



STANDARD COMMERCIAL DRUG FORMULARY
PRIOR AUTHORIZATION GUIDELINES

FENFLURAMINE

| Generic | Brand | HICL | GCN | Medi-Span | Exception/Other |
|--------------|----------|-------|-----|------------------------|-----------------|
| FENFLURAMINE | FINTEPLA | 02116 | | GPI-10 (7260002810) | |

GUIDELINES FOR USE

INITIAL CRITERIA (NOTE: FOR RENEWAL CRITERIA SEE BELOW)

1. Does the patient have a diagnosis of seizures associated with Dravet syndrome and meet **ALL** of the following criteria?
 - The patient is 2 years of age or older
 - Therapy is prescribed by or given in consultation with a neurologist
 - The patient had a trial of or contraindication to clobazam **AND** valproic acid derivatives

If yes, **approve for 12 months by HICL or GPI-10 with a quantity limit of #11.8mL 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 **FENFLURAMINE (Fintepla)** requires the following rule(s) be met for approval:

- A. You have seizures associated with Dravet syndrome (severe type of seizure disorder that begins during the first year of life)
- B. You are 2 years of age or older
- C. Therapy is prescribed by or given in consultation with a neurologist (doctor who specializes in the brain, spine, and nerves)
- D. You had a previous trial of clobazam AND valproic acid derivatives, unless there is a medical reason why you cannot (contraindication)

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.

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RENEWAL CRITERIA

1. Does the patient have a diagnosis of seizures associated with Dravet syndrome **AND** meet the following criterion?
 - The patient has shown continued clinical benefit (e.g. reduction of seizures, reduced length of seizures, seizure control maintained)

If yes, **approve for 12 months by HICL or GPI-10 with a quantity limit of #11.8mL 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 **FENFLURAMINE (Fintepla)** requires the following rule(s) be met for approval:

- A. You have seizures associated with Dravet syndrome (severe type of seizure disorder that begins during the first year of life)
- B. You have shown continued clinical benefit (such as reduction of seizures, reduced length of seizures, seizure control maintained) while on therapy

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 Fintepla.

REFERENCES

Fintepla [Prescribing Information]. Emeryville, CA: Zogenix, Inc., June 2020.

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|---------|------------|-----|
| Library | Commercial | NSA |
| Yes | Yes | No |

Part D Effective: N/A

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P&T Approval: 04/20