



ASX RELEASE

11 July 2014

QRxPHARMA PROVIDES FEEDBACK ON END-OF-REVIEW MEETING WITH THE FDA ON MOXDUO®

Sydney, Australia and Bedminster, New Jersey – QRxPharma Limited (ASX: QRX and OTCQX: QRXPY) today provided feedback on its 9 July End-of-Review (EOR) meeting with the United States Food and Drug Administration (FDA). The meeting was held to discuss the feasibility and requirements for approving Moxduo, an immediate release Dual Opioid®, for the treatment of moderate to severe acute pain.

In advance of the meeting, QRxPharma outlined several questions to discuss with FDA to ensure the Company receives clear direction for the Moxduo program. The questions addressed the overall approach for registration of Moxduo, potential study design and the number of clinical studies.

“We were encouraged by the extent of engagement on clinical issues by the Food and Drug Administration, and we found the meeting to be constructive and helpful,” said Dr. Edward Rudnic, CEO of QRxPharma. “We are evaluating the path forward and will utilise this guidance to determine the appropriate next steps.”

###

For more information please contact:

Edward M Rudnic, Ph.D.
Chief Executive Officer
Tel: +1 301 538 7080
Email: ed.rudnic@qrxpharma.com

Chris J Campbell
Chief Financial Officer and Company Secretary
Tel: +61 2 8404 4131
Email: chris.campbell@qrxpharma.com

About QRxPharma

QRxPharma Limited is an Australian based, specialty pharmaceutical company focused on the development and commercialisation of new pain management and abuse prevention products. The Company’s product portfolio includes both late and early stage clinical drug candidates with the potential for reduced risks and improved patient outcomes. QRxPharma has entered into strategic agreements with Actavis Inc., Paladin Labs Inc., Aspen Group and Teva Pharmaceuticals for the commercialisation of immediate release Moxduo in the US, Canada, Australia (including New Zealand and Oceania), South Africa and Israel. The Company’s clinical pipeline includes an intravenous (IV) and controlled release (CR) formulation of Moxduo. QRxPharma is also collaborating with Aesica Formulation Development Limited, for the worldwide promotion of QRxPharma’s proprietary Stealth Beadlets™ abuse deterrence technology. For more information, visit www.qrxpharma.com.



Forward Looking Statements

This release contains forward-looking statements. Forward-looking statements are statements that are not historical facts; they include statements about our beliefs and expectations. Any statement in this release that states our intentions, beliefs, expectations or predictions (and the assumptions underlying them) is a forward-looking statement. These statements are based on plans, estimates and projections as they are currently available to the management of QRxPharma. Forward-looking statements therefore speak only as of the date they are made, and we undertake no obligation to update publicly any of them in light of new information or future events. By their very nature, forward-looking statements involve risks and uncertainties. A number of important factors could therefore cause actual results to differ materially from those contained in any forward-looking statement. Such factors include risks relating to the stage of products under development; uncertainties relating to clinical trials; dependence on third parties; future capital needs; and risks relating to the commercialisation of the Company's proposed products.