FY 2020 results webcast

19 February 2021





This presentation contains forward-looking statements, including (without limitation) statements concerning the progress of our clinical pipeline, our expectations regarding commercial sales of Jyseleca, the global R&D collaboration with Gilead, the amendment of our arrangement with Gilead for the commercialization and development of Jyseleca, the timing and/or outcome of the strategic re-evaluation and the cash burn guidance 2021, statements relating to interactions with regulatory authorities, the potential approval process for filgotinib in RA, UC and additional indications, such additional regulatory authorities requiring additional studies, the outcome of pricing and reimbursement interactions, statements relating to the build-up of our commercial organization for filgotinib, statements regarding data from Galapagos' clinical research programs with ziritaxestat which may not support registration or further development due to safety, efficacy or other reasons for IPF, SSc or any other indication, the expected impact of COVID-19, and our strategy, business plans and focus, the slides captioned "ISABELA discontinued," "Deep R&D portfolio based on novel targets," "Broad pipeline," "Building our European commercial footprint," "Outlook," statements regarding the expected timing, design and readouts of ongoing and planned clinical trials (i) with filgotinib in RA, UC, IBD, and other potential indications (ii) with GLPG1205 and GLPG4716 in IPF, (iii) with the Toledo program (iv) with GLPG3667 in Pso, (v) with GLPG0555 in OA, expectations regarding the commercial potential of our product candidates, and our strategy, business plans and focus. When used in this presentation, the words "anticipate," "believe," "can," "could," "estimate," "expect," "intend," "is designed to," "may," "might," "will," "plan," "potential," "possible," "predict," "objective," "should," and similar expressions are intended to identify forward-looking statements.

Except for filgotinib's approval for the treatment of RA by the European Commission and Japanese Ministry of Health, Labour and Welfare, our other drug candidates mentioned in this presentation are investigational; their efficacy and safety have not been fully evaluated by any regulatory authority and they are not yet approved for any use outside of clinical trials.

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Galápagos

ISABELA discontinued

- Dose dependent mortality
- Efficacy below expectations
- All activities with ziritaxestat discontinued
- IPF pipeline: 4 development programs aimed at fibrosis

We remain committed to IPF and fibrosis



>>> 2020 in review

	Q2	Q3	Q4			
	Jyseleca positive UC Ph3 results	Jyseleca approved & launched in EU & JP in RA	ROCCELLA in OA fails	OncoArendi	Updated agreement Jyseleca EU	
		CRL filgotinib RA	Filing Jyseleca in UC EU	Fidelta sold		
			Toledo Roundtable	Start 3 PoC SIK2/3 GLPG3970		
Inflammation	Inflammation Fibrosis Corporate development			PINTA positive IPF		
4						

Deep R&D portfolio based on novel targets

programs in LO

13 10 clinical stage programs

27 validated targets

* LO: Lead optimization

Broad pipeline

Asset	Target	Preclinical	Phase 1	Phase 2	Phase 3	Approval				
Filgotinib	JAK1	CD Ph3 ongoing, submitted UC in EU, approved for RA in EU & Japan								
`3970	SIK2/3	Toledo, PoCs in 5	indications							
`3667	TYK2	Ph1b Pso								
`555	JAK1	Ph1b OA								
`4399	SIK3	Toledo								
`3121	JAK1/TYK2									
`4876	SIK2/3	Toledo								
Other	>10 novel									
`1205	GPR84	Preparing for Ph2	o in IPF							
`4716	Chitinase	Preparing for Ph2	in IPF							
`4586	Undisclosed					Inflammation				
'4605	SIK2/3	Toledo				Fibrosis				
Other	7 novel					Kidney diseases				
GLPG2737	CFTR	PCKD				Other				
GLPG4059	Novel	Metabolic								

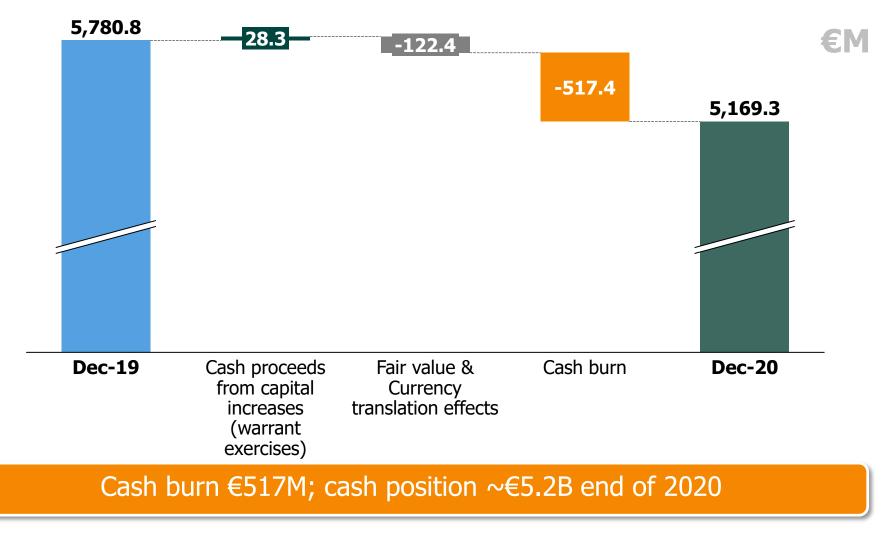
Building our European commercial footprint

- NICE (UK) recommendation moderate RA
- Germany launch on track
- Reimbursement process other countries ongoing
- Complete transition by YE





Cash & current financial investments





Key financials FY '20



(*) Continuing operations (excluding Fidelta)





Filgotinib

- Filing UC Japan
- Outcome MANTA/RA-y
- EU approval decision UC
- DIVERSITY recruited CD

Readouts

- Toledo POCs Pso/RA/UC
- `3667 (TYK2) Ph1b Pso
- `555 (JAK1) Ph1b OA

Cash burn guidance 2021 under review







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