VIRsiRNAdb: a curated database of experimentally validated viral siRNA/shRNA

Nishant Thakur, Abid Qureshi and Manoj Kumar*

Bioinformatics Centre, Institute of Microbial Technology, Council of Scientific and Industrial Research (CSIR), Sector 39-A, Chandigarh-160036, India

Received August 15, 2011; Revised October 4, 2011; Accepted November 9, 2011

ABSTRACT

RNAi technology has been emerging as a potential modality to inhibit viruses during past decade. In literature a few siRNA databases have been reported that focus on targeting human and mammalian genes but experimentally validated viral siRNA databases are lacking. We have developed VIRsiRNAdb, a manually curated database having comprehensive details of 1358 siRNA/shRNA targeting viral genome regions. Further, wherever available, information regarding alternative efficacies of above 300 siRNAs derived from different assays has also been incorporated. Important fields included in the database are siRNA sequence, virus subtype, target genome region, cell type, target object, experimental assay, efficacy, off-target and siRNA matching with reference viral sequences. Database also provides the users with facilities of advance search, browsing, data submission, linking to external databases and useful siRNA analysis tools especially siTarAlign which align the siRNA with reference viral genomes or user defined sequences. VIRsiRNAdb contains extensive details of siRNA/shRNA targeting 42 important human viruses including influenza virus, hepatitis B virus, HPV and SARS Corona virus. VIRsiRNAdb would prove useful for researchers in picking up the best viral siRNA for antiviral therapeutics development and also for developing better viral siRNA design tools. The database is freely available at http://crdd.osdd.net/servers/virsiRNadb.

INTRODUCTION

Viral diseases remain one of the public health problems due to emerging and reemerging nature of viruses such as influenza, hepatitis, Human Immunodeficiency Virus (HIV), Human Papillomavirus (HPV) & Severe Acute Respiratory Syndrome (SARS) etc. (1). Combating majority of these viruses is compromised due to lack of effective vaccines and antiviral drugs (2). Besides, development of new vaccines and antiviral drugs, there are continuous efforts to search for alternative therapeutic interventions. Lately, RNA interference has emerged as a potential approach in the battle against pathogenic viruses (3,4) and other human diseases (5,6).

RNAi was first reported by Fire et al. (7) when authors showed a potent gene silencing effect after injecting double stranded RNA into *C. elegans*. In RNA silencing pathway, long dsRNA is processed by RNase III family member, dicer, to a 19–21 nucleotide long double stranded siRNA, with 2-nucleotide unphosphorylated 3' overhangs (8). The double stranded siRNA is composed of a guide (antisense) strand and a passenger (sense) strand. Unwinding of the siRNA duplex is catalyzed by argonaute. After the unwinding step, the guide strand is incorporated into the RNA Induced Silencing Complex (RISC), while the passenger strand is released. Using the antisense strand RISC targets the complementary mRNA resulting in the cleavage of the latter (9).

Using RNA silencing mechanism, researchers have reported considerable decrease in the expression of targeted viral genes (10,11). For example, siRNAs directed against the influenza virus nucleocapsid (NP) and RNA transcriptase (PA) genes inhibited its transcription and replication (12). Similarly, siRNAs against the hepatitis B virus polymerase (PA), precore (PreC) and surface (S) regions inhibited the viral replication (13). In another study, siRNAs synthesized to target the E, M and N genes of SARS-CoV effectively down regulated the target genes expression by over 80% in a dose-dependent manner (14). Inhibition of virus replication for several human viruses using RNAi strategy has been reviewed (3,15,16).

RNAi approach offers several advantages for antiviral therapeutics development. It has ability to target all types

---

*To whom correspondence should be addressed. Tel: +91 172 6665453; Fax: +91 172 2690632; Email: manojk@imtech.res.in; manojkumardelhi@yahoo.co.in

The authors wish it to be known that, in their opinion, the first two authors should be regarded as joint First Authors.

© The Author(s) 2011. Published by Oxford University Press.
This is an Open Access article distributed under the terms of the Creative Commons Attribution Non-Commercial License (http://creativecommons.org/licenses/by-nc/3.0), which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.
of viral genomes [ssDNA, dsDNA, RNA(+), RNA(−) and dsRNA] which makes this versatile mechanism to be harnessed as broad-spectrum antiviral therapy (4). Further, RNAi targets a short stretch of viral nucleic acids instead of a functional domain of a viral protein, therefore, even a small viral genome offers many potential targets (11). Even more, multiple antiviral siRNAs can be expressed simultaneously or pooled in a way similar to current drug combination anti-viral therapy of infected individuals to sustain prolonged effect (17,18).

In the past decade, a number of RNAi therapeutic programs with focus on cancer, metabolic diseases, respiratory disorders, retinal degeneration, dominantly inherited brain and skin diseases and infectious diseases have entered the clinical practice (6,19,20). Simultaneously, several RNAi based antiviral therapeutic projects have also reached at clinical trial stages (21), for example, RSV (Phase II) (22), HBV (Phase I) (23), HCV (Phase II) (24) and HIV (Phase I) (25). Ongoing clinical trials further emphasize the need for development of the viral RNAi resources.

There is no dedicated viral siRNA database, except HIVSirDB, an HIV specific siRNA database (26). However, there are a few other siRNA databases reported in literature like HuSiDa (27) and siRNAdb (28) which provide sequences of published functional siRNA targeting human genes while siRecords (29) focused on siRNA data of mammalian RNAi experiments and DSTTHO (30) on human onco genes. VIRsiRNAdb is an attempt to provide comprehensive details of the experimentally validated viral siRNA targeting the diverse genome regions of as many as 42 important human viruses at one platform to help researchers working in the field of siRNA based antiviral therapeutic development.

**DATABASE CONTENT**

**Data acquisition**

Exhaustive literature search was carried out to extract the relevant articles from PubMed. This was accomplished by searching queries having combination of two keywords: (i) terms most commonly used for gene silencing viz: RNA interference, RNAi, silencing, siRNA(s), shRNA(s), small interfering RNA(s), short interfering RNA(s), small hairpin RNA(s) and short hairpin RNA(s) and (ii) virus names including their common names, aliases & abbreviations (like Severe acute respiratory syndrome, SARS, Corona Virus, SARS-CoV). Full text search using the above two keywords combinations was performed for each of the human viruses individually. The search results are given in Supplementary Table S1. Around 4000 abstracts were screened so as to select the articles likely to contain relevant viral siRNA information. Reviews, general methodological and non-English articles were not considered. After initial screening, around 1000 remaining potential articles were examined in detail to retrieve the viral siRNA information. Articles of siRNAs targeting the host genome regions were excluded. Further, articles that did not have individual siRNA sequence or its experimental efficacies were also not included. After this extensive filtering, 221 research articles were shortlisted to collect the siRNA data. In our database, complete siRNA data of almost all human viruses reported in the literature have been included.

**Database architecture**

The database provides comprehensive information of experimentally validated viral siRNAs which includes: (i) siRNA sequence, (ii) family of virus, (iii) virus subtype, (iv) target gene, (v) siRNA location, (vi) GenBank accession, (vii) design algorithm, (viii) production method, (ix) siRNA concentration, (x) cell type, (xi) transfection method, (xii) incubation time, (xiii) PubMed ID, (xiv) object used (i.e. mRNA, protein, virus load etc), (xv) efficacy, (xvi) efficacy assay (e.g. Western blot, PCR, plaque number, ELISA) and (xvii) references. Further, wherever available, extended information regarding alternative efficacy assays has also been provided. Architecture of the database is depicted in the Figure 1. Structure of each siRNA predicted by Mfold (31) was also displayed in the data. In addition, we have also provided information of viral siRNA off-targets in human and the siRNA sequence matching with the reference viral genome sequences.

**Database statistics**

VIRsiRNAdb database provides information of 1358 experimentally validated siRNAs pertaining to 42 important human viruses belonging to 19 different virus families and targeting as many as 150 different viral genome regions. For HBV, HCV, SARS and Coxsackievirus many genome regions were being targeted by siRNAs as given in Table 1. The database entries contain siRNA experiments based on 71 different cell lines but Huh-7, 293T, MDCK, HepG2.2.15 and HeLa cell lines were mostly used (Figure 2a). In the database, 45% of the total siRNAs were highly effective with >70% inhibition efficacy and 9% siRNA have >90% efficacy. siRNAs (23%) have moderate efficacy of 50–70% whereas 32% of siRNAs were less effective with efficacy rating <50% (Figure 2b).

One of the major hindrances in RNAi based therapeutics is the lack of siRNA specificity. Besides, directly affecting the expression of the desired genes, a siRNA may affect regulation of unintended transcripts which possess complementarity to the siRNA sequence. siRNA off-target effect was initially reported in 2003 (32) and later Amanda Birmingham et al. (33) reported that off-targeting is associated with the presence of one or more perfect 3’ untranslated region (UTR) matches with the hexamer or heptamer seed region (positions 2–7 or 2–8) of the antisense strand of the siRNA. Seed based siRNA off-target was experimentally demonstrated by others also (34,35). The impact of non-specific siRNA off-target effect in therapeutic application was further reviewed (36).

We have predicted the off-targets in human for all the siRNAs present in our database, using three algorithms: (i) BLAST (37), (ii) Seed Locator (33) and (iii) SpecificityServer (38). Result outputs of each algorithm
are given against respective siRNA record as link under off-target column. BLAST algorithm was commonly used to detect possible off-target effects of a siRNA by searching it against the human Unigene or transcriptome database (28). We have also used BLAST (37) with \( -e 1000; -q -4; -r 5 \) parameters and found that around 13% of siRNA having off-targets in the human genome. Seed Locator output include total genes with at least one seed match and multiple seed matches in the 3'-UTR. Finally, results of SpecificityServer which is designed to identify potential non-specific matches to siRNA showed that 113 siRNAs are not specific for both siRNA strands while 101 siRNA have off-targets for the sense strand and remaining does not have any off-target.

As we know that viruses exhibit greater genetic variability, therefore it is important to know that in how many viral genome sequences, siRNA sequence is matching. This analysis is helpful for users in selecting such siRNA which is having high matching with maximum reference viral strains. Significance of selection of conserved regions targeted by siRNA in HIV-1 has been discussed by Naito et al. (39,40). We have checked the siRNA sequence matching with the respective viral reference genome sequences available at NCBI. For this purpose, we have used ALIGN0 algorithm (41), which computes the alignment of two DNA sequences without penalizing for end-gaps. Pie chart result displayed the number of nucleotide differences or mismatches (0, 1, 2, 3, >3) between of each siRNA and respective viral reference genome sequences in the alignment. Cumulative results of all the siRNA showed that 2% of siRNAs were fully (100%) matching with respective viral genome sequences and 16% matched with 90–99% viral genomes while 61% were having <50% matching as shown in Figure 2c.

There are reports of escape mutants generated by the virus in the siRNA target site to overcome the effect of RNAi. These escape mutations in the target sequence decreases the potency of siRNA gene silencing (42). Wilson (43) observed maximum escape mutations at 12th and 18th residues for HCV NS5B while Konishi (44) reported appreciable mutation at the 15th residue for HCV NS5A gene. In another study, Jun (45) recorded changes in Coxsackie virus at positions 10 and 13. We have collected such 57 siRNA escape sequences having 52 substitutions; 2 deletions; 1 insertion and 2 substitution/deletion mutations. Position of these escape substitutions mutations among 57 escape sequences are shown in Figure 2d.

**Tools**

Viral siRNA database allows the users to take advantage of useful tools like siTarAlign, siRNAmap and siRNAblast. siTarAlign aligns the siRNA sequence with the respective virus/family reference genomes sequences using either BLAST (37) or Smith–Waterman algorithm from EMBoss suite (46) The output shown below displays a list of flaviviruses and influenza A viruses targeted by respective siRNA (Figure 3). Viral/family reference genomes were taken from the NCBI viral genome resources as summarized in the Supplementary Table S2. In siTarAlign, user defined viral genome sequences can also be uploaded to align the siRNA sequence with user provided sequences also.

The ‘siRNAmap’ is a simple tool to display the perfectly matching siRNA available in our database to the user provided viral sequence. So, it helps the user to know that against the user provided viral sequence, how many siRNAs are available in VIRsiRNAdb. Additionally, the

---

**Figure 1.** VIRsiRNAdb database architecture.
<table>
<thead>
<tr>
<th>Virus</th>
<th>Target gene</th>
<th>No. of siRNAs</th>
<th>Virus</th>
<th>Target gene</th>
<th>No. of siRNAs</th>
</tr>
</thead>
<tbody>
<tr>
<td>BK polyomavirus</td>
<td>T-Ag(2)</td>
<td>2</td>
<td>Influenza A virus</td>
<td>M(33), NP(26), PB(19), PA(11), NS(6), C(1)</td>
<td>96</td>
</tr>
<tr>
<td>Chikungunya virus</td>
<td>E1(1), NSP3(1)</td>
<td>2</td>
<td>Influenza B virus</td>
<td>PB(33), NP(17), M(11), NS(10), PA(10)</td>
<td>81</td>
</tr>
<tr>
<td>Dengue virus [DENV]</td>
<td>E(6), NS(5), 5'-UTR(5), 5'-UTR(2), C(1), NS(1), PreM(1)</td>
<td>22</td>
<td>Japanese encephalitis virus [JE]</td>
<td>NS(14), E(2)</td>
<td>6</td>
</tr>
<tr>
<td>Ebola virus [EBOV]</td>
<td>ZNP(2), ZT(2), ZL(1)</td>
<td>5</td>
<td>John Cunningham virus [JCV]</td>
<td>T-Ag (1)</td>
<td>1</td>
</tr>
<tr>
<td>Encephalomyocarditis virus</td>
<td>EMCV-IRES(1)</td>
<td>1</td>
<td>Junin virus</td>
<td>Z(4)</td>
<td>4</td>
</tr>
<tr>
<td>Enterovirus [EV]</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>3D Pol(1), VP1(2), 3C(2), 3C pro(2), 5'NTR(2), 3'-UTR(1), MET-2C(1), VP2(1)</td>
<td>22</td>
<td>La Crosse virus</td>
<td>M(G2)(7), L(4), S(3)</td>
<td>14</td>
</tr>
<tr>
<td>Epstein–Barr virus [EBV]</td>
<td>EBNA1(25), LMP1(10), PR(4), BKRF3(2), LMP2A(1), Zta(1)</td>
<td>43</td>
<td>Lassa virus</td>
<td>GPC(1), L(1), NP(1), Z(1)</td>
<td>4</td>
</tr>
<tr>
<td>Hazara nairovirus</td>
<td>L(4), M(4), S(4)</td>
<td>12</td>
<td>Lymphocytic choriomeningitis virus</td>
<td>L(1), Z(1)</td>
<td>2</td>
</tr>
<tr>
<td>Hendra virus</td>
<td>N(2)</td>
<td>2</td>
<td>Marburg virus</td>
<td>VP30(2), NP(1)</td>
<td>3</td>
</tr>
<tr>
<td>Hepatitis A virus [HAV]</td>
<td>L(4), N(4)</td>
<td>8</td>
<td>Measles virus</td>
<td>N(16), L(8)</td>
<td>24</td>
</tr>
<tr>
<td>Hepatitis B virus [HBV]</td>
<td>S(60), X(48), C(25), P/S(24), P(17), C/P(11), ORF-C(3), ORF-S(3), P/X(3), NLS(2), PRE(2), preS/P(2), PA(1), PreS1(1), HB-Ag(7)</td>
<td>227</td>
<td>Polio virus</td>
<td>Capsid(1), 5NC(1)</td>
<td>2</td>
</tr>
<tr>
<td>Hepatitis C virus [HCV]</td>
<td>5'-UTR(38), 3'-UTR(26), NSS(8), Core(19), E(11), NS(8), E2(7), IRES(5'-UTR(6), NS5(3), NS4B(2), NAA(1), N5A(1)</td>
<td>145</td>
<td>Reovirus</td>
<td>µNS(7), σNS(4), µ2(2)</td>
<td>13</td>
</tr>
<tr>
<td>Hepatitis delta virus [HDV]</td>
<td>Delta Ag(16)</td>
<td>16</td>
<td>Rotavirus</td>
<td>NSP(2)</td>
<td>2</td>
</tr>
<tr>
<td>Hepatitis E virus [HEV]</td>
<td>ORF2(4), RdRp(4), Helicase(2), Replicase(2), 3CAE region(1)</td>
<td>13</td>
<td>SARS coronavirus</td>
<td>ORF(2a), N-protein(31), ORF5, M-protein(23), ORF4, E-protein(22), ORF2, Spike(18), Replicase(16), RDRP(11), ORF(1a)(8), ORF(1b)(7), ORF(3a)(7), (6), 3A(3), NSP1(3), ORF7(3), 3'-UTR(2), 5'-UTR(1), Leader(1), TRS(1)</td>
<td>163</td>
</tr>
<tr>
<td>Herpes simplex virus [HSV]</td>
<td>U51(4), UL39(4), UL40(4), DNA polymerase(3), gB(3), UL29(3), UL5(3), Vp1(6), gB(2), UL27(2), UL38(2), gE(1), K13(1), ORF75(1)</td>
<td>36</td>
<td>Semliki forest virus</td>
<td>Cold(7), Hot(7)</td>
<td>14</td>
</tr>
<tr>
<td>Human metapneumovirus</td>
<td>L(26), N(11), M(9), F(8), P(4)</td>
<td>58</td>
<td>St. Louis encephalitis [SLE]</td>
<td>E(1), C(2), NS(5)</td>
<td>5</td>
</tr>
<tr>
<td>Human papillomavirus [HPV]</td>
<td>E6(42), E7(39), E6/E7(8)</td>
<td>89</td>
<td>Vaccinia</td>
<td>E3L(4)</td>
<td>4</td>
</tr>
<tr>
<td>Human respiratory syncytial virus [HRSV]</td>
<td>NS1(4), P(4), NS2(1)</td>
<td>9</td>
<td>West Nile virus [WNV]</td>
<td>E(18), NS(17), Core(7), C(4), NS(14), NSH(4), P/ M(3), 3'-UTR(2), NS(2), CAP(1), NS2A(1), NS4A/B(1)</td>
<td>64</td>
</tr>
<tr>
<td>Human rhinovirus</td>
<td>3D(4), 2C(3), 5'-UTR(3), VP3(3), 3C(2), VP1(2), VP2(2), 2A(1), 3A(1), VP4(1)</td>
<td>22</td>
<td>Yellow fever virus</td>
<td>NS(2), E(1), NS(1)</td>
<td>4</td>
</tr>
</tbody>
</table>
siRNAblast allows alignment of a user provided siRNA sequence against all the siRNA sequences available in our database. This helps the user to confirm whether a given siRNA sequence or similar one has already been reported or not.

Data retrieval

It is possible to perform a quick search based on various database fields i.e. Virus name, siRNA sequence, target region, cell line and Pubmed ID. We have included a separate search option to retrieve siRNA with efficacy; greater than, equal to and lower than for a given value. Database also has qualitative efficacy of some siRNAs (where numerical values were not available) in three categories viz. ‘High’ (>70%), ‘Medium’ (50–70%) and ‘Low’ (<50%). The efficacy search will also fetch siRNAs with qualitative efficacies.

In the search output we have implemented the sorting and filtering functions. By clicking the heading of the given field, user can sort the displayed data. Simultaneously, by entering the desired keyword in the designated field, user can filter the siRNA data. Multiple filtering can be accomplished by entering desired keyword in different fields one after another. ‘Advanced Search page’ allows for more flexible queries using logical operators (AND, OR). These options enable the user to readily find appropriate siRNA data. External links pointing to the GenBank accession of the siRNA target sequence, Pubmed ID and International Committee on Taxonomy of Viruses (ICTV) are given for each siRNA record.

Data submission

Authors generating experimental viral siRNA data are encouraged to submit the data directly into viral siRNA database. For this purpose, a web form for data submission is provided. Submitted information will be included in the database update after ascertaining its authenticity.

Implementation

VIRsiRNAdb database is implemented on Red Hat Linux with MySQL (5.0.51b) and Apache (2.2.17) in back-end and front-end of web interface is implemented with PHP (5.2.14).

Future developments

As increasing number of articles are being published in the area of viral RNAi, therefore, in future our main priority would be to update the existing viral siRNA data as well as to include siRNA information for new viruses once appropriate data is available. We would also include virus specific siRNA design tool to further help the researchers.

Figure 2. Database statistics (a) Cell line used (b) siRNA efficacy (c) siRNA sequence matching with reference viral genomes (d) Positions of the escape mutations.
SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online: Supplementary Tables 1 and 2.

FUNDING

Council of Scientific and Industrial Research, and Department of Biotechnology, Government of India. Funding for open access charge: Institute of Microbial Technology (CSIR), Sector 39-A, Chandigarh, India.

Conflict of interest statement. None declared.

REFERENCES