Plant grafting and its application in biological research

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Grafting is widely used for various aspects of plant biological research. We reviewed grafting methods and graft development processes for herbaceous plants. We introduced also methods of investigating the development of graft union and compatibility, and the application of grafting in understanding translocation of substances and long-distance signaling in plants.

grafting, translocation of substances, long-distance signaling

Plant grafting originated during the ‘Warring States’ period of China’s history, and is thus more than two thousand years old. During its long history, grafting techniques have been continuously developed and improved. Today, grafting is commonly used as a routine method of asexual reproduction for the agricultural production of many types of plant, including fruit trees, vegetables and flowers. It may be used to enhance resistance, improve quality, and increase production. It has therefore brought enormous economic benefits to agricultural production.

Grafting is the method of purposively combining the buds or branches of one plant with the roots and base of another plant. The upper component is the ‘scion’, and the lower is the ‘rootstock’ or ‘stock’. ‘Autografting’ is grafting between the same plants of the same species; ‘homografting’ is grafting between different plants of same species; ‘heterografting’ is grafting between plants of different species [1]. The grafting combinations used in agricultural production are commonly compatible, as both scion and rootstock survive after grafting and grow normally until flowering [2]. Most heterografting combinations are incompatible as either scion or rootstock, and both may die after grafting.

Grafting has been widely used to study transport of substances within plant tissues, flowering regulation, and signal transduction mechanisms. We reviewed grafting methods, graft union development, and grafting applications in plant biological research.

1 Grafting methods

Grafting is commonly performed at plant stems using flat, cleft, and approach methods, among which cleft grafting is the most commonly used. In cleft grafting, the scion base is cut into a wedge shape. The rootstock is first sliced horizontally and then cut into a cleft. The scion is inserted into the cleft in the rootstock. Flat grafting is commonly used in studies of graft union development. In flat grafting, both the scion and rootstock are cut into flat surfaces, and are grafted together. In approach grafting, the cambium of the scion and rootstock are exposed by cutting the epidermis and cortex. The surfaces are then combined. To successfully perform grafting, the precise docking of anatomical structures between scion and rootstock is very important. Dislocation of vascular system and cambium often leads to graft failure or delayed development [3].

For different research purposes, a variety of other grafting methods have been introduced, including callus grafting, grafting of an explanted stem segment in vitro, and micrografting.

To study cytological events during incompatible grafting,
Moore and Walker [4] performed in vitro callus grafting between Solanum pennellii and Sedum telephoides; they found that the cells near the graft union had dilated endoplasmic reticulum and thickened cell walls. A co-suspension culture of pear (Pyrus communis) and quince (Cydonia oblonga) callus resulted in significant reduction in quince growth [5].

Gebhardt and Goldbach [6] established a micrografting method using plantlets obtained from tissue culture of Prunus shoot-tips under sterile conditions, and cultured the grafted plantlets in vitro in liquid medium. Russo and Slack [7] introduced a method to screen and analyze virus-resistant potato cultivars by grafting potato plantlets in vitro.

Parkinson and Yeoman [8] introduced an in vitro grafting method using explanted stem segments. They cut a part of a plant stem into segments, grafted these together, and cultured them in sterile MS medium. To simulate normal physiological conditions, the medium was divided into two parts. The medium in contact with the scion was supplemented with sucrose, auxin, and cytokinin at particular concentrations, while the medium in contact with the rootstock was only supplemented with cytokinin. Since it is easy to change media composition and culture conditions, this method has been used to explore the impacts of different factors and conditions on graft compatibility and physiological function [9].

Arabidopsis thaliana has been widely used as a model plant in the study of plant genetics, developmental biology, and molecular biology. The grafting of A. thaliana using different combinations has been used to investigate transport of substances, signal transduction, floral induction, systemic resistance, and abiotic stress response [10,11]. A. thaliana has rosette leaves, and its most suitable grafting positions are the inflorescence stem and the hypocotyl. Grafting of an inflorescence stem has usually been performed using cleft and flat grafting methods [10,12]. A few days after bolting, the inflorescence stem is cut with a razor. The grafting site is rapidly covered with silicone from a tube, to ensure that the scion is closely connected with the rootstock. Turnbull et al. [11] first established a hypocotyl grafting system in Arabidopsis thaliana using seedlings grown in petri dishes. They cut the hypocotyl of seedlings under a stereomicroscope, and used these for autografting or homografting. Grafting during bolting is convenient because the inflorescence stem diameter is relatively thick. However, during this period, A. thaliana has completed most of its important physiological development including the induction of flower organs. Though hypocotyl grafting is relatively difficult, this grafting system is more suitable for the study of substance and signal exchanges between roots and shoots. Rhee and Somerville [12] performed preliminary studies of the histology of the graft union from inflorescence stems. The detailed development process of Arabidopsis thaliana grafting has not been well studied, however. Determining when the vascular tissue connection between scion and rootstock takes place, is very important in understanding how signals are transducted and substances are transported.

2 Graft union development in herbaceous plants

The developmental process in compatible grafting of herbaceous plants roughly includes: (1) the formation of the isolation layer and the initial adhesion of scion and rootstock, (2) the formation of the wound callus, and (3) reconnection of the vascular bridge between scion and rootstock [13,14].

2.1 Isolation layer formation and initial adhesion between scion and rootstock

Cutting during the grafting process can lead to the formation of an isolation layer, also known as a necrotic layer. Observation under an electron microscope has indicated that the isolation layer is composed of the remaining cell walls of cutting-induced dead cells and surrounding substances with a high electron density [15,16]. The initial adhesion between scion and rootstock has been observed to occur 1 d after grafting, and is related to a pronounced dictyosome activity in cells near the graft interface [16]. During the early stages of grafting, the isolation layer is distributed over the whole graft interface. With the formation of the callus, the isolation layer is disrupted in the vascular area, and is then gradually extended. Thus, scion and rootstock are able to directly connect. Lastly, the isolation layer disappears, mainly because of growth pressures caused by the callus dividing at both sides of the graft union, and by callus absorption [17].

2.2 Formation of the wound callus

Two to three days after grafting, the wound callus is formed by division of the parenchyma cells from cambium, phloem, xylem, and pith near the wounded surface. In herbaceous plants, callus cells are first formed at vascular bundles and cortex, with only a few in the pith [14,15,17]. Mixing of the wound callus cells of both graft partners can lead to further connection between scion and rootstock.

With the formation of the callus, plasmodesmata are formed between scion and rootstock cells in the thinning region of the isolation layer. Because the plasmodesmata are formed in non-dividing cell walls, they are considered secondary plasmodesmata. Kollmann and Glockmann [18] accurately distinguished the contacting cell walls in a Vicia faba/Helianthus annuus graft based on ultrastructural features, and confirmed the presence of secondary plasmodesmata between rootstock and scion. The secondary plasmodesmata are able to connect cells between rootstock and scion to form a continuous symplast. Thus, substance exchange between adjacent cells and cell communication can be achieved. These may play important roles in the further
development of the graft union. Wang [9] found that no plasmodesmata were formed between scion and rootstock in incompatible *Vicia faba/Helianthus annuus* autograft, and the phloem of scion and rootstock were not connected.

### 2.3 Formation of wound repair vascular bridges between scion and rootstock

Formation of wound repair vascular bridges between scion and rootstock is a critical developmental event in compatible grafting [2]. In herbaceous plant grafting, differentiation of xylem elements can be observed in the wound callus 3 d after grafting. These newly formed xylem elements are irregular in shape and generally directly differentiate from callus cells at the graft union or parenchyma cells around vascular bundles [17,19]. Four days after a pea (*Pisum sativum*) autograft, xylem elements were differentiated from the callus cells of both scion and rootstock. Seven days after grafting, wound repair xylem bridges connecting vascular bundles of scion and rootstock were observed [17]. Similarly, formation of wound xylem bridges was also observed in *Vicia faba* and *Nicandra physaloides* autografts of explanted stem segments, 5 d after grafting [20]. Monzer and Kollmann [21] identified xyary connections within a graft based on species-specific secondary wall thickenings in a *Lophopora williamsii/Trichocereus spechianna* heterograft.

Before differentiating into wound sieve elements, callus and parenchyma cells at the graft union normally divide one to several times to form sieve elements and companion cells [15,20,22]. In a pea autograft 5 d after grafting, callus cells of scion and rootstock differentiated into sieve elements. Eight days after grafting, wound repair phloem bridges connecting scion and rootstock were observed [17]. In a *Cucumis sativus/Cucurbita ficifolia* heterograft, wound phloem bridges were established at the graft union 7 d after grafting. The average number of sieve tube connections in a heterograft of *Cucumis sativus/Cucurbita ficifolia* was significantly lower than in the respective homograft, *Cucumis/Cucumis* [15]. In an explanted stem grafting system, wound phloem bridges in autografts of *Vicia faba* and *Nicandra physaloides* were observed 5 d after grafting. Those within a heterograft of *Nicandra physaloides/Lycopersicon esculentum* were observed 7 d after grafting [20]. In a *Vicia faba/Helianthus annuus* heterograft, specific cell markers were used to distinguish sieve elements of scion and rootstock. Sieve elements of *Vicia faba* were found to contain P-type plastids with starch and protein, while those of *Helianthus annuus* contained S-type plastids with only starch. These cell markers were used to accurately identify the interspecific sieve plates between *Vicia faba* and *Helianthus annuus* graft partners [22].

As wound repair xylem and phloem elements further differentiate within the graft union, a newly formed wound cambium can eventually develop between them and give rise to new phloem and xylem strands [14,16]. In herbaceous plants, wound cambium is often formed two to three weeks after grafting. The formation of wound cambium crossing the graft union is often considered a sign of the completion of the graft union [14,22].

Studies of autografts of explanted stem segments of *Nicandra physaloides* and *Vicia faba* have shown that the development of grafting *in vitro* is similar to that *in vivo* [9,20]. Given the consistency of culture conditions, explanted stem segment grafting is more suitable for biochemical and physiological research than grafting *in vivo*. A new way to study compatible or incompatible mechanisms of grafting is thus available.

Grafting success is influenced by genetics, structure, growth characteristics, physiological, and biochemical factors. Among these, genetic factors are critical for compatibility. In general, the closer the taxonomic relationship between scion and rootstock, the higher the grafting success rate. In most incompatible graft combinations, vascular continuity is not established between scion and rootstock [23]. Physiological factors may be important reasons for graft incompatibility. Gur et al. [24] studied pear/quince graft combination, and found that cyanogenic glycosides produced by the quince moved across the graft interface, where they were broken down by β-glicosidase produced by the pear, to liberate cyanide, resulting in the death of graft.

### 3 Examining methods for graft union development and compatibility

#### 3.1 Measurement of breaking weight

Roberts and Brown [25] developed a technique of measuring the force required to break the graft, at different stages after grafting. This can provide a basis for analysis of the progress of the graft over time. Tensile strength is expressed as breaking weight (g/mm²). Studies have showed that during the formation of the graft union, tensile strength can be divided into three stages: (1) At 1–3 d from grafting, tensile strength is little and increases slowly, by 1 g/mm² daily, in both compatible and incompatible grafts; (2) tensile strength increases sustainably at 3–11 d, 28 fold in a compatible autograft, reaching a maximum of 56 g/mm². In an incompatible graft it increases only at 2–5 d, reaching a maximum of 12 g/mm²; (3) tensile strength in a compatible graft stabilizes and becomes similar to that of an ungrafted internode, while that of an incompatible graft gradually decreases from day 5. Changes in tensile strength are related to histological and cytological changes within the graft joint [26].

#### 3.2 Measurements of electrical resistance

Yang et al. [27] measured changes in electrical resistance at the graft junction in a compatible *Lycopersicon esculentum* autograft. They found that resistance changes were related to the developmental process of the graft union. Two to
three days after grafting, the electrical resistance increased rapidly with an isolation layer forming and becoming thicker. In the next 3–8 d, electrical resistance steadily decreased with callus proliferation, the breakage of the isolation layer, formation of secondary plasmodesmata between contacting cells of scion and rootstock, and the differentiation of vascular elements in the callus. Finally, resistance began to drop to the level of the intact stem. However, for an incompatible *Amaranthus tricolor/Lycopersicon esculentum* heterograft, the electrical resistance at the graft junction was persistently high because of the persistence of the isolation layer.

3.3 Measurements of hydraulic conductance

Root hydraulic conductance \( (L_0) \) is a measure of nutrient and water absorption in grafted plants; it has therefore been used to determine the graft developmental process [28]. Fernández-García et al. [29] determined the \( L_0 \) value of a *Lycopersicon esculentum* autograft, 4, 8, 12 and 15 d after grafting. They found that the \( L_0 \) was zero at day 4 after grafting, then linearly increased and reached a maximum between days 12 and 15. Meanwhile, histological examination found xylem conduits between scion and rootstock at the graft union, and fully developed xylem at day 15, suggesting that changes in the \( L_0 \) value reflected the xylem developmental process within the graft.

4 Application of grafting in plant biological research

4.1 Application of grafting in studies of substance transport

The formation of vascular bridges between rootstock and scion provides an important developmental index for compatible grafts [2,15]. Successful grafting, however, is reflected not only in the structure, but more importantly, in the physiological functions of vascular bridges in the exchange of substances between scion and rootstock. Substance exchange is reciprocal: from scion to rootstock and from rootstock to scion. Assimilate and mineral transport have been commonly used to examine the vascular development of the graft, and the transport functions at the graft union.

Parkinson et al. [30] utilized iron particles with different diameters to detect the transport function of xylem conduits in an explanted stem graft. They found that iron particles were transported with water from rootstock to scion in the compatible autograft of *Lycopersicon esculentum* and *Nicandra physaloides*, but not in an incompatible heterograft of *Lycopersicon esculentum/Nicandra physaloides*. This indicates that graft incompatibility is related to the transport capacity of the xylem at the graft union.

\( ^{14} \text{C} \)-labeled assimilates are commonly used to examine the transport function of wound phloem connections between scion and rootstock. De Stiger [31] used \( ^{14} \text{C} \)-labeled CO\(_2\) to study assimilate transport in a *Silene armeria* autograft. They found that transmission from one partner to the others occurred 7 d after grafting and took place at a high rate by 16 d. The transport of assimilates has been explained in terms of the source-sink concept. Rachow-Brandt and Kollmann [32] studied a compatible heterograft of *Lycopersicon esculentum* on *Solanum tuberosum*, and a less compatible heterograft of *Vicia faba* on *Helianthus annuus*. *Lyco- persicon esculentum*, *Vicia faba*, and *Solanum tuberosum* homografts served as controls. The transport of \( ^{14} \text{C} \)-labeled assimilates across the graft union started 5–7 d after grafting, in compatible hetero- and homograft combinations. Subsequently, the number of sieve tubes in the graft union and the transport rate of assimilation increased, indicating that assimilate transport is via phloem. For a less compatible graft of *Vicia faba* on *Helianthus annuus*, although wound phloem bridges were established between scion and rootstock, only 2% of assimilates were transported 11 d after grafting. In comparison, 40% and 30% of assimilates in *Vicia faba* and *Helianthus annuus* autografts were transported, respectively. Kollmann and Glockmann [22] found that heavy callose was deposited at the sieve pores of an interspecies graft of *Vicia faba* and *Helianthus annuus*. The sieve pores thus became narrow; this may, to some extent, have been related to incompatibility. In incompatible grafts, assimilate transport from scion to rootstock is often blocked.

Wang and Kollmann [20] determined the assimilate transport of different explanted stem grafts, at different developmental stages, using \( ^{14} \text{C} \)-labeled sucrose. They found that 3 d after grafting, only a small amount of labeled sucrose was transported from scion to rootstock, possibly by diffusion through secondary plasmodesmata. Five to seven days after grafting, the assimilate transport rate had increased significantly because of the formation of wound phloem bridges between graft partners. In incompatible grafts, because no wound phloem bridge formed between scion and rootstock, the rate of assimilate transport was low, and most assimilates were blocked at the graft union.

\( ^{6} \text{S} \text{Carboxyfluorescein (CF)} \) has been commonly used to trace substance transport in sieve tubes. Like assimilates, it is also transported from source to sink [33]. Schöning and Kollmann [34] used CF to examine the transport function of wound phloem in an explanted stem autograft of *Lycopersicon esculentum* and *Helianthus annuus*, and found that CF transport was carried out in wound phloem.

Tiedemann and Carstens-Behrens [35] collected phloem exudate from heterografts of *Cucumis sativus* (scion)/*Cucurbita maxima* (rootstock) 5–7 weeks after grafting, and examined protein patterns using SDS-PAGE. Compared with a *Cucurbita* control plant, the phloem exudate from the *Cucumis* scion of the heterografts had at least 4 novel protein bands with molecular weights exactly complementing the *Cucurbita* protein pattern, includ-
4.2 Application of grafting in studies of long-distance signaling in plants

Grafting is an ideal method for studying long-distance signaling in plants. Using different combinations of mutants, transgenic plants, and wild-type plants as scion or rootstock, it is possible to analyze reciprocal long-distance signaling mechanisms.

(i) Long-distance transport of RNA through plant phloem. In recent years, studies have found that some RNAs function as non-cell-autonomous signaling molecules, are transported within plants, and control organ development or defense responses. Plant plasmodesmata provide intercellular transport channels for proteins or RNA-protein complexes. The vascular system, the specialized phloem in particular, provides long-distance transport pathways for non-cell-autonomous protein and RNA-protein complexes [38]. Xoconostle-Cázares et al. [39] grafted cucumber (<i>Cucurbita maxima</i>) onto pumpkin (<i>Cucurbita maxima</i>), collected sieve exudates from the scion, and analyzed their molecular composition. Their research provided direct evidence that the <i>CmPP16</i> mRNA of the rootstock was transported to the scion, thus crossing the graft union. Using <i>in situ</i> RT-PCR, Ruiz-Medrano et al. [40] found that <i>CmNACP</i> mRNA was transported via phloem from pumpkin rootstock into apical tissues of the cucumber scion. <i>CmNACP</i> belongs to NAC domain gene family. Some members of this family are involved in apical meristem development. This experiment confirmed the transport of specific mRNAs to shoot apices in plants. Kim et al. [41] investigated a dominant leaf mutant of tomato called Mouse ears (<i>Mae</i>), that is caused by chromosomal rearrangements resulting in a fusion of <i>LeT6</i> and <i>PEP</i>. If a normal scion is grafted onto a <i>Mae</i> mutant rootstock, new leaves that initiate on the scion have the <i>Mae</i> morphology. In addition, <i>in situ</i> PCR detected <i>Mae</i> transcripts in the scion apex. These experiments confirmed that long-distance transport of regulatory mRNAs in plants could control plant morphology.

(ii) Long-distance transport of signaling protein in plants. Wounding caused by insect or herbivore attack is able to induce systemic responses in higher plants. Specific signals can be delivered from wounds to undamaged sites, causing expression of proteinase inhibitors (PIs). The wound-signaling polypeptide is systemin, composed of 18 amino acids. Systemin is released from its precursor, pro-systemin, composed of 200 amino acids [49]. McGurl et al. [50] found that proteinase inhibitors I and II were constitutively expressed subunits of the PP1 and PP2 proteins within <i>Cucurbita</i>. Golecki et al. [36] further showed that in a <i>Cucurbita maxima</i> scion grafted onto <i>Cucurbita sativus</i> rootstock, at least 9 additional proteins were transported from rootstock to scion, 9–11 d after grafting; indicating that protein exchange occurred in the sieve tubes of scion and rootstock, after phloem reconstruction at the graft union. That the PP1 and PP2 mRNA of the <i>Cucurbita maxima</i> or <i>Cucurbita ficifolia</i> rootstock were undetectable in the <i>Cucurbita sativus</i> scion suggests that proteins rather than mRNA were transported. <i>Cucurbita maxima</i> PP1 was immunolocalized in sieve elements of the extrafascicular phloem of the <i>Cucurbita sativus</i> scion, whereas <i>Cucurbita maxima</i> PP2 was found in both the sieve elements and companion cells [37].
pressed in the leaves of prosystemin transgenic tomato, but only expressed under wounding in non-transgenic plants. Granting of non-transgenic tomato plants as scions onto prosystemin transgenic rootstock was able to induce constitutive expression of proteasome inhibitor proteins in the leaves of both scion and rootstock. These results support the proposed role of systemin as the mobile wound signal in plant wound responses.

The transition from vegetative to floral meristems in higher plants depends on the interaction between internal and environmental signals. Florigen is produced in the leaf after inductive photoperiods and transported to the shoot apex where it triggers floral morphogenesis. Genetic studies of *Arabidopsis* have shown that *FLOWERING LOCUS T* (*FT*) can induce long-day and short-day plant flowering [51]. Heterografting has shown that the effective florogenic signal was transported from flowering *Cucurbita maxima* rootstock to the *Cucurbita moschata* scion, under long-day conditions. Although real-time RT-PCR detected no *FT* transcript in the phloem sap collected from the *Cucurbita maxima* rootstock, *Cm-FT L1* and *Cm-FT L2* proteins were detectable by mass spectrometry of phloem sap proteins, revealing that *Cucurbita maxima* *FT*, and not *FT* mRNA, is transferred across the graft union in the phloem translocation stream [52]. *FT* was expressed in the cotyledons and leaves of *Arabidopsis* under long-day conditions, and induced the target genes such as *APETALA1 (AP1)* in the shoot apices to initiate floral morphogenesis. Granting experiments indicated that *FT* protein was transported from scions to the apical region of the rootstock, suggesting that *FT* protein rather than its mRNA, is the essential component of florogen [53].

In summary, grafting technology has made important contributions to agriculture for thousands of years. As a unique research system, grafting has been widely used in plant scientific research. Its application has solved some basic theoretical issues such as substance transport and signaling transduction. To resolve new theoretical problems there is a need to select appropriate mutants or transgenic plants, and establish different grafting combinations. However, some issues in grafting still need to be solved. One is how to overcome graft incompatibility, and thus widen its application in agriculture. Another is how to advance monocot grafting technology and utilize it in research involving monocotyledons as model plants.

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