



# Louisiana

## Soanz<sup>®</sup> (torsemide tablets)

Policy # 00794

Original Effective Date: 07/11/2022

Current Effective Date: 07/11/2022

*Applies to all products administered or underwritten by Blue Cross and Blue Shield of Louisiana and its subsidiary, HMO Louisiana, Inc. (collectively referred to as the "Company"), unless otherwise provided in the applicable contract. Medical technology is constantly evolving, and we reserve the right to review and update Medical Policy periodically.*

## When Services May Be Eligible for Coverage

*Coverage for eligible medical treatments or procedures, drugs, devices or biological products may be provided only if:*

- *Benefits are available in the member's contract/certificate, and*
- *Medical necessity criteria and guidelines are met.*

Based on review of available data, the Company may consider Soanz<sup>®</sup> (torsemide tablets) to be **eligible for coverage\*\*** when the patient selection criteria are met.

### Patient Selection Criteria

Coverage eligibility for Soanz (torsemide tablets) will be considered when the following criteria are met:

- Requested drug is being used for the treatment of edema associated with heart failure or renal disease; AND
- Patient is 18 years of age or older; AND
- There is clinical evidence or patient history that suggests the use of GENERIC torsemide 5 mg, 10 mg, 20 mg, or 100 mg tablets will be ineffective or cause an adverse reaction to the patient.

*(Note: This specific patient criterion is an additional Company requirement for coverage eligibility and will be denied as not medically necessary\*\* if not met.)*

## When Services Are Considered Not Medically Necessary

Based on review of available data, the Company considers the use of Soanz (torsemide tablets) when there is an absence of clinical evidence or patient history that suggests the use of GENERIC torsemide 5 mg, 10 mg, 20 mg, or 100 mg tablets will be ineffective or cause an adverse reaction to the patient to be **not medically necessary.\*\***

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## **When Services Are Considered Investigational**

*Coverage is not available for investigational medical treatments or procedures, drugs, devices or biological products.*

Based on review of available data, the Company considers the use of Soanz (torsemide tablets) for non-FDA approved indications OR for patients under 18 years of age to be **investigational**.\*

## **Background/Overview**

Soanz is a loop diuretic indicated in adults for the treatment of edema associated with heart failure or renal disease. It is available in 20 mg, 40 mg, and 60 mg tablets. Soanz was not studied in clinical trials. It was only studied in kinetic studies versus the generic torsemide tablets. Generic torsemide tablets are available in 5 mg, 10 mg, 20 mg, and 100 mg tablet strengths. There is no advantage of using Soanz over generic torsemide tablets, which offer a safe, efficacious, and economically advantageous option for therapy.

## **FDA or Other Governmental Regulatory Approval**

### **U.S. Food and Drug Administration (FDA)**

Soanz is indicated in adults for the treatment of edema associated with heart failure or renal disease.

## **Rationale/Source**

This medical policy was developed through consideration of peer-reviewed medical literature generally recognized by the relevant medical community, U.S. Food and Drug Administration approval status, nationally accepted standards of medical practice and accepted standards of medical practice in this community, technology evaluation centers, reference to federal regulations, other plan medical policies, and accredited national guidelines.

The intent of this policy is to ensure that Soanz is being used per its FDA approved indication. Additionally, the most economical option for therapy should be used as there are no safety or efficacy advantages by using Soanz over generic torsemide 5 mg, 10 mg, 20 mg, or 100 mg tablets.

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## **References**

1. Soanz [package insert]. Sarfez Pharmaceuticals, Inc. Vienna, Virginia. Updated December 2021.

## **Policy History**

Original Effective Date: 07/11/2022

Current Effective Date: 07/11/2022

06/02/2022 Medical Policy Committee review

06/08/2022 Medical Policy Implementation Committee approval. New policy.

Next Scheduled Review Date: 06/2023

\*Investigational – A medical treatment, procedure, drug, device, or biological product is Investigational if the effectiveness has not been clearly tested and it has not been incorporated into standard medical practice. Any determination we make that a medical treatment, procedure, drug, device, or biological product is Investigational will be based on a consideration of the following:

- A. Whether the medical treatment, procedure, drug, device, or biological product can be lawfully marketed without approval of the U.S. Food and Drug Administration (FDA) and whether such approval has been granted at the time the medical treatment, procedure, drug, device, or biological product is sought to be furnished; or
- B. Whether the medical treatment, procedure, drug, device, or biological product requires further studies or clinical trials to determine its maximum tolerated dose, toxicity, safety, effectiveness, or effectiveness as compared with the standard means of treatment or diagnosis, must improve health outcomes, according to the consensus of opinion among experts as shown by reliable evidence, including:
  1. Consultation with technology evaluation center(s);
  2. Credible scientific evidence published in peer-reviewed medical literature generally recognized by the relevant medical community; or
  3. Reference to federal regulations.

\*\*Medically Necessary (or “Medical Necessity”) - Health care services, treatment, procedures, equipment, drugs, devices, items or supplies that a Provider, exercising prudent clinical judgment, would provide to a patient for the purpose of preventing, evaluating, diagnosing or treating an illness, injury, disease or its symptoms, and that are:

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- A. In accordance with nationally accepted standards of medical practice;
- B. Clinically appropriate, in terms of type, frequency, extent, level of care, site and duration, and considered effective for the patient's illness, injury or disease; and
- C. Not primarily for the personal comfort or convenience of the patient, physician or other health care provider, and not more costly than an alternative service or sequence of services at least as likely to produce equivalent therapeutic or diagnostic results as to the diagnosis or treatment of that patient's illness, injury or disease.

For these purposes, “nationally accepted standards of medical practice” means standards that are based on credible scientific evidence published in peer-reviewed medical literature generally recognized by the relevant medical community, Physician Specialty Society recommendations and the views of Physicians practicing in relevant clinical areas and any other relevant factors.

‡ Indicated trademarks are the registered trademarks of their respective owners.

**NOTICE:** If the Patient’s health insurance contract contains language that differs from the BCBSLA Medical Policy definition noted above, the definition in the health insurance contract will be relied upon for specific coverage determinations.

**NOTICE:** Medical Policies are scientific based opinions, provided solely for coverage and informational purposes. Medical Policies should not be construed to suggest that the Company recommends, advocates, requires, encourages, or discourages any particular treatment, procedure, or service, or any particular course of treatment, procedure, or service.

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