Increased RNA Editing May Provide a Source for Autoantigens in Systemic Lupus Erythematosus

Highlights

- SLE patients have elevated RNA editing in their blood
- In SLE, elevated editing at coding regions can potentially generate autoantigens
- Elevated RNA editing may promote SLE progression by increasing autoantigenic load

In Brief

Roth et al. show that SLE patients have elevated RNA editing, a process that modifies an RNA sequence from the sequence encoded in the genome. This is manifested as both increased numbers and various altered coding sequences, which may promote autoimmune progression by increasing autoantigenic load.
Increased RNA Editing May Provide a Source for Autoantigens in Systemic Lupus Erythematosus

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https://doi.org/10.1016/j.celrep.2018.03.036

SUMMARY

RNA-editing mechanisms, which induce nucleotide substitution in the RNA, increase transcript and protein diversities. Editing dysregulation has been shown to lead to grave outcomes, and transcriptome-wide aberrant RNA editing has been found in tumors. However, little is known about the involvement of editing in other diseases. Systemic lupus erythematosus (SLE) is a multisystemic autoimmune disease characterized by a loss of tolerance for autoantigens from various tissues and the production of multiple autoantibodies. Here, we show that blood samples from individuals with SLE have abnormally high levels of RNA editing, some of which affect proteins and potentially generate novel autoantigens. We suggest that elevated RNA editing, either by ADARs or APOBECs, may be involved in the pathophysiology of SLE, as well as in other autoimmune diseases, by generating or increasing the autoantigen load, a key requisite for the progression of autoimmunity.

INTRODUCTION

Adenosine-to-inosine (A-to-I) editing is a prevalent form of post-transcriptional modification (Bazak et al., 2014a; Ramaswami and Li, 2014), catalyzed by the adenosine deaminase acting on RNA (ADAR) enzymes (Bass, 2002; Nishikura, 2010; Savva et al., 2012). Because inosine is interpreted by most cellular machineries as a guanosine (G), there are several possible outcomes to this deamination (Speyer et al., 1962). Hence, editing can alter splice consensus elements, microRNA seeds, and binding sites, as well as protein coding sequences (Burns et al., 1997; Ekdahl et al., 2012; Higuchi et al., 2000; Rueter et al., 1999). The reaction is catalyzed in humans by the ADAR1 (ADAR) and ADAR2 (ADARB1) enzymes. ADAR2 is confined to the nucleus, whereas ADAR1 has two isoforms—the p110 isoform that is localized in the nucleus and the p150 interferon (IFN)-inducible isoform that shuttles between the nucleus and the cytoplasm (Eckmann et al., 2001; George and Samuel, 1999).

A-to-I RNA editing is a constitutive and finely tuned process that is involved with several systems in the body and requires a delicate balance of activity. On the one hand, insufficient editing has detrimental and even lethal effects, partly due to the accumulation of endogenous double-stranded RNAs (dsRNAs), which triggers an innate immune response (Liddicoat et al., 2015a; Mannion et al., 2014; Pestal et al., 2015). ADAR1 editing deficiency is involved in autoimmune disorders such as Aicardi-Goutières syndrome (AGS) and dyschromatosis symmetrica hereditaria (DSH) (Liu et al., 2006; Rice et al., 2012). Both show some phenotypic similarities to systemic lupus erythematosus (SLE), and AGS patients, like SLE patients, are characterized by elevated type I IFN production (Crow and Manel, 2015; Lee-Kirsch et al., 2014; Rice et al., 2012).

On the other hand, excessive ADAR activity may be harmful. Studies show that elevated levels of ADAR1 usually correlate with higher editing activity, manifested both as higher rates of editing and as editing of non-typical editing sites (Han et al., 2015; Paz-Yaakov et al., 2015). Elevated levels of editing have been shown to have significant effects on the transcriptomic diversity of cancers and, in several cases, even correlate with patient survival (Han et al., 2015; Paz-Yaakov et al., 2015).

The other type of RNA editing in humans is the deamination of cytidine to uridine (C-to-U) by several members of the APOBEC protein family of cytidine deaminases, mainly APOBEC1 and APOBEC3A (Sharma et al., 2015; Smith et al., 2012). Similar to ADAR1, APOBEC3 paralogs are upregulated by IFN (Peng et al., 2006). Overexpression of APOBEC3A was shown to affect thousands of sites that have the potential to recode more than a thousand proteins (Sharma et al., 2017).

SLE is a multisystemic heterogenic autoimmune disease. It is characterized by autoantibodies to a variety of autoantigens derived from various tissues, as well as antinuclear antibodies (Lipsky, 2001; Tsokos, 2011). The causes and pathogenesis of the disease are not fully understood. SLE patients usually have
elevated levels of circulating type I IFN and increased expression of both type I and type II IFN-stimulated genes (ISGs) (Chiche et al., 2014; Crow and Manel, 2015; Hagberg and Rönnblom, 2015). Thus, inhibitors of IFN and downstream ISGs are attractive and promising therapeutic targets for treatment (Furie et al., 2017; Hagberg and Rönnblom, 2015; Kirou and Gkrouman, 2013; Niewold, 2000). Moreover, high artificially induced levels of IFN can cause SLE-like symptoms (Ho et al., 2008; Niewold, 2008). Altered levels of ISGs and RNA modifications have been marked as potential contributors to SLE pathogenesis in several previous studies (Graham and Utz, 2005; Hueber et al., 2004). This includes increased rates of alternative splicing (Ng et al., 2004) and indications of aberrant RNA editing in a few genes, as well as elevated expression of ADAR1 in lymphocytes (Laxminarayana et al., 2002, 2007; Orlowski et al., 2008). However, these early studies focused only on small subset of editing targets.

Here, by analyzing wide-scale, high-throughput sequencing data (Hung et al., 2015), we show that SLE patients have elevated global RNA-editing levels. In addition, we use computational predictions to demonstrate that this excessive editing can potentially generate autoantigens, which may then be presented on major histocompatibility complex (MHC) molecules and induce an immune response. Editing in coding regions, which alters the amino acid sequence (recoding), may prove immunogenic. Because proteasomal degradation is presumably unaffected by RNA editing, an edited form of a protein could be degraded and eventually be presented by the MHC system. We previously showed that medullary thymic epithelial cells (mTECs), which mediate the presentation of self-epitopes in the thymus, routinely express many edited forms of proteins and thereby forestall the induction of a dangerous immune response against edited peptide-derived autoantigens (Danan-Gotthold et al., 2016). However, possibly due to incomplete presentation and/or confinement of the process to selected edited versions, not all edited forms are presented by the mTECs. As a consequence, T cells specific for edited epitopes may escape negative selection and react to cells that present an editing-derived neo-autoantigen (Danan-Gotthold et al., 2016). Although the main activity of editing is to inhibit the innate immune response to endogenous dsRNA (Liddicoat et al., 2015b; Mannion et al., 2014; Pestal et al., 2015), our results suggest that RNA editing can trigger an immune response directed against editing-originated neo-autoantigens, exacerbating autoimmunity. We therefore suggest it is involved in the etiology and pathogenesis of SLE and possibly other autoimmune diseases.

RESULTS

Enhanced A-to-I Editing in SLE Patients

Many studies of editing in various diseases concentrated on previously detected editing sites. Here, we used RNA sequencing (RNA-seq) data to evaluate changes in global levels of editing, without prior knowledge of specific sites. To compare the RNA-editing levels in SLE patients with those in healthy individuals, we first used two complementary approaches to estimate the global editing rate using RNA-seq samples from Hung et al. (2015) (99 from SLE patients and 18 from healthy individuals) (Table S1); the Alu editing index (AEI), which measures the global rate of editing in Alu repeats (Bazak et al., 2014b), and the global rates of clustered editing sites in each sample, according to the normalized number of hyper-edited (HE) sites (see Experimental Procedures) (Porath et al., 2014). Both types of analysis revealed that global editing was significantly elevated in the blood of SLE patients compared to that of controls (Wilcoxon p value = 4.96e–6 and 8.27e–7 for AEI and HE analyses, respectively). Because higher editing levels may be associated with higher levels of ISGs, we grouped the patients according to the original dataset of interferon signature metric (ISM) division, which measures the presence of the IFN-inducible genes’ expression signature. As expected, ISM-high SLE patients had significantly higher editing levels than controls (Wilcoxon p value = 4.16e–7 and 2.48e–8 for AEI and HE analyses, respectively) (Figures 1A and 1B). However, even ISM-low patients had significantly higher editing levels than controls (Wilcoxon p value = 3.28e–2 and 3.93e–2 for AEI and HE analyses, respectively) (Figures 1A and 1B). Overall, the editing signal was clean. A-to-G mismatches comprised most mismatches (82.79%), and the ADAR deamination motif (5’ neighbor preference A = U > C > G) (Eggington et al., 2011; Riedmann et al., 2008) was observable (Figures 1C and 1D), indicating that most of the A-to-G substitutions detected in the HE analysis resulted from ADAR editing. Similar results were obtained from another, but smaller, independent dataset from Rai et al. (2016) (Figure S1).

As a complement approach, we investigated the level of RNA editing at known sites that have been identified and characterized by previous studies (see Experimental Procedures). Among the sites at which editing levels were found to differ significantly between the controls and the ISM-high patients, most had increased editing levels. This was also the general trend in sites without significant changes (Figure 1E). These results further support the observation that RNA editing is elevated in the blood of ISM-high patients.

To determine whether ADAR1, which is itself an ISG, is responsible for the increased editing levels, we examined its expression levels. As expected, its expression was significantly higher in ISM-high patients compared to controls (2.24-fold change, Wilcoxon p value = 3.77e–6) (Figure 1F). However, in contrast to the results of the global RNA-editing levels, no significant differences were observable between ISM-low patients and controls (Wilcoxon p value = 0.54). This may indicate that ADAR1 expression is regulated directly by IFN, while RNA-editing levels remain relatively high in these patients even without IFN induction. In contrast, the expression of ADAR2 was lower in ISM-high patients compared to controls (0.75-fold change, Wilcoxon p value = 0.035) (Figure 1G). Altogether, our results show significantly elevated ADAR1 expression and editing activity in ISM-high patients.

Enhanced C-to-U Editing in SLE Patients

Proteins from the additional deaminase family, the APOBEC, were also significantly upregulated in ISM-high patients compared to controls (Figure 2A; Table S2). This may lead to
the accumulation of C-to-U mismatches, which can serve as a potential source for neo-autoantigens. A dataset of putative C-to-U editing sites in monocytes and macrophages has been published (Sharma et al., 2015). Of the sites, 252 were detected in the examined dataset (96.5% of the expressed sites), and most of them (75%) were edited in more than 20% of the samples. Out of these, 26 had significantly elevated editing rates in ISM-high patients (false discovery rate [FDR] < 0.1), and a similar global trend was observable (Figure 2B). Because the editing probably occurs only in specific cell types (e.g., monocytes and macrophages), very low editing rates were observed in the total cell population (90% of the sites had a mean rate of 0%–2%). To further assess the C-to-U editing levels in SLE patients and healthy individuals, we measured the global rates of HE C-to-U clusters in each sample (see Experimental Procedures). As was the case for the A-to-I editing, ISM-high patients had significantly more clustered C-to-U sites per sample (Wilcoxon p value = 4.87e−3) (Figure 2C), and their numbers correlated well with APOBEC3A levels (r = 0.70) (Figure 2D) indicating that this is the main deaminase driving this elevation.

Elevated Levels of Recoding Events in SLE Patients

Only a small portion of A-to-I RNA editing results in the recoding of proteins. To identify recoding sites associated with SLE, we systematically searched for differentially edited sites with a non-synonymous outcome. Because the current approaches for de novo detection of recoding sites without a matched DNA sequences from the same individual perform poorly, we limited the analyses to high-credibility sites found within the HE regions or to the previously verified ones.

We first examined putative recoding sites detected by the HE analysis, in which the potentially editable adenosine is found on the transcribed strand. Excluding known SNPs and sites within highly polymorphic genomic regions, we identified 624 putative recoding events in SLE patients.
recoding sites. As expected, most of these sites (95%) were edited in SLE samples, and their prevalence was higher than in controls (an average of 1.100 sites per million mapped reads in SLE patients versus 0.967 in controls, Wilcoxon p value = 0.018) (Figure 3A). The obtained signal was noisy (Figure S2A), indicating that a portion of these sites are not bona fide ADAR targets. Nevertheless, the neighbor preferences obtained fairly agree with the ADAR motif (Figure S2B), supporting the contribution of genuine editing by ADAR. Many of these sites were stochastically edited (545 of 624 sites were edited in fewer than five samples). We therefore compared the relative number of samples that expressed the edited version of the transcript between the groups. Two such sites, in the genes IFITM1 and ODF3B, were found to have higher editing levels in ISM-high patients (Wilcoxon, FDR < 0.05) (Figures 3B and 3C), with no significant expression difference in SLE patients and controls in the genes SH3BP2 and ARL6IP4 (at chr4:2,835,556 and chr12:123,466,262, respectively).

We also examined the editing rates at recoding sites identified in other studies (see Experimental Procedures). Two sites were found to have higher editing levels in ISM-high patients (Wilcoxon, FDR < 0.05) (Figures 3B and 3C), with no significant expression difference in SLE patients and controls in the genes SH3BP2 and ARL6IP4 (at chr4:2,835,556 and chr12:123,466,262, respectively).

These results support the hypothesis that the high global editing levels in SLE patients can give rise to a variety and higher levels of the edited versions of proteins.

DISCUSSION

The results of this study suggest the possibility of a connection between autoimmunity and excessive RNA editing. We surmise that the latter may facilitate the generation of autoantigens in peripheral tissues. Because these autoantigens might not necessarily be expressed in the thymus, reactive T cells may escape the negative selection and recognize these recoded proteins as non-self. The elevated global editing activity in SLE patients, one of the manifestations of the inflammatory condition, results...
in increased variety and higher levels of edited forms of proteins. Similarly, conditions that lead to editing alterations can result in new recoding events (Daniel et al., 2012). These have the potential to be processed into autoepitopes that may then be subsequently presented on the MHC molecules, thus stimulating an autoimmune response (Figure 4). Our results therefore enrich knowledge about the recently discovered role of ADAR1 and RNA editing in regulating the innate immune system and support the connection between imbalance of RNA editing and immune dysfunction.

We hypothesize that the elevated editing may be involved in positive feedback, aggravating autoimmunity (Figure 4). Inflammatory cytokines produced by an immune response triggered by elevated editing levels may maintain or even further increase the editing levels by stimulating the IFN-induced deaminases, resulting in the production of still more, potentially immunogenic, editing-recoded peptides. In addition, the edited epitope not only may prove immunogenic but also may initiate a process of epitope spreading (Lehmann et al., 1992; Sercarz, 1998; Vanderlugt and Miller, 2002). Moreover, SLE has several characteristics that may promote the immunogenicity of RNA editing, such as the upregulation of epitope presentation pathway by IFNs and the accumulation of dead cell debris (Boehm et al., 1997; Lipsky, 2001; Rusinova et al., 2013; Schroder et al., 2004).

Although the results presented here are promising and demonstrate the potential of editing to create neo-autoantigens, there are several limitations to this type of research, leading to incomplete detection of the editing-derived proteome and assessment of its immunogenicity. First, from an analytical perspective, while the HE analysis enables the discovery of recoding editing sites, the HE reads comprise only a minority (~1%) of the total reads, and presumably a small fraction of the total editing. Moreover, the recoding sites detected here have a relatively low signal-to-noise ratio, so the credibility of each site is not high enough, even though most of them are genuine and altogether the sites are credible. Second, from a biological perspective, several limitations arise. The RNA was extracted mainly from living cells; thus, it is possible that editing sites generating particularly immunogenic peptides may be underrepresented in the data because of their elimination by a fiercer immune reaction against such cells. Another potential drawback is introduced by the RNA-seq data used for the analysis being derived from whole blood, which contains heterogeneous cell types, which may dampen the signal and add noise. Thus, it is probable that the number of recoding sites detected here is an underestimation of the actual number of recoding editing sites in SLE patients.

In summary, we have shown significantly elevated RNA editing in SLE patients and revealed its potential to give rise to neo-autoantigens, implying a role for RNA editing in the etiology and progress of SLE. These findings provide another link between RNA editing and autoimmune diseases.

**EXPERIMENTAL PROCEDURES**

The Supplemental Experimental Procedures includes more details of the applied analyses.

**Datasets**

A whole-blood (Paxgene) RNA-seq dataset of 18 healthy individuals and 99 active SLE patients (GEO: GSE72509) from Hung et al. (2015) and a smaller dataset of 12 SLE patients and 4 controls (GEO: GSE60183) from Rai et al. (2016) were downloaded. Details are provided in Table S1.

The reads were aligned to the human genome (hg19) using STAR 2.4.2 (Dobin et al., 2013).

**HE**

We used a recently described pipeline (Porath et al., 2014) that enables the measurement of editing of heavily edited reads, which standard schemes fail to align correctly (Carmi et al., 2011).

**AEI**

To measure the editing in Alu elements, we used a previously described method (Bazak et al., 2014b). Due to the magnitude of the effect and the specific occurrence of editing in Alu elements (such as editing in clusters), this...
measurement results in a clean signal with a minimal false-positive rate (Bazak et al., 2014a).

**Known Sites**

RNA editing levels for a list of pre-known editing sites were calculated using REDIToolKnown, which is part of the REDItools package (Picardi and Pesole, 2013). The list of editing sites was compiled from previous studies (Khermesh et al., 2016; Pinto et al., 2014; Ramaswami et al., 2012).

**Expression Analysis**

The DESeq package (Anders and Huber, 2010) in R was used for analyzing differential gene expression in all control and SLE patient samples.

**Statistical Analysis**

The statistical analysis was done using R (R Project for Statistical Computing, http://www.r-project.org/). Unless otherwise specified, the statistically significant difference (5% confidence level) among two groups was tested using the Wilcoxon rank test. The tests were performed using the default parameters and in a two-sided manner. Where appropriate, p values were corrected for multiple testing.

**Recoding Editing Sites Analyses**

To provide the peptide sequence for the analyses and find recoding, HE and pre-known sites filtered for SNPs and HLA genes were annotated using wANNOVAR (Chang and Wang, 2012; Wang et al., 2010). HLA alleles for each sample were derived using OptiType (Szolek et al., 2014). For each sample, recoding sites (detected by the HE analysis) inside peptides that have at least one window (see Supplemental Experimental Procedures) with strong affinity to the sample-predicted HLA alleles were selected according to the NetMHCPan4.0 (Jurtz et al., 2017) predictions. A recoding event was considered to have an affinity-enhancing effect if the edited version of the peptide, for all windows on average, had a higher affinity (of at least 10%). The number of epitopes per peptide chain was calculated as previously described in Agranovich et al. (2013).

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Supplemental Experimental Procedures, two figures, and two tables and can be found with this article online at https://doi.org/10.1016/j.celrep.2018.03.036.

**ACKNOWLEDGMENTS**

We thank Eli Eisenberg, Amos Schaffer, Binyamin A. Knisbacher, Lea Shallev, and various Levanon lab members for the helpful discussions. We thank Ziv Paz and George Tsokos for the stimulating discussions at the early stages of the project. We thank Tamar Roth-Fenster for assisting in editing and drafting the manuscript. M.D.-G. is grateful to the Azrieli Foundation for the award of an Azrieli Fellowship. E.Y.L. was supported by the European Research Council.
(311257) and the Israel Science Foundation (1380/14). C.J.C. was supported by the Israel Science Foundation (1422/15).

AUTHOR CONTRIBUTIONS
E.Y.L., S.H.R., and M.D.-G. designed the research. S.H.R. and M.D.-G. developed and performed the computational analyses under the guidance of E.Y.L. M.B.-I. and Y.L. performed the epitope counting analysis. E.Y.L., S.H.R., and M.D.-G. wrote the manuscript with input from C.J.C. and G.R. All authors read and approved the final manuscript.

DECLARATION OF INTERESTS
The authors declare no competing interests.

Received: July 5, 2017
Revised: November 21, 2017
Accepted: March 9, 2018
Published: April 3, 2018

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Supplemental Information

Increased RNA Editing May Provide a Source for Autoantigens in Systemic Lupus Erythematosus

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Figure S1. Editing is significantly increased in SLE patients (dataset taken from (Rai et al., 2016)), Related to Figure 1. The compared cohorts are healthy controls (blue) and SLE patients (red). Paired comparisons were evaluated using the Wilcoxon test. The global levels of editing were assessed by determining (A) the editing in Alu elements (Alu Editing Index - AEI) and (B) the number of hyper edited (HE) sites (normalized by the number of mapped reads) per sample.
Figure S2. The signal cleanness of recoding sites at hyper edited regions, Related to Figure 3.
(A) Percentage of all mismatches and (B) the nucleotide frequencies of the neighboring nucleotides around the A-to-G sites
### Supplemental Tables

#### Table S2. Elevated APOBEC proteins levels in SLE, Related to Figure 2.

| Gene name   | Fold Change between ISM-high patients and controls | Wilcoxon p-value |
|-------------|---------------------------------------------------|------------------|
| APOBEC3A    | 2.58                                              | 5.25e-08         |
| APOBEC3B    | 2.48                                              | 0.00237          |
| APOBEC3F    | 1.80                                              | 1.96e-08         |
| APOBEC3G    | 1.48                                              | 4.28e-08         |
| APOBEC3H    | 1.84                                              | 0.000885         |
Supplemental Experimental Procedures

Reads alignment
The reads were aligned to the human genome (hg19) using STAR 2.4.2 (Dobin et al., 2013), using alignment parameters that limit the number of mismatches to 0.1 of the mapped length and accept only uniquely aligned reads (default parameters except alignIntronMax - 1000000, alignMatesGapMax - 1000000, outFilterMismatchNmax - 999, outFilterMismatchNoverReadLmax - 0.1, outFilterMultimapNmax - 1, outReadsUnmapped - Fastx, quantMode – GeneCounts). This resulted in three output files for each sample (an alignment file, a fastq file containing all the unmapped reads, and a file containing the number of reads per gene) that were used in the next analyses.

Hyper Editing
RNA editing by ADAR1 usually happens in clusters. Thus, many heavily edited reads may differ from the DNA to the extent that standard schemes fail to align them correctly (Carmi et al., 2011). A recently described pipeline enables the measurement of such heavily edited reads (Porath et al., 2014). It was observed that HE is closely correlated with ADAR1 activity (Paz-Yaacov et al., 2015; Porath et al., 2014). In this approach, the reference genome and the reads are transformed, changing a mismatch dominance (i.e. that the relative part of A to G mismatches, which was set to 0.8. The numbers of hyper-edited sites were normalized by the number of the mapped reads in each sample. Two samples in which we identified a very high rate of clustered A-to-C substitutions (more than 70% of all clustered substitutions found in the sample) were excluded from this analysis and from all further calculations using sites detected by it, since A-to-C clustered substitutions were previously shown to result from an Illumina machine artifact (Dohm et al., 2008).

Alu editing index
As most of the editing sites in the human genome (over 90%) are concentrated in Alu genomic regions, AEI gives a good approximation of the global editing level and ADAR1 editing efficiency in the sample (Bazak et al., 2014a).

Reads that were aligned to Alu elements were collected and mismatches between them and the reference genome (hg19) were identified. To get a reliable collection of mismatches, mismatches with a Phred score lower than 30 or those reported as SNPs in dbSNP (SNP build 135) were filtered out (Eisenberg et al., 2005). To eliminate putative sequence errors the results were further filtered using a probabilistic model assuming an a priori sequencing error rate of 0.001 (corresponding to Phred score of 30). This was achieved by applying Benjamini-Hochberg multiple testing correction for all Alu adenosines, with a false discovery rate (FDR) of 0.05. Over 90% of all Alu evaluated had A-to-G mismatch dominance (i.e. that the relative part of A-to-G mismatches is higher than all others combined), which further supports the cleanness of the signal. However, for the calculations of the editing levels we used all Alu elements (to avoid biases caused by choosing dominant A-to-G Alus only) as described below.

A straightforward comparison of editing levels, based on the editing rates at all sites, is quite complicated and requires ultra-high coverage due to the fact that most adenosines in Alu elements are edited to at least some extent (though usually to a low degree, less than 1% of adenosines) (Bazak et al., 2014a, 2014b). Hence, any set of detected sites in a given sample may represent only a small random fraction of the editing sites. Thus, the number of reads supporting a specific mismatch is not a good measure of the effect in the whole sample. To apply all the above to our data set, we used the AEI (Alu Editing Index). This index gives the average editing level across all adenosines in Alu elements weighted by their expression, which is the ratio of A-to-G mismatches (presumably due to deamination to inosines), over the total number of nucleotides aligned to an Alu repeat (both adenosines and presumed inosines). This index averages over millions of adenosines, which as previously described, (Bazak et al., 2014b) makes it rather robust to statistical noise.

Recoding Editing Sites Analyses
The examination of editing rates at non-synonymous, pre-known editing sites was limited to 60 sites, which were expressed by at least 10 samples in each group, so that n ≥ 10 for each group of the Wilcoxon test.
Protein prediction was assessed using a list of edited sites filtered from known SNPs (dbSNP135 (Sherry et al., 2001) and highly polymorphic HLA genes, but with conserved editing sites (Pinto et al., 2014) included). The amino acid sequence (as found in the NCBI database) was queried and the "mutated" AA was planted in the right place of the sequence generating peptides of 21 AA long (where the edited AA is in the middle). The affinity of each peptide in the couple (edited and non-edited form) for the predicted HLA alleles of the samples were evaluated using NetMHCpan 4.0 (Jurtz et al., 2017) for K-mers (window) of between 8-11 AA long.

Expression analysis
The gene annotations for the expression analysis were obtained from the UCSC gene tables (Karolchik et al., 2003). The number of reads aligned to each gene was calculated by STAR.

Number of Epitopes Prediction
The number of epitopes per peptide chain was calculated by combining three algorithms: a proteasomal cleavage algorithm (Ginodi et al., 2008), a TAP binding algorithm (Peters et al., 2003), and the MLVO MHC binding algorithm (Vider-Shalit and Louzoun, 2011) as previously described in (Agranovich et al., 2013).

Since the used algorithms only apply to the 39 most common alleles in Caucasian population, the epitopes were computed for the 35 HLA alleles of the relevant samples. The results were weighted according to the allele frequency in the global human population. The efficiency of these algorithms and their quality has been systematically validated (previously) vs. epitope databases and was found to induce low false positive (FP) and false negative (FN) error rates (Maman et al., 2011a, 2011b). The number of epitopes was calculated for both the sequence and the average of three randomly scrambled versions of it, in order to assess whether the number of epitopes is affected by the sequence of the peptide or the AA composition (a biochemical effect).
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