

The VirF₂₁:VirF₃₀ protein ratio is affected by temperature and impacts *Shigella flexneri* host cell invasion

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One sentence summary: The ratio of the two virulence-regulatory proteins VirF₂₁:VirF₃₀ increases at low temperature and can dictate the invasive properties of *Shigella flexneri*.

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Abstract.

Shigella spp., the etiological agents of bacillary dysentery in humans, have evolved an intricate regulatory strategy to ensure fine-tuned expression of virulence genes in response to environmental stimuli. A key component in this regulation is VirF, an AraC-like transcription factor, which at the host temperature (37°C) triggers, directly or indirectly, the expression of > 30 virulence genes important for invasion of the intestinal epithelium. Previous work identified two different forms of VirF with distinct functions: VirF₃₀ activates virulence gene expression, while VirF₂₁ appears to negatively regulate *virF* itself. Moreover, VirF₂₁ originates from either differential translation of the *virF* mRNA or from a shorter leaderless mRNA (l1mRNA). Here we report that both expression of the *virF*₂₁ l1mRNA and the VirF₂₁:VirF₃₀ protein ratio are higher at 30°C than at 37°C, suggesting a possible involvement of VirF₂₁ in minimizing virulence gene expression outside the host (30°C). Ectopic elevation of VirF₂₁ levels at 37°C indeed suppresses *Shigella*'s ability to infect epithelial cells. Finally, we find that the VirF₂₁ C-terminal portion, predicted to contain a Helix-Turn-Helix motif (HTH2), is required for the functionality of this negative virulence regulator.

Keywords: *Shigella*, virulence genes, regulation, infection, cell invasion, Shigellosis

Introduction

Enterobacterial pathogens coordinate the expression of virulence factors through complex regulatory networks in order to colonize and disseminate in the host gut epithelium. *Shigella flexneri* bacteria, facultative intracellular microbes causing bacillary dysentery in humans, have become paradigmatic for the study of virulence gene regulation. The expression and combined action of numerous virulence factors, mainly encoded on a large virulence plasmid (pINV), ultimately result in the invasion of colonic epithelial cells in the lower gut (Mattock and Blocker 2017). Subsequently, the bacteria multiply intracellularly and spread to adjacent cells, resulting in cell death and inflammatory destruction of the gut mucosa (Schroeder and Hilbi 2008, Arena et al. 2015). A crucial regulator of the *Shigella* infection process is VirF, an AraC-like transcription factor, responsible for the invasive phenotype (Di Martino et al. 2016a). The synthesis of VirF occurs when *Shigella* senses the transition from the external environment to the human host (Falconi et al. 1998, Prosseda et al. 2004). VirF then triggers a regulatory cascade involving the expression of *virB* and *icsA* genes (Tobe et al. 1993, Tran et al. 2011). VirB activates a second wave of virulence genes involved in the assembly of a type 3 secretion system (T3SS), its effectors (the *ipa*-*spa* operons), and a second AraC-like transcriptional activator, *mxiE* (Le Gall et al. 2005, Parsot 2005, Schroeder and Hilbi 2008). *IcsA*, on the other hand, promotes *Shigella* dissemination across adjacent cells through host actin polymerization

(Bernardini et al. 1989, Lett et al. 1989). Finally, the master regulator VirF also activates some chromosomally located genes (e.g. the spermidine excretion complex MdtJl; the chaperones IbpA, HtpG, DnaK and the protease Lon) whose expression may optimize *Shigella*'s intracellular life style (Barbagallo et al. 2011, Leuzzi et al. 2015).

The activation of *virF* is a key event for the successful invasion and dissemination of *Shigella* within the host epithelium, and is therefore stringently regulated. A multitude of environmental signals (e.g. temperature, pH, osmolarity) affect *virF* expression through several regulatory mechanisms. Some of these mechanisms have been described at the molecular level, as the temperature-dependent expression of the *virF* gene (Falconi et al. 1998, Prosseda et al. 2004). At temperatures below 32°C (non-permissive), the nucleoid-associated protein H-NS tightly binds two sites within the *virF* promoter. This prevents access of the RNA polymerase and therefore leads to *virF* transcriptional silencing. At the permissive host temperature (37°C), relaxation of an intrinsically-curved DNA region, located between the two H-NS binding sites, hampers H-NS binding and favors access of the nucleoid associated protein FIS to one of its binding sites (Falconi et al. 2001, Prosseda et al. 2004). This results in activation of *virF* transcription. VirF subsequently acts as an anti-silencer, counteracting H-NS-mediated repression on for example *virB* and *icsA* promoters. Besides direct binding to the *icsA* promoter, VirF also stimulates *icsA* expression by lowering the intracellular concen-

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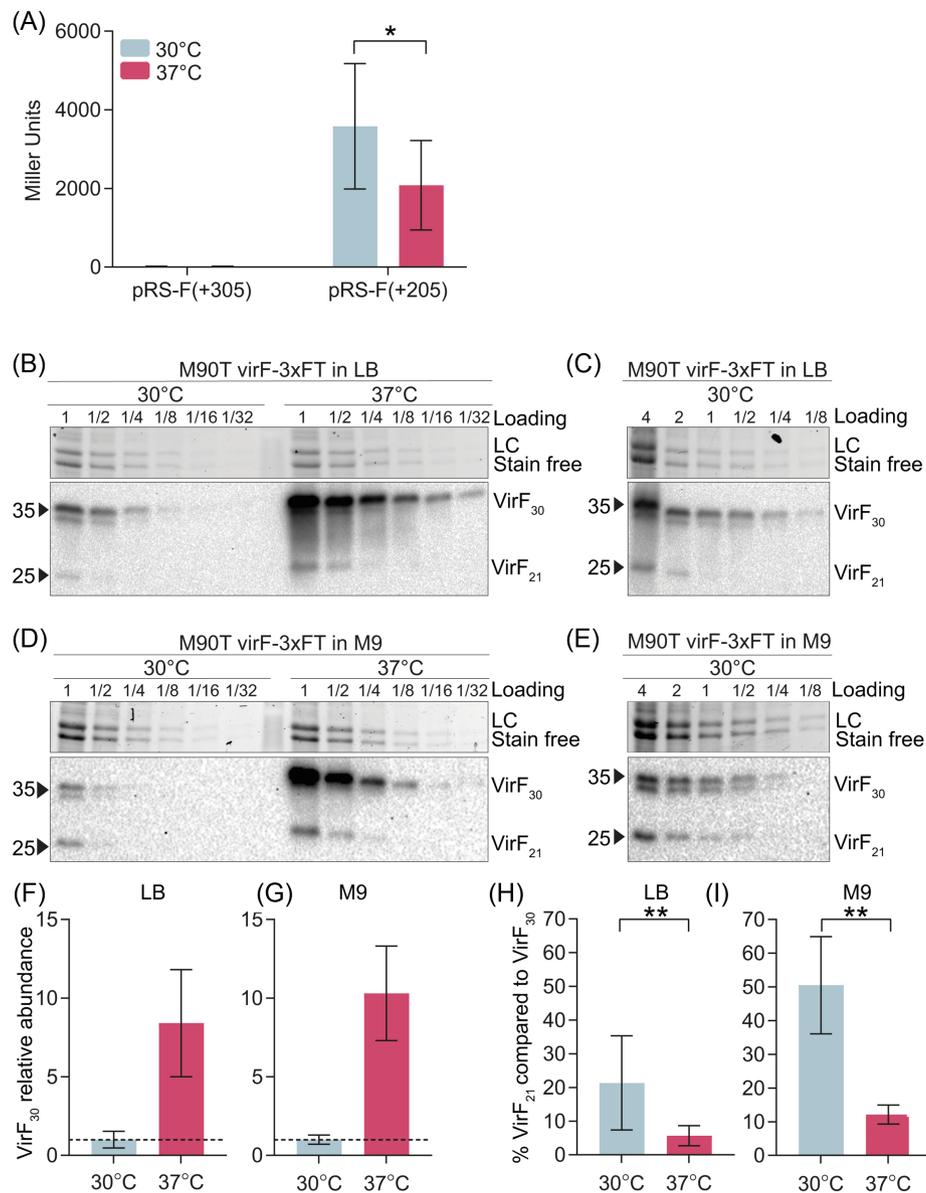


Figure 1. Low temperature (30°C) stimulates *virF₂₁* llmRNA expression and an increased *VirF₂₁*:*VirF₃₀* protein ratio. **(A)** β -Galactosidase activity of the *virF-lacZ* transcriptional fusion pRS-F(+205) containing the internal promoter for the leaderless mRNA. The analysis was performed in *E. coli* DH10b. pRS-F(+305) was used as a negative control. The β -Galactosidase activity was determined after subculture at 30 or 37°C. The activity is reported in Miller Units and represents the mean and standard deviation of 7 (pRS-F(+305)) and 13 (pRS-F(+205)) biological replicates from 3 different experiments. Statistical significance, comparing the β -Galactosidase activity of the pRS-F(+205) fusion at 30 or 37°C, was determined by Mann-Whitney U test, * $P < 0.05$. **(B)** Detection of *VirF₃₀* and *VirF₂₁* at 30° and 37°C in protein extracts of the *Shigella* M90T strain carrying *virF-3xFT* grown in LB medium. A representative western blot with serial dilutions of the protein extracts is shown. **(C)** Detection of *VirF₃₀* and *VirF₂₁* at 30°C in protein extracts of the *Shigella* M90T strain carrying *virF-3xFT* grown in LB medium. A representative western blot in which protein extracts were concentrated to facilitate quantification is shown. **(D)** Detection of *VirF₃₀* and *VirF₂₁* at 30° and 37°C in protein extracts of the *Shigella* M90T strain carrying *virF-3xFT* grown in M9 medium. A representative western blot with serial dilutions of the protein extracts is shown. **(E)** Detection of *VirF₃₀* and *VirF₂₁* at 30°C in protein extracts of the *Shigella* M90T strain carrying *virF-3xFT* grown in M9 medium. A representative western blot in which protein extracts were concentrated to facilitate quantification is shown. **(F)** The relative *VirF₃₀* content in the *Shigella* M90T strain carrying *virF-3xFT* grown in LB at 30° and 37°C was determined by quantification of western blots of serially diluted samples. *VirF₃₀* level at 30°C was set as 1. Shown is the mean and standard deviation of 3 independent experiments. **(G)** The relative *VirF₃₀* content in the *Shigella* M90T strain carrying *virF-3xFT* grown in M9 medium at 30° and 37°C was determined by quantification of western blots of serially diluted samples. *VirF₃₀* level at 30°C was set as 1. Shown is the mean and standard deviation of 4 independent experiments. **(H)** *VirF₂₁* levels in the *Shigella* M90T strain carrying *virF-3xFT* grown in LB at 30°C and 37°C were quantified from western blots of concentrated samples. *VirF₂₁* content was quantified in comparison with the *VirF₃₀* content. Shown is the mean and standard deviation of 8 (30°C) and 6 (37°C) independent experiments. Statistical significance was determined by a Mann-Whitney U test, ** $P < 0.01$. **(I)** *VirF₂₁* levels in the *Shigella* M90T strain carrying *virF-3xFT* grown in M9 medium at 30°C and 37°C were quantified from western blots of concentrated samples. *VirF₂₁* content was quantified in comparison with the *VirF₃₀* content. Shown is the mean and standard deviation of 5 independent experiments. Statistical significance was determined by Mann-Whitney U test, ** $P < 0.01$.

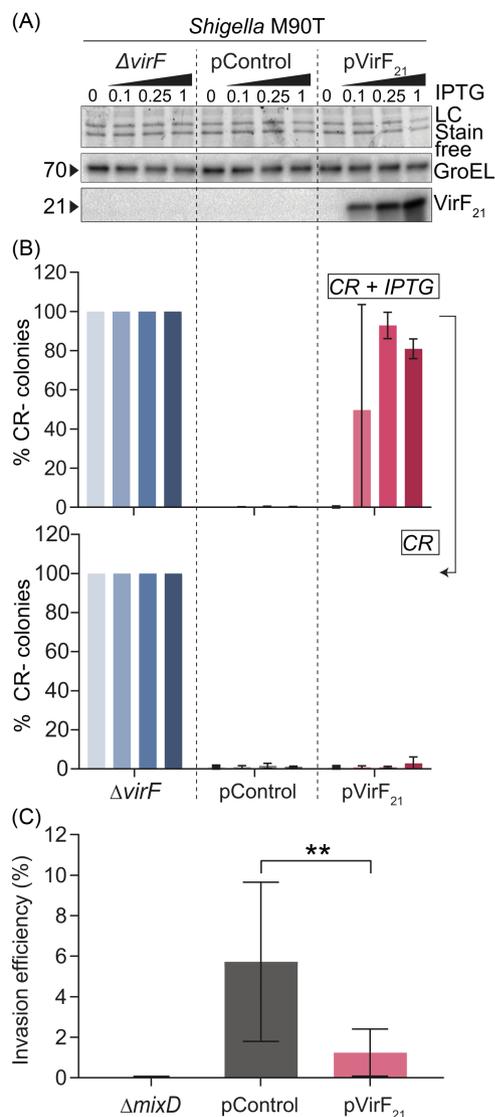


Figure 2. Elevated expression of VirF₂₁ suppresses *Shigella* virulence at 37°C. **(A)** Western blot with VirF antibodies on extracts from a *Shigella* M90T Δ*virF* mutant and M90T strains harbouring pControl (empty vector) or pVirF₂₁, a plasmid carrying the *virF*₂₁ coding sequence under an inducible pTaq promoter. The strains were grown in the presence of increasing concentration of IPTG (0, 0.1, 0.25, 1 mM) to induce VirF₂₁ expression. GroEL protein was detected and used as internal loading control. A loading control using the Stain free method is also shown. **(B)** (upper panel) % CR-colonies upon spreading of the indicated strains on CR plates containing increasing concentrations of IPTG (0, 0.1, 0.25, 1 mM) to induce VirF₂₁ expression. (bottom panel) % CR-colonies upon scraping and re-plating of the colonies obtained on the previous plates, onto new CR plates without IPTG selection. Data come from at least three replicates from two independent experiments. ~200–600 colonies/replicate were examined for the CR phenotype. **(C)** Invasion efficiency of the indicated *Shigella* M90T strains in sub-confluent Caco-2 cell layers. Cells were infected at MOI 100 for 1h, and analysed by selective plating of intracellular bacteria. Shown are CFU data expressed as the percentage of the inoculum retrieved in the intracellular population. Shown are CFU data for 7 (M90T pControl) and 9 (M90T Δ*mixD*, M90T pVirF₂₁) biological replicates from 3 independent experiments. Bars represent mean and standard deviation. Statistical significance was determined by Mann Whitney U test, ***P* < 0.01.

tration of the antisense rna RnaG, known to cause premature *icsA* transcriptional termination (Tran et al. 2011).

In addition to temperature-dependent regulation of *virF* expression, several other regulatory mechanisms have been described, involving e.g. IHF (Porter and Dorman 1997), CpxA/R (Nakayama and Watanabe 1998), and EnvZ/OmpR (Bernardini et al. 1990), as well as specific post-transcriptional tRNA modifications (Durand et al. 1997, 2000, Hurt et al. 2007). More recently, a chromosomal LysR-like transcriptional regulator, YhjC, has been shown to activate *virF* expression, suggesting further cross-talk between chromosomal and pINV located genes (Li et al. 2021). Altogether, these factors contribute to reach a fine-tuned VirF threshold concentration, sufficient to activate *virB* and *icsA* expression and thereby trigger the *Shigella* host cell invasive program (Adler et al. 1989, Dagberg and Uhlin 1992, Prosseda et al. 1998).

Previously, the existence of an additional layer in the regulation of *virF* expression was discovered. The *virF* mRNA itself, through differential translation, in fact gives rise to two forms of VirF protein: VirF₃₀ (30 kDa) and VirF₂₁ (21 kDa). VirF₃₀ acts as primary activator of the virulence gene cascade, whereas VirF₂₁ appears to negatively autoregulate *virF* expression through direct promoter binding (Di Martino et al. 2016b). In addition, VirF₂₁ can originate also from a shorter, leaderless mRNA (llmRNA), transcribed from a gene-internal promoter (Di Martino et al. 2016b). While the molecular interactions of this regulatory loop were defined in the previous study, it had remained unknown which conditions affect VirF₂₁ expression and how this can impact *Shigella*'s host cell invasive phenotype.

Here we characterized the conditions governing *virF*₂₁ llmRNA expression and the overall VirF₂₁:VirF₃₀ protein ratio. We found that at 30°C, a common condition *Shigella* encounters during its extracellular (non-invasive) lifestyle, transcription of the *virF*₂₁ llmRNA from the internal promoter is favored and the VirF₂₁:VirF₃₀ protein ratio is elevated, as compared to the permissive host temperature of 37°C. Moreover, ectopically elevating VirF₂₁ levels at 37°C resulted in a marked and reversible block of the *Shigella* host cell invasive phenotype. The C-terminal part of VirF₂₁ was found to be required for this suppression. We discuss the possible connections between environmental sensing, the fitness costs of virulence gene expression, and VirF₂₁-dependent suppression of the host cell invasive program.

Material and methods

Bacterial strains and general procedures

Strains and plasmids used in this study are listed in Table S1. M90T is a *S. flexneri* serotype 5 strain (Sansonetti et al. 1982). Strain M90T Δ*virF* carries a deletion of the *virF* gene (Leuzzi et al. 2015). Strain M90T *virF*-3xFT carries the 3xFLAG tag sequence at the C-terminus of the pINV-encoded *virF* gene (Leuzzi et al. 2015). Strain M90T Δ*mixD* carries a deletion of the *mixD* gene and has been constructed using the one-step gene inactivation method (Datzenko and Wanner 2000), transforming M90T pKD46 with the PCR product obtained using plasmid pKD4 as template and the oligo pairs *mixD*_delF/*mixD*_delR (Table S2). The plasmids pControl (previously named pGIP7), pVirF₂₁ (previously named pAC-21), pRS-F(+205) and pRS-F(+305) were described previously (Di Martino et al. 2016b, Falconi et al. 2001).

The plasmids pVirF₂₁_I97N, pVirF₂₁_V108A, pVirF₂₁_V145T and pVirF₂₁_Y141stop were obtained by Gibson Assembly (Gibson et al. 2009), ligating *in vitro* synthesized PCR products (purchased from Genewiz) into the BamHI restriction site in pControl using the Gib-

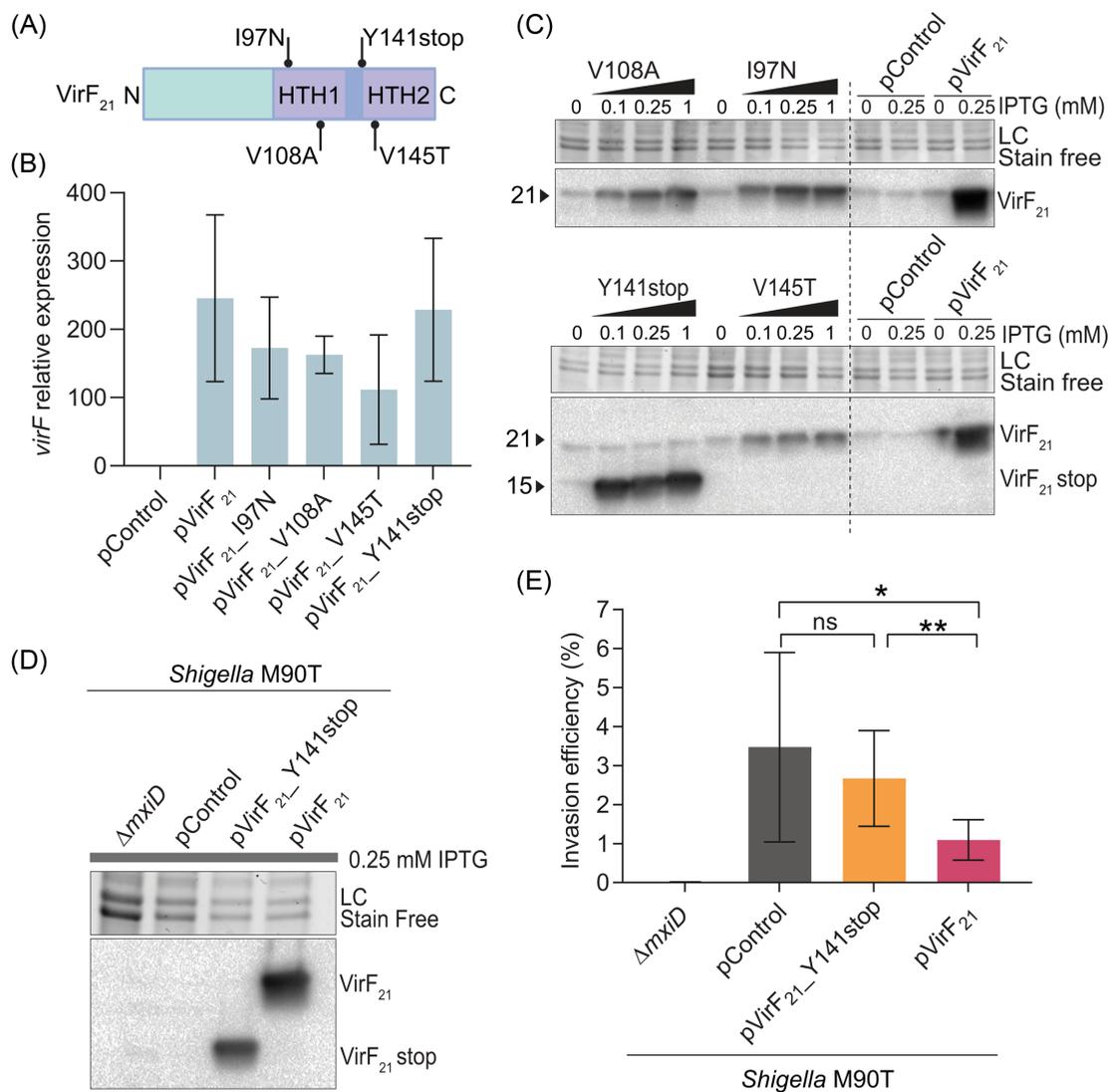


Figure 3. The C-terminal HTH2 motif is required for VirF₂₁ function. **(A)** Schematic representation of the VirF₂₁ protein sequence, with relevant mutagenized amino acid positions indicated. **(B)** *virF* mRNA expression levels ($2^{-\Delta\Delta Ct}$) as a function of protein induction with 0.25 mM IPTG in *Shigella* M90T strains harbouring pControl, pVirF₂₁, pVirF₂₁_I97N, pVirF₂₁_V108A, pVirF₂₁_V145T, or pVirF₂₁_Y141stop. Data come from 6–7 biological replicates from 3 independent experiments and were normalized to the *virF* expression in the M90T pControl strain. **(C)** VirF₂₁ protein levels detected by western blot as a function of IPTG induction (increasing concentration: 0, 0.1, 0.25, 1 mM) in *Shigella* M90T strains harbouring pControl, pVirF₂₁, pVirF₂₁_I97N, pVirF₂₁_V108A, pVirF₂₁_V145T, or pVirF₂₁_Y141stop. VirF₂₁_Y141stop produces a smaller protein (~15kDa), since the last 39 aa in the C-terminal part are deleted. **(D)** VirF₂₁ protein levels detected by western blot in a *Shigella* M90T Δ *mxiD* mutant and in *Shigella* M90T strains harbouring pControl, pVirF₂₁, or pVirF₂₁_Y141stop plasmids in the presence of 0.25 mM IPTG. **(E)** Invasion efficiency of the indicated *Shigella* M90T strains in sub-confluent Caco-2 cell layers. Cells were infected at MOI 100 for 1h, and analysed by selective plating of intracellular bacteria. Shown are CFU data expressed as the percentage of the inoculum retrieved in the intracellular population. Shown are CFU data for 4 (Δ *mxiD*) and 6 (*Shigella* M90T strains harbouring pControl, pVirF₂₁, or pVirF₂₁_Y141stop plasmids) biological replicates from 2 independent experiments. Bars represent mean and standard deviation. Statistical significance was determined by Mann Whitney U test, ns = non significant, *P < 0.05, **P < 0.01.

son Assembly Cloning kit from NEB (#E5510S). Amino acid substitutions were adjusted according to the *Shigella* codon usage. The resulting plasmids were transformed into the wild-type M90T strain. Sequences of the PCR products used are listed in Table S2. Bacteria were routinely grown in LB medium at 37°C, unless otherwise specified. When required, strains were grown in M9 complete medium (M9 minimal medium supplemented with 10 mg/ml thiamine, 0.2% glucose, 0.5% casamino acids and 10 mg/ml nicotinic acid). When necessary, antibiotics were supplemented at the following concentrations: ampicillin, 50 μ g/ml; chloramphenicol, 15 μ g/ml; kanamycin, 50 μ g/ml; streptomycin, 50 μ g/ml. Plasmid DNA extraction, DNA transformation, electrophoresis, purification of DNA fragments and sequencing were performed as de-

scribed previously (Green and Sambrook 2012). PCR reactions were performed using DreamTaq DNA polymerase (Thermo Fisher Scientific, #EP0702) or Phusion DNA polymerase (Thermo Fisher Scientific, #F-530L). All oligonucleotide primers used in this study are listed in Table S2.

β -galactosidase assays

β -galactosidase assays were performed as previously described (Miller 1992) on sodium dodecyl sulphate-chloroform-permeabilized cells grown in LB or M9 medium (to OD₆₀₀ 0.5–0.6). β -galactosidase activity of the pRS-F (+205) and pRS-F (+305) transcriptional fusions was assessed under different conditions. For temperature shift, ON cultures grown at 30°C in LB were

subcultured 1:100 in LB at 30°C or 37°C. To mimic intestine-like conditions, ON cultures grown at 30°C were subcultured 1:100 at 37°C in M9 medium supplemented with the following compounds: Sodium deoxycholate (0, 2.5, 5 mg/ml), Bile salts (0, 6, 9 mg/ml), NaCl (0, 0.1, 0.2M), Hydrogen peroxide (0, 10, 50, 100 mM). To screen different pH conditions, bacteria were subcultured 1:100 in M9 medium at pH 5, 6 and 7.

Immunodetection of VirF proteins

VirF protein levels were detected by western blot through enhanced chemiluminescence. In brief, equal amount of proteins was extracted from strains grown at OD600 ~0.6, separated on Any kD™ Mini-PROTEAN® TGX Stain-Free™ Protein Gels (Biorad, #4568126) and transferred onto Trans-Blot Turbo Mini 0.2 µm PVDF Transfer Packs (Biorad, #1704156). The stain-free method was used to obtain the loading control (Colella et al. 2012). The method is based on the fluorescent detection of tryptophan residues in the protein sequence, as a result of the presence of a trihalo compound in the gel. After protein separation by electrophoresis, each gel was imaged upon exposure to UV-light for 5 min and the same region was selected as loading control for all western blots. Immunodetection was performed as described in Di Martino et al. 2016b using polyclonal halon anti-VirF, anti-FLAG (Sigma, #F1804) and anti-GroEL (Sigma, #A8705) antibodies. Quantification by Western blots were obtained by serial dilution of protein extracts, with the relative amounts calculated from a standard curve. For the protein extracts derived from cultures grown at 30°C, concentrated samples were used to calculate the standard curves.

Congo red binding assay

Congo Red (CR) plates were prepared adding 0.108 mg/ml of Congo Red dye (Sigma, #C6277) to Trypticase Soy Agar (Sigma, #22091) and supplemented with 0, 0.1, 0.25, or 1 mM IPTG (Sigma, #I6758). The indicated *Shigella flexneri* M90T strains were grown ON at 30°C with appropriate antibiotics, diluted 1:40 and subcultured at 37°C for ~2 h (OD600 ~0.7). Subcultures were serially diluted and plated on CR plates containing increasing concentrations of IPTG. The following day Congo Red positive (CR+) and negative (CR-) colonies were enumerated. Subsequently, colonies were scraped from each plate, serially diluted and plated on new CR plates without the IPTG selection.

Epithelial cell culture and infections

Caco-2 cells were grown in DMEM GlutaMAX (Gibco, #31966-021) supplemented with 10% heat-inactivated fetal bovine serum (FBS; Gibco, #10270106) and 0.1 mM Non Essential Amino Acids (Gibco, #11140035) at 37°C with 10% CO₂. Cultures were passaged two to three times/week in the presence of 100IU/ml penicillin and 100 µg/ml streptomycin, but antibiotics were omitted during infection experiments. Caco-2 cells were seeded in 12-well plates 24–48 h before infection. The indicated *S. flexneri* strains were grown ON at 30°C in LB with appropriate antibiotics, diluted 1:50 in the presence of 0.25 mM IPTG to allow the induction of VirF₂₁. The subcultures were further incubated for 2h at 37°C or for 1 h at 30°C before shifting for 1.5 h to 37°C (OD600 ~0.7). Upon infection, bacteria were centrifuged on top of the cultured epithelial cells for 15 min at 700 g, followed by 45min incubation at 37°C and 10% CO₂. The culture medium was replaced with fresh medium containing 200 µg/ml Gentamicin (Sigma, #G1914) and the cells were further incubated for 2 h. At 3h post-infection (p.i.) cells were washed and lysed adding 0.1% Sodium deoxycholate, the lysates

were then diluted and plated on LB agar plates with appropriate antibiotics, followed by enumeration of the number of colony-forming units (CFUs).

qRT-PCR

Total RNA purification and cDNA synthesis were performed as previously described (Di Martino et al. 2016b). qRT-PCR was performed using Maxima SYBR green/ROX qPCR master mix (2X) (Thermo Fisher Scientific, #K0222) on a CFX384 Touch Real-Time PCR Detection System (Biorad). The levels of *virF*, *virF*₃₀, *virB*, *mxiE* and *icsA* transcripts were analysed using the 2^{-ΔΔCT} (cycle threshold [CT]) method and results are reported as the fold increase relative to the reference (Livak et al. 2001). The house-keeping gene *nusA* was used for normalization. The following oligonucleotide primers were used (see Table S2): *nusAQF/nusAQR*, *virFQF/virFQR*, *virF30QF/virF30QR*, *virBQF/virBQR*, *mxiEQF/mxiEQR* and *icsAQL/icsAQR*.

Results and discussion

Expression levels of the *virF*₂₁ l1mRNA and the *VirF*₂₁:*VirF*₃₀ protein ratio are both elevated at non-permissive temperature (30°C)

Previous work identified a *virF*₂₁ translationally capable l1mRNA, whose transcription is dependent on the presence of an internal promoter located two nucleotides upstream the *VirF*₂₁ translational start site (Di Martino et al. 2016b). To determine how and when expression of the *virF*₂₁ l1mRNA occurs, *E. coli* strains harbouring a *virF*₂₁(l1mRNA)-*lacZ* transcriptional fusion construct (pRS-F (+205)), or a control fusion (pRS-F (+305)) (Di Martino et al. 2016b) were grown under different conditions, mimicking either the environment *Shigella* encounters during host cell invasion in the gut (i.e. exposure to sodium deoxycholate, bile salts, high osmolarity, oxidative stress, low pH) or the environment outside the host (non-permissive temperature: 30°C) (Marteyn et al. 2012). No difference in the pRS-F (+205) β-galactosidase activity was observed during osmotic, oxidative and pH-stress, as compared to the untreated control (Fig S1A-B-C). Increasing concentrations of either sodium deoxycholate alone or a bile salt mixture, hence mimicking the biliary secretions encountered in the human intestinal tract (Faherty et al. 2012, de Buy Wenniger et al. 2013, Di Ciaula et al. 2017, Nickerson et al. 2017, Chanin et al. 2019), resulted in ~6 and ~3-fold increase in β-galactosidase activity, respectively (Fig S1D-E). This suggests an increase in *virF*₂₁ l1mRNA transcription under these two conditions. However, the relative abundance of *VirF*₂₁ protein, measured in a *Shigella flexneri* M90T strain harbouring a 3xFlag-tagged version of the *VirF* proteins, did not increase accordingly in response to these stimuli (Fig S1F-G-H). This may imply that under the conditions tested here, the majority of *VirF*₂₁ protein originates from alternative translation of the *virF* full length mRNA. The existence of some other unknown post-transcriptional regulatory mechanisms, hampering *VirF*₂₁ translation, can also not be ruled out. In either case, typical conditions encountered by *Shigella* within the host may influence *virF*₂₁ l1mRNA transcription, but do not seem to significantly alter *VirF*₂₁ protein levels.

Interestingly, we found that the β-galactosidase activity of the pRS-F (+205) fusion was significantly higher at 30°C than at the permissive host temperature of 37°C (Fig. 1A). Translation of both *VirF* forms in *Shigella flexneri* M90T was observed at both temperatures (Fig. 1B-E). In agreement with the positive regulation of *virF* expression at 37°C, *VirF*₃₀ protein abundance was ~8–10-

fold higher at 37°C than at 30°C both in LB (Fig. 1B-C-F) and in M9 medium (Fig. 1D-E-G). Notably, the VirF₂₁:VirF₃₀ protein ratio differed dramatically between the two temperatures. While VirF₂₁ represented ~5–10% of the VirF₃₀ protein content at 37°C (~5% in LB; ~10% in M9), it reached peaks of ~20–50% at 30°C (~20% in LB; ~50% in M9) (Fig. 1C-E-H-I).

Taken together these results suggest that while *virF*₂₁ lmrRNA transcription may be affected by several different stimuli, a high VirF₂₁:VirF₃₀ protein ratio is favoured at environmental temperature, rather than under host-like conditions. In this context, VirF₂₁ might function as a molecular brake to minimize fitness costs when the *Shigella* virulence program is not required or undesired.

The *virF* genetic arrangement leading to the transcription and translation of two proteins from a single gene is not an isolated example. The *E. coli* copper chaperone CopA and the *Salmonella* LysR-type regulator LtrR also display transcription of two mRNA molecules and the translation of two protein forms under specific environmental conditions (Drees et al. 2017, Rebollar-Flores et al. 2020). Furthermore, computational analysis aimed at discovering overlooked regulatory elements showed that gene internal promoters are often associated with horizontally transferred genes, both in *E. coli* and in some archaeal species (Ten-Caten, Vêncio et al. 2018). This is particularly relevant here, since *virF* was horizontally acquired on the pINV during *Shigella*'s evolution towards pathogenicity (Yang et al. 2005). Altogether, this suggests the existence of a widespread adaptation strategy in bacteria to expand and diversify the protein repertoire and thereby optimize the response to changing external conditions.

Ectopic expression of VirF₂₁ at the permissive temperature reversibly suppresses the *Shigella* host cell invasive program

VirF₃₀ activity governs the transition of *Shigella* between non-invasive and invasive states. When switching from 30°C to 37°C, already a modest increase in *virF* transcription is sufficient for full activation of the downstream virulence cascade and a host cell invasive phenotype (Le Gall et al. 2005). To test the hypothesis that VirF₂₁ expression can prevent switching to the invasive phenotype, we explored the consequences of elevating VirF₂₁ protein levels under the permissive temperature (37°C).

First, we investigated the effect of ectopic VirF₂₁ expression on the ability of *Shigella* to bind Congo Red (CR), a phenotype linked to virulence and the presence of *virF*, resulting in red colonies (CR+) on solid medium (Sakai et al. 1986a). *Shigella flexneri* M90T was transformed with either a plasmid that allowed IPTG-inducible expression of VirF₂₁ (pVirF₂₁; carries the Ptac promoter; reported as pAC-21 in Di Martino et al. 2016b), or the corresponding empty vector (pControl; previously named pGIP7 in Falconi et al. 2001). Ectopic expression of VirF₂₁ was detected in the presence of graded concentrations of IPTG (0.1–0.25–1 mM), but not in the absence of IPTG (Fig. 2A). We plated dilutions of exponential cultures of M90T pControl, M90T pVirF₂₁ and M90T Δ*virF* on CR plates containing increasing concentrations of IPTG and incubated at 37°C. Fig. 2B and figure S2 show that IPTG-induced ectopic expression of VirF₂₁ led to the appearance of a high percentage (~50–90%) of white (CR-) colonies, reaching comparable levels as the non-virulent M90T Δ*virF* mutant, a strain known to exhibit a completely CR- phenotype (Sakai et al. 1986b). In particular, a robust CR- phenotype (>80% white colonies) was observed on CR plates containing 0.25 and 1 mM IPTG, while the percentage white colonies was somewhat variable in the presence of 0.1 mM IPTG. This observation suggests a borderline VirF₂₁ expression at

IPTG concentrations lower than 0.25 mM. Importantly, when the CR- M90T pVirF₂₁ colonies were collected from the plates supplemented with IPTG and re-plated on new CR plates devoid of the IPTG inducer, the bacteria reverted back to virtually exclusively CR+ colonies (Fig. 2B; <3% CR-). These results suggest that CR binding is subjected to a VirF₂₁-driven reversible switch, linked also to a decrease in virulence gene expression (Fig. S3A-B; and Di Martino et al. 2016b).

Next, we infected human colonic epithelial Caco-2 cells with the M90T pControl and M90T pVirF₂₁ strains, to test how ectopic VirF₂₁ expression impacts *Shigella* host cell invasion. M90T Δ*mxid* (lacking the outer membrane ring MxiD protein, resulting in a nonfunctional T3SS) was used as a non-invasive control strain. To ensure robust VirF₂₁ expression with minimal side effects, we induced VirF₂₁ expression with the intermediate concentration of IPTG (0.25 mM, as informed by the CR binding assay; Fig. 2A and B). As expected, the M90T Δ*mxid* strain failed at infecting Caco-2 cells (Fig. 2C). Notably, ectopic expression of VirF₂₁ (M90T pVirF₂₁) led to a ~4–5-fold decrease in *Shigella*'s ability to infect Caco-2 cells, as compared to the M90T pControl strain (Fig. 2C).

Taken together, these results show that elevated VirF₂₁ expression at 37°C can suppress the *Shigella* invasive program, signified by a CR-phenotype on plates, reduced virulence gene expression, and hampered capacity to infect epithelial cells. Considering that VirF₂₁ makes up a larger fraction of the total VirF protein pool at 30°C than at 37°C (Fig. 1), the above results reinforce the hypothesis that VirF₂₁ may negatively tune *Shigella* virulence gene expression, when this is undesirable. It is known that the pINV virulence plasmid is subjected to high counter-selective pressure at 37°C. With increasing number of generations, mutations, insertions of IS sequences and/or complete loss of the virulence cascade top regulators *virF* or *virB* occur at 37°C (Sasakawa et al. 1986, Schuch and Maurelli 1997, Pilla et al. 2017). Occasionally, the selective pressure can escalate leading even to the loss, or integrational silencing, of the entire pINV (Zagaglia et al. 1991, Pilla et al. 2017). At 30°C, virulence gene expression is silenced and therefore the selective pressure on the pINV is relieved, resulting in minimal loss or mutations (Schuch and Maurelli 1997). In this context, it is tempting to speculate that VirF₂₁ expression represents an additional regulatory layer to minimize virulence gene expression leakage and therefore promote overall pINV stability under certain environmental conditions.

The C-terminal region of VirF₂₁ is required for negative regulation of *Shigella* host cell invasion

The experimental setting based on ectopic VirF₂₁ expression precludes assessment of the impact of endogenous VirF₃₀ and VirF₂₁ levels expressed from their native genetic context. Despite significant efforts, we have however been unsuccessful at generating a *Shigella flexneri* scarless mutant expressing VirF₃₀ protein only from the endogenous locus. This may suggest that the native *virF* locus sequence is unusually intolerant to perturbation, although we cannot formally rule out other technical explanations.

To better understand the relationship between VirF₂₁ sequence and function, and to verify the specificity of the above results, we therefore opted for a site directed mutagenesis approach, targeting the untagged *virF*₂₁ gene cloned into the pVirF₂₁ plasmid. VirF₃₀ and VirF₂₁ belong to the family of AraC-like transcriptional regulators (Cortés-Avalos et al. 2021). This group comprises both positive and negative transcriptional regulators, which often control virulence systems across different gram-negative bacterial species (i.e. *Salmonella*, *Yersinia*, *Vibrio cholera*, (Gallegos et al. 1997, Cortés-

Avalos et al. 2021)). The mechanisms governing AraC-like protein expression and regulation have been successfully studied in many cases. However, their biochemical and structural properties have been less well characterized, since AraC-like proteins are often difficult to purify (Cortés-Avalos et al. 2021). The DNA sequences targeted by VirF₃₀ have nevertheless been identified in some cases (i.e. within *icsA*, *RNAG*, and *virB* promoters) (Tobe et al. 1993, Giangrossi et al. 2010, Tran et al. 2011), and the VirF₂₁ binding site within the *virF* promoter was also previously mapped (Di Martino et al. 2016b). VirF₂₁ and VirF₃₀ share the C-terminal portion, which contains the two typical AraC-like Helix-Turn-Helix (HTH) DNA binding motifs, separated by an alpha helix linker. VirF₂₁ lacks the N-terminal domain of VirF₃₀, which is believed to have oligomerization properties. Some of the amino acids likely involved in the interaction between VirF₃₀ and its DNA targets have been identified by a combined random and site directed mutagenesis approach (Porter and Dorman 2002).

In an attempt to obtain a non-functional VirF₂₁, we transplanted an assortment of mutations shown to affect VirF₃₀ function in the prior study (Porter and Dorman 2002). The following mutations were introduced onto the pVirF₂₁ plasmid: I97N, V108A, V145T and Y141stop, here reported considering VirF₃₀-Met84 (Porter and Dorman 2002) = VirF₂₁-Met1 (Fig. 3A; Di Martino et al. 2016b; previously reported in Porter and Dorman 2002 as I180N, V191A, V228T, and Y224Och respectively). The substitutions I97N and V108A target the HTH1 DNA binding motif, while the V145T substitution is located within the HTH2 DNA binding motif. Finally, the deletion of the HTH2 motif was achieved by introducing a stop codon at position 141 (Y141stop; deletion of 39 aa in the C-terminus). Upon induction with IPTG in *Shigella flexneri* M90T, the wild type (wt) and the mutated versions of virF₂₁ showed broadly similar transcriptional levels (Fig. 3B). However, VirF₂₁ protein levels (monitored by a halon anti-VirF antibody) were markedly lower for the VirF₂₁_I97N, VirF₂₁_V108A and VirF₂₁_V145T mutant constructs than in the VirF₂₁_wt carrying strain (Fig. 3C), suggesting a possible impact of these amino acid substitutions on protein stability. Only the truncated VirF₂₁_Y141stop construct generated protein levels comparable to the strain harbouring the pVirF₂₁_wt plasmid (Fig. 3C-D). Next, we infected Caco-2 cells with the strains ectopically expressing either VirF₂₁_wt (pVirF₂₁) or VirF₂₁_stop (pVirF₂₁_Y141stop). As evident from Fig. 3E, the *Shigella* pVirF₂₁_Y141stop strain retained the ability to infect Caco-2 cells at similar levels as the control strain (*Shigella* pControl), while the *Shigella* pVirF₂₁_wt strain again showed a ~3 fold lower invasion capacity (Fig. 3E; compare also with Fig. 2C). These results show that the VirF₂₁_stop protein, lacking the predicted DNA binding HTH2 motif, can be expressed to similar levels as full-length VirF₂₁, but is non-functional. This validates the specificity of the VirF₂₁ suppressive effects observed in the above experiments (Fig. 2), and reveals a key role of the C-terminal portion for VirF₂₁ functionality.

Conclusions

Pathogenic bacteria are masters at adapting to fast-changing environmental cues. *Shigella* encounters many different environmental conditions and switches flexibly between extracellular and intracellular lifestyles. While the activation of the *virF* regulatory cascade is a crucial event for *Shigella* expression of the invasive program (Schroeder and Hilbi 2008), it also constitutes a significant fitness cost for the bacterial population (Schuch and Maurelli 1997). Thus, it is not surprising that *Shigella* employs a multi-layered regulatory arsenal to ensure expression of the viru-

lence genes only when these are needed. VirF₂₁ has been identified as a possible negative autoregulator of VirF₃₀, but the impact on the invasive *Shigella* phenotype had not been addressed (Di Martino et al. 2016b). Our results expand on these previous findings, by illustrating that virF₂₁ mRNA expression and the VirF₂₁:VirF₃₀ protein ratio is enhanced at 30°C, a common condition *Shigella* encounters outside of the host. In this context, it seems plausible that VirF₂₁ serves to suppress virulence gene expression when not desired. Indeed, when ectopically expressed, VirF₂₁ is capable of suppressing the *Shigella* virulence program at the permissive temperature 37°C, resulting in a CR- phenotype on plates, lowered levels of virulence gene transcripts, and an impaired ability to infect host cells. This suppressive activity requires the HTH2-motif-containing C-terminus of the VirF₂₁ protein.

While the physiological impact of VirF₂₁ remains to be completely explored under the multitude of possible environmental conditions, the findings presented here highlight the interconnected mechanisms that ensure fine-tuned regulation of virulence properties across the *Shigella* life cycle.

Supplementary data

Supplementary data are available at [FEMSLE](https://femsle.onlinelibrary.wiley.com/doi/10.1111/femsle.13691) online.

Author contribution

Conceptualization: M.E.S., M.L.D.M. Methodology: E.S., M.E.S., M.L.D.M. Investigation: E.S., M.L.D.M. Formal analysis: E.S., M.L.D.M. Resources: B.C., G.P., M.E.S., M.L.D.M. Supervision: M.E.S., M.L.D.M. Project administration: M.E.S., M.L.D.M. Funding acquisition: M.E.S., M.L.D.M. Visualization: E.S., M.L.D.M. Writing—Original Draft: M.L.D.M. Writing—Reviewing & Editing: E.S., M.E.S., M.L.D.M. All authors read, commented on, and approved the final manuscript.

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