 438 during development of potato plants (*Solanum tuberosum* L.). *Plant Growth Regulation*.36:121-
 439 126.

- 440 **Ahn YO, Kim S.H, Kim CY, Lee JS, Kwak S, Lee H. 2010.** Exogenous sucrose utilization and
441 starch biosynthesis among sweetpotato cultivars. *Carbohydrate Research*. 345: 55-60.
- 442 **Akihiro T, Mizuno K, Fujimura F. 2005.** Gene expression of ADP-glucose pyrophosphorylase and
443 starch contents in rice cultured cells are cooperatively regulated by sucrose and ABA. *Plant Cell*
444 *Physiology*. 46:937-946.
- 445 **Aksenova NP, Konstantinova TN, Golyanovskaya SA, Sergeeva LI, Romanov GA. 2012.**
446 Hormonal regulation of tuber formation in potato plants. *Russian Journal of Plant Physiology*.
447 59:451-466.
- 448 **Castro LAB. 1992.** Spatial and temporal gene expression patterns occur during corm development.
449 *Plant Cell*. 4:1549-1559.
- 450 **Cenzano A, Vigliocco A, Kraus T, Abdala G. 2003.** Exogenously applied jasmonic acid induces
451 changes in apical meristem morphology of potato stolons. *Annals of Botany*. 91:915-919.
- 452 **Chang L, Wang X. 2019.** Overview of Development Status of Taro Industry in the World. *Modern*
453 *Agricultural Science and Technology*. 2:57-59.
- 454 **Dai L, Qiu C, Xiong L, Sun Q. 2015.** Characterisation of corn starch-based films reinforced with
455 taro starch nanoparticles. *Food Chemistry*. 174:82-88.
- 456 **Durbak A, Yao H, McSteen P. 2012.** Hormone signaling in plant development. *Current Opinion in*
457 *Plant Biology*.15:92-96.
- 458 **Falade KO, Okafor CA. 2013.** Physicochemical properties of five cocoyam (*Colocasia esculenta* and
459 *Xanthosoma sagittifolium*) starches. *Food Hydrocolloids*. 30:173-181.
- 460 **Garcia MNM, Stritzler M, Capiati DA. 2014.** Heterologous expression of Arabidopsis ABF4 gene in
461 potato enhances tuberization through ABA-GA crosstalk regulation. *Planta*. 239:615-631.
- 462 **Gukasyan IA, Golyanovskaya SA, Grishunina EV, KKonstantinova TN, Romanov GA. 2005.**
463 Effect of Rol Transgenes, IAA, and Kinetin on Starch Content and the Size of Starch Granules in
464 Tubers of In Vitro Potato Plants. *Russian Journal of Plant Physiology*. 52:809-813.
- 465 **Han X, Zhang DX, Wang L, Li Q. 2018.** Research Progress on the Nutrition Components and
466 Processing and Utilization of Taro. *China Fruit Vegetable*. 38:9-13.
- 467 **Hu Y, Zhan S, Ku WZ, Wang R, Xiao L. 2017.** The Roles of ABA and GA3 in the Joint Regulation of
468 Potato Tuber Formation and Starch Accumulation in Vitro. *Mol. Plant Breeding*.15:4210-4214.
- 469 **Huang XF, Peng J, Ke WD, Liu YM, Sun YL, Dong HX, Li F, Liu YP, Zhu HL, Li SM, Ye YY,**
470 **Huang LC, Li MH, Wang Y, Zhong L, Zhou K. 2014.** Analysis on Quality Characters of 206
471 Taro [*Colocasia esculenta* (Linn.)Schott] Germplasm Resources. *Journal of Plant Genetic*
472 *Resources*. 15:519-525.
- 473 **Huang XF, Ke WD, Sun Y. 2016.** High yield and high efficiency cultivation of high quality taro, 1st
474 ed.; China Agriculture Press Co, China, pp.22-30.
- 475 **Jing YP, Li DL, Liu DT, Yu XR, Hu ML, Gu YJ, Wang Z. 2013.** Anatomical Structure of the
476 Tuberous Root Growth and Its Amyloplast Development in Sweet Potato. *Acta Botanica Boreali-*
477 *Occidentalia Sinica*.33:2415-2422.
- 478 **Jung JKH, McCouch S. 2013.** Getting to the roots of it: genetic and hormonal control of root
479 architecture. *Frontiers Plant Science*.4:186.
- 480 **Kim KJ, Kim KS. 2005.** Changes of endogenous growth substances during bulb maturation after
481 flowering in *Lilium oriental* hybrid 'Casa Blanca'. *Acta Horticulturae*. 570:661–667.
- 482 **Kim T, Goo Y, Lee C, Lee B, Bae, J, Lee S. 2009.** The sweetpotato ADP-glucose pyrophosphorylase
483 gene (ibAGP1) promoter confers high-level expression of the GUS reporter gene in the potato
484 tuber. *Comptes Rendus Biologies*. 332:876-885.
- 485 **Kloosterman B, Koeyer DD, Griffiths R, Flinn B, Bachem C. 2008.** Genes driving potato tuber
486 initiation and growth: identification based on transcriptional changes using the POCI array.
487 *Functional and Integrative Genomics*. 8:329-340.
- 488 **Krauss A, Marschner H. 1982.** Influence of nitrogen nutrition d aylength and temperature on
489 contents of gibberellic and abscisic acid and on tuberization in potato plants. *Potato Research*.
490 25:13-21.
- 491 **Li LJ, Zhang XD, Pan EC, Sun L, Xie K, Li G, Cao BS. 2006.** Relationship of Starch Synthesis with
492 It's Related Enzymes' Activities During Rhizome Development of Lotus. *Scientia Agricultura*
493 *Sinica*. 39:2307-2312.
- 494 **Liu J. 2001.** Formation Mechanism of Potato Tuber in Vitro and Regulation of Tuber Formation.
495 Doctor's Thesis, Huazhong Agric. University, Wuhan, Hubei, China.
- 496 **Liu D, Xu M, Hu Y, Wang R, Tong J, Xiao L. 2019.** Dynamic Changes of KeyPlant Hormones
497 during Potato Tuber Development. *Mol. Plant Breeding*. 17:1998-2003.
- 498 **Lian Y, Zou Y, Dong HR, Jin LP, Lin H. 2002.** The Changes of Several Endogenous Phytohormones
499 during Microtuber Formation in vitro in *Solanum tuberosum* L. *Acta Horticulturae Sinica*,
500 29:537-541.
- 501 **Liang RF, Li CZ, Zhang J, He LF, Wei BH, Gan XQ, He HY. 2011.** Changes of Matter
502 Accumulation and Relative Enzymatic Activity during Yam Tuber Development. *Acta Agronomica*
503 *Sinica*. 37:903-910.
- 504 **Matsuo T, Mitsuzono H. 1988.** Variation in the levels of major free cytokinins and free abscisic acid
505 during tuber development of sweet potato. *J. Plant Growth Regul.*7:249-258.

- 506 **Min Y, Yao Y, Wang J, Hu XW, Guo JC. 2010.** Observation on the structure and development of
507 amyloplast in early stage of cassava storage root with scanning electronic microscope. Journal of
508 Chinese Electron Microscopy Society. 29:379-384.
- 509 **Miao YY, Zhu ZB, Guo QS, Yang XH, Liu L, Sun Y, Wang CL. 2016.** Dynamic changes in
510 carbohydrate metabolism and endogenous hormones during *Tulipa edulis* stolon development
511 into a new bulb. Journal of Plant Biology. 59,121-132.
- 512 **Njintang YN, Scher J, Mbofung CMF. 2008.** Physicochemical, thermal properties and
513 microstructure of six varieties of taro (*Colocasia esculenta* L. Schott) flours and starches. Journal
514 of Food Engineering. 86:294-305.
- 515 **Otani M, Hamada T, Katayama K, Kitahara K, Kim S, Takahata Y, Suganuma T, Shimada T.
516 2007.** Inhibition of the gene expression for granule-bound starch synthase I by RNA interference
517 in sweet potato plants. Plant Cell Reports. 26:1801-1807.
- 518 **Prajapati R, Kalariya M, Umbarkar R, Parmar S, Sheth N. 2011.** *Colocasia esculenta*: A potent
519 indigenous plant. International Journal of Nutrition Pharmacol Neurological Diseases. 1:90-96.
- 520 **Qi HL, Yin ZY, Zheng XX. 2015.** Study Progress on Starches of Root and Tuber Plants. Journal of
521 Chongqing Technol Business University. (Nat Sci Ed).32:68-71.
- 522 **Quan F, Zhang AX, Cao XW. 2002.**The Role of Plant Hormones in the Formation and Development
523 of Potato Tubers. Chinese Potato Journal. 1:29-32.
- 524 **Roumeliotis E, Visser RG, Bachem CW. 2012b.** A crosstalk of auxin and GA during tuber
525 development. Plant Signaling Behavior. 7:1360-1363.
- 526 **Roumeliotis E, Kloosterman B, Oortwijn M, Kohlen W, Bachem C. 2012a.** The effects of auxin
527 and strigolactones on tuber initiation and stolon architecture in potato. Journal of Experimental
528 Botany.63:4539-4547.
- 529 **Roitsch T, Ehneß R. 2000.** Regulation of source/sink relations by cytokinins. Plant Growth
530 Regulation. 32:359-367.
- 531 **Romanov GA. 2009.** How do cytokinins affect the cell. Russian Journal of Plant Physiology. 56:268-
532 290.
- 533 **Sarkar D, Pandey SK, Sharma S. 2006.** Cytokinins antagonize the jasmonates action on the
534 regulation of potato (*Solanum tuberosum*) tuber formation in vitro. Plant Cell, Tissue and Organ
535 Culture. 87:285-295.
- 536 **Sefa-Dedeh S, Agyir-Sackey EK. 2002.** Starch structure and some properties of cocoyam
537 (*Xanthosoma sagittifolium* and *Colocasia esculenta*) starch and raphides. Food Chem. 79:435-
538 444.
- 539 **Sebnem S, Sedef NE. 2012.** Production of resistant starch from taro (*Colocasia esculenta* L. Schott)
540 corm and determination of its effects on health by in vitro methods. Carbohydrate Polymers.
541 90:1204-1209.
- 542 **Sheng JY. 2021.** Corm Development of Taro and Its Response to Exogenous Hormones. Master's
543 Thesis, Yangzhou University, China.
- 544 **Shu K, Zhou WG, Yang WY. 2017.** APETALA 2-domain-containing transcription factors: focusing on
545 abscisic acid and gibberellins antagonism. New Phytologist. 217:977-983.
- 546 **Song CF, Xu K. 2004.** Absorption and distribution of nitrogen, phosphorus and potassium in taro.
547 Plant Nutrition and Fertilizing Science.10:403-406.
- 548 **Song BT, Xie CH, Liu J. 2005.** Expression of potato sAGP gene and its effects on contents of starch
549 and reducing sugar of transgenic potato tubers. Entia Agricultura Sinica. 38:1439-1446.
- 550 **Sun JD, Sun XM, Yuan ZZ. 2014.** Taro Cultivation, 1st ed.; China Agricultural Press, China, pp.34-
551 36.
- 552 **Tao GQ, Letham DS, Yong JWH, Zhang K, John PCL, Schwartz O, Wong C, Farquhar GD.
553 2010.** Promotion of shoot development and tuberisation in potato by expression of a chimaeric
554 cytokinin synthesis gene at normal and elevated CO₂ levels. Functional Plant Biology.37:43-54.
- 555 **Tang HL. 2015.** The Activity Changes of Key Potato Starch Synthase Effects on the Starch Content of
556 Tuber. Northeast Agric. Univ. China.
- 557 **Tang W, Li Q, Zhang YG, Wang X, Hou M, Ma DF. 2011.** Advances on Structure and Function of
558 Key Enzymes and Genes Involved in Starch Biosynthesis in Sweetpotato. Mol. Plant Breeding.
559 9:1479-1485.
- 560 **Takahata Y, Tanaka M, Otani M, Katayama K, Kitahara K, Nakayachi O, Nakayama H,
561 Yoshinaga M. 2010.** Inhibition of the expression of the starch synthase II gene leads to lower
562 pasting temperature in sweetpotato starch. Plant Cell Reports. 29:535-543.
- 563 **Takahashi K, Fujino K, Kikuta Y, Fujino K. 1995.** Involvement of the accumulation of sucrose and
564 the synthesis of cell wall polysaccharides in the expansion of potato cells in response to jasmonic
565 acid. Plant Sci. 111:11-18.
- 566 **Vreugdenhil D, Struik PC. 1989.** An integrated view of the hormonal regulation of tuber formation
567 in potato (*Solanum tuberosum*). Physiologia plantarum. 75:525-531.
- 568 **Vreugdenhil D, Sergeeva LI. 1999.** Gibberellins and tuberization in potato. Potato Research.
569 42:471-481.
- 570 **Wang WH, Li M, Song BC, Liu CL, Zhang XX. 2001.** Effect of Different Culture Way on Growth
571 and Yield of *Colocasia esculenta* Schott. Journal of Laiyang Agricultural College. 18:12-14.

- 572 **Wang QM, Zhang LM, Guan YA, Wang ZL. 2006.** Endogenous Hormone Concentration in
573 Developing Tuberous Roots of Different Sweet Potato Genotypes. *Agric. Sci. China.* 5:919-927.
- 574 **Wei CX, Lan SY, Xu ZX. 2002.** Relation between envelope and proliferation of amyloplast in
575 endosperm cell of rice. *Journal of Chinese Electron Microscopy Society.* 21:123-128.
- 576 **Wei CX, Zhang J, Zhou WD, Chen YF, Xu RG. 2008.** Ultrastructural Observation on the
577 Development of Small Starch Granule (Compound Starch Granule) of Wheat Endosperm. *Journal*
578 *of Triticeae Crops.* 25:804-810.
- 579 **Wu W, Chang QT, Wang A. 2021.** Research Progress of *Colocasia esculenta* Germplasm Resources
580 in China. *Anhui Agricultural Science.* 49:4-7.
- 581 **Xu RC, Yang MY, Xu JP, Liu SD. 2020.** Preliminary Study on Dynamic Characteristics of Biological
582 Characters in Growth and Development of Taro (*Colocasia esculenta*). *Acta Agricultural. Jiangxi.*
583 32: 44-48.
- 584 **Xu X, Lammeren AM, Vermeer E, Vreugdenhil D. 1998.** The role of gibberellin, abscisic acid, and
585 sucrose in the regulation of potato tuber formation in vitro. *Plant Physiol.* 117:575-584.
- 586 **Xu Z, Zhen LW, Zhang Y, Zhang SS, Guo DP. 2022.** Regulatory network of tuberization in potato
587 (*Solanum tuberosum* L.). *Plant Physiology Journal.* 58:109-118.
- 588 **Xie ZJ, Jiang D, Cao WX, Dai TB, Jing Q. 2003.** Relationships of endogenous plant hormones to
589 accumulation of grain protein and starch in winter wheat under different post-anthesis soil water
590 statuses. *Plant Growth Regul.* 41:117-127.
- 591 **Yang LJ. 2005.** Plant Growth Substances and Their Application in Potato. *Heilongjiang Agricultural*
592 *Science.* 1:49-50.
- 593 **Zang YW, Jiang FL, Cheng YQ, Kong XY, Wu Z. 2016.** Study on Dynamic Changes of the Major
594 Carbohydrate Content and the Related Enzyme Activities during the Microcorm Development of
595 *Colocasia esculenta*. *Acta Botanica Boreali-Occidentalia. Sinca.* 36:700-705.
- 596 **Zhang GM, Zhang DX. 1990.** The relationship between geographic distribution and ploidy level of
597 taro. *Euphytica.* 47:25-27.
- 598 **Zhang LL, Zhu YZ, Li DP, Jiang Y, Dong B, Li F. 2020.** Effect of Heat-Modified Starch from
599 Laiyang Taro on the Formation and Stability of Emulsions. *Food Sci.* 41:51-57.
- 600 **Zheng F. 2008.** Effects of Different Sowing Dates and Fertilization Conditions on the Growth and
601 Yield of Red Bud Taro. *Shanghai Vegetables.* 5:69-71.
- 602 **Zhou J, Li DF, Xu J, Wang LL, Qiu YX. 2018.** Adsorption of Cationic Taro Starch for Removal of C.
603 I. Direct Violet 1 and C. I. Disperse Blue 56 in Wastewater. *Natural Product Research and*
604 *Development.* 30:1078-1084.
- 605 **Zhu XW, Cui WX, Zhang EJ, Wang LL, Yu XR, Xiong F. 2018.** A Comparative Study on
606 Morphological and Physicochemical Properties of Starches Isolated from Mother and Filial Bulbs
607 of 'Binglang' and 'Xiangsha'. *Acta Horticulturae Sinica.* 45:1314-1326.
- 608 **Zhu XW, Cui WX, Zhang EJ, Sheng JY, Xiong F. 2017.** Morphological and physicochemical
609 properties of starches isolated from three taro bulbs. *Starch-Strke.* 70:1-2.
- 610 **Zhu YM, Luo XL, Fan WJ. 2016.** Effects of endogenous cytokinin content on starch accumulation in
611 root tuber of cassava. *Journal of Southern Agriculture.* 4:1279-1284.

Figure 1

three classifications of taros in agricultural production of China

(A) Kui taro; (B) Multi-cormels taro; (C) Multi-head taro.

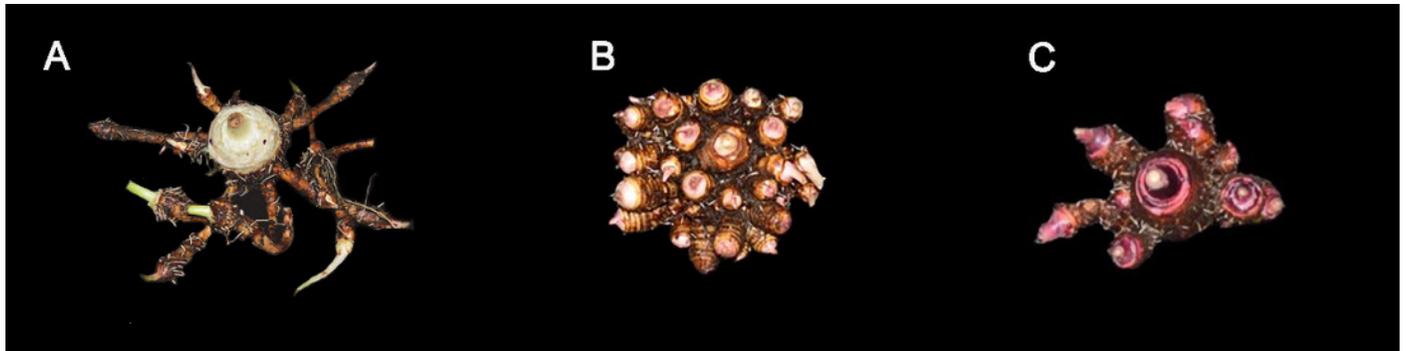


Figure 2

The development of taro according to the diameter

The taro bulb is developed by the bud and the diameter increases with the development process.



Figure 3

The taro bulb's mother taro, sub taro and grandson taro.

(A) Multi-head taro; (B) Multi-cormels taro. Mother taro: it is developed by the top bud of the seed taro. Sub taro: it is developed from the lateral bud on the mother taro. Grandson taro: it is developed from the lateral bud of the sub taro.

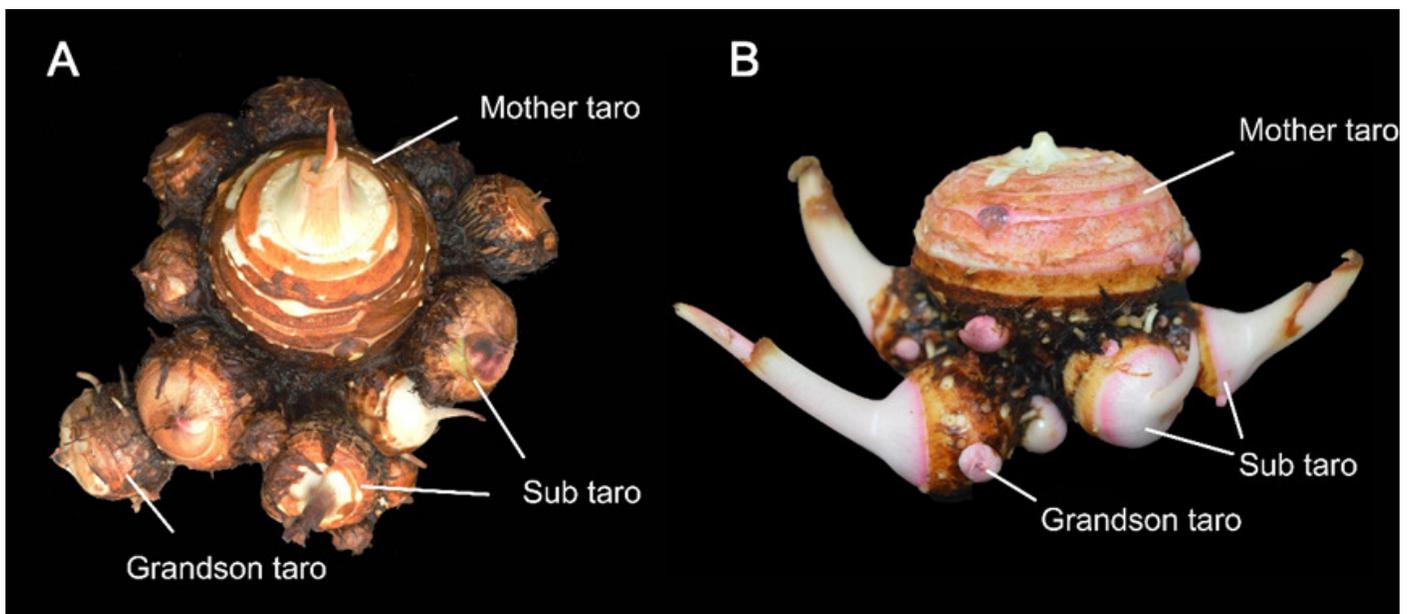


Figure 4

The development process of amyloplasts in the parenchyma cells of Kui taro bulbs.

(A,E) Bulb diameter 1 cm; (B,F) Bulb diameter 5 cm; (C,G) Bulb diameter 9 cm; (D,H) Bulb diameter 13 cm; (Am) amyloid; (Cy) cytoplasm.

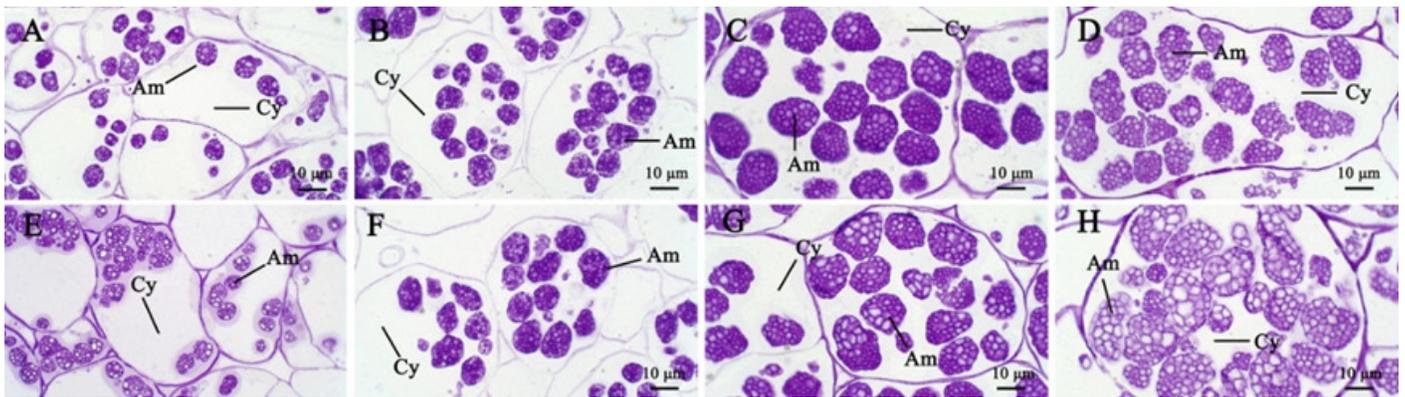


Figure 5

Transmission electron microscope pictures of starch grains of Kui taro bulbs

(A) Bulb diameter 1 cm; The proliferation of amyloplasts, the red box shows the small starch granules after the overflow and shrinkage, and the shape has changed from irregular to spherical. (B) Bulb di-iameter 5 cm; The arrangement of small starch granules in large starch bodies, red asterisks represent large starch granules, blue asterisks represent small starch granules, two types of starch granules are distributed on both sides of starch bodies.

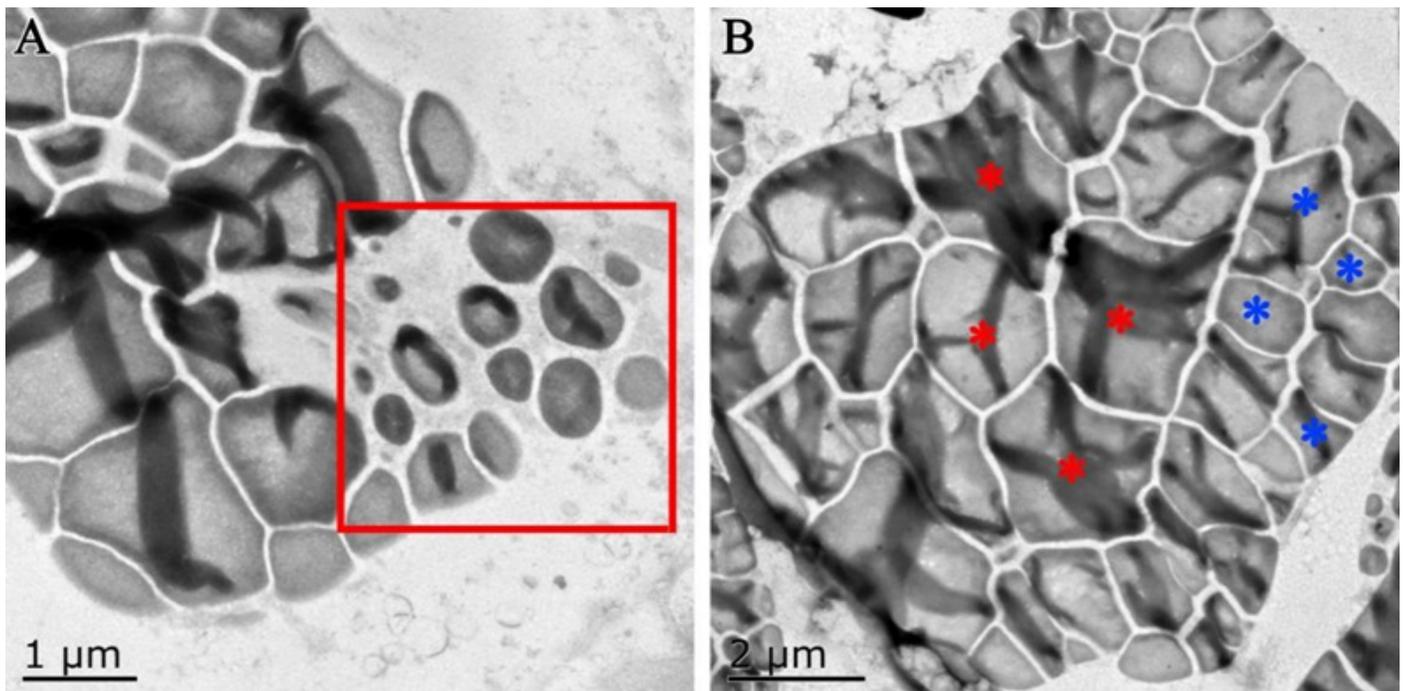


Figure 6

Effects of external environmental factors and internal factors on taro bulb expansion and starch enrichment.

