



Company Overview September 2012

Disclaimer

This presentation, including information contained in this disclaimer, is given to you in strict confidence. By attending the presentation, you agree that no part of this presentation or disclaimer may be disclosed, distributed or reproduced to any third party without the consent of QRxPharma Limited ("QRxPharma").

This presentation is being provided for the sole purpose of providing the recipients with background information about QRxPharma's business. This presentation, including the information contained in this disclaimer, does not constitute an offer, invitation or recommendation to subscribe for or purchase any security and neither the presentation, disclaimer nor anything contained in them forms the basis of any contract or commitment. This presentation does not purport to summarize all information that an investor should consider when making an investment decision. It should be read in conjunction with QRxPharma's other continuous disclosure announcements lodged with the ASX which are available at www.asx.com.au. Before making an investment decision you should consider whether it is suitable for you in light of your own investment profile and objectives and financial circumstances and the merits and risk involved.

No representation, express or implied, is made as to the fairness, accuracy, completeness or correctness of information, opinions and conclusions contained in this presentation, including the accuracy, likelihood of achievement or reasonableness of any forecasts, prospects, returns or statements in relation to future matters contained in the presentation ("forward-looking statements"). Such forward-looking statements are by their nature subject to significant uncertainties and contingencies and are based on a number of estimates and assumptions that are subject to change (and in many cases are outside the control of QRxPharma and its Directors) which may cause the actual results or performance of QRxPharma to be materially different from any future results or performance expressed or implied by such forward-looking statements. Forward-looking statements are provided as a general guide only and should not be relied upon as an indication or guarantee of future performance.

To the maximum extent permitted by law, neither QRxPharma nor its related corporations, directors, employees or agents, nor any other person, accepts any liability, including, without limitation, any liability arising from fault or negligence, for any loss arising from the use of this presentation or its contents or otherwise arising in connection with it.

You represent and confirm by attending and/or retaining this presentation, that you accept the above conditions.





Morphine + Oxycodone



QRxPharma Snapshot

- Australian based pain drug developer with offices in Sydney and Bedminster, NJ (ASX:QRX, OTCQX:QRXPY)
- Comprehensive portfolio around MOXDUO product line
- Blockbuster potential: global opioid market estimated at \$14 billion annually¹
- U.S. commercialisation partnership with Actavis, Inc.
- MOXDUO Complete Response Letter (CRL) issued by the U.S. FDA in June 2012; Productive Review Meeting in August 2012
 - Additional data package to FDA 2H, 2012; NDA refiling with FDA decision anticipated mid-2013



Solid Foundation for Growth

- Multiple formulations for complete "hospital-to-home" care; expected patent exclusivity through 2029
- MOXDUO delivers equal or better pain relief with fewer side effects than current treatments
- Potential changes in US regulatory policy are favourable for commercialisation of MOXDUO IR
- Double digit royalties on US sales of MOXDUO IR
- Experienced management team and Board of Directors
- KOL confidence in MOXDUO IR as a potential therapeutic option; Company commitment to bringing product to market



MOXDUO Product Portfolio From Hospital to Home

| | MOXDUO® IR | MOXDUO [®] CR | MOXDUO [®] IV |
|--------------|--|---|---|
| Delivery | Immediate Release | Controlled Release | Intravenous |
| Target | Moderate to severe acute pain | Chronic pain (i.e. osteoarthritis, back, neuropathic) | Hospital based: moderate to severe acute pain |
| Formulation | Oral Capsule | Oral tablet w/abuse deterrent | Injectable |
| Partnerships | Actavis, Inc. US commercialisation | | Aoxing Pharmaceuticals Strategic Alliance |
| Status | Additional Data Package and NDA re-filing | Phase 1 Complete | Phase 2; concurrent formulation development |





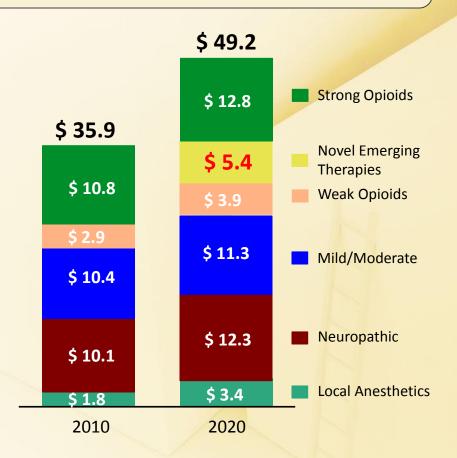
THE PAIN MARKET



Current Global Pain Market

Drug Class Sales for Pain in Major Pharmaceutical Markets, 2010 – 2020 (US\$ billions) ¹

- Large market opportunity: US\$14 billion¹
- Opioids are the "gold standard"
- Limited product innovation
- Strong opioids forecasted to maintain sales dominance through 2020 (aging population)
- Payors and KOLs: "need for better pain relief with fewer side effects"



Acute Pain Market AT-A-GLANCE



US Market Opportunity

- \$2B+ US Market with 210M+ annual Rxs (CAGR of 5-6%)
- Acute pain affects 75M Americans
- Limited product innovation; regulatory hurdles for new therapies
- Limited branded competition expected near-term

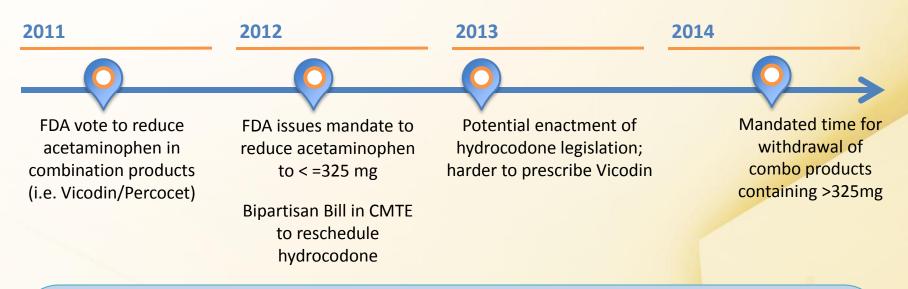
Clinical Unmet Needs

- Inadequate postoperative pain management
- Reduction in opioid-related AEs, specifically GI & CNS that limit their use
- Existing acute pain drugs associated with hepatic and GI toxicities

In the US, 6+ prescriptions are written for an acute opioid every second.



Proposed U.S. Regulatory Changes



- FDA mandated lower strength opioid/APAP combos will decrease efficacy and increase number of patients needing acute pain medicine
- Potential rescheduling of Vicodin to Schedule 2 will make it harder to prescribe and decrease number of prescriptions
- Creates void of about 50% of acute pain market (100 million Rx's)



Disrupted Acute Pain Market Provides MOXDUO IR Opportunity

2010 US Prescription Market Share for Acute Pain Opioids







IMMEDIATE RELEASE



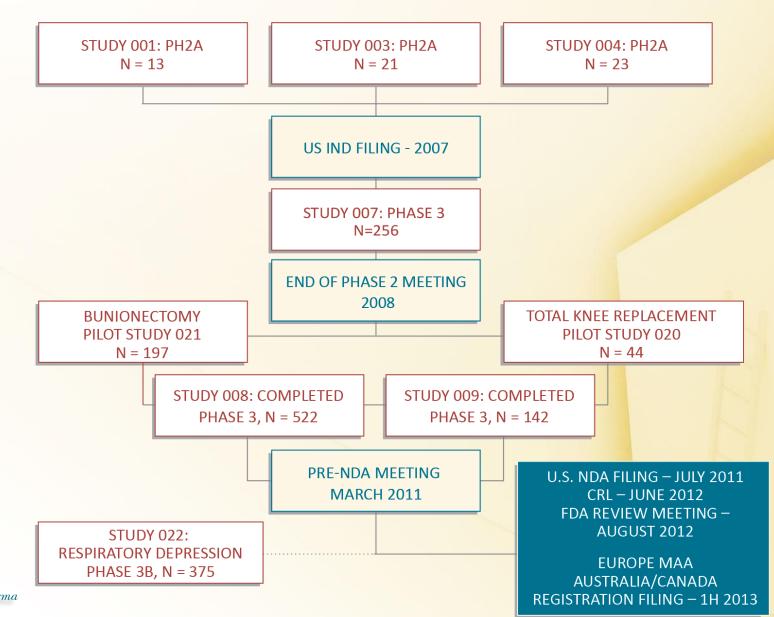
MOXDUO IR Product Profile

Key Features

- First line therapy for the treatment of moderate to severe acute pain
- Only opioid-opioid combination product available
- Immediate release formulation of morphine and oxycodone in a fixed
 3:2 ratio in capsules of the strengths:
 - 3 mg/2 mg
 - 6 mg/4 mg
 - 9 mg/6 mg
 - 12 mg/8 mg
- Four to six hourly dosing
- Demonstrated reduction in the occurrence and intensity of clinically significant opioid-related side effects compared to morphine, oxycodone and Percocet[®]



MOXDUO IR Clinical Development Path



Key Trial Conclusions

- Bunionectomy Trials: Pilot 021 & Pivotal 008 (n=719)
 - Met primary analgesic efficacy endpoint vs. morphine and oxycodone
 - MOXDUO IR proven superior to components on efficacy measures
 - Consistent safety advantage of MOXDUO IR
 - Pilot: 50-75% lower frequency of moderate to severe nausea, vomiting & dizziness compared to equi-analgesic doses of morphine or oxycodone
 - Phase 3: Despite higher dose and better pain relief of MOXDUO than morphine or oxycodone, AE rate and duration not statistically different

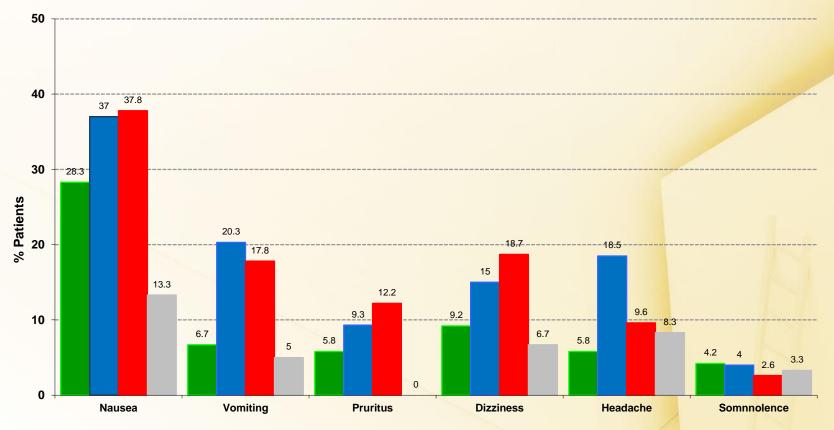


Key Trial Conclusions

- Total Knee Replacement Trials: Pilot 020 & Pivotal 009 (n=186)
 - Met all primary analgesic efficacy endpoints vs. Percocet
 - Pilot: MOXDUO superior to Percocet
 - Pivotal: MOXDUO High Dose better pain relief than low dose
 - Frequency of AEs much lower than Percocet
 - Significant pharmacoeconomic benefit: improved time to walk, sleep, etc.



Equi-analgesic doses of MOXDUO, Morphine, Oxycodone, vs. Placebo – All Studies

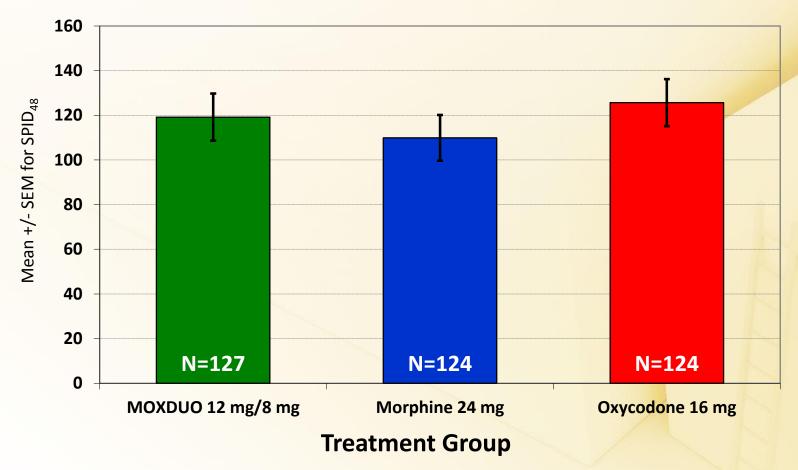


Opioid-Associated Adverse Event

MOXDUO Morphine Oxycodone Placebo



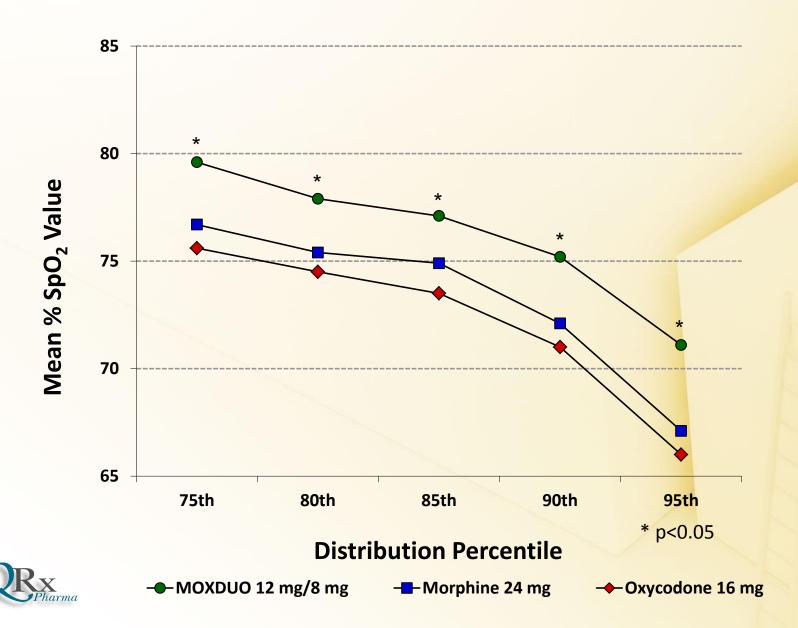
Respiratory Study 022 SPID₄₈ Equi-analgesic Doses





Source: Table 14.2.1.4

MOXDUO Produces Significantly Less Severe Oxygen Desaturation Study 022 Mean SpO₂ Values



MOXDUO Safety Advantage

STUDIES OF COMBINED OPIOIDS CONSISTENTLY PROVIDE EQUIVALENT EFFICACY WITH A SIGNIFICANT REDUCTION IN OPIOID-RELATED MODERATE TO SEVERE ADVERSE EVENTS

| QRx STUDIES | | |
|---|---|---|
| MOXDUO IR Study 022 (bunionectomy) | > | Oxygen desaturation less severe and of shorter duration compared to equianalgesic doses of Morphine or Oxycodone |
| MOXDUO IR Study 021 (bunionectomy) | > | 50-75% reduction in moderate to severe nausea, vomiting and dizziness compared to equianalgesic doses of Morphine and Oxycodone |
| MOXDUO IR Study 020 (knee replacement) | > | 100% reduction in moderate to severe nausea and emesis compared to the Percocet |
| MOXDUO IV Study (hip replacement) | > | 35% reduction in nausea and 38% reduction in emesis Compared to IV Morphine |
| MOXDUO Two Phase 2 trials in Australia (chronic pain) | | 34-40% decrease in the amount of drug to achieve equianalgesia compared to oral morphine. Decreased rate of drowsiness, dizziness, constipation and nausea. |

INDEPENDENT STUDIES¹

| Blumenthal et al 2007 (Spinal discectomy) | > | 80-100% reduction in nausea and emesis compared to PCA Morphine |
|---|---|--|
| Jamison et al 1998 (Chronic low back pain) | > | 17-49% reduction in intensity of a range of adverse events compared to Oxycodone |
| Lauretti et al 2004 (Cancer pain) | > | 86% reduction in nausea and 100% reduction in emesis compared to Morphine. |



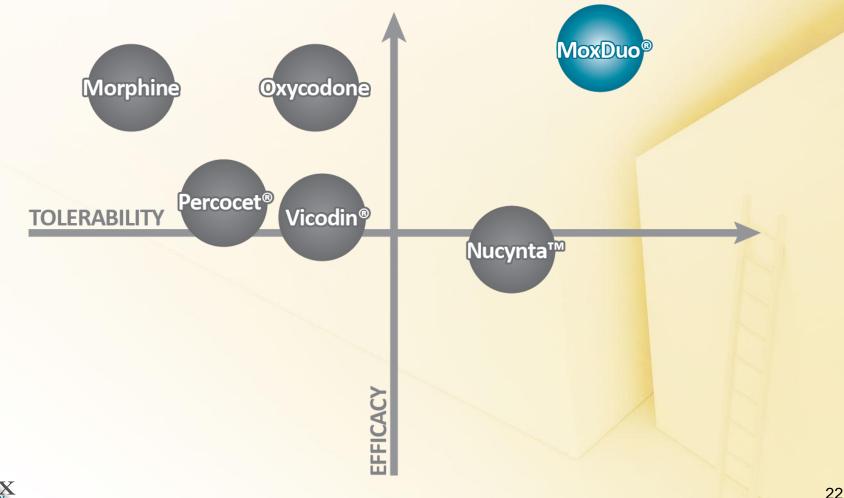
MOXDUO Pharmacoeconomic Benefits

- Versus Percocet[®] TKR study, MOXDUO IR patients were out of bed faster, walked and slept better
- US study finds \$4,880 \$36,152¹ incremental costs in patients suffering GI side effects following treatment with IR opioids
 - Extended hospitalization, increased nursing care and re-admissions
- Reimbursers, Managed Care and KOL feedback
 - Significant benefit from decreasing hospitalization by as little as 4 hours or recovery room time by 20 minutes

MOXDUO's advantageous safety profile may improve patient recovery, decrease hospital time and lower total cost of care.



MOXDUO IR Value Proposition = Greater Tolerability + Equal/Better Analgesia





Immediate Release Commercialisation Plan



Actavis Strategic Partnership

- Exclusive US commercialisation and development rights for MOXDUO IR
 - Actavis pays all product launch, marketing and sales costs
- 10%-30% royalties based on net sales thresholds from launch
 - Except 50% royalty on \$150m of cumulative sales (starting from 3-6 months following product launch)
- QRxPharma retains a right to co-promotion/profit-share
 - Option to create sales force and provide up to 25% of the effective selling effort to US prescribers after first 12 months of launch
- QRxPharma retains ownership of MOXDUO IR outside US





- European based Actavis Group hf is a leading manufacturer of branded and generic opioids worldwide
 - Revenues of ~EUR 1.8 billion & 10,000 employees worldwide
 - US sales highlighted by Kadian[®] (extended release morphine)
- Watson Pharmaceuticals acquisition proposed in April 2012
 - Creates 3rd largest global generics company
 - ~\$8.0 billion projected 2012 pro forma combined revenue
 - Maintaining commitment to branded marketplace
- MOXDUO IR commercialisation preparation ongoing



Mitigating Reimbursement Risk

- Appropriate pricing, contracting and patient pull-through
- Advantaged market other acute pain products (Vicodin[®] and Percocet[®]) being reduced due to safety issues and potential rescheduling
- Potential for significant pharmacoeconomic benefits recognized by payers/KOLs
- Reimbursement strategy = Tier 3 Formulary
 - Insurance companies will not have to pay more
 - Customer co-pays are manageable





Pipeline Potential



MOXDUO CR (Controlled Release)

- Sustained release formulation to provide at least 12 hours of analgesia for moderate to severe chronic pain
- Abuse deterrent and tamper resistant features
- Phase 1 results showed:
 - High bioavailability and complete absorption
 - One fifth the variability of OxyContin
 - Lower peaks and higher troughs should lead to: better safety & lower side effects; better tolerability at higher doses
 - Will provide very stable plasma levels when given twice daily
 - Should be an effective once or twice daily treatment
- Current formulation will progress to Phase 2



MOXDUO: US Peak Sales Potential

| | MOXDUO IR | MOXDUO CR | MOXDUO IV |
|----------------------------|--|---|--|
| Market Size | ■ ~200 mm Rx (2012) | ■ ~34 mm Rx (2015) | ■ ~29 mm Rx (2014) ¹ |
| | Annual market growth 1.0% | Annual market growth 3.0% | Annual market growth 1.0% |
| | • QRx targets ~ 50% of market | • QRx targets 100% of market | • QRx targets 100% of market |
| Market Penetration | Initial share: 1.0% (2012) | Initial share: 1.4% (2015) | Initial share: 1.5% (2014) |
| | Peak share: 5.0% (2015) | Peak share: 13.9% (2020) | Peak share: 13.0% (2018) |
| Pricing | Initial price: \$112 based on 4 doses per day and 14 days of therapy | Initial Rx Price: \$180 based on 2 doses per day and 30 days of therapy | Initial price: \$32 based on 4 vials per day and 2 days of therapy |
| | Annual price increase: 5.0% | Annual price increase: 5.0% | Annual price increase: 5.0% |
| | ■Peak sales: ~\$680 mm | Peak net sales: ~\$1,300 mm | Peak net sales: ~\$150 mm |
| Blockbuster Opportunity | Paracetamol Limitation -Peak sales: ~\$1,350 mm | Oxycontin - \$3 billion/year - off patent in 2013, opening market | |
| | plus Vicodin Rescheduling - Peak sales: ~\$2,000 mm | for MOXDUO CR in 2015 | |





Company Overview



Leadership Team

Senior Management

- John Holaday, PhD (CEO)
- Ed Rudnic, PhD (COO)
- Chris Campbell (CFO)
- Richard Paul, MD (EVP Drug Development)
- Warren Stern, PhD (Clinical Consultant)
- Janette Dixon, PhD (VP Global BD)
- Patricia Richards, MD, PhD (CMO)

Board of Directors

- Peter Farrell, PhD Chairman (ResMed)
- Michael Quinn (Innovation Capital)
- Peter Campbell (Sonic Healthcare)
- Gary Pace, PhD (ResMed, founder QRxPharma)
- John Holaday, PhD (CEO)

Scientific Advisory Board

- Solomon Snyder, MD (Chair)
- Lester Crawford, DVM, PhD
- Robert Lenox, MD
- Michael J Cousins, MD, AM
- Horace H Loh, PhD
- Gavril Pasternak, MD, PhD
- Richard Payne, MD



Financial Summary (4 September 2012)

Shares on issue:

Market cap:

Cash on hand:

30 June 2012

Cash burn:

Share registry:

Listing:

145 million (ordinary)

A\$101 million

A\$23 million (last reported) CY2013 +80% institutional / HNW ASX: QRX / OTCQX: QRXPY



MOXDUO IR Key Milestones

| DATE | MILESTONE |
|-----------------|---|
| ✓ July 2011 | NDA submission to US FDA |
| ✓ December 2011 | Signed strategic partnership with Actavis |
| × 25 June 2012 | NDA PDUFA Date; CRL Received |
| ✓ August 2012 | FDA Review Meeting |
| • 2H, 2012 | Submission of additional data package requested by FDA |
| • Mid-2013 | Anticipated decision from FDA on a refiled MOXDUO NDA |
| • 1H, 2013 | Submit additional regulatory filings: Europe, Australia & Canada |



Investment Highlights

- **Comprehensive Portfolio:** MOXDUO delivers equal or better pain relief with fewer side effects than current treatments
- Actavis commercialisation partnership in the US: MOXDUO IR to be flagship branded product
- Advantaged market: favourable US regulatory and potential prescription scheduling changes
- Blockbuster potential: global opioid market estimated at \$US14bn¹
- **Strong IP:** Expected patent exclusivity through 2029
- Expanded pipeline: Further progress MOXDUO CR and MOXDUO IV products



Contact Information

Australia

United States

QRxPharma Limited Level 1, 194 Miller Street North Sydney, NSW 2060 +61 2 9492 8021 +61 2 8920 0314 (fax) 1430 US Highway 206 Suite 230 Bedminster, NJ 07921 +1 908 506 2900 +1 908 506 2918 (fax)

