

POSILAC- sometribove suspension **Union Agener Inc**

Posilac™

Posilac (sometribove zinc suspension)

DESCRIPTION: Sterile, prolonged-release injectable formulation of a recombinant DNA-derived bovine somatotropin analog in single-dose syringes each containing 500 mg of sometribove zinc.

USE: To increase production of marketable milk in healthy lactating dairy cows.

CAUTION: Federal law prohibits extra-label use of this drug to enhance food and/or fiber production in animals.

DOSAGE: Inject one syringe of Posilac every 14 days. Start during the 9th or 10th week (57-70 days) after calving and continue until the end of lactation.

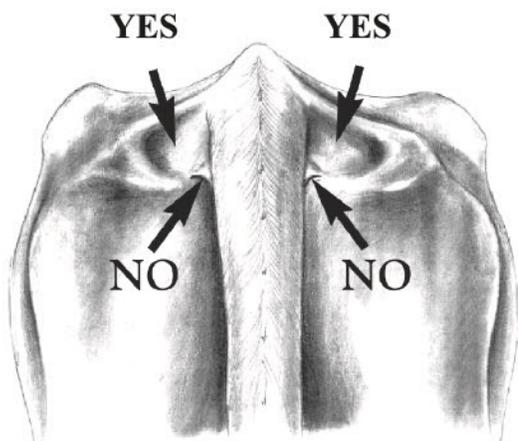
ADMINISTRATION: Allow syringes to warm to room temperature (15° to 30° C, 59° to 86° F) before use.

INJECTION TECHNIQUE: Inject Posilac subcutaneously (under the skin). Recommended injection sites are *neck area, behind the shoulder or in the depression on either side of the tailhead* (see diagrams below). Alternate between the cow's left and right side on consecutive injections. Remove surface dirt from the injection site area before injecting. Inject entire contents of the syringe subcutaneously. Do not reuse syringes.

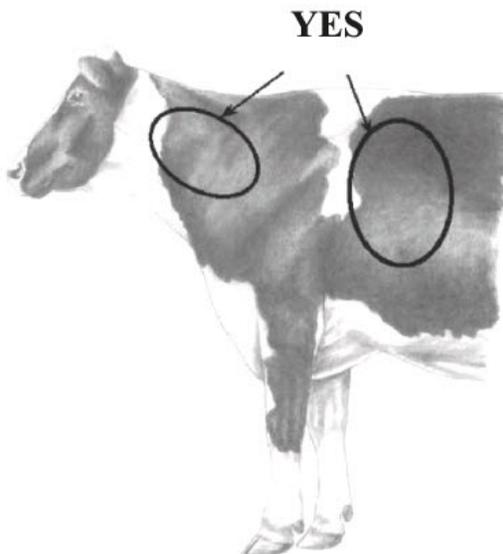
Inject directly into the deepest depressions on either side of the tailhead (marked "Yes"). Avoid the bone, muscles, tendons and ligaments of the tail and the rectal and anal muscles. **Do NOT inject into the caudal fold (marked "No") because this may invalidate USDA tuberculosis testing.** Locate the caudal fold by raising the tail.

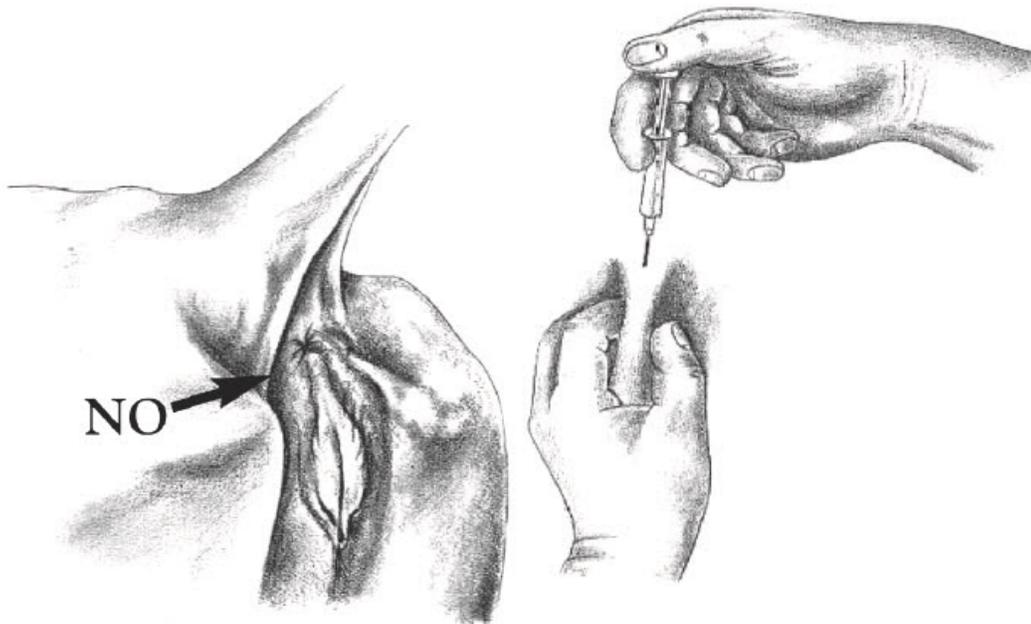
INJECTION SITES:

Tailhead Depression



Neck & Behind The Shoulder





Gather skin and inject between skin and muscle layers.

WITHDRAWAL PERIODS:

No milk discard time and no withdrawal period is required when used according to labeling.

USER SAFETY WARNINGS: Not for use in humans, Keep out of reach of children, Avoid prolonged or repeated contact with Posilac with eyes and skin, Posilac is a protein. Frequent skin contact with proteins may produce an allergic reaction in some people. Always wash hands and skin exposed to Posilac with soap and water after handling. Clothing soiled with the product should be laundered before reuse.

ANIMAL SAFETY WARNINGS:

- Use in lactating dairy cows only.
- Safety to replacement bulls born in dairy cows injected with Posilac has not been established.
- Avoid injecting within 2 weeks of slaughter to minimize injection site blemishes on carcass.
- **Nutritional Management:** Cows injected with Posilac increase voluntary feed intake over several weeks following the start of supplementation. This increase occurs sooner for first lactation cows than for second lactation or older cows. The increased feed intake continues during supplementation and may continue through the dry period and the following early lactation. However, cows treated with Posilac tend to maintain lower body condition than untreated cows. This effect is more pronounced for second lactation or older cows.
- **Feed diets formulated to meet or exceed the nutritional requirements recommended by the National Research Council. Consider milk yield, stage of lactation, and body condition when making dietary changes. Manage the feeding program to optimize milk yield and to have cows in**

appropriate body condition, particularly during late lactation and the dry period. Increasing the energy density of diets fed to cows treated with Posilac is normally not required. Avoid sudden dietary changes.

- **Reproduction:** Cows injected with Posilac may have reduced pregnancy rates and increased days open. **Have a comprehensive and ongoing herd reproductive health program in place on your dairy before using Posilac.**
- **Mastitis:** Cows injected with Posilac are at an increased risk of mastitis (visibly abnormal milk) and may have higher somatic cell counts. **Have comprehensive mastitis management practices in place on your dairy before using Posilac.**
- **General Health:** Cows injected with Posilac may require more therapeutic drug treatment for mastitis and other health problems. Cows injected with Posilac may experience periods of increased body temperature unrelated to illness. To minimize the effect, take appropriate measures during periods of high environmental temperature to reduce heat stress. Use care to differentiate whether increased body temperature is caused by illness or use of Posilac. Cows injected with Posilac may have more enlarged hocks and disorders of the foot region. Posilac treatment may reduce hemoglobin and hematocrit values.
- **Injection Site Reactions:** A mild temporary swelling of 3-5 cm (1-2 inches) in diameter may occur at the injection site beginning about 3 days after injection and may persist up to 6 weeks following injection. Larger swellings may occur in cows injected in the neck area compared to the behind the shoulder or in the depression on either side of the tailhead. Some cows may experience swellings up to 10 cm (4 inches) in diameter that remain permanent but are not associated with animal health problems. However, if permanent blemishes are objectionable to you, stop supplementation of these cows. Also stop using Posilac in cows with injection site swellings that repeatedly open and drain.
- **Udder Edema:** Posilac is approved for use starting during the 9th or 10th week of lactation. Risk of udder edema may increase if injections start later in lactation.

ADDITIONAL INFORMATION:

- Milk production response during each 14-day injection period is cyclic and will be greatest during the middle of each period.

STORAGE: Store under refrigeration (2° to 8° C; 36° to 46° F). DO NOT FREEZE. Allow syringes to warm to room temperature (15° to 30° C; 59° to 86° F) before use. Avoid prolonged exposure to excessively high temperature and sunlight. Expiration dates are stated on syringes and box labeling.

ENVIRONMENTAL SAFETY: Dispose of used syringes in a leak-resistant, puncture-resistant container in accordance with applicable Federal, state, and local regulations.

HOW SUPPLIED: Single-dose syringes in 25 or 100 count boxes.

QUESTIONS/COMMENTS?

Contact Union Agener, Inc. at +1 844-952-0330. To report side effects, contact Union

Agener, Inc. at +1 844-952-0330. For additional information about reporting side effects for animal drugs, contact FDA at + 1 1-888-FDA-VETS or <http://www.fda.gov/reportanimalae>

Approved by FDA under NADA # 140-872
Manufactured by
Union Agener, Inc.
1788 Lovers Lane, Augusta, Georgia 30901, U.S.A.
4024774

PRINCIPAL DISPLAY PANEL - 25 Syringe Carton

Posilac (sometribove zinc suspension)

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25 Single-dose syringes

Use in healthy lactating dairy cows to increase production of marketable milk. Have comprehensive and ongoing herd nutritional management, mastitis control, and reproductive health programs in place on your dairy before using Posilac. See panels and accompanying package insert for complete directions for use, warnings, precautions, side effects, and additional information.

CAUTION: Federal law prohibits extra-label use of this drug to enhance food and/or fiber production in animals.

CONTENTS: 25 single-dose syringes each containing 500 mg sometribove zinc and directions for use.

DESCRIPTION: Sterile, prolonged-release injectable formulation of a recombinant DNA-derived bovine somatotropin analog in single-dose syringes each containing 500 mg of sometribove zinc.

UNION AGENER

Posilac™
sometribove zinc suspension

25 Count

ENVIRONMENTAL SAFETY: Dispose of used syringes in a leak-resistant, puncture-resistant container in accordance with applicable Federal, state, and local regulations.

ADDITIONAL INFORMATION:

- See package insert.
- Avoid contact with eyes and skin.
- Not for use in humans. Keep out of reach of children.
- Avoid contact with eyes and skin.

ANIMAL SAFETY WARNING:

- See package insert.
- Use in lactating dairy cows only.
- Safety to replacement bulls born in dairy cows injected with Posilac has not been established.
- Avoid injecting within 2 weeks of slaughter to minimize injection site blemishes on carcasses.

WITHDRAWAL PERIODS: No milk discard time and no withdrawal period is required when used according to labeling.

USER SAFETY WARNINGS:

ENVIRONMENTAL SAFETY: Dispose of used syringes in a leak-resistant, puncture-resistant container in accordance with applicable Federal, state, and local regulations.

ADDITIONAL INFORMATION:

- See package insert.
- Avoid contact with eyes and skin.
- Not for use in humans. Keep out of reach of children.
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Approved by FDA under NADA # 140-872

4033329

LOT NO. EXP. DATE

1 8 4 4 - 9 5 2 - 0 3 3 0

1 7 8 8 L O V E R S L A N E A U G U S T A G E O R G A

Questions or comments, please call
(Preguntas y comentarios, por favor llame al)

+1 844-952-0330

UNION AGENER

Posilac™



25 Jeringas de dosis individuales

USO: Use en vacas lecheras sanas en lactancia para aumentar la producción de leche comercializable. Tenga un control alimenticio del ganado más amplio y continuo, control de mastitis y salud reproductiva antes de usar Posilac. Vea los paneles laterales y el prospecto dentro del paquete para obtener informaciones más completas sobre indicaciones de uso, advertencias, precauciones, efectos secundarios y informaciones adicionales.

PRECAUCIÓN: La ley federal prohíbe el uso de este producto para mejorar la producción de alimento y/o fibra en animales.

CONTENIDO: 25 jeringas de dosis individuales, cada una con 500 mg de sometribova suspensión de zinc y prospecto con indicaciones de uso.

DESCRIPCIÓN: Formulación inyectable estéril de liberación prolongada, de un análogo de somatotropina bovina derivada por medio de ADN recombinante, en jeringas con una dosis individual cada una con 500 mg de sometribova zinc.

TIEMPO DE RETIRO: Cuando se usa de acuerdo con el etiquetado, no se requiere tiempo de descarte de la leche y no es necesario un tiempo de retiro.

ADVERTENCIAS DE SEGURIDAD DEL USUARIO:

- Para uso exclusivo en animales.
- Mantenga este y todos los medicamentos fuera del alcance de los niños.
- Evite el contacto con los ojos y la piel.

ADVERTENCIAS DE SEGURIDAD ANIMAL:

- Lea el prospecto.
- Use únicamente en vacas lecheras en lactancia.
- No ha sido establecida la seguridad para toros de reemplazo nacidos de vacas lecheras tratadas con Posilac.
- Evite inyectar dentro de las 2 semanas antes de enviar al matadero para minimizar marcas en la canal en el sitio de inyección.

INFORMACIONES ADICIONALES:

- Lea el prospecto.

SEGURIDAD AMBIENTAL: Deseche las jeringas usadas en contenedores sellados resistentes a la perforación, de acuerdo a las regulaciones aplicables a nivel federal, estatal y local.

UNION AGENER

Posilac™
sometribova suspensión de zinc

Caja con 25

PRODUCTOS DE LA HUILLA DE CARBONO E60

Questions or comments, please call
(Preguntas y comentarios, por favor llame al)

+1 844-952-0330

UNION AGENER

Posilac™



Posilac (sometribova suspensión de zinc)

Mantener en refrigeración (2 °C a 8 °C; 36 °F a 46 °F). NO CONGELAR. Permita que las jeringas alcancen la temperatura ambiente (15 °C a 30 °C; 59 ° a 86 °F) antes de usar. Evite la exposición prolongada a temperaturas excesivamente altas y a la luz solar.

Elaborado por Union Agener, Inc. 1788 Lovers Lane, Augusta, Georgia 30901, E.U.A.

PREGUNTAS/COMENTARIOS? Contacte a Union Agener, Inc. al +1 844-952-0330. Para reportar efectos secundarios, contacte Union Agener, Inc. al +1 844-952-0330. Para obtener informaciones adicionales sobre cómo reportar efectos secundarios para animales, contacte FDA al +1 888-FDA-VETS o <http://www.fda.gov/reportanimal>

Aprobado por la FDA bajo número NADA # 140-872

sometribove suspension

Product Information

Product Type	OTC ANIMAL DRUG	Item Code (Source)	NDC:86106-0225
Route of Administration	SUBCUTANEOUS		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
sometribove (UNII: PBK5EQG5CQ) (sometribove - UNII:PBK5EQG5CQ)	sometribove	500 mg

Inactive Ingredients

Ingredient Name	Strength
Sesame Oil (UNII: QX10HYY4QV)	
Aluminum Monostearate (UNII: P9BC99461E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:86106-0225-3	25 in 1 CARTON		
1	NDC:86106-0225-1	1 in 1 SYRINGE		
2	NDC:86106-0225-2	100 in 1 CARTON		
2	NDC:86106-0225-1	1 in 1 SYRINGE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NADA	NADA140872	03/01/2020	

Labeler - Union Agener Inc (116587901)

Establishment

Name	Address	ID/FEI	Business Operations
Union Agener Inc.		116587901	ANALYSIS, API MANUFACTURE, LABEL, MANUFACTURE, PACK

Revised: 3/2025

Union Agener Inc