Translation error clusters induced by aminoglycoside antibiotics

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Aminoglycoside antibiotics target the ribosome and induce mistranslation, yet which translation errors induce bacterial cell death is unclear. The analysis of cellular proteins by quantitative mass spectrometry shows that bactericidal aminoglycosides induce not only single translation errors, but also clusters of errors in full-length proteins in vivo with as many as four amino acid substitutions in a row. The downstream errors in a cluster are up to 10,000-fold more frequent than the first error and independent of the intracellular aminoglycoside concentration. The prevalence, length, and composition of error clusters depend not only on the misreading propensity of a given aminoglycoside, but also on its ability to inhibit ribosome translocation along the mRNA. Error clusters constitute a distinct class of misreading events in vivo that may provide the predominant source of proteotoxic stress at low aminoglycoside concentration, which is particularly important for the autocatalytic uptake of the drugs.
minoglycoside antibiotics (AGAs) are positively charged oligosaccharides that target bacterial ribosomes. AGAs induce misreading of the mRNA, resulting in the incorporation of incorrect amino acids into protein, and/or inhibit translocation, the movement of the ribosome along the mRNA. Many AGAs, such as paromomycin (Par), tobramycin (Tob), neomycin (Neo), gentamicin (Gen), kanamycin (Kan), or amikacin (Amk), are clinically important as broad-spectrum antibiotics. On the other hand, their propensity to induce translation errors can be utilized to alleviate the symptoms of human genetic diseases (e.g., Duchenne muscular dystrophy) by increasing readthrough of premature stop codons. The negative side effect of AGA treatment is its unfavorable effect on translation in mitochondria. In particular, the high oto- and nephrotoxicity due to targeting of mitoribosomes often hampers systemic administration of AGAs.

Although misreading is the key element of the AGA action, we know surprisingly little which types of errors are induced and how they affect the proteome and the fitness of bacterial cells. AGAs preferentially induce a subset of errors that result from a misreading of the third position of the mRNA codon, but errors in the first and second position can also occur. The type of most prevalent errors depends on the drug, e.g., streptomycin (Str) is less efficient than other AGAs in inducing U-G mismatches in the second position or G-U mismatches at any position of the codon–anticodon complex. Severe misreading leads to growth arrest or cell death in bacteria. However, the AGA-induced error load alone cannot explain their bactericidal effects. In fact, ribosomal ambiguity (ram) mutants, with alterations in ribosomal protein S4, also cause increased misreading, but grow at almost wild-type (wt) rates. This led to the notion that there is something particular to the bactericidal effect of AGAs, but what causes their detrimental effect on bacterial fitness and cell viability remains unclear.

AGAs are polyatomic molecules that bind to the negatively charged outer membrane of bacteria and enter the periplasm via porin channels. Once in the cytosol, AGAs bind to the ribosome and induce misreading by stabilizing an error-prone conformation of the decoding center of the ribosome. Accumulation of translation errors in membrane proteins leads to the disintegration of membrane structures, renders the membrane permeable for small molecules, and allows for a massive influx of AGAs. However, how sub-lethal intracellular AGA concentrations and the associated mild increase in mistranslation cause damage of membrane proteins is unknown. At bactericidal concentrations, proteotoxic stress induces the heat shock response, aggregation of specific sub-proteomes, including membrane and metabolic proteins, protein oxidation and carboxylation, and inclusion body formation. Further downstream effects of AGA treatment are dramatic changes of the cell metabolism, oxidative stress, DNA damage, and ultimately cell death. In contrast, sub-lethal AGA concentrations may even be beneficial for bacteria, as they help cells to adapt to stress conditions, change to a more drug-resistant lifestyle (e.g., biofilm formation), or acquire antibiotics resistance.

The majority of AGAs affect both decoding and translocation steps of translation. The efficacy of AGAs is often attributed to their ability to induce misreading, rather than to their effect on translocation. In fact, Str, which is bactericidal and induces misreading, has no effect on translocation. Vice versa, spectinomycin (Spc), an AGA that causes a translocation defect without inducing misreading, is bacteriostatic. One potential exception is apramycin (Apr) which is bactericidal, although recent studies suggested that it does not induce misreading. This is important, because Apr is less ototoxic than other AGAs in model organisms, presumably because its antibacterial and anti-mitoribosomal activities are uncoupled. Moreover, due to its unique structure, Apr is not susceptible to many prevalent resistance mechanisms and is thus a promising drug for the treatment of multiresistant bacteria.

Somewhat paradoxically, the bactericidal effect of AGAs is lost when a translation is blocked by bacteriostatic antibiotics, such as chloramphenicol.

Here we used quantitative mass spectrometry to correlate the AGA-dependent cell growth inhibition, miscoding burden, and stress responses, and to obtain insights into the bactericidal effects of AGAs. We found that AGA binding induces not only single errors, but clusters of errors with two, three or four amino acid substitutions located close to each other in the protein sequence. We characterize this type of misreading events for a variety of AGAs and show that their prevalence depends not only on the misreading propensity of a given AGA but also on its ability to inhibit translocation. We also show that Apr, an AGA which is thought not to cause misreading, induces frequent translation errors. These results reveal an additional, unexpected aspect of AGA action in bacteria, suggest how antibiotics cause proteotoxic stress, and provide a simple explanation as to why some misreading-inducing antibiotics are bactericidal.

**Results**

**AGA-induced missense errors and cellular fitness.** We first asked what causes the strong detrimental effect of AGAs on bacterial cell viability. To distinguish the AGA-specific response from general stress effects caused by errors, we compared the error load and fitness of *Escherichia coli* cells after AGA treatment with those of untreated *ram* (ribosomal ambiguity) *E. coli* cells that are error-prone due to a C-terminal truncation in ribosomal protein S4. *Ram* cells are viable and have near-wt growth rates with only ~30% longer doubling time. For the initial experiments, we used Str, because it does not impede translation. Likewise, the translation rate in *ram* cells is not affected, which allows us to attribute potential fitness defects to errors in drug levels. Cell viability was estimated using changes in cell density. Proteotoxic stress was assessed by monitoring proteome responses and translational fidelity was determined by data-dependent and targeted mass spectrometry. Hierarchical clustering shows that Str treatment induces the heat-shock response (Supplementary Fig. 1a–e), in agreement with previous reports. The upregulation of heat-shock proteins, such as chaperones (e.g., GroEL or DnaK) and proteases (e.g., Lon), is indicative of protein misfolding. The small chaperones IbpA and IbpB, which guide unfolded proteins into protein aggregates for their subsequent disaggregation, are induced at higher Str concentrations compared to other heat-shock proteins, and their accumulation coincides with the onset of growth inhibition. Also, *ram* cells have increased levels of heat-shock proteins, compared to the untreated *wt* cells, but the response is mild and comparable to that in *wt* cells treated with Str at sub-lethal concentrations, about 2–4 μM (Fig. 1a; dashed line).

To determine the error frequency (*E*) in the same samples, we analyzed translation errors by data-dependent acquisition (DDA) mass spectrometry. We first searched for errors in peptides from elongation factor Tu (EF-Tu), which was chosen because it is constitutively expressed and highly abundant in the *E. coli* lysate, thus allowing us to detect rare misreading events. We quantified peptides that carry single amino acid substitutions and calculated *E* as a ratio of an incorrect peptide to the corresponding correct parental peptide (Supplementary Fig. 1g). As expected, there is a correlation between Str-induced translation errors and the stress response (Fig. 1b). At 2–4 μM Str, *E* in the Str-treated cells is lower than in the *ram* cells, whereas the heat-shock response is similar. At 6–8 μM Str, the error load becomes similar, but at a
cost of a much stronger heat-shock response and lower fitness in the Str-treated cells. Notably, some *ram* strains have even higher error loads than the one used in our study. Thus, translation errors per se do not explain the strong stress response and growth inhibition of Str-treated cells.

We next asked why similar apparent error loads that are relatively harmless for *ram* cells are proteotoxic at sub-lethal Str concentrations in *wt* cells. To some extent, this can be a result of the adaptation of *ram* cells to high miscoding levels, which we cannot exclude (Supplementary Fig. 1f). Alternatively, the differences might be attributed to the type of errors, e.g., some types of amino acid substitutions may result in preferential misfolding and aggregation, whereas others may be neutral. Indeed, certain types of amino acid substitutions (e.g., Y → H, G → C, I → F, G → R) are more abundant in Str-treated cells, whereas *ram* cells have more errors involving G-U mismatches in the codon-anticodon complex, such as R → H or E → K substitutions (Fig. 1c). To test whether Str-induced errors result in preferential protein aggregation, we analyzed peptides in aggregated vs. soluble EF-Tu fraction and found that the protein in aggregates indeed contains more errors (Fig. 1d). We then classified the errors as either Str-specific, which we define as more than fivefold more abundant in Str-treated than in the *ram* cells, or common errors, and quantified the distribution between soluble and aggregated fractions for each of the two groups. The error enrichment in protein aggregates is similar for Str-specific and common errors (Fig. 1e); hence, Str-specific errors per se do not cause excessive protein aggregation. However, we note that there is a subgroup of Str-specific errors that appear only at high Str concentrations and are preferentially found in aggregates (Fig. 1d–f and Supplementary Fig. 2a). These errors tend to destabilize proteins, as evident from their reduced thermal stability in vitro (Fig. 1g). Moreover, they become abundant concomitantly with the growth arrest and upregulation of IbpA and IbpB, which indicates an overload of the quality control machinery (Supplementary Fig. 2b, c). Thus, while these destabilizing substitutions start to accumulate at conditions where cells lose viability, they cannot account for the proteotoxic effects seen at sub-lethal AGA concentrations. To understand which types of errors cause early stress responses, we then analyzed the Str-induced missense peptides in greater depth and, surprisingly, identified peptides that had more than one amino acid substitution in the sequence. As such error clusters have not been described before and their appearance might explain the enhanced proteotoxicity of AGAs, in the following we focus our analysis on this particular type of AGA-induced errors.

**Error clusters as a hallmark of AGA action.** To identify further peptides with two or more amino acid substitutions, we first searched the Str-treated samples for peptides with more than one amino acid substitution by DDA (Fig. 2a and Supplementary Fig. 4a). Next,
we predicted potential clusters based on the experimentally measured frequencies of single amino acid substitutions ('Methods'). Guided by these predictions, we enriched the expected candidate peptides by QRAS, a method developed for the Quantiﬁcation of Rare Amino acid Substitutions10. QRAS relies on the chromatographic enrichment of missense peptides using synthetic reference peptides as standards, resulting in the improvement of the signal-to-noise ratio and high conﬁdence identiﬁcations. We found a large number of peptides with two amino acid substitutions, which were either adjacent or interspersed by several correct amino acids (Supplementary Fig. 4b). Error clusters appear only in Str-treated samples and are absent in ram cells (Fig. 2b). Error clusters are enriched in aggregates, but the effect depends on the type of errors (Fig. 2c). While in...
Fig. 2 Error clusters as a hallmark of AGA action.  

**a** EF-Tu-derived peptides with one (gray) or two (red) errors in Str-treated wt cells and in untreated ram cells. Shown are means of three technical replicates (n = 3). **b** Error clusters induced by Str are absent in the ram strain. Str concentration (4 \( \mu \text{M} \)) was chosen to achieve the same \( E_f \) in wt and ram cells. Peptides with error clusters were detected by QRAS and SRM (dtop = 1, fragment ions color coded).

**c** Enrichment of error clusters in aggregates. After treating cells with Str (8 \( \mu \text{M} \)), EF-Tu was isolated from the soluble cell fraction and the insoluble aggregates and peptides analyzed by DDA. Shown are the means of three technical replicates (n = 3). **d** Time course (left panel) and concentration dependence (right panel) of Str treatment for D208E-F211L error cluster detected by PRM. Shown are means ± SD of three technical replicates (n = 3).

**e** Str-induced error clusters with three and four substitutions. Target peptides were enriched by QRAS and detected by PRM. **f** Examples of PRM validation for error clusters in different proteins induced by Str (8 \( \mu \text{M} \)). **g** Types of misreading events in error clusters. Errors with ambiguous classification of the misreading position are shown in gray; in cases where misreading can arise from mismatches at different positions, the errors were classified as originating from the 3rd > 1st > 2nd position mismatch, according to their response to AGA treatment. **h** Effect of different AGAs. The error cluster D208E-F211L in EF-Tu detected by PRM upon treatment with spectinomycin (Spc), streptomycin (Str), dihydrostreptomyacin (DH5), paromomycin (Par), apramycin (Apr), sisomycin (Sis), gentamicin (Gen), gentamicin (G418), hygromycin B (HygB), neomycin (Neo), tobramycin (Tob), kanamycin A (KanA), kanamycin B (KanB), amikacin (Amk), or neamine (Nea) at concentrations that gave rise to maximum misreading (‘Methods’). Shown are means ± SD of three technical replicates (n = 3). See also Supplementary Figs. 1-3 and Source data file.

many cases both single errors and their combinations are well tolerated (e.g., D208E and F211L, Fig. 1f, g, and D208E-F211L, Fig. 2c (blue symbols)), in other cases the increased error load leads to enhanced aggregation (Fig. 2c, red symbols).

Although the prevalence of error clusters depends on Str concentration and treatment duration, they are clearly detectable already after short incubation times and at low, sub-lethal Str concentrations, which shows that their occurrence is not related to cell death (Fig. 2d). Moreover, we also found error clusters with three or four amino acid substitutions in one peptide sequence (Fig. 2e and Supplementary Fig. 4b). To ensure that error clusters are prevalent in the proteome, we also analyzed different full-length proteins in the *E. coli* lysate (Fig. 2f and Supplementary Fig. 3a). Overall, we validated 96 error clusters with errors originating from mismatches in the first, second, or third position of the codon–anticodon complex (Fig. 2g, Source data file).

To study whether error cluster formation is caused by AGAs other than Str, we used AGAs from different structural classes (Fig. 2h). All bactericidal AGAs tested give rise to error clusters; spectinomycin (Spc), which is bacteriostatic, does not induce error clusters. Surprisingly, also Apr, which was reported not to cause significant misreading, induces single errors and error clusters. The frequency of error clusters (\( E_f^{\text{cluster}} \)) is in the same range as \( E_f \) for most single amino acid substitutions. Overall, this shows that the appearance of error clusters is a hallmark of the action of bactericidal AGAs.

**Apr effect on misreading and translation.** The surprising finding that Apr induces errors prompted us to study the effects of Apr on misreading and translation in more detail. We validated that all Apr batches purchased from different vendors induced significant misreading (Supplementary Fig. 5a). Most errors that are induced by Str are induced by Apr as well, including the error clusters (Fig. 3a, b, and Supplementary Fig. 5b). Str and Apr induce very similar error loads and proteotoxic stress response at concentrations that inhibit growth (Fig. 3c, d). To better understand why recent in-vitro studies came to a different conclusion, we tested how Apr affects protein synthesis in a fully reconstituted translation system. As mRNAs we used a natural mRNA coding for the model protein SlyD and a homopolymeric mRNA (poly(U)) coding for polyphenylalanine. At Apr concentrations of 0.1–1 µM translation is impaired (Fig. 3e–g). For SlyD, both the end level (Fig. 3e and Supplementary Fig. 5c) and the rate (Fig. 3f, g and Supplementary Fig. 5c) of translation are affected. The very low translation efficiency, which was also noted in the previous reports, might have masked the occurrence of the misreading events in those experiments. Apr also reduces the rate of poly(U) translation with Phe-tRNA^Phe^ (Fig. 3g and Supplementary Fig. 5d), i.e., at conditions where no misreading can occur due to the lack of near-cognate tRNA, supporting the suggested effect of Apr on translation (the rate of peptide bond formation is not affected by the AGA).

**Probability of making successive errors.** Given that all bactericidal AGAs including Apr induce error clusters, we next asked whether consecutive errors in the same peptide can be explained by their stochastic occurrence. For errors that occur stochastically and are independent of each other, the expected \( E_f^{\text{cluster}} \) is a product of individual \( E_f \) values for each position, which would be \( <10^{-6}-10^{-5} \) in most cases. In contrast, the observed error clusters can be almost as frequent as most single amino acid substitutions, indicating that consecutive misreading events occur non-randomly (Figs. 3a, 3b). To estimate how non-random these events are, we determined by QRAS the \( E_f^{\text{cluster}} \) values for clusters with two or three errors, as well as the \( E_f \) values for the respective single amino acid substitutions, and compared the experimental and the estimated stochastic values (Fig. 3a and Supplementary Fig. 4c). The measured error frequency for clusters is up to 60,000-fold higher than that predicted for a combination of random misreading events at the same positions, largely independent of the detection method (SRM, PRM vs. DDA) or the protein (EF-Tu or other model proteins, Supplementary Fig. 3b). Due to the vectorial nature of translation, the appearance of the first error is independent of all subsequent misreading events, which implies that after the first error is made, all subsequent errors are much more frequent than expected. Analysis of the Str data shows that the \( E_f^{\text{cluster}} \) value depends linearly on the \( E_f \) value of the first misreading event (\( E_f^{\text{ist}} \)) and the slope of the plot is independent of the type of experiment (Fig. 4b). The slope of 0.08 in the example of Fig. 4b provides an estimate for the error frequency of the second misreading event, which we denote as \( E_f^{\text{ext}} \), \( E_f^{\text{ist}} \times E_f^{\text{ext}} = E_f^{\text{cluster}} \). For a given type of error clusters, \( E_f^{\text{ext}} \) values are largely independent of the protein tested and the position of the cluster in the protein sequence (Supplementary Fig. 3c). \( E_f^{\text{ext}} \) for the second and third substitutions are very similar (Fig. 4c). The propensity to make consecutive errors depends on the AGA, e.g., for a given peptide the \( E_f^{\text{ext}} \) is 0.09 and 0.05 for Par and Str, respectively, but is much higher, 0.25, for Apr (Fig. 4d). The \( E_f^{\text{ext}} \) values provide a lower-limit estimate for the intrinsic error frequency of decoding on those ribosomes
where an AGA is bound, i.e., at conditions where misreading is independent of the intracellular concentration of an AGA.

AGAs preferentially induce misreading in the third codon position\(^9\)-\(^11\). To test whether this tendency applies to consecutive errors, we inspected misreading preferences in clusters for a fixed amino acid substitution, such as F211L or D208E, both likely induced by third-position misreading. Indeed, the majority of misreading events in the clusters can be explained by third-position misreading, whereas substitutions due to the second position mismatches are clearly less abundant (Fig. 4e and Supplementary Fig. 6a, b). If a particular error is frequent as a single error, it is also frequent as an error in the cluster (Fig. 4f), suggesting that the preference for a particular type of misreading is conserved in error clusters.

### Distance dependence of successive errors

Finally, we determined the frequency of successive errors \(E_f^{\text{next}}\) in different error clusters for different AGAs (Figs. 4d, 5a, and Supplementary Fig. 7a). \(E_f^{\text{next}}\) with Str or Par shows only slight variation, \(E_f^{\text{next}} = 0.04-0.18\) and \(0.02-0.09\), respectively. However, with Apr the \(E_f^{\text{next}}\) value is highly variable, as in some cases no error clusters are detected, whereas in other cases \(E_f^{\text{next}}\) reaches 0.25. We then noticed that with Str the frequency of the successive error is almost independent of the distance between the incorrect amino acids in a peptide. In contrast, with Apr the probability of the next misreading event decreases dramatically with distance; with Par the dependence is very weak and the \(E_f^{\text{cluster}}\) value for the two distant misreading events is close to the stochastic one (Fig. 5a, b, left panel). We then analyzed the distance dependence of error

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**Fig. 3 Apramycin induces proteotoxic misreading and inhibits translocation.** a Single amino acid substitutions induced by Apr and Str. Errors induced predominantly by Apr (16 \(\mu\)M) are shown in cold colors; those induced by Str (8 \(\mu\)M) in warm colors. Shown are means ± SD of three technical replicates (\(n=3\)). b Error clusters in Apr-treated cells (red) compared to single errors (gray). Shown are means ± SD of three technical replicates (\(n=3\)). c Induction of the heat shock response. The chaperones IbpA/IbpB are significantly induced upon Apr (16 \(\mu\)M) and Str (8 \(\mu\)M) treatment as quantified by SRM. Shown are means ± SD of three technical replicates (\(n=3\)). d Induction of the heat shock response. The chaperones IbpA/IbpB are significantly induced upon Apr (16 \(\mu\)M) and Str (8 \(\mu\)M) treatment as quantified by SRM. Shown are means ± SD of three technical replicates (\(n=3\)). e Products of the in-vitro translation of SlyD mRNA. Translation products are separated by SDS PAGE and detected using the fluorescence of the N-terminal BODIPY-Met. f Time courses of in-vitro translation of SlyD mRNA at different Apr concentrations. Translation products were detected as in (e). g Rate of in-vitro translation (aa/s) at different Apr concentrations for SlyD mRNA (white bars) and poly(U) mRNA (gray bars). Shown are means ± SD of three technical replicates (\(n=3\)). See also Supplementary Fig. 5 and Source data file.
Misreading preferences in error clusters. Str-induced error clusters were quantified by PRM and LFQ. Str titration of the distance dependencies suggests that the AGA dissociates from the ribosome within several rounds of elongation. The most likely explanation for the distance dependence is that the AGA decreases with distance (Fig. 5b, middle panel). The most likely explanation for the distance dependence is that the AGA dissociates from the ribosome within several rounds of elongation. Exponential fitting of the distance dependencies suggests that the probability to continue translation with the AGA still bound is about 0.3–0.7 for the cases shown in Fig. 5, middle panel, suggesting that the rates of translation and AGA drop-off are comparable. In contrast, with other AGAs, such as Par, Str, dihydrostreptomycin (DHS), and neamine (Nea), there is very little distance dependence (Fig. 5b, right panel). A replot of $E_{\text{next}}$ vs. $E_{\text{1st}}$ (Fig. 5c) suggests that there are two groups of AGAs, those that induce moderate misreading for single amino acid substitutions, but once bound—induce frequent local consecutive errors (Apr, Gen, Kan, Sis, Tob), and those that have a strong misreading effect on a single position, but are less efficient in inducing error clusters (DHS, Nea, Par, Str). These data show

Fig. 4 Frequencies of consecutive errors. a Absolute quantification of error clusters by QRAS. Left panels: $E_f$ of one, two, or three amino acid substitutions after AGA treatment. Dash is a median value shown as a visual guide. Right panels: Error amplification over stochastic level. Shown are means ± SD of at least three independent biological replicates (n = 3). b Correlation between $E_{\text{1st}}$ and $E_{\text{cluster}}$. Peptides were quantified by PRM and LFQ assuming identical ionization and fragmentation properties. Shown are means ± SD of three technical replicates (n = 3). c $E_{\text{next}}$ for the 2nd and 3rd errors in the cluster. $E_{\text{next}}$ is calculated as $E_{\text{next}} = \frac{E_{\text{next}}}{E_{\text{1st}} \times E_{\text{2nd}}}$, where $E_{\text{1st}} \times E_{\text{2nd}}$ represents the expected stochastic occurrence of error cluster. Shown are means ± SD of at least three independent biological replicates (n = 3). d Data for the probability of single errors is derived from Supplementary Fig. 1g; the probability to occur in error cluster is based on Fig. 4e and Supplementary Fig. 6. Pearson correlation coefficients as insets. See also Supplementary Fig. 6 and Source data file.
that, although all AGAs that induce misreading can induce clusters of missense errors, the prevalence of such errors and the distance between consecutive errors in a cluster depends on the nature of the AGA.

**Discussion**

The analysis of AGA-induced translation errors in cellular proteins by mass spectrometry reveals the existence of prevalent clusters of translation errors, which is a distinct class of misreading events in vivo and likely explains why most AGAs are bactericidal. The comparison of AGA-induced errors in *wt E. coli* cells with those in an error-prone *ram* strain shows that the error frequency or the type of errors alone do not account for the strong stress response and fitness loss upon mild AGA treatment. Although single errors induced at low AGA concentrations do not cause excessive protein aggregation, heat-shock proteins are upregulated to a much higher extent than in the *ram* strain, which has a much higher error level, but shows only a moderate heat-shock response. At high AGA concentrations, the high error load results in the accumulation of destabilized protein variants, formation of protein aggregates, upregulation of the chaperones IbpA and IbpB, and a concomitant loss of cell viability. This “kiss-of-death” response is much more dramatic than a mild heat-shock response detected in the *ram* cells at a comparable error load. These observations show that AGA causes specific defects that induce an early heat-shock response and compromise cell fitness. Error clusters with multiple translation errors appear early upon AGA treatment and at low AGA concentrations, but are not found in the *ram* proteome. Regardless of the frequency of the first error in the string, all subsequent errors are much more abundant than if they were stochastic. Furthermore, the probabilities of the second and third errors in a cluster are very similar. These two observations strongly suggest that the AGA binds to the ribosome over a number of elongation cycles, thereby inducing consecutive errors. This increases the misreading potential at low AGA concentrations by orders of magnitude and makes all the downstream errors in a cluster independent of the AGA concentration. Error clusters can thus constitute the predominant source of proteotoxic effects, in particular at low AGA concentrations or in the initial phase of AGA treatment where the median error frequency for single errors is low. Moreover, our data indicate that proteins with error clusters are somewhat more prone to aggregation than those with single errors. While the effect of multiple amino acid substitutions on stability is often additive, the combinatorial effect on cellular fitness is often stronger, which is known as negative epistasis. The threshold model explains this behavior assuming that first mutations are buffered by the excess of stability, whereas subsequent mutations reduce the stability beyond the allowed stability margin.

The error clusters described here are fundamentally different from the spontaneous consecutive misreading events that can occur in the absence of the AGA and cause premature translation termination. Termination produces incomplete proteins that are likely to be degraded by the cellular quality control machinery. Peptides with adjacent amino acid substitutions at the C terminus that likely represent the premature termination products were recently detected by mass spectrometry upon translation of model constructs. In contrast to such spontaneous misreading events, AGA-induced error clusters are found in full-length proteins, although we cannot exclude that a fraction of nascent peptides with two adjacent errors is degraded via the termination editing pathway. Furthermore, AGA-induced translation errors are not necessarily sequential; rather, individual misreading events can be separated by as much as 13 correct
The AGA-induced error clusters are induced by all bactericidal AGAs tested, underscoring the importance of this phenomenon for understanding the mode of AGA action.

We provide the first characterization of error clusters induced by different AGAs. In general, they emerge already at low AGA concentrations and early after the beginning of the AGA treatment. Although the probability to make the first error varies with the nature of the AGA, its concentration, and the time of treatment, the probability of a subsequent error depends only on the nature of the AGA. The preference for a particular type of misreading event (e.g., a third codon position mismatch) is common for the first and subsequent substitutions. These observations suggest that the propensity of the second and subsequent errors depends on the error-prone conformation of the ribosome and the residence time of the particular AGA on the ribosome once it is bound.

Based on their efficacy in inducing single and multiple errors, bactericidal AGAs can be subdivided into two groups, (I) those that are potent misreading agents for single substitutions, but have a moderate efficacy in inducing error clusters, and (II) those that have a moderate effect on single errors, but induce very strong misreading in a narrowly spaced cluster sequence. One characteristic example of the group I AGA is Str, which is a potent misreading agent that does not affect translocation. Analysis of the error clusters suggests that Str remains bound to the ribosome for at least ten elongation cycles, thereby inducing long error clusters with a relatively moderate error frequency for each consecutive error (Fig. 6). The error frequency of consecutive errors in the cluster reaches 0.18, which is close to the maximum misreading value of 0.25 measured in vitro at saturating Str concentration. Nea, which also belongs to group I, is inefficient in inducing error clusters probably because it inhibits translocation and dissociates rapidly from the ribosome.

The majority of AGAs belong to group II. They induce a medium-low error frequency for the first position in a cluster, but the frequency of the next error is very high, in particular for KanA and Sis, reaching 0.36 and 0.40, respectively, which means that when the antibiotic is bound, up to 36% or 40% of the ribosomes can misread the next codon. These AGAs affect translocation in addition to misreading. The rate of translocation on the ribosomes that have an AGA bound is comparable to that of the AGA dissociation, which results in a gradual loss of the AGA with each elongation cycle and explains the strong distance dependence of the error frequency in the cluster (Fig. 6). This is consistent with in-vitro single-molecule FRET data showing that AGAs can stay on the ribosome for several elongation cycles. Together, these results indicate that the rates of translocation in the presence of AGA and of AGA dissociation, as well as the ribosome conformation induced by AGA binding, define the prevalence and the size of error clusters. Apr induces pronounced error hot spots, whereas Str introduces milder, but far longer error clusters, some of which are likely to supersede the length of average tryptic peptides.

The notion that misreading alone does not explain the bactericidal effects of AGA has important clinical implications. Noteworthy, most of the global protein synthesis inhibitors are bacteriostatic. For example, chloramphenicol and viomycin induce translation errors, but are bacteriostatic. This is because after binding the antibiotic, the ribosome that made a mistake stops translation. In fact, peptides with an incorrect C-terminal amino acid were identified by mass spectrometry after chloramphenicol treatment of E. coli cells. The appearance of such aberrant protein fragments, which may be removed by the quality control machinery, may be less harmful for the cell than the accumulation of full-length proteins with error clusters. Similarly, chloramphenicol rescues the bactericidal effect of AGAs probably because the inhibition of translation by chloramphenicol would preclude the AGA-induced formation of error clusters and thus avoid synthesizing aberrant proteins.

**Fig. 6 Model of AGA-induced error cluster formation.** The efficacy of an AGA in inducing proteotoxic error clusters depends on its ability to bind and induce the 1st error, its residence time on the ribosome, and the inhibitory effect on translocation. Str, which does not affect translocation, and its derivative DHS, induce frequent single errors and remain bound to the ribosome for many elongation cycles, resulting in formation of long error clusters. AGAs that have a moderate inhibitory effect on translocation induce short clusters, because they tend to dissociate after only a few slow translation rounds. Nea has a high misreading potential, but dissociates rapidly from the ribosome, resulting in low levels of presumably stochastic error clusters which are thus not distance dependence. Viomycin stops translocation which terminates translation by ribosomes that have the antibiotic bound and hence prevents formation of the error clusters.
The balance between effects on decoding and translocation might be species-dependent\(^6\). Bacteriostatic drugs might become bactericidal in some organisms if the residual translocation speed in the presence of the antibiotics is higher due to alterations in the ribosome structure. For Apr, its strong effect on misreading and translocation, combined with specific features of AGA in mitochondria, might explain why Apr is less oto- and nephrotoxic than other AGAs.

In summary, our results suggest that bactericidal AGAs can stay bound to the ribosome for several elongation cycles resulting in a high probability of multiple consecutive misreading events. While the error frequency of single amino acid substitutions depends mostly on the stochastic binding of AGA to the ribosome, subsequent errors are determined (i) by the propensity of the AGA-induced conformation of the ribosome to incorporate near-cognate amino acids and (ii) on the residence time of the AGA on the ribosome compared to the translocation rate. In a sequence context where many codons allow for third position mismatches, consecutive errors result in aberrant proteins whose abundance is much higher than expected from the stochastic combination of individual error frequencies at low AGA concentrations. This potentiation of the misreading effect for consecutive errors in the cluster and the higher likelihood of error clusters to induce aggregation would explain how very low concentrations of AGAs initiate the autocatalytic uptake of the drugs, presumably by inducing the formation of defective membrane proteins. Moreover, the aminoglycoside-induced synthesis of highly aberrant aggregation-prone proteins could explain how bacteria sense proteotoxic stress at low aminoglycoside concentrations to develop drug resistance or communicate with other bacteria in the population\(^31\,62\). Future studies will reveal to which extent and which error clusters lead to the precipitation of specific subproteomes, destabilize the membrane integrity in the initial phase of AGA uptake, or promote mitochondrial proteotoxic stress in eukaryotes. Notably, such errors might contribute to side effects when AGAs are used to suppress premature stop codons in humans. As a distinct class of AGA-induced responses, error clusters are of major importance for the understanding of AGA-induced responses in humans. As a distinct class of AGA-induced responses, error clusters are of major importance for the understanding of AGA-induced responses in humans.
Trypsin (Trypsin Gold, Promega) was added to the protein at a substrate/protein ratio of ~200:1, and EF-Tu was proteolyzed for 16 h at room temperature. Tryptic peptides were identified by centrifugation through a 30-kDa cut-off filter and the concentrate was washed with H2O. Samples from both elution steps were pooled.

For proteolytic EF-Tu prepared under native conditions, 3–100 nmol of EF-Tu was precipitated overnight with five volumes of ice-cold acetone at −20 °C. Protein was collected by centrifugation, washed with ice-cold 80% ethanol, and the pellet dried. EF-Tu was reconstituted in 1% Rapigest (in Water) at 25 mM NH4HCO3, and incubated for 10 min at 37 °C. Disulfide bonds were reduced by addition of 20 mM DTT (in 25 mM NH4HCO3) in two incubation steps, at 60 °C for 10 min, and at 37 °C for 20 min. Alkylation of thiols was performed in 30 mM iodoacetamide (in NH4HCO3) and incubating the sample at RT for 30 min in the dark. Rapigest (in Water) in the sample was diluted to 0.1% with 25 mM NH4HCO3. Trypsin (1 µg/µl) (Trypsin Gold, Promega) was added to the sample (final concentration 0.01 µg/µl) and EF-Tu proteolyzed overnight at 37 °C. In-gel proteolysis was carried out according to the standard protocol.

Proteome analysis. Cells (1 OD660) were opened in SDS loading buffer. Proteins (corresponding to 0.1 OD660 of cells) were desalted by a 10% Mini-PROTEAN TGX StainFree PAGE (GE Healthcare). Proteins were stained with Coomassie and in-gel proteolysed with trypsin.

Samples were analyzed on an Ultimate 3000 RSLC system coupled to a QExactive HF Quadrupole-Orbitrap mass spectrometer. Tryptic peptides were loaded on a C18 precolumn (2.5 cm length, 150 µm inner diameter, Reprosil-Pur 120 Å, 5 µm (Dr Maisch GmbH, Germany)). Bound peptides were eluted and separated on a C18 capillary column (31 cm, 75 µm ID, Reprosil-Pur C18-AQ 120 Å C18, 1.9 µm) at a flow rate of 300 nL/min, with a 45 min linear gradient from 5 to 45% acetonitrile (ACN) in 0.1% formic acid. Analysis was performed in positive ion mode using a 30 DDA method. MS survey spectra were acquired at a resolution setting of 60,000 at 400 m/z (Full Width at Half Maximum) in the range of 350–1600 m/z. Automatic gain control (AGC) target values and maximum injection times for MS runs were 1e6 and 50 ms. Precursors with charge states z > 2–5 above the intensity threshold of 1.2 e4 were selected at an isolation width of 1.6 m/z for fragmentation by higher energy collision dissociation (HCD) with a normalized collision energy setting of 35%. Ions of unassigned charge states were excluded from fragmentation selection, and the masses of fragmented precursors were dynamically excluded for 25 s. MS/MS transients were acquired at a resolution setting of 15,000 FWHM using AGC target values and maximum injection of 1e4 and 50 ms, respectively.

Data were processed using MaxQuant software (version 1.6.5.0).66 Technical replicates (n = 3) were merged, while biological replicates were kept separate for statistical analysis. If not mentioned otherwise, standard software settings were kept. The database contained all proteins of the E. coli proteome (MG1655, uniprot database) annotated with known lab contaminants. Carbamidomethylation was used as a fixed modification, and methionine oxidation and N-terminal acetylation as variable modifications. Identifications were matched between the runs and were filtered using a target decoy approach at a false discovery rate of 0.01. Biological replicates were compared by LFQ using the MaxQuant LFQ algorithm.67 To identify Str-induced changes in the proteome the LFQ values were further analyzed in Perseus (version 1.6.5.0) removing peptides with a presence of < 30% and a cut-off filter was applied and only proteins that were found in ≥60% of all runs were kept for analysis. Missing values were replaced(931,943),(999,992) by random numbers that are drawn from a normal distribution which is modeled on the distribution of the experimental data (default values: width 0.3, downshift 1.8). Significantly altered proteins were identified applying a permutation-based ANOVA statistic applied to a permutation of the FDR (50 = 4; FDR 0.05, 250 randomizations). Intensities of significantly regulated proteins were normalized to their z-score and biological replicates were averaged. Co-regulated proteins were identified by hierarchical clustering according to their Euclidian distance. The enrichment of biological processes in the individual clusters was determined by applying Fisher Exact test. One cluster contained proteins, which were significantly upregulated with increasing Str concentrations comprising chaperones and proteases of the heat-shock response. The LFQ-values of these co-regulated chaperones and proteases were scaled to an interval (0–1).

Quantification of peptides with single amino acid substitutions by DDA. Cells (1 OD600) were opened in an SDS loading buffer. Proteins (corresponding to 0.1 OD600 cells) were separated by 10% Criterion TGX SDS PAGE (BIORAD). Protein bands of interest were excised from the gel, and the proteins were in-gel proteolysed with trypsin. Samples were analyzed on an Ultimate 3000 RSLC system coupled to a QExactive HF-X quadrupole Orbitrap mass spectrometer. Tryptic peptides were loaded on a C18 precolumn (2.5 cm, 150 µm ID, Reprosil-Pur 120 Å, 5 µm). Bound peptides were eluted and separated on a C18 capillary column (31 cm, 75 µm ID, packed with Reprosil-Pur 120 Å, 1.9 µm) at a flow rate of 300 nL/min, with a 76 min linear gradient from 5 to 42% ACN in 0.1% formic acid.

Acquisition was performed using two acquisition schemes to maximize identifications while keeping consistent quantifications. Quantification runs were performed in positive ion mode using a ten top DDA method with two micro scans per MS spectrum. MS survey spectra were acquired at a resolution setting of 120,000 FWHM in the range of 350–1600 m/z, using AGC target values and maximum injection of 3e6 and 100 ms, respectively. Precursors with charge states z = 2–7 above threshold intensity of 3.0 e4 were selected at an isolation width of 1 m/z. Precursors were isolated using a combination of a DDA cut-off filter and a normalized collision setting of 35%. Ions of unassigned charge states were excluded from fragmentation selection, and the masses of fragmented precursors were dynamically excluded for 15 s. MS/MS transients were acquired at a resolution setting of 15,000 FWHM using AGC target values and maximum injection of 1e4 and 50 ms, respectively.

To gain additional misreading peptide identifications that are aligned to the quantification runs, two additional acquisition schemes were applied using the same chromatography setting. First, a top 30 DDA method using survey scans at a resolution setting of 60,000 FWHM was applied for the analysis of 8 µM and 16 µM Str-treated cell cultures. Second, we applied DDA with MS gas-phase fractionation (350–600; 650–850; 850–1600 m/z). These additional acquisition schemes, however, provided not more than 5% additional unique misreading peptide identifications.

Acquired data were processed using MaxQuant software (Version 1.6.5.0).66 We constructed two sequence databases to allow for the systematic detection of sequence variants. In a first search, a database containing the UniProtKB E. coli MG1655 reference proteome (4.491 entries) augmented with known lab contaminants was used. In the main search, EF-Tu peptide sequences (24 cognate peptides) with all possible amino acid substitutions were added to the database.

Identification of error clusters by DDA-MS. Peptides with error clusters were initially identified using the PEAKS software (version 10.5) applying the SPIDER algorithm. The database contained the UniProtKB E. coli MG1655 reference proteome.
proteome and 314 potential modifications in-built in the software were taken into account. The precursor mass tolerance was set to 10 ppm and the fragment mass tolerance to 0.02 Da. Peptide identifications were filtered using a target decoy approach with decoy fusion at a false discovery rate of typically 2% and a typical mutation ion intensity ≥2%. Typical AGA-induced misreading events were weighted by adjusting the mutation-weight matrix; i.e., the likelihood of R → C, N → K, D → E, E → D, Q → H, K → N, M → I, F → L, F → S, S → T, T → S, Y → N, and Y → H was set to 1.882, the value of V → I substitutions. Cluster candidates identified by PEAKS and error clusters that were considered likely to occur because they include combinations of frequent single errors, and the respective peptides with single substitutions, were included in the MaxQuant database described above. MaxQuant results were further analyzed in Skyline69. Missense clusters that were induced by Str were manually validated by inspecting the mass accuracy of the precursor, the ion-don product of the precursor isotope distribution, and the consistency of the highest-scoring MS/MS spectrum with predicted PROSIT spectra71 (see Supplementary Fig. 3a). The existence of error clusters was confirmed in ambiguous cases. In some peptides, the same amino acid substitution can occur at different positions in a peptide. This can result in regioisomers that often co-elute (see Supplementary Fig. 3c: spectra with red asterisks). These isobaric peptides also fragment resulting in chimeric spectra with lower-scoring peptide-spectral matches, which hampers quantification. Therefore, we excluded samples with high amounts of sample were analyzed and controls of non-treated cells were performed. In parallel reaction monitoring (PRM), correct peptides, peptides with single substitutions, and peptides with error clusters were targeted in their highest populated charge state. For the individual experiments and the performance of the mass spectrometer, and for the analysis of error clusters, we applied the QRAS software72,73 and abundance differences were estimated based on the sum of the individual integrated fragment intensities.

**Selection and quantification of error clusters by targeted MS.** A comprehensive proteome-wide analysis of error clusters is not possible due to dynamic range restrictions and the limited sensitivity of the approach. Error clusters reported here were selected for targeting if the peptides for all individual misreading events in the cluster were detected by DDA and these errors were reduced by AGA treatment. Only error clusters with net delta mass = 0 and different from known PTMs and peptide derivatizations were considered. In DDA experiments, apparent error clusters that were not induced by Str were excluded from the analysis. For targeted experiments, subsets of error clusters were chosen based on sequence distance and physico-chemical properties of the respective peptides (e.g., hydrophobicity). All error clusters that were selected for targeting and that could be unambiguously identified were included in the analysis. The frequency of error clusters was estimated by LFQ and absolute quantification (see Figure captions and the Source data file) in the following way.

In LFQ, correct peptides, peptides with single substitutions, and peptides with error clusters were targeted by PRM in the same sample. Peptides with single substitutions and correct peptides were identified by the high sequence coverage of their co-eluting fragment ions and their high dot product derived from the comparison to spectra identified by search engine runs. In ambiguous cases, AQUA peptides were used to validate the identity of peptides with single substitutions. For the detection of peptides with error clusters, we spiked-in isotope-labeled reference peptides (estimated ±10 fmol on column, I/P-L). These peptides, which are not used for quantification, help to select interference-free pseudo-transitions and to identify the missense peptides by their close-to-one ratio-doped product. Identical amounts of sample were analyzed and controls of non-treated cells were performed to exclude false-positive identifications. LFQ data were only used to estimate error frequencies if the substitutions did not dramatically alter the physico-chemical properties of the peptides (e.g., D → E, E → D). If clusters involved amino acid substitutions that led to changes in the charge state distribution (e.g., Y → H) or changes in their tryptic cleavage pattern (e.g., R → C, N → K, K → N), measured intensities were not directly interpreted as abundances (Supplementary Figs. 7b (panel 2) and 8). To include such data in the analysis of the distance dependency of error frequencies and to normalize the impact of fragmentation differences, error values for various AQUA were normalized to the εcorr of Str. For the absolute quantification of error clusters, we applied the QEXAMS workflow18 with minor changes. QEXAMS relies on the chromatographic enrichment of target missense peptides in different, partially orthogonal chromatographic steps. After enrichment, target missense peptides are detected and quantified by SRM or PRM. In detail, chromosome-encoded EF-Tu with a C-terminal His-tag was subcloned and expressed in E. coli. For the analysis of the tryptic digest was determined and 3–4 tryptic EF-Tu peptides were used to quantify the absolute amount of proteolyzed EF-Tu using AQUA peptides19. Sub-quantification of AQUA peptides corresponding to missense peptides with one, two, three, or four substitutions were spiked into the tryptic digest, typically at a ratio of 1:1000–1:10,000. Subsequently, peptides in the digest were separated in three chromatographic steps: (1). By size-exclusion chromatography (SEC) on a Superdex Peptide 10/300 GL column in an isocratic HPLC run (20% ACN in 0.1% formic acid, (2). By reversed-phase chromatography at neutral pH using a LiChrospher WP300 RP-18 (5 µm) column in an ACN gradient (2–85% ACN) at a flow rate of 0.2 mL/min (3). By reversed-phase chromatography at acidic pH using a LiChrospher WP300 RP-18 (5 µm) column in an ACN gradient (0–85% ACN in 0.1% trifluoroacetic acid in 65 min). In each chromatographic dimension, fractions were screened by SRM to identify target missense peptides. The tryptic fractions were pooled, concentrated, and applied to the next chromatographic step. After the QEXAMS workflow, targeted missense peptides are typically ~1000-fold enriched and often among the most abundant peptides in the fraction.

Peptides with error clusters were quantified within the dynamic range of the mass spectrometer by SRM and were further validated by PRM (Supplementary Fig. 4c). To avoid false-positive identifications the potential contamination of AQUA peptides with unlabeled peptides was controlled (in all cases ≤0.1%). In few cases were unlabeled peptide was detected in AQUA peptides, this contamination was taken into account by considering only those quantifications where the signal intensity of the sample exceeded the background level by at least threefold. To illustrate the dependence of error clusters on AGA treatment and to further exclude quantification errors due to light peptide contaminations, the missense peptides derived from treated and untreated cells were enriched by QEXAMS (Figs. 2b, e and Supplementary Fig. 4b). To allow for a quantitative comparison the loaded sample was normalized by the intensity of the enriched AQUA peptides. In general, in the analysis of error clusters, the spillover between different chromatographic runs was controlled and was not detectable within the dynamic range of the mass spectrometer.

**Selected reaction monitoring (SRM).** Samples were analyzed on an Ultimate 3000 RSLC system coupled to a TSQ Quantiva triple quadrupole mass spectrometer (Thermo Fisher Scientific). Peptides were separated using an in-house packed column (14 cm, 50 µm ID, packed with Reprosil-Pur C18-AQ 120 Å, 3 µm) with 0.1% formic acid at 0.3 µl/min flow rate. Q1 and Q3 were set to unit resolution (0.7 FWHM). A spray voltage of 2100 V was used with a heated ion transfer tube setting of 325 °C. The Chromatofitting setting was kept at 3. Scheduled transitions were recorded in a 5 min window and a cycle time of 1 s was applied. Skyline software (Version 3.5) was used for the evaluation of the acquired data22,23. The predominant charge state of each peptide was targeted and 3–5 transitions per peptide were monitored. To ensure high confidence identification, only peptides with ratio dot products ≥0.95 were considered. The light/heavy ratio of each peptide was used for the absolute quantification of correct and missense peptides, which were used to calculate the error frequencies.

**Parallel reaction monitoring (PRM).** Samples were analyzed on an Ultimate 3000 RSLC system coupled to QExactive HF-X hybrid Quadrupole-Orbitrap, QExactive HF hybrid Quadrupole-Orbitrap, or QExactive Plus Hybrid Quadrupole-Orbitrap mass spectrometers (Thermo Fisher Scientific). Acquisition parameters were tuned for the individual experiments and the performance of the mass spectrometer, and can be found in the Source data file. For LFQ, correct peptide ions and peptide ions with amino acid substitutions were targeted in their highest populated charge state. Sets of interference-free fragment chromatograms were extracted using the Skyline software22,23 and abundance differences were estimated based on the sum of the individual integrated fragment intensities. Quantification and statistical analysis. Error frequencies were estimated as the ratio of missense peptide and correct parental peptide. The expected stochastic frequencies of error clusters was estimated as the product of the error frequencies of the individual misreading events. The amplification over the stochastic level was calculated as the ratio of cluster error frequency and expected stochastic error frequency. The probability of the next misreading event ($E_{\text{next}}$) was determined in three different ways (see Supplementary Fig. 7b). ($E_{\text{next}}$) was derived from time courses and AGA titrations as the slope when the frequency of error clusters was analyzed as a function of the first misreading event (Figs. 4b, d, 5a, and Supplementary Fig. 7a). In addition, $E_{\text{next}}$ was estimated from point measurements at a fixed time and AGA concentration. Here, $E_{\text{next}}$ was estimated as the ratio of the abundances of the error cluster and the first error. (2) For data generated by absolute quantification and QEXAMS the ratio of the error frequencies of the error cluster and the first error were used (Figs. 4a, c and Supplementary Fig. 7b, panel 3). (3) Alternatively, peptides with error clusters and their first error were quantified by LFQ and $E_{\text{next}}$ was calculated as the ratio of the integrated intensities of peptides with error cluster to peptides with the single first error (Fig. 5b middle and right panels). The QEXAMS workflow (Fig. 8). Notably, the pass criteria for the detection of error clusters, for the analysis of error amplification, and for the analysis of $E_{\text{next}}$ differ somewhat (see...
Supplementary Fig. 3a–c): For the analysis of error amplification correct peptides, peptides with error clusters and with the first and the second error alone have to be quantified. None of the errors should introduce significant changes in physico-chemical properties of peptides, such as charge state, cleavage pattern, or ionizability. For the analysis of the pass, criteria are slightly relaxed and only the second error should not introduce significant changes in physico-chemical properties.

Experiments were repeated as indicated in figure legends. Technical replicates are derived from repeats in the analysis of the same sample. Biological replicates derived from analysis of separate, biologically distinct samples produced independently of each other starting from culture inoculation from different clones, and followed by individual cell growth, drug treatment, sample processing, and data acquisition. Statistical analyses were performed with Perseus, Excel, and GraphPadPrism as described in ‘Methods’ and the figure legends. Reported p values are based on unpaired t tests (two-tailed (Supplementary Fig. 1), and one-tailed (Fig. 1d, f)). When representative results are shown, the experiment has been repeated three times with similar results.

Reporting summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability
Data supporting the findings of this manuscript are available from the corresponding authors upon reasonable request. A reporting summary for this article is available as a Supplementary Information file. Data that support the findings of this study have been deposited to the ProteomeXchange Consortium via the PRIDE and PanoramaPublic repositories. All proteomics data were uploaded to PRIDE repository (in general: PXD019328, data related to Supplementary Fig. 3: PXD022989). Replicates of IDA-based analysis of amino acid substitutions and targeted MS data that supports all key conclusions was deposited to PanoramaPublic (PXD019328) (https://panoramaweb.org/ Error-cluster.url). Spectral evidence for error clusters is presented in the Supplementary Figures. A detailed report on the data structure and availability is given in the Source Data file. Source data are provided with this paper.

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Author contributions

I.W. designed the project and carried out the mass spectrometry experiments with contributions from R.G. and A.N.G. E.S. carried out in-vitro translation experiments. C.L., H.U., and M.V.R. contributed to experimental design and data interpretation. I.W. and M.V.R. wrote the manuscript with contributions of all authors. Funding was acquired by I.W., M.V.R., and H.U.

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Competing interests

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Additional information

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