The Early Asexual Development Regulator *fluG* Codes for a Putative Bifunctional Enzyme

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*FluG* is a long recognized early regulator of asexual development in *Aspergillus nidulans*. *fluG* null mutants show profuse aerial growth and no conidial production. Initial studies reported sequence homology of FluG with a prokaryotic type I glutamine synthetase, but catalytic activity has not been demonstrated. In this study, we conducted an in-depth analysis of the FluG sequence, which revealed a single polypeptide containing a putative N-terminal amidohydrolase region linked to a putative C-terminal γ-glutamyl ligase region. Each region corresponded, separately and completely, to respective single function bacterial enzymes. Separate expression of these regions confirmed that the C-terminal region was essential for asexual development. The N-terminal region alone did not support conidial development, but contributed to increased conidial production under high nutrient availability. Point mutations directed at respective key catalytic residues in each region demonstrated that they were essential for biological function. Moreover, the substitution of the N- and C-terminal regions with homologs from *Lactobacillus paracasei* and *Pseudomonas aeruginosa*, respectively, maintained functionality, albeit with altered characteristics. Taken together, the results lead us to conclude that FluG is a bifunctional enzyme that participates in an as yet unidentified metabolic or signaling pathway involving a γ-glutamylated intermediate that contributes to developmental fate.

**Keywords:** FluG, *Aspergillus nidulans*, enzyme, development, amidohydrolase, γ-glutamyl ligase

**INTRODUCTION**

Surface cultures of the model ascomycete *Aspergillus nidulans*, present a complex developmental pattern which combines vegetative hyphae with asexual (Adams et al., 1998) and sexual structures (Dyer and O’Gorman, 2012). This pattern is modulated in response to environmental factors, such as the composition of the substrate, exposure to the gas phase and light.

The genetic regulation that governs the primal transition from vegetative hyphae to asexual development has been examined in considerable detail. A set of genes, which are expressed in vegetative hyphae, are required to initiate the morphogenetic process. They are collectively known as Upstream Developmental Activators (UDAs). These, in turn, activate a second set of the genes, known as the Central Developmental Pathway (CDP), which are only expressed in conidia-bearing structures (Yu et al., 2006).
One of the earliest acting UDA factors is fluG (AN4819). Its deletion results in colonies which fail to produce conidiophores and accumulate aerial vegetative hyphae resulting in a raised mycelial morphology, commonly known as fluffy (Lee and Adams, 1994). In addition, ΔfluG mutants show defects in autolysis (Emri et al., 2005) and secretion (Wang et al., 2015).

A remarkable feature of FluG signaling involves its extracellular transmission. A wild type colony can induce the sporulation of an adjacent fluG null mutant separated by a membrane (Lee and Adams, 1994). The search for the signal conducted by Rodriguez-Urra et al. (2012) identified an adduct of two excreted secondary metabolites [diorcinol and dehydroactinol (DHO)] that was capable of inducing a partial sporulation response when added onto a fluG null mutant colony. Interestingly, Márquez-Fernández et al. (2007) had earlier discovered that the deletion of a phosphopantetheinyl transferase (PPTase) cfwA/npgA, an enzyme that acting upstream of DHO biosynthesis, exhibited a phenotype with severely reduced growth and asexual development. Rodriguez-Urra et al. (2012) pointed out, however, that an earlier report by Nielsen et al. (2011) described the phenotype of a null mutant of the gene coding for the synthesis of the austinoI (ausA, AN8383), the precursor in the DHO biosynthesis, as conidial. Hence, a clarification on the connection between FluG signaling and DHO-induced conidiation is pending.

Early studies on the FluG protein showed that it is expressed in the cytosol throughout all stages of development, and that the C-terminal region (aa387 to aa865) is necessary for induction of conidiophore development. Interestingly, this region showed sequence similarity with prokaryotic type I glutamine synthetase (GS) (Lee and Adams, 1994; D’Souza et al., 2001). Later investigations showed, however, that this region did not display GS activity (Margelis et al., 2001). Other reports indicated that it resembles gene SCO6962 from Streptomyces coelicolor (Rexer et al., 2006) which has been attributed to a γ-glutamyl ligase (GGL) (Krysenko et al., 2017). The N-terminal region has no attributed functional role, but has been reported to bear similarity with an amidohydrolase (SCO6961) from Streptomyces coelicolor (Rexer et al., 2006).

Although fluG-like sequences (containing a N-terminal amidohydrolase and a GS) have been reported in the Aspergilli (Mah and Yu, 2006; Ogawa et al., 2010; Chang et al., 2012; Wang et al., 2015), they are also present in genomes of diverse fungal genera (Schumacher et al., 2015; Li et al., 2017), bacteria (Strong et al., 2003; Rexer et al., 2006) and higher plants, where the proteins have attributed roles in nodulation and biotic stress signaling (Mathis et al., 1999, 2000; Trevaskis et al., 2002; Doskočilová et al., 2011; Silva et al., 2015).

In this investigation, we confirm that an ausA null mutant has a conidiating phenotype, thus showing that FluG-related signaling is not directly connected to DHO-induced conidiation. In addition, an updated structural analysis of the FluG protein revealed that the N-terminal region presented structural similarity to a prokaryotic amidohydrolase from Lactobacillus paracasei, and the C-terminal region presented structural similarity to a γ-glutamyl aromatic monoamine ligase from Pseudomonas aeruginosa. Alanine substitutions of the predicted key catalytic residues in each region yielded loss of function phenotypes which were comparable to those of the respective null mutants. Finally, the replacement of both N- and C-terminal regions by their respective homologs, yielded functional phenotypes. Our findings situate FluG as a putative bifunctional enzyme that may regulate the balance between vegetative growth and asexual development through the levels of a γ-glutamylated metabolite.

**MATERIALS AND METHODS**

**Fungal Strains and Culture Conditions**

The strains used are shown in **Supplementary Table 1**. Wild type (veA+) strains were used for FluG mutagenesis and phenotypic analysis. For the analysis of phenotypic complementation experiments, and ausA-related assays, veA1 strains were used, in order to establish comparisons with previously published experiments (Lee and Adams, 1994; Yager et al., 1998; D’Souza et al., 2001; Nielsen et al., 2011). Strains were cultured as previously described by Pontecorvo et al. (1953) in Aspergillus minimal medium (MMA), using trace elements as described by Käfer (1965), or complete medium (CMA; MMA + 5 g/L yeast extract) with a pH 6.8. Ammonium tartrate (5 mM) and D-glucose (2% w/v) were added as nitrogen and carbon sources, respectively, in solid or liquid form with the appropriate supplements. All cultures where incubated at 37°C in white light (Sera DayLight Brilliant T8 15 W/m²; light source at 30 cm from the cultures). Phenotypic complementation experiments were carried out in CMA medium, incubating for a further 72 h after the strains contacted each other (Lee and Adams, 1994). In fluorescence microscopy experiments, strains were grown in adequately supplemented MMA containing 0.1% D-glucose, 5 mM ammonium tartrate and 25 mM monobasic sodium phosphate, similar to watch minimal medium (WMM) (PénaIva, 2005).

**Generation of Mutant, Tagged, and Overexpression Strains**

The oligonucleotides used are listed in **Supplementary Table 2**. Genomic cassettes were amplified through the fusion-PCR technique (Yang et al., 2004; Korbie and Mattick, 2008). The A. nidulans transformation technique used was an adaptation of the protoplast generation procedure described by Szewczyk et al. (2006), followed by the transformation protocol described by Tilburn et al. (1983). The strains coding for separate regions of FluG, point mutants of both regions and homologous regions of Lactobacillus paracasei and Pseudomonas aeruginosa genes were generated by transforming protoplasts of a ΔfluG strain with the genomic cassettes bearing the desired modifications plus the 5′- and 3′-UTR regions. The selection of transformants was done using 5-fluoroorotic acid monohydrate (5-FOA; 2 mg/mL; Apollo Scientific, Stockport, United Kingdom).

N- and C-terminally GFP-tagged fluG strains were obtained using the method for GFP-tagging described by Yang et al. (2004). The strain carrying the inducible alcA expression promoter of the N-terminally GFP-tagged fluG was constructed.
using plasmid pNT5 (Takeshita et al., 2008). 

\[ \text{alcAp}::\text{GFP}::\text{fluG} \]

expressing plasmid was obtained by cloning fluG (PCR amplified with oligonucleotides AscI-fluG(–ATG)–F and BamHI-fluG–R, 1 kb on the N-terminal sequence of fluG with no starting ATG sequence; see Supplementary Table 2) into the AscI and BamHI sites of pNT5, yielding to pM12. The plasmid was then transformed into the wild type (BD824) strain. For localization studies, strains coexpressing the GFP-tagged fluG under native or inducible promoter and the histone H1 (hhoa) fused C-terminally to mRFP (mRFP; red fluorescent protein) and mCherry (mCh; cherry red) as nuclear markers, respectively, were constructed (Markina-Iñarrairaegui et al., 2011).

The homologous recombination of the constructs was confirmed by Southern blots. This experiment, using 5′- and 3′-UTR regions as probes, was done as described previously (Garzia et al., 2009). All the mutant strains were sequenced to prove there were no undesired mutations.

**Spore and Dry Cell Mass Measurements**

The procedure of obtaining spore and dry cell mass (DCM) ratios was as follows: conidia counts (millions/cm²) were obtained after culturing strains in MMA and CMA at 37°C for 72 h. Colony areas were measured using Digimizer Version 4.6.1 (MedCalc Software bvba). Spores were collected by excising the entire colony, soaking the agar in Tween 0.02% (diluted from Tween 20), vortexing twice for 1 min and quantified using a hemocytometer. DCM (mg/cm²) from each strain was obtained by growing colonies on a Spectra/Per® 6 Dialysis Membrane with a 3.5 kDa molecular weight cut off (Spectrum Laboratories) and culturing at 37°C for 72 h. The mycelium was then removed from the membrane and dried at 80°C overnight.

The values obtained with the WT and the ΔfluG strains were used as references to analyze mutant strains.

**Statistical Analysis of the Results**

Quantitative results were obtained as the mean with standard deviation of at least three independent counts, each consisting of three or four biological replicates per strain and condition. The results were subjected to the Grubbs’ test (Grubbs, 1969) using the R software version 3.4.1 (R Core Team, 2008) in order to detect and discard outliers with a confidence level of 95%. A PERMANOVA pair-wise test using an Euclidean distance concentration and purity were calculated using a NanoDropTM 2000c Spectrophotometer (Thermo Fischer Scientific, Waltham, MA, United States). All samples were verified by electrophoresis using a 1.2% (w/v) agarose gel.

**Biinformatics**

The fluG DNA sequence (AN48192.2) was retrieved from Ensembl Fungi (Kersey et al., 2018) and homologs in the Aspergilli were identified by Fungal Compara (in Ensembl Fungi). Multiple sequence alignments were performed using the Clustal Omega application provided by EBI (Li et al., 2015). Alignments visualization and analysis were performed with Jalview version 2.10.3b1 (Waterhouse et al., 2009) or GeneDoc version 2.6.002 (Nicholas et al., 1997). FluG protein residue conservation scores among all FluG homologs in the Aspergilli were obtained using the Jensen–Shannon divergence (Capra and Singh, 2007).

Identity among all FluG proteins was obtained as an average value of all FluG sequence identities in the Aspergilli compared with the *Aspergillus nidulans* FluG.

**Biinformatic investigations of the FluG sequence were made using the following predictors:**

- **SignalIP 4.1** (Nielsen, 2017) for signal-peptide prediction;
- **NLStradamus** (Nguyen Ba et al., 2009) and NetNES 1.1 (la Cour et al., 2004) for nuclear localization and export signals prediction, respectively;
- **TMHMM server** v2 (Sonnhhammer et al., 1998) for prediction of transmembrane helices in protein; epstfind (Rechsteiner and Rogers, 1996) to find PEST motifs as potential proteolytic cleavage sites.

Crystal structure templates for each region were identified using SWISS-MODEL (Biasini et al., 2014), and those with high QMEAN values were further analyzed (Benkert et al., 2009). The highest scoring structures in terms of sequence identity and coverage were used as templates for FluG structural modeling. Protein structures were obtained from the Protein Data Bank (PDB) (Berman et al., 2002). Prior to modeling, the X-ray data was preprocessed and curated (e.g., removal of water, addition of explicit hydrogens) using the Protein Preparation Wizard from the Schrödinger Suite, version 2017-1 (Sastry et al., 2013). Missing side chains were generated using Prime (Jacobson et al., 2004) while the protonation states of each side chain were generated using EPIK at pH = 7 (Shelley et al., 2007). Protein minimization was performed using the OPLS3 force field (Shivakumar et al., 2010; Harder et al., 2016). Homology models were built using program Prime based on the curated reference crystal structure and applying the SWISS-MODEL alignment for each region. The structural and spatial distribution consistency of the FluG site-directed mutant models was validated overlapping these models against the wild type FluG model, using Quick Align and Superimpose tools.

**RNA Isolation**

RNA was isolated from the mycelium of strains cultivated in a flask for 18 h at 37°C. Mycelia was collected by filtration through Miracloth (Calbiochem®), squeezed to dry and frozen in liquid nitrogen. Total RNA was isolated using the NucleoSpin® RNA Kit (Macherey-Nagel GmbH & Co. KG) and RNA concentration and purity were calculated using a NanoDropTM 2000c Spectrophotometer (Thermo Fischer Scientific, Waltham, MA, United States). All samples were verified by electrophoresis using a 1.2% (w/v) agarose gel.

**Quantitative RT-PCR**

The oligonucleotides employed for quantitative RT-PCR (qPCR) experiments are detailed in Supplementary Table 2. Briefly, cDNA from each investigated sample was synthesized using PrimeScriptTM RT reagent Kit (Perfect Real Time) (Takara Bio USA Inc.) from 500 ng of total RNA, following manufacturer’s instructions. The final 20 μL PCR reaction included 1.5 μL of 1:2 diluted cDNA as template, 4 μL of 5x PyroTaq EvaGreen® qPCR Mix Plus with ROX (Cultek Molecular Bioline, Madrid, Spain), and transcript-specific forward and reverse oligonucleotides at a 0.5 μM final concentration. Real-time PCR was carried out in a 7500 Real-Time PCR System (Applied Biosystems®) according to the manufacturer’s directions. The reaction consisted of 15 min at
95°C of initial denaturation, and 40 cycles of DNA amplification (15 s at 95°C of denaturation, 15 s at 60°C of annealing and 30 s at 72°C of elongation). After each PCR, we performed a melting curve analysis to confirm the specific amplification of a single DNA segments and the absence of non-specific amplified DNA.

The fluorescent signal obtained for each gene was normalized to that obtained with the β-tubulin gene (benA) to correct for sampling errors.

**Protein Isolation and Western Blot**
Protein extraction from lyophilized samples was performed by the alkaline-lysis extraction, using lysis buffer (0.2 M NaOH, 0.2% β-mercaptoethanol), as described by Hervás-Aguilar and Peñalva (2010).

Tagged protein expression was analyzed by Western blotting. Proteins were resolved in 8% SDS-polyacrylamide gels and electrotransferred to Immun-Blot® PVDF membranes by Trans-Blot® TurboTM Transfer System (Bio-Rad). Once transference was completed, Ponceau staining was used as validation for loading control. GFP-FluG was detected using mouse anti-GFP (1/5,000; Roche). Peroxidase-conjugated goat anti-mouse IgG immunoglobulin (1/4,000; Jackson ImmunoResearch Lab) cocktail was used as secondary antibody. Peroxidase activity was detected using ClarityTM Western ECL Substrate (Bio-Rad). Chemiluminescence was observed using a ChemiDocTM XRS+ System (Bio-Rad) and signal intensity was measured with Image LabTM version 5.2 software (Bio-Rad).

**Fluorescence Microscopy**
Conidia of strain BD1181 were cultured in uncoated glass-bottom dishes (Ibidi® GmbH, Germany; 2.5 mL of medium per well) for 16 h at 25°C (Peñalva, 2005). After this period, the medium was replaced with fresh medium supplemented with 100 mM L-threonine (Sigma) in place of D-glucose to induce alcA(p)::GFP::fluG expression.

Fluorescence images were acquired using a Zeiss Axios Observer Z1 inverted microscope equipped with a 63x Plan Apochromat 1.4 oil immersion Lens, Axiocam MRm Rev.3 camera, a Zeiss HXP 120C external light source for epifluorescence excitation and fitted with filter set 38HE for green fluorescence (Ex BP 470/40; FT 495; Em BP 525/50) and filter set 43HE for red fluorescence (Ex BP 545/25; FT 570; Em BP 605/70). Numerous samples were observed before taking representative images. Fluorescence levels were measured using Fiji software (Schindelin et al., 2012).

**RESULTS**
**DHO-Induced Asexual Development Is Independent of the FluG (UDA) Pathway**
In order to verify previous results by observation that an ausA null mutant strain is unaffected in asexual development (Nielsen et al., 2011), an ΔausA (veA1; BD688) strain was generated and phenotypically characterized in comparison to a parent wild type (WT, veA1; TN02A3) strain, as shown in Figure 1A. The values

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**FIGURE 1 | (A) Phenotypes and C/area (millions/cm²) ratios of the wild type (WT; TN02A3) (reference value) and ausA null mutant (ΔausA; BD688) in MMA and CMA. Scale bar = 1 cm. (B) Phenotypic complementation of a WT and ΔausA mutant in contact with a fluG null (ΔfluG; TTA127.4) and a cfwA2 (CRO1) mutant. White arrows (✓) indicate the contact area where yellow conidia are visible. Scale bar = 5 mm.**
for C and DCM for each strain are provided in Supplementary Table 4. In MMA and CMA, the C-values recorded for both WT and ΔausA mutant strains were statistically similar. These results indicated that asexual development was not significantly affected in the ΔausA mutant.

In addition, a phenotypic complementation experiment confirmed that both WT (veA1) and ΔausA (veA1) strains were able to efficiently complement the sporulation defects of a ΔfluG (veA1; TTA127.4) and cfwA2 (veA1; CRO1) mutant, as can be observed in Figure 1B. Thus, regardless of the induction effect caused by DHO on asexual development, it cannot be attributed as part of the FluG-dependent UDA pathway. This led to the conclusion that an in-depth study of the FluG protein was required to gain insight on its function.

The FluG Sequence Codes for a Putative Bifunctional Enzyme
A protein BLAST of FluG homologs revealed that they were present in all 14 sequenced species of the Aspergilli (Figure 2), with a considerable degree of identity (70 ± 4%) (Supplementary Table 3). Earlier sequence analyses referred to in the introduction (Lee and Adams, 1994; Mathis et al., 1999) had indicated that the fluG sequence contained two regions which coded for two putative prokaryotic enzymes. We therefore proceeded to undertake a protein sequence and structure analysis using current applications and databases. A search of available crystal structures using the sequence of the FluG protein showed that the N-terminal region bears similarity with a range of amidohydrolases. The highest sequence coverage and structural homology score was obtained with a putative metal-dependent hydrolase of the TIM-barrel fold (pdb entry 2QPX, coded by gene LSEI_0440) present in Lactobacillus paracasei ATCC 334. The 2QPX crystal structure corresponded to the 4–406 amino acid region of FluG, with an identity of 25% and a similarity of 45%. Conjointly, the C-terminal region exhibited the highest scores with a GSI (pdb entry 4LNN, coded by gene flna) from Bacillus subtilis (strain 168) (Murray et al., 2013) and a GGL (pdb entry 4HPP, coded by gene PA5508) from Pseudomonas aeruginosa PAO1 (Ladner et al., 2012). The 4LNN crystal structure coincided with the 429–865 amino acid region of FluG with a sequence identity of 24% and a similarity of 38%, whereas the 4HPP crystal structure covered the 443–865 amino acid region of FluG, with sequence identity of 22% and a similarity of 33% (Table 1).

In order to uncover new features of the protein, we conducted additional studies. Sequence analyses performed using different predictors indicated that the sequence contained neither signal peptides, nuclear localization or export signals nor putative transmembrane domains. However, the PEST predictor marked two potential PEST domains in the C-terminal region (aa571 to aa590 with a score of 14.62 and aa529 to aa541 with a score of 6.84, respectively). Analyses of the genomic DNA and mRNA of fluG sequence revealed two predicted introns between nucleotides 1126–1176, and 2420–2472. A previous RNA-seq study (Garzia et al., 2013) reported a single mRNA transcript of 2,598 nucleotides, in which both introns are spliced out.

Localization of FluG in vivo
A GFP-tagged version of the protein was used for in vivo localization. Strains tagged in the N- and C-terminal regions of the protein were grown in MMA and CMA for 72 h, and their phenotype observed (Supplementary Figure 1). The strain tagged in the C-terminal presented a fluffy phenotype, indicative of loss of function, while the N-terminally tagged strain presented a wild type phenotype. Hence, the latter was chosen to perform localization experiments. In order to determine the incubation time at which the protein was sufficiently expressed for observation, the pattern of expression in liquid vegetative incubation time at which the protein was expressed was controlled under the alcA promoter, a shift to a medium containing threonine was conducted at time zero. The results revealed that GFP::FluG protein could only be detected in the strain alcA(p)::GFP::fluG 2 h after medium shift to threonine containing medium (Figure 3A).

Consistent with the results obtained, no GFP–FluG fluorescence was detected in the strain bearing the native fluG promoter (not shown). In the strain with the alcA inducible promoter, GFP–FluG was localized 2 h after the medium shift in the cytoplasm throughout the length of the hypha. Nuclei were stained by tagging histone H1 tagged with mCherry fluorescence, but no GFP signal was detected in the nuclear compartment as shown in Figure 3B. In addition, the protein appeared to be absent from the lumen of vacuoles (white arrows). Taken together, our microscopic observations indicated that FluG was evenly distributed throughout the cytoplasm of vegetative cells.

![Figure 2](https://example.com/image2.png)

**Figure 2** | Protein sequence conservation score among all FluG homologs in the Aspergilli. The catalytic residues for both regions are labeled (residue numbers are listed in Tables 2, 3). These residues are conserved in all Aspergillus species, and they are all located in highly conserved areas. Red asterisks mark the residues that were mutated.

**Table 1** | Crystal structure homologs to the N- and C-terminal region.

|          | N-terminal region | C-terminal region |
|----------|------------------|------------------|
| PDB entry| 2QPX             | 4LNN             |
| Coverage (aa) | 4–406           | 429–865          | 443–865          |
| Seq. identity (%) | 25              | 24               | 22               |
| Seq. similarity (%) | 43              | 38               | 33               |

The coverage is shown using the amino acid numbers of FluG. Sequence identity and amino acid positive matches were calculated by Maestro (Schrödinger Suite).
Computational Analysis of the N- and C-Terminal Regions

The closest homolog to the N-terminal region of FluG, 2QPX, was used as a template to model structural features. The residues shared by both sequences and their predicted functions are shown in Table 2. In 2QPX, residues H17, H19, and D317 coordinate to the first Zn$^{2+}$ (Mα), whilst H227 and H263 coordinate to the second Zn$^{2+}$ (Mβ). In addition, a lysine (K166) undergoes a post-translational carbamylation to bridge both Zn$^{2+}$ ions, and a tyrosine (Y171) is predicted to interact with the substrate in certain amidohydrolases. The N-terminal region of FluG conserves all the above-mentioned residues (H20, H22, K192, H264, H298, and D354), along with an appropriate spatial arrangement to bind and bridge both metals, together with the tyrosine (Y197) that could interact with the substrate (Figure 4). These residues are conserved through all the Aspergilli FluG homologs, as marked in Figure 2.

On the other hand, the GSI crystal structure (4LNN) contains a glutamate residue (E304) that abstracts a proton from the transition state to form glutamine and ADP. It is worth noting, however, that the GGL crystal structure and the FluG (C-terminal) model each contain a tryptophan residue in the same position (W296 and W726, respectively), which cannot fulfill the function of glutamate. Furthermore, the conserved residue D53 in GSI that removes a proton from the ammonium to the first Zn$^{2+}$ (Mα), whilst H227 and H263 coordinate to the second Zn$^{2+}$ (Mβ). In addition, a lysine (K166) undergoes a post-translational carbamylation to bridge both Zn$^{2+}$ ions, and a tyrosine (Y171) is predicted to interact with the substrate in certain amidohydrolases. The N-terminal region of FluG conserves all the above-mentioned residues (H20, H22, K192, H264, H298, and D354), along with an appropriate spatial arrangement to bind and bridge both metals, together with the tyrosine (Y197) that could interact with the substrate (Figure 4). These residues are conserved through all the Aspergilli FluG homologs, as marked in Figure 2.

Structural alignment of the C-terminal region using the crystal structures of a GSI (4LNN) and a GGL (4HPP), also showed homology in key catalytic residues (Table 3). The amino acids that coordinate to the metal ions in both GGL (one Mg$^{2+}$ [Mα] coordinated to E133, E180, and E187) (Figure 5A) and GSI (two Mg$^{2+}$; Mα coordinated to E134, E189 and E196, and Mβ coordinated to E132, H245, and E333) (Figure 5B) were structurally conserved in the C-terminal region of FluG (E566, E626, E633, H682, and E752). Moreover, the residues that bind to the glutamate ligand (R290 in 4HPP; R298 in 4LNN) and interact with ATP (H238, R308, and R313 in 4HPP; N247, R316, and R321 in 4LNN) were present in FluG (R720 for the glutamate; H682, R739, and R744 for the ATP, respectively). The residues listed in Table 3 are fully conserved in FluG for all the Aspergilli (Figure 2).

On the other hand, the GSI crystal structure (4LNN) contains a glutamate residue (E304) that abstracts a proton from the transition state to form glutamine and ADP. It is worth noting, however, that the GGL crystal structure and the FluG (C-terminal) model each contain a tryptophan residue in the same position (W296 and W726, respectively), which cannot fulfill the function of glutamate. Furthermore, the conserved residue D53 in GSI that removes a proton from the ammonium

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**TABLE 2** | Predicted catalytic residues in 2QPX, and their FluG equivalents.

| Residues | Predicted role |
|----------|----------------|
| 2QPX     | FluG           |
| H17      | H20 Coordinates with the Mα metal |
| H19      | H22 Coordinates with the Mα metal |
| K166     | K192 Once carbamylated (KCX), bridges both Mα and Mβ |
| Y171     | Y197 Coordinates with the substrate to aid the cleavage of the bond |
| H227     | H264 Coordinates with Mβ |
| H263     | H298 Coordinates with Mβ |
| D317     | D354 Coordinates with Mα |

Metals labeled as Mα and Mβ correspond to the first and the second Zn$^{2+}$ atoms. The lysine residue (K166) undergoes a post translational modification that results in a carbamylation.

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**FIGURE 4** | Structural alignment of 2QPX (green) refined model, with the FluG modeled structure (gray), based on 2QPX template. The Mα is coordinated to H20, H22, K192, and D354; the Mβ is coordinated to K192, H264, and H298. A water molecule is coordinated to both metals and D354 (blue dashed line).
ion to create ammonia is aligned with V42 and S481 in GGL and FluG, respectively; neither of the aforementioned amino acids can perform the role of a GSI.

An important aspect that differentiates GGL and GSI enzymes is the fact GGL forms hexameric rings and GSI enzymes form two hexameric subunits that assemble into a dodecameric unit. The formation of the dodecamer depends on the conformation of the C-terminal end of these enzymes adopts. Structural alignments performed with SWISSMODEL using 4LNN and 4HPP crystals revealed that the C-terminal end of FluG exhibited higher QMEAN values with the 4HPP crystal in this region (not shown).

Phenotypic Characterization of WT and ΔfluG Mutant

The WT and ΔfluG strains were established as references for comparisons with strains containing modifications, and their phenotypes were characterized in minimal (MMA) and complete medium (CMA). The conidial production in millions per centimeter squared of colony surface (C) in 72 h old colonies of both strains is presented in Figure 6A. Numerical values for C and DCM are separately provided in Supplementary Table 5. The WT colonies presented a C-value of 4.1 in MMA, distinctly greater than the ΔfluG strain, 0.02, but significantly lower DCM values. In CMA, the WT strain presented increased C and DCM indices with respect to MMA. The ΔfluG strain showed even lower conidial counts than in MMA and increased DCM values. Taken together, the results indicate that, under nutrient limitation (MMA), the wild type strain distributes resources between conidiation and growth, while the ΔfluG mutant destines resources exclusively to growth. However, under nutrient sufficiency (CMA), both strains attained apparently maximum growth levels, and accentuated their differences in conidial production. It is due to these clearly discernible differences, that the C-value was considered as the principal phenotypic indicator of functionality when examining genetically modified strains.

The N- and C-Terminal Regions of FluG Perform Distinct Functions

Each of the two FluG regions was expressed separately under the control of the native promoter. The results shown in Figure 6A indicate that in MMA, the mutant expressing only
the C-terminal region presented a C-value significantly greater to that of the WT strain, while in CMA, it was one-fifth of the WT C-value. These results indicate that the C-terminal region was capable of fulfilling the role of the complete FluG polypeptide, but the absence of the N-terminal region limited its efficacy in CMA.

The strain exclusively expressing the N-terminal region, presented a C-value which was significantly similar to that of the ΔfluG mutant in MMA, but no so in CMA media. These results support the interpretation that the C-terminal region performs the core function of the protein in asexual development, but that the N-terminal region may provide a supporting function under conditions of high nutrient availability.

A strain expressing both N- and C-terminal regions as two separate polypeptides, each under the control of the native promoter of fluG was analyzed (N+C separated) (Figure 6A). The recorded C-value in MMA was significantly similar to that recorded for the WT and C-terminal region strains. In CMA, the value was at an intermediate point between both strains (Supplementary Table 5). These results not only indicate that both regions are functional when separately expressed, but support the view that the N-terminal region performs a relevant supportive role in conidial production, especially when both regions are expressed in the same polypeptide and under nutrient rich conditions.

In order to ascertain that the phenotypes observed were not affected by altered gene expression levels, strains were examined by qPCR and compared to that of the WT type strain, as shown in Supplementary Figure 2. With the exception of the ΔfluG mutant, all mutant strains presented transcript levels equal or higher than those observed in the WT strain.
Targeted Amino Acid Substitutions Affect Biological Function

Previously reported mutations on residues that were homologous to H20, H22, and D354 in the N-terminal region of FluG had been shown to be essential for amidohydrolase activity (Kim and Kim, 1998; Li et al., 2006; Lohkamp et al., 2006; Nguyen et al., 2009). Therefore, double (H20–H22), single (D354), and triple (H20–H22–D354) alanine substitution mutants were constructed. A similar strategy was followed for the C-terminal region, where residues E566, E626, H682, R720, R739, R744, and E752 were chosen based on previous reports (Kurihara et al., 2008; Ladner et al., 2012; Murray et al., 2013). The targeted residues are marked with asterisks in Figure 2. All these amino acid substitutions tested negative for propensity to disrupt structure using homology modeling (see “Materials and Methods” section). The observed phenotypes for the full set of mutants are shown in Figure 6B.

All the strains carrying mutations in the N-terminal region exhibited significantly lower C-values with respect to the WT reference in both media. In MMA, the C-values obtained for mutants H20–H22, D354, and H20–H22–D354 were significantly similar. In CMA, however, significant differences between the H20–H22, D354, and H20–H22–D354 mutants were observed with the triple mutant showing the maximum reduction in C-value (Figure 6B). This indicates an additive effect of the mutations, only under high nutrient availability.

In the C-terminal region, residues E566, E626, H682, R720, R739, R744, and E752 were separately substituted by alanine. All the mutant strains presented statistically lower C-values in MMA when compared to the WT. These low values were nevertheless significantly higher than those obtained in the ΔfluG strain, indicating that single amino acid modifications resulted in a partial inactivation of the activity. In CMA, the C-values were negligible for the majority of mutants in comparison to the WT strain. When compared to the ΔfluG strain, the majority of the mutants were statistically similar. Three mutants however (H682A, R739A, and R744A) were statistically different, indicating a partial inactivation (Figure 6B). Strain R739A, presented a diverging phenotype yielding statistically higher C-values in CMA when compared to the other mutants of this region.

The quantification of fluG expression in all the tested mutants indicated that the transcript was at least as abundant as in the WT strain (Supplementary Figure 2).

Taken together, the results obtained by these substitutions support the view that residues predicted to perform catalytic activity in both N- and C-terminal regions are involved in the biological functions earlier recognized for their corresponding regions.

Heterologous Expression of N- and C-Terminal Region Homologs

Given the sequence and structural homologies observed between both N- and C-terminal regions with those of Lactobacillus paracasei (LSEI_0440) and Pseudomonas aeruginosa (PA5508), we constructed strains in which the exogenous sequences were expressed under the control of the fluG promoter in combinations that included three protein chimeras (LSEI_0440+C-terminal; N-terminal+PA5508; LSEI_0440+PA5508) and one single polypeptide (PA5508).

The phenotypes of the mutant strains are shown in Figure 6C. The C-values obtained in the LSEI_0440+C-terminal and N-terminal+PA5508 strains were significantly higher than the WT strain in MMA. In the same conditions, the PA5508 and LSEI_0440+PA5508 exhibited statistically similar C-values when compared to the WT strain. On the other hand, the C-values of the three protein chimeras (LSEI_0440+C-terminal; N-terminal+PA5508; LSEI_0440+PA5508) were statistically similar to the WT reference in CMA. However, the strain expressing the PA5508 polypeptide alone exhibited a significantly lower value compared to the WT in the same medium.

The results clearly show that the protein encoded by gene LSEI_0440 could functionally replace the N-terminal region of FluG; similarly, the protein encoded by PA5508 fulfilled the role of the C-terminal region of FluG, either as a single polypeptide (PA5508 mutant) or fused to the N-terminal region of FluG (N-terminal+PA5508 chimera). In line with the aforementioned results, the fact that the chimera expressing a fusion protein of LSEI_440 and PA5508 exhibits a statistically comparable phenotype to a WT strain confirms that both bacterial enzymes can work jointly in replacing both FluG regions.

DISCUSSION

This investigation initially attempted to clarify the role of DHO as a conidiation-inducing secondary metabolite in relation to FluG function. Previous evidence that secondary metabolites (such as DHO) may play a role in the induction of sporulation (Márquez-Fernández et al., 2007; Rodriguez-Urra et al., 2012), was in conflict with observations describing a null ausA mutant, unable to synthesize DHO, as conidial (Nielsen et al., 2011). This investigation moreover showed that such mutants can produce and transmit the conidiation-inducing factor to an adjacent ΔfluG colony. Thus, DHO-induced conidiation is not functionally related to the FluG pathway. This scenario led us to conduct an updated analysis of the FluG sequence to obtain a molecular outlook on its function.

Preliminary bioinformatic approaches indicated that FluG comprises two putative catalytic units, connected in the same polypeptide by a linker region, which likely function as a bifunctional enzyme performing two independent reactions. In vivo fluorescence microscopy experiments confirmed that FluG is localized throughout the cytoplasm, indicating that the protein may participate in a metabolic process that is ubiquitous to the whole mycelium. However, considerable evidence shows that fungal mycelia are spatially and functionally structured. For example, in gene expression (Levin et al., 2007;
In this regard, the role of FluG might be hypothesized as the mycelium-wide biosynthesis of a metabolic product or signal specifically required for conidiophore biogenesis. Conidiophores are produced at specific points in the network, likely acting as metabolic sinks. This is in accordance with reported cytosolic streaming directed at the conidiophore (Bleichrodt et al., 2013). Thus, further molecular studies involving FluG function should take the mycelial context into consideration.

The C-terminal region was confirmed as necessary and sufficient for the induction of development and presented clear sequence similarities with both GSI and GGL enzyme sequences. Close examination of the differences between these two types of enzymes revealed that FluG more closely matches the latter (Kurihara et al., 2008; Ladner et al., 2012; Takeo et al., 2013). To address this point, a site-directed mutational approach was planned to determine the importance of the predicted catalytic residues present in GSI and GGL. However, GSI catalytic activity involves the interaction between two residues (a glutamate and an aspartate) that shield the γ-glutamyl phosphate intermediate and deprotonate ammonium (forming ammonia) to attack the γ-glutamyl phosphate (Murray et al., 2013). Interestingly, these key acidic residues are not conserved in the sequences of GGL, FluG or those GSI homologs that reportedly do not catalyze the synthesis of glutamine, but γ-glutamylamidation reactions (Kurihara et al., 2008; Ladner et al., 2012; Takeo et al., 2013; Krysenko et al., 2017). On the other hand, residues acting on metal binding and glutamate orientation are usually conserved among these enzymes (Ladner et al., 2012), as was confirmed in FluG. The attachment of glutamate has been attributed to exert a scaffolding or protective role on substrates destined to specific metabolic fates (Walker and van der Donk, 2016), as will be discussed below. The strain in which the GGL coded by PA5508 replaced the C-terminal region of FluG recovered the functionality of the WT strain in all tested mutants and conditions. Earlier reports associated this gene with the polyamine utilization below. The strain in which the GGL coded by PA5508 replaced fates (Walker and van der Donk, 2016), as will be discussed.

Although details of the reaction purportedly catalyzed by the N-terminal region of FluG remain unresolved at this point, the predicted structural features point to the hydrolysis of a bond, excluding deaminase (elimination of ammonia from aromatic bases) activity. Alignments based exclusively on proteins with solved crystal structures, showed a notable residue conservation between the N-terminal region of FluG and LSEI_0440. Furthermore, both enzymes exhibited strong similarities with cyclic amidohydrolases (Nam et al., 2005), such as dihydropyrimidinases (Lohkamp et al., 2006; Hsieh et al., 2013) and hydantoinsases (Cheon et al., 2002) that take part in the pyrimidine degradation pathway. Unfortunately, although LSEI_0440 could functionally replace the amidohydrolase region of FluG, the actual activity of this enzyme remains unknown. Previous reports on the occurrence of promiscuous enzymes among the amidohydrolase superfamily (Baier et al., 2016) should be taken into consideration, when evaluating the functional replacement of the N-terminal of FluG by LSEI_0440. Bearing in mind that the amidohydrolase superfamily has emerged from a common ancestor (Seibert and Raushel, 2005) and that ancestral enzymes may have multifunctional (non-specialized) or promiscuous catalytic activity (Baier et al., 2016), LSEI_0440 might as well retain enzymatic characteristics that confer substrate or catalytic promiscuity, compared to a more specialized role of the N-terminal of FluG. On the other hand, the inherent Michaelis constant
in order to feed alternative pathways (Walker and van der Donk, 2016). Furthermore, nitrogen acquisition through glutamylated intermediates in bacteria has been reported to occur in the biosynthesis of polyamines (Kurihara et al., 2008; Krysenko et al., 2017) and certain catecholamines (Ladner et al., 2012; Takeo et al., 2013). Regardless of the predictions obtained for the C-terminal GSI enzymatic nature to date, phenotypic evidence obtained in this investigation sustains that FluG performs the γ-glutamylation reaction of yet unknown compounds, assisted by the N-terminal region. Given this scenario, it would seem that the purported substrate of FluG could undergo at least two metabolic fates, namely the promotion of growth, as observed in mutants affected in C-terminal activity, or asexual development. The intracellular pool of that substrate may be much greater under high nutrient availability and scavenging by the N-terminal amidohydrolase activity may be an important factor in balancing the ratio of growth and asexual development within the colony.

**AUTHOR CONTRIBUTIONS**

MI-S conducted and designed the experimental work, analyzed the data, and wrote the manuscript. LT-G conducted and designed the experimental work, carried out the statistical treatment of the results, and revised the manuscript. MC contributed to the bioinformatic approach, validated the final predictions, and revised the manuscript. UU co-conceived the work, ensured the integrity of the work, contributed to writing the manuscript, and revised and approved the final version for submission.

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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.2019.00778/full#supplementary-material
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**Conflict of Interest Statement:** 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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