

Silymarin + Multivitamins

Liveraide® Pro

500 mg / Softgel Capsule
HEPATIC PROTECTOR / VITAMIN

PATIENT INFORMATION LEAFLET

NAME OF PRODUCT:

LIVERAIDE PRO SOFTGEL CAPSULE

DESCRIPTION OF THE PRODUCT:

Yellow to yellowish-brown colored oblong shaped softgel capsule containing yellowish-brown mass.

WHAT IS IN THE MEDICINE?

Each capsule contains

Silymarin.....	150 mg
Thiamine HCl (Vitamin B1).....	2.1 mg
Folic Acid (Vitamin B9).....	2 mg
Pyridoxine HCl (Vitamin B6).....	1.95 mg
Cyanocobalamin (Vitamin B12).....	3.6 mcg

WHAT IS THIS MEDICINE USED FOR?

Use as support for various chronic inflammatory disease of liver as an aid in the treatment. This includes alcoholic and non-alcoholic fatty liver, viral hepatitis, liver cirrhosis and liver damage caused by certain drugs, for the management, protection, and maintenance of liver health. For the prevention & treatment of Vitamin B1, B6, B12 and Folic Acid deficiencies.

Silymarin in for gastrointestinal and hepatobiliary disorders. Hepatoprotectant. Used in various liver disorders (fatty liver), as well as to prevent hepatotoxicity associated with poisoning. (Matindale 36th edition)

Thiamine (Vitamin B1) deficiency develops when the dietary intake is inadequate, severe deficiency leads to the development of a syndrome known as beri-beri. Chronic "dry" beri-beri is characterized by peripheral neuropathy, muscle weakness and paralysis. Acute "wet" beri-beri is characterized by cardiac failure and oedema. Wernicke-Korsakoff syndrome (demyelination of the CNS) may develop in severe cases of thiamine deficiency notably in association with chronic alcoholism. (Matindale 36th)

Pyridoxine (Vitamin B6), a water soluble vitamin, is involved mainly in amino acid metabolism, but is also involved in carbohydrate and fat metabolism. It is also required for the formation of hemoglobin. (Matindale 36th)

Folic Acid (Vitamin B9) together with an adequate amount of Cyanocobalamin (B12) for the treatment of a diagnosed megaloblastic anaemia due to its deficiency. (Matindale 36th)

HOW MUCH AND HOW OFTEN SHOULD YOU USE THIS MEDICINE?

Orally (By mouth), Once a day after meal or as prescribed by a physician.

WHEN SHOULD YOU NOT TAKE THIS MEDICINE?

If you are allergic to any component of the product.

CARE THAT SHOULD BE TAKEN WHEN TAKING THIS MEDICINE:

- Ask the doctor before use if you are on a sodium restricted diet.
- Do not take more than the recommended dose.
- Do not use after the expiry date on the label.

UNDESIRABLE EFFECTS:

A mild laxative action has been observed in occasional instances.

WHAT OTHER MEDICINE OR FOOD SHOULD BE AVOIDED WHILE TAKING THIS MEDICINE?

The amount of nutrients in the product is not expected to interact with food or other medicines.

WHAT SHOULD YOU DO IF YOU MISSED A DOSE?

Take the missed dose as soon as you remember. Skip the missed dose if it is almost time for your next scheduled dose. Do not take extra medicine to make up the missed dose.

SIGNS AND SYMPTOMS OF OVERDOSE:

Diarrhea, weakness and feeling tired, or trouble concentrating. Numbness, mouth and tongue pain. Seek medical attention immediately at the first sign of any suspected drug overdose.

WHAT TO DO WHEN YOU HAVE TAKEN MORE THAN THE RECOMMENDED DOSAGE?

If you take more than the recommended dosage, consult a doctor.

HOW SHOULD YOU KEEP THIS MEDICINE?

Store at temperature not exceeding 30°C.

Keep out of reach of children.

Avoid exposure to direct sunlight.

WHEN SHOULD YOU CONSULT YOUR DOCTOR?

If any undesirable effects occur.

If also taking other medications.

Pregnant and lactating mothers.

ADVERSE EFFECT:

Gastrointestinal disturbances and hypersensitivity reactions have been reported rarely.

INTERACTIONS:

If taking other medications, consult a physician before taking the drug for any advice of drug interactions.

AVAILABILITY:

In Alu/Alu Strips x 6 softgel capsules. Tradebox by 60's.

ADR REPORTING STATEMENT:

- For suspected adverse drug reaction, report to the FDA: www.fda.gov.ph
- Patient should seek medical attention immediately at the first sign of any adverse drug reaction.

REGISTRATION NUMBER:

FDA Reg. No. DR-XY49370

DATE OF FIRST AUTHORIZATION:

April 3, 2025

DATE OF REVISION:

N/A

Manufactured by:

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