Antibody Heavy Chain CDR3 Length-dependent Usage of human IGHJ4 and IGHJ6 Germline Genes

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STATEMENT OF SIGNIFICANCE: Therapeutic antibody discovery using synthetic diversity has been proved productive. The quality of diversity design limits the developability of synthetic hits. Here we quantitatively determined the CDR-H3 length-dependent usage of human DH/JH germline genes and the resulting spatial paratope landscape, which sheds light on rational design of synthetic diversity with improved probability of success.

ABSTRACT: Background: Therapeutic antibody discovery using synthetic diversity has been proved productive, especially for target proteins not suitable for traditional animal immunization-based antibody discovery approaches. In recent years, many lines of evidences suggest that the quality of synthetic diversity design limits the development success of synthetic antibody hits. The aim of our study is to understand the quality limitation and to properly address the challenges with a better design. Methods: Using VH3-23 as a model framework, we analyzed the naturally and productively rearranged CDR-H3 diversity in human immune repertoire. With homology modeling, we further built VH3-23-based structural models to understand the spatial paratope and its influencing parameters. Results: We observed and quantitatively mapped CDR-H3 loop length-dependent usage of human IGHJ4 and IGHJ6 germline genes in the natural human immune repertoire. Skewed usage of DH2-JH6 and DH3-JH6 rearrangements was quantitatively determined in a CDR-H3 length-dependent manner in
natural human antibodies with long CDR-H3 loops. Such CDR-H3 length dependent usage of human germline genes was not impacted by the choices of VH in the V(D)J recombination, ethnic background and health conditions. Structural modeling suggests choices of JH help to stabilize antibody CDR-H3 loop and JH only partially contributes to the paratope. **Conclusions:** We quantitatively determined the CDR-H3 length-dependent usage of human germline genes, which makes it possible to design synthetic diversity fully mimicking that of natural immune repertoire. Our observations shed light on the design of next generation synthetic diversity with improved probability of success.

**KEYWORDS:** CDR-H3; Diversity; Synthetic antibody library; JH4; JH6

**INTRODUCTION**

Therapeutic antibodies can be discovered via *in vivo*, *in vitro* or *in silico* approaches (1, 2). *In vivo* approach, relying on the immunization of wild type or transgenic animals carrying human antibody gene segments, while *in vitro* approach employs the selection power by display technologies to pan large and diverse antibody libraries (3). Both approaches have been proved very successful and have generated best-selling antibody therapeutics like Keytruda and Humira, respectively (4, 5). Recent advances in artificial intelligence and machine learning also facilitated *in silico* rational antibody design (1).

*In vitro* selection of synthetic antibody libraries complements *in vivo* immunization-based approaches by providing unique, non-natural antibody paratopes and escaping the limitations of self-tolerance (6, 7). One of challenges in the rational design of man-made antibody diversity lies in the complementarity determining region (CDR) H3 loop (CDR-H3) of antibody heavy chain variable region. Traditionally, CDR-H3 loop was randomly diversified by degenerate codons or parsimonious degenerate codons (8), which limited the therapeutic potential of antibody hits. Although TRInucleotide technology and Slonomics technology helped to reduce dysfunctional motifs and to precisely control the codon biases, the CDR-H3 still could not fully mimic that of natural diversity generated by V(D)J recombination and somatic hypermutation process (9, 10). This limits the translational success of therapeutic antibody candidates derived from *in vitro* selection of phage or yeast libraries (11). Recently, massive datasets of human immune repertoires by Briney, Soto and others made it possible for us to survey billions of antibody heavy chain sequences for a better understanding of CDR-H3 natural diversity at immune repertoire level (12-14).

In this study, we aim to understand natural immune diversity of the CDR-H3 loop in a precisely controlled manner. From antibody engineers’ perspective, we observed CDR-H3 length-dependent usage of human IGHJ4 and IGHJ6 gene segments in the natural human immune repertoire. We also observed the biased usage of DH3-JH6 rearrangements in antibodies with long CDR-H3 loops. Inspired by our observations, we also conducted spatial analysis of the VH3-23
antibody paratopes and defined the parameters influencing an antibody’s solvent accessible surface area (SASA). This knowledge sheds light on the design of next generation synthetic diversity.

MATERIALS AND METHODS

Data source
All antibody sequences were obtained from previous studies, including the IGHV3-23*01 dataset (European Molecular Biology Laboratory accession nos. AM076988–AM083316) (15,16) and the Dengue virus (DENV) acutely infected patients dataset (the BioProject accession number PRJNA205206) (17). The DENV dataset was processed as what’s described in the publication (17).

Antibody homology modelling and germline usage analysis
The Fv models were generated using Rosetta following the authors’ instructions (18,19). In brief, a template was selected. Template complementarity determining regions were grafted onto the template frameworks, the frameworks were then assembled according to a template VH-VL orientation. The CDR-H3 of the top ranked model was then de novo modeled and the relative VH-VL orientation was refined via local docking. Kabat numbering was used to number the CDR loops and the SASA was measured using Rosetta. Germline usage analysis was performed by aligning the variable domain sequences to the IMGT reference germline genes using IgBlast (20), the output was parsed using Change-O (21).

Statistical analysis
Statistical analysis was performed using OriginPro 2021. If the dataset conforms to normal distribution, one-way analysis of variance was used (22). Significance level was set to 0.05. Turkey test was selected for the mean comparison, and the Levene test was selected for the homogeneity of variance test. P value was obtained by means comparison and overall analysis of variance. If the dataset did not conform to the normal distribution, then non-parametric test method was used. The significance level was set to 0.05. Since the data were not repeated measures, the Kruskal-Wallis ANOVA multi-sample independent non-parametric test was selected (23). P value was obtained at the significance level of 0.05. If P>0.05, there is no significant difference between the two groups of data. If 0.05>P>0.01, there is a significant difference between the two sets of data, marked as *. If 0.01>P>0.001, there is a significant difference between the two sets of data, marked as **. If P<0.001, there is a significant difference between the two sets of data, marked as ***.

RESULTS
Antibody heavy chain CDR3-dependent usage of human IGHJ4 and IGHJ6

Antibody complementarity determining region (CDR) H3 loop (CDR-H3) is highly variable. The diversity of CDR-H3 is mainly generated by V(D)J recombination via different mechanisms (24). Skewed usage of human germline genes like IGHJ4 and IGHJ6 were previously reported in CDR-H3 (25,26). Briney B.S. et al discovered that a skewed small subset of IGHD and IGHJ gene segments were particularly used to encode long CDR-H3s in human peripheral blood antibodies (26). It, however, remains a still poorly understood correlation. To further understand the exact correlation between CDR-H3 length and skewed human germline gene usage, we analyzed the same set of VH3-23*01 sequences (European Molecular Biology Laboratory accession # AM076988–AM083316) that were extensively validated and studied in the field (15,16,27,28). This enabled us to elucidate the correlation on a pre-defined antibody framework and genetic background.

Analysis of the same set of unique, productively rearranged human VH3-23*01 gene sequences revealed strong correlation between CDR-H3 length and IGHJ germline gene usage (Fig.1a). All antibody sequences analyzed in this study encode unique CDR-H3s to eliminate bias introduced by redundancy in the human immune repertoire. As shown in Fig.1a, when CDR-H3 length increases from 7aa to 23aa, the germline usage of IGHJ4 decreased gradually from 67.57% at 7aa length to 6.11% at 23aa length, while the usage of human IGHJ6 germline gene increased proportionally from 27.03% at 7aa length to 94.90% at 23aa length. At a shorter CDR-H3 length like 7aa, IGHJ4 was preferentially used (67.57%) over other IGHJ germline genes. At a relatively longer CDR-H3 length like 23aa, IGHJ6 (94.90%) was predominantly used while other IGHJ germline genes were very rarely used. CDR-H3 length below 7aa or longer than 23aa was not analyzed in this study due to inadequate number of IGHV3-23*01 sequences available.

The dataset used in Fig.1a analysis contains antibody sequences isolated from a cohort of healthy and viral infection-naive adults of Danish background. These IGHV3-23*01 sequences were productively rearranged predominantly to IGHJ4 or IGHJ6. To avoid biases incurred from data source, we asked ourselves: (1) is genetic background or health conditions playing a role? And (2) is this observation reproducible for other human IGHV germline genes? To properly address above concerns, we analyzed a dataset (the BioProject accession number PRJNA205206) containing antibody sequences sampled from 44 Nicaraguan individuals acutely exposed to Dengue virus (DENV) during acute symptomatic dengue at 2-5 days post-symptom onset (17). Regardless of the germline usage of IGHJ1, IGHJ2, IGHJ3 and IGHJ5, the same germline usage patterns of IGHJ4 and IGHJ6 were observed with antibody sequences derived from a large cohort of patients acutely infected with DENV (Fig. 1c). This suggests that, as CDR-H3 loops go longer, the increased germline usage of IGHJ6 is an intrinsic feature of V(D)J recombination and it is independent of health conditions or ethnic backgrounds. To answer the question if this observation
is unique to human IGHV3-23\(^{*}01\) germline gene, we dissected the antibody sequences based on IGHV germline usage from the acute DENV cohort (n=44) and the accompanying healthy control cohort (n=8). Fig. 1b, 1d shows the same exact pattern of IGHJ4 and IGHJ6 germline usage in 68,067 and 8,539 antibody sequences derived from the acute DENV cohort and the accompanying healthy cohort, respectively.

Above data suggest CDR-H3 length-dependent usage of human IGHJ4 and IGHJ6 germline. Such skewed usage is an intrinsic feature of human heavy chain CDR-H3 diversity generation. It is not impacted by ethnic background, health conditions or the usage of human IGHV germline genes.

We then combined the two datasets and dissected antibody sequences based on human IGHJ germline gene usage. Total 68,023 antibody sequences were analyzed, regardless of health conditions, ethnic background and human IGHV germline gene usage. Fig. 2 showed the CDR-H3 length distribution of each IGHJ group, combined (Fig. 2a) or individualized (Fig. 2b). Of the 68,023 antibody sequences, 29,873 (43.92\%) belong to the IGHJ4 germline group, while 21,283 (31.29\%) belong to the IGHJ6 germline group. Even though only 889 and 991 antibody sequences were obtained in the IGHJ1 and IGHJ2 group, respectively, the Gaussian distribution pattern of the CDR-H3 length distribution of all IGHJ groups suggest the combined dataset is diverse and scientifically sound for further analysis. Fig. 2b showed the average length of CDR-H3 in natural human antibodies rearranged to human IGHJ6 germline gene is 16.40 amino acids (lower right, n=21,283). The IGHJ6 antibodies CDR-H3 length distribution peaked at length of 17 amino acids. The average length of CDR-H3 in natural human antibodies rearranged to human IGHJ4 germline gene is 12.43 amino acids (lower left, n=29,873). The IGHJ4 antibodies CDR-H3 length distribution peaked at length of 12 amino acids. This observation is consistent with what’s been described in Fig. 1, that is, human antibodies with shorter CDR-H3 tend to be preferentially rearranged to human IGHJ4 germline genes, while antibodies with longer CDR-H3 tend to be preferentially rearranged to human IGHJ6 germline genes. CDR-H3 amino acid usage frequency was further determined in each IGH antibody group. No statistically significant difference was observed on the overall amino acid usage patterns across all IGH antibody groups (Pearson’s Chi-squared test, p>0.05), albeit some amino acid residues like tyrosine were differentially used across IGH antibody groups (Fig. S1).

**CDR-H3 length-dependent preference in DH-JH recombination**

Recent study by Sankar K. et al showed that in natural human antibody repertoire the antibody CDR-H3 diversity was dynamically shaped by VH, VL and JH germline segment use (29). Previously we discovered CDR-H3 length-dependent usage of human IGHD germline genes (28). Such CDR-H3 length-dependent usage of non-canonical cysteines was extensively characterized.
in human VH repertoires (30) and in other animals like chicken (31), bovine (32), shark and camelid (33). Intrigued by the roles of human IGHD gene segments in shaping human antibody CDR-H3 diversity, we analyzed the human IGHD germline gene usage of natural antibodies derived from 8 healthy donors and 44 DENV acutely infected patients – the same dataset we used in section 3.1. Not surprisingly, we observed CDR-H3 length-dependent usage of human IGHD3 and IGHD2 germline genes. As CDR-H3 goes longer, the usage of IGHD3 and IGHD2 increased from 24.87% and 14.60% at 7 amino acids length to 57.52% and 30.70% at 23 amino acids length, respectively (Fig. 3a). Such CDR-H3 length-dependent usage of human IGHD2 and IGHD3 is independent of health conditions, as we observed the same germline usage patterns in both healthy donors (Fig. 3a, upper panel, n=8,359) and DENV acutely infected patients (Fig. 3a, lower panel, n=68,067).

When CDR-H3 is of 7 amino acids length, in approximately 17% of the antibodies in the immune repertoire, the CDR-H3 diversity is likely generated by direct rearrangement of IGHV to IGHJ. We could not rule out the possibility that IgBlast might fail to identify the IGHD region, especially when the IGHD is short and extensively mutated. As CDR-H3 loop length increases, the percentage of direct VH-JH recombined sequences gradually decreased to 0%. At 14 amino acids CDR-H3 length or longer, direct rearrangement of IGHV to IGHJ was very rarely observed. This indicates that in antibodies with longer CDR-H3 loops, IGHD gene segments are critical components of CDR-H3 diversity. More intriguingly, human IGHD7 germline gene was rarely used in the human immune repertoire (Fig. 3a).

Given the previous observation of CDR-H3 length-dependent usage of IGHJ4 and IGHJ6 germline genes, we were wondering about potential biases in human IGHD rearrangements to human IGHJ. To answer this question, we turned to the IGHV3-23*01 dataset of Danish origin. As the human IGHV germline usage is fixed to human IGHV3-23*01, the DH-JH recombination analysis is not impacted by the choices of IGHV. We observed CDR-H3 length-dependent preferential rearrangement of IGHD3/IGHD2 gene segments to human IGHJ6 germline gene (Fig. 3b). Of 2,125 VH3-23-DH3/DH2 antibody sequences that don’t use IGHJ4, a strong correlation was observed between CDR-H3 length and DH3/DH2 germline usage (Fig. 3b, lower right). In contrast, of 1,116 VH3-23-DH3/DH2 antibody sequences that don’t use IGHJ6, a strong negative correlation was observed between CDR-H3 length and DH3/DH2 germline usage (Fig. 3b, upper right). These data suggest that DH3/DH2 was preferentially rearranged to IGHJ6 as CDR-H3 goes longer. Such a biased preference was not observed with other human IGHD germline genes. Our observation is consistent with a previous study by Briney B.S. et al in the analysis of a B-cell receptor repertoire of a HIV infected patient cohort (26). This suggests the preferential rearrangement of DH3/DH2 to JH6 is universal in CDR-H3 diversity generation. Such a preference is likely required by the intrinsic need to stabilize the long CDR-H3 loops.
Choice of JH only partially contributes to the solvent accessible surface area of an antibody’s paratope

We then moved on to determine if preferentially rearranged DH/JH combinations can be used to guide the design of rationally designed antibody libraries. An essential question to answer is what parameters we can control to finely tune the size of the surface area and the physicochemical property of an antibody’s paratope. The IGHV3-23*01 dataset was again used in this analysis. To avoid bias introduced by irregular framework, all antibody sequences used in this analysis were manually validated to be free of framework insertions or deletions. Similarly, to avoid bias introduced by skewed CDR-H3 amino acid usage, all IGHV3-23*01 sequences were manually validated on the diversity to remove redundant sequences or sequences with irregular CDR loops, framework deletions or insertions. First, we determined the relationship between CDR-H3 length and the solvent accessible surface area (SASA) of the antibody paratope. Simulated models were built by pairing human IGHV3-23*01 variable region, of various CDR-H3 length ranging from 7aa to 23aa, with human IGKV1-39*01/IGKJ2 variable region. IGKV1-39 was chosen because the VH3-23/VK1-39 pairing was considered a very productive VH/VL pairing with good drug like properties (34,35). The SASA values were calculated based on Kabat numbering of the antibody CDR loops. In this analysis, the only variables are the CDR-H3 length and the choices of various JH. As shown in Fig. 4a, a positive correlation was observed. As CDR-H3 went longer, the SASA of antibody paratope grew bigger from 3008.88±116.10 Å² of 7aa length to 3666.42±224.78 Å² of 23aa length. Interestingly, when CDR-H3 is longer than 22 amino acids, the SASA remained stagnant. This observation needs to be further validated with more IGHV3-23*01 sequences of extra CDR-H3 length. In above VH3-23/VK1-39 pairings, the VK1-39 framework was fixed to JK2. We were wondering if the choice of JK would change the SASA of VH3-23/VK1-39 paired antibodies. In this analysis, we chose a dataset with a fixed 10aa length of CDR-H3 and IGKV1-39*01 rearranged to various IGKJ germline genes. Fig. 4b showed the statistical analysis of VH3-23/VK1-39 antibody paratopes with various human JKs. No statistically significant difference was observed among the five JK groups (p>0.05). This indicates that choices of JK minimally impacted the SASA of an antibody’s paratope. We then paired a set of VH3-23*01 of 14aa length with various kappa light chain germline genes rearranged to JK2. Fig. 4c showed that VK pairing statistically significantly impacted the SASA of antibody paratopes (p<0.001). Choices of distal or proximal VK germline also statistically significantly impacted the SASA of antibody paratopes (p<0.001). In a pairwise analysis, only statistically significant difference was observed between VK3-11, VK3-20 and their distal counterparts (Fig. 4d, p<0.001). To determine the impact of JH on the SASA of antibody paratope, we paired VH3-23*01 of 14aa length, rearranged to various JH, with VK1-39*01/JK2. As shown in Fig. 4e, in a pairwise analysis,
statistically significant difference was observed between JH4/JH5 and JH6 (Fig. 4e, p<0.001). This indicates that JH6 significantly contributed to the SASA of antibody paratope.

In this analysis, using a set of precisely controlled data with minimal variables, we confirmed the commonly accepted knowledge in the field that antibody paratope is mainly impacted by the CDR-H3 length and VH/VL germline pairings. We also observed that JK, but not JH, minimally impacted the overall SASA of antibody paratope.

**DISCUSSION**

The CDR-H3 assembly mechanism, regulation and influence on antibody diversity were extensively reviewed in the field (24,36,37). Recent advances in next generation sequencing and immune repertoire mining helped us to understand the antibody diversity to a degree to enable man-made variable genes for antibody discovery (7). In this study, our goal is to understand the components of CDR-H3 diversity in a precisely controlled manner, hoping the knowledge gained can help rational design of next generation antibody libraries. Unlike many other studies mining immune repertoire, we utilized a well validated IGHV3-23*01 dataset (15,16,27,28) to keep variables minimal in our study.

We observed CDR-H3 length-dependent usage of IGHJ4 and IGHJ6 germline genes in the immune repertoire. The increased usage of IGHJ6 in antibodies with long CDR-H3 loops validated a well recognized feature of tyrosine enrichment at the junction of DH-JH recombination (38). Among all human JH gene segments, JH6 is the longest (39), which explained the increased usage of IGHJ6 germline genes in antibodies with long CDR-H3. Likely, tyrosine residues were introduced repeatedly for the continued diversification at the DH-JH junction as CDR-H3 loops become longer. Tyrosine, on one side, is one of the dominant CDR-H3 residues critical for antibody binding specificity (28,38). On the other side, tyrosine-rich motifs were frequently observed in amyloidogenic proteins like β2-microglobulin (40,41), suggesting an important role of tyrosine in controlling the integral structure of proteins. Tyrosine motifs at the DH-JH junction likely are not involved in antigen-antibody interactions. In antibodies with long CDR-H3 loops, we hypothesize that tyrosine residues at the DH-JH junction serve a structural role to help assembling and stabilizing the CDR-H3 loop. Such role is currently under-appreciated in the field and it needs to be tested in the future.

A highly skewed DH3-JH6 rearrangement was observed in this study. It correlates well with CDR-H3 loop length but it is independent of ethnic background and health conditions. As both DH3 and JH6 are the longest in their respective families, it is not surprising to observe the biased DH3-JH6 rearrangements in antibodies with long CDR-H3 loops. Interestingly, biased DH3-JH6 rearrangements were also observed in the comparative immune repertoire analysis of a
humanized rodent model (39), even though the humanized rodents tend to prefer short DH genes. These suggest DH3, as well as DH2 (28,30), serves to stabilize the rigid and protruding CDR-H3 loops by rearrangement with JH6 in antibodies with long CDR-H3 loops. Antibodies with non-canonical cysteine residues in the CDR-H3 were frequently observed in human broadly neutralizing antibodies against HIV-1, human hepatitis C virus (HCV), human cytomegalovirus (HCMV) and human influenza virus (42-45). This suggests the stabilized CDR-H3 loop may contribute to unique binding attributes like high affinity or broader binding specificity towards difficult to reach epitopes on the virions (28). Such unique structural and binding properties in antibodies with longer CDR-H3 loops could be of interest in the next generation synthetic human antibody libraries to help fight emerging pathogens like COVID-19.

Above findings inspired us to perform the spatial analysis of the antibody paratope. By computational simulations, in a precisely controlled manner, we defined the spatial patterns of antibody paratope and the influencing parameters like CDR-H3 length, pairing light chain germline genes, choices of JK and JH etc. The conformations of CDR-H3 loops are pivotal elements in CDR-H3 diversity design. In this study, the CDR-H3 conformation was not considered a key parameter due to the lack of well-recognized predicative tools. We’ll keep improving the analysis when it is technically possible and will expand the analysis to all human VH germline genes. The knowledge obtained will help to guide the rational design of next generation antibody libraries. In the new library design, IGHJ gene fragments will be mixed together proportionally to faithfully reflect the CDR-H3 length-dependent germline usage. CDR-H3 diversity synthesis can be achieved via TRInucleotide technology so that the amino acid distribution pattern follows that of natural immune repertoire. We hope this will properly address the poor antibody developability issue the synthetic diversity field is facing. Targeting a pre-defined antigenic epitope, a high quality antibody library of minimal size and diversity can be designed to facilitate antibody discovery on demand.

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**DISCLOSURE OF INTEREST**

The authors declare no conflict of interest.
AUTHOR CONTRIBUTIONS
Conceptualization, L.C., C.Y. and Y.S.; methodology, L.C. and Y.Y.; validation, K.Y. and H.W.; formal analysis, L.C., K.Y and H.W.; data curation, K.Y. and H.W.; writing—original draft preparation, L.C., R.W., H.W. and Y.K; writing—review and editing, L.C., C.Y., Y.Y. and Y.S.; visualization, H.W.; supervision, R.W., Y.Y. and L.C.; All authors have read and agreed to the published version of the manuscript.

DATE AVAILABILITY STATEMENT
The following data were used in this study: the set of VH3-23*01 sequences (European Molecular Biology Laboratory accession nos. AM076988–AM083316) and DENV cohort sequences (the Bi-oProject accession number PRJNA205206).

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**FIGURE LEGENDS**

**Figure 1.** Antibody heavy chain CDR-H3 length-dependent usage of human IGHJ4 and IGHJ6 germline genes in antibody sequences derived from (A) healthy and viral-infection naive cohort of Danish origin; (B) and (C) Nicaraguan individuals acutely exposed to Dengue virus; (D) the accompanying healthy cohort of the DENV study.
Figure 2. The CDR-H3 length distribution of different human IGHJ antibodies. (A) Combined; (B) individualized.

Figure 3. (A) The CDR-H3 length-dependent usage of human IGHD2 and IGHD3 germline genes; (B) IGHD3/IGHD2 were preferentially rearranged to human IGHJ4/IGHJ6 germline genes as CDR-H3 length goes longer.
**Figure 4.** (A) The SASA of an antibody paratope is determined by many factors, including CDR-H3 length; (C)(D) VH/VL pairing; (B)(E) Rearrangement to JK or JH only partially impact the size of SASA.