

Digital Sequence Information : An Overview

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DSI and the Nagoya Protocol

- What we now call Digital Sequence Information was raised repeatedly at various times during the negotiations (EU supported a submission of independent research on this in UNEP/CBD/WG-ABS/3/INF/4). But, the discussion did not go anywhere. It has now come back.
- During the course of the negotiations an increasing number of organisms had their genomes sequenced and technology for sequencing at scale radically improved (next generation sequencing or NGS). As of April GenBank contained in excess of 2 trillion DNA bases in approx. 652 million sequences (strings of bases of varying lengths). In 2003 there were only 30 million sequences.
- This presentation will walk through some of the basics of DSI before turning to trends, costs, and geographic distribution. It will conclude with discussion of key issues arising and potential options.

Key Issues

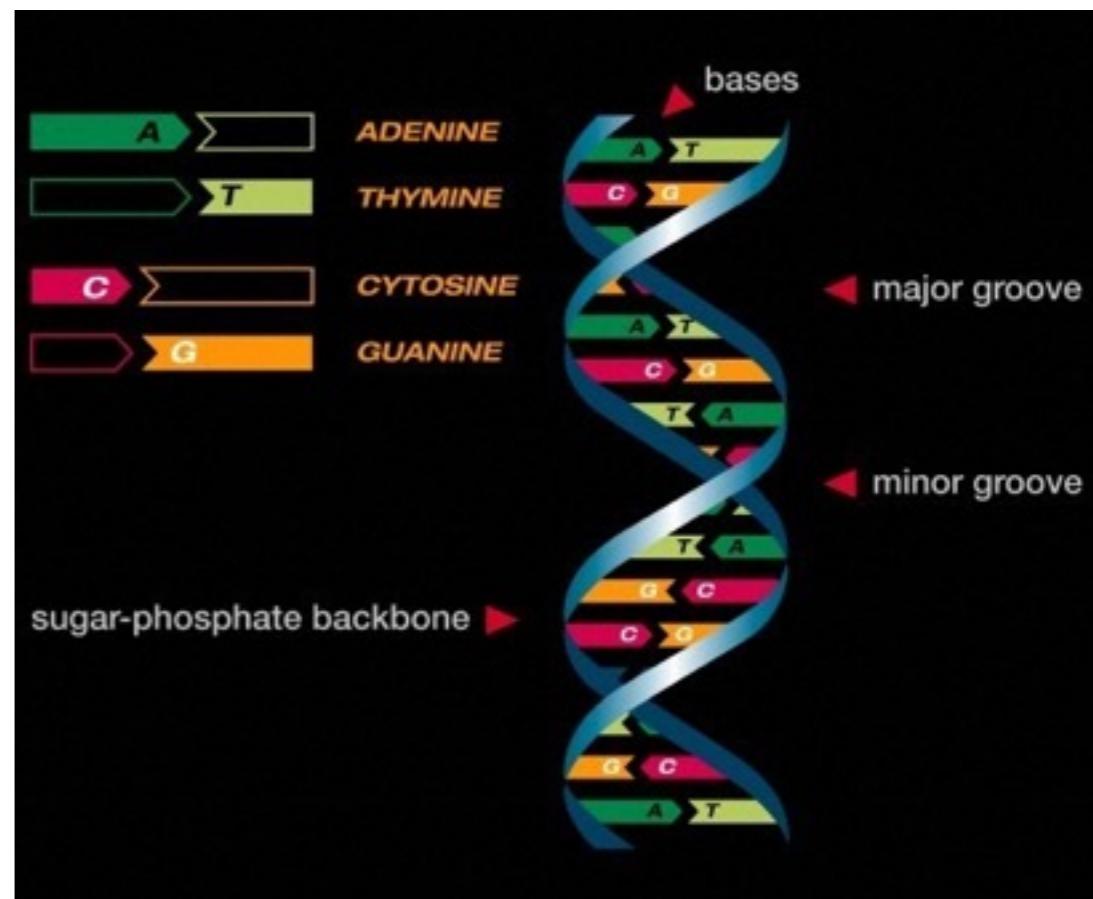
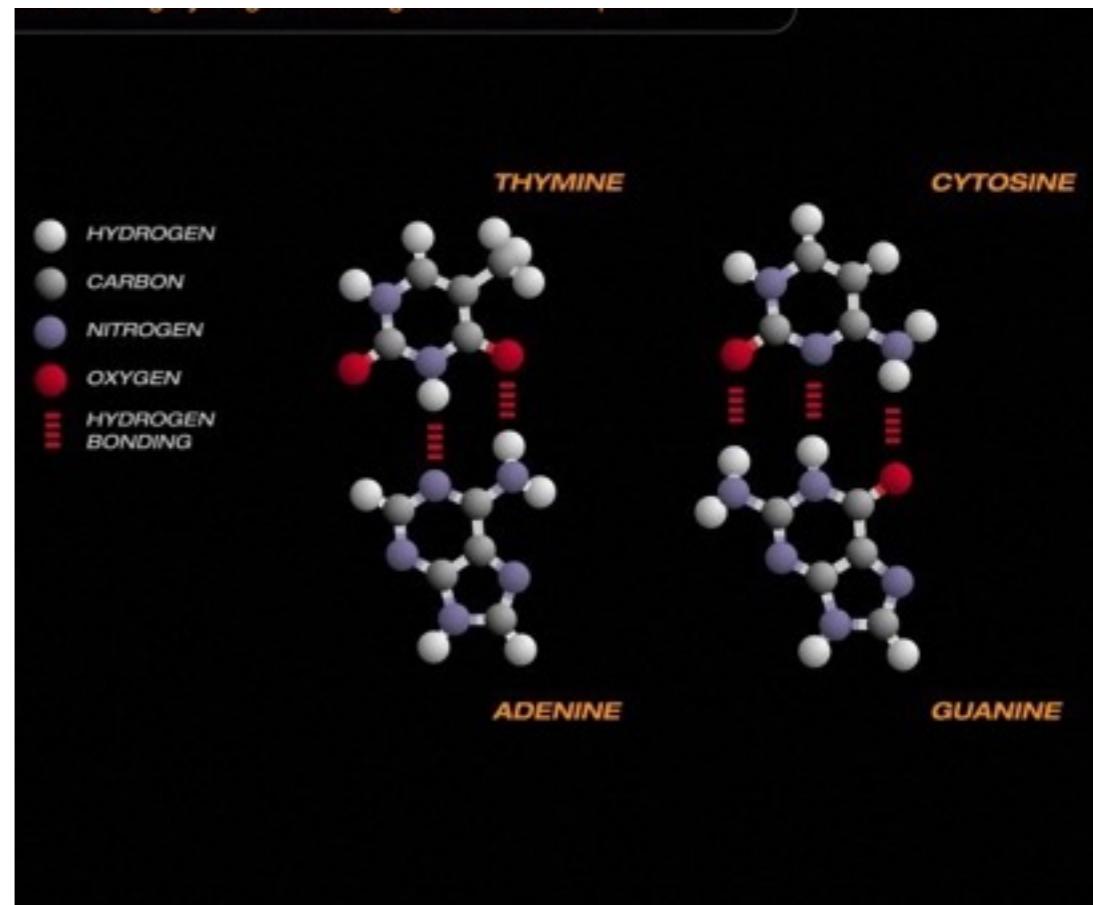
- Clarifying Trends
- Clarifying the who, what, where (and where from)
- Actual and Potential Uses
- Questions of Value
- Terms and Conditions of Databases
- Options for Parties and Potential Consequences
- What else needs to be established?

Key Questions

- What are the terms and conditions under which international electronic transfers are made?;
- Should electronic transfers be regulated?;
- What are the potential costs and benefits of the regulation of electronic transfers?;
- What forms of regulation of electronic transfers might be appropriate? (UNEP/CBD/WG-ABS/3/INF/4 at 15)

DNA & RNA

- Deoxyribonucleic acid (DNA) & Ribonucleic acid (RNA) molecules are the chemical foundations of cells and organisms.
- DNA molecules consist of four bases (A, C, T & G) that bind to each other in an ordered way (A & T, C & G) described as base pairs. RNA = Uracil instead of Thymine.
- There are different types of DNA and RNA molecules that are described in terms of sources such as mitochondrial DNA (mDNA) or functions (e.g. messenger or transfer = mRNA, tRNA)



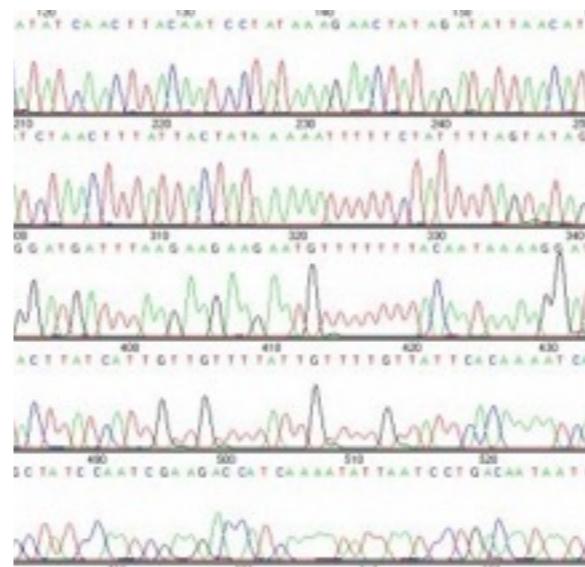
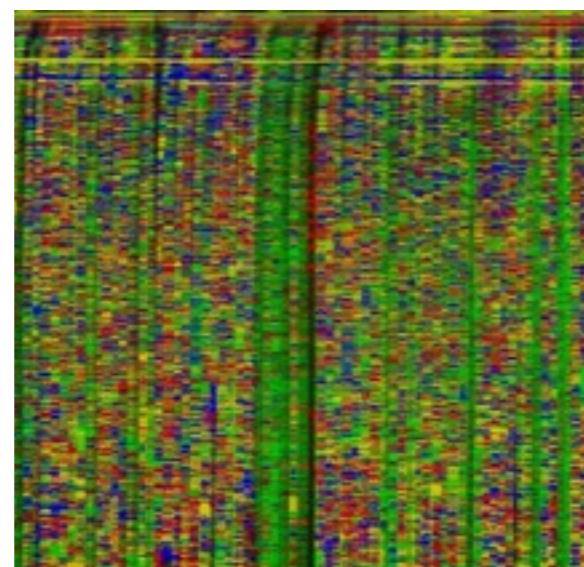
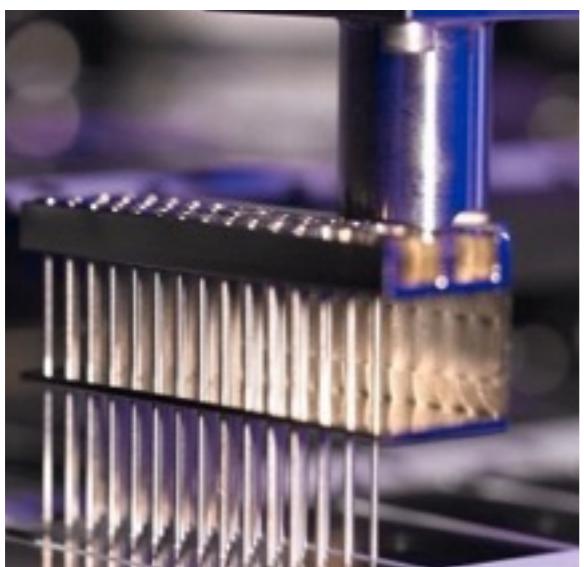
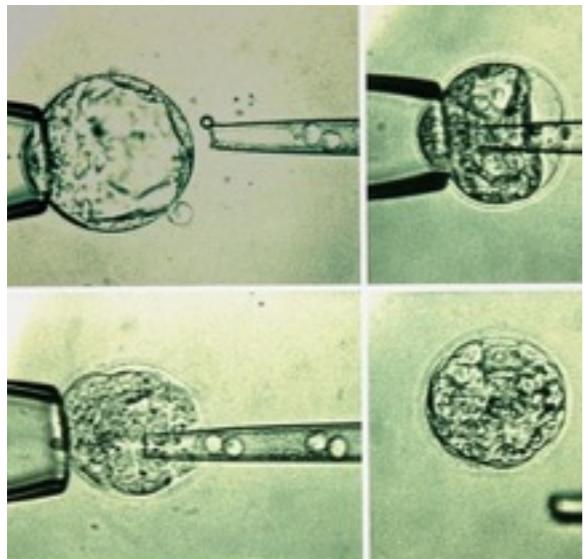
Amino Acids

- The ordering of DNA codons (arrangements of four bases) are associated with the expression of amino acids that form the basis for building proteins
- There are 20 main amino acids that are expressed through codons.
- So the TTTC codon forms Leucine or Leu. While TTCG forms Serine or Ser.
- DNA is transcribed into amino acids and structured into proteins through bonding with RNA as the messenger that triggers gene expression in the cell.



Sequencing

From Sanger Sequencing to Next Generation
Sequencing (NGS)



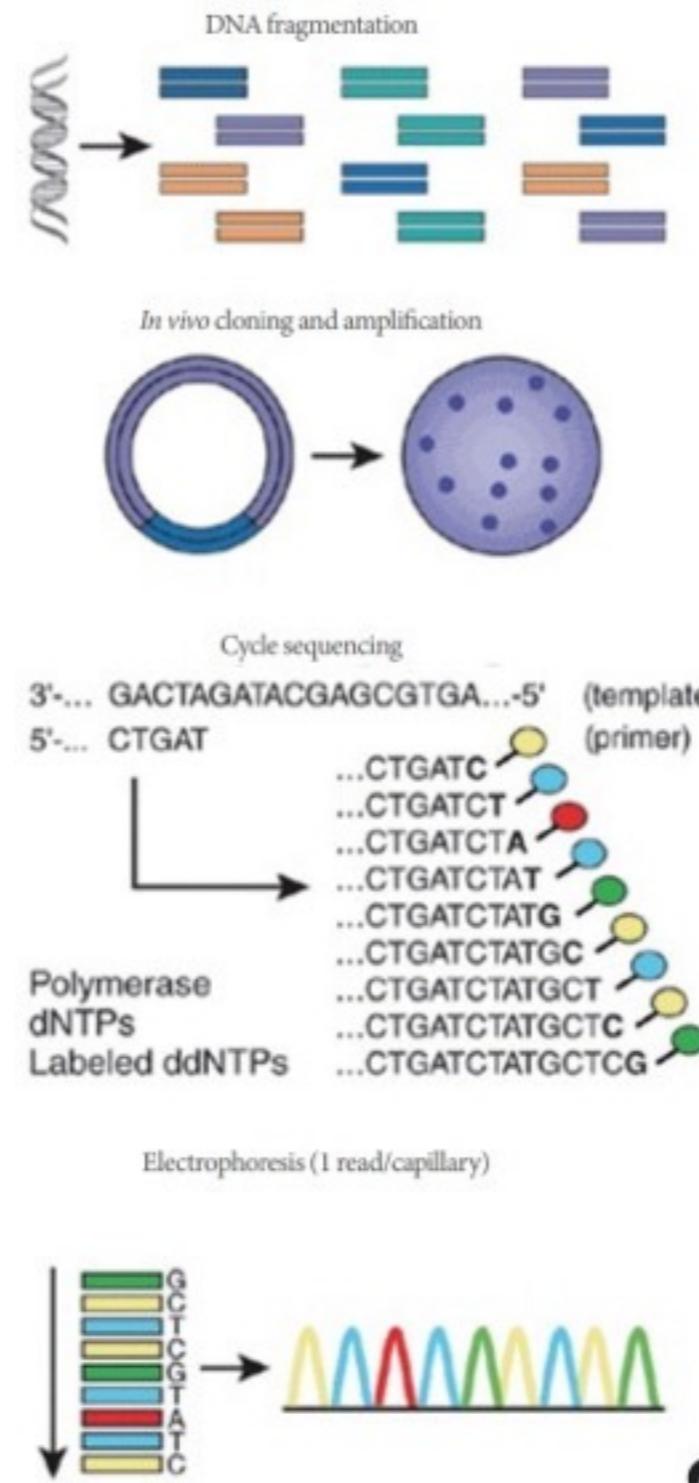
A photograph of a computer screen displaying a sequence of DNA bases. The sequence is written in a large, bold font, with each base color-coded (A=green, T=red, C=blue, G=black). Below the sequence, a smaller, standard text font displays the same sequence for comparison.

Sanger Sequencing (1977). extract into plasmids, culture colonies, extract & clean, sequence (gel), map

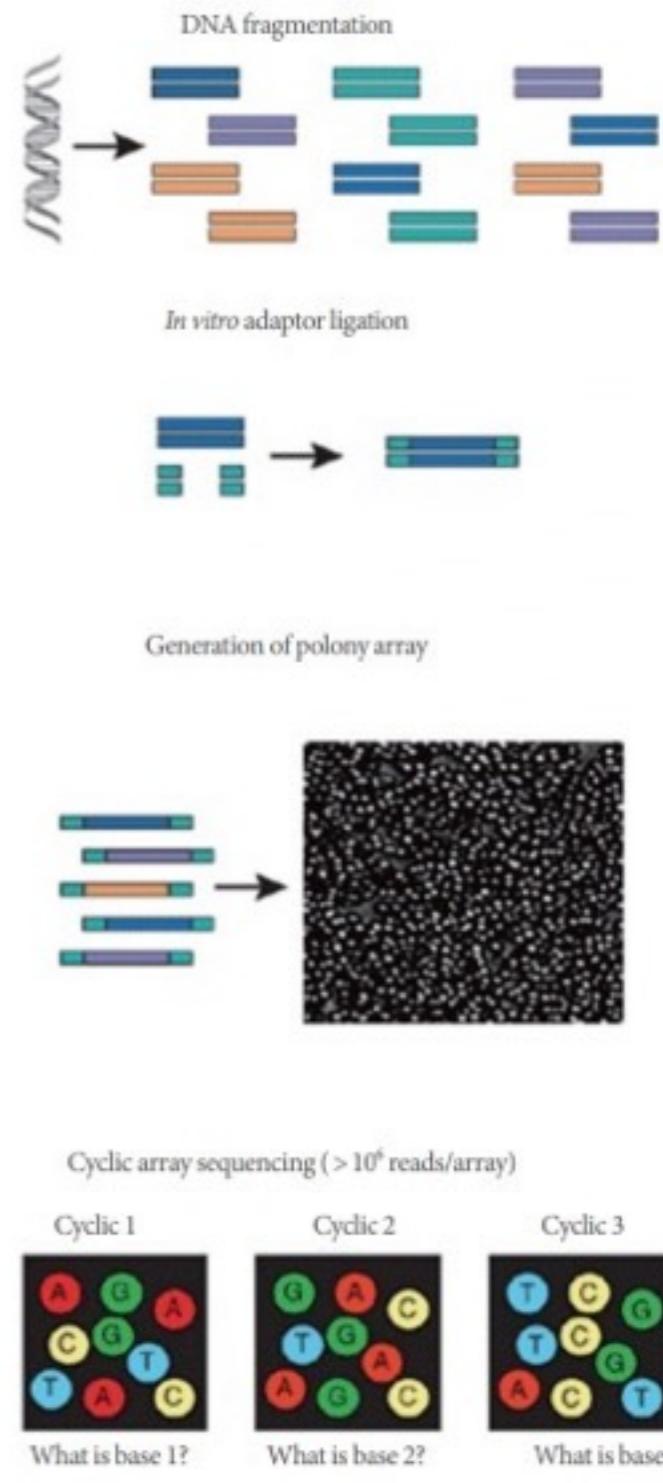
NGS

- Sanger Sequencing is accurate but slow. Next Generation Sequencing:
- Allows for the construction of libraries;
- No *in vivo* cloning and colony picking. Done *in vitro*;
- Can be organised in arrays and highly parallel so can sequence faster and on a larger scale;
- Approaches include: pyrosequencing, sequencing by synthesis, ligation and phospholinked real time sequencing;
- Key companies include Roche, Illumina, Oxford Nanopore, Qiagen, Life Technologies, Complete Genomics, Helicos Biosciences, Pacific Biosciences.

Sanger Sequencing



Next Generation Sequencing



Trends

Deposits, Costs, Organisms and Actors

International Nucleotide Sequ X

www.insdc.org

INSDC International Nucleotide Sequence Database Collaboration

ABOUT INSDC POLICY ADVISORS DOCUMENTS

International Nucleotide Sequence Database Collaboration

- The International Nucleotide Sequence Database Collaboration (INSDC) is a long-standing foundational initiative that operates between [DDBJ](#), [EMBL-EBI](#) and [NCBI](#). INSDC covers the spectrum of data raw reads, through alignments and assemblies to functional annotation, enriched with contextual information relating to samples and experimental configurations.

Data type	DDBJ	EMBL-EBI	NCBI
Next generation reads	Sequence Read Archive	European Nucleotide Archive (ENA)	Sequence Read Archive
Capillary reads	Trace Archive		Trace Archive
Annotated sequences	DDBJ		GenBank
Samples	BioSample		BioSample
Studies	BioProject		BioProject

- The INSDC advisory board, the [International Advisory Committee](#), is made up of members of each of the databases' advisory bodies. At their most recent meeting, members of this committee unanimously endorsed and reaffirmed the existing data-sharing policy of the three databases that make up the INSDC, which is stated below.
- Individuals submitting data to the international sequence databases should be aware of [INSDC policy](#).

How to submit data

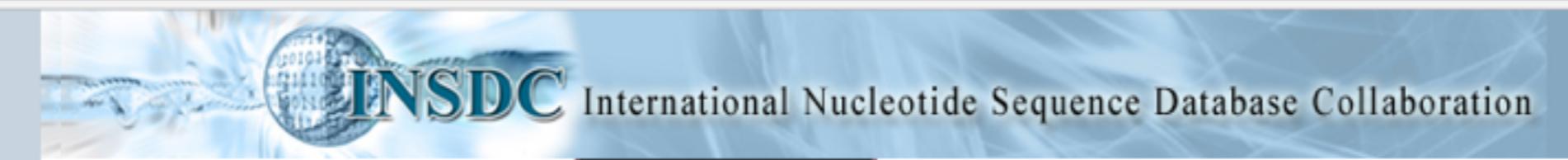
- For full details of how to submit data to the databases, please select a collaborating partner.
- [DDBJ](#), [ENA](#), [GenBank](#)
- The INSDC Feature Table Definition Document is available [here](#).

INSDC
International Nucleotide Sequence Database Collaboration

Site maintained by the External Services team at [EMBL-EBI](#) | [Terms of Use](#) | [Privacy](#) | [Cookies](#)

Key database collaboration

INSDC is made up of EMBL-EBI (EU), GenBank (US) & DDBJ (Japan)



ABOUT INSDC

POLICY

ADVISORS

DOCUMENTS



International Nucleotide Sequence Database Collaboration Policy

Soren Brunak, Antoine Danchin, Masahira Hattori, Haruki Nakamura, Kazuo Shinozaki, Tara Matise, Daphne Preuss (2002)
Nucleotide Sequence Database Policies
Science 298 (5597): 1333 15 Nov 2002

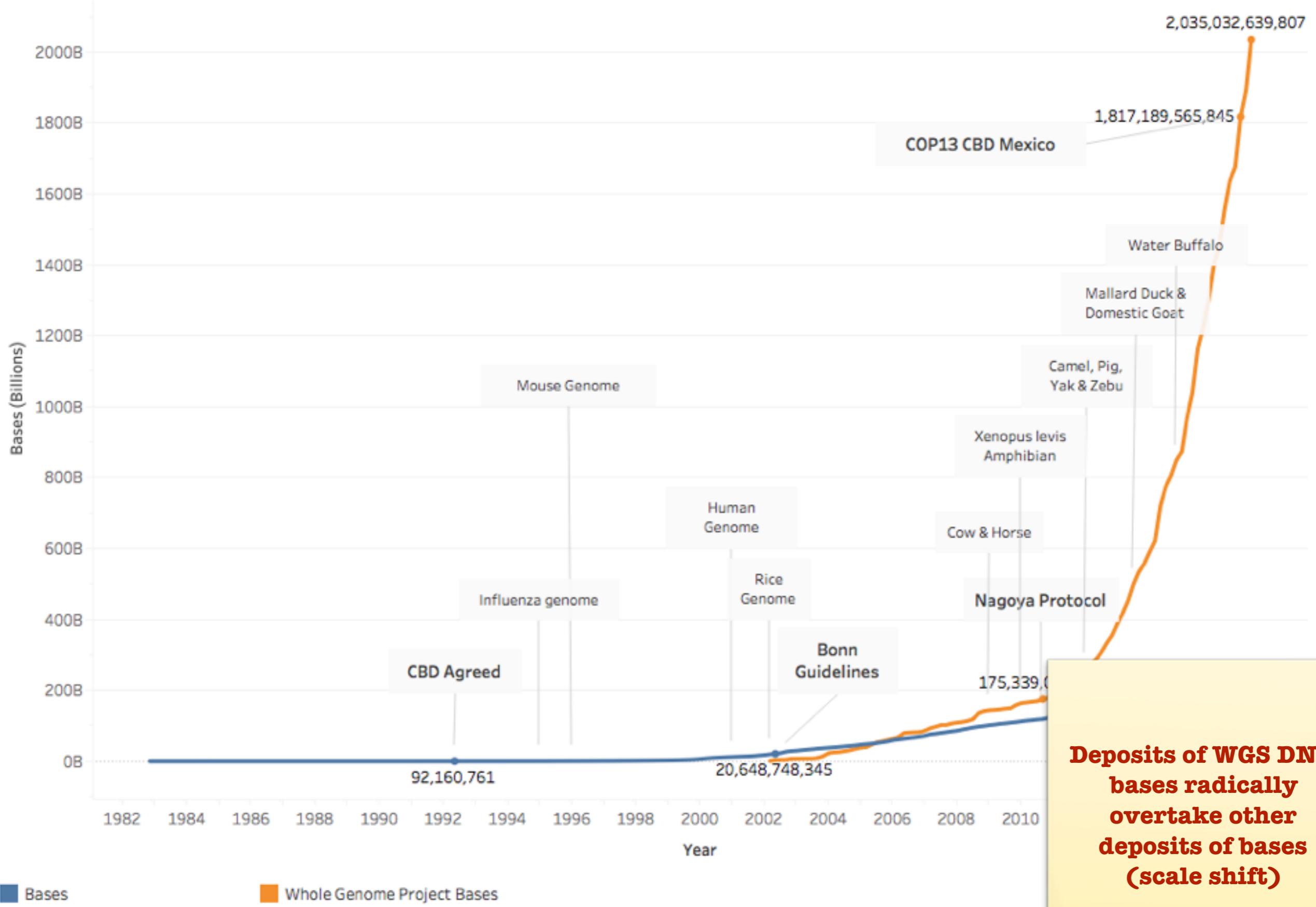
1. The INSD has a uniform policy of free and unrestricted access to all of the data records their databases contain. Scientists worldwide can access these records to plan experiments or publish any analysis or critique. Appropriate credit is given by citing the original submission, following the practices of scientists utilizing published scientific literature.
2. The INSD will not attach statements to records that restrict access to the data, limit the use of the information in these records, or prohibit certain types of publications based on these records. Specifically, no use restrictions or licensing requirements will be included in any sequence data records, and no restrictions or licensing fees will be placed on the redistribution or use of the database by any party.
3. All database records submitted to the INSD will remain permanently accessible as part of the scientific record. Corrections of errors and update of the records by authors are welcome and erroneous records may be removed from the next database release, but all will remain permanently accessible by accession number.
4. Submitters are advised that the information displayed on the Web sites maintained by the INSD is fully disclosed to the public. It is the responsibility of the submitters to ascertain that they have the right to submit the data.
5. Beyond limited editorial control and some internal integrity checks (for example, proper use of INSD formats and translation of coding regions specified in CDS entries are verified), the quality and accuracy of the record are the responsibility of the submitting author, not of the database. The databases will work with submitters and users of the database to achieve the best quality resource possible.

INSDC

International Nucleotide Sequence Database Collaboration

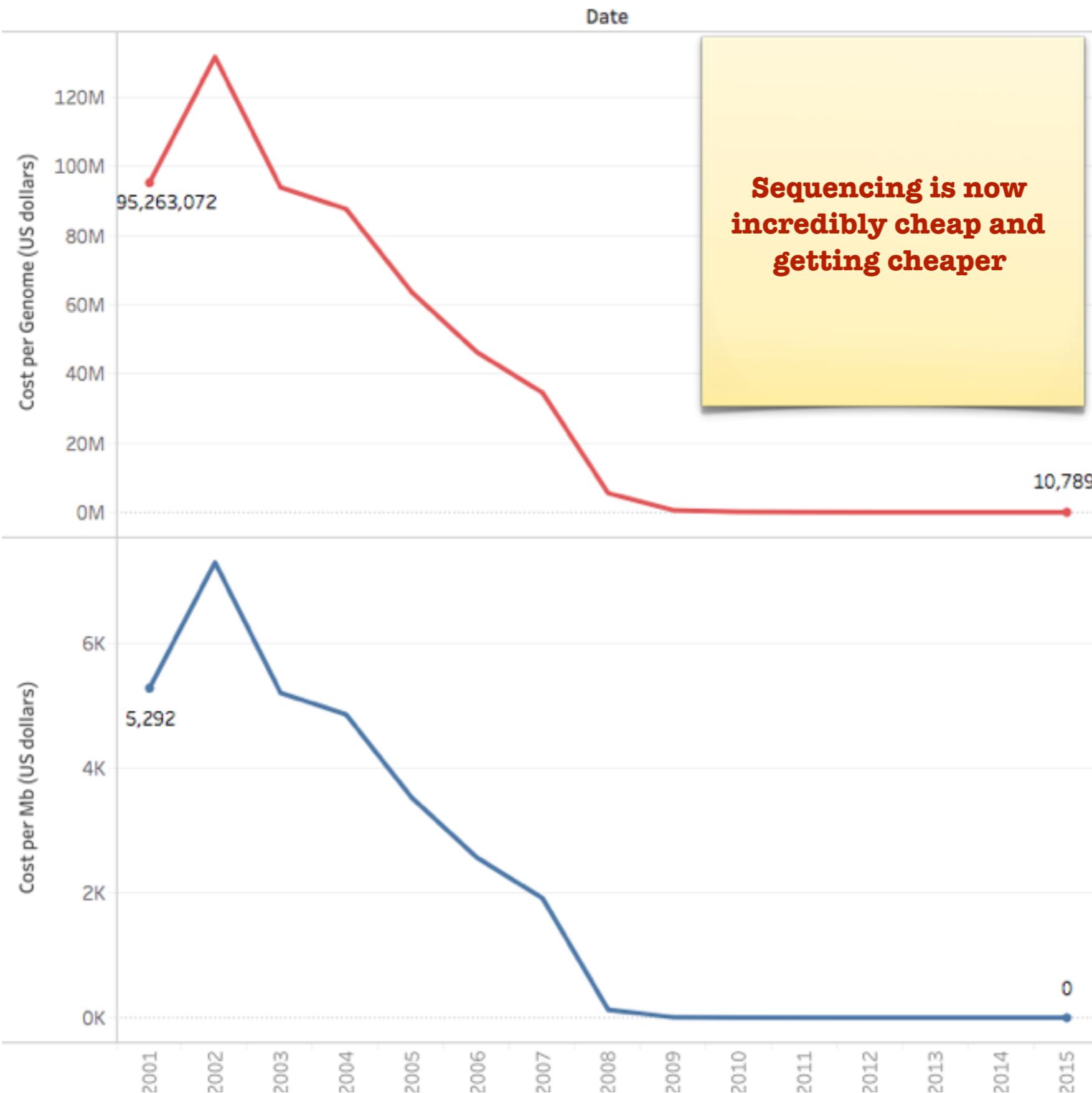
**Open Access Policy
with no conditions**

Genbank: Trends in Bases & Whole Genome Project Bases (cumulative) with landmarks

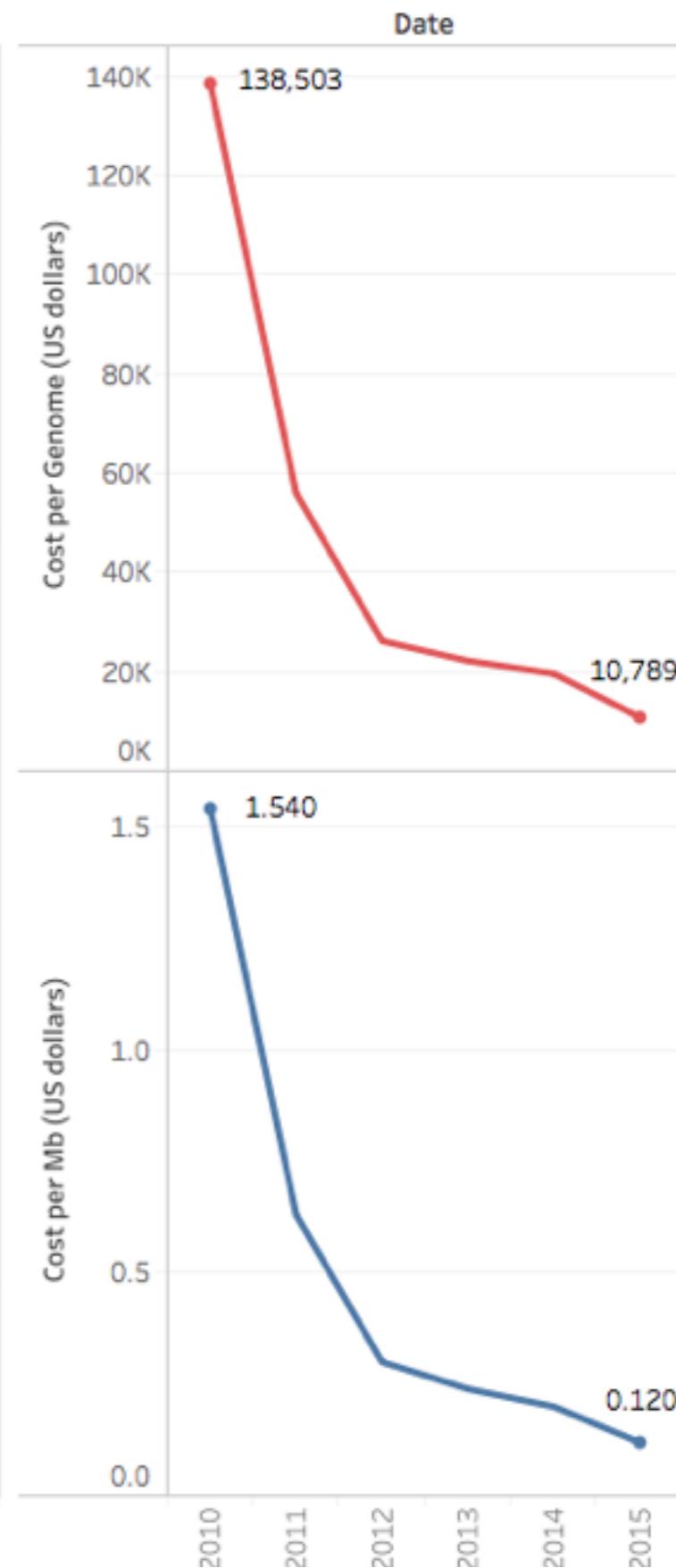


Deposits of WGS DNA bases radically overtake other deposits of bases (scale shift)

Trends in Sequencing Costs (US dollars)

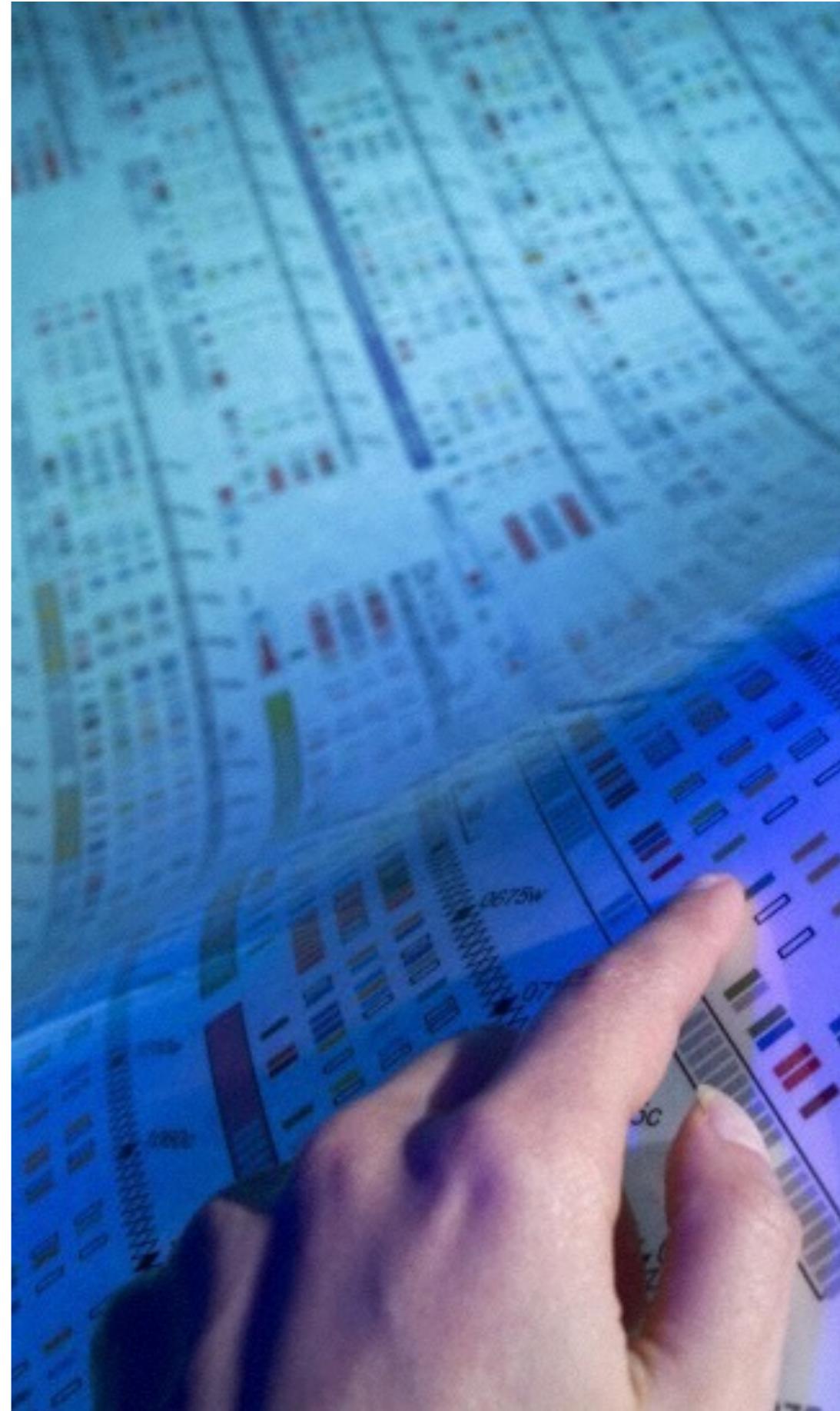


Trends 2010 to 2015

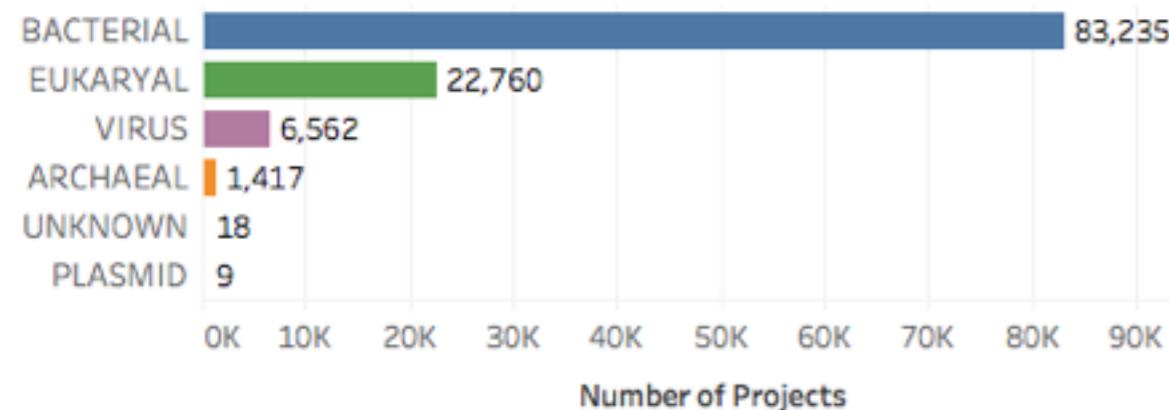


Whole Genome Sequencing Projects

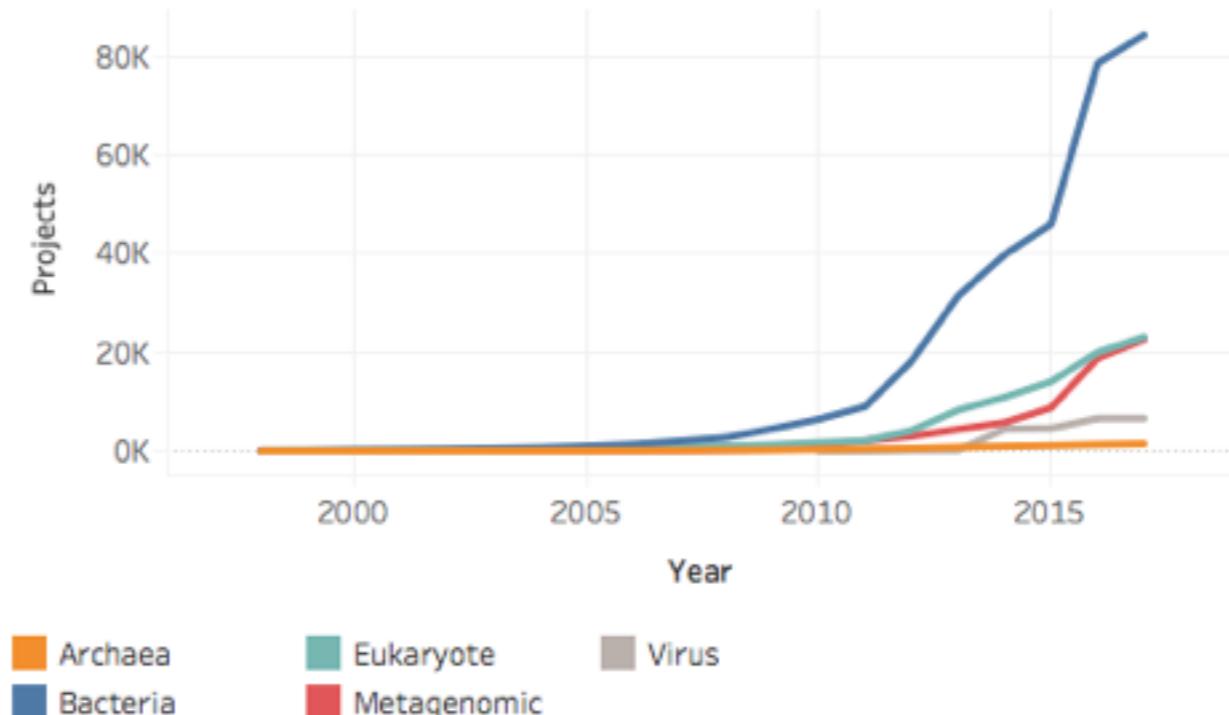
- A growing trend towards whole genome sequencing as sequencing costs fall;
- The Genomes Online Database (GOLD) is the main source of data on global projects;
- Data on projects is submitted voluntarily and may be incomplete. Data fields on funding are incomplete and others require further clean up... with those caveats...



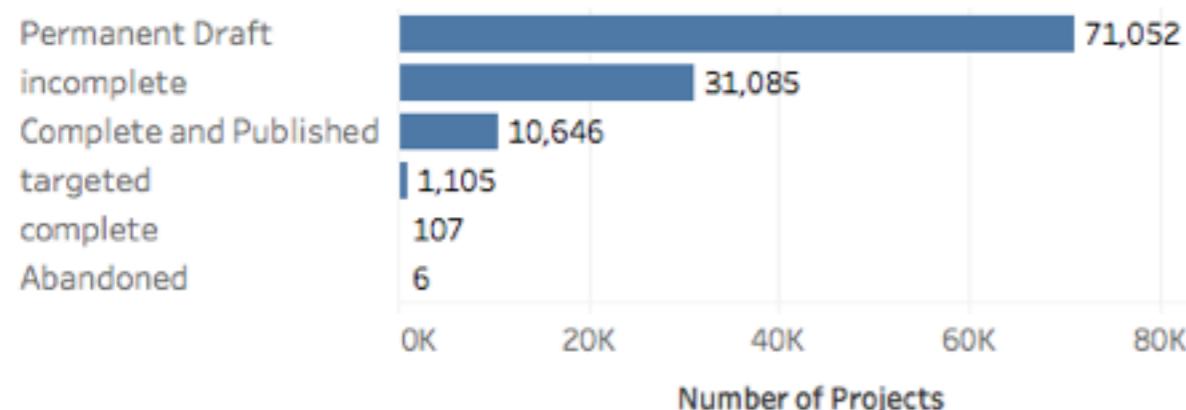
Whole Genome Projects by Domain



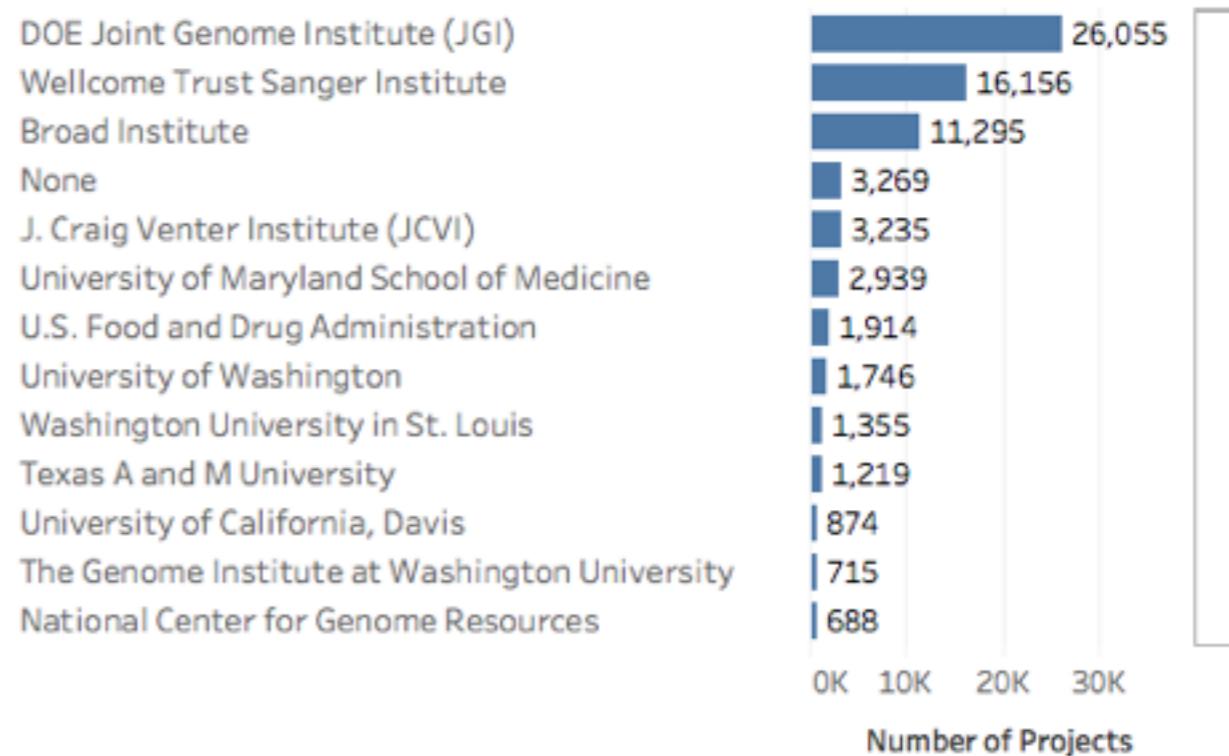
Genome Project Trends by Domain

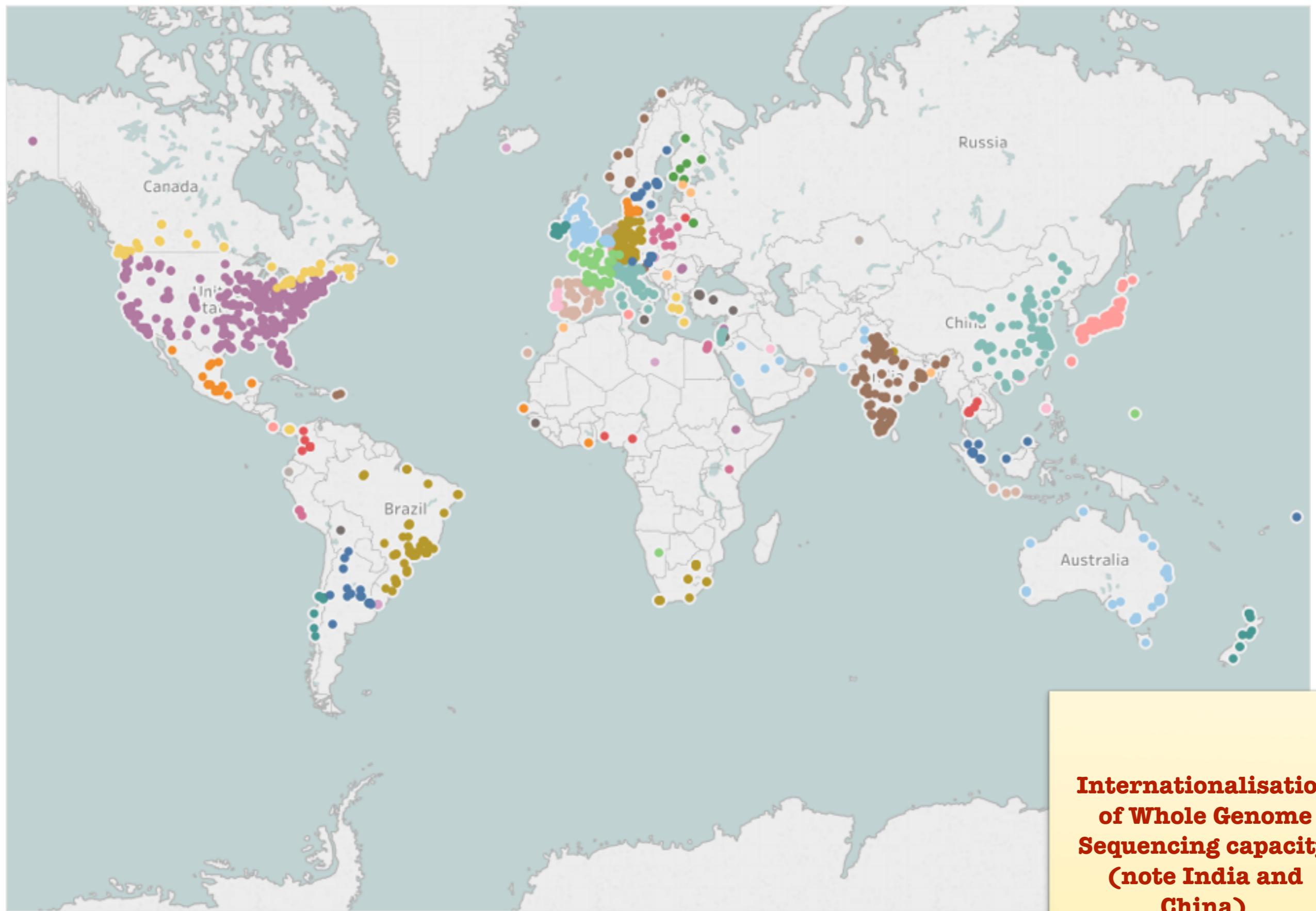


Project Status

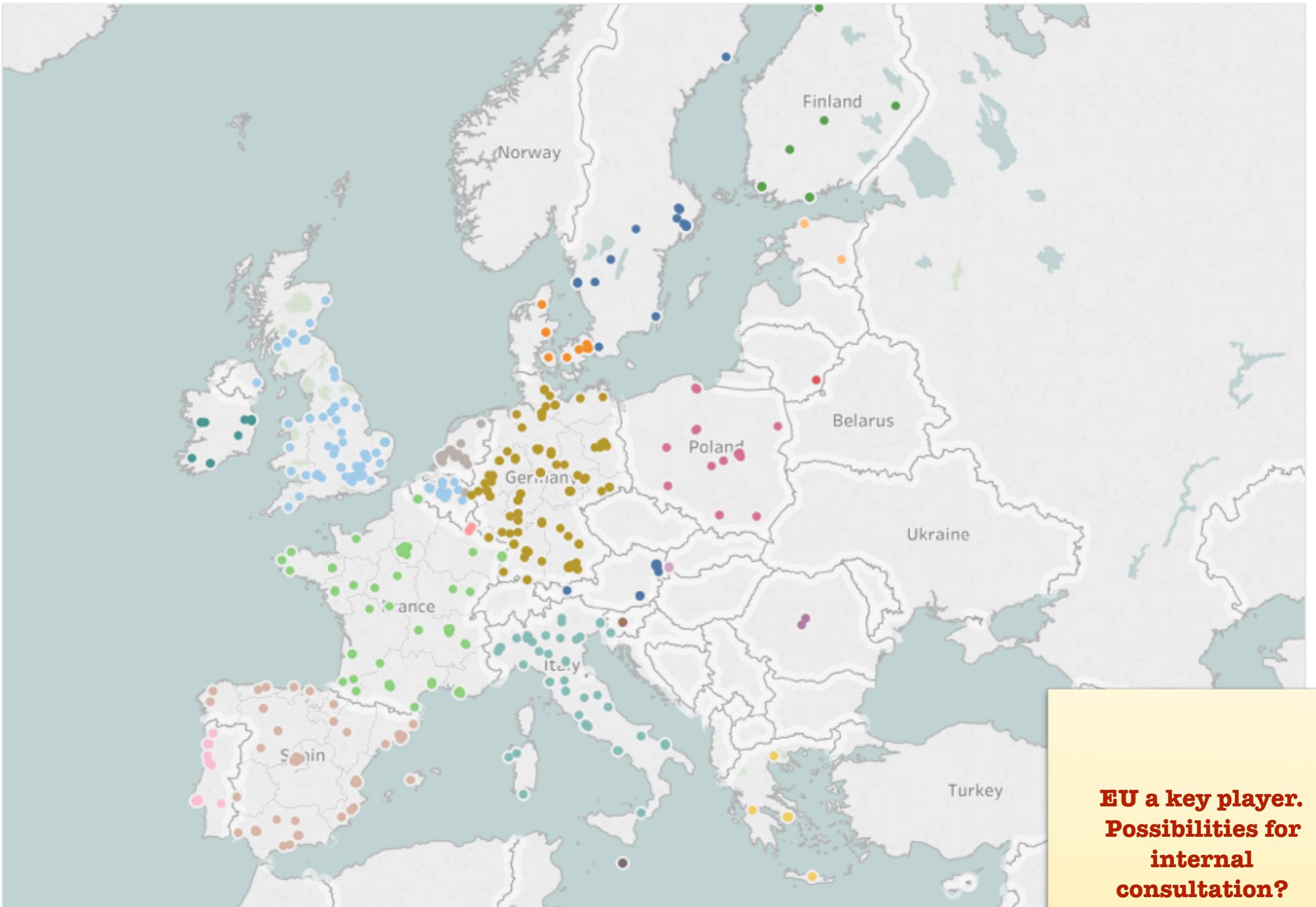


Genome Sequencing Centres (raw)





**Internationalisation
of Whole Genome
Sequencing capacity
(note India and
China)**



Organisations with Whole Genome Projects in the EU

**EU a key player.
Possibilities for
internal
consultation?**

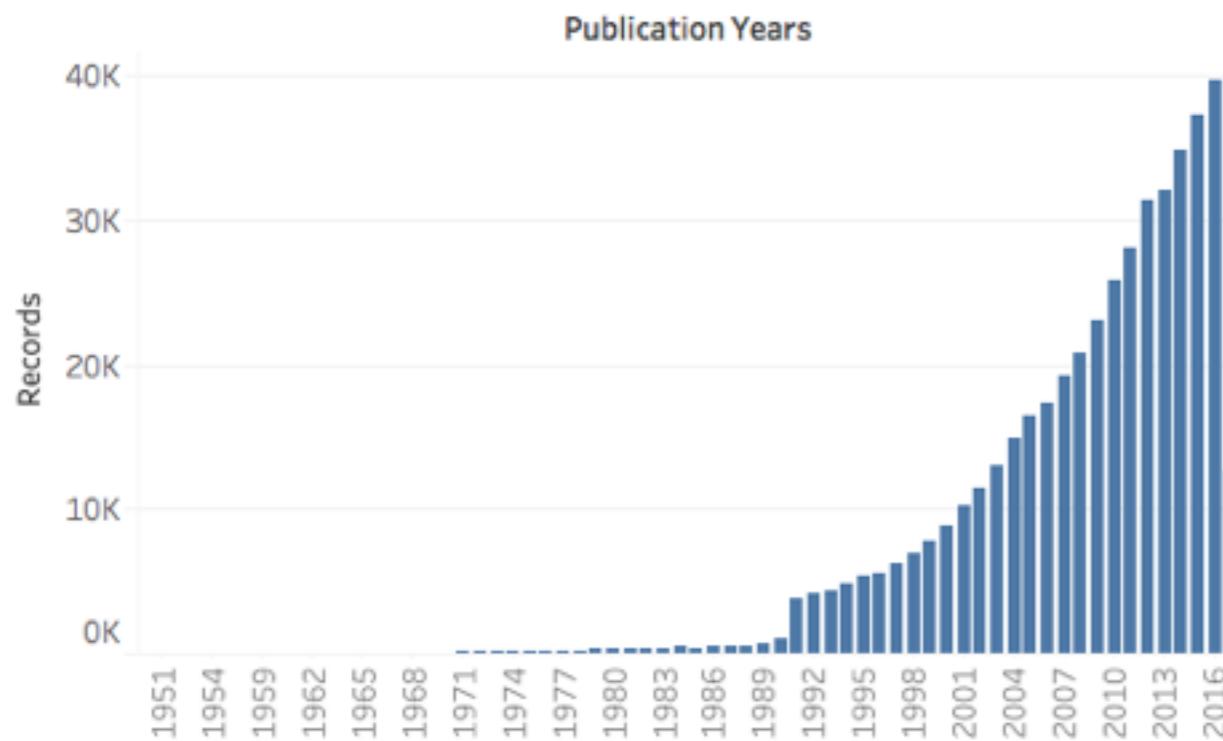
Scientific Publications in Genomics

Scientific outputs from DSI research can be illuminated using publication data for subjects such as genomics. This can also assist with identifying key areas of research and key actors.

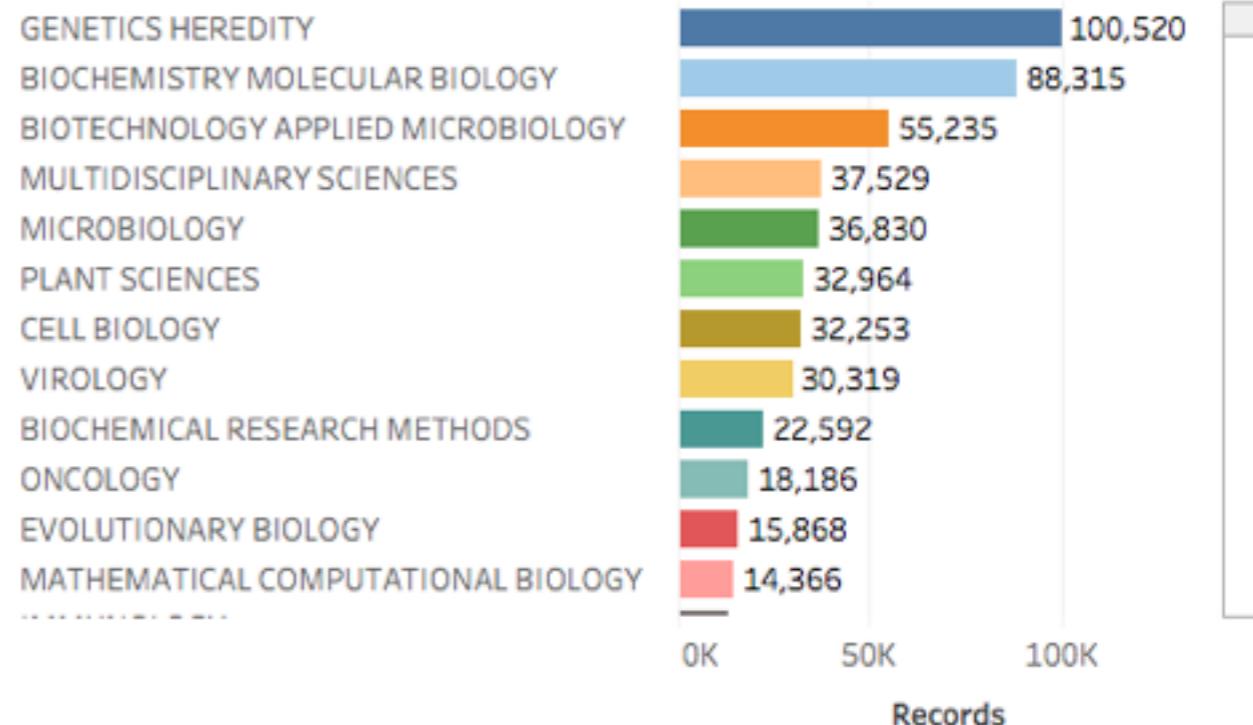
The Omics

- Functional Genomics (transcription, translation, protein - protein interactions)
- Structural Genomics (description of all proteins encoded by a genome)
- Epigenomics (factors influencing phenotypes)
- Metagenomics (sequencing environmental samples for taxonomy etc.)
- Synthetic Genomics (synthetic biology, engineering new genetic components and organisms from scratch)
- Conservation genomics (informing conservation decision making)
- Proteomics (understanding the protein complement of a cell or organism)
- Molecular Taxonomy, Cladistics and DNA barcoding
- ... yet more omics

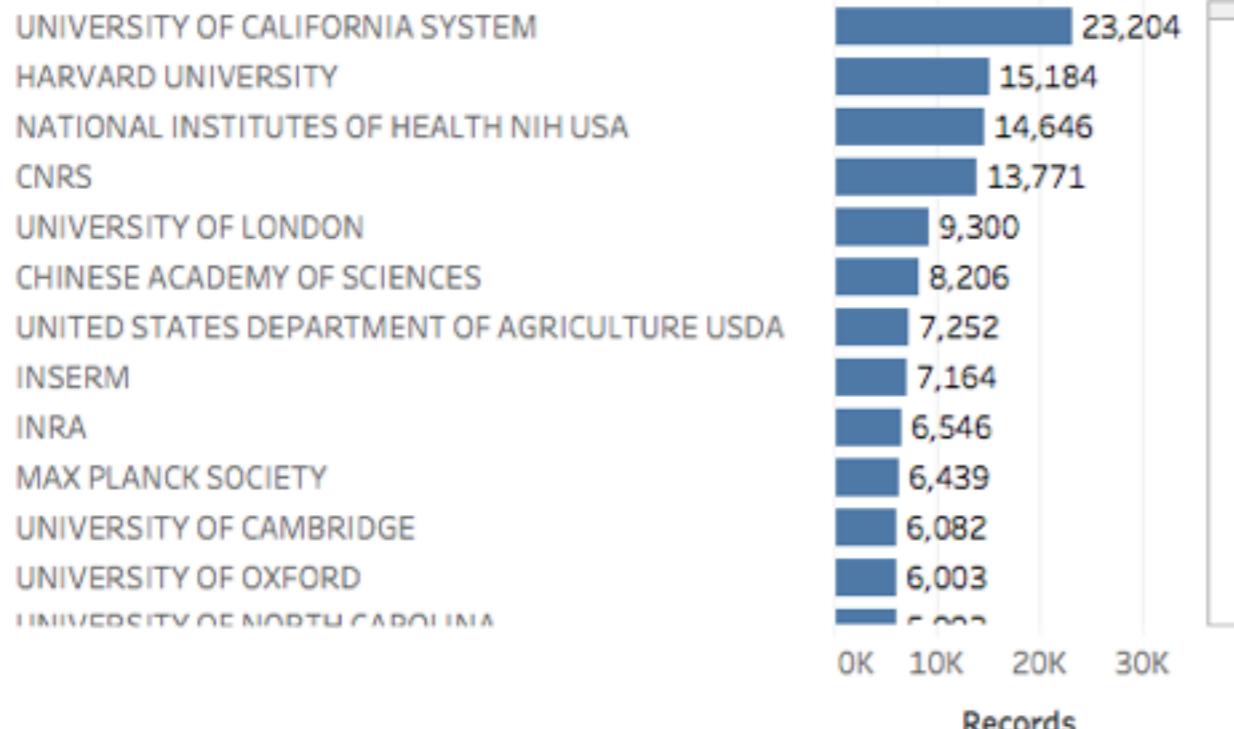
Genomics Scientific Publications



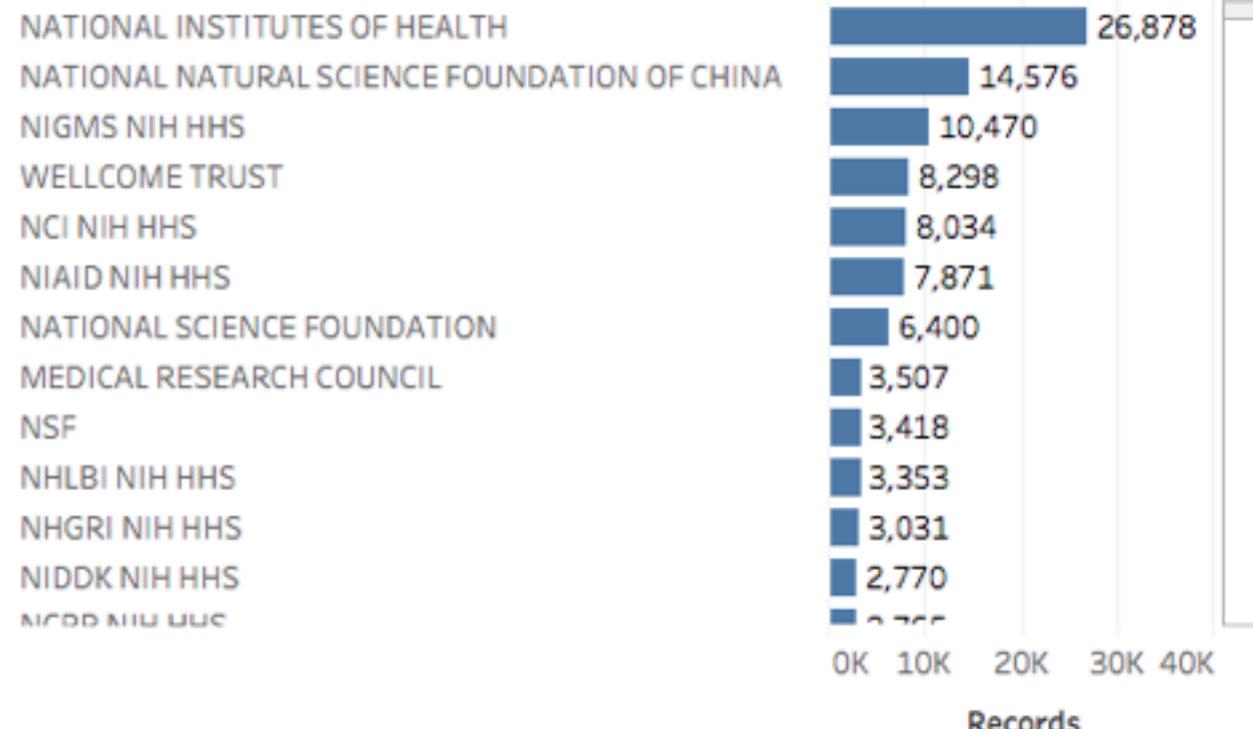
Genomics by Subject Area



Genomics Top Organisations (Raw)



Genomics Funding (Raw)



Source: Search of Web of Science Core Collection topic field for genome, genomes and genomics 29/05/2017. Organisation and Funding agency data has not been cleaned and is classified as raw. This will affect rankings.

Genomics Publications Rank Country



Genomics Publications



“...it is conceivable that technological innovation may one-day permit the *in situ* extraction of genetic material and transfer of data to electronic form without the necessity of the collection, taxonomic identification and storage of field samples.”

- Oldham 2004: UNEP/CBD/WG-ABS/3/INF/4 at 17

VolTRAX W List of open-source bioinform... New Tab

Secure https://nanoporetech.com/products/voltrax

Rapid, programmable, portable, disposable sample processor

VolTRAX VolTRAX Introduction Programme

About VolTRAX

Oxford Nanopore offers a range of options for converting your original biological sample to a form ready for application into a nanopore sensing device.

Oxford Nanopore has developed VolTRAX – a small device designed to perform library preparation automatically, so that a user can get a biological sample ready for analysis, hands free.

[VolTRAX Introduction Programme >](#)



Do you need any help? 

Portable Sample Preparation US\$2300

About MinION

MinION is the only portable, real time device for DNA and RNA sequencing.

Each consumable flow cell can now generate 5-10Gb of DNA sequence data. Ultra-long read lengths are possible (hundreds of kb) as you can choose your fragment length. The MinION streams data in real time so that analysis can be performed during the experiment and workflows are fully versatile.

The MinION weighs under 100g and plugs into a PC or laptop using a high speed USB 3.0 cable. No additional computing infrastructure is required. Not constrained to a laboratory environment, it has been used up a mountain, in a jungle, in the arctic and on the International Space Station.

The MinION is commercially available, simply by paying a starter pack fee of \$1,000. The MinION starter pack includes materials you need to run initial sequencing experiments, including a MinION device, flow cells, kits and membership of the Nanopore Community.



Real Time Portable DNA sequencing. US \$100 basic pack, US\$4999

MinION W List of open-source bioinform... New Tab

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PRODUCTS HOW IT WORKS APPLICATIONS GET STARTED PUBLICATIONS

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Do you need any help? 

Sequencing links to cloud based analytics

Intellectual Property Issues

- Issues around the implications of patent rights have been widely debated (and patent regulations have increasingly been restricted). However, it is important to bear in mind that in addition to patents DSIs as data involves
- Copyright (in sequences)
- Database Rights (in applicable jurisdictions)

Implications for ABS

- Developing countries are likely to incorporate articles into national legislation and ABS contracts on sequence data. This is logical but the question is the impact relative to the gain.
- The International Nucleotide Sequence Database Collaboration effectively asserts that DNA, RNA and amino acid sequence data belongs in the public domain (unrestricted use). That could be a good thing...but...
- Provider countries are likely to question the legitimacy of this assertion and may turn to countries that are not part of the INSDC that will meet requirements such as renewed PIC & MAT for the use of sequence data.
- If providers do introduce regulations on DSI (which looks fairly inevitable at present) the question would be how to operationalise that?

Examples: BN000065, histone

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[ENA](#) > [Submit & Update](#) > Species BARCODE checklist

Minimum information about species barcode nucleotide sequence

The Species BARCODE Data Standard is a biodiversity standard formulated by the Consortium for the Barcode of Life (CBOL) for reporting minimum information about species barcode nucleotide sequences. The CBOL specifies requirements on reporting sample provenance information and on sequence quality with the aim to create a reference library of barcode DNA sequences integrated with related biodiversity information, such as taxonomy, specimen vouchers or geo-reference. Ultimately, DNA barcoding shall serve as a global standard for species identification.

The International Barcode of Life project (iBOL) develops a DNA barcode reference library that will serve as DNA-based identification system for multi-cellular life.

The Barcode of Life Data Systems (BOLD) is the central informatics platform for DNA barcoding providing acquisition, storage, analysis and publication of DNA barcode records.

A suitable species barcode marker has to meet several criteria. Ideally, the barcode marker (1) can be easily amplified in one read following a standardised protocol, (2) is on both sides flanked by a highly conserved region for reliable primers annealing, (3) is capable of organism identification on a species level.

Currently, the CBOL approves as effective barcodes the following loci:

- for metazoa, the cytochrome c oxidase 1 (cox1) gene region
- for land plants, a two-locus barcode, the ribulose-bisphosphate carboxylase (rbcL) and maturaseK (matK) gene regions (with recommendation to collect also non-coding regions, such as the chloroplast trnH-psbA spacer region)
- for fungi, the ribosomal internal transcribed spacer (ITS) region

INSDC records that meet the criteria of Species BARCODE Data Standard have the keyword 'BARCODE'.

The MIMARKS includes the Species BARCODE Data Standard, which means that a MIMARKS-compliant dataset is also Species BARCODE compliant.

Species BARCODE data submission

The Species BARCODE reporting requirements are divided into mandatory (available [here](#)), highly recommended (available [here](#)) and optional (available [here](#)) irrespective of the sequenced marker locus.

Submit & Update

- ▶ Data formats
- Uploading data files
- ▶ Reads
- ▶ Sequences
- ▶ Genome assemblies
- Taxonomy
- ▶ Sample checklists
- Environmental
- Epigenomic
- Species BARCODE
- Metadata model
- Register submission account
- ▶ Programmatic XML submissions

Popular

- Submit and update
- Sequence submissions
- Genome assembly submissions
- Submitting environmental sequences
- Citing ENA data
- Rest URLs for data retrieval
- Rest URLs to search ENA

Latest ENA news

27 Apr 2017: [New ENA discover](#)

ENA has launched a new API to search across all data types: <https://www.ebi.ac.uk/ena/>

ENA has a range of submission forms with requirements.

This is for DNA barcodes

sequences now available

Mandatory Species BARCODE checklist

Field	Description	Example
Organism name;	Formal taxonomic name of this metozoan organism or informal name if unpublished/unidentified.	Arabidopsis thaliana
Bio-repository data	Reference to physical specimen from which the sequence was obtained (e.g. curated museum collection, living specimen), can be structured or unstructured.	structured YMUK:12345 unstructured ABCD-12345
Country	Political name of country or ocean in which a sequenced sample or isolate was collected.	France, Mediterranean Sea
Translation table	Mitochondrial translation table for this organism. Choose between vertebrate (table 2) and invertebrate (table 5) codes.	2
Codon Start (required to determine reading frame)	The codon start for the reading frame which should be translated is the coordinate of the base for the first complete codon.	3
Forward Primer Name	Name of the forward direction PCR primer.	ArthFW1
Forward Primer Sequence	Sequences should be given in the IUPAC degenerate-base alphabet, except for the modified bases; those must be included within angle brackets.	GACATTGKG<I>T
Reverse Primer Name	Name of the reverse direction PCR primer.	ArthRV1
Reverse Primer Sequence	Sequences should be given in the IUPAC degenerate-base alphabet, except for the modified bases; those must be included within angle brackets.	CATGRTTAGAC

Highly recommended Species BARCODE checklist

Field	Description	Example
Latitude/Longitude	Geographical coordinates of the location where the specimen was collected, in decimal degrees (to 2 places).	47.94, -12.45
Identified by	The person that identified the organism/sample.	John White
Collector	Name of the person that originally collected the sample/organism	John White
Collection Date	Date of collection of the original sample/organism	12-Apr-2013

Mandatory disclosure of Country and additional voluntary options available

**Previous work
explored the use of
creative commons
style licensing
models for ABS**



Convention on Biological Diversity

Distr.
GENERAL

UNEP/CBD/WG-ABS/8/INF/3
30 July 2009

ENGLISH ONLY

AD HOC OPEN-ENDED WORKING GROUP
ON ACCESS AND BENEFIT-SHARING

Eighth meeting
Montreal, 9-15 November 2009
Item 3 of the provisional agenda*

THE ROLE OF COMMONS/OPEN SOURCE LICENCES IN THE INTERNATIONAL REGIME ON ACCESS TO GENETIC RESOURCES AND BENEFIT-SHARING

Note by the Executive Secretary

1. The Executive Secretary is pleased to circulate herewith, for the information of participants in the eighth meeting of the Ad Hoc Open-ended Working Group on Access and Benefit-sharing, a discussion paper on the role of commons/open source licences in the international regime on access to genetic resources and benefit-sharing, ESRC Centre for Economic and Social Aspects of Genomics (Cesagen), University of Lancaster and the Peruvian Society for Environmental Law (SPDA). This paper is referred to in the first paragraph of the suggestions on operational text submitted by Cesagen, which is also available at <https://www.cbd.int/abs/submissions/abswg-08-cesagen-en.pdf>.
2. The paper is being circulated in the form and language in which it was received by the Secretariat.

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- ▶ Licensing types
- ▶ Public domain
- ▶ Places to share
- ▶ For developers

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The regulatory challenge

- The use of Creative Commons style licences for Sequence Data would run straight into the no restriction requirements of the INSDC. My understanding is that this policy originated from the efforts by companies to use restrictive licensing on genome sequence data. Science would have been the loser.
- However. providers will be confronted with the challenge of how to protect their interest and at the same time promote scientific research and cooperations and innovation for genetic resources. A pure public domain argument is unlikely to gain traction...

Conclusions

- If provider countries go down the route of introducing requirements on digital sequence data into their legislation and MAT, this is likely to have significant consequences for scientific research (notably taxonomy) and over the longer term for innovation.
- At the same time a pure public domain argument is unlikely (in my view) to succeed because it will not address provider concerns.
- A middle ground may be possible but it would need to be simple (in terms of options) in order to address scale measured in terms of billions and trillions of bases from organisms distributed around the world. Models for this already exist but would need to be adjusted for ABS needs.
- The alternative may possibly be fragmentation of DSI into multiple silos depending on the willingness of database providers to accept provider country conditions.

The global public goods dimension

- Genomes (notably the human genome) have been treated as a global public good. In the context of the Nagoya Protocol it is important to emphasise the opportunities that may exist for international cooperation in taxonomy, conservation genomics, to address human health issues (e.g. neglected diseases) or identify strategies for adaptation to climate change that are enabled by genomics and DSI.
- The investments and international collaboration that exists in genome sequencing and genomic research are valuable in themselves in terms of knowledge and technology transfer and capacity-building. Above all perhaps they have value in advancing knowledge and understanding of biodiversity and genetic resources. More could be made to highlight this at the expense of the perils of the pursuit of hyperownership in the context of the promises of biotechnology.