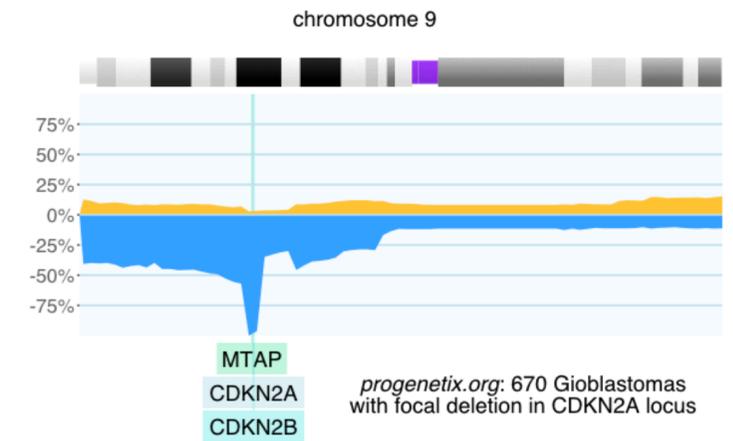


ELIXIR hCNV

Updates and future plans

Michael Baudis | ELIXIR Human Data Day | 2021-06-11



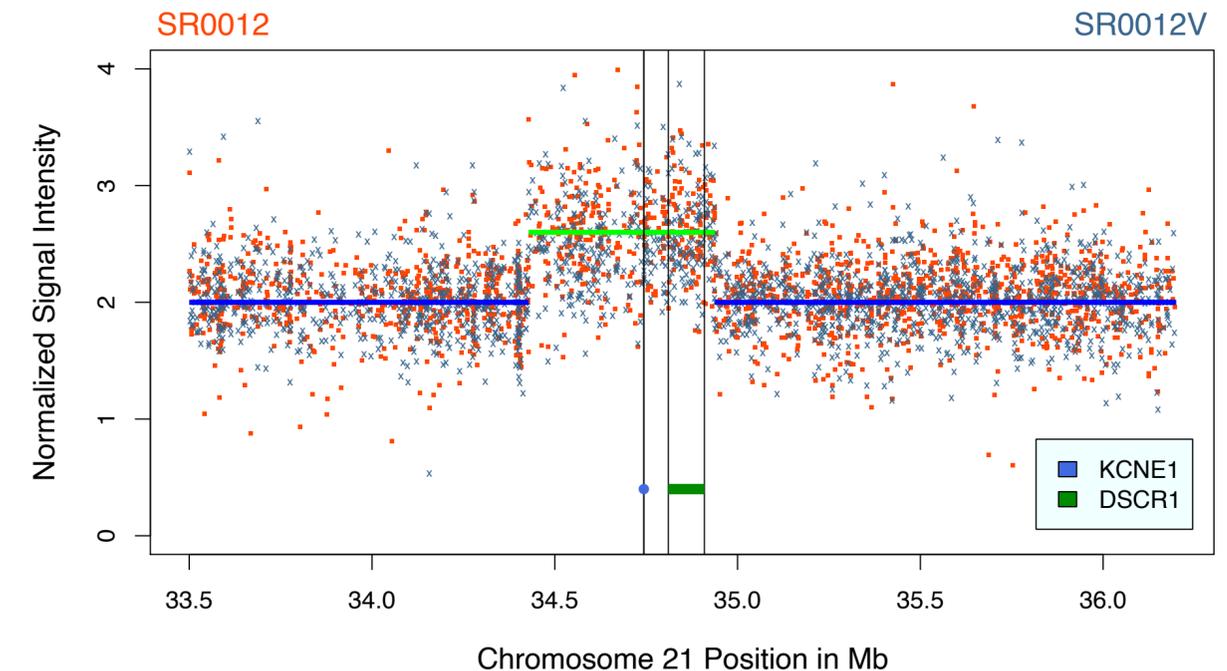
Why hCNV Community?

Structural Genome Variation Data :: Resources and Technologies

- structural genome variations are a major contributor to genetic diseases and cancer
- knowledge about and standards for copy number variations / aberrations (CNV/CNA) has not been in step with NGS & GWAS driven SNV/SNP assessment

Mission statement

Despite the fact that **Copy Number Variations** are the **most prevalent genetic mutation type**, identifying and interpreting them is still a major challenge. The ELIXIR human Copy Number Variation (hCNV) Community aims to implement processes to make the **detection**, **annotation** and **interpretation** of these variations easier



CNV with unknown clinical impact in a case of Silver-Russel Syndrome

Local Affymetrix Genotyping 6 signal distribution pattern and segmentation result in patient SR12 (SR0012, orange data) and his father (SR0012V, steelblue data). In both samples a duplication in the DSCR can be observed, affecting the whole KCNE1 and DSCR1/RCAN coding regions. In contrast, DYRK1A lays ~2.5 Mb distal of the duplication. Only the genes discussed in this article are shown.

RESEARCH ARTICLE

AMERICAN JOURNAL OF
medical genetics PART A

Identification of a 21q22 Duplication in a Silver–Russell Syndrome Patient Further Narrows Down the Down Syndrome Critical Region

Thomas Eggermann,^{1*} Nadine Schönherr,¹ Sabrina Spengler,¹ Susanne Jäger,¹ Bernd Denecke,² Gerhard Binder,³ and Michael Baudis⁴

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Received 26 June 2009; Accepted 6 November 2009

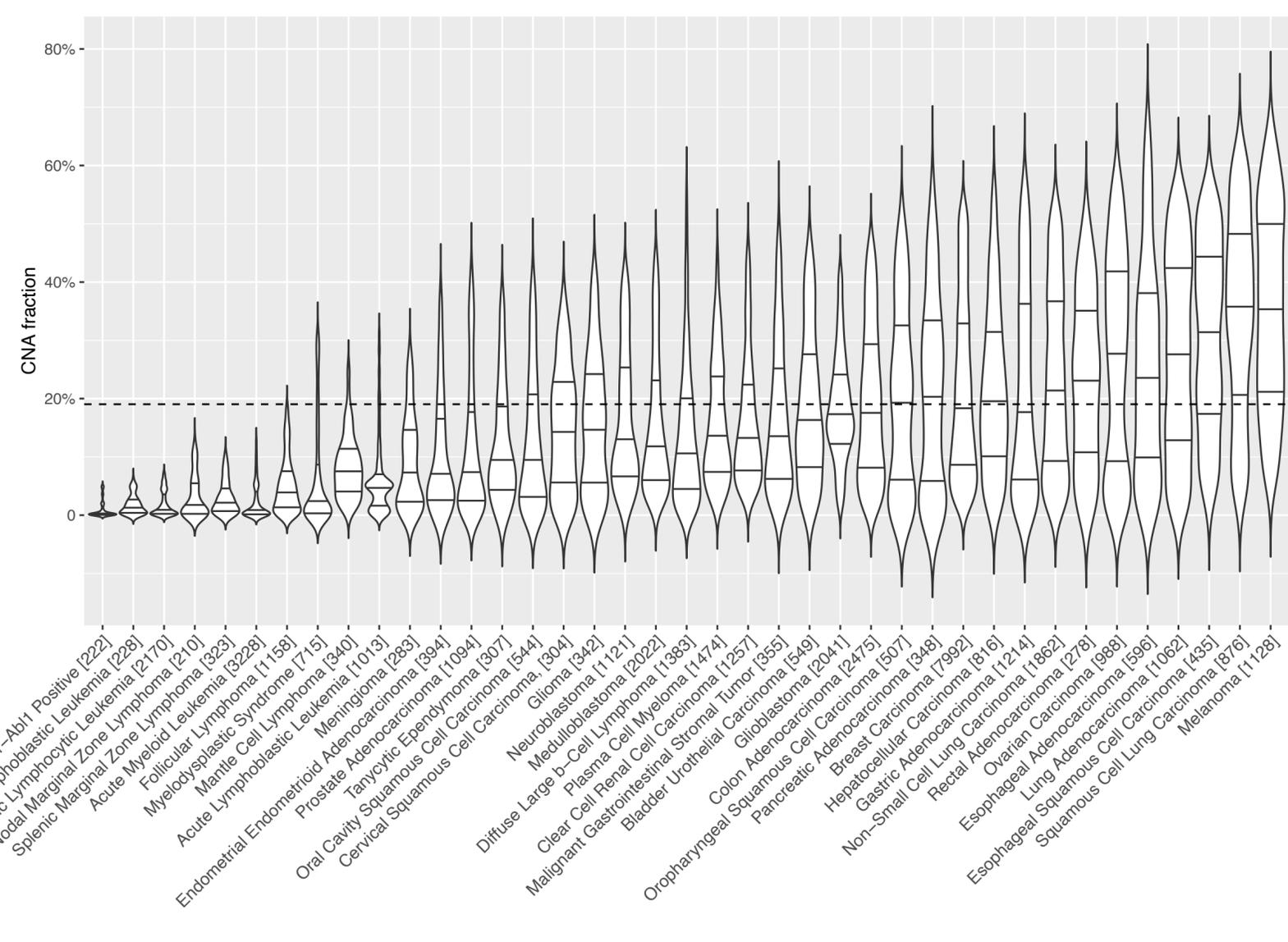
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Genome CNV coverage in Cancer Classes

- 43654 out of 93640 CNV profiles; filtered for entities w/ >200 samples
- Single-sample CNV profiles were assessed for the fraction of the genome showing CNVs (relative gains, losses)
- range of medians 0.001 (CML) - 0.358 (malignant melanomas) knowledge about and standards for copy number variations / aberrations (CNV/CNA) has not been in step with NGS & GWAS driven SNV/SNP assessment



ELIXIR hCNV Community

Structural Genome Variation Data :: Resources and Technologies

- First meeting of group in 2018
- ELIXIR Human Copy Number Variation (hCNV) approved in 2019
- initial implementation study (2019-2021) for community set-up, gap analysis and exploration of technical deliverable

Mission statement

Despite the fact that **Copy Number Variations** are the **most prevalent genetic mutation type**, identifying and interpreting them is still a major challenge. The ELIXIR human Copy Number Variation (hCNV) Community aims to implement processes to make the **detection**, **annotation** and **interpretation** of these variations easier

Purpose

The human CNV community (h-CNV) has been officially created in December 2018. It aims to address the major challenge of NGS data interpretation in the era of whole genome sequencing for the most frequent mutation type: Copy Number Variation. Seven topics have been identified during the kick-off meeting and further refined with all h-CNV partners. This ultimately led to the proposal described in this implementation study.

Node	Name of PI
ELIXIR-FR	Christophe Bérout, David Salgado, Marc Hanauer, Victoria Dominguez
ELIXIR-CH	Michael Baudis
ELIXIR-DE	Jan Korbel
EMBL-EBI	Thomas Keane, Fiona Cunningham
ELIXIR-ES	Joaquin Dopazo, Alfonso Valencia, Salvador Capella, Sergi Beltran, Steven Laurie, Gemma Bullich, Laura I. Furlong, Janet Piñero
ELIXIR Hub	John Hancock, Gary Saunders, Kathi Lauer, Leyla Garcia
ELIXIR-NL	Bauke Ylstra, Daoud Sie, Leon Mei, Morris Swertz (UMCG), Lennart Johansson
ELIXIR-NO	Eivind Hovig, Pubudu Samarakoon
ELIXIR-HU	Attila Gyenesei, Katalin Monostory
ELIXIR-SI	Brane Leskošek, Polonca Ferk, Marko Vidak
ELIXIR-UK	Krzysztof Poterlowicz
Delivery	Starting from June 2019 for a period of 24 months.



Christophe Bérout
(ELIXIR France)



David Salgado
(ELIXIR France)



Gary Saunders
(Human Data Coordinator,
ELIXIR Hub)



Michael Baudis
(ELIXIR Switzerland)



hCNV Implementation Study 2019-2021

Setting the Scope | Solidifying the Community | First Deliveries

- challenge participants and define the wider landscape as well as future directions
- set of 7 work packages
 - ➔ landscape analysis
 - ➔ technical products
 - ➔ resource improvement
 - ➔ community building & outreach
- regular meetings, website, hackathons...
- ▶ WP1 - Optimal CNV detection pipelines for research and diagnostics
- ▶ WP2 - Definition of reference datasets
- ▶ WP3 - Improvement of community formats for CNV exchange
- ▶ WP4 - Enabling CNV data discovery in diagnostic and phenotypic context
- ▶ WP5 - Creation of innovative tools
- ▶ WP6 - FAIRification of h-CNV databases and datasets
- ▶ WP7 - Dissemination



hCNV Implementation Study 2019-2021

Setting the Scope | Solidifying the Community | First Deliveries

- highly ambitious goals, beyond available support
 - especially reference / benchmarking dataset generation and pipeline development
- emerging interactions and collaborations with ELIXIR platforms & communities and beyond
 - Galaxy
 - GA4GH / ELIXIR Beacon project
- ▶ WP1 - Optimal CNV detection pipelines for research and diagnostics
- ▶ WP2 - Definition of reference datasets
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hCNV Implementation Study 2019-2021

Some Achievements and Deliveries

- Benchmarking tools and OpenEbench TransBioNet testing event
- demonstration of CNV detection tools in clinical (cancer) setting
- amending *bio.tools* for extensive list of CNV related analysis tools
 - ➔ <https://bio.tools/t?domain=elixir-hcnv>
- updating / registering shared hCNV resources at fairsharing.org
- consensus collection of perceived requirements for efficient and effective CNV file and data exchange formats



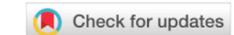
hCNV Implementation Study 2019-2021

Some Achievements and Deliveries

- HGVS satellite meeting – Human CNV – June 14th 2019 – Göteborg Sweden
- hCNV community workshop ELIXIR All-Hands Lisbon – June 2019
- survey of data annotation formats, including comments on VCF development
- start FAIRification of CNV national / reference databases (BANCCO, Progenetix)
- Community white paper published
- Biohackathon Paris 2019
- in 2021 start of shared meetings of subgroup with Beacon variants scout team

F1000Research

F1000Research 2020, 9(ELIXIR):1229 Last updated: 01 JUN 2021



OPINION ARTICLE

The ELIXIR Human Copy Number Variations Community: building bioinformatics infrastructure for research [version 1; peer review: 1 approved]

David Salgado ¹, Irina M. Armean², Michael Baudis ³, Sergi Beltran^{4,5}, Salvador Capella-Gutierrez ^{6,7}, Denise Carvalho-Silva ^{2,8}, Victoria Dominguez Del Angel ⁹, Joaquin Dopazo ¹⁰, Laura I. Furlong ¹¹, Bo Gao ³, Leyla Garcia ^{2,12,13}, Dietlind Gerloff¹⁴, Ivo Gut^{4,5}, Attila Gyenesi¹⁵, Nina Habermann¹⁶, John M. Hancock ¹³, Marc Hanauer¹⁷, Eivind Hovig ^{18,19}, Lennart F. Johansson²⁰, Thomas Keane², Jan Korbel¹⁶, Katharina B. Lauer ¹³, Steve Laurie⁴, Brane Leskošek²¹, David Lloyd ¹³, Tomas Marques-Bonet²², Hailiang Mei²³, Katalin Monostory²⁴, Janet Piñero ¹¹, Krzysztof Poterlowicz ²⁵, Ana Rath¹⁷, Pubudu Samarakoon²⁶, Ferran Sanz¹¹, Gary Saunders ¹³, Daoud Sie²⁷, Morris A. Swertz²⁰, Kirill Tsukanov ², Alfonso Valencia^{6,7,28}, Marko Vidak²¹, Cristina Yenyxe González², Bauke Ylstra²⁹, Christophe Bérout^{1,30}

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²⁰Department of Genetics, University of Groningen, University Medical Center Groningen, Groningen, The Netherlands

²¹Faculty of Medicine - ELIXIR Slovenia, University of Ljubljana, Ljubljana, Slovenia

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²⁴Institute of Enzymology, Research Centre for Natural Sciences, Budapest, Hungary

²⁵Centre for Skin Sciences, University of Bradford, Bradford, UK



hCNV Implementation Study 2019-2021

Some Achievements and Deliveries

- survey about genomic variation file formats and their use, suitability for representing CNV data
- part of the survey was focused specifically on VCF, a key GA4GH standard at the intersection of human and computer readable formats
- Results
 - ➔ BED-like formats are frequently used, but the better defined flavours are not optimal for CNVs and other SVs
 - ➔ JSON w/ schema has potential, but still misses finalized GA4GH schemas (VRS emerging) and suffers "readability" issues for non-bioinformatics customers
 - ➔ VCF was considered as a/the variant standard file format, but not "CNV-friendly" in v4.2 and in the existing tools for I/O handling of CNV data

ELIXIR hCNV 2019-21 Deliverable D3.2

Project Title:	First hCNV Community Implementation Study
Deliverable title:	Create a consensus collection of perceived requirements for efficient and effective CNV file and data exchange formats
WP No.	3
WP Title	Improvement of community formats for CNV exchange
Contractual delivery date:	30.11.2019
Actual delivery date:	12.12.2019
WP leads:	Thomas Keane
Partner(s) contributing to this deliverable:	EMBL-EBI

Report authors: Kirill Tsukanov¹, Sundararaman Venkataraman, Giselle Kerry, Thomas Keane (EMBL-EBI)

2. Results	3
2.1. Feedback overview	3
2.2. Terminology	4
2.3. Existing file formats	4
2.3.1. VCF (Variant Call Format)	4
2.3.2. BED and related tab-separated formats	5
2.3.3. JSON with a schema	5
2.3.4. Other formats	5
2.4. Opinion on CNV representation in VCF	6
2.5. Requirements for CNV formats of the future	6
2.6. Conclusions. Note about use cases	7
3. Impact	8

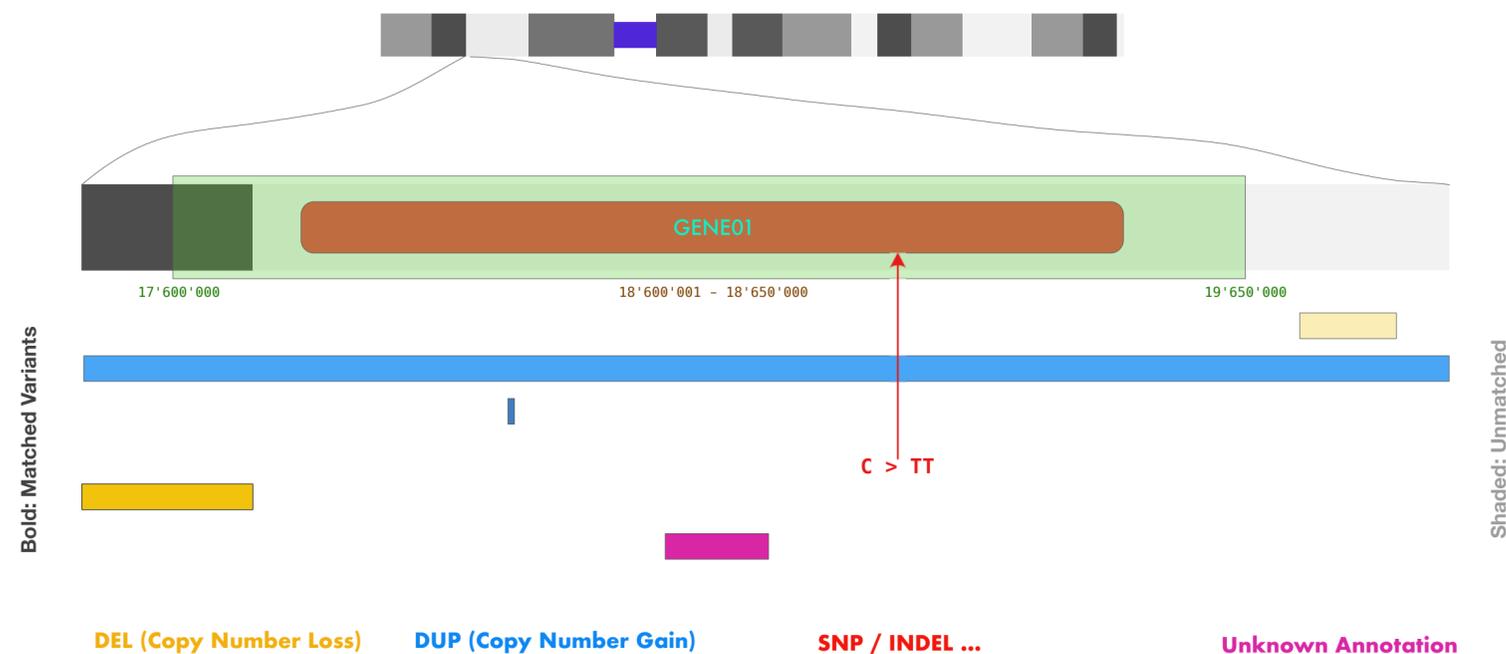


Beacon v2: Extended Variant Queries



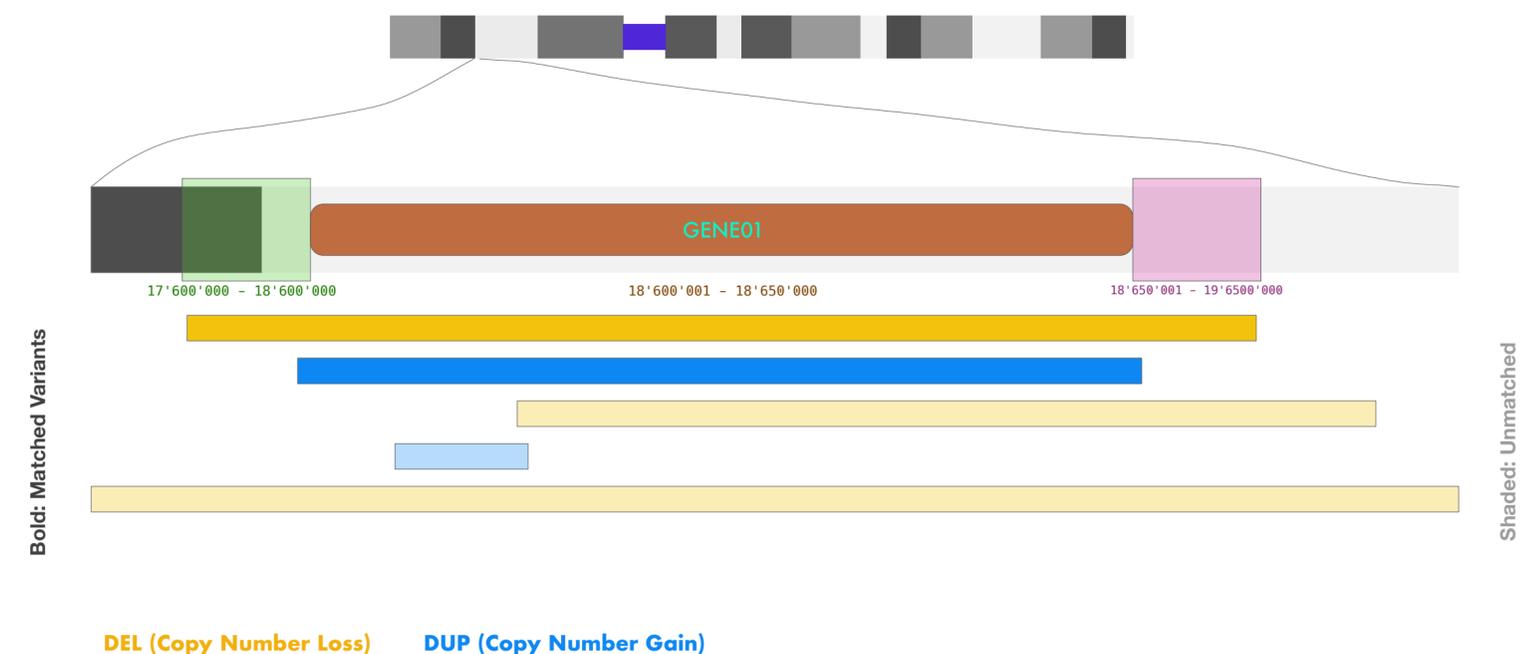
Range and Bracket queries enable positional wildcards and fuzziness

Genome Range Query (matching variants in a region)



- Genome Range Queries provide a way to "fish" for variants overlapping an indicated region, e.g. the CDR of a gene of interest
- Additional parameters (e.g. variant type, reference or alternate bases) limit the scope of the responses
- new Beacon v2 size parameters to limit structural variants (e.g. "focal" CNVs)

Genome Bracket Query (full match)



- Genome Bracket Queries allow to search for structural variants with start and end positions falling into defined sequence ranges
- allows to query any contiguous genomic variant (and in principle also can step in for range queries)
- typical use case is e.g. the query for variants such as duplications covering the whole CDR of a gene, while limiting the allowed start or end regions

2x hCNV Implementation Studies 2021-2023

Reference hCNV datasets, use-case workflows and benchmarking

The ELIXIR human CNV Community (hCNV) was created in December 2018. In two years contributions to the field have been numerous (ELIXIR IS, Rare Diseases, Federated Human Data, Beacons, GA4GH, EJP-RD and Beyond 1 Million Genomes - B1MG). The Community now aims to address the major challenge of NGS data interpretation in the era of whole genome sequencing: Copy Number Variation. During the first commissioned service offered as a starting grant, the Community has identified various gaps to proceed with CNV tools benchmarking and in particular for Exome and targeted sequencing, which are by far the most widely used technologies in diagnostic laboratories and in research. Within this implementation study we want to provide solutions and bioinformatic infrastructure solutions to fill identified gaps, and to make these biomedical reference materials available (i.e. via Open Science) to the various communities and platforms.

Interactions and utility to other projects

ELIXIR platforms:

Data, Tools, Interoperability, Training

ELIXIR Communities:

hCNV, Galaxy, Rare diseases, Federated Human Data

National and International projects:

EJP-RD, B1MG, EOSC-Life, EOSC-Pillar

Beacon and beyond – Implementation-driven standards and protocols for CNV discovery and data exchange

The initial 2019-2021 hCNV community implementation study employed a set of perceived needs to a) deliver first community standards and procedures; b) identify intersections with other ELIXIR communities and stakeholders in ELIXIR connected organizations, such as GA4GH; and c) to streamline priorities for relevant, achievable deliveries of hCNV community projects.

This proposal for an hCNV implementation study focuses on those potential high-value targets for data access and delivery, using reference resources and community stakeholder engagement to directly implement and test hCNV resources aligned with ELIXIR ecosystems.

The main target here will be the empowerment of the Beacon protocol, to act as standard for federated hCNV discovery and data delivery, in conjunction with additional GA4GH derived standards.

Intersecting ELIXIR Platforms, Communities and Projects:

- ELIXIR Galaxy Community
- ELIXIR AAI Infrastructure Service
- ELIXIR Compute Platform
- ELIXIR Training Platform
- ELIXIR FHD Community
- ELIXIR Health Data Focus Group
- ELIXIR Beacon Strategic Implementation Study
- ELIXIR Interoperability Platform

External Projects and Partners:

- EJP-RD
- GA4GH (Discovery, Genomic Knowledge Standards, Phenopackets)



hCNV Implementation Studies 2021-2023 No. 1

Reference hCNV datasets, use-case workflows and benchmarking

- only limited datasets exist to test and benchmark tools for the analysis of CNV and structural variations
 - recent datasets focused on high-quality Whole Genome Sequencing (WGS) analyses but not on the most commonly used Whole Exome Sequencing (WES) or genomic array technologies
 - generation of publicly accessible reference sets (raw and interpreted CNV data) for a variety of technological platforms will allow the hCNV community to generate the mandatory material
 - creation of “control datasets” required by many detection tools
 - complement standardization and benchmarking efforts such as the “Genome in a Bottle” initiative
 - integrate with Galaxy community & platforms
- ▶ WP1 - Dataset selection and generation
 - ▶ WP2 - Analyse and Compare CNV with other Benchmarking initiatives
 - ▶ WP3 - Exploitation of the datasets by the Galaxy Community
 - ▶ WP4 - Training and dissemination



hCNV Implementation Studies 2021-2023 No. 2



Beacon and beyond – Implementation-driven standards and protocols for CNV discovery and data exchange

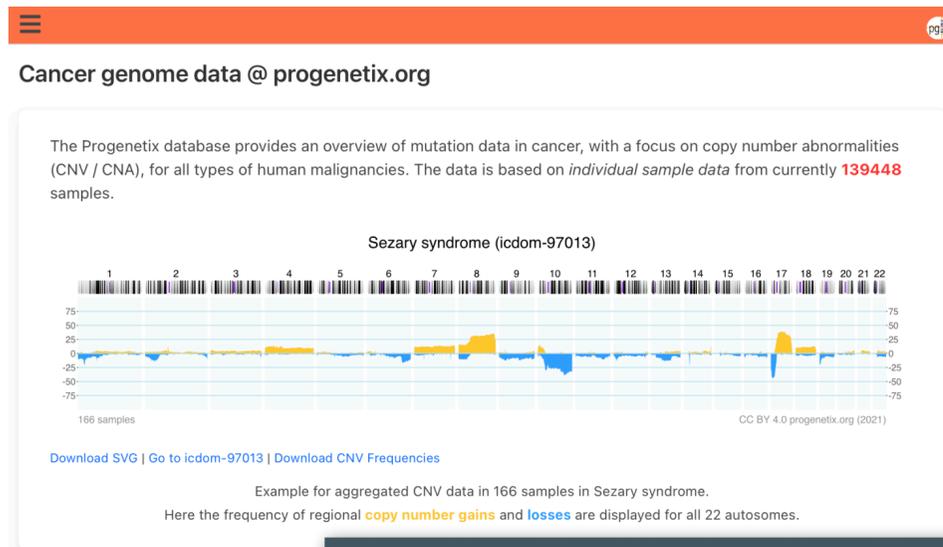
- reinforce work on priority areas established in the current hCNV Implementation Study
 - extend collaborations with the Rare Diseases and Galaxy Communities, EJP-RD and GA4GH
 - Expected outcomes
 - ➔ shared CNV resources testing advanced versions of the Beacon protocol
 - ➔ integration of GA4GH standards such as Phenopackets in such resources
 - ➔ tools for data ingestion and export for standard formats (e.g. VCF, Phenopackets) and CNV-specific improvements of such standards
 - ➔ ELIXIR AAI demo on clinical and research hCNV resources
 - ➔ demonstration of Galaxy pipeline adoption for real-world hCNV data analysis projects
 - connecting to international partners, e.g. Cancer Genomics Consortium (U.S.)
- ▶ WP1 - hCNV community reference resources
 - ▶ WP2 - hCNV Resources and Beacon
 - ▶ WP3 - Galaxy Community Intersection and Data Exchange
 - ▶ WP4 - Workflows and Tools for hCNV Data Exchange Procedures
 - ▶ WP5 - Training and dissemination



hCNV Implementation Studies 2021-2023 No. 2



Beacon and beyond – Implementation-driven standards and protocols for CNV discovery and data exchange



- ▶ WP1 - hCNV community reference resources
- ▶ WP2 - hCNV Resources and Beacon
- ▶ WP3 - Galaxy Community Intersection and Data Exchange
- ▶ WP4 - Workflows and Tools for hCNV Data Exchange Procedures
- ▶ WP5 - Training and dissemination

"Galaxify", "Beaconize" & "Phenopack"
Progenetix & RD-CNVdb prototypes



Beacon & Phenopackets



Data *discovery* and *delivery* using standardized GA4GH formats and schemas

- modern standards and protocols such as Beacon **v2** & Phenopackets **v2** are essential for federation and exchange of biomedical data
 - emerging / established principles are the use of hierarchical coding systems and with widespread use of CURIEs
 - other formats based on international standards, e.g.
 - ISO (ISO 8601 time & period, ISO 3166 country codes ...)
 - IETF (GeoJSON ...)
 - W3C (CURIE ...)
 - these standards become pervasive throughout GA4GH's ecosystem
- ➡ Beacon query **filters** correspond well to Phenopackets data
- ➡ Phenopackets as supported protocol for Beacon data delivery

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  "label" : "neoplastic sample"
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  "label" : "Medulloblastoma"
},
```

hCNV Implementation Studies 2021-2023

Focus on Integration with ELIXIR Platforms and Communities - and beyond

- original 2019-2021 implementation study provided visibility and established connections for new studies
- instrumental were Biohackathons, use case & standards surveys and co-participation of group members
- future work plans to leverage the resources of participants through pre-established interactions and synergies
- 2 independent studies provide clearer definitions of deliverables and individual scopes

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Björn Grüning	DE
Christophe Béroud	FR
David Salgado	FR
Jordi Rambla	ES
Kirill Tsukanov	EMBL-EBI
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