Importance of symplasmic communication in cell differentiation

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Abbreviations: CFDA – 5-(and-6), Carboxyfluorescein Diacetate; ER, endoplasmic reticulum; GFP, green fluorescent protein; HPTS, 8-Hydroxyppyrene-1,3,6-trisulfonic acid; MP, movement protein; NCAP, non-cell-autonomous proteins; PD, plasmodesmata; PM, plasma membrane; TNT, tunneling nanotubules

Symplasmic communication via plasmodesmata (PD) is part of the system of information exchange between plant cells. Molecules that pass through the PD include ions, some hormones, minerals, amino acids, and sugars but also proteins, transcription factors, and different classes of RNA, and as such PD can participate in the coordination of plant growth and development. This review summarizes the current literature on this subject and the role of PD in signal exchange, the importance of symplasmic communication and symplasmic domains in plant cell differentiation, and highlights the future prospective in the exploration of PD functions in plants. Moreover, this review also describes the potential use of barley root epidermis and non-zygotic embryogenesis in study of symplasmic communication during cell differentiation.

The presence of PD, channels within the walls between neighboring cells, can be interpreted as the main reason determined at least 2 continuous systems in plants: apoplast and symplasm.¹ The apoplast is an extraprotoplasmic region composed of cell walls with microspaces within the cell wall, some intercellular spaces, and the lumina of dead cells.² The symplasm is a system consisting of all protoplasts interconnected by PD and bounded by continuous plasma membrane.³ Terminology used for the description of these systems in not consistent in the literature and because of it, the terminology proposed by Romberger et al.¹ will be used. Namely, in this review we will use the term “symplasm,” and appropriate derivative terms as “symplasmic” or “symplastically,” as we agree with Romberger et al.¹ that the term “symplastic” has another meaning referring to growth of cells within the plant tissue.⁴ From functional point of view, PD are the main part of the symplast, but we would like to point to another possible spatial system which can be distinguished because of PD structure. Namely, it is worth introducing into the nomenclature the term “endoplasmic symreticulum” as suggested previously,³ that would comprise a third continuous spatial system within a plant body. This system is composed of the endoplasmic reticulum traversing through PD thus creating the unity of the endoplasmic reticulum within the plant body.

Plasmodesmata structure and permeability—present knowledge

PD are structures present in plants,⁶ algae,⁷,⁸ and fungi,⁹ which comprise cytoplasmic channels that connect the neighboring cells, and allow the communication between individual cells (cell-to-cell communication).¹⁰-¹⁴ The first report describing these “channels” across neighboring plant cell walls, deduced from observation with the use of conventional microscopy, comes from 1880,⁶ and name “plasmodesmen” was used for the first time in 1901 by Strasburger.¹⁵ During the last few years detailed ultrastructure of PD was described in review papers,¹⁶,¹⁷ so here we only mention that cytoplasmic channels are surrounded by plasma membrane (PM) internally lined with an appressed ER membrane, termed the desmotubule (Fig. 1).¹⁸,¹⁹

The diameter of PD is not constant and may be reduced until the complete closure inter alia by the deposition of callose (β-1,3-glucan) in the neck regions of PD.²⁰,²¹ Deposition of this polysaccharide depends on the activity of 2 enzyme groups: β-1,3-glucan synthase that produces callose, and β-1,3-glucanase responsible for callose degradation.²²,²³ The diameter and permeability of PD may be modified during cell development or in response to the external conditions, like temperature, pathogen attack, or wounding.²⁴,²⁵ The permeability of PD is also limited, by the diameter of microchannels (Fig. 1), and the value of SEL (size exclusion limit), described in units of mass, is used in most cases to determine which molecules can pass through the PD, what is an indicator of the maximal molecular size of the molecule/molecules traversed through PD.²⁶ Many studies on the communication via PD are based on the transport of low molecular fluorochromes, fluorescent labeled dextrans, or green fluorescent protein (GFP), which allows to compare PD permeability for molecules of different sizes.²⁷-²⁹ Sometimes to determine the maximum size of
molecule, that may migrate through PD, GFP molecules, and complexes of 2 or 3 GFPs molecules (2xGFP/3xGFP) are being used.\textsuperscript{30,31} It is important to take into consideration that in such cases the SEL can be between 27–81 kDa. However, it must be understood not as a diameter of microchannels participating in GFP movement, but the parameter describing the molecule size, including its length, which can influence the movement of the molecules in question. The correlation between increasing size of GFP complexes and the reduced permeability of PD is obvious,\textsuperscript{30} but it cannot be excluded that 3, connected in series, molecules of GFP, and one single GFP may move through PD with the same diameter of microchannels (\textbf{Fig. 1}). Moreover molecules with a lower molecular weight may have a larger diameter than the molecules of larger weight (\textbf{Table 1}).\textsuperscript{32,33} This explains why the description of PD microchannel diameter using of the radius of molecules – MEL (molecular exclusion limit) is more accurate than molecule weight.\textsuperscript{34-38}

**Symplasmic transport—route for molecules including macromolecules**

Initially it was postulated that PD are an intracellular channels for the diffusion of small molecules, such as ions or sugars.\textsuperscript{6,39} However, subsequent studies on the PD described these structures as dynamic gateways actively transporting or blocking transport of macromolecules: proteins and RNAs.\textsuperscript{37,40,41} The first information regarding macromolecules transported through PD was based on the studies on movement protein (MP) encoded by \textit{Tobacco mosaic virus}.\textsuperscript{24,42} MPs interact with the PD and may increase their diameter, break symplasmic isolation of cells, and spread viruses through the plant tissues.\textsuperscript{43,44} Further it was discovered that plant proteins may pass via PD, using a similar mechanism as MPs, and all proteins transported symplasmically between cells have been called as non-cell-autonomous proteins (NCAPs).\textsuperscript{45} To NCAPs are included for example: transcription factors, like \textit{KNOTTED1} (KN1), which plays a crucial role in maize meristem organization,\textsuperscript{46} or cell-fate-deciding proteins, involved for example in the root hair development, like \textit{TRANSPARENT TESTA GLABRA1} (TTG1),\textsuperscript{47} \textit{CAPRICE} (CPC),\textsuperscript{48} or \textit{CAPRICE-LIKE MYB3} (CPL).\textsuperscript{49} The movement of NCAPs related to cell differentiation is regulated by as yet unknown mechanisms, that precisely guide proteins to the specific individual cells (like TTG1, CPC, CPL) or cell layers (like KN1; for a review see refs. 16,17). Interactions between NCAPs and PD are not well understood, although the motifs crucial for symplasmic transport of different proteins have been described.\textsuperscript{35,50,51}

A second group of molecules that may be symplasmically transported between neighboring cells are RNAs including small RNA (sRNA) involved in RNA silencing.\textsuperscript{52} sRNAs are classified as short (21–24 nucleotides), non-coding molecules that have an influence on transcriptional and post-transcriptional regulation of the target sequences.\textsuperscript{53} Transport of sRNA between plant cells and their role in the cell differentiation or in the response to the viruses infection is well understood (for a review see refs. 54,55). One of the best documented examples is the description of 2 microRNA molecules: miR165a and miR166b, involved in the development of vascular tissue in \textit{Arabidopsis thaliana} root.\textsuperscript{56,57} Both miRNAs expressed in root endodermis, non-cell-autonomously suppress the expression of PHABULOSA (PHB), class III HD-ZIP transcription factor. And this suppression of PHB in the peripheral root stele is required for the xylem differentiation.\textsuperscript{56} Also the gradual distribution of PHB among the root stele, due to the miR165a/ miR166b silencing, is crucial for the differentiation of pericycle and ground tissue patterning in \textit{Arabidopsis} roots.\textsuperscript{57} Moreover, the expression of MIR165a/MIR166b is activated in the endodermis by \textit{SHORT-ROOT} (SHR) transcription factor, that is also transported via PD,\textsuperscript{56,58} these data indicate

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**Table 1.** Comparison of the molecular weight and diameter of some of the molecules used in the analysis of symplasmic communication.

| Chemicals                                  | Molecular weight [Da] | Molecule diameter [nm] | Ref. |
|--------------------------------------------|-----------------------|------------------------|------|
| 5-(and-6)-Carboxyfluorescein Diacetate (CFDA) | 376.21                | 1.3                    | 35   |
| 8-Hydroxypyrene-1,3,6-trisulfonic acid (HPTS) | 524.39                | 0.9                    | 33   |
| Green fluorescent protein (GFP)            | 27 000                | 2.8                    | 34   |
| Dextran                                    | 10 000                | 2.3                    | 32   |

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![Figure 1](image-url). Schematic representation of the molecular structure of PD and movement of molecules. Description is given in the text.
that NCAPs play a role in cell differentiation at multiple levels and may interact with others NCAPs or key cell-fate deciding proteins.

Symplasmic communication/isolation—basic definition

The discovery that the plant body is divided into regions consisting of cells which are not connected by PD, or in which such connections are temporally closed or diminished, resulted in the terms “symplasmic domains” and “subdomains” or “symplasmic fields” being used. A symplasmic domain is a cell or group of cells which are connected by PD between each other, but on the border of a domain is not connected by functional PD with the neighbor cells or connection is diminished. If such a lack of connection by PD is permanent the domain is called “permanent symplasmic domain” and the best example is stomata cells. More interesting are the “temporary symplasmic domains,” which consist of cells, or group of cells which only temporally closed PD on the domain border or the movement of molecules through PD is diminished quantitative or qualitative. Sometimes within the domain, subdomains can be distinguished. Temporal domains are more interesting because the analysis of their appearance/disappearance and the mechanisms involved in their function can provide answers to the role of symplasmic domains in plant growth and development and the spatio-temporal correlation between symplasmic domains and cell differentiation.

It must be noticed that apart from “symplasmic domain,” the term “symplasmic field” exists. This term was introduced for the description of symplasmic isolated areas present in the apical meristem. The authors proposed such a nomenclature because apical meristem is different from mature tissues which term symplasmic domain was introduced in the way that apical meristem is continuously renewing its cellular composition.

Correlation between symplasmic communication/isolation and cell differentiation

It is worthy of note that changes in gene expression are correlated with the changes in symplasmic communication based on the number of PD, or their permeability as demonstrated during crown gall development, where no PD exists on the border between the tumor and the rest of the plant, in Chara vulgaris closing of PD is the necessary for proper spermatogenesis or in the case of Onoclea sensibility where disturbing of the connection by PD resulted in possessing of the cells totipotency.

As noted previously, the role of PD in cell differentiation is based on the discovery that plant cells can communicate by PD with neighboring cells within the symplasmic domain and share with each other common components and behavior. It is known that cell differentiation is correlated with the formation of symplasmic domains or subdomains (for a review see refs. 17,70,71). It is also known that cells in symplasmic connection (within symplasmic domain) are characterized by the same direction and frequency of cell division, that the more advanced the state of cell differentiation the lower symplasmic connection, and disturbance in symplasmic transport causes changes in normal plant growth and development. It is postulated, that changes in the direction of cell differentiation are correlated with the changes in symplasmic communication. For example, in callus of Cocos nucifera embryogenic cells were separated symplasmically from the explant at the globular stage of somatic embryo development suggesting that the acquisition of embryogenic competence is preceded by closing of PD. Another example from studies on somatic embryogenesis of symplasmic isolation as a main factor allowing changes in cell fate comes from studies on Quercus suber L. callus where dividing cells (future somatic embryos) were symplasmically isolated from the explant.

Zygotic and non-zygotic embryogenesis—a good model for symplasmic communication studies

The hypothesis about the role of the symplasmic isolation of cells during zygotic embryogenesis was first postulated after an analysis of the ultrastructure of developing embryos in different plant species. It was shown that during the early stages of zygotic embryogenesis, the embryo sac-forming meiocyte is not connected to adjacent cells by PD. Moreover, for the embryo sac of Capsella bursa-pastoris it was shown that the egg is connected by PD to the synergids at first but that these connections disappear after fertilisation, and from that moment the young zygote is not connected to other cells by PD and additionally that the sac is isolated from the nucellar cells by the absence of PD. A detailed analysis of the symplasmic communication using fluorescent tracers before and after fertilisation of a Torenia fournieri embryo sac confirmed that the functional connection between the central cell and the egg apparatus cells by PD decreased after fertilisation. These results point not only to the problem of cell-to-cell communication during morphogenesis, but also to the important role of the symplasmic isolation of different genotypes and generations.

The best-described correlation between symplasmic communication and development is the zygotic embryogenesis of Arabidopsis. These studies showed that the zygotic embryo is a single symplasmic domain during all of the stages of development only in the case of low molecular weight molecules (0.5 kDa). However, their movement through PD decreased with the more advanced developmental stages in the case of larger molecules (10 kDa F-dextran). The use of transgenic lines with GFP allowed the symplasmic subdomains that are correlated with the embryo tissues and organ development to be determined. It is important to note that other phenomena such as the unidirectional movement through PD during zygotic embryogenesis have also been described. It has been shown that substances move between the suspensor and embryo in both directions in the case of globular stage zygotic embryos. When embryos shift in symmetry from radial to bilateral symmetry, communication by PD was only from the embryo to the suspensor. Detailed information about symplasmic isolation/communication during the zygotic embryogenesis has been well described in the literature recently.

During zygotic embryogenesis, it is difficult to study the mechanism underlying the regulation of this developmental process for many reasons. That is why non-zygotic embryogenesis (including somatic embryogenesis and androgenesis) has become a very good experimental system for studies on the
mechanisms that control embryogenesis and embryo development at different organization levels including symplasmic communication between the explant and embryogenic cells and within non-zygotic embryos during histio- and organogenesis.

Somatic embryogenesis can be divided into at least 2 phases: 1) changes that take place within the explant that lead to the resumption of totipotency, thus allowing somatic embryo development, and 2) within the somatic embryo during the development of the embryo tissues and organs. In the case of explant cells, the main question is what influences the cells within the explant to develop in different directions and why will only some of them develop into somatic embryos? Many factors/reasons have been postulated and among them symplasmic communication seems to play an important role in the regulation of explant cell differentiation during somatic embryogenesis.85 It has been postulated that such isolation is a prerequisite for changes in the cell developmental program.86 The information that is available in the literature concerns an ultrastructural analysis of the explant with attention being paid to the presence of PD between groups of explant cells with different features of their walls and cytoplasm. However, it must be noted that the presence of PD does not mean that intercellular connections are involved in the exchange of “information” because PD can be not functional, but the lack of PD on the “borders” between different groups of cells within the explant indicates a lack of symplasmic communication.

Some information about the role of cell-to-cell communication by PD within the explant comes from experiments using plasmolysis as a factor that disturbs the cytoplasmic connection by PD. In the case of Panax ginseng,87 pretreatment with sucrose caused plasmolysis, and the number of somatic embryos increased 4-fold in comparison to untreated explants.87,88 Moreover, such results also showed that somatic embryos that developed from a preplasmolyzed explant were of a single-cell origin in comparison to untreated explants. Similar results were obtained for Eleutherococcus senticosus, Oncidium Sharry Baby, Panax japonicas, or Daucus carota where plasmolysis of an explant caused an increase in the frequencies of somatic embryos.89-91 Plasmolysis as a factor that breaks the PD connection between explant cells may suggest that each explant cell behaves as a single cell, which allows these cells to develop in different directions and in this case allows the reprogramming of explant cells into embryogenically competent cells and finally induces somatic-embryo development.90 According to these results, changes in the cell-to-cell communication between the explant cells are a prerequisite to changing cell fate.

An analysis of the morphology of Theobroma cacao L. explants showed that the frequency of PD influences somatic embryogenesis.92 Namely, that in the epidermal cells within the staminodes, tissues that are involved in primary somatic embryogenesis, the frequencies of PD were higher in comparison to the tissue explants for secondary embryo production. The authors suggested that it is possible that the higher level of intercellular connections that is seen in staminode tissues is in part responsible for the higher degree of cellular co-ordination that is seen in primary embryogenesis as compared with secondary embryogenesis.92 In the callus of Panax quinquefolium in which the authors distinguished a few kinds of cells such as parenchymatous, initial embryogenic, grouped embryogenic cells, and embryoid cells, only in the last were many PD detected.93

An ultrastructural analysis of embryogenic cell clusters of Cephalotaxus harringtonia showed differences in the outer cell wall thickness of embryogenic masses, and the frequencies of PD, which were abundant in the walls between embryogenic cells.94 Similar results were obtained earlier in callus that was derived from Citrus sinensis, which showed that cells that were destined to form new proembryoids became surrounded by greatly thickened cell walls without PD.95 An analysis of androgenesis in Zea mays and Triticum aestivum also showed that cells that realize the same developmental program are connected by numerous intercellular connections.36,97

To date, the only information about cell-to-cell movement through PD between explant cells and the somatic embryo comes from analyses using low-molecular weight fluorochrome on the example of Arabidopsis somatic embryogenesis, which showed that at least for some time, the somatic embryo is symplasmically isolated from the explant.86,87 These results also showed that the physical isolation of cells by the deposition of cutin may be a factor that allows the changes in the direction of cell differentiation within the explant.85 An analysis of the distribution of the low-molecular weight symplasmic tracers (HPTS and uncaged fluorescein) during barley androgenesis indicates the symplasmic isolation of the protodermis from the underlying cells of the late globular embryo stage onwards, and the embryonic organs at the mature stage of development, which allows the conclusion that in androgenic embryos in Hordeum vulgare the existence of symplasmic domains correlates with tissue and organ development.85,98,99 An analysis of symplasmic communication was performed using various fluorescent tracers with different molecular weights and cytoplasm soluble GFP being expressed under the control of an STM promoter.99 An analysis of symplasmic communication showed that at the beginning of the culture, PD connectivity between the cells of explants increased. Further changes in symplasmic communication were observed after the next few days of the culture and were correlated with the cells divisions of explants. The changes manifested themselves as a restriction of symplasmic communication on the borders between the dividing and non-dividing cells, which indicated that the parts of the explants that were engaged in morphological processes were isolated from those that were not involved in cell fate changes. Moreover, symplasmic isolation between the somatic embryos and explants was detected. Isolation of the dividing cells from the rest of the plant body was also observed in the case of 35S::BBM transgenic seedlings. Meristematic cells at the cotyledons were isolated from the remaining cells. Furthermore, small groups of dividing cells, which give rise to somatic embryos, were symplasmically isolated from the neighborhood.99 These results suggest that the symplasmic connection of embryogenic competent cells is restricted at this stage of embryogenesis. The results that were obtained showed a correlation between the decrease in symplasmic communication and changes in cell fate and somatic embryogenesis.
Studies using low-molecular symplasmic transport fluorochromes in both species, Arabidopsis and barley, showed that symplasmic domains correlated with the differentiation in the first place, the protodermis, and afterwards other embryo tissues and organs. In general, these data showed a correlation between the decrease in symplasmic communication and changes in the cell differentiation of non-zygotic embryos.99 The data obtained from these studies of somatic embryogenesis and androgenesis indicate that in this type of embryogenesis, the general rules related to the correlation between symplasmic communication and plant development are the same or similar as in the case of zygotic embryogenesis. These data also suggest that cell-to-cell communication is a universal mechanism that is involved in regulation of cell differentiation.

On the whole, therefore, the critical evidence supports the view that resumption of totipotency that leads to embryogenesis in cultured cells is preceded by cell isolation and that differentiation of somatic embryo cells into tissues and the formation of a somatic embryo organ is correlated with changes in symplasmic communication.

Barley root epidermis: a new model for symplasmic communication studies

The recent data has suggested that barley root epidermis may be a useful model for investigation of the role of symplasmic communication in plant cell differentiation.73 The root epidermis is composed of 2 types of cells: those producing root hairs (trichoblasts) and non-root hair cells (atrichoblasts).100 In the meristicmatic zone of root all epidermal cells are similar in size, shape, composition, and number of organelles, amount of cytoplasm, or number of molecules localized in cytoplasm.101 Later, in the differentiation zone of root, cells start to differentiate into trichoblasts and atrichoblasts,102 and 2 neighboring cells can realize different developmental programmes.103 From the previous studies, it is known that in this situation the neighboring cells are usually symplasmically isolated.74 Similar results were obtained in studies on Arabidopsis root epidermis, where after microinjection into the individual root hair or non-root hair cell, fluorescent tracers (Lucifer Yellow and carboxyfluorescein) were not transported to the surrounding cells.104 It was also shown, that during the differentiation of barley root epidermal cells, symplasmic communication is restricted in wild type plants and in the mature root hair zone epidermal cells are still symplasmically isolated.73 Moreover, for 2 allelic root hairless mutants rhl1.a and rhl1.b isolation of epidermal cells was not present. Immunogold detection of callose with the use of the transmission electron microscope revealed that in PD localized between epidermal cells in the differentiation zone of wild type root higher number of callose molecules were deposited, in comparison to the comparable zone of root in root hairless mutant. In mutant plants only single callose molecules were present, and their number was similar to this in the meristicmatic zone of root.73 Results obtained for Arabidopsis and barley clearly indicate a strong correlation between differentiation of root epidermal cells and restriction of symplasmic communication.103,104 However, there are a few alternative mechanisms of establishing the root epidermal patterns in plants: symmetrical or asymmetrical division of mother cell, asymmetrical expansion of daughter cells, or some unknown factor that induce root hair development of individual cells.105 In the Arabidopsis root, epidermal cells are arranged in files of trichoblasts and atrichoblasts, and in the early stage of development, in the meristicmatic region, the fate of cells is determined.101 Whereas in barley the shootward-last division of epidermal cell (division localized the furthest from the root meristem) is symmetric and both daughter cells are identical, but afterwards cells start to expand asymmetrically (Fig. 2).106 The rootward cell (localized closer to the root meristem) is shorter, contains more dense cytoplasm, as well as higher number of mitochondria in comparison to the longer shootward cell. During the differentiation process only shorter cells produce root hairs, what indicates that asymmetric cell expansion plays a key role in root epidermal patterning in barley.106 Additionally, in barley root hairless mutants rhl1.b107 and brb (bald root barley)108 all epidermal cells are similar in size and cytoplasm density, what confirms the hypothesis about importance of asymmetrical expansion in root hair development.106 Current data indicate that the root epidermis of barley may be a new useful model for the studies of symplasmic communication, because in this case, 2 identical daughter cells start asymmetric expansion, which is related to the restriction of symplasmic communication. Disturbances in symplasmic isolation and/or asymmetric cell expansion are responsible for root hairless phenotype – trait that is easily observed during phenotypic analysis.

Future Perspectives

Since plant development primarily depends upon positional information, it would be useful to know the influence of this factor on PD function. Our knowledge regarding the mechanism by which some molecules can pass through the PD and others cannot is still obscure. Currently, information about the proteins connected with the PD structure and function is limited. However, recent data indicate that intracellular connections are not just conduits for macromolecular transport, but also represent sites for glycan synthesis and, potentially, post translational modification.109 Recent studies of grafting have shown that cells exchange chloroplasts by PD between

![Figure 2. Illustration of the early stage of cell differentiation of barley root epidermal cells, which is correlated with the restriction of symplasmic isolation.](Image 239x627 to 563x732)
genetically distinct cell types,10,11 which points to an open question regarding the similarity between PD and tunneling nanotubes (TNT) discovered in animal cells in the correlation of growth and development. The future direction of studies must answer the question as to how PD function and formation are integrated with intracellular signaling pathways and plant physiology."
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