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Characterization of the impact of GMP/GDP synthesis inhibition on replicative lifespan extension in yeast

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Abstract

Slowing down aging-associated accumulation of molecular damage or its prevention represents a promising therapeutic paradigm to combat aging-related disease and death. While several chemical compounds extend lifespan in model organisms, their mechanism of action is often unknown, reducing their therapeutic potential. Using a systematic approach, here we characterize the impact of the GMP pathway on yeast lifespan and elucidate GMP synthesis inhibition as a lifespan extension mechanism. We further discover that proteasome activation extends lifespan in part through the GMP pathway. GMP synthesis inhibition exerts its lifespan extension effect independently of the canonical nutrient-sensing pathway regulating lifespan. Exposing longitudinally aging yeast cells to GMP pathway inhibition in an age-dependent manner, we demonstrate that the lifespan extension is facilitated by slowing, rather than reversing, the aging process in cells. Using a GUK1 mutant with lower GMP-to-GDP conversion activity, we observe lifespan extension, suggesting that reduced GDP level by itself can also extend yeast lifespan. These findings elucidate the involvement of nucleotide metabolism in the aging process. The existence of clinically-approved GMP pathway inhibitors elicits the potential of a new class of therapeutics for aging-related disorders.

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Author Contributions

PL, EAS, and MA contributed to project planning, and design and preparation of the manuscript. EAS and PL contributed to strain construction, data collection, data analysis, and preparation of the figures. TZO contributed to strain construction, data collection, and data analysis. TZY contributed to data collection. PL, EAS, TZO, TZY, and MA interpreted the data and results, and read and approved the manuscript.

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Competing Financial Interests

The authors declare competing financial interests: EAS and MA have filed a PCT International patent application.

Keywords

aging; yeast; mycophenolic acid; GMP; GDP; proteasome; replicative lifespan

Introduction

Aging is the primary risk factor for human morbidity and mortality throughout the developed world^{1–3}. Prevention or elimination of the molecular damage that causes aging therefore represents a promising therapeutic paradigm to combat the plethora of diseases that manifest late in life. Target-based drug discovery has advanced several therapeutic candidates to human trials, including metformin⁴ and mTOR inhibitors⁵. While these treatments hold promise, additional interventions are necessary to fully minimize the health effects of advanced age.

In recent years, phenotype-based discovery approaches^{6,7} have identified many compounds capable of extending lifespan in model organisms¹. These compounds are a starting point for therapeutic development, but the mechanism of action for many of them remains unknown, reducing their therapeutic potential. One such compound is mycophenolic acid (MPA), an FDA-approved immunosuppressant that extends the replicative lifespan (RLS) of the yeast *Saccharomyces cerevisiae*⁶.

MPA's primary, but not exclusive, effect is to reduce cellular GMP pools through inhibition of inosine monophosphate dehydrogenase (IMD), the rate-limiting enzyme in *de novo* GMP synthesis^{8,9} (Fig. 1A). The *de novo* GMP synthesis involves the production of xanthosine 5'-phosphate (XMP) from inosine 5'-phosphate (IMP) via the action of IMP Dehydrogenase (IMD). Conversion of IMP to XMP is accompanied by the reduction of NAD⁺ to NADH¹⁰. This reaction is followed by the production of GMP from XMP through the action of GMP synthase GUA1. Conversion of XMP to GMP is accompanied by the conversion of ATP to AMP^{11,12}. GMP can also be synthesized via a salvage pathway in the presence of exogenous guanine^{8,13}. This reaction is catalyzed by the dimeric hypoxanthine-guanine phosphoribosyltransferase HPT1. GMP is converted to GDP by the guanylate kinase GUK1.

Yeast RLS is the most rapid aging model: most cells die within several days after birth. Due to the evolutionarily-conserved aging mechanisms, yeast replicative lifespan is closely related to healthspan and lifespan in humans. Several interventions^{14,15} that extend yeast RLS promise to improve general health in humans. For example, a rapamycin analog improves immune response in the elderly¹⁶. Moreover, dietary restriction reduces biomarkers for diabetes, cardiovascular disease, and cancer¹⁷.

RLS is quantified by the number of mitotic division events experienced by a newborn yeast cell until its death; it is measured by counting the daughter cells budded off from the tracked/aging mother cell. Differently from RLS, a second commonly-used lifespan metric for yeast, chronological life span (CLS), measures the chronological time a cell can remain viable in a post-replicative state.

Aging and lifespan are multifactorial processes whose comprehensive quantitative characterization at the whole-genome level is still missing. For example, the full range of pathways and how their dynamic interactions regulate lifespan are not definitely known, yet. Despite this reality, three major lifespan-extension pathways (nutrient sensing pathway, proteasome pathway and sirtuin pathway) have been previously identified¹⁸ for *S. cerevisiae*. These pathways may not be fully orthogonal to each other, as the sirtuin pathway has been thought¹⁹ to act downstream of the nutrient-sensing pathway.

Excluding our previous work⁶ which elucidated MPA's lifespan-extending effect on yeast, no previous study directly and empirically implicated MPA as a longevity-enhancing chemical compound. Therefore, MPA's action mechanism has been unknown. Here, we seek to understand the mechanism behind MPA's ability to extend replicative lifespan. We find that guanine reverses MPA's pro-longevity effect, suggesting that reduction in cellular GMP, or its downstream metabolites, is responsible for the lifespan extension. We then investigate if MPA's RLS extension mechanism overlaps with two of the previously-described aging pathways (nutrient sensing and proteasome pathways), elucidating that it partially overlaps with the effect exerted by activated proteasome. Using a GUK1 mutant with lower guanylate kinase activity, we again observe lifespan extension, suggesting that reduced GDP level by itself contributes to the lifespan extension in a major way.

Results

Inhibition of GMP synthesis extends yeast RLS

We hypothesized that the reduction in cellular GMP pools was responsible for MPA's pro-longevity effect. Using our microfluidic "yeast replicator" platform²⁰, we quantified the number of daughter cells produced by mother cells individually tracked throughout their replicative lifespan. These single-cell RLS measurements were made in minimal media environments containing MPA (at standard 10 μ M) with or without supplemental guanine.

While the average lifespan of the cells grown in the guanine-supplemented environment was slightly reduced compared to DMSO control, the difference was not statistically significant (Table 1). Importantly, we found that the longevity effect of MPA was prevented by the supplementation of exogenous guanine (Fig. 1B). These results confirm that MPA extends replicative lifespan in *S. cerevisiae* through inhibition of GMP synthesis; however, further experimental characterizations are needed to determine if it is decreased levels of GMP and/or its downstream metabolites that cause the lifespan extension. In this study, for the determination of statistical significance, we used the log-rank test which is a hypothesis test to compare two survival distributions. It is important to note that the log-rank test takes into account the entire survival distribution when assessing significance, instead of simply comparing means from the two distributions.

Inhibition of GMP synthesis extends lifespan independent of the nutrient sensing pathway

We next asked how lifespan extension via GMP-synthesis inhibition could be placed among the known lifespan pathways. To answer this question, we used a systematic approach to categorize longevity interventions to genetic regulators of lifespan. A longevity intervention

can act either within or independent from any known longevity pathway. If within a longevity pathway, the intervention must act upon, upstream from, or downstream of a given pathway component. We aimed to conclusively identify the placement of the lifespan extension mechanism through GMP-synthesis inhibition relative to the known genetic lifespan pathways using the two-step Longevity Placement Test (LPT) (Fig. 2A, Fig. S1, Supplementary Information). In Step 1, we ask if the longevity intervention extends lifespan in a strain lacking a critical lifespan pathway component, the probe gene. In Step 2, we ask whether an epistatic agent that prevents lifespan extension from the longevity intervention can also prevent lifespan extension conferred by modulation of the probe gene. By combining this information, one can definitively classify the relationship of a longevity intervention to a known lifespan regulation pathway.

We used the LPT to determine the relationship of GMP synthesis inhibition to two major lifespan-extension pathways known for *S. cerevisiae*^{18,19}. For this purpose, MPA was used as the longevity intervention, and guanine was used as the epistatic agent. The first pathway we tested was the nutrient sensing pathway²¹, which encompasses dietary restriction¹⁵ and target of rapamycin (TOR) inhibition²². As the LPT probe genes, we chose TOR1 and HXK2, whose individual deletion is known to extend yeast lifespan. TOR1 is a protein kinase subunit of the TORC1 complex that controls cell growth in response to nutrient availability. HXK2, on the other hand, is a hexokinase whose deletion provides a genetic mimicry of nutrient limitation because it phosphorylates intracellular glucose as part of glucose metabolism. MPA further extended lifespan in the long-lived *TOR1* ($p=0.000634$) and *HXK2* ($p=0.0000269$) strains, while guanine did not suppress gene-deletion-caused lifespan extension in these strains (Fig. 2B–E, Table 1). These results indicate that inhibition of GMP synthesis exerts its longevity effect independent of the nutrient-sensing pathway.

Proteasome activation extends lifespan in part through the GMP pathway

Next, we asked whether inhibition of GMP synthesis extended RLS through the second major lifespan-extension pathway, the proteasome pathway. UBR2 is a ubiquitin ligase subunit, and its deletion activates the proteasome by stabilizing RPN4, a transcription factor that promotes expression of proteasome subunits²³. Activation of the proteasome via UBR2 deletion extends RLS independent of the nutrient sensing pathway²⁴. Therefore, we chose UBR2 as the LPT probe gene. MPA led to extension ($p = 0.012$) of yeast RLS in a *UBR2* strain (Fig. 3A, Table 1), compared to an extension ($p = 1.43 \times 10^{-9}$) in wild-type cells (Fig. 1B, Table 1). On the other hand, guanine supplementation, at a saturating concentration, partially but significantly ($p = 0.015$) suppressed lifespan extension from UBR2 deletion (Fig. 3B, Table 1). These results indicate that UBR2 deletion activates, if not exclusively, the same lifespan extension mechanism as MPA without saturating the target; in other words, UBR2 deletion extends lifespan in part through the GMP pathway. A role for the GMP pathway in the phenotypic effects of UBR2 deletion is reinforced by the observation that IMD proteins are among the most highly upregulated proteins in a *UBR2* strain²⁴. We therefore conclude that, downstream of UBR2, activated proteasome exerts its lifespan-extension effect partially through the GMP pathway, as supplementing *UBR2* strain's growth media with excess guanine significantly but only partially reverses the lifespan extension.

Since multiple steps exist between UBR2 deletion and proteasome activation, it was possible that MPA acted downstream of UBR2, but upstream of proteasome activation. We sought to differentiate these possibilities. We measured proteasome activity in wild-type cells in the presence and absence of MPA (Fig. 3C, Fig. S2), and found that MPA does not activate the proteasome. Furthermore, guanine does not alter proteasome activity in *UBR2* cells (Fig. 3C). This demonstrates that MPA regulates lifespan without modulating the proteasome. We further demonstrated that MPA extends RLS in a *PRE9* strain (Fig. 3D, Table 1), in which proteasome activation is not able to increase RLS due to the absence of the proteasome subunit PRE9². These results indicate that MPA acts to extend lifespan downstream of proteasome activation. Since deletion of UBR2 extends lifespan exclusively²⁴ through proteasome activation and excess guanine supplementation only partially reverses the lifespan-extension effect of UBR2 deletion, we conclude that proteasome activation extends lifespan in part through the GMP pathway (Fig. 3E). While the other mechanism(s) through which activated proteasome extends lifespan is still unknown, a previous hypothesis^{2,3} suggests that proteasome activation leads to lifespan extension via more efficient removal of misfolded and damaged proteins. Future studies will show which proteins are more prone to aging-associated damage and how their removal through activated proteasome impacts cellular lifespan.

MPA slows, rather than reverses, the aging process

Interventions that extend lifespan may mechanistically act to slow the accumulation of aging factors or aging-related damage, reverse aging-related damage, or suppress its effects. We sought to determine through which mechanism MPA extended the lifespan. For this purpose, we assessed the RLS of yeast cells treated with MPA only for the first 24 hours of a Replicator experiment (Fig. 4A), only after the first 24 hours of a Replicator experiment (Fig. 4B), or with a 6-hour pulse treatment between the 24th and 30th hour of the experiment (Fig. 4C). We found that MPA treatment for only part of the lifespan, either early or late, resulted in reduced lifespan extension compared to whole-lifespan treatment (Fig. 4A–B, Table 1). Furthermore, pulse treatment resulted in little to no lifespan extension (Fig. 4C, Table 1). These results indicate that MPA slows, rather than reverses, the accumulation of aging factors in cells.

Throughout this study, we used MPA at the 10 μ M concentration because of the “standard” nature of this concentration based on many compound-screening assays. However, a dose-response characterization we performed showed that 10 μ M was indeed the most efficient MPA concentration in terms of its RLS-extending effect (Fig. S3).

Measuring the impact of lowered guanylate kinase activity on replicative lifespan

Is it GMP or one of its downstream metabolites whose reduced production under MPA treatment causes lifespan extension? While we could not directly measure intracellular GMP, GDP, and GTP levels in aging cells due to lack of experimental probes or their integration to our microfluidic platform, we sought to answer this question by focusing on the synthesis of GDP from GMP. Yeast GUK1 gene encodes a guanylate kinase which converts GMP to GDP. Knocking out GUK1 makes cells inviable, however a previous study²⁵ produced a viable variant of the GUK1 gene with lower guanylate kinase activity.

Using its own upstream and downstream intergenic regions, we first cloned one copy of the mutated GUK1 gene into our strain background, followed by the deletion of the endogenous GUK1 gene from yeast genome. Sequencing the open reading frame together with the upstream/downstream intergenic regions showed us the presence of a single G-to-A mutation at the 34th base from the start of the gene, leading to a Glycine to Serine amino acid change.

Next, we measured the replicative lifespan of this *guk1-1* strain in the absence of MPA in our microfluidic platform and observed an average lifespan of 28.3 generations (Fig. 5A, Table 1), corresponding to a significant ($p = 2.83 \times 10^{-5}$) extension of yeast lifespan compared to wild-type. The lifespan extension facilitated by the *guk1-1* mutant was very close ($p = 0.1155$) to the extension caused by the MPA treatment (Fig. 5B, Table 1).

While these results strongly suggest a major role for reduced GDP (and/or its downstream metabolites) to cause lifespan extension, they are not contradictory to the earlier results we observed from yeast cells treated with MPA, as inhibition of GMP synthesis by MPA would already be expected to indirectly lead to a reduction in GDP levels. Moreover, reduced GMP, by itself, may still have a partial effect on lifespan extension. Enabled by the capability to make single-cell level measurements of intracellular GMP and GDP concentrations in aging cells, future studies will show the relative contributions of GMP and GDP reduction on the lifespan extension through the GMP pathway.

Discussion

In this study, we showed how MPA facilitates lifespan extension in yeast. Since MPA has a well-characterized role on the GMP pathway, a direct mechanism for its lifespan extension effect emerged through GMP synthesis inhibition. This was further confirmed by the observation that adding excess guanine into growth media reverses the RLS extension effect of MPA. Here we also showed that activated proteasome extends lifespan in part through the GMP pathway. Furthermore, we found that MPA exerted its lifespan-extending effect by slowing, rather than reversing, the aging process. As part of our characterizations to locate the pathway through which MPA impacts lifespan, we found that the nutrient-sensing pathway was not involved in lifespan extension from the use of MPA. Performing experiments with a GUK1 mutant with lower guanylate kinase activity, we observed lifespan extension, suggesting that reduced GDP levels can extend yeast lifespan in the absence of MPA. Taken together, our work elucidates the involvement of nucleotide metabolism in the aging process.

MPA is an FDA-approved immunosuppressant that is used in the clinic to prevent graft rejection after organ transplant operations. MPA inhibits the production of T-cell cytokines²⁶. This inhibition is facilitated by preventing T-cells to progress to the S-phase of the cell cycle²⁶. It is also known²⁷ that MPA inhibits DNA replication by inducing GTP depletion, which suppresses the synthesis of RNA-primed DNA intermediates. Through which potential mechanism could the inhibition of DNA replication extend lifespan? In general terms, lifespan extension for a cell can be realized by either slowing the accumulation of molecular damage or through its elimination at old age. Our results indicate

that MPA exerts its RLS-extending effect through slowing down the aging process. Slower aging, in turn, could be realized by more efficient continuous repair^{28,29} alleviating accumulation of damage. We therefore propose that MPA extends yeast replicative lifespan via more efficient repair of molecular damage. Among the potential mechanisms directly or indirectly leading to repair pathway activation is stress caused by MPA-induced inhibition of DNA replication. In support of this potential mechanism, genes involved in DNA-damage repair have been shown^{30,31} to be induced in response to environmental stress mediated by Msn2/4^{32,33}. As another layer of support, a recent study showed that a transient pharmacological endoplasmic reticulum stress, imposed early in development on *Caenorhabditis elegans*, enhanced proteostasis and increased adult lifespan³⁴. Mimicking the transient unfolded protein response up-regulation using the glycosylation inhibitor tunicamycin, the authors saw increased lifespan³⁴. MPA is also known to induce inhibition of glycosylation of proteins through depletion of guanosine nucleotides³⁵.

Our work uncovers the possibility of a new class of therapeutics targeting aging-related disease. Future studies will determine the clinical viability of GMP/GDP synthesis inhibitors for delaying the onset of or for the prevention of aging-related disease. The finding that mycophenolic acid extends lifespan in a *TOR1* genetic background promotes the idea of combination therapy with the TOR inhibitors currently in clinical trials. Characterization of new targets through which cellular aging³⁶⁻⁴³ can be regulated represents an important milestone towards the ultimate goal of fully understanding aging and lifespan determinants at single-cell resolution.

Materials And Methods

Yeast strains and media conditions used for determining lifespan characteristics and proteasome activity

All experiments in *S. cerevisiae* were conducted in a BY4741 haploid strain background (TransOMIC TKY0002). Haploid deletion strains were purchased from the Yeast Knockout Collection maintained by GE Dharmacon, carrying the KanMX selection marker for the deletions. Complete supplement mixture (CSM 2% glucose) media was used for all experiments, with cells maintained in aerobic conditions at 30 °C in 50 mL conical tubes (Becton Dickinson F2070) or sterile 250 mL Erlenmeyer flasks. Cultures were performed in an Innova-42 shaker (New Brunswick Scientific) set to 225 rpm.

To construct the *guk1-1* mutant strain in the BY4741 genetic background, we requested from Dr. Bertrand Daignan-Fornier the strain “129” (a strain in the PLY121 genetic background) published in a previous study²⁵. This strain contained the *guk1-1* allele²⁵. We PCR-amplified and transformed (using *spHIS5* as the selection marker) into the *ho* locus of the BY4741 strain one copy of the *guk1-1* ORF (564bp) and its entire intergenic regions (upstream and downstream of *guk1-1* ORF, 1258bp of total length). *spHIS5* ORF was flanked by *TEF1* promoter and terminator from *Ashbya gossypii*. Next, in another transformation we performed, the endogenous *GUK1* ORF was deleted using KANMX4 selection marker whose ORF was flanked by *TEF1* promoter and terminator. PCR checks and local sequencing performed on the isogenic colony collected after the second yeast transformation showed us that the endogenous *GUK1* ORF was deleted and the *guk1-1*

allele was integrated in the *ho* locus. DNA sequencing also revealed that there was a single G-to-A mutation at the 34th base position of the *guk1-1* ORF, leading to a Glycine-to-Serine amino acid change. There were no mutations (compared to the BY4741 genetic background) on the upstream and downstream intergenic regions (PCR-amplified from the strain “129”) around the *guk1-1* ORF.

Replicative lifespan measurements for haploid yeast in the Replicator

RLS measurements were performed in the Replicator device for haploids, as described in our previous publication²⁰. Briefly, cells were grown overnight in CSM 2% glucose for 24 hours to an OD₆₀₀ of 0.1, and then loaded to the microfluidic Replicator device. Media was then flowed through the device, containing DMSO (American Bio AB00435), 10 μM mycophenolic acid (Sigma M5255), and/or guanine (Sigma G11950) at a saturating concentration of 20 mg/L. An automated microscope was used to collect time-lapse images of individual mother cells every 10 min for 120 hours, and RLS was determined by counting the number of daughters produced before death. RLS was scored manually by observing the time-lapse image series produced in a Replicator experiment. Cells were included for measurement if they entered the trap prior to the 10th hour of the experiment. Passage into each generation was denoted by the appearance of a bud.

Protein extraction and proteasome assay

For protein extraction, 50 mL of cells were grown for 18 hours to an OD₆₀₀ of approximately 0.8, then transferred to a 50 mL conical tube and centrifuged at 4255 xg for 5 minutes at room temperature. The supernatant was discarded, and the cells were resuspended in 150 μL of cold lysis buffer (50 mM Tris-HCl, pH 7.5, 0.5 mM EDTA, 5 mM MgCl₂, with cOmplete™ ULTRA mini protease inhibitor tablets, EDTA free) and transferred to a 1.5 mL tube. A ¼ volume of 500–750 μm glass beads (Acros Organics 397641000) was added to each tube. For 10 rounds, the tubes were chilled in ice water for 1 minute, then vortexed at maximum speed for 30 seconds to physically rupture the cells, and returned to the ice water. Samples were then spun for 3 minutes at 2500 xg and 4 °C, and the supernatant was transferred to a fresh tube. The solution was further clarified by centrifugation at 8000 xg for 10 minutes at 4 °C, and the supernatant was transferred to a fresh tube. Protein concentration was measured using a Nanodrop measuring the absorbance of the sample at 280 nm.

The proteasome assay used was previously described²⁴. The assay was performed in a 96-well clear-bottom plate (Costar 3603) with 50 μg of total protein in 200 μL of lysis buffer. The fluorogenic proteasome substrate Suc-LLVY-AMC (Bachem I-1395) was added to a final concentration of 100 μM. Fluorescence intensity with an excitation wavelength of 380 nm and an emission wavelength of 460 nm was recorded at 5-minute intervals using a Neo2 plate reader (BioTek) set to mix constantly and maintain 30 °C. Negative control reactions were performed in the presence of 50 μM MG132 (Sigma 474787), a proteasome inhibitor.

Statistical methods for lifespan characteristics and proteasome activity

Differences in lifespan characteristics were assessed through Log-Rank test using MATLAB with a cut-off value of P = 0.05. A summary of the results is provided in Table 1. The script

for Log-Rank test was downloaded from MATLAB central: <http://www.mathworks.com/matlabcentral/fileexchange/20388-logrank>. Differences in proteasome activity were assessed using the unpaired, two-tailed, parametric t-test function.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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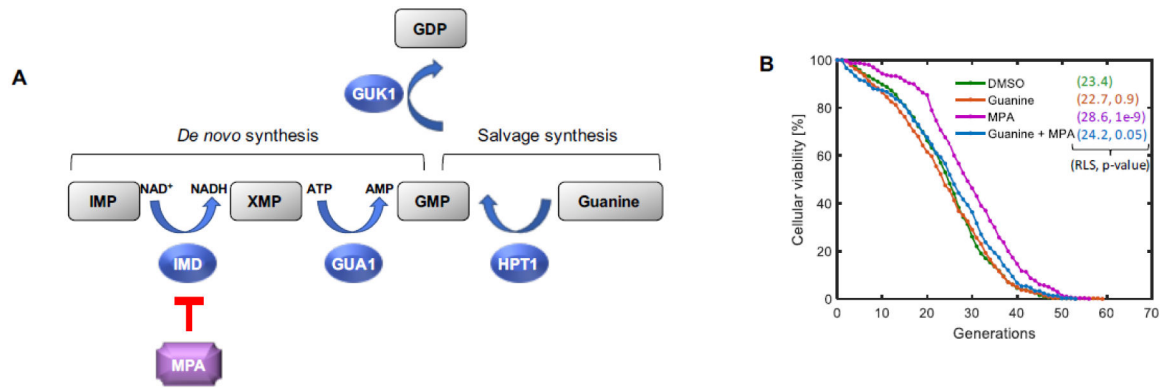


Figure 1. Inhibition of GMP synthesis extends yeast RLS. A.

Simplified schematic representation of GMP synthesis pathways in *S. cerevisiae*. MPA limits *de novo* GMP synthesis via inhibition of IMD genes. Conversion of IMP to XMP is accompanied by the reduction of NAD^+ to NADH. Conversion of XMP to GMP is accompanied by the conversion of ATP to AMP. GMP is also synthesized via the salvage pathway in the presence of exogenous guanine. GMP is converted to GDP by the guanylate kinase GUK1. **B.** Lifespan curves for wild-type, haploid yeast (BY4741) in the presence or absence of MPA and guanine. $N = 400$ cells each for the DMSO and guanine conditions, pooled from four independent experiments of 100 cells each; $N = 300$ cells each for the MPA and guanine&MPA conditions, pooled from three independent experiments of 100 cells each. Average RLS values associated with all growth conditions are reported in colored numbers in parentheses. Next to each average RLS value reported, p-value obtained from log-rank test is reported; each lifespan curve is compared to the curve obtained in the DMSO condition.

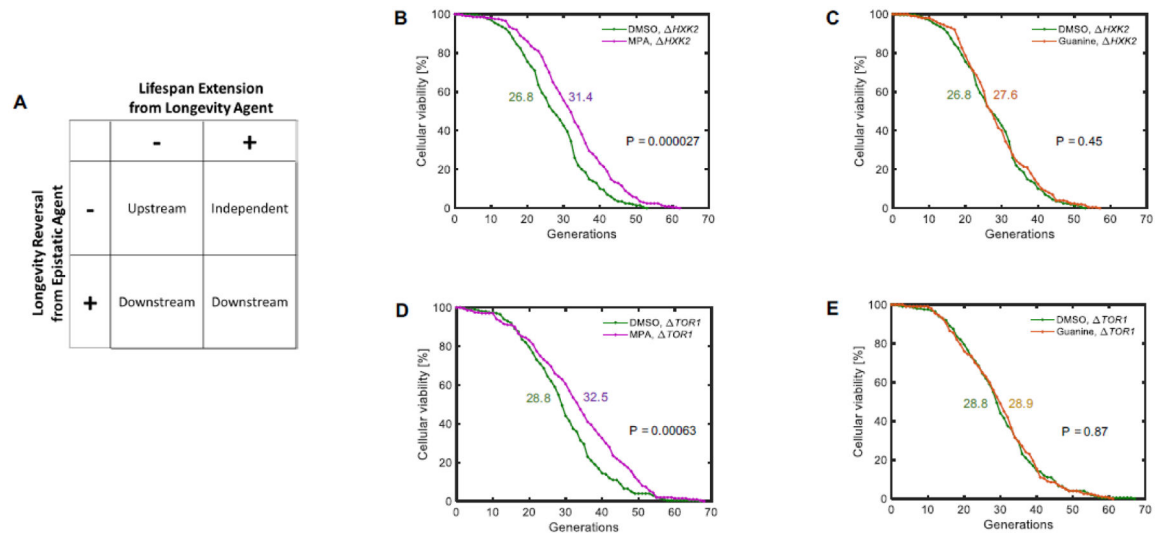


Figure 2. Inhibition of GMP synthesis extends lifespan independent of the nutrient sensing pathway.

A. Schematic representation of the LPT test and its interpretation. **B-E.** Lifespan curves corresponding to Step 1 (**B, D**) or Step 2 (**C, E**) of the LPT test for the nutrient sensing pathway, including a dietary restriction mimetic (**B, C**) and TOR inhibition (**D, E**). $N = 200$ cells for each condition, pooled from two or more independent experiments. Colored numbers next to the lifespan curves indicate the average replicative lifespan values corresponding to the lifespan curve matching its color.

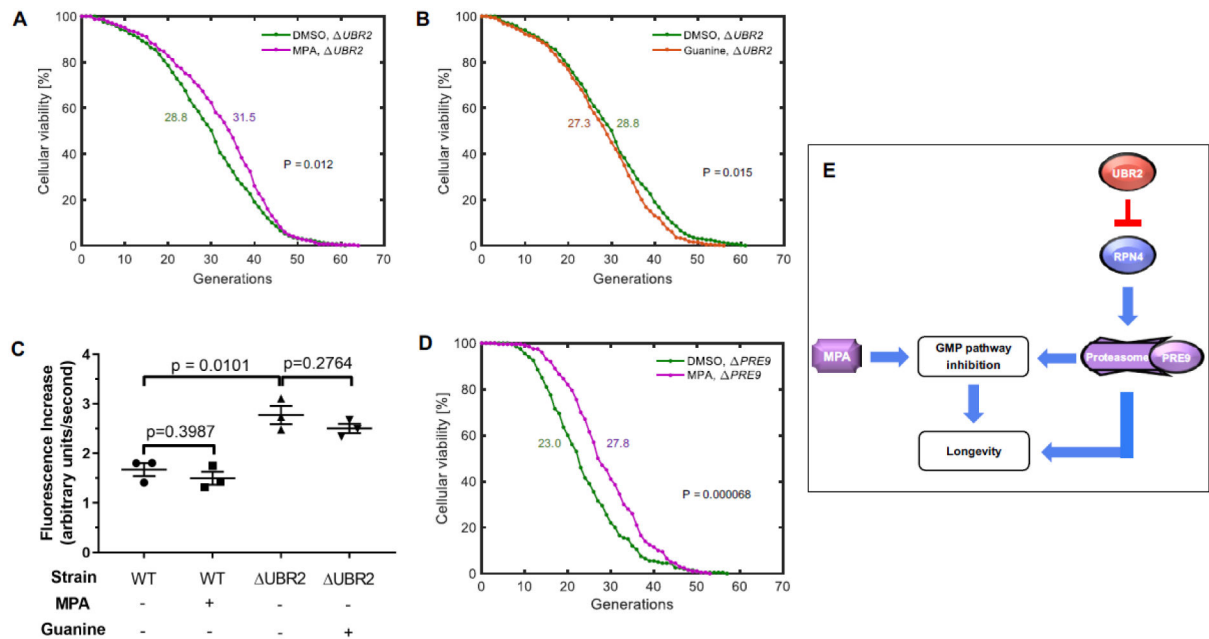


Figure 3. Proteasome activation extends lifespan in part through the GMP pathway.

A-B. Lifespan curves corresponding to Step 1 (**A**) and Step 2 (**B**) of the LPT test for the proteasome pathway of lifespan extension. $N = 300$ cells for the MPA condition, pooled from three independent experiments of 100 cells each; $N = 400$ cells for the DMSO condition, pooled from four independent experiments of 100 cells each; $N = 400$ cells for the guanine condition, pooled from four independent experiments of 100 cells each. **C.** Proteasome activity for wild-type (BY4741) cells, or $\Delta UBR2$ cells, in the presence or absence of MPA or guanine. $N = 3$ biological replicates for each condition. Errors bars are standard error of the mean. **D.** Lifespan curve for a $\Delta PRE9$ strain in the presence or absence of MPA. $N = 200$ cells for each condition, pooled from two independent experiments of 100 cells each. In (**A**, **B**, **D**), colored numbers next to the lifespan curves indicate the average replicative lifespan values corresponding to the lifespan curve matching its color. **E.** A schematic model presenting the relationship of longevity interactions described in this paper. The actions of MPA converge on the actions of the proteasome at the level of GMP pathway inhibition. Inhibition of GMP pathway extends lifespan. Proteasome activation also extends lifespan independently of GMP pathway inhibition.

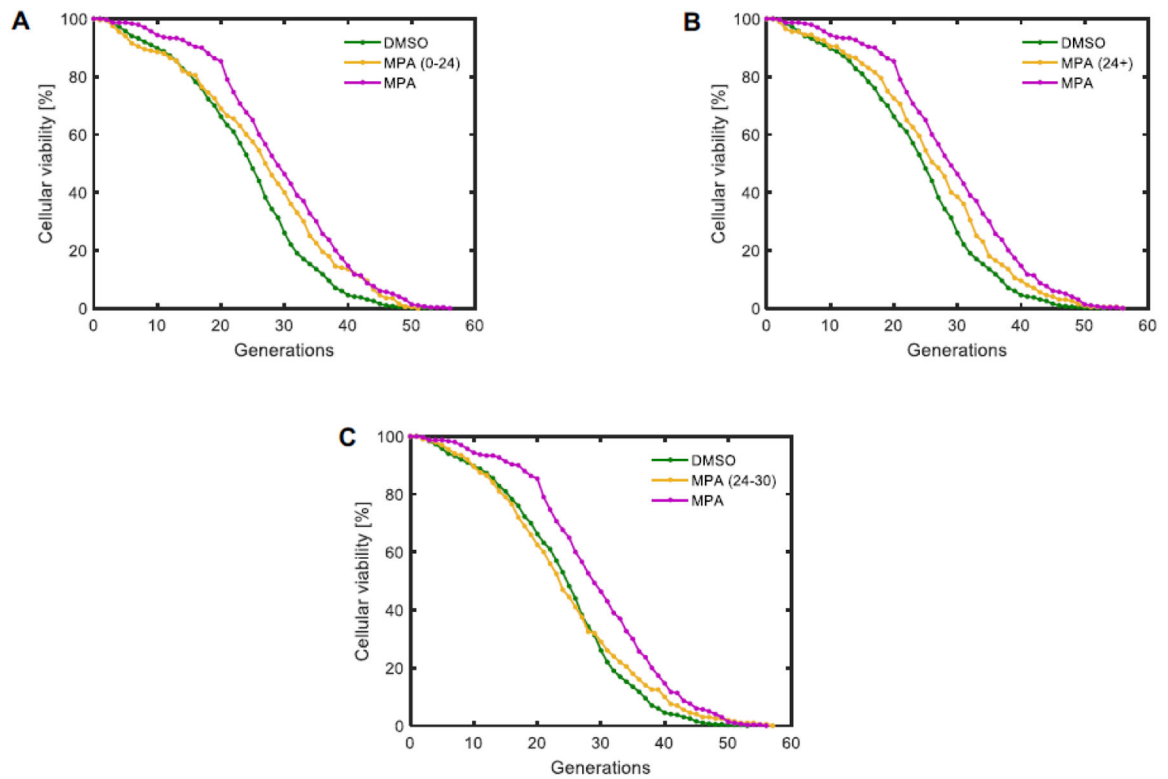


Figure 4. MPA slows, rather than reverses, the aging process.

A-C. Replicative lifespan curves for *S. cerevisiae* treated with 10 μ M MPA only during the first 24 hours of a Replicator experiment (**A**), only after the first 24 hours of a Replicator experiment (**B**), or between the 24th and 30th hour of a Replicator experiment (**C**). N = 400 cells for the DMSO condition, pooled from four independent experiments of 100 cells each; N = 300 cells for the full-time MPA condition, pooled from three independent experiments of 100 cells each; N = 200 cells for the partial-time MPA conditions, pooled from two independent experiments of 100 cells each.

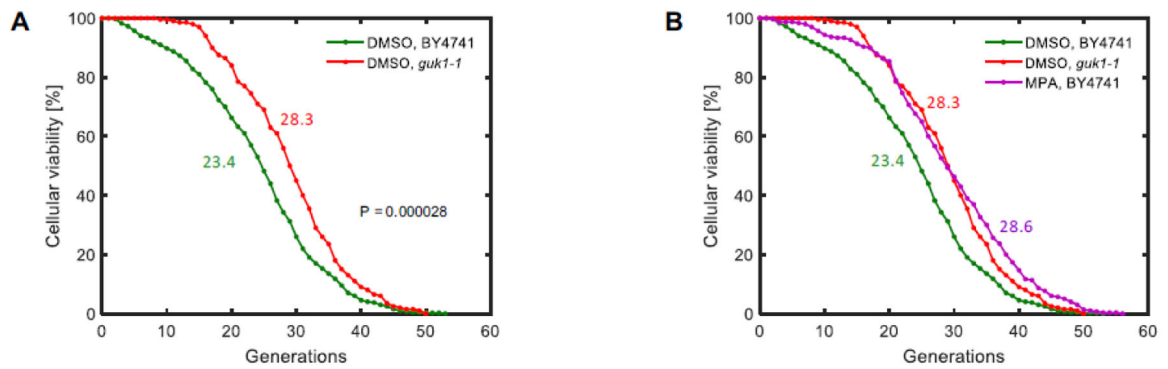


Figure 5. Impact of lowered guanylate kinase activity on replicative lifespan.

A. Lifespan curves for haploid yeast (BY4741) in wild-type or mutated *guk1-1* genetic background. **B.** Lifespan curves for haploid yeast (BY4741) in wild-type or mutated *guk1-1* genetic background, together with wild-type strain under MPA treatment. $N = 400$ cells for the wild-type DMSO condition, pooled from four independent experiments of 100 cells each; $N = 200$ cells for the mutated *guk1-1* strain in DMSO condition, pooled from two independent experiments of 100 cells each; $N = 300$ cells for wild-type strain under MPA treatment, pooled from three independent experiments of 100 cells each. Colored numbers next to the lifespan curves indicate the average replicative lifespan values corresponding to the lifespan curve matching its color.

Table 1.

Summary of lifespans and log-rank test results.

Condition 1	Condition 1 Mean Lifespan (Generations)	Condition 2	Condition 2 Mean Lifespan (Generations)	Percent Change	p-value obtained from log-rank test
BY4741-DMSO	23.39	BY4741-Guanine	22.7	-2.949978623	0.9034
BY4741-DMSO	23.39	BY4741-MPA	28.553	22.0735357	1.43E-09
BY4741-DMSO	23.39	BY4741-MPA-Guanine	24.183	3.390337751	0.0463
BY4741-DMSO	23.39	HXK2-DMSO	26.845	14.77126977	6.60E-04
HXK2-DMSO	26.845	HXK2-Guanine	27.6	2.812441795	0.45
HXK2-DMSO	26.845	HXK2-MPA	31.35	16.78152356	0.0000269
BY4741-MPA	28.553	HXK2-MPA	31.35	9.795818303	0.0039
BY4741-DMSO	23.39	TOR1-DMSO	28.805	23.1509192	4.27E-08
TOR1-DMSO	28.805	TOR1-Guanine	28.935	0.451310536	0.87
TOR1-DMSO	28.805	TOR1-MPA	32.525	12.91442458	0.000634
BY4741-MPA	28.553	TOR1-MPA	32.525	13.91097258	2.40E-06
BY4741-DMSO	23.39	UBR2-DMSO	28.8	23.12954254	2.32E-13
UBR2-DMSO	28.8	UBR2-Guanine	27.28	-5.277777778	0.0147
UBR2-DMSO	28.8	UBR2-MPA	31.5	9.375	0.0122
BY4741-MPA	28.553	UBR2-MPA	31.5	10.32115715	2.35E-04
BY4741-Guanine	22.7	UBR2-Guanine	27.28	20.17621145	7.99E-08
BY4741-DMSO	23.39	PRE9-DMSO	23.005	-1.646002565	0.5019
PRE9-DMSO	23.005	PRE9-MPA	27.84	21.01717018	0.0000684
BY4741-MPA	28.553	PRE9-MPA	27.84	-2.497110636	0.1524
BY4741-DMSO	23.39	BY4741-MPA(0-24)	24.37	4.189824711	0.0013
BY4741-DMSO	23.39	BY4741-MPA(24+)	25.55	9.23471569	0.0066
BY4741-DMSO	23.39	BY4741-MPA(24-30)	23.755	1.560495938	0.2429
BY4741-MPA	28.553	BY4741-MPA(0-24)	24.37	-14.64994922	0.0436
BY4741-MPA	28.553	BY4741-MPA(24+)	25.55	-10.51728365	0.0122
BY4741-MPA	28.553	BY4741-MPA(24-30)	23.755	-16.80383848	4.83E-04
BY4741-MPA(0-24)	24.37	BY4741-MPA(24-30)	23.755	-2.523594584	0.15
BY4741-MPA(0-24)	24.37	BY4741-MPA(24+)	25.55	4.842018876	0.66
BY4741-MPA(24+)	25.55	BY4741-MPA(24-30)	23.755	-7.025440313	0.279
BY4741-DMSO	23.39	guk1-1-DMSO	28.3	20.99187687	2.83E-05
BY4741-MPA	28.553	guk1-1-DMSO	28.3	-0.886071516	0.1155