

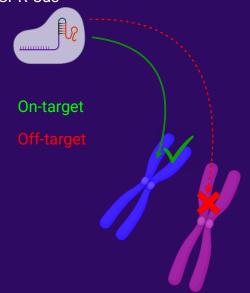
ADVANCING CELL & GENE THERAPY SAFETY

Felix Neumann
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2,200+ ongoing gene therapy programs, **but safety** remains a major concern

CRISPR-Cas



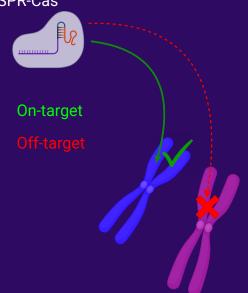
Gene editing requires thorough analysis from research to post-surveillance

Kalter, N., Fuster-García, C., et al. (2025). Off-target effects in CRISPR-Cas genome editing for human therapeutics: Progress and challenges. PubMed, 36(3), 102636. https://www.fda.gov/regulatory-information/search-fda-guidance-documents/human-gene-therapy-products-incorporating-human-genome-editing (since January 2024)

Future of Healthcare: Cell & Gene Therapies

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CRISPR-Cas



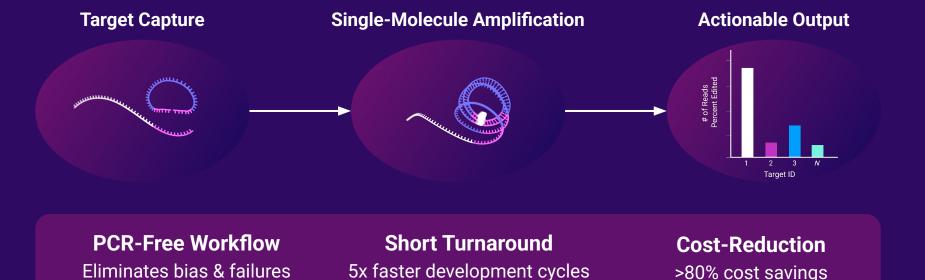
Gene editing requires thorough analysis from research to post-surveillance

FDA mandates comprehensive on- & off-target validation



Gold standard platform to satisfy all development stages is missing

Countagen solves the problem with a novel © PCR-free methodology



Proven technology through validation across segments & shared white papers

Countagen AB

Building a complete technology suite to serve gene editing needs



Imaging-Based

GeneAbacus

Targeted Validation

- Consumable reagent kit
- In-house analysis
- Early access customers

MDAnderson Cancer Center



Sequencing-Based

LockSeq

Comprehensive Analysis

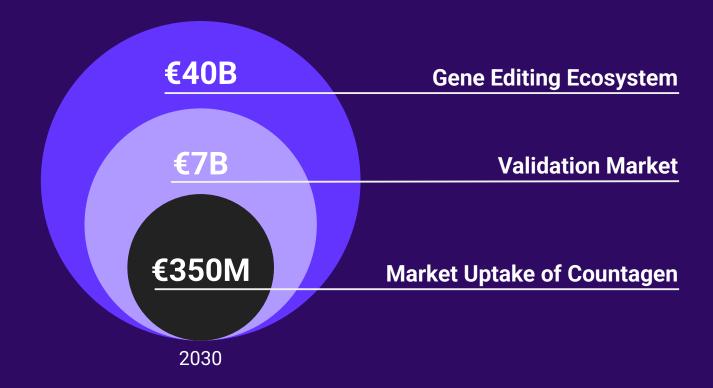
- Sample-to-report
- Full service
- Pilot collaborations

National Institute of Standards and Technology



Rapidly growing billion € opportunity





Regulatory tailwinds create mandatory demand

World-class team & assets







Mats Nilsson Professor at Stockholm University & Wellcome Sanger



Mårten Winge Serial Entrepreneur



Malte Kuhnemund R&D Director at 10X Genomics



Iván Hernández-Neuta CSO at Countagen



Felix Neumann CEO at Countagen

Scientific Advisors



Sören Turan Scientist at Bayer Pharmaceuticals



Bernhard Schmierer Head at the CRISPR Functional Genomics Unit











STRIKE Pharma

Countagen

Strong Intellectual Property

Probing & Signal Generation	Readout	Data & Technology
Gene editing analysis methods with padlocks and RCA GB2604872B1 P91668GB4	Methods and devices for single molecule DNA counting W02020212531A1 ² W02023075663A1 ³	Proprietary Software Market Intelligence & Network Unique Tech Know-How/Expertise

National phase: Seeking worldwide protection. Granted in UK. (1)

(4) Priority application filed in UK (Dec 2024).

⁽²⁾ National phase: Seeking protection in Europe. Granted in US.

⁽³⁾ National phase: Seeking worldwide protection.





€2M to Capture the US & EU Market

- 50% LockSeq platform optimization & service establishment
- 20% Market entry & expansion (US + Europe simultaneously)
- 15% GeneAbacus integration & partnership
- 15% Team growth & regulatory compliance

Key Milestones

- 2025: LockSeq KOL endorsement
- 2026: LockSeq service pilots; GeneAbacus partnership; €1.5M ARR
- 2027: €10M ARR; established player for comprehensive on & off-target validation

Our vision is to make Countagen the gold standard for cell & gene editing safety assessment



Let's Connect!

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