GigaByte – A New Workflow for Rapid Dissemination of Datasets & Tools -Bringing Papers to Life

eResearch 2021, Wellington, NZ

Nicole Nogoy Executive Editor, ORCID: 0000-0002-5192-9835 (GIGA) by te Publishing at the Speed of Research

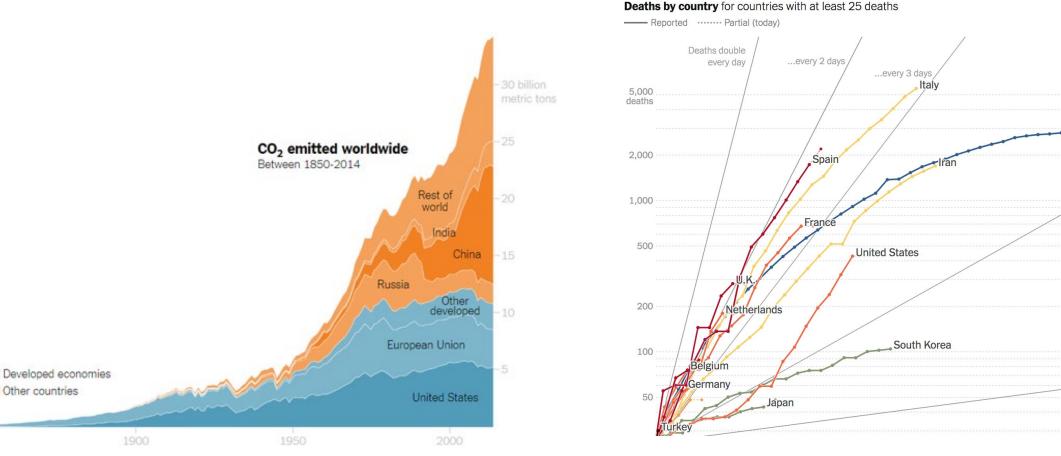




Urgent Challenges Research Needs to Address

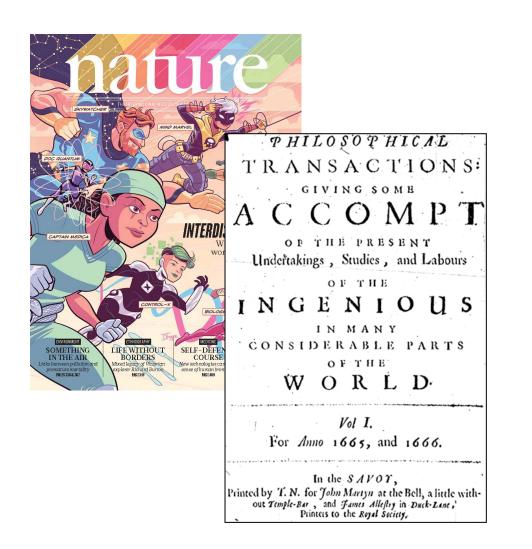
Climate change

Disease pandemics (e.g., COVID19)



https://www.nytimes.com/2019/02/28/learning/teach-about-climate-change-with-these-24-new-york-times-graphs.html https://www.nytimes.com/interactive/2020/03/21/upshot/coronavirus-deaths-by-country.html

Urgent Challenges: Publishing to the Rescue?



- Need to disseminate & communicate information openly to global community
- Expensive. Information is held back by paywalls, APCs, and other barriers
- Need to share this quickly and in a trusted form (peer reviewed)
- Laborious, archaic tech, and untransparent processes
- Research data, software and underlying methods and results need to be shared for scrutiny and re-use
- Hard work and little incentive to share
- Needs to be understandable by policy makers, public
- Barriers of language, jargon, and lack of interaction

$(GIGA)^n$ Attempt to Address This: *GigaScience* + GigaDB (2012)



GigaScience, 9, 2020, 1–11 doi: 10.1093/gigascience/giaa100 Data Note





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Construction of a chromosome-scale long-read

reference genome assembly for potato

Gina M. Pham ^{(©1,1}, John P. Hamilton ^{(©1,1}, Joshua C. Wood ^(©1), Joseph T. Burke ^(©1), Hainan Zhao¹, Brieanne Vaillancourt ^(©1), Shujun Ou ^(©2), Jiming Jiang ^{(©1,3,4} and C. Robin Buell ^(©1,4,5,*)

¹Department of Plant Biology, Michigan State University, 612 Wilson Road, East Lansing, MI 48824, USA; ²Department of Ecology, Evolution, and Organismal Biology, Iowa State University, 2200 Osborne Dr, Arnes, IA S0011, USA; ³Department of Horticulture, Michigan State University, 1066 Bogue St, East Lansing, MI 48824, USA; ⁴MSU AgBioResearch, Michigan State University, 446 W. Circle Drive, East Lansing, MI 48824, USA and ⁵Plant Resilience Institute, Michigan State University, 4109 Milson Road, East Lansing, MI 48824, USA

*Correspondence address. C. Rohin Buell, Department of Plant Biology, Michigan State University, 612 Wilson Road, East Lansing, MI 48824, USA. E-mail: bael@msu.edu@http://orcid.org/0000-0002-6727-4677 These authors contributed equally to this work.

Abstract

Background: Worldwide, the cultivated potato, Solanum tuberosum L., is the No. 1 vegetable crop and a critical food security crop. The genome sequence of DM1-3 516 R44, a doubled monoploid clone of S. tuberosum Group Phureja, was published in 2011 using a whole-genome shotgun sequencing approach with short-read sequence data. Current advanced sequencing technologies now permit generation of near-complete, high-quality chromosome-scale genome assemblies at minimal cost. Findings: Here, we present an updated version of the DMI-3 516 R44 genome sequence (v6.1) using Oxford Nanopore Technologies long reads coupled with proximity-by-ligation scaffolding (Hi-C), yielding a chromosome-scale assembly. The new (v6.1) assembly represents 741.6 Mb of sequence (87.8%) of the estimated 844 Mb genome, of which 741.5 Mb is non-gapped with 731.2 Mb anchored to the 12 chromosomes. Use of Oxford Nanopore Technologies full-length complementary DNA sequencing enabled annotation of 32,917 high-confidence protein-coding genes encoding 44,851 gene models that had a significantly improved representation of conserved orthologs compared with the previous annotation. The new assembly has improved contiguity with a 595-fold increase in N50 contig size, 99% reduction in the number of contigs, a 44-fold increase in N50 scaffold size, and an LTR Assembly Index score of 13.56, placing it in the category of reference genome quality. The improved assembly also permitted annotation of the centromeres via alignment to sequencing reads derived from CENH3 nucleosomes. Conclusions: Access to advanced sequencing technologies and improved software permitted generation of a high-quality, long-read, chromosome-scale assembly and improved annotation dataset for the reference genotype of potato that will facilitate research aimed at improving agronomic traits and understanding genome evolution.

Keywords: long-read; chromosome-scale; reference genome; potato

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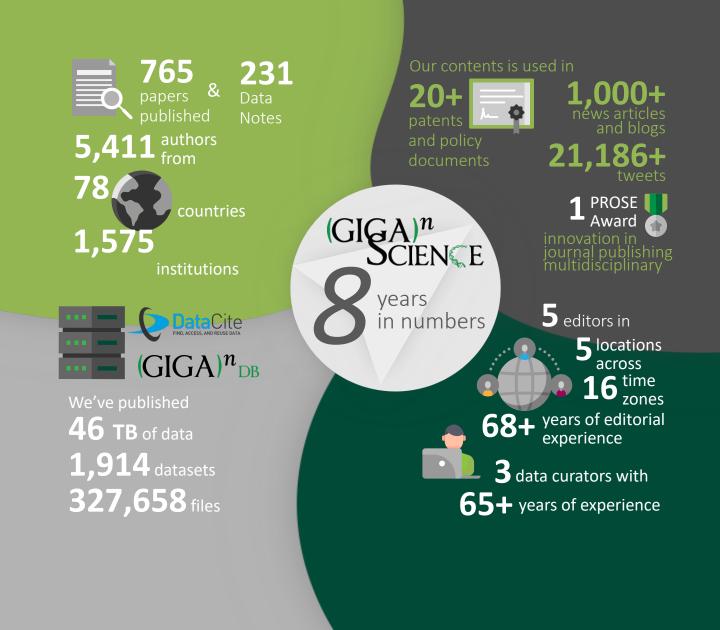




http://gigadb.org/

 $(GIGA)^{n}_{DB}$

http://gigasciencejournal.com



Lessons Learned

- Technology is really the bottleneck
- Traditional publishing process is too slow & expensive
- Much too focused on narrative and static "version of record"



Guillaume Devailly @G_Devailly

When you discover that **@GigaScience** exists, but GigaNature doesn't.

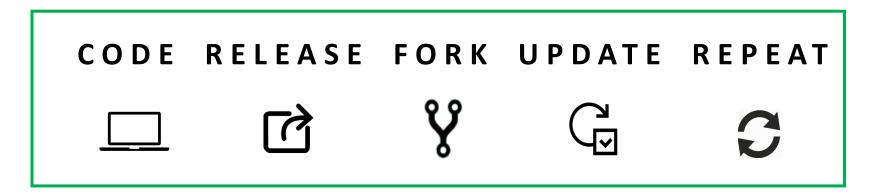
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1:33 PM · Jul 22, 2020 · Twitter Web App

A New Approach

Follow the Software Paradigm?



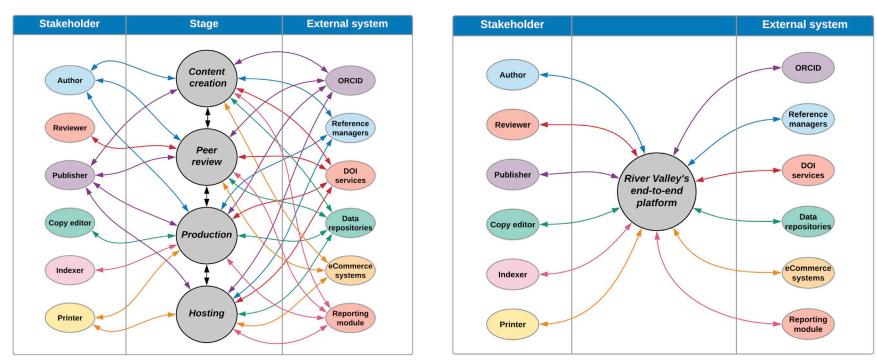
Deconstruct the "Version of Record"?



Move To New XML End-to-End Pipeline



Custom end-to-end workflow makes integrations simpler with one integration point



(GIGA) byte A New Approach With New Tech

Main advantage of workflow is XML from start to end

Several modules acting as one platform: no import/export of files, so fast and accurate

Cutting out production allows huge time & cost saving (currently 4-8 hours NOT 2 months+)

Any number of versions can be published instantly, including typographic quality PDF or updates/forks –minimal effort

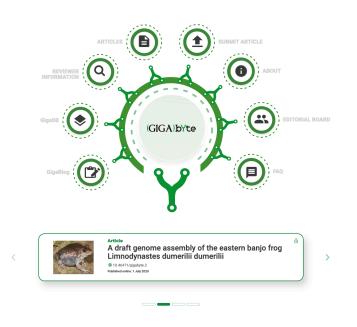
Allows instantaneous switch of views - push of a button

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User flag	Select some options
Must change password at first logon	0
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DATA RELEASE (GIGA)byte **Chromosome-level genome** assembly of a benthic associated Syngnathiformes species: the common dragonet, Callionymus lyra Sven Winter^{1,2,*}, Stefan Prost^{3,4}, Jordi de Raad^{1,2,3}, Raphael T. F. Coimbra^{1,2}, Magnus Wolf^{1,2}, Marcel Nebenführ¹, Annika Held¹, Melina Kurzawe¹, Ramona Papapostolou¹, Jade Tessien¹, Julian Bludau¹, Andreas Kelch¹ Sarah Gronefeld¹, Yannis Schöneberg¹, Christian Zeitz¹, Konstantin Zapf¹, David Prochotta¹, Maximilian Murphy¹, Monica M. Sheffer⁵, Moritz Sonnewald⁶, Maria A. Nilsson^{2,3} and Axel Janke^{1,2,3} 1 Institute for Ecology, Evolution and Diversity, Goethe University, Frankfurt am Main, Germany 2 Senckenberg Biodiversity and Climate Research Centre, Frankfurt am Main, Germany 3 LOEWE-Centre for Translational Biodiversity Genomics, Frankfurt am Main, Germany 4 South African National Biodiversity Institute, National Zoological Garden, Pretoria, South Africa 5 Zoological Institute and Museum, University of Greifswald, Greifswald, Germany 6 Senckenberg Research Institute, Department of Marine Zoology, Section Ichthyology, Frankfurt am Main, Germany ABSTRACT Background: The common dragonet, Callionymus lyra, is one of three Callionymus species inhabiting the North Sea. All three species show strong sexual dimorphism. The males show strong morphological differentiation, e.g., species-specific colouration and size relations, while the females of different species have few distinguishing characters. Callionymus belongs to the 'benthic associated clade' of the order Syngnathiformes. The 'benthic associated clade' so far is not represented by genome data and serves as an important outgroup to understand the morphological transformation in 'long-snouted' syngnatiformes such as seahorses and pipefishes Findings: Here, we present the chromosome-level genome assembly of C. lyra. We applied Oxford Submitted: 03 September 2020 Nanopore Technologies' long-read sequencing, short-read DNBseq, and proximity-ligation-based Revised: 09 October 2020 Accepted 19 October 2020 scaffolding to generate a high-quality genome assembly. The resulting assembly has a contig Published 20 October 2020 N50 of 2.2 Mbp and a scaffold N50 of 26.7 Mbp. The total assembly length is 568.7 Mbp. of which over 538 Mbp were scaffolded into 19 chromosome-length scaffolds. The identification of 94.5% Corresponding author E-mail complete BUSCO genes indicates high assembly completeness. Additionally, we sequenced and sven.winter@senckenberg.de assembled a multi-tissue transcriptome with a total length of 255.5 Mbp that was used to aid the Published by GigaScience Press. annotation of the genome assembly. The annotation resulted in 19.849 annotated transcripts and Preprint submitted at https identified a repeat content of 27.7%. //doi.org/10.1101/2020.09.08.287078 Conclusions: The chromosome-level assembly of C. lyra provides a high-quality reference This is an Open Access article genome for future population genomic, phylogenomic, and phylogeographic analyses. distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org licenses/by/4.0/), which permits unrestricted reuse, distribution, and Subjects Genetics and Genomics, Evolutionary Biology, Marine Biology reproduction in any medium

provided the original work is properly cited. *Gigabyte*, 2020, 1–10

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