Square Pharma gets USFDA's approval

USFDA (United States Food and Drug Administration) has released Establishment Inspection Report (EIR) on recent Pre Approval Inspection (PAI) to Square Pharmaceuticals Ltd. The report states that Square had a satisfactory cGMP inspection covering manufacturing of solid dosage form.

Manufacturing and Quality Assurance system of the oral solid dosage facilities of Square Pharma was audited in accordance with the Code of Federal Regulations (21 CFR: Food & Drugs) of United States of America.
The audit ended without issuing any 483 form. Square Pharma facilities have the track record of complying with the most stringent cGMP audits of highly regulated agencies like UK and Australia.

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