Mgm101p Is a Novel Component of the Mitochondrial Nucleoid That Binds DNA and Is Required for the Repair of Oxidatively Damaged Mitochondrial DNA

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Abstract. Maintenance of mitochondrial DNA (mtDNA) during cell division is required for progeny to be respiratory competent. Maintenance involves the replication, repair, assembly, segregation, and partitioning of the mitochondrial nucleoid. MGM101 has been identified as a gene essential for mtDNA maintenance in S. cerevisiae, but its role is unknown. Using liquid chromatography coupled with tandem mass spectrometry, we identified Mgm101p as a component of highly enriched nucleoids, suggesting that it plays a nucleoid-specific role in maintenance. Subcellular fractionation, indirect immunofluorescence and GFP tagging show that Mgm101p is exclusively associated with the mitochondrial nucleoid structure in cells. Furthermore, DNA affinity chromatography of nucleoid extracts indicates that Mgm101p binds to DNA, suggesting that its nucleoid localization is in part due to this activity. Phenotypic analysis of cells containing a temperature-sensitive mgm101 allele suggests that Mgm101p is not involved in mtDNA packaging, segregation, partitioning or required for ongoing mtDNA replication. We examined Mgm101p’s role in mtDNA repair. A strain compared with wild-type cells, mgm101 cells were more sensitive to mtDNA damage induced by UV irradiation and were hypersensitive to mtDNA damage induced by gamma rays and H2O2 treatment. Thus, we propose that Mgm101p performs an essential function in the repair of oxidatively damaged mtDNA that is required for the maintenance of the mitochondrial genome.

Key words: mtDNA • nucleoid • maintenance • DNA repair • yeast

An increasing number of human diseases are being identified that are associated with mitochondrial DNA (mtDNA)-linked mutations (Brown and Wallace, 1994; Sherratt et al., 1997). To understand the etiology of these diseases, the details of the basic molecular mechanisms underlying the maintenance of mtDNA must first be understood. The accurate maintenance of mtDNA requires several distinct processes. A case is the case for nuclear DNA, mtDNA must be replicated and packaged into nucleoids. The nucleoids then must be segregated from one another and partitioned into daughter cells. The repair of damaged mtDNA is also likely to be important for mtDNA maintenance, especially considering the highly oxidative environment of the organelle. Defects in any one of these processes might result, at a minimum, in nonfunctional mtDNA, and may cause the complete loss of mtDNA, a lethal event in humans. The budding yeast Saccharomyces cerevisiae has been used to analyze these processes in more detail. In addition to being tractable to genetic, biochemical and cytological approaches, yeast are facultative anaerobes. Thus, S. cerevisiae cells that lose mtDNA can still be propagated and studied, when a fermentable carbon source such as glucose is provided.

In S. cerevisiae, there are an average of 50–100 of the 80-kb mitochondrial genome copies, depending on growth conditions and ploidy (Williamson, 1976). The genome is known to encode seven proteins that include subunits of energy transducing inner membrane complexes, one mitochondrial ribosomal protein, several minor proteins involved in gene splicing and conversion, ribosomal and transfer RNAs, and one RNA involved in tRNA processing (Pon and Schatz, 1991). In vivo, mtDNA is organized

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1. Abbreviations used in this paper: MBP, maltose binding protein; mtDNA, mitochondrial DNA.
into punctate nucleoid structures that are distributed in a somewhat regular pattern within the mitochondrial reticulum. Each nucleoid has been estimated to contain ~4–5 copies of mtDNA and is packaged in a manner that makes it more resistant to digestion by exogenously added DNases (Williamson, 1976; Miyakawa et al., 1987, 1995; Newman et al., 1996). These structures have been shown to be the unit of inheritance for mtDNA (Lockshon et al., 1995).

Genetic studies in yeast have suggested that mtDNA partitioning into daughter cells is a nonrandom process (Coen et al., 1970; Birky, 1978; Strausberg and Perlman, 1978; Zinn et al., 1987; Aziproz and Butow, 1993). More recently, this model has been substantiated by direct observation and comparison of the behavior of mtDNA and mitochondrial proteins in yeast zygotes formed by mating rho+ (containing mtDNA) cells (Aziproz and Butow, 1993; Nunnari et al., 1997; Okamoto et al., 1998). Specifically, it has been observed, that while haploid-derived mitochondrial proteins are able to freely diffuse within yeast zygotic mitochondrial reticulum, haploid mtDNA remains segregated in the zygote (Nunnari et al., 1997). Although haploid-derived mtDNA does not move throughout the zygote, it preferentially moves into the first zygotic bud (Nunnari et al., 1997). Similar conclusions have been recently drawn from cytological experiments using zygotes formed by mating rho+ to rho− (lacking mtDNA) cells (Okamoto et al., 1998). These observations suggest that mtDNA is linked to putative partitioning machinery that insures accurate inheritance of mtDNA. To date, components of this proposed partitioning apparatus remain unidentified. Interestingly, a subset of mutations that cause abnormal mitochondrial morphology such as mdm10, mmm1, and fzo1 also result in mtDNA loss, suggesting that these components may be important for mtDNA inheritance (Burgess et al., 1994; Sogo and Yaffe, 1994; Hermann et al., 1998). No yeast proteins have been identified with convincing sequence similarity to eubacterial nucleoid segregation or division proteins, suggesting that novel mechanisms and components for these processes have arisen for mitochondria in eucaryotes.

In contrast to the partitioning apparatus, components directly involved in the replication, repair, and recombination of mtDNA have been described. Replication of mtDNA in S. cerevisiae is mediated by Mip1p, a pol-γ DNA polymerase that has been shown to possess both polymerizing and 3′ to 5′ exonuclease activity (Foury, 1989). This exonuclease activity, together with Msh1p, a MutS homologue required for mitochondrial DNA mismatch repair, mediates proofreading and thus the high fidelity replication of mtDNA (Vanderstraeten et al., 1998). Additional proteins likely to be involved in the replication, recombination, and repair of mtDNA include Rlm1p, the mitochondrial homologue of eubacterial SSB, Pif1p, a DNA helicase, Abf2p, a DNA binding protein in the HMG family, (Lahaye et al., 1991; Diffany and Stillman, 1992; Kao et al., 1993), Mgt1p, a cruciform-cutting endonuclease present in the mitochondrion and the genetic locus, MHR1, which corresponds to an unknown gene identified in a genetic screen for mtDNA recombination-deficient mutants (Ling et al., 1995; Lockshon et al., 1995). Although many genes important for mtDNA metabolism have been described, surprisingly little is known about the molecular mechanisms involved in mtDNA replication, recombination and repair.

**MGM101** (for mitochondrial genome maintenance) was isolated in a genetic screen for mutants that caused temperature-sensitive loss of mtDNA and a null mutation in MGM101 indicates that it is essential for the maintenance of mtDNA (Chen et al., 1993). However, the functional role that Mgm101p plays in mtDNA maintenance is unknown. Sequence analysis of Mgm101p indicates it is a novel, highly basic 30-kD protein (Chen et al., 1993). In addition to the S. cerevisiae MGM101, genes encoding the K. lactis and S. pombe MGM101 homologues have been identified (Chen et al., 1993; Clark-Walker and Chen, 1996; see Fig. 1). These gene products have an overall high degree of identity and similarity (Corpet, 1988). The most conserved region is contained in the COOH-terminal half of Mgm101p, where the majority of the basic residues are found, suggesting that this region is important for Mgm101p function. Deletion of a portion of the COOH terminus of S. cerevisiae Mgm101p causes a complete loss of function (Chen et al., 1993). The high degree of conservation between the evolutionarily divergent S. cerevisiae and S. pombe suggests that a metazoan Mgm101p homologue may exist. Given the likely importance of this novel protein in mtDNA maintenance, our goal was to characterize Mgm101p and, through the phenotypic analysis of a temperature-sensitive mgm101 allele, determine its essential role in the maintenance of the mitochondrial genome.

**Materials and Methods**

**Yeast Genetic Techniques**

*Yeast strains used in this study are listed in Table I. Media preparations and genetic techniques were performed as described (Guthrie and Fink, 1991). Either glucose (D at 2%) or glycerol (G at 2%) was used as a carbon source as specified.*

**Isolation and Characterization of mgm101-2 Cells**

*W303 haploid cells were mutagenized using either EMS, nitrosoguanidine or UV light to 10% viability and screened for temperature sensitive growth on YPG. Cells from these colonies were grown at 37°C on YPD for ~100 generations and were screened for loss of mtDNA using DAPI and fluorescence microscopy.*

**Complementation analysis using a recessive truncated loss of function allele of MGM101 (strain CS6-5D, kindly provided by G.D. Clark-Walker) revealed that one mutant from this screen was allelic to MGM101 (JNY131[mgm101-2]). The Australian National University. Sporulation and tetrad analysis of this cross confirmed the mutation was at the MGM101 locus.*

To characterize JNY131(mgm101-2), cultures of W303 or JNY131 (mgm101-2) were grown at permissive temperature (22°C) to log phase in YPG, washed into YPD and were incubated at either permissive or non-permissive temperatures. At various time intervals, aliquots were taken and cells were analyzed for respiratory competence, the presence of mtDNA and mitochondrial morphology. To analyze for respiratory competence, cells were plated onto YPD plates and colony color was assessed and quantitated. Because all strains analyzed contained the ade2 mutation, on YPD, white and sectored colonies were classified as respiratory deficient and red colonies were classified as respiratory competent. To visualize mtDNA, cells were fixed either in 70% ethanol containing 10 ng/ml DAPI or with 3.7% formaldehyde for 1 h, followed by spheroplasting and incubation in PBS containing 1 μg/ml DAPI. Fixed, stained cells were washed into PBS and imaged using standard epifluorescence microscopy. To analyze mitochondrial morphology, aliquots of cells were also processed for indirect immunofluorescence and imaged using confocal microscopy (see below).
Sequence Analysis of mgm101-2

The mutation in mgm101-2 was determined by double-stranded DNA sequence analysis. Yeast genomic DNA strains J NY 131 and W 303 was prepared as described (Guthrie and Fink, 1991) and used as template for polymerase chain reaction (PCR) with the following sets of primers corresponding to the MGM 101 gene: 5'-CGGAATTCCTCCTTGGACAACACTTTCGT-3' and 5'-GCTCTAGACATTTTAAAGGATATTCCAAAC-3', 5'-GGAGATTCTCCTTGAGCACAACCTTTGCTGT-3' and 5'-CGGAGATCCCGCACCATTCTTTCGATAGGATCCAG-3' (BRL-Custom Primers Inc.). The PCR products from five independent reactions with a given set of primers were pooled, gel purified, and sequenced using the above listed primers by the Division of Biological Sciences automated DNA Sequencing Facility at University of California, Davis.

Fusion Protein Expression and Antibody Production

For the production of anti-M-glm101p and anti-A-bf2p antibodies, maltose binding protein (MBP) fusions were created. M GM 101 and ABF2 were cloned into p MAL-c2 (New England Biolabs, Inc.) using an in-frame 5' EcoRI site and a 3' Xbal site introduced using PCR. Fusion proteins were expressed in E.cherichia coli (DH5α) at 25°C and purified using amylase affinity chromatography following the methods outlined by New England Biolabs, Inc. Rabbit polyclonal antibodies were raised against fusion proteins by injecting 1 mg of purified protein subcutaneously every 2 wk. An nti-M-glm101p antibody was affinity purified as described (Pringle et al., 1991).

Preparation of Mitochondria

Highly enriched yeast mitochondria were prepared essentially as described (Glick and Pon, 1995) with the following modifications. Cultures of either W 303 or J NY 131 were grown in Y PG media to log phase. Before harvesting, yeast cell walls were removed using yeast lytic enzyme (ICN). Isolated crude mitochondrial pellets from lysed cells were resuspended in 50% w/v optiprep (Nycoderm, 20 mM HEPES, pH 7.4, osmotically balanced with sorbitol to 600 m osmolar). This mixture was used as the bottom layer of optiprep step gradients formed in SW28 centrifuge tubes (Beckman); the layers atop it had densities of 1.1 g/ml and 1.16 g/ml, and were similarly osmotically balanced with sorbitol. Gradients were spun at 80,000 g for a minimum of 3 h in an SW28 rotor; mitochondria floated to the interface between the 1.1 and 1.16 g/ml. This interface was collected and diluted approximately fivefold with mitochondrial isolation buffer (MIB: 0.6 M sorbitol, 20 mM Hepes, pH 7.4) and spun at 100,000 g for 30 min to pellet the mitochondria. This pellet was resuspended, centrifuged at 3000 g for 5 min to pellet aggregated material and the supernate was centrifuged again at 100,000 g for 10 min and the pellet was resuspended in a minimal volume of MIB.

Isolation of Mitochondrial Nucleoids and Fractions Enriched in mtDNA binding Proteins

Mitochondrial nucleoids were prepared using a modification of a technique described by Miyakawa et al. (1987). Mitochondrial lysate was prepared by diluting purified mitochondria to a protein concentration of 2.5 mg/ml with nucleoid extraction buffer (NXB: 20 mM Hepes, pH 7.4, 100 mM sucrose, 20 mM KCl, 1% NP-40, 1 mM spermidine, 100 ng/ml DAPI, 1 mM PMSE, 0.5 mM DTT) followed by an incubation on ice for 1 h. This lysate was layered onto a 37.5%/60%/80% sucrose step gradient. In these gradients, the 37.5% layer contained NXB, and the 60% layer contained 20 mM Hepes, pH 7.4, 20 mM KCl, 1% M ega-8 (Calbiochem), 1 mM spermidine, 100 ng/ml DAPI, and 0.5 mM DTT. Gradients were spun at 100,000 g for 90 min; mitochondrial DNA -containing structures banded at the 37.5%/60% interface as assessed by fluorescence microscopy of DAPI stainable structures and enrichment of both DNA assayed using the picogreen reagent (Molecular Probes) and the nucleoid-associated protein Abf2p (see below). This material was collected and diluted twofold with NXB containing 2% NP-40 and reband over another 37.5%/60%/80% step gradient.

LC/MS/MS Analysis of Nucleoid-associated Proteins

To extract nucleoid-associated proteins from DNA, an equal volume of NXB containing 2 M KCl was added to the diluted nucleoid fraction. This mixture was incubated for 1–2 h on ice to produce a high salt extract enriched for nucleoid-associated proteins. This mixture was layered atop a 30% sucrose cushion containing NXB with 1 M KCl, and centrifuged at 100,000 g for 90 min. The top fraction, not including the top/30% interface, was collected. Protein in this fraction was precipitated by the addition of trichloroacetic acid to 15% wt/vol followed by centrifugation at 20,000 g for 30 min. The resulting pellet was rinsed twice with cold 80% ethanol and then resuspended in freshly prepared 8 M urea, 100 mM ammonium bicarbonate, pH 8.0, at room temperature. A filter 12 h at 4°C, the mixture was centrifuged at 20,000 g for 30 min. The protein concentration in the supernatant from this spin, termed HE 5 , was determined using the Nano-orange reagent kit (Molecular Probes). DTT was added to HSE to 4 mM, and the mixture was incubated for 30 min at 37°C then diluted fourfold with 50 mM Tris-HCl, pH 8.0, 20 mM KCl, 1 mM CaCl2, containing 1 μg of modified trypsin (Boehringer) for every 25 μg of protein. This mixture was incubated for 12 h at 37°C and then frozen in liquid nitrogen.

Nano-column high performance liquid chromatography coupled to electrospray ionization tandem mass spectrometry was used to identify the mixture of proteins present in the HSE fraction. The sample was loaded onto a 100 μm by 200 μm fused silica capillary (Polymerics, Inc.) that was packed to a length of ~15 cm with 10 μm POROS 10 R 2 reverse phase material (Perseptive Biosystems; K. Kennedy and Jorgeson, 1989). The fritted end of the column was inserted into the needle of the Finnigan TSQ-700 electrospray ion source. The sample was directly loaded onto the micro column by helium pressurization of the sample in a stainless steel bomb (K. Kennedy and Jorgeson, 1989).

Solvent flow from a dual syringe pump (Applied Biosystems) was split 100:1 pre-column to deliver a final flow rate of 1 to 1.5 μl/min through the column. By using a mobile phase consisting of 0.5% acetic acid (solvent A) and 80:20 acetonitrile/water containing 0.5% acetic acid (solvent B) liquid chromatography was performed by ramping from 0% B to 60% B over 90 min. Electrospray ionization was carried out at a voltage of 4.6 kV and tandem mass spectra were acquired automatically during the entire gradient run (Link et al., 1997). Each tandem mass spectrum was searched, using the SEQUEST program (Yates et al., 1996) against the yeast orfs protein database obtained from the Saccharomyces Genome Database (Stanford University). Sequences for bovine trypsin and human keratin were included to facilitate identification of potential contaminants. Each high scoring peptide sequence, and the corresponding tandem mass spectrum, was manually inspected to insure the match was correct.

DNA Cellulose Chromatography of Nucleoid-associated Proteins

Nucleoid-associated proteins were isolated from mitochondria and extracted from mitochondrial DNA according to the methods described above. Nucleoid-extracted proteins were dialyzed for 3 h at 4°C against a buffer containing 25 mM Tris, pH 7.5, 100 mM NaCl, 1 mM EDTA, 1 mM beta-mercaptoethanol. After dialysis, the extract was fractionated using native DNA cellulose chromatography as described (Alberts and Herrick, 1971).

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Table I. Yeast Strains

| Strain | Genotype | Source |
|--------|----------|--------|
| W303   | ade2-1, leu2-3, his3-11, 15, trpl-1, ura3-1, can1-100, Mata or Mata | R. Rothstein |
| CS6-1D | leu2, ura3, his4, ade2, ade3, MGM101::URA3, Mata | Chen et al., 1993 |
| JNY131 | same as W303, except mgm101-2, Mata, or Mata | This study |
| W303TK | same as W303, except can1::GPD-HSVTK, Mata | Numairi et al., 1997 |
| JNY131TK | same as JNY131, except can1::GPD-HSVTK, Mata | This study |
Rapid Preparation of Extracts for Western Blotting

Cultures were grown either in YPD or YPG to early log phase. 5 OD600 units were harvested by centrifugation and cells were washed once in distilled water. Cell pellets were resuspended in 75 μl SDS-PAGE sample buffer (2% SDS, 80 mM Tris-HCl, pH 6.8, 10% glycerol (vol/vol), 5% 2-mercaptoethanol, 1 mM phenylmethylsulfonyl fluoride, 3 mM benzamidine). The mixture was boiled for 10 min at 100°C, and centrifuged for 15 min at 13,000 g. The supernatant was used for SDS-PAGE and immunoblot analysis.

SDS-PAGE and Western Blotting

Proteins were analyzed using 17.5% SDS-PAGE, prepared according to (Anderson et al., 1973). For Western blotting, anti-Mgm101p, anti-Abf2p, and anti-Por1p (Molecular Probes) antibodies were used at dilutions of 1:1000, 1:100, and 1:500, respectively. Primary antibodies were incubated for a minimum of 1 h at 25°C, followed by incubation for a minimum of 30 min with secondary antibody at a dilution of 1:1000 (goat anti-rabbit HRP antibody and goat anti-mouse HRP antibody; Southern Biotechnology Associates, Inc.). Western blots were developed using ECL reagents (Amersham).

Indirect Immunofluorescence

To analyze mitochondrial morphology in strain JNY131, cells were shifted to nonpermissive temperature (37°C) and samples were taken at various time intervals and processed for indirect immunofluorescence. To localize Mgm101p, W303 was transformed with a 2μ plasmid containing MGM101 (kindly provided by G.D. Clark-Walker) and was grown in YPG at 30°C to log phase. At least 5 A600 units of cells were fixed and processed for rapid preparation of extracts for Western blotting (kindly provided by G.D. Clark-Walker) and was grown in YPG at 30°C for 8 min at 13,000 g.

BrdU Incorporation and Detection

Thymidine kinase gene from Herpes virus was introduced into JNY131(mgm101-2) and W303 by crossing and sporulation. These strains, JNY131TK and W303TK, were cultured overnight in YPG to 0.2–0.3 OD600 unit/ml. For BrdU incorporation, overnight cultures of cells were washed twice in 0.9% NaCl, placed on ice and exposed to a Cs source for various times corresponding to 0–600 grays. For H2O2 treatment, overnight cultures of cells were washed once in YPD, and resuspended in YPD containing 0–20 mM H2O2 and incubated at 25°C for 1 h. Cells were then harvested and washed twice in 20 mM potassium phosphate, pH 7.4. Immediately after exposure, cells were plated on YPD plates. A fiber exposure to each mutagen, colonies were formed at a semi-permissive temperature of 34°C (4-d incubation) and respiratory competency was determined by colony color as described.

UV, Gamma Ray, and H2O2 Exposure

W303 and JNY131(mgm101-2) strains were grown at 25°C overnight in YPG to 0.2–0.3 OD600 unit/ml. For UV-irradiation, both JNY131 (mgm101-2) and W303 cells were washed into YPD and preincubated at 37°C for 2 h before irradiation. Cells were then plated onto YPD plates and exposed to UV radiation (Stratalinker; Stratagene) at doses ranging from 0–200 J/m2. For gamma-irradiation, overnight cultures of cells were washed twice, resuspended in 0.9% NaCl at 2 × 106 cells/ml, placed on ice and exposed to a Cs source for various times corresponding to 0–600 grays. For H2O2 treatment, overnight cultures of cells were washed once in YPD, and resuspended in YPD containing 0–20 mM H2O2 and incubated at 25°C for 1 h. Cells were then harvested and washed twice in 20 mM potassium phosphate, pH 7.4. Immediately after exposure, cells were plated on YPD plates. A fiber exposure to each mutagen, colonies were formed at a semi-permissive temperature of 34°C (4-d incubation) and respiratory competency was determined by colony color as described.

Results

Independent Genetic and Biochemical Approaches Indicate That Mgm101p’s Function Is Nucleoid-specific

In an effort to understand the mechanism underlying mitochondrial nucleoid organization and function, we have started to identify the proteins found in highly enriched (70-fold from total extracts) preparations of nucleoids using the technique of liquid chromatography coupled with tandem mass spectrometry (see M aterials and M ethods; Y ates et al., 1996). U sing this technique, many individual proteins can be identified from a heterogeneous mixture of proteins, making it ideal for analyzing constituents of biological complexes (Y ates et al., 1996). Tandem mass spectrometry analysis of trypsin-digested nucleoid-enriched fractions consistently identified several peptides contained in the protein encoded by the MGM101 gene (Fig. 1, see overlines). These biochemical data suggest that Mgm101p plays a direct role in mtDNA maintenance through an association with the mitochondrial nucleoid.

We have also isolated a temperature sensitive allele of M G M101, JNY131(mgm101-2), in a genetic screen for mutants that are unable to maintain mtDNA (see M aterials and M ethods; J. Wagner, E.D. Wong, and J. Nunnari, unpublished observations). To determine the specific mutation in mgm101-2, we sequenced the mutated gene. Sequencing of the mgm101-2 allele revealed a single point mutation, resulting in a change of the conserved A sp131 residue to A sn (Fig. 1, asterisk). This residue is invariant in Mgm101p among all species and is found at the beginning of the highly conserved COOH-terminal region of the protein. Interestingly, this alteration is a relatively conservative change in Mgm101p’s primary structure. The fact that it causes the catastrophic loss of mtDNA under non-permissive conditions reinforces the importance of the COOH-terminal region in Mgm101p function.

To gain more insight into the role of MGM101 in mtDNA maintenance, we have characterized the kinetics of mtDNA loss in JNY131(mgm101-2) cells onto YPD media at fixed time points after shifting to nonpermissive temperature. A fiber exposure to nonpermissive conditions, colonies were formed at permissive temperature and respiratory competence was assessed by examining colony color. Respiratory competent cells containing an ade2 mutation accumulate a red pigment when
adenine is limiting in the medium. Thus, on YPD plates, red colonies are classified as respiratory competent and white and sectored colonies are classified as respiratory deficient. This classification was confirmed by analyzing the growth of colonies replica plated on medium containing the nonfermentable carbon source, glycerol. Under permissive conditions, no difference in the generation of respiratory incompetent colonies was observed in JNY131 (mgm101-2) and W303, indicating that mgm101-2 allele is functional under these conditions (Fig. 2 A, t 50 h and Table II). After shifting to nonpermissive temperature, however, JNY131(mgm101-2) cells began to lose respiratory competence after 4 h (two doublings). With continued exposure to nonpermissive temperature, the number of nonrespiratory colonies increased until the entire population of JNY131(mgm101-2) cells were respiratory deficient at 12 h (Fig. 2 A and Table II). In contrast, a substantially lower percentage (overall 10%) of W303 cells became respiratory deficient at nonpermissive conditions (Fig. 2 A).

The relatively rapid loss of respiratory competence in the JNY131(mgm101-2) strain under nonpermissive conditions is similar to that observed previously for mgm101-1 cells (Chen et al., 1993) and reflects the importance of Mgm101p in mtDNA maintenance.

To determine the nature of the nonrespiring phenotype in JNY131(mgm101-2), cells from nonrespiring colonies were crossed to wild-type cells lacking mtDNA. Diploids resulting from this cross were also respiratory deficient, indicating that loss of respiratory function was the result of a mutation (rho2) or complete loss (rho8) of mtDNA.

Table II. Correlation of mgm101-2 Phenotypes

| Time, hrs at 37°C | Respiratory deficient | Rho8 | Unlabeled by DAPI | Unlabeled by BrdU |
|------------------|----------------------|-----|------------------|------------------|
| 0                | 2/91 (2%)            | N/D | 0/50 (0%)        | 1/50 (2%)        |
| 2                | 4/66 (6%)            | N/D | 0/50 (0%)        | 3/50 (6%)        |
| 4                | 10/89 (11%)          | N/D | 2/50 (4%)        | 9/50 (18%)       |
| 6                | 56/110 (51%)         | N/D | 23/50 (46%)      | 12/50 (24%)      |
| 8                | 90/113 (80%)         | N/D | 0/12 (0%)        | 16/50 (32%)      |
| 10               | 137/141 (97%)        | N/D | 4/18 (22%)       | 21/50 (42%)      |
| 12               | 68/68 (100%)         | N/D | 7/15 (47%)       | 30/50 (60%)      |
| 14               | (100%)               | N/D | 13/17 (77%)      | N/D              |
| 16               | (96%)                | N/D | 23/24 (96%)      | N/D              |
To distinguish between rho− and rho+ cells, mtDNA was examined directly in cells from nonrespiratory JNY 131 (mgm101-2) colonies by staining with the DNA-specific fluorescent probe, DAA PI. Interestingly, after a nonpermissive exposure time that resulted in 100% nonrespiring colonies, only ~50% of the nonrespiring colonies consisted of rho+ cells (Fig. 2 A and Table II). The remaining half of the colonies consisted of rho− cells that still possessed mitochondrial nucleoids. However, after a more prolonged exposure of JNY 131(mgm101-2) cells to nonpermissive temperature (16 h), 100% of nonrespiring colonies consisted of cells that were rho+ (Fig. 2 A and Table II), consistent with the colony phenotype observed in the screen that identified JNY 131(mgm101-2) (see Materials and Methods). These observations indicate that a transition occurs where nonfunctional mtDNA (rho−) is formed in JNY 131(mgm101-2) cells before the complete loss of mtDNA (rho−). This phenotype is consistent with an early deficiency in JNY 131(mgm101-2) cells that causes the degradation of mtDNA or results in a block in mtDNA replication.

Because mtDNA loss is also observed in a sub-group of mutants that affect mitochondrial morphology, such as mgm1, mdm10, mdm12, and fzo1, mitochondrial structure in JNY 131(mgm101-2) cells was assessed by indirect immunofluorescence using antibodies directed against Por1p, a mitochondrial outer membrane protein (Burgess et al., 1994; Hermann et al., 1998; Sogo and Yaffe, 1994). Mitochondrial morphology was unchanged and was indistinguishable from the cortical, reticular structure of wild-type mitochondria in S. cerevisiae. Transmission of the mitochondrial organelle also was unaffected, even after exposure to nonpermissive temperature for 12 h (Fig 2 B). This indicates that loss of respiratory function and mtDNA in JNY 131(mgm101-2) is not associated with mitochondrial morphology and transmission defects. Together with the biochemical identification of Mgm101p as a component of enriched nucleoid preparations, these data suggest that Mgm101p plays a more direct role in mtDNA maintenance.

**Mgm101p Is a Nucleoid-associated Protein**

Both genetic and biochemical analyses suggest that M gm101p is localized to the mitochondrial nucleoid. To determine the subcellular localization of Mgm101p, we raised polyclonal antibodies against a MBP-M gm101p fusion protein. Western blot analysis of W 303 whole cell extracts with anti-M gm101p antibodies detected a band at 30 kD, the predicted molecular mass of M gm101p (Fig. 3 A, lane 1). In cell extracts made from CS6-1D, which contains a recessive inactive truncated copy of *MGM101*, a band at ~20 kD was detected, corresponding to the predicted M gm101p product formed by the chromosomal disruption in this strain (Fig. 1, open triangle, Fig. 3 A, lane 2).

To determine whether Mgm101p was localized to the mitochondria, we fractionated whole cell extracts from W 303(pep4) cells by differential centrifugation. Samples from each fraction were analyzed by SDS-PAGE and immunoblotted with anti-Por1p, a mitochondrial outer membrane protein, anti-A bf2p, a known mitochondrial nucleoid-associated protein, and anti-M gm101p (Fig. 3 B).

**Figure 3. M gm101p cofractionates with mitochondrial nucleoids.** (A) Cultures of strain W 303(pep4) and CS6-1D were grown in YPD. Total cell extract was prepared by lysing cells using the rapid boiling method (see Materials and Methods). A constant amount of total cell extract (0.2 OD600 units) from W 303(pep4) (lane 1) and CS6-1D (lane 2) was analyzed by SDS-PAGE and immunoblotted with anti-M gm101p antibodies. (B) Cultures of strain W 303(pep4) were grown in YPG. Cells were spheroplasted, lysed and fractionated as described in Materials and Methods. A constant amount (50 µg) of the total cell extract (T, lane 1), the 3,000 g supernate (LS, lane 2) and pellet fractions (LP, lane 3) and the 9,000 g supernate (lane 4) and pellet fractions (M, lane 5) was analyzed by SDS-PAGE and immunoblotted with anti-Por1p, anti-A bf2p, and anti-M gm101p antibodies. Mitochondria were lysed and mitochondrial nucleoids were enriched by sucrose gradient centrifugation according to Materials and Methods. A constant amount (50 µg) of the M fraction (M, lane 6), the top fraction from the sucrose gradient (TF, lane 7) and the 60/80% interface fraction containing mitochondrial nucleoids (NUC, lane 8) was analyzed as described above.

After low speed centrifugation (3000 g) of total cell lysate (Fig. 3 B, lane 1), mitochondria were recovered in both the supernatant (Fig. 3 B, lane 2) and pellet (Fig. 3 B, lane 3), as indicated by the detection of Por1p and A bf2p. The presence of mitochondria in the low speed pellet was probably due to incomplete cell lysis. A similar fractionation pattern was observed for M gm101p. When the low speed supernatant was centrifuged at 9,000 g, Por1p and A bf2p again cofractionated and were recovered in the high speed pellet (Fig. 3 B, M, lane 5), consistent with the mitochondrial localization of these proteins. M gm101p was also recovered and enriched in the pellet fraction, indicating that it is a mitochondrial protein (Fig. 3 B, M, lane 5).

To determine whether M gm101p was associated with mitochondrial nucleoids, the mitochondrial-enriched pellet (Fig. 3 B, M, lane 5) was subjected to equilibrium density centrifugation and detergent extraction (Fig. 3 B, M, lane 6). Mitochondrial nucleoids were purified further by
fractionation of the mitochondrial detergent extract on two successive sucrose density gradients (Fig. 3 B, lanes 7 and 8). Samples from the second sucrose gradient were subjected to SDS-PAGE and immunoblotted with antibodies against Por1p, Abf2p, and Mgm101p. As indicated by the fractionation behavior of the nucleoid marker Abf2p, mitochondrial nucleoids migrated to the 37.5%/60% sucrose interface, consistent with previously published results (Fig. 3 G, NUC, lane 8; Newman et al., 1996). In contrast, Por1p did not migrate into the sucrose gradient, as indicated by its presence in the gradient’s top fraction, confirming its nonnucleoid association and demonstrating the observed enrichment of nucleoid-associated proteins at the 37.5%/60% sucrose interface (Fig. 3 B, TF, lane 7). Mgm101p cofractionated with Abf2p and was quantitatively recovered in the fraction representing the 37.5%/60% interface, suggesting that it is tightly associated with nucleoids (Fig. 3 B, TF, lane 8).

To confirm the biochemical fractionation data, we localized Mgm101p by tagging the protein at the NH\textsubscript{2} and the COOH terminus with GFP. Fusion of mito-GFP to the NH\textsubscript{2} terminus of Mgm101p (mito-GFP-Mgm101p, see Materials and Methods) created a protein that when expressed from a constitutive promoter (ADH) in JNY131(mgm101-2) cells could partially complement the temperature-sensitive growth phenotype on glycerol, indicating that it was functional. In addition, DAPI staining of nonrespiring JNY131(mgm101-2) cells expressing mito-GFP-Mgm101p revealed that they were rho\textsuperscript{−}, in contrast to the rho\textsuperscript{+} phenotype that results from mgm101-2. This also indicates that mito-GFP-Mgm101p is able to partially rescue the defect in JNY131(mgm101-2) cells. Expression of a COOH-terminal fusion of GFP to Mgm101p failed to complement growth of JNY131(mgm101-2) on nonfermentable carbon sources. However, JNY131(mgm101-2) cells expressing Mgm101p-GFP were also 100% rho\textsuperscript{−} under nonpermissive conditions. This indicates that Mgm101p-GFP also retains some function. Interestingly, overexpression of either mito-GFP-Mgm101p or Mgm101p-GFP, but not wild-type Mgm101p, in W303 cells resulted in a significant increase in the formation of nonrespiratory cells (~50-fold increase), suggesting that fusion of GFP at either terminus interferes with some process required for mtDNA maintenance.

Visualization of either live wild-type or JNY131(mgm101-2) cells expressing GFP-Mgm101p (Fig. 4 A) and Mgm101p-GFP (data not shown) revealed that GFP was localized to punctate structures at the cortex of the cell that completely overlap with vitally DAPI-labeled mtDNA. Thus, in vivo, Mgm101p/GFP fusion proteins are localized to the mitochondrial nucleoids. Localization of Mgm101p in cells was also determined

![Figure 4](image-url)
using indirect immunofluorescence with anti-Mgm101p antibodies. No staining above background was observed with anti-Mgm101p in wild-type cells (data not shown). However, in cells overexpressing Mgm101p, anti-Mgm101p labeled punctate structures at the cortex of the cell (Fig. 4 B). These punctate structures were also labeled with DAPI, indicating that they contained mtDNA. As stated above, overexpression of Mgm101p in wild-type cells had no observable effect on mtDNA maintenance or inheritance as compared with wild-type. It is important to note that because we could only detect Mgm101p when overexpressed in cells, we cannot rule out the possibility that, when present at wild-type levels, Mgm101p may localize to only a subset of nucleoids. However, these data are in agreement with data obtained from both the biochemical fractionation behavior of Mgm101p and the localization of the Mgm101p/GFP fusions and indicate that Mgm101p is localized specifically and uniquely to the mitochondrial nucleoid.

**Mgm101p Binds DNA**

The COOH-terminal region of Mgm101p is highly basic suggesting that Mgm101p has the ability to bind DNA. To determine if Mgm101p has DNA binding activity, nucleoids were isolated from W303 and associated proteins were extracted with high salt containing buffer, separated from mtDNA by sucrose gradient centrifugation and subjected to DNA-cellulose chromatography. Both nucleoid proteins Abf2p and Mgm101p were quantitatively recovered in this high salt nucleoid extract, indicating that their association with this structure is salt-sensitive. Given that this may reflect their association with mtDNA, the salt concentration in the extract was lowered to 200 mM before DNA-cellulose chromatography. Under these conditions, both Abf2p and Mgm101p were soluble and sedimented as monomers as determined by sucrose gradient sedimentation (data not shown). Abf2p bound to the DNA-cellulose column and could be eluted only at high salt concentrations (1 M), consistent with its known high affinity interaction with DNA (Fig. 5, compare lanes 1 and 5). Mgm101p also bound and eluted quantitatively from the DNA cellulose at relatively high salt concentrations (500 mM, Fig. 5, lane 4). This tight association of Mgm101p with DNA-cellulose suggests that Mgm101p specifically interacts with DNA. Furthermore, both Mgm101p and Abf2p were significantly enriched, 60- and 100-fold, respectively, in DNA-cellulose column eluates. Taken together, these data suggest that the localization of Mgm101p to the nucleoid is at least in part due to direct interaction with mtDNA.

**Mgm101p Content Decreases in mgm101-2 Cells at Nonpermissive Temperature**

To gain more insight into Mgm101 function, we examined Mgm101p under both permissive and nonpermissive conditions in JNY131(mgm101-2). Total extracts from JNY131(mgm101-2) and W303 cultures grown under both conditions were analyzed by SDS-PAGE and immunoblotting with anti-Mgm101p (Fig. 6 A, compare lanes 1 and 2). This faster migrating product is unlikely to be an in vitro isolation artifact because the identical migration behavior was observed when Mgm101-D131Np was examined from denaturing immunoprecipitation of Mgm101p from S35-Met pulse-labeled cells (data not shown). The faster migration of Mgm101p observed under permissive conditions is thus likely the result of an in vivo proteolytic modification that may contribute to the phenotype observed at nonpermissive temperature (see below).

As a function of time at nonpermissive temperature, the amount of Mgm101-D131Np observed was reduced dramatically in JNY131(mgm101-2) cells (Fig. 6 B, lanes 1–5). Specifically, reduction of Mgm101-D131Np content occurred after 4 h and Mgm101-D131Np was not detectable above background after 6 h of exposure to nonpermissive temperature. In contrast, the steady state level of Mgm101p was unaffected under these conditions (data not shown).

**Figure 5.** Mgm101p binds DNA. Mitochondrial nucleoids were isolated as described in Materials and Methods. Nucleoid-associated proteins were dissociated and separated from mitochondrial DNA by incubation for 2 h in 1 M KCl followed by centrifugation where the salt extract was layered over a 30% sucrose cushion. The supernatant containing high salt extracted proteins was dialyzed into Tris-EDTA buffer, pH 7.5, 200 mM NaCl (S, lane 1) and adsorbed to a native DNA cellulose column. Proteins bound to the column were eluted with 250 mM, 500 mM, and 1 M NaCl.

**Figure 6.** Mgm101p content in mgm101-2 cells decreases under nonpermissive conditions. A. Cultures of strain W303(pep4), JNY131(mgm101-2), and CS6-1D were grown in YPD. Total cell extracts were prepared by the rapid boiling method as described in Materials and Methods. 0.33 OD600 units of W303, pep4 (lane 1), JNY131 (lane 2) and CS6-1D (lane 3) was analyzed by SDS-PAGE and immunoblotted with anti-Mgm101p antibody. B. JNY131(mgm101-2) cells were grown to log phase in YPG and cells were harvested, split and washed into YPD medium at 24°C or YPD medium prewarmed at 37°C. These two cultures of JNY131(mgm101-2) were grown in YPD at 24°C and 37°C, respectively, and samples were collected every 2 h. Total cell extracts were prepared and analyzed as described in Part A.
This observed thermolability of Mgm101p-D131Np slightly precedes and correlates with loss of respiratory function and mtDNA in JNY 131(mgm101-2) observed at nonpermissive temperature (Fig. 2 A) and therefore is likely the primary cause of these phenotypes.

Gross Alterations in Packaging, Segregation, and Partitioning of mtDNA Are Not Observed in mgm101-2

We investigated the function of Mgm101p in mtDNA maintenance by characterizing changes in nucleoid morphology in JNY 131(mgm101-2) during loss of mtDNA under nonpermissive conditions. Failure of nucleoid segregation, as observed in ΔMGT1 cells, causes the accumulation of fewer and brighter-stained nucleoid structures (Lockshon et al., 1995). Defects in mtDNA packaging, as observed in ΔABF2 cells, cause nucleoid staining to become more diffuse (Newman et al., 1996). We reasoned that such analysis would help distinguish between these possible explanations of loss of mtDNA in JNY 131(mgm101-2).

We visualized nucleoids using DAPI in fixed JNY-131(mgm101-2) and W303 cells collected at fixed time points after exposure to nonpermissive temperature. Up to 4 h of exposure to nonpermissive temperature, the majority of JNY 131(mgm101-2) cells contained mitochondrial nucleoids that were indistinguishable from nucleoids observed in W303 cells (Fig. 7 A, 4 h and Table II). This is consistent with the fact that the majority of JNY 131(mgm101-2) cells retained respiratory competence during this time (Fig. 2 A and Table II). In JNY 131(mgm101-2) cells exposed to greater than 4 h of nonpermissive temperature mitochondrial nucleoid morphology remained punctate, but the staining of the nucleoids with DAPI became less intense and fewer nucleoids were detected (Fig. 7 A, 6 h, see arrows and Table II). No staining of nucleoids with DAPI was detected after 8 h of exposure time (Table II). In contrast, no change in nucleoid intensity was observed in either JNY 131(mgm101-2) cells cultured under permissive conditions or W303 cells exposed to nonpermissive temperature, indicating that the effect was specific to the temperature-sensitive mgm101-2 allele (Fig. 7 A). This phenotype correlates with and is likely the result of the temperature-dependent decrease in Mgm101-D131Np levels observed in JNY 131(mgm101-2) at nonpermissive temperature. The lack of significant change in nucleoid morphology in JNY 131(mgm101-2) suggests that segregation and packaging of mtDNA are not significantly affected. Furthermore, nucleoid partitioning into daughter buds of JNY 131(mgm101-2) cells was also unaffected (Fig. 7 A, 6 h, see arrows).

Although segregation, packaging and partitioning of mtDNA do not seem to be grossly affected in JNY 131(mgm101-2) cells, we did observe a dramatic decrease in the intensity of DAPI nucleoid staining over time at nonpermissive temperature. Interestingly, DAPI staining of nucleoids became undetectable before complete mtDNA loss in JNY 131(mgm101-2) as assessed by examining mtDNA in cells from nonrespiratory JNY 131(mgm101-2) colonies (Fig. 2 A and Table II), consistent with our interpretation that mgm101-2 may cause mtDNA loss as a
result of a block in mtDNA replication or by causing mtDNA degradation.

**Mgm101p Is Not Required for the Maintenance of mtDNA Replication**

To determine whether loss of mtDNA in JNY131 (mgm101-2) was the result of a block in mtDNA replication, we monitored the incorporation of the thymidine analogue, 5-bromodeoxyuridine (BrdU) into mtDNA by indirect immunofluorescence as previously described (Nunnari et al., 1997). Towards this goal, we constructed JNY131 (mgm101-2) and W303 strains that contained a chromosomal copy of an exogenous thymidine kinase (JNY131TK and W303TK, respectively). Mitochondrial DNA replication was monitored in JNY131TK and W303TK cells grown under both permissive and nonpermissive conditions by pulse labeling with BrdU. Punctate staining in both JNY131TK and W303TK cells was detected at the cell cortex by indirect immunofluorescence of labeled cells with anti-BrdU antibodies (Fig. 7 B). This punctate staining was not detected in either W303 rho8 or W303 cells that had not been labeled with BrdU, indicating that it was specifically the result of incorporation of BrdU into mtDNA (Nunnari et al., 1997 and data not shown). Under permissive conditions, no significant differences in BrdU incorporation into mtDNA between JNY131TK and W303TK were observed (data not shown).

Exposure of cells to nonpermissive temperature up to 4 h did not cause a significant reduction in the number of JNY131TK cells containing BrdU-labeled mtDNA or the intensity of mtDNA labeling as compared with W303 cells (Fig. 7 B, 4 h and Table II). This nonpermissive exposure time correlates with the onset of loss of respiratory function and the decrease in nucleoid staining intensity observed with DAPI in JNY131 (mgm101-2) (Fig. 7 A and Table II). Thus, the lack of an observable change in BrdU incorporation into mtDNA in JNY131 (mgm101-2) before these phenotypes suggests that a block in mtDNA replication in JNY131TK is not the cause of these defects. Even after exposure times where Mgm101p-D131Np content was greatly reduced by Western analysis, we observed BrdU-labeled mtDNA in JNY131 (mgm101-2) cells of equal intensities (measured by pixel intensity, n = 20) to BrdU-labeled mtDNA in W303 cells (Fig. 7 B, 6 h, see arrows and Table II). However, the percentage of JNY131TK cells that contained any detectable BrdU-labeled mtDNA decreased with time (Table II). Not surprisingly, this unlabeled population correlates with the JNY131 (mgm101-2) respiratory deficient colonies that contained rho8 cells (Fig. 2 A and Table II), indicating that lack of BrdU-labeled mtDNA in this population of cells was a secondary consequence of mtDNA loss. Taken together, these data suggest that under nonpermissive conditions, loss of respiratory function and mtDNA in JNY131 is not caused by a block in ongoing mtDNA replication.

**mgm101-2 Cells Are Sensitive to UV-induced mtDNA Mutations and Hypersensitive to Both Gamma Ray and H2O2-induced mtDNA Mutations**

Given that ongoing mtDNA replication is unaffected in JNY131 (mgm101-2), we reasoned that the decrease in nucleoid staining by DAPI observed in JNY131 (mgm101-2) might be the result of mtDNA degradation. Consistent with this interpretation is that, after exposure to nonpermissive temperature, JNY131 (mgm101-2) cells recovered rho-mtDNA under permissive conditions before complete mtDNA loss (Fig. 2 A). Observations made in other systems, such as E. coli, indicate that damaged DNA is susceptible to DNA degradation. Thus, one explanation for these phenotypes is that JNY131 (mgm101-2) cells are deficient in the repair of damaged mtDNA.

To determine if Mgm101p plays a role in mtDNA repair, we examined the sensitivity of mtDNA to DNA damage in JNY131 (mgm101-2) and W303 strains by monitoring the percent formation of respiratory deficient colonies. To examine the role of Mgm101p in the repair of UV-damaged mtDNA, we compromised Mgm101 function by pretreating JNY131 (mgm101-2) cells for 2 h at nonpermissive temperature before irradiation. After UV irradiating JNY131 (mgm101-2) and W303 cells at various doses, colonies were formed at the semi-permissive temperature of 34°C. Under these conditions, in the absence of irradiation, we observed only a slight increase (2%) in the percentage of nonrespiratory colonies formed by JNY131 (mgm101-2) cells as compared with W303 cells (Fig. 8 A, time 0). Upon UV-irradiation, however, a significant dose-dependent increase in the percentage of nonrespiratory colonies was observed in JNY131 (mgm101-2) as compared with W303 (Fig. 8 A), indicating a defect in UV-induced mtDNA repair in JNY131 (mgm101-2) cells. In contrast to the sensitivity of JNY131 (mgm101-2) cells to mtDNA damage, nuclear DNA damage, as assessed by DAPI damage-induced cell death, is unaffected (Fig. 8 A), indicating that Mgm101p functions specifically in the repair of mtDNA and not nuclear DNA. Both pretreatment at nonpermissive temperature and incubation at the semi-permissive temperature of 34°C were necessary to observe significant mgm101-2-dependent sensitivity to UV-induced mtDNA damage, consistent with the fact that Mgm101-131Np is functional under permissive conditions.

To determine if the mtDNA repair defect in JNY131 (mgm101-2) cells was specific to UV-induced types of DNA damage, such as the formation of pyrimidine dimers, we examined the sensitivity of JNY131 (mgm101-2) to gamma ray–induced mtDNA damage. In contrast to UV-irradiation, no preincubation at nonpermissive temperature before irradiation. After UV irradiating JNY131 (mgm101-2) and W303 strains by monitoring the percent formation of respiratory deficient colonies. To examine the role of Mgm101p in the repair of UV-damaged mtDNA, we compromised Mgm101 function by pretreating JNY131 (mgm101-2) cells for 2 h at nonpermissive temperature before irradiation. After UV irradiating JNY131 (mgm101-2) and W303 cells at various doses, colonies were formed at the semi-permissive temperature of 34°C. Under these conditions, in the absence of irradiation, we observed only a slight increase (2%) in the percentage of nonrespiratory colonies formed by JNY131 (mgm101-2) cells as compared with W303 cells (Fig. 8 A, time 0). Upon UV-irradiation, however, a significant dose-dependent increase in the percentage of nonrespiratory colonies was observed in JNY131 (mgm101-2) as compared with W303 (Fig. 8 A), indicating a defect in UV-induced mtDNA repair in JNY131 (mgm101-2) cells. In contrast to the sensitivity of JNY131 (mgm101-2) cells to mtDNA damage, nuclear DNA damage, as assessed by DAPI damage-induced cell death, is unaffected (Fig. 8 A), indicating that Mgm101p functions specifically in the repair of mtDNA and not nuclear DNA. Both pretreatment at nonpermissive temperature and incubation at the semi-permissive temperature of 34°C were necessary to observe significant mgm101-2-dependent sensitivity to UV-induced mtDNA damage, consistent with the fact that Mgm101-131Np is functional under permissive conditions.
Most importantly, JNY131(mgm101-2) cells were more sensitive to gamma ray-induced mtDNA damage than to UV-induced mtDNA damage (compare Fig. 8, A and C). This observation suggests that MGM101 functions primarily in the repair of oxidatively damaged mtDNA selectively caused by gamma ray irradiation.

Ionizing radiation in the form of gamma rays results in a range of DNA damage, but the predominant form is oxidative damage to bases and oxidative damage to sugars, which results in single strand breaks in the DNA (Ward, 1988). The severe defect of JNY131(mgm101-2) cells in the repair of ionizing-induced mtDNA damage is consistent with a model in which MGM101 functions in the repair of oxidative mtDNA damage caused by the oxidant H$_2$O$_2$. As shown in Fig. 8, E and F, JNY131(mgm101-2) cells were hypersensitive to H$_2$O$_2$-induced mtDNA damage as compared with wild-type. As in the case of gamma ray-induced damage, no preincubation at nonpermissive temperature was required before treatment to observe the mgm101-2-dependent hypersensitivity to mtDNA damage, but after treatment colonies were formed at the semi-permissive temperature of 34°C. The hypersensitivity to both gamma ray- and H$_2$O$_2$-induced mtDNA damage in JNY131(mgm101-2) cells indicates that MGM101 function is required selectively for the repair of oxidatively damaged mtDNA. Indeed, this repair defect in JNY131(mgm101-2) cells may be the cause of the catastrophic loss of mtDNA in JNY131(mgm101-2) cells at 37°C that have not been treated with DNA damaging agents.

**Discussion**

The mitochondrial nucleoid is a complex nucleoprotein structure that functions not only to package and store the genetic material, but also to organize the metabolic and segregational activities associated with its maintenance and inheritance. It is surprising that, with a few exceptions, both the constituents and the mechanisms involved in the assembly of this structure are relatively uncharacterized to date (Cho et al., 1988; Van Dyk et al., 1992; Newman et al., 1996; Okamoto et al., 1998). The MGM101 gene was previously shown to be required for the maintenance of mtDNA, but its precise role remained unidentified (Chen et al., 1993).
MGM101 Has a Nucleoid-specific Function

Our analysis of Mgm101p and the mgm101-2 mutant strain indicates that MGMT101's function is specific to the nucleoid structure and mtDNA. Loss of MGMT101 function initially causes cells to rapidly lose respiratory function due to irreversible mutations in mtDNA (accumulation of rho− genomes) and ultimately causes the complete and catastrophic loss of mtDNA. This loss of mtDNA in mgm101-2 cells is not associated with any changes in the structure or organization of the mitochondrial organelle. This is in contrast to another group of mutants that cause mtDNA loss as a result of a primary defect in the maintenance of mitochondrial morphology (Burgess et al., 1994; Sogo and Yaffe, 1994; Hermann et al., 1998). Sing a number of independent approaches, we have also observed that Mgm101p is exclusively localized to nucleoids in cells. The nucleoid-specific localization of M gm101p is probably due to its ability to bind tightly and specifically to DNA, suggesting that Mgm101p's function in mtDNA maintenance requires this DNA binding activity. Consistent with this is the fact that deletion of a small portion of the COOH terminus composed of 23% of positively charged residues results in loss of MGM101 function (Chen et al., 1993).

MGM101 Is Not Involved in mtDNA Packaging, Segregation, or Partitioning

What is the nature of Mgm101p's nucleoid-specific function? In vivo, nucleoid-specific functions required for mtDNA maintenance include the replication, repair, assembly, segregation, and partitioning of the mitochondrial nucleoid. From our analyses of mgm101-2 cells, it is unlikely that Mgm101p is required for mtDNA packaging or segregation because in mgm101-2 cells, nucleoid morphology remained punctate and only a decrease in DAPI staining intensity of nucleoids under nonpermissive conditions was observed. It is important to note that our analysis of nucleoid morphology using DAPI might not have identified a subtle change in packaging or segregation of mtDNA. However, one would not expect a subtle change to cause the catastrophic loss of mtDNA observed in mgm101-2 cells. In addition, alterations in both the packaging and segregation of mtDNA have been previously observed using this technique. In cells lacking ABF2, a nuclear gene that encodes a mtDNA packaging protein, nucleoids lose their punctate structure and instead possess a more diffuse morphology within the organelle (Newman et al., 1996). In cells lacking GT1, a gene that encodes a cruciform-cutting endonuclease, fewer nucleoids are observed and each contains a greater amount of mtDNA, as assessed by DAPI staining intensity (Lockshon et al., 1995). Furthermore, loss of MGMT101 function did not cause a defect in the partitioning of nucleoids to daughter buds. To date, no mutants have been found that block the movement of mtDNA within the organelle. The observed decrease in nucleoid staining intensity with DAPI during loss of MGMT101 function suggests instead that MGMT101 is involved in another, more central process required for mtDNA maintenance, such as DNA metabolism.

MGM101 Is Required for the Repair of Oxidatively Damaged mtDNA

Based on our data, the most likely model is that MGMT101 functions in the repair of oxidatively damaged mtDNA. Most consistent with this interpretation, is that mtDNA in mgm101-2 cells is hypersensitive to damage caused by gamma ray irradiation and H2O2. These two agents induce DNA damage predominantly in the form of oxidative damage to bases and to sugars (Ward, 1985, 1988). In contrast, mgm101-2 cells were less sensitive to damage caused by UV treatment that induces oxidative DNA damage to a lesser extent (Ward, 1988). In addition, at the time of DNA damage by gamma ray and H2O2 treatments, Mgm101p was functional in mgm101-2 cells, indicating that Mgm101p functions in the repair of damaged mtDNA and not in the protection of mtDNA from damage. In light of the probable role of MGMT101 in mtDNA repair, the decrease in nucleoid DAPI staining intensity observed in mgm101-2 cells likely results from a transition where damaged mtDNA is degraded and ultimately lost, a phenomenon observed in other systems in response to unrepaired, damaged DNA (Michel et al., 1997). Our data also indicate that MGMT101 functions exclusively in the repair of DNA contained in the mitochondrial organelle.

It is interesting to note that mtDNA in wild-type cells is more resistant to treatments that produce oxidative damage, suggesting that yeast cells possess a greater ability to repair this type of mtDNA lesion (Fouy, 1982). Thus, the repair pathway for oxidatively damaged mtDNA might be one of the major repair pathways in the mitochondrial organelle. Indeed, given the highly oxidative environment of the mitochondrial organelle it is possible that MGMT101 functions exclusively in oxidative damage repair and that
this function is essential for mtDNA maintenance. Alternatively, it is also possible that MGM101 function is not restricted to mtDNA repair and that its requirement for mtDNA maintenance reflects its role in a more fundamental DNA metabolic activity.

What type of mtDNA repair pathway is MGM101p involved in? A S is the case for nuclear DNA, several different repair pathways function in the mitochondria. In yeast, there are at least two known repair pathways for mtDNA. One direct pathway is responsible for the repair of UV-induced pyrimidine dimers and is dependent on the PHR1 gene, which encodes a photolyase that functions in both the nucleus and the mitochondria (Sancar, 1985; Branda and Isaya, 1995). Given that MGM101 functions selectively in the repair of oxidatively damaged mtDNA, it is unlikely that it is involved in the direct PHR1-dependent photoreactivation repair pathway. Furthermore, the effects of UV-induced mtDNA damage in mgm101Δ cells were assessed in the dark and the PHR1 pathway is known to be dependent on light for activation (Sancar, 1985). A mtDNA mismatch excision repair pathway has also been characterized in yeast and is dependent on the MSH1 gene, which encodes an ATPase that is homologous to eubacterial MutS mismatch repair protein (Chi and Kolodner, 1994a,b). It is also unlikely that MGM101 is involved specifically in this pathway, because, while loss of MGM101 function results only in the accumulation of mutated copies of the mitochondrial genome, a substantially less severe phenotype (Chi and Kolodner, 1994). Thus, if MGM101 is involved in mismatch repair, it is likely to be required additionally for a process more central to mtDNA maintenance.

Modification of nucleotide bases is the predominant form of DNA damage by oxidants (Ward, 1985, 1988). Thus, it is possible that MGM101p functions in base excision and/or nucleotide excision repair pathways, the major pathways responsible for the repair of this type of DNA lesion. Mitochondrial base excision repair pathways have been characterized in other organisms, making it likely that such pathways will also exist in yeast mitochondria (Pinz and Bogenhagen, 1998). However, to date, there is no evidence to suggest that mitochondria from any organism possess nucleotide excision repair pathways. Indeed, from a number of studies it has been suggested that mitochondria lack mechanisms to repair more bulky DNA lesions (Foury, 1982). Oxidative damage to sugar moieties in DNA also result in single strand breaks (Ward, 1985, 1988). If multiple single strand breaks occur in a localized area, the accurate repair of such a lesion would require a recombinational process (Ward et al., 1985). Thus, it is also possible that MGM101p functions in such a repair pathway. Recombination-mediated repair pathways have not been described in detail in mitochondria (Foury and Kolodny, 1983; Ling et al., 1995). However, it is known that mtDNA is highly recombinogenic (Thomas and Wilkie, 1968), suggesting that damaged mtDNA can be repaired in a manner dependent on this process.

What molecular role does MGM101p perform in mitochondrial DNA repair and maintenance? Clues to the molecular function of mitochondrial proteins can often be gained by searching for similarity among their origins in eubacterial proteins. However, it is interesting to note that structural homologues of mitochondrial proteins involved in DNA metabolism are also found in such evolutionarily diverse systems as eukaryotes and phage, suggesting that mtDNA metabolism has evolved significantly from its eubacterial ancestors. MGM101p falls into a potentially novel class of evolutionarily distinct proteins. Although MGM101p shows a high degree of conservation with other putative MGM101 homologues in organisms as diverse as S. pombe, it exhibits no significant homology to any known proteins, nor does it possess any obvious enzymatic domains or motifs. Based on our findings that MGM101p is essential for both mtDNA maintenance and repair and is able to bind directly to DNA, it seems probable that MGM101p is directly involved in mtDNA metabolism.

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