INTRODUCTION

Sclerotinia sclerotiorum is one of the most aggressive and widespread necrotrophic plant pathogens, infecting more than 400 plant species. The fungus causes substantial economic losses of many economically important crops, such as oilseed rape, soybean, sunflower, and various vegetables (Bolton et al., 2006; Derbyshire et al., 2017; Navaud et al., 2018). The fungus can produce many cell...
wall-degrading enzymes and oxalic acid as virulence factors to facilitate its colonization of host plants, which involves numerous physiological processes (Cessna et al., 2000; Kabbage et al., 2015; Nuhse et al., 2000; Riou et al., 1991). Recently, it was suggested that the pathogen might have a rather short biotrophic lifestyle in the early infection stage (Kabbage et al., 2015).

Plant defence against S. sclerotiorum is complex and exhibits a distinct quantitative trait locus disease response (Badet et al., 2019; Perchepied et al., 2010). To tackle the menace of this destructive pathogen, it is critical to understand the mechanisms of plant–S. sclerotiorum interaction. In general, plant immunity is classified into two types: (a) pathogen-associated molecular pattern (PAMP)-triggered immunity (PTI)/microbe-associated molecular pattern (MAMP)-triggered immunity and (b) effector-triggered immunity (ETI) (Jones & Dangl, 2006). To date, ETI has not been observed in the plant–S. sclerotiorum pathosystem (Wang et al., 2019a; Yang et al., 2018). In contrast, an increasing body of evidence suggests that PTI plays a predominant role in plant resistance to S. sclerotiorum. Several elicitors are involved in evoking PTI, such as chitin (Le et al., 2014), Sclerotinia culture filtrate elicitor1 (SCFE1) (Zhang et al., 2013), cutinase (ScCut) (Zhang et al., 2014), SsSm1 (a cerato-platanin family protein) (Pan et al., 2018), and Vme02 (Nie et al., 2019).

In general, PTI contributes to plant basal resistance and plays a vital role in a plant’s resistance to numerous potential pathogens. PTI can be further divided into early and late response processes. The early reactions of PTI include calcium influx, production of reactive oxygen species (ROS), and activation of mitogen-activated protein kinases (MAPKs) to trigger transcriptional processes regulated by plant transcription factors (Tena et al., 2011). Callose deposition and PTI marker gene transcription are observed in later responses (Zipfel & Robatzek, 2010).

After the initiation of PTI and ETI, defence signals are transduced through a complex signalling network, resulting in changes in the expression of thousands of host genes and defensive responses (Lewis et al., 2015; Zipfel, 2014; Zipfel et al., 2006). Phytohormones such as jasmonic acid (JA), salicylic acid (SA), and ethylene (ET) function as signalling molecules to regulate signal transduction. Recent studies revealed that SA, JA, ET, auxin, and abscisic acid (ABA) are all involved in regulating resistance against S. sclerotiorum (Guo & Stotz, 2007; Nováková et al., 2014; Perchepied et al., 2010; Stotz et al., 2011; Wang et al., 2012, 2019b, 2020; Xu et al., 2014, 2018; Zeng et al., 2020; Zhou et al., 2015). Among these pathways, the JA signalling pathway plays an essential role in plant resistance to S. sclerotiorum invasion. Wang et al. (2012) and Oliveira et al. (2015) found that exogenous application of methyl jasmonate (MeJA) could induce plant resistance against S. sclerotiorum. Furthermore, JA production was also found to increase after infection by S. sclerotiorum (Ranjan et al., 2019; Wang et al., 2012; Wei et al., 2010). Transcriptomic and proteomic analyses have uncovered many genes and proteins involved in the JA signalling pathway, which is associated with S. sclerotiorum infection (Joshi et al., 2016; Liang et al., 2009; Wu et al., 2016; Zhao et al., 2007, 2009). JA has also been established to regulate plant resistance to S. sclerotiorum (Guo & Stotz, 2007; Takahashi et al., 2010). A large body of evidence indicates that altered expression of some defence-related genes could induce higher expression of JA-regulated genes (Chen et al., 2013; Le et al., 2014; Liu et al., 2018; Stotz et al., 2011; Wang et al., 2014b; Yang et al., 2019), including MAPK genes (Wang et al., 2009, 2019a).

To identify the genes involved in plant defence against S. sclerotiorum, we screened T-DNA-inserted homozygous mutants from the Arabidopsis Biological Resource Center by determining the phenotypes of resistance to S. sclerotiorum 1980. We obtained a mutant line, SALK_026489C, which was named an-t1 and has a T-DNA insertion that causes a null allele of the gene ANGUSTIFOLIA (AN, AT1G01510) (Gachomo et al., 2013). AN encodes a plant homologue of mammalian C-terminal-binding protein/brefeldin A-ADP ribosylated substrate (CtBP/BARS) (Kim et al., 2002). AN is involved in regulating a myriad of plant growth processes, development, and abiotic stress responses, including control of leaf and cell shapes, root formation, and resistance to pathogen infection. It also mediates the organization of microtubules, the alignment of actin filaments, and the production of ROS in Arabidopsis (Bai et al., 2013; Bhasin & Hulskamp, 2017; Dang et al., 2018; van Dop et al., 2020; Folkers et al., 2002; Gachomo et al., 2013; Iwabuchi et al., 2019; Kim et al., 2002; Tsuge et al., 1996). In this study, we found that AN negatively regulates Arabidopsis immunity against the necrotrophic pathogen S. sclerotiorum through the PTI and JA signalling pathways at the early infection stage.

2 | RESULTS

2.1 | ANGUSTIFOLIA negatively regulates Arabidopsis resistance to S. sclerotiorum at the early infection stage

To establish the role of AN in resistance/susceptibility to S. sclerotiorum, Arabidopsis an-t1 mutant leaves were detached and inoculated with mycelial plugs of S. sclerotiorum 1980. At 24 h postinoculation (hpi), an-t1 developed weaker symptoms than wild-type Col-0 (Figures 1a,b and S1a–c). To confirm the function of AN, an-t2 (another knockout mutant of AN that has a different insertion site from that of an-t1) was similarly infected with S. sclerotiorum 1980. Like an-t1, the an-t2 mutant also showed stronger resistance to the pathogen than wild-type plants at 24 hpi (Figures 1a,b and S1a). However, the mutant showed similar susceptibility to the pathogen at the late infection stage (48 hpi) when compared to wild-type plants (Figure 1b,c). These results demonstrate that AN functions as a negative regulator of Arabidopsis resistance to S. sclerotiorum at the early stages of infection (at 24 hpi).

To further confirm the role of AN at the early stages of S. sclerotiorum infection, invasion and invasive hyphal development in infected leaf tissue were visualized by staining with trypan blue. As shown in Figure 1c, S. sclerotiorum formed extensive hyphae that ramified throughout the mesophyll in Col-0 leaves. In contrast, hyphae were significantly reduced in the an-t1 and an-t2 mutant leaf tissues.
We used *Botrytis cinerea* strain 05.10, a related necrotrophic pathogen, to challenge the *an* mutants and the wild-type Col-0. Like *S. sclerotiorum* infection (Figure S1b,c), the *an*-t1 mutant was significantly more resistant than wild-type Col-0 before 72 hpi, but the resistance vanished in the *an*-t1 mutant at 84 hpi (Figure S1d,e). These results suggest that AN is involved in immunity to necrotrophic pathogens such as *S. sclerotiorum* and *B. cinerea* at the early stage of infection.

### 2.2 | *Arabidopsis* lines overexpressing AN are highly susceptible to *S. sclerotiorum*

To explore the function of AN in regulating defence responses to necrotrophic pathogen infection, a complemented line (*an*-t1 COM#) with the native promoter in the *an*-t1 background was inoculated with *S. sclerotiorum*. We found a comparable resistance between *an*-t1 COM# and Col-0 (Figures 2a,b and S1a). In addition, we analysed a transgenic *Arabidopsis* line overexpressing AN in the *rdr6*-11 background (AN-OE). The mutation of *RDR6* prevents silencing of the exogenous transgene, because *RDR6* is complementary to certain transcripts of silenced transgenes, which hence serve as templates for synthesis of double-stranded RNA to induce RNA silencing in plants (Betti et al., 2021; Harmoko et al., 2013). AN-OE plants were more susceptible to *S. sclerotiorum* than *rdr6*-11 plants (Figures 2c,d and S1a). Moreover, microscopic examination and trypan blue staining revealed that the level of invasion was comparable between *an*-t1 COM# and wild-type plants (Figure 2e,f), and AN-OE plants had more abundant hyphal colonization than *rdr6*-11 (Figure 2g,h). These data are consistent with a negative role of AN in antifungal immunity.

We hypothesized that loss of function of AN might also confer immunity against bacterial pathogens. To test this hypothesis, *an* mutants were inoculated with the hemibiotrophic bacteria *Pseudomonas syringae* pv. *tomato* (Pst) DC3000 and *P. syringae* pv. *maculicola* (Psm) ES4326. At 3 days postinoculation (dpi), the *an*-t1 mutant developed weaker disease symptoms (chlorosis and necrosis) than wild-type plants (Figure S2a). The bacterial titres were significantly lower in the *an*-t1 and *an*-t2 mutants and markedly higher in AN-OE than in control plants at 3 dpi with *P. syringae* (Figure S2b). These results are consistent with some previous studies (Gachomo et al., 2013; Xie et al., 2020) that demonstrated that AN regulates biotic and abiotic stress responses by steady-state regulation of H$_2$O$_2$ accumulation and resistance gene expression.

Together, these results indicate that AN regulates resistance to *S. sclerotiorum* and *P. syringae*, revealing that AN may serve as a regulator in the plant defence responses to both pathogens.

### 2.3 | AN suppresses *S. sclerotiorum*- and chitin-induced ROS production

Gachomo et al. (2013) and Dang et al. (2018) reported that H$_2$O$_2$ accumulated in AN knockout mutants of *Arabidopsis*. *S. sclerotiorum* releases oxaloacetic acid to suppress host ROS production to facilitate colonization at the early stage of infection (Kabbage et al., 2015). This prompted us to investigate ROS production in AN knockout mutants upon PAMP or MAMP treatment. First, we analysed ROS production by histochemical staining. The nitroblue tetrazolium (NBT) staining results revealed that the *an*-t1 and *an*-t2 mutants accumulated a higher level of ROS at the infection sites.
than wild-type plants (Figure 3a,b). In contrast, the AN-OE plants showed a visible reduction in ROS accumulation compared to rdr6-11 (Figure 3a,b). Furthermore, 3,3'-diaminobenzidine (DAB), another ROS (hydrogen peroxide) indicator, was used for staining and showed that an-t1 and an-t2 mutants accumulated higher levels of ROS at the infection sites than the wild-type plants at 24 hpi (Figure 3c,d). However, the AN-OE plants had a significantly reduced level compared with rdr6-11 (Figure 3c,d). We also measured PAMP-triggered oxidative burst in the an mutants. AN knockout mutants exhibited a more pronounced level of oxidative burst than the wild-type plants, while AN-OE plants demonstrated a strongly compromised PAMP-induced oxidative burst compared with the rdr6-11 background (Figures 3e,f and S3a,b). These results indicate that AN negatively regulates ROS accumulation upon MAMP or PAMP triggers.

2.4 | AN negatively regulates MAPK activation and late PTI responses

ROS production is a typical early event of PTI. To test whether AN is required for broad PTI events, we analysed chitin- and flg22-induced MAPK activity in the an mutants. Consistent with previous studies, we detected a significant activation of MPK6 and MPK3 within 5 and 10 min upon chitin or flg22 elicitation in the wild-type seedlings (Figures 4a,b and S3c,d) (Asai et al., 2002; Galletti et al., 2011; Yeh et al., 2016). The an-t1 seedlings demonstrated a stronger activation of MPK3 and MPK6 than the wild type at 5 and 10 min after treatment (Figures 4a,b and S3c,d). However, MPK3 and MPK6 activation was weaker in AN-OE than in the wild type at 5 min after chitin treatment or at 10 min after flg22 treatment (Figures 4a,b, and S3c,d), suggesting that AN also negatively regulates chitin- or flg22-induced MPK activation.

Furthermore, we analysed the late PTI responses in the AN knockout mutants. After treatment with chitin or flg22 for 12 h, callose deposition in the an-t1 and an-t2 mutants reached higher levels...
FIGURE 3 AN suppressed reactive oxygen species (ROS) accumulation. (a) Nitroblue tetrazolium (NBT) staining was used to visualize the oxidative burst in response to Sclerotinia sclerotiorum 1980 infection. The experiments were repeated twice with similar results. (b, c) Quantitative analysis of ROS accumulation by ImageJ software after NBT staining (b) and 3,3′-diaminobenzidine (DAB) staining (c). Values represent the maximum, upper quartile, median, lower quartile, and minimum from two independent experiments, each consisting of eight leaves ($n=8$). Different letters denote significant differences based on two-way analysis of variance followed by Tukey post hoc tests ($p<0.05$). (d) DAB staining was used to visualize the oxidative burst in response to S. sclerotiorum 1980 infection. The experiments were repeated twice with similar results. (e, f) Monitoring of ROS production in $an$ mutants treated with 10 $\mu$g/ml chitin for 60 min. After elicitation, relative ROS levels in Arabidopsis leaf discs were quantified using relative light units (RLUs). Values are the mean ± SD of three independent experiments, each with 16 leaf discs ($n=16$). Different letters indicate significant differences between mutants and wild-type control based on two-way analysis of variance followed by Tukey post hoc tests ($p<0.05$).
than in wild-type leaves (Figures 4c,d and S3e,f). In contrast, less callose deposition was observed in the AN-OE lines (Figures 4c,d and S3e,f). To further evaluate the role of AN in late PTI responses, we monitored the expression levels of the PTI marker genes pathogenesis-related protein 1 (PR1) and plant defensin 1.2 (PDF1.2) after inoculation with S. sclerotiorum or flg22 treatment. PR1 was induced at a higher level in the an-t1 mutant than in wild-type plants upon S. sclerotiorum infection or flg22 treatment (Figures 4e and S3g), whereas PDF1.2 was expressed at lower levels before 12 h and subsequently up-regulated at 24 hpi with S. sclerotiorum (Figure 4e).

These results suggest that AN is an essentially negative regulator of pathogen- or PAMP-induced PTI responses.

### 2.5 AN affects JA accumulation after S. sclerotiorum infection

PR1 and PDF1.2 are marker genes for the SA and JA signalling pathways, respectively. Considering the expression patterns of these two genes, we monitored the levels of SA and JA in response to S. sclerotiorum infection. SA and JA accumulation in the an-t1 plants was comparable to that in wild-type plants in the absence of pathogen (Figures 5a and S4). Interestingly, in contrast to SA, the JA content significantly increased to higher levels in the an-t1 mutant from 12 to 24 hpi with S. sclerotiorum compared to wild-type plants (Figures 5a and S4), while it drastically reduced at 30 hpi, probably due to the extension of lesions in the an-t1 mutant at the late stage of infection. Moreover, the expression of AN was rapidly suppressed by MeJA treatment, reaching a minimum at 3 h after treatment (Figure 5b). These results strongly suggest that the involvement of AN in the response to S. sclerotiorum at the early stage of infection is related to the JA signalling pathway.

Previously, Dang et al. (2018) identified allene oxide cyclase 2 (AOC2), an essential enzyme in JA biosynthesis, as an AN-interacting candidate protein and also reported that AN did not interact with catalase 2 (CAT2). Therefore, to understand how AN regulates JA accumulation, we first verified the interaction between AOC2 and AN by luminometry in Nicotiana benthamiana, using CAT2 as a negative control. Strong fluorescence was only detected when AN and AOC2 were coexpressed (Figure 5c), indicating AN interacts with AOC2. Next, we performed a bimolecular fluorescence complementation assay by transiently coexpressing YFP-AN and YFP-N-AOC2 in N. benthamiana. YFP fluorescence was observed only upon coexpression of YFP-AN with YFP-N-AOC2 but not in the controls (Figure 5d). Our coimmunoprecipitation assay results in Arabidopsis protoplasts support this interaction (Figure 5e). Our in vitro pull-down analysis showed that His-tagged AN interacted with glutathione S-transferase (GST)-tagged AOC2 (Figure 5f), indicating AN directly interacts with AOC2.

In Arabidopsis, the AOC family has four members (AOC1, AOC2, AOC3, and AOC4), which have remarkable functional redundancy (Pollmann et al., 2019; Stenzel et al., 2003, 2012). We investigated the interaction between AN and the other AOC family members using luminometry and coimmunoprecipitation assays. The results showed that AN interacted with all AOC family members (Figure S5a,b). These data illustrate that AN may be involved in JA synthesis through direct interaction with AOCs in response to necrotrophic pathogen infection.

### 2.6 AOC1 and AOC2 positively regulate plant resistance to S. sclerotiorum

To further determine the function of AOCs in the defence response to S. sclerotiorum, we obtained two T-DNA insertion lines from AraShare: SALK_054568 (named aoc3-1), which has an insertion in the first exon of AOC3, and SALK_124897 (named aoc4-1), which has an insertion in the second exon of AOC4 (Figure S6a). Reverse transcription-quantitative PCR results showed that the expression of AOC3 and AOC4 was significantly reduced in the aoc3-1 and aoc4-1 mutants, respectively (Figure S6b). To further explore the function of AOC3 and AOC4 in the Arabidopsis immune response against S. sclerotiorum, we performed a detached-leaf inoculation assay using S. sclerotiorum 1980. Twenty-four hours later, the susceptibility phenotype of aoc3-1 and aoc4-1 was similar to that of wild-type plants (Figure S6c,d).

We failed to obtain T-DNA insertion mutants of AOC1 and AOC2 from AraShare. We generated loss-of-function mutants of the two...
genes in a Col-0 background by the clustered regularly interspaced short palindromic repeats (CRISPR)/CRISPR-associated protein 9 (Cas9) system (Figure S7a,b). Two independent edited lines of each gene, aoc1cas9-1, aoc1cas9-2, aoc2cas9-1, and aoc2cas9-2, were selected for S. sclerotiorum inoculation. Similar to aoc3-1 and aoc4-1, the aoc1cas9-1, aoc1cas9-2, aoc2cas9-1, and aoc2cas9-2 mutants demonstrated a wild-type resistance to S. sclerotiorum 1980 (Figure S7c,d). These results suggest that AOCs are functionally redundant in the immune response against S. sclerotiorum infection or might not be involved in the response to S. sclerotiorum.

To further disclose the functions of AOCs in immune responses, we generated aoc3-1 aoc4-1, aoc1cas9-1 aoc2cas9-1, and aoc1cas9-2 aoc2cas9-1 double mutants by genetic crossing. At 24 hpi with S. sclerotiorum 1980, the aoc3-1 aoc4-1 double mutant was as susceptible as
wild-type plants (Figure 6a,b). Notably, the aoc1 cas9-1 aoc2 cas9-1 and aoc1 cas9-2 aoc2 cas9-1 double mutants developed larger lesions than wild-type plants (Figure 6a,b), suggesting that AOC1 and AOC2 play more crucial roles than AOC3 and AOC4 in the immune response against S. sclerotiorum infection.

3 | DISCUSSION

AN was cloned from A. thaliana and identified as the first member of the CtBP family 20 years ago (Kim et al., 2002). AN has extensive homology in the N-terminus with human CtBP. Mutations in AN lead to multiple morphological phenotypes, including narrow leaves, underbranched trichomes, wider tip angles of conical cells, and increased resistance to various stresses (Bai et al., 2013; Dang et al., 2018; Kim et al., 2002; Xie et al., 2020). Our data reveal that mutations in the AN gene result in increased resistance to necrotrophic fungal pathogen infection in the early stages as well as hemibiotrophic bacteria (Figures 1 and 2, Figures S1 and S2). Plants have developed different defence mechanisms against infection with necrotrophic and hemibiotrophic pathogens, primarily through the JA/ET and SA signalling pathways, respectively. Hence, the elevated resistance of an mutants to different types of pathogens may be the result of perturbations in broad-spectrum immunity, such as the PTI signalling network.

Plants can use PTI as a basal defence mechanism to ward off diverse pathogens (Boller & He, 2009; Boller & Felix, 2009; Huang & Zimmerli, 2014). Perturbation of PTI compromises plant defence...
against fungal and bacterial pathogens (Kim et al., 2014; Tsuda et al., 2009). Our phenotypic studies show that AN negatively regulates broad PTI responses, including callose deposition, MPK3/6 activation, ROS accumulation, and the expression of PTI-responsive genes triggered by MAMPs/PAMPs (Figures 3 and 4, Figure S3). However, the molecular mechanisms by which AN suppresses PTI responses require further exploration. A recent study reported that AN exhibits occasional nuclear localization and transcriptional repressor activity (Xie et al., 2020). AN might suppress the expression of the MAMP-specific marker genes PR1 and PDF1.2 through the transcriptional suppression of PTI-related genes. AN has plant-specific motifs that can be phosphorylated by casein kinase, although leaf shape and trichome branching do not appear to require phosphorylation (Minamisawa et al., 2011). It may be worthwhile to determine in the future whether the phosphorylation is necessary for AN function in plant immunity.

ROS have been considered detrimental to cells because they are toxic and cause damage to proteins, lipids (membranes), and nucleic acids, thereby releasing nutrients for necrotrophic pathogens (Temme & Tudzynski, 2009; Wang et al., 2019b). Other reports have shown that the relationship between the buildup of ROS and decreased resistance to necrotrophic pathogens is more intricate than anticipated and cannot be taken as a general rule (Finiti et al., 2014; Petriacq et al., 2016). Previous studies have shown that the release of oxalate by S. sclerotiorum could enhance fungal pathogenicity by favouring an oxidative burst and maintaining a reducing environment in host cells at the early stage of infection (Williams et al., 2011). Hence, ROS production is essential for governing the outcome of S. sclerotiorum-plant interactions at the early stage of infection (Kabbage et al., 2015). Once the infection is established, necrotrophic S. sclerotiorum benefits from induction of plant ROS generation, leading to programmed cell death of the host tissue involved (Wang et al., 2019b; Williams et al., 2011). Consistent with this observation, we found the temporal resistance of an mutants to the necrotrophic pathogens S. sclerotiorum and B. cinerea, which might be caused by the higher level of ROS production. Despite the destructive activity that causes progressive oxidative damage and ultimately cell death in the host cell, ROS are well-described second messengers involved in regulating a variety of cellular processes, including MAPK activation (Wang et al., 2019a), defence gene re-programming (Kovtun et al., 2000; Wang et al., 2014a, 2015), and defence-related phytohormone SA and JA production (Jagodzik et al., 2018; Wu & Ge, 2004). In this study, we have provided multiple lines of evidence revealing AN’s role in regulating PTI responses, including MAPK activation, transcriptional regulation of defence-related genes, and JA accumulation. These PTI responses were probably initiated by ROS, which are negatively regulated by AN. It may be worthwhile to examine the requirement of ROS production for AN’s function in regulating other PTI responses by reducing the level of ROS in an mutants through toxicology and genetic approaches. However, the possibility of feedback regulation of ROS production through other PTI responses that are mediated by AN cannot be ruled out. The interplay between ROS and defence-related hormones plays an essential role in controlling plant defence responses (Xia et al., 2015), indicating the reciprocal regulation of ROS and SA and JA/ET signalling pathways. The detailed role of ROS production and the relevance to pathogen infection of AN require further exploration by further characterization of the AN-associated proteins, which may provide a fundamental understanding of ROS production in necrotrophic pathogen infection.

JA is a vital phytohormone that regulates plant resistance to S. sclerotiorum (Sheard et al., 2010). JA is produced from α-linolenic acid (18:3), which is converted to the intermediate cis-(+)-12-oxophytodienoic acid by enzymes including lipoxygenase, allene oxide
synthase, and AOC in the chloroplast (Wasternack & Hause, 2013). In the peroxisome, cis-(+)-12-oxophytodienoic acid is converted to JA (Bussell et al., 2013). In this study, AN was shown to regulate the JA biosynthesis pathway through direct interaction with AOC family proteins (Figures 5c–f and 5i). It has been reported that heteromerization controls the activity of AOCs, and the perturbation of these protein–protein interactions results in activity reduction in vivo and in vitro (Otto et al., 2016). AN probably suppresses JA biosynthesis through interaction with AOCs, which might disturb the heteromerization of AOCs, leading to the restricted enzyme activity of AOCs. It will be worth further investigating how AN temporally and spatially regulates AOC enzyme activity in response to pathogen infection.

An increasing body of evidence suggests that the SA signalling pathway antagonizes the JA signalling pathway to regulate plant defense responses (Glazebrook, 2005; Vlot et al., 2009). Due to the mutual inhibitory relationship between the SA and JA signalling pathways, it is thought that plants select either pathway based on the lifestyle of the invading pathogen (Glazebrook, 2005). Biotrophic pathogens activate the SA signalling pathway, while necrotrophic pathogens up-regulate the JA and ET signalling pathways (Pieterse et al., 2012). Xie et al. (2020) reported that AN antagonistically regulates plant resistance to the hemibiotrophic pathogen P. syringae and the necrotrophic pathogen B. cinerea through the transcriptional coregulation of MYB46-mediated SA signalling and WRKY33-mediated JA/ET signalling, respectively. In this study, we also found apparent up-regulation of PR1, a marker gene of the SA signalling pathway and resistance to the hemibiotrophic pathogen P. syringae in the an mutants (Figures S2 and S3g). The pathogen infection assays also demonstrated that AN positively regulates the resistance to infection with necrotrophic pathogens at late stages (Figure S1b–e), which is consistent with Xie’s report that the an mutant was more susceptible to B. cinerea infection at 72 hpi. This infection period was considered as a late infection stage. It is possible that the resistance to necrotrophic pathogens in AN knockout mutants at the early stage of infection, which is also characterized by high JA content upon the pathogen infection stage, and up-regulate JA signalling through WRKY33-mediated transcriptional regulation in the nucleus at the late infection stage. Hence, it is likely that the dual function of AN in regulating defence responses to necrotrophic pathogens through JA signalling depends on the spatial and temporal subcellular localization, which is determined by the function of interacting proteins. Visualizing the spatiotemporal localization dynamics of AN during pathogen infection may help to reveal the different functional mechanisms of AN in the defence response.

In summary, our study provides new insights into the involvement of AN in plant immunity against biotrophic and necrotrophic pathogens via broad modulation of PTI responses and the JA signalling pathway (Figure 7). Further studies are needed to elucidate the precise molecular function of AN in regulating the plant defence response.

4 | EXPERIMENTAL PROCEDURES

4.1 | Plant materials and growth conditions

The T-DNA insertion mutants an-t1 (SALK_026489c) and an-t2 (CS851381) and transgenic plant lines an-t1 COM# and AN-OE were obtained from Dr Deshu Lin (Fujian Agriculture and Forestry University) and were described previously (Dang et al., 2018). The aoc3-1 (SALK_054568) and aoc4-1 (SALK_124897) T-DNA insertion lines in a Col-0 background were obtained from AraShare. The aoc3-1 aoc4-1 double mutant was obtained by genetic crossing. The aoc1cas9-1 and aoc2cas9-1 mutants in a Col-0 background were generated by CRISPR/Cas9 (Castel et al., 2019). The aoc1cas9-1 aoc2cas9-1 and aoc1cas9-2 aoc2cas9-1 double mutants were obtained by genetic crossing of the aoc1cas9-1, aoc1cas9-2, and aoc2cas9-1 single mutants. Arabidopsis plants were grown at 22°C on half-strength Murashige and Skoog (1/2 MS) medium (pH 5.7) with 12% (wt/vol) agar or in commercial potting soil/perlite (3:1) by placing in the dark at 4°C for 3 days, followed by a 12 h light/12 h dark photoperiod with 150 μmol·m⁻²·s⁻¹ light and 65% relative humidity.

Bacterial strains Pst DC3000 and Psm ES4326 were provided by Dr Baomin Feng (Fujian Agriculture and Forestry University). All bacteria were cultured at 28°C in King’s B medium with 50 mg/L rifampicin and kanamycin for Pst DC3000 or 50 mg/L rifampicin for Psm ES4326. The fungi B. cinerea 05.10 and S. sclerotiorum 1980 were obtained from Martin Dickman (Texas A&M University) and cultured at 24°C on potato dextrose agar plates in a light incubator (Zimmerli et al., 2001).

4.2 | Pathogen infection assays

For S. sclerotiorum inoculation, rosette leaves from 4-week-old Arabidopsis plants were detached and placed in Petri dishes containing 0.8% agar, with the petiole embedded in the medium. After inoculating the hyphal plugs on the leaves, the plates were covered with a transparent plastic lid and incubated at 22°C with a 12-h photoperiod.

For B. cinerea infection, spores were diluted to 5 × 10⁵ spores/ml in potato dextrose broth medium. Five microlitres of spore suspension was dropped on the leaf surfaces of 4-week-old plants (three leaves per plant). Disease symptoms and lesion diameters were evaluated for each independent experiment (six plants). For the bacterial infection assay, the leaves of 4-week-old Arabidopsis plants were infiltrated with 2 × 10⁶ cfu/ml Pst DC3000 or 1 × 10⁶ cfu/ml Psm ES4326 in 10 mM MgCl₂. After infiltration, the plants were kept at 100% relative humidity, and symptoms were evaluated 3 days later. Bacterial titres were determined as previously described (Zimmerli et al., 2000). Briefly, the leaves of 4-week-old Arabidopsis plants were infiltrated with 2 × 10⁴ cfu/ml Pst DC3000 and 5 × 10⁴ cfu/ml Psm ES4326 in 10 mM MgCl₂. At the indicated time, the bacterial growth in the leaves was determined by grinding 0.4 cm² leaf discs in 500 μl 10 mM MgCl₂. Serial dilutions were plated on agar plates containing the corresponding antibiotics.
4.3 | Trypan blue staining

Trypan blue staining was performed as described previously (Koch & Slusarenko, 1990). Briefly, fungus-challenged leaves were excised and placed in a six-well microtitre plate. The staining solution (0.05% trypan blue) was added to cover the leaves, and the plate was incubated at room temperature for 8 h or overnight in the dark. Then the staining solution was removed and the leaves were further incubated in chloral hydrate solution (2.5 g/ml) for 24 h at room temperature. After that, the solution was discarded and the leaves were dipped in 70% glycerol. All samples were observed for blue staining using white light and a microscope (S9i; Leica).

4.4 | NBT staining for O$_2^-$ detection

Leaves were detached at 24 hpi and immersed in 10 mM potassium phosphate buffer, pH 7.5, containing 0.1% NBT (Sigma) for 8 h in the dark at room temperature (Sang et al., 2001). After that, the leaves were removed from the NBT solution, washed three times with double distilled water, and immersed in 100% ethanol to destain the tissues for 2 h. Tissues not sufficiently clear were further boiled in 95% ethanol for 10 min in a water bath. Subsequently, the leaves were maintained in 40% glycerol, and microscopic images were captured using a stereomicroscope (S9i; Leica). The dark blue colour indicates the presence of O$_2^-$.

4.5 | DAB staining for H$_2$O$_2$ detection

Leaves were collected at 24 hpi and immersed in 1 mg/ml DAB solution (pH 3.8) (Sigma) for 8 h or overnight in the dark at room temperature. Then the DAB solution was replaced with 100% ethanol and kept for 3 h. Tissues not sufficiently clear were boiled in acetic acid/glycerol/ethanol (1:1:1 vol/vol/vol) for 10 min in a water bath. Subsequently, the leaves were maintained in 50% glycerol and were examined under a microscope. H$_2$O$_2$ was detected as a brown precipitate in the inoculated leaves.

4.6 | ROS burst assay

ROS assays were conducted as previously described (Chinchilla et al., 2007). Briefly, 4-week-old plant leaves were cut into 2-mm$^2$
small leaf discs. Sixteen leaf discs (four discs/plant) were incubated in double distilled water in 96-well plates overnight. The following day, the water was replaced by 200 nM flg22 or 1 mg/ml chitin containing 2 mM luminol (Sigma-Aldrich) and 10 mg/ml peroxidase (Sigma-Aldrich). Luminescence was recorded every 1 min for 1 h using a CentroLIAPc LB 692 plate luminometer.

4.7 | MAPK assay

The activity of MAPKs was detected as described previously (Chung & Sheen, 2017). Briefly, 10-day-old seedlings grown on 1/2 MS agar medium were treated with 200 nM flg22 or 1 mg/ml chitin for 5 and 10 min or with water only as a control. The treated seedlings were rapidly pooled for harvest and frozen in liquid nitrogen. Total protein was extracted by incubating the sample powder in extraction buffer (50 mM Tris-HCl, pH 7.5, 5 mM EDTA, 5 mM EGTA, 2 mM dithiothreitol, 10 mM β-glycerophosphate, 10 mM NaF, 1 mM Na3VO4, 10 mM NaF, 1 mM PMSF, 1x protease inhibitor cocktail) at 4°C for 15 min. MAPK activity was detected by western blot using monoclonal anti-phospho-p44/42 MAPK (Erk1/2) (Th202/Tyr204) from Cell Signaling Technology (dilution 1:5000) according to the manufacturer’s protocol. An anti-actin antibody (Merck, dilution 1:3000) was used to verify equal loading. The experiments were repeated at least three times.

4.8 | Callose deposition

Arabidopsis seedlings were grown on 1/2 MS medium with 12% (wt/vol) agar for 2 weeks at 22°C with 150 µmol-m⁻²-s⁻¹ light in a 12/12 h dark/light photoperiod and 65% relative humidity. The seedlings were transferred into liquid MS medium in six-well plates for 12 h and then treated with 500 nM flg22 or 1 mg/ml chitin for 12 h. Control plants were treated with sterile water only. Seedlings were cleared overnight by shaking in 100% ethanol at room temperature and then washed three times (2 h for each washing step) with 70 mM NaH₂PO₄ (pH 8.0). Cleared seedlings were stained overnight at room temperature with 70 mM NaH₂PO₄ (pH 8.0) solution containing 0.01% aniline blue. Callose deposits were visualized under UV illumination using an SP8 confocal microscope (Leica) as described previously (Singh et al., 2012). Callose deposits were calculated using ImageJ software.

4.9 | Reverse transcription-quantitative PCR

Arabidopsis seedlings grown in 1/2 MS medium with 12% (wt/vol) agar for 10 days were transferred into liquid 1/2 MS medium for one night and then treated with 100 nM flg22 or 1 mg/ml chitin. Samples were collected at the indicated time points. The seedlings were ground into a fine powder using a sterile mortar and pestle in liquid nitrogen. RNA was extracted using TRIzol reagent (Invitrogen) and treated with RNase-free DNase (Solarbio). First-strand cDNA synthesis was carried out using the PrimeScript RT reagent kit with gDNA Eraser according to the manufacturer’s instructions (Takara). AT1G49240 (ACTIN8) was used for normalization. Quantitative PCR analysis was performed using an Applied Biosystems 7500 system with SYBR Green (Takara). The primer sequences used are listed in Table S1.

4.10 | Measurement of SA and JA content

Four-week-old Arabidopsis plants were inoculated with S. sclerotiorum. The samples were collected and ground into a fine powder with a mortar and pestle in liquid nitrogen. Then, 0.1 g of the powder was sent to Jiangxi LiKon Science and Technology Company for SA and JA content analyses as described previously (Sacks et al., 2018).

4.11 | Luciferase complementation assay

An Agrobacterium-mediated transient luciferase expression assay was conducted as described previously (Zhou et al., 2018). Briefly, agrobacteria with the desired split luciferase constructs were cultured in liquid Luria–Bertani (LB) medium containing 50 µg/ml kanamycin at 28°C with constant shaking at 220 rpm overnight. The cells were centrifuged at 5000 × g for 5 min and resuspended with induction medium (10 mM MgCl₂, 200 mM acetosyringone, 10 mM MES, pH 5.6). Subsequently, the bacterial suspensions were adjusted to a final concentration of OD₆₀₀ = 1 and placed in the dark at room temperature for 4 h. Then mixtures of Agrobacterium cultures expressing pNluc and pCluc and its derivatives at a ratio of 1:1 (vol/vol) were infiltrated into the leaves of N. benthamiana by a 1-ml needleless syringe. At 40 to 48 h postinfiltration, the leaves were sprayed with 1 mM d-luciferin and then kept in the dark for 5 min. Images were captured using a CCD imaging system.

4.12 | Bimolecular fluorescence complementation assay

The YFP₅⁺-AOC2 and YFP₇⁻-AN constructs or YFP₅⁺ and YFP₇⁻ empty vectors were co-transformed into N. benthamiana by Agrobacterium-mediated transformation. Forty-eight hours later, YFP fluorescence was detected by confocal microscopy.

4.13 | In vitro pull-down assay

AN and AOC2 were cloned into pET-28 (with a His tag) and pGEX-4T-1 (with a GST tag), respectively. Proteins expressed in Escherichia coli were purified using Ni-IDA Sefinose resin (Sangon Biotech) and glutathione agarose beads (Sangon Biotech) according to the manufacturer’s instructions. For GST pull-down, 10 µg of AN-His was
incubated with GST or GST-AOC2 in 1 ml pull-down buffer (10 mM HEPES, pH 7.5, 100 mM NaCl, 1 mM EDTA, 10% [vol/vol] glycerol, 1% [vol/vol] Triton X-100) for 2 h at 4°C with gentle shaking. Input and pulled down proteins were separated by 10% SDS-PAGE and detected by immunoblotting using appropriate antibodies.

### 4.14 Coimmunoprecipitation assay

A coimmunoprecipitation assay was performed as described previously ([Yoo et al., 2007](#)). Positive transformants of AOC1-HA, AOC2-HA, AOC3-HA, AOC4-HA, and GFP-AN or GFP were lysed in 1 ml of lysis buffer (50 mM Tris-HCl, pH 7.4, 75 mM NaCl, 5 mM EDTA, 10% glycerol, 0.5% Triton X-100, 1x protease inhibitor cocktail, 1x phosphatase inhibitor cocktail) and then centrifuged at 2,400 × g for 10 min. For immunoprecipitation, the protein extracts were incubated with GFP agarose beads at 4°C for 6 h. The immunoprecipitated proteins were washed four times with washing buffer (50 mM Tris-HCl, pH 7.4, 75 mM NaCl, 5 mM EDTA, 10% glycerol, 0.17% Triton X-100) before separation by SDS-PAGE and detection with the respective antibodies (anti-GFP, Abmart; anti-HA, Abmart).

### 4.15 Statistical analysis

All box plots and bar graphs were generated using GraphPad Prism v. 5.0 software. Statistical significance was determined by Student’s t test, one-way analysis of variance, or two-way analysis of variance followed by Tukey post hoc tests using IBM SPSS Statistics.

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### DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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