I. Reversible transformation of plant degradation of reserve oil via Glyoxysomes, which are peroxisomes engaged in the mitochondria acquire the ability to oxidize glycine. Organelles. Etioplasts differentiate into chloroplasts while are the functional transformations of many constitutive seed germination. Accompanying these metabolic changes heterotrophic to autotrophic growth occur in the greening of pumpkin cotyledons, the reverse transition of several crucial steps of photorespiration. After the functional cycle, are transformed into leaf peroxisomes that function in dramatic metabolic changes that underlie the shift from heterotrophic to autotrophic growth occur in the greening of seed germination. Accompanying these metabolic changes are the functional transformations of many constitutive organelles. Etioplasts differentiate into chloroplasts while mitochondria acquire the ability to oxidize glycine.

Glyoxysomes, which are peroxisomes engaged in the degradation of reserve oil via β-oxidation and the glyoxylate cycle, are transformed into leaf peroxisomes that function in several crucial steps of photosynthesis. After the functional transition of glyoxysomes to leaf peroxisomes during the greening of pumpkin cotyledons, the reverse transition of leaf peroxisomes to glyoxysomes occurs during senescence. Gene expression, alternative splicing, protein translocation and protein degradation control the functional transformation between glyoxysomes and leaf peroxisomes.

II. Transcriptomics, proteomics and phenomics of plant peroxisomes

Enzymes localized in plant peroxisomes are synthesized in the cytosol and function after their post-translational transport into peroxisomes. Almost all of the peroxisomal matrix proteins contain one of two targeting signals (PTS1 and PTS2) within the molecules. PTS1 is a unique tripeptide sequence found in the carboxyl terminus of the mature proteins. In contrast, PTS2 is involved in a cleavable amino terminal presequence of peroxisomal proteins that are synthesized as a precursor protein with larger molecular mass.

We identified 256 gene candidates of PTS1- and PTS2-containing proteins and another 30 genes of non-PTS-containing proteins from the Arabidopsis genome. Custom-made DNA microarrays covering all these genes were used to investigate expression profiles of the peroxisomal genes in various organs. They revealed that peroxisomes in root cells play a pivotal role in polyamine catabolism. We also made a two-dimensional protein map of glyoxysomes and leaf peroxisomes isolated from Arabidopsis and soybean. Peptide MS fingerprinting analyses allowed us to identify novel peroxisomal membrane proteins, i.e., voltage-dependent anion-selective channel and adenine nucleotide carrier 1 (PNC1). We also found that peroxisomal membrane ATP-binding cassette transporter promotes seed germination by inducing pectin degradation under the control of abscisic acid signaling. The overall results provide us with new insights into plant peroxisomal functions.

Bioinformatic analysis of the Arabidopsis genome predicted the presence of 15 kinds of genes, called PEX genes, for peroxisomal biogenesis factors. We demonstrated that PEX5 and PEX7 form a cytosolic receptor complex and recognize PTS1- and PTS2-containing proteins, respectively. PEX14 is a peroxisomal membrane docking protein that captures the receptor-cargo complex. We also comprehensively investigated whether or not these predicted PEX genes function in peroxisome biogenesis by generating knockdown mutants that suppress PEX gene expression by RNA-interference. Phenotypes of these mutants allowed us to identify the functional PEX genes, which can be classified into two groups: PEX genes regulating for peroxisomal protein import and PEX genes regulating for peroxisomal morphology. We continue to investigate the detailed molecular functions of other PEX genes. Of these, we proposed that PEX10 is essential for the maintenance of ER morphology and for biosynthesis of cuticular wax.

III. Identification of novel components essential for peroxisome biogenesis

To better understand peroxisome biogenesis, we isolated a number of Arabidopsis mutants having aberrant peroxisome morphology (apem mutants) based on them having a
different pattern of GFP fluorescence from the parent plant, GFP-PTS1, in which peroxisomes with normal sizes and numbers can be visualized with GFP.

Of these apem mutants, APEM1 gene (whose defect causes the elongation of peroxisomes and mitochondria) encodes dynamin-related protein 3A, one member of the dynamin family. APEM2 and APEM4 (whose defects cause a decrease in the efficiency of protein transport) were revealed to encode proteins homologous to PEX13 and PEX12, respectively, and both proteins are components of the protein-translocation machinery on peroxisomal membranes.

In addition, we reported on characterization of other apem mutants, apem3 and apem9. APEM3 encodes Peroxosomal membrane protein 38, and its defect causes enlargement of peroxisomes. APEM9 is the plant-specific PEX that has a role in tethering the PEX1-PEX6 complex on peroxisomal membranes (Figure 1).

We are currently analyzing the functions of other APEM proteins such as APEM10 and APEM11. From these analyses, we will be able to identify the components responsible for peroxisome biogenesis, and to address the mechanism at the molecular level.

IV. ER derived organelles for protein storing and defense strategy

Plant cells develop various types of endoplasmic reticulum (ER)-derived structures with specific functions. ER bodies are ER-derived compartments observed in Arabidopsis. They are rod-shaped structures surrounded by ribosomes, and are widely distributed in the epidermal cells of whole seedlings. Undamaged rosette leaves have no ER bodies, but accumulate ER bodies after wounding or jasmonic acid treatment. This suggests that ER bodies function in the defense against herbivores. ER bodies in seedlings include PYK10, a β-glucosidase with an ER retention signal. When plant cells are damaged, PYK10 forms large protein aggregates that include other β-glucosidases (BGLUs), GDSL lipase-like proteins (GLLs) and cytosolic jacalin-related lectins (JALs). The aggregate formation increases glucosidase activity, possibly producing toxic products (Figure 2). Arabidopsis nai1 mutants have no ER bodies in the entire plant and do not accumulate PYK10. NAI1 encodes a transcription factor that has a basic-helix-loop-helix (bHLH) domain and regulates the expression of PYK10 and NAI2. The Arabidopsis nai2 mutant has no ER bodies and reduced accumulation of PYK10. NAI2 encodes a unique protein that localizes to the ER body. We found that the membrane protein of ER body 1 (MEB1) and MEB2 are integral membrane proteins of the ER body. NAI2 deficiency relocates MEB1 and MEB2 to the ER network. These findings indicate that NAI2 is a key factor that enables ER body formation. We are now investigating the function of NAI2 on ER body formation by heterologously expressing it in onion and tobacco cells.

V. Vacuoles responsible for programmed cell death in plants

The vacuolar processing enzyme (VPE) belongs to the cysteine protease family found in higher plants and animals. VPE is responsible for the maturation of various types of vacuolar proteins. We revealed a novel function of VPE in various instances of programmed cell death (PCD) in plants. VPE is identified as the proteinase that exhibits caspase-1
activity in plants. The plant hypersensitive response (HR), a type of defense strategy, constitutes well-organized PCD. No HR occurs on the tobacco mosaic virus-infected leaves of VPE-deficient tobacco plants. These results suggest that VPE is involved in vacuolar collapse, which triggers PCD.

Using inhibitors for caspase-3 and the proteasome (also known to affect animal cell death), we found that the activities of both are required for bacterium-induced cell death in plants. RNA interference-mediated silencing confirmed that one of the three Arabidopsis proteasome catalytic subunits, PBA1, is required for the fusion of the vacuolar and plasma membranes, which triggers PCD.

Plants evolve a death strategy mediated by vacuolar systems, which are not seen in animals. Interestingly, vacuoles are the key players in the plant-specific cell death system.

VI. Roles of molecular chaperones on cell differentiation

Molecular chaperones are cellular proteins that function in the folding and assembly of certain other polypeptides into oligomeric structures. To clarify the roles of molecular chaperones on cell differentiation, we have purified and characterized chaperonin and HSP70s and analyzed their roles in the translocation of proteins into chloroplasts.

We found that HSP90 inhibitor induced genes with heat shock response element (HSE) motifs in their promoters, suggesting that heat shock transcription factor (HSF) is involved in the response. Arabidopsis HSFs interacted with HSP90.2. Thus, it appears that in the absence of heat shock, HSP90 actively suppresses HSF function. During heat shock, HSP90 is transiently inactivated, which leads to HSF activation. This data indicates that HSP90 regulates correct gene expression for heat acclimatization in plants. We also observed that HSP90 is involved in hormone responses in Arabidopsis. The evolutionary and functional characterizations are now being investigated.

VII. Update of The Plant Organelles Database 2 (PODB2) and release of Plant Organelles World

The Plant Organelles Database 2 (PODB2) was built to promote a comprehensive understanding of organelle dynamics. This public database is open to all researchers. PODB2 consists of four individual units: the organelles movie database, the organellome database, the functional analysis database, and external links. The organelles movie database contains time-lapse images and 3D structure rotations. The organellome database is a compilation of static image data of various tissues of several plant species at different developmental stages. The functional analysis database is a collection of protocols for plant organelle research. The amount of included data is increasing day by day. We will add new content, which is dedicated to organelle movement and morphology in response to environmental stimuli, soon. It is expected that PODB2 will contribute to systems biology through the combination of the included data with other ‘omics’ data and computational analyses. In addition, we released a new website, Plant Organelles World, which is based on PODB2 as an educational tool to engage members of the non-scientific community. We expect that PODB2 and Plant Organelles World will enhance the understanding of plant organelles among researchers and the general public who want to explore plant biology.

![Figure 3. The graphical user interface of the organelles movie database in PODB2 (http://podb.nibb.ac.jp/Orgenellome).](image)

Publication List

(Original papers)


(Review Article)