ARS2 Plays Diverse Roles in DNA Damage Response, Fungal Development, and Pathogenesis in the Plant Pathogenic Fungus Fusarium graminearum

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Arsenite-resistance protein 2 (Ars2) is an important nuclear protein involved in various RNA metabolisms in animals and plants, but no Ars2 ortholog has been studied in filamentous fungi. Although it is an essential gene in most model eukaryotes, FgARS2 null mutants were viable in the plant pathogenic fungus Fusarium graminearum. The deletion of FgARS2 resulted in pleiotropic defects in various fungal developmental processes. Fgars2 mutants were irregular in nuclear division, and conidial germination was significantly retarded, causing the fungus to manifest its hypersensitive phenotypes under DNA damage stress. While FgARS2 deletion caused abnormal morphologies of ascospores and defective ascospore discharge, our data revealed that FgARS2 was not closely involved in small-non-coding RNA production in F. graminearum. The dominant nuclear localization of FgArs2-green fluorescent proteins (GFP) and abnormal nuclear division in FgARS2 deletion mutant implicated that FgArs2 functions in the nucleus.

Intriguingly, we found that FgArs2 established a robust physical interaction with the cap binding complex (CBC) to form a tertiary complex CBC-Ars2 (CBCA), and disruption of any CBCA complex subunit drastically attenuated the virulence of F. graminearum. The results of the study indicate that Ars2 regulates fungal development, stress response, and pathogenesis via interaction with CBC in F. graminearum and provide a novel insight into understanding of the biological functions of Ars2 in filamentous fungi.

Keywords: Fusarium graminearum, arsenite-resistance protein 2, DNA damage response, fungal reproduction, fungal virulence

INTRODUCTION

Genome maintenance is a constant concern for the identity and function of eukaryotic cells. However, DNA is continuously damaged by endogenous and environmental agents, posing a serious threat to the faithful transmission of genetic information (Rouse and Jackson, 2002). Defects caused by un-repaired or mis-repaired DNA damages result in genomic instability, cell cycle arrest, and lethality, underlining the importance of these processes in the cell and whole...
to the proliferative state of the cell, and a conditional
caused increased leaf production and earlier flowering (Prigge
abnormalities during shoot development (Clarke et al., 1999;
A. thaliana ARS2 2010). In addition, it is likely that
knockout in hematopoietic tissues leads to decreased cellularity
in bone marrow (Gruber et al., 2009). However, the precise
biological and biochemical functions of
ARS2 biological and biochemical functions of
in fission yeast, plants, insects,
and mammals (Golling et al., 2002; Oh et al., 2003; Amsterdam et al., 2004; Lobbes et al., 2006; Wilson et al., 2008; Kim et al., 2010). In addition, it is likely that ARS2 plays a critical role as a developmental regulator in cell proliferation. In plants, mutation of the A. thaliana ARS2 ortholog resulted in pleiotropic abnormalities during shoot development (Clarke et al., 1999; Prigge and Wagner, 2001), while the overexpression of AtSE caused increased leaf production and earlier flowering (Prigge and Wagner, 2001). In human cells, HsARS2 expression is linked to the proliferative state of the cell, and a conditional HsARS2 knockout in hematopoietic tissues leads to decreased cellularity in bone marrow (Gruber et al., 2009). However, the precise biological and biochemical functions of ARS2 and the CBC in pathogenic fungi are largely unknown.

The ascomycete Fusarium graminearum is an important plant pathogen causing Fusarium head blight (FHB) in cereal
crops worldwide (Sutton, 1982; Goswami and Kistler, 2004). The serious consequences of F. graminearum infection are not only the losses of yield and quality, but contamination of the grains by mycotoxins, such as trichothecenes and zearalenone, that threaten human and animal health (Desjardins, 2006). Although F. graminearum reproduces both sexual (ascospores) and asexual (conidia) spores, ascospores are thought to have essential roles as primary inocula (Sutton, 1982; Guenther and Trail, 2005; Leslie and Summerell, 2006). In addition, the initial structures or associated hyphae of the perithecia also function as survival structures for overwintering, indicating its indispensable part in the fungal life cycle (Sutton, 1982; Guenther and Trail, 2005). Since previous biological and chemical strategies to control FHB have not been effective in preventing disease outbreaks, a comprehensive understanding of the cellular and molecular mechanisms of fungal development and pathogenesis as well as the host stress/defense responses is necessary to develop effective and sustainable methods to manage FHB (Goswami and Kistler, 2004).

Our previous studies identified sixteen putative transcription factors (TFs) involved in DDR and further demonstrated that cell cycle regulation and DDR were closely related to sexual reproduction and virulence in F. graminearum (Son et al., 2011c; Min et al., 2014; Son et al., 2016). In this study, we attempted to functionally analyze another gene involved in DDR, FgARS2, using cellular, genetic, and biochemical approaches. FgARS2 was not the essential gene in F. graminearum, in contrast to the lethality of its orthologs in model eukaryotes. Therefore, the objectives of the study were to uncover the genetic and biological functions of FgARS2 in DDR, fungal development, and pathogenesis in F. graminearum. Our findings shed light on the novel role of the CBCA complex and DDR in the fungal pathogenesis. To our knowledge, this study is the first to characterize the ARS2 ortholog in filamentous fungi and provides new insights into its diverse roles in fungal developmental processes and pathogenesis.

MATERIALS AND METHODS

Fungal Strains and Media

The F. graminearum wild-type strain Z-3639 (Bowden and Leslie, 1999) and the mutants used in this study are listed in Supplementary Table S1. Standard laboratory methods and culture media for Fusarium species were used (Leslie and Summerell, 2006). For fungal sporulation, the conidia of all strains were induced on yeast malt agar (YMA) (Harris, 2005) or in carboxymethyl cellulose (CMC) medium (Cappellini and Peterson, 1965). The growth temperature for the fungal strains was 25°C. The wild-type and transgenic strains were stored as mycelia and conidia in 20% glycerol at −80°C.

Nucleic Acid Manipulation and Genetic Modifications

The genomic DNA was extracted following the standard protocol (Leslie and Summerell, 2006). Restriction endonuclease digestion, agarose gel electrophoresis, gel blotting, and
DNA blot hybridization were performed using standard techniques (Sambrook and Russell, 2001). The PCR primers (Supplementary Table S2) used in this study were synthesized by an oligonucleotide synthesis facility (Bionics, Seoul, South Korea).

To complement the Fgars2 deletion mutants, 5′ and 3′ flanking region including open reading frame (ORF) of the Fgars2 allele was amplified using the Fgars2-5F com/Fgars2-3N com primer pair from F. graminearum strain Z-3639 (Supplementary Table S2). The hygromycin resistance cassette (HYG) was amplified from the pBCATPH vector using the pBCATPH-comp 5′For/pBCATPH-comp 3′Rev primer pair (Min et al., 2012). The resulting amplicons were fused via double-joint (DJ) PCR as previously described (Yu et al., 2004). The final PCR constructs were transformed into the Fgars2 deletion mutants as described previously (Han et al., 2007).

To generate the Fgcbp20 and Fgcbp20 deletion mutants, the 5′- and 3′-flanking regions of each gene and a genetic resistance cassette (GEN) were amplified from Z-3639 and pH99, respectively, and were fused by a second round DJ PCR. The subsequent procedures for the third round of PCR and transformation were the same as those used for complementation of the Fgars2 gene.

To construct mutants overexpressing a green fluorescent protein (GFP)-tagged Fgars2, the 5′-flanking region of the Fgars2 and Fgars2 ORFs were amplified using primer pairs Fgars2-5F OEG/Fgars2-5R OEG and Fgars2-3F OEG/Fgars2-3R OEG, respectively. A GEN-PEF1α -GFP fragment carrying elongation factor 1α promoter (PEF1α) of F. verticillioides followed by the GFP ORF was amplified from pSKGEN (Lee et al., 2011) with the primers Neo-for new and eGFP-P1. Three amplicons were fused as described above. A final product was amplified using the nested primers Fgars2-5N OEG and Fgars2-3N OEG.

### Stem-Loop Reverse Transcription- and Quantitative Real-Time (qRT)-PCR

To detect small RNAs, stem-loop RT-PCR analyses of siRNAs and miRNAs were performed as previously described (Chen et al., 2005; Varkonyi-Gasic et al., 2007). Briefly, RNAs highly enriched for small RNA species (<200 nt) were isolated using a mirVana™ miRNA isolation kit (Invitrogen, Carlsbad, CA, United States). Each 100 ng of RNA samples and the stem-loop RT primers that hybridized to the small RNA molecules were used for the reverse transcription reactions, followed by amplification of RT products using miRNA-specific forward primer and the universal reverse primer from 25 to 30 cycles (Supplementary Table S2). The amplification products were visualized on 3% agarose gels.

Total RNA for qRT-PCR was extracted from mycelia that were grown in CM for 3 days or perithecia after 7 days of sexual induction using an Easy-Spin total RNA extraction kit (iNtRON Biotechnology, Seongnam, South Korea). First-strand cDNA was synthesized with a SuperScript III First-Strand Synthesis System (Invitrogen) using oligo(dT)20 according to the manufacturer’s instructions. qRT-PCR was performed using an SYBR Green Master Mix (Bio-Rad, Hercules, CA, United States) and a CFX real-time PCR system (Bio-Rad). The endogenous housekeeping gene ubiquitin C-terminal hydrolase (FgUBH1; FGS0_01231) was used for normalization (Kim and Yun, 2011). The PCR assays were repeated three times with two biological replicates. The transcript level relative to that of the housekeeping gene was expressed as 2−ΔΔCT (Livak and Schmittgen, 2001).

### Sexual Development and Virulence Tests

Mycelia grown on carrot agar for 5 days were mock-fertilized with sterile 2.5% Tween 60 solution to induce sexual reproduction as previously described (Leslie and Summerell, 2006). Eight days after induction, the perithecia from each strain were dissected in a drop of 15% glycerol, and the ascospores within the perithecia were observed under a DE/Axio Image A1 microscope (Carl Zeiss, Oberkochen, Germany). For ascospore discharged observation, a semicircular agar block (11 mm in diameter) that was covered with mature perithecia was placed on a coverslip and incubated in the chamber for 24 h (Trail et al., 2005).

A virulence test of the fungal strains was performed using the wheat cultivar Eunpamil as previously described (Son et al., 2011a). Briefly, 10 µL of a conidial suspension (1 × 10^5 conidia mL^-1) of each strain was point-inoculated onto a spikelet of the wheat head at early anthesis. Inoculated plants were incubated in a humidified chamber for 3 days and subsequently transferred to a greenhouse. After 21 days, the number of spikelets showing disease symptoms was counted.

### Microscopic Observation

Microscopic observation was performed using a DE/Axio Imager A1 microscope (Carl Zeiss) equipped with the filter set 38HE (excitation 470/40; emission 525/50) for GFP and the filter set 15 (excitation 546/12; emission 590) for red fluorescent protein (RFP).

Wheat heads inoculated with the GFP expressing strains were sampled 6 days after inoculation. Freehand longitudinal sectioned across the center of the spikelets were prepared using a clean scalpel (Baldwin et al., 2010). Sectioned wheat heads were observed under reflected light and GFP-fluorescent light (470-nm excitation and 525-nm emission wavelength filters) using a SteREOMar Lumar V12 microscope (Carl Zeiss).

### Yeast Two-Hybrid Assay

A yeast two-hybrid assay was conducted using the DUALhunter System (Dualsystems Biotech, Zurich, Switzerland) according to the manufacturer’s instructions. To construct plasmids, the coding sequence of each tested gene was amplified from the cDNA of Z-3639 using primers with a Saccharomyces cerevisiae GAL4 expression site (Supplementary Table S2). Full-length Fgars2 and Fgcbp20 cDNAs were cloned into the pDH81 vector, a Cub-based bait vector, while full-length Fgcbp80 and Fgcbp20 cDNAs were cloned into pRN3-N, a NubG-based prey vector.

Saccharomyces cerevisiae strain NMY51 (MATa his3Δ200 trpl-901 leu2-3, 112 ade2 lys2-(lexAop)4-HIS3 ura3-(lexAop)8-lacZ ade2:(lexAop)8-ADE2 GAL4) was grown in YPD medium
(1% Bacto yeast extract, 2% Bacto peptone, 2% glucose, and 0.004% adenine sulfate). Bait and prey plasmids were co-transformed using the lithium acetate method (Gietz and Schiestl, 2007). Transformants were selected after 3–4 days of growth at 30°C on synthetic dextrose (SD) medium lacking histidine, leucine, tryptophan, and adenine (SD-Leu/Trp, SD-Leu/Trp/His, and SD-Leu/Trp/His/Ade). Empty vector and pDL2-Alg5 (−) were included as negative controls, and pA1-Alg5 (+) was included as a positive control.

RESULTS

FgARS2 Is Required for Vegetative Growth and Maintenance of Cell Cycle Progression

In silico analysis of the protein sequence revealed that FgArs2 (locus ID: FGSG_01106) contains 870 amino acids, and its domain architecture is similar to those of other representative orthologs in some model eukaryotes (Figure 1A). However, FgArs2 showed a low similarity and identity in protein sequence compared to other representative orthologs; 12–16% in the overall sequence, 18–31% in DUF3546 domain, and 15–39% in ARS2 domain, respectively (Supplementary Figure S1). In addition, the nuclear localization signals predicted by NLStradamus (Ba et al., 2009) displayed two sequences (NLS1, 50 – RRDRRERSSPAADVRYEPPRR – 72; and NLS2, 134 – MKEEKERARTGRRREPRTGPDREKKA – 163) at its N-terminus, suggesting that FgArs2 might function as a nuclear protein in F. graminearum.

To carry out genetic complementation, the construct containing the FgARS2 ORF fused with a hygromycin B resistance gene cassette was introduced into protoplasts of the Fgars2 strain, resulting in DCB101 strains (Supplementary Figure S2). Fgars2 exhibited reduced radial growth, and the aerial mycelia of Fgars2 tended to be denser than those of the wild-type and DCB101 strains on both complete medium (CM) and minimal medium (MM) (Figure 1B). The deletion of FgARS2 also significantly retarded conidial germination compared to the wild-type and DCB101 strains (Figure 1C). In particular, approximately 92 and 88% of conidia in the wild-type and DCB101 strains germinated 5 h after inoculation in CM, respectively, while only about 40% of conidia of the Fgars2 strains germinated. However, all of conidia from wild-type, Fgars2, and DCB101 strains were eventually germinated at 8 h after incubation, indicating their normal viability (Figure 1C).

To decipher the mechanism underlying defective vegetative growth, the DCB102 (Fgars2 hH1-GFP) strains were generated by an outcross between the mat1g (mat1 hH1-GFP) and Fgars2 strains (Supplementary Table S1). Microscopic observation revealed that the deletion of FgARS2 caused a defective phenotype of conidia during nuclear division but did not affect the conidial production, size, or septum number. As a result, 24% of the conidia of DCB102 bore more than two nuclei within a cell, while 10% of these conidia with two nuclei within a cell were scored in the wild-type strain (Figures 1D,E). The mitosis assay further revealed that the lack of FgARS2 resulted in marked irregularity in nuclear division during germination, while the nuclear division was arrested in both wild-type and Fgars2 strains when treated with a high concentration of hydroxyurea (HU) (Figure 1E). Fgars2 strains showed sensitive phenotypes and were significantly inhibited the growth rate against both HU and bleomycin (BLM) which are DNA-damaging agents (Figures 1F,G). The expression level of FgARS2 in the wild type during HU treatment was significantly higher than that of Fgars2 without HU treatment (Figure 1H), corroborating the bona fide role of FgARS2 during DDR in F. graminearum. Taken together, these results demonstrated that FgARS2 is required for vegetative growth and the maintenance of the cell cycle progression in F. graminearum.

FgARS2 Is Important for Pathogenesis

Since cell cycle regulation and DDR are closely related to pathogenesis in plant pathogenic fungi, we evaluated the pathogenicity of the Fgars2 in flowering wheat heads. As expected, the wild-type strain caused typical head blight symptoms of wheat heads (Figure 2A). In contrast, the Fgars2 mutants were unable to spread from the inoculated spikelet to adjacent spikelets on the heads, leading to a significant decrease in the disease index compared to that in the wild-type strain (Figure 2A). This avirulent feature of the Fgars2 mutants was fully restored to the wild-type level in DCB101, the Fgars2-complemented strain, suggesting the important role of this gene during wheat infection of F. graminearum.

We generated an Fgars2 strain constitutively expressing cytosolic GFP, the DCB103 (Fgars2 GFP-HYG), to observe mycelial movement during wheat head infection via an outcross between KM19 (mat1 GFP-HYG) and Fgars2. The HK12 strain (Son et al., 2011b), F. graminearum wild-type strain expressing GFP and carrying the Fgars2 allele, readily colonized the injected spikelet approximately 6 days after inoculation and began to infect adjacent spikelets via rachis nodes (Figure 2B). Although Fgars2 successfully colonized the injected spikelets as HK12, the hyphae of Fgars2 could not spread to the neighboring spikelets from those that were inoculated. In addition, hyphal growth via rachis was rarely observed in Fgars2-infected plants.

FgARS2 Is Involved in Sexual Reproduction

The sexual development of the F. graminearum strains was observed on 7-day-old sexually induced cultures. All F. graminearum strains tested including the Fgars2 deletion strains produced mature perithecia successfully, however, the perithecium numbers of Fgars2 were significantly reduced compared to those of the wild type (Figure 3A). The deletion of Fgars2 also resulted in severely defective forcible ascospore discharge compared to the wild type (Figure 3B). All of these defective phenotypes of the DCB101 strain were restored to the wild-type level during sexual reproduction.
To further unravel the defective phenotypes of *Fgars2* in sexual development, we dissected mature perithecia to observe rosette asci (Figure 3C). The *F. graminearum* wild-type strain typically produced eight spindle-shaped ascospores per ascus. In contrast, some asci of the *Fgars2* mutants that were defective in ascospore discharge contained abnormally shaped ascospores. Further microscopic observation clarified that the abnormally shaped ascospores varied in their morphologies, as some are larger, smaller, and broken (Figure 3C). These data suggest that the defective forcible ascospore discharge of *Fgars2* mutants was derived from abnormalities in ascospore production.

**Fgars2** Is Not Closely Involved in Small Non-coding RNA Production

ARS2 and its orthologs have been known to be required for miRNA and siRNA production. Also, our previous study reported that the exonic small interference RNA (ex-siRNA)-mediated RNA interference (RNAi) mechanism fine-tunes the
FIGURE 2 | Pathogenesis of Fgars2 mutants. (A) Virulence on wheat heads. The center spikelet of each wheat head was injected with 10 µL of a conidial suspension. The disease index (diseased spikelets per wheat head) is denoted below the picture. The asterisks indicate that the data differ significantly ($P < 0.05$) as determined by Tukey's test. The pictures were taken 21 days after inoculation. Mock, negative control mock-inoculated with 0.01% Tween 20. (B) Micrographs of manually generated sections after infection of wheat. Wheat spikelets were inoculated with conidial suspensions from strains expressing GFP in the cytoplasm. Infected wheat heads were longitudinally dissected 6 days after inoculation and examined under a fluorescence microscope. GFP fluorescence represented hyphae spreading from the inoculation points. Arrowheads marked the inoculated spikelets. Reflected, reflected light.

transcriptome during ascospore formation in F. graminearum (Son et al., 2017). In order to investigate whether Fgars2 plays an important role for siRNA biogenesis, we first examined the production of ex-siRNA candidates in the wild-type and Fgars2 strains (Figure 4 and Supplementary Figure S2). Abundances of the siRNAs were easily detectable in both the wild-type and Fgars2 strains after 20, 25, or 30 cycles of stem-loop PCR (Figure 4). Furthermore, there was an indistinguishable change in the amount of putative ex-siRNAs between the wild-type and Fgars2 mutant strains (Figure 4).

Although several studies have attempted to characterize microRNA-like RNAs (milRNAs), little is known about the functional milRNA in filamentous fungi (Torres-Martínez and Ruiz-Vázquez, 2017). Considering the conserved function of ARS2 during the miRNA biogenesis process shown in plants and mammals, we predicted milRNAs in F. graminearum based on in silico prediction tools that are commonly used for miRNA prediction in model organisms (Figure 5 and Supplementary Figure S3). Using the previously reported miRNA prediction pipeline (Hwang et al., 2013), 1095 miRNA candidates were obtained, reduced to 208 by manual curation, and finally grouped into 109 families in 112 precursors (Supplementary Table S3). No conserved milRNA candidates were identified in a search against the mature miRNAs in the miRBase (Kozomara and Griffiths-Jones, 2013) or even among previously predicted milRNA candidates of F. oxysporum or F. graminearum (Chen et al., 2014, 2015). Most identified milRNAs were 20 nt or 21 nt long with 5’-U, lacked RNAi (Fgdicer1 Fgdicer2 and Fgago1 Fgago2) greatly abolished milRNA production (Figure 5A). Intriguingly, the predicted milRNAs that were 20 nt and 21 nt long were exclusively produced by Fgdicer2 and Fgago1, while both Fgdicers and Fgago2 were important for global sRNA production in F. graminearum (Figures 5B,C).

We further determined the amounts of milRNAs candidates in the wild-type and Fgars2 strains. Similar to the siRNAs, the amounts of putative milRNAs were unaffected by the disruption of Fgars2 compared to the wild type (Figure 4). Taken together, our results suggest that Fgars2 is not closely involved in the milRNA and siRNA production in F. graminearum.

Subcellular Localization of Fgars2

Because two nuclear localization signal sequences were observed at the Fgars2 N-terminus, we examined the subcellular localization of Fgars2, which might provide insights to understand its biological functions. To achieve this, we generated
strains overexpressing the FgARS2 gene fused with GFP, resulting in DCB104 strains (Supplementary Figure S4). It is noteworthy that the DCB104 strains possessed indistinguishable phenotypes compared to the wild-type strain (data not shown). To confirm the nuclear localization of FgArs2-GFP, the DCB104r (FgARS2:P<sub>EF1a</sub>-GFP-FgARS2-GEN; hH1-RFP-GEN) strains were generated via outcrossing of the mat1r (Son et al., 2011a) and DCB104 strains. Localization was observed in cultures grown on CM (for hyphae), YMA (for conidia), and carrot agar (for ascospores). FgArs2-GFP in the DCB104r strains co-localized with hH1-RFP (Figure 6) in all of the observed developmental stages, validating the distinct nuclear localization of FgArs2.

**FgArs2 and Cap Binding Complex Proteins Are Important for Pathogenesis**

Since the Ars2 protein physically interacts with the CBC proteins in model organisms, we questioned whether the interaction is still conserved in *F. graminearum*. In model eukaryotes, the CBC consists of 20 and 80 kDa proteins, denominated Cbp20 (cap-binding protein 20) and Cbp80, respectively (Gonatopoulos-Pournatzis and Cowling, 2014). The plant Cbp20/80 protein sequences were utilized to identify CBC orthologs in *F. graminearum* using the BLAST algorithm\(^1\). There is only one orthologous gene for each

\(^1\)http://fungidb.org
CBC subunit found in the F. graminearum genome, assigned as FgCBP20 (locus ID: FGSG_05488) and 80 (locus ID: FGSG_07328), respectively. Accordingly, FgCbp20 encoded a 173-amino-acid (aa) protein, and FgCbp80 was a long protein with 798 aa. We used the yeast two-hybrid system to validate the interaction between FgCbp20, FgCbp80, and FgArs2.

Our results of yeast two-hybrid assay revealed that FgCbp80 physically interacted with FgCbp20, suggesting that they form the heterodimeric cap-binding complex (Figure 7). FgArs2 interacted with both FgCbp80 and FgCbp20, although the strength of the FgArs2-FgCbp20 interaction was weaker than that of FgArs2-FgCbp80, which is consistent with those in plants and mammals. These results suggest that the FgArs2-FgCbp80-FgCbp20 complex may involve some conserved functions or might play novel functions in filamentous fungi, including F. graminearum.

To further explore the genetic and biological functions of the module FgArs2-FgCbp80-FgCbp20 in F. graminearum, we constructed and characterized the single deletion mutants of FgCBP80 and FgCBP20 in detail (Supplementary Table S1 and Supplementary Figure S5). Similar to FgARS2, FgCBP80 and FgCBP20 were not essential for somatic growth in this fungus. In addition, single deletions of both FgCBP80 and FgCBP20 produced morphologically indistinguishable phenotypes compared to the wild-type strain (data not shown). Intriguingly,
FIGURE 6 | Subcellular localization of FgArs2. A DCB104r strain containing FgArs2-GFP and hH1-RFP was grown on CM (mycelia), carrot agar medium (ascospores), or YMA (conidia) for microscopic observation. DIC, differential interference contrast. Scale bar = 10 µm.

FIGURE 7 | Yeast two-hybrid analysis of the interaction between FgArs2 and cap binding complex. The pairs of plasmids pDH1-ARS2/pAI-Alg5 and pDH1-ARS2/pDL2-Alg5 as well as pDH1-CBP20/pAI-Alg5 and pDH1-CBP20/pDL2-Alg5 served as positive and negative controls, respectively. The growth of transformed yeast was assayed on synthetic dextrose medium lacking His, Leu, Trp, and Ade. Columns in each panel display the serial decimal dilution.

both the Fgcbp80 and Fgcbp20 mutants drastically attenuated pathogenicity on wheat heads, although these defects were milder than that of Fgars2 (Figure 8). These results strongly fulfilled the hypothesis that CBC proteins are required for the pathogenesis of F. graminearum and suggested that the robust physical interaction of FgArs2 with FgCbp80-FgCbp20 complex might be specifically required for virulence in this fungus.

DISCUSSION

We identified sixteen putative TFs involved in DDR in our previous study and further demonstrated that the cell cycle regulation and DDR were closely related to sexual reproduction and virulence in F. graminearum (Min et al., 2014; Son et al., 2016). In this study, we uncovered the novel cellular function of the ARS2 ortholog in DDR as well as pathogenesis in the plant pathogenic fungus F. graminearum. Our results showed its diverse roles in the fungal developmental processes, while the molecular function of FgArs2 in the sRNA production did not conserved in F. graminearum. In addition, we provided genetic evidences demonstrating that FgArs2 physically interacts with CBC proteins to form a robust CBCA complex, and the stability of this tertiary complex governs fungal pathogenesis.

ARS2 takes part in global post transcriptional regulation and therefore ARS2 orthologs have been shown to be essential in fission yeast, plants, insects, and mammals (Golling et al., 2002; Oh et al., 2003; Amsterdam et al., 2004; Lobbes et al., 2006; Wilson et al., 2008; Kim et al., 2010). In contrast, Fgars2 deletion mutants were viable in F. graminearum (Son et al., 2011c). Our in silico analysis indicated that FgArs2 preserved the functional domains, and the deletion of FgARS2 resulted in pleiotropic defects in the fungal developmental processes of F. graminearum (Figures 1, 3). HsARS2 knockdown disrupted the replication-dependent histone mRNA processing and caused a reduction in histone expression, leading to disruption of the cell cycle (Kiriyama et al., 2009; Gruber et al., 2012) and AtSE has also been shown to control leaf development, meristem activity, and inflorescence architecture in A. thaliana (Machida et al., 2011). Combined, Ars2 likely plays conserved roles in cell cycle progression and proliferation from mammals, insects, and plants to fungi and other eukaryotes.

The growth rate of Fgars2 appears to be sensitive during treatment with DNA-damaging agents, and the transcript level of Fgars2 is increased after treatment with DNA-damaging agent (Son et al., 2016). Our results revealed that Fgars2 was...
irregular in nuclear division, and its conidial germination was significantly retarded, presumably due to its hypersensitive phenotypes during DNA-damage stress (Figure 1). Similarly, myogenic cells deficient in HsARS2 resulted in a slow cell cycle and spent an increased time in the S phase (O’sullivan et al., 2015). Thus, our findings further deciphered the important role of Ars2 in the proper mitotic cell division and DDR in filamentous fungi and possibly in other eukaryotes.

The physical interaction between ARS2 and CBC proteins through the 7-methylguanosine cap structure of nuclear RNA polymerase II transcripts has been extensively documented (Laubinger et al., 2008; Gruber et al., 2009; Sabin et al., 2009). In addition, the vast majority of ARS2 functions involved in its interactions with CBP20/80 is conserved in model eukaryotes (O’sullivan and Howard, 2016). In our study, FgArs2 physically established a robust interaction with both FgCbp80 and FgCbp20 proteins, indicating that this complex structure is evolutionally conserved in eukaryotes. Similarly, NcCbp20 and NcCbp80 directly interact to form the CBC, and this heterodimeric complex is predominantly nuclear in N. crassa (Decker et al., 2017). Genetic observation also showed that FgCBP20 and 80 were not essential for somatic growth as previously reported in N. crassa (Decker et al., 2017). While the disruption of NcCBC causes a drastic reduction in ascospore production, the single deletion of FgCBP20 or FgCBP80 resulted in morphologically indistinguishable phenotypes compared to the wild-type strain. Instead, our results clarified that as is the case with Fgars2, both Fgcbp20 and Fgcbp80 mutants attenuated their virulence on wheat heads, suggesting a role of the CBCA theme in fungal pathogenesis.

The Fgars2 mutants produced reduced amounts of trichothecene, however, they grew similarly under oxidative stress condition compared to the wild-type (Son et al., 2011c). Moreover, Fgars2 strains showed sensitive phenotypes against several DNA-damaging agents, HU and BLM (Figures 1F,G). Because HU inhibits DNA replication by inhibiting ribonucleotide reductase and BLM induces DNA double-strand breaks, we suspected that FgArs2 is involved in the DDR in F. graminearum (Ehrenberg and Reichard, 1972; Stubbe and Kozarich, 1987). Given that plant defenses can also elicit damages to DNA, there should be a link between virulence and DDR. Collectively, based on our achieved data, the attenuated virulence of Fgars2 mutant is likely regulated by combination of different factors, including reduced trichothecene production, DDR defects, and dysregulation of CBCA theme. Further investigations would be necessary to decode this sophisticated interaction.

In last years, studies on model eukaryotes revealed that ARS2 is required for proper miRNA biogenesis. AtSE acts as a scaffold to stimulate the efficiency and accuracy of pri-miRNA processing by the AtDCL-1/AtHYL-1 complex (Dong et al., 2008; Machida et al., 2011); therefore, the disruption of AtSE resulted in decreased levels of mature miRNAs and increased levels of pri-miRNA transcripts (Grigg et al., 2005; Lobbes et al., 2006; Yang et al., 2006). In mammalian systems, the decreased levels of mature miR-21, let-7 miRNAs, as well as primary miR-21 transcripts, were the consequence of HsARS2 depletion (Gruber et al., 2009). Both DmARS2 and DmCBC are required for miRNA- and siRNA-mediated silencing as well as antiviral defense in D. melanogaster (Sabin et al., 2009).

Most eukaryotes except some yeast species possess conserved RNAi mechanism. While higher eukaryotes such as plants and animals have evolved a miRNA-mediated RNAi as an important post-transcriptional gene regulation mechanism, endogenous ex-siRNA-mediated transcriptional regulation play a crucial role in fungal development and stress responses in lower eukaryotic fungi such as a basal fungus Mucor circinelloides (Torres-Martinez and Ruiz-Vázquez, 2016). F. graminearum only utilize ex-siRNA-mediated RNAi during late stages of sexual development, but not for other developmental stages or genome defenses (Son et al., 2017). This study uncovered that FgArs2 is critically required for ascosporogenesis but dispensable for the production of putative miRNAs and siRNAs. In line with this, some functions of Ars2, such as the role for miRNA or
siRNA production, seem to have been lost in *F. graminearum*. However, Ars2-dependent cellular processes have been evolved to govern sexual reproduction, DDR, and virulence in this fungus. In particular, formation of Ars2-CBC complex was revealed to be important for fungal virulence.

**CONCLUSION**

Our data proved that *FgARS2* is required for diverse roles in DDR, fungal progressions, and pathogenesis, but not for sRNA production in *F. graminearum*. In addition, *FgArs2* physically interacted with the CBC, and this robust theme, together with trichothecene and DDR, seemed to be essential factors governing fungal pathogenesis in *F. graminearum*. Taken together, our study provides new insights into the diverse roles of ARS2 ortholog in pathogenic fungi during fungal development and pathogenesis.

**DATA AVAILABILITY STATEMENT**

All datasets generated for this study are included in the manuscript/Supplementary Files.

**AUTHOR CONTRIBUTIONS**

D-CB, J-EK, Y-WL, J-AS, and HS conceived and designed the experiments. D-CB, J-EK, JS, GC, and HS performed the experiments. D-CB, J-EK, JL, and J-AS analyzed the data. GC, Y-WL, J-AS, and HS contributed the reagents, materials, and analysis tools. D-CB, J-EK, JS, Y-WL, J-AS, and HS wrote the manuscript. All authors read and approved the final version of manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2019.02326/full#supplementary-material

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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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