Human Gene and Protein Database (HGPD): a novel database presenting a large quantity of experiment-based results in human proteomics

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ABSTRACT

Completion of human genome sequencing has greatly accelerated functional genomic research. Full-length cDNA clones are essential experimental tools for functional analysis of human genes. In one of the projects of the New Energy and Industrial Technology Development Organization (NEDO) in Japan, the full-length human cDNA sequencing project (FLJ project), nucleotide sequences of approximately 30,000 human cDNA clones have been analyzed. The Gateway system is a versatile framework to construct a variety of expression clones for various experiments. We have constructed 33,275 human Gateway entry clones from full-length cDNAs, representing to our knowledge the largest collection in the world. Utilizing these clones with a highly efficient cell-free protein synthesis system based on wheat germ extract, we have systematically and comprehensively produced and analyzed human proteins in vitro. Sequence information for both amino acids and nucleotides of open reading frames of cDNAs cloned into Gateway entry clones and in vitro expression data using those clones can be retrieved from the Human Gene and Protein Database (HGPD, http://www.HGPD.jp). HGPD is a unique database that stores the information of a set of human Gateway entry clones and protein expression data and helps the user to search the Gateway entry clones.

INTRODUCTION

In 2003, complete sequences of the human genome were decoded by the human genome sequencing project (1). In postgenomic research, one of the most essential subjects is the functional and structural analysis of gene products (proteins). As access to full-length cDNA clones is critical for such work, many projects, such as the FLJ project (2,3), the Kazusa cDNA project (4), the US Mammalian Gene Collection (MGC) program (5), German (6), Chinese (7) and other cDNA projects have been executed to isolate as many full-length cDNAs with as high quality as possible. For comprehensive and high-throughput expression of human proteins, both full-length cDNA clones and a versatile system for using these clones are essential. For functional analysis of proteins, one often needs to fuse various tags at either the N- or C-termini, to adjust the reading frames of the open reading frame (ORF) and tags or to locate adequate regulatory sequences [promoters, enhancers, internal ribosomal entry sites (IRESes), etc.] close to the ORF. These manipulations can be extremely difficult when a large number of clones are being handled. The Gateway cloning system (Invitrogen, CA, USA) is based on versatile expression...
vectors and has the potential to overcome these barriers (8). We have therefore adopted Gateway technology and constructed 33,275 human Gateway entry clones that will serve as key resources for this versatile system. Sequence information of Gateway entry clones can be retrieved from the Human Gene and Protein Database (HGPD, http://www.HGPD.jp or http://HGPD.lifesciencedb.jp/) (Figure 1). ORFDB (http://orf.invitrogen.com/) (9) and the ORFeome collaboration (http://www.orfeomecollaboration.org/html) have been published as similar resources. Entry clones in ORFDB are only N-types which have a stop codon at the end of the ORF and are primarily dedicated for native or N-terminal fusion proteins, although one could produce native or C-terminal fused protein with suppression technology (10). Lamesch et al. (11) have reported on the construction of 12,212 entry clones with which the ORFeome collaboration was formed. Since a large fraction of ORFeome clones are F-types that delete the stop codon for C-terminal fusion proteins, proteins that possess a functional domain at the C-terminus might not have full biological activity when expressed based on these clones. Therefore, one might need both N- and F-type entry clones. In our collection, both types have been prepared for 11,774 cDNAs, which means that our collection may have more flexibility for various in vitro and in vivo experiments.

Utilizing these clones with a highly efficient cell-free protein synthesis system featuring wheat germ (12), we have produced and analyzed 13,364 human proteins in vitro. The expression data can be retrieved from HGPD (Figure 1). HGPD manages and stores primary protein expression data, which differs from other databases, such as the Human Protein Reference Database (HPRD, http://www.hprd.org/) (13), Gene Ontology database (GO, http://www.geneontology.org/) (14), Universal Protein Resource (UniProt, http://www.pir.uniprot.org/) (15) or NCBI Entrez Gene (http://www.ncbi.nlm.nih.gov/) (16).

DATABASE CONTENTS

In HGPD, biological data such as in vitro expression data of human proteins are presented on the frame of cDNA clusters. To build the basic frame, sequences of FLJ cDNAs and others deposited in public databases (Human ESTs, RefSeq, Ensembl, MGC, etc.) are assembled onto the genome sequences (Table 1). Information for human Gateway entry clones is presented with the source cDNAs. The specific features of the HGPD that we would like to emphasize are that it contains (i) the world largest collection of Gateway entry clones, (ii) arrangement of both N- and F-type entry clones, and especially, (iii) SDS–PAGE patterns of proteins expressed in the cell-free wheat germ extract system (‘PE’ shown in Figure 1).

Gateway entry clones

To facilitate utilization of full-length cDNA clones, we have adopted the versatile Gateway expression system which offers high-throughput gene transfer technology for functional gene analysis and protein expression. For conversion to entry clones, we selected an ORF region in each cDNA meeting one of the following criteria: (i) ORFs-encoding products ≥150 aa [although the longest ORF starting with an AUG codon has highest priority, the selected ORF is finally determined by taking into consideration homology search results of shorter ORFs with BLASTX(nr) and BLASTP against SwissProt and RefSeq databases]; (ii) both ‘149 aa ≥ORF ≥100 aa’ and ‘ORF with an ATGpr value (17) ≥0.4’; (iii) both ‘100 aa >ORF’ and ‘known gene’. Those ORF regions were PCR amplified with attB sequences of the Gateway system at both ends. Then those fragments were recombined with attP sequences of the Gateway donor vector pDONR201 (Invitrogen). Eventually, we constructed 33,275 Gateway entry clones utilizing FLJ clones as major cDNA resources. Sequence information, such as amino acid and nucleotide sequences of ORF regions and sequence differences of Gateway entry clones from source cDNAs are presented in the ‘GW: Gateway Summary’ page (for help, see http://hgpd.hinv.jp/sys_info/help.html#w120_gw). The details for construction and usage of entry clones will be published elsewhere (18).

Gateway entry clones are available from NITE Biological Resource Center (NBRC), Department of Biotechnology, National Institute of Technology and Evaluation. Distribution of clones by NBRC requires the signing of an MTA by both private companies and academic institutions. Distribution charges will be 30,000 and 15,000 Yen (JPN; approx. US$300 and $150, respectively) per clone for private companies and academic institutions, respectively. More information is available through the ‘clone inquiry’ page (http://hgpd.hinv.jp/sys_info/order_clone.html) of HGPD or the notice page (http://www.nbrc.nite.go.jp/e/hgentry-e.html) of NBRC.

SDS–PAGE patterns of human proteins synthesized in vitro

The Gateway system is a versatile expression vector system that is adequate for handling large numbers of clones. For expression of large numbers of human proteins, we adopted the wheat germ cell-free protein synthesis system. In addition, we devised a new procedure to prepare template DNA for transcription, which makes the step simpler and more efficient. By applying those protocols, we expressed 13,364 human proteins with a C-terminal V5 or His tag and analyzed them using SDS–PAGE. Expression patterns of proteins in both the total and supernatant fractions are displayed in the ‘PE: Protein Expression’ page (for details, see http://hgpd.hinv.jp/sys_info/help.html#w120_pe). Essentially all of the human proteins analyzed in our work were shown to be expressed. This implies that in vitro cell-free systems using wheat germ extract offer a very efficient system for protein production.

Computational analysis of individual cDNA sequences with BLAST, Pfam, PROSITE, PSORT, SignalP, SOSUI and GO

Functional motifs and domains, subcellular localization information, leader sequences and transmembrane
Figure 1. Search flow in HGPD. Each representative page of HGPD is shown: Top, top page. After entering a proper ID, such as ‘DDBJ/EMBL/GenBank Accession No.’, ‘Ensembl Transcript No.’, ‘Gene Symbol’, ‘FLJ ID’ and ‘Sequence ID’, the ‘Information Overview’ window will emerge. It presents a summary of all information on the cluster to which the queried cDNA clone belongs. Search results for ‘AK092682’ (DDBJ/EMBL/GenBank Accession No., AK092682; FLJ ID, FLJ35363; Sequence ID, C-SKMUS2000679) are presented as an example. A ‘PE’ button opens a ‘Protein Expression’ window through a GW page, which is indicated by dotted arrows. An L2 window which is linked with an L2 window and can be used for ‘search by chromosome coordinates’ is not shown (for details, see http://hgpd.hinv.jp/sys_info/help.html#l2b).
Table 1. Entries of HGPD

<table>
<thead>
<tr>
<th>Dataset</th>
<th>Number of data</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gateway Entry Clones _ N-type(\text{a})</td>
<td>12 754</td>
</tr>
<tr>
<td>Gateway Entry Clones _ F-type(\text{b})</td>
<td>20 521</td>
</tr>
<tr>
<td>In vitro Protein Expression (SDS–PAGE) Patterns(\text{c})</td>
<td>13 364</td>
</tr>
<tr>
<td>FLJ cDNAs(\text{d})</td>
<td>35 083</td>
</tr>
<tr>
<td>Public Database cDNAs (including FLJ cDNAs)</td>
<td>112 992</td>
</tr>
<tr>
<td>FLJ ESTs(\text{e})</td>
<td>1 430 438</td>
</tr>
<tr>
<td>Public ESTs</td>
<td>3 862 807</td>
</tr>
</tbody>
</table>

\(\text{a}\)Entry clones with a naturally occurring stop codon.
\(\text{b}\)Entry clones without a stop codon for adding a tag at the C-terminal end.
\(\text{c}\)As the number of FLJ cDNAs.
\(\text{d}\)Number of published clones: 30 063, unpublished clones: 5020.
\(\text{e}\)Deposited by the FLJ project.

domains were inferred using BLAST, Pfam (http://www.sanger.ac.uk/Software/Pfam/), PROSITE (http://www.expasy.ch/prosite/), PSORT (http://psort.hgc.jp/), SignalP (http://www.cbs.dtu.dk/services/SignalP/), SOSUI (http://bp.nuap.nagoya-u.ac.jp/sosui/sosuiimenu0.html) and GO.

Mapping and clustering of cDNA clones

Local alignments between human cDNAs and human genome sequences (UCSC hg17 NCBI Build 35) were calculated using megablast (http://www.ncbi.nih.gov/blast). Initially, the alignment with the highest score was selected and a single locus was assigned for each cDNA. Those cDNAs with sequences overlapping not less than 1 base at the same locus and strand were defined as constituting the same cluster. All entries cataloged in HGPD are presented in Table 1.

WEB INTERFACE

The search flow of HGPD is illustrated in Figure 1. The top page (http://hgpd.hinv.jp/sys_info/help.html#id_search) of the HGPD viewer is represented in the upmost part of Figure 1. To begin the search, the ID number (in a definitive or degenerated form) such as DDBJ/EMBL/GenBank accession number, Ensembl transcript number, Gene Symbol, FLJ ID or Sequence ID is entered into the text box. When a query hits the data in HGPD, an ‘Information Overview’ page comes out. It shows all data concerning all members clustered with a queried sequence. In addition, all information stored in HGPD for searched clusters and cDNA clones is documented on the page. The ‘Locus’ column represents the cluster ID obtained by genome mapping of all the cDNA sequences, including expressed sequence tags. Buttons ‘L1’ and ‘L2’ are linked with ‘L1: Locus View 1’ (http://hgpd.hinv.jp/sys_info/help.html#w015) and ‘L2: Locus View 2’ (http://hgpd.hinv.jp/sys_info/help.html#w022), respectively. The ‘Gene Symbol’ column represents the official symbol appearing in the Entrez Gene database for each cDNA clone. cDNA clones that have not been assigned a ‘Gene Symbol’ are designated as ‘-’. The ‘Accession No.’ column represents the registered ID in the public database for each cDNA clone. Buttons ‘C1’ and ‘C2’ in the ‘cDNA Info’ column are linked to ‘C1: cDNA Summary 1’ and ‘C2: cDNA Summary 2’ (for details, see http://hgpd.hinv.jp/sys_info/help.html#w013 and http://hgpd.hinv.jp/sys_info/help.html#w014 for C1 and C2, respectively). Information on cDNA clones, including sequences and homology search results, is presented on the ‘cDNA Summary 1’ and ‘cDNA Summary 2’ pages. The ‘FLJ ID’ column indicates the FLJ ID number of the FLJ cDNA clone. Any cDNA clone that has not been assigned an FLJ number is designated as ‘-’. FLJ clones were eventually found to have three kinds of IDs: ‘DDBJ/EMBL/GenBank Accession No.’, ‘FLJ ID’ and ‘Primary Clone ID’. The ‘Sequence ID’ column shows the ID of a sequence of a cDNA clone. For sequences of cDNAs other than FLJ cDNAs, an accession number for DDBJ/EMBL/GenBank is depicted. The column ‘Protein Info’ is linked to information on expressed proteins using Gateway entry clones. A ‘GW’ button is linked with sequence information on entry clones and a ‘PE’ button is linked with protein expression through a ‘GW: Gateway Summary’ page.

In the search flow of HGPD, some links open new windows and other links load in the current window (http://hgpd.hinv.jp/sys_info/help.html#search_flow). Windows that show various data (C1, C2, GW and PE) focusing on a single cDNA clone open in the current window, as translocation can be essentially reversible (one versus one). Other windows which display multiple clones or clusters (‘Information Overview’, L1, L2 and L2) will in principle open a new window when transferred, as translocation is usually irreversible (one versus multiple).

Data for amino acid and nucleotide sequences of ORFs cloned into Gateway entry clones, summary of protein expression and others can be downloaded from the top page of HGPD (http://hgpd.hinv.jp/sys_info/download.html).

FUTURE DEVELOPMENTS

Several modifications in browser interface will be done. (i) The database will be updated by next spring to correspond to UCSC hg18/NCBI build 36. (ii) Various search interfaces will be introduced in a future version.

Information on about 18 000 more human entry clones will be included shortly, which will put the cumulative number of our collection at 50 000. Fourteen thousand entries on protein expression data in Escherichia coli will also be presented in HGPD. Additionally, data for subcellular localization for 14 000 expressed human proteins, which have been examined in HeLa cells, are being processed for publication.

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REFERENCES


Conflict of interest statement. None declared.