Melatonin and Its Homologs Induce Immune Responses via Receptors trP47363-trP13076 in *Nicotiana benthamiana*

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Melatonin (N-acetyl-5-methoxytryptamine), a naturally occurring small molecule, can protect plants against abiotic stress after exogenous treatment with it. It is not known if melatonin homologs, such as 5-methoxytryptamine and 5-methoxyindole, that are easy and more cost-effective to synthesize can stimulate the plant immune system in the same manner as melatonin. In the present study, we assessed the biological activity of the melatonin homologs, 5-methoxytryptamin and 5-methoxyindole. The results showed that melatonin and its homologs all induced disease resistance against *Phytophthora nicotianae* in *Nicotiana benthamiana* plants. The application of all three compounds also induced stomatal closure and the production of reactive oxygen species. Gene expression analysis indicated that the expression of genes involved in hydrogen peroxide (H₂O₂), nitric oxide (NO) production, and salicylic acid (SA) biosynthesis was significantly upregulated by all three compounds. Four homologs of the melatonin receptors were identified by blasting search with the phytomelatonin receptor in *Arabidopsis*. Molecular docking studies were also used to identify four putative melatonin receptors in *N. benthamiana*. Further experimentation revealed that silencing of the melatonin receptors trP47363 and trP13076 in *N. benthamiana* compromised the induction of stomatal closure, *PR-1a* gene expression and SA accumulation by all three compounds. Collectively, our data indicate that the induction of defense responses in *N. benthamiana* by melatonin, 5-methoxytryptamine, and 5-methoxyindole involves the melatonin receptors trP47363 and trP13076.

**Keywords:** melatonin, 5-methoxytryptamin, 5-methoxyindole, disease resistance, *Nicotiana benthamiana*, stomatal closure, salicylic acid, transmembrane receptor

**INTRODUCTION**

Plants have evolved two innate immunity systems that respond to a broad range of microorganisms (Zuppini et al., 2004; Zhang et al., 2010; Jian et al., 2021). In the early phase of the plant immunity response, pathogen-associated molecular patterns (PAMPs) and pathogen elicitors can directly induce a defense response in plants (Chen et al., 2015).
Elicitors may be secreted by the microorganism attempting to infect a host plant or substances released by hydrolytic enzymes produced by the pathogen or plant. Elicitors include a variety of compounds secreted by phytopathogens, including proteins, glycol-proteins, synthetic molecules, glycans, and lipids (Zhang et al., 2009). Plant cells respond to the elicitors by an influx in calcium ions, as well as the production of active oxygen species (AOS) and nitric oxide (NO). These induced molecules regulate many processes and interconnect different pathways to amplify and generate a physiological response by the host plant, such as a hypersensitive response and stomatal closure, through the induction of transcriptional and metabolic changes (Garcia-Brugger et al., 2006; Zhang et al., 2010). The various downstream responses that are triggered are not independent but rather overlap, including the induced expression of pathogenesis-related (PR) genes, phytohormone homeostasis regulation, synthesis of reactive oxygen species (ROS), and secondary metabolite accumulation (La Camera et al., 2004; Pieterse et al., 2009; Feng and Shan, 2014).

Melatonin (N-acetyl-5methoxytryptamine), originally discovered in the 1950s in the pineal gland, represents biological molecules that have been extensively studied. Melatonin (MT) has been shown to possess a broad spectrum of biological functions in both animals and plants (Lerner et al., 1958; Arnano and Hernández-Ruiz, 2006; Reiter et al., 2010; Liang et al., 2015; Zhang et al., 2015). For example, MT is considered a indole that can be potentially used to combat viral diseases, such as severe acute respiratory syndrome and West Nile virus (Bonilla et al., 2004), and has been shown to regulate circadian rhythms, the immune system, and ROS signaling, as well as sleep, food intake, mood, and body temperature in humans (Dollins et al., 1994; Rodriguez et al., 2004; Brainard et al., 2011; Carrillo-Vico et al., 2013). MT can also induce host resistance in plants to pathogens, such as Alternaria spp., Pseudomonas syringae pv. tomato DC3000, and Fusarium spp. (Lee et al., 2014; Arnano and Hernández-Ruiz, 2015). MT was first identified in plants in 1995 (Dubbelts et al., 1995; Hattori et al., 1995) and has since received increased attention due to its common presence and versatile functions in the plant kingdom. For example, MT has been reported to alleviate abiotic stresses (cold, heat, drought, salt, and heavy metals) in plants (Shi et al., 2015b; Wei et al., 2015; Li et al., 2016b; Ding et al., 2018). MT in plants has also been reported to function as a growth regulator (Murch and Saxena, 2002; Arnano and Hernández-Ruiz, 2006) regulate seed germination, root development, and provide photoprotection, as well as regulate flowering, leaf senescence, seed yield, and fruit ripening (Wang et al., 2012; Byeon and Back, 2014; Byeon et al., 2014a; Zhang et al., 2014; Sun et al., 2015). The role of MT in plant immunity has been investigated but only limited information is available on the activity of MT-related homologs and the identification of MT receptors in plants.

The biosynthetic pathway of MT in many different organisms, including amphibians, reptiles, and mammals is well established (Reiter, 1999; Falcón et al., 2009) and differs in several respects from MT biosynthesis in plants. In the MT biosynthesis pathway in vertebrates, tryptophan is converted into either 5-hydroxytryptophan by tryptophan 5-hydroxylase or tryptamine by a tryptophan decarboxylase. While the 5-hydroxytryptophan pathway is predominant in animals, the tryptamine pathway is dominant in rice (Oryza sativa; Park et al., 2012). The MT biosynthetic pathway is not regulated in the same manner in plants and animals. The biosynthetic pathway for MT in plants is more complex, relative to vertebrates; however, only limited data are available on MT biosynthesis in plants (Tan et al., 2013). All the enzymes in the MT biosynthetic pathway in rice (O. sativa) have been recently characterized and localized (Byeon et al., 2014b). In contrast to biosynthesis, the production of synthetic MT is a common practice and results in the synthesis of high purity MT, which serves as the primary source of MT for therapeutic use in humans. MT is commercially synthesized in a four-step reaction, using 5-methoxyindole as the main starting material (Cheng-Hong et al., 2009).

In the present study, we investigated the effect of 5-methoxytryptamine and 5-methoxyindole on plant defense response. Results indicated that 5-methoxytryptamine and 5-methoxyindole can induce a resistance response in Nicotiana benthamiana against Phytophthora nicotiana similar to melatonin. The synthetic compounds 5-methoxytryptamine and 5-methoxyindole also induced stomatal closure and ROS production and activated the SA signaling pathway, which has not been investigated in other species. We also identified four putative melatonin receptors, trp47363, trp13076, trp49122, and trp40966 in N. benthamiana through bioinformatics prediction and molecular docking analyses. Finally, we found that the homologs receptors trp47363 and trp13076 act downstream of MT, 5-methoxytryptamine, and 5-methoxyindole in N. benthamiana through VIGS experiments. Collectively, the data suggest that the transmembrane receptors trp47363 and trp13076 function as receptors of MT, 5-methoxytryptamine and 5-methoxyindole to mediate plant defense response in N. benthamiana.

MATERIALS AND METHODS

Fungal and Plant Growth Conditions
The original culture of P. nicotiana was provided by professor Daolong Dou (Nanjing Agricultural University, Jiangsu, China). A 0.6-cm-diameter plug containing P. nicotiana mycelium was placed at the center of V8 plates (200 ml V8 juice and 3.0 g CaCO₃ were boiled in 1 l of water and solidified with 1.5% agar), and the plates were incubated at 25°C for 4 days. N. benthamiana seeds were surface sterilized in 95% (v/v) ethanol for 5 min, followed by a 5% (w/v) solution of sodium hypochlorite for 5 min. This was followed by five washes with double distilled water (ddH₂O). Seeds were allowed to germinate in Petri plates containing solid (Murashige and Skoog 1962) medium. Seedlings were then transferred to pots containing sterilized vermiculite at a density of one per pot. Seedlings were incubated in a controlled environment growth chamber at 25°C under a 16 h/8 h light/dark cycle.
The Effect of MT on Disease Severity in *Nicotiana benthamiana* Caused by *Phytophthora nicotianae*

Chitosan, MT, and its related compounds were dissolved in 99.7% (v/v) ethanol, then, diluted all the compounds to 50 μM using the sterilized water. The effect of 50 μM solutions of either MT, 5-methoxytryptamine, 5-methoxyindole, N-acetyltryptamine, tryptamine, indole, or chitosan of disease severity in *N. benthamiana* caused by *P. nicotianae* was assessed. Six-week-old *N. benthamiana* plants were treated with 50 μM solutions of MT, MT-homologs, and chitosan by watering the soil of seedlings with the prepared solutions. Control seedlings were treated with 0.005% (v/v) ethanol/water. Four hours after the treatments were applied, nine leaves from three plants were detached and inoculated with a 7 mm × 7 mm hyphal plug of *P. nicotianae* placed on the surface of the right side of each leaf, after inoculated samples were placed at 25°C. For disease symptoms were recorded after 48 h of incubation. Leaves were fixed in 95% ethanol, and the disease severity was assessed by measuring the diameter of the *P. nicotianae* lesion. Each assay was repeated three times.

Effect of Treatments on Stomatal Aperture

Stomatal apertures were measured as described by Chen et al. (2004). *N. benthamiana* plants were watered with MT, or its homologs after seedlings had fully opened their stomata in the light for 2–3 h. This was done to minimize the effect of extraneous factors on stomatal response. Plants were then watered with either 50 μM solutions of either MT, 5-methoxytryptamine, 5-methoxyindole, N-acetyltryptamine, tryptamine, indole, or chitosan. Measurements of stomatal apertures were recorded 3 h after the treatments were applied. Images of different stomatal aperture leaf strips were captured with an Olympus BX43 microscope and then measured using CellSens Standard Software (Tokyo, Japan). The diameter of 50 randomly selected stomata, which got from nine leaves and three plants, was measured. Each assay was repeated three times.

3,3’-Diaminobenzidine Staining for Hydrogen Peroxide

Leaves collected 6 h after treatment were incubated in PBS buffer (pH 7.4) containing 0.5% (w/v) diaminobenzidine (DAB) for 8 h at 25°C in the light as described in Samuel et al. (2005). Leaves were then boiled in 96% ethanol for 10 min to remove chlorophyll and any unreacted dye. The remaining brown precipitates results from the reduction of the DAB by hydrogen peroxide were then assessed. The intensity and pattern of H₂O₂ staining in the leaves were analyzed using Quantity One software (Bio-Rad, Milan, Italy).

RNA Isolation and Reverse Transcription-Quantitative PCR

Total RNA was extracted using a Plant RNA Kit (Omega Bio-Tek, Norcross, GA, United States) according to the manufacturer’s instructions. First-strand cDNA was synthesized using Reverse Transcriptase (TaKaRa Bio Inc., Dalian, China) with oligo dT primers, and the resulting cDNA was used as a template for subsequent PCR amplification. RT-PCR products were resolved on an agarose gel to determine qualitative expression levels of target genes. RT-qPCR was performed using SYBR Premix Ex Taq (TaKaRa Bio Inc.) on a 7,500 Fast Real-Time PCR Detection System (Applied Biosystems, Foster, CA United States). Expression of the *N. benthamiana* *EF1a* gene was used to normalize gene expression in each sample.

Virus-Induced Gene Silencing

The first phytomelatonin receptor (CAND2/PMTR1) was recently identified in *Arabidopsis thaliana* and was demonstrated to regulate the stomatal closure induced by MT (Jian et al., 2018). Four membrane proteins (trP47363, trP13076, trP49122, and trP40966) were identified in *N. benthamiana* by BLASTP analysis using *Arabidopsis* CAND2/PMTR1 as a query. Specific, short fragments (~400 bp) of *NbtrP49122-trP40966, NbtrP47363-trP13076*, or *NbtrP47363-trP13076-trP49122-trP40966* were designed using the Sol Genomics VIGS tool.2 These fragments were PCR-amplified with primers introducing 5’end EcoRI and 3’ end XhoI sites and were cloned into the pTRV2 vector between EcoRI and XhoI sites (Li et al., 2002). Cotyledons of two-week-old *N. benthamiana* seedlings were infiltrated with a mixture of *Agrobacterium tumefaciens* GV3101 harboring pTRV1 (OD₆₀₀ = 0.5) and pTRV2: Gene to be silenced (OD₆₀₀ = 0.5) into cotyledons and grown for 4–5 weeks under short-day conditions (22°C, 11 h light/13 h dark). Gene silencing was confirmed by semi-quantitative amplification of the silenced target gene. Total RNA of *N. benthamiana* was extracted from a single leaf disk (8 mm in diameter) using a Plant RNA Kit (Omega Bio-Tek, Norcross, GA, United States). First-strand cDNA was synthesized using Reverse Transcriptase (TaKaRa Bio Inc., Dalian, China) with oligo dT primers, and the resulting cDNA was used as a template for subsequent PCR amplification.

*Agrobacterium*-Mediated Transient Expression Analysis

*Agrobacterium tumefaciens* GV3101 was transformed with binary vector effector constructs by electroporation (2.2 kV/6 ms/ Bio-Rad). Transformed cells were grown on L-agar plates with selective antibiotics for 2 days then inoculated and grown in liquid L-media with selective antibiotics. Overnight cultures were centrifuged at 5,000 rpm for 4 min, resuspended in agro-infiltration solution (10 mM MgCl₂/10 mM MES) and adjusted to an OD₆₀₀ = 0.4 before leaf infiltration with a blunt-end syringe.

Measurement of Free SA Accumulation

Free SA was extracted from leaves and roots and quantified according to the method of Schuhegger et al. (2006).
Statistical Analysis
Each experiment was conducted with a minimum of three independent replications. The data were analyzed by ANOVA, followed by Duncan's multiple range test \((p < 0.01)\), using SPSS software (SPSS Inc., Chicago, IL, United States).

RESULTS

Effect of Melatonin and Its Homologs on Disease Severity in Nicotiana benthamiana Caused by Phytophthora nicotianae

MT plays an important role in plant protection against biotic stresses (Regodon et al., 2005; Carrillo-Vico et al., 2013; Vielma et al., 2014; Nabavi et al., 2019); however, it is expensive to produce, limiting its potential use in agriculture. Therefore, the impact of several MT-homologs on disease resistance in N. benthamiana against P. nicotianae was investigated. Water was used as a negative control and chitosan was used as a positive control. The experiment was designed to determine if 50 \(\mu\)M solutions of the treatment compounds could induce defense responses against P. nicotianae in N. benthamiana. Results indicated that the pretreatment of N. benthamiana plants with MT, 5-methoxytryptamine, and 5-methoxyindole significantly inhibited \((p < 0.01)\) lesion size in inoculated leaves of N. benthamiana by 16.56–23.31\%. Chitosan, used as a positive control, reduced lesion size by 34.36\%, relative to the untreated control group. The other MT-homologs, N-acetyltryptamine, tryptamine, and indole, did not affect lesion size, relative to the untreated control (Figures 1A,B). Since MT, 5-methoxytryptamine, and 5-methoxyindole are based on a 5-methoxyindole backbone, while N-acetyltryptamine, tryptamine, and indole are based on indole, we postulated that 5-methoxyindole represents the functional backbone of MT.

Melatonin, 5-methoxytryptamin, and 5-methoxyindole Induce Stomatal Closure in Nicotiana benthamiana Leaves

Guard cells exhibit a classic innate immune response to both PAMP compounds and pathogens (Zhang et al., 2009). Elicitors, such as chitosan, have been reported to induce stomatal closure (Lee et al., 1999). Therefore, we hypothesized that MT and MT-homologs would also induce stomatal closure. To test this hypothesis, the aperture of guard cells in epidermal peels of N. benthamiana was examined and measured under a microscope after being treated with a buffer, or 50 \(\mu\)M solutions of MT or MT-homologs compared with positive control treated with Chitosan. Results indicated that 3 h after the treatment MT, 5-methoxytryptamin, and 5-methoxyindole induced stomatal closure, while N-acetyltryptamine, tryptamine, and indole had no effect (Figures 2A,B).

Melatonin and Its Homologs Induce the \(\text{H}_2\text{O}_2\) Accumulation in Nicotiana benthamiana Leaves

Several studies have demonstrated that \(\text{H}_2\text{O}_2\) is involved in elicitor-inhibited stomatal opening and elicitor-enhanced stomatal closure (Lee et al., 1999). Therefore, \(\text{H}_2\text{O}_2\) accumulation was measured in N. benthamiana leaves treated with MT, 5-methoxytryptamine, or 5-methoxyindole to determine if \(\text{H}_2\text{O}_2\) was involved in the effect of these chemical compounds on stomatal closure. \(\text{H}_2\text{O}_2\) accumulation was measured by DAB staining. Figure 3A is representative photographs showing the presence of DAB–\(\text{H}_2\text{O}_2\) reaction product in N. benthamiana leaves at 8 h after treatment. Heavy brown precipitates, indicating \(\text{H}_2\text{O}_2\) accumulation, were observed in leaves of plants treated with MT and MT-homologs, relative to the untreated control (Figure 3A). These observations were consistent with the measurement of relative DAB staining presented in Figure 3B.

Effect of Melatonin, 5-methoxytryptamin, and 5-methoxyindole to regulate on the Expression of Genes Involved in \(\text{H}_2\text{O}_2\) and NO Production

Nitric oxide regulates innate immunity by functioning as a signaling molecule in a wide range of organisms including plants, especially in the regulation of stomatal guard cells (Ali et al., 2007; Asai and Yoshioka, 2009; Zhang et al., 2010). Rapid production of AOS, including \(\text{H}_2\text{O}_2\), superoxide, and the highly reactive hydroxyl radical, is a characteristic component of the resistance response in plants, where AOS is believed to function as second messengers in the activation of resistance reactions, such as defense-related genes (Lee et al., 1999; Pasqualini et al., 2007). We hypothesized that the induction of disease resistance by MT, 5-methoxytryptamine, and 5-methoxyindole in plants is correlated with the upregulation of genes involved in \(\text{H}_2\text{O}_2\) and NO production. Our results indicated that two genes involved in \(\text{H}_2\text{O}_2\) production, \(\text{rohbA}\) and \(\text{rohbB}\), were both significantly upregulated by MT, 5-methoxytryptamine, and 5-methoxyindole (Figures 4A,B). The expression of nitrate reductase genes, \(\text{NIA1}\) and \(\text{NIA2}\), was also affected by the three tested compounds (Figures 4C,D). \(\text{NIA1}\) was upregulated by all three compounds, while the expression of \(\text{NIA2}\) was differentially impacted by the three compounds, being slightly induced by MT, more upregulated by 5-methoxyindole, and downregulated by 5-methoxytryptamine.

Melatonin and Its Homologs Activate an SA-Mediated Signaling Pathway

Plants utilize a broad range of defense mechanisms to respond and prevent invasion by pathogens. Systemic acquired resistance is associated with the accumulation of the hormone, salicylic acid (SA), while induced systemic resistance is mediated by jasmonic acid (JA) and ethylene (ET)-regulated pathways (Schuhegger et al., 2006; Zhu et al., 2014; Jian et al., 2019). To determine which signaling pathway is involved in the resistance of N. benthamiana to pathogens induced by MT and its homologs, the expression of the SA-responsive gene \(\text{PR-1a}\), the JA synthesis-related gene \(\text{LOX}\), and the ET-responsive gene \(\text{ERF1}\) in response to
the application of these compounds. Results revealed that the expression of *PR-1a* is dramatically increased in *N. benthamiana* leaves 3 h after the application of MT, 5-methoxytryptamine, or 5-methoxyindole and that there

**FIGURE 1** | Effect of melatonin and its homologs on disease severity in *N. benthamiana* caused by *P. nicotianae*. **(A)** Disease symptoms in *N. benthamiana* leaves following pretreatment with 50 μM solutions of MT and MT-homologs and inoculate with *P. nicotianae*. Fully expanded *N. benthamiana* leaves were detached from plants 4 h after being watered with the test compounds. The leaves were placed in Petri dishes containing water-saturated filter paper and then inoculated with *P. nicotianae*. Photographs of the lesions were taken at 48 h post-inoculation. Water (control) was used as a negative control and chitosan was used as positive control. Ethanol-bleached *N. benthamiana* leaves were also photographed to more clearly illustrate the region of dead cells. **(B)** Average lesion diameter (red circles) in *N. benthamiana* leaves treated with different test compounds. Data represent the mean ± standard deviations (SDs) of three replicates (experiments). Columns with different letters indicate significant differences in lesion size as determined by a Duncan’s multiple range test.
was increased accumulation of free SA (**Figures 5A,D**), while *ERF1* was repressed (**Figure 5B**). Notably, the expression of *LOX* was not induced by any of the three tested compounds (**Figure 5C**).
Bioinformatics Prediction of Four Membrane Receptors Potentially Involved in Binding Melatonin in *Nicotiana benthamiana*

MT was identified in plants as early as 1995; however, the function and signaling pathway of this putative phytohormone are largely undetermined due to the lack of identification of its receptor. The first phytomelatonin receptor (CAND2/PMTR1), however, was recently identified in *A. thaliana* and was demonstrated to regulate the stomatal closure induced by MT (Jian et al., 2018). In the current study, four membrane receptors were identified in *N. benthamiana* by BLASTP analysis (see footnote 1) using *Arabidopsis* CAND2/PMTR1 as a query. The amino acid sequence homology between the MT receptor...
in *Arabidopsis* (CAND2/PMTR1) and the four predicted receptors in *N. benthamiana* (trP47363, trP13076, trP49122, and trP40966) was high (Figure 6A). Sequence analysis indicated that the highest homology was between *Arabidopsis* CAND2/PMTR1 and *N. benthamiana* trP49122 and trP40966, with approximately 62% identity at the amino acid level. The sequence similarity between CAND2/PMTR1 and the *N. benthamiana* proteins trP47363, trP13076 was approximately 51%. A phylogenetic analysis of the MT receptor from *Arabidopsis* (CAND2/PMTR1) and the four transmembrane proteins revealed two distinct evolutionary branches (Figure 6B). Notably, however, the MT receptor (CAND2) in *Arabidopsis* has a shorter branch length distance from the membrane proteins trP49122 and trP40966 in *N. benthamiana* than from the proteins trP47363 and trP13076. A comparison of the predicted transmembrane structure of CAND2 in *Arabidopsis* with the putative MT receptors in *N. benthamiana* indicated that the structure of CAND2 was more similar to trP49122, trP40966, and trP47363 than to trP13076 (Figure 6C).

**In silico** Molecular Docking Analysis of the *Nicotiana benthamiana* Membrane Receptors and Melatonin and 5-methoxyindole

To further evaluate the molecular interaction between the putative transmembrane receptors identified in *N. benthamiana* and MT and 5-methoxytryptamine and 5-methoxyindole, a molecular docking study was conducted. Molecular docking
analysis was conducted in silico using the surflex-dock program within SYBYL-X 2.0 software. The protein structure of the evaluated proteins was obtained from PDB. The scores obtained from the analysis indicate that MT can bind more efficiently to the receptor proteins than 5-methoxyindole, which lacks an amide bond. Since the structure of 5-methoxyindole is smaller than MT, the former molecule can enter the binding pocket formed by the receptor protein more flexibly and all of the crash scores for 5-methoxyindole are closer than of the crash scores of MT when comparisons are made for docking with the same receptor protein. The putative receptor proteins trP49122 and trP13076 do not exhibit an effective binding pocket, suggesting that even small molecules may not be able to interact with these proteins. CAND2 was confirmed to be a putative receptor protein for MT in Arabidopsis (Jian et al., 2018). The docking analysis resulted in a crash score of 3.61 for CAND2, which is considered to represent a good receptor protein for MT. As illustrated in Figure 7A, CAND2 can form an active bowl-like pocket and the oxygen atom at the amide group of ASN100 has one H-bonding interaction with the active hydrogen at the amide bond of MT (–NHCO– ------ H2NOC-ASN100).

**Figure 7A** also illustrates that trP47363 most likely represents the best receptor protein for MT in *N. benthamiana*, with a

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**FIGURE 5** | Melatonin and MT-homologs activate an SA-mediated signaling pathway. The effect of MT, 5-methoxytryptamine, and 5-methoxyindole on the expression of SA-, ethylene-, and JA-dependent marker genes in *N. benthamiana* leaves. *N. benthamiana* leaves were treated with 0.005% (v/v) ethanol/water (control) or 50 μM solutions of MT, 5-methoxytryptamine, or 5-methoxyindole. The relative expression of PR-1a (A), ERF1 (B), and LOX (C) by RT-qPCR. Expression values were normalized to the levels of EF-1a. Data are the mean ± standard deviation (SDs; n = 9). Columns with different letters indicate a significant difference (p < 0.01) between the treatment groups as determined by a Duncan's multiple range test. (D) Level of free SA in *N. benthamiana* leaves after treatment with MT, 5-methoxytryptamine, or 5-methoxyindole. Columns with different letters indicate a significant difference (p < 0.01) between the treatment groups as determined by a Duncan's multiple range test. FW, fresh weight.
C score of 4.61 for docking with MT, which is the highest score indicated among the five receptor proteins. The binding of trP47363 with MT is illustrated in Figure 7B. Visual inspection indicates that MT is tightly embedded in the hole-shaped active binding pocket (Figure 7B), indicating that MT has a higher probability of interacting with trP47363 than with the other receptor proteins. The oxygen atom of the methoxy at the fifth position of the indole ring in MT provides one H-bonding interaction with the hydrogen atom of the hydroxyl group in TYR253 (-OCH ------HO-TYR253).

Melatonin, 5-methoxytryptamin, and 5-methoxyindole Induce Stomatal Closure in *Nicotiana benthamiana* Leaves Involve the Transmembrane Receptors trP47363 and trP13076

The bioinformatics and molecular docking analyses provide fundamental information on MT and MT-homologs induced resistance in *N. benthamiana* to *P. nicotianae*. Further molecular and genetic evidence is needed, however, to support the role of the transmembrane receptors in MT and its homologs induced disease resistance. In this regard, knocking down the expression of key genes involved in plant immunity through virus-induced gene silencing (VIGS) has been widely used to evaluate the involvement of various defense components in plant immunity (Liu et al., 2002; Peart et al., 2002; Oh et al., 2014). We utilized VIGS to determine the need for the identified transmembrane receptors in MT, 5-methoxytryptamin, and 5-methoxyindole induced disease resistance in *N. benthamiana*. *N. benthamiana* plants were generated that were silenced for ΔNbtrP49122-ΔNbtrP40966 (pTRV2:NbtrP49122-trP40966), ΔNbtrP47363-ΔNbtrP13076 (pTRV2:NbtrP47363-trP13076), and ΔNbtrP47363-trP13076-trP49122-trP40966 (pTRV2:NbtrP47363-trP13076-trP49122-trP40966) and were utilized together with control plants (pTRV2:EV). To first validate our experimental conditions, stomatal closure induced by MT, 5-methoxytryptamin, and 5-methoxyindole was evaluated in control and the VIGS mutants of *N. benthamiana*. Results indicated that MT, 5-methoxytryptamin, and 5-methoxyindole induced stomatal closure in EV- and
ΔNbtrP49122-trP40966-plants but not in ΔNbtrP47363-trP13076- or in ΔNbtrP47363-trP13076-trP49122-trP40966-silenced N. benthamiana plants (Figures 8A,B). Collectively, the results indicate that the transmembrane receptors trP47363 and trP13076 are key components in the regulation of stomatal closure in N. benthamiana leaves by MT, 5-methoxytryptamin, and 5-methoxyindole.

The Transmembrane Receptors trP47363 and trP13076 Play a Key Role in Melatonin, 5-methoxytryptamin, and 5-methoxyindole Activation of the SA Signaling Pathway

Our results indicated that MT, 5-methoxytryptamin, and 5-methoxyindole can significantly upregulate PR-1a gene expression in wild N. benthamiana plants and the mutant line ΔNbtrP49122-trP40966 (Figures 9A,B). Therefore, to determine which transmembrane receptor is involved in the expression of the SA marker gene, PR-1a, induced by MT, 5-methoxytryptamin, and 5-methoxyindole, the expression of PR-1a in different receptor mutants created by VIGS was assayed. Interestingly, MT- and 5-methoxytryptamine-induced PR-1a expression was significantly lower than in the control in ΔNbtrP447363-trP13076 and ΔNbtrP47363-trP13076-trP49122-trP40966 (Figures 9C,D). Additionally, 5-methoxyindole-induced expression of PR-1a was only slightly higher than MT-induced expression, or unchanged, in ΔNbtrP447363-trP13076 and ΔNbtrP47363-trP13076-trP49122-trP40966 (Figures 9C,D). Collectively, the data indicate that MT, 5-methoxytryptamin, and 5-methoxyindole activate the SA signaling pathway through the receptors trP47363 and trP13076.
Melatonin, 5-methoxytryptamin, and 5-methoxyindole induce stomatal closure in *N. benthamiana* leaves involve the transmembrane receptors trP47363 and trP13076. MT-, 5-methoxytryptamin-, and 5-methoxyindole-induced stomatal closure in leaves of VIGS mutants of *N. benthamiana* revealing the role of the transmembrane receptors, trP47363 and trP13076. Two-week-old *N. benthamiana* plants were infiltrated with a mixture of *A. tumefaciens* Gv3101 strains carrying pTRV1 or pTRV2 constructs as described in the Materials and Methods. Five weeks later, plants were watered with 0.005% (v/v) ethanol/water (control) or 50 μM solutions of MT, 5-methoxytryptamine, or 5-methoxyindole. Stomata were observed under a microscope 3 h after the application of the treatment solutions. (A) Representative low- and high-magnification images. (B) Measurements of stomatal aperture. Data are the mean ± standard deviation (SD; n = 150). Columns with different letters indicate significant differences between the different treatment groups in the different VIGS lines as determined by a Duncan’s multiple range test.
The Transmembrane Receptors trP47363 and trP13076 Are Involved in Melatonin-, 5-methoxytryptamin-, and 5-methoxyindole-Induced Accumulation of SA

To determine if SA levels are altered in leaves of VIGS mutants of *N. benthamiana*, SA levels were measured in the different receptor mutants that were treated with MT, 5-methoxytryptamine, and 5-methoxyindole. Results indicated all three compounds significantly induced SA accumulation in wild-type plants and the ΔNbrtrP49122-trP40966 mutant (Figures 10A,B) but not in ΔNbrtrP47363-trP13076 or ΔNbrtrP47363-trP13076-trP49122-trP40966 mutants (Figures 10C,D). These results suggest that the transmembrane receptors trP47363 and trP13076 recognize MT, 5-methoxytryptamine, and 5-methoxyindole which are involved in the regulation of SA accumulation.
DISCUSSION

The role of melatonin (MT) in a variety of biological processes has been extensively investigated in animal systems and demonstrated to play a role in the regulation of circadian rhythms, the immune system, induction of ROS, sleep, food intake, mood, and body temperature (Dollins et al., 1994; Rodriguez et al., 2004; Brainard et al., 2011; Carrillo-Vico et al., 2013). In plants, MT has been reported to play a role in the regulation of seed germination, root development, photoprotection, flowering, leaf senescence, seed yield, and fruit ripening (Wang et al., 2012; Byeon and Back, 2014; Byeon et al., 2014a; Zhang et al., 2014; Sun et al., 2015). Studies have also implicated MT in plant immunity (Yin et al., 2013; Lee et al., 2015; Shi et al., 2015c,d; Mandal et al., 2018). While a variety of MT-homologs exist, which are more cost-effective to produce, it is not known if these MT-homologs, such as 5-methoxytryptamin and 5-methoxyindole can induce a defense response in plants. In the present study, we demonstrated that both 5-methoxytryptamin and
5-methoxyindole can induce the same defense response as MT in leaves of *N. benthamiana*. The results also indicate that the MT-homolog-induced defense response occurs through the activation of SA signaling. MT, 5-methoxytryptamine, and 5-methoxyindole induced disease resistance against *P. nicotianae* in leaves of *N. benthamiana* plants, and also induced stomatal closure through the activation of H$_2$O$_2$ production and SA accumulation. Based on these results, we suggest that both 5-methoxytryptamine and 5-methoxyindole can serve as cost-effective alternatives to the use of MT to induce a disease resistance response in plants.

NO and H$_2$O$_2$ as important components of signal transduction pathways, regulate a large number of physiological functions in biological systems. Stomatal closure occurs in response to physiological and stress stimuli (Bright et al., 2006; Garcia-Brugger et al., 2006), and numerous studies have shown that various elicitors can induce reactive oxygen bursts to limit the expansion of invasive pathogens (Zhang et al., 2009, 2010). MT has also been shown to induce increases in H$_2$O$_2$ production in *Arabidopsis* (Jian et al., 2018). In the present study, we found that H$_2$O$_2$, one of the most well-characterized plants signaling molecules, is involved in the MT, 5-methoxytryptamine, and 5-methoxyindole induction of systemic resistance in *N. benthamiana* against *Phytophthora parasitica var. nicotianae* (Figure 1). Our study also demonstrated that guard cells respond to MT, 5-methoxytryptamine, and 5-methoxyindole generate H$_2$O$_2$ and undergo a narrowing of their stomatal apertures (Figure 2). We also confirmed that MT, 5-methoxytryptamine, and 5-methoxyindole can stimulate the accumulation of both H$_2$O$_2$ and SA in *N. benthamiana* leaves (Figures 3, 5). These observations suggest that MT, 5-methoxytryptamine, and 5-methoxyindole could be considered as potential candidates for a new group of general elicitors of plant defense.

Disease resistance in plants is a highly regulated phenomenon that depends on several hormone-regulated signaling pathways, including salicylic acid (SA), jasmonic acid (JA), and ethylene (ET), each of which is activated by a different set of biotic and abiotic stimuli (Schu hegger et al., 2006). Pretreatment of plants with different elicitors can render a plant more resistant to subsequent pathogen attack, and elicitors have been demonstrated to induce various plant defense responses, such as the hypersensitive response, pathogenesis-related (PR) gene expression, and cell wall stabilization (Lee et al., 1999). Peanut (*Arachis hypogaea*) leaves treated with chitosan, a well-documented elicitor, exhibit an increase in endogenous SA levels, as well as an increase in intercellular chitinase and glucanase activity (Sathiyabama and Balasubramanian, 1998). In the present study, SA, a well-characterized signaling molecule in plants, was shown to be involved in the systemic resistance induced by MT, 5-methoxytryptamine, and 5-methoxyindole, as evidenced by the upregulation of PR-1a and SA accumulation, both of which are considered as markers of the SA-dependent defense pathway (SAR; Yoshioka et al., 2009; Figure 5).

MT functions as a neurohormone in vertebrates. Four families of MT receptors have been identified in animals (Reppert et al., 1994, 1995; Nosjean et al., 2000; Levoye et al., 2006). Phytomelatonin was first identified in plants in 1995 (Dubbels et al., 1995; Hattori et al., 1995; Tassel et al., 1995). Relative to its role in vertebrates, the designation of phytomelatonin as a hormone in plants remains controversial partially due to the lack of identification of an MT receptor or a receptor-mediated signaling pathway, as well as the function of such a pathway. The first phytomelatonin receptor (CAND2/PMTR1) was recently identified in *A. thaliana* and shown to mediate MT-induced stomatal closure (Jian et al., 2018). In our present study, CAND2/PMTR1 was used as a query in a BLASTP analysis to identify four putative transmembrane receptors (trP47363, trP13076, trP49122, and trP40966) in *N. benthamiana*. The resulting putative receptors identified in *N. benthamiana* exhibited a high amino acid sequence homology with the CAND2/PMTR1 of MT receptor in *A. thaliana* (Figure 6A). A phylogenetic tree and transmembrane structure prediction between the MT receptor in *A. thaliana* (CAND2/PMTR1) and the four transmembrane proteins in *N. benthamiana* were also similar (Figures 6B, C).

An *in silico* molecular docking analysis was also conducted to further evaluate the molecular interaction between the transmembrane receptors and MT, 5-methoxytryptamine, and 5-methoxyindole in *N. benthamiana*. As shown in Figure 7, trP47363 appears to be the best MT receptor protein in *N. benthamiana* based on its C score (4.61) for docking with melatonin, which was the highest among the five analyzed proteins and revealed that MT and MT-homologs could be tightly embedded into a hole-shaped active binding pocket. We speculate that trP47363 represents a putative transmembrane protein that is most likely to recognize MT, 5-methoxytryptamine, and 5-methoxyindole to trigger the induction of an induced immunity response in *N. benthamiana*. We then measured the expression of the plant defense gene, PR-1a, and SA accumulation and their association with stomatal closure in leaves VIGS mutants of *N. benthamiana* treated with MT, 5-methoxytryptamine, or 5-methoxyindole. The VIGS mutants were constructed to silence the expression of the four transmembrane receptors. The collective results indicated that the putative transmembrane receptors trP47363 and trP13076 appear to be involved in the induced immunity response in *N. benthamiana*. This includes stomatal closure, PR-1a expression and SA accumulation induced by MT, 5-methoxytryptamine, and 5-methoxyindole (Figures 8–10). Based on these results, it appears that the transmembrane receptor trP47363 plays an important role in recognizing MT, 5-methoxytryptamine, and 5-methoxyindole, which then induces a plant defense response by activating the SA-mediated signaling pathway.

In conclusion, we evaluated the ability of MT and MT-homologs to induce a resistance response in *N. benthamiana* and the underlying mechanisms associated with the resistance response. We demonstrated that 5-methoxyindole represents the functional backbone in MT and can induce the same response in *N. benthamiana* as MT. We also identified putative MT receptors in *N. benthamiana* and demonstrated that MT, 5-methoxytryptamine, and 5-methoxyindole are recognized by the same putative receptors. Our study provides direct evidence that
5-methoxytryptamine and 5-methoxyindole can induce resistance in *N. benthamiana*, in the same manner as MT. The three compounds trigger stomatal closure, induce the production of ROS, as well as SA accumulation. Notably, the membrane receptors trP47363 and trP13076 recognize MT, 5-methoxytryptamine, and 5-methoxyindole in *N. benthamiana*. We provide evidence that 5-methoxytryptamine and 5-Methoxyindole can be substituted for MT as an inducer of plant resistance. Thus, 5-methoxytryptamine and 5-Methoxyindole have excellent commercial potential for their use in agricultural production due to their ability to induce a resistance response, and the low cost of production relative to synthetic MT.

**DATA AVAILABILITY STATEMENT**

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

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**AUTHOR CONTRIBUTIONS**

XG, JC, and JLiu conceived and designed the study and revised the manuscript. MK designed the study, analyzed the data, and wrote and revised the manuscript. TS and JLia performed the experiments, processed the data, and provided the suggestions. QA, QG, and HW revised the manuscript. All authors discussed the results and commented on the manuscript.

**FUNDING**

This work was supported by the National Key R&D Project (2018YFE0206000), the Key Project of NSFC Regional Innovation and Development Joint Fund (U20A2039), and the Joint Foundation of Scientific Research Think Tank of Biological Manufacturing Industry in Qingdao (QDSWZK201902).
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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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