UvKmt6-mediated H3K27 trimethylation is required for development, pathogenicity, and stress response in *Ustilaginoidea virens*

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**ABSTRACT**

Polycomb repressive complex 2 (PRC2) is responsible for the trimethylation of lysine 27 of histone H3 (H3K27me3)-mediated transcriptional silencing. At present, its biological roles in the devastating rice pathogenic fungus *Ustilaginoidea virens* remain unclear. In this study, we analyzed the function of a putative PRC2 catalytic subunit UvKmt6. The results showed that disruption of UvKmt6 resulted in reduced growth, conidiation and pathogenicity in *U. virens*. Furthermore, UvKmt6 is essential for establishment of H3K27me3 modification, which covers 321 genes in the genome. Deletion of UvKmt6 led to transcriptional derepression of 629 genes, 140 of which were occupied with H3K27me3 modification. Consistent with RNA-seq and ChIP-seq analysis, UvKmt6 was further confirmed to participate in the transcriptional repression of genes encoding effectors and genes associated with secondary metabolites production, such as PKSs, NRPSs and *Cytochrome P450s*. Notably, we found that UvKmt6 is involved in transcriptional repression of oxidative, osmotic, cell wall and nutrient starvation stresses response-related genes. From the perspective of gene expression and phenotype, in addition to the relatively conservative role in fungal development, virulence and production of secondary metabolites, we further reported that UvKmt6-mediated H3K27me3 plays a critical role in the response to various stresses in *U. virens*.

**KEYWORDS**

*Ustilaginoidea virens*; virulence; PRC2; Kmt6; epigenetic regulation

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**INTRODUCTION**

Epigenetic regulation is dependent on modifications of genomic DNA or histone, but not on the changes of DNA sequences [1]. It is known that the lysine methylation of histone H3 plays an important role in epigenetic regulation. In general, di-/tri- methylation of H3 lysine 4 and H3 lysine 36 are responsible for transcriptional activation, while di-methylation of H3 lysine 9 and tri-methylation of H3 lysine 27 (H3K27me3) are involved in transcriptional silencing [2]. Among these modifications, H3K27me3 is conserved from fungi to higher organisms such as *Arabidopsis*, *Drosophila* and mammals [3]. The H3K27me3 modification was first found in *Drosophila* and mediated by polycomb repressive complex 2 (PRC2) [4]. The PRC2 complex consists of core subunits Ezh (Kmt6), Su(z)12, Esc (Eed) and additional subunits RbAp48/Nurfl55 (P55), among which Ezh is a key catalytic subunit of PRC2 [5]. Subsequent studies have shown that Ezh (Kmt6)-mediated H3K27me3 modification is involved in the development and differentiation of various organisms. For example, Ezh2, homologue of Ezh, is up-regulated in many cancer cells and associated with tumorigenesis of breast, pancreatic, colorectal and endometrial cancer [6,7]. In addition, Ezh2 is essential for maintaining intestine integrity and caudal fin regeneration in zebrafish [8] and early gastrulation development in mouse [9]. In model plant *Arabidopsis*, accumulation of H3K27me3 modification on *Flowering Locus C* (FLC) regulates developmental transition induced by vernalization [10].

H3K27me3 modification on chromatin plays distinct roles in regulation of differentiation and development in filamentous fungi. In *Neurospora crassa*, H3K27me3 modification mediated by PRC2 covers 6.8% of genome. Deletion of *SET7* (Ezh/KMT6 homologues) caused loss of H3K27me3, but did not result in defects of growth and sexual development in *N. crassa* [3].
Similarly, absence of H3K27me3 modification does not lead to obvious phenotypic changes and gene transcriptional activation in pathogenic fungus Zymoseptoria tritici, although it reduces the loss of accessory chromosomes [11]. It was further found that H3K27me3 is involved in spatiotemporal expression regulation of virulence-related effectors in Z. tritici [12]. In contrast, in the wheat head blight fungus Fusarium graminearum, growth and the production of various metabolites are regulated by Kmt6-mediated repression [13]. Knockdown of KMT6 also induced derepression of secondary metabolite genes in clusters in the rice bakanae pathogen Fusarium fujikuroi, which led to increased synthesis of secondary metabolites, such as beauvericin [14]. Furthermore, Kmt6-mediated H3K27me3 modification is required for fungal growth and development in F. fujikuroi [14]. In addition to plant pathogens, in a beneficial endophyte Epichloë festucae, H3K27me3 plays an important role in building symbiotic interaction with Lolium perenne by regulating specific expression of alkaloid biosynthesis-related genes [15]. In the rice blast fungus Magnaporthe oryzae, MoKmt6 catalyzes H3K27me3 and is required for growth, conidiation, pathogenicity and orchestration of effectors transcription [16,17]. A recent report in the fungus Podospora anserina revealed that loss of Kmt6 also resulted in severe defects in growth, and differentiation by causing genome-wide loss of H3K27me3 modification [18]. Therefore, PRC2-mediated H3K27me3 modification in fungi is conservative, but has multiple functions, which is worthy of further study.

Ustilaginoidea virens, the causal agent of the rice false smut disease, is an economically important fungal pathogen posing threats to rice production [19,20]. The infection of U. virens not only reduces the quality and yield of rice, but also contaminates the rice seeds and straw with toxins, such as ustiloxins and ustilaginoïdins [21–23]. Recently, a great progress has been made in depicting infection process of U. virens. Briefly, at the booting stage, the U. virens conidia germinate and enter into the rice spikelet through the space between lemma and palea to infect stamens [24]. Subsequently, invasive hyphae spread into stigmas and fill the floral organs, finally develop into false smut balls, which are typical and visible symptom of the rice false smut disease [25]. In order to complete the infection, U. virens needs to adapt to inevitable stresses from host and environment in variety of ways, such as secreting a lot of effectors to modulate rice immunity [26–29] or regulations of stress response-related pathways [30–33]. Although a great progress has been made in the research of rice false smut fungus, the infection mechanism of U. virens is still largely unclear. Revealing its pathogenic mechanism is of great significance for formulating effective control strategies.

In this study, we set out to investigate the biological roles of H3K27me3 modification in U. virens. By disruption of UvKMT6, we found that it is essential for H3K27me3 modification and required for growth, consolidation and virulence. Furthermore, in combination with ChIP-seq, RNA-seq, and phenotypic analysis, we revealed that H3K27me3-mediated transcriptional repression kept tight correlation with effector transcription, secondary metabolism, and stress responses in U. virens.

Results

Identification of UvKMT6 in U. virens

By BLASTp with F. graminearum Kmt6 (XP_011316633.1) protein sequence as a query, Kmt6 homologues including U. virens Kmt6 (KDB18066.1), Metarhizium robertsii Kmt6 (XP_007816866.1), M. oryzae Kmt6 (XP_003718975), N. crassa Set7 (XP_965043.2), A. thaliana Mea (NP_563658.1), Mus musculus Ezh1 (XP_006532241.1) and Caenorhabditis elegans Mes-4 (NP_506333.1) were identified. All of these Kmt6 homologues contain a SET domain predicted by the SMART tool (http://smart.embl-heidelberg.de) (Figure 1a). Sequence alignment of eight Kmt6 homologues showed that Kmt6 homologues displayed high sequence similarity, ranging from 47% (M. musculus) to 75% (M. robertsii) (Figure S1). Furthermore, a phylogenetic tree of Kmt6 homologues constructed by the MEGA 7.0 software showed that UvKmt6 was closely related to M. robertsii Kmt6 (Figure 1b). These results indicated that Kmt6 is highly conserved in different fungi, and may play an important role in U. virens as other fungi (Figure 1b).

Next, expression level of UvKMT6 at different infection stages including 3, 5, 6, 7 and 9 dpi (days post inoculation) was monitored by qRT-PCR (quantitative Real-time PCR). The results showed that the expression level of UvKMT6 during the infection process increased more than two folds comparing with that of mycelial stage except 7 dpi (Figure 1c), implicating a possible role of UvKMT6 during infection stage in U. virens.

Targeted gene deletion and complementation of UvKMT6

To investigate biological roles of UvKMT6 in U. virens, UvKMT6 deletion mutants were constructed based on the principle of homologous recombination (Figure 2a). Then, Southern blotting and qRT-PCR assays were used to verify that UvKMT6 gene was replaced by the
**UvKMT6 is required for growth, conidiation, and virulence in U. virens**

Because the colonies of ΔUvkmt6 mutants appeared to be smaller than that of the WT strain, hyphal growth was evaluated by inoculating mycelial plugs...
of the WT, ΔUvkmt6-16, –18 and ΔUvkmt6-C strains on the PSA (potato sucrose agar medium) plates. As the Figure 3a and Figure S2 displayed, the colony diameter of ΔUvkmt6-16 and –18 were slightly reduced than WT during the cultured process, indicating that UvKmt6 facilitates hyphal growth. This role was further confirmed by measuring mycelial weight in liquid cultures. When cultured in liquid PS (potato sucrose medium) culture for 7 d (days), mycelial dry weights of the ΔUvkmt6-16 and –18 strains were also less than those of WT (Figure 3c).

In contrast, the vegetative growth was rescued in the ΔUvkmt6-C strain (Figure 3), suggesting that UvKmt6 plays important roles in mycelial growth in U. virens.

Kmt6 homologues were reported to be involved in asexual development in M. oryzae and F. graminearum [13,16], but not in N. crassa [3]. To determine whether UvKmt6 was required for conidiation in U. virens, the same amount of mycelial plugs of the WT, ΔUvkmt6 and ΔUvkmt6-C strains were cultured in equal volume of liquid PS medium. After culture for 7 days in a shaker, conidiation were measured by determining conidial concentration in the PS liquid medium. Conidia in the liquid media were counted with hemocytometer and photographed under microscope. The results showed that deletion of UvKMT6 caused a significant reduction (approximate 90%) in conidiation compared with those of WT and ΔUvkmt6-C strains (Figure 3d, e), suggesting that UvKMT6 is involved in asexual development in U. virens.

To examine whether UvKmt6 is required for the pathogenicity of U. virens, the mixtures of mycelia and conidia of WT, ΔUvkmt6-16, –18 and complemented strains were injected into booting-stage panicles of susceptible rice cultivar Wanxian 98 (Oryza sativa L. indica), respectively. Three weeks after inoculation, it was found that the ΔUvkmt6-16 and –18 developed less smut balls on inoculated panicles than WT (Figure 3f and g). In contrast, the reintroduction of UvKMT6 restored the virulence to the WT level.
Figure 3. Disruption of UvKMT6 resulted in reduced growth, conidiation, and pathogenicity in U. virens. (a) Colonies of the WT, ΔUvkm6 and complemented strains. Mycelial plugs of the WT, ΔUvkm6 and complemented strains were cultured on PSA plates for 14 d, and then photographed. (b) The colony diameters of the indicated strains. (c) UvKmt6 plays important roles in hyphal growth in U. virens. Mycelia of indicated strains cultured in PS media for 7 d were harvested and weighed after drying. (d and e) Knockout of UvKMT6 gene resulted in decreased conidation in PS medium. The mycelial plugs of the WT, ΔUvkm6 and ΔUvkm6-C strains were cultured in equal volume of liquid PS medium. At 7 dpi, conidiation were measured by determining conidial concentration in the PS liquid medium. Conidia in the liquid media were counted with hemocytometer and photographed under microscope. Bar = 5 μm. (f and g) The number of smut balls formed on the inoculated rice panicles with the ΔUvkm6 strain was less than those of the WT and complemented strains. Bar, 1 cm. Values represent the mean ± SD from three independent repeats. * or *** indicate P value <0.01 or <0.001 compared to the WT.
These results suggested that UvKmt6 contributes to fungal virulence in *U. virens*.

**UvKmt6 is essential for the establishment of histone modification H3K27me3**

To explore whether UvKmt6 is indeed required for H3K27me3 modification, nucleic proteins of WT, ΔUvkmt6-16, −18 and complemented strains were extracted and subjected to immunoblot with specific H3K27me3 antibodies. In the ΔUvkmt6-16 and −18 mutants, the levels of H3K27me3 modification were almost undetectable compared to obvious bands in the WT and complemented strain ΔUvkmt6-C (Figure 4a). In contrast, trimethylation of H3K36 and H3K4 modification, which are required for activation of transcription, did not show obvious changes after disruption of *UvKMT6* (Figure 4a). These results indicated that UvKmt6 is specifically required for H3K27me3 modification.

To further analyze the deposition of H3K27me3 on the chromatin, ChIP-seq assay was carried out with WT and ΔUvkmt6-16 strains. In the WT, H3K27me3 modification showed high enrichment and marked specific chromosomal regions (Figure 4b and supplemental table S2). In contrast, H3K27me3 peaks were nearly undetectable in the ΔUvkmt6 mutant, which was consistent with the aforementioned immunoblotting results. These results further indicated that UvKmt6 is essential for H3K27me3 modification in *U. virens*.

**UvKmt6-mediated H3K27me3 plays a vital role in transcriptional repression**

It is well known that Kmt6-mediated H3K27me3 modification plays vital roles in gene silencing [5]. To investigate the roles of H3K27me3 in the transcriptional regulation in *U. virens*, RNA-seq analysis was performed using vegetative mycelia of WT and ΔUvkmt6 cultured in the liquid PS medium for 7 d. Compared with the WT strain, a total of 1120 genes were differentially expressed in ΔUvkmt6 mycelia (adjusted p value < 0.01), with 629 genes up-regulated (log2 > 1) and 491 down-regulated (log2 < −1). Gene ontology (GO) analysis of differential expression genes (DEGs) showed that DEGs were involved in various important biological processes: cellular process (165/1120), development process (30/1120), positive and negative regulation of biological process (27/1120 and 39/1120), metabolic process (137/1120) and cellular component organization or biogenesis (71/1120), response to stimuli (56/1120) (Figure 5a). These data implied the involvement of UvKmt6-mediated H3K27me3 modification in extensive biological processes in *U. virens*.

![Image](image-url)

**Figure 4. UvKmt6 is essential for establishing H3K27me3 modification in *U. virens*.** (a) Deletion of *UVKMT6* resulted in loss of H3K27me3 modification. Total nuclear proteins of WT, ΔUvkmt6 and complemented strain ΔUvkmt6-C were isolated and separated by 15% SDS-PAGE (Sodium dodecylsulphate polyacrylamide gel electrophoresis) gel electrophoresis. Protein blots with specific H3, H3K27me3, H3K4me3 and H3K36me3 antibodies indicated that UvKmt6 is specifically involved in H3K27me3 modification. (b) The distribution of H3K27me3 modification in the WT and ΔUvkmt6 strains. H3K27me3-marked genomic sequences in the WT and ΔUvkmt6 strains were immunoprecipitated with H3K27me3 antibodies, and then sequenced. There was enrichment of H3K27me3 modification in the WT, but not in the ΔUvkmt6 strain. Two repeated biological experiments were carried out with similar results.
To determine the relationship between H3K27me3 enrichment and transcriptional regulation, comparative analysis was performed with three gene sets among H3K27me3-occupancy genes, up- and down-regulated genes in ΔUvkmt6. As shown in Figure 5b and c, the 140 out of 629 up-regulated genes in the ΔUvkmt6 mutant were H3K27me3 enriched genes, while only 12 down-regulated genes showed H3K27me3 enriched genes. The other 489 up-regulated genes may be indirectly regulated by H3K27me3 modification. These results suggested that the de-repressed genes in the ΔUvkmt6 are highly correlated with the absence of H3K27me3 occupancy in U. virens.

**Uvkmt6-mediated H3K27me3 modification regulates transcription of effectors in U. virens**

Recently, fungal histone modification H3K27me3 was demonstrated to repress transcription of effectors, which are specific expressed during in planta growth stage to contribute to pathogenicity [12,15,17]. In U. virens, there are at least 193 putative effectors [26]. Among them, transcriptional levels of 55 putative effectors are up-regulated during the vegetative growth stage when UvKMT6 was disrupted (Figure 6a and b). Notably, there were five effector genes occupied with H3K27me3, including Uv8b_6470, Uv8b_2964, Uv8b_2286, Uv8b_3638, and Uv8b_562, which have

![Figure 5. Uvkmt6-mediated H3K27me3 is mainly associated with transcriptional repression.](image-url)

(a) Gene Ontology (GO) analysis of differentially expressed genes (DEGs) between the ΔUvkmt6 mutant and WT strain. DEGs were significantly enriched in biological processes including cellular development and differentiation process, metabolic process and response to stimuli. (b) Venn diagrams show the overlap between H3K27me3-marked genes and up-regulated genes in the ΔUvkmt6 mutant when compared with WT strain. (c) Venn diagrams display the overlap of 12 genes between H3K27me3-marked genes and down-regulated genes in the ΔUvkmt6 mutant.
been reported to be involved in suppression of host immunity [26]. Furthermore, qRT-PCR and ChIP-qPCR analysis confirmed that these 5 effectors increased their expression for 2.5 to 45 folds (Figure 6c) and decreased their H3K27me3-occupancy for more than 90% (Figure 6d) in the ∆Uvkmt6 in comparison with those in WT. These results indicated that UvKmt6-mediated H3K27me3 modification participated in the transcriptional repression of effectors during vegetative growth stage in U. virens.

**Secondary metabolites-related genes are regulated by UvKmt6**

Genes encoding PKS (polyketide synthases), NRPS (nonribosomal peptide synthases), and Cytochrome P450 are involved in secondary metabolites synthesis and virulence in pathogenic fungi [22,26]. In F. graminearum, most of the genes encoding PKSs, NRPSs, and P450s are enriched with H3K27me3 modification, and are derepressed by deletion of FgKMT6.

Figure 6. UvKmt6-mediated H3K27me3 modification suppresses the expression of a large number of effectors at vegetative growth stage in U. virens. (a) The overlapping between U. virens 193 predicted effectors, up-regulated genes in ∆Uvkmt6, and H3K27me3-marked genes was present by venn diagrams. The association significance of two gene sets was calculated by Fisher’s exact test with P values labeled. (b) Heatmaps showing up-regulated expression levels of 55 effectors in ∆Uvkmt6 compared with those of WT during vegetative growth stage. (c) The relative transcriptional levels of 5 representative effectors were determined by qRT-PCR. (d) ChIP-qPCR verified the enrichment of H3K27me3 modification on the chromatin of representative effectors in the WT strain. Data represents mean ± SD of three independent replicates. ** or *** represent P value < 0.005 or < 0.001 compared to that of WT.
[13]. To comprehensively understand the transcriptional regulation of secondary metabolites synthesis genes by H3K27me3 modification, expression levels of 14 PKS, 17 NRPS and 39 Cytochrome P450 genes were compared in the WT and ΔUvkmt6 strains using the RNA-seq data. In contrast to WT, 11 of 14 PKSs, 9 out of 17 NRPSs and 20 of 39 Cytochrome P450 genes were significantly up-regulated in the ΔUvkmt6 strain (Figure 7a-c). To verify RNA-seq results of these PKSs, NRPSs and Cytochrome P450s genes, qRT-PCR assays were carried out. Consistent with RNA-seq data, deletion of UvKMT6 resulted in more than 6 folds up-regulation of 4 PKS genes including Uv8b_6010, Uv8b_6111, Uv8b_7563 and Uv8b_6705 (Figure 7d). When UvKMT6 was disrupted, transcriptional levels of 4 NRPS genes including Uv8b_3148, Uv8b_745, Uv8b_5435 and Uv8b_6701 also increased 3–70 folds in the ΔUvkmt6 mutants compared with those in the WT strain (Figure 7e). In addition, expression levels of 7 selected Cytochrome P450 genes including Uv8b_5604, Uv8b_3220, Uv8b_6892, Uv8b_2642, Uv8b_1536, Uv8b_4515 and Uv8b_5418 increased at various degrees from 6 to 25 folds (Figure 7f). Furthermore, ChIP-qPCR assay of the qRT-PCR tested up-regulated genes showed that most of them (12/15) were occupied with H3K27me3 modification except genes Uv8b_745, Uv8b_3220 and Uv8b_4515 (Figure S4). These results suggested that UvKmt6 participates in regulation of various secondary metabolites-related genes by H3K27me3-mediated repression in U. virens.

The toxic compounds and/or metabolites produced by the U. virens cultured in liquid PS media could inhibit germination of rice seeds [31,32,34]. Due to the significantly decreased H3K27me3-occupancy and increased relative transcription level of many secondary metabolic genes when UvKMT6 was deleted, it is suggested that the toxic compounds and/or metabolites produced by U. virens may be changed. Therefore, the toxicity of culture filtration of strains WT, ΔUvkmt6-16, −18 and ΔUvkmt6-C to rice seed germination was evaluated. The results showed that the length of shoots treated with filtration of ΔUvkmt6-16 and −18 was significantly shorter than those of WT and ΔUvkmt6-C strains (Figure S3), further indicating that disruption of UvKMT6 led to secondary metabolic disorder.

**UvKMT6 is involved in response to various stresses**

In order to survive and complete infection, U. virens fungus needs to adapt to various stresses from the host and environment [35]. By RNA-seq analysis, we found that a lot of stress response-related genes were affected extensively when UvKMT6 was deleted (Table S3 and S4). Cell wall stress response-related genes, including Uv8b_4757 (chitin synthase 4) and Uv8b_3908 (class 2 chitin synthase), were confirmed to be up-regulated by qRT-PCR assay (Figure 8a). Similar transcriptional up-regulation was also found among osmotic stress response-related genes, including Uv8b_1888 (UvHOG1) and Uv8b_4551 (UvSKN7), and oxidative stress response-related genes, including Uv8b_7732 (mono-oxygenase), Uv8b_7398 (ligninase h2 precursor), Uv8b_2 (WSC domain containing protein), Uv8b_1100 (cytochrome P450) and Uv8b_7112 (laccase-like protein) (Figure 8b and c). ChIP-qPCR assay of these genes showed that most of them (6/9) were occupied with H3K27me3 modification except genes Uv8b_4757, Uv8b_1888 and Uv8b_7732 (Figure S3) in the WT strain. To test if UvKmt6 plays a role in adaptation to stresses, the WT, Δ Uvkmt6, and Δ Uvkmt6-C strains were inoculated on the PSA medium and PSA plates amended with oxidative stress reagent H2O2, osmotic stress reagent NaCl or sorbitol, or cell wall stress reagents SDS (sodium dodecyl sulfate), CFW (calcofluor white) and CR (Congo red) for 15 d. Compared with WT or complemented strain ΔUvkmt6-C, the colonies of the ΔUvkmt6-16 and −18 strains were bigger and displayed less sensitivity to all these stress-mimicking reagents including H2O2, NaCl, sorbitol, SDS, CFW and CR (Figure 8d and e). Therefore, it was indicated that UvKmt6 regulated the response to oxidative, osmotic and cell wall stresses in U. virens.

In addition, RNA-seq analysis also showed that genes related to nutrient utilization were regulated by UvKmt6 (Figure 5a, Table S3 and S4). qRT-PCR analysis further confirmed that the expression levels of genes related to carbon utilization, including Uv8b_2924 (putative beta-glucosidase), Uv8b_7070 (glucan 1,3-beta-glucosidase) and Uv8b_4862 (putative glucosidase), were increased by 2 to 14 folds in the ΔUvkmt6 mutants (Figure 9a). Moreover, ChIP-qPCR assay of the three genes showed that both Uv8b_2924 and Uv8b_7070 were occupied with H3K27me3 in the WT (Figure S3), suggesting a regulatory role in response to nutrient condition by H3K27me3 modification in U. virens. Therefore, the growth of ΔUvkmt6-16 and −18, WT and complemented strains were compared on PSA, SD (synthetic dropout medium) and SD-G (synthetic dropout medium without glucose) plates. When cultured for 7 d, the colony size of ΔUvkmt6-16 and −18 mutants were smaller than that
of WT or complemented strains on the PSA and SD plates. In contrast, the colony diameter of the ΔUvkmt6-16 and ΔUvkmt6-18 were larger than that of WT or complemented strains on SD-G plates (Figure 9b and c), suggesting that UvKMT6 is involved in the response to nutrient stress.
Discussion

Epigenetic modification of H3K27me3 established by PRC2 complex plays important roles in transcriptional regulation and development in various organisms [1]. *U. virens* caused rice false smut disease is a threat to rice production and food safety due to the formation of smut balls and mycotoxins [19]. In order to grasp the biological roles of PRC2-mediated H3K27me3 modification in *U. virens*, we investigated the functions of the uncharacterized homologue of methyltransferase Kmt6, which is a core component of PRC2 complex, in this study. We noted that, unlike essential roles of Kmt6 in *F. fujikuroi* [36], UvKMT6 gene was knocked out successfully. Using the ΔUvkmt6 mutants, we found that the UvKmt6-mediated H3K27me3 modification plays significant roles in development and pathogenicity in *U. virens*. In addition, transcriptome profiling, ChIP-seq assay in combination with phenotypic analysis revealed that UvKmt6 participates in extensive regulation of a wide range of genes including effectors, secondary metabolites synthesis genes, and stress response-related genes.
The characterization of virulence-related factors will better help us to reveal the infection mechanism of *U. virens*. In this study, an epigenetic regulator UvKmt6 was found to contribute to the virulence of *U. virens* in several ways. First, disruption of *UvKMT6* resulted in drastic decrease in conidiation, which usually leads to reduced pathogenicity in *U. virens* [35]. For example, UvAtg8 is an autophagy marker, and its gene deletion mutant is significantly reduced in pathogenicity due to the decrease of conidia production [32]. Moreover, transcriptional factor UvCom1 and UvHox2 involved in conidiation are also necessary for the rice false smut balls formation [37,38]. The reduction in conidiation could be caused by the transcriptional dysregulation of conidiation-related genes in the deletion mutant of *UvKMT6* (Table S3 and S4). Second, approximately a quarter of putative effectors were abnormal expressed when *UvKMT6* was deleted. As we know, effectors secreted by pathogens are employed to suppress host immunity and contribute to virulence [28,29,39]. The fact that downregulation of *UvKMT6* during *U. virens* infection at 7 dpi also

**Figure 9. The UvKmt6 is related to carbon sources starvation response.** (a) Expression levels of carbon utilization-related genes increased in the ΔUvkmt6-18 strains were verified by qRT-PCR assay. (b and c) UvKmt6-mediated H3K27me3 modification plays a role in response to nutrient starvation stress. The WT, ΔUvkmt6 and ΔUvkmt6-C strains were grown on the PSA, SD, and SD-G plates for 14 d, and then photographed. The colony of ΔUvkmt6 cultured on PSA and SD plates was smaller than WT, but bigger on the SD-G plates than WT. Data represents mean ± SD of three independent replicates. Asterisks highlight statistically significant difference (**, P < 0.005; ***, P < 0.001).
suggests that it contributes to releasing transcriptional suppression of H3K27me3-repressed genes such as effectors and promote infection. Therefore, we infer that the abnormal expression of effector proteins may be an important reason for the decrease of virulence in ∆Uvkmt6 mutants. Third, the deletion of UvKMT6 resulted in alteration of stress sensitivity in U. virens, which may also be one of the reasons for the reduced virulence of ∆Uvkmt6 mutants. In previous reports, it has been implicated that alteration of stress sensitivity links to virulence of U. virens. For instance, absence of the cAMP signaling pathway component UvAc1 and UvPdeH or MAPK UvPmk1 displayed differential sensitivity to stress-mimicking reagents and reduced virulence [30,33]. As H3K27me3 is a hallmark of facultative heterochromatin regions of the genome, which has been reported to be associated with genome stability [11], it is very possible that deletion of UvKmt6 may also affect the genome stability in U. virens. In conclusion, UvKmt6 plays an important role in the virulence of U. virens likely by regulating the condensation, maintaining spatial and temporal specific expression of effectors, genome integrity and responding to stresses.

Enrichment of H3K27me3 on chromosomes depends on the Kmt6 methyltransferase in PRC2 complex [11,13,16]. In U. virens, UvKmt6 is also essential for marking genomic regions with H3K27me3, which is demonstrated by immunoblot and ChIP-seq (Figure 4). It was known that part of H3K27me3-marked genes tend to be silenced in different filamentous fungi, such as in N. crassa, M. oryzae, F. graminearum and F. fujikuroi [3,13,17,36]. Notably, genes involved in secondary metabolism in F. graminearum and F. fujikuroi are marked with H3K27me3, and disruption of KMT6 also derepresses expression of PKS, NRPS and Cytochrome P450 genes [13,36]. Here, we found that the transcriptional repression activity was also associated with local enrichment of H3K27 methylation in U. virens. This is supported by the fact that 140 genes of 321 H3K27me3-occupation genes were up-regulated in the UvKMT6 deletion mutant, while only 12 down-regulated genes overlapped with 321 H3K27me3-occupation genes. Furthermore, disruption of UvKMT6 induced expression of effectors, secondary metabolism-related genes including PKS, NRPS and Cytochrome P450 genes and stress response-related genes, which were occupied by H3K27me3 modification in the WT. Thus, our results further implied that the roles of H3K27me3 in transcriptional repression are relatively conserved.

In addition to the relatively conservative regulation of effectors and secondary metabolic gene expression, we further found that UvKmt6 appears to be a negative factor in responding to cell wall stress, osmotic stress, and oxidative stress in U. virens. The increased tolerance to various stresses in the UvKMT6 deletion mutant might be caused by UvKmt6-mediated repression of stress response-related genes such as chitin synthases-encoded genes Uv8b_3908 and Uv8b_4757, osmotic stress responding genes UvHOG1 and UvSKN7, and oxidative stress responding genes including laccase and cytochrome P450s. In U. virens, knock-out of MAPK HOG1 encoding gene UvHOG1 resulted in increased sensitivity to osmotic and cell wall stresses, and down-regulated transcriptional expression of stress response-related gene UvSKN7 [31]. Therefore, the increased expression of UvHOG1 and UvSKN7 may be one of the explanations for the tolerance of ∆Uvkmt6 mutants to osmotic and cell wall stresses. Except cell wall, osmotic and oxidative stresses, ∆Uvkmt6 mutant also showed a better growth than WT strain under carbon source starvation stress. These phenotypes may be associated with increased expression of glucosidase-encoded genes in ∆Uvkmt6 mutant. It has been reported that glucosidases catalyze efficient hydrolysis of β-glucan to produce glucose in M. oryzae [40]. The underlying mechanism of UvKmt6-mediated H3K27me3 modification on expression of these stress response-related genes need to be further studied.

Taken together, our results shed light on important roles of UvKmt6-mediated H3K27me3 modification in transcriptional repression of effectors during vegetative growth stage, secondary metabolites, stress response, development, and pathogenicity in U. virens. As far as we know, our study is the first report on depicting roles of epigenetic modification in U. virens, which extends current understanding of epigenetic regulation in pathogenic fungi and supports the link between H3K27me3 modification and infection-related biological processes in U. virens.

Materials and methods

Strains and growth condition

The U. virens WT strain HWD-2 was a kind gift from Prof. Junbin Huang of Huazhong Agriculture University (China). The strains were cultured on potato sucrose agar (PSA, potato 200 g/L, sucrose 20 g/L and agar 20 g/L) plates in constant dark chamber at 28°C. To evaluate vegetative growth, mycelial plugs were cut from 14-d-old PSA culture and transferred to new PSA plates for 14 d. To collect mycelia and conidia, mycelial plugs grew in liquid potato sucrose media shaking at 180 rpm for 7 d. Conidia were harvested from the filtrate by filtering the liquid cultures with three layers of gauze to determine conidial concentrations with a hemocytometer.
**The construction of vectors and transformation**

To generate the deletion mutants of *UvKMT6*, a gene replacement strategy was used [32]. Briefly, approximate 1 Kb of 5’ UTR and 3’ UTR regions were amplified from the genomic DNA of the HWD-2 strain using the primer pairs UvKMT6-5 F/R and UvKMT6-3 F/R (Supplementary Table S1). Then, these flanking fragments were ligated sequentially to the flanking of *hygromycin resistance gene* cassette in the pFGL821 (Addgene, 58,223). The sequence of resultant plasmid was confirmed by sequencing and subsequently introduced into the HWD-2 strain by ATMT. To generate the complementary vector, the *UvKMT6* fragment containing 1.5 Kb of promoter and coding region was amplified (primers listed in Table S1) and ligated into the vector pFGL823 [34]. The sequence of plasmid was confirmed by sequencing, and then introduced into the deletion mutant by ATMT to obtain the complemented strains. All the correct transformants were confirmed by PCR, qRT-PCR and Southern blot assay (primers listed in Table S1) [32].

**Stresses treatments**

To determine sensitivity to various stresses, mycelial plugs were inoculated on the PSA plates supplemented with 0.015% H₂O₂, 0.3 M NaCl, 0.5 M sorbitol, 0.03% SDS, 120 mg/mL CFW or 120 mg/mL CR, respectively. Stress sensitivity was assessed by measuring the diameter of the plates after cultured for 14 d at 28°C. The formula of relative inhibition rate was calculated as follow: Growth inhibition rate = (diameters of strain colony on the PSA – diameters of strain colony on the PSA amended with different chemicals)/diameters of the strain colony on the PSA × 100%. For the nutrient starvation assay, the SD (yeast nitrogen base without amino acids 1.7 g/L, ammonium sulfate 5 g/L, glucose 20 g/L and agar 20 g/L) medium and SD-G (yeast nitrogen base without amino acids 1.7 g/L, ammonium sulfate 5 g/L and agar 20 g/L) plates were used. Pictures were taken to present the vegetative growth under stress conditions. All these assays were repeated three times with at least three copies each time.

**Inoculation assay**

The inoculation assay was performed as previously described [25,34,41]. Briefly, seeds of the rice cultivar Wanxian 98 (a susceptible rice cultivar, *Oryza sativa* L. *indica*), were cultivated in field for approximate three months and transferred to greenhouse until the booting stage. The WT, ∆Uvkmt6 and complemented strain ∆Uvkmt6-C were cultured in PS (potato 200 g/L, sucrose 20 g/L) media for 7 d. Then, the resultant cultures were broken into mycelial pieces to make a mixed suspension of mycelia with conidia, and the mixtures were normalized by adjusting conidia concentration to 1 × 10⁶/mL. 2 mL of the mixture was injected into panicles and all rice plants were cultivated at the relative humidity of 95% and 22°C for 2 d in the dark following 28°C for 3 weeks. The disease symptoms were represented by the number of false smut balls and the photo of infected panicles. Toxicity assays with culture filtrates were performed as described using Wanxian 98 seeds (Zheng et al., 2016). All experiments in this part were repeated three times in each test.

**Western blotting**

For detecting histone modification, 0.2 g mycelia were collected from 7 dpi in the liquid PS and ground in liquid nitrogen. The ground mycelial powders were suspended with buffer I (20 mM Tris pH 7.5, 20 mM KCl, 2 mM MgCl₂, 25% Glycerol, 250 mM Sucrose, 0.1 mM PMSF (Phenylmethanesulfonyl fluoride), 5 mM beta-mercaptoethanol, 1× proteinase inhibitors). The resulting solutions were filtered through one layer of Miracloth (Millipore) and centrifuged. Then, the pellets were suspended with buffer II (50 mM Tris-HCl pH 7.4, 150 mM NaCl, 1 mM EDTA, 1% Triton100 and 1× protein inhibitor) to obtain total nucleus proteins. Total nucleus proteins were subject to 15% SDS-PAGE and transferred to PVDF membrane, which immunoblotted with antibodies including anti-H3 (Huabio, M1309-1), anti-H3K27me3 (Active motif, 39,155), anti-H3K4me3 (Abcam, ab1012), anti-H3K36me3 (Abcam, ab9050) antibody, respectively. The results of immunoblotting were captured with an imaging system using a chemiluminescence kit (Bio-Rad).

**Chromatin immunoprecipitation (ChIP) and sequencing**

The ChIP experiments were performed according to previous methods with minor modification [42,43]. Briefly, 7-d cultured mycelia cultured in the PS media were cross-linked with 1% formaldehyde for 20 mins. After glycine solution (125 mM) was added to stop crosslink, mycelia was harvested with three layers of gauze, ground and suspended with the nuclei extracting buffer (10 mM Tris pH 8.0, 10 mM Sodium butyrate, 400 mM Sucrose, 0.1 mM PMSF, 5 mM β-
mercaptoethanol and 1 × proteinase inhibitors). The resultant nuclei were then lysed with 1 mL nuclei lysis buffer (50 mM HEPES pH 7.5, 150 mM NaCl, 1 mM EDTA, 10 mM Sodium butyrate, 0.1% deoxycholate, 1% Triton X-100, 0.1% SDS, 1 mM PMSF and 1× Roche protease inhibitor cocktail). The nuclear DNA solution was broken into DNA fragments whose lengths ranged from 200 to 500 bp using a sonication machine Diagenode Bioruptor (Figure S3). After pre-treating with protein A beads (Thermofisher, 10001D), the supernatant was incubated with anti-H3K27me3 antibody (Active motif, 39,155) overnight at 4°C. Then another piece of protein A beads was added into the above reaction system to bind anti-H3K27me3 antibody and then washed three times. The immunoprecipitated DNA was recycled and used for ChIP-seq and ChIP-qPCR. For ChIP-seq assay, the recycled DNA was sequenced using a high-throughput sequencing platform Illumina Hiseq-PE150 by the Novogene Corporation (China). The resulting clean read pairs were mapped to the genome of the Uv8b strain with Bowtie2 (Version 2.2.8) [44]. The peak calling was performed with MACS2 (Version 2.1.1) [45], and visualized with the integrative genomics viewer [46]. The ChIP-seq assay was conducted with two biological repeats. To verify whether genes were occupied by H3K27me3, ChIP-qPCR assays with two independent repeats were conducted with gene TUB as internal reference. The qPCR assay was performed by using the immunoprecipitated DNA with anti-H3K27me3 antibody from the wild-type strain and ΔUvkmt6 as templates with the tubulin gene (Uv8b_900) as internal control. The statistical analysis was performed with t test. The PCR primers used in these experiments were listed in the Supplementary Figure S6 and Table S1.

qRT-PCR and RNA sequencing

Total RNA was isolated from mycelia cultured for 7 d in liquid PS with three independent biological repeats using the TRizol (Invitrogen) reagent. Subsequently, cDNA were synthesized with a reversely transcribed kit (TAKARA). The cDNA was subjected to qRT-PCR assay with SYBR Green qPCR Master Mix (TAKARA) using the tubulin gene (Uv8b_900) as internal control. The primers used in the experiments were listed and described in Supplementary Table S1.

RNA sequencing and comparison of differentially expressed genes of indicated strains was performed as previously described [47]. Briefly, purified RNA concentration was measured with Qubit RNA assay kit and the RNA integrity was determined by Bioanalyzer 2100 RNA-6000 Nano Kit (Agilent Technologies, CA, USA). A total amount of 3 μg RNA was used for RNA sample preparations. The mRNA-seq libraries were constructed using NEBNext UltraTM RNA Library Prep Kit (New England Biolabs, California, America) according to the manufacturer’s instructions followed by sequencing on an Illumina Hiseq X-Ten with Hiseq-PE150 strategy at Novogene. The resulting clean reads after removing adapter, low quality reads and reads containing ploy-N with fastp were mapped to the U. virens reference genome using Hisat2 v2.0.5. Then the featureCounts v1.6.4 were used to generate raw counts. The gene transcript abundances were normalized and differentially expressed genes analysis was analyzed by DESeq2 software. Only the genes’ different expression with log2|FC| > 1 and P-adjust values < 0.05 were considered significantly differential expressed. GO enrichment analysis was carried out under the Bonferroni-corrected P ≤ 0.05 compared with the whole-transcriptome background. GO functional enrichment analysis was accomplished on the website (https://www.omicshare.com/tools/). Reference genome and gene model annotation files were downloaded from genome website directly (https://ftp.ncbi.nlm.nih.gov/genomes/all/GCA/000/687/475/GCA_000687475.1_Assembly_for_version_1_of_the_Villosiclava_virens_genome/). Three biological replicates were performed for each strain/mutant. To determine the significance of the association between two sets of genes, P value was obtained by Fisher’s exact test using TBtools [48].

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Disclosure statement

No potential conflict of interest was reported by the authors.

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Data availability statement

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

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