RESEARCH ARTICLE

BGD: A Database of Bat Genomes

Jianfei Fang, Xuan Wang, Shuo Mu, Shuyi Zhang, Dong Dong*

Institute of Molecular Ecology and Evolution, SKLEC & IECR, East China Normal University, Shanghai, China

* ddong.ecnu@gmail.com

Abstract

Bats account for ~20% of mammalian species, and are the only mammals with true powered flight. For the sake of their specialized phenotypic traits, many researches have been devoted to examine the evolution of bats. Until now, some whole genome sequences of bats have been assembled and annotated, however, a uniform resource for the annotated bat genomes is still unavailable. To make the extensive data associated with the bat genomes accessible to the general biological communities, we established a Bat Genome Database (BGD). BGD is an open-access, web-available portal that integrates available data of bat genomes and genes. It hosts data from six bat species, including two megabats and four microbats. Users can query the gene annotations using efficient searching engine, and it offers browsable tracks of bat genomes. Furthermore, an easy-to-use phylogenetic analysis tool was also provided to facilitate online phylogeny study of genes. To the best of our knowledge, BGD is the first database of bat genomes. It will extend our understanding of the bat evolution and be advantageous to the bat sequences analysis. BGD is freely available at: http://donglab.ecnu.edu.cn/databases/BatGenome/.

Introduction

Bats are mammals of the order Chiroptera, representing about 20% of all classified mammalian species worldwide [1]. Bats have long been regarded as one of the most unusual and specialized animals. They have long been regarded as special animals for the sake of being mysterious flyers of the night, and they are actually the only mammalian group with true flight capability. Furthermore, most of the bats are masters of echolocation, which allows bats to detect, localize, and even classify their prey in the complete darkness.

For the sake of these specialized phenotypic traits, many researches have been devoted to explore the underlying molecular mechanisms of bats at the sequence level [2]. For example, the 'hearing gene' Prestin was recently reported to have undergone sequence convergence between echolocating bats and dolphins [2]. Energy metabolism genes were reported to be targets of natural selection and allowed adaptation to the energy demand during the origin of flight [3]. Recently, several bat genomes have been sequenced and assembled, and these data provided us valuable resources for further scientific researches on the biology and conservation of bats [4–6]. The prevailing theory is that flying vertebrates (bats and birds) tends to have smaller genomes than other vertebrates due to metabolic constraints on cell sizes and genome
Consistent with this finding, the bat genomes (~2 Gb) are relatively smaller than other mammals. Up to date, there's no specialized and comprehensive database that focuses on storage of bat genomes. To conveniently access the bat genomes, a uniform database for the bat genomes is necessary. In this work, we collected the genome sequences of six bats (including two megabats, Pteropus alecto, Pteropus vampyrus and four microbats, Myotis davidii, Myotis brandtii, Myotis lucifugus, Eptesicus fuscus, Fig 1) from various databases, and uniformly in silico annotated these genome sequences. BGD was developed as a public database for readily accessing the bat genomes and genes, and a platform for extensive biological interpretations.

**Materials and Methods**

The genome and gene sequences of Pteropus vampyrus (Ptevap1.0) and Myotis lucifugus (Myoluc2.0) were downloaded from Ensembl database (http://www.ensembl.org/index.html) [8], and other four bat genomes (Pteropus alecto, ASM32557v1; Myotis brandtii, ASM41265v1; Myotis davidii, ASM32734v1; Eptesicus fuscus, EptFus1.0) were downloaded from NCBI genome database (http://www.ncbi.nlm.nih.gov/genome/). All the genomes have not been assembled into chromosomes, and sequence scaffolds were obtained. Accurately prediction of protein-coding genes is the most important task of genome annotation. The bat genomes were sequenced separately, and bat genes were annotated using different pipelines. For example, the genes of Pteropus vampyrus and Myotis lucifugus were predicted based on homology searching method in Ensembl database. Moreover, the genome of Eptesicus fuscus is still not well annotated. So, a uniform pipeline is necessary for gene annotation. In this work, we uniformly annotated the bat genomes using a combination of homology-based and de novo method according to previously published pipeline [4]. Because human, mouse and dog proteins are well annotated in mammals. For the homology-based method, human, mouse and dog proteins were collected and mapped on the genomes using tblastn. Then, homologous genome sequences were aligned against the matching proteins using Genewise (version 2.2.0) [9]. For the de novo prediction method, Augustus [10] and Genescan v1.0 [11] were employed. The RNA-seq data of Myotis davidii, Myotis brandtii and Pteropus alecto were also downloaded to help annotate the genomes. Finally, all lines of evidences were combined together using EVM.
(r2012-06-25) software (evidence_modeler.pl –genome bat_genome.fa –gene_predictions –weights/weight.txt –protein_alignments/bat_genblastg.gff –transcript_alignments –exec_dir 50). At last, 21237, 16956, 21593, 22125, 19496, 18366 protein coding genes were obtained from the genomes of *Pteropus alecto*, *Pteropus vampyrus*, *Myotis davidii*, *Myotis brandtii*, *Myotis lucifugus*, *Eptesicus fuscus*, respectively. We compared our predicted genes with previous annotated genes, and the result showed that these results are very similar (S1 Fig). Then, a serious of annotation works were performed in order to obtain comprehensive genomic functional information. First, the prediction of gene function domains was performed using InterproScan v5 [12] software against InterPro database [13], which integrates together predictive information about protein function from a number of resources and provides an overview of protein functions. Second, full-length cDNA sequences of bats were mapped to genomes using BLAT [14]. Then, we performed BLASTP (E-value 1e-5) against NCBI RefSeq and UniRef databases to find the best hit for each gene. The statistics of six bat genomes and annotated information are shown in Table 1.

### Results and Discussion

We stored and managed data for BGD using MySQL on a Linux system. BGD uses several common gateway interface scripts to process user’s input to search the database. A schematic diagram of BGD organization is shown in Fig 2.

### Retrieve data

The searching engine can be used to acquire the annotated gene information. In the current version, BGD has been designed with simple search and batch search engines and can be accessed with gene symbols or BGD ID. BGD can return a list of bat genes, coupled with biological implications, Gene ontology information and nucleotide or amino acid sequences.

### Genome browser

BGD utilizes a genome browser, implemented with GBrowse v2.0 [15], to navigate gene annotation along the bat genome assemblies. GBrowse is a well-known browser that combines database and interactive web pages to display the annotation of the genome. The genome browser connecting to a MySQL backend is used, and researchers can view the genomic features aligned to the genome.

### Synteny browser

A six-way genomes comparison among the bat species was performed, and we used OrthoCluster v1 [16] for the detection of synteny blocks among bat genomes. The result can be visualized using GBrowse (version 1.69). It can be used to compare co-linear regions of

| Species            | Number of scaffolds | Scaffold N50 (bp) | Number of contigs (bp) | Contig N50 (bp) | Number of genes |
|--------------------|---------------------|-------------------|------------------------|-----------------|-----------------|
| *Pteropus vampyrus*| 96,944              | 124,060           | 388,808                | 8,527           | 16,956          |
| *Pteropus alecto*  | 65,598              | 15,954,802        | 170,164                | 31,841          | 21,237          |
| *Myotis brandtii*  | 169,750             | 3,225,832         | 325,414                | 23,289          | 22,125          |
| *Myotis lucifugus* | 11,654              | 4,293,315         | 72,785                 | 64,330          | 19,496          |
| *Myotis davidii*   | 101,769             | 3,454,484         | 325,280                | 15,182          | 21,593          |
| *Eptesicus fuscus* | 6,789               | 13,454,942        | 167,058                | 21,392          | 18,366          |

| Species            | Number of contigs (bp) | Contig N50 (bp) | Number of genes |
|--------------------|------------------------|-----------------|-----------------|
| *Pteropus vampyrus*| 16,956                 | 8,527           | 16,956          |
| *Pteropus alecto*  | 21,237                 | 31,841          | 21,237          |
| *Myotis brandtii*  | 22,125                 | 23,289          | 22,125          |
| *Myotis lucifugus* | 19,496                 | 64,330          | 19,496          |
| *Myotis davidii*   | 21,593                 | 15,182          | 21,593          |
| *Eptesicus fuscus* | 18,366                 | 21,392          | 18,366          |

*doi:10.1371/journal.pone.0131296.t001*
multiple genomes using the familiar GBrowse-style web page. The 'hearing gene' Prestin was recently widely reported in bats [2, 17], which play important role in bat echolocation. Here, we showed an example of comparative synteny of Prestin gene (S2 Fig) in BGD synteny browser.

**Phylogeny server**

To better understand the evolution of bat genes, BGD provides an online phylogeny tool. Considering the accuracy and efficacy, neighbor-Joining method was implemented, and only 50 or 100 bootstrap replicates can be selected. In the current version, BGD have employed phyloXML (version 1.10) software [18] for the online phylogenetic tree visualization. Mozilla Firefox or Safari web browser are highly recommended, and Sun Java 1.5 or higher version is needed. An example of Prestin genes were provided in BGD, and the online result was shown in S3 Fig.

**BLAST server**

BLAST is one of the most useful entrance site for genome database. At BGD, researchers can search against a variety of genomic sequences. We packed all bat gene sequences to facilitate search for homologs of other mammalian species.

**Future directions**

Other bat genome sequences and population genomic studies for bats should be forthcoming. It will be very useful for analyzing bat genome and gene sequences to explore the bat evolution.
Future directions include an incorporation of more bat genome data to provide a richer source of comparative implementation of bat sequence analysis.

Conclusion
We presented an easily accessible database, offering access to the genome of bat species. The integration of annotated genome can enhance the role of BGD as an essential resource for bat evolution analysis. The BGD enables use of genomic data toward facilitating further understanding of the fundamental biology of bat species, and the adaptation of specialized traits. To the best of our knowledge, BGD is the first repository centralizing the genomes and genes of bat species. The database not only provides a large resource for the bat researches, but also supplies a platform for comparative genomic analysis.

Supporting Information
S1 Fig. Comparison of identified genes between our findings and original results.
(JPG)
S2 Fig. Comparative synteny of Prestin gene in bat genomes.
(JPG)
S3 Fig. Phylogenetic tree of Prestin gene generated in BGD database.
(JPG)

Author Contributions
Conceived and designed the experiments: DD. Performed the experiments: DD. Analyzed the data: JF XW SM SZ DD. Contributed reagents/materials/analysis tools: JF DD. Wrote the paper: DD.

References
1. Solari S, Baker RJ. Mammal species of the world: a taxonomic and geographic reference. Journal of Mammalogy. 2007; 88(3):824–30. doi:10.1644/06-mamm-r-422.1
2. Liu Y, Cotton JA, Shen B, Han X, Rossiter SJ, Zhang S. Convergent sequence evolution between echolocating bats and dolphins. Current biology: CB. 2010; 20(2):R53–4. Epub 2010/02/05. doi:10.1016/j.cub.2009.11.058 PMID: 20129036.
3. Shen YY, Liang L, Zhu ZH, Zhou WP, Irwin DM, Zhang YP. Adaptive evolution of energy metabolism genes and the origin of flight in bats. Proceedings of the National Academy of Sciences of the United States of America. 2010; 107(19):8666–71. Epub 2010/04/28. doi:10.1073/pnas.0912613107 PMID: 20421465; PubMed Central PMCID: PMC2889356.
4. Zhang G, Cowled C, Shi Z, Huang Z, Bishop-Lilly KA, Fang X, et al. Comparative analysis of bat genomes provides insight into the evolution of flight and immunity. Science. 2013; 339(6118):456–60. Epub 2012/12/22. doi:10.1126/science.1230835 PMID: 23258410.
5. Lindblad-Toh K, Garber M, Zuk O, Lin MF, Parker BJ, Washietl S, et al. A high-resolution map of human evolutionary constraint using 29 mammals. Nature. 2011; 478(7370):476–82. Epub 2011/10/14. doi:10.1038/nature10530 PMID: 21993624; PubMed Central PMCID: PMC3207357.
6. Seim I, Fang X, Xiong Z, Lobanov AV, Huang Z, Ma S, et al. Genome analysis reveals insights into physiology and longevity of the Brandt’s bat Myotis brandtii. Nature communications. 2013; 4:2212. Epub 2013/08/22. doi:10.1038/ncomms3212 PMID: 23962925; PubMed Central PMCID: PMC3753542.
7. Smith JDL, Gregory TR. The genome sizes of megabats (Chiroptera: Pteropodidae) are remarkably constrained. Biol Letters. 2009; 5(3):347–51. doi:10.1098/rsbl.2009.0016. ISI:000266144300017.
8. Flicek P, Amode MR, Barrett D, Beal K, Billis K, Brent S, et al. Ensembl 2014. Nucleic acids research. 2014; 42(Database issue):D749–55. Epub 2013/12/10. doi:10.1093/nar/gkt1196 PMID: 24316576; PubMed Central PMCID: PMC3964975.
9. Birney E, Clamp M, Durbin R. GeneWise and Genomewise. Genome research. 2004; 14(5):988–95. Epub 2004/05/05. doi: 10.1101/gr.1865504 PMID: 15123596; PubMed Central PMCID: PMC479130.

10. Stanke M, Morgenstern B. AUGUSTUS: a web server for gene prediction in eukaryotes that allows user-defined constraints. Nucleic acids research. 2005; 33(Web Server issue):W465–7. Epub 2005/06/28. doi: 10.1093/nar/gki458 PMID: 15980513; PubMed Central PMCID: PMC1160219.

11. Burge C, Karlin S. Prediction of complete gene structures in human genomic DNA. Journal of molecular biology. 1997; 268(1):78–94. Epub 1997/04/25. doi: 10.1006/jmbi.1997.0951 PMID: 9149143.

12. Zdobnov EM, Apweiler R. InterProScan—an integration platform for the signature-recognition methods in InterPro. Bioinformatics. 2001; 17(9):847–8. Epub 2001/10/09. PMID: 11590104.

13. Hunter S, Apweiler R, Attwood TK, Bairoch A, Bateman A, Binns D, et al. InterPro: the integrative protein signature database. Nucleic acids research. 2009; 37(Database issue):D211–5. Epub 2008/10/23. doi: 10.1093/nar/gkn785 PMID: 18940856; PubMed Central PMCID: PMC2686546.

14. Kent WJ. BLAT—the BLAST-like alignment tool. Genome research. 2002; 12(4):656–64. Epub 2002/04/05. doi: 10.1101/gr.229202 Article published online before March 2002. PMID: 11932250; PubMed Central PMCID: PMC187518.

15. Stein LD. Using GBrowse 2.0 to visualize and share next-generation sequence data. Briefings in bioinformatics. 2013; 14(2):162–71. Epub 2013/02/05. doi: 10.1093/bib/bbt001 PMID: 23376193; PubMed Central PMCID: PMC3603216.

16. Vergara IA, Chen N. Using OrthoCluster for the detection of synteny blocks among multiple genomes. Current protocols in bioinformatics / editorial board, Baxevanis Andreas D [et al]. 2009; Chapter 6:Unit 6 10 6 1–8. doi: 10.1002/0471250953.bi0610s27 PMID: 19728289.

17. Li G, Wang J, Rossiter SJ, Jones G, Cotton JA, Zhang S. The hearing gene Prestin reunites echolocating bats. Proceedings of the National Academy of Sciences of the United States of America. 2008; 105(37):13959–64. Epub 2008/09/09. doi: 10.1073/pnas.0802097105 PMID: 18776049; PubMed Central PMCID: PMC2544561.

18. Han MV, Zmasek CM. phyloXML: XML for evolutionary biology and comparative genomics. BMC bioinformatics. 2009; 10:356. Epub 2009/10/29. doi: 10.1186/1471-2105-10-356 PMID: 19860910; PubMed Central PMCID: PMC2774328.