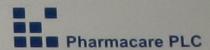


Sitagliptin / Metformin

50/850



Each film coated tablet contains 50 mg Sitagliptin (as phosphate monohydrate) and 850 mg Metformin HCI. Store tablets at temperature not exceeding 25°C.

Keep out of reach of children.

PHARMACARE PLC
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SITAGLU-MET® 50/500, 50/850

COMPOSITION:

SITAGLU-MET 50/500: Each tablet contains 50 mg Sitagliptin (as p. monohydrate) and 500 mg Metformin HCI.

SITAGLU-MET 50/850: Each tablet contains 50 mg Sitagliptin (as phosp monohydrate) and 850 mg Metformin HCI.

SITAGLU-MET 50/1000: Each tablet contains 50 mg Sitagliptin (as phosp monohydrate) and 1000 mg Metformin HCI.

THERAPEUTIC ACTIVITY: Sitagliptin belongs to DPP-4 (Dipeptidyl peptida enzyme inhibitor and Metformin belongs to Biguanide group.

APPROVED INDICATIONS: Sitaglu-Met is a medicine contains ingredients: Sitagliptin and Metformin, which lower blood sugar. together to control blood sugar levels in patients with type 2 diabete in whom this combination is appropriate. Sitaglu-Met alon recommended diet and exercise program is intended to lower bloo patients with type 2 diabetes.

WHEN SHOULD THE PREPARATION NOT BE USED?

- · If you have type 1 diabetes mellitus.
- · If you have kidneys which are not working properly.
- If you have diabetic ketoacidosis (increased ketone levels in the urine: diabetes complication which includes high blood sugar, raloss, nausea or vomiting).
- If you are going to get an injection of dye or contrast agents to procedure, Sitaglu-Met will need to be stopped for a short time, ta doctor about when you should stop Sitaglu-Met and when you st again.
- If you are allergic to any of the active ingredients. Symptoms of allergic reaction may include: rash, raised red patches on your skin swelling of the face, lips, tongue, and throat that may cause d breathing or swallowing.
- Sitaglu-met is not intended for administration to children under 1
 age.



