UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 8-K

CURRENT REPORT
Pursuant to Section 13 or 15(d)
of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): January 13, 2025

Foghorn Therapeutics Inc.

(Exact name of registrant as specified in its charter)

Delaware	**001-39634**	**47-5271393**
(State or other jurisdiction of incorporation)	**(Commission File Number)**	**(IRS Employer Identification No.)**

500 Technology Square, Ste 700
Cambridge, MA **02139**
(Address of principal executive offices) (Zip Code)

(Registrant's telephone number, including area code): (617) 586-3100

Not Applicable
(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:

☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)

☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12)

☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b))

☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	**Trading Symbol(s)**	**Name of each exchange on which registered**
Common Stock, $0.0001 par value per share	**FHTX**	**The Nasdaq Global Market**

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company ☒

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

Item 2.02 Results of Operations and Financial Condition.

On January 13, 2025, Foghorn Therapeutics Inc. (the "Company") disclosed certain preliminary financial information for the fiscal year ended December 31, 2024 ahead of the 43rd Annual J.P. Morgan Healthcare Conference. Specifically, the Company disclosed that for the fiscal year ended December 31, 2024, the Company's current expectation with respect to its cash, cash equivalents and investments in marketable securities (unaudited) is $243.8 million.

On January 13, 2025, the Company issued a press release announcing these preliminary results and other developments. The press release is attached as Exhibit 99.1 hereto and incorporated by reference herein. Additionally, the Company is furnishing as Exhibit 99.2 to this Current Report on Form 8-K a presentation, dated January 2025, containing these preliminary results, which the Company intends to use in meetings with or presentations to investors.

The information in this Item 2.02 is unaudited and preliminary, and does not present all information necessary for an understanding of the Company's results of operations for the fiscal year ended December 31, 2024 or financial condition as of December 31, 2024. The audit of the Company's financial statements for the year ended December 31, 2024 is ongoing and could result in changes to the information in this Item 2.02.

The information in this Item 2.02 (including Exhibits 99.1 and 99.2 attached hereto) is being furnished and shall not be deemed "filed" for purposes of Section 18 of the Exchange Act, or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference into any filing by the Company under the Securities Act or the Exchange Act, except as expressly set forth by specific reference in such filing.

Item 7.01 Regulation FD Disclosure.

The information provided in Item 2.02 above is incorporated herein by reference.

The information in this Item 7.01 (including Exhibits 99.1 and 99.2 attached hereto) is being furnished and shall not be deemed "filed" for purposes of Section 18 of the Exchange Act, or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference into any filing by the Company under the Securities Act or the Exchange Act, except as expressly set forth by specific reference in such filing.

Forward-Looking Statements

This Current Report on Form 8-K "forward-looking statements." Forward-looking statements include, but are not limited to, statements regarding the Company's preliminary financial statements, and other statements identified by words such as "could," "may," "might," "will," "likely," "anticipates," "intends," "plans," "seeks," "believes," "estimates," "expects," "continues," "projects" and similar references to future periods. Forward-looking statements are based on our current expectations and assumptions regarding capital market conditions, our business, the economy and other future conditions. Because forward-looking statements relate to the future, by their nature, they are subject to inherent uncertainties, risks and changes in circumstances that are difficult to predict. As a result, actual results may differ materially from those contemplated by the forward-looking statements. Important factors that could cause actual results to differ materially from those in the forward-looking statements include factors set forth under the heading "Risk Factors" in the Company's Annual Report on Form 10-K for the year ended December 31, 2023, as filed with the Securities and Exchange Commission.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description
99.1	Press release issued on January 13, 2025
99.2	Investor Presentation dated January 2025

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

FOGHORN THERAPEUTICS INC.

By: /s/ Kristian Humer
Kristian Humer
Chief Financial Officer

Date: January 13, 2025

Foghorn Therapeutics Highlights Program Progress and Strategic Objectives for 2025

First-in-class oral selective SMARCA2 (BRM) inhibitor, FHD-909 (LY4050784), advancing in Phase 1 trial for SMARCA4 mutated cancers, with non-small cell lung cancer (NSCLC) as the primary target patient population

FHD-909 preclinical combination data with pembrolizumab and KRAS inhibitors to be presented at the AACR Annual Meeting (April 25-30, 2025)

Selective degradation of ARID1B achieved with expected update in 2025; continued progress of Selective CBP degrader and Selective EP300 degrader

Strong balance sheet with cash, cash equivalents, and marketable securities of $243.8 million provides cash runway into 2027*

CAMBRIDGE, Mass. -- (GLOBE NEWSWIRE) – January 13, 2025 -- Foghorn® Therapeutics Inc. (Nasdaq: FHTX), a clinical-stage biotechnology company pioneering a new class of medicines that treat serious diseases by correcting abnormal gene expression, today announced its strategic objectives for 2025.

“In 2025, we expect important progress across our inhibitor and degrader programs. Our highly selective SMARCA2 inhibitor, FHD-909, continues to enroll and dose patients in a Phase 1 trial for SMARCA4 mutated cancers, with NSCLC as the primary patient population. In addition, for FHD-909, preclinical combination data with pembrolizumab and KRAS inhibitors will be presented at the AACR Annual Meeting in April, with partner Lilly,” said Adrian Gottschalk, President and Chief Executive Officer of Foghorn. “We are also excited with the progress of our preclinical pipeline. We have successfully achieved selective degradation of ARID1B, a major synthetic lethal target implicated in up to 5% of all solid tumors, and plan to provide a program update in 2025. We are continuing to advance our Selective CBP degrader and our Selective EP300 degrader, which are both implicated in a wide range of cancers. With our strong balance sheet and pipeline programs advancing, we look forward to an impactful 2025.”

**Unaudited, estimated as of December 31, 2024*

Program Overview and Upcoming Milestones

FHD-909 (LY4050784). FHD-909 is a first-in-class oral SMARCA2 selective inhibitor that has demonstrated in preclinical studies to have high selectivity over its closely related paralog SMARCA4, two proteins that are the catalytic engines across all forms of the BAF complex. Selectively blocking SMARCA2 activity is a promising synthetic lethal strategy intended to induce tumor death while sparing healthy cells. SMARCA4 is mutated in up to 10% of NSCLC alone and implicated in a significant number of solid tumors.

- **Advancing Phase 1 trial.** First patient dosed in October 2024 in the Phase 1 trial for FHD-909 in SMARCA4 mutated cancers, with NSCLC as the primary target population.
- **Preclinical combination data to be presented.** In 2025, preclinical data for FHD-909 in combination with pembrolizumab or KRAS inhibitors will be presented at the AACR Annual Meeting (April 25-30, 2025).

Ongoing strategic collaboration with Lilly. Collaborating with Lilly to create novel oncology medicines that includes a U.S. 50/50 co-development and co-commercialization agreement for Foghorn's selective SMARCA2 oncology program, agreements for a selective inhibitor and a selective degrader, and an additional undisclosed oncology target. The collaboration also includes three discovery programs from Foghorn's proprietary Gene Traffic Control® platform.

Selective CBP degrader program. Selectively targets CBP in EP300 mutated cancer cells found in many types of cancer, including bladder, gastric and endometrial cancers. CBP and EP300 are highly similar acetyltransferases that create a synthetic lethal relationship when EP300 is mutated. Attempts to selectively drug CBP have been challenging due to the high level of similarity between the two proteins, while dual inhibition of CBP/EP300 has been limited by hematopoietic toxicity.

- **Identified potent and selective CBP protein degraders.** Pharmacodynamic and pharmacokinetic preclinical data demonstrate:
    - Deep and sustained CBP degradation significantly inhibited tumor growth in mouse xenograft solid tumor models.
    - Robust monotherapy preclinical anti-tumor activity that was not associated with significant body weight loss, thrombocytopenia or anemia.
    - Long-acting injection formulation that resulted in tumor regression from a single dose in a mouse xenograft efficacy study.

Selective EP300 degrader program. Selective degradation of EP300 for the treatment of hematopoietic malignancies and prostate cancer. Attempts to selectively drug EP300 have been challenging due to the high level of similarity between EP300 and CBP, while dual inhibition of CBP/EP300 has been limited by hematopoietic toxicity. EP300 lineage dependencies are established in multiple myeloma, and diffuse large B cell lymphoma.

- **Identified potent and selective EP300 degraders and advancing oral degrader efforts.** Pharmacodynamic and pharmacokinetic preclinical data demonstrate candidates:
  - Are well tolerated *in vivo* with no observed decrease in platelet levels, and no effects on megakaryocyte viability at pharmacologically relevant concentrations in *ex vivo* studies.
  - Have robust anti-tumor activity in solid tumors and hematological malignancies, including prostate cancer, multiple myeloma, and diffuse large B cell lymphoma.

Selective ARID1B degrader program. Selectively targets and degrades ARID1B in ARID1A-mutated cancers. ARID1A is the most mutated subunit in the BAF complex and amongst the most mutated proteins in cancer. These mutations lead to a dependency on ARID1B in several types of cancer, including ovarian, endometrial, colorectal and bladder. Attempts to selectively drug ARID1B have been challenging because of the high degree of similarity between ARID1A and ARID1B and the fact that ARID1B has no enzymatic activity to target.

- **ARID1B is a major synthetic lethal target implicated in up to 5% of all solid tumors.**
- **Developed highly potent and selective binders.** Preclinical data demonstrated potent and selective small molecule binders to ARID1B.
- **Selective degradation of ARID1B achieved.** Foghorn has successfully selectively degraded ARID1B and expects to provide an update on the Selective ARID1B degrader program in 2025.

Chromatin Biology and Degrader Platform

Foghorn continues to advance its chromatin biology and degrader platform with investments in novel ligases, long-acting injectables, oral delivery and induced proximity.

Strong Balance Sheet and Cash Runway. As of December 31, 2024, the Company had $243.8 million cash, cash equivalents and marketable securities (unaudited), providing cash runway into 2027.

About Foghorn Therapeutics

Foghorn® Therapeutics is discovering and developing a novel class of medicines targeting genetically determined dependencies within the chromatin regulatory system. Through its proprietary scalable Gene Traffic Control® platform, Foghorn is systematically studying, identifying and validating potential drug targets within the chromatin regulatory system. The Company is developing multiple product candidates in oncology. Visit our website at www.foghorntx.com for more information on the Company, and follow us on X (formerly Twitter) and LinkedIn.

Forward-Looking Statements

This press release contains “forward-looking statements.” Forward-looking statements include, but are not limited to, statements regarding the Company’s initiation, timing, progress and results of research and development programs and pre-clinical studies and clinical trials, including with respect to the Phase 1 dose escalation trial of FHD-909 with Lilly, and other statements identified by words such as “could,” “may,” “might,” “will,” “likely,” “anticipates,” “intends,” “plans,” “seeks,” “believes,” “estimates,” “expects,” “continues,” “projects” and similar references to future periods. Forward-looking statements are based on our current expectations and assumptions regarding capital market conditions, our business, the economy and other future conditions. Because forward-looking statements relate to the future, by their nature, they are subject to inherent uncertainties, risks and changes in circumstances that are difficult to predict. As a result, actual results may differ materially from those contemplated by the forward-looking statements. Important factors that could cause actual results to differ materially from those in the forward-looking statements include regional, national or global political, economic, business, competitive, market and regulatory conditions, including risks relating to our clinical trials and other factors set forth under the heading “Risk Factors” in the Company’s Annual Report on Form 10-K for the year ended December 31, 2023, as filed with the Securities and Exchange Commission. Any forward-looking statement made in this press release speaks only as of the date on which it is made.

Contact:
Karin Hellsvik, Foghorn Therapeutics Inc.
khellsvik@foghorntx.com

Exhibit

FOGHORN®
THERAPEUTICS

Unique biology

Precision therapeutics

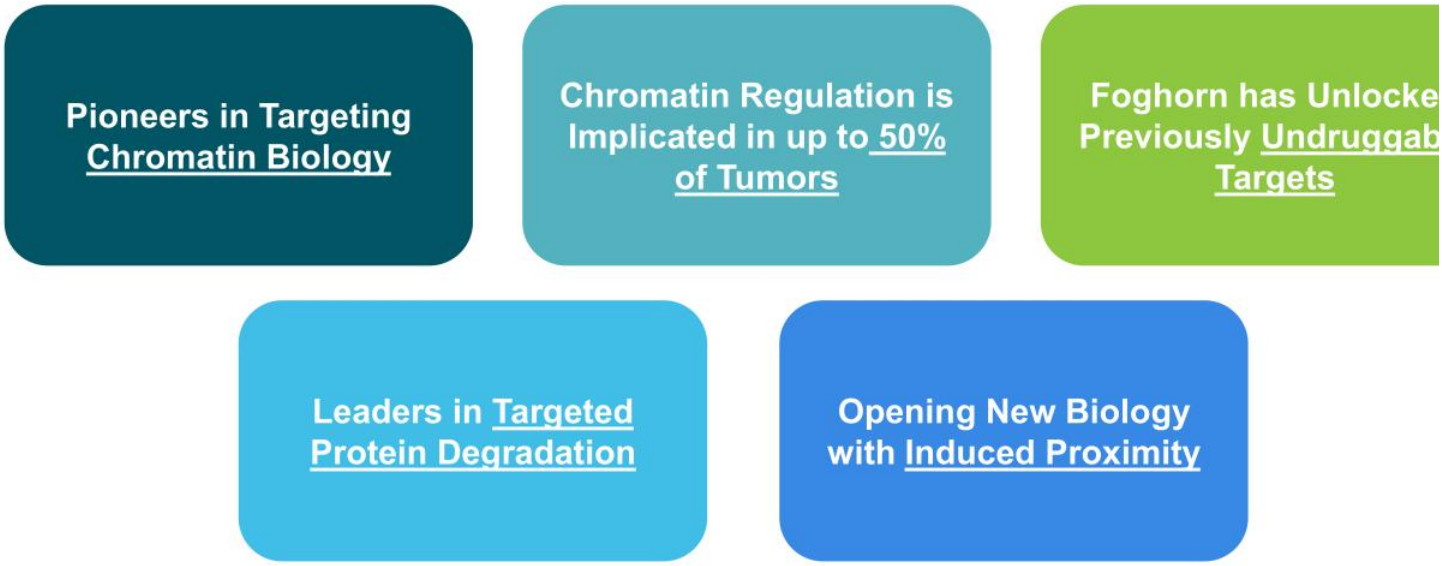
Broad impact

January 2025

Forward Looking Statements

This presentation contains forward-looking statements that are based on management's beliefs and assumptions and on infor currently available to management. All statements other than statements of historical facts contained in this presentation are fo looking statements. In some cases, you can identify forward-looking statements by terms such as “could,” “may,” “might,” “will,” “anticipates,” “intends,” “plans,” “seeks,” “believes,” “estimates,” “expects,” “continues,” “projects” or the negative of these te other similar expressions, although not all forward-looking statements contain these words. Forward-looking statements inclu are not limited to, statements concerning: the potential outcomes from our collaboration agreement with Lilly; the initiation, progress and results of our research and development programs and pre-clinical studies and clinical trials, including with respec Phase 1 dose escalation trial of FHD-909 with Lilly; our ability to advance product candidates that we may develop and to succe complete preclinical and clinical studies; our ability to leverage our initial programs to develop additional product candidates us Gene Traffic Control Platform®; the impact of exogeneous factors, including macroeconomic and geopolitical circumstances, and our collaborators’ business operations, including our research and development programs and pre-clinical studies; develo related to our competitors and our industry; our ability to expand the target populations of our programs and the availability of p for clinical testing; our ability to obtain regulatory approval for FHD-909 and any future product candidates from the FDA an regulatory authorities; our ability to identify and enter into future license agreements and collaborations; our ability to continue to our CDMOs and CROs for our manufacturing and research needs; regulatory developments in the United States and foreign co our ability to attract and retain key scientific and management personnel; the scope of protection we are able to establish, maint enforce for intellectual property rights covering FHD-909, our future products and our Gene Traffic Control Platform; and our proceeds from capital-raising transactions, estimates of our expenses, capital requirements, and needs for additional financi should, therefore, not rely on these forward-looking statements as representing our views as of any date subsequent to the date presentation. Additional important factors to be considered in connection with forward-looking statements are described Company's filings with the Securities and Exchange Commission, including withing the section entitled "Risk Factors" in the Com Annual Report on Form 10-K for the fiscal year ended December 31, 2023. Any forward-looking statements represent the Com views only as of the date of this presentation and should not be relied upon as representing its views as of any subsequent da Company explicitly disclaims any obligation to update any forward-looking statements. The Company’s business is sub substantial risks and uncertainties.

Foghorn is a Leader in Chromatin Biology, Successfully Drugging Challenging Targets



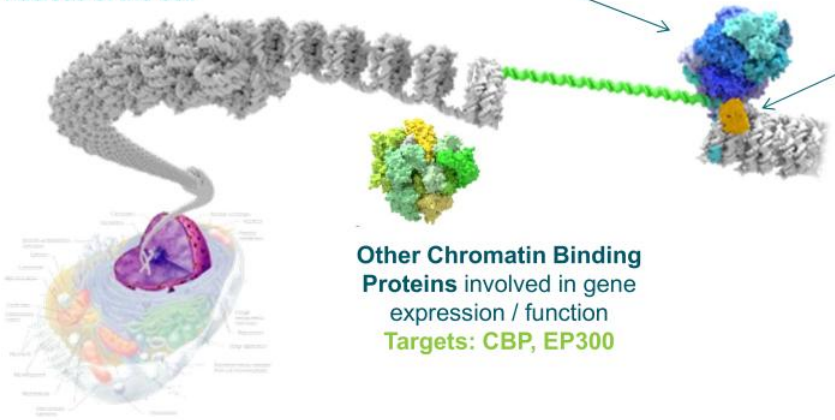
Chromatin Regulatory System Orchestrates Gene Expression: Multiple Opportunities for Targets and Therapeutics

Chromatin Regulatory System genes are **implicated** across a **wide range of cancers**

Chromatin – compacted form of DNA inside the nucleus of the cell

Chromatin Remodeling Complexes – specialized multiprotein machines that allow access to DNA
Targets: SMARCA2, ARID1B

Transcription Factors – proteins that help turn specific genes "on" o "off" by working in concert with the chromatin remodeling complex to bind to DNA
Targets: Multiple TFs



Other Chromatin Binding Proteins involved in gene expression / function
Targets: CBP, EP300

Leveraging **synthetic lethality** and **lineage dependencies**

Foghorn Has Progressed Multiple Programs Against Challenging Targets

SMARCA2: Potential in up to 10% of NSCLC and up to 5% of all solid tumors
Challenge: Industry has failed to develop a selective inhibitor

CBP: Role in bladder, colorectal, breast, gastric, lung cancers
Challenge: Toxicities with dual inhibition, difficulty engineering selectivity

EP300: Role in both solid and heme malignancies
Challenge: Toxicities with dual inhibition, difficulty engineering selectivity

ARID1B: Role in ovarian, endometrial, colorectal cancer. Potential in up to 5% of all solid tumors
Challenge: Industry has had no success with selective target engagement

FHD-909
First selective inhibito
in the clinic

Selective CBP Degrade
IND expected H1 2026

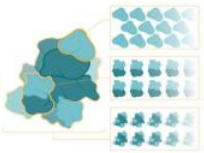
Selective EP300 Degrad
IND expected in H2 202

Selective ARID1B Degra
Degradation achieved; Pro
update expected in 202

… and more.

SMARCA2 = BRM

Foghorn's Gene Traffic Control® Platform Designed to Deliver Precision First-in-Class Therapeutics: Integrated, Scalable, Efficient, Repeatable

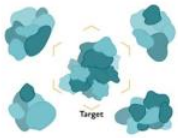


2. Assays & Biochemistry Capabilitie

Engineering selectivity via unique assays and protein capabilities

- Protein purification, production & interrogati
- High fidelity, difficult to make proteins
- In silico modeling and computational chemis

"Where to Drug"



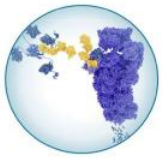
1. Chromatin Biology

Deep mechanistic understanding of chromatin regulatory system

- Bioinformatics
- Genomics
- Epigenomics

Identify Dependencies

"What to Drug"



3. Chemistry & Drugging

Biology first, small molecule modality agnostic

- Selective, small molecules (inhibitors, protein degraders, TF disruptors)
- Protein degradation platform
- Formulation & long-acting delivery

"How to Drug"

Legend: Patents | Know How / Trade Secret

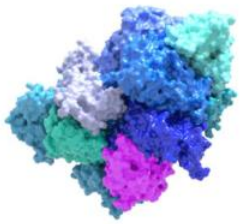
Foghorn's Unique Platform Capabilities Evolved From Drugging a Specific Chromatin Remodeling Complex (BAF)*

Challenge: produce, manipulate, study, and drug a 1.5 megadalton multi-protein complex

Assays and Biochemistry Capabilities

- Purification & recombinant production of large proteins and protein complexes
- Biochemistry & biophysics of intrinsically disordered proteins
- High throughput screening for binders and inhibitors

BAF Chromatin Remodeling Complex



Challenge: drug highly si proteins that have no enz function

Protein Degrader Platfo

- Proprietary linker library
- Suite of assays specific to degradati synthesis kinetics, degradation kineti
- Optimal E3 ligase pairing
- Ternary complex modeling
- Long-acting formulation technology

Current and Future Applications

- **Selectively drugging highly similar proteins / hard to drug proteins**
- **Disease area expansion**
- **Going beyond chromatin – novel biology with complex proteins**
- **Payloads for ADCs***

**Brahma-Associated Factor (BAF). Antibody Drug Conjugates (ADCs)*

Platform of Proprietary Tools and Unique Know-How Enables Development of Best-in-Class Degraders

Proprietary E3 Ligase Libraries

- Proprietary E3 ligase binders
  - ~ 15 -17 in internal library
  - UBR5

Computational Chemistry
(ML, AI, Virtual Screens)

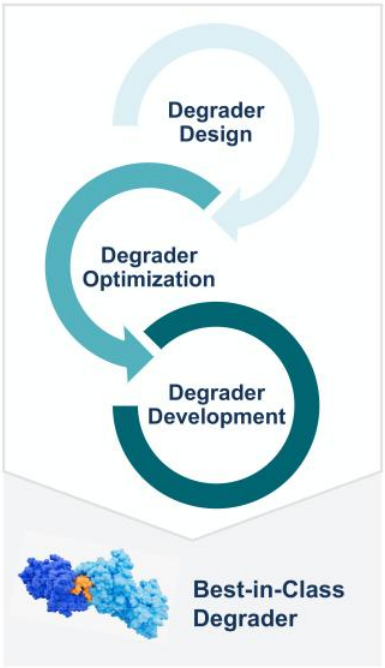
- Structural prediction/ docking

Linker Toolkit

- Well-developed toolkit
- Flexibility-rigidity
- Charge
- Hydrophobicity

Binder Discovery

- Commercial and proprietary covalent and fragment screening libraries
- Target binders



Degrader Mechanics

- Specificity / Selectivity (global proteomics)
- Kinetics (Dmax/DC50)
- Cooperativity (e.g. ternary complex)
- Permeability
- MoA characterization
- Cellular and permeability assays

Exploratory Biology

- *In Vitro* efficacy
- Indication selection
- Combinations
- Resistance mechanisms

***In Vivo* Vali**
(Efficacy and

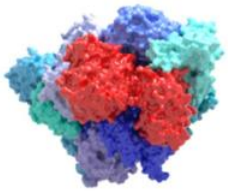
- *In Vivo* mod
- PK/PD dete modeling an prediction

Proprietary Formulation & Delivery Platform

- Oral, IV and LAI formulation for drug deliver

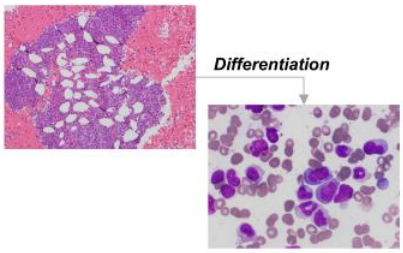
The Next Foghorn Chapter: Delivering Multiple Potential Blockbusters Into the Clinic

Pioneering BAF and Chromatin Biology
(2016 – 2020)



- ✓ Built platform and developed deep understanding of biology
- ✓ Producing BAF and transcription factors at scale
- ✓ **Demonstrated druggability of chromatin regulatory system**

POC, Platform & Pipeline Expansion
(2021 – 2024)

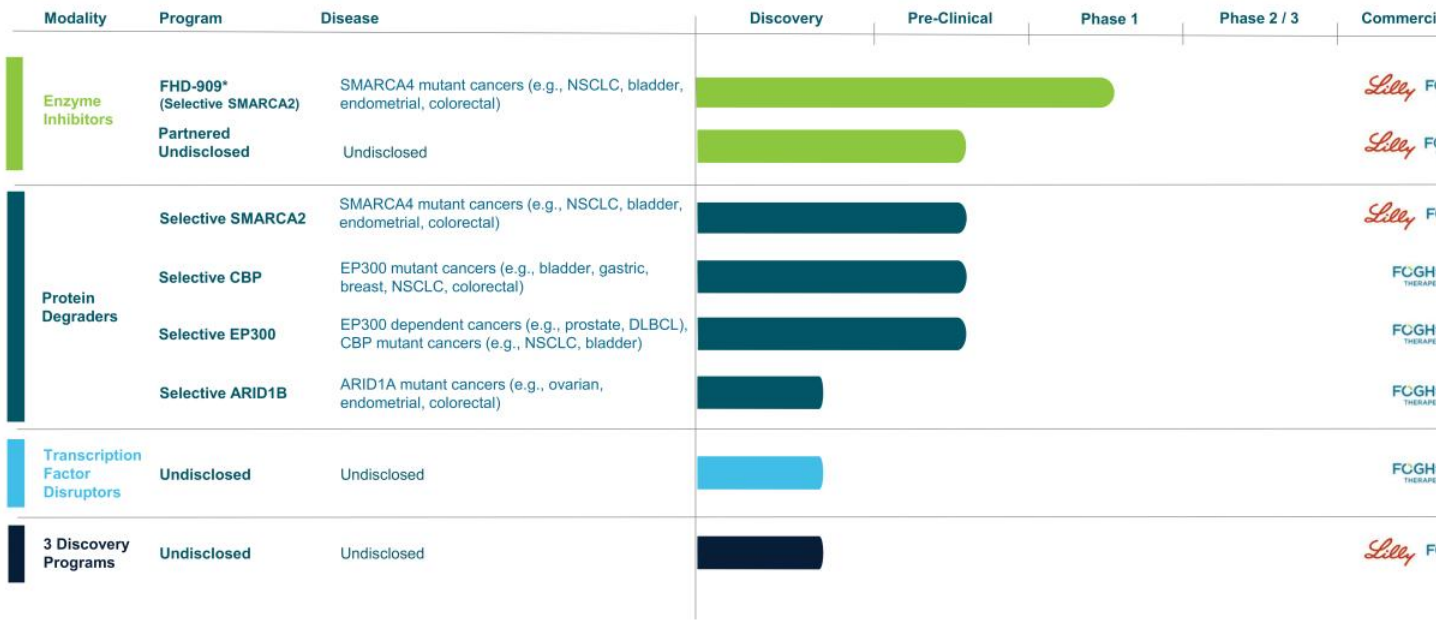


- ✓ **Lilly strategic collaboration**
- ✓ **Developed first-in-class Selective SMARCA2 inhibitor, FHD-909**, and SMARCA2 degrader
- ✓ Advanced Selective CBP degrader, Selective EP300 degrader and Selective ARID1B degrader
- ✓ Expanded protein degrader platform, new applications for Induced Proximity

Progress Multiple High Value Assets into the Clin
(2025 – 2027)

- **FHD-909 advancing in Phase 1 trial**
- **ARID1B degradation achieved – pr update expected in 2025**
- **Potential for 5 additional INDs; Pip platform, disease area expansion**

Foghorn is Advancing a Pipeline of First-in-Class Precision Therapeutics With Potential for Broad Application in Oncology…



*LY4050784
SMARCA2 = BRM

…With Multiple Near-Term Value Inflection Points Through 2026

FHD-909 (LY4050784) (Selective SMARCA2 Inhibitor)	Preclinical Combination Data	April 2025 (AACR)
	Phase 1 Dose Escalation Data	Confidential
Selective SMARCA2 Degrader	IND / Phase 1 Initiation	Confidential
Selective CBP Degrader	IND / Phase 1 Initiation	2026
Lilly Target #2	Target Disclosure and IND	Confidential
Selective EP300 Degrader	IND / Phase 1 Initiation	2026
Selective ARID1B Degrader	Selective Degradation Achieved	Program Update 2025

SMARCA2 = BRM
SMARCA4 = BRG1

Potential Multi-Billion Dollar Opportunities in Oncology

Foghorn Owned

Partnered w Lilly

Platform has potential for therapeutic area expansion (e.g., immunology and inflammation)

**LY4050784*

FOGHORN THERAPEUTICS

Clinical & Preclinical Programs

- FHD-909 (LY4050784) – Selective SMARCA2 Inhibitor
- Selective CBP Degrader
- Selective EP300 Degrader
- Selective ARID1B Degrader

Selective SMARCA2 Inhibitor and Degrader

For SMARCA4 Mutated Cancers

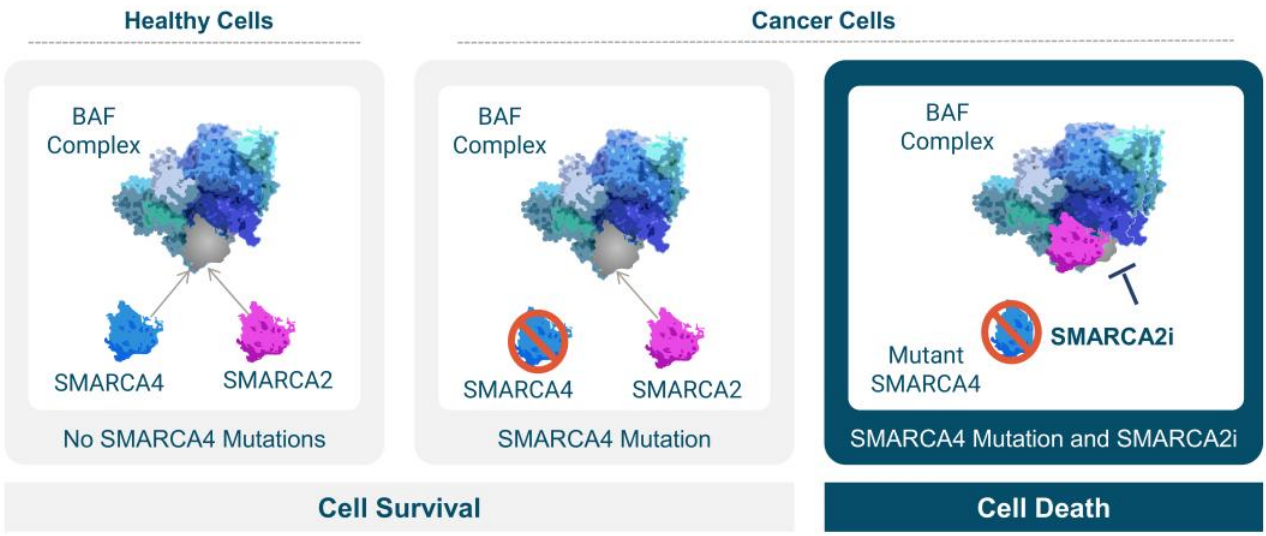
SMARCA2 = BRM
SMARCA4 = BRG1

FHD-909 Selective SMARCA2 Inhibitor in Phase 1 Trial; Preclinical Combination Data at AACR 2025

	Selective SMARCA2 Inhibitor FHD-909*	Selective SMARCA2 Degrader
Biology	Exploit the synthetic lethal relationship between SMARCA2 and mutated SMARCA4	
Stage / Next Milestone	Phase 1 dose escalation trial ongoing; Preclinical combination data with pembrolizumab, KRAS and chemo at AACR	Advancing through late preclinical development
Opportunity	SMARCA4 mutated cancer including ~10% of NSCLC and up to 5% of all solid tum	
Lilly Partnership	50/50 global R&D cost share | 50/50 U.S. economics | tiered ex-U.S. royalties sta in the low double-digit range and escalating into the twenties	

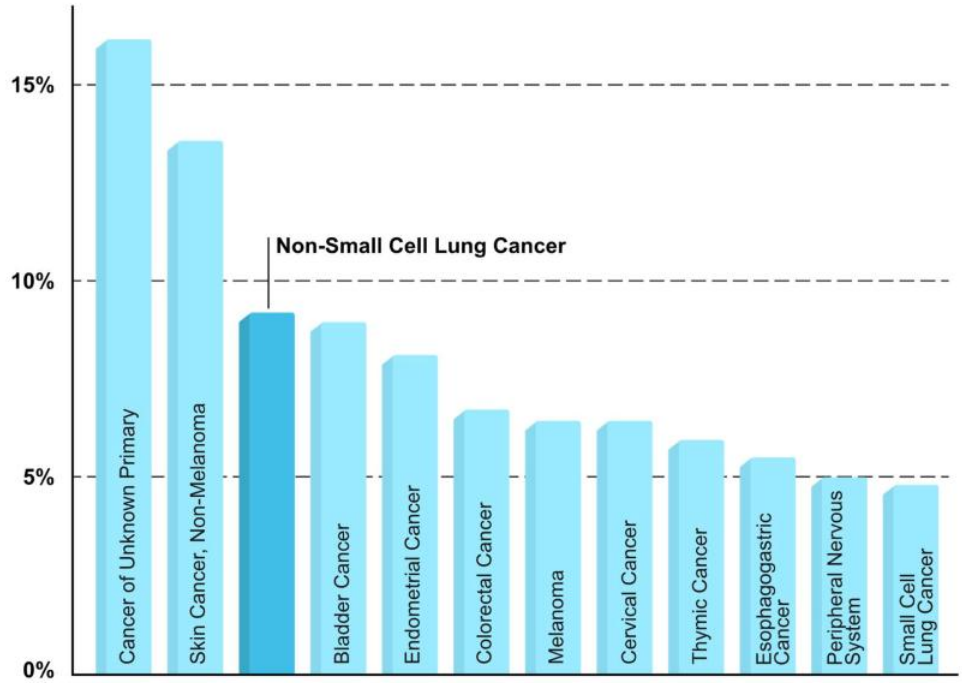
*LY4050784

Selective SMARCA2 Inhibition: Promising Strategy to Exploit Synthetic Lethal Relationship Between SMARCA2 and Mutant SMARCA4



Precision medicine targeting synthetic lethal relationships is a proven clinical approach now used in multiple cancers (e.g., PARP inhibitors)

SMARCA4 is Mutated in Up to 10% of NSCLC; Up to 5% of Solid Tumors



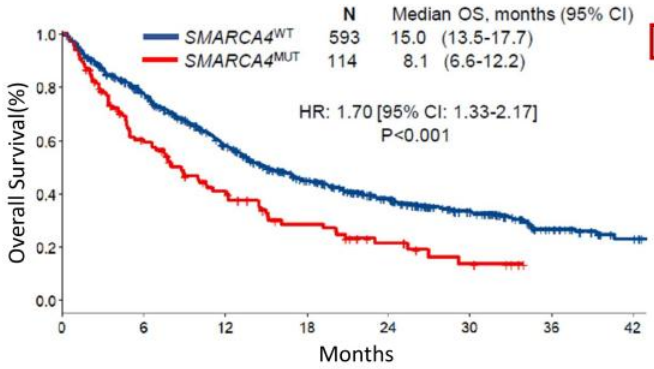
SMARCA4 mutated acr
a broad range of tumor

Accounts for ~5% of solid tumors

AACR GENIE via cBioPortal

Patients With NSCLC Harboring SMARCA4 Mutations Have Significantly Worse Clinical Outcomes and Define a High Unmet Need Patient Population

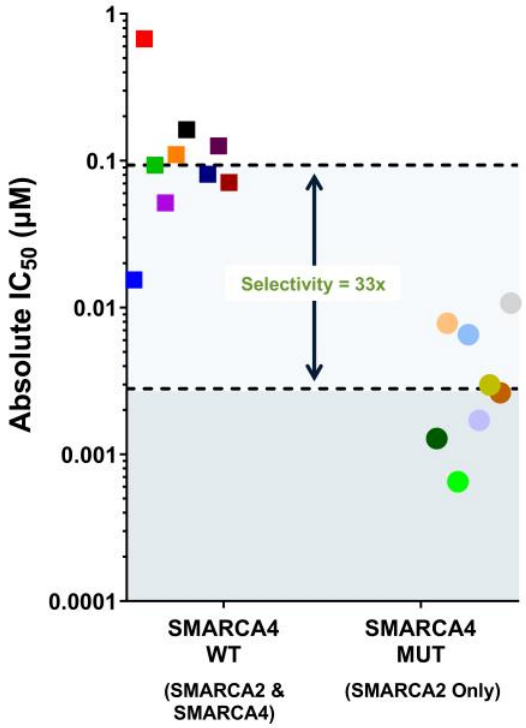
Overall Survival for SMARCA4wt vs SMARCA4mut[1]

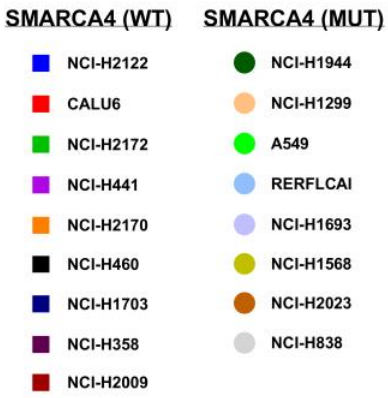


SMARCA4 Mutated in Up to 10% of NSCLC Tumors, Minimal Overlap With Other Mutations[2]

1. Alessi JV, et al., 2021; 2. TCGA via cBioPortal

FHD-909 Demonstrated Approximately 33-fold Selectivity Across 17 SMARCA4 Mutant and Wild-Type Cell Lines *In Vivo*





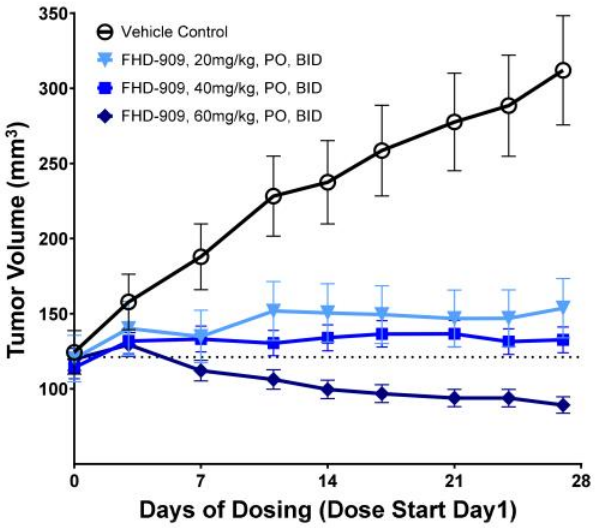
	Median IC_{50} (µM)
SMARCA4 WT	0.0932
SMARCA4 MUT	0.0028

Spread in potency for w type versus mutant cell indicates

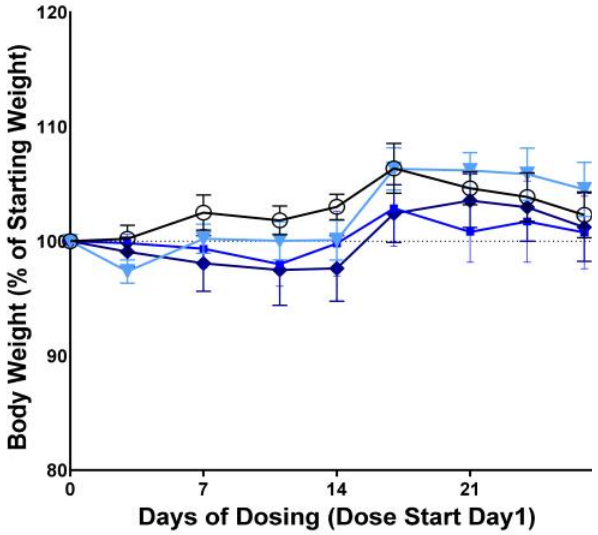
33-fold selectivity observed

FHD-909 Monotherapy Demonstrated Regression *In Vivo* in H2126 SMARCA4 Mutant NSCLC Model and Was Well Tolerated

H2126 Reduction in Tumor Volume



H2126 Body Weight

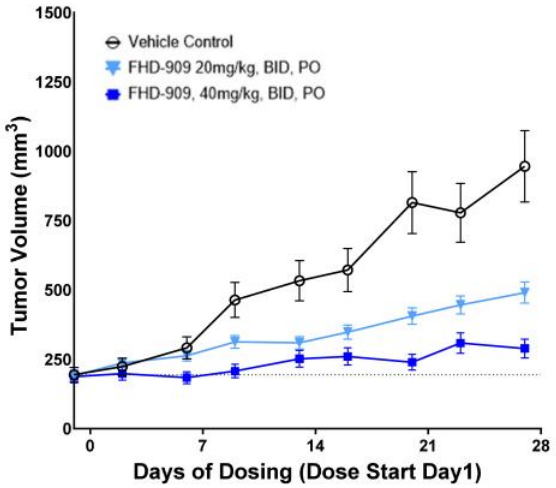


Genetic Background: SMARCA4 W764R, TP53 E62*, STK11-/-, CDKN2A-/-, KEAP1 R272C

NOTE: All doses were well tolerated. Dosing holidays were applied at the high dose, as appropriate.

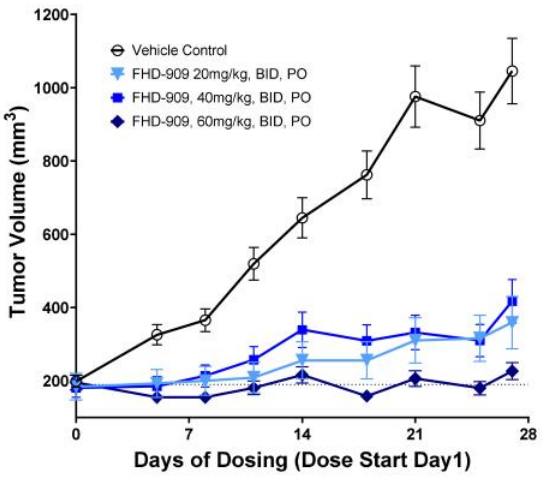
FHD-909 Monotherapy Demonstrated 96% TGI in A549 and Tumor Stasis in RERF-LC-AI Mutant NSCLC Models

A549 Model



Genetic Background: SMARCA4, Q729fs / H736Y, KRAS G12S, STK11-/-, CDKN2A-/-, KEAP1 G333C

RERF-LC-AI Model

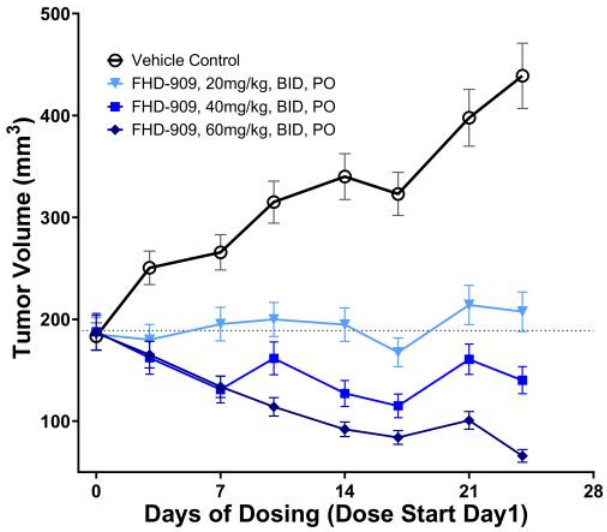


Genetic Background: SMARCA4 mut p.E1496*, TP5 p.Q104*, NF1 p.E1699*

NOTE: All doses were well tolerated. Dosing holidays were applied at the high dose, as appropriate.

FHD-909 Monotherapy Demonstrated Regression in H1793 SMARCA4 Mutant NSCLC Model

H1793 Model



Genetic Background: SMARCA4, E514*, TP53 R209* R273H, ARID1A C884*

- **FHD-909** delivered across range of SMARCA4 mut xenograft models
- Results ranging from impressive TGI to regression as monotherapy
- All doses across all four models were well tolerated

NOTE: All doses were well tolerated. Dosing holidays were applied at the high dose, as appropriate.

FHD-909 (LY4050784) Trial Design

Dose Escalation

- Restricted to SMARCA4 mutated tumors
- SMARCA4 mutant status confirmed by standard NGS panel
- Further enrichment for NSCLC patients as trial progresses
- Tumor histology agnostic
- Enrolling in US and Japan

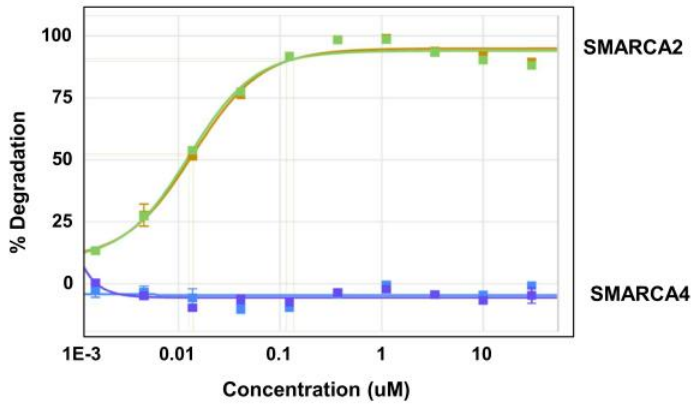
Dose Expansion

- Arm 1: SMARCA4 mutant NSCLC
- Arm 2: Other SMARCA4 mutant tumors (e.g., bladder, endometrial, colorectal)
- Potential for combination arm(s)

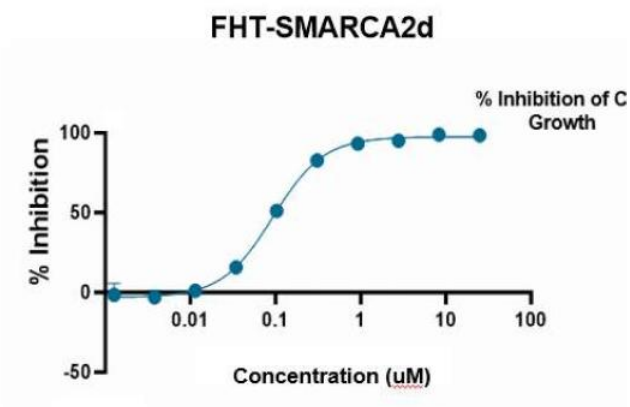
FHD-909 preclinical combination data with pembrolizumab, KRAS inhibitors and chemo to be presented at the AACR Annual Meeting, April 2025

Selective SMARCA2 Degrader Achieved Complete SMARCA2 Degradation and Cell Growth Inhibition *In Vitro*

SMARCA2 / SMARCA4 HIBIT Data



A549 Ten-Day Proliferation Assay



Degraders Caused Time- and Dose-Dependent SMARCA2 Degradation
Antiproliferative Effects in A549 Mutant NSCLC Model

NOTE: Data as of Q4 2021.

Selective CBP Protein Degrader

For EP300 Mutated Cancers

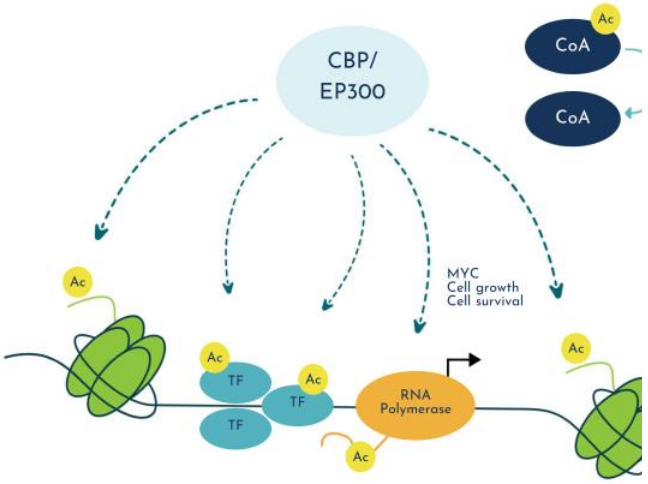
CBP and EP300 Proteins – A Decades Long Challenge in Selectivity

CBP and EP300 Biology

- CBP and EP300 are highly homologous, paralog histone acetyltransferases regulating enhancer-mediated transcription and protein stability
- Dysregulation of CBP and EP300 has been implicated in multiple cancers
- Dual targeting has revealed tolerability and safety issues

Foghorn's Solution... Highly Selective Degradation

- Achieved selective targeting which results in improved tolerability and efficacy
- Advancing two separate programs with defined dependencies and patient populations



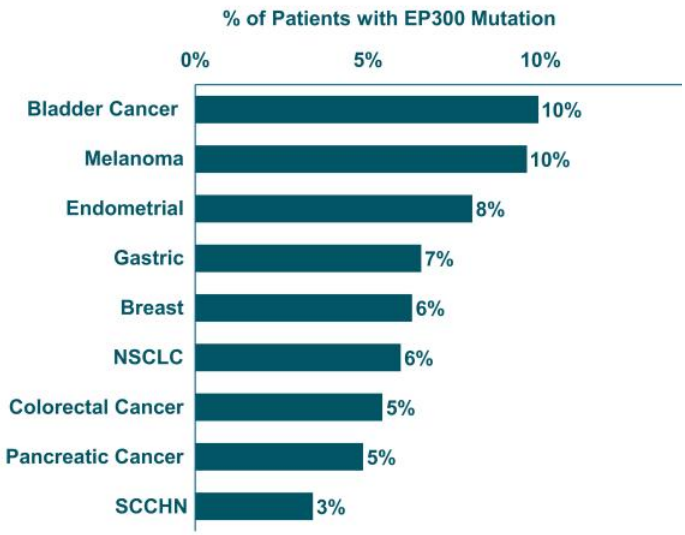
EP300 Degrader Approach
Focus on EP300 Lineage Dependent Cancers

CBP Degrader Appr
Focus on EP300 Mu
Cancers via Synthetic L

Summary: Selective CBP Protein Degrader for EP300 Mutated Cancers

Target / Approach	• CREB binding protein (CBP) • Targeted protein degrader
Initial Indication	• EP300 mutated cancers (e.g., subsets of bladder, colorectal, breast, gastric and lung cancers)
Mutation / Aberration	• EP300 mutated cancers
Stage / Next Milestone	• Preclinical • IND planned for 2026
New Patients Impacted / Year*	• Up to 10% of patients have an EP300 mutation across solid tumors representing ~ 100K addressable population
Key Differentiation	• Highly selective and potent • Increased tolerability relative to non-selective compounds • Long-acting formulation targets Q2-4W dosing • Compelling combination potential

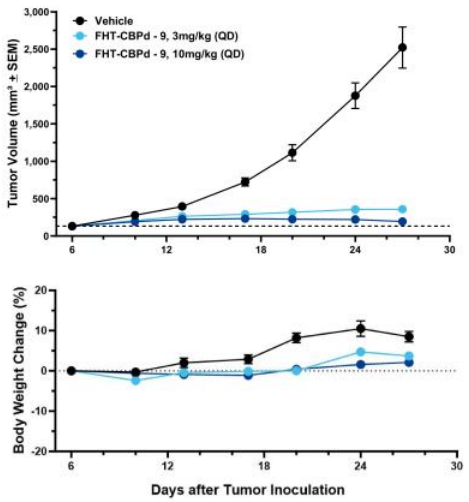
Commercial Opportunity



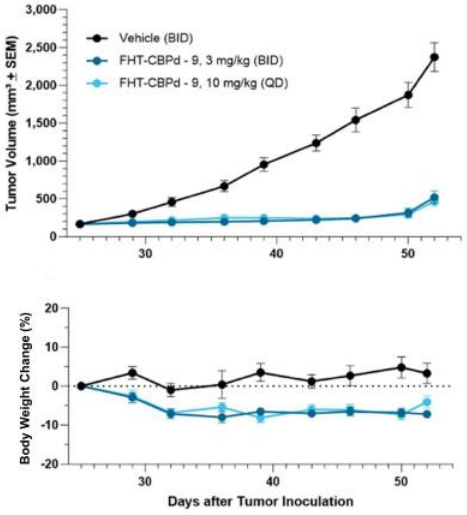
** Per year incidence in the U.S., EU5, Japan . Source: Clarivate DRG Mature Markets Data.*

Selective CBP Degradation Results in Significant Anti-Tumor Activity in EP300mut Solid Tumor Models

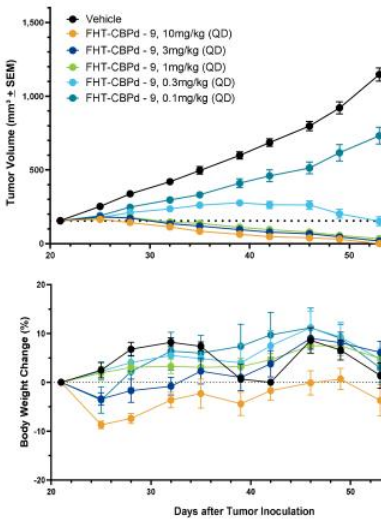
EP300mut Lung Cancer CDX (LU99)



EP300mut Bladder Cancer CDX (639V)



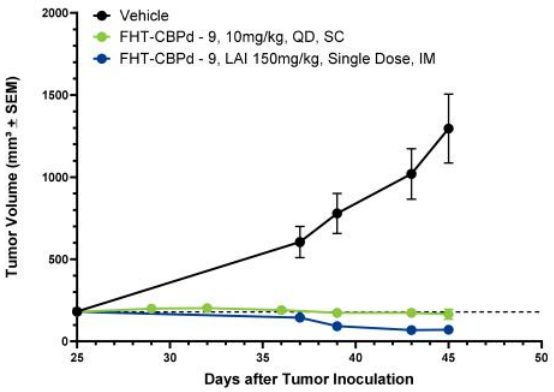
EP300mut Gastric Cancer CD



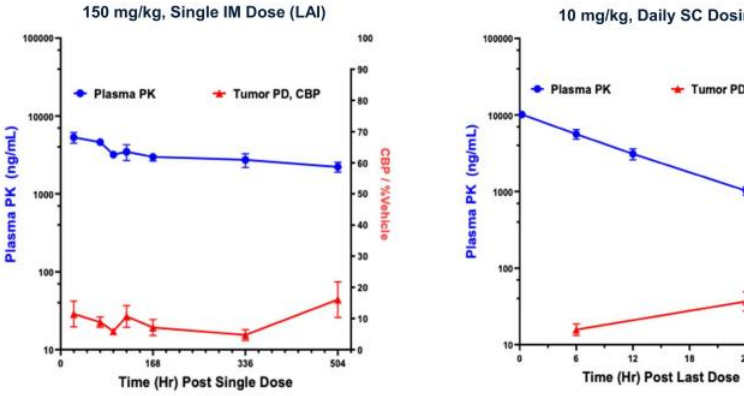
Degrader Selectivity	CBP	EP300	Fold Selectivity
FHT-CBPd-9 DC_{50} *@24h*	0.05 nM	64 nM	>1000x

Long-Acting Injectable (LAI) Formulation Provides Sustained Target Coverage, Anti-Tumor Activity and Tolerability With a Single Injection

EP300mut (639V) Bladder CDX
(Daily Dosing and Single Injection of LAI Formulation)



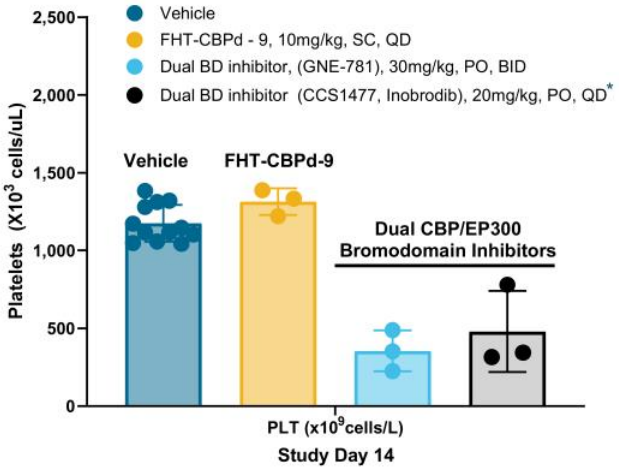
Plasma PK and Tumor PD (CBP) Measured From Bladder CDX Efficacy



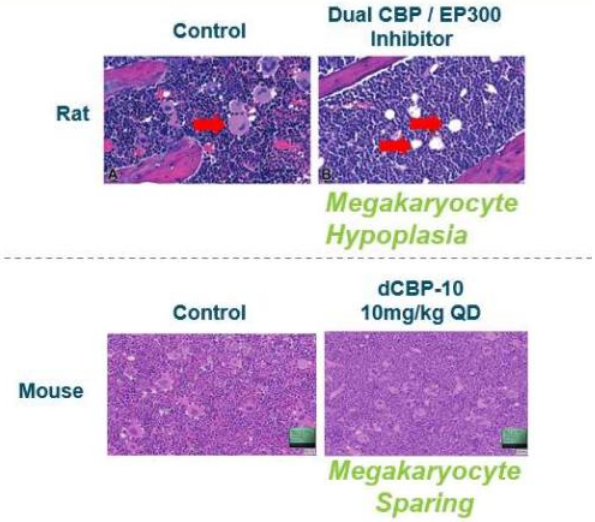
- Single injection of FHT-CBPd-9 in a LAI resulted in sustained plasma exposure (PK) and CBP degradation (PD) throughout the 3-week experiment, ultimately leading to tumor regressions
- No thrombocytopenia was observed through end of study

Preclinical Studies Indicate Selective CBP Degradation Did Not Show Thrombocytopenia and Spared Megakaryocytes *In Vivo*

Platelet Counts Post Two Weeks of Dosing
(In Vivo Mice Models*)*



Bone Marrow Staining
(In Vivo)



**Dual CBP/EP300 inhibition study used 3 weeks of dosing.*

Selective EP300 Protein Degrader

For CBP Mutated and EP300 Dependent Cancers

Summary: Selective EP300 Protein Degrader for CBP Mutant & EP300 Dependent Cancers

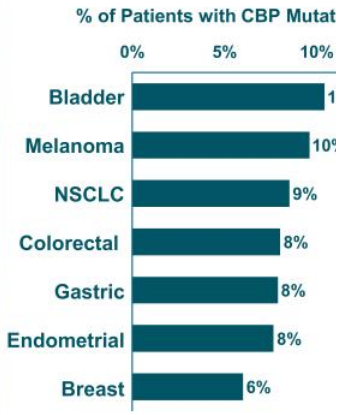
Target / Approach	• E1A binding protein p300 (EP300) • Targeted protein degrader
Indications	• AR+ Prostate • Broad range of heme malignancies • Bladder, melanoma, others
Mutation / Aberration	• EP300 dependent cancers • CBP mutant cancers
Stage / Next Milestone	• Preclinical • IND planned for 2026
New Patients Impacted / Year*	• Over 100,000
Key Differentiation	• Deeper efficacy response with selective degrader vs non-selective molecules • Improved tolerability profile vs non-selective molecules • Patient selection biomarker for Diffuse Large B-Cell Lymphoma (DLBCL)

Commercial Opportunity

EP300 Dependent Cancers

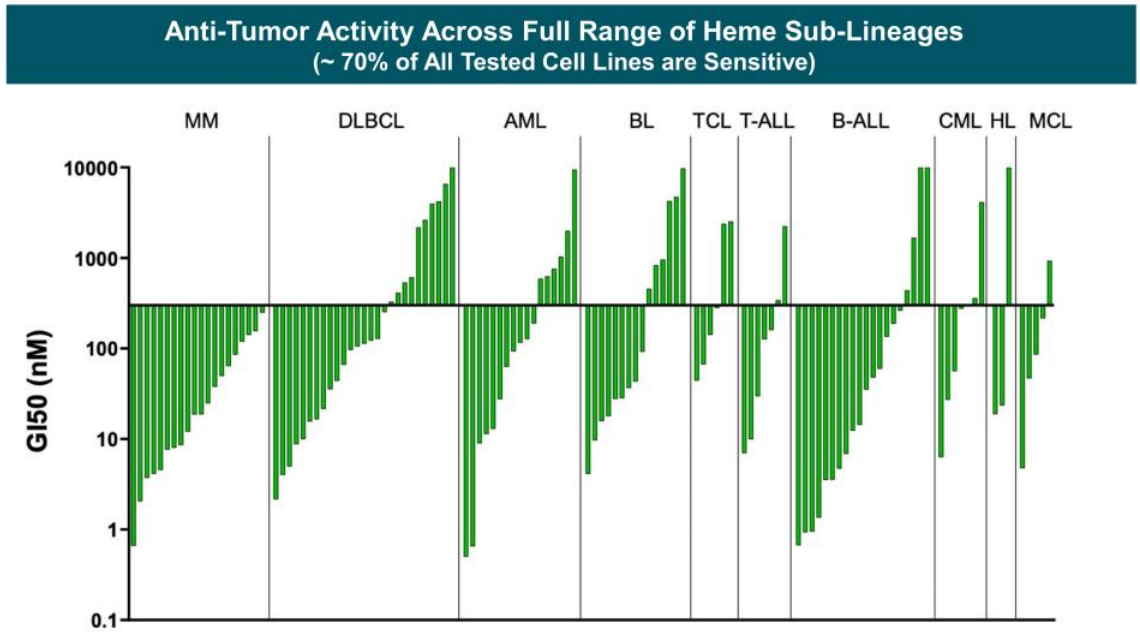
- **Solid Tumors**
  - AR+ mCRPC
  - HR+ breast
- **Hematologic malignancies**
  - DLBCL
  - Multiple Myeloma

CBP Mutant Cancers



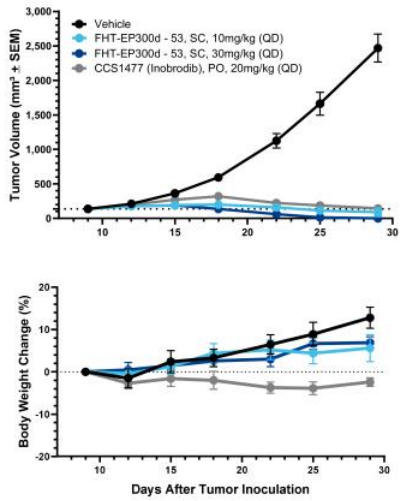
** Per year incidence in the U.S., EU5, Japan. Source: Clarivate DRG Mature Markets Data.*

EP300 Degradation Shows Anti-Proliferative Activity in Broad Range of Hematological Malignancies



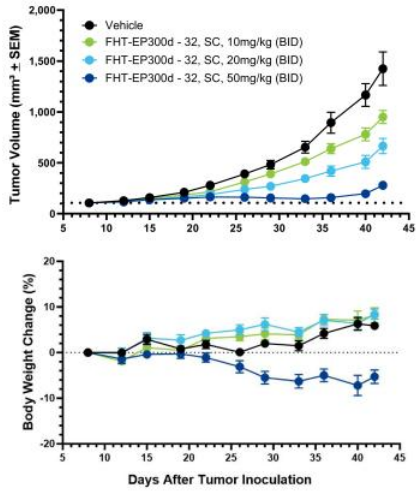
EP300 Degradation Results in Significant Tumor Growth Inhibition in Multiple Myeloma, DLBCL and Prostate Models

Multiple Myeloma (MM1S) CDX



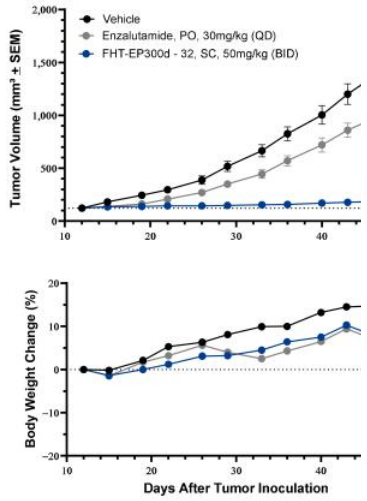
Degrader Selectivity	CBP	EP300	Fold Selectivity
FHT-EP300d-53 DC_{50} *@24h*	>1 uM	0.7 nM	>1000x

DLBCL (KARPAS422) CDX



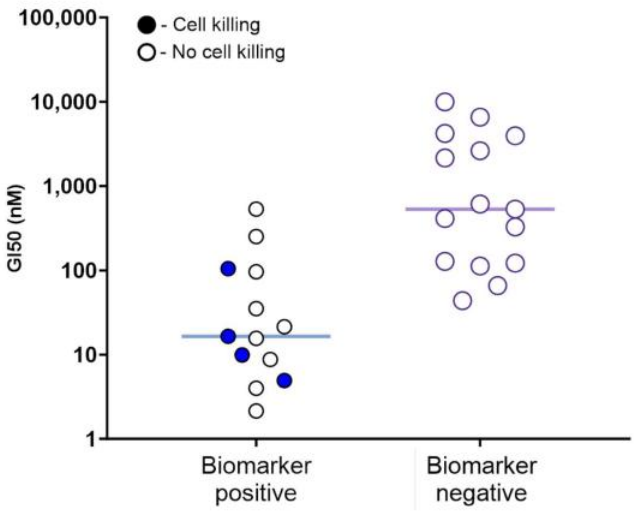
Degrader Selectivity	CBP	EP300	Fold Selectivity
FHT-EP300d-32 DC_{50} *@24h*	>1 uM	23 nM	>40x

AR+ Prostate (VCaP) CDX



Identification of a Patient Selection Biomarker in DLBCL

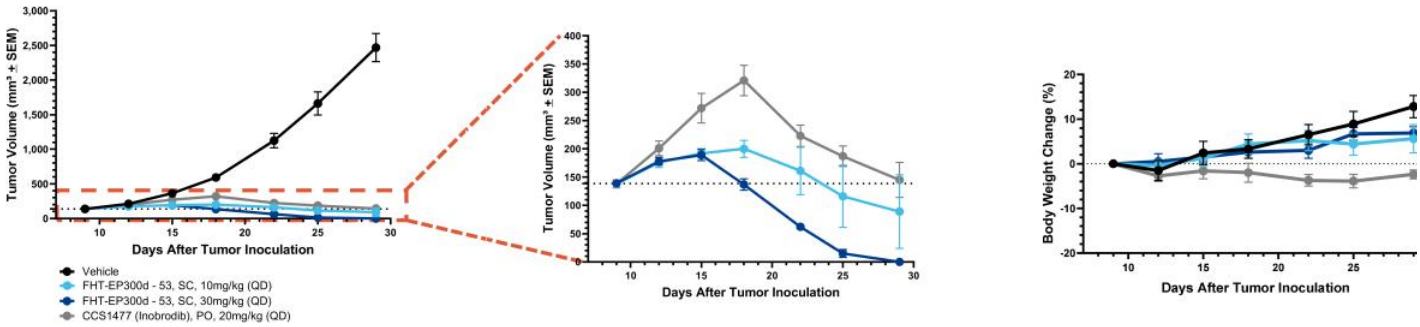
In Vitro Anti-Proliferation Screen Using Selective EP300 Degrader



- Screened 27 DLBCL cell lines; ~60% are sensitive
- Two-step biomarker of sensitivity:
  - EP300 present (no high-impact mutations in EP300) and
  - One of two other mutations
- Mechanistic hypothesis being further validated

Selective EP300 Degrader Demonstrated Complete Response (Tumor Regression) in Multiple Myeloma Model

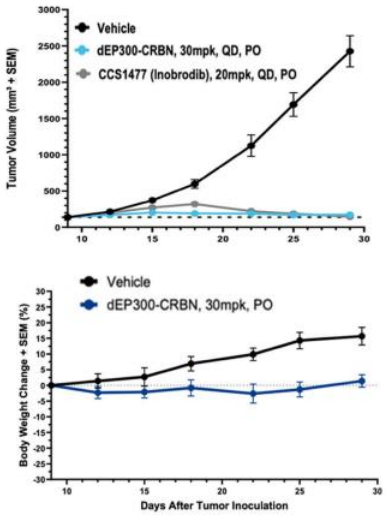
Multiple Myeloma CDX Treated With dEP300 – VHL

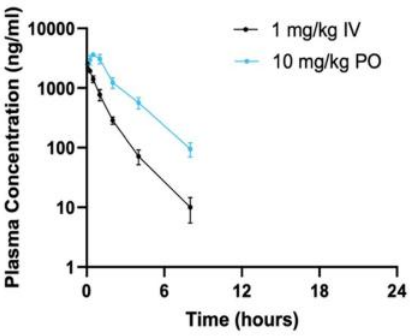


- **Non-selective dual CBP/EP300 inhibitor** shows tumor stasis, but clinical safety (i.e., **thrombocytopenia**) resulted in dosing holida
- **Selective EP300 degrader** can achieve **deeper responses** (complete tumor regression) with **no thrombocytopenia**
- Selective EP300 degrader with improved therapeutic window enables **sustained target coverage and improved efficacy**

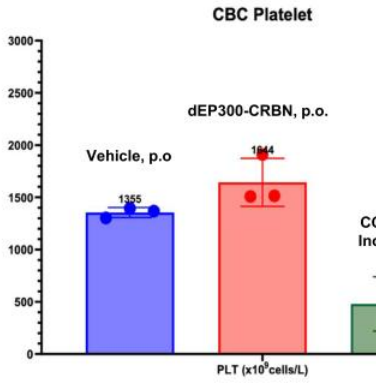
Oral EP300 Selective Degrader Shows Promising Efficacy; Well Tolerated With No Thrombocytopenia

Multiple Myeloma CDX Treated With dEP300 – CRBN





dEP300-CRBN demonstrates 32% oral bioavailability



Degrader Selectivity		CBP	EP300	Fold Selectivity
FHT-EP300d-CRBN	*DC50 @24h*	50 nM	0.2 nM	250x

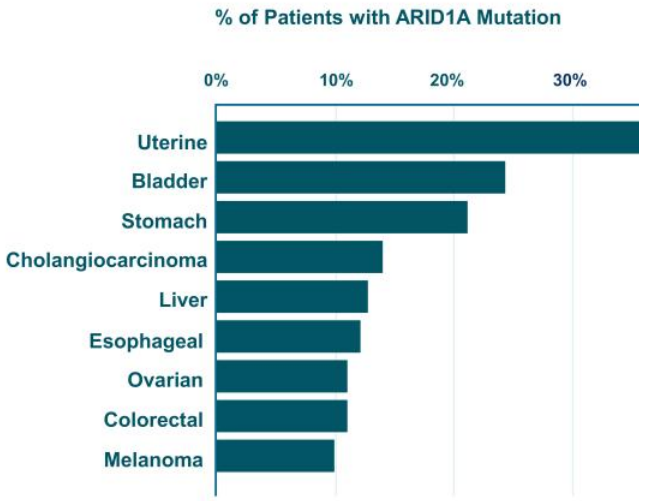
Selective ARID1B Protein Degrader

For ARID1A Mutated Cancers

ARID1B is a Major Synthetic Lethal Target With Potential in Up To 5% of All Solid Tumors; Degradation Achieved, Program Update in 2025

Target / Approach	• ARID1B • Targeted protein degrader
Initial Indication	• ARID1A mutated cancers (e.g. ovarian, endometrial, colorectal, bladder and other cancers)
Mutation / Aberration	• ARID1A mutations
Stage / Next Milestone	• Preclinical • Program update in 2025
New Patients Impacted / Year*	• ARID1A is one of the most mutated protein in cancers (~ 5% of all solid tumors) representing > 175K addressable patients across solid tumors
Key Differentiation	• Multiple ARID1B binders with nM affinity and selectivity • ARID1B degradation achieved

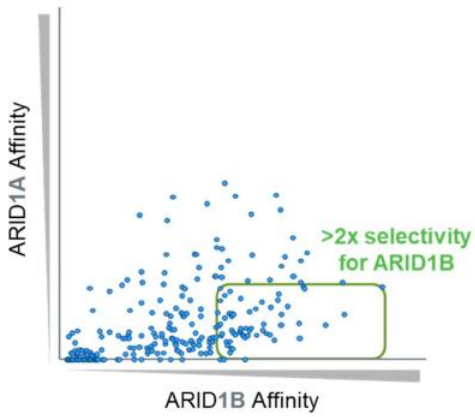
Commercial Opportunity



** Per year incidence in the U.S., EU5, Japan. Source: Clarivate DRG Mature Markets Data.*

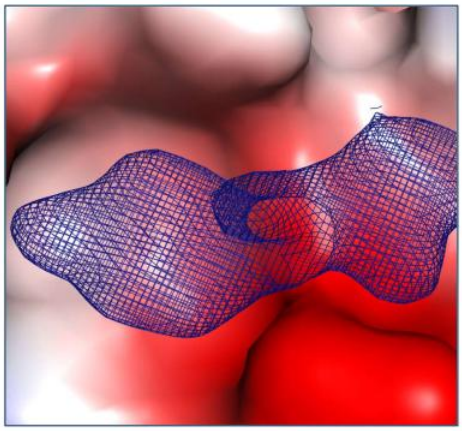
Compound Screening and Structure-Based Optimization Yielded Selecti ARID1B Binders

Identification of Selective ARID1B Binders



- Mapped and purified several potential ligandable regions of ARID, which were then screened against various compound libraries
- Characterized binding using multiple biochemical and biophysical techniques: e.g. DSF, ASMS, NMR, and SPR

X-Ray Crystal Structures Detail Selective ARID1B Binding

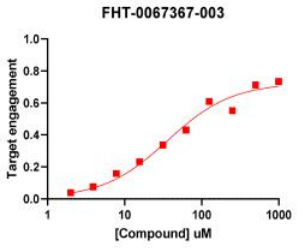


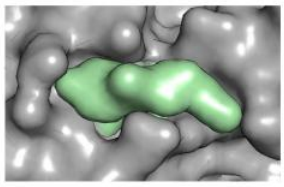
- Determined X-ray crystal structure of ARID ligandable domains with specific binders
- Leveraged these structures to drive binding affinities and expand binding chemotypes

Structure-Based Optimization Drove Improved ARID1B Binding Affinity From 100 uM to Less Than 200 nM

Gen 1: Screening Hit

ARIDb-1
ARID1B Kd: **100 uM**

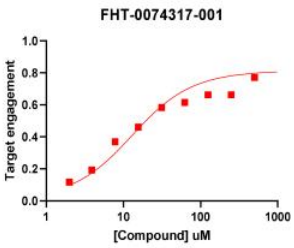


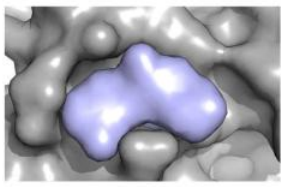


1.4 Å co-xtal structure

Gen 2: Early Optimization

ARIDb-2
ARID1B Kd: **15 uM**

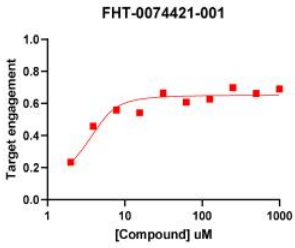


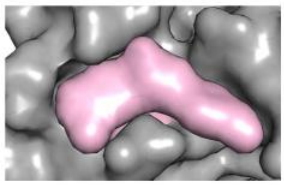


2.0 Å soak structure

Gen 3: Sub-uM Affinity

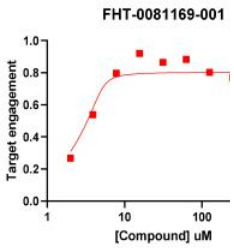
ARIDb-3
ARID1B Kd: **0.5 uM**

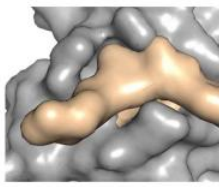




1.9 Å co-xtal structure

ARIDb-9
ARID1B Kd: **0.2**



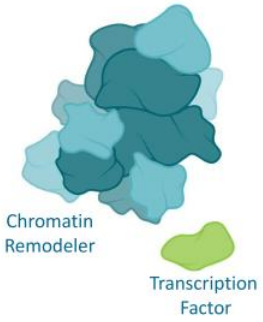


1.7 Å soak structure

Drugging Transcription Factors

Multiple Approaches

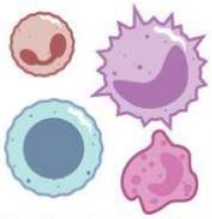
Foghorn's Transcription Factor Platform: 4 Approaches



Modulating TF activity via Protein-Protein Interaction Disruptors

Degrading TFs via recruitment of the native E3 ligase using molecular glue

= MG (Molecular

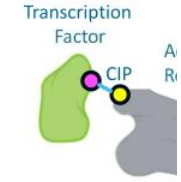


Disease relevant cell lines
(change image – copyright?)

Transcription Factor

Leveraging Phenotypic Screening to reduce TF levels in disease relevant settings

Re-wiring transcription through Induced Proximity with TFs



= CIP (Chemi
of Proximity)

Multiple Near-Term Value Inflection Points Through 2026

FHD-909 (LY4050784) (Selective SMARCA2 Inhibitor)	Preclinical Combination Data	April 2025 (AACR)
	Phase 1 Dose Escalation Data	Confidential
Selective SMARCA2 Degrader	IND / Phase 1 Initiation	Confidential
Selective CBP Degrader	IND / Phase 1 Initiation	2026
Lilly Target #2	Target Disclosure and IND	Confidential
Selective EP300 Degrader	IND / Phase 1 Initiation	2026
Selective ARID1B Degrader	Selective Degradation Achieved	Program Update 2025

SMARCA2 = BRM
SMARCA4 = BRG1

Developing First-in-Class Precision Medicines Targeting Major Unmet Needs in Cancer

Leader in Unique Area of Cancer Biology

Foghorn is a **leader in targeting chromatin biology,** which has the potential to address underlying dependencies of many genetically defined cancers

Platform with initial focus in oncology, **therapeutic area expansion potential**

Large Market Potential

Chromatin biology is implicated in up to **50% of tumors**, potentially impacting **~2.5 million patients**

Foghorn's current pipeline potentially addresses **more than 500,000** of these patients

Broad pipeline across a range of targets and small molecule modalities

Well-Funded

$243.8 million in cash and equivalents (unaudited)

(as of 12/31/2024)

Cash runway into 2027

Shares outstanding: approximately 62.5M*

Value Drivers

Selective SMARCA2 Inhibitor, FHD-909, partnered with Lilly, in **Phase 1 trial**

Advancement of preclinical assets (Selective SMARCA2 Degrader, CBP, EP300, ARID1B) towards INDs

Protein degrader platform with expansion into induced proximity

Major Strate Collaborati

Strategic collaborat Lilly; **$380 million** 50/50 U.S. econ split on two lead pr

**Includes common shares outstanding and pre-funded warrants as of 12/31/2024.*

FOGHORN®
THERAPEUTICS

Unique biology

Precision therapeutics

Broad impact

January 2025

Appendix

Lilly Collaboration Validates Foghorn Approach: Significant Upfront and Deal Economics

$380 Million Up-front

$300 million cash

$80 million in Foghorn common stock at a price of $20 per share

50/50 U.S. Economics on Two Programs

50/50 U.S. economic split on SMARCA2-Selective and another undisclosed program

Tiered ex-U.S. royalties starting in the low double-digit range and escalating into the twenties based on revenue levels

Three Undisclosed Discovery Programs

Option to participate in a percentage of the U.S. economics

Tiered ex-U.S. royalties from the mid-single digit to low-double digit range

$1.3 billion in potential milestones