GENEOSCOPY

The colorectal cancer **prevention** company



Geneoscopy's proprietary GI health platform...



Stool instead of Blood





Stool: Direct source of epithelial cells, valuable GI biomarkers



Blood: Distant and diffuse signal, limited to advanced disease

RNA instead of DNA





RNA: Phenotypic, quantitative and dynamic information



DNA: Genotypic and qualitative, limited functionality and insight



Our unique approach has minimal direct competition and is protected via exclusive rights to 3 filed utility patents

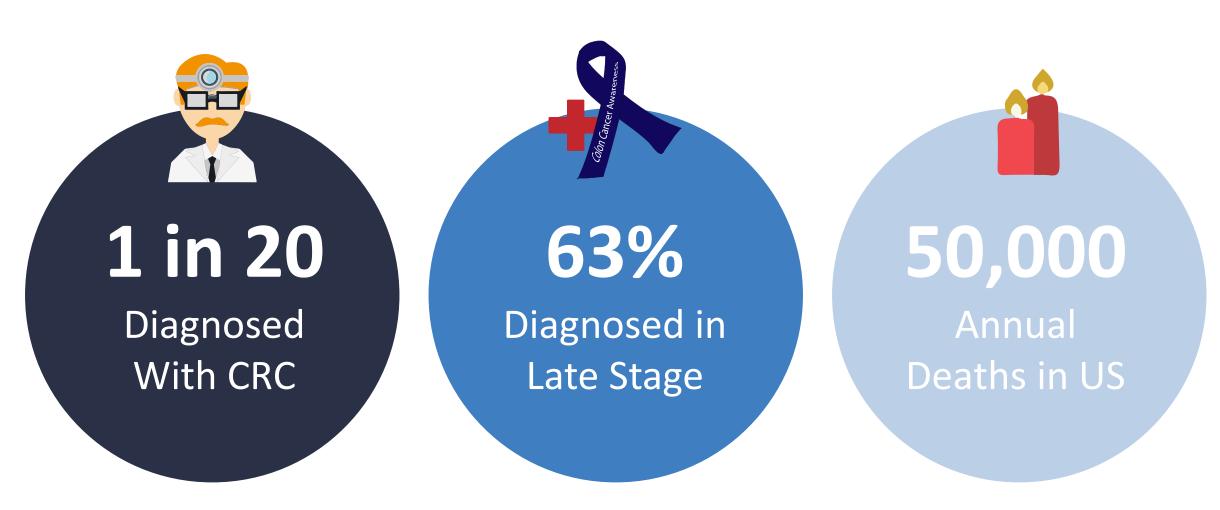
...offers a solution for a number of unmet clinical needs



					Current	ipeline Future	
	Screening	Diagnosis	Treatment	Surveillance	Recurrence	Drug Dev	
Colorectal Cancer							
Crohn's Disease							
Ulcerative Colitis							
Infectious Disease							
Necrotizing Enterocolitis							
Celiac Disease							
Irritable Bowel Syndrome							
Diabetic Gastroparesis							

Colorectal cancer is the 2nd deadliest cancer worldwide





Patient aversion to colonoscopy drives low compliance...



Bowel Prep

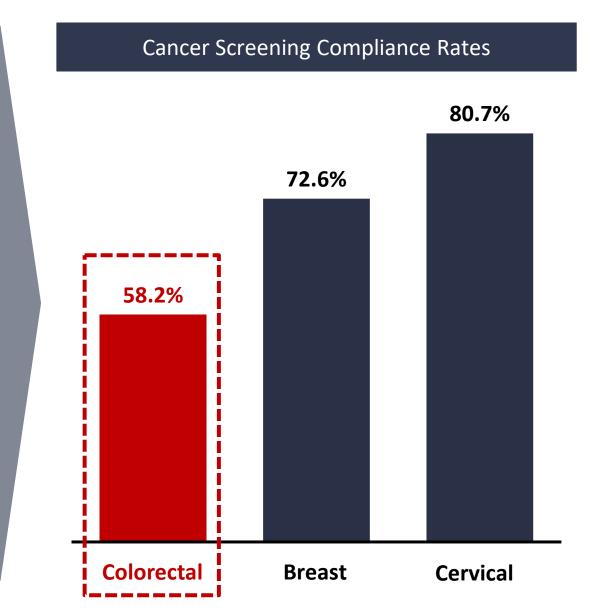
Sedation

Discomfort







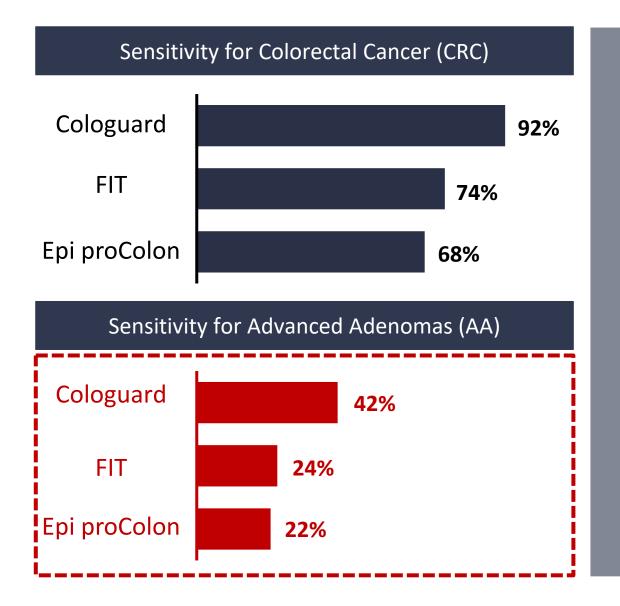


Noninvasive screening landscape



Test	Sample Type	Detection Method	Method Limitations	
Epi proColon	Blood	DNA biomarkers	CRC sensitivity AA sensitivity	
FIT Fecal Immunochemical Test Color-cuta Cance Cance Color-cuta Cance Color-cuta Cance Color-cuta Cance Color-cuta Cance Color-cuta Cance Color-cuta Cance Cance Color-cuta Cance Color-cuta Cance	Stool	Hemoglobin	AA sensitivity Compliance	
Cologuard	Stool	FIT + DNA biomarkers	AA sensitivity	
Geneoscopy	Stool	FIT + RNA biomarkers	RNA extraction/ preservation	

...and noninvasive tests are suboptimal at preventing CRC



"In this update of each organization's guidelines, screening tests are grouped into those that primarily detect cancer early and those that can detect cancer early and also can detect adenomatous polyps, thus providing a greater potential for prevention through polypectomy...It is the strong opinion of these 3 organizations that colon cancer prevention should be the primary goal of screening."





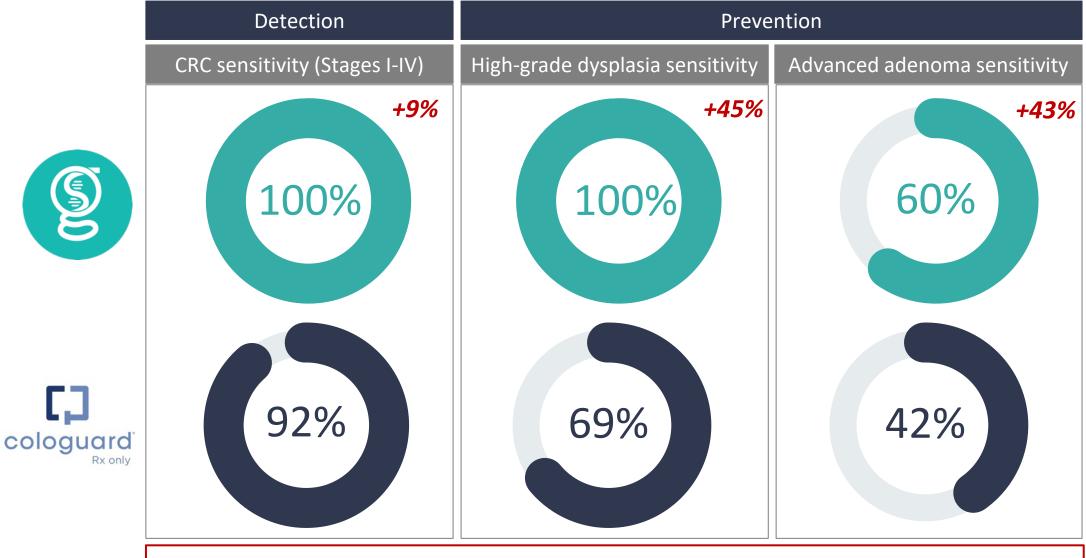






Geneoscopy prevents CRC through adenoma detection...





Further validation in >9,000-patient pivotal clinical study (June 2021)

...translating to an attractive product for stakeholders



Physician Choice Modeling



- ROSA modeled physician behavior to discern PCP/ GI preferences for CRC screening tests
- Simulator estimates a 28%-35% preference share for Geneoscopy (\$4.3B-\$5.4B annual revenue)
- 4 out of 6 top test attributes are directly related to prevention via adenoma detection

PCP Top Attributes			GI Top Attributes			
1.	Patient out-of-pocket cost	1.	Colorectal cancer sensitivity			
2.	High grade dysplasia sensitivity	2.	Advanced adenoma sensitivity			
3.	Advanced adenoma sensitivity	3.	High grade dysplasia sensitivity			

Health Economics Analysis



- PRECISIONheor has completed a costeffectiveness evaluation for CRC screening
- Geneoscopy's prevention-based approach improves health outcomes vs. Cologuard:
 - 15.8% reduction in CRC cases through CRC prevention eliminating 8,400 cases of CRC per year
 - ➤ 16.3% reduction in CRC mortality through CRC prevention and early detection saving 3,000 lives per year
- Geneoscopy's test fits a value-based system driven by quality measures (e.g., Star ratings)

Third-party research confirms that physicians and payers prefer prevention capabilities

Colorectal cancer prevention is an attractive market...







Large Total Addressable Market

108 Million Screening Population (Americans ages 45-75)

÷

3.0 Year
Screening
Interval
(Cologuard
intended use)

X

85.0%
Screening
Compliance
(Long-run
societal
target)

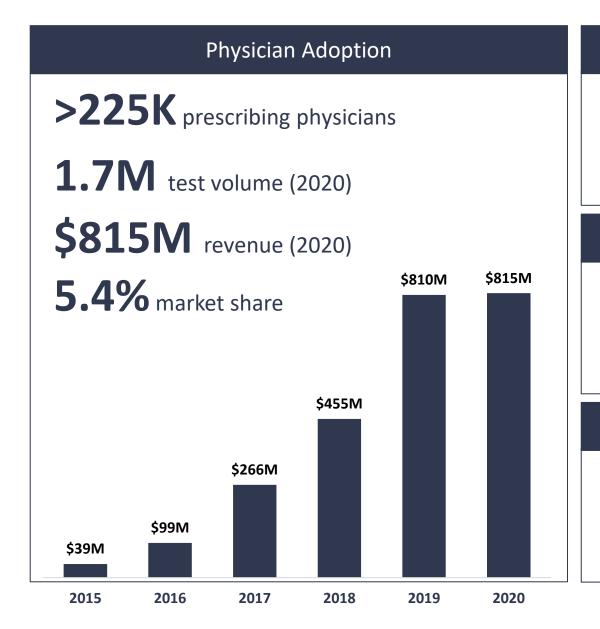
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\$500 Revenue Per Test (Cologuard level)

\$15.3 Billion
Target
Market

...validated by Cologuard's commercial traction





Patient Experience

\$0 patient out-of-pocket cost

88% rated experience to be very positive

Payor Support

97% insurance coverage

\$509 CMS reimbursement

Investor Interest

\$21.3B market cap (April 20, 2021)

\$3.0B+ capital raised in public markets

Core team with strong foundation in precision medicine





Andrew Barnell, MBA Chief Executive Officer







Erica Barnell, PhD Chief Science Officer





Management Team



Vince Wong Chief Commercial Officer





Gary Gallimore VP, Software & IT





Julie LaRocca VP, Quality / Regulatory





Ann Zuniga *Sr. Director, Product Dev*







Sr. Director, Product Mamt









Advisors / Board Members



Jim Merselis **Diagnostics Executive**





Katherine Tynan, PhD Diagnostics Consultant





Don Hardison Former Exact Sciences CEO





Appendix

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Pillars of Geneoscopy's standalone launch strategy













DTC Marketing

Digital patient activation via DTC prevention messaging

Channel Partnerships

Partnered sales force with key strategic

Telehealth Ordering

Telehealth-driven prescribing model via captive physicians

KOL Support

GI-focused education and engagement strategy Preferred
Payer Status

Health economics and outcomes research with robust data

Why It Works

Key

Activity

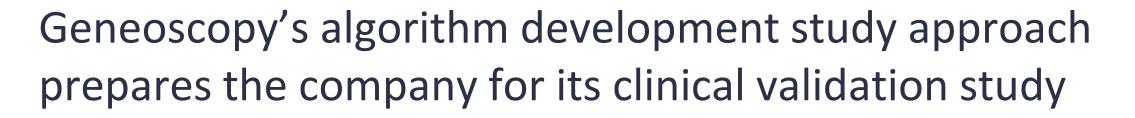
Low fixed cost infrastructure to drive consumer demand

Low fixed cost infrastructure to educate physicians

Rapid scalability
via growing
telehealth market
penetration

GI thought leadership influences PCPs and OB/GYNs Reduction in population health costs via prevention

"Prevention" is the critical medical marketing differentiator with all stakeholders





Real-World Study Design

- **Population:** United States
- > Sites: Hundreds of endoscopy sites
- > Collection Method: 99.8% prospective
- > Patients: Intended use population
- > Handling: Samples shipped in mail for up to 4 days
- ➤ Instrumentation: Final FDA cleared platforms
- > Reproducibility: Varied lots across the test system
- > Features: 14 available features
- > Thresholds: Selected within training folds

Reproducible Model Development

- ➤ Positive Patients: CRC, advanced adenomas, other precancerous adenomas (41% of patients)
- ➤ Negative Patients: Benign / hyperplastic polyps, healthy patients (59% of patients)
- Model Features: 10 markers (FIT, smoking status, 8 RNA biomarkers)
- ➤ Model: Ordinal regression
- ➤ Evaluation Technique: 5-fold Internal crossvalidation, hold-out testing set
- ➤ Threshold Setting: Targeted 85% specificity for patients with no findings on colonoscopy

Competitors: Prospective Study vs. Case Control Results



		cologuard	[] 2.0 cologuard	GRAIL	freenome	() GUARDANT	Training	Testing
Prospective	CRC Sens	92%					100% (n=3)	100% (n=4)
	AA Sens	42%					59% (n=66)	60% (n=50)
	OPA Sens	17%					22% (n=279)	25% (n=139)
	Specificity	88%					85% (n=591)	84% (n=175)
Case Control	CRC Sens	98%	92% ¹	79%²	91%³	79% ⁴		
	AA Sens	57%	46% ¹	No Data	41%³	No Data		
	OPA Sens	No Data	No Data	No Data	No Data	No Data		
	Specificity	90%	92%1	99%²	90%³	98%4		

¹ Partially prospective. Small adenomas were included, but no benign polyps and samples were weighted toward CRC and large advanced adenomas

² Analysis only included healthy patients and patients with cancer. Stage I-III sensitivity = 70%

³ Analysis only included healthy patients and patients with cancer selected from a larger patient cohort. AA Sens includes over representation of high-grade dysplasia and/or villous architecture, and was derived from a separate data set and algorithm than the cancer sensitivity

⁴ Analysis only included healthy patients and patients with cancer. Stage I sensitivity = 64%

Liquid biopsy assays do not represent near-term threats

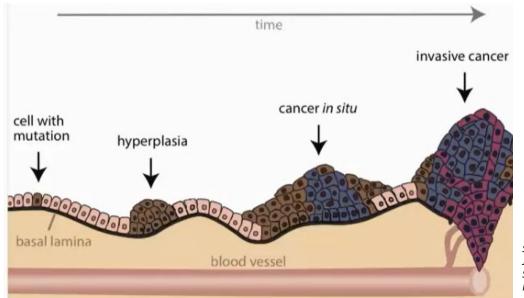


Limitations of Liquid Biopsy Research

- Study Design: Retrospective, case-control studies have limited reproducibility and largely fail to address prevention through adenoma detection
- Scalability: Studies currently report on <50% of enrolled patients implying either selection bias or assay failure due to sample input requirements
- Early Stage / Tumor Origin Detection: Lack of high-quality tissue of origin predictions combined with low early-stage sensitivity (<25%) limits clinical actionability</p>
- Large Feature Pools: Whole genome / methylome assays are incredibly expensive and will face reimbursement pushback from payers

Blood Has Limited Access to Early-Stage / Precancerous Biomarkers

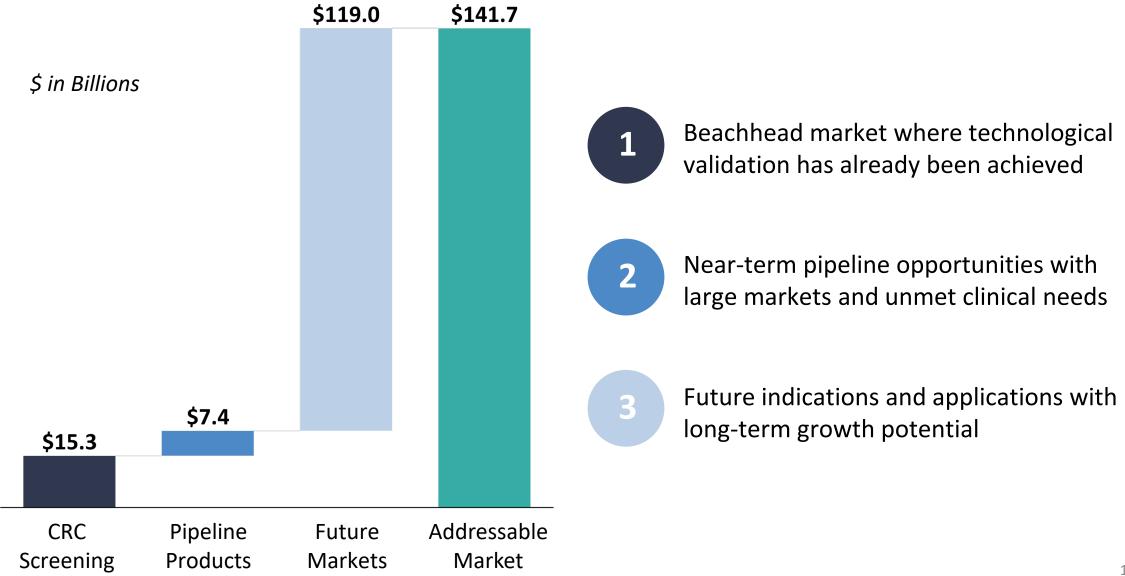
- Liquid biopsy approaches rely on biomarkers that are physically segregated from blood via biological barriers:
 - Lamina propria separates colonic cells from submucosa
 - Adhesion molecules prevent basement membrane penetration
 - Connective tissue is comprised of sticky stromal cells
 - Endothelial basement membrane separates plasma from colonic cells / biomarkers



Source: Johns Hopkins School of Medicine

GI represents large, established, and growing markets





Thank You

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