Brand name: ZENOL XUM Liquid

RISK CATEGORIES
Second-class OTC drug

CHARACTERISTICS
- Topical analgesic anti-inflammatory agent

Felbinac helps relieve shoulder and low back pain.
Long body and wide face
This agent is thick and liquid gel so that it is less likely to drip.
It spreads smoothly without causing sticky feeling after drying.
Less odor

ACTIVE INGREDIENTS
Per 100 g

ACTIVE INGREDIENTS	Content
Felbinac	3.0 g
L-Menthol	2.0 g

Inactive ingredients: Hydroxypropylcellulose, Polyoxyethylene Hardened Caster Oil, Flavor, Triethanolamine, Alcohol

For further information:

Distributed by:	TAIHO PHARMACEUTICAL CO., LTD.
Manufactured and Distributed by:	Mikasa Seiyaku Co., Ltd.
Dosage form	Gel
Packaging unit	52 mL x 1 box
Manufacturer's suggested retail price	JPY 1,898 (JPY 1,725 excluding tax)
JAN Code	4987117372304
Expiration	3 years

INDICATIONS
Shoulder pain due to stiff shoulder, low back pain, muscle pain, joint pain, bruise, sprain, tenosynovitis (pain and swelling in hands, wrists, and ankles), elbow pain (tennis elbow, etc.)

DOSAGE AND ADMINISTRATION
Apply an appropriate amount to the affected area twice to four times daily.

<Precautions>
(1) The sponge surface may be damaged if used when dry.Before applying the agent, check that the sponge surface is fully wet with the solution.
(2) Strictly follow the dosing instructions.
(3) Be careful not to get the agent into the eyes. If the agentgets into the eyes, immediately wash with water or lukewarm water. If symptoms are severe, consult an ophthalmologist.
(4) For external use only. Do not take the agent orally.
(5) Do not cover the affected area with a poorly-ventilated material such as wrap film, etc. after applying the agent.

PRECAUTIONS FOR USE
☒ The following precaution should be observed.
(If not observed, it may aggravate the current symptoms or increase the risk of adverse reactions.)
1. The following persons should not use the agent.
(1) Persons with a history of allergic symptoms to the agent or any of the ingredients of the agent
(2) Persons with a history of asthma
(3) Pregnant women or possibly pregnant women
(4) Children under 15 years

2. Do not apply to the following areas.
(1) Around the eyes, mucous membranes, etc.
(2) Eczema, contact dermatitis, wounds
(3) Athlete's foot (tinea pedis), tinea cruris, etc. or purulent areas

Consultation

1. The following persons should consult a physician, pharmacist or registered salesperson before using the agent.
(1) Persons under treatment by a physician
(2) Persons with a history of allergic symptoms to any drug

2. The following symptoms may be adverse reactions to the agent. If any of these symptoms occur, discontinue the use of the agent immediately, and consult a physician, pharmacist or registered salesperson with this leaflet.

Affected body system	Symptoms
Skin	Rash, redness, itching, swelling, tingling sensation, contact dermatitis

The following serious symptom may rarely occur. If the symptom occurs, consult a physician immediately.

Symptom	Symptoms
Shock (anaphylaxis)	Skin itching, urticaria, hoarse voice, sneezing, larynx itching, difficulty breathing, palpitation, consciousness clouding, etc. occur immediately after

3. If symptoms do not improve after using the agent for about one week, discontinue the use of the agent, and consult a physician, pharmacist or registered salesperson with this leaflet.

CONTACT

For inquiries, please contact the shop where you purchased the product or the following:
Distributor Contact: Customer Relations Office, Taiho Pharmaceutical Co., Ltd.
1-27 Kandanishiki-cho, Chiyoda-ku, Tokyo, 101-8444, Japan
Phone: 0120-4527-66
Business hours: 9:00-17:00 (Excluding weekends and holidays)
Website: https://www.taiho.co.jp/

Manufacturer and distributor : MIKASA SEIYAKU CO., LTD