Plants can be infected by a variety of pathogens, most of which can cause severe economic losses. The plants resist the invasion of pathogens via the innate or acquired immune system for surviving biotic stress. The associations between plants and pathogens are sophisticated beyond imaging and the interactions between them can occur at a very early stage after their touching each other. A number of researchers in the past decade have shown that many biochemical events appeared even as early as 5 min after their touching for plant disease resistance response. The early molecular interactions of plants and pathogens are likely to involve protein phosphorylation, ion fluxes, reactive oxygen species (ROS) and other signalling transduction. Here, we reviewed the recent progress in the study for molecular interaction response of fungal pathogens and host plant at the early infection stage, which included many economically important crop fungal pathogens such as cereal rust fungi, tomato 

Cladosporium fulvum, rice blast and so on. By dissecting the earlier infection stage of the diseases, the avirulent/virulent genes of pathogen or resistance genes of plant could be defined more clearly and accurately, which would undoubtedly facilitate fungal pathogenesis study and resistant crop breeding.

1. Introduction

One of the differences between plants and animals is that plants are unable to move. They rely on the immune system to perceive and identify a pathogen, and then make a series of response mechanisms [1–3]. Most interactions between plants and pathogens start from genetic and molecular aspects. The ‘gene-for-gene’ hypothesis was first proposed by Harold Henry Flor via investigating flax and flax rust race-specific resistance in 1955 [4]. The biochemical basis of this hypothesis is the interaction between resistance (R) gene products and avirulence (Avr) gene products. Plants have developed multiple mechanisms to recognize pathogen invasion and trigger immune responses directly or indirectly [5].

So far, it has been reported that dozens of plant diseases have been caused by pathogenic interaction systems. Many studies have shown that the interactions between plants and pathogens associated with protein phosphorylation. Phosphatases and protein kinases play a vital role in the activation of the early-stage disease resistance responses [6,7]. Rapid phosphorylation response to pathogens and other elicitors includes some downstream mitogen-activated protein kinases (MAPKs) [8–11], calmodulin protein kinases [12] and syntaxin-like proteins [13].

Knowledge of phosphorylation events and their regulation is crucial to understand the mechanisms of plant and pathogen interactions [14]. Protein phosphorylation, a common regulation mode in vivo, plays an important role in cellular signal transduction process. Phosphorylation of proteins occurs mainly on serine (included threonine) and tyrosine, and the enzymes and functions of these two types of amino acids are different.

A decade ago, plant innate immune response regulatory mechanisms were analysed via quantitative phosphoproteomic [15]. The interactions occurred earlier...
2. The early responses of host plants to pathogens

‘Immunity’ is a protective reaction in which the organism maintains their own physiological balance and stability by identifying and eliminating antigenicity foreign body. The plant immune system formed two divergent branches in the long-term evolution and development process: the sophisticated and specific adaptive immunity and the more universal innate immunity. The adaptive immune system can specifically recognize and selectively remove invading pathogens. However, it would take several weeks to form a sustained response and the majority of organisms lack this acquired immune system [23,24]. Compared with the adaptive immune system, the innate immune system does not need specialized immune cells to develop a protective response [24].

The innate immune system, on the other hand, involves a population of cells and signalling pathways that constitutively function to respond rapidly to pathogens at the site of infection [24,25]. The cells of the innate immune system detect pathogen-associated molecular patterns (PAMPs) and microbe-associated molecular patterns (MAMPs) via their pattern recognition receptors (PRRs) [24,26]. PRRs include NOD-like receptors (NLRs), Toll-like receptors (TLRs), RIG-I-like receptors (RLRs) and C-type lectin receptors (CLR) [27]. The plant has formed two kinds of innate immune mechanism—PAMP-triggered immunity (PTI) and effector-triggered immunity (ETI)—that lead to a rapid disease response in the process of long-term cooperative coevolution with pathogen [28–31]. The two branches of the plant immune system can be described as a ‘zigzag’ model (figure 1) [28]. ETI was first discovered in plants that are dependent on the plant resistance proteins (R proteins) to identify pathogen-secreted proteins directly or indirectly and activate a strong resistance reaction inhibiting pathogen infection [24]. PTI is a relatively weak resistance reaction activated by PRR identifying conservative pathogen PAMPs [32]. Currently, the molecular mechanisms of PTI and ETI defence response have been deeply investigated in Arabidopsis thaliana [33]. The immune system is the foundation of the interaction between fungi and host.

Some scholars have found that the early response of crops’ cells or tissues was triggered by elicitors. For example, fungal elicitor specifically induces the proteins transient, rapid and consecutive phosphorylation in the parsley (Petroselinum crispum) cells, and the phosphorylation
activates some pathogen resistance-related genes. In the microsome and part of the cytoplasm, a neutral 45 kDa protein phosphorylated as early as 1 min after being treated with elicitor and the increase of a 26 kDa nuclear protein phosphorylation starts also at the earliest stage [34].

### 3. The signal transduction processes in the early response

The initial touching of pathogen and plant would rapidly trigger the signal transduction process on the plasma membrane and cytoplasm of plant cells [35]. The involved signal transduction in the early response covered many pivotal channels which can set subsequent responses at a multi-level of gene expression patterns. Among them, the signal components of ion flux, salicylic acid (SA) and other hormones were mostly investigated in recent reports [35–37].

The phosphorylation reaction was associated with the presence of Ca\(^{2+}\) involved in the signal transduction processes [34]. Protein phosphorylation events occurred in vivo within minutes when elicitor treated tobacco cells. The function of elicitor was completely blocked by the protein kinase inhibitors K-252a and staurosporine. The protein kinase inhibitors can also inhibit the early biochemical responses induced by elicitors [38].

The ion flux events of plant responses to MAMPs occurred within approximately 0.5–2 min [39,40]. These changes include increased influx of Ca\(^{2+}\) and efflux of K\(^{+}\), and an efflux of anions, particularly of nitrate [41]. The ion fluxes lead to membrane depolarization [42]. Even though little is known about the ion channels, MAMPs were evident to stimulate an influx of Ca\(^{2+}\) within approximately 0.5–2 min [39,40]. These changes include increased influx of Ca\(^{2+}\) within approximately 0.5–2 min [39,40].

The RPG1 protein is a functional kinase located in the plasma membrane, endomembranes and cytosol. The resistance protein RPG1 disappeared rapidly (within 5 min) when barley seedling leaves were inoculated by avirulent and viable stem rust fungus pathotype MCCF (figure 2). The appearance of the RPG1 protein is due to phosphorylation and the phosphorylated status sustained for 20 h after inoculation. It is suggested that RPG1 protein phosphorylation is essential for disease resistance. The reciprocal responses of barley and the phosphorylated status sustained for 20 h after inoculation. It is suggested that RPG1 protein phosphorylation is essential for disease resistance. The reciprocal responses of barley and the phosphorylated status sustained for 20 h after inoculation.

### 4. Plant disease resistance genes involved in the early response

A few important studies of host resistance genes were initiated at early infection stage (table 1). The milestone report for the early interaction was published in 2011 for barley stem rust [51]. It was a devastating disease caused by *Puccinia graminis f. sp. tritici* (Pgt) for barley production in most areas of North America until the barley cultivars (cvs.) were first announced in 1942. Henceforth, *Rpg1* gene has protected barley cultivars from severe stem rust losses for over 70 years. The *Rpg1* located in the short arm of barley chromosome 1(7H) is a novel resistance gene homology with receptor kinases [58–60]. The highly susceptible cultivar Golden Promise became a high resistance disease due to transformed the *Rpg1* gene by genetic engineering [61]. Although the resistance of the *Rpg1* gene is wide, it cannot function to all virulent types of Pgt [62]. The *Rpg1* gene encodes a constitutively expressed protein containing two tandem kinase domains: the protein kinase 1 (pK1) domain and protein kinase 2 (pK2) domain. The pK1 is a pseudokinase, whereas the pK2 domain is catalytically active, and both domains are required for stem rust resistance. The pseudokinase pK1 domain is associated with disease resistance and the pK2 domain is involved in protein phosphorylation [63,64].

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<table>
<thead>
<tr>
<th>plant</th>
<th>gene</th>
<th>hpi</th>
<th>pathogen</th>
</tr>
</thead>
<tbody>
<tr>
<td>barley</td>
<td><em>Rpg1</em></td>
<td>5 min</td>
<td><em>Puccinia graminis f. sp. tritici</em> (Pgt) [51]</td>
</tr>
<tr>
<td>tomato</td>
<td><em>CI-9</em></td>
<td>5 min</td>
<td><em>Cladosporium fulvum</em> [12]</td>
</tr>
<tr>
<td>maize</td>
<td>CCT7</td>
<td>3</td>
<td><em>Curvularia lunata</em> [52]</td>
</tr>
<tr>
<td>rice</td>
<td>OsAAE3</td>
<td>12</td>
<td><em>Magnaporthe oryzae</em> [53]</td>
</tr>
<tr>
<td>wheat</td>
<td>TaCDPK2</td>
<td>4</td>
<td><em>P. triticina</em> (Pt) [54]</td>
</tr>
<tr>
<td>wheat</td>
<td>TaGAMTA4</td>
<td>4</td>
<td><em>Pr</em> [55]</td>
</tr>
<tr>
<td>wheat</td>
<td>Lr57</td>
<td>24</td>
<td><em>P. striiformis f. sp. tritici</em> (Pst) [57]</td>
</tr>
<tr>
<td>wheat</td>
<td>TaCERK1, TaCEBiP, TaRboh</td>
<td>24</td>
<td><em>P. striiformis f. sp. tritici</em> (Pst) [57]</td>
</tr>
</tbody>
</table>

### Table 1. Related plants genes response to pathogen at early plant–pathogen interaction. hpi: hours post inoculation; gene: the gene of host plant involved in the early interaction significantly.
The process of the activation is involved in Ca\(^{2+}\) kinases achieved through post-translational mechanisms, and responses and tyrosine phosphorylation [9,10,66]. Serine/threonine. SIPK activation was involved in resistance.

Cf-9 responded to race-specific elicitor Avr9, but not by race-specific elicitor g22P.aer [13]. Increased at 24 h, which were triggered by the race-specific signalling pathway. Finally, the first 454 sequencing was performed at 4 days after peanut inoculation of S. rolfsii. Further studies of possibly related resistance genes and avirulence genes would be useful to the research of early host–pathogen interaction.

The early interactions of blast fungus (M. oryzae) and rice occur at the apoplastic pathogen genes involved in the early interaction (table 2).

<table>
<thead>
<tr>
<th>Pathogen genes involved in the early response</th>
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<tbody>
<tr>
<td>Very few studies and discoveries were published on fungal pathogen genes involved in the early interaction (table 2).</td>
</tr>
</tbody>
</table>
Table 2. Pathogen genes involved in the early response to plant. hpi: hours post inoculation; gene: the genes of fungal phytopathogen involved in the early interaction significantly.

<table>
<thead>
<tr>
<th>pathogen</th>
<th>gene</th>
<th>hpi</th>
<th>interaction plant</th>
</tr>
</thead>
<tbody>
<tr>
<td>Puccinia graminis f. sp. tritici</td>
<td>RGD-binding gene, VPS9 gene</td>
<td>5 min</td>
<td>barley [51]</td>
</tr>
<tr>
<td>Cladosporium fulvum</td>
<td>Avr9</td>
<td>3–5 min</td>
<td>tomato [11,67]</td>
</tr>
<tr>
<td>Phytophthora sojae</td>
<td>Avr1b</td>
<td>24</td>
<td>soya bean [74]</td>
</tr>
<tr>
<td>Colletotrichium higginsianum</td>
<td>ChMK1</td>
<td>no clear</td>
<td>cruciferous crops [75]</td>
</tr>
<tr>
<td>Magnaporthe oryzae</td>
<td>MGS0074, MGS0274, MGS0338, MGS0718,</td>
<td>24</td>
<td>rice [76]</td>
</tr>
<tr>
<td></td>
<td>MGS0997, MGS1242, and MGS1460 in Y99-63</td>
<td></td>
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</tr>
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</table>

Also, the mechanisms of some known avirulent genes (e.g. stem rust genes: the RGD-binding gene and the VPS9 gene) have been unknown.

In their research of molecular interaction between stem rust and barley, Nirmala et al. found that the arginine-glycine-aspartic acid peptide loops can prevent the formation of adhesion structures for spore attachment, the germination of the dynamic spores and the phosphorylation of Rpg1 [51]. They purified and identified two proteins: arginine-glycine-aspartic acid (RGD)-binding protein with fibronectin type III and breast cancer type 1 susceptibility protein domains, and vacuolar protein sorting-associated protein 9 (VPS9 protein) with a coupling of ubiquitin to endoplasmic reticulum degradation domain from the ungerminated avirulent rust spores via the arginine-glycine-aspartic acid affinity chromatography. RGD-binding protein and VPS9 protein together induce hypersensitive response (HR) in vitro, phosphorylation and degradation of Rpg1 in barley with a functional Rpg1 gene. The RGD-binding gene and the VPS9 gene are constitutively expressed in almost all cells of the avirulent race of stem rust fungus MCCF [58]. So far, the understanding of these two genes is not comprehensive. There is still a lot of work to do describing their functional network.

The avirulent gene Avr9 of tomato leaf mildew is another disease-related gene involved in the early interaction. The research results for Avr9 genes was initially compared even with those of VPS9 gene and RGD-binding gene of rust fungus. Avr9 gene encodes a preprotein that contains 63 amino acids which could function during the interaction between the fungus Cladosporium fulvum and tomato. Avr9 generated K$^+$ outward-rectifying by 2.5-fold to threefold and almost completely suppressed inward-rectifying of K$^+$ within 3–5 min. The K$^+$ channel reactions were specific and irreversible [11,68].

Another important research was reported for oomycetes. Shan et al. first cloned the avirulence gene A1r1b of oomycete pathogen Phytophthora sojae by fine structural genetic mapping [74]. The A1r1b gene contains two genes: the A1r1b-1 gene and the A1r1b-2 gene. The A1r1b-1 gene was localized to a single 60 kb bacterial artificial chromosome (BAC). The A1r1b-1 gene is polymorphic and encodes a small, hydrophilic secreted protein that is a specific elicitor. The A1r1b-1 protein triggered a specific and systemic HR in soya bean leaves (carrying the Rps1b resistance gene). A1r1b-1 protein entering the soya bean leaf cells need RXLR (Arg-X-Leu-Arg, × is any amino acid) and dEER (Asp-Glu-Glu-Arg) [77,78]. Experiments revealed that the A1r1b-2 gene required the accumulation of the mRNA of the A1r1b-1 gene. That is to say that A1r1b-2 controlled the accumulation of A1r1b-1 mRNA. One of the advantages of the A1r1b-1 gene is that it could cause the hypersensitive response to spread to the whole plant. But its maximum expression was 24 h and 48 h after inoculation. It is clear that A1r1b-1 gene works later than Avr9 gene, VPS9 gene and RGD-binding gene. And early research of this gene are not clear [74].

In the interaction between Colletotrichium higginsianum and cruciferous crops, Wei et al. investigated a Fus3/Kss1-related MAPK gene (ChMK1) from Colletotrichium higginsianum [75]. The ChMK1 is essential to pathogenicity, appressorium formation, conidiation production, cell wall integrity, growth rate and melanin formation for C. higginsianum. That is to say ChMK1 gene plays an essential role in the early tomato infection.

This kind of early response also could be affected by nutrition limitation [76]. The expression of seven genes (MGS0074, MGS0274, MGS0338, MGS0718, MGS0997, MGS1242 and MGS1460) in Y99-63 (one strain of rice blast fungus), encoding cysteine-rich proteins, were upregulated to different extents in the early M. oryzae–rice interaction under nitrogen limitation. Cysteine-rich proteins might be enrolled in the cross-talking of nitrogen limitation and the early infection response [76].

6. Conclusion and discussion

Here, we reviewed the recent progress in the study for molecular interaction response of fungal pathogens and host plant at the early infection stage, which included some economically important crop fungal pathogens such as cereal rust fungi, tomato Cladosporium fulvum, M. oryzae and so on. According to the research so far, the distinct mechanisms for the early molecular interactions of plants and pathogens are likely to involve protein phosphorylation, ion fluxes, reactive oxygen species (ROS) and other signalling transduction.

The barley leaves start responses–protein phosphorylation and then trigger the disease resistance mechanism within 5 min of inoculating the stem rust pathogen avirulent ureidiospores MCCF. During the interaction between tomato and leaf mildew, the product of the fungus Cladosporium fulvum avirulence gene Avr9 resulted in K$^+$ salt loss by 2.5-fold to threefold and almost complete suppression of K$^+$ salt within 3–5 min. The Rpg1 gene and the Cf-9 gene can resist the invasion of the VPS9 gene, the RGD-binding gene and the Avr9 gene rapidly via the trigger of disease-resistant mechanisms. The examples of these two interactions showed that plants and pathogens recognize each other rapidly after touching and then trigger the signal pathways respectively to achieve molecular interactions of plants and pathogens.
the purpose of disease resistance or infection. While we know that there are interactions between R gene and Avr gene, the disease resistance mechanisms and a series of signalling pathways remain to be studied.

Research into molecular responses at early infection stage for fungal pathogen and host plant interactions are of significance for pathogenesis study and resistant crop breeding. Understanding of the molecular events occurring at the early interaction stage would be an essential step for describing the initial mechanism of pathogen–host interactions in many important agricultural disease systems and medical pathogen systems. The candidate genes revealed by the study of early infection could bring out the target genes for crop improvement by transgenic methods or genome editing. The manifest conclusions obtained in this field are still limited. Further investigations and new technologies should be employed in this direction, which will attract more and more researchers to join this creative area and contribute to the agriculture.

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