



Improving treatments for
patients with severe and
chronic diseases



Company Presentation
January 2023

Forward looking statements

This presentation contains forward-looking statements that provide our expectations or forecasts of future events such as new product developments and regulatory approvals and financial performance.

Camurus is providing the following cautionary statement. Such forward-looking statements are subject to risks, uncertainties and inaccurate assumptions. This may cause actual results to differ materially from expectations and it may cause any or all of our forward-looking statements here or in other publications to be wrong. Factors that may affect future results include currency exchange rate fluctuations, delay or failure of development projects, loss or expiry of patents, production problems, unexpected contract, patent, breaches or terminations, government-mandated or market-driven price decreases, introduction of competing products, Camurus' ability to successfully market products, exposure to product liability claims and other lawsuits, changes in reimbursement rules and governmental laws and interpretation thereof, and unexpected cost increases.

Camurus undertakes no obligation to update forward-looking statements.

Camurus snapshot



Rapidly growing commercial stage company

Leader in opioid dependence treatment with Buvidal weekly and monthly depots



Advancing late-stage pipeline with blockbuster potential

Prospects for multiple new approvals in coming years in CNS and rare disease indications



Strong financial performance

Entering profitability in 2022



Unique FluidCrystal® technology platform

Commercially validated, with a broad range of applications

LISTED ON NASDAQ STOCKHOLM
TICKER **CAMX**; EMPLOYEES: ~175

Significant recent progress



Strong financial performance

- ✓ High double-digit year-on-year revenue growth
- ✓ Entering profitability in 2022
- ✓ Robust cash position ~SEK 520 m – no debt



Commercialization execution

- ✓ Leader in long-acting opioid dependence treatment in the EU and Australia
- ✓ Strong sales growth supported by an expanding evidence base
- ✓ Further potential through label and geographic expansion

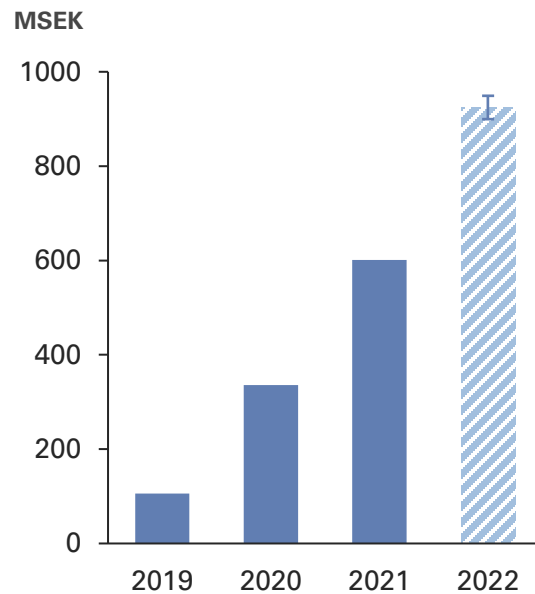


Pipeline advancement

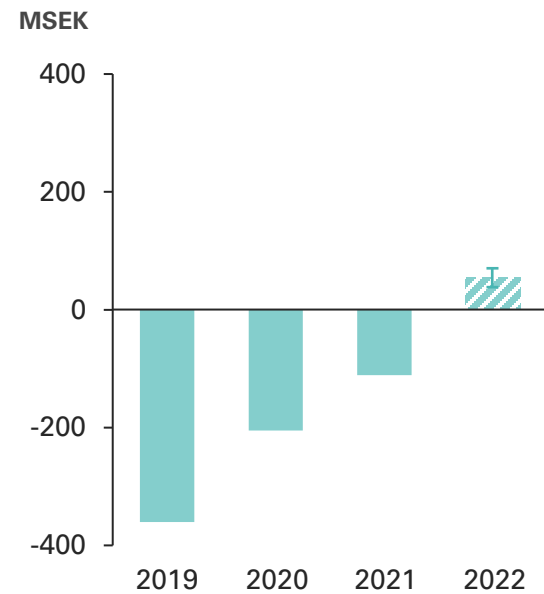
- ✓ Successful life-cycle management
- ✓ Key programs in registration phase
- ✓ Four Phase 3 studies in rare disease indications
- ✓ Significant potential in early-stage programs and technology platform

On track for full year profitability

Revenues¹



Operating results¹



FY 2022 outlook

Total revenue
SEK 900 to 950 million

Product sales
SEK 875 to 925 million

Operating results
SEK 40 to 70 million
(increased from SEK -20 to 40 million)

¹Forecasted 2022 revenue and operating results

Opioid dependence – escalating global health crisis

Largest society burden of all drugs¹

- 61 million opioid users worldwide¹
- Opioid crisis worsened during COVID-19 pandemic

High need for better access to care and new treatment alternatives

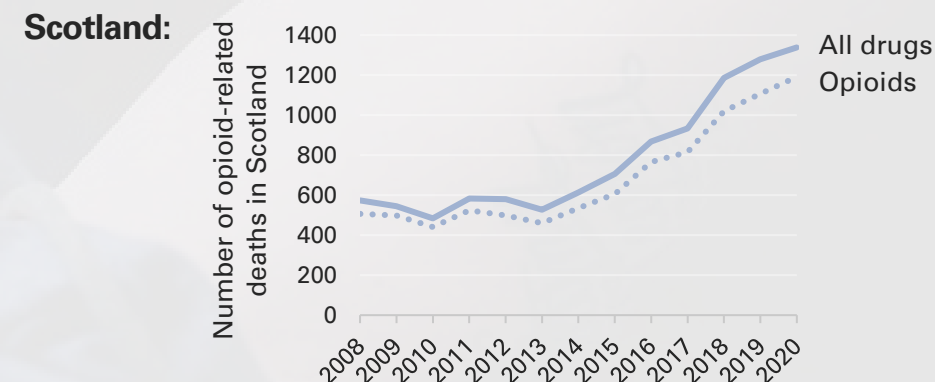
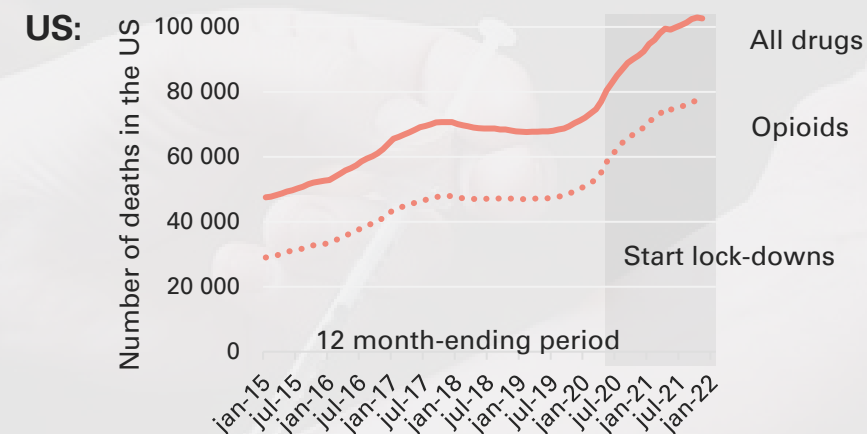
- Long-acting injections a new paradigm in opioid dependence treatment

Significant limitation with current daily medications

- Diversion, misuse, risk of overdose, poor retention, burdens and stigma of daily medications

¹United Nations: World drug report 2022²SAMSHA;³EMCDDA;⁴www.cdc.gov/hchs/nvss/vsrr/drug-overdose-data.htm
⁵<https://www.nrscotland.gov.uk/statistics-and-data/statistics/statistics-by-theme/vital-events/deaths/drug-related-deaths-in-scotland/2020>

Escalating opioid overdose deaths



Buvidal – game changing opioid dependence treatment

Weekly and monthly, subcutaneous buprenorphine for individualized treatment of opioid dependence within a framework of medical, social and psychological treatment in adults and adolescents 16 years or over¹

Demonstrated benefits to patients and society

- Superior treatment outcome and patient satisfaction²⁻⁵
- Blockade of subjective opioid effects from first dose³
- Reduced treatment burden and improved quality of life^{5,6}
- Decreased risk of diversion, misuse and pediatric exposure^{7,8}
- Reduced treatment costs⁹

**“Buvidal became
my way out”**

Justin, Buvidal patient in
Australia

¹ SmPC Buvidal May 2021; ²Lofwall et al. JAMA Int. Med. 2018;178(6): 764-773; ³Walsh et al, JAMA Psychiatry 2017;74(9):894-902; ⁴Frost, M., et al. Addiction. 2019;114(8):1416-1426. doi: 10.1111/add.14636; ⁵Lintzeris, N., et al. JAMA Network Open. 2021;4(5):e219041. doi:10.1001/jamanetworkopen.2021.9041; ⁶Barnett et al Drug and Alcohol Dependence 2021; <https://doi.org/10.1016/j.drugalcdep.2021.108959>; ⁷EPAR for Buvidal; ⁸Dunlop, A. J., et al. Addiction. 2021. <https://doi.org/10.1111/add.15627>; ⁹Dunlop, A. Oral presentation at CPDD June 2020.

Buvidal sales growth underscores potential

Leadership in opioid dependence treatment

- High double-digit year-on-year sales growth
- Buvidal available in 18 countries in Europe, Australia and the Middle East
- Est. >32,000 patients in treatment at the end of Q3
- Passed milestone of >1 million sold Buvidal units since launch

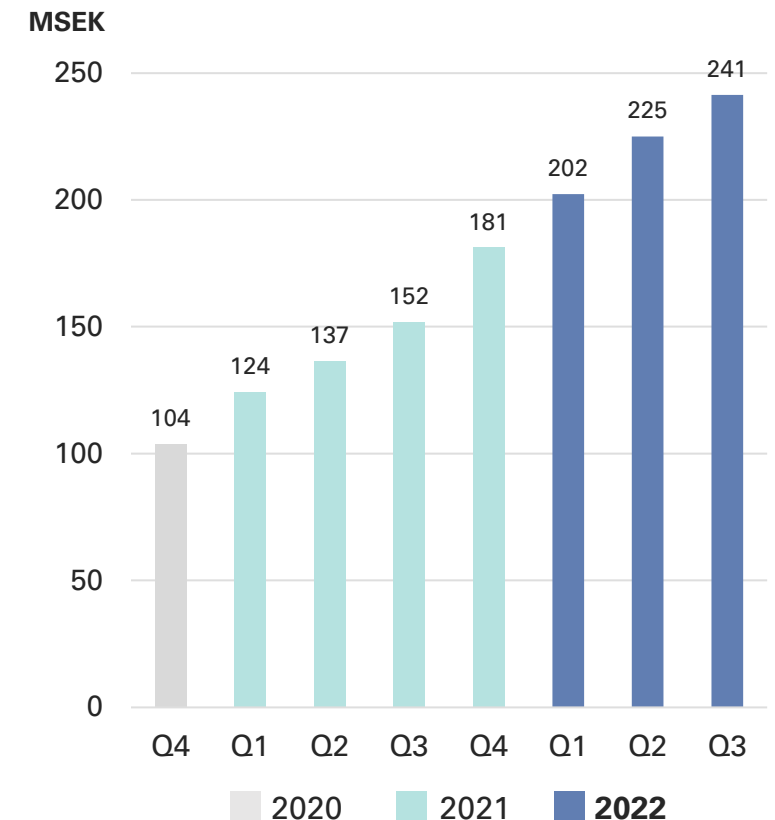
Significant additional potential in geographic expansion

- Recent market approvals for Buvidal in Egypt and Saudi Arabia
- Tentative approval of Brixadi¹ in the US. NDA resubmitted by Braeburn for final approval
- Additional five national regulatory applications by Camurus under review

Indication expansion to chronic pain

- Market authorization applications under review in EU and Australia

Quarterly product sales



¹Product rights to Brixadi™ (CAM2038) in North America licensed to Braeburn Inc.

Substantial growth potential in MENA

Large unmet need¹

- Large untreated populations
- Future challenging problems to MENA countries
- Estimated >3 million opioid users² and 0.3 – 1 million injection drug users³

Buvidal in MENA

- Early access programs three markets
- New approvals in Saudi Arabia and Egypt – **first approved treatment of opioid dependence**
- Significant untreated populations



Market expansion to the US

Brixadi¹ NDA status

- ✓ Tentative NDA approval of Brixadi by FDA
- ✓ Resubmission of Brixadi NDA on 23 November 2022 by Camurus' US licensee Braeburn
- ✓ NDA acceptance with PDUFA date 23 May 2023
- ❑ Brixadi NDA approval decision

High unmet medical need and market potential

- US opioid crisis continues with est. 80,000 overdose deaths annually
- Long-acting injectable (LAI) market size ~US\$ 800m with ~3-4% patient share²
- LAI buprenorphine market growth ~70% YoY

Ongoing US investigator sponsored studies

STUDY	EST. PATIENTS
MOMs³ Medication Treatment for OUD in Expectant Mothers	200 (Brixadi 100)
ED-INNOVATION⁴ Emergency Department-Initiated buprenorphine and Validation Network Trial	2000 (Brixadi 1000)
RDD⁵ Optimizing Retention, Duration and Discontinuation Strategies for Opioid Use Disorder Pharmacotherapy	2630 (Brixadi 650)
FRIENDS⁶ A comparative effectiveness trial of XR Naltrexone vs XR-BUP with individuals leaving jail	240 (Brixadi 120)
EXHIT ENTRE Exemplar Hospital Initiation Trial to Enhance Treatment Engagement	314 (Brixadi 157)

¹BrixadiTM is the US trade name for Buvidal[®]; ²Company estimates based on Indivior and Alkermes sales data.

³Winhusen et al. *Contemp Clin Trials*. 2020; 93:106014. <https://doi.org/10.1016/j.cct.2020.106014>;

⁴D'Onofrio et al. *Contemp Clin Trials*. 2021; 104:106359. <https://doi.org/10.1016/j.cct.2021.106359>;

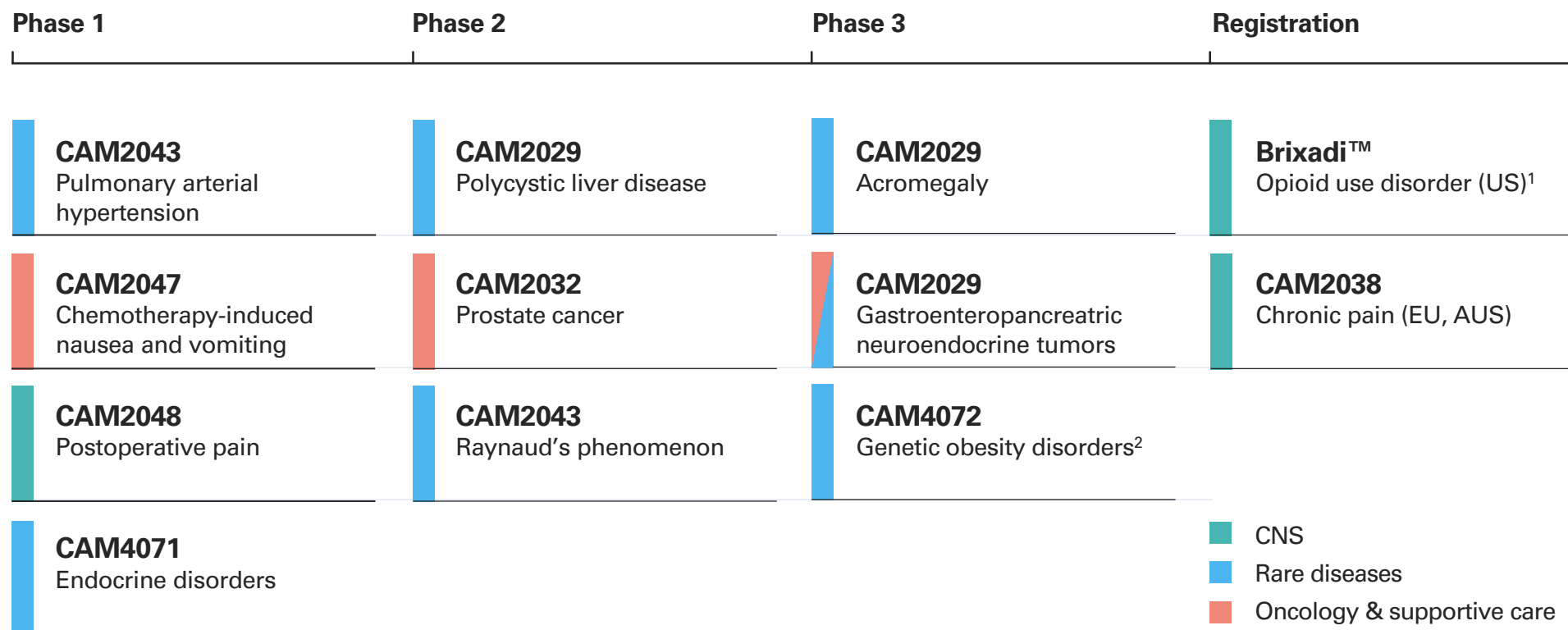
⁵Schulman et al. *Addict Sci Clin Pract*. 2021; 16: 15. <https://doi.org/10.1186/s13722-021-00223-z>;

⁶Gordon et al. *J Subst Abuse Treat*. 2021; 128:108241. <https://doi.org/10.1016/j.jsat.2020.108241>

Brixadi well positioned against competition

Long-acting injectables features	ONCE-MONTHLY 		Weekly/Monthly 
Weekly dosing	—	—	✓
Monthly dosing	✓	✓	✓
Multiple doses	—	—	✓
Choice of inj. sites	—	—	✓
Smallest needle	(19G)	(20G)	✓ (23G)
Lowest dose volume	0.5–1.5mL	3.4mL	✓ 0.16–0.64mL
Room temp. storage	—	—	✓
Day one initiation	—	—	✓
Clin. Data vs active control*	—	—	✓
Launched	US, CAN, AUS, IL	US	EU, UK, AUS

Broad and diversified mid- to late-stage pipeline



¹Licensed to Braeburn in North America; ²Licensed to Rhythm Pharmaceuticals worldwide

CAM2029 – octreotide subcutaneous depot in Phase 3 development

Octreotide SC depot under assessment in three, serious rare disease indications

- Acromegaly
- Gastroenteropancreatic neuroendocrine tumors (GEP-NET)
- Polycystic liver disease (PLD)

Designed for enhanced efficacy and patient convenience



CAM2029 targeting 3-billion dollar SSA market

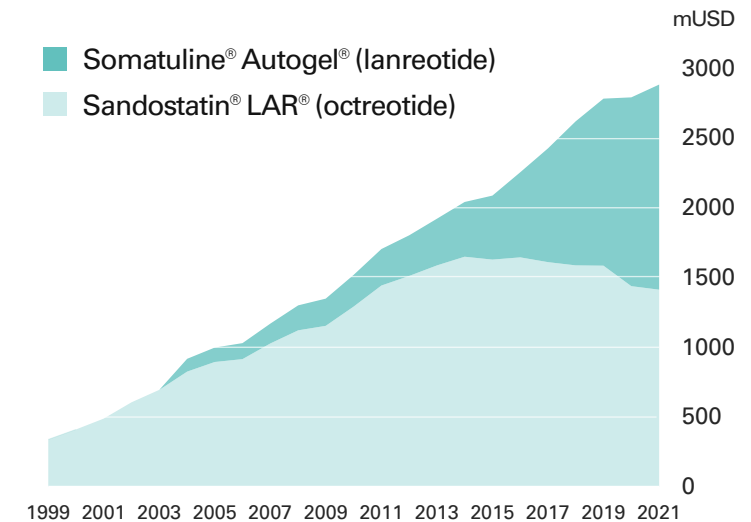
SSAs established treatment with limitations

- First-line treatment of acromegaly and neuroendocrine tumors (NET)
- Established safety and efficacy profile
- Potential for significant improvements of efficacy and patient convenience

CAM2029 best-in-class treatment potential

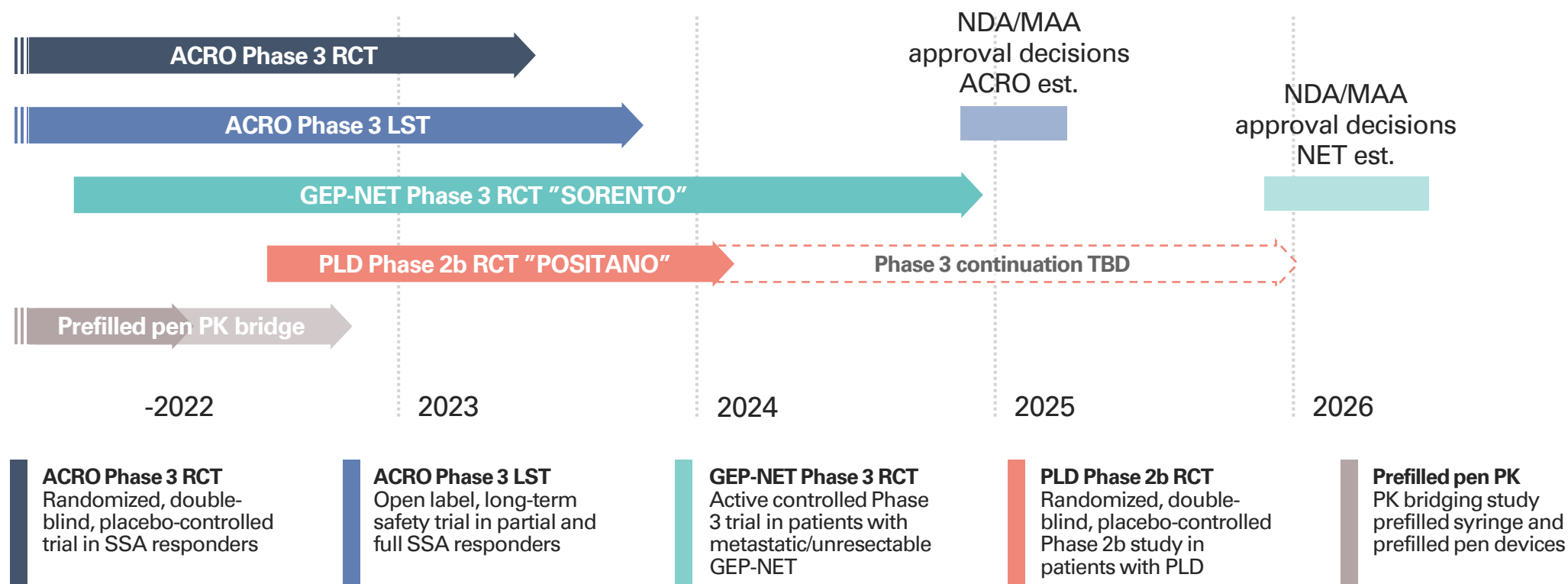
- Convenient self-administration with state-of-the-art pen device
- Enhanced SSA exposure (500% bioavailability increase)
- Potential for improved disease control and treatment outcomes

Annual sales of first generation SSAs¹



¹GlobalData 2022

CAM2029 extensive clinical program



Timelines are indicative. PK – pharmacokinetic; PD – pharmacodynamic; RCT – Randomized control trial; LST – Long-term safety trial; ACRO – acromegaly, GEP-NET – gastroenteropancreatic neuroendocrine tumors; PLD – polycystic liver disease; OLE – open label extension

Phase 3 acromegaly program under completion

Pivotal randomized, placebo-controlled Phase 3 trial

- Rigorous, 24-week, randomized, double-blind, placebo-controlled trial
- Primary endpoint biochemical response ($\text{IGF-1} \leq 1 \times \text{ULN}$)
- Filling regulatory requirement for efficacy

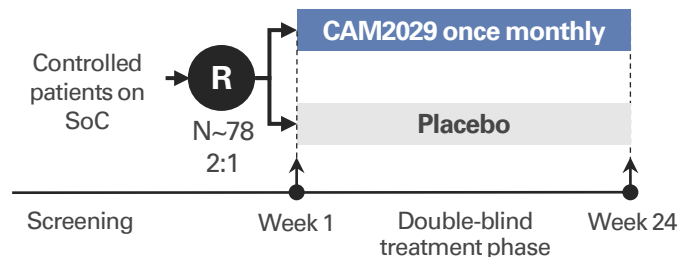
Long-term safety Phase 3 trial

- 52-week long-term safety, switch and extension trial
- Endpoints include safety (primary), IGF-1, GH and PROs (QoL)
- Filling regulatory requirements for safety exposure

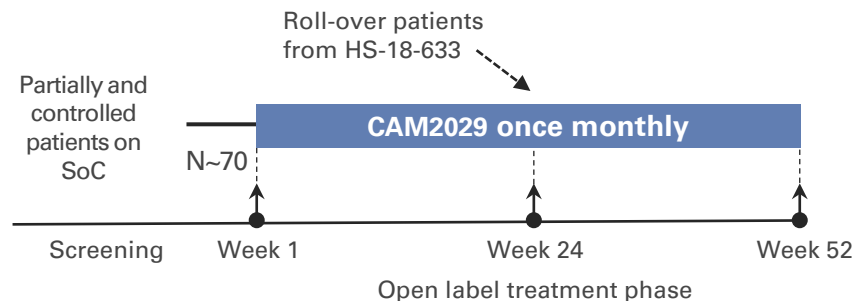


- ✓ Two Phase 3 trials under completion
- ✓ Recruitment completed in pivotal trial
- ✓ Long-term safety trial extended with additional 12-month period
- ❑ **Phase 3 efficacy results July 2023**
- ❑ **Phase 3 long-term results H2 2023**
- ❑ Est. NDA and MAA submissions 2023/24

AcrolInnova 1



AcrolInnova 2

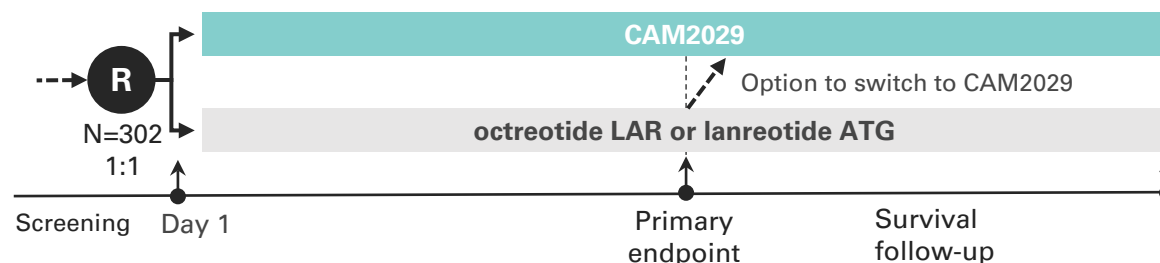


SORENTO largest RCT in NET

Randomized, active-controlled Phase 3 trial

- Primary endpoint is superiority in progression free survival, PFS, vs. standard of care (first-line medical treatment)
- Assessed after 194 progression events
- Multiple patient reported outcomes included in study
- Single, large trial fulfilling regulatory requirements for safety and efficacy
- Broad GEP-NET population of grade 1 to grade 3

SORENTO



PFS – progression free survival; LAR – long-acting release; GEP-NET – gastroenteropancreatic neuroendocrine tumors

SORENTO™

Subcutaneous Octreotide Randomized
Efficacy in Neuroendocrine Tumors

- ✓ SORENTO Phase 3 trial ongoing
- ✓ >30% patients enrolled
- ❑ Enrollment completion est. mid-2023
- ❑ Completion SORENTO efficacy part after 194 PFS events
- ❑ **Est. topline results 2024**
- ❑ **NDA/MAA submissions 2025**

POSITANO program for CAM2029 in PLD

Significant unmet need with no approved treatment

- PLD is a rare, genetic and chronic disorder
- Progressive growth of cysts in the liver, can cause severe symptoms
- No approved medical treatment – increased scientific evidence for SSA's
- Estimated ~30,000 patients with symptomatic PLD¹

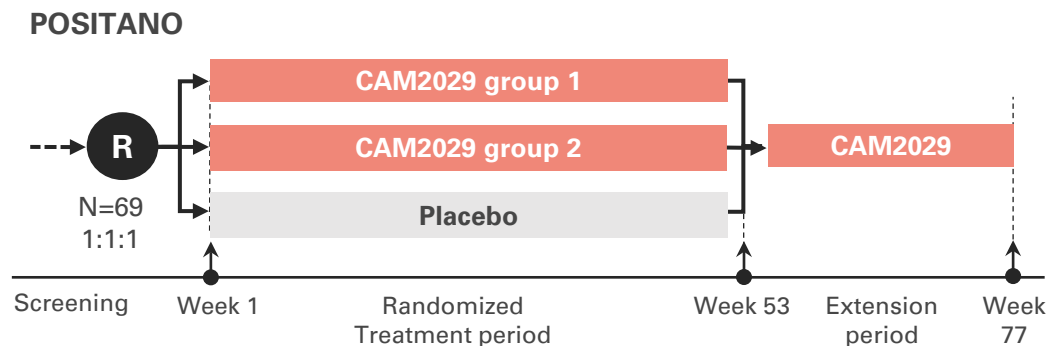
POSITANO trial to assess efficacy and safety

- 52-week randomized, placebo-controlled, three-arm trial
- Primary endpoint is liver volume change
- Key secondary endpoint Camurus' developed PROs, PLD-S

positano™

POlycystic liver Safety and efficacy TriAl
with subcutaNeous Octreotide

- ✓ Orphan drug designation (US)
- ✓ New PROs developed and aligned with FDA
- ✓ Phase 2b trial started June 2022
- ❑ Est. enrollment completion mid-2023
- ❑ **Topline results est. H2 2024**



PLD – polycystic liver disease, SSAs – somatostatin analogues; PRO – patient reported outcome; PLD-S – PLD symptoms
¹Globe Life Science 2020,

Significant market potential for CAM2029

Attractive opportunity

- Highly concentrated target audiences
- Differentiated product properties
- Switch opportunity from established first-line treatments

CAM2029 peak sales estimates from third party market research¹⁻⁴

	TERRITORY	PATIENT POPULATION	EST. PEAK PATIENT SHARE	EST. PEAK SALES
ACRO ¹	EU/AUS	16,500 ⁴	20 – 35%	€30 – 65 million
	US	10,000	25 – 40%	\$150 – 280 million
NET ¹	EU/AUS	68,000 ⁴	30%	€300 – 400 million
	US	37,000	40%	\$1,200 – 1,500 million
PLD ¹	EU/AUS	15-18,000 ⁴	30 – 40%	€80 – 100 million
	US	12-13,000	30 – 40%	\$200 – 300 million

GlobalData report⁵



”Top selling drug to enter the market will be Camurus’ Octreotide LA”

Estimates CAM2029 sales of **US\$210m** US+EU5 sales in 2029 in acromegaly

¹Globe Life Science Aug 2022, data on file; ²Globe Life Science 2020, data on file; ³Assuming €10-12.5k (EU/AUS) and \$60-70K (US) per year net pricing in acromegaly, €15-20k (EU/AUS) and \$80-100K (US) per year net pricing in NET, and €17.5k (EU/AUS) and \$60K (US) per year net pricing in PLD; ⁴Patient numbers extrapolated from 5EU estimates by assuming same prevalence across European countries and Australia

Other rare disease opportunities

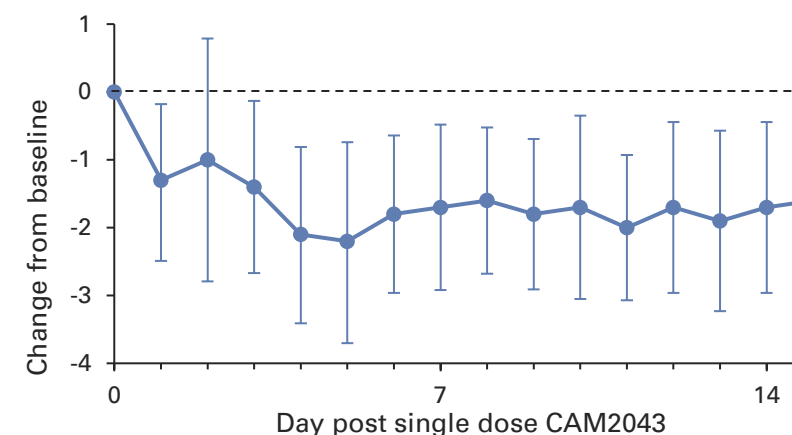
Setmelanotide SC depot, CAM4072

- Developed by license partner Rhythm
- Positive PK and PD results in Phase 2a MAD study
- Phase 3 trial ongoing in switch patients with genetic obesity disease, e.g. Bardet Biedl Syndrome (BBS)
- ☐ Topline Phase 3 results expected in 2023
- ☐ Second Phase 3 trial in naïve patients planned to start in H1 2023
- ☐ Camurus eligible to milestones and royalty payments

Treprostinil SC depot, CAM2043

- Targeting high medical need in treating Raynaud's Phenomenon and PAH
- Recent Phase 2a results indicate efficacy in Raynaud's Phenomenon¹
- ☐ Evaluations of next steps ongoing

Significant change in Raynaud's condition score (95% CI)



¹Camurus' Interim Report Second Quarter 2022. ²Clinical Trial Report HS-18-638, September 2022. PAH – Pulmonary Arterial Hypertension

Preparing for CAM2029 commercialization

Using existing infrastructure in Europe and Australia

- Concentrated audience and significant market potential (est. > \$0.4 billion) across indications^{1,2}
- Established scalable commercial infrastructure

Establishing own US commercial infrastructure

- Single largest market with high market potential (est. > \$1.5 billion) across indications^{1,2}
- Positive payor response to CAM2029 with recognition of benefits and likelihood to be on formularies
- Commercial organization can be built stepwise as new indications are approved
- Medical science and market access functions in 2023
- Full commercial team in 2024 ready for launch

¹Globe Life Science Aug 2022, data on file; ²Globe Life Science 2020, data on file

Key priorities in 2023

-  Grow and strengthen market leading position of Buvidal
-  Expand to new geographies and indications
-  Phase 3 results and NDA preparations for CAM2029
-  Business development and inorganic growth opportunities

Q&A

Camurus AB | Ideon Science Park, SE-223 70 Lund, Sweden
P +46 46 286 57 30 | info@camurus.com | camurus.com



Experienced and committed management team



Fredrik Tiberg, PhD
President & CEO, CSO
In Company since: 2002
Holdings: 1,680,000 shares,
15,000 subscription warrants
& 102,000 employee options

Education: M.Sc. in Chem. Eng., Lund Institute of Technology, PhD and Assoc. Prof. Physical Chemistry, Lund University.
Previous experience: More than 20 years leadership experience from the pharmaceutical industry. Professor Physical Chemistry at Lund University, Sect. Head Institute Surface Chemistry, Visiting Professor at Oxford University



Jon Garay Alonso
Chief Financial Officer
In Company since: 2022
Holdings: 1,450 shares &
57,750 employee options

Education: Bachelor in Business Administration by Universidad Comercial de Deusto. Executive MBA by IESE Business School.
Previous experience: More than 20 years experience from Finance within pharmaceutical and medtech companies, incl. Baxter, Gambro, Convatec, Bristol Myers Squibb.



Maria Lundqvist
Head of Global HR
In Company since: 2021
Holdings: 1,000 subscription
warrants and 38,500
employee options

Education: B.Sc. in Business and Economics, Uppsala University
Previous experience: More than 20 years of experience of leadership roles within Human Resources, including HR Director Nordics at Teva Pharmaceuticals and HR positions at Tetra Pak, Vestas and AstraZeneca.



Richard Jameson
Chief Commercial Officer
In Company since: 2016
Holdings: 29,193 shares, 8,000
subscription warrants and
57,750 employee options

Education: B.Sc. in Applied Biological Sciences from University West of England
Previous experience: General Manager, UK & Nordics for Reckitt Benckiser (2010 – 2013) and Area Director Europe, Middle East and Africa for Indivior (2013 – 2016).



Peter Hjelmström, MD, PhD
Chief Medical Officer
In Company since: 2016
Holdings: 38,500 employee
options

Education: MD, PhD and Assoc. Prof. Karolinska Institutet, Postdoc. Yale University
Previous experience: More than 15 years of experience from the pharmaceutical industry, including as Medical Director at Orexo and Head of Clinical Science at Sobi



Fredrik Joabsson, PhD
Chief Business Dev. Officer
In Company since: 2001
Holdings: 50,070 shares &
38,500 employee options

Education: M.Sc. in Chemistry, PhD in Physical Chemistry, Lund University
Previous experience: More than 20 years of experience in pharmaceutical R&D, business development and alliance management.



Torsten Malmström, PhD
Chief Technical Officer
In Company since: 2013
Holdings: 46,858 shares &
38,500 employee options

Education: M.Sc. in Chemistry, PhD in Inorganic Chemistry, Lund University
Previous experience: More than 20 years of experience from pharmaceutical R&D including Director Pharmaceutical Development at Zealand Pharma, Director of Development at Polypeptide, Team Manager at AstraZeneca.



Annette Mattsson
VP Regulatory Affairs
In Company since: 2017
Holdings: 2,004 shares and
38,500 employee options

Education: Bachelor of Pharmacy, Uppsala University and Business Economics, Lund University
Previous experience: More than 25 years of experience within regulatory affairs, including European RA Director/Global RA Lead at AstraZeneca and Global RA Lead at LEO Pharma.



Agneta Svedberg
VP Clinical & Regulatory Dev.
In Company since: 2015
Holdings: 22,987 shares &
38,500 employee options

Education: M.Sc. In Radiophysics and B.Sc. In Medicine from Lund University, Executive MBA from Executive Foundation Lund
Previous experience: More than 25 years of experience in drug development, incl. as COO at Zealand Pharma, CEO of Cantargia, Senior VP Clinical Development at Genmab.



Markus Johnsson
Senior VP R&D
In Company since: 2003-2017,
2019-
Holdings: 21,000 shares &
23,500 employee options

Education: Ph.D. in physical chemistry and M.Sc. in chemistry from Uppsala University.
Previous experience: More than 20 years of experience from pharmaceutical development and project management

Shareholders and analyst coverage

Shareholders as of 31 December 2022	Number of shares	% of capital	% of votes
Sandberg Development AB	21,875,692	39.5	39.5
Fjärde AP-fonden	3,070,571	5.5	5.52
Didner & Gerge Fonder	2,286,697	4.1	4.1
Avanza Pension	2,067,525	3.7	3.7
Fredrik Tiberg, CEO	1,680,000	3.0	3.0
State Street Bank and Trust	1,130,265	2.0	2.0
JP Morgan Chase Bank	937,574	1.7	1.7
Backahill Utveckling	826,491	1.5	1.5
Svenskt Näringsliv	800,000	1.4	1.4
Lancelot Avalon	625,000	1.1	1.1
Öhman Fonder	587,940	1.1	1.1
Afa Försäkring	564,560	1.0	1.0
Camurus Lipid Research Foundation	495,250	0.9	0.9
COJ Service AB	425,000	0.8	0.8
Handelsbankens fonder	422,249	0.8	0.8
Other shareholders	17,628,229	31.9	31.9
In total	55,383,447	100.0	100.0

Analysts

Carnegie

Erik Hultgård

DNB

Patrik Ling

Handelsbanken

Suzanna Queckbörner

Mattias Häggblom

Jefferies

James Vane-Tempest

Nordea

Viktor Sundberg

Pareto

Peter Östling

Bryan Garnier

Alex Cogut

CAM2029 recent and upcoming milestones

ACRO

- ✓ Two Phase 3 trials ongoing
- ✓ Recruitment completed in pivotal efficacy trial (RCT)
- ❑ **Topline results mid 2023**
- ❑ Long-term safety results H2 2023
- ❑ Est. NDA/MAA submissions 2023/24



NET

- ✓ Phase 3 SORENTA trial ongoing, largest randomized NET study
- ❑ Est. completion of patient recruitment mid 2023
- ❑ **Est. topline results H2 2024** – after 194 PFS events
- ❑ Est. NDA/MAA submissions 2025

SORENTA™

Subcutaneous Octreotide Randomized
Efficacy in Neuroendocrine Tumors

PLD

- ✓ Orphan drug designation (US)
- ✓ New PROs developed and aligned with FDA
- ✓ Phase 2b POSITANO trial ongoing
- ❑ Est. completion of patient enrollment in H1 2023
- ❑ **Topline efficacy results H1 2024**

positano™

Polycystic liver Safety and efficacy Trial
with subcutaneous Octreotide