



Synact Pharma AB

A new medicine to treat inflammatory diseases

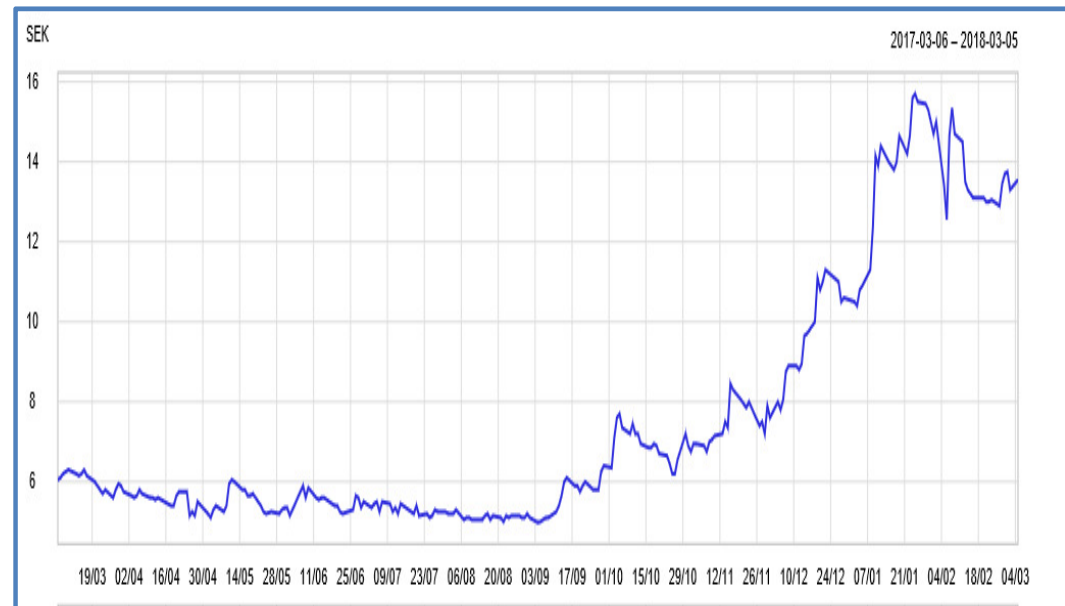
Jeppe Øvlesen, MBA
CEO

Thomas Jonassen, MD
CSO

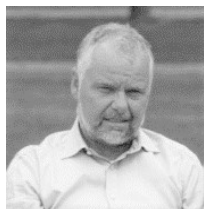
SYNACT PHARMA – OVERVIEW



- Synact Pharma AB listed at Aktietorget in July 2016
- Ticker SYNACT; Aktietorget
- Introduction price at 6.40 SEK per share
- Current (March 6) 14.35 SEK per Share
- Market cap (March 6) of 178.0 SEK



SYNACT PHARMA TEAM



Jeppe Øvli Øvlesen, MBA

- CEO
- > 15 years of CEO experience
- Investor and founding Board Member at a number of Biotech/Medtech companies
- Co-founder of TXP Pharma
- Former CFO & VP BD of Action Pharma



Thomas Jonassen, MD

- Co-founder and CSO,
- Member, Board of Directors
- Associate Professor, KU in Denmark
- Honorable Professor, WHRI, United Kingdom
- Co-founder of TXP Pharma
- Co-founder and former CSO of Action Pharma



Henrik Stage, MSc

- CFO
- Former CEO and CFO at Santaris Pharma
- >25 years experience from Biotech and financial industry



Torbjørn Bjerke, MD

- Chairman, Board of Directors
- Co-founder and Chairman in TXP PHarma
- > 25 years track record from Pharma industry as Head R&D and CEO.
- Co-founder of Action Pharma A/S
- Member, BoD for DBV Technologies



Lars Adellsson

- Member, Board of Directors
- Former GM, GlaxoSmithKline Sweden; Austria
- Former CEO, Medivir
- Member, BoD for Evolan Pharma; Swedish Pharmaceutical Manufacturers Association

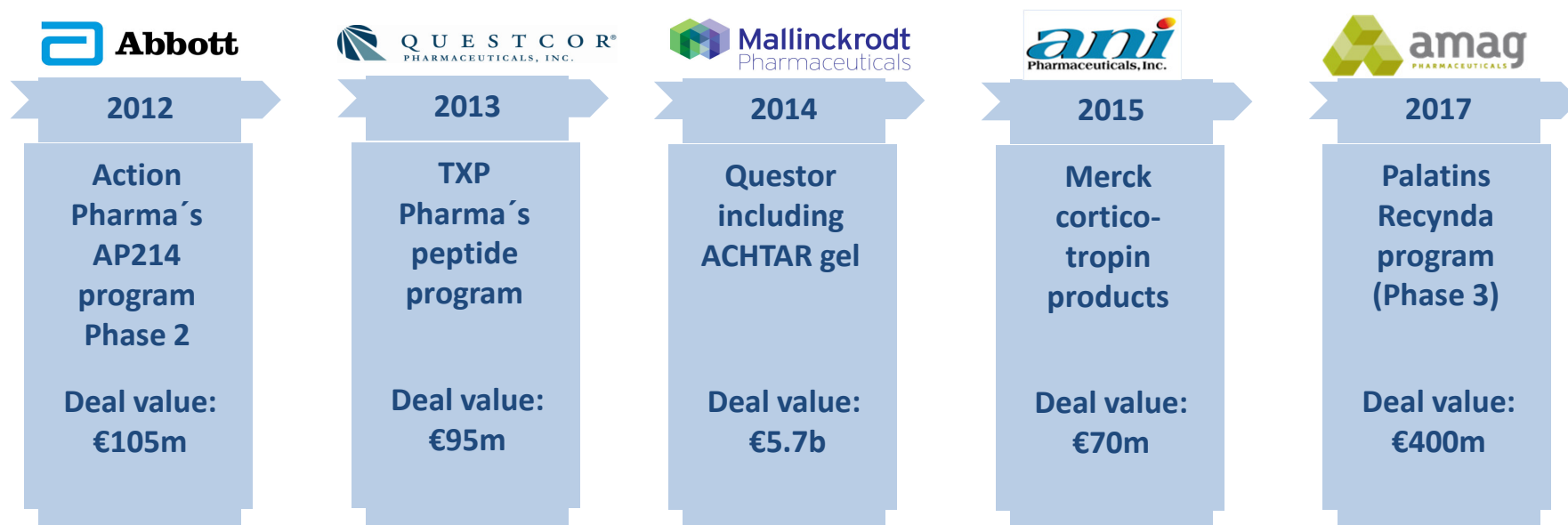


Charlotte Edenius, MD, PhD

- Member, Board of Directors
- Former Executive Vice President for R&D, Medivir AB; Senior Vice President for R&D Orexo AB, Vice President and CSO Biolipox AB and various roles AstraZeneca Clinical R&D
- Member, BoD for Kancera, Immunicum and Gesynta

MELANOCORTIN THERAPY – RECENT DEALS

SYNACT  PHARMA

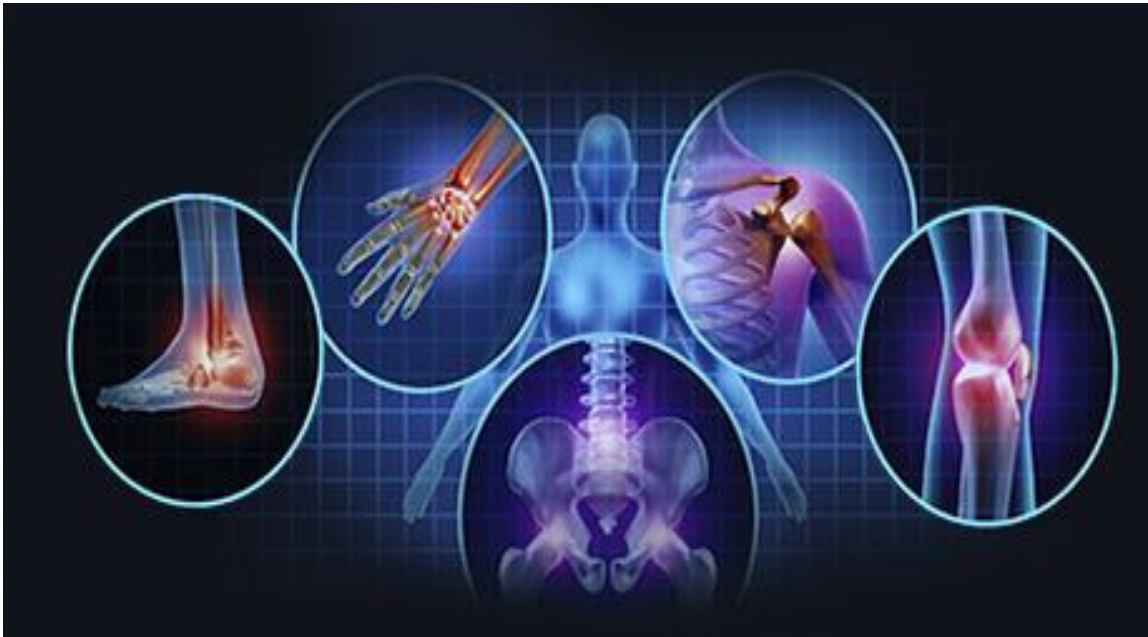


Also worth mentioning is Rhythm Pharmaceuticals - targeting the MC4r in POMC defiance Obesity patients- who recently went public-current market cap: in excess of 700 M US\$

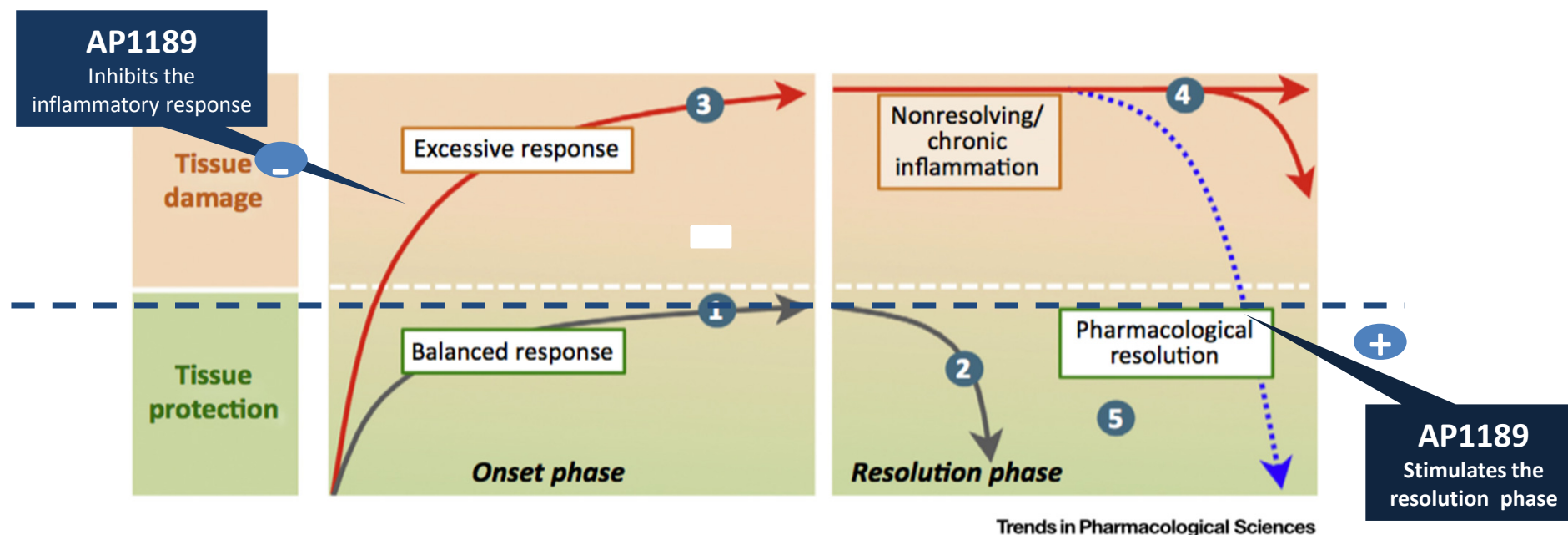
All specialized programs focused on peptide derived therapy

- Main strategy is to develop lead compound AP1189 into Phase II clinical development in inflammatory joint diseases
- Investigate opportunities in additional indications with the possibility to setup development path in second indication
- Strong IP position
- Business model to sell or out-license the project to bio pharma or big biotech
- Current market for lead indication is several Bn \$ annually and growing





- AP1189 SynAct Pharma's lead project is in clinical development for active joint disease – ie rheumatoid and psoriatic arthritis
- Status from ongoing Phase 1 studies shows
 - Well tolerated with attractive safety profile
 - Promising pharmacokinetics profile, supporting once daily oral dosing
 - Tablet for once daily dosing is currently tested in repeat dosing regiment
- Filing of Phase IIa clinical application is scheduled for Q2 2018
- The potential of the compound in additional high-value indications -includes systemic lupus (SLE), Intestinal bowel disease (IBD), multiple Sclerosis (MS) and nephrotic syndrome (NS)- is exploited in ongoing experimental studies.



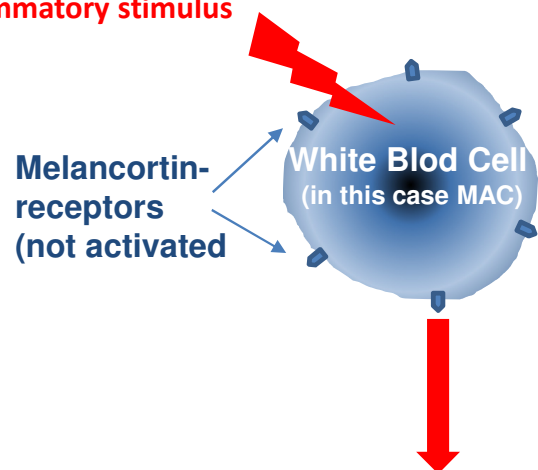
- Resolution therapy holds promise to correct overshooting/ongoing inflammation typical of several pathological settings.

AP1189- STIMULATION OF MELANOCORTIN RECEPTORS ON WHITE CELLS -

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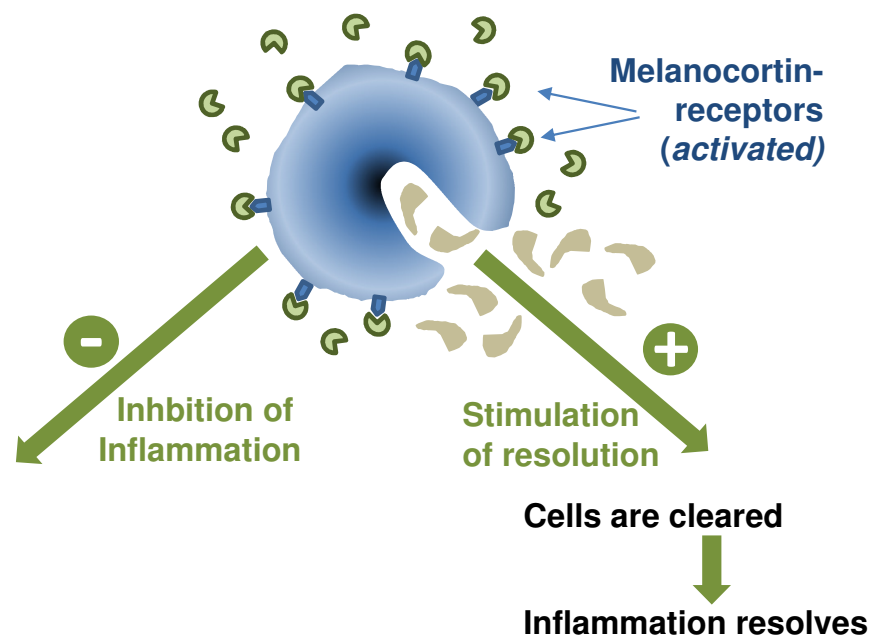
The inflammatory proces:

Inflammatory stimulus



- Release of Inflammatory mediators
- Infiltration with white cells

+ AP1189 



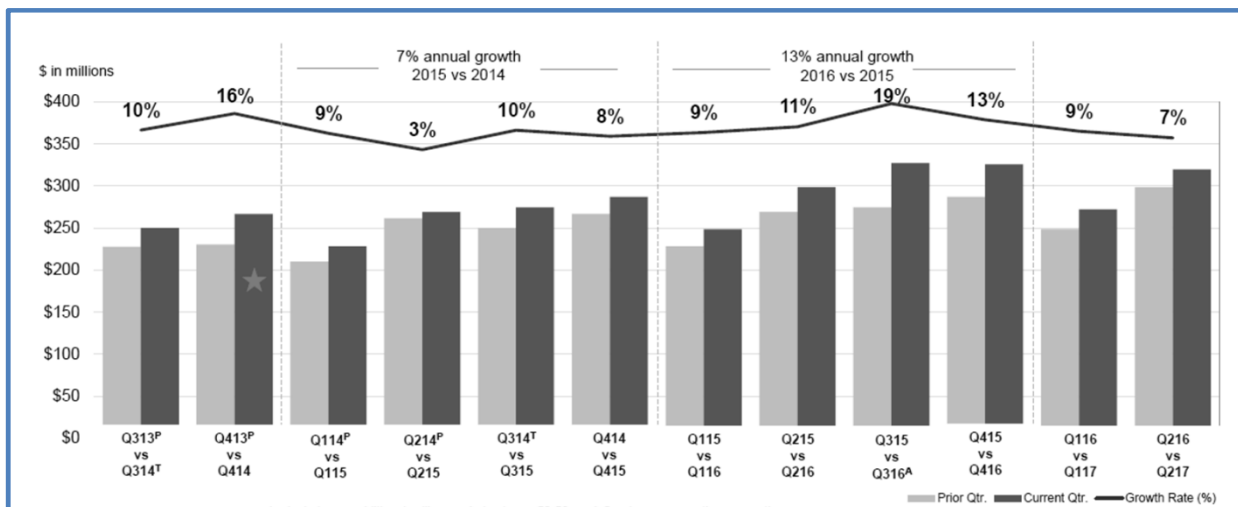
AP1189 stimulates melanocortin-receptors and facilitates thereby resolution of inflammation

MELANOCORTIN DERIVED THERAPY- CURRENT MARKED

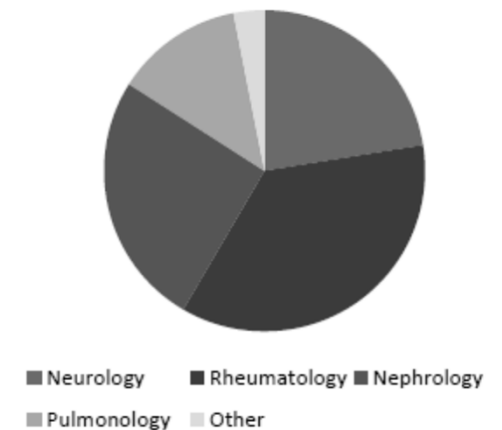
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- ACTH (ACTHAR gel) is a potent anti-inflammatory compound that mediates its anti-inflammatory effects through melanocortin receptor activation
- Current annual sale of around \$ 1.25 Bn -

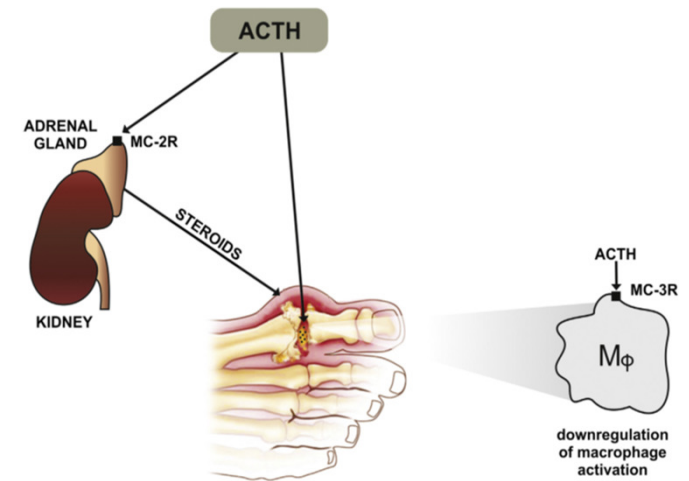


2016 Net Sales by TA



Source: Mallinckrodt Pharmaceuticals
Investor Briefing October 4, 2017

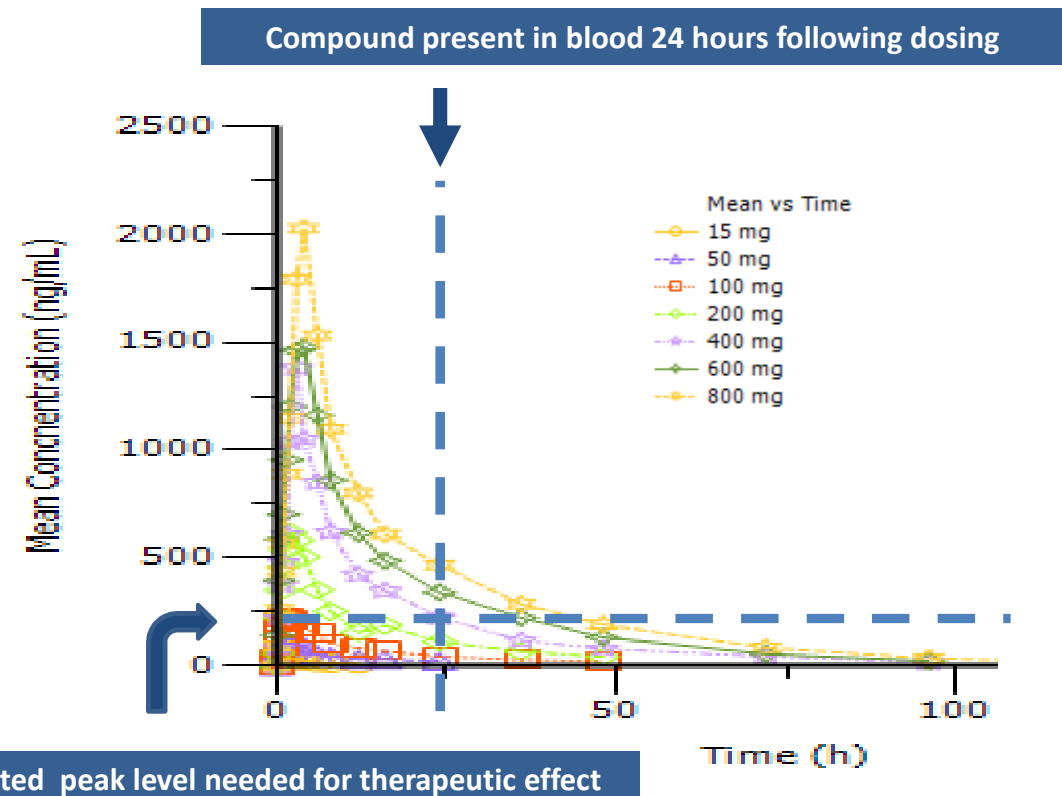
- ACTH (Achartar gel) has potent anti-inflammatory effects in a variety of severe autoimmune and inflammatory disorders
- Two-way mode of action:
 1. Melanocortin receptor activation (non-selective)
 2. Release of steroid hormones
- Route of administration: subcutaneous injections



Limited for difficult to treat patients, due to side-effect profile with unwanted steroid hormone effects and stimulation of dermal pigmentation

AP1189 has the beneficial therapeutic effects related to melanocortin receptor stimulation- AP1189 does not have steroid hormone effects and does not induce dermal pigmentation – in addition AP1189 is oral available

- Dosing completed in 64 healthy volunteers (male as well as post menopausal women)
- In general well tolerated up to highest dose level, where peak exposures reached > 10 times the efficacious concentrations
- Long elimination half life support once daily administration
- Repeat dosing with newly developed tablet is ongoing



MARKTET OPPORTUNITY IN ACTIVE JOINT DISEASE

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- PsA and RA patients present with comparable clinical symptoms and respond to similar treatment modalities, despite different etiologies of disease
- Both the RA and PsA treatment markets are dominated by high priced biologics providing feasibility for novel oral entrants with moderate-high price levels
- Both the RA and PsA markets are growing at double digit rates due to aging populations and novel market entrants
- The RA market is roughly 10x larger than the PsA market in \$ terms



- Patients with active joint disease are currently treated with Methotrexate (MTX)
- Around 40% of these patients have an inadequate response to MTX-
- Next line compounds are expensive and often associated with unwanted side effects
- Even among patients treated with the expensive biologics, up to 30% of the patients still suffer from inadequate treatment effects



A compound given as early add-on to MTX that could reduce the need for second line treatment and/or reduce the dose levels of MTX to option remission of symptoms would have major impact in active joint disease



AP1189

- excellent oral PK and can be developed as a once-daily tablet formulation
 - very well tolerated in ongoing Phase 1 clinical testing
 - robust efficacy in preclinical models of arthritis, IBD and peritonitis
 - Ongoing studies evaluate additional development opportunities
 - potential to be a first-in-class treatment option for inflammatory diseases with big medical needs
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Phase I

- Part 1 and 2 completed
- Part 3 to be completed followed by reporting

Phase II

- Preparation of Phase II Clinical trial in lead indication
- Trial application to be filed in Q2
- Study to be initiated in Q3-

Pre-Clinical

- Testing of compound in additional disease models
- Evaluate potential new business opportunities emerged from new data

Continuous business development discussions with potential partners



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