Filgotinib and our commercial ambition

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Pioneering for patients

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Except for filgotinib's approval for the treatment of rheumatoid arthritis by the European Commission and Japanese Ministry of Health, Labour and Welfare, our drug candidates 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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Outcomes FDA discussion

No path forward for 200mg in RA

- GILD decided not to move forward with RA in the U.S.
- Insufficient opportunity now in PsA, AS & uveitis

MANTA/RA-y: up to 52 weeks follow-up required

- for any patients who do not recover fully by week 26 in H1 ’21

IBD opportunity in US remains

- Positive Ph3 read-out in UC
- CD trial continues with data expected in H1 '22

New agreement for filgotinib

- GLPG responsible for all commercial activities for all indications in Europe
- Transition to GLPG commercial organization in Europe by end '21

- 50/50 P&L share for European commercialization until end '21
- All commercial economics to GLPG as of 1 Jan '22, subject to royalty 8% - 15% starting in 2024
- No more EU milestones to GLPG
- GILD to pay GLPG €160M

- GILD retains commercial rights ex-Europe
- Milestones & royalties to GLPG (20-30%) still applicable outside Europe

Broader R&D collaboration not impacted

Realizing our commercial vision

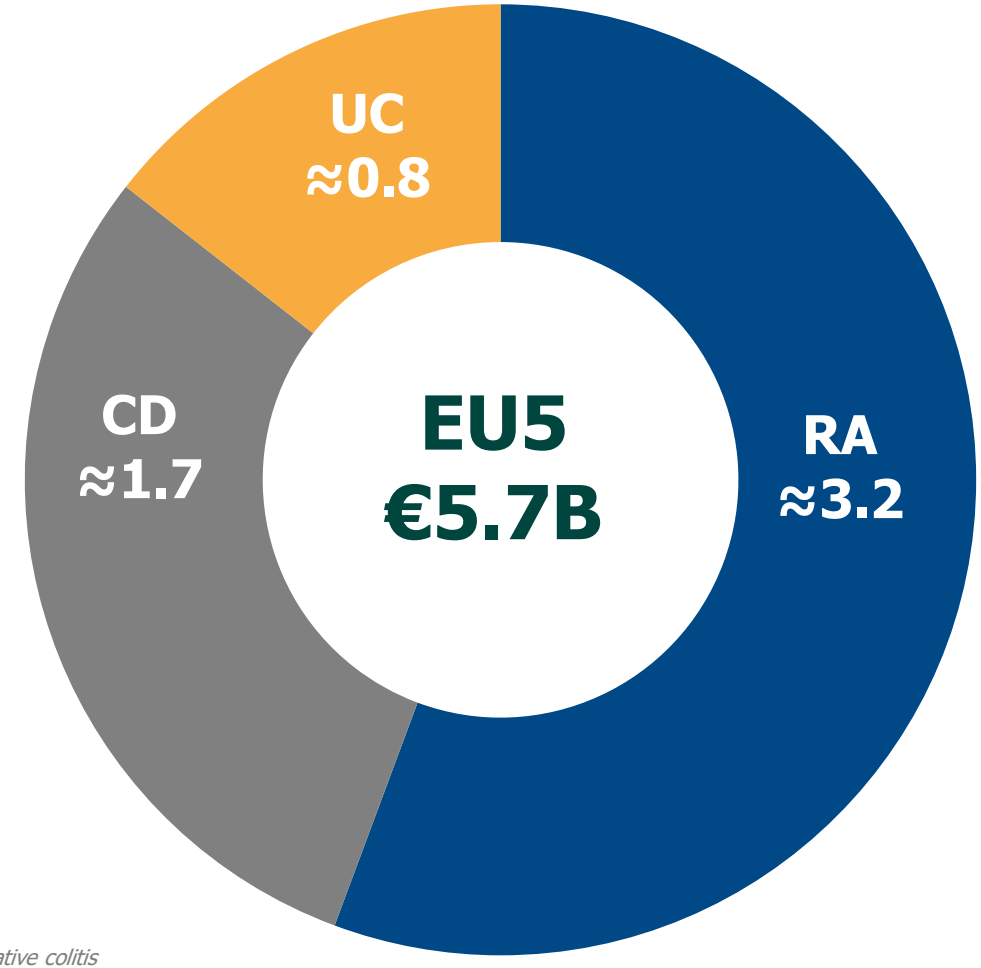
Value creation

Acceleration commercial presence across Europe

Preparation for future launch opportunities

Alignment with overall R&D collaboration with Gilead

EU5 inflammation market today*



Ambition: ≈€0.5B peak sales

8-12% market share for Jyseleca

RA: rheumatoid arthritis; CD: Crohn's disease; UC: ulcerative colitis
Source: IQVIA Analytic Link (MAT to Q2 2020) – est value by disease at ex mfr list prices. All biologics and tsDMARDs.

** U5 inflammation market accounts for approximately 68% of total EU market*

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Jyseleca in RA

Lasting activity

Differentiated safety profile

Fast onset

Two doses approved in Europe and Japan

Convenience of oral

Rapid, robust responses

Monotherapy

Filgotinib is approved for RA in the EU and Japan and not approved for use in any other indication nor any other region.

See the European Summary of Product Characteristics (SmPC) for Jyseleca, which includes contraindications and special warnings and precautions, available at www.ema.europa.eu.

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Transition path

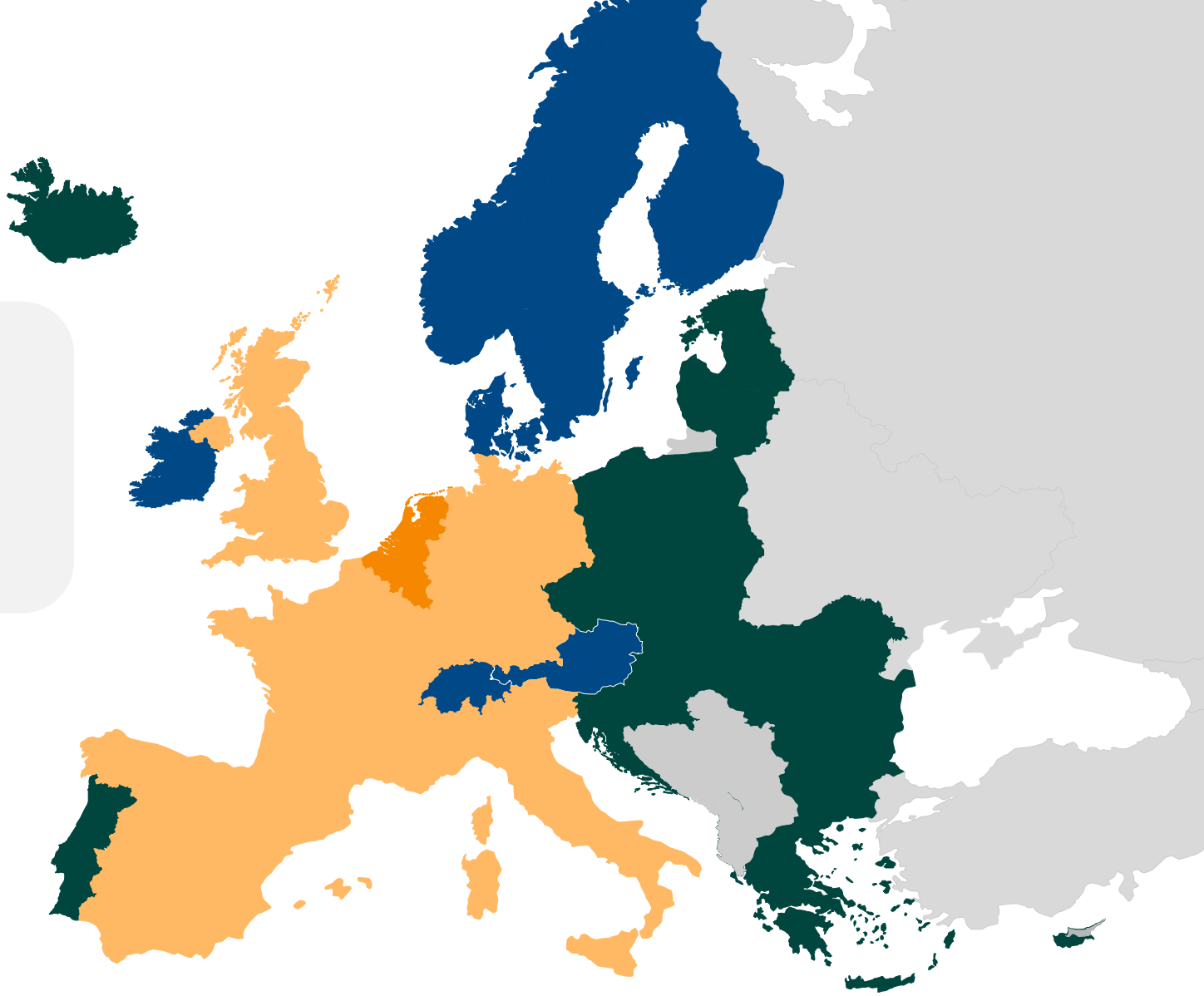
Market size

≈10% No change to Belgium & NL

≈70% EU5: Transfer of full business asap in '21

≈15% Alpine, Nordics & Ireland: transfer by YE '21

≈5% Rest of Europe: rights to GLPG



Transition to full European coverage by end 2021

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Filgotinib timeline

H1 '21	H2 '21	H1 '22	H2 '22
MANTA/RAy week 26 results	UC approval decision EC	DIVERSITY topline (CD)	Potential CD submission EMA
UC submission Japan	European commercial transition complete	UC approval decision Japan	Potential CD submission Japan
CHMP opinion UC			
Commercial transition			