



Advanced Therapies Opportunities and Challenges

London, November 2017

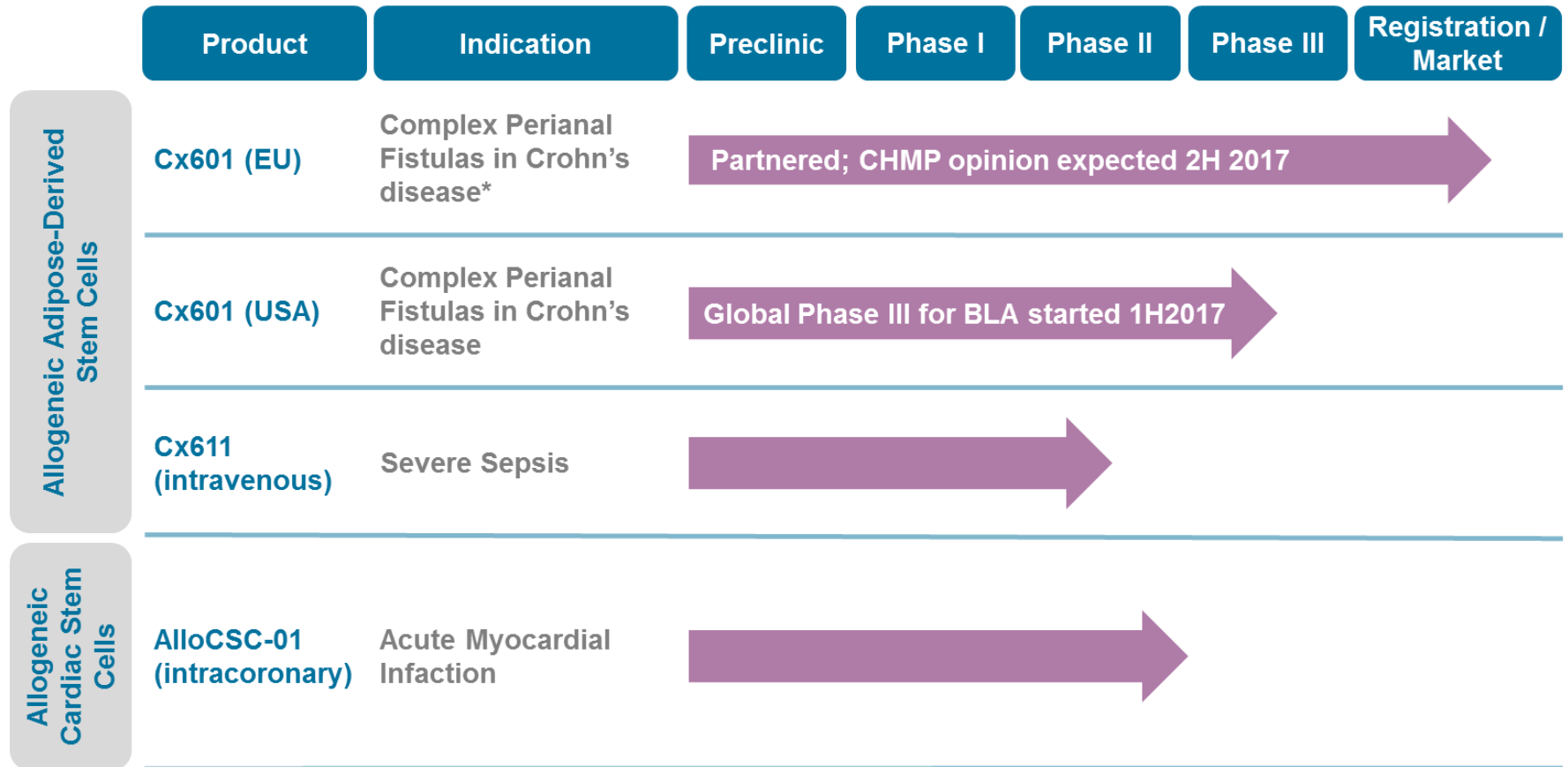
Forward-Looking Statements

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Pipeline

Pre-commercial stage company; Pipeline offering future upside



* Potential label in EU: treatment of complex perianal fistulas in patients with Crohn's disease, who have had an inadequate response to at least one conventional or biologic therapy.



**Novel, locally administered treatment for complex perianal
fistulas in Crohn's disease**

CHMP opinion expected 2H2017

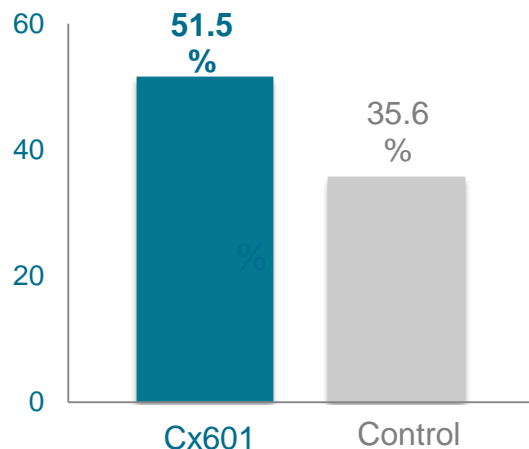
Primary Endpoint Met at Week 24

Benefit sustained, lower relapse¹ rate at week 52

Combined Remission² at W24

(mITT³ Population n= 204)

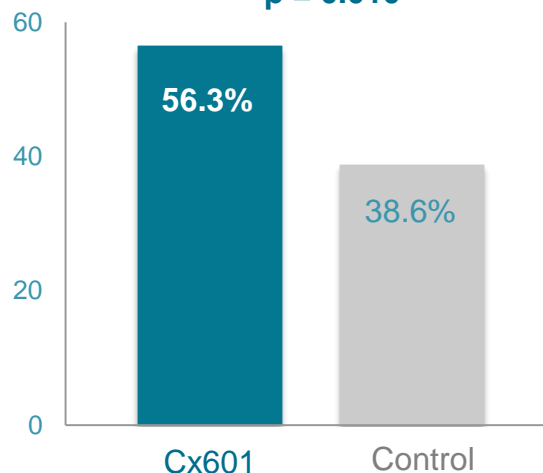
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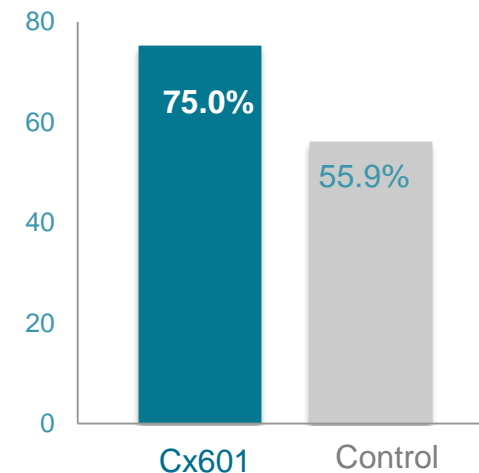
Combined Remission at W52

(mITT Population n= 204)

p = 0.010



Relapse Rate at W52



- Cx601 patients had a 44% greater probability of achieving **Combined Remission** and a **shorter median time** to Clinical Remission (6.7 vs. 14.6 weeks)
- > 50% of patients receiving Cx601 had all treated fistulas in Combined Remission one year after a single administration of the product
- 75.0% of Cx601 patients in combined remission at W24 did not relapse,

¹ Relapse: reopening of any of the treated external openings with active drainage as clinically assessed, or development of perianal collection ≥ 2 cm of the treated perianal fistula confirmed by centrally blinded MRI assessment in patients with clinical remission at any previous visit

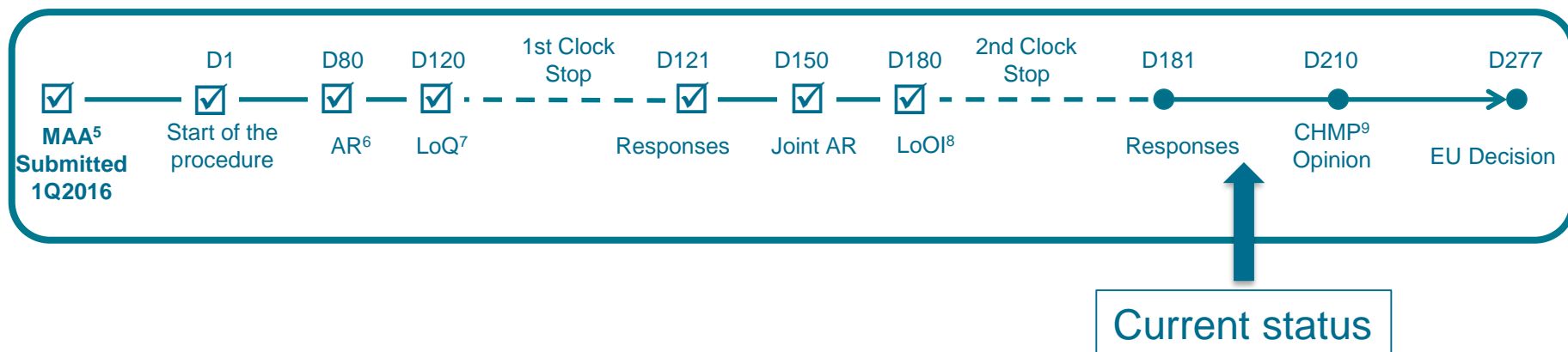
² Closure of all treated external openings draining at baseline despite gentle finger compression, and absence of collections > 2cm by MRI (Magnetic Resonance Imaging)

³ mITT: modified Intention To Treat i.e. patients randomized, treated and with ≥ 1 post-baseline assessment. Efficacy results are consistent across all statistical populations

CHMP Opinion Expected 2H2017

Clear and fast pathway to the market built on a solid regulatory strategy

- Team with previous experience in obtaining MA¹ of cell therapy product
- Orphan Designation received 2009
- 5 Scientific Advice Meetings held with EMA² (2 pre-clinical, 2 CMC³, 1 clinical)
- Approved PIP⁴ with 20 patients to be started not before 2020
- GMP license for commercial manufacturing granted
- CHMP opinion expected 2H2017



Ex-US Rights of Cx601 Licensed to Takeda

TiGenix keeps significant upside potential

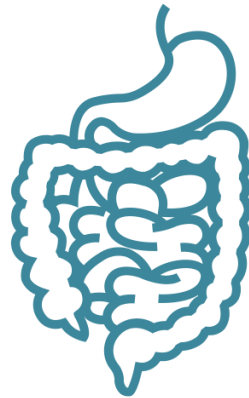
- Exclusive ex-US development and commercialization rights to Cx601 for the treatment of complex perianal fistulas in Crohn's disease patients
- TiGenix retains full US rights as well as the right to develop Cx601 in new indications
- EUR 25M up front plus EUR 10M equity investment
- TiGenix eligible to receive potentially up to EUR 355M in regulatory and sales milestones, including a EUR 15M EU marketing approval milestone
- Double-digit royalties on net sales, tiered to reimbursement price
- Takeda will assume manufacturing responsibilities for Cx601 after an initial period of product supply by TiGenix for the EU

Cx601: Pipeline Expansion Under Evaluation

Potential for Cx601 growth beyond complex perianal fistulas



Other gastrointestinal fistulas



Gastrointestinal indications
other than fistulas

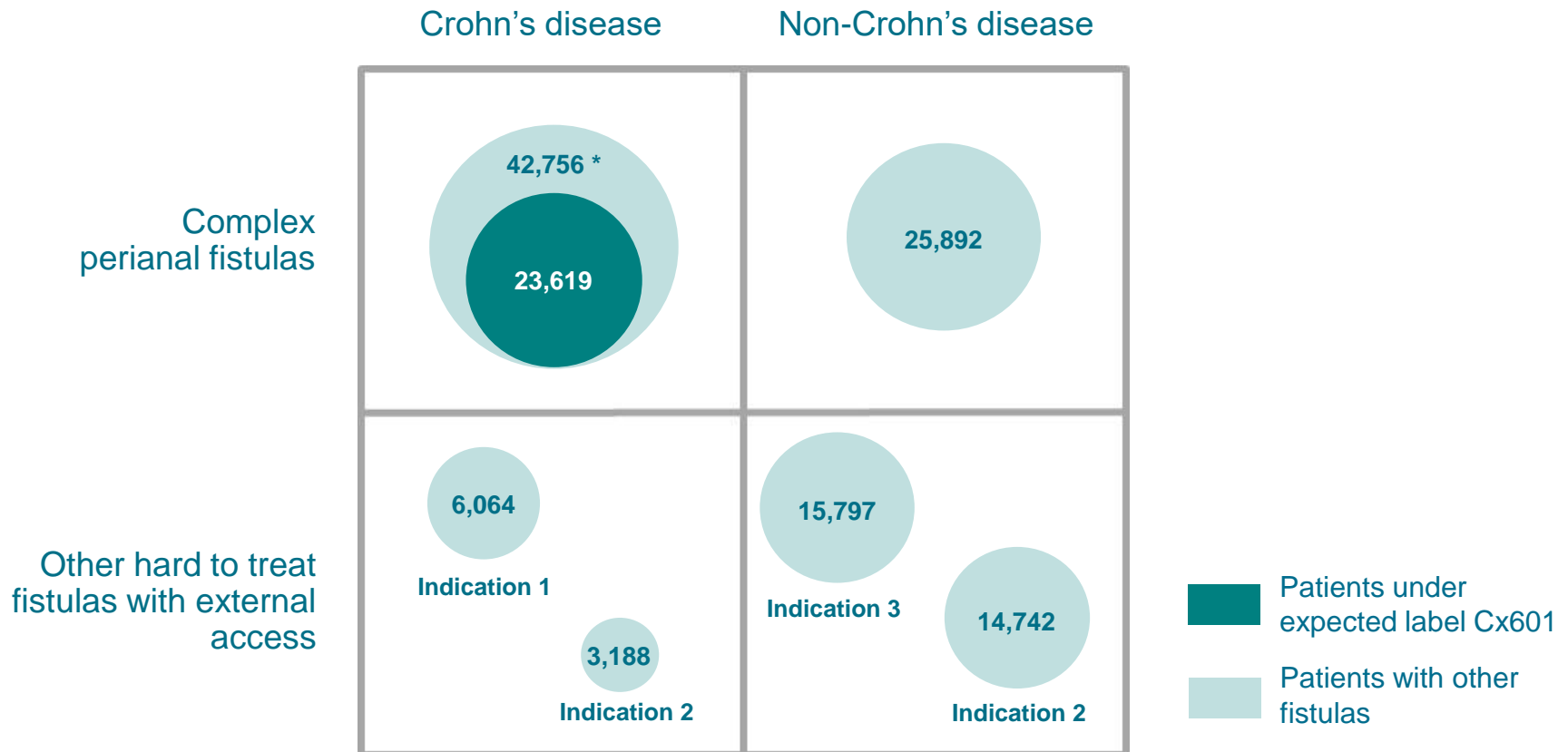


Other indications

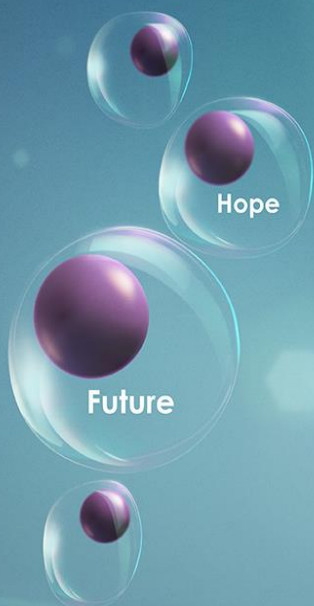
Cx601: Significant Potential in Other Gastrointestinal Fistulas

Addressable population could be four times larger

Estimated Patient Populations in US (2014)



Source: Truven MarketScan® database¹



Stem cells are **full of possibilities**. We make them a **reality**.



Working hard to turn hope into reality

TiGenix is a leading company in **biotechnology** specialized in **stem cells research**, providing breakthrough therapies in order to improve patients' lives.

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