**V<sub>H</sub> replacement footprint analyzer-I, a Java-based computer program for analyses of immunoglobulin heavy chain genes and potential V<sub>H</sub> replacement products in human and mouse**

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**METHODS ARTICLE**

**INTRODUCTION**

Antibodies are the effective molecules in the adaptive immune system to recognize specific antigens and combat bacterial and viral infections, as well as malignant cells (1). To recognize almost unlimited numbers of antigens, a tremendously diversified repertoire of antibody specificities is generated through V(D)J gene rearrangement (1, 2). V(D)J recombination is catalyzed by the recombinase activating gene products (RAG1 and RAG2) that recognize recombination signal sequences (RSS) (3–5). Functional RSS consists of a heptamer (CAGTGTG), a nonamer (GTTTTTTGT), and a non-conserved spacer region of 12 or 23 base pairs in between (6, 7). Efficient recombination occurs only when a pair of RSSs with 12- and 23-bp spacers, known as the 12/23 rule (7, 8). During V(D)J recombination, the RAG1 and RAG2 complexes first nick between the heptamer and the coding sequence, leaving a blunt signal end and a hairpin sealed DNA coding end (7–9). The two signal ends are usually fused to form a signal junction activating gene products (RAG1 and RAG2) that recognize recombination signal sequences (RSS) (3–5). Functional RSS consists of a heptamer (CAGTGTG), a nonamer (GTTTTTTGT), and a non-conserved spacer region of 12 or 23 base pairs in between (6, 7). Efficient recombination occurs only when a pair of RSSs with 12- and 23-bp spacers, known as the 12/23 rule (7, 8). During V(D)J recombination, the RAG1 and RAG2 complexes first nick between the heptamer and the coding sequence, leaving a blunt signal end and a hairpin sealed DNA coding end (7–9). The two signal ends are usually fused to form a signal junction.

The coding region, a short stretch of nucleotides from the previous rearranged V<sub>H</sub> gene can be retained in the newly formed V<sub>H</sub>–D<sub>H</sub> junction as a “footprint” of V<sub>H</sub> replacement. Such footprints can be used as markers to identify Ig heavy chain (IgH) genes potentially generated through V<sub>H</sub> replacement. To explore the contribution of V<sub>H</sub> replacement products to the antibody repertoire, we developed a Java-based computer program, V<sub>H</sub> replacement footprint analyzer-I (V<sub>H</sub>RFA-II), to analyze published or newly obtained IgH genes from human or mouse. The V<sub>H</sub>G RFA-I program has multiple functional modules: it first uses service provided by the IMGT/TV-QUEST program to assign potential V<sub>H</sub> replacements in correlation with publications, keywords, or V<sub>H</sub> J<sub>H</sub> gene usages, and mutation status; it can further analyze the amino acid usages encoded by the identified V<sub>H</sub> replacement footprints. In summary, this program provides a useful computation tool for exploring the biological significance of V<sub>H</sub> replacement products in human and mouse.

**Keywords:** V<sub>H</sub> replacement, RAG, B cell, IgH gene, IGH sequencing, VDJ rearrangement

V<sub>H</sub> replacement occurs through RAG-mediated secondary recombination between a rearranged V<sub>H</sub> gene and an upstream unrearranged V<sub>H</sub> gene. Due to the location of the cryptic recombination signal sequence (cRSS, TACTGTG) at the 3′ end of V<sub>H</sub> gene coding region, a short stretch of nucleotides from the previous rearranged V<sub>H</sub> gene can be retained in the newly formed V<sub>H</sub>–D<sub>H</sub> junction as a “footprint” of V<sub>H</sub> replacement. Such footprints can be used as markers to identify Ig heavy chain (IgH) genes potentially generated through V<sub>H</sub> replacement. To explore the contribution of V<sub>H</sub> replacement products to the antibody repertoire, we developed a Java-based computer program, V<sub>H</sub> replacement footprint analyzer-I (V<sub>H</sub>RFA-II), to analyze published or newly obtained IgH genes from human or mouse. The V<sub>H</sub>G RFA-I program has multiple functional modules: it first uses service provided by the IMGT/TV-QUEST program to assign potential V<sub>H</sub> replacements in correlation with publications, keywords, or V<sub>H</sub> J<sub>H</sub> gene usages, and mutation status; it can further analyze the amino acid usages encoded by the identified V<sub>H</sub> replacement footprints. In summary, this program provides a useful computation tool for exploring the biological significance of V<sub>H</sub> replacement products in human and mouse.

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Immunoglobulin (Ig) gene V(D)J recombination occurs in a step-wised manner during early B cell development (2, 11, 12). Normally, D<sub>H</sub> to J<sub>H</sub> rearrangement occurs before V<sub>H</sub> to DJ<sub>H</sub> rearrangement on one of the Ig heavy chain (IgH) alleles, followed by V<sub>k</sub> to J<sub>k</sub> and then V<sub>κ</sub> to J<sub>κ</sub> rearrangement on the Ig light chain (IgL) loci (2, 11, 12). Due to the random nature of RAG-mediated rearrangements, approximately two thirds of the rearranged Ig genes may be out of the reading frame, which cannot produce functional Ig peptides (13). Functionally rearranged IgH genes may produce IgH peptides that fail to pair with surrogate or functionally rearranged conventional IgL chains (13). Moreover, functional Ig genes may encode self-reactive antibodies (14–16). In order for these B cells to survive, early B lineage cells retain the ability to reinitiate RAG-mediated secondary recombination.
to alter the rearranged Ig genes, a process known as receptor editing (14–16). Receptor editing of the IgL genes would be easy to envision because the organization of the mouse and human Igκ locus enables continuous secondary recombination by joining an upstream Vκ gene segment with a downstream Jκ gene segment, leading to the deletion of the previously formed VκJκ joint (14, 15). B cells also have a default option to delete the entire Igκ locus and initiate de novo rearrangement of the Igκ locus (14, 15). Secondary rearrangement on the Igκ locus is conceptually difficult, because the primary rearrangement deletes all DJκ gene segments flanked by 12-bp RSSs. The remaining upstream VH and downstream JH gene segments are flanked by 23-bp RSSs, which are difficult to recombine (17). Nevertheless, secondary Igκ rearrangement to generate functional Igκ genes from non-functional Igκ rearrangements was observed in mouse pre-B cell lines even before the discovery of the RAG genes (18, 19). Comparison of the non-functional and newly formed functional Igκ rearrangements led to the identification of a cryptic RSS (cRSS), TACTGTTG motif, embedded at the 3′ end of the rearranged VH genes (18–20). Based on these observations, a novel VH to VDJH recombination mechanism was proposed as VH replacement (18–20). Subsequent studies demonstrate that VH replacement is employed to rescue pro B cells with two alleles of non-functional Igκ rearrangements (17, 21), to edit Igκ genes encoding anti-DNA antibodies (22–24), and to change the knocked-in Igκ gene encoding monoclonal anti-NP antibodies and to generate a diversified antibody repertoire (25, 26).

VH replacement changes almost the entire VH coding region (27). However, due to the location of the cRSS, a short stretch of nucleotides from the previously rearranged VH gene may be remained at the newly formed V-D junctions after each round of VH replacement (16, 27, 28). Such remnants can be used as footprints to trace the occurrence of VH replacement and to identify potential VH replacement products (16, 27, 28). Our previous analysis of 417 human Igκ sequences indicated that VH replacement contributes to the diversification of the primary human antibody repertoire (27). This conclusion was supported or argued by subsequent analyses of Igκ genes from human or mouse (29–32). Most of these analyses were based on relatively small number of Igκ gene sequences or sequences from few individuals. A comprehensive analysis of large numbers of Igκ gene sequences is required to fully address the biological significance of VH replacement in antibody repertoire diversification.

Analysis of Igκ gene sequences obtained from B cells of different developmental stages or in different disease states provided tremendous information regarding the development and selection of the antibody repertoire. Currently, there are about 61,000 human and 17,000 mouse Igκ gene sequences available at the NCBI database. With the advanced next generation sequencing (NGS) technology, millions of Igκ gene sequences can be easily obtained (33–35). To identify potential VH replacement products in a large number of Igκ gene sequences and to explore the biological significance of VH replacement products in different diseased subjects in human and mouse, we developed a Java-based computer program, named VH replacement footprint analyzer-I (VH RFA-I).

**MATERIALS AND METHODS**

**COMPUTER HARDWARE AND SOFTWARE REQUIREMENTS**
The VH RFA-I program can be operated on any desktop computer with Microsoft Windows, Mac OS X, or different Linux operating system. It requires Java runtime environment (jre) 1.6 or higher version for operating and Microsoft Excel 2007 or higher version for data export.

**SOFTWARE DEVELOPMENT**
The VH RFA-I program was developed using the NetBeans 7.01 IDE with Java development kit (JDK) and tested under Windows, Mac OS X, and Ubuntu Linux. Two free Java libraries were used, a csv parser library1 and an Excel parser library2.

**REFERENCE HUMAN AND MOUSE V_H GENE SEQUENCES**
The reference human and mouse VH germine gene sequences used for generating the VH replacement footprint libraries were downloaded from the IMGT database and listed in Table S1A,B in Supplementary Material.

**DESCRIPTION OF THE HUMAN AND MOUSE IgH GENE SEQUENCE TRAINING DATA SETS**
Two sets of IgH gene sequences, one from human and the other from mouse, were used in the initial testing and training of the VH RFA program. The 417 human Igκ gene sequences were from a study that examined whether peripheral blood B cells of preterm infants show similar restrictions as fetal liver B cells (36). These sequences had been used in our previous analysis to manually identify potential VH replacement products (27). These sequences are referred as the Z417 test sequences in this study and the results of Z417 test sequences are shown at each step of the analysis.

**RESULTS**

**AN OVERVIEW OF THE V_H RFA-I PROGRAM AND FUNCTIONAL MODULES**
As shown in the workflow of the VH RFA-I program (Figure 1), the VH RFA-I program consists of multiple functional modules for the analysis of IgH genes and for the identification and analysis of VH replacement products in published or newly generated IgH gene sequences from human or mouse. The VH RFA-I program is a single executable Jar file, which can be operated on any computer operating platform. The VH RFA-I program can be launched by double click of the executable Jar file, VH Replacement Analyzer-I, which opens the main interface of the VH RFA-I program (Figure 2). All the functional modules are listed as clickable bars in the main interface. The detailed functions of these modules are discussed below.

**THE FASTA FORMAT CONVERTER**
The FASTA Format Converter was designed to convert GenBank files to FASTA files. It can be operated by clicking the first functional bar, I have a GeneBank File and would like to convert it into FASTA format (Figure 2). This function module converts IgH gene sequences downloaded from the NCBI database from GenBank

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1http://opencsv.sourceforge.net/
2http://jexcelapi.sourceforge.net/
format to FASTA format, which can be used for subsequent analysis. This file converter differs from other converters in that it will eliminate entries that do not contain actual sequence data. You can specify the locations of the input GenBank file and the output FASTA file in the pop-up window.

RETRIEVE \( V_H \), \( D_H \), AND \( J_H \) GENE ASSIGNMENT RESULTS FROM IMGT
The \( V_H \)RFA-I program uses the IMGT/V-QUEST program to assign the potential \( V_H \), \( D_H \), and \( J_H \) germline genes. In order to handle a large number of IgH gene sequences, we designed the IMGT Downloader functional module (Figure 3) to automatically send IgH sequences in batches of 50 sequences in FASTA format to the IMGT/V-QUEST program for analysis and export the \( V_H \), \( D_H \), and \( J_H \) gene assignment results as Excel files to a user specified location (Figure 3). The HTTP requests are sent to “http://imgt.org/IMGT_vquest/vquest.” Dependent on the speed of the internet, the \( V_H \)RFA-I program can analyze every 50 IgH sequences within 1 min.

For each analysis, the user can specify the species of IgH sequences (Figure 3A), number of accepted \( D_H \) germline gene segments (Figure 3B), number of accepted mutations within the 3′ \( V_H \) gene (Figure 3C), \( D_H \) gene (Figure 3D), and 5′ of \( J_H \) gene (Figure 3E). To be analyzed, IgH sequence files can be selected from a local computer and the downloaded result files can be directed to a local computer (Figures 3F, G, respectively). The process will be started after clicking the functional bar: upload sequences and start downloading Excel Files (Figure 3H). The downloading process will be indicated in the Download Progress window (Figure 3I). If there is any mistake during the file uploading and downloading process, a note will be posted on the Message Board (Figure 3J). In the test run of the Z417 test IgH sequences, the V-QUEST analysis results were deposited at a user specified local hard drive with 50 sequences per file (Figure 3K). The results contain all the information from the V-QUEST (Figure 3L). After
FIGURE 3 | The IMGT downloader. Diagram shows the interface of the IMGT Downloader. The IMGT Downloader allows users to use the IMGT/V-QUEST program to analyze large numbers of IgH gene sequences by uploading IgH sequences and downloading V-QUEST analysis results to a local computer. The user can specify human or mouse sequences (A), numbers of D\(\beta\)H genes (default = 1) (B), number of accepted mutations in the 3\' V\(\beta\) region (C), D\(\beta\) region (D), and 5\' J\(\beta\) region (E). After these settings, the user can upload the IgH sequences (in FASTA file) (F) and specify the directory where the downloaded V-QUEST analysis Excel files can be stored (G). The analysis can be started by clicking the Upload sequences and start downloading Excel Files bar (H). The analysis progress (I) and message during the analysis (J) will also be shown. The V-QUEST analyses results of the test sequences are downloaded to a user specified location (K). The detailed results of sequence 1–50 are shown in the V-QUEST format (L).

IDENTIFICATION OF V\(\beta\) REPLACEMENT FOOTPRINTS

The footprint analyzer module uses the sequence analysis results retrieved from the IMGT/V-QUEST program to identify potential V\(\beta\) replacement products. Basically, it searches for potential V\(\beta\) replacement footprint motifs within the N1 and N2 regions of each IgH sequence and export all the analysis results in a single CSV file. The user can specify the species of sequences to be analyzed (Figure 4A, with the Z417 test sequence files), uploaded the files to the program (Figure 4B), select the different V\(\beta\) replacement footprint library (Figure 4C), and specify the minimum length of the V\(\beta\) replacement footprints (Figure 4D).
The Footprint Analyzer module is built into the program. It does not have a graphic user interface (GUI) but gets its parameters from and is invoked by the Footprint Analyzer (Figure 4C). It loads IMGT germline references (Table S1A,B in Supplementary Material), extracts nucleotide sequences after the cRSS (TACTGTG motif) to generate a library of potential VH replacement footprints with different length. The user has five options to choose the source of the VH replacement footprints library by selecting “only functional genes,” “only non-functional genes,” “all genes,” “functional less non-functional genes,” or “non-functional less functional genes” (Figure 4C). Potential VH replacement footprints for both human and mouse are listed in Table S2 in Supplementary Material, as grouped by lengths. During the primary recombination, the 3′ end of VH genes can be
trimmed off by exonuclease activities after processing the coding end hairpin structure. During the V<sub>H</sub> replacement process, the 5′ end of such footprints could also be trimmed off by exonuclease. The Footprint generator can generate a library of potential V<sub>H</sub> replacement footprints with 3–12 bp in length according to the user’s selection of the Minimum Signature Length in the combo box (Figure 4D).

The Footprint Analyzer starts to search the longest motifs and then to the shorter motifs based on the user’s selection. The user can specify the location of the output result file (Figure 4E) and also save the footprint library used for each analysis (Figures 4F, G). The analysis progress will be indicated in the Analyzer Progress window (Figure 4K). The user also has the option to exclude GGG sequences by checking the checkbox (Figure 4H). The results will be saved in Excel format. As shown in Figure 4L, potential V<sub>H</sub> replacement footprint with user specified length (5-mer) were identified in both N1 regions (N1 signatures) or N2 regions (N2 signatures) together with the V<sub>H</sub>, D<sub>H</sub>, and J<sub>H</sub> gene assignment results.

THE PUBLICATION ANALYZER
All the IgH gene sequences deposited at the NCBI database are linked with their original publications with all the information. To explore the biological significance of the identified V<sub>H</sub> replacement products, we designed a special Publication Analyzer functional module. The Publication Analyzer groups IgH sequence analysis results according to their PubMed identifications (PMID). To do so, the user needs to select the original GenBank file (Figure 5A) and the V<sub>H</sub> replacement analysis results to start the analysis (Figure 5B). In the output results, the V<sub>H</sub> replacement products results will be linked with the PubMed ID of the original IgH sequence (Figure 5C). Under the GenBank ID pull down manual, the user can open the Abstract pages of selected PubMed IDs (maximum of five) (Figure 5D); copy the GenBank IDs from selected publications to the clipboard (Figure 5E); save GenBank records of selected publications (Figure 5F); and save the V<sub>H</sub> replacement footprint analysis results of selected publication, as generated by the Footprint Analyzer (Figure 5G). The Publication Analyzer can also provide the original footprint result file for the selected publications (Figure 5H).

THE KEYWORD ANALYZER
The Keyword Analyzer groups sequence IDs according to their linked keywords from the GenBank files. The Keyword Analyzer will use the footprint analysis result file (Figure 6A), GenBank file containing the original sequences to generate the footprint analysis
result file (Figure 6B), keyword analysis result file (Figure 6C). After starting the analysis (Figure 6D), the program will parse the DEFINITION, KEYWORDS, and FEATURES sections of the GenBank record for each IgH gene sequence. An ID will be assigned to a keyword if the GenBank entry contains the keyword. Depending on the availabilities of all VDJ assignments, N1 footprints, or N1 footprints, it also assigns IDs to these bins within each keyword. Same as the File Format Converter, the Keyword Analyzer ignores GenBank records without actual sequence data. As such analysis takes substantial amount of time when the GenBank file is complex, a log window is provided to monitor the process (Figure 6E). For examples, all the keywords associated with the Z417 test sequences from the NCBI database are listed in Column A, Keyword (Figure 6F).

ASSEMBLE THE KEYWORD GROUP

The Keyword Group Picker visualizes results from keyword analysis and footprint analysis, allowing the user to select group of keywords of interest and output the related footprint analysis results. This functional module analysis provides the user an opportunity to manually inspect a subset of sequences for particular studies. After selecting the footprint analysis result file (Figure 7A) and choosing the keyword analysis result file (Figure 7B), the results ordered by keywords ascending alphabetically and case insensitive
FIGURE 7 | The keyword group picker. Diagram shows the interface of the Keyword Group Picker. (A) Textbox to select the footprint analysis result file. (B) Textbox to select the keyword analysis result file. (C) Button to move selected rows from (F) to (J). (D) Textbox for entering search string to locate keywords in (F). (E) Button to start locating keywords containing string in (D). (F) Window area containing contents of the keyword analysis result file. (G) Button to move selected rows from (J) to (F). (H) Button to select a keyword analysis result file so that keywords can be isolated, to repeat a previous pick. (I) Button to select keywords associated with entered GenBank ID. (J) Window area displaying the selected keywords. (K) Combo box to select the type of sequences to output. (L) Checkbox to indicate intention to dump footprint analysis result into a single sheet. (M) Textbox for entering the sheet name if (L) is selected. (N) Textbox for choosing the output file. (O) Button to start the pick/isolation process.

will be shown in the table below (Figure 7F). Typing inside the table with the first letter of any keyword will allow quick location of the keywords. The user can also select specific keywords (Figure 7C) to move them from the upper window (Figure 7F) to the lower window (Figure 7J) for further analysis or deselect the keywords (Figure 7G). Pressing Enter (Figure 7D) or clicking the functional bar (Figure 7E) will select all keywords containing strings. The user can also select keywords from a picked file (Figure 7H) or select keywords according to their sequence IDs (Figure 7I). The user needs to specify the name and location of the output result file (Figure 7N). There are four options for the output results, which can be specified by the user (Figure 7K): “all sequences” will select footprint analysis results in all the keywords listed in the lower window (Figure 7J); “Screened Sequences” will select those with all V, D, and J assignments; “N1 Sequences” will select those with footprints in the N1 region; “N2 Sequences” will select those with footprints in the N2 region. The format of the output results can also be specified by checking the checkbox (Figure 7L) and providing a name (Figure 7M), in which the results will be exported as an Excel file in which the first sheet contains statistics, the second sheet contains the merged footprint analysis results, and the third sheet contains the results as...
shown in the lower window (Figure 7). Otherwise, the footprint analysis results will be exported in separate sheets according to keywords. The analysis can be started by clicking the Start Output bar (Figure 7O).

THE AMINO ACID CONTRIBUTION ANALYZER

The Amino Acid Contribution Analyzer analyzes the IgH CDR3 amino acid sequences and identifies the amino acids contributed by the identified V_H replacement footprints in the N1 or N2 regions. If the input file is an Excel file, it iterates through all footprint analysis result sheets and generates four sheets: "N1-" sheet contains sequences with N1 footprint; "N2-" sheet contains sequences with N2 footprints; "N1AAs-" contains results with amino acids contributed by N1 regions; "N2AAs-" contains results with amino acids contributed by N2 regions. An amino acid is considered to be contributed by a V_H replacement footprint if the first or second nucleotide of its codon is encoded by the footprint. The user can select the Input Files (Figure 8A) from all the analyzed results, such as Excel files generated by the Keyword Group Picker, or CSV files generated by the Footprint Analyzer. The user also needs to specify the location of the output file (Figure 8B). The analysis can be started by clicking the "Start Amino Acid Usage Analyzer" bar (Figure 8C). As an example, the amino acids contributed by the identified footprints in Z417 test sequences are listed following the N1 signature (Figure 8D).

THE AMINO ACID USAGE CALCULATOR

The Amino Acid Usage Calculator analyses the usages of amino acid within the N1 regions. The user can select the input files to be analyzed (Figure 9A) and the results will be shown in the window (Figure 9B) or copied to the clipboard (Figure 9C). The user needs to specify a location for the output result file (Figure 9D). The analysis can be started by clicking the "Calculate" bar (Figure 9E). As an example, the results of amino acids usage in the N1 region of the Z417 test sequences are shown in Excel format (Figure 9F). Such results can be easily converted to different type of displays for

FIGURE 8 | The amino acid contribution analyzer. Diagram shows the interface of the Amino Acid Contribution Analyzer. (A) Textbox for selecting the footprint analysis result file. (B) Textbox for selecting the output file. (C) Button for starting the analyzer. (D) A sample result showing the V_H replacement footprints and amino acid residues encoded by the identified V_H replacement footprints the test sequences.

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The amino acid usage calculator

The amino acid usage calculator is shown in Figure 9G. The amino acid usage is presented in a bar graph in Figure 9G.

The VDJ Frequency Calculator

The VDJ Frequency Calculator calculates the frequencies of V, D, J gene usages and IgH gene CDR3 length. Input Files can be selected from VH replacement footprint analysis result file in either CSV format or Excel format, as output by the Footprint Analyzer or the Keyword Group Picker, respectively. If the input files are in Excel format, it will populate the combo box with names of sheets containing VH replacement footprint analysis results or copied to the clipboard. The user needs to specify the location of the output result file. The output results can be ranked according to the VH gene family or the VH gene name. The analysis can be started by clicking the Calculate bar. As an example, the results of the usages different VH genes in the Z417 test sequences were calculated; the frequencies of VH replacement footprints in the N1 or N2 regions of IgH genes using each VH germline gene are also listed in the output file (not shown); and the distribution of IgH genes with different CDR3 length was also calculated.

The Clonal Stripper

To focus on analysis of the unique IgH sequences in any dataset, we designed the Clonal Stripper functional module. The Clonal Stripper removes redundant sequences based on their identical CDR3 regions. Input files can be selected from the results of either the Footprint Analyzer or the Keyword Group Picker, in CSV or Excel format, respectively. The name of the analyzed result files will be shown in the window or copied to the clipboard. The user needs to specify a location for the output result file. After stripping, the results will be saved as a CSV file in the same format as the output result by the Footprint Analyzer. Within the Z417 test sequences, there are three repeated sequences, which can be identified and eliminated by the clonal striper function (data not shown).

The GenBank File Tailor

After stripping off IgH sequences with identical CDR3 regions, the GenBank File Tailor function module reanalyze the GenBank files according to stripped sequence files to get rid of the repeated sequences from the GenBank record IDs and save the rest unique sequences into a new FASTA file.

The Mutation Analyzer

The Mutation Analyzer uses the results retrieved from the IMGT/V-QUEST program by the IMGT Downloader to calculate the number of mutations within the VH region and mutation rate. The analysis can be started by clicking the “Start Analyzer” bar, and the progress will be indicated in...
FIGURE 10 | The VDJ frequency calculator. Diagram shows the interface of the VDJ Frequency Calculator. (A) Button to select the input footprint analysis result file. (B) Combo box for selecting the sheet for processing, when an Excel file is selected as the input file. (C) Button to copy the value in (B) to clipboard. (D) Button to choose the output file. (E) Radio button group to select the sorting criterion for the output results. (F) Button to start the calculator. (G) The output results of VH gene usage in the test sequences were presented as a bar graph. (H) Distribution of the Z417 test IgH gene sequences with different CDR3 lengths.

FIGURE 11 | The clonal stripper. Diagram shows the interface of the Clonal Stripper. (A) Button to choose the input footprint analysis result file, which can be CSV file generated by the footprint analyzer or Excel file generated by the Keyword Group Picker. (B) Combo box for selecting the sheet for analysis, if an Excel file is selected in (A). (C) Button to copy the name of selected sheet to the clipboard. (D) Button to choose the output file. (E) Button to start the stripping process.
Footprint Analyzer

the window in Figure 13F. As an example of the output results, the position of the mutation within the V<sub>H</sub> gene, the length of the V<sub>H</sub> gene, the mutation number, and the mutation rate of each IgH gene are listed in the Excel file (Figure 13G).

THE MUTATION MATCHER

The Mutation Matcher recalculates the mutation analysis results of a subgroup of V<sub>H</sub> replacement analysis results according to the results obtained from the Mutation Analyzer. Input file can be selected from the result files from the Footprint Analyzer or the Keyword Group Picker (Figure 14A). For the latter, names of sheets containing footprint analysis results will populate the combo box (Figure 14B) or copied to the clipboard (Figure 14C). The mutation file should contain the mutation results for all the sequences (Figure 14D). The user needs to specify a location for the output result file (Figure 14E) and a maximum mutation rate (Figure 14F). Analysis can be started by clicking the Calculate bar (Figure 14G). An example of the output result is shown in the Excel format (Figure 14H).

THE FOOTPRINT RESULT SPLITTER

The Footprint Result Splitter reanalyzes the footprint analysis results according to their V<sub>H</sub>, D<sub>H</sub>, or J<sub>H</sub> genes. The input files (Figure 15A) should be in CSV format, as generated by the Footprint Analyzer. The user needs to specify the location of the output result files (Figure 15B). The results can be split based on the V<sub>H</sub> genes, D<sub>H</sub> genes, or the J<sub>H</sub> genes (Figure 15C) and the analysis can be started by clicking the Split bar (Figure 15D). The results will be saved as individual files for each germline V<sub>H</sub> gene in user specified location, as shown in Figure 15E. For example, the IGHV1–69 file contains the results of all the IgH genes using the V<sub>H</sub>1–69 germline gene (Figure 15F).

DISCUSSION

In summary, we have developed a Java-based computer program, V<sub>H</sub>RFA-I, to analyze large number of IgH gene sequences from human or mouse origin and to identify and analyze potential V<sub>H</sub> replacement products. The different functions of the V<sub>H</sub>RFA-I program are described in this report along with the results at each step of analysis using the Z417 test sequences. This program will be especially useful to explore the biological significance of V<sub>H</sub> replacement products in human and mouse. Currently, there is no such program available.

We have included multiple functional modules in this program to analyze the frequencies of V<sub>H</sub> replacement products according to their publication, keywords, V<sub>H</sub>, D<sub>H</sub>, J<sub>H</sub> gene usages, and mutation status. Using such functions, we can determine the distribution of V<sub>H</sub> replacement products in IgH genes derived from different diseased subjects. The V<sub>H</sub>RFA-I program can also identify the amino acids contributed by the potential V<sub>H</sub> replacement footprints and calculated the usages of different amino acids. The V<sub>H</sub>RFA-I program can correlate the mutation status of the identified potential V<sub>H</sub> replacement products, which will provide information regarding the selection of such V<sub>H</sub> replacement products during immune response. Another advantage of the V<sub>H</sub>RFA-I program is that it can quickly identify potential V<sub>H</sub> replacement footprints at different lengths, such as 3-, 4-, 5-, 6-, and 7-mer. Such analysis cannot be done without computer help. Clearly, with shorter length of footprint motifs, there are higher frequencies of V<sub>H</sub> replacement products. Unfortunately, there is no experimental approach to determine whether the 3-, 4-, or 5-mer of V<sub>H</sub> replacement footprints are more representative of the true occurrence of V<sub>H</sub> replacement. For all the data analyses, we arbitrarily chose 5-mer footprint motifs to calculate the frequencies of V<sub>H</sub> replacement products. Using the V<sub>H</sub>RFA-1 program, we have finished analyses of the 17,000 murine IgH gene sequences (32) and the 60,000 human IgH gene sequences available from the NCBI database (results will be published in separate studies). The results obtained in these studies revealed a significant contribution of V<sub>H</sub> replacement products to the antibody repertoires in human and mice.

Like any other sequence analysis based method, the V<sub>H</sub>RFA-1 program also has its limitations. The V<sub>H</sub>RFA-1 program can search for the existence of V<sub>H</sub> replacement footprints purely based

![Figure 12](image-url) | The GenBank file tailor. Diagram shows the interface of the GenBank File Tailor. (A) Button to choose the footprint analysis result file. (B) Button to choose the input GenBank file for tailoring. (C) Button to choose the output file. (D) Button to start the tailoring process.
on sequence analysis. It can identify V_{H} replacement footprints in the N1 regions as well as the N2 regions. Clearly, V_{H} replacement can only contribute footprints to the N1 regions. The identified “footprints” in the N2 regions can only be generated by random nucleotide addition. Statistical analysis results indicated that the frequencies of V_{H} replacement footprints with different lengths in the N1 regions are significantly higher than that in the N2 regions (32), which supports the sequence analysis based method to the identification of potential V_{H} replacement products. The V_{H}RFA-1 program relies on the IMGT/V-Quest online service to assign the potential V_{H}, D_{H}, and J_{H} gene usage, which is a critique step for subsequent identification of V_{H} replacement footprints.
in the $V_H$–$D_H$ junction. In certain IgH sequence analysis, we do notice that the IMGT $V_H$, $D_H$, or $J_H$ gene assignment might not be correct, which leads to the mistake in the identification of potential $V_H$ replacement footprints. Another issue that also affects the identification of $V_H$ replacement footprints is the potential existence of multiple $D_H$ gene segments within IgH genes. Although it is still under debate, the latest version of the IMGT/V-Quest program has already included the option to assign up to three potential $D_H$ gene segments within the $V_H$ to $J_H$ regions based on the standard stringency. Surprisingly, there are many IgH genes that contain multiple potential $D_H$ gene segments (explored in separate studies). The existence of multiple $D_H$ gene segments will change the assignment of the N1 and N2 regions and thus affect the identification of $V_H$ replacement footprints. The current version of the $V_H$RFA-1 program only works with the default setting in the IMGT/V-Quest program, which identifies one $D_H$ gene segment for each IgH genes. The multiple $D_H$ gene segments assignment results have a different output format, which is not suitable for the $V_H$RFA-1 program.

In our previous studies, we considered both the 5-mer $V_H$ replacement footprint (5-0 method) and the 6-mer $V_H$ replacement footprint with one nucleotide mismatch (6-1 method) to identify potential $V_H$ replacement products (27, 37). The current version of the $V_H$RFA-1 program only use the non-mutated
potential V<sub>H</sub> replacement footprint motif library derived from V<sub>H</sub> germline genes. In this setting, mutated V<sub>H</sub> replacement footprint motif within the V<sub>H</sub>–D<sub>H</sub> junction cannot be identified by the current program. We are still developing the next version of computer program to tolerate one nucleotide mismatch within a 6-mer of V<sub>H</sub> replacement footprint motif.

In summary, the V<sub>H</sub>RFA-I program offers a computational tool to analyze large numbers of IgH gene sequences to identify and analyze potential V<sub>H</sub> replacement products in human and mice.

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SUPPLEMENTARY MATERIAL
The Supplementary Material for this article can be found online at http://www.frontiersin.org/Journal/10.3389/fimmu.2014.00040/abstract

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