ArgenTAG

Next-generation single-cell sequencing.







Cells are the basic unit of life

- The human body has 37 trillion cells.
- Understanding how cells work could reveal hidden mechanisms of critical human diseases.



Single-cell methods aim to reveal the state of cells

- Cells combinatorially turn on/off genes.
- Genes that are "turned on" make RNA molecules.
- DNA sequencing machines read RNA molecules.







The state of a single cell fits in about 400 paper reams!

Current single-cell methods are Low Definition

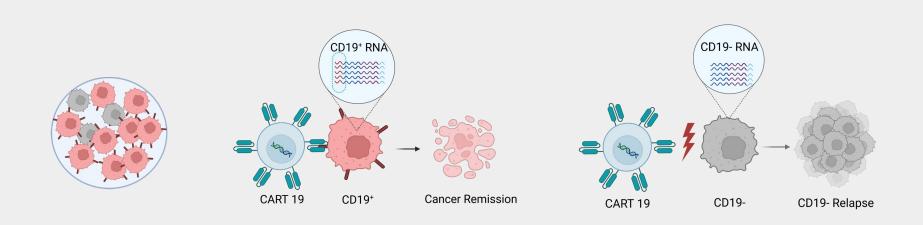
- 80% of human genes make +1 RNA type.
- Short-read sequencers only read RNA starts/ends.
- RNA differences per gene go beyond starts/ends.



Complete RNA information could make the difference.

Use Case: Current prediction of a cancer relapse in CAR T-cell therapy is difficult

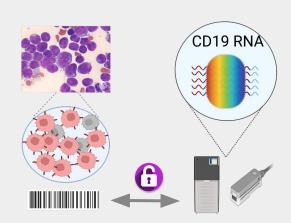
B-cell Acute Lymphoblastic Leukemia: 60-70% remission followed by 30-50% relapse.



Rabilloud T. et al. Single-cell profiling identifies pre-existing CD19-negative subclones in a B-ALL patient with CD19-negative relapse after CAR-T therapy. Nat Commun. 2021 Feb 8;12(1):865.

How to extend cancer relapse prediction with the next generation of single-cell methods

- Long reads to uncover true RNA diversity.
- High throughput to catch rare cells.
- Clean design to reach clinical research.



Millions of DNA barcodes on top of noisy long-read sequencers.

ArgenTAG high-definition single-cell kits to unlock the hidden 80% of human RNA landscape

Proprietary barcoding technology allows a device-free, automation-friendly protocol, on top of long-read (LR) sequencers.

Exploratory
Expert
Production



ArgenTAG's unique barcoding technology

- DNA barcodes safely carry cell IDs on top of noisy LR sequencers.
- DNA barcode composition follows
 Digital Communication principles to withstand LR sequencing noise.
- Scalable design, millions of robust DNA barcodes >>> enabling technology.



PATENT APPLICATION
US - FU



Ezpeleta, J. et al. Robust and scalable barcoding for massively parallel long-read sequencing. Sci Rep 12, 7619 (2022).

Early movers in the fast-growing single-cell market

- Enter LR sequencing with small, medium and large scale single-cell kits
- Consolidate LR sequencing with an instrument
- Expand LR sequencing market with solutions for the Global DNA sequencing market: target sequencing, metagenomics, microbial genome sequencing.

US\$ 40 B

Global DNA Sequencing Market Size, 2030

US\$ 2,9 B

TAM > Academic, Biotech, Pharma and Gov that could benefit from single-cell, 2030

US\$1B

SAM, Long-read seq customers by 2030

US\$ 100 M som - 10%

A family of kits for all kind of customers

> Pharmas, Government

Purpose: Cell therapies, population genomics Avg. expenditure: US\$ 729,640

Total customers 2021: 782 (PacBio + Nanopore)

US\$10,000

Production Large-scale Kit

> Biotechs, Core facilities

Purpose: Drug discovery Avg. expenditure: US\$ 78,416

Total customers 2021: 430 (Nanopore)

US\$ 5,000

Expert
Medium-scale Kit

Collaborations, Custom projects, Services

> Academic Labs, Pls

Purpose: Basic Research Ava. expenditure: US\$ 6,728

Total customers 2021: 5,501 (Nanopore)

US\$ 2,500

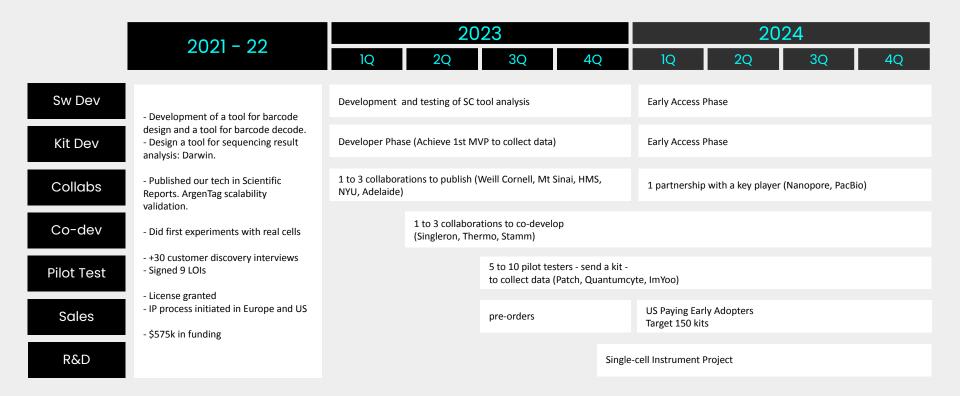
Exploratory Small-scale Kit

First single-cell high definition solution in the market

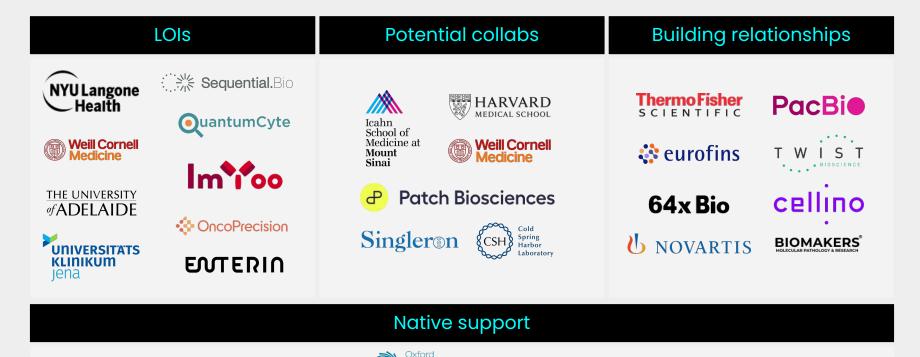
	ArgenTAG	Parse Bio	Fluent Bio	10x Genomics
Product	Kits	Kits	Kits	Instrument
Scalable	Yes	Yes	No	No
High Definition	Yes	No	No	No
Quick results	Yes	No	No	No
Price (50k reference cells)	\$5k per kit	\$10k per kit	N/A	Capex: \$65k, \$10k per run



Raising \$2,5 M to build MVP and kick off 3 collabs



Key players waiting for our solution





A team pushing the limits of biotechnology

> TEAM



CFO **ELIZABETH TAPIA, Ph.D.** Engineering



Dir. Process Dev **SOFIA LAVISTA, Ph.D.** Molecular Biology

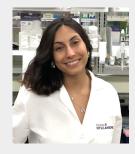




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Scientific Advisor **GEORGE CHURCH** Lead, Wyss Institute at Harvard University

INVESTORS









ArgenTAG

> First high-definition single-cell solution that will enable researchers, biotechs and pharmas to reveal the hidden RNA landscape of the human cells - to positively impact with a new generation of applications in oncology, immunology and neuroscience.





Learnings from customer discovery/interviews

Use case - Expert validation:

Dr. Joaquín Martinez Lopez - UCM, Spain Dr. Luis Lombardía - Molecular diagnostics, CNIO, Spain

Technology proposal - Expert validation:

Luciano Martelotto - Tech center, Adelaide University, AU Hagen - Lab, Weill Cornell, USA

Potential partners - Business

Thermofisher
PacBio
Singleron
Biomakers

Stamm

Novartis - email

Investors

Pre-work for Seed Round 15 investors waiting kick off Kick off: Oct 3 to Oct 21 Data room: Oct 24 to Nov 4 Term sheets: Nov 7 to Nov 18

WIP: Market size and data room

Proof of Concept

70% progress

Early adopter program

3 projects, 30% progress

Appendix

Customer and buying process

3500 existing single-cell customers using 10x12 experiments per year\$120,000 pull-through per year

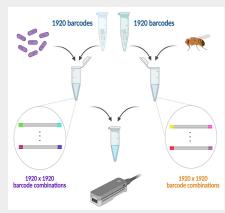
Decision maker: PI, Director level

Sell-in period: 3 months

ArgenTAG core tech validation

4,000 barcodes on top of 15% MinION sequencing errors

- 85% read recovery.
- 0.2% barcode missassignments.
- Illumina, 384 barcodes and 0.1% errors.



scientific reports
May 10, 2022 nature portfolio

Unlocks long read single-cell and more...

ArgenTAG (AT) single-cell pipeline



3. AT Protocol

2. AT Barcodes

4. LR Sequencing

5. AT Analysis

1. Cells in Suspension