Substrate-Specific Effects of Natural Genetic Variation on Proteasome Activity

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Abstract

The bulk of targeted cellular protein degradation is performed by the proteasome, a multi-subunit complex consisting of the 19S regulatory particle, which binds, unfolds, and translocates substrate proteins, and the 20S core particle, which degrades them. Protein homeostasis requires precise, dynamic control of proteasome activity. To what extent genetic variation creates differences in proteasome activity is almost entirely unknown. Using the ubiquitin-independent degrons of the ornithine decarboxylase and Rpn4 proteins, we developed reporters that provide high-throughput, quantitative measurements of proteasome activity *in vivo* in genetically diverse cell populations. We used these reporters to characterize the genetic basis of variation in proteasome activity in the yeast *Saccharomyces cerevisiae*. We found that proteasome activity is a complex, polygenic trait, shaped by variation throughout the genome. Genetic influences on proteasome activity were predominantly substrate-specific, suggesting that they primarily affect the function or activity of the 19S regulatory particle. Our results demonstrate that individual genetic differences create heritable variation in proteasome activity and suggest that genetic effects on proteasomal protein degradation may be an important source of variation in cellular and organismal traits.

Introduction

Protein homeostasis requires continuous degradative protein turnover. The vast majority of targeted protein degradation occurs through the concerted actions of the ubiquitin system and the proteasome (collectively, the ubiquitin-proteasome system [UPS]). The ubiquitin system is a series of enzymes that mark proteins for degradation by covalent attachment of the small protein ubiquitin. The proteasome is a multi-subunit complex that binds and degrades ubiquitinated cellular proteins^{1–5}. UPS protein degradation influences virtually all aspects of cellular physiology^{2,3,6}, necessitating precise, dynamic control of proteasome activity. Accordingly, a diverse array of regulatory mechanisms control proteasome activity by altering the abundance, function, or composition of the cell’s complement of proteasomes⁷. Imbalances between proteasome activity and the proteolytic needs of the cell adversely impact cellular viability and are implicated in numerous human diseases, including metabolic, immune, and neurodegenerative disorders^{8–13}.

We recently established individual genetic differences as an important source of variation in UPS protein degradation¹⁴. Quantitative trait locus (QTL) mapping revealed widespread genetic influences on the activity of the ubiquitin system. DNA variants in ubiquitin system genes cause altered processing, recognition, and ubiquitination of UPS substrates¹⁴. However, whether genetic

influences on the activity of the proteasome are equally numerous is unknown.

The proteasome consists of two functionally distinct modules. The 19S regulatory particle selects, unfolds, and translocates substrates, which are then processively degraded by the 20S core particle. DNA sequence differences may differentially affect the 19S regulatory particle and the 20S core particle. Genetic influences on the 19S regulatory particle could manifest as substrate-specific changes in protein degradation. In contrast, genetic influences on 20S core particle activity should alter turnover similarly across substrates. To date, however, there have been no efforts to characterize genetic influences on proteasome activity, in part, because of a lack of suitable methods. Comprehensively characterizing genetic influences on proteasome activity with high statistical power requires assays that can provide high-throughput, quantitative, *in vivo* measurements of proteasomal protein degradation. Several high-throughput reporter systems have been developed to measure UPS activity^{14–17}. However, these systems are responsive to changes in both ubiquitin system and proteasome activity.

A handful of endogenous cellular proteins are proteasomally degraded without ubiquitination¹⁸. These proteins contain short peptide sequences, termed ubiquitin-independent degrons, that promote rapid proteasomal degradation^{19–23}. Ubiquitin-independent degrons simultaneously function as proteasome recognition elements that engage the 19S regulatory particle and unstructured initiation regions for 20S core particle degradation^{18,20,22–26}. These peptides are transferrable; conjugating a ubiquitin-independent degron to a heterologous protein converts it to a short-lived, ubiquitin-independent proteasome substrate^{22–24,26,27}. This property has been leveraged to create genetically encoded, high-throughput reporters of proteasome activity^{20,28}. However, in genetically diverse populations, the output of these systems is confounded by differences in reporter expression caused by individual genetic differences between cells^{29–32}.

Here, we used the ubiquitin-independent degrons from ornithine decarboxylase (ODC)^{19,20} and Rpn4^{21–23} to build high-throughput reporters for measuring proteasome activity in genetically diverse cell populations. We used these reporters to perform statistically powerful QTL mapping of the genetics of proteasome activity. Our results reveal that proteasome activity is a polygenic trait shaped by variation throughout the genome. Proteasome activity QTLs were largely substrate-specific, suggesting that genetic variation shapes proteasome activity primarily through effects on the 19S regulatory particle. Individual genetic differences are an important source of variation in proteasome activity that may contribute to the complex genetic basis of the many cellular and

organismal traits influenced by proteasomal protein degradation.

Results

Single-Cell Measurements Reveal Genetic Differences in Proteasome Activity

We sought to develop a reporter system capable of measuring proteasome activity with high throughput and quantitative precision without confounding from genetic influences on reporter expression. To do so, we built a series of tandem fluorescent timers (TFTs), fusions of two fluorescent proteins with distinct spectral profiles and maturation kinetics^{33,34}. Our TFTs contained a faster-maturing green fluorescent protein (GFP) and a slower-maturing red fluorescent protein (RFP) (Figure 1A). The two fluorophores in the TFT mature at different rates. As a result, the RFP / GFP ratio changes over time. If the TFT's degradation rate is faster than the RFP's maturation rate, the negative $\log_2$ RFP / GFP ratio is directly proportional to the TFT's degradation rate (Figure 1B). The TFT's RFP / GFP ratio is independent of the reporter's expression level¹⁷, making it possible to use TFTs in genetically diverse cell populations without confounding from genetic influences on reporter expression^{14,17}.

To relate the RFP / GFP ratio to proteasome activity, we fused the ubiquitin-independent degrons from the mouse ODC and yeast Rpn4 proteins to our TFTs. The ubiquitin-independent proteasomal degradation of mouse ODC is conserved in yeast and, unlike yeast ODC, does not require expression of antizyme for efficient proteasomal degradation^{20,24}. We fused amino acids 410 through 461 of mouse ODC to the C-terminus of the sfGFP-mCherry TFT, consistent with the geometric requirements of the ODC degron²⁰, to create the ODC TFT (Figure 1C). The Rpn4 protein contains a ubiquitin-independent degron in amino acids 1 to 80^{22,23}. We fused this sequence to the N-terminus of the sfGFP-mCherry TFT to create the Rpn4 TFT (Figure 1C). We reasoned that the distinct recognition mechanisms, sequences, and geometric requirements of the ODC and Rpn4 degrons would allow us to distinguish substrate-specific genetic effects on the 19S regulatory particle from substrate-independent genetic effects on the 20S core particle.

We characterized the ODC and Rpn4 TFTs in single cells by flow cytometry. We first evaluated the sensitivity and dynamic range of each TFT by comparing the RFP / GFP ratio in the BY laboratory strain, which is closely related to the S288C reference strain, and a BY strain lacking the *RPN4* gene (hereafter “BY *rpn4*Δ”). *RPN4* encodes a transcription factor for proteasome genes, and deleting *RPN4* reduces proteasome activity^{21,25,35}. Deleting *RPN4* strongly reduced

the negative $\log_2$ RFP / GFP output from the ODC and Rpn4 TFTs (t-test $p = 1.4\text{e-}6$ and $1.6\text{e-}13$, respectively; Figure 1D/E), showing that our TFTs provide sensitive measurements of proteasome activity. Consistent with previous reports^{18,26,27}, in the BY strain the ODC TFT was more rapidly degraded than the Rpn4 TFT (t-test $p = 6.9\text{e-}10$, Figure 1D/E). Taken together, our results show that our TFTs provide quantitative, substrate-specific readouts of proteasome activity.

To understand how natural genetic variation affects proteasome activity, we measured the output of the ODC and Rpn4 TFTs in two genetically divergent *S. cerevisiae* strains, BY and RM, a genetically divergent vineyard strain. The RM strain showed higher proteasome activity towards the ODC and Rpn4 TFTs than BY (t-test $p = 1.9\text{e-}4$ and $1.2\text{e-}8$, respectively; Figure 1D/E). We observed a significant interaction between strain background and proteasome substrate such that the magnitude of the BY/RM strain difference was greater for the Rpn4 TFT than the ODC TFT (two-way ANOVA interaction $p = 0.013$). Together, these results show that individual genetic differences create heritable, substrate-dependent variation in proteasome activity.

Figure 1: *Design and characterization of proteasome activity reporters. A. Schematic of the production and maturation of a TFT. B. A bar plot created with hypothetical data shows how differences in a TFT's degradation rate influence the reporter's RFP, GFP, and negative log₂ RFP / GFP levels. C. Diagram of mouse ODC and yeast Rpn4 showing the location of each protein's ubiquitin-independent degron. "AZB" = antizyme binding site, "AS" = active site, "AD" = transcriptional activation domain, "C₂H₂" = C₂H₂ zinc finger DNA binding domain. D. Density plots of the negative log₂ RFP / GFP ratio from 10,000 cells for each of 8 independent biological replicates per strain per reporter for the indicated strains and TFTs. Thin, opaque lines show individual biological replicates and thicker, transparent lines show the group average for the indicated strains. E. The median from each biological replicate in D. is plotted as a stripchart. t-test p-values are shown for the indicated strain versus BY.*

Bulk Segregant Analysis Identifies Complex, Polygenic Influences on Proteasome Activity

To map genetic influences on proteasome activity, we used our ODC and Rpn4 TFTs to perform bulk segregant analysis, a statistically powerful genetic mapping method that compares large numbers of individuals with extreme values for a trait of interest selected from a genetically diverse population^{14,32,36–38}. In our implementation, the method identifies proteasome activity QTLs, regions of the genome containing one or more DNA variants influencing proteasome activity. We created genetically diverse cell populations by mating BY strains harboring either the ODC or Rpn4 TFT with RM and sporulating the resulting diploids (Figure 2A). We collected cells from the 2% tails of the proteasome activity distribution using fluorescence-activated cell sorting (FACS) (Figure 2B-E) and used whole-genome sequencing to determine the allele frequency difference between the high and low UPS activity pools at each DNA variant. At QTLs affecting proteasome activity, the allele frequencies will be significantly different between pools, while at unlinked loci the allele frequencies will be the same. We called significant QTLs using a logarithm of the odds (LOD) threshold previously determined to produce a 0.5% false discovery rate for TFT-based genetic mapping¹⁴ (see "Methods") and retained only QTLs detected at genome-wide significance in both of two independent biological replicates. To visualize the direction and magnitude of QTL effect sizes, we plotted the allele frequency difference along each chromosome (Supplementary Figures 1 and 2). We identified 11 QTLs for the ODC TFT and 7 QTLs for the Rpn4 TFT (Figure 3, Table 1). The distribution of proteasome activity QTL effect sizes, as measured by the allele

frequency difference between pools, was continuous and consisted predominantly of QTLs with small effects (Figure 3, Table 1). Together, our mapping results demonstrate that proteasome activity is a polygenic trait, shaped by variation throughout the genome.

Figure 2: Mapping genetic influences on proteasome activity using bulk segregant analysis. *A.* Schematic of the experimental approach. *B./C.* Proteasome activity distributions for the ODC TFT (*B.*) and Rpn4 TFT (*C.*). Vertical lines show the gates used to collect cells with extreme high or low proteasome activity. *D./E.* Backplot of cells collected using the gates in *B./C.* onto a scatter plot of GFP and RFP for the ODC (*D.*) and Rpn4 (*E.*) TFTs.

Figure 3: Proteasome activity QTLs for the ODC and Rpn4 TFTs. A. Line plot of the RM allele frequency difference between the high and low UPS activity pools at the peak position for statistically significant proteasome activity QTLs. B. As in A., but for the LOD score for proteasome activity QTLs. In A./B. asterisks denote shared QTLs detected with the ODC and Rpn4 TFTs.

Reporter	Chromosome	LOD	AFD	Peak Position	Left Index	Right Index
ODC TFT	IIa	9.76	0.099	69800	32850	107100
ODC TFT	IIb	7.13	-0.121	418100	358850	462650
ODC TFT	IVa	5.64	-0.101	85150	30400	127400
ODC TFT	V	12.83	-0.149	291350	247700	325650
ODC TFT	VIIa	8.14	-0.150	20000	0	52800
ODC TFT	VIIb	28.74	0.225	409000	390050	431700
ODC TFT	X	16.36	0.182	666850	649350	691550
ODC TFT	XII	8.13	0.111	768150	666200	846700
ODC TFT	XIIIa	18.96	0.192	47800	25200	75850
ODC TFT	XIIIb	7.96	0.127	410900	377350	450100
ODC TFT	XIVa	8.81	-0.110	441750	381400	501600
Rpn4 TFT	IVb	12.64	-0.133	240600	213200	309150
Rpn4 TFT	V	10.09	-0.127	259650	218250	294900
Rpn4 TFT	VIIa	10.21	-0.148	88550	53550	141350
Rpn4 TFT	VIIc	6.80	-0.111	882500	840650	926150
Rpn4 TFT	XII	40.11	0.233	672850	661800	685750
Rpn4 TFT	XIVb	16.58	0.145	544150	497300	574600
Rpn4 TFT	XV	30.00	-0.224	167400	142600	186200

Table 1: *Proteasome activity QTLs. The table lists all QTLs detected, sorted first by reporter, then by chromosome. Lowercase letters following chromosome numbers are used to distinguish QTLs on the same chromosome. “LOD”, logarithm of the odds; “AFD”, RM allele frequency difference (high proteasome activity pool minus low proteasome activity pool) at the QTL peak position. Each number is the average value calculated from two independent biological replicates for a given QTL. “Peak Position”, “Left Index”, and “Right Index” refer to base pair positions on the indicated chromosome.*

Genetic Influences on Proteasome Activity are Predominantly Substrate-Specific

We evaluated the extent to which QTLs detected with the ODC and Rpn4 TFTs overlapped, reasoning that shared QTLs may reflect substrate-independent genetic influences on proteasome activity, such as altered 20S core particle activity. Only 3 of 18 QTLs (on chromosomes V, VII, and XII; Figure 3, Table 1) were shared between the ODC and Rpn4 TFTs. Because QTLs

that influence the 20S core particle would be expected to affect both TFTs, the high degree of observed QTL specificity between reporters suggests that variant effects on proteasome activity predominantly influence substrate selection by the 19S regulatory particle.

We recently mapped genetic effects on the UPS N-end Rule pathway¹⁴, in which ubiquitinated N-terminal amino acids function as 19S regulatory particle recognition elements⁶. We compared the overlap of the 149 UPS N-end rule activity QTLs (obtained earlier using 20 N-end rule reporters¹⁴) and the 18 proteasome activity QTLs to better understand the substrate specificity of genetic influences on proteasome activity. Six proteasome activity QTLs did not overlap any of the 149 UPS activity QTLs (Figure 4, Supplementary Table 1). Even though the ODC/Rpn4 degrons and poly-ubiquitin chains each act as proteasome binding tags, these substrate-specific QTLs suggest that genetic variation may differentially alter the processing, function, or recognition of these elements. The remaining 12 proteasome activity QTLs overlapped with varying numbers of the 20 UPS N-end rule reporters, with a range of 1 to 10 and a median of 3 (Figure 4, Supplementary Table 1). The low extent of overlap in proteasome activity QTLs and UPS N-End Rule QTLs further highlights the high degree of substrate specificity in genetic influences on proteasome activity.

Figure 4: *Overlap of proteasome activity QTLs with UPS N-end Rule QTLs. The barplot shows the number of UPS N-end Rule reporters (of a possible 20) for which a given proteasome activity QTL was detected. QTLs on the y-axis are labeled as in Table 1.*

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176 We previously used QTLs detected with multiple UPS N-end rule reporters to nominate and validate
177 candidate causal QTL genes¹⁴. We applied this approach to our proteasome activity QTLs to gain
178 insights into the mechanisms by which genetic variation may shape proteasome activity. A QTL on
179 chromosome XII was detected with the ODC and Rpn4 TFTs and 5 N-end Rule reporters (Figure
180 4, Supplementary Table 1). The QTL's median peak position at base pair 665,600 occurs in the
181 immediate vicinity of *HAP1*, a transcription factor at which BY carries a transposon insertion that
182 influences a variety of traits in this cross of yeast strains^{31,39,40}. A QTL on chromosome XIV was
183 detected with the ODC TFT and 9 UPS N-end rule reporters and its peak at base pair 464,275 was
184 located near the *MKT1* gene (Figure 4, Supplementary Table 1). *MKT1* encodes an RNA-binding
185 protein that also influences numerous traits in the BY/RM cross^{31,31,32,37,41}. These QTLs thus
186 likely shape proteasome activity through indirect mechanisms. The chromosome VIIb QTL was
187 detected using the ODC TFT and 10 UPS N-end rule TFTs (Figure 4, Supplementary Table 1).

The median peak position of this QTL (at base pair 409,450) occurs in the immediate vicinity of *RPT6*, an ATPase of the 19S regulatory particle. The RM allele of this QTL increases proteasome or UPS activity for all reporters for which the QTL was detected. *RPT6* expression is increased by the RM allele at this locus³¹, making *RPT6* a strong candidate causal gene for this QTL. This region was not linked to the output of the Rpn4 TFT, suggesting that the effect of variants at *RPT6* are likely to affect the 19S regulatory particle in a substrate-specific manner.

Discussion

Protein homeostasis depends on precise, dynamic control of proteasome activity. However, the factors that create physiological and pathological variation in proteasome activity remain incompletely understood. Using a statistically powerful genetic mapping method, we have established individual genetic differences as an important source of heritable variation in proteasome activity. Genetic influences on proteasome activity are numerous and, similar to other polygenic traits^{40,42,43}, comprise a continuous distribution of effect sizes, with many QTLs of small effect and few QTLs of large effect. Our results add to the emerging picture of the complex effects of genetic variation on the UPS, which include widespread effects on the activity of both the ubiquitin system¹⁴ and, as we show here, the proteasome.

The proteasome consists of two functionally distinct modules. The 19S regulatory particle binds, unfolds, and translocates proteins to the 20S core particle, which processively degrades them²⁻⁵. By mapping genetic influences on two ubiquitin-independent proteasome substrates with distinct sequences, geometries, and recognition mechanisms^{18-23,44}, we observed that genetic influences on proteasome activity were primarily substrate-specific (Figure 3, Table 1). Because all proteasome substrates are processed by the 20S core particle, DNA variants that influence the 20S core particle would be expected to affect all substrates, even if some effects may be missed due to incomplete power to detect QTLs. In contrast, the high degree of substrate-specificity we observed suggests that genetic influences on the 19S regulatory particle are more numerous than those affecting the 20S core particle. This notion is consistent with the multiple functions served by the 19S core particle. DNA sequence differences may influence the 19S regulatory particle's substrate binding, unfolding, or translocation to the 20S core particle^{2,4}. The 20S core particle is highly evolutionarily conserved and, unlike the eukaryote-specific 19S regulatory particle, is found in eukaryotes, bacteria, and archaea^{45,46}. Thus, variation in 20S core particle genes or in genes with *trans*-acting influences on 20S core particle function or activity may be rapidly purged by purifying

selection.

The numerous proteasome activity QTLs we have identified add new insight into how genetic variation shapes the molecular effectors of cellular protein degradation. We recently mapped the genetics of the UPS N-end rule pathway and discovered multiple DNA variants that alter the activity of several components of the ubiquitin system¹⁴. Here, we show that genetic effects on proteasome activity are similarly numerous. However, our analysis of QTL sharing suggests relatively little overlap in the sources of heritable variation in the ubiquitin system and the proteasome (Figure 4). Although many stimuli, such as protein misfolding or heat shock, cause similar changes in the activity of the ubiquitin system and the proteasome, recent work shows that these systems can also be regulated independently and function autonomously of one another^{7,47}. For example, ubiquitination can initiate events besides proteasomal protein degradation, including lysosomal protein degradation, altered protein subcellular localization, and signaling cascade activation^{47–49}. Likewise, a number of cellular proteins are bound and degraded by the proteasome without modification by the ubiquitin system¹⁸. Thus, predicting how genetic variation shapes the degradation of UPS substrates will require consideration of genetic effects on both the ubiquitin system and the proteasome.

We have developed a reporter system that provides high-throughput, quantitative, *in vivo* measurements of proteasome activity in genetically diverse cell populations. The elements in our reporters function in many other eukaryotic organisms^{17,22,24} and applying this system to other species may clarify how genetic effects on proteasome activity shape cellular and organismal traits. Alleles of proteasome genes confer beneficial traits in plants, including heat tolerance⁵⁰ and resistance to multiple pathogens^{51,52}. Mutations in proteasome genes have been linked to a strikingly diverse array of human diseases, including coronary artery disease⁵³, ischaemic stroke⁵⁴, intellectual disability^{55,56}, deafness⁵⁷, morphological abnormalities⁵⁷, and Graves' disease^{58,59}. However, the vast majority of studies linking variation in proteasome genes to organismal phenotypes have not tested whether and how the identified mutations alter proteasome activity. Applying our reporter system to proteasome gene variants would provide insight into the mechanisms and magnitude of the effects of these variants. Heritable variation in traits regulated by the proteasome may also result from the cumulative contributions of many alleles with small effects. Applying the mapping strategy developed here or using our reporters with other recently-developed mapping approaches⁶⁰ may provide new insights into the genetic basis of a variety of cellular and organismal traits, including the many diseases marked by aberrant proteasome activity.

Materials and Methods

Tandem Fluorescent Timer (TFT) Reporters of Proteasome Activity

We used TFTs, fusions of two fluorescent proteins with distinct spectral profiles and maturation kinetics, to measure proteasome activity. The most common TFT implementation consists of a faster-maturing green fluorescent protein (GFP) and a slower-maturing red fluorescent protein (RFP)^{17,33,34,61}. Because the two fluorescent proteins mature at different rates, the RFP / GFP ratio changes over time. If the TFT's degradation rate is faster than the RFP's maturation rate, the negative log₂ RFP / GFP ratio is directly proportional to the TFT's degradation rate^{17,33}. The RFP / GFP ratio is also independent of the TFT's expression level,^{17,33} enabling high-throughput, quantitative measurements of TFT turnover in genetically diverse cell populations^{14,17}. All TFTs in the present study contained superfolder GFP (sfGFP)⁶² and mCherry⁶³ separated by an unstructured 35 amino acid peptide sequence to minimize fluorescence resonance energy transfer¹⁷.

To measure proteasome activity with our TFTs, we fused the ubiquitin-independent degrons from the mouse ornithine decarboxylase (ODC) and yeast Rpn4 proteins to our sfGFP-mCherry TFTs. Ornithine decarboxylase (an enzyme in polyamine biosynthesis) contains a ubiquitin-independent degron in its C-terminal 37 amino acids^{19,20,27,44}. Rpn4 (a transcription factor for proteasome genes) contains a ubiquitin-independent degron in its N-terminal 80 amino acids^{21,22,25}. Both degrons are recognized and bound by the 19S regulatory particle without ubiquitin conjugation and function as unstructured initiation regions⁶⁴ for 20S core particle degradation. Attaching either degron to a heterologous protein converts it into a short-lived proteasomal substrate with half-lives of approximately 5 minutes for the ODC degron and 20 minutes for the Rpn4 degron^{18,26,27}. The ODC and Rpn4 degron sfGFP-mCherry TFTs thus provide direct, quantitative, substrate-specific readouts of proteasome activity (Figure 1).

We used a previously described approach¹⁴ to construct TFT reporters and yeast strains harboring TFTs. Each TFT contained the constitutively active *TDH3* promoter, the *ADH1* terminator, sfGFP, mCherry, and the Kanmx selection module⁶⁵. TFTs were constructed so that the ubiquitin-independent degron was immediately adjacent to mCherry (Figure 1C), consistent with established guidelines for optimizing TFT function³⁴. We used BFA0190 as the plasmid backbone for all TFT plasmids. BFA0190 contains 734 bp of sequence upstream and 380 bp of sequence downstream of the *LYP1* ORF separated by a *SwaI* restriction site. We inserted TFT reporters into BFA0190 by digesting the plasmid with *SwaI* and inserting TFT components between the *LYP1* flanking

sequences using isothermal assembly cloning (Hifi Assembly Cloning Kit; New England Biolabs [NEB], Ipswich, MA, USA). The 5' and 3' *LYP1* flanking sequences in each TFT plasmid contain natural SacI and BglII restriction sites, respectively. We produced linear DNA transformation fragments by digesting TFT-containing plasmids with SacI and BglII and gel purifying the fragments (Monarch Gel Purification, NEB). Genomic integration of each linear transformation fragment results in deletion of the *LYP1* gene, allowing selection for TFT integration at the *LYP1* locus using the toxic amino acid analogue thialysine (S-(2-aminoethyl)-L-cysteine hydrochloride)^{66–68} and G418⁶⁵.

Yeast Strains and Handling

Yeast Strains

We used two genetically divergent *Saccharomyces cerevisiae* yeast strains for characterizing our proteasome activity TFTs and mapping genetic influences on proteasome activity. The haploid BY strain (genotype: MATa *his3Δ hoΔ*) is a laboratory strain that is closely related to the *S. cerevisiae* S288C reference strain. The haploid RM strain is a vineyard isolate with genotype MATα *can1Δ::STE2pr-SpHIS5 his3Δ::NatMX AMN1-BY hoΔ::HphMX URA3-FY*. BY and RM differ, on average, at 1 nucleotide per 200 base pairs, such that approximately 45,000 single nucleotide variants (SNVs) between the strains can serve as markers in a genetic mapping experiment^{32,36,37,40}. We also engineered a BY strain lacking the *RPN4* gene (hereafter “BY *rpn4Δ*”) to characterize the sensitivity and dynamic range of our TFT reporters. We replaced the *RPN4* gene with the NatMX cassette, which confers resistance to the antibiotic nourseothricin⁶⁵. To do so, we transformed BY with a DNA fragment created by PCR amplifying the NatMX cassette from plasmid from Addgene plasmid #35121 (a gift from John McCusker) using primers with 40 bp of homology to the 5' upstream and 3' downstream sequences of *RPN4* using the transformation procedure described below. Strain genotypes are presented in Table 2.

Short Name	Genotype	Antibiotic Resistance	Auxotrophies
BY	<i>MATa his3Δ hoΔ</i>		histidine
RM	<i>MATα can1Δ::STE2pr-SpHIS5 his3Δ::NatMX hoΔ::HphMX</i>	clonNAT, hygromycin	histidine
BY <i>rpn4Δ</i>	<i>MATa his3Δ hoΔ rpn4Δ::NatMX</i>	clonNAT	histidine

Table 2: Strain genotypes

The media formulations for all experiments are listed in Table 3. Synthetic complete media powders (SC -lys and SC -his -lys -ura) were obtained from Sunrise Science (Knoxville, TN, USA). We added the following reagents at the following concentrations to yeast media where indicated: G418, 200 mg/mL (Fisher Scientific, Pittsburgh, PA, USA); ClonNAT (nourseothricin sulfate, Fisher Scientific), 50 mg/L; thialysine (S-(2-aminoethyl)-L-cysteine hydrochloride; MilliporeSigma, St. Louis, MO, USA), 50 mg/L; canavanine (L-canavanine sulfate, MilliporeSigma), 50 mg/L.

Media Name	Abbreviation	Formulation
Yeast-Peptone-Dextrose	YPD	10 g/L yeast extract 20 g/L peptone 20 g/L dextrose
Synthetic Complete	SC	6.7 g/L yeast nitrogen base 1.96 g/L amino acid mix -lys 20 g/L dextrose
Haploid Selection	SGA	6.7 g/L yeast nitrogen base 1.74 g/L amino acid mix -his -lys -ura 20 g/L dextrose
Sporulation	SPO	1 g/L yeast extract 10 g/L potassium acetate 0.5 g/L dextrose

Table 3: Media Formulations

Yeast Transformations

We used the lithium acetate/single-stranded carrier DNA/polyethylene glycol (PEG) method for all yeast transformations⁶⁹. In brief, yeast strains were inoculated into 5 mL of YPD liquid medium for overnight growth at 30 °C. The next day, we diluted 1 mL of each saturated culture into 50 mL of fresh YPD and grew cells for 4 hours. Cells were washed in sterile ultrapure water and then in transformation solution 1 (10 mM Tris HCl [pH 8.0], 1 mM EDTA [pH 8.0], and 0.1 M lithium acetate). After each wash, we pelleted the cells by centrifugation at 3,000 rpm for 2 minutes in a benchtop centrifuge and discarded supernatants. After washing, cells were suspended in 100 μ L of transformation solution 1 along with 50 μ g of salmon sperm carrier DNA and 300 ng of transforming DNA and incubated at 30 °C for 30 minutes with rolling. Subsequently, 700 μ L of transformation

solution 2 (10 mM Tris HCl [pH 8.0], 1 mM EDTA [pH 8.0], and 0.1 M lithium acetate in 40% PEG) was added to each tube, followed by a 30 minute heat shock at 42 °C. Transformed cells were then washed in sterile, ultrapure water, followed by addition of 1 mL of liquid YPD medium to each tube. Cells were incubated in YPD for 90 minutes with rolling at 30 °C to allow for expression of antibiotic resistance cassettes. We then washed the cells with sterile, ultrapure water and plated 200 μ L of cells on solid SC -lys medium with G418 and thialysine, and, for strains with the NatMX cassette, clonNAT. We single-colony purified multiple independent colonies (biological replicates) from each transformation plate for further analysis as indicated in the text. Reporter integration at the targeted genomic locus was verified by colony PCR⁷⁰.

Yeast Mating and Segregant Populations

We used a modified synthetic genetic array (SGA) methodology^{67,68} to create populations of genetically variable, recombinant cells (“segregants”) for genetic mapping. BY strains with either ODC or Rpn4 TFTs were mixed with the RM strain on solid YPD medium and grown overnight at 30 °C. We selected for diploid cells (successful BY/RM matings) by streaking mixed BY/RM cells onto solid YPD medium containing G418, which selects for the KanMX cassette in the TFT in the BY strain, and clonNAT, which selects for the NatMX cassette in the RM strain). Diploid cells were inoculated into 5 ml of liquid YPD and grown overnight at 30 °C. The next day, cultures were washed with sterile, ultrapure water, and resuspended in 5 mL of SPO liquid medium (Table 3). We induced sporulation by incubating cells in SPO medium at room temperature with rolling for 9 days. After confirming sporulation by brightfield microscopy, we pelleted 2 mL of cells, which were then washed with 1 mL of sterile, ultrapure water, and resuspended in 300 μ L of 1 M sorbitol containing 3 U of Zymolyase lytic enzyme (United States Biological, Salem, MA, USA) to degrade ascus walls. Asci were digested for 2 hours at 30 °C with rolling. Spores were then washed with 1 mL of 1 M sorbitol, vortexed for 1 minute at the highest intensity setting, and resuspended in sterile ultrapure water. We confirmed the release of cells from asci by brightfield microscopy and plated 300 μ L of cells onto solid SGA medium containing G418 and canavanine. This media formulation selects for haploid cells with (1) a TFT via G418, (2) the *MATa* mating type via the *Schizosaccharomyces pombe* *HIS5* gene under the control of the *STE2* promoter (which is only active in *MATa* cells), and (3) replacement of the *CAN1* gene with *S. pombe* *HIS5* via the toxic arginine analog canavanine^{67,68}. Haploid segregants were grown for 2 days at 30 °C and harvested by adding 10 mL of sterile, ultrapure water and scraping the cells from each plate. Each segregant population cell suspension was centrifuged at 3000 rpm for 10 minutes and resuspended in 1 mL of SGA medium. We added 450 μ L of 40% (v/v) sterile glycerol solution to 750 μ L to each segregant culture and

stored this mixture in screw cap cryovials at -80°C . We stored 2 independent sporulations each of the ODC and Rpn4 degron TFT-containing segregants (derived from our initial matings) as independent biological replicates.

Flow Cytometry and Fluorescence-Activated Cell Sorting

Flow Cytometry

We characterized our proteasome activity TFTs using flow cytometry. For all flow cytometry experiments, we inoculated yeast strains into 400 μL of liquid SC -lys medium with G418 for overnight growth in 2 mL 96 well plates at 30°C with 1000 rpm mixing on a MixMate (Eppendorf, Hamburg, Germany). The next day, 4 μL of each saturated culture was inoculated into a fresh 400 μL of G418-containing SC -lys media and cells were grown for an additional 3 hours prior to flow cytometry. We performed all flow cytometry experiments on an LSR II flow cytometer (BD, Franklin Lakes, NJ, USA) equipped with a 20 mW 488 nm laser with 488/10 and 525/50 filters for measuring forward and side scatter and sfGFP fluorescence, respectively, as well as a 40 mW 561 nm laser and a 610/20 filter for measuring mCherry fluorescence. Table 4 lists the parameters and settings for all flow cytometry and fluorescence-activated cell sorting (FACS) experiments. We recorded 10,000 cells each from 8 independent biological replicates per strain for the ODC and Rpn4 degron TFTs.

Parameter	Laser Line (nm)	Laser Setting (V)	Filter
forward scatter (FSC)	488	500	488/10
side scatter (SSC)	488	275	488/10
sfGFP	488	500	525/50
mCherry	561	615	610/20

Table 4: *Flow cytometry and FACS settings.*

All flow cytometry data was analyzed using R (R Foundation for Statistical Computing, Vienna Austria) and the flowCore R package⁷¹. We filtered each flow cytometry dataset to exclude all events outside of $10\% \pm$ the median forward scatter (a proxy for cell size). This gating approach captured the central peak of cells in the FSC histogram and removed cellular debris, aggregates of multiple cells, and restricted our analyses to cells of the same approximate size¹⁴.

For the statistical analysis of flow cytometry results, we used the negative $\log_2$ RFP / GFP ratio as a measure of proteasome activity. We computed the negative $\log_2$ RFP / GFP ratio for each of 10,000 cells per strain per reporter and extracted the median from each sample for subsequent statistical analyses. We used a one-way ANOVA with Tukey's HSD post-hoc test for all between groups comparisons of proteasome activity.

Fluorescence-Activated Cell Sorting

We used FACS to collect pools of segregant cells for genetic mapping by bulk segregant analysis^{32,37}. We thawed and inoculated segregant populations into 5 mL of SGA medium containing G418 and canavanine for overnight growth at 30 °C with rolling. The following morning, we diluted 1 mL of cells from each segregant population into a fresh 4 mL of SGA medium containing G418 and canavanine. Diluted segregant cultures were grown for 4 hours prior to sorting on a FACSAria II cell sorter (BD). Plots of side scatter (SSC) height by SSC width and forward scatter (FSC) height by FSC width were used to remove doublets from each sample and cells were further filtered to contain cells within $\pm 7.5\%$ of the central FSC peak. We empirically determined that this filtering approach excluded cellular debris and aggregates while retaining the primary haploid cell population. We also defined a fluorescence-positive population by retaining only those TFT-containing cells with sfGFP fluorescence values higher than negative control BY and RM strains without TFTs. We collected pools of 20,000 cells each from the 2% high and low proteasome activity tails (Figure 2) from two independent biological replicates (spore preparations) for each TFT. Pools of 20,000 cells were collected into sterile 1.5 mL polypropylene tubes containing 1 mL of SGA medium that were grown overnight at 30 °C with rolling. After overnight growth, we mixed 750 μ L of cells with 450 μ L of 40% (v/v) glycerol and stored this mixture in 2 mL 96 well plates at -80°C .

Genomic DNA Isolation, Library Preparation, and Whole-Genome Sequencing

To isolate genomic DNA from sorted segregant pools, we first pelleted 800 μ L of each pool by centrifugation at 3,700 rpm for 10 minutes. Supernatants were discarded and cell pellets were resuspended in 800 μ L of a 1 M sorbitol solution containing 0.1 M EDTA, 14.3 mM β -mercaptoethanol, and 500 U of Zymolyase lytic enzyme (United States Biological) to digest cell walls. Zymolyase digestions were carried out by resuspending cell pellets with mixing at 1000 rpm for 2 minutes followed by incubation for 2 hours at 37 °C. After completing the digestion reaction, we pelleted and resuspended cells in 50 μ L of phosphate buffered saline. We then used the Quick-DNA 96 Plus kit (Zymo Research, Irvine, CA, USA) to extract genomic DNA according to the manufacturer's protocol, including an overnight protein digestion in a 20 mg/mL

proteinase K solution at 55 °C prior to loading samples onto columns. DNA was eluted from sample preparation columns using 40 μ L of DNA elution buffer (10 mM Tris-HCl [pH 8.5], 0.1 mM EDTA). DNA concentrations for each sample were determined with the Qubit dsDNA BR assay kit (Thermo Fisher Scientific, Waltham, MA, USA) in a 96 well format using a Synergy H1 plate reader (BioTek Instruments, Winooski, VT, USA).

We used genomic DNA from our segregant pools to prepare a short-read library for whole-genome sequencing on the Illumina Next-Seq platform using a previously-described approach^{14,32,37}. The Nextera DNA library kit (Illumina, San Diego, CA, USA) was used according to the manufacturer's instructions with the following modifications. We fragmented and added sequencing adapters to genomic DNA by adding 5 ng of DNA to a master mix containing 4 μ L of Tagment DNA buffer, 1 μ L of sterile molecular biology grade water, and 5 μ L of Tagment DNA enzyme diluted 1:20 in Tagment DNA buffer and incubating this mixture on a SimpliAmp thermal cycler using the following parameters (Thermo Fisher Scientific): 55 °C temperature, 20 μ L reaction volume, 10 minute incubation. We PCR amplified libraries prior to sequencing by adding 10 μ L of the tagmentation reaction to a master mix containing 1 μ L of an Illumina i5 and i7 index primer pair mixture, 0.375 μ L of ExTaq polymerase (Takara), 5 μ L of ExTaq buffer, 4 μ L of a dNTP mixture, and 29.625 μ L of sterile molecular biology grade water. To multiplex samples for sequencing, we generated all 96 possible index oligo combinations using 8 i5 and 12 i7 index primers. Libraries were PCR amplified on a SimpliAmp thermal cycler using the following parameters: initial denaturation at 95 °C for 30 seconds, then 17 cycles of 95 °C for 10 seconds (denaturation), 62 °C for 30 seconds (annealing), and 72 °C for 3 minutes (extension). The DNA concentration of each reaction was quantified using the Qubit dsDNA BR assay kit (Thermo Fisher Scientific). We pooled equimolar amounts of each sample, ran this mixture on a 2% agarose gel, and extracted and purified DNA in the 400 bp to 600 bp region using the Monarch Gel Extraction Kit (NEB) according to the manufacturer's instructions.

The pooled library was submitted to the University of Minnesota Genomics Center (UMGC) for quality control assessment and Illumina sequencing. UMGc staff performed three quality control (QC) assays prior to sequencing. The PicoGreen dsDNA quantification reagent (Thermo Fisher Scientific) was used to determine library concentration, with a concentration ≥ 1 ng/ μ L required to pass. The TapeStation electrophoresis system (Agilent Technologies, Santa Clara, CA, USA) was used to determine library size, with libraries in the range of 200 to 700 bp passing. Finally, the KAPA DNA Library Quantification kit (Roche, Penzberg, Germany) was used to determine library

functionality, with libraries requiring a concentration $\geq$ than 2 nM to pass. The submitted library passed each QC assay. The library was sequenced on a Next-Seq 550 instrument in mid-output, 75 bp paired-end mode, generating 153,887,828 reads across all samples, with a median of 9,757,090 and a range of 5,994,921 to 14,753,319 reads per sample. The mean read quality for all samples was > 30 . The median read coverage of the genome was 21, with a range of 16 to 25 across all samples. Data will be deposited into the NIH Sequence Read Archive following publication.

QTL Mapping

We used a previously-described approach to identify QTLs from our whole-genome sequencing data^{14,32,37}. We initially filtered our raw reads to retain only those with a mean base quality score greater than 30. Filtered reads were aligned to the *S. cerevisiae* reference genome (sacCer3) with the Burroughs-Wheeler alignment tool⁷². We used samtools⁷³ to remove unaligned reads, non-uniquely aligned reads, and PCR duplicates and then to produce vcf files containing coverage and allelic read counts at each of 18,871 high-confidence, reliable SNPs^{36,74}, with BY alleles as reference and RM alleles as alternative alleles.

QTLs were called from allele counts using the MULTIPPOOL algorithm⁷⁵. MULTIPPOOL estimates a logarithm of the odds (LOD) score by calculating a likelihood ratio from two models. In the noncausal model, the locus is not associated with the trait and the high and low proteasome activity pools have the same frequency of the BY and RM alleles. In the causal model, the locus is associated with the trait, such that the BY and RM allele frequencies differ between pools. QTLs were defined as loci with a LOD $\geq$ to 4.5. In a previous study¹⁴, we empirically determined that this threshold produces a 0.5% false discovery rate (FDR) for TFT-based genetic mapping by bulk segregant analysis. We used the following MULTIPPOOL settings: bp per centiMorgan = 2,200, bin size = 100 bp, effective pool size = 1,000. As in previous studies^{32,37}, we excluded variants with allele frequencies higher than 0.9 or lower than 0.1^{14,32,37}. QTL confidence intervals were defined as a 2-LOD drop from the QTL peak (the QTL position with the highest LOD value). We also computed the allele frequency difference (Δ AF) at each allele as a measure of each QTL's effect size and to visualize QTLs. Because allele frequencies are affected by random counting noise, we smoothed allele frequencies along the genome using loess regression prior to calculating Δ AF for each sample.

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485 **Competing Interests**

486 The authors declare that they have no competing interests.

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489 **Supplementary Materials**

490 The following supplementary materials are included with this manuscript:

491 Supplementary Figures 1-2

492 Supplementary Table 1

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Supplementary Materials

Supplementary Figure 1. ODC TFT QTL mapping results. Top. The loess smoothed RM allele frequency difference (high proteasome activity pool minus low proteasome activity pool) is plotted across the *S. cerevisiae* genome. The dashed red lines mark the empirically determined 99.9% quantile. Bottom. LOD scores for ODC TFT QTL mapping. The dashed red line indicates the empirically determined LOD score significance threshold (4.5), which results in a 0.5% FDR. For both panels, asterisks denote QTLs, colored by independent biological replicate.

Supplementary Figure 2. *Rpn4* TFT QTL mapping results. *Top.* The loess smoothed RM allele frequency difference (high proteasome activity pool minus low proteasome activity pool) is plotted across the *S. cerevisiae* genome. The dashed red lines mark the empirically determined 99.9% quantile. *Bottom.* LOD scores for *Rpn4* TFT QTL mapping. The dashed red line indicates the empirically determined LOD score significance threshold (4.5), which results in a 0.5% FDR. For both panels, asterisks denote QTLs, colored by independent biological replicate.

Reporters	Chromosome	LOD	AFD	Peak Position	Left Index	Right Index
ODC TFT/N-end Rule (3)	IVa	7.12	-0.108	78825	5600	110075
ODC TFT/Rpn4 TFT/N-end Rule (8)	VIIa	9.24	-0.149	125200	91475	166375
ODC TFT/N-end Rule (10)	VIIb	16.21	0.173	409450	389250	436750
Rpn4 TFT/N-end Rule (3)	VIIc	7.42	-0.115	873575	839950	903725
ODC TFT/N-end Rule (3)	X	11.19	0.141	626100	588400	660250
ODC TFT/Rpn4 TFT/N-end Rule (5)	XII	10.21	0.113	665600	636800	690250
ODC TFT/N-end Rule (2)	XIIIa	16.95	0.170	47800	25200	75850
ODC TFT/N-end Rule (1)	XIIIb	15.345	0.1573	341525	366875	394975
ODC TFT/N-end Rule (9)	XIVa	32.50	-0.226	464275	448675	478825
Rpn4 TFT/N-end Rule (8)	XV	18.30	-0.176	163350	138600	186200

Supplementary Table 1. *Overlap of proteasome activity QTLs with UPS N-end rule QTLs. Numbers in parentheses in the “Reporters” column indicate how many of 20 possible UPS N-end Rule reporters the corresponding proteasome activity QTL was detected for. Numeric columns display the median value calculated from the set of proteasome activity and UPS N-end Rule reporters for which the QTL was detected. “Peak Position”, “Left Index”, and “Right Index” refer to base pair positions on the indicated chromosome.*