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Form 425
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Legal Matters IRISH LAW RESTRICTIONS ON CERTAIN INFORMATION Mylan N.V.'s ("Mylan") offer for Perrigo Company plc ("Perrigo") is governed by the Irish Takeover Panel Act, 1997, Takeover Rules 2013 (the "Irish Takeover Rules"). Under the Irish Takeover Rules, Mylan management is prohibited from discussing any material information or significant new opinions which have not been publicly announced. Any person interested in shares of Mylan or Perrigo is encouraged to consult their professional advisers. FORWARD-LOOKING STATEMENTS This communication contains "forward-looking statements." These statements are made pursuant to the safe harbor provisions of the Private Securities Litigation Reform Act of 1995. Such forward-looking statements may include, without limitation, statements about the proposed acquisition of Perrigo by Mylan (the "Perrigo Proposal"), Mylan's acquisition (the "EPD Transaction") of Mylan Inc. and Abbott's non-U.S. developed markets specialty and branded generics business (the "EPD Business"), the benefits and synergies of the Perrigo Proposal or EPD Transaction, future opportunities for Mylan, Perrigo, or the combined company and products, any other statements regarding Mylan's, Perrigo's, or the combined company's future operations, anticipated business levels, future earnings, planned activities, anticipated growth, market opportunities, strategies, competition, and other expectations and targets for future periods, and statements about Mylan remaining very well-positioned to supply the anticipated customer and patient demand for the EpiPen® Auto-Injector and that Mylan sees an additional \$0.25 benefit to adjusted diluted EPS in 2016 with respect to the EpiPen® Auto-Injector. These may often be identified by the use of words such as "will," "may," "could," "should," "would," "project," "believe," "anticipate," "expect," "plan," "estimate," "forecast," "potential," "intend," "continue," "target" and variations of these words or comparable words. Because forward-looking statements inherently involve risks and uncertainties, actual future results may differ materially from those expressed or implied by such forward-looking statements. Factors that could cause or contribute to such differences include, but are not limited to: uncertainties related to the Perrigo Proposal, including as to the timing of the offer and a compulsory acquisition, whether Perrigo will cooperate with Mylan and whether Mylan will be able to consummate the offer and a compulsory acquisition, the possibility that competing offers will be made, the possibility that the conditions to the consummation of the offer will not be satisfied, and the possibility that Mylan will be unable to obtain regulatory approvals for the offer or be required, as a condition to obtaining regulatory approvals, to accept conditions that could reduce the anticipated benefits of the offer; the ability to meet expectations regarding the accounting and tax treatments of a transaction relating to the Perrigo Proposal and the EPD Transaction; changes in relevant tax and other laws, including but not limited to changes in healthcare and pharmaceutical laws and regulations in the U.S. and abroad; the integration of Perrigo and the EPD Business being more difficult, time-consuming, or costly than expected; operating costs, customer loss, and business disruption (including, without limitation, difficulties in maintaining relationships with employees, customers, clients, or suppliers) being greater than expected following the Perrigo Proposal and the EPD Transaction; the retention of certain key employees of Perrigo and the EPD Business being difficult; the possibility that Mylan may be unable to achieve expected synergies and operating efficiencies in connection with the Perrigo Proposal and the EPD Transaction within the expected time-frames or at all and to successfully integrate Perrigo and the EPD Business; expected or targeted future financial and operating performance and results; the capacity to bring new products to market, including but not limited to where Mylan uses its business judgment and decides to manufacture, market, and/or sell products, directly or through third parties, notwithstanding the fact that allegations of patent infringement(s) have not been finally resolved by the courts (i.e., an "at-risk launch"); any regulatory, legal, or other impediments to our ability to bring new products to market; success of clinical trials and our ability to execute on new product opportunities; any changes in or difficulties with our inventory of, and our ability to manufacture and distribute, the EpiPen® Auto-Injector; the scope, timing, and outcome of any ongoing legal proceedings and the impact of any such proceedings on financial condition, results of operations, and/or cash flows; the ability to protect intellectual property and preserve intellectual property rights; the effect of any changes in customer and supplier relationships and customer purchasing patterns; the ability to attract and retain key personnel; changes in third-party relationships; the impact of competition; changes in the economic and financial conditions of the businesses of Mylan, Perrigo, or the combined company; the inherent challenges, risks, and costs in identifying, acquiring, and integrating complementary or strategic acquisitions of other companies, products, or assets and in achieving anticipated synergies; uncertainties and matters beyond the control of management; and inherent uncertainties involved in the estimates and judgments used in the preparation of financial statements, and the

providing of estimates of financial measures, in accordance with accounting principles generally accepted in the United States ("GAAP") and related standards or on an adjusted basis. For more detailed information on the risks and uncertainties associated with Mylan's business activities, see the risks described in Mylan's Quarterly Reports on Form 10-Q for the quarters ended March 31, 2015 and June 30, 2015 and our other filings with the Securities and Exchange Commission ("SEC"). These risks, as well as other risks associated with Mylan, Perrigo, and the combined company are also more fully discussed in the Registration Statement on Form S-4 (which includes an offer to exchange/prospectus and was declared effective on September 10, 2015, the "Registration Statement") in connection with the Perrigo Proposal. You can access Mylan's filings with the SEC through the SEC website at www.sec.gov, and Mylan strongly encourages you to do so. Mylan undertakes no obligation to update any statements herein for revisions or changes after the date of this communication.

Legal Matters RESPONSIBILITY STATEMENT The directors of Mylan accept responsibility for the information contained in this communication, save that the only responsibility accepted by the directors of Mylan in respect of the information in this communication relating to Perrigo, Perrigo's subsidiaries and subsidiary undertakings, the Perrigo board of directors and the persons connected with them, which has been compiled from published sources, has been to ensure that such information has been correctly and fairly reproduced or presented (and no steps have been taken by the directors of Mylan to verify this information). To the best of the knowledge and belief of the directors of Mylan (who have taken all reasonable care to ensure that such is the case) the information contained in this communication is in accordance with the facts and does not omit anything likely to affect the import of such information.

DEALING DISCLOSURE REQUIREMENTS Under the provisions of Rule 8.3 of the Irish Takeover Rules, if any person is, or becomes, 'interested' (directly or indirectly) in, 1% or more of any class of 'relevant securities' of Perrigo or Mylan, all 'dealings' in any 'relevant securities' of Perrigo or Mylan (including by means of an option in respect of, or a derivative referenced to, any such 'relevant securities') must be publicly disclosed by not later than 3:30 pm (New York time) on the 'business' day following the date of the relevant transaction. This requirement will continue until the date on which the 'offer period' ends. If two or more persons co-operate on the basis of any agreement, either express or tacit, either oral or written, to acquire an 'interest' in 'relevant securities' of Perrigo or Mylan, they will be deemed to be a single person for the purpose of Rule 8.3 of the Irish Takeover Rules. Under the provisions of Rule 8.1 of the Irish Takeover Rules, all 'dealings' in 'relevant securities' of Perrigo by Mylan or 'relevant securities' of Mylan by Perrigo, or by any party acting in concert with either of them, must also be disclosed by no later than 12 noon (New York time) on the 'business' day following the date of the relevant transaction. A disclosure table, giving details of the companies in whose 'relevant securities' 'dealings' should be disclosed, can be found on the Irish Takeover Panel's website at www.irishtakeoverpanel.ie. Interests in securities arise, in summary, when a person has long economic exposure, whether conditional or absolute, to changes in the price of securities. In particular, a person will be treated as having an 'interest' by virtue of the ownership or control of securities, or by virtue of any option in respect of, or derivative referenced to, securities. Terms in quotation marks are defined in the Irish Takeover Rules, which can also be found on the Irish Takeover Panel's website. If you are in any doubt as to whether or not you are required to disclose a dealing under Rule 8, please consult the Irish Takeover Panel's website at www.irishtakeoverpanel.ie or contact the Irish Takeover Panel on telephone number +353 1 678 9020 or fax number +353 1 678 9289. Goldman Sachs, which is authorized by the Prudential Regulation Authority and regulated by the Financial Conduct Authority and the Prudential Regulation Authority in the United Kingdom, is acting for Mylan and no one else in connection with the Perrigo Proposal and will not be responsible to anyone other than Mylan for providing the protections afforded to clients of Goldman Sachs, or for giving advice in connection with the Perrigo Proposal or any matter referred to herein. Goldman Sachs does not accept any responsibility whatsoever for the contents of this communication or for any statement made or purported to be made by them or on their behalf in connection with the offer. Goldman Sachs accordingly disclaims all and any liability whether arising in tort, contract, or otherwise which it might otherwise have in respect of this communication or any such statement.

Legal Matters ADDITIONAL INFORMATION In connection with the Perrigo Proposal, Mylan has filed certain materials with the SEC (and anticipates filing further materials), including, among other materials, the Registration Statement. In connection with the Perrigo Proposal, Mylan also filed with the SEC on September 14, 2015 a Tender Offer Statement on Schedule TO, which includes the offer to exchange / prospectus (the “Offer to Exchange / Prospectus”), form of letter of transmittal and other related offer documents. Mylan has mailed the Offer to Exchange / Prospectus to Perrigo shareholders in connection with the Perrigo Proposal. This communication is not intended to be, and is not, a substitute for such filings or for any other document that Mylan may file with the SEC in connection with the Perrigo Proposal. **INVESTORS AND SECURITYHOLDERS OF MYLAN AND PERRIGO ARE URGED TO READ THE DOCUMENTS FILED WITH THE SEC CAREFULLY AND IN THEIR ENTIRETY (IF AND WHEN THEY BECOME AVAILABLE) BEFORE MAKING AN INVESTMENT DECISION BECAUSE THEY WILL CONTAIN IMPORTANT INFORMATION ABOUT MYLAN, PERRIGO AND THE PERRIGO PROPOSAL.** Such documents will be available free of charge through the website maintained by the SEC at www.sec.gov or by directing a request to Mylan at 724-514-1813 or investor.relations@mylan.com. Any materials filed by Mylan with the SEC that are required to be mailed to shareholders of Perrigo and/or Mylan will also be mailed to such shareholders. This communication has been prepared in accordance with U.S. securities law, Irish law, and the Irish Takeover Rules. A copy of this communication will be available free of charge at the following website: perrigotransaction.mylan.com. Such website is neither endorsed, nor sponsored, nor affiliated with Perrigo or any of its affiliates. **PERRIGO®** is a registered trademark of L. Perrigo Company. **NON-SOLICITATION** This communication is not intended to, and does not, constitute or form part of (1) any offer or invitation to purchase or otherwise acquire, subscribe for, tender, exchange, sell, or otherwise dispose of any securities, (2) the solicitation of an offer or invitation to purchase or otherwise acquire, subscribe for, sell, or otherwise dispose of any securities, or (3) the solicitation of any vote or approval in any jurisdiction pursuant to this communication or otherwise, nor will there be any acquisition or disposition of the securities referred to in this communication in any jurisdiction in contravention of applicable law or regulation. No offer of securities shall be made except by means of a prospectus meeting the requirements of Section 10 of the Securities Act of 1933, as amended. **FURTHER INFORMATION** The distribution of this communication in certain jurisdictions may be restricted or affected by the laws of such jurisdictions. Accordingly, copies of this communication are not being, and must not be, mailed or otherwise forwarded, distributed or sent in, into, or from any such jurisdiction. Therefore, persons who receive this communication (including, without limitation, nominees, trustees and custodians) and are subject to the laws of any such jurisdiction will need to inform themselves about, and observe, any applicable restrictions or requirements. Any failure to do so may constitute a violation of the securities laws of any such jurisdiction. To the fullest extent permitted by applicable law, Mylan disclaims any responsibility or liability for the violations of any such restrictions by any person.

Legal Matters Non-GAAP Financial Measures This communication includes the presentation and discussion of certain financial information that differs from what is reported under GAAP. These non-GAAP financial measures, including, but not limited to, adjusted diluted EPS, adjusted cash provided by operating activities, adjusted third party net sales from Europe, adjusted Generics segment third party net sales, adjusted third party net sales, adjusted total revenues, adjusted gross profit, adjusted gross margins, adjusted net earnings attributable to Mylan, adjusted constant currency total revenues, adjusted constant currency third party net sales, EBITDA, adjusted EBITDA, adjusted R&D, adjusted SG&A, and adjusted effective tax rate, are presented in order to supplement investors' and other readers' understanding and assessment of the Company's financial performance. Mylan has also presented certain non-GAAP financial measures for Perrigo, including, but not limited to, adjusted diluted EPS and adjusted EBITDA margin, which have been taken from published sources. Management uses these measures internally for forecasting, budgeting and measuring its operating performance. In addition, primarily due to acquisitions, Mylan believes that an evaluation of its ongoing operations (and comparisons of its current operations with historical and future operations) would be difficult if the disclosure of its financial results were limited to financial measures prepared only in accordance with GAAP. In addition, Mylan believes that including EBITDA and supplemental adjustments applied in presenting adjusted EBITDA pursuant to our debt agreements is appropriate to provide additional information to investors to demonstrate the Company's ability to comply with financial debt covenants (which are calculated using a measure similar to adjusted EBITDA) and assess the Company's ability to incur additional indebtedness. We also report sales performance using the non-GAAP financial measure of "constant currency" total revenues, adjusted total revenues, third party net sales, and adjusted third party net sales. This measure provides information on the change in net sales assuming that foreign currency exchange rates had not changed between the prior and current period. The comparisons presented as constant currency rates reflect comparative local currency sales at the prior year's foreign exchange rates. We routinely evaluate our third party net sales performance at constant currency so that sales results can be viewed without the impact of foreign currency exchange rates, thereby facilitating a period-to-period comparison of our operational activities, and we believe that this presentation also provides useful information to investors for the same reason. The "Summary of Adjusted Revenues by Segment" table in the Appendix compares adjusted third party net sales on an actual and constant currency basis for each reportable segment and the geographic regions within the Generics segment for the three and nine months ended September 30, 2015 and 2014. Also, set forth below, Mylan has provided reconciliations of such non-GAAP financial measures to the most directly comparable GAAP financial measures (which in the case of Perrigo's reconciliations, have been taken from published sources), other than Perrigo's adjusted diluted EPS company guidance and Thomson Reuters consensus estimates of adjusted EBITDA and adjusted diluted EPS which cannot be reconciled as they are from a third party source. Mylan does not endorse or adopt Thomson Reuters consensus estimates. Investors and other readers are encouraged to review the related GAAP financial measures and the reconciliations of the non-GAAP measures to their most directly comparable GAAP measures set forth in the Appendix, and investors and other readers should consider non-GAAP measures only as supplements to, not as substitutes for or as superior measures to, the measures of financial performance prepared in accordance with GAAP.

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Robert J. Coury - Executive Chairman Heather Bresch - CEO Rajiv Malik - President John Sheehan - CFO Today's Presenters

Q3 2015 Highlights Adjusted constant currency revenues grew 36%* compared to the prior year Mylan legacy adjusted constant currency revenues grew 14%*, reflecting continued strength in our legacy business Generics segment adjusted constant currency revenues grew 48%* and the legacy Mylan Generics segment grew 19%*, with positive growth across all regions Adjusted diluted EPS grew 23%* compared to the prior year *Adjusted metrics and constant currency measures are non-GAAP financial measures. Please see the Appendix for reconciliations of such non-GAAP financial measures to the most directly comparable GAAP financial measures.

EpiPen® Auto-Injector With 85% market share, we continue to see positive script volume on a year-to-date basis. Mylan remains very well-positioned to supply the anticipated customer and patient demand, as we have consistently done for over 25 years. Based on recent events, we see EpiPen® Auto-Injector contributing an additional \$0.25 to \$0.30 benefit to adjusted diluted EPS in 2016.¹ Illustrative impact of recent long term landscape changes in the EpiPen® Auto-Injector market on Mylan adjusted diluted EPS in 2016. This illustrative EPS growth is not intended to constitute a profit forecast for any period nor should it be interpreted to mean that earnings or earnings per share will necessarily be greater or lesser than those for the relevant preceding financial periods of Mylan.

2015 Financial Guidance 2015 adjusted diluted EPS is now expected to be at the upper end of the \$4.15 to \$4.35 guidance range *The 20% year-on-year growth rate referred to in this slide is illustrative only, and is intended to convey Mylan's year-on-year adjusted diluted EPS growth rate in the event that Mylan achieves 2015 adjusted diluted EPS which is marginally above the midpoint of its existing \$4.15 to \$4.35 guidance range. This represents, once again, greater than 20%* year-over-year growth

Perrigo's Decelerating Core Highlights Weakness in Standalone Strategy Mylan Transaction Creates a Stronger and More Competitive Platform to Drive Growth Source: Public filings, Wall Street Research (08-Sep-2015 Jefferies Biogen report), Thomson Reuters consensus estimates as used by Perrigo in its September 17, 2015 Investor Presentation and Perrigo Investor Presentation "Creating Value for Shareholders: Now and for the Long Term", filed October 22, 2015. Mylan does not endorse or adopt Thomson Reuters consensus estimates, which are used for illustrative purposes only. Nothing on this slide is intended to be a profit forecast or a target. 1 2016E Adjusted diluted EPS Thomson Reuters Consensus Estimate as used by Perrigo as of September 16, 2015. 2 Calculated by taking Thomson Reuters 2016E adjusted diluted EPS consensus estimate of \$9.30 and subtracting \$0.64, calculated using post-tax income of \$94mm (as per slide 32 of Perrigo's investor presentation dated October 22, 2015) assuming 147mm of Perrigo diluted shares outstanding. 3 Based on Perrigo and Omega public filings except for Q1 2015 Branded Consumer, which is based on Wall Street Research. Pro forma for Omega acquisition assuming a January 1, 2014 close; actual transaction close occurred on March 30, 2015. 4 As used by Perrigo in its September 17, 2015 Investor Presentation titled "Responding to Mylan's Inadequate Tender Offer," page 29. 5 Reflects estimated U.S. MS market based upon Wall Street Research. Figures in thousands. 6 Assumes ~\$314mm net income impact of Specialty Sciences in CY2015. Net income impact calculated as post tax adjusted EBITDA assuming 1% tax rate of Specialty Sciences and FY15 revenue contribution of Specialty Sciences applied to CY15 revenue for Perrigo based on midpoint of company guidance. Adjusted diluted EPS impact of Specialty Sciences assumes Perrigo standalone adjusted diluted EPS of \$7.75 (midpoint of 2015E adjusted diluted EPS company guidance) and Perrigo share count of 144mm (based on company guidance). Nothing on this slide is intended to be a profit forecast or a target.

Significant Underperformance Deceleration of Consumer Facing Business Concerns Regarding Tysabri Sustainability
 Sustainability
 5 New EPS Guidance Relies on One-Time Cost Cuts, Financial Engineering, and Masks
 Underperformance Declining Consumer Facing Business Comprises ~75%
 4 of Perrigo's Revenue Tysabri, Which Goes Off-Patent By 2023, Comprises ~28%
 6 of Perrigo's Adj. Diluted EPS U.S. Patients on Tysabri (000s) 2010-2014 CAGR: (4.2)%
 Perrigo Revenue (Incl. Acquisitions) 2014 YTD 2015 YTD % Change
 Consumer Healthcare \$ 2,108 \$ 2,107 (0.1)%
 Branded Consumer 1,277 1,088 (15)%
 Rx (Incl. Acquisitions) 671 790 18 %
 Specialty Sciences 231 250 8 %
 Other and API 89 75 (16)%
 Previous Guidance / Consensus Revised Guidance
 2015 Sales (Mgmt. Guidance) (\$bn) \$5.4-\$5.7 \$5.3-\$5.45
 Adjusted Diluted Consensus 2016E EPS, Ex-new Initiatives \$8.901 \$8.662
 Declining Tysabri usage Failed phase 3 trial for SPMS Any off-label SPMS usage may be at risk

Q3 2015 Highlights By Region *Total Mylan growth reflects the quarter-over-quarter comparison of total adjusted revenues. Segment and region growth reflects the quarter-over-quarter comparison of adjusted third-party net sales. All growth rates are stated on a constant currency basis. Adjusted metrics and constant currency growth rates are non-GAAP financial measures. Please see the Appendix for reconciliations of such non-GAAP financial measures to the most direct comparable GAAP financial measures. **EPD Business growth reflects the quarter-over-quarter comparison under Mylan in Q3 2015 versus Q3 2014 under Abbott Laboratories, on a constant currency basis. Q3 2014 Q3 2015 \$ in millions Adjusted Revenue* +36% +95% +29% +47% -5% Adjusted Revenue* Growth Legacy Mylan EPD Business** North American Generics +24% +3% European Generics +6% +3% ROW Generics +21% +14% Specialty -5% N/A Total Mylan +14% +5%

Continued Execution of Our Growth Drivers *CCG = clinic commission group Sirdupla - Sirdupla™ pMDI continues to perform and has received majority endorsement from UK CCG* Revefenacin - Entered into Phase III clinical program expected to complete in Q3 2016 Generic Flovent® / Flixotide® - Completed pilot pharmacokinetic studies for US program. Progressing with manufacturing activities; generic Flixotide® is on track to be approved in Q1 2016 in the EU Generic Advair® / Seretide® Formulation developed to be qualitatively and quantitatively the same as RLD Completed our clinical endpoint study and have shown product to meet all draft guideline criteria Human Factors Studies demonstrate our DPI device can be used as successfully with new patients as those on RLD device Two PK studies complete with final PK study ongoing and will support our December ANDA submission Recently completed a collaborative pre-ANDA meeting with the FDA Respiratory Biologics & Insulin Analogs Robust portfolios with six biosimilars and three insulin analogs in active development, in collaboration with Biocon Plan to file three biosimilar applications and an application for an interchangeable glargine (US) during 2016 Trastuzumab - Completed enrollment of our Phase III study. Currently commercialized Hertraz™ in ten countries with multiple new launches planned in 2016 Pegfilgrastim - Completed Phase I clinical trial and enrollment in our Phase III trials Adalimumab - Phase I clinical trial completed and have initiated our Phase III clinical program Glargine - Completed recruitment for both the Type 1 and the Type 2 diabetes trials. Continue to pursue our discussions with FDA regarding interchangeability. Completion and qualification of state-of-the-art facility in Malaysia and activities to transfer product into that facility

Source: Perrigo Annual Reports on Form 10-K filed August 13, 2015 and August 14, 2014, Perrigo Company (“Perrigo Co”) Annual Reports on Form 10-K filed on August 15, 2013 and August 16, 2012, and Perrigo investor presentation dated September 17, 2015, titled “Responding to Mylan’s Inadequate Tender Offer” Mylan Well Positioned to Integrate Complementary Businesses, Aligned With Its Core Competencies Private Label Manufacturing (Consumer Healthcare) US consumer healthcare contract manufacturing portfolio Mylan’s global supply chain and manufacturing platform represents a core competency of Mylan with the highest level of operational excellence serving the same customer base Omega (Branded Consumer Healthcare) Legacy Omega operations Mylan’s established commercial platform in Europe (Rx and Gx) in both the physician and retail channels allows Mylan to optimize Omega’s OTC product portfolio Generics (Rx Pharmaceuticals) Prescription generic pharmaceuticals business Mylan has been a generics leader for decades and is well-equipped to enhance Perrigo’s prescription portfolio and its specialty sales force Non-Core Royalty Asset (Specialty Sciences) Primarily the Tysabri royalty stream from the Elan acquisition Mylan could maximize the use of cash from this asset by better reinvesting in the business API (Other) Can be effectively integrated with Mylan’s existing API business, which includes sourcing within our internal network and external customers Perrigo Segment Overview:

Mylan Today: Global and Scalable Supply Chain Ability and agility to respond to large and small volume orders with short lead times Manufacturing & Supply Capabilities Serving Market Needs Global Supply Chain and Operational Excellence More than 40 internal manufacturing sites more than 1,400 3P suppliers and CMOs Broad range of dosage forms: tablets, capsules, powders, injectables, aerosols, patches, gums, creams, liquids and ointments Manufacturing strategically located to support markets Vendor managed inventory for select accounts Ship to more than 35,000 customers Delivery of 56 billion doses to patients annually Direction of 55+ distribution centers Packaging, labeling and artwork meeting local requirements (language, design) 1,400+ products 15,000+ SKUs 145 markets ~40 languages

Effectively Leveraging our API Capabilities it's about how you deploy and leverage these technologies and assets and strategically focus on opportunities CAPACITIES More than 50% of our pipeline and key products are vertically integrated DMFs (molecules) Pipeline Molecules Total Molecules 249 42 291 ~3,700 Volume (in kiloliters) ~4,400 Volume (in metric tons) ~2,700 Volume (in kiloliters) ~8,400 Volume (in metric tons) ~7,300 Total Volume (in kiloliters) 12 Contract Manufacturing Facilities (for APIs) 24 Contract Manufacturing Facilities (for intermediates) 9 API Sites ~900 Volume (in kiloliters) ~1,200 Volume (in metric tons) ~14,000 Total Volume (in metric tons)
PRODUCTS VERTICAL INTEGRATION OPPORTUNITIES

Operational Excellence with Our Network of 33 Internal Finished Dosage Form Sites 12 3 Complex Products Sites 18 Oral Solid Dose Sites* Injectables Sites It's not just having the capabilities, it's about how we leverage and how we deploy these assets Tablets Capsules Injectables Ointments Creams Scientists & Regulatory Professionals Product Pipeline ~3,000 ~4,100 STRONG R&D PLATFORM* DIVERSE TECHNOLOGIES OPPORTUNITIES Transdermals Inhalers Nasals Extruder/ Beads 65B capacity Ability to produce up to 500m units Capacity to manufacture 260M patches, 15M creams gels, building DPI capacity *Including FamyCare (pending deal closure)

60 Day Plan Ranjan Chaudhuri will join Mylan as the commercial lead for our OTC business. Ranjan has extensive experience in this space, most recently in the smoking reduction and cessation category, leading the recent divestiture of this business to Perrigo. Created a Governance team including an interim CFO as well as Commercial, Operations, HR, OTC, Compliance, Legal, Security, IT, and Communications Leads Third party advisors have been identified to partner with and support the integration and OTC commercial execution on Day 1 Leverage Mylan's existing Integration Office First 60 Days Pre-Close Activities Underway If needed, management team prepared to be deployed to Allegan, MI immediately Get to know key talent and engage Perrigo and Omega management and employees in our due diligence efforts and share our vision for the combined entities and their role in the success as well as opportunities for their future growth. Further due diligence to assess additional opportunities for the combined company Begin to develop integration and synergy realization roadmaps

Q3 2015 Financial Results & 2015 Guidance \$ millions, except EPS Q3 2015 Q-o-Q Growth 2015 Guidance Y-o-Y Growth*** Total Revenue* \$2,712 36% (cc)** \$9,600-\$10,100 33% (cc)** Gross Margin* 58% +400 bps 53%-55% +160 bps R&D as % of Revenue* 6.4% (110) bps 6.5%-7.5% (30) bps SG&A as % of Revenue* 18.2% (80) bps 19%-21% +70 bps EBITDA* \$987 34% \$2,900-\$3,300 31% Net Earnings* \$734 59% \$2,075-\$2,175 50% Diluted EPS* \$1.43 23% \$4.15-\$4.35 23% (cc)** Operating Cash Flow* \$1,125 139% \$1,600-\$1,800 40% Capital Expenditures \$85 27% \$350-\$450 23% Effective Tax Rate* 17% (800) bps 18% (700) bps Diluted Share Count 514 29% 495-500 25% *Adjusted metrics ** (cc) refers to constant currency ***Year-on-year growth is calculated from the midpoint of the 2015 guidance ranges Adjusted metrics and constant currency measures are non-GAAP financial measures. Please see the Appendix for reconciliations of such non-GAAP financial measures to the most directly comparable GAAP financial measures. Note: Quarter-over-Quarter (Q-o-Q) growth compares Q3 2015 actual results to Q3 2014 actual results.

Perrigo Implied Hypothetical Unaffected Stock Price Based on: Perrigo's Implied Hypothetical Unaffected Stock Price Mylan's Offer Has Been Supporting The Perrigo Share Price Perrigo Undisturbed 2016E P/E Multiple on April 7, 2015 18.3 x Change in Selected Peer Average 2016E P/E since April 7, 2015 (28.6)% Implied Perrigo Unaffected 2016E P/E 13.1 x Perrigo 2016E Adjusted Diluted EPS³ as of October 28, 2015 \$ 9.30 Implied Hypothetical Unaffected Perrigo Share Price \$121.83 Implied 2015E EV / EBITDA⁴ 13 x Perrigo Undisturbed 2016E P/E Multiple on April 7, 2015 18.3 x Change in Perrigo Proxy Peer Average 2016E P/E since April 7, 2015 (9.8)% Implied Perrigo Unaffected 2016E P/E 16.5 x Perrigo 2016E Adjusted Diluted EPS³ as of October 28, 2015 \$ 9.30 Implied Hypothetical Unaffected Perrigo Share Price \$ 153.45 Implied 2015E EV / EBITDA⁴ 15 x Perrigo Undisturbed Share Price (as of April 7, 2015) \$ 164.71 S&P Pharmaceuticals Index (since April 7, 2015) (19.0)% Implied Hypothetical Unaffected Perrigo Share Price \$ 133.42 Implied 2015E EV / EBITDA⁴ 14 x Change in Perrigo's Proxy Peers Average² 2016E P/E Multiple Share Price Indexed to S&P Pharmaceuticals Index Change in Selected Peer Average¹ 2016E P/E Multiple A B C Source: Thomson Reuters as of October 28, 2015 Note: Average hypothetical share price is based on the average of abovementioned three calculation methods outlined on this slide. Undisturbed share price and P/E are as of April 7, 2015. Average change in share price and changes in average P/E multiplies shown in A,B,C above are for the period starting April 7, 2015 and ending October 28, 2015. Other factors also impact Perrigo's share price and Mylan's offer should not be considered to be the sole factor impacting Perrigo's share price. Mylan does not endorse or adopt Thomson Reuters consensus estimates, which are used for illustrative purposes only. Nothing on this slide is intended to be a profit forecast or a target. 2016 P/E multiple calculated as share price divided by 2016E Thomson Reuters consensus estimate adjusted diluted EPS as of October 28, 2015. Adjusted diluted EPS is a non-GAAP measure. 1 Based on Selected Peer Average 2016E P/E Multiple to Thomson Reuters's Current 2016E Adjusted Diluted EPS Estimate for Perrigo. Selected peers consist of Valeant, Mallinckrodt, Endo, Jazz, Teva, Akorn and Mead Johnson. 2 Based on Perrigo's public Proxy Peers Average 2016E P/E Multiple to Thomson Reuters's Current 2016E Adjusted Diluted EPS Estimate for Perrigo. Perrigo's Peers per Perrigo's definitive proxy statement, filed September 25, 2015, and consist of Abbvie, Mallinckrodt, Actavis, Mead Johnson, Allergan, Mylan, Bristol-Myers Squibb, Regeneron, Celgene, Cubist, Shire, United Therapeutics, Endo, Valeant, Hospira, Zoetis, and Jazz Pharmaceuticals. Excludes Allergan due to sale of generics business to Teva as well as recently acquired Actavis, Cubist and Hospira. 3 Thomson Reuters consensus estimate as of October 28, 2015. Mylan does not endorse or adopt Thomson Reuters consensus estimates, which are used for illustrative purposes only. Nothing on this slide is intended to be a profit forecast or a target. Adjusted diluted EPS and adjusted EBITDA are non-GAAP financial measures. 4 Calculated as $[(HUSP * \text{diluted shares outstanding}) + \text{net debt}] / 2015E \text{ adjusted EBITDA}$. Reflects CY2015E Perrigo adjusted EBITDA of \$1.6bn per Thomson Reuters consensus estimates as of April 7, 2015 and October 28, 2015. Mylan does not endorse or adopt Thomson Reuters consensus estimates. Adjusted EBITDA is a non-GAAP financial measure. Cash and debt as of latest Perrigo Annual Report on Form 10-K filed August 13, 2015 of \$0.8bn and \$5.3bn respectively. Assumes 147mm diluted Perrigo shares outstanding.

Transaction is Immediately Accretive to Earnings and Value for the Perrigo Shareholders Perrigo Standalone Mylan / Perrigo (2.30 Shares + \$75 Cash) Mylan / Perrigo (3.97 Shares + \$0 Cash)⁴ 2016 Phased-in¹ 2016 Run-rate¹ 2016 Phased-in 2016 Run-rate 2016 Phased-in 2016 Run-rate Mylan Standalone Adj. Diluted EPS² \$ 4.72 \$ 4.72 \$ 4.72 \$ 4.72 Acc. / Dil. (Phased-In Synergies per Perrigo Presentation page 26) (12)% 0%⁵ (12)% 0%⁵ Implied Mylan PF Adj. Diluted EPS \$ 4.15 \$ 4.71 \$ 4.15 \$ 4.71 Exchange Ratio 2.30 x 2.30 x 3.97 x 3.97 x Adjusted Diluted EPS to Perrigo Shareholders \$ 9.45 \$ 9.83 \$ 9.55 \$ 10.83 \$ 16.48 \$ 18.70 Accretion to Perrigo 2016E Adjusted Diluted EPS per Consensus (\$9.30)³ 3 % 16 % 77 % 101 % 16.0 x \$ 151 \$ 157 \$ 228 \$ 248 \$ 264 \$ 299 14.0 x \$ 132 \$ 138 \$ 209 \$ 226 \$ 231 \$ 262 12.0 x \$ 113 \$ 118 \$ 190 \$ 205 \$ 198 \$ 224 Illustrative Value to Perrigo Shareholders at Various CY2016E P/E Multiples Source: Perrigo public filings and Thomson Reuters consensus estimates as used by Perrigo in its September 17, 2015 Investor Presentation and Perrigo Investor Presentation “Creating Value for Shareholders: Now and for the Long Term”, filed October 22, 2015. Mylan does not endorse or adopt Thomson Reuters consensus estimates, which are used for illustrative purposes only. This is a Pro-Forma estimate and indicative only and not a target or profit forecast. Nothing in this slide is intended to be a profit forecast. Pro-Forma values are illustrative only and any references to value per share, adjusted diluted EPS, share price and P/E should not be treated as targets or profit forecasts. Adjusted diluted EPS is a non-GAAP financial measure. Implied Mylan Pro-Forma adjusted diluted EPS is based on accretion / (dilution) and phasing in of synergies per Perrigo’s presentation released September 17, 2015, titled “Responding to Mylan’s Inadequate Tender Offer.” Assumes phased-in synergies for the combined company per September 17, 2015 Perrigo Investor presentation. Full run-rate synergies for the combined company are not expected to be realized until the end of year four following the consummation of the offer. Implied Mylan / Perrigo PF adjusted diluted EPS assumes that Mylan acquires 100% of Perrigo ordinary shares in the offer and that Mylan shareholders realize the benefit of all synergies realized in the transaction. ¹ Per Perrigo Investor Presentation “Creating Value for Shareholders: Now and for the Long Term”, filed October 22, 2015. ² Thomson Reuters Consensus Estimate as Used by Perrigo as of September 16, 2015 ³ Accretion to Perrigo shareholders calculated comparing pro forma adjusted diluted EPS to Perrigo shareholders to standalone Perrigo adjusted diluted EPS per Thomson Reuters as of October 28, 2015. ⁴ Assumes \$75 / share is received by Perrigo shareholders is reinvested at Mylan current share price of \$44.89 as of October 28, 2015. ⁵ Run-rate synergies implied based on difference between “No Synergies” and “Ramped Synergies” of 25% per Perrigo’s presentation filed September 17, 2015. With no synergies, 16% dilution on \$4.72 of 2016E EPS implies \$3.96 PF adjusted diluted EPS. With ramped synergies, 12% dilution on \$4.72 of 2016E adjusted diluted EPS implies \$4.15 PF adjusted diluted EPS. Thus the 25% synergy ramp produced $(\$4.15 - \$3.96) = \$0.19$ of earnings. If \$0.19 of earnings is 25% of run-rate synergies, then implied 2016E run-rate synergies is $(4 * \$0.19) = \0.74 of earnings. 2016E adjusted diluted EPS with run-rate synergies then are $(\$3.96 + \$0.74) = \$4.71$ 2016E adjusted diluted EPS.

Bottom Line: The MATH is a Clear and Compelling Value Proposition Relative to Standalone Plan 1 No Deal Mylan Offer 2 Mylan / Perrigo (2.30 Shares + \$75 Cash) Mylan / Perrigo (3.97 Shares + \$0 Cash)³ Illustrative Standalone P/E 14 x 16 x Standalone Value Per Perrigo Share¹ \$9.45 \$ 132 \$ 151 Illustrative Mylan / Perrigo 2016 P/E 12 x 14 x Total Value Per Perrigo Share² \$9.55 \$ 190 \$ 209 Illustrative Mylan / Perrigo 2016 P/E 12 x 14 x Total Value Per Perrigo Share² \$16.48 \$198 \$231 Perrigo Standalone Source: Perrigo public filings and Thomson Reuters consensus estimates as used by Perrigo in its September 17, 2015 Investor Presentation and Perrigo Investor Presentation “Creating Value for Shareholders: Now and for the Long Term”, filed October 22, 2015. Mylan does not endorse or adopt Thomson Reuters consensus estimates, which are used for illustrative purposes only. This is a Pro-Forma estimate and indicative only and not a target or profit forecast. Nothing in this slide is intended to be a profit forecast. Pro-Forma values are illustrative only and any references to value per share, adjusted diluted EPS, share price and P/E should not be treated as targets or profit forecasts. Adjusted diluted EPS is a non-GAAP financial measure. Mylan Pro-Forma adjusted diluted EPS based on accretion / (dilution) and phasing in of synergies per Perrigo’s presentation released September 17, 2015, titled “Responding to Mylan’s Inadequate Tender Offer.” Assumes phased-in synergies per Perrigo presentation. See slide 17 for more information on anticipated synergies from the transaction. 1 Calculation is based on 2016E adjusted diluted EPS of \$9.45 for Perrigo assuming phased-in synergies as used by Perrigo in Perrigo Investor Presentation “Creating Value for Shareholders: Now and for the Long Term”, filed October 22, 2015. Please refer to slide 31 of Perrigo’s investor presentation dated October 22, 2015. 2 Calculation is based on 2016E adjusted diluted EPS Thomson Reuters Consensus Estimate as used by Perrigo as of September 16, 2015. 3 Assumes \$75 / share is received by Perrigo shareholders is reinvested at Mylan current share price of \$44.89 as of October 28, 2015.

Appendix

Reconciliation of Non-GAAP Metrics

Reconciliation of Non-GAAP Metrics Below is a reconciliation of GAAP net earnings attributable to Mylan N.V. to EBITDA and adjusted EBITDA for the three and nine month period compared to the respective prior year period:

Reconciliation of Non-GAAP Metrics Below are reconciliations of key non-GAAP financial metrics for the three and nine month period compared to the respective prior year period. The non-GAAP financial metrics are presented in order to supplement investors and other readers' understanding and assessment of company financial performance.

Reconciliation of Non-GAAP Metrics

Reconciliation of Non-GAAP Metrics

Summary of Adjusted Total Revenues by Segment

Perrigo: Reconciliation of Adjusted EBITDA Margin Source: Perrigo Annual Reports on Form 10-K filed August 13, 2015 and August 14, 2014, Perrigo Co.'s Annual Reports on Form 10-K filed August 15, 2013 and August 16, 2012, and Perrigo investor presentation dated September 17, 2015, titled "Responding to Mylan's Inadequate Tender Offer" ¹
Amortization of acquired intangible assets related to business combinations and asset acquisitions

Reconciliation of Forecasted Guidance

