A Ras GTPase associated protein is involved in the phototropic and circadian photobiology responses in fungi

Silvia Polaino1, José M. Villalobos-Escobedo2, Vilplenda P. S. Shakya3, Alejandro Miralles-Durán3, Suman Chaudhary1, Catalina Sanz4, Mahdi Shahriari4, Eva M. Luque3, Arturo P. Eslava6, Luis M. Corrochano3, Alfredo Herrera-Estrella2 & Alexander Idnurm1,5

Light is an environmental signal perceived by most eukaryotic organisms and that can have major impacts on their growth and development. The MadC protein in the fungus Phycomyces blakesleeanus (Mucoromycotina) has been postulated to form part of the photosensory input for phototropism of the fruiting body sporangiophores, but the madC gene has remained unidentified since the 1960s when madC mutants were first isolated. In this study the madC gene was identified by positional cloning. All madC mutant strains contain loss-of-function point mutations within a gene predicted to encode a GTPase activating protein (GAP) for Ras. The madC gene complements the Saccharomyces cerevisiae Ras-GAP ira1 mutant and the encoded MadC protein interacts with P. blakesleeanus Ras homologs in yeast two-hybrid assays, indicating that MadC is a regulator of Ras signaling. Deletion of the homolog in the filamentous ascomycete Neurospora crassa affects the circadian clock output, yielding a pattern of asexual conidiation similar to a ras-1 mutant that is used in circadian studies in N. crassa. Thus, MadC is unlikely to be a photosensor, yet is a fundamental link in the photoresponses from blue light perceived by the conserved White Collar complex with Ras signaling in two distantly-related filamentous fungal species.

Eukaryotic life harnesses its energy from the sun, which cycles through a 24 hour period. Light represents a potent environmental signal available to organisms. In addition to sensing light, many species have also evolved circadian regulation to anticipate the time of day, using changes in light and temperature to set their clocks. The molecular mechanisms behind light-sensing and circadian clocks are being sought in ongoing molecular genetic investigations in a suite of organisms, including prokaryotes, animals, plants and fungi. Most species of this latter group respond to light. The fungi offer experimental benefits because they are non-photosynthetic and have haploid and unicellular growth stages.

Starting in the 1960s, two fungal species emerged as models for photobiology. Of these, the ascomycete Neurospora crassa became the predominant species for light and clock studies in the fungi. It was in N. crassa that the components to sense light, that is the White Collar (WC) complex, as well as to establish the circadian feedback loops, were first identified1,2. A key strain isolated in the 1960s was the band (bd) mutant, which provides a tight banding pattern of the asexual conidia approximately every 24 hours as part of the circadian output, whereas wild type strains have a diffuse pattern of conidiation3,4. The bd mutation was incorporated into the strains used in almost every subsequent circadian clock experiment. The mutation was only relatively recently identified by gene mapping approaches5. Band is an unusual allele of the gene encoding the small GTPase protein Ras6. The

1Division of Cell Biology and Biophysics, School of Biological Sciences, University of Missouri-Kansas City, Kansas City, USA. 2Laboratorio Nacional de Genómica para la Biodiversidad, CINVESTAV Sede Irapuato, Irapuato, Guanajuato, Mexico. 3Departamento de Genética, Universidad de Sevilla, Sevilla, Spain. 4Departamento de Microbiología y Genética, Universidad de Salamanca, Salamanca, Spain. 5School of BioSciences, University of Melbourne, Australia. Correspondence and requests for materials should be addressed to A.I. (email: alexander.idnurm@unimelb.edu.au)
**Figure 1. Identification of the madC gene of *P. blakesleeanus* by positional cloning.** (A) Phenotypes of a wild type and five representative madC mutant strains. The effect of light, illuminated from the left side, on the sporangiophores of the wild type (WT) NRRL1555 and the allelic series of madC strains (1–5) grown in potato dextrose agar medium. (B) Segregation of the madC gene with molecular markers. Progeny (*n* = 93) from a cross between strains UBC21 × B2 were analyzed using PCR-RFLP markers. Marker names indicate the primers used for the amplification of the regions that are polymorphic between the two parents. Distances are in map units. (C) Position of mutations in the madC gene on a diagram of the gene, in which blue boxes indicate exons. 19 independent madC strains were examined: all bear a mutation along the gene. Numbers 1–5 indicate the alleles identified in the gene. The mutant strains have reduced phototropism and have allele 1: strains A202, A914, B2, B3, B4, S193, S196 and S205; allele 2: strains C148, L1, L72 and S5; allele 3: strains C39, C54 and C93; allele 4: strains A491 and A492; allele 5: strain A905. (D) Nucleotide changes in the madC mutants compared to the wild type sequence. Lower-case letters are intron sequences. All the alleles would produce an impaired protein: alleles 1 and 2 affect the 5′ G of the intron, alleles 3 and 4 introduce premature stop codons (underlined), and allele 5 introduces a new 3′ splicing site.

*P. blakesleeanus* produces asexual spore-forming structures, the sporangiophores, that are several cm long and about 0.1 mm wide and that bend towards blue light (Fig. 1A). The mutants with impaired phototropism led to the identification of ten different genes, named *mad* genes, defined by complementation tests and genetic linkage analyses.6,8,10 The *mad* genes were subdivided into two classes, one involved in the photoreceptor function and the other in other tropic responses (known as the “stiff” mutants). Between two and five different photosensors were predicted for the species11. The *madA* and *madB* genes were identified as homologs of the *N. crassa* *wc-1* and *wc-2* genes, respectively, based on the hypothesis that these types of proteins would also regulate the blue light responses of *P. blakesleeanus*.12,13 The phenotype in two *madI* mutants was recently identified as being caused by a situation in which the strains are heterokaryons, with a wild type copy and mutated copy of the *madI* gene14. The identity of the remaining seven *mad* genes could not be predicted based on the current knowledge of photobiology in fungi.

Based on the *madC* phenotype by itself and through the analysis of strains combining *madC* mutations with mutations in other *mad* genes, MadC was predicted to be a photosensor or to interact with MadA or MadB. The core domains for sensing light that are present in photoreceptor proteins are limited in number and few interacting components are known for the White collar complex in fungi. Hence, the discovery and characterization of *madC* could provide an advance in understanding photobiology, and was the impetus to identify the *madC* gene.

**Results**

**Mapping-based identification of the madC gene.** To position the *madC* gene within the context of the genome sequence14, crosses were established between strain UBC21 (wild type, +) and two *madC* phototropism mutant strains, A914 and B2, of the (−) sex. Of the two crosses, B2 × UBC21 yielded a larger set of progeny, with 93 progeny isolated from 93 different zygospores: 35 were wild type and 58 were *madC* mutants. 56 PCR-RFLP markers were used on a subset of progeny to identify markers linked to the phototropism phenotype. The lowest recombination frequencies were found for markers on scaffolds located in linkage group IV16. Refined mapping with additional markers and using all 93 progeny indicated that the *madC* gene lies between RFLP marker amplified with primer combination ALID0403-ALID0404 in scaffold 33 and primer combination ALID0391-ALID0392 in scaffold 5 (Fig. 1B). The phototropic phenotype was less than 1 cM from the ALID0403-ALID0404 RFLP marker.
Taking the genetic linkage map of *P. blakesleeanus* and aligning to the genome sequence the markers that gave low recombination frequencies in the UBC21 × A914 and UBC21 × B2 crosses defined a small region in the genome, at the end of the scaffold 33, in which the madC gene could lie (Fig. 1B). The mutations in genes in madC strains were sought within this interval. Point mutations were identified in one gene (encoding protein id: 200804), initially in strain A914 and then in the two madC strains (A905 and B2) with available Illumina sequence data. In comparison, 17 other mad mutants whose genome sequences are available did not bear mutations in this gene.

All madC strains have loss-of-function alleles in a gene encoding a Ras GTPase activating protein. By BLAST comparisons to the NCBI and *Saccharomyces cerevisiae* genome database, the protein impaired in madC strains is most similar to Ras GTPase activating proteins (GenBank KY522796). The candidate madC gene was sequenced from a larger collection of wild type and madC strains (Table S1). All madC strains carried mutations while no wild type or other strains did. Five different alleles were identified in the madC strains (Fig. 1C). All are predicted to produce non-functional proteins. Two mutations affect the most 5’ G nucleotides of introns, two introduce premature stop codons, and one introduces a new 3’ splicing site (Fig. 1D); the use of this new 3’ splice site for the madC allele in strain A905 was confirmed by sequencing the cDNA in that region. 19 of the madC mutants are independently-derived strains, suggesting hot spots for mutations, especially in intron splice sites as was observed previously with the madB mutants that all contain the identical mutation within an intron splice site.

The mutations in the candidate madC gene were further linked to the impairment in phototropism by genetic segregation data using additional crosses (Data set S1). This is best illustrated in the UBC21 × A202 cross. The mutation, which in this case conveniently changes a restriction enzyme site, perfectly co-segregates with reduced phototropism, while recombination is seen between the phototropism phenotype and adjacent markers.

**MadC of *P. blakesleeanus* functions in Ras signaling.** The madC gene encodes a putative Ras GTPase activating protein (GAP). Ras GAPs participate in the regulation of Ras activity by increasing the rate of hydrolysis of GTP to the inactive GDP-bound form. It is not possible to produce stably transgenic *P. blakesleeanus* strains, limiting the possible experiments to perform in this fungus to further characterize MadC function. Instead, we used two approaches to provide evidence that the *P. blakesleeanus* MadC protein is a Ras GAP.

The first approach was the yeast two-hybrid interaction method to examine if MadC of *P. blakesleeanus* could physically interact with Ras homologs in its genome. Based on best reciprocal BLAST matches with the *S. cerevisiae* Ras1/Ras2 proteins, 10 candidate Ras homologs are present in *P. blakesleeanus*. Two genes encoding Ras homologous proteins were selected (id: 188551 and id: 177348), most closely related to the *S. cerevisiae* homologs and both of which are expressed, and their cDNAs isolated and cloned into yeast two-hybrid vectors along with the madC cDNA. Yeast strains expressing the *P. blakesleeanus* Ras homologs or MadC fused to the Gal4 domains grew on media without adenine or without histidine, and had higher β-galactosidase activity. These growth patterns and enzymatic activity indicate that MadC interacts physically with both RasA and RasB (Fig. 2). To explore further the relationship of MadC with other light-sensing components, the MadA and MadB homologs were cloned and expressed in the same yeast two-hybrid system. While MadA and MadB interacted in this system, as previously reported, there was no evidence of interaction between MadC with either MadA or MadB.

The second approach was to explore whether MadC is a functional Ras GAP by cross-species complementation. *S. cerevisiae* encodes two Ras GAP homologs, Ira1 and Ira2, that exhibit different phenotypes in mutants depending on the parent strain background. These proteins, as the homologs of the NF1 protein of humans, have been extensively studied and the Ras GAP proteins have a high degree of functional conservation: for instance, human NF1 can replace the functions of the two IRA genes in *S. cerevisiae*. We deleted the IRA1 homolog in *S. cerevisiae* strain W303, and then expressed the *P. blakesleeanus* madC cDNA clone under the control of the constitutive ADH1 promoter in the ira1Δ strain. Deletion of IRA1 in this *S. cerevisiae* genetic background renders the strains sensitive to exposure to heat shock. Expression of the *P. blakesleeanus* madC gene in the ira1Δ strain restored in large part its heat shock sensitivity to wild type levels (Fig. 3). This result suggests that MadC is sufficiently conserved to function in regulating Ras function in this distant relative.

**Transcriptional changes caused by mutation of madC.** A potential interaction between the light sensing White Collar complex and the Ras signaling system would be at the transcriptional level. The transcript abundances of madA, madB and madC, and the cryA gene encoding a cryptochrome as a photoregulated gene control, were assessed in wild type and mad mutant strains grown under darkness or exposed to light, using quantitative reverse transcriptase PCR. No major differences were observed in the transcript levels of the mad genes in response to light (Fig. S1).

A global analysis of transcript changes was therefore undertaken using an RNA-sequencing approach. The transcriptional response of the wild type and madC mutant strains (Fig. 4A) indicated that overall the number of light regulated genes is reduced in the mutant, with 553 genes up-regulated and 352 genes down-regulated in the wild type (Supplemental Dataset 2), whereas there are 497 and 28 in the mutant (Supplemental Dataset 3), respectively. Based on their pattern of expression in the two strains, the set of light responsive genes can be separated into six clusters. Cluster I contains the 330 induced genes that are shared by the two strains (Fig. 4A,B). Notably, this cluster contains the paralogs of the genes madA and madB (i.e. wcoA, wcoB, wctB, wctC, wctD) and the cryA gene, representing all genes with a potential role in photoreception in *P. blakesleeanus*, and whose response to light is not affected by the mutation of the madC gene. Cluster IV encompasses 78 down-regulated genes that are shared between the two strains (Fig. 4A,B). Clusters III and VI contain the differentially expressed genes detected only in the madC mutant (167 up-regulated and 28 down-regulated), while clusters II and V contain those genes that appear to be responsive only in the wild type strain (Fig. 4A).
Interestingly, in general light responsive genes display a stronger response in the wild type than in the madC mutant (Fig. 4B), which is reflected in an important number of genes that pass the cut-off established [false discovery rate (FDR) < 0.05; fold-change ≥ 2] in the wild type, and have the same tendency in the mutant but do not pass this filter. As highlighted by the intersection of the reference diagonal lines (crossing the 0 of log2FC) and the cut-off lines of log2FC, many of these genes are close to the cut-off that was established (Fig. 4A). This set of genes actually passes the FDR filter but does not fulfill the fold-change criteria (green dots in Fig. 4A). In this regard, we detected 94 up-regulated genes (FDR < 0.05; fold-change ≥ 2) in the wild type that in the madC mutant show a FC ≥ 1.5. Similarly, among the down-regulated genes we found 133 fulfilling the cut-off criteria only in the wild type.

Figure 2. *P. blakesleeanus* MadC and Ras homologs physically interact in a yeast two-hybrid assay. *S. cerevisiae* strain PJ69-4A has the ADE2, HIS3 and β-galactosidase genes under the control of galactose-inducible promoters. Physical interactions between proteins fused to the Gal4 activation domain (in plasmid pGADT7) and DNA binding domain (in plasmid pGBK7) trigger the expression of these three reporters. (A) Growth of strains co-transformed with two plasmids on –leucine –tryptophan (–leu –trp) to maintain the plasmids, or on media to measure interactions (–histidine + 0.5 mM 3-aminotriazole, or –adenine). (B) The average β-galactosidase activities of the strains, with standard errors from two biological replicates. (C) Arrangement of the cDNA clones in the plasmids with the Gal4 activation and DNA binding domains. EV; empty vector.

Figure 3. The *P. blakesleeanus* madC gene complements the heat shock sensitivity of the *S. cerevisiae* ira1Δ mutant. The wild type strain of *S. cerevisiae* and an ira1Δ strain were transformed with an empty vector or with a vector expressing the madC gene. Ten-fold serial dilutions of the strains were plated on YPD medium. One plate was heat shocked at 50 °C for 120 min. The plates were then cultured 2 days at 30 °C.
type, and that in the madC strain show a FC ≤ −1.5. These comparisons allow us to conclude that most of the differentially expressed genes of the wild type are shared with the madC mutant, and that the corresponding gene is most likely involved in the modulation of the light response. It is, however, noteworthy that although both strains seem to activate genes related to signal transduction processes, in the mutant strain this GO category as well as that of lipid metabolic process is not significantly enriched (Fig. 4C; Table S2). Furthermore, there is no GO-term enrichment in the genes repressed in the mutant, whereas aspartate and alanine metabolic processes are enriched in the set of genes repressed in the wild type.

The volcano plots in Fig. 5 show the differentially-expressed genes for both wild type (Fig. 5A) and madC (Fig. 5B) strains. These graphs illustrate a clear tendency for induction of gene expression in response to light rather than to repression in both strains. A set of very strongly induced genes (fold change higher than 16) called our attention since they could be guiding the main light response. In the wild type strain 50 genes have an expression level of this magnitude, and in the mutant only 30. Among these, 24 genes do not reach this level of expression in the madC strain. Within this group of transcripts affected by the madC mutation there are two Rho GTPases (id: 76783 & id: 152853). These proteins participate in the regulation of cytoskeletal rearrangements.

Figure 4. Comparison of the transcriptional response to light of wild type and madC strains. (A) Plot of gene transcript levels of the wild type vs madC strains by log2FC. Red circles represent differentially expressed genes in response to light for the wild type, and blue dots those differentially expressed in the madC strain (FDR <0.05; log2 fold-change ≥ 1). Grey dots represent genes that are not differentially expressed, and green dots are genes that do not pass the fold-change ≥1 but have a significant change in the madC strain as determined by FDR. Diagonal lines (crossing the 0 of log2FC for both strains) serve as a reference to identify genes in the borderline of the cut-off criteria. (B) Venn diagrams showing the overlap of the repressed and induced genes in response to light between the wild type and madC strains. (C) Category enrichment in differentially expressed genes (*FDR <0.2; **FDR <0.05). Color intensity represents the percentage of genes belonging to each category and includes GO terms for Biological Process and Molecular Function, which contain less than 900 and more than 5 genes. The dendrogram highlights clusters based on percentages of genes contained in each gene ontology.
and vesicular traffic in eukaryotes and are part of the central elements that participate in the shift from isotropic to polarized growth in filamentous fungi, confirming that pathway signaling mediated by GTPases in the madC strain is affected\(^2\)\(^1\)\(^{–}\)\(^2\)\(^3\)\(^\text{.} \) PAC2 (id: 6945) is another gene highly expressed in the wild-type strain and affected in madC; Pac2 participates in alpha-tubulin and beta-tubulin folding to be functional and null mutants of pac2 are super-sensitive to benomyl, a microtubule depolymerizing drug\(^2\)\(^4\)\(^\text{.} \) These observations suggest that defects in vesicular transport may exist in the madC mutant. Another gene affected in the madC mutant encodes a dia-cylglycerol kinase (DGK, id: 175731) that catalyzes the conversion of diacylglycerol (DAG) to phosphatidic acid (PA), suggesting that phospholipid signaling in the light response is affected by the absence of madC. On the other hand, the cpcC gene (id: 121290) encoding the Clock-Interacting Protein (CIPC), a negative-feedback regulator of the mammalian circadian clock\(^2\)\(^5\)\(^\text{.} \) is strongly induced in the wild type strain with a fold-change greater than 19, whereas in the madC mutant this gene is not differentially expressed in response to light (see inset in Fig. 5A), and shows a significant decrease in expression even in the dark.

The MadC homolog can impact photoresponses in other fungi. MadA and MadB (or WC-1 and WC-2, as named in N. crassa) are conserved in many fungi\(^2\)\(^6\)\(^–\)\(^2\)\(^7\)\(^\text{.} \) Likewise, homologs of MadC are found across the fungi and also in other eukaryotes (Figure S2). To address whether or not these homologs may also play roles in photobiology responses in other fungi, we tested their functions in a basidiomycete, Cryptococcus neoformans, and an ascomycete, N. crassa, which are both model species for photobiology research within their respective phyla.

Figure 5. Gene expression profile in response to light of wild type and madC strains. Volcano plot of gene expression profile in response to light in the wild type (WT; A) and madC (B) strains, annotated as the proteins encoded by those genes. Plots of FDR versus log2 fold change for light/dark comparison in each strain. The two blue vertical lines in each chart mark two-fold change. Labels indicate representative genes with highly significant fold changes (log2 FC >4). In A, the expression in counts per million (CPM) of the cpcC (homolog of clock-interacting protein CIPC) gene is shown.
The madC homolog (named IRA1; CNAG_06929, a single copy gene) in *C. neoformans* was deleted, and the strains tested for the three phenotypes associated with light-sensing in this fungus. In contrast to the loss of the white collar homologs *BWC1* and *BWC2*, there was no effect of deleting the madC homolog on UV sensitivity or repression of mating by light (Fig. 6). A reduction in virulence was observed using an insect model of disease, like that seen for the wc-1 mutant. A double *bwc1* *ira1* mutant was isolated through genetic crosses, and its virulence tested. This strain showed an additional decrease in virulence, suggesting the actions of two independent pathways on virulence. *Ira1* likely alters Ras signaling, known to be required for virulence in *C. neoformans* and *P. blakesleeanus*, as a gene encoding a Ras GTPase activator.

The function of the madC homolog in *N. crassa* (*ira-1*; NCU06122, a single copy gene) was examined. Previous studies have shown that Ras impacts the circadian output in *N. crassa*. Homokaryon mutants without a copy of *ira-1* were isolated, by crossing a heterokaryon strain obtained from the *N. crassa* gene deletion project. Double *bd* *ira-1* mutants were isolated by performing additional crosses. The wild type, *ira-1::hph* mutant, *bd* strain, and double *ira-1::hph* *bd* mutant were grown in light and placed in races tubes in constant darkness. Strikingly, the *ira-1::hph* mutant produced rhythmic conidia approximately every 24 h like the *bd* mutant (Fig. 7A).

The *ira-1::hph* mutant grows slower than the *bd* mutant (Fig. 7A). Extending the incubation time to 14 days shows that the mutant continues to make bands of conidia up to about 10 days (Fig. 7B). The double *ira-1::hph* *bd* mutant exhibited additive effects, with slower growth than either the *ira-1::hph* or *bd* single mutants. Despite the slow growth, the double mutant was capable of forming bands in a circadian manner in constant darkness. The strength gradually diminished, but could still be faintly observed after 28 days (Fig. 7B). The reason why the RAS-1^bd^ protein isoform causes banding is unknown, as the *bd* mutant has little change in GDP/GTP ratio as compared to the wild type, but has a small decrease in the GDP exchange rate.

Discussion

Two of the genes needed for phototropism in *P. blakesleeanus*, *madA* and *madB*, are the homologs of the *N. crassa* *wc-1* and *wc-2* genes, and were identified by predicting they were homologs of this conserved light-sensing complex. The *madC* gene, encoding the other main photosensory component in the phototropism response, was unknown since the 1960s. In this study, we identify *madC* in *P. blakesleeanus* as a gene encoding a Ras GTPase activating protein.

The genomes of Mucoromycotina species often feature extensive gene duplication, either from whole genome and/or segmental duplication events. For instance, the *madA* and *madB* genes of *P. blakesleeanus* and the

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**Figure 6. Phenotypes of the *C. neoformans* *ira1* mutant are not associated with light sensing.** (A) Light represses mating in crosses between wild type or *ira1* mutant parents, but does not on the *bwc1* parents. Yeast cells were mixed on Murashige-Skoog medium and incubated in the dark or light. Mating is represented by the filaments emerging from the edges of the yeast colonies. (B) The *ira1* mutants have a wild type level of UV sensitivity in contrast to the *bwc1* mutants. 10-fold serial dilutions of yeast cells were plated onto media, one plate was irradiated with 120 J/m² UV, and strains grown 2 days to form colonies. (C) The *ira1* mutant has a reduction in virulence, with an additive effect when combined with the *bwc1* mutation. Wax moth larvae were inoculated for each strain, as indicated, in two independent experiments and survival monitored over time. Left graph *n* = 17; the wild type is more virulent than the two other strains (P < 0.001). Right graph *n* = 12; the differences between *ira1*Δ and *ira1*Δ + IRA1 (P < 0.05) or *ira1*Δ and *bwc1*Δ *ira1*Δ (P < 0.05) are statistically significant.
homologous genes in other species in the Mucoromycotina are members of gene families with three or four
genes, respectively. The same situation is observed for homologs of the madC gene identified here, since
*P. blakesleeanus* or related species have three or more madC homologs in their genomes (Figure S2). Considering
the Ras pathway in *P. blakesleeanus*, the situation is even more complex, with ten Ras homologs in this species.
However, the duplication of the Ras GAPs in *P. blakesleeanus* may have aided the discovery of this function
in the phototropic response. This is because mutation of the homologs in other fungi can result in deleterious
effects on growth. This includes the extreme phenotype of being essential for viability in some *S. cerevisiae* strain
backgrounds, and mutations causing growth or developmental defects in the basidiomycete *Schizophyllum
commune* or ascomycetes *Aspergillus nidulans*, *Colletotrichum orbiculare* and *Schizosaccharomyces pombe*.
The *P. blakesleeanus* madC mutants have no defects in growth, development (e.g. mating or asexual sporulation)
or other tropisms (gravity, avoidance response, photoinduction of sporangiophores or photoinduction of carotene
synthesis) although a recent report suggests that one strain carrying a madC mutation may be hypersensitive to
gravity. The madC gene may have specialized its function after gene duplication to be specific to phototropism
allowing the other Ras GAP proteins to perform their basic cellular roles.

The interactions between Ras signaling and photobiology are potentially widespread in eukaryotes. We tested
the function of the homolog of the *P. blakesleeanus* madC gene in two divergent fungal species. While we did
not see an effect on the responses to light in *C. neoformans*, this may reflect the possible life style of this species
which is predominantly as a yeast. In contrast, both the Ras GAP homolog of MadC and Ras itself are involved
in the circadian output in *N. crassa*. The additive effect in *N. crassa* of the double mutant supports the hypothesis
proposed by Belden et al. that the bd mutation impacts the interaction of Ras with guanine exchange factors:
decreased growth rates in the bd ira-1 double mutants are consistent with impaired growth seen for dominant
negative and dominant active forms of Ras.

Our transcriptome analysis of the *madC* mutant indicates that the function of this gene is not directly related
to the transcriptional activation of the specific light responsive genes. However, congruently with the role in
signaling processes of a GTPase, genes important for signal transduction are affected in response to the stimulus,
since some of these genes are strongly induced in response to light and are no longer responsive in the mutant.

In *N. crassa*, the Rho-GTPase CDC-42 has been reported to participate in vesicular transport to the hyphal
tip via the actin cytoskeleton and microtubules to carry out a thigmotropic response: this response depends
on calcium-regulated supply of vesicles to the extreme hyphal apex via the actin cytoskeleton. In the wild
type strain of *P. blakesleeanus*, the phospholipase C (PLC; id: 121739) and diacylglycerol kinase DGK genes are
induced, the latter is affected in *madC*. In both *N. crassa* and *C. albicans* PLC activity increases intracellular Ca^{2+}
concentrations at the tip via inositol 1,4,5-triphosphate (IP_3)-mediated calcium release from vesicles, and has
been related to the thigmotropic response through a calcineurin-related mechanism and DGK is an essential
enzyme in phosphatidic acid-dependent signaling pathways. The signaling cascade mediated by PLC and DGK
is also involved in the phototropic movement of *Arabidopsis* chloroplasts and protection against high-intensity
UV-B irradiation. Thus, a light response network is described, which induces cytoskeletal modification and
transport of vesicles to the tip of the sporangiophore, where there is an increase in calcium that can lead to
phototropism in *P. blakesleeanus*.

One future direction is to explore if aspects of light sensing in *N. crassa* are altered in the *ira-1* mutant. Striking
also is the role of the Ras GAP homolog in fruit fly *Drosophila melanogaster*, in which loss-of-function causes
arrhythmic clock functions. A potential mammalian link is the role of the Ras family member, Dextras1, in the

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**Figure 7. The *N. crassa* homolog of madC impacts circadian output.** (A) The wild type strain, *bd* strain, the
mutant in the madC homolog *ira-1*:hph and double *ira-1*:hph *bd* mutant were cultured in races tubes, growing
from left to right, in light for 24 h at 25 °C and then transferred to constant darkness for 5 d at 25 °C. (B) Using
similar culturing and initial illumination conditions as in (A), subsequently the *ira-1*:hph mutant was grown in
darkness 14 days and the double *ira-1*:hph *bd* mutant grown in darkness for 28 days. Tubes are 16 mm in
diameter. The * symbols indicate the positions of the growing fronts of the mycelia when photographed.
light input into the circadian clock in mice\(^4^6\). Other studies continue to implicate an essential role for Ras signaling in circadian rhythms in animals\(^4^7\)–\(^4^9\).

Interestingly, although in *P. blakesleeanus* a role for *madC* in circadian rhythms has not been established, the finding of the strongly affected expression level of the CIPC homolog correlates with the observed phenotype in the *ira-1* mutant in *N. crassa*. Thus, *madC* may have a highly conserved regulatory role in circadian rhythms.

In summary, we provide evidence that MadC is involved in Ras signaling in the life style of *P. blakesleeanus* to provide its ability to respond to light. It is remarkable that in two highly-diverged species of fungi, *P. blakesleeanus* and *N. crassa*, mutations that affect the same signaling pathway to alter photobiology were isolated in the 1960s, then remained unknown for over four decades. Future directions are to establish how MadC and Ras, or other Ras regulators interplay with the White Collar components to control the outputs of light signaling.

**Materials and Methods**

***P. blakesleeanus* strains and crosses.** Details about the wild type and *mad* mutant *P. blakesleeanus* strains that were used are provided in Table S1. Crosses were established on V8 juice agar medium, and progeny isolated from zygospores two-four months later\(^1^6\).

**Mapping *madC*.** DNA was isolated from the progeny from crosses between *mad* mutants [A914 and B2 (−)] with a wild type parent [UBC21 (+)] and used as the templates for using molecular markers developed previously to make a genetic linkage map of *P. blakesleeanus*\(^1^6\). Phototropism was scored by tropic reactions of sporangio-phores growing on potato dextrose agar (PDA) medium in a homemade dark box with a glimmer window. The molecular marker and phototropism data were analyzed for linkage in JoinMap 4.0 software\(^5^0\).

**Yeast two hybrid analysis.** The 5′ and 3′ ends of the *madC* gene were identified using rapid amplification of cDNA ends (GeneRacer kit, Invitrogen, Carlsbad, CA). A full-length cDNA clone was amplified with primers ALID1628–ALID1629 from reverse transcribed mRNA (SuperScript III First-Strand Synthesis System for RT-PCR, Invitrogen) and directly cloned using the EcoRI site into the pGADT7 and pGBK7 plasmids. The *madA* cDNA was subcloned as an EcoRI fragment into the EcoRI sites of these two plasmids. The *madB* cDNA was subcloned as an NcoI-Sall fragment into the NcoI-Xhol sites of pGADT7 plasmid or NcoI-Sall sites of pGBK7 plasmid. The *rasA* and *rasB* cDNAs were amplified with primers ALID1734–ALID1735 and ALID1736–ALID1737, respectively. The amplicons were recombined directly with the pGBK7 plasmid in *S. cerevisiae*. The *rasA* and *rasB* cDNAs were amplified from these plasmids with primers ALID1829–ALID1830 and ALID1831–ALID1832, respectively; these amplicons were recombined directly with the pGADT7 plasmid in *S. cerevisiae*. The cDNA clones were sequenced to identify those without errors.

The plasmids and controls were co-transformed into *S. cerevisiae* strain PJ69-4A using the lithium acetate/PEG protocol\(^5^1\). Assessing the interactions of proteins followed the methods in James et al.\(^5^2\).

**Deletion of *S. cerevisiae* *ira1A* and complementation with the *P. blakesleeanus* *madC* gene.** The *IRA1* gene was deleted in the *S. cerevisiae* strain W303 by replacing it with the selectable marker KanMX6, from the plasmid pFA6a-GFP (S65T)-KanMX6, amplified with primers SP98 and SP99. After transformation, the cells were plated onto YPD medium containing G-418. A gene replacement strain was identified and confirmed with different combination of primers, SP100 and KanB, KanC and SP101, SP100 and SP102, SP103 and SP101.

The *madC* cDNA was amplified with primers ALID1628 and ALID1629 and cloned into the EcoRI site of plasmid pTH19, which has expression from the constitutive *ADH1* promoter\(^1^6\). The W303 strain of *S. cerevisiae* and an *ira1::KanMX6* mutant were transformed with the control plasmid pTH19 and with the plasmid expressing the *madC* gene. The strains were inoculated in 5 ml YNB + glucose + leucine + tryptophan + adenine + histidine medium and grown overnight at 25 °C in shake cultures. Ten-fold serial dilutions of the strains were plated on YPD medium and exposed to 50 °C for up to 120 min, then allowed to grow two days at 30 °C.

**Disruption of the *madC* homolog gene (IRA1) in *C. neoformans* and characterization of mutant strains.** The MadC homolog was identified by BLASTp. The corresponding gene was replaced with a construct conferring nourseothricin resistance by homologous recombination. The construct was made by overlap PCR using primers listed in Table S3, and transformed into strain KN99a\(^1^4\) by biolistic delivery, with subsequent selection on YPD medium containing nourseothricin (100 μg/ml). A MATa version was isolated by crossing this strain to congeneric strain KN99a\(^1^4\), and a double *bwc1Δ ira1Δ* mutant was isolated by crossing. A complemented strain was generated by amplifying a wild type copy of *IRA1*, cloning it adjacent to a marker conferring G-418 resistance, and transforming this construct by *Agrobacterium* -mediated transformation in the *C. neoformans* *ira1Δ* mutant with selection on G-418 (100 μg/ml) and cefotaxime (200 μg/ml) for counter-selection of the bacteria.

The phenotypes of the strains were examined for inhibition of mating by light, UV sensitivity and virulence. Mating was tested by mixing strains of a and a mating types on Murashige-Skoog medium, and keeping one set under white light. UV sensitivity was tested by plating ten-fold serial dilutions onto YPD medium, and irradiating the plates with a UV transilluminator. The virulence of the strains were tested in the *Galleria mellonella* insect model\(^5^3\). Larvae were purchased from Vanderhorst Wholesale, Inc. (Saint Marys, OH). In brief, 1 x 10⁶ cells, from overnight cultures were washed and then resuspended in phosphate buffered saline (PBS), and then injected into the rear proleg of 12–17 larvae. Animal survival was monitored daily, and survival between groups compared using log-rank tests.

**Characterization of the *madC* homolog (ira-1) in *N. crassa*.** *N. crassa* strains were obtained from the Fungal Genetics Stock Center (FGSC, http://www.fgsc.net)\(^5^6\). *N. crassa* genomic DNA was isolated using the ZR Fungal/Bacterial DNA kit (Zymo Research, Irvine, CA), and gene replacement validated by PCR. Strain
manipulation and growth media preparation followed standard procedures and protocols. Race tubes were inoculated, and incubated in constant light for 24 h at 25 °C before transfer to constant darkness at 25 °C (Advanced Intellus Environmental Controller Incubator, Percival Scientific, Perry, IA) for 5, 14 or 28 days, depending on the strain.

The strains ira-1::hph bd and ira-1::hph were generated by crossing the ira-1::hph strain FGSC 22825, from the N. crassa knockout project, with the bd strain FGSC 1858. Mutations were confirmed by PCR and/or sequencing of amplicons. The phenotypes of the ira-1::hph mutants segregating from the cross were not like their parents; the ira-1::hph mutants (FGSC 22824 and FGSC 22825) are heterokaryons based on PCR analysis. The ira-1::hph homokaryotic strains were crossed with the wild type strain FGSC 4200, and strains of both mating type isolated, and deposited to the FGSC.

Light induction, RNA isolation, quantitative RT-PCR and RNA-sequencing. 104 activated spores (48 °C, 15 min) of each P. blakesleeanus strain were grown at 22 °C in minimal medium. P. blakesleeanus dark-grown mycelia were exposed to blue light (2782.10 J/m2) at the age of 48 h for 30 min. After light exposure mycelia were collected and frozen in liquid nitrogen. Mycelia were disrupted by two 0.5-min pulses in a cell homogenizer (Fastprep-24, MP Biomedicals) with 1.5 g of zirconium beads (0.5 mm diameter) using the RNeasy Plant Mini Kit (Qiagen) following the manufacturer procedure. RNA samples were treated with DNase I prior to use in RT-PCR experiments. Total RNA was used for cDNA synthesis and amplification (primers in Table S3). Quantitative RT-PCR experiments were performed in a LightCycler 480 II (Roche, Madrid, Spain) using the One-Step SYBR PrimeScript RT-PCR kit (Takara Bio, Japan), 0.2 μl of each primer and 50 ng of RNA in 10 μl reactions. The reactions consisted of 5 min at 42 °C, followed by 10s at 95 °C, and then 40 cycles of DNA amplification (5 s at 95 °C and 20 s at 60 °C). The results for each gene were normalized to the corresponding results obtained with the actin gene (act-1) to correct for sampling errors. Then, the results obtained with each sample were normalized to the RNA sample from the corresponding mycelia after illumination.

Libraries for RNAseq were prepared with RNA from mycelia of the wild type and the A905 madC mutant strains using the TruSeq kit protocol (Illumina). Three biological replicates for each condition were sequenced using a HiSeq 2500 sequencer with the 1 × 100 format. 100-base-pair-long reads were obtained and mapped to the P. blakesleeanus genome V2 using bowtie, searching for end-to-end hits with at most three mismatches. Alignment results were recorded in BAM format for further downstream analysis. Read counts per gene were calculated for each library using a shell script and collapsed into a table.

Differential expression analysis of gene expression in the wild type and madC strains in response to light was performed. Only genes with at least three reads per million were considered for differential expression analyses; this was done using the edgeR package. Normalization of the data was performed using the Cox-Reid adjusted likelihood method. For determining differential expression between the comparisons, we used the Generalized Linear Model likelihood ratio test. False discovery rates (FDR) were calculated and genes with a FDR < 0.05 and fold-change ≥ 2 were considered differentially expressed. Subsequently, we compared the differentially expressed genes of both strains in the plot of Figs 4A and 5, to highlight the genes with similar or different profiles between the two sets of genes.

A GO-term enrichment analysis was performed with the hypergeometric distribution method. GO terms with FDR ≤ 0.2 were considered significantly enriched in each comparison (Table S2). All RNAseq data have been deposited in the GEO database with accession number GSE93056.

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