

Analysts' Meet

4 February 2020

Agenda



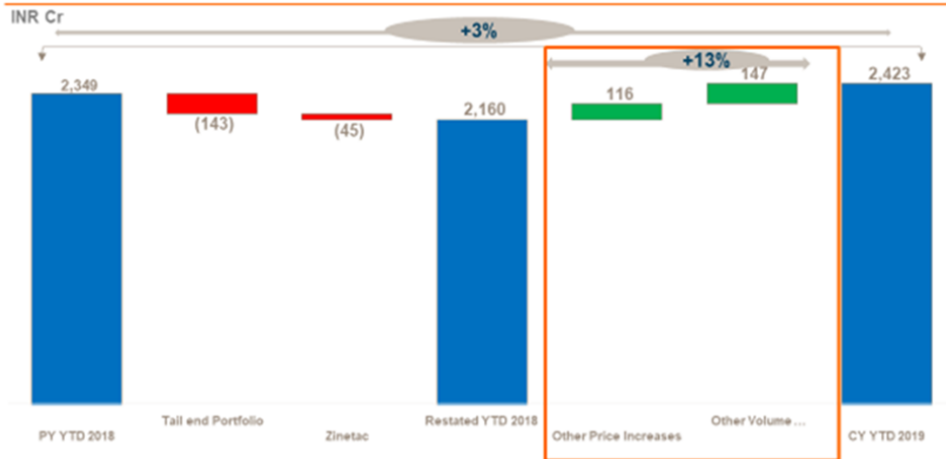
Performance YTD / Q3 19-20



YTD EBIDTA margin +1%; YTD EPS +11%; GP margin +3%

Performance YTD

Headline Growth 3%; Underlying 13% (7% Volume + 6% Price)



Performance Q3

Q3 Sales

Sales 770 Cr / -6%

Underlying growth +6%

Portfolio

	Value	Growth
Key Brands	479Cr	+12%
Rx Key Brands	304Cr	+8%
Vx Key Brands	176Cr	+20%

Portfolio	Value	Growth
Key Brands	1479Cr	+21%
Rx Key Brands	963Cr	+18%
Vx Key Brands	516Cr	+26%

External Growth

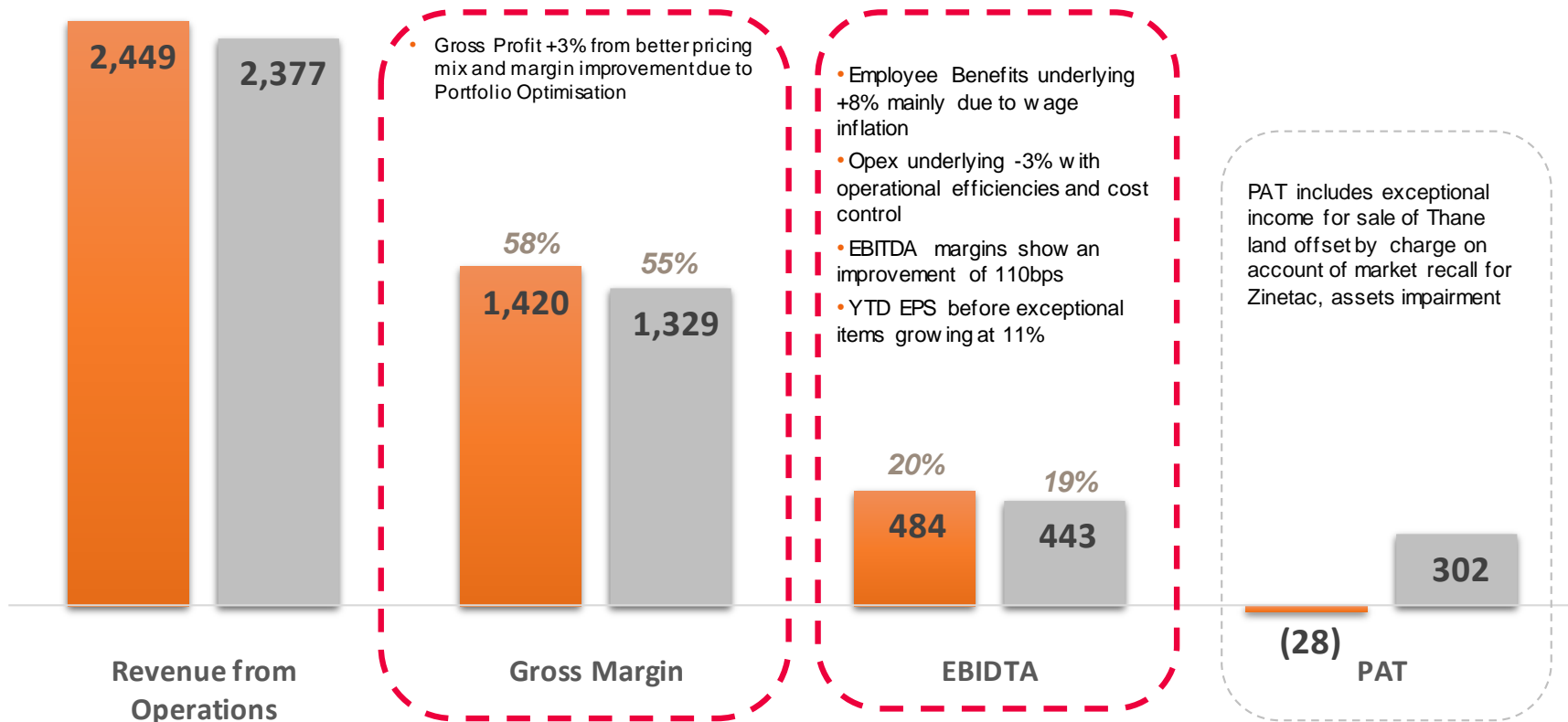
- 2x the market for Promoted brands with EI at 106
- Underlying external growth is 12% with EI of 102

YTD 2019-20 – Performance v PY YTD



Headline: Sales +3% , PAT before exceptional +11%

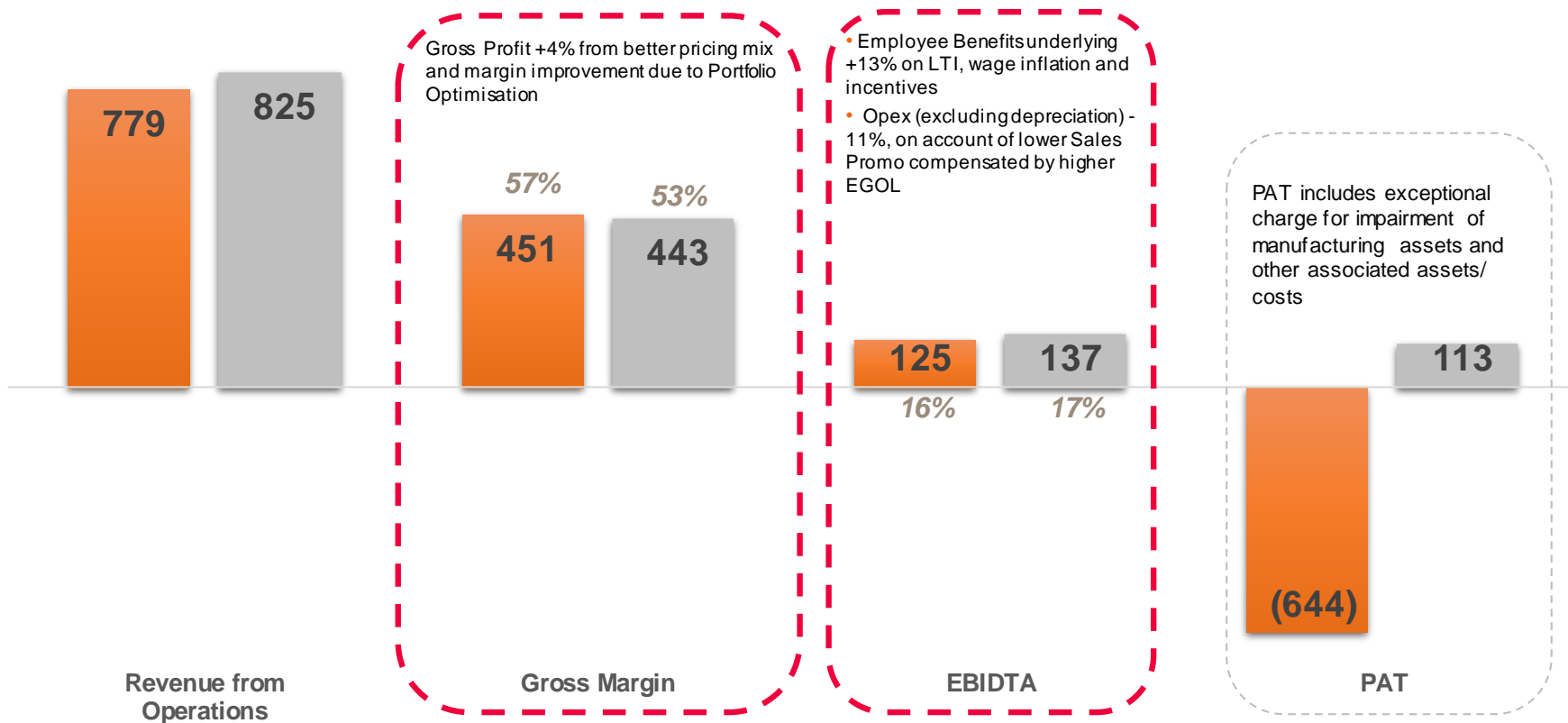
Exceptional Charge 336 Cr from impairment and related costs



Q3 2019-20 – Performance v PY

Headline: Sales - 6% , PAT (before exceptional) @ 9%

Exceptional Charge 737 Cr from impairment and related costs



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Background of Vemgal



Situation

- ❑ Nashik – multi dose form facility; severely under invested. Need for remediation to maintain the Global GSK standards
- ❑ Need to increase capacity to cope up with strong business growth ambition ; maintain market share and maintain commitment to patient safety

Response / Objective

- Planned robust futuristic supply chain for India -
 - ❑ Seamless supply & business continuity to support strong business growth ambition; reduce reliance on Nashik
 - ❑ Compliance to GSK Global standards and maintain commitment to patient safety
- Proposal for a high volume Greenfield site for OSD (Oral Solid Dose)**

Investment

- ❑ Greenfield project with 9bn dose capacity with INR 1000 cr investment – 60% capacity dedicated for Zinetac
- ❑ Focus on high volume low complexity products
- ❑ 2013: Initial project approval
- ❑ 2014: Vemgal project initiation
- ❑ 2019: Manufacturing license received with commercialization planned in 2019-20
- ❑ Sep 2019: Voluntary global recall of ranitidine products, including Zinetac in India.

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Financial Performance YTD / Q3 2019-20

Vemgal Update

Unexpected Development: Ranitidine

India Growth Story

Innovation Focus

Unexpected Global Development: Ranitidine



- **NDMA** (N-nitrosodimethylamine) issue highlighted in mid Sept 2019 leading to voluntary recall of Zinetac
- **GSK continues to respond to queries from global regulatory authorities**
 - The EMA has initiated an investigation into the potential root causes for the formation of NDMA in Ranitidine containing products
- **The investigation into the causes for the formation of NDMA is highly complex**
 - Root cause analysis is on-going & multiple options are being explored
 - Currently testing a significant representation of batches of finished product (tablets, effervescent tablets, IV solution, and syrup dose forms)
 - 20 scientists working full time have analysed over 360 batches to date
- **Patient safety remains a top priority and it is fundamental to maintaining trust**

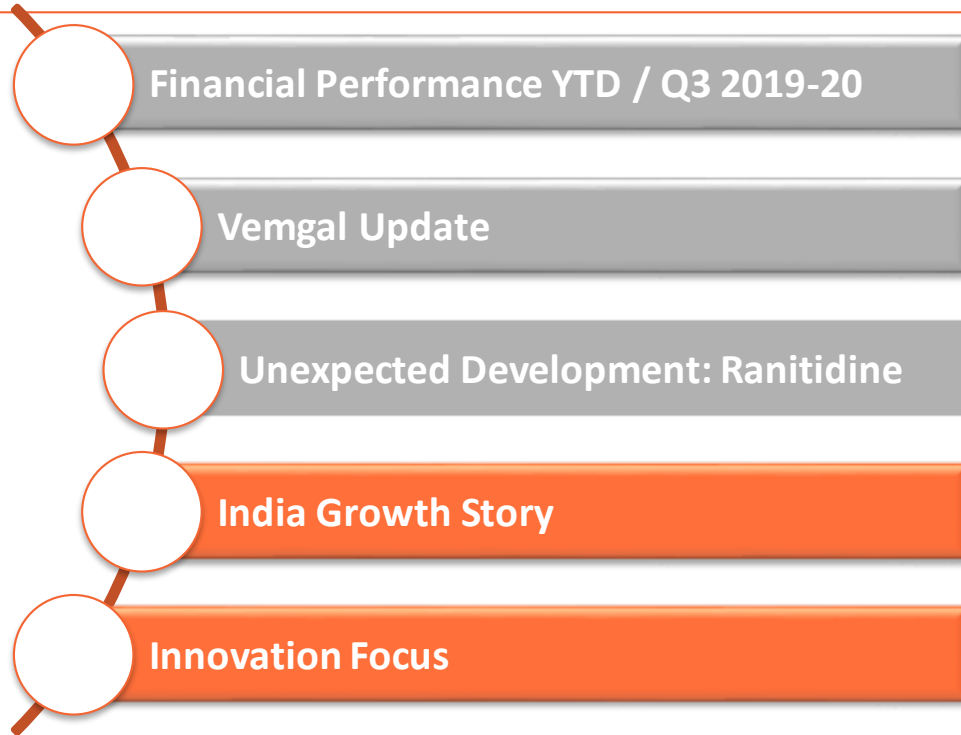
Options explored

- ❑ Alternate SKU production at site; Additional capital investment required hence low financial viability;
- ❑ Low volumes for alternate products leading to significant under utilization of capacity at Vemgal

Financial Governance

- ❑ Ranitidine issue resulted in under-utilization and triggered an impairment
- ❑ Recognize financial impairment of INR 640 cr connected to the underutilization of GSK's manufacturing facilities and INR 97 cr on account of other related assets / cost.

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2019-2021: Accelerating India Rx profitable growth



Winning in right TAs



Increased SoV on key brands to drive growth in AI, Derma and Vaccines

Redeployment of 400 FTEs and increment of 200 FTE. Increase reach by 140K new HCPs

Strengthen specialty pipeline with launch of Nucala and Menveo. Initiate registration process for Shingrix

Winning in channels



Build trust with channels
Acquire/ Build capability to engage with channels

Build technology platforms to commercially transact with 100K pharmacies

+100 FTE

Order fulfilment of distributed brands (Non-promoted)

Winning with HCPs virtually



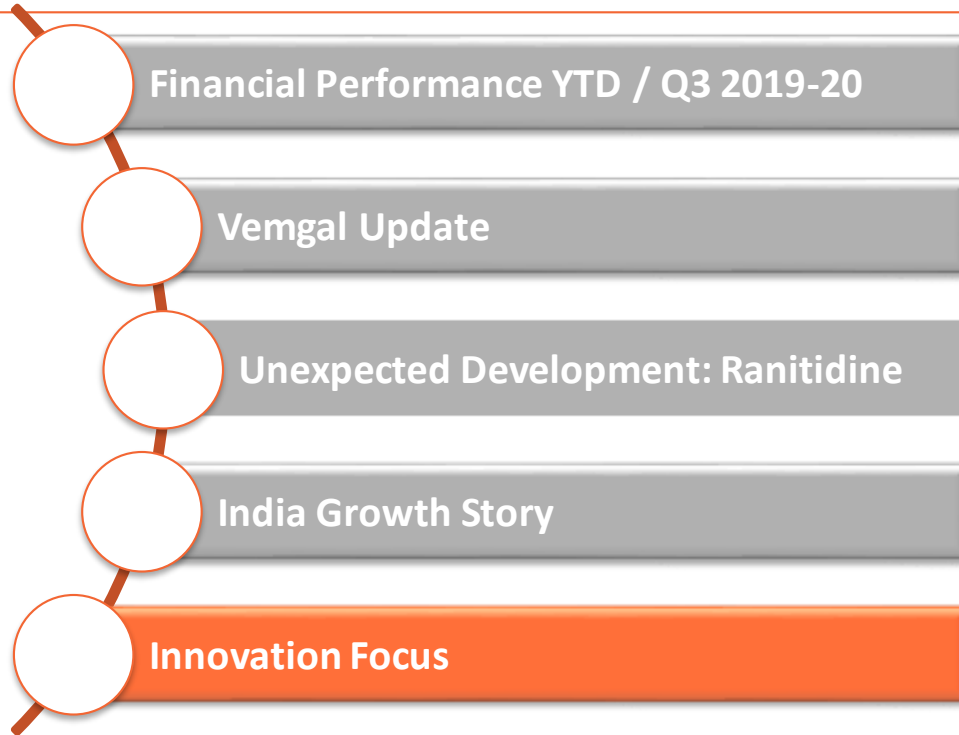
Established a model for digital only promotion for 6 identified brands with agency onboard

Enhance customer experience through increased touchpoint (1.5x) enabled by MCM

Focus to drive higher Qualified Reach

Improved supply efficiency with focus on OTIF/ Inventory

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Innovation

- Continued success of Nucala (Mepolizumab) which is a humanised monoclonal antibody and indicated as an add-on treatment for severe refractory eosinophilic asthma in adult patients
- Launch of Menveo vaccine for protection against meningococcal disease.
- Continued commitment on introducing new therapies / specialities in the market both in Pharma and Vaccines with the initial success of Nucala and Menveo.

