Neurospora Importin α Is Required for Normal Heterochromatic Formation and DNA Methylation

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Abstract

Heterochromatin and associated gene silencing processes play roles in development, genome defense, and chromosome function. In many species, constitutive heterochromatin is decorated with histone H3 tri-methylated at lysine 9 (H3K9me3) and cytosine methylation. In Neurospora crassa, a five-protein complex, DCDC, catalyzes H3K9 methylation, which then directs DNA methylation. Here, we identify and characterize a gene important for DCDC function, dim-3 (defective in methylation-3), which encodes the nuclear import chaperone NUP-6 (Importin α). The critical mutation in dim-3 results in a substitution in an ARM repeat of NUP-6 and causes a substantial loss of H3K9me3 and DNA methylation. Surprisingly, nuclear transport of all known proteins involved in histone and DNA methylation, as well as a canonical transport substrate, appear normal in dim-3 strains. Interactions between DCDC members also appear normal, but the nup-6dim-3 allele causes the DCDC members DIM-5 and DIM-7 to mislocalize from heterochromatin and NUP-6 dim-3 itself is mislocalized from the nuclear envelope, at least in conidia. GCN-5, a member of the SAGA histone acetyltransferase complex, also shows altered localization in dim-3, raising the possibility that NUP-6 is necessary to localize multiple chromatin complexes following nucleocytoplasmic transport.

Author Summary

The epigenetic information contained in chromatin is essential for development of higher organisms, and if misregulated, can lead to the unregulated growth associated with human cancers. Chromatin is typically classified into two basic types: gene-rich ‘euchromatin’, and gene-poor heterochromatin, which is also rich in repeated DNA and ‘repressive chromatin marks’. As in humans and eukaryotes generally, heterochromatin in Neurospora crassa is decorated with DNA methylation and histone H3 lysine 9 (H3K9) methylation,
Introduction

The densely staining regions of eukaryotic chromosomes, referred to as heterochromatin, typically contain repetitive, A:T rich DNA, and are characterized by low gene density, reduced genetic recombination, di- or tri-methylation of lysine 9 on histone H3 (H3K9me2 or H3K9me3), and DNA methylation [1–4]. Heterochromatin is critical for centromere and telomere function, and is largely responsible for the silencing of transposable elements [1,4]. Unlike the situation in animals and plants, which require DNA methylation for normal development [5,6], the fungus *Neurospora crassa* does not require DNA methylation for viability [4,7]. Neurospora has characteristics of heterochromatin found in higher eukaryotes and is convenient for genetic and biochemical studies. These traits have led to the identification of genes involved in establishing, maintaining, and regulating DNA methylation and other features of heterochromatin [8].

A single DNA methyltransferase (DNMTase), DIM-2, is responsible for all DNA methylation in vegetative tissue of Neurospora [9]. DIM-2 directly interacts with heterochromatin protein-1 (HP1) [10], which binds to H3K9me3 [11,12]. The histone methyltransferase (HMTase) DIM-5 [13,14] is responsible for trimethylation of H3K9. *In vivo*, DIM-5 activity depends on all members of the five protein complex, DCDC (DIM-5/-7/-9, CUL4, DDB1 Complex) but DIM-7 alone appears sufficient to target DIM-5 to incipient heterochromatin regions [8,15]. DCDC resembles Cullin-4 E3 ubiquitin ligase complexes, with the WD-40 protein DIM-9 being the putative DCAF (DDB1/CUL4 associated factor), which is normally expected to recognize substrates destined for ubiquitination. However, results of recent studies indicate that DCDC does not function as a canonical ubiquitin ligase [16]. Thus, important questions regarding how DCDC and other members of the heterochromatin/DNA methylation machinery function and are controlled remain unanswered.

To improve our understanding of the control of DNA methylation and heterochromatin formation, we characterized the *Neurospora dim-3* strain, which shows a substantial loss of DNA methylation [7]. Genetic mapping, whole genome sequencing, and complementation tests identified the causative mutation in the *nup-6* gene, resulting in a critical change in the eighth ARM repeat of NUP-6 (Importin α). This protein (also known as Srp1p in yeast and karyopherin α in humans) is the canonical nucleocytoplasmic transport adaptor. Importin α binds “cargo” proteins to be transported into the nucleus, complexes with Importin β, and then is shuttled through the nuclear pore complex to the nucleoplasm [17–19]. We found that *dim-3* strains have a drastic reduction in global H3K9me3, indicating that NUP-6 is required for...
proper DCDC function. Nuclear transport and interactions of critical DCDC components appear normal in dim-3 strains but at least two DCDC components (DIM-5 and DIM-7) are mis-localized from heterochromatin. Curiously, the SAGA histone acetyltransferase, but not DCDC components, showed increased localization in euchromatin of a dim-3 strain. Altogether, our results reveal a nuclear transport-independent role of NUP-6 in localizing chromatin complexes to sub-nuclear targets.

Results
Loss of DNA methylation in dim-3 strains is caused by a mutation in the nup-6 gene

The dim-3 gene was genetically identified in a brute-force screen for methylation defects following N-methyl-N’-nitro-N-nitrosoguanidine mutagenesis [7]. Southern blots probed for the representative interspersed heterochromatic regions 8G3, 8A6, and 2B3 [20] in a histidine auxotrophic dim-3 strain illustrate the substantial DNA methylation reduction caused by the allele (Fig. 1A). Genome-wide bisulfite sequencing (BS-Seq; Rountree and Selker, in preparation) demonstrated that the residual DNA methylation in a dim-3 mutant is distributed normally, or largely normally, to heterochromatic regions (Figs. 1B and S1).

Dim-3 was mapped to the right arm of Linkage Group V, and high-throughput sequencing identified two point mutations in the open reading frame of gene NCU01249, E396K and R469H; both were confirmed by Sanger sequencing. NCU01249 encodes NUP-6 (NIH GenBank accession EAA31416.1), which is predicted to form a structure similar to yeast Importin α [21], a protein that is highly conserved in eukaryotes [19]. Neurospora NUP-6 includes an N-terminal ~80 amino acid Importin β-binding domain (IBB), followed by ten ~40 amino acid ARM repeats (Fig. 1C), each of which should fold into a triple-alpha helical bundle to form a binding pocket for the nuclear localization signals (NLS) of cargo proteins (S2A Fig.; [22]). ARM repeats 1–9 are thought to recognize the NLS [22] while cargo release factors interact with the less-conserved 10th ARM repeat [23]. The E396K and R469H changes in nup-6\textsuperscript{dim-3} are in the putative 8th and 10th ARM repeats, respectively (Figs. 1C and S2A), and should not destabilize the NUP-6\textsuperscript{dim-3} protein nor impact its nuclear shuttling (S2B-S2C Fig.). Interestingly, dim-3 strains have a minor growth defect (S2D Fig.) and are homozygous sterile, indicating that one or both of the amino acid substitutions compromise an important cellular process.

To confirm that nup-6\textsuperscript{dim-3} causes the observed DNA methylation loss in dim-3 strains, we introduced a wild type (WT) nup-6 gene (nup-6\textsuperscript{+}) at an ectopic locus (his-3) of a dim-3 strain. Ectopic nup-6\textsuperscript{+} restored global DNA methylation to WT levels at all the representative heterochromatic loci tested (Fig. 1A), suggesting that nup-6\textsuperscript{dim-3} was indeed responsible for the Dim phenotype. To determine if one or both of the mutations cause the phenotype, we replaced the endogenous nup-6\textsuperscript{+} allele with engineered mutant alleles. As expected, the strain with the reintroduced nup-6\textsuperscript{dim-3} allele showed reduced DNA methylation, although somewhat less so than the original dim-3 isolate (Figs. 1D and S3). Reintroduction of the R469H mutation did not affect DNA methylation, whereas the reintroduced E396K mutation caused a loss equivalent to that observed with the reintroduced nup-6\textsuperscript{dim-3} allele, implicating this change as the causative mutation (Figs. 1D and S3). As expected, the nup-6 gene appears essential, because strains with the gene deleted are only viable as heterokaryons, containing both nup-6\textsuperscript{+} and Δnup-6::hph nuclei [24]. Thus, there is no reason to expect that the E396K creates a null allele.
Histidine exacerbates the dim-3-mediated DNA methylation loss

While comparing DNA methylation levels in various dim-3 strains, we noticed that the histidine-requiring dim-3 strains had an exacerbated reduction in DNA methylation, leading us to
systematically test the possible effect of histidine on DNA methylation in \textit{dim-3} and \textit{dim-3}\textsuperscript{+} strains. We found that histidine supplementation decreased DNA methylation at all heterochromatic regions tested in a \textit{dim-3} strain, as demonstrated by Southern blotting (Fig. 2A) and genome-wide bisulfite sequencing (BS-Seq; Figs. 1B and S1), but has no marked effect on DNA methylation levels in a \textit{dim-3}\textsuperscript{+} strain.

We tested several possibilities to determine the source of this histidine effect. The “cross pathway control” system causes de-repression of several amino acid biosynthetic pathways concomitant with changes in the level of an individual amino acid [25]. We tested its involvement in the histidine effect by checking for possible consequences of added arginine, lysine, and tryptophan, but found they neither caused loss of DNA methylation nor restored it when added with histidine (S4 Fig.). RNAi components are not required for DNA methylation [11], but histidine supplementation and exposure to other DNA damaging agents, induces expression of the Neurospora Argonaute QDE-2 (EAA31129.2) and increases the levels of qiRNAs (QDE-2-interacting small RNAs; [26]). We tested the possible involvement of DNA damage and RNAi in the histidine effect by growing a \textit{dim-3} strain in medium containing histidine (his), hydroxyurea (HU), or ethyl methane sulfonate (EMS). Only histidine addition reduced DNA methylation in \textit{dim-3} strains (S5A Fig.), and this effect also occurred in a \textit{Δqde-2} strain (S5B Fig.), indicating the histidine effect in \textit{dim-3} strains involves neither DNA damage nor qiRNAs.

Considering that NUP-6 is critical for nuclear transport and that histidine does not further deplete H3K9me3 levels (below), we investigated whether the histidine effect results from altered nuclear transport and/or sub-nuclear localization of DIM-2 (AF348971.1) or HP1 (AY363166.1), which operate downstream of H3K9me3. We found no significant difference in levels of DIM-2-3xFLAG in nuclei from a \textit{dim-3} strain grown in the presence or absence of histidine relative to WT (Fig. 2B), suggesting that neither histidine nor the \textit{nup-6}\textsuperscript{-}\textit{dim-3} allele impacts DIM-2 nuclear shuttling. In addition, cytological analyses of HP1-GFP (S6 Fig.; [11]) and DamID analyses of HP1-DAM (below) revealed no evidence of a transport defect in \textit{dim-3} strains.

We next tested if histidine perturbs sub-nuclear targeting of DIM-2 or HP1 in a \textit{dim-3} strain. The localization of many heterochromatin-specific proteins, including DIM-2, are not readily detected by standard chromatin immunoprecipitation (ChIP), perhaps due to transient chromatin interactions, but can be detected by DamID [8]. Therefore, we fused the DNA adenine methyltransferase (\textit{dam}) gene to the downstream ends of the \textit{dim-2} [27] and \textit{hpo} (encoding HP1) genes, expressed these constructs in \textit{dim-3} and \textit{dim-3}\textsuperscript{+} strains grown in minimum medium (Figs. 2C and S7A). DIM-2-DAM was found to localize to heterochromatin in both \textit{dim-3}\textsuperscript{+} and \textit{dim-3} strains grown in minimum medium (Figs. 2C and S7A). In contrast, in a \textit{dim-3} strain grown with histidine, DIM-2-DAM showed reduced localization to heterochromatin, while histidine did not alter DIM-2-DAM localization in a \textit{dim-3}\textsuperscript{+} strain (Figs. 2C and S7A). Unlike the situation with DIM-2, histidine did not reduce, and in fact slightly increased, the heterochromatic localization of HP1-DAM (Figs. 2D and S7B). These findings suggest that in \textit{dim-3} strains, histidine may compromise the direct interaction between DIM-2 and HP1 that is necessary for DNA methylation in Neurospora [10].

The \textit{dim-3} strain has a global reduction of H3K9me3

To determine whether the reduced DNA methylation observed in \textit{dim-3} strains reflects a loss of H3K9me3, we assessed global H3K9me3 levels by western blotting. H3K9me3 was greatly
Fig 2. Histidine supplementation exacerbates loss of DNA methylation in dim-3 strains by displacing DIM-2 from heterochromatin. [A] Southern blot assay (as in Fig. 1A) of DNA methylation in dim-3+ (WT) or dim-3 prototrophic strains grown in minimal or histidine supplemented medium. [B] (left) α-FLAG western blots showing DIM-2-3xFLAG levels in WT and dim-3 nuclei grown in medium with or without histidine compared with a loading control (histone H3, [hH3]); (right) Quantification of DIM-2-3xFLAG levels from three independent experiments, normalized to hH3 levels. [C-D] Southern blot assay of DamID experiments with WT and dim-3 strains containing [C] DIM-2-DAM or [D] HP1-DAM, grown with or without histidine, and probed for the heterochromatic region 2:B3. Genomic DNA was digested with DpnI (DI), specifically cutting GA^mTC sequences or with the A^m-insensitive isoschizomer DpnII (DII) to monitor complete digestion.

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reduced in dim-3 strains compared to that in a WT strain, and was reestablished after introduction of an ectopic nup-6+ gene (Fig. 3A), indicating that NUP-6 is required for normal levels of H3K9me3. Considering that added histidine reduced DNA methylation levels in dim-3 strains (Fig. 2), we checked if histidine exacerbates the H3K9me3 loss. Western blots showed no additional loss of H3K9me3 (Fig. 3B), supporting the notion that histidine reduces DNA methylation by compromising DIM-2 localization in a dim-3 background.

To determine if the residual H3K9me3, like DNA methylation, is found in normal heterochromatic regions in dim-3 strains, we carried out H3K9me3-specific ChIP with high throughput sequencing of associated DNA (ChIP-seq), and despite low signal, found an apparently equivalent distribution of this chromatin mark as in a dim-3+ strain, indicating that the remaining H3K9me3 is correctly localized to heterochromatin in a dim-3 strain (Figs. 3C and S8). Presumably, decreased DNA methylation in dim-3 strains is due to the dramatic reduction of H3K9me3 in heterochromatin.

The dim-3 mutation causes loss of telomeric silencing

To determine if the H3K9me3 loss in dim-3 strains might compromise heterochromatin-associated silencing, we tested the expression of drug resistance markers integrated at telomeric (telVR::hph) and centromeric (cenVIR::bar) sites, which are silent even in the absence of DNA methylation, i.e. in a dim-2 mutant [27,28]. The dim-3 allele de-repressed the telVR::hph
marker as evidenced by growth in the presence of hygromycin (Fig. 3D), indicating that heterochromatin in dim-3 is compromised, although this de-repression is not as striking as in an hpo mutant, which fully de-represses the marker [28]. The dim-3 allele did not de-repress the cen-VIR:bar marker (Fig. 3D), consistent with the existence of a DNA methylation-independent mechanism for silencing centromeric heterochromatin [27].

**Nuclear transport of the H3K9me3 machinery is not deficient in dim-3 strains**

Considering that the canonical function of NUP-6 is to transport proteins into the nucleus, we tested if a dim-3 strain has diminished nuclear transport of proteins required for heterochromatin formation and of a protein known to depend on the Importin α/β import system. Our observations that DIM-2 and HP1 localize normally in the nucleus (Figs. 2B and S6) and that dim-3 strains have only a minor growth defect (S2C Fig.) did not support the idea that dim-3 strains are defective in nuclear shuttling. Nevertheless, because the dim-3 mutant shows a striking loss of H3K9me3, we tested whether nuclear transport of DCDC components (DIM-5 [AF419248.1], DIM-7 [AL513463.1], DDB1 [EAA33111.1], DIM-9 [XP_956278.2], and CUL4 [XP_957743.2]) is impaired. To monitor nuclear transport, nuclei were isolated from dim-3+ and dim-3 strains bearing DCDC members individually 3xFLAG-tagged at their endogenous loci [8,15] and protein levels were assessed by western blotting (Figs. 4A and S9). Nuclei preparations were shown to be clean of cytoplasmic contamination by probing western blots for phosphoglycerate kinase (α-PGK, EAA33194.1; Fig. 4B).

We found that the nuclear level of every component of DCDC was undiminished by the dim-3 mutation (Figs. 4A and S9). Indeed, some of the components actually showed increased nuclear abundance. DIM-5 and DIM-7 showed equivalent levels in dim-3+ and dim-3 strains while the nuclear levels of DDB1, DIM-9 and CUL4 were higher in the dim-3 strain (Figs. 4A and S9). Moreover, an examination of the distribution of DCDC components between the nuclear and cytoplasmic fractions revealed no increase in the relative amount of the proteins in the cytoplasmic fraction (Fig. 4B). DIM-7, which is an exclusively nuclear protein, showed no change in abundance or localization. Similarly, while the nuclear level of DIM-5 increased slightly in the dim-3 strain (Fig. 4A), the relative nuclear/cytoplasmic distribution of the protein was equivalent in dim-3+ and dim-3 strains. DIM-9, which is predominantly nuclear in dim-3+ strains, showed an increase in both the cytoplasmic and nuclear fractions of the dim-3 strain (Fig. 4B). It is interesting that the nuclear levels of DDB1, DIM-9, and CUL4 were elevated in dim-3 strains. The increase in CUL4 only occurred when this strain is supplemented with histidine (compare Figs. 4B and S9), which is consistent with the induction of a DNA damage response, as previously documented [26]. The basis for increased DDB1 and DIM-9 levels in dim-3 strains is not clear but may reflect a feedback mechanism to regulate the amount of functional DCDC. Results of qRT–PCR analyses of dim-7, ddb1, and dim-9 transcripts in dim-3+ and dim-3 strains did not reveal RNA differences that could account for the differences in protein levels (S10 Fig.), suggesting the effect is at the translational or posttranslational level.

To test if the dim-3 mutation impacted nuclear transport of a protein with a nuclear localization signal (NLS) known to be bound and transported by Importin α, we overexpressed a GFP reporter construct with an N-terminal SV40 monopartite NLS [22,29–32] in dim-3+ and dim-3 strains and examined nuclear GFP signal by fluorescence microscopy. We found that 91.6% of dim-3+ cells had strong GFP signal inside their nuclei, and this result was mirrored in a dim-3 strain, where 92.5% of cells had nuclear GFP (Fig. 4C). Thus, NUP-6dim-3 effectively transports canonical nuclear cargo.
Interactions within the DCDC are not decreased in \textit{dim-3} strains

Given that all DCDC members, as well as DIM-2 and HP1, are transported into the nucleus in \textit{dim-3} strains, we next considered the possibility that NUP-6\textsubscript{\textit{dim-3}} somehow interferes with DCDC assembly. To test this hypothesis, we monitored DCDC component interactions in

\begin{align*}
\text{Fig 4. Nuclear transport of DCDC components is unaffected by the \textit{dim-3} mutation.} & \\
\text{[A] \(\alpha\)-FLAG and \(\alpha\)-histone H3 (hH3; loading control) western blots of \textit{dim-3}+ (WT) or \textit{dim-3} nuclei expressing individually FLAG-tagged DCDC components. All strains were grown in the presence of histidine. Average and standard deviation of nuclear FLAG-tagged protein levels of three experiments are indicated below the representative images shown.} & \\
\text{[B] \(\alpha\)-FLAG western blots of the total (T) proteins, cytoplasmic (C) fraction, and nuclear (N) fraction of nuclei preparations from \textit{dim-3}+ (WT) and \textit{dim-3} strains expressing \textit{DIM-5-3xFLAG}, \textit{DIM-7-3xFLAG}, or \textit{DIM-9-3xFLAG}. Representative \(\alpha\)-histone H3 (hH3) and \(\alpha\)-PGK blots from the \textit{DIM-5-3xFLAG} nuclei are shown as nuclear and cytoplasmic fraction controls, respectively; equivalent results were obtained for all nuclei preparations.} & \\
\text{[C] Representative differential interference contrast (DIC) and GFP fluorescent images of (left) \textit{dim-3}+ or (right) \textit{dim-3} strains expressing an overexpressed NLS\textsuperscript{\textit{SIV40}}-GFP reporter construct (pCCG::NLS\textsuperscript{\textit{SIV40}}::LexADBD::GFP). While just one representative, multinucleate cell is displayed in the figures here and elsewhere in the paper, we visualized numerous vegetative cells, including both conidia and hyphal cells [62]. Percentages of conidia exhibiting each pattern are listed on the right (P value = 0.43, \(X^2\)-test). Scale bar indicates 5\(\mu\)m.} & \\
\end{align*}

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dim-3 and dim-3+ nuclei by co-immunoprecipitation (co-IP) assays with 3xFLAG-tagged DCDC members. We began by analyzing the interaction between DIM-7 and DIM-5. Equivalent levels of DIM-7-3X-FLAG were recovered from dim-3 and dim-3+ nuclei (Fig. 5A), consistent with our finding that DIM-7 levels in the nucleus are not affected by the dim-3 mutation (Figs. 4A and S9). Equivalent levels of DIM-5 were found in co-IPs of both backgrounds, implying that DIM-7 binding to DIM-5 is not compromised in dim-3 cells (Fig. 5A). Since the DIM-5 interaction with the DCDC members appears mediated through DIM-7 [15], we assessed DIM-5 binding to DIM-9-3XFLAG in dim-3+ and dim-3 nuclei. Because more DIM-9-3XFLAG was recovered from dim-3 nuclei than from dim-3+ nuclei (Fig. 5A), we normalized

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the amount of DIM-5 purified to the DIM-9-3xFLAG bait levels. The DIM-5 level directly correlated with the DIM-9-3xFLAG level (Fig. 5A), indicating that DIM-5 is not limiting and this interaction is not compromised in dim-3 cells.

We also monitored DIM-9-3XHA binding to DDB1-3xFLAG. Equal levels of DDB1-3xFLAG were purified from dim-3 and dim-3 nuclei, and the DIM-9-3xHA interaction appeared normal (Fig. 5B), suggesting no disruption in DCAF-substrate adaptor binding by the dim-3 mutation. We then monitored the nuclear interaction between DIM-7-3xFLAG and DIM-9-3xHA, which were previously shown to directly interact [15]. DIM-9-3xHA showed an increased interaction with DIM-7-3xFLAG in dim-3 nuclei (Fig. 5C), potentially because DIM-9 levels are increased in dim-3 strains (Fig. 5C, input). In summary, all examined DCDC members showed normal or increased interactions in dim-3 nuclei, implying that the H3K9me3 loss is not due to impaired DCDC assembly in dim-3 strains.

**DIM-5 and DIM-7 are mislocalized from heterochromatin in dim-3 strains**

It remained possible that the DCDC is not properly localized to heterochromatin in dim-3 strains. To investigate this possibility, we examined the localization of DIM-5, DIM-7, and DIM-9 in dim-3 strains by DamID. DIM-5-DAM was previously shown to localize to heterochromatin, and this localization was dependent on DIM-7 [8]. We found that the dim-3 mutation caused substantially reduced localization of DIM-5-DAM at the three representative heterochromatic regions tested (8:A6, 2:B3, and 8:G3; Figs. 6A and S11A). This reduction was not exacerbated by histidine (S11B Fig.), in contrast to the case for DIM-2 localization (Fig. 2). Probing for euchromatin regions (hH3 and pan-1) revealed that DIM-5-DAM is not misdirected to euchromatin (Figs. 6A and S11A).

The C-terminus of DIM-7 is critical for normal function [8], such that DAM-tagged DIM-7 does not fully complement dim-7 mutations. Nevertheless, as with DIM-5-DAM, we found that association of DIM-7-DAM with the heterochromatin regions (8:A6, 2:B3, and 8:G3) was also markedly reduced in the dim-3 background (Figs. 6B and S11C), suggesting that NUP-6dim-3 impacts targeting of DIM-7-DAM to heterochromatin. In contrast, DamID of DIM-9-DAM did not reveal marked differences between the dim-3 and dim-3 strains (Figs. 6B and S11C). We note that DIM-7-DAM and DIM-9-DAM may show a greater association with the euchromatic gene tested (hH3) than did DIM-5-DAM, perhaps reflecting an unknown role of these proteins outside of heterochromatin. The interaction of DIM-7-DAM, but not DIM-9-DAM, with the euchromatin marker was reduced in the dim-3 strain (Figs. 6A and S11C).

To confirm that DIM-7-DAM is mislocalized, we expressed DIM-7-GFP in dim-3+ and dim-3 vegetative tissue. The level of DIM-7-GFP produced from the native dim-7 promoter was not cytologically detectable, leading us to express it under the control of the stronger cgg-1 promoter. Despite overexpression, DIM-7-GFP co-localized with HP1-mCherry (S12 Fig.), at least in the cells examined (conidia), suggesting that DIM-7-GFP behaves normally to mark heterochromatic regions in vivo despite the importance of the DIM-7 C-terminus [8]. In 93.5% of dim-3+ cells DIM-7-GFP formed compact foci, typically at or near the nuclear periphery (Figs. 6C and S12), consistent with DIM-7-GFP marking heterochromatic regions. Interestingly, approximately half of dim-3 nuclei examined also showed such foci (Fig. 6D). It would be interesting to learn whether the observed difference between dim-3 and dim-3+ cells reflect differences in cell cycles of these strains. Unfortunately, because no genetic method to synchronize Neurospora has been developed, our studies were limited to unsynchronized cells. DIM-7-GFP appears equivalently expressed and transported into nuclei of dim-3+ and dim-3 strains (S13 Fig.). Thus, the dim-3 mutation seems to partially perturb normal localization of DIM-7 fusion proteins within the nucleus.
NUP-6$^{\text{dim-3}}$ is predominantly delocalized from the nuclear periphery.

The cytological mislocalization of DIM-7-GFP in a $\text{dim-3}$ mutant led us to examine the distribution of NUP-6 and NUP-6$^{\text{dim-3}}$ as well. Our expectation, based on work in other systems...
was that NUP-6 would be largely associated with the nuclear periphery. Indeed, at least in conidia, the majority of NUP-6-GFP was nuclear, but consistent with its role in nuclear transport, a small percentage was observed in the cytoplasm. Most (75.3\%) dim-3\+ cells showed NUP-6-GFP localization near the nuclear membrane (NM), forming either foci or a contiguous ring surrounding the genomic DNA (Fig. 7A, upper panel; additional examples of NUP-6-GFP localized at the nuclear periphery are provided in S14 Fig.; Z-stack fluorescent images in S1 Movie). We note that the vast majority of the NM-associated NUP-6 does not co-localize with HP1-marked heterochromatin (S15 Fig.) although some overlap of fluorescent signals was occasionally observed (yellow arrows) leaving open the possibility that NUP-6 is transiently associated with heterochromatin. A minority of dim-3\+ cells (24.7\%) show diffuse NUP-6-GFP in the nucleus (Fig. 7A, lower panel). We do not know the basis for the variability in nuclear localization of NUP-6\+GFP but it is noteworthy that, because of technical limitations, our study used unsynchronized cells, leaving open the possibility that the protein is differentially distributed in the cell cycle and that the dim-3 mutation affects the cell cycle.

In contrast to the case in dim-3\+ cells, a majority of dim-3 conidia (73.1\%) showed diffuse localization of NUP-6\+dim-3\-GFP in the nucleus (Fig. 7B, lower panel; additional examples of dispersed, nuclear NUP-6\+dim-3\-GFP are provided in S16 Fig.; also evident in Z-stacks of NUP-6\+dim-3\-GFP, S2 Movie). Nevertheless, 26.9\% of dim-3 conidia still showed compact structures at the nuclear periphery, similar to those observed with wild type NUP-6, consistent with the partial phenotype of dim-3 strains. The NUP-6\-GFP foci were found at the nuclear envelope, as shown by co-localization with a tagged component of the nuclear pore complex, NUP-84-mCherry ([35,36]; XP_964074.1; Fig. 7C); NUP-6\+dim-3\-GFP was not seen to localize at the nuclear envelope (Fig. 7D). Thus, both NUP-6 and DIM-7 appear to be partially mislocalized from the nuclear envelope in the dim-3 background, raising the possibility that mislocalization of NUP-6 leads to mislocalization of DIM-7 and other members of DCDC.

GCN-5 has increased euchromatic localization in a dim-3 strain

The results presented above suggest that NUP-6 plays a role in the normal localization of DCDC at heterochromatin. To explore the possibility that NUP-6 is also involved in localizing proteins associated with euchromatin, we examined the histone acetyltransferase complex SAGA, which is normally associated with a subset of gene promoters [37,38]. We built C-terminal DAM fusions for two components of the complex, GCN-5 (EDO65389.2), the acetyltransferase subunit, and TAF-5 (XP_961292.2), a structural subunit, and tested their localization by DamID at the euchromatic am (amination deficient) gene (EAA32325.1) and the hpo gene. Relative to the situation in the wildtype strain, the dim-3 strain, GCN-5-DAM showed increased association to am and hpo, while TAF-5-DAM showed no marked change in its association with these genes (Fig. 7E). As expected, these SAGA members were not found at heterochromatin (Fig. 7F). We conclude that NUP-6 differentially influences the localization of members of DCDC and SAGA, representatives of chromatin complexes normally found in heterochromatin and euchromatin, respectively.

Discussion

Our characterization of the Neurospora dim-3 mutation, which causes a partial loss of DNA methylation [7], led us to discover an unexpected function for the nuclear transport protein NUP-6 (Importin α). We mapped dim-3 to a region of Linkage Group V and determined that a mutation causing a glutamic acid to lysine substitution (E396K) in nup-6 is responsible for the methylation defect. NUP-6 is the Neurospora homolog of the Xenopus / Drosophila Importin α, yeast Srp1p, and human Karyopherin α. As first demonstrated with Xenopus
Fig 7. Abnormal localization of NUP-6^{dim-3} and increased euchromatic localization of SAGA. [A-B] Representative differential interference contrast (DIC), GFP fluorescent, Hoechst33342 stained DNA, and merged images of [A] dim-3^{+} cells expressing NUP-6-GFP or [B] dim-3 cells expressing NUP-6^{dim-3}-GFP. Each panel displays one conidium, with one to three nuclei visualized. Top panels show examples of localization at nuclear periphery and bottom panels show examples of dispersed nuclear localization. Percentages of conidia exhibiting each pattern are listed on the right; the P value for loss of membrane localization is \(<1\times10^{-124}\) (\(X^2\)-test). Densely staining DNA foci are thought to represent centromeric heterochromatin [11]. Scale bar indicates 5 \(\mu m\). [C-D] Differential interference contrast (DIC), GFP fluorescence, mCherry fluorescence, and merged images of strains expressing the nuclear pore complex member NUP-84-mCherry and [C] NUP-6-GFP or [D] NUP-6^{dim-3}:GFP. Scale bar indicates 5 \(\mu m\). For C, one conidium with three nuclei (one out of the focal plane) is displayed. For D, a conidium with one nucleus is displayed. [E-F] DamID experiment, as in Fig. 3, of SAGA complex members GCN-5-DAM and TAF-5-DAM in dim-3^{+} (WT) and dim-3 strains, probed for [E] euchromatic regions (am, hpo) or [F] a heterochromatic region (8:A6). [G] Illustrations depicting how heterochromatin and DNA methylation depend, at least in part, on a nuclear transport-independent role of NUP-6. In wild type strains (left), DCDC components are hypothetically transported into the nucleus by the NUP-6/Importin \(\beta\) dimer through the Nuclear Pore Complex (NPC), and NUP-6 facilitates formation of active DCDC, which
Importin α [39], this protein serves as an adaptor for classical nucleocytoplasmic transport, shuttling proteins from the cytoplasm to the nucleus through the nuclear pore complex [17–19]. We found that in addition to its defect in DNA methylation (Fig. 1A-B), dim-3 strains have an even more pronounced deficiency in H3K9me3 (Fig. 3A-B) and show reactivation of the silenced telomeric telVR::hph marker (Fig. 3D), indicating that heterochromatin is compromised in dim-3 strains. Interestingly, results of DamID suggest that both HP1 and DIM-2 can appropriately localize to heterochromatin in dim-3 strains grown in minimal medium (Fig. 2C-D). Apparently the reduced H3K9me3 levels in dim-3 strains still allow nucleation of near-wild type levels of HP1, which in turn recruits DIM-2, resulting in only moderately reduced DNA methylation (Fig. 2A). Histidine supplementation exacerbates loss of DNA methylation in the dim-3 mutant (Fig. 3A) and results in mislocalization of DIM-2 (Fig. 2C) without causing further reduction of H3K9me3 or loss of HP1 association (Figs. 3B, 2D and S7B). This curious histidine effect, which is not found with wild type strains and is independent of induction of DNA repair processes by histidine [26] and other agents, likely results by compromising the DIM-2—HP1 interaction or by perturbing DIM-2 activity.

Considering that H3K9me3 depends on a five protein complex, DCDC (DIM-5/-7/-9, CUL4, DDB1 Complex), it seemed likely that the reduced tri-methylation of H3K9 caused by the dim-3 mutation resulted from defective nuclear transport of at least one member of the DCDC. Surprisingly, we observed normal, or increased, nuclear levels of all DCDC components (Figs. 4A-B and S9). Similarly, HP1 and DIM-2, which are downstream of H3K9me3 in the DNA methylation pathway, showed no reduction in nuclei (Figs. 2B and S6). Likewise, a canonical NLS-containing import substrate was effectively transported into dim-3 nuclei (Fig. 4C). Together, these observations suggest that NUP-6\textsubscript{dim-3} has no nucleocytoplasmic transport defect.

Because NUP-6\textsubscript{dim-3} does not appear to compromise nucleocytoplasmic transport of the DNA methylation machinery, we investigated whether NUP-6 is involved in targeting DCDC to heterochromatin. DamID experiments revealed that DIM-7 and DIM-5 association with heterochromatin is reduced in a dim-3 strain (Fig. 6A-B). Moreover, cytological examination of DIM-7-GFP showed a marked reduction in foci in the dim-3 background (Fig. 6C-D) despite being equivalently expressed and transported in the different backgrounds (S13 Fig.), suggesting that NUP-6 is necessary for proper localization of one or more members of the DCDC. This mislocalization presumably results in other defects noted in dim-3 strains, including the large reduction in H3K9me3 (Fig. 3A-C), the reduced HP1-GFP localization (S6 Fig.), and the loss of telVR::hph silencing (Fig. 3D). Interestingly, despite being mislocalized, the DIM-5 and DIM-7 interactions with other DCDC components appear normal (Fig. 5), suggesting that formation of this complex does not depend on localization at its normal sub-nuclear site. Since the remaining H3K9me3 in dim-3 is only found at its normal locations (Fig. 3C), it is conceivable that the mislocalized DCDC does not tri-methylate H3K9 because the DCDC requires some signal from the sequences that underlie heterochromatin for its activity. The observation that DIM-9 localization to heterochromatin was only minimally disrupted in a dim-3 strain (Fig. 6B), raises the possibility that DIM-9, DDB1, and CUL4 use a different mechanism from that of DIM-5 and DIM-7 for heterochromatin targeting to form the DCDC.
We conclude that dim-3 strains are not defective in nuclear transport, but are partially defective in localizing some of the heterochromatin machinery. We were curious whether euchromatin machinery might also be affected, either positively or negatively. DamID analyses of two members of the SAGA histone acetyltransferase complex revealed increased localization of one of the proteins (GCN-5) to euchromatin loci in the dim-3 background (Fig. 7C). Interestingly, studies in Drosophila demonstrated an interaction between Importin α and the chromatin insulator protein CTCF [40].

There have been previous clues from S. cerevisiae and Xenopus of nuclear transport-independent roles of Importin α homologs. In yeast, srp1 was originally isolated as a dominant suppressor of ribosomal RNA-specific Polymerase I mutations [33], yet two of the characterized srp1 alleles had markedly different phenotypes that showed intragenic complementation [41,42]. The srp1-31 allele is defective in the nuclear transport of mitotic control proteins [43,44], whereas the srp1-49 allele is proficient for nuclear transport, but shows defects in nucleolus formation and proteasome function [41,42]. In Xenopus, Importin α inhibits the onset of mitosis by binding the mitotic-promotion factor TPX2 in non-mitotic cell cycle stages [32].

Nuclear transport-independent functions of NUP-6 may require a stable localization to specific sub-nuclear regions. In Xenopus, nuclear membrane (NM) associated Importin α promotes nuclear envelope formation independent of the nuclear transport components Importin β, Ran, or CAS [45], and in yeast, Srp1p is in proximity of the NM [31,33,34]. We observed a similar localization for NUP-6*-GFP in Neurospora conidia (Fig. 7A), while NUP-6dim-3-GFP was predominantly found away from the NM (Fig. 7B); the localization of NUP-6-GFP at the NM may vary with stages of the cell cycle and might be differentially controlled in individual cells, possibilities we cannot currently address. The apparent association of NUP-6 with the NM might be mediated through an interaction with the nuclear pore complex (NPC), as we found that NUP-6*-GFP co-localized with the NPC member NUP-84 (Fig. 7C). An association of NUP-6 with the inner NM could provide a platform to target chromatin complexes (Fig. 7G, left). We speculate that the dim-3 change in NUP-6 compromises this targeting (Fig. 7G, right). Of possible relevance to our findings suggesting a nuclear transport-independent function of NUP-6 / Importin α for heterochromatin formation and DNA methylation, the yeast NPC member Nup170p was recently reported to directly interact with chromatin, including the nucleolar-ribosomal DNA repeats and sub-telomeric regions to repress transcription [46]. Other nucleocytoplasmic-independent roles for the nuclear trafficking machinery have been suggested, including targeting Argonaute proteins to cytoplasmic P bodies for miRNA-mediated gene silencing by the human Importin β homolog, Importin-8 [47]. It is also noteworthy that NPC members, including Nup98, Sec13, and Nup50, have been found in NPC-independent nucleoplasm "pools" that directly interact with promoters of developmental or cell cycle genes to activate transcription and increase epigenetic memory through a transport-independent mechanism [48–50]. It will interesting to learn if NUP-6, nucleoporins, or other components of nucleocytoplasmic transport machinery or the NPC mediate heterochromatin formation, DNA methylation, and other functions through transport-independent processes in higher eukaryotes.

Materials and Methods

Detailed protocols (S1 Text), list of strains (S1 Table), and oligonucleotide sequences (S2 Table) are available online.
Neurospora growth, strain building, and plasmid construction

*Neurospora crassa* strains were grown, maintained, and crossed following standard protocols [51]. All nucleic acid manipulation protocols are given in S1 Text. Gene replacement constructs were generated as described [52] with modifications detailed in S1 Text.

Nucleic acid and protein manipulations

Genomic DNA isolation (modified from [53]) and Southern blotting [54] were performed as previously described. Primers to amplify DNA from methylated or control regions are listed in S2 Table or [20]. Whole genome sequencing was performed as previously described [55].

ChIP-seq was performed as previously described [56]. To display ChIP-seq data, sequencing reads were mapped to the Neurospora genome using Bowtie 2 [57]. Bam files were converted to read density tdf files spanning 200bp windows using the count function of Integrative Genomics Viewer’s (IGV) IGVtools, and were displayed using the IGV (www.broadinstitute.org/igv/; [58]). H3K9me3 ChIP-seq from a wild type strain has been previously described [56]; for the H3K9me3 ChIP-seq from *dim-3*, 11,662,061 reads (77.63% of total) mapped at least once to the Neurospora genome. The Y-axis shows the number of mapped reads at each position multiplied by $10^6$ and divided by the total number of mapped reads [58], providing normalization between samples.

For bisulfite sequencing, genomic DNA was isolated from strains grown at 32°C with shaking for ~48 hours in Vogel’s liquid medium, and 50ng was treated with bisulfite using the EZ DNA Methylation Lightning kit (ZYMO Research). Libraries were prepared for sequencing with the EpiGnome Methyl-Seq Kit and EpiGnome Index PCR Primers (Epicentre), purified with Agencourt AMPure XP beads (Beckman Coulter), Qubit HS Assay quantified (Life Technologies), visualized on the Fragment Analyzer (Advanced Analytical), and sequenced on an Illumina HiSeq 2000 (www.illumina.com) at the UO Genomics Core Facility as single-end 100nt reads. The BRAT-BW software package (compbio.cs.ucr.edu/brat/; [59]) was used to prepare and map the reads to the *N. crassa* OR74A (annotation NC12) genome, which was converted to a four stranded reference genome to permit bisulfite mapping. BRAT-BW acgt-count “-B” option cytosine-only files produced for the forward and reverse strand reads were merged. A python script (bidensity) was written to calculate the average 5mC level over a specified sliding window across the genome, producing a .wig file for IGV display.

For qRT-PCR analyses, RNA was isolated as previously described [60]. Qubit-RNA assay quantified (Life Technologies), and equal levels of RNA were DNase treated (Invitrogen) and the first strand cDNA was synthesized with Superscript III Kit (Life Technologies) following the manufacturers’ protocol. Diluted cDNA was used for triplicate quantitative real-time PCR experiments using FAST SYBR Green master mix (KAPA) with the primers 4701/4702 (*dim-7*), 4661/4662 (*ddb-1*), and 4663/4664 (*dim-9*), and normalized to *actin* (Accession number U78026.1; primers 3209, 3210) (S2 Table) on a Step One Plus Real Time PCR System (Life Technologies).

Immunoprecipitation and western blotting was performed as previously described [52] using antibodies listed in S1 Text. Extraction of histones was performed as previously described [61].

Accession numbers

Data files of H3K9me3 ChIP-seq and Bisulfite-seq have been deposited to GEO (http://www.ncbi.nlm.nih.gov/geo/) under the accession numbers (GSE61173; ChIP-seq) and (GSE61174; BS-seq). The accession number GSE61175 reports both data sets.
Supporting Information

S1 Fig. dim-3 has a genome wide loss of DNA methylation which is exacerbated by histidine. Bisulfite sequencing (BS-seq) demonstrating the genome-wide cytosine methylation in dim-3 (wildtype; “WT”, black track) and dim-3 strains (grown in minimal medium = red track; grown in medium containing histidine = blue track) displayed by Integrative Genomics Viewer, with y-axis denoting the number of normalized mapped reads (reads’x10^6/total number of mapped reads; [58]). Genes are displayed on the x-axis below the DNA methylation peaks as green vertical lines, while distances (in Megabases) from the left end of each Linkage Group (LG) are displayed above the graph. Due to the reduced cytosine methylation in the dim-3 samples, the signal-to-noise ratio is lower, which renders the background more prominent.

S2 Fig. NUP-6 structure; NUP-6^{dim-3} has normal expression and nuclear transport; dim-3 strains exhibit slower growth rates. [A] Structural model of NUP-6 (Importin α; [22] shown in “ribbons” graphic, with N- and C-termini labeled. Residues found to be mutated in the dim-3 strain are colored red (E396K, the causative mutation) and orange (R469H) and displayed as “ball and stick” graphics on the structure. Modeling was done with Pymol (www.pymol.org). [B] Total (T) extract, as well as the cytoplasmic (C) and nuclear (N) fractions from strains expressing NUP-6-T3xFLAG or NUP-6^{dim-3}-3xFLAG, as well as a strain lacking any 3xFLAG-tagged protein, were analyzed by standard western procedures, blotting for the FLAG tagged protein as well as control cytoplasmic (phosphoglycerate kinase; α-PGK) and nuclear (histone H3) proteins. Quantification of three separate experiments, with error shown, presented as the percent of cytoplasmic and nuclear NUP-6-T3xFLAG are graphed below. [C] Isolated nuclei from strains expressing NUP-6-T3xFLAG and NUP-6^{dim-3}-3xFLAG, as well as a strain lacking any 3xFLAG-tagged protein, were analyzed by standard western procedures, blotting for the FLAG tagged protein or the histone H3 loading control. The reproducible higher mobility of NUP-6^{dim-3}-3xFLAG might be indicative of post-translational modification(s), as Xenopus Importin α has been reported to have multiple phosphorylation sites [45]. Quantification of three separate experiments, with error shown, presented as the ratio of NUP-6 protein normalized to the nuclear level of NUP-6-T3xFLAG are shown below. [D] Linear growth rate (centimeters per hour; cm/hr) of four wild type Neurospora strains (N5622-N5625, black; average growth rate = 0.36 ±0.004 cm/hr, average R^2 = 0.99) and four dim-3 strains (N5628-N5631, red; average growth rate = 0.20 ±0.01 cm/hr, average R^2 = 0.99), as measured by race tube growth.

S3 Fig. Reintroduction of the dim-3 allele or the E396K mutation cause a loss of DNA methylation. Heterochromatic (8:G3 or 2:B3) or euchromatic (am) region Southern blot assays (as in Fig. 1A) of indicated mutations reintroduced into a dim-3 (WT) strain, along with methylation mutant (dim-5, dim-3) and WT controls. “dim-3” strain has both mutations reintroduced while “WT” control shows the effect of just the hygromycin targeting cassette.

S4 Fig. Alanine and methionine, but not other amino acids, negate the histidine effect on DNA methylation. Southern blot of genomic DNA digested with the 5mC-sensitive enzyme AvaI to assess DNA methylation levels in a dim-3 (WT) strain supplemented with histidine, or dim-3 strains supplemented with additional amino acids, as listed, or those same amino acids in tandem with histidine. Supplemented methionine presumably restores DNA methylation by increasing internal S-adenosylmethionine (SAM) levels, as shown previously [63].
Southern blotting is done as Fig. 1A.

S5 Fig. Histidine specifically causes a loss of DNA methylation in dim-3. (A) Southern blot of genomic DNA digested with the 5mC-sensitive enzyme AvaII to assess DNA methylation levels in dim-3” (WT) or dim-3 prototrophic strains grown in the absence of chemicals (-) or in the presence of histidine (HIS), hydroxyuridine (HU) or ethylmethylsulfanate (EMS). Note that all of these chemicals induce QDE-2 interacting RNAs (qiRNAs), which reportedly are indicative DNA damage or a compromised genome [26]. Southern blotting is done as Fig. 1A. (B) Southern blot of genomic DNA digested with the 5mC-sensitive enzyme AvaII to assess DNA methylation levels in WT or dim-3 strains deleted of the Neurospora Argonaut gene, qde-2. Southern blotting is done as Fig. 1A.

S6 Fig. HP1-GFP is effectively transported into the nucleus in dim-3. Differential Interference Contrast (DIC) and fluorescent microscopy images of HP1-GFP expressed in dim-3” (WT) and dim-3 conidia (one pair shown for each) demonstrating that HP1 is transported into nuclei. Scale bar indicates 5μm. HP1-GFP concentrates as green foci in WT cells (top), which co-localizes with DNA [11], but appears mislocalized (single focus) in dim-3 nuclei (bottom) as observed with Δdim-7 [8], presumably due to reduced H3K9me3 levels [8,11,15,61,64].

S7 Fig. Histidine displaces DIM-2-DAM, but not HP1-DAM from heterochromatin. Southern blot assay of DamID experiments with dim-3” (WT) and dim-3 strains containing [A] DIM-2-DAM or [B] HP1-DAM grown with or without histidine and probed for the heterochromatic region 8:A6 or the euchromatic region pan-1. Genomic DNA was digested with DpnI (DI), specifically cutting GA^mTC sequences or with the A^m- insensitive isoschizomer DpnII (DII) to monitor complete digestion.

S8 Fig. dim-3 has a genome-wide loss of H3K9me3. Chromatin Immunoprecipitation sequencing (ChIP-seq) of H3K9me3 from dim-3” (wildtype, “WT”, black track) [56] and a dim-3 strain (grown in minimal medium, light gray track) displayed by Integrative Genomics Viewer, with y-axis denoting the approximate number of normalized mapped reads (reads x10^6/total number of mapped reads; [58]). Genes are displayed on the x-axis below the H3K9me3 peaks as vertical lines, while distances (in Megabases) from the left end of each Linkage Group (LG) are displayed above the graph. Due to the vastly reduced trimethylation of H3K9 in the dim-3 strain, the signal-to-noise ratio is reduced, rendering the background more prominent.

S9 Fig. Nuclear transport of DCDC components is unaffected by the dim-3 mutation. α-FLAG and α-histone H3 (hH3; loading control) western blots of dim-3” (WT) or dim-3 nuclei expressing individually FLAG-tagged DCDC components, as in Fig. 4A, except non-auxotrophic strains were grown in minimum medium. Average and standard deviation of nuclear FLAG-tagged protein levels of three experiments are indicated below. Representative images are shown.

S10 Fig. Transcription of dim-7, ddb-1, and dim-9 is minimally altered in a dim-3 strain. Fold change of mRNA levels in dim-3” (WT) and dim-3 strains, as assessed by quantitative RT-PCR using the primers oAK262/oAK263 (dim-7), oAK222/223 (ddb-1), and oAK224/225 (dim-9), with signal normalized to actin (primers 3209, 3210), is displayed as a bar chart.
Standard deviation of three biological replicates, each with at least three technical replicates, is shown. Inset shows a representative ethidium bromide-stained agarose gel image showing comparable levels of input RNA used for the qPCR reaction (doublet of ribosomal RNA shown). (EPS)

S11 Fig. DIM-5-DAM and DIM-7-DAM are mislocalized from heterochromatin, and histidine does not further displace DIM-5-DAM. [A] DamID Southern blot experiments of the heterochromatic region 8:G3, as in Fig. 3 and 6, with two independent samples of dim-3- (WT) and dim-3 strains expressing DIM-5-DAM. [B] DamID experiments with dim-3- (WT) and dim-3 expressing DIM-5-DAM grown in minimum medium (min) or medium containing histidine (+HIS). Δdim-7; DIM-5-DAM sample indicates a complete loss of DIM-5-DAM targeting [8]. Southern blots from three heterochromatic regions (8:A6, 2:B3, and 8:G3) and one euchromatic region (pan-1) are shown. [C] DamID experiments with DIM-7-DAM and DIM-9-DAM. (EPS)

S12 Fig. DIM-7-GFP co-localizes with HP1-mCherry marked heterochromatin and is near the inner nuclear periphery. (A) Representative differential interference contrast (DIC), GFP fluorescent, mCherry fluorescent, and merged images of P_crg::DIM-7-GFP and HP1-mCherry in a WT strain. One conidium with two nuclei is displayed. Scale bar indicates 5μm. (B) Representative differential interference contrast (DIC), GFP fluorescent, mCherry fluorescent, and merged images of P_crg::NCA-1-GFP (CAB65295.1), which marks the nuclear and cell membranes [65] and P_crg::DIM-7-mCherry in a WT background. One conidium with one nucleus is displayed. (EPS)

S13 Fig. P_crg::DIM-7-GFP exhibits equal expression and nuclear transport between dim-3+ and dim-3 strains. [A] α-GFP western blots of whole cell extract of either a dim-3+ (WT) strain, or dim-3+ or dim-3 strains expressing P_crg::DIM-7-GFP. Top blot shows the levels of DIM-7-GFP fusion protein (predicted molecular weight: 102.5KDa), while the bottom blot shows the position of free GFP (predicted molecular weight: 26.0KDa). PonceauS-stained membrane serves as a loading control. [B] α-GFP western blot showing the total culture, cytoplasmic fraction, and nuclear fraction levels of overexpressed DIM-7-GFP found in nuclei isolated from dim-3+ (WT) or dim-3 strains expressing this construct. PonceauS-stained membrane of proteins present at the dye front serves as a loading control. (EPS)

S14 Fig. Punctate localization NUP-6-GFP. [A-B] Additional examples, relative to Fig. 7A, of differential interference contrast (DIC), GFP fluorescent, Hoechst33342 stained DNA, and merged images of [A] dim-3+cells expressing NUP-6-GFP from the native promoter, or [B] overexpressed NUP-6-GFP using the ccr promoter in dim-3 cells. Each panel displays one conidium, with one to three nuclei. Densely staining DNA foci are thought to represent centromeric heterochromatin [11]. Scale bar indicates 5μm. For overexpressed NUP-6-GFP, 83.6% of conidia (n = 544) exhibited a nuclear periphery localization while 16.4% of conidia (n = 107) had a nuclear localization; the P value for loss of membrane localization is <1x10^-33 (X²-test). (EPS)

S15 Fig. NUP-6-GFP minimally co-localizes with HP1-mCherry-marked heterochromatin. Differential interference contrast (DIC), GFP fluorescent, mCherry fluorescent, and merged images of NUP-6-GFP and HP1-mCherry in a WT strain background. Yellow arrows mark
potential co-localization sites. Four independent examples are shown, and scale bar indicates 5μm. For each panel, one conidium showing one or two nuclei is displayed.

(S16 Fig. Dispersed localization of NUP-6\textsuperscript{dim-3}-GFP. [A-B] Additional examples of differential interference contrast (DIC), GFP fluorescent, Hoechst33342 stained DNA, and merged images of [A] \textit{dim-3} cells expressing NUP-6\textsuperscript{dim-3}-GFP from the native promoter, as in Fig. 7B, or [B] overexpressed NUP-6\textsuperscript{dim-3}-GFP using the \textit{ccg} promoter in \textit{dim-3} cells. Each panel displays one conidium, with one to three nuclei. Densely staining DNA foci are thought to represent centromeric heterochromatin [11]. Scale bar indicates 5μm. For overexpressed NUP-6\textsuperscript{dim-3}-GFP, 54.8% of conidia (n = 583) exhibited a nuclear periphery localization while 45.2% of conidia (n = 481) had a nuclear localization; the P value for loss of membrane localization is <1x10\textsuperscript{-33} (X\textsuperscript{2}-test).

(AEPS)

S1 Movie. NUP-6\textsuperscript{dim-3}-GFP forms INM structures inside the nucleus. Z-stack image movie of NUP-6-GFP constructed with fluorescent images and compiled using ImageJ. One conidium with three nuclei is displayed.

(AVII)

S2 Movie. NUP-6\textsuperscript{dim-3}-GFP is mislocalized from the INM. Z-stack image movie of NUP-6\textsuperscript{dim-3}-GFP constructed with fluorescent images and compiled using ImageJ. One conidium with two nuclei is displayed.

(AVII)

S1 Table. Strains used in this study.

(DO CX)

S2 Table. Oligonucleotides used in this study.

(DO CX)

S1 Text. Detailed Materials and Methods.

(DO CX)

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Author Contributions

Conceived and designed the experiments: ADK MRR EUS SMH KKA. Performed the experiments: ADK MRR SMH KKA PLG. Analyzed the data: ADK MRR EUS SMH KKA. Wrote the paper: ADK MRR EUS SMH KKA PLG.

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