



# Announces Third Quarter 2022 Earnings and Acquisition of Fulgent Pharma

November 7, 2022

# Disclaimer

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## Forward-Looking Statements and Market Data

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This presentation also includes market data and forecasts with respect to the industry in which the Company operates. In some cases the Company relies upon and refers to market data and certain industry forecasts that have been obtained from third-party surveys, market research, consultant surveys, publicly available information and industry publications that the Company believes to be reliable. These data and estimates involve a number of assumptions and limitations, and you are cautioned not to give undue weight to such estimates.

## Non-GAAP Financial Measures

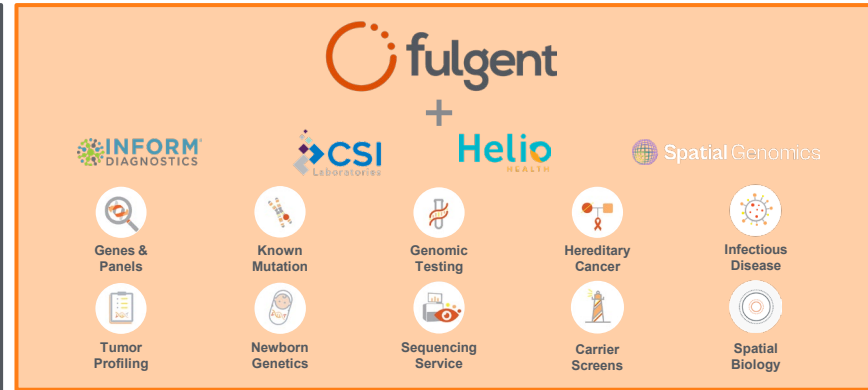
This presentation contains certain supplemental financial measures that are not calculated pursuant to U.S. generally accepted accounting principles (“GAAP”). These non-GAAP measures are in addition to, not a substitute for or superior to, measures of financial performance prepared in accordance with GAAP. A reconciliation of non-GAAP measures to GAAP measures is contained in this presentation.



**Ming Hsieh**  
Chairman, CEO, Founder

# Transaction Overview

- Fulgent Genetics acquired all outstanding capital stock of Fulgent Pharma at an enterprise value of \$100 million, in a combination of Fulgent common stock and cash
- Closing was November 7, 2022
- Fulgent gains access to novel nano-drug delivery platform with US FDA DMF
  - Lead drug candidate ready for Phase II/III clinical trials
  - Strong oncology pipeline using the same delivery platform with shortened development time
  - 32 issued and 4 pending patents
- Acquired a talented scientific team with proven track record
- Transforms FLGT from a genomic testing/service business into a fully integrated precision medicine company to address continuum of cancer care

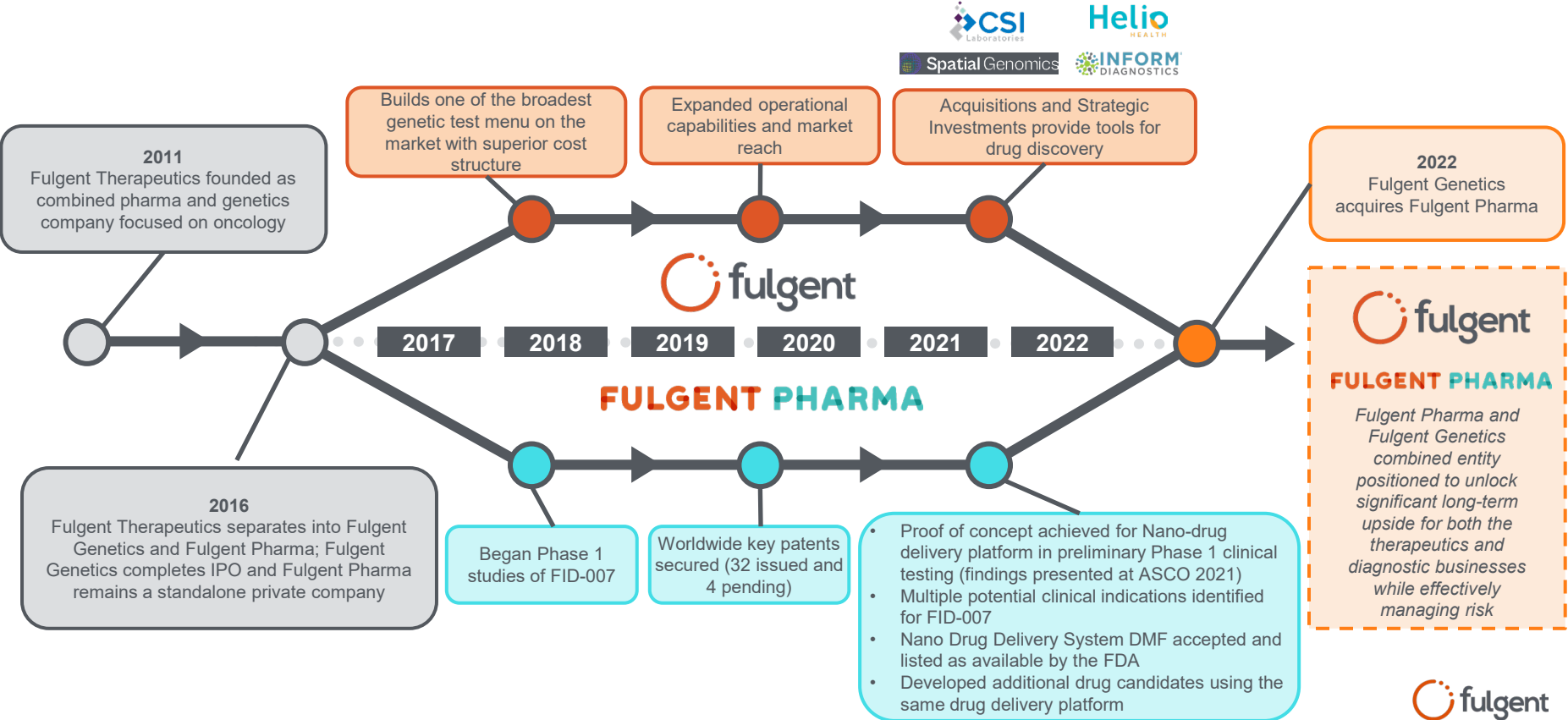


**FULGENT PHARMA**

**Nanoencapsulation Platform Technology  
with Lead Asset FID-007**

Initial Target Indications: H&N, Pancreatic, Breast, and Lung Cancer

# Fulgent Genetics + Fulgent Pharma History



# Strategic Vision – A One-stop Solution for Cancer Care



*To build a vertically integrated solution to combat cancer*  
early detection | clinical diagnostics | post treatment monitoring | drug discovery and cancer treatment



*Leading Genetic Testing Company Offering Tech-Enabled Diagnostic Solutions*



**Nano-Drug  
Delivery Platform**

**FULGENT PHARMA**

*Exciting Cancer Therapeutic Opportunity  
Realizing Precision Medicine Potential*

## Therapeutic and Diagnostic Entity Providing Comprehensive Solutions Across the Cancer Care Continuum

- Vertically integrated “one-stop” solution across the healthcare chain following the CSI, Inform Diagnostics, and Pharma acquisitions
- Proprietary nano drug delivery technology platform serves as an underpinning technology between diagnostic and pharm to help create a more sustainable and profitable business model in precision medicine for years to come
- Addition of a talented scientific team creates a strong synergy and competitive advantage that may be leveraged across the combined business
- Potential near-term opportunity includes shortened 505(b)(2) drug development and commercialization timelines and potential long-term opportunity leverages large data insights and novel analytical tools from diagnostics business to enable additional precision medicine pipeline through organic or partnered development strategies
- Commitment to continue growing diagnostic and therapeutic opportunities through organic investments and M&A
- Seasoned management team along with strong cash position allow Fulgent to enter therapeutic opportunities while managing risk

# Long-Term Vision: Fulgent Continuum of Care

Diagnosis

Therapy

Database + Drug Discovery  
+ Patient Care



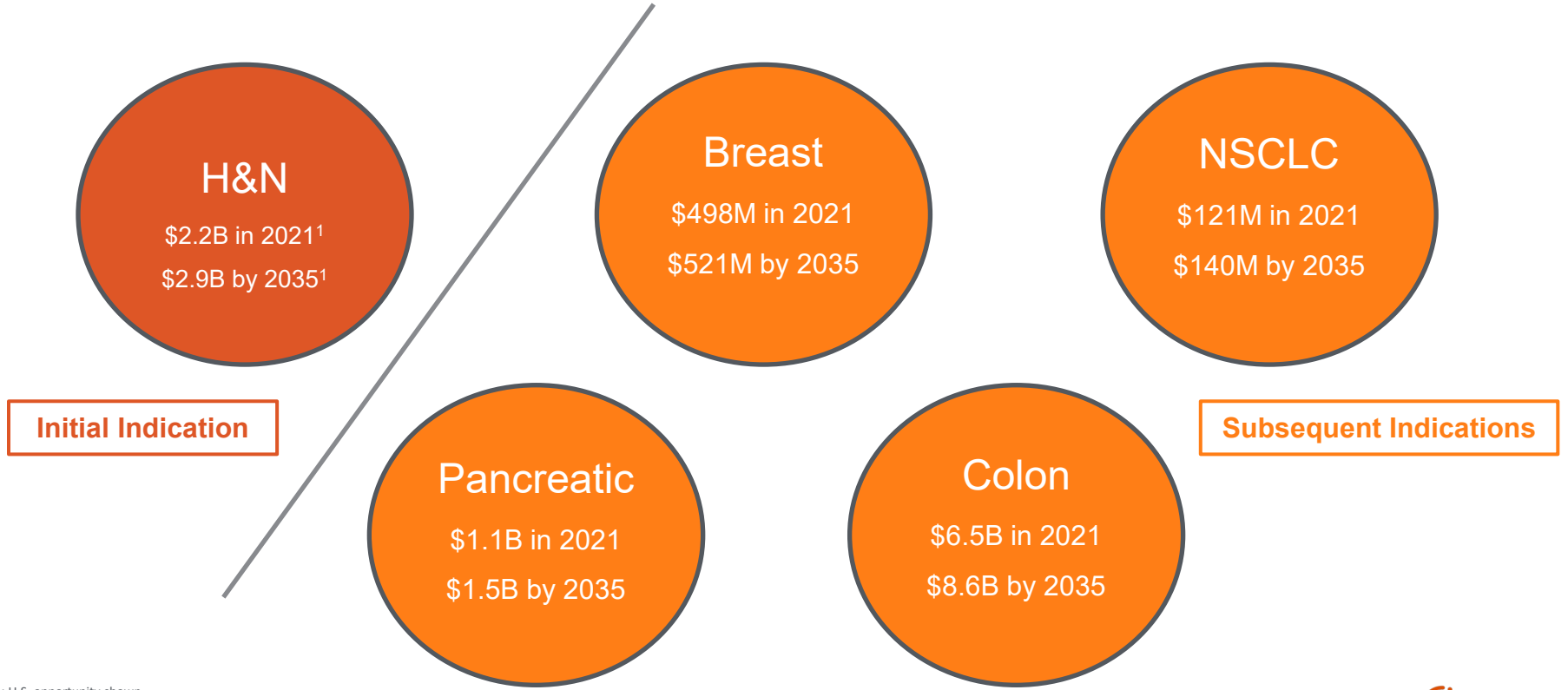
- Transforms from a service company to a combined diagnostic and therapeutic company focused on precision medicine
- Large oncology market opportunity
- Careful pipeline management will responsibly engage therapeutic opportunities while managing potential risks
- Valuable, diverse assets and future sustainable revenue and margins



## FULGENT PHARMA

- Well-funded by diagnostics to advance H&N therapy trials and commercialization process
- Genomics data accelerates pipeline therapeutics development, e.g. spatial biology for tumor micro-environment profiling
- Long-term opportunity to leverage data insights from diagnostics to enable precision medicine through proprietary or partnered development strategies
- Manufacturing capability to aid drug development

# Potential Market Opportunity



Note: U.S. opportunity shown  
Sources: Evaluate Pharma and Wall Street research estimates  
1. H&N market opportunity for both 2<sup>nd</sup> line and 3<sup>rd</sup> line therapy

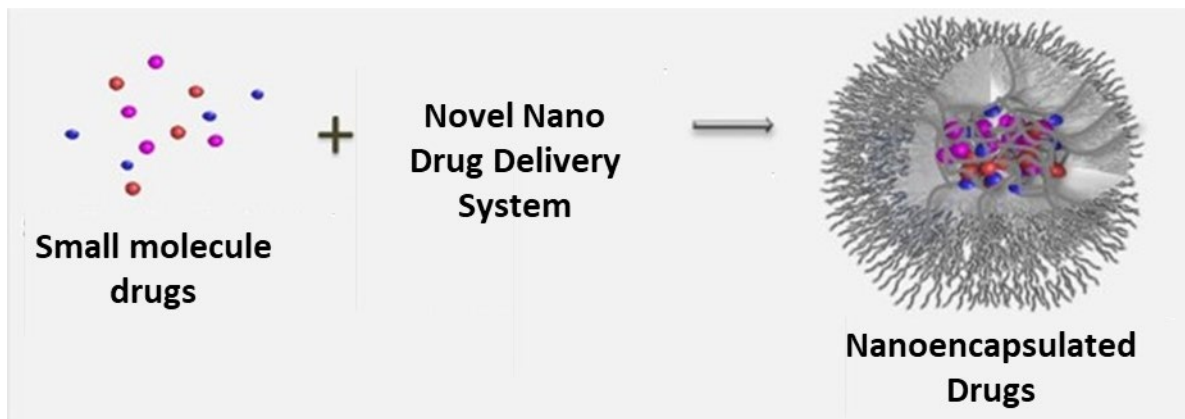




**Dr. Ray Yin, Ph.D.**  
**Co-Founder of Fulgent Therapeutics**  
**President of Fulgent Pharma**

- Founder & CEO, ANP Technologies, Inc.
- Former Team Leader of Nanobiotechnology for Chem/Bio Defense, U.S. Army Research Laboratory
- Holder of 46 drug delivery/detection patents

# Nano-Drug Delivery Platform Overview



## Novel Nano-Drug Delivery Platform

**Soluble in both water and various organic solvents and capable of hot melt mixing with APIs**

- Many drug candidates failed during preclinical and clinical development and testing due to poor water solubility
- Nanoencapsulation produces amorphous drug candidates with improved solubility and potentially enhanced absorption, drug PK profiles, safety and efficacy
- Broadly applicable to both IV and oral drug delivery formulations
- Potentially shortened development time
- Plug and play drug delivery platform provides multiple shots on goal
- Simple and low-cost production process

# FID-007 Program Overview

## FID-007 Phase I First in Human Clinical Trial – Preliminary Findings

- Dose levels up to 125 mg/m<sup>2</sup>/week with manageable safety profile, without yet reaching MTD
  - Dosing at 160 mg/m<sup>2</sup>/week is ongoing
- There is preliminary evidence of anti-tumor activity in heavily pre-treated patients across different tumor types
- Partial clinical data presented at ASCO 2021

## FID-007 Phase I Preliminary Highlights (as of 6/10/22):

### H&N Cancer

- 100% Disease Control Rate (DCR<sup>1</sup>) and 33% Overall Response Rate (ORR) were observed in 6 heavily treated H&N patients

### Ampullary/Pancreatic

- 75% DCR and 50% ORR were seen in 4 heavily treated ampullary and pancreatic patients

### Immune Checkpoint Inhibitors (ICIs) Resistant Patients

- 67% DCR and 33% ORR were seen in 6 heavily treated patients of different types of cancer with PD-1 or PD-L1 antibody treatment as the last line prior to enrollment in FID-007 trial

**Anticipate more data to be published in 2023**

## A Phase 1 Trial of FID-007, a Novel Nanoparticle Paclitaxel Formulation, in Patients with Solid Tumors

Jacob Thomas<sup>1</sup>, Diane Habib<sup>1</sup>, Diana Hanna<sup>1,2</sup>, Irene Kang<sup>1</sup>, Syma Iqbal<sup>1</sup>, Jorge Nieva<sup>1</sup>, Denise Tsao-Wei<sup>1</sup>, Francisco Acosta<sup>1</sup>, Ming Hsieh<sup>3</sup>, Yilong Zhang<sup>3</sup>, Anthony El-Khoueiry<sup>1</sup>

<sup>1</sup>University of Southern California, Norris Comprehensive Cancer Center; <sup>2</sup>Hoag Memorial Hospital; <sup>3</sup>Fulgent Pharma



Note: all findings are preliminary

1. DCR includes Stable Disease (SD), Partial Response (PR), Complete Response (CR)

# FID-007 and Other Candidates Using our Nano Delivery Platform

- Wholly-owned drug candidate focused on Head & Neck (H&N), Pancreatic, Lung, and Breast Cancer
  - Seeking initial therapeutic indication for 2<sup>nd</sup> or 3<sup>rd</sup> line treatment of H&N cancer
  - Exploring potential ampullary or ICI resistant
- Small molecule therapy uses proprietary nanoencapsulation technology, which may help mitigate toxicity while maintaining tumor reduction efficacy
- Potential FDA approval strategy uses 505(b)(2) studies, which may shorten clinical trial process and accelerate timeline to commercialization

| Drug Candidates | Target        | Indication                             | Pre-Clinical | Clinical P1 | Clinical P2 | Clinical P3 | Milestones   |
|-----------------|---------------|--|--------------|-------------|-------------|-------------|--|
| FID-007         | Cytotoxic     | Potential BE to Abraxane (505(b)(2))   | ▶            |             |             |             | Present P1 Data 2023<br>Begin P2/3 Enrollment 2023 |
|                 |               | Head and Neck (H&N) (505(b)(2))        | ▶            |             |             |             | Begin P2 Enrollment 2024                           |
|                 |               | Ampullary or ICI Resistant (505(b)(2)) | ▶            |             |             |             | Go/No-go Based on BE Study                         |
| FID-022         | Cytotoxic     | Colon and others                       | Pre-IND ▶    |             |             |             | IND-enabling Study in 2023<br>IND Submission 2024  |
| FPS-002         | STING Agonist | Vaccine Adjuvant                       | Pre-IND ▶    |             |             |             | Potential Partnership                              |

# Preclinical Pipeline

*Robust Pipeline Focused on Unmet Needs in Oncology*

## *505(b)(2) Approach*

| Drug Candidates | Target      | Indication      |
|-----------------|-------------|-----------------|
| FID-021         | Undisclosed | Multiple Cancer |
| FID-023         | Undisclosed | Leukemia        |
| FID-025         | Undisclosed | Brain Cancer    |

## *NCE Approach*

| Drug Candidates | Target                 | Indication                                  |
|-----------------|------------------------|---|
| FPT-020         | Multi kinase inhibitor | Gastric, Colon, Bladder, Endometrial Cancer |
| FPT-006         | Multi kinase inhibitor | Leukemia                                    |
| FPB-001         | BMI1 inhibitor         | Brain Cancer                                |

**Genomic Database Fuels Development and Addresses Issues of Drug Resistance**



**Ming Hsieh**  
Chairman, CEO, Founder

# Key Takeaways

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- Transaction reinforces Founder vision and potentially transforms the existing business into a new paradigm, generating a creative and sustainable business model in precision medicine for years to come
- “One-stop” shop verticalized across the healthcare chain and Fulgent Genetics’ increased customer base following the CSI, Inform Diagnostics, and Pharma acquisitions
- Initial therapeutic indication for 2<sup>nd</sup> or 3<sup>rd</sup> line treatment of Head & Neck (H&N) cancer has potential to provide an attractive entry point, rapid commercialization track, and a path to profitability in the therapeutic segment
- Long-term opportunity to leverage data insights from diagnostics business to enable precision medicine through proprietary or partnered development strategies
- Commitment to continue growing diagnostic and therapeutic opportunities through organic investments and M&A



**FULGENT PHARMA**

# THIRD QUARTER 2022 FINANCIAL RESULTS







**Brandon Perthuis**  
Chief Commercial Officer



**Paul Kim**  
Chief Financial Officer

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**Q&A**



