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Lupin receives US FDA approval for kidney ailment tabs

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The product will be manufactured at Lupin’s facility in Nagpur

Lupin has received approval for its Sevelamer Hydrochloride Tablets, 400 mg and 800 mg from the US FDA to market a generic equivalent of Renagel tablets, 400 mg and 800 mg, of Genzyme Corporation. The product will be manufactured at Lupin’s facility in Nagpur

Sevelamer hydrochloride tablets are indicated for the control of serum phosphorus in patients with chronic kidney disease (CKD) on dialysis. Sevelamer Hydrochloride Tablets (RLD: Renagel) had estimated annual sales of $80 million in the US.