

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

LEVOTHYROXINE-TIROSINT

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
LEVOTHYROXINE	TIROSINT,	02849		GPI-10	FORM = CAPSULE
SODIUM	LEVOTHYROXINE			(2810001010)	

GUIDELINES FOR USE

1. Does the patient have a diagnosis of congenital or acquired hypothyroidism?

If yes, continue to #3. If no, continue to #2.

- 2. Does the patient have a diagnosis of thyrotropin-dependent well-differentiated thyroid cancer **AND** meet the following criterion?
 - The requested medication is being used as an adjunct to surgery and radioiodine therapy

If yes, continue to #3. If no. do not approve.

DENIAL TEXT: See the denial text at the end of the guideline.

- 3. Does the patient meet **ALL** of the following criteria?
 - The patient is 6 years of age or older
 - The patient had a trial and failure of generic levothyroxine tablets
 - There is documentation of rationale for not using generic levothyroxine tablets

If yes, approve for 12 months by GPID or GPI-14 with a quantity limit of #2 per day. If no, do not approve.

DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **LEVOTHYROXINE-TIROSINT** requires the following rule(s) be met for approval:

- A. You have ONE of the following diagnoses:
 - 1. Congenital (present from birth) or acquired hypothyroidism (low thyroid function)
 - 2. Thyrotropin (a type of thyroid hormone)-dependent well-differentiated thyroid cancer
- B. You are 6 years of age or older
- C. You had a trial and failure (drug did not work) of generic levothyroxine tablets
- D. There is documentation of rationale (reason) for not using generic levothyroxine tablets
- E. If you have thyrotropin-dependent well-differentiated thyroid cancer, approval also requires:
 - 1. The requested medication will be used as an adjunct (add-on) to surgery and radioiodine therapy (a type of radiation therapy)

(Denial text continued on next page)

CONTINUED ON NEXT PAGE

Copyright © 2023 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Revised: 12/2/2022 Page 1 of 2



STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

LEVOTHYROXINE-TIROSINT

GUIDELINES FOR USE (CONTINUED)

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

RATIONALE

For further information, please refer to the Prescribing Information and/or Drug Monograph for Tirosint.

REFERENCES

• Tirosint [Prescribing Information]. Pambio-Noranco, Switzerland: IBSA Institut Biochimique SA; June 2022.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Created: 07/21

Commercial Effective: 01/01/23 Client Approval: 12/22 P&T Approval:10/21

Copyright © 2023 MedImpact Healthcare Systems, Inc. All rights reserved. This document is proprietary to MedImpact. MedImpact maintains the sole and exclusive ownership, right, title, and interest in and to this document.

Revised: 12/2/2022 Page 2 of 2