DDBJ Updates in 2024

Yaz Nakamura

中村保一

The 9th Big Data Forum for Life and Health Sciences & The 21st Asian Bioinformatics Consortium Symposium Oct.. 16-18, 2024 Beijing

Center Village Keep Number one ドイト I J Nakamura Yasukazu



mountain river sun/day





National Institute of Genetics

National Institute of Genetics: www.nig.ac.jp

Research Departments Department of Informatics Department of Genomics and Evolutionary Biology ARITA, Masanori AKASHI, Hiroshi **Biological Networks** ► Lab HP Evolutionary Genetics ► Lob HP ▶ KAWAMOTO, Shoko IKEO, Kazuho Genetic Informatics ► Lob HP DNA Data Analysis KUROKAWA, Ken ▶ INOUE, Ituro Genome Evolution ► Lab HP Human Genetics MORI, Hiroshi KITANO, Jun Genome Diversity ► Lob HP Ecological Genetics ▶ NAKAMURA, Yasukazu KURAKU, Shigehiro Genome Informatics Molecular Life History ▶ Lob HP OKUBO, Kousaku Naruva Gene-Expression ▶ Lob HP Analysis Population Genetics Vutaka Plant Genetics omparative Genomics Department of Gene Function Department of Chromosome and Phenomics HIRATA, Tatsumi ▶ KANEMAKI, Masato ► Lob HP Brain Function Molecular Cell ▶ Lob HP Engineering IWASATO, Takuji KIMURA, Akatsuki Mammalian Neural ▶ Lob HP ▶ Lab HP Cell Architecture Circuits KAWAKAMI Koichi MAESHIMA, Kazuhiro ► Lab HP Molecular and Genome Dynamics ► Lab HP Developmental Biology SAITO, Kuniaki KOIDE, Tsuyoshi Invertebrate Genetics ► Lob HP Mouse Genomics ► Lob HP SHIMAMOTO, Yuta Resource Physics and Cell Biology ▶ Lob HP MIYAGISHIMA, Shin-ya Laboratory Symbiosis and Cell ► Lob HP Evolution NIKI, Hironori Microbial Physiology ► Lab HP NONOMURA, Ken-ichi ► Lob HP Plant Cytogenetics ODA. Yoshihisa Cell Dynamics and ► Lob HP Signaling SAGA, Yumiko Mammalian ► Lab HP Development SAKAI, Noriyoshi Model Fish Genetics ► Lab HP SAWA Hitoshi Multicellular ► Lab HP Organization YONEHARA, Keisuke Multiscale Sensory ► Lob HP Structure Center for Frontier Research

Center for Frontier Research KUBO, Fumi Systems Neuroscience Lob IP MURAYAMA, Yasuto Chromosome Biochemistry

Intellectual Infrastructure Center









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Japanese Studies International Research Center for Japanese Studies

Japanese History National Museum of Japanese History

Statistical Science

The Institute of Statistical

Mathematics

Q

National Institute of Japanese Literature

Reset



Particle and Nuclear Physics

Institute of Particle and Nuclear

Studies



Molecular Science

Institute for Molecular Science

Basic Biology

National Institute for Basic Biology





Physiological Sciences

National Institute for Physiological

Sciences

Institute of Materials Structure

Science





Genetics National Institute of Genetics







A





Japanese Language

Sciences

National Institute for Japanese

Accelerator Science Accelerator Laboratory / Applied Research Laboratory



Astronomical Science National Astronomical Observatory

Informatics

National Institute of Informatics

Fusion Science

National Institute for Fusion Science

Space and Astronautical Science

蜜

Institute of Space and Astronautical Science



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Materials Structure Science **Global Environmental** Studies Research Institute for Humanity and Nature

Polar Science National Institute of Polar Research

DDBJ and the INDSC

Sequencing cost





Sequence Read Archive growth (2024.2.25)



Open access hases

DDBJ

DNA DataBank of Japan



INSDC

The international nucleotide sequence database collaboration



Global Participation Đ

About INSDC 🔿

Technical Specifications Đ



History

1995	EMBL data library was organized, and asked international cooperation for nucleotide sequence data bank to Japan.	
1982	EMBL and GenBank started international cooperation, and invited Japan to participate their data bank.	
1983	Aimed at contribution for international data bank to collect, to evaluate and to provide nucleotide sequence data, trial data loading was started.	
1984	NIG; the National Institute of Genetics was reorganized as an Inter-University Research Institute. DDBJ began to work at NIG.	
1986	DNA Database Advisory Committee organized.	
1987	DDBJ release 1 was provided. By this release, we regard this year as official start of DDBJ operation.	
1995.04	To operate DDBJ more efficiently, CIB; the Center for Information Biology was established in NIG.	
2001.04	CIB was reorganized as CIB-DDBJ; the Center for Information Biology and DNA Data Bank of Japan	DDBJ
2004.04	NIG was reorganized as a member of ROIS; Research Organization of Information and Systems. DDBJ has also belonged to ROIS.	INSDC
2005.05	DDBJ, EMBL-Bank and GenBank agreed to call their collaboration INSDC; International Nucleotide Sequence Database Collaboration; and to call the unified nucleotide sequence database INSD; the International Nucleotide Sequence Database.	NCBI ENA
2007.04	DBCLS; Database Center for Life Science was newly founded in ROIS	
2009	DDBJ faculty staff have greatly been reshuffled. DDBJ collaborates with DBCLS more closely. INSDC added a collaborative meeting to deal with huge sequence data produced by the next generation sequencers (Sequence Read Archive) and traces produced by traditional sequencers (Trace Archive).	
2012.04	DDBJ, expanding its DNA databank activities, was restructured as one of the Intellectual Infrastructure Project Centers of NIG, being separated from CIB.	
2013.10	Collaborating with NBDC; National Bioscience Database Center, DDBJ Center started to operate the archive for all types of individual-level genetic and de- identified phenotypic data from human subjects, JGA; Japanese Genotype- phenotype Archive.	

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Jun Mashima, Kazunori Aoki, Hideo Aono, Yuji Ashizawa, Yukino Dobashi, Mayumi Ejima, Masahiro Fujimoto, Asami Fukuda, Tomohiro Hirai, Michiaki Hiramatsu, Naofumi Ishikawa, Kenji Kato, Aimi Kawasaki, Yuichi Kodama, Junko Kohira, Takehide Kosuge, Kyungbum Lee, Mika Maki, Fujitaka Matsumori, Kimiko Mimura, Hideki Mochizuki, Naoko Murakata, Yoshiyuki Nogi, Toshihisa Okido, Yoshihiro Okuda, Maki Ono, Katsunaga Sakai, Yukie Sakon, Makoto Sato, Rie Sugita, Kimiko Suzuki, Takahiro Suzuki, Daisuke Takagi, Yaeko Takiguchi, Toshiaki Tokimatsu, Haru Tsutsui, Koji Watanabe, Tomohiko Yasuda, Emi Yokoyama, Masanori Arita, Takeshi Kawashima, Osamu Ogasawara, Kosaku Okubo, Nozomu Sakurai, Yasuhiro Tanizawa, Toshihisa Takagi, and Yasukazu Nakamura

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The INSDC meeting at EBI, May 2019



The INSDC meeting online, May 2021



International Nucleotide Sequence Database Collaboration

Global Participation



To achieve its mission and vision, INSDC is establishing and implementing a plan that incorporates new Members. Diversifying participation through new membership will advance open science and data sharing and, in turn, drive innovation.

	expand its membership to include regional collaborators and/or organisations representing all continents covering all organisms and all environments
	ensure its operation reflects emerging trends in scientific discovery and societal needs for the benefits that result from sequencing information
INSDC aims to:	build equitable systems that enable the global benefit from sequence information
	ensure that diverse perspectives of genetic sequence generators and managing repositories are reflected in its operations
Global participation	To formalise the INSDC collaboration and to foster new membership, the INSDC developed a Founders Arrangement and a Membership Arrangement.
Membership requirements	INSDC Members are required to demonstrate appropriate capacity, commitment, and activity in areas such as governance, technical infrastructure, data operations, and communications and engagement as described in the Membership Acceptance and Performance Guidelines.

MEMBERSHIP ARRANGEMENT

FOR PARTICIPATION IN AND CONTRIBUTION TO THE INTERNATIONAL NUCLEOTIDE SEQUENCE DATABASE COLLABORATION (INSDC)

This Membership Arrangement specifies terms for institutional participation in and contribution to the International Nucleotide Sequence Database Collaboration (INSDC).

Whereas the INSDC is a global collaboration of independent government or non-profit organizations that manage nucleotide sequence databases that capture, preserve, and present comprehensive nucleotide sequence information and annotations to preserve the scientific record and enable broad sharing of such data; and

Whereas Members see value in broad international collaboration to improve global coverage of genomic sequences from all regions of the world; and

Whereas, Members commit to the collection and distribution of nucleotide sequence data and related analyses; and

Whereas, Members adhere to the principles of free and unrestricted access and sharing of genomic sequence data; and

Whereas, Members commit to facilitate access to the data collected by INSDC Members; and

Whereas, Members have the scientific and technical capabilities and facilities needed to participate; and

Whereas, Members agree that this arrangement provides the general framework for collaborating on the collection and distribution of nucleotide sequence data and associated metadata.

NOW, THEREFORE, the Member agrees to the following:

1. Scope of INSDC The International Nucleotide Sequence Database Collaboration (INSDC) is a global collaboration of

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NLM-NCBI

Activities on the databases

Databases at Bioinformation and DDBJ Center § DDBJ

	Annotated sequences	Capillary reads	NGS reads	Study	Sample	Assembly	Functional genomics	Variation	Genotype and phenotype	Metabolomics
NCBI	GenBank	Trace Archive	Sequence Read Archive	BioProject	BioSample	Assembly	GEO	dbSNP/dbVar	dbGaP	
EBI		Euro	pean Nucleotide	e Archive (EN		ArrayExpress	EVA/DGVa	EGA	MetaboLights	
DDBJ	DDBJ	Trace Archive	Sequence Read Archive	BioProject	BioSample	Assembly	GEA	JVar-SNP/SV	JGA	MetaboBank

INSDC: International Nucleotide Sequence Databank Collaboration

Databases at Bioinformation and DDBJ Center § DDBJ

	Annotated sequences	Capillary reads	NGS reads	Study	Sample	Assembly	Functional genomics	Variation	Genotype and phenotype	Metabolomics
NCBI	GenBank	Trace Archive	Sequence Read Archive	BioProject	BioSample	Assembly	GEO	dbSNP/dbVar	dbGaP	
EBI		Euro	European Nucleotide Archive (ENA)					EVA/DGVa	EGA	MetaboLights
DDBJ	ddbj -	race Archive	Sequence Read Archive	BioProject	BioSample	Assembly	GEA	JVar-SNP/SV	JGA	MetaboBank

INSDC: International Nucleotide Sequence Databank Collaboration

An example for the "trad" database

LOCUS DEFINITION	AB091058 2109 bp DNA linear BCT 02-SEP-2003 Gluconacetobacter xylinus cmcase, ccp genes for endo-beta-1,4-glucanase, cellulose complementing protein, complete	CDS
ACCESSION VERSION KEYWORDS	cds. AB091058 AB091058.1	
SOURCE	Gluconacetobacter xylinus <u>Gluconacetobacter xylinus</u> Bacteria; Proteobacteria; Alphaproteobacteria; Rhodospirillales; Acetobacteraceae; Gluconacetobacter.	
REFERENCE AUTHORS	1 (bases 1 to 2109) Kawano,S., Tajima,K., Uemori,Y., Yamashita,H., Erata,T.,	
TITLE	Munekata,M. and Takai,M. Direct Submission	BASE COUNT ORIGIN
JOURNAL	Submitted (28-AUG-2002) to the DDBJ/EMBL/GenBank databases. Contact:Kenji Tajima Hokkaido University, Graduate School of Engineering; N13W8,	1 cgt 61 gcg 121 tat
REFERENCE	Kita-ku, Sapporo, Hokkaido 060-8628, Japan 2	181 ggg 241 atg
AUTHORS	Kawano,S., Tajima,K., Uemori,Y., Yamashita,H., Erata,T., Munekata,M. and Takai,M.	301 ttc 361 ctg
TITLE	Cloning of Cellulose Synthesis Related Genes from Acetobacter xylinum ATCC23769 and ATCC53582: Comparison of Cellulose Synthetic Ability Between ATCC23769 and ATCC53582	421 gac 481 gtc 541 tcc
JOURNAL COMMENT	Unpublished (2002)	601 cgt 661 ctg
FEATURES source	Location/Qualifiers 12109	721 tgg 781 atg
	/db_xref=" <u>taxon:28448</u> " /mol_type="genomic DNA"	841 gcc 901 gga
	<pre>/note="synonym:Acetobacter xylinum" /organism="Gluconacetobacter xylinus" /ateria="among 53500"</pre>	961 gat 1021 gag
CDS	/strain="ATCC 53582" 101038 /codon start=1	1081 gaa 1141 cct 1201 ggc
	/gene="cmcase" /product="endo-beta-1,4-glucanase"	1261 aaa 1321 aaa
	/protein_id=" <u>BAC82540.1</u> " /transl_table=11	1381 ccg 1441 ccg
	/translation="MSVMAAMGGAQVLSSTGAFADTAPDAVAQQWAIFRAKYLRPSGR VVDTGNGGESHSEGQGYGMLFAASAGDLASFQSMWMWARTNLQHTNDKLFSWRFLKGH	1501 tgc 1561 cgc
	QPPVPDKNNATDGDLLIALALGRAGKRFQRPDYIQDAMAIYGDVLNLMTMKAGPYVVL MPGAVGFTKKDSVILNLSYYVMPSLLQAFDLTADPRWRQVMEDGIRLVSAGRFGQWRL PPDWLAVNRATGALSIASGWPPRFSYDAIRVPLYFYWAHMLAPNVLADFTRFWNNFGA	1621 agg 1681 ctt 1741 atg
	NALPGWVDLTTGARSPYNAPPGYLAVAECTGLDSAGELPTLDHAPDYYSAALTLLVYI ARAEETIK"	1741 atg 1801 atc 1861 tgg
		1921 aca

/codon start=1 /gene="ccp" /product="cellulose complementing protein" /protein id="BAC82541.1" /transl table=11 /translation="MSASGSDEVAGGGQAGSPQDFQRVLRSFGVEGGQYSYRPFVDRS FDVTGVPEAVERHFDQAEHDTAVEEQVTPAPQIAVAPPPPPVVPDPPAIVTETAPPPP VVVSAPVTYEPPAAAVPAEPPVOEAPVOAAPVPPAPVPPIAEOAPPAAPDPASVPYAN VAAAPVPPDPAPVTPAPQARVTGPNTRMVEPFSRPQVRTVQEGATPSRVPSRSMNAFP RTSASSISERPVDRGVADEWSPVPKARLSPRERPRPGDLSFFFQGMRDTRDEKKFFPV ASTRSVRSNVSRMTSMTKTDTNSSQASRPGSPVASPDGSPTMAEVFMTLGGRATELLS PRPSLREALLRRRENEEES" 343 a 661 c 661 g 444 t tteettta tgteggteat ggeggegatg ggaggggege aggtgettte atecaeeggt

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gttegeag acacegeeee egatgeggte gegeageaat gggeeatett eegegeeaag tottogto coagoggaog tgtogtggat acgggcaatg gtggcgaato coatagtgag gcagggct atggcatgct ctttgccgcg tcggcggggg accttgcgtc gttccagtcg gtggatgt gggcgcgcac caacctgcag cataccaatg acaagctgtt ttcctggcgg cctcaagg ggcatcagcc cccggtgccc gacaagaaca atgccacaga tggcgacctg gategege ttgegettgg tegtgeggge aagegtttee agegeeeega ttacatteag legecatgg ceatttatgg egatgtgetg aacetgatga egatgaagge gggaeegtat cgtcctca tgcccggtgc tgtcggcttt accaagaagg acagcgtgat cctcaacctg ctattacg tcatgccctc gctgctgcag gcgttcgacc ttacggccga cccgcgctgg tcaggtga tggaagacgg gattcgcctt gtttccgccg gccgtttcgg gcagtggcgc gccccccg actggctggc ggtgaatcgc gccaccggtg cgctgtcgat cgcatcggga geegeege getttteeta tgatgegatt egggtgeege tttattttta ttgggegeat gctggcgc cgaacgtgtt ggctgatttc acccgattct ggaataattt cggggctaat cctgccag gatgggttga tctgacaaca ggggcgcgtt cgccgtacaa cgccccgcct atatettg etgttgeega atgeaegggg ettgattetg eeggggaaet eeegaeaetg tcatgcgc ccgattatta ttccgcagcg ttgacgctgc tcgtttacat cgcgcgggcg ggagacta taaagtgagt gcttcagggt ctgatgaggt ggctggggga gggcaggctg agteegea ggatttteag egggteetge gttettttgg tgtegaaggt gggeagtatt taccggcc gtttgttgac cgttcctttg atgtgacagg cgtgcccgag gctgttgaaa cacttega teaggeggag catgaeacgg eggttgagga geaggteact ecegegeeae atcgcggt cgcaccgcca ccgccgccag tcgttcctga cccgcccgcc atcgtgacgg nacegegee ceegeegeet gtegtggtea gegeteeggt caegtatgaa eceeeggetg gccgtgcc ggcagagcct cccgttcagg aagcccccgt gcaggcggcg ccggttcccc gegeetgt geeeegatt geggageagg eteeteege ggegeeggae eeggeateeg prograting gaacgroups grageaccor treeaccross record gradesets ccgcagge gegegtgaeg gggeegaaea eeegtatggt ggageeettt teeegeeege gtccgcac ggtgcaggag ggggcaaccc cgtcacgtgt accttcgcgt tcaatgaacg ttcccccg cacatcagca tcgtccataa gtgagcgtcc ggtggacagg ggtgttgccg gaatggag teetgtteeg aaggeaegee teageeegeg ggagegteeg egteeegeg ctgagett tttettteag gggatgegeg acaecegtga tgaaaagaag ttettteeeg gcgtccac gcgatcagtt cgttctaatg tttccaggat gaccagcatg accaagacag 1981 ccacaatggc cgaagtgttc atgacgctgg gtggtcgtgc gacggaactc ctcagccccc 2041 gtccttcgct gcgggaggcg ctgttgcgtc gtcgtgaaaa cgaagaagaa tcctaaggcc 2101 ctatattca

Trad in DDBJ





The total number of registrations related to DDBJ entries in 2023 were ca. 6,000; about a half of submissions made by web interface and a half were pre-format MSS submittions. (The rapid increase in the number of MSS submissions in 2021 was due to a large amount of MAG (Metagenome-Assembled Genome) from the University of Tokyo.) As of December 2023, the total number of bases is about 27 trillion, and the total number of sequences is about 4.0 billion.

Databases at Bioinformation and DDBJ Center § DDBJ

	Annotated sequences	Capillary reads	NGS reads	Study	Sample	Assembly	Functional genomics	Variation	Genotype and phenotype	Metabolomics
NCBI	GenBank	Trace Archive	Sequence Read Archive	BioProject	BioSample	Assembly	GEO	dbSNP/dbVar	dbGaP	
EBI	Eurc pean Nucleotide Archive (ENA)					ArrayExpress	EVA/DGVa	EGA	MetaboLights	
DDBJ	DDBJ	Trace Archive	Sequence Read Archive	BioProject	BioSample	Assembly	GEA	JVar-SNP/SV	JGA	MetaboBank

INSDC: International Nucleotide Sequence Databank Collaboration

SRA in DDBJ (DRA)





DDBJ SRA (DRA) registrations in 2023 were 2,361 (116TB). The total SRA and fastq file sizes published were 16.5PB and 1.4PB, respectively. (From April 2017 to May 2019, the amount did not grow due to the cessation of mirroring of EBI/NCBI data.)

Databases at Bioinformation and DDBJ Center § DDBJ

	Annotated sequences	Capillary reads	NGS reads	Study	Sample	Assembly	Functional genomics	Variation	Genotype and phenotype	Metabolomics
NCBI	GenBank	Trace Archive	Sequence Read Archive	BioProject	BioSample	Assembly	GEO	dbSNP/dbVar	dbGaP	
EBI		Euro	pean Nucleotide	e Archive (EN		ArrayExpress	EVA/DGVa	EGA	MetaboLights	
DDBJ	DDBJ	Trace Archive	Sequence Read Archive	BioProject	BioSample	Assembly	GEA	JVar-SNP/SV	JGA	MetaboBank

INSDC: International Nucleotide Sequence Databank Collaboration

GEA: Genomic Expression Archive



Samples



GEA release (total data volume)



GEA's submissions and samples numbers by year.

Number of relased experiments and samples from GEA.

GEA is a database of gene expression data from microarrays and NGS in DDBJ. In 2023, we had 79 data submissions, and 240 Experiments and ca. 4,000 Samples in total, were publicly available.

Japanese Genotype-phenotype Archive (JGA) § DDBJ



A control-access database for personal genome phenotype/genotype information operated jointly with NBDC. Registration and use of the database are subject to NBDC review. In 2023, 87 data and 97 TB were submitted. As of the end of 2023, 358 studies, 344,702 samples, and 895 TB of data were released. The number of applications approved by NBDC in 2022 totaled 49 (domestic: 17, overseas: 18). A cumulative total number is 263 applications approved.

MetaboBank



Samples

1,200

1,000

800

600

400

200

0

2023



MetaboBank submissions and sample numbers by year.

The number of released studies and samples from MetaboBank.

MetaboBank Version 2 uses an IDF/SDRF registration form as a metadata input format equivalent to GEA and EBI's MetaboLight. Web content, including the registration form and guidelines and the workflow from data registration to publication, have been prepared and publicized. The data registration for 2023 is 12 Studies, and 25 Studies were open at the end of 2023.

NIG Supercomputer: an infrastructure for lifescience



Osamu Ogasawara, PhD Project Associate Professor Head of HPC Division



- Singularity/Docker containers are available for users.
- As of last year (December 31, 2023), there were 1,953 total users.
- The large-capacity archive storage system dedicated to the database was a hierarchical storage system with 15 PB of disk and 15 PB of tape. It was replaced with a Lustre file system with an effective capacity of 40 PB (DDN, Hitachi, Ltd.) As a result, it is possible to mount DRAs and other files from all computation nodes of the supercomputer at a higher speed. The 15 PB tape part was purchased and continues to be used as an offline backup.

Capacity Buildings: Vietnam, Indonesia and Thailand (2023)







for Life and Health Sciences

National Institute of Genetics: www.nig.ac.jp

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Center for Frontier Research KUBO, Fumi Systems Neuroscience Lob IP MURAYAMA, Yasuto Chromosome Biochemistry

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Genome Informatics lab.

- ¥Yaz Nakamura, Professor
- ¥Yasuhiro Tanizawa, Assist. Professor
- Takako Mochizuki, Project Researcher
- Mika Sakamoto, Project Researcher
- Satomi Asano, Project Researcher
- Sakamoto, Technical Assistant
- Fumi Hayashi, Technical Assistant
- Se Misato Godo, Technical Assistant
- Dauyey Kaisar, D5 Student [coral fish evolution]
- Hanjie Mao, D5 Student [cat genetic diseases]
- Mohamed Elmanzalawi, D1 Student [human rare diseases]

Takatomo Fujisawa, Researcher (DDBJ)





Nakamura lab's genome works



Citrus unshiu An orange **@**農研機構



Gryllus *bimaculatus* A cricket 東京大学



Marchantia polymorpha A liverwort

Nitzschia spp. A non-photosynthetic diatom





京都大学 KYOTO UNIVERSITY

et al.





Tea tree (2,500)





Strawberry (400)

Felis catus







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information obtained from Hi-C and ontical manning data. Homology-based and ab initio gene

bioRxiv is receiving many new papers on coronavirus SARS-CoV-2. A reminder: these are preliminary reports that have not been peer-reviewed. They should not be regarded as conclusive, guide clinical practice/health-related behavior, or be reported in news media as established information.

New Results	Previous	Next 🗲
AnAms1.0: A high-quality chromosome-scale assembly of a domestic cat	Posted May 19, 2020.	
 Felis catus of American Shorthair breed Sachiko Isobe, D Yuki Matsumoto, Claire Chung, D Mika Sakamoto, D Ting-Fung Chan, D Hideki Hirakawa, Genki Ishihara, D Hon-Ming Lam, Shinobu Nakayama, Shigemi Sasamoto, D Yasuhiro Tanizawa, Akiko Watanabe, Kei Watanabe, Masaru Yagura, D Yasukazu Nakamura doi: https://doi.org/10.1101/2020.05.19.103788 	Download PDF	 Email Share Citation Tools
This article is a preprint and has not been certified by peer review [what does this mean?].	🎔 Tweet 🚺 いいね! 1	
Abstract Full Text Info/History Metrics Preview PDF	COVID-19 SARS-CoV	/-2 preprints from
Abstract	medRxiv and bioRxiv	• •
The domestic cat (<i>Felis catus</i>) is one of the most popular companion animals in the world.	Subject Area	
Comprehensive genomic resources will aid the development and application of veterinary	Genomics	
medicine including to improve feline health, in particular, to enable precision medicine which is		
promising in human application. However, currently available cat genome assemblies were	Subject Areas	
mostly built based on the Abyssinian cat breed which is highly inbred and has limited power in representing the vast diversity of the cat population. Moreover, the current reference assembly	All Articles	
remains fragmented with sequences contained in thousands of scaffolds. We constructed a	Animal Behavior and Cog	nition
reference-grade chromosome-scale genome assembly of a domestic cat, Felis catus genome	Biochemistry	
of American Shorthair breed, Anicom American shorthair 1.0 (AnAms1.0) with high contiguity	Bioengineering	
(scaffold N50 > 120 Mb), by combining multiple advanced genomic technologies, including	Bioinformatics	
PacBio long-read sequencing as well as sequence scaffolding by long-range genomic	Discharter	

Table 3. Statistics of the AnAms1.0 and felcat genomes.

	AnAm	s1.0	felC	at9
	All scaffolds	Scaffolds correspond to nucleic chromosomes	All scaffolds	Scaffolds correspond to nucleic chromosomes
No. of sequences	20	19	4,508	19
Total length, bp	2,493,141,6	2,493,140,262	2,521,863 \5	2,460,251,910
Avg. length, bp	124,657	131,217,909	55	129,486,943
Max. length, bp	243,50	243,504,654	242,1	242,100,913
Min. length, bp	1,3	41,750,578	1 9	44,648,284
N50 length, bp	151,107,6	151,107,676	149,751 9	149,751,809
A	718,322,616	718,322,255	721,302,099	703,154,254
Т	719,032,844	719,032,444	721,735,169	703,748,672
G	517,632,441	517,632,143	516,811,973	504,031,439
С	517,568,802	517,568,480	516,603,929	503,993,730
Ν	0.83	0.83	1.8	1.84
Total, ATGC	2,472,556,703	2,472,555,322	2,476,453,170	2,414,928,095
GC%, ATGC	41.9	41.9	41.7	41.7

Expansion of Cats-I for vet information infrastructure



Search



Complete genome project for Big dogs [new]

- For Dogs and Human Healthcare



Canis lupus

Aiming to make T2T (376 scaffolds \Rightarrow 39!)

Pacbio Revio Hifi reads

	<u> </u>				•
	ASM1132765v1	GCA_011327655.1	Canis lupus familiaris (dog)	Chihuahua long coa	•
	ASM1163472v1	GCA_011634725.1	Canis lupus familiaris (dog)	Chihuahua long coa	• • •
	UniMelb_Wolf_Refassem_1	GCA_007922845.1	Canis lupus (gray wolf)	Chinese wolf (breed)	• • •
	ASM864105v3	GCA_008641055.3	Canis lupus familiaris (dog)	German Shepherd (• • •
	UU_Cfam_GSD_1.0	GCA_011100685.1 GCF_011100685.1	Canis lupus familiaris (dog)	German Shepherd (NCBI RefSeq	• • •
	ASM544666v1	GCA_005446665.1	Canis lupus familiaris (dog)	Great Dane (breed)	•
	UMICH_Zoey_3.1	GCA_005444595.1 GCF_005444595.1	Canis lupus familiaris (dog)	Great Dane (breed)	• • •
	TAMU_N220234	GCA_031771975.1	Canis lupus familiaris (dog)	Irish Wolfhound (br	• • •
	TAMU_N210636	GCA_040939265.1	Canis lupus familiaris (dog)	Irish Wolfhound (br	• • •
$\left(\right)$	ASM1204487v1	GCA_012044875.1	Canis lupus familiaris (dog)	Labrador retriever (:
	ASM1204501v1	GCA_012045015.1	Canis lupus familiaris (dog)	Labrador retriever (* *
	Yella_v2	GCA_031165255.1	Canis lupus familiaris (dog)	Labrador retriever (• •
	ROS_Cfam_1.0	GCA_014441545.1 GCF_014441545.1	Canis lupus familiaris (dog)	Labrador retriever (NCBI RefSeq	:
	ASM18141v1	GCA_000181415.1	Canis lupus familiaris (dog)	poodle (breed)	• • •
	ASM325472v2	GCA_003254725.2 GCF_003254725.2	Canis lupus dingo (dingo)	Sandy (isolate) NCBI RefSeq	• •
	Clu-1	GCA_034620435.1	Canis lupus (gray wolf)	wolf_2809A (isolate)	•



precision medicine for companion animals



"Personal" Medicine for companion animals

In traditional veterinary medicine, the same treatment is given to any individual where some cats are not responsive to specific treatments. With the genomic approach, individually suited treatments can be applied based on individual traits predicted from genome information for more effective and efficient veterinary medicine.

Acknowledgments: DDBJ members





https://www.ddbj.nig.ac.jp/about/staff-e.html

