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Intracellular Action of a Secreted Peptide Required for Fungal Virulence

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SUMMARY

Quorum sensing (QS) is a bacterial communication mechanism in which secreted signaling molecules impact population function and gene expression. QS-like phenomena have been reported in eukaryotes with largely unknown contributing molecules, functions, and mechanisms. We identify Qsp1, a secreted peptide, as a central signaling molecule that regulates virulence in the fungal pathogen *Cryptococcus neoformans*. *QSP1* is a direct target of three transcription factors required for virulence, and *qsp1* mutants exhibit attenuated infection, slowed tissue

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The accession number for the ChIP-seq and RNA-seq data reported in this paper is GEO: GSE73203.

SUPPLEMENTAL INFORMATION

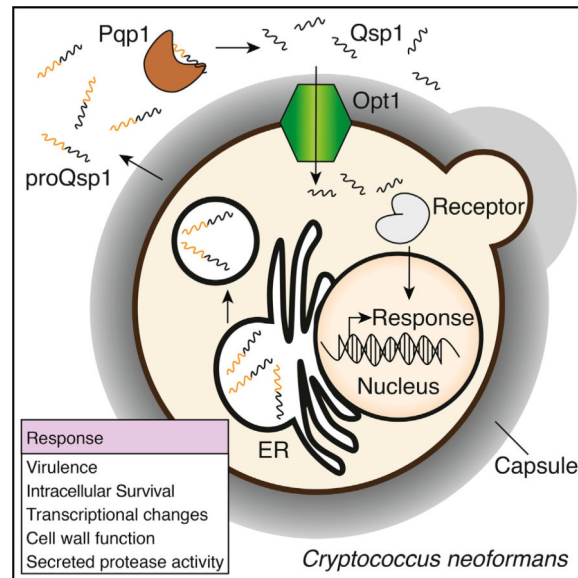
Supplemental Information includes Supplemental Experimental Procedures, six figures, and eight tables and can be found with this article online at <http://dx.doi.org/10.1016/j.chom.2016.05.001>.

AUTHOR CONTRIBUTIONS

C.M.H. and H.D.M. designed the study. D.K.S. performed experiments in Figures 7 and S3B. A.I.G. performed experiments in Figures 3D, 4H, 5B–5D, S1F, and S6C and Table S3. A.I.G., I.C., and S.P. created knockouts for screening. S.C.C. and C.S.C. performed experiments in Figure 3E and provided protocols for assays in Figure 5G. D.W. and K.N. performed experiments in Figures 2C and 2D. J.K.D., J.J.M., and J.R.Y. performed MS analyses in Figure 5. D.T. and J.P. performed the experiment in Figure S2C. R.U. and J.K.L. performed the experiment in Figure S4D. C.A.C. performed sequence and phylogeny analysis in Figure S6I and Table S5. V.P. provided protocols and guidance for the assay in Figure S2G. C.M.H. performed experiments in Figures 1, 2, 3, 4, 5, and 6 and S1–S6. C.M.H. and H.D.M. wrote the manuscript. All authors contributed to editing the manuscript.

accumulation, and greater control by primary macrophages. Qsp1 mediates autoregulatory signaling that modulates secreted protease activity and promotes cell wall function at high cell densities. Peptide production requires release from a secreted precursor, proQsp1, by a cell-associated protease, Pqp1. Qsp1 sensing requires an oligopeptide transporter, Opt1, and remarkably, cytoplasmic expression of mature Qsp1 complements multiple phenotypes of *qsp1*. Thus, *C. neoformans* produces an autoregulatory peptide that matures extracellularly but functions intracellularly to regulate virulence.

Graphical Abstract



INTRODUCTION

Cryptococcus neoformans is the most common cause of fungal meningitis. This species causes 950,000 cases and 625,000 deaths annually, an estimated 40% of all HIV-related mortality (Armstrong-James et al., 2014). *C. neoformans* is a member of a pathogen species complex that includes *C. neoformans* var. *grubii* (serotype A) and *C. neoformans* var. *neoformans* (serotype D). We study *C. neoformans* var. *grubii*, which is responsible for ~90% of infections in HIV/AIDS patients. Investigations of *C. neoformans* pathogenesis have focused on three virulence traits: melanin production, growth at human body temperature, and production of a polysaccharide capsule.

Many bacterial pathogens require autoregulatory quorum sensing (QS) systems for virulence, which are regarded as attractive therapeutic targets. In classical bacterial QS paradigms, cells secrete a signal and monitor its concentration to sense cell number and/or diffusion. Bacteria use QS to regulate diverse biological functions (Bassler and Miller, 2013). In Gram-negative bacteria, the most common secreted signals are acyl homoserine lactones (AHLs) (Schuster et al., 2013). Gram-positive bacteria generally utilize peptides (Ng and Bassler, 2009).

QS-like phenomena in fungi have been proposed based on the effects of conditioned media (e.g., Albuquerque and Casadevall, 2012; Albuquerque et al., 2014). However, QS systems with specialized, species-specific biosynthetic pathways and specific reception mechanisms akin to those of bacteria have not yet been elucidated in any eukaryote. A candidate for a bona fide QS signal in *C. neoformans* was identified in studies of a mutant *C. neoformans* serotype D strain lacking the co-repressor Tup1 (Lee et al., 2007). The authors observed that *tup1* strains, but not wild-type strains, failed to grow when a low number of cells were plated. Purification of the activity from *tup1* culture supernatants revealed an 11 amino acid peptide that was able to rescue this defect. They suggested that this peptide could be part of a QS system in *C. neoformans* and named it Qsp1 (Quorum Sensing-Like Peptide 1). Unfortunately, no phenotype for a *qsp1* strain was reported, and the reason *tup1* strains require Qsp1 is not known. A similar low-density plating defect was not observed in a serotype A strain lacking *TUPI* (Lee et al., 2009). Thus, the normal function of Qsp1 remained unknown.

RESULTS

QSP1 Is a Direct Target of Three Virulence Regulators

We reasoned that virulence factors would be enriched in the regulatory targets of Gat201, a key transcription factor required for virulence (Chun et al., 2011; Liu et al., 2008). Our previous ChIP-on-chip studies of Gat201 identified important targets (Chun et al., 2011). To develop an extended network, we employed ChIP-seq (Figure 1A). Previous studies suggested genes whose promoters are bound by more than one transcription factor regulating a particular process often share that function (Beyhan et al., 2013; Nobile et al., 2012; Pérez et al., 2013). Thus, we selected Gat204 and Liv3, transcription factors previously identified as Gat201 targets and implicated in virulence (Chun et al., 2011; Liu et al., 2008), for ChIP-seq in tissue culture conditions.

Gat201, Gat204, and Liv3 control a transcription factor network, sharing characteristics of published regulatory networks (Hughes and de Boer, 2013; Sorrells and Johnson, 2015). Gat201, Gat204, and Liv3 targets (Table S1) overlap significantly ($p < 10^{-300}$ by chi-square test), with 225 “central targets” shared by all three regulators (Figure 1B). RNA-seq analysis indicates 42% of binding events in the network cause a change in target transcript levels (Figures S1A and S1B and Table S1). Each transcription factor binds its own promoter and the promoter of the other two regulators (Figure 1C). De novo motif finding using the ChIP-seq data (Figure 1D) shows that Gat201 and Gat204 have typical GATA family transcription factor specificities (Ko and Engel, 1993) (Figure S1). Liv3, a Wor1-ortholog, produced a DNA binding motif reminiscent of the *C. albicans* Wor1 motif (Lohse et al., 2010) (Figure S1C).

A prominent central target, *QSPI* is a gene encoding a small secreted peptide (Lee et al., 2007) (Figures 1E and S1D). RNA-seq analysis revealed that *QSPI* produces the most abundant transcript in cells grown in tissue culture conditions (Figure 1F). *QSPI* is predicted to encode a peptide precursor containing a signal sequence and a 24 amino acid pro-peptide (proQsp1). The C-terminal 11 amino acids of proQsp1 encode the mature Qsp1 peptide (Lee et al., 2007) (Figure 1G). To determine whether high amounts of the peptide are produced,

we raised polyclonal rabbit antisera against Qsp1 (Figure S1E). Qsp1 is detected in wild-type culture supernatants by immunoblotting, producing a signal that comigrates with the synthetic peptide (Figure S1F). We developed a quantitative ELISA, which revealed that the peptide accumulates to micromolar concentrations in culture supernatants as a function of cell density (Figure 1H).

QSP1 Is Required for Virulence

As a highly expressed, central target of the Gat201-Gat204-Liv3 network, *QSP1* was a candidate virulence factor. We assessed whether *qsp1* mutants display a growth defect in vitro and observed no defects in the growth of *qsp1* cells on media containing oxidative, nitrosative, pH, iron starvation, osmotic, or cell wall stressors (Figures S1G and S1H). Next, we assessed virulence of two independent *qsp1* mutants using an intranasal infection model in A/J mice (Chun and Madhani, 2010; Lim et al., 1980). We found that *qsp1* mutants are significantly attenuated (Figure 2A). Since A/J mice have a C5 complement deficiency (Nilsson and Müller-Eberhard, 1967), we tested whether *qsp1* mutants were also attenuated in C57Bl/6 mice (Huffnagle et al., 1998). We obtained similar results in this mouse background (Figure S2A).

In *C. neoformans*, virulence-attenuated mutants fall into several classes. One group comprises mutants deficient in early pulmonary accumulation. Another group corresponds to mutants that accumulate normally in the lung but are deficient in dissemination. To differentiate between these two groups, we quantified the cryptococcal viable cell counts (CFUs) in lung homogenates obtained from mice infected with either wild-type or *qsp1* strains. *qsp1* mutants accumulated to approximately 3-fold fewer CFUs at day 4 after infection and to 20-fold fewer CFUs at day 13 (Figure 2B). These defects were not due to a deficiency in initial seeding of the tissue (Figure S2B). We examined CNS infection using an intracisternal infection model in immunosuppressed rabbits (Perfect et al., 1980). We detected no difference in cerebrospinal fluid CFUs between wild-type and *qsp1* (Figure S2C).

Cryptococcal infections that result in a Th2-dominant response tend to favor *C. neoformans* (Herring et al., 2004; Piehler et al., 2011). Th1-dominant responses more effectively clear the fungus (Zhang et al., 2009). We therefore examined whether there is an altered inflammatory response during *qsp1* infection. We quantified leukocytes recruited to lungs after infection with wild-type or *qsp1* strains using flow cytometry-based profiling (Figure 2C). Consistent with the lower pulmonary burden found after infection with *qsp1* mutants, there were fewer leukocytes recruited. The majority were eosinophils, indicating a Th2-polarized infection (Akuthota et al., 2008). We assessed the level of cytokines in lungs and found no differences between wild-type and mutant infections (Figure 2D). Type 2 cytokines (IL-4, IL-5, and IL-13) (Herring et al., 2004) dominated at 3.5 days after infection. IL-13 was the most abundant cytokine 13 days after infection. These data are consistent with studies of T cell polarization during wild-type *C. neoformans* infections (Qiu et al., 2013; Zhang et al., 2010).

We examined the interaction between the *qsp1* mutant and macrophages, the most abundant immune cells in the alveolar space (Gordon and Read, 2002). Hosts susceptible to

cryptococcal infection display decreased survival when depleted of macrophages (Osterholzer et al., 2009; Shao et al., 2005). Macrophages also contribute to cryptococcal dissemination (Coelho et al., 2014). We measured phagocytosis rates of wild-type and *qsp1* strains using bone marrow-derived macrophages (BMDMs) as described previously (Chun and Madhani, 2010; Nicola and Casadevall, 2012) (Figure S2D). We observed no difference between wild-type and *qsp1* mutants when cryptococcal cells were unopsonized (Figures 2E and S2E) or opsonized with a monoclonal antibody against the capsule (Figures 2F and S2E). We reasoned that *qsp1* cells may have altered fitness after phagocytosis in the high-density environment of the phagolysosome and measured the ability of the *qsp1* mutant to accumulate within macrophages after phagocytosis (Figure S2F). Wild-type cells accumulate intracellularly significantly faster than *qsp1* cells, which displayed approximately 2-fold fewer CFUs 24 and 48 hr after infection (Figure 2G). This observation raised the possibility that phagosome maturation may be altered in macrophages infected with the *qsp1* mutant. However, colocalization studies revealed no impact of *QSP1* on the level of association between cryptococcal cells and acidic macrophage compartments (Figure S2G).

Intercellular Signaling by Qsp1

We serendipitously observed that *qsp1* mutants form dry, wrinkled colonies at room temperature, in contrast to smooth, mucoid colonies produced by wild-type (Figure 3A). This colony phenotype provided an opportunity to assay Qsp1 function. We developed a confrontation assay in which recipient strains were grown as colonies near a patch of donor strain cells. When *qsp1* cells are confronted by a wild-type patch, they grow to produce smooth colonies (Figure 3B). We reasoned that this effect was due to Qsp1 diffusing from the donor. Indeed, synthetic Qsp1 was sufficient to cause *qsp1* cells to produce smooth colonies (Figure S3A). To determine whether this effect was specific, we synthesized and tested the predicted proQsp1 and pro region peptides. We also synthesized a peptide in which the 11 amino acids of Qsp1 are scrambled (Scrambled Qsp1). We found that proQsp1 is sufficient to complement the *qsp1* colony phenotype, but the pro region peptide and scrambled Qsp1 are not (Figure S3A). To assess structure-function relationships in Qsp1, we synthesized mutant peptides, including all possible single amino acid substitutions to alanine/glycine for large side chains or to tryptophan for small side chains and N- and C-terminal truncations of Qsp1 (Table S8). None of the truncations display colony phenotype-altering activity (data not shown). All but three positions in Qsp1 are sensitive to substitution (Figure S3B).

Modulation of Canonical Virulence Factors by Qsp1

Other cryptococcal mutants have been reported to form dry colonies, mostly mutants lacking capsule (Chang and Kwon-Chung, 1994, 1998) or mutants that are hyphal (Walton et al., 2006). However, *qsp1* cells grow as yeasts and stain normally with an anti-capsular monoclonal antibody (mAb339) (data not shown). Therefore, we subjected wild-type and *qsp1* cells to capsule-inducing conditions and quantified capsule size. At 37°C, *qsp1* mutants formed capsules that were similar to wild-type. At room temperature, *qsp1* mutant cells had larger capsules than wild-type (Figures 3C and S3C).

Cell wall-associated melanin is another well-studied virulence factor of *C. neoformans*. The *lac1 lac2* mutant lacks both laccase enzymes responsible for producing melanin and therefore remains light-colored on L-DOPA plates. We found that *qsp1* mutants displayed altered melanization at 37°C, and the phenotype depended on the condition of cultures used to inoculate the assay. Using cells from saturated cultures, we observed that *qsp1* cells are hypomelanized. This defect could partially be rescued by addition of synthetic Qsp1. Conversely, *qsp1* cells grown on filters on solid medium and then transferred to L-DOPA plates are hypermelanized (Figure 3D). *lac1* cells are attenuated for virulence, and mouse lungs infected with this mutant show altered leukocyte recruitment and cytokine levels (Qiu et al., 2012). As the *qsp1* mutant does not display such changes, it is unlikely that *qsp1*'s attenuated virulence is due solely to changes in melanization.

Qsp1 Impacts Multiple Secreted Endoprotease Activities

A secreted metalloprotease was shown to be required for cryptococcal invasion of the central nervous system (Vu et al., 2014). We have recently identified a secreted aspartyl protease required for cryptococcal virulence (S.C.C., P.A. Dumesic, C.M.H., A.J. O'Donoghue, L. Pallova, P. Majer, H.D.M., and C.S.C., unpublished data). In work to be described elsewhere, we developed fluorogenic peptide substrates (Figure S3D) to assay secreted serine, aspartyl, and metallo endoprotease activities in *C. neoformans*. We identified the genes encoding each activity (S.C.C., P.A. Dumesic, C.M.H., A.J. O'Donoghue, L. Pallova, P. Majer, H.D.M., and C.S.C., unpublished data). Then, we examined the impact of Qsp1 on these activities in *C. neoformans*. In wild-type culture supernatants from tissue culture conditions, secreted metalloprotease and serine endoprotease activities dominate. Conversely, secreted aspartyl endoprotease activity dominates when cultures are grown in minimal media. *qsp1* mutants have decreased secreted aspartyl endoprotease activity in minimal media and increased secreted serine endoprotease and metalloprotease activities in tissue culture conditions compared to wild-type. Growing cultures with synthetic Qsp1 restored serine and aspartyl endoprotease activities (Figure 3E).

Qsp1 Regulates the Transcriptome at High Cell Density

To determine whether Qsp1 regulates gene expression, we performed a low-coverage RNA-seq experiment comparing wild-type and *qsp1* cells grown in log-phase and saturated culture conditions. We observed 100 genes differentially expressed in *qsp1* compared to wild-type saturated cultures but only five differentially expressed genes in log-phase cultures (Figure 4A and Table S2). This indicates a density-dependent impact on the transcriptome. We performed a high-coverage RNA-seq experiment comparing saturated wild-type and *qsp1* cultures and identified 1,749 differentially expressed genes. This set of Qsp1-regulated genes is enriched for genes that encode proteins of the cell wall/extracellular proteome of *C. neoformans* (Figure 4B) (Eigenheer et al., 2007). Also, many *QSP1*-regulated genes are known or predicted to be involved in cell wall biogenesis or regulation (Figure S4A).

qsp1 Mutants Display Density-Dependent Cell Wall Defects

We assessed cell wall structure of wild-type and *qsp1* cells by electron microscopy as described previously (O'Meara et al., 2013) (Figure 4C). We observed that *qsp1* cells

display significantly thinner cell walls than wild-type (Figure 4D). Previous studies have shown that wild-type cell walls are organized into distinct layers, typical of fungi (Doering, 2009; Free, 2013). The number and width of layers is dynamic and condition dependent (Sakaguchi et al., 1993; Shepardson and Cramer, 2013). We observed distinct layers in 90% of the wild-type cell walls examined. In contrast, we observed layers in about 60% of *qsp1* cell walls (Figure 4E).

As described, *qsp1* mutants do not display plating defects on agar containing the cell wall perturbants Calcofluor White, caffeine, or SDS. However, we reasoned that this assay would not capture phenotypes that are specific to high cell density, since single cells are deposited. Instead, we analyzed cell wall function in dense conditions by incubating saturated cultures with cell wall stressors and examining the viability of cells after incubation by plating. *qsp1* mutants display a loss of viability relative to wild-type when exposed to high concentrations of SDS and caffeine (Figure 4F). This phenotype could be rescued if cells were grown with synthetic Qsp1 before the assay. In contrast, exponentially growing cultures display no difference in viability between the two genotypes. We did not observe increased killing of saturated *qsp1* cultures upon acidified nitrite or hydrogen peroxide treatment (Figures S4B and S4C). Since the *qsp1* capsule and cell wall phenotypes are reminiscent of mutants lacking chitin or chitosan, we quantified chitin and chitosan content of cell walls but found that wild-type and *qsp1* mutants display similar profiles (Figure S4D).

A Transcription Factor Functions Downstream of Qsp1

Of the 100 most-changed transcripts in *qsp1* cells, we observed a significant enrichment for regulatory targets of the transcription factor Liv3 (Figure S4E). To test whether Liv3 mediates some or all of the Qsp1 response, we constructed a *qsp1 liv3* deletion mutant. We performed RNA-seq of *qsp1* and *qsp1 liv3* both grown to saturation in either the presence or absence of synthetic Qsp1. We identified 851 differentially expressed genes in *qsp1* mutants grown in the presence of synthetic Qsp1 compared to *qsp1* mutants grown in the absence of synthetic Qsp1 (Figure 4G and Table S2). These strongly overlap with genes identified when wild-type and *qsp1* cells are compared. In contrast, we observed only 282 differentially expressed genes in the *qsp1 liv3* mutant grown with synthetic Qsp1 compared to without (Figure 4G). Thus, Liv3 mediates a majority (66%) of the transcriptional response to Qsp1.

We assessed whether *liv3* mutants share the dry colony phenotype of *qsp1* and found that *liv3* mutants have a dry colony phenotype at room temperature (data not shown). The phenotype is most pronounced at 30°C (Figure S4F). *liv3* mutants form smooth colonies when grown at 37°C (Figure S4F). Supporting a downstream role for Liv3, *liv3* mutants cannot be complemented by a nearby patch of wild-type cells (Figure S4G) or by the synthetic Qsp1 peptides (Figure S4H). In addition, expression of Liv3 using the copper-regulated p*CTR4* promoter (Ding et al., 2013) bypasses the dry colony phenotype of *qsp1* cells (Figure 4H).

Forward Genetic Screen for Mutants in the QSP1 Pathway

We sought additional components of the Qsp1 signaling pathway. We attempted to construct a deletion mutant for each of the 1,010 direct targets of Gat201, Gat204, and/or Liv3. We screened the 705 mutants successfully created for a dry colony phenotype. We profiled melanin, capsule, and cell morphology for the nine mutants identified (Table S3). One of the mutants is *liv3*. We screened an additional 1,407 mutants produced while constructing a deletion collection for *C. neoformans* (2,112 strains constructed and screened). Of these, we identified six additional mutants with the dry colony phenotype (Table S3). These include three known acapsular mutants (*cap10*, *cap60*, and *cap64*; Chang and Kwon-Chung, 1998, 1999; Chang et al., 1996) and one mutant defective in capsule attachment (*pbx1*; Kumar et al., 2014; Liu et al., 2007).

Pqp1 Is a Protease Required for Qsp1 Processing

Among the mutants with a dry colony phenotype was a predicted secreted subtilisin-like serine protease encoded by *CNAG_00150* (Figure 5A). Only a peptide corresponding to the C-terminal 11 amino acids of the predicted proQsp1 peptide was detected in culture supernatants in prior studies. Therefore, we hypothesized that *CNAG_00150* encodes the protease responsible for cleaving the predicted proQsp1 species. We refer to this protease as proQsp1 Protease 1 (Pqp1). The colony phenotype of a *qsp1 pqp1* mutant can be complemented by synthetic Qsp1 but not by synthetic proQsp1 (Figure 5B). Thus, cells lacking *PQP1* are defective in processing synthetic proQsp1 into Qsp1 but are able to respond to Qsp1 itself.

Using synthetic Qsp1 and proQsp1, we determined that while Qsp1 antisera recognize both peptides, they cannot be distinguished by SDS-PAGE and immunoblotting. Therefore, we used one-dimensional isoelectric focusing gel electrophoresis (IEFGE). In both rich and minimal media, we identified a band with identical mobility as synthetic Qsp1 in wild-type supernatants but not in *pqp1* or *qsp1* supernatants. In *pqp1* supernatants from rich media, we observed a band migrating at a more basic pI, but its mobility differed from synthetic proQsp1 (Figure 5C). The larger band appearing in *pqp1* supernatants may be a modification to the predicted precursor. Since synthetic proQsp1 complements the colony phenotype of *qsp1* cells, this modification may not be necessary. To test this explicitly, we added synthetic proQsp1 to *qsp1* cultures and analyzed the products by IEFGE and immunoblotting (Figure S5A). Consistent with processing, we observed the appearance of a band that comigrated with synthetic Qsp1. This band did not appear when synthetic proQsp1 was added to *qsp1 pqp1* cultures (Figure 5D).

We analyzed peptide species in wild-type culture supernatants by LC-MS/MS and identified peaks (Figure 5E) and spectra (Figure 5F) corresponding to the Qsp1 species. However, we did not observe peaks or spectra corresponding to Qsp1 in *qsp1* or *pqp1* culture supernatants (Figure 5E), supporting the model that Pqp1 is required for maturation of the endogenous precursor. No peak corresponding to proQsp1 was identified in culture supernatants from any genotype examined, again consistent with a modification of this putative precursor.

Pqp1 was previously identified as an extracellular, but surface-associated, protein (Eigenheer et al., 2007). We quantified proQsp1 processing activity by intact cells and cell-free supernatants using a proQsp1-like substrate (Figure S5B). Approximately 90% of protease activity against this substrate is cell associated (Figure 5G). A small amount of residual cleavage (10%–20% of wild-type) was found in *pqp1* mutant cells, presumably due to one or more unknown proteases that cleave at some position in the substrate. We analyzed nine additional mutants lacking predicted secreted proteases (Table S4); none display a dry colony phenotype.

A Predicted Oligopeptide Transporter Required for Peptide Sensing

We searched downstream components in the *QSP1* signaling pathway by analyzing published mutants in G protein-coupled receptors and two-component receptors (Table S4). None showed a dry colony phenotype (data not shown). However, our screen identified a gene encoding a predicted oligopeptide transporter, which we named *OPT1* (Figure S6A). The *QSP1* and *OPT1* genes are located next to each other and both display binding of Gat201, Gat204, and Liv3 to their promoters (Figure 6A). Significantly, *opt1* mutants show nearly all of the phenotypes of *qsp1* mutants, including dry colonies and increased capsule size at room temperature (Figures 6B, 6C, and S6B). *opt1* mutants display the same melanization pattern as *qsp1* mutants (Figure S6C). Additionally, *opt1* mutants show viability defects in saturated cultures exposed to SDS and caffeine that are indistinguishable from those of *qsp1* mutants (Figures 6D and S6D). Likewise, *opt1* mutants have no plating defects on any stress conditions tested (Figures S6E and S6F). Finally, *opt1* mutants display no differences in unopsonized (Figure 6E) or opsonized (Figure 6F) uptake by BMDMs compared to wild-type. *opt1* mutants have a 2-fold intracellular proliferation defect after phagocytosis by BMDMs, similar to *qsp1* (Figure 6G).

Although the phenotypes of *opt1* mutants closely mirror those of *qsp1* mutants, *opt1* mutants differ in two important respects: (1) they can produce Qsp1 and (2) they fail to respond to Qsp1. The dry colony phenotype of *opt1* mutants cannot be complemented by wild-type patches (Figure 6H). However, *opt1* patches can complement nearby *qsp1* colonies, indicating that *opt1* mutants still produce Qsp1. By ELISA, we found that *opt1* mutants produce wild-type amounts of Qsp1 in culture supernatants at cell densities up to OD 4.5. At higher densities, we observed less accumulation (Figure S6G). However, the colony phenotype of *opt1* mutants and the sensitivity of saturated *opt1* cultures to cell wall stressors are not complemented by addition of synthetic Qsp1 (Figures 6I and S6H). Therefore, these phenotypes are not due to a defect in Qsp1 production.

Qsp1 Functions Intracellularly

The requirement of Opt1, a predicted oligopeptide transporter, for the response to Qsp1, but not for its production, suggests that Qsp1 may be imported to act within cells. Thus, we replaced the *QSP1* gene with an N-terminal fusion of ubiquitin to the mature Qsp1 peptide. This construct expresses a precursor (iUbi-Qsp1) predicted to be cleaved by endogenous ubiquitin proteases and release the mature Qsp1 peptide cytoplasmically (Varshavsky, 2000) (Figure 7A). We controlled the expression of this precursor using the *pCTR4* promoter. These strains did not produce detectable extracellular Qsp1 when assayed using

confrontation assays (data not shown). Strikingly, expression of the iUbi-Qsp1 construct by growth on inducing media produces a wild-type colony phenotype. Repression of the gene recapitulates the dry colony phenotype of the *qsp1* mutant (Figure 7B). The media conditions used to regulate the p*CTR4* promoter do not alter the colony phenotypes of wild-type or *qsp1* cells (Figure 7B). Furthermore, we observed that induction of iUbi-Qsp1 also complements the capsule size changes (Figure 7C) and SDS/caffeine sensitivity (Figure 7D) of the *qsp1* mutant. The requirement of *OPT1* for Qsp1 action and the ability of iUbi-Qsp1 to complement multiple *qsp1* phenotypes indicates an intracellular site of action for Qsp1.

DISCUSSION

We describe a peptide-based cell-cell signaling system required for the virulence of *Cryptococcus neoformans*, the most common cause of fungal meningitis and a major driver of mortality in HIV/AIDS. This work provides clear evidence that autoregulatory signaling is required for fungal virulence. In addition, our studies reveal that the signaling peptide, Qsp1, is formed by extracellular proteolysis of a precursor but that sensing of Qsp1 occurs intracellularly, a signaling mechanism without precedent in eukaryotes.

A Secreted Peptide Required for Virulence in a Eukaryotic Pathogen

We found that *qsp1* cells are substantially attenuated in a murine inhalation model. We ruled out a number of possibilities for the attenuated virulence phenotype of *qsp1* cells, including stress sensitivity and general growth defects. Additionally, our studies did not reveal any detectable change in the dysfunctional Th2 polarization that characterizes cryptococcal disease. As macrophages are key determinants of the outcome of *C. neoformans* infection, we compared the interactions of wild-type and *qsp1* cells with primary macrophages and observed a defect in intracellular accumulation of *qsp1* cells compared to wild-type. The intracellular accumulation defect of *qsp1* cells likely contributes to its virulence defect as other cryptococcal mutants with decreased accumulation inside macrophages are attenuated in an intranasal model of infection (Evans et al., 2015; Liu and Xue, 2014; Ma et al., 2009). An intriguing possibility is that confinement in the phagolysosome leads to the accumulation of high local concentrations of Qsp1 (Carnes et al., 2010), thereby triggering an adaptive program. We observed no changes in the phagosomal acidification of macrophages infected with *qsp1* mutants when compared to those infected with wild-type, indicating that phagosome maturation per se is not blocked. Any number of antimicrobial mechanisms that act after phagosome acidification may be interdicted by Qsp1 signaling. Alternatively, the Qsp1 system may simply enable fungal mechanisms that resist killing such as cell wall remodeling.

Qsp1 Mediates Autoregulatory Signaling

Our identification of phenotypes for *qsp1* cells enabled us to demonstrate that the 11 amino acid mature Qsp1 peptide is a cell-cell signaling molecule produced by *C. neoformans* that acts on cryptococcal cells. RNA-seq analysis confirms this view that Qsp1 acts on cryptococcal cells themselves and that the impact of *QSP1* on the transcriptome is density dependent. These observations are consistent with our finding that only saturated cultures show sensitivity to stress conditions. We observed Qsp1-dependent changes in all three

major endoprotease activities. Qsp1 promotes an aspartyl protease activity that we have found in other studies to be required for virulence (S.C.C., P.A. Dumesic, C.M.H., A.J. O'Donoghue, L. Pallova, P. Majer, H.D.M., and C.S.C., unpublished data). In contrast, Qsp1 limits metalloprotease and serine protease activities, which may be deleterious at high cell densities. The outer surface of fungi is the cell wall, a dynamic and highly organized protective organelle that impacts virtually all transactions with the environment. It was therefore notable that genes encoding extracellular/cell wall proteins were significantly enriched among those regulated by *QSP1* at high cell density. In addition, *PLB1*, a gene encoding a phospholipase required for cell wall integrity and virulence, was among genes activated by Qsp1 (Cox et al., 2001; Siafakas et al., 2007). Indeed, our studies of cell wall ultra-structure revealed changes in thickness and organization in *qsp1* mutants. Furthermore, we observed a concomitant sensitivity of saturated *qsp1* cultures to killing by cell wall stressors. Our studies provide a clear example in fungi of a mutant that does not synthesize a signaling molecule and exhibits a density-dependent phenotype.

Production of the Signal Requires a Cell-Associated Serine Protease

We used the *qsp1* dry colony phenotype to perform a forward genetic screen for additional components of the pathway. The analysis identified a predicted subtilase-like serine protease, *PQP1*. Our data suggest that Pqp1 is the protease that cleaves a Qsp1 precursor to produce the 11 amino acid mature Qsp1 peptide. (1) *pqp1* cells fail to produce Qsp1. (2) While *qsp1*'s colony phenotype can be complemented by either Qsp1 or a predicted precursor, proQsp1, only Qsp1 can complement *pqp1* cells. (3) An exogenously added predicted precursor (proQsp1) can be processed by a cell-associated activity present in wild-type and *qsp1* strains but not mutants lacking Pqp1. (4) Deletion of genes coding for all other predicted secreted endoproteases did not yield any mutants with a dry colony phenotype.

Reception Requires a Predicted Oligopeptide Transporter: Intracellular Action of Qsp1

Additional mutants isolated based on their dry colony phenotypes were not complemented by synthetic Qsp1, making them candidates for factors involved in the response to Qsp1. Among these, the *opt1* mutant shares the full constellation of *qsp1* phenotypes. The notable difference is that, rather than being defective in producing the signal, the *opt1* mutant is defective in responding to the signal. We reasoned that Opt1 functions in the response to Qsp1. Opt1 is a member of a family of oligopeptide transporters (Gomolplitinant and Saier, 2011), leading us to hypothesize that Opt1 transports Qsp1 into cells.

We tested this hypothesis by constructing a strain to conditionally express the mature Qsp1 peptide intracellularly (iUbi-Qsp1). As expected, the iUbi-Qsp1 strain does not secrete Qsp1. Nonetheless, it displays a wild-type colony phenotype upon induction, indicating that expression of mature Qsp1 intracellularly complements the dry colony phenotype. Significantly, we observed that iUbi-Qsp1 also complements additional phenotypes of the *qsp1* mutant: capsule size alterations and sensitivity to cell wall stressors. Together with the requirement of a predicted oligopeptide transporter for the cellular response to Qsp1, these data indicate that Qsp1 functions intracellularly after being imported.

Evolution of Peptide-Based Autoregulatory Signaling

QSPI and *OPT1* form a two-gene cluster in *C. neoformans*, raising the possibility that they were recently acquired by horizontal gene transfer. However, phylogenetic analysis (Figure S6I) indicates that *OPT1* was present in the ancestor of present day *Tremellaceae*, the basidiomycete family that harbors the *Cryptococcus* species complex. Moreover, genes encoding candidate Qsp1 precursors lie adjacent to *OPT1* in *Filobasidiella depauperata*, *Tsuchiyaea wingfieldii*, and *Cryptococcus amylolentus* (Table S5). Thus, the Qsp1 system likely evolved prior to the emergence of the pathogenic *Cryptococcus neoformans*/*Cryptococcus gattii* species complex. This conclusion is germane because, while bacterial QS molecules are typically sensed by cell surface receptors, in a handful of Gram-positive species the response to a QS peptide requires a multisubunit oligopeptide permease (Lazazzera and Grossman, 1998; Rutherford and Bassler, 2012). The permease enables import of the peptide and its binding to intracellular receptors. Furthermore, in some of these species, the mature peptide is produced by C-terminal cleavage of a precursor by a cell wall-associated peptidase. As none of the components involved appear to have a proximal bacterial ancestor, this eukaryotic system arose via convergent evolution.

EXPERIMENTAL PROCEDURES

Yeast strains

Yeast strains used in this study are listed in Table S6. All *C. neoformans* strains were derived from the described parent using published procedures (Chun and Madhani, 2010). Gat201, Gat204, and Liv3 strains are derived from the CM18 parent for consistency with previous publications. The rest of the strains reported are derived from the KN99 α wild-type since that isolate is more closely related to the H99 clinical isolate (Janbon et al., 2014).

Chromatin Immunoprecipitation and RNA-Seq Cultures

C. neoformans cultures of 50 ml were grown overnight. 50 ODs were washed 1 \times with water and resuspended in 15 ml DMEM. This suspension was transferred to 150 mm \times 25 mm tissue culture-treated plates already containing 5 ml DMEM. Cultures were incubated sitting at 37°C, 5% CO₂ for 8 hr for ChIP-seq and 24 hr for RNA-Seq.

Chromatin Immunoprecipitation

C. neoformans cultures were grown and immunoprecipitation was performed as previously published (Chun et al., 2011). Libraries were prepared for high-throughput sequencing and analyzed using Bowtie1 (Langmead, 2010), SAMtools (Li et al., 2009), and the MACS2 peak caller (Zhang et al., 2008) as described in the Supplemental Experimental Procedures. Overlap between biological replicates was high (Table S7), but for subsequent analysis we only included genes whose promoters had at least one ChIP-seq peak in each of the three biological replicates analyzed. Binding motifs were analyzed using Gimsan (Ng and Keich, 2008).

RNA Expression Profiling

Total RNA was isolated and libraries prepared as described previously (Dumesic et al., 2015). Analysis was performed using Tophat (Trapnell et al., 2012) and the TMM algorithm (Oshlack et al., 2010) or DESeq (Anders and Huber, 2010) as described in the Supplemental Experimental Procedures.

Mass Spectrometry

Peptides were enriched in samples from culture supernatants by precipitation with 90% methanol and 1% acid. LC-MS/MS identification was performed as described in the Supplemental Experimental Procedures.

Colony Phenotype Patch Complementation Assays

All assays were performed at room temperature unless otherwise specified. Peptides for Qsp1 mutational analysis (Table S8) were synthesized by Peptide 2.0.

ELISA and Immunoblotting Assays

Cell-free supernatants were analyzed by ELISA or immunoblotting as described in the Supplemental Experimental Procedures using 1:1,000 diluted anti-Qsp1 serum.

Fluorescent Substrate Protease Activity Assay

Protease assays were performed with the specified substrates as previously described (Jambunathan et al., 2012).

Growth, Viability, Capsule, and Melanin Assays

Growth on various medias, capsule, and melanin were analyzed as previously described (Liu et al., 2008) with minor changes described in the Supplemental Experimental Procedures.

Cell Wall Analysis

24 hr DMEM cultures were analyzed as previously described (Reese et al., 2007).

Intranasal Infection Model

Mouse lung infections were performed as previously described (Chun and Madhani, 2010), under the supervision of the UCSF IACUC Protocol AN091509.

Rabbit Meningitis Model

Rabbit intrathecal infections were performed as previously described (Perfect et al., 1980), according to the Duke University IACUC protocol A003-14-01.

Leukocyte Recruitment and Cytokine Analysis

Lung leukocytes and cytokines were isolated and analyzed as previously described (Wiesner et al., 2015), according to University of Minnesota IACUC protocols 1010A91133 and 1207A17286.

Macrophage Interaction Analysis

Macrophage uptake and intracellular proliferation assays were performed as previously described (Nicola and Casadevall, 2012) with minor changes described in the Supplemental Experimental Procedures.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Highlights

- Qsp1 is an autoregulatory signaling peptide required for fungal virulence
- Qsp1 controls intracellular growth, cell wall, and protease activities
- Opt1 is a peptide transporter required for Qsp1 action in receiving cells
- Intracellular expression of mature Qsp1 complements *qsp1* phenotypes

In Brief

Bacterial quorum sensing (QS) often controls virulence. Homer et al. reveal a peptide-based QS system in the fungal pathogen *C. neoformans*. This system requires signaling mechanisms in which the exported peptide is imported and acts intracellularly to control aspects of virulence, including growth, cell surface properties, and secreted protease activities.

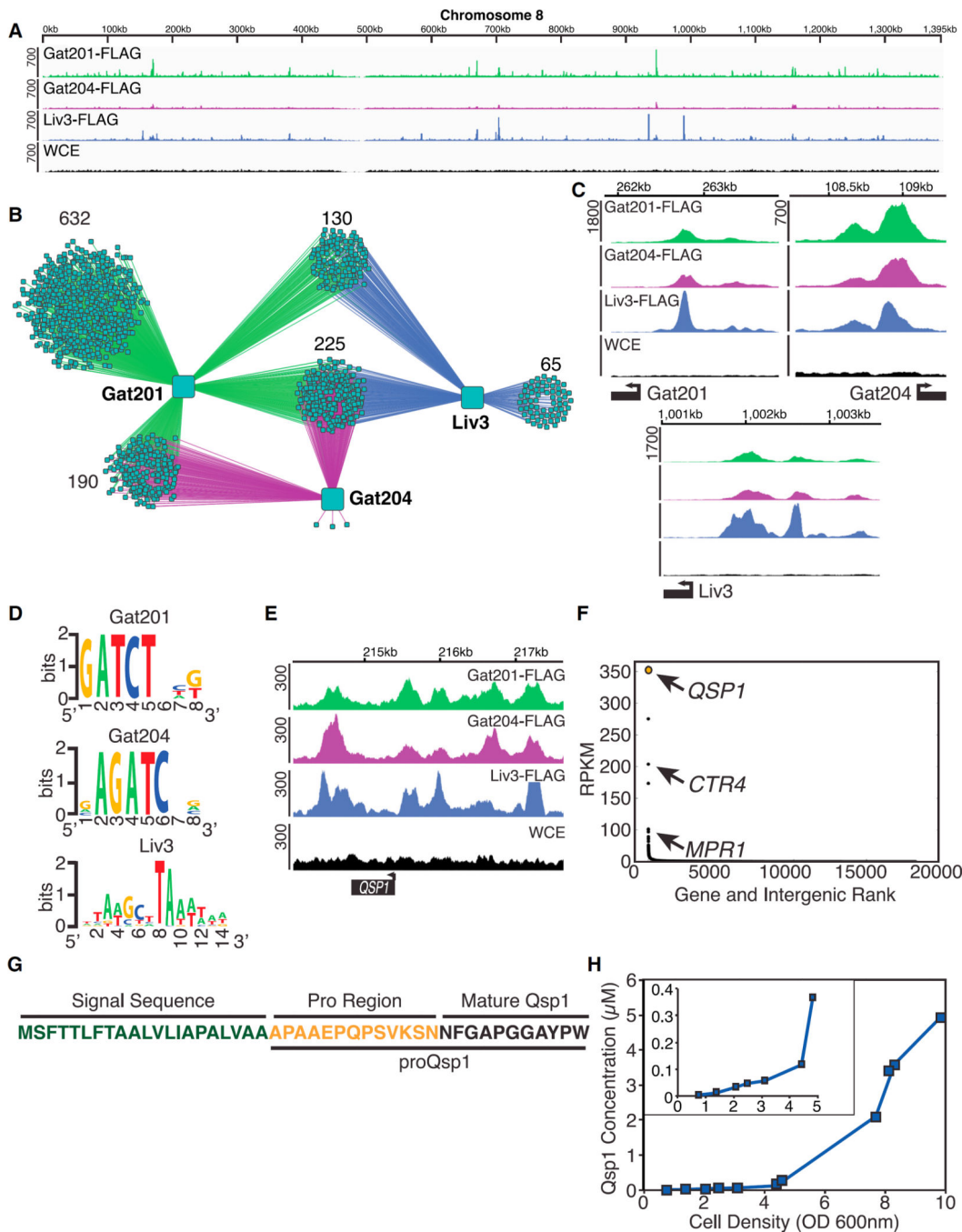


Figure 1. *QSP1* Is a Direct Target of the Gat201-Gat204-Liv3 Regulatory Network

- (A) ChIP-seq data from chr8. y axis: number of reads.
- (B) Network diagram.
- (C) Gat201, Gat204, and Liv3 binding at each other's promoter.
- (D) Binding motifs derived from ChIP-seq datasets.
- (E) ChIP-Seq data at the *QSP1* promoter.
- (F) Rank and RPKM for each gene in a wild-type strain grown in tissue cultures conditions.
- (G) Predicted Qsp1 precursor.

(H) Qsp1 accumulation in wild-type culture supernatants. Inset: Qsp1 accumulation at lower cell densities.

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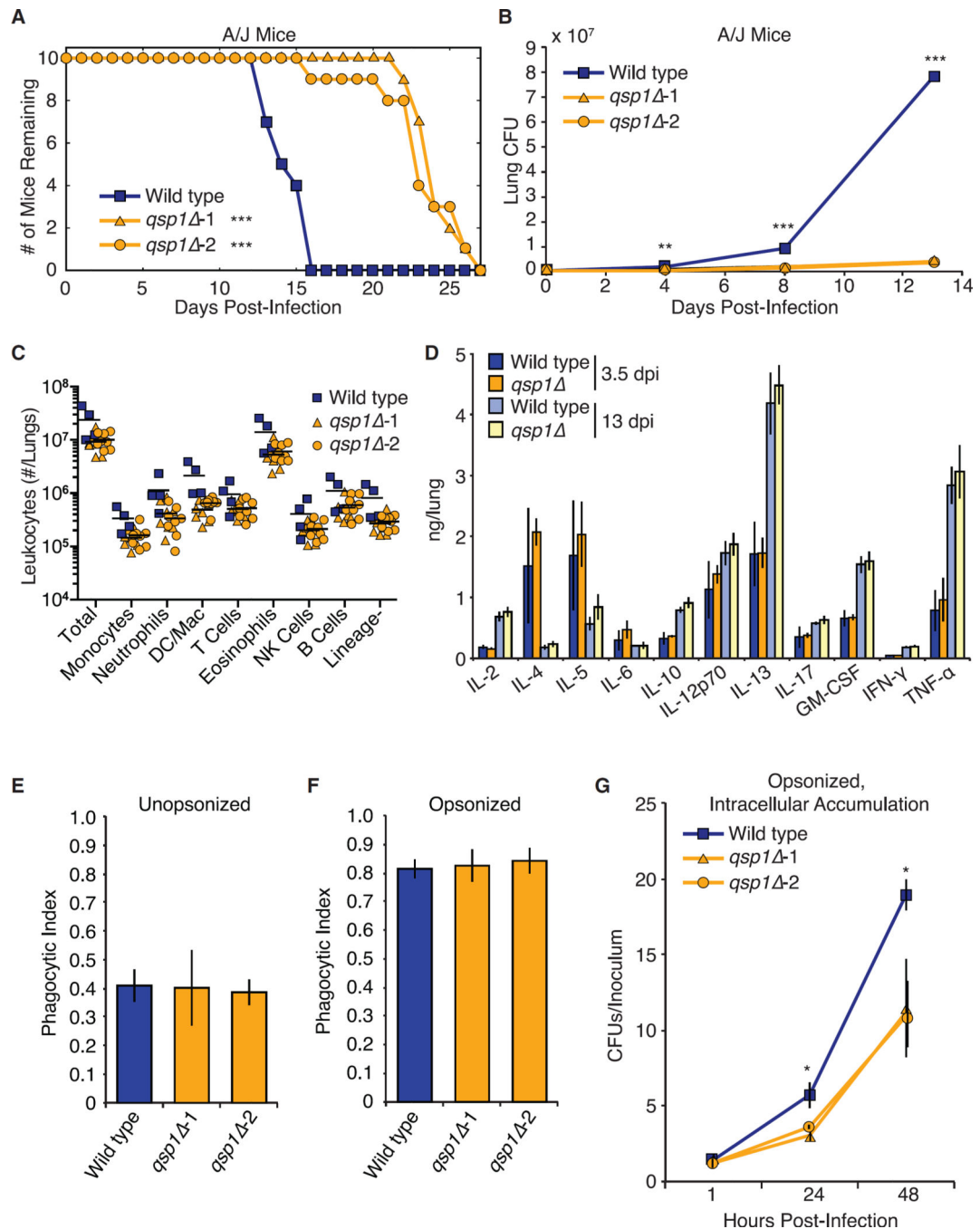


Figure 2. Qsp1 Is Required for Virulence and Accumulation within Macrophages

- (A) Survival analysis. 10 A/J mice infected per genotype. **p < 10⁻⁴ (log-rank test).
 (B) Lung burden analysis. 9 A/J mice infected per genotype. **p < 10⁻³, ***p < 10⁻⁴ (t test).
 (C) Leukocyte recruitment. 4 C57Bl/6 mice infected with wild-type, 8 C57Bl/6 mice infected per *qsp1* strain.
 (D) Cytokine analysis. 10 C57Bl/6 mice infected per genotype. Error bars are SD.

(E and F) Phagocytic index of unopsonized (E) and opsonized (F) *C. neoformans*. >1,200 BMDMs quantified per genotype. Error bars are SD.

(G) Intracellular accumulation. CFUs from BMDM cell lysates, normalized to starting inoculum. Error bars represent 95% confidence intervals constructed by bootstrapping. * $p < 0.05$ (bootstrap).

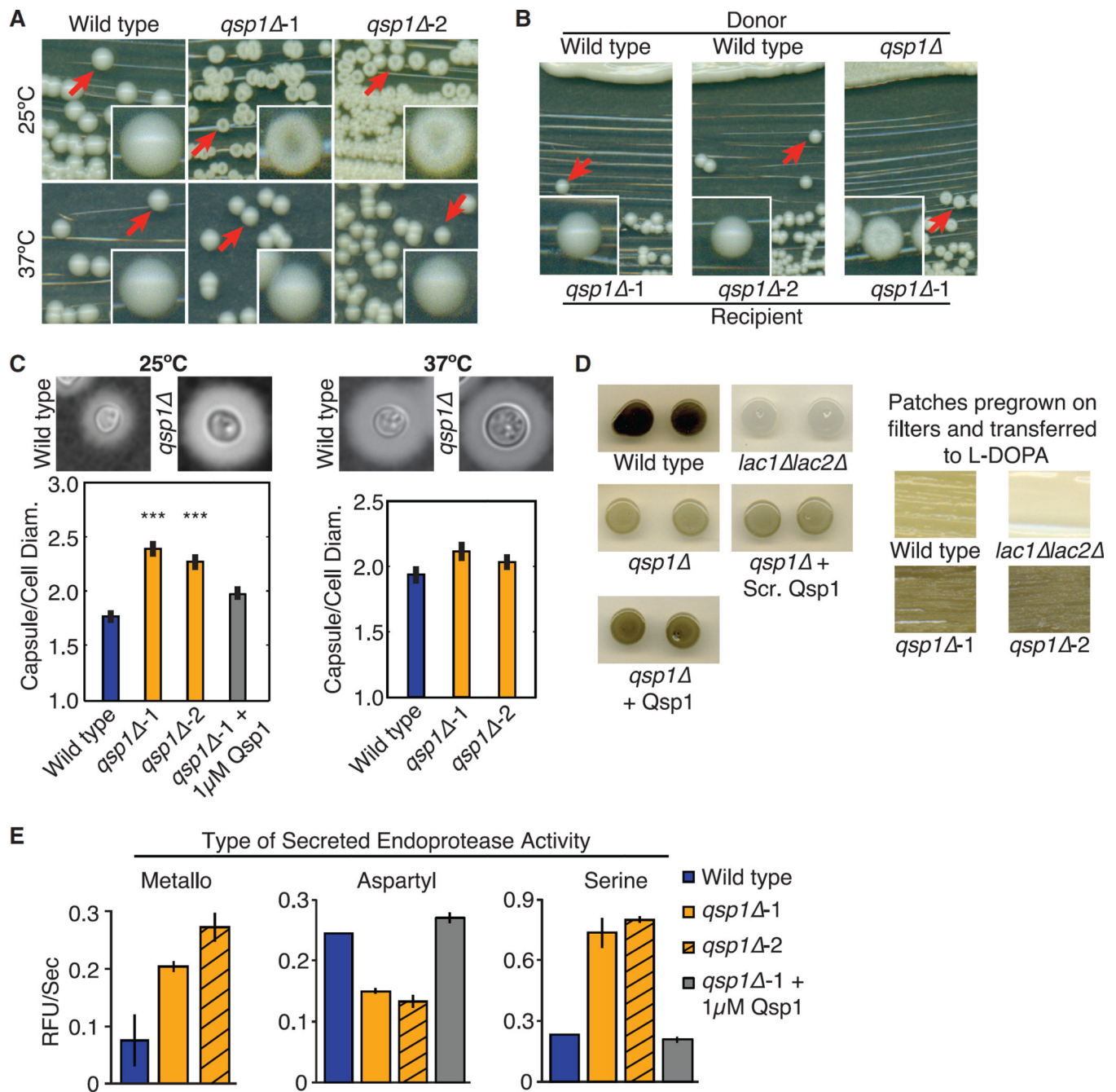


Figure 3. Impact of Qsp1 on Virulence-Related Phenotypes

(A) Colony morphology.

(B) Complementation assays.

(C) Capsule quantification. Cryptococcal capsule assessed using India ink. *** $p < 10^{-5}$ (t test). Error bars are SE > 100 cells per genotype quantified.

(D) Melanin assay. Left: cells grown in liquid culture with or without synthetic peptides before spotting. Right: patches grown on filters and transferred to L-DOPA plates.

(E) Secreted protease activities. Error bars are SD.

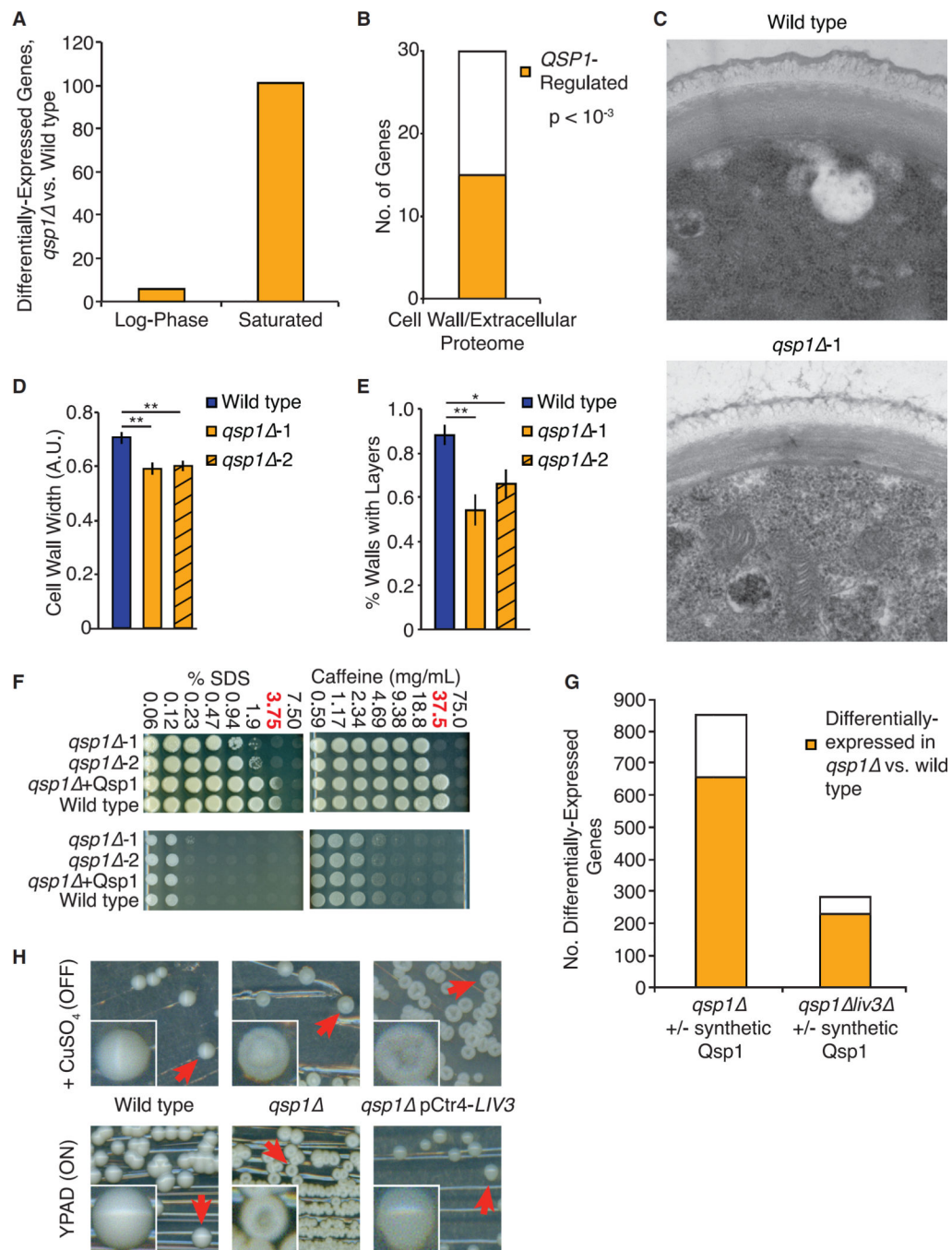


Figure 4. Qsp1 Has Density-Dependent Impacts on Gene Expression and Cell Wall Integrity
 (A) Density-dependent RNA-seq. Expression profiling for *qsp1* and wild-type log-phase and saturated cultures. The number of differentially expressed genes ($p_{adj} < 0.05$ by DESeq) is shown.
 (B) Overlap of *QSP1*-regulated genes with published cell wall and secreted proteome (Eigenheer et al., 2007). *QSP1*-dependent genes are 2.3-fold enriched ($p < 10^{-3}$ by hypergeometric distribution).
 (C) Transmission electron microscopy.
 (D) Bar chart showing cell wall width (A.U.) for Wild type, *qsp1Δ-1*, and *qsp1Δ-2*. ** indicates statistical significance.
 (E) Bar chart showing % Walls with Layers for Wild type, *qsp1Δ-1*, and *qsp1Δ-2*. ** indicates statistical significance.
 (F) Spot assays showing growth of *qsp1Δ-1*, *qsp1Δ-2*, and *qsp1Δ+Qsp1* on media containing increasing concentrations of % SDS and Caffeine (mg/mL).
 (G) Bar chart showing the number of Differentially-Expressed Genes for *qsp1Δ* +/- synthetic Qsp1 and *qsp1Δliv3Δ* +/- synthetic Qsp1. The orange portion represents genes differentially expressed in *qsp1Δ* vs. wild type.
 (H) Micrographs showing cell wall integrity on YPAD (ON) and + CuSO₄ (OFF) for Wild type, *qsp1Δ*, and *qsp1Δ* pCtr4-LIV3. Red arrows indicate cell wall defects.

(D and E) Quantification of cell wall width (D) and percentage of cells with visible cell wall layers (E). ** $p < 0.001$, * $p < 0.01$ (t test). Error bars are SD.

(F) Stress assays. Cultures were incubated with stressors and plated for viability. The concentration where *qsp1* viability is most different from wild-type is shown in red.

(G) *LIV3*-dependent genes. Expression profiling *qsp1* and *qsp1 liv3* saturated cultures grown with or without synthetic Qsp1. Differentially expressed genes shown, with genes differentially expressed in *qsp1* versus wild-type in yellow.

(H) *LIV3* suppression of the *qsp1* dry colony phenotype.

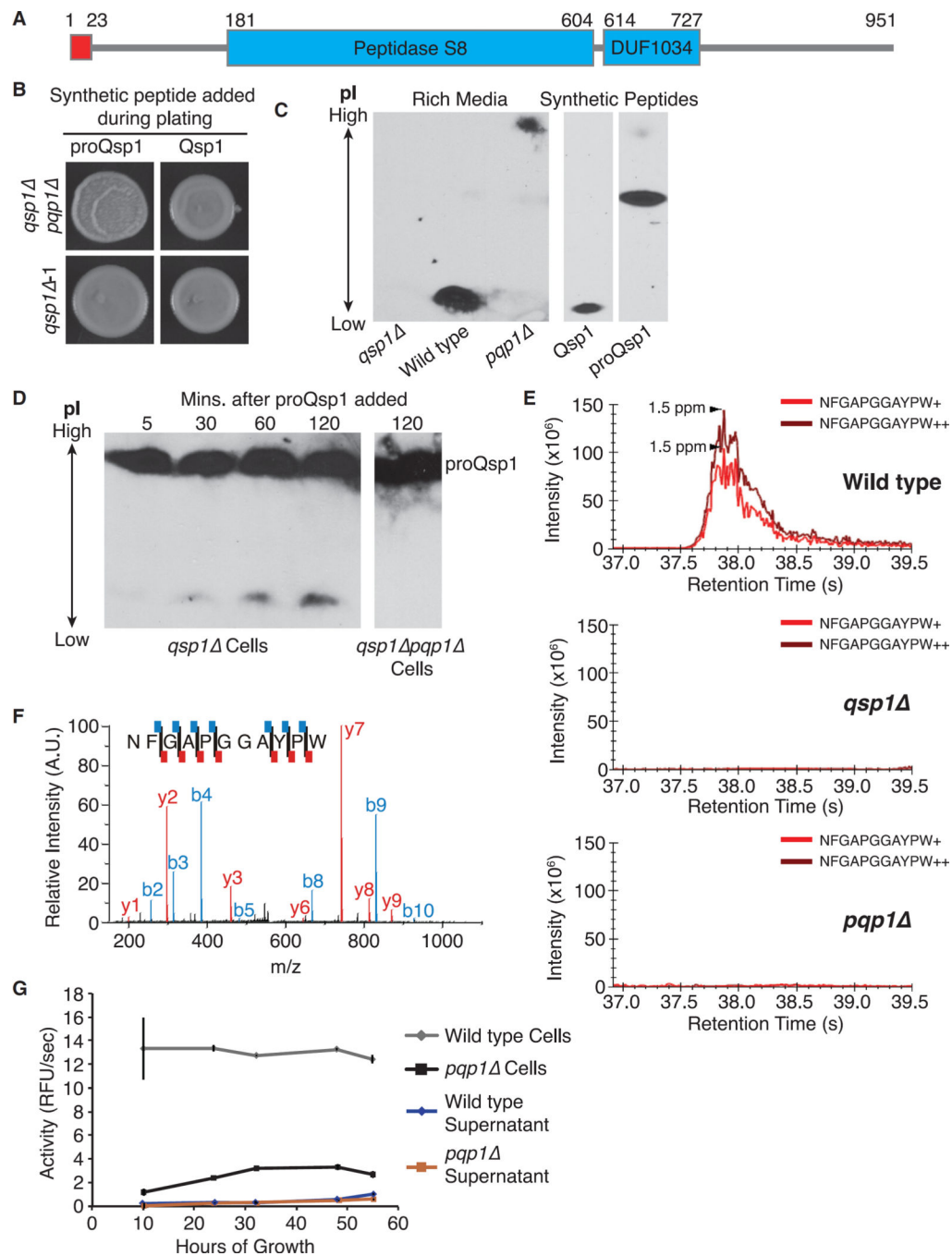


Figure 5. Pqp1 Is Required for Qsp1 Precursor Processing

(A) Pqp1 protein structure.

(B) Spotting assays.

(C) Secretion assays. IEFGE and immunoblotting against Qsp1/proQsp1 in culture supernatants and Qsp1/proQsp1 synthetic peptides. Middle and right panels displaying mobilities of synthetic peptides are from the same blot, with irrelevant lanes removed.

(D) Predicted Qsp1 precursor cleavage assays. Synthetic proQsp1 was incubated with cultures. Supernatants were isolated and then analyzed for cleavage by IEFGE and

immunoblotting against Qsp1/proQsp1. Extracted ion chromatogram of Qsp1 from LC/MS of culture supernatants.

(E) Qsp1 MS2 spectra. Each fragment detected is shown with red (y ions) or blue (b ions) lines on the peptide sequence.

(F) Pqp1 activity assay. Error bars are SD.

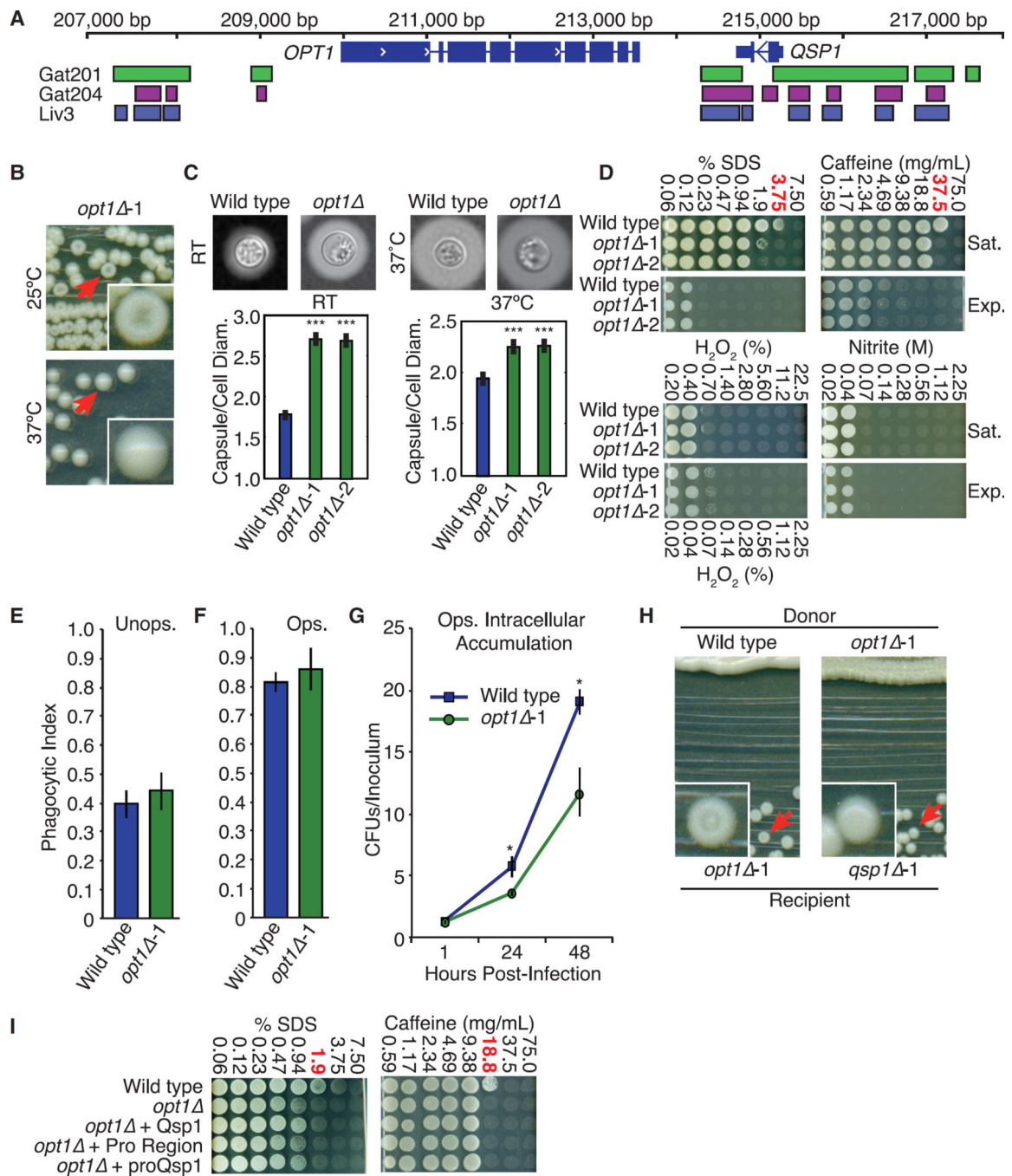


Figure 6. A Predicted Oligopeptide Transporter Is Required for Cells to Respond to Qsp1

(A) Transcription factor binding sites near *QSP1* and *OPT1*.

(B) Colony morphology.

(C) Capsule sizes. Cryptococcal capsule stained with India ink. >100 cells per genotype quantified. *** $p < 10^{-5}$ (t test). Wild-type data from Figure 3 included for comparison. Error bars are SE.

(D) Stress assays. The concentration where *qsp1* viability is most different from wild-type is shown in red. Wild-type data from Figure 4 included for comparison.

(E and F) Phagocytic index of unopsonized (E) and opsonized (F) *C. neoformans*. >1,200 BMDMs quantified per genotype. Wild-type data from Figure 2 included for comparison. Error bars are SD.

(G) Macrophage intracellular accumulation assay. CFUs isolated from BMDMs, normalized to starting inoculum. Wild-type data from Figure 2 included for comparison. Error bars are 95% confidence intervals. * $p < 0.05$ (bootstrap).

(H) Complementation assays.

(I) Stress assays as described in (D).

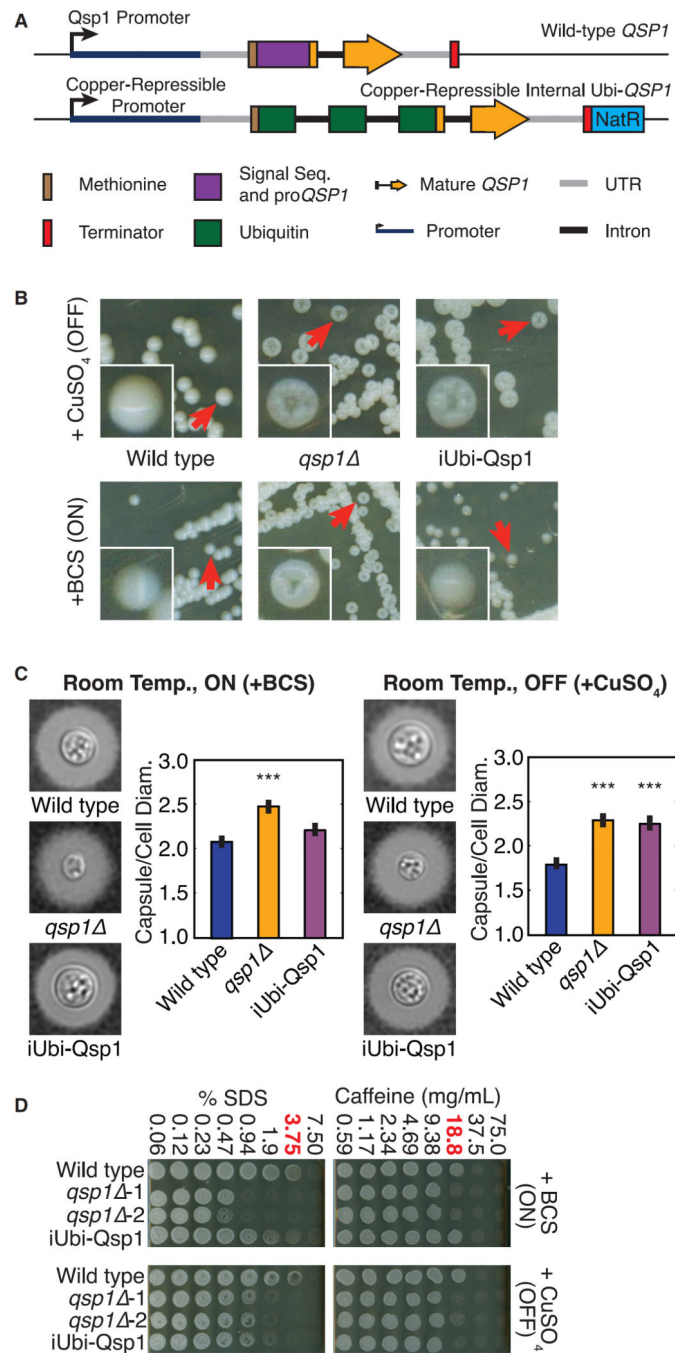


Figure 7. Cytoplasmic Expression of Mature Qsp1 Complements Multiple *qsp1* Phenotypes (A) Schematic of wild-type *QSP1* and iUbi-Qsp1.

(B) Colony morphologies.

(C) Capsule quantification. Cryptococcal capsule stained with India ink. >100 cells per genotype/condition quantified. *** $p < 10^{-5}$ (t test). Error bars are SE.

(D) Stress assays.