

Mallinckrodt Pharmaceuticals

Investor Briefing 2017 Presentation Hospital Portfolio

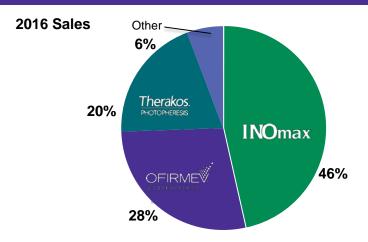
Ron Lloyd EVP and President, Hospital Therapies

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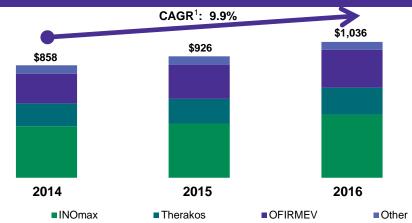


Mallinckrodt is becoming major hospital pharmaceutical player

Mallinckrodt's Hospital Products



MNK Hospital Net Sales† Have Grown Steadily (\$MM)



Note: All data in calendar years

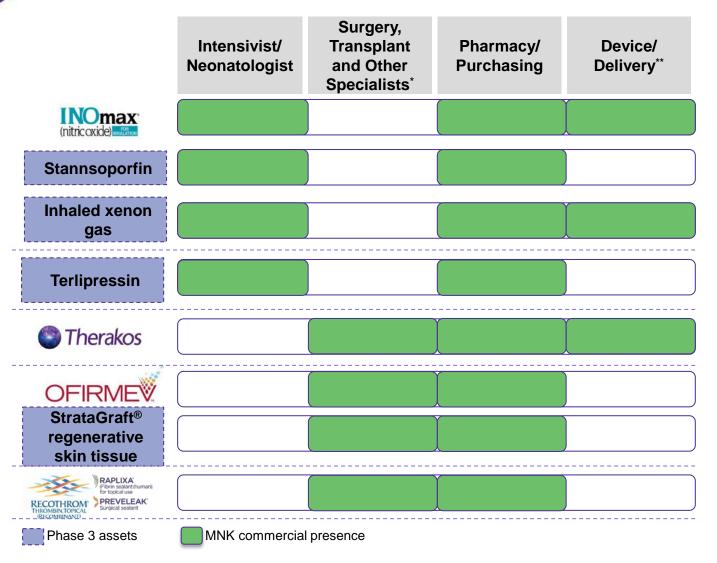
† Pro forma sales; historical net sales includes revenues prior to MNK ownership Net sales include \$663 million in 2014, \$322 million in 2015 and \$3 million in 2016, representing net sales of acquired products prior to Mallinckrodt's ownership

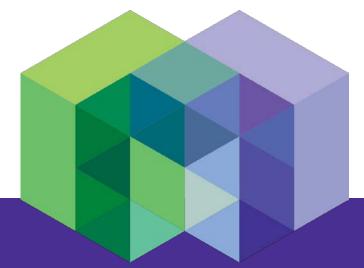
Mallinckrodt's Hospital Focus

- Growing presence in fragmented hospital market; one of few companies with hospital focus
- Category-leading products; hospital net sales >\$1 billion
- Emerging pipeline of first-in-class products to address areas of high unmet needs
- Reach and experience to effectively develop and commercialize hospital products
 - Established commercial presence in U.S., EU, Japan, Australia and Canada
 - Unique drug/device expertise, service infrastructure
 - Decision-maker access with deep relationships and proven ability to execute unique, value-based contracts

Source: IMS sales of non-biologic brands and branded generics; MNK net sales include OFIRMEV® (acetaminophen) injection, INOMAX® (nitric oxide) gas, for inhalation, Therakos® immunology platform and other hospital products

Portfolio of complementary, innovative products and established hospital touch points





- Broad presence and hospital focus makes
 Mallinckrodt a compelling partner for companies looking to license out developmental or marketed assets
- Strong HEOR¹ capabilities and hospital/GPO² contracting experience allows MNK to maximize value of transformational therapies

Therakos approved for use in transplant and other indications in EU

^{**} Includes respiratory therapists, specialty nurses and apheresis specialists



is gold standard for treatment of HRF¹ in term/near-term neonates



- INOmax® is Mallinckrodt's 2nd largest product with sales of ~\$500 million
- >55% of 22,000² U.S. babies born annually with HRF treated with INOmax
- Ex-U.S. footprint covers Japan, Canada, Australia and other markets
- Multi-pronged durability based on:
 - Regulatory requirements compliance
 - Intellectual property
 - Strong, 15-year market acceptance built on patient safety dedication, unmatched customer service and continuous investment in innovation
- INOmax DS_{IR} ® Plus delivery device is most advanced system in the world
 - Designed with key safety features needed for fragile infants
 - Installed base of over 4,000 devices in hospitals across the U.S.
- INOmax offered as a total service package





offering is a high-touch, full-service model

Broad customer base in Field teams to engage and 100% of U.S. Level IIIC train neonatologists, nurses and Level IV NICUs1 and respiratory therapists ~800 Hospitals in U.S. 5 U.S. Regional 24/7 Customer Care **Service Centers** ~200 Hospitals ex-U.S. **Customer Care** team understands and responds to customer needs **Manufacturing**

Sales and Marketing

INOMAX SERVICE MODEL INCLUDES:

- Drug
- Device use (+ backups)
- Training
- Emergency delivery (≤4 hours)
- 24-hour customer service
- Device servicing
- Free upgrades
- Unlimited supplies
- Reimbursement support

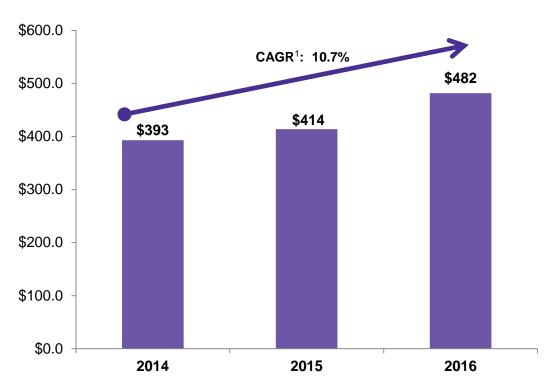


INOmax growth drivers: execute on strengthening value proposition to customers

Calendar Year Net Sales[†] (\$MM)



Growth Drivers



Enhance customer experience Continue to deliver and ensure customers understand the full value of our comprehensive service model

Provide flexible contracting

Allow customers flexibility to choose pricing option that best fits their needs

Advance device technology Develop next-generation, compact, portable device aimed at further enhancing safety and simplifying use, with flexibility to use in a number of settings



Note: All data in calendar years

† Pro forma sales; historical net sales may include revenues prior to MNK ownership

Includes net sales prior to the acquisition of INOmax (on April 16, 2015) of \$118 million for calendar 2015 that are excluded from Mallinckrodt's historical financial information.



Therakos photopheresis platform is only U.S.-approved ECP² system for CTCL^{1*} patients



- Non-Hodgkin's CTCL presents as red, scaly patches or thickened plaques of skin
- Therakos can effectively treats skin manifestations of CTCL
 - Available >30 years
 - Over 1 million treatments delivered worldwide
- ECP² generally used in patients whose disease has progressed beyond early stages
 - Therakos' estimated use in ~10% of ~7,500 eligible U.S. patients with systemic CTCL
 - Available primarily at major MSA³ academic centers
- Therakos has broad European label covering CTCL, GVHD⁴ and other immune-modulating diseases
- Pursuing opportunities to expand into other geographies, including Japan



¹ Cutaneous T-Cell Lymphoma

² Extracorporeal Photopheresis

³ Metropolitan Statistical Area

⁴ Graft versus Host Disease

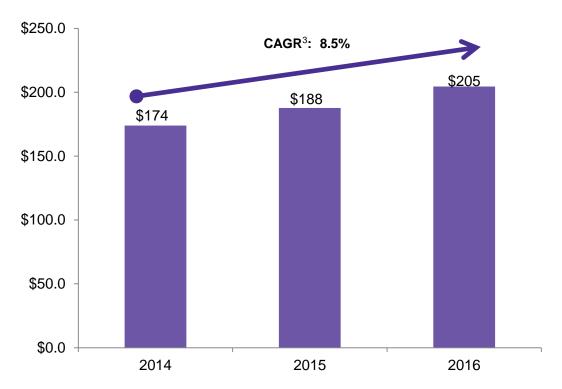


Therakos growth drivers: new device placements, strong clinical and patient focus, and expanded access

Calendar Year Net Sales[†] (\$MM)



Growth Drivers



Conversion from XTS® to **CELLEX®** device

- Complete customer conversion to advanced CELLEX devices
- Normalized kit inventory levels

Drive use in key indications Increase U.S. CTCL^{2*} use via clinical focus and patient education campaign

Expand access in key markets

Enhance European reimbursement across broad label with HEOR1 and clinical data

Note: All data in calendar years

† Pro forma sales; historical net sales may include revenues prior to MNK ownership

Includes net sales prior to the acquisition of Therakos (on September 28, 2015) of \$138 million for calendar 2015 that are excluded from Mallinckrodt's historical financial information.

- 1 Health Economics and Outcome Research
- 2 Cutaneous T-Cell Lymphoma
- 3 Compound Annual Growth Rate
- * All references to CTCL are intended to refer to skin manifestations of CTCL.



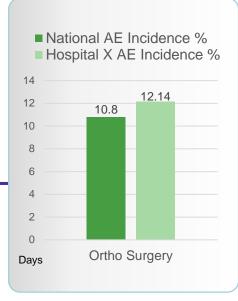
is leading branded IV analgesic in U.S.

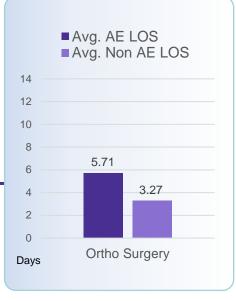


- Offers physicians non-opioid analgesic as part of multimodal analgesia (MMA) protocol
 - MMA widely recommended by nationally recognized organizations including The Joint Commission, American Society of Anesthesiologists and American College of Surgery
- ~15% market share of patients in targeted surgical procedures¹
- Can reduce length of stay and hospital costs per HEOR² data
 - Potential \$4.7MM annual savings in medium-sized hospital³
- Unmet need still exists: hospital-level Medicare data shows link between opioid-associated adverse events and longer hospital stays⁴









¹ Includes patients with analgesics undergoing bariatric, colorectal, joint replacement, spine, cardiothoracic, hysterectomy, C-sections, and general surgery; Source: Company analysis of Premier DB based on 2015 Inpatient Discharges, HCUP NIS Data

² Health Economics and Outcomes Research

³ Shaffer, Advanced Therapy, Nov. 2016

⁴ Use of Ofirmev has an attendant decrease in opioid consumption, the clinical benefit of which was not demonstrated in clinical trials

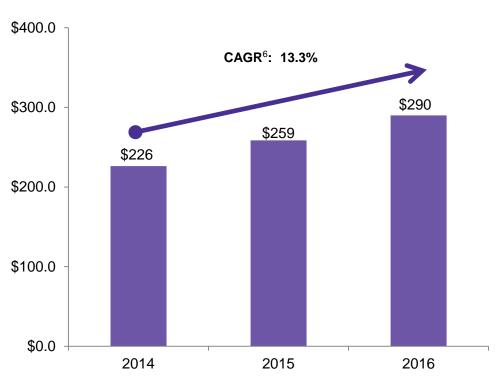


growth drivers: educate customers on OFIRMEV value proposition

Calendar Year Revenue[†] (\$MM)



Growth Drivers



Highlight true IV opioid costs with CMS¹ data

 Focus on hospital-level, Medicare data showing link between adverse events associated with opioids and longer hospital stays⁵

Demonstrate
OFIRMEV's
value
proposition

 Demonstrate reduced length of stay, reduced discharges to skilled nursing facilities and improved hospital costs through HEOR² data

Drive innovative contracting

- Leverage innovative contracting to pull-through OFIRMEV's value proposition
- Major contract wins with GPOs³ and IDNs⁴

Note: All data in calendar years

Includes net sales prior to the acquisition of Ofirmev (on March 19, 2014) of \$31 million for calendar 2014 that are excluded from Mallinckrodt's historical financial information.

[†] Pro forma sales; historical net sales may include revenues prior to MNK ownership

¹ Centers for Medicare and Medicaid Studies

² Health Economics and Outcome Research

³ Group Purchasing Organization

⁴ Integrated Delivery Network

⁵ Use of Ofirmev has an attendant decrease in opioid consumption, the clinical benefit of which was not demonstrated in clinical trials 6 Compound Annual Growth Rate

Longer-term growth: build pipeline of first-in-class products for critically ill patients in areas with high unmet need

Stannsoporfin

- Treatment of severe bilirubin (jaundice) when phototherapy is insufficient
- Extended levels of high bilirubin can impact neurological development
- No currently approved pharmacological therapies; exchange blood transfusion is last-resort option



Neonate with high bilirubin counts manifested as jaundice



StrataGraft



Autograft

StrataGraft regenerative skin tissue

- Treatment of burn patients with off the shelf treatment
- Helps avoid autografts the removal and reuse of skin from another part of patient's body
- Among first products designated a Regenerative Medicine Advanced Therapy (RMAT) by the FDA

Terlipressin

- Treatment of hepatorenal syndrome (HRS) Type 1, a serious consequence of cirrhosis with high mortality rates
- No U.S.-approved therapies; terlipressin is approved in several other countries for HRS¹ and other conditions



HRS-Type 1 patient depicting liver failure with ascites



Patient undergoing resuscitation after cardiac arrest

Inhaled xenon gas drug-device

- Treatment of resuscitated cardiac arrest patients' to improve survival and functional outcomes
- No approved pharmacological therapies; hypothermia most commonly used therapy

Expected near-term launches can contribute significantly to strategic plan growth

| Product | Proposed Indication | Market Overview | Expected Launch | Loss of Exclusivity | Est. Global Peak Net Sales ¹ |
|-------------------|--|---|--------------------|------------------------|---|
| Stannsoporfin | Reduction of bilirubin levels in neonates | ~70,000 to 125,000 babies annually in the U.S. with severe jaundice and resistance to phototherapy or readmitted | 2018 | 2032 | >\$125mm |
| StrataGraft | Treatment of partial thickness and full thickness burns | ~10,000 hospitalized burn patients annually in U.S. require autografting | 2020 | 2032 | >\$125mm |
| Terlipressin | Reduce mortality and time to discharge in Hepatorenal Syndrome (HRS) Type1 | ~10,000 to as many as ~30,000 in U.S. are impacted by HRS Type 1, with a larger population impacted by HRS Type 2 | 2020 | 2027 | >\$300mm |
| Inhaled xenon gas | Improve neurological function after cardiac arrest | ~200,000 patients annually in U.S. are resuscitated after cardiac arrest | 2020 | 2027 | >\$125mm |

Mallinckrodt has built strong hospital presence – well positioned for the future

 Building a growing product portfolio, with over \$1 billion in hospital net sales

 Developing and expanding unique capabilities required to succeed: deep relations across key hospital decision makers, unique drug-device expertise, value-based contracting, and global scale

 Capitalizing on deep pipeline with the potential for four launches in next three years

 Becoming a leading hospital pharmaceutical company for critical care patients, and the leader in the neonatology and NICU¹ space



Thank You

