



STANDARD COMMERCIAL DRUG FORMULARY  
PRIOR AUTHORIZATION GUIDELINES

ABROCITINIB

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
ABROCITINIB	CIBINQO	47767		GPI-10 (9027200500)	

**GUIDELINES FOR USE**

**INITIAL CRITERIA (NOTE: FOR RENEWAL CRITERIA SEE BELOW)**

1. Does the patient have a diagnosis of refractory, moderate to severe atopic dermatitis and meet **ALL** of the following criteria?

- The patient is 12 years of age or older
- Therapy is prescribed by or in consultation with a dermatologist, allergist, or immunologist
- The patient has atopic dermatitis involving at least 10% of body surface area (BSA) OR atopic dermatitis affecting the face, head, neck, hands, feet, groin, or intertriginous areas
- The patient has TWO of the following: intractable pruritus, cracking and oozing/bleeding of affected skin, impaired activities of daily living
- The patient had a trial of or contraindication to ONE preferred agent: Dupixent (dupilumab), Rinvoq (upadacitinib)
- Cibinqo will NOT be used concurrently with other systemic biologics (e.g., Adbry [tralokinumab-ldrm], Dupixent [dupilumab]) for atopic dermatitis or other JAK inhibitors (e.g., Xeljanz [tofacitinib], topical Opzelura [ruxolitinib]) for any indication

If yes, continue to #2.

If no, do not approve.

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

2. Does the patient have a trial of or contraindication to **TWO** of the following?

- High or super-high potency topical corticosteroid (e.g., triamcinolone acetonide, fluocinonide, clobetasol propionate)
- Topical calcineurin inhibitor (e.g., tacrolimus, pimecrolimus)
- Topical PDE-4 inhibitor (e.g., crisaborole)
- Topical JAK inhibitor (e.g., ruxolitinib)
- Phototherapy

If yes, **approve for 6 months by HICL or GPI-10 with a quantity limit of #1 per day.**

If no, do not approve.

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

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ABROCITINIB

INITIAL CRITERIA (CONTINUED)

**INITIAL 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 **ABROCITINIB (Cibinqo)** requires the following rule(s) be met for approval:

- A. You have refractory, moderate to severe atopic dermatitis (a type of skin condition)
- B. You are 12 years of age or older
- C. Therapy is prescribed by or in consultation with a dermatologist (a type of skin doctor), allergist (a type of allergy doctor), or immunologist (a type of immune system doctor)
- D. You have atopic dermatitis involving at least 10% of body surface area (BSA) OR atopic dermatitis affecting the face, head, neck, hands, feet, groin, or intertriginous areas (between skin folds, the hands, feet, etc.)
- E. You have TWO of the following: intractable pruritus (severe itching), cracking and oozing/bleeding of affected skin, impaired activities of daily living
- F. You had a trial of or contraindication (harmful for) to TWO of the following:
  - 1. High or super-high potency topical corticosteroid (such as triamcinolone acetonide, fluocinonide, clobetasol propionate)
  - 2. Topical calcineurin inhibitor (e.g., tacrolimus, pimecrolimus)
  - 3. Topical PDE-4 inhibitor (Phosphodiesterase-4 Inhibitors such as crisaborole)
  - 4. Topical Janus kinase (JAK) inhibitor (Janus kinase inhibitor such as ruxolitinib)
  - 5. Phototherapy (light therapy)
- G. You had a trial of or contraindication (harmful for) to ONE preferred medication: Dupixent (dupilumab), Rinvoq (upadacitinib)
- H. You will NOT use Cibinqo concurrently (at the same time) with other systemic biologics (such as Adbry [tralokinumab-ldrm], Dupixent [dupilumab]) for atopic dermatitis or other Janus kinase (JAK) inhibitors (such as Xeljanz [tofacitinib], topical Opzelura [ruxolitinib]) for any indication

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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ABROCITINIB

RENEWAL CRITERIA

1. Does the patient have a diagnosis of refractory, moderate to severe atopic dermatitis and meet **ALL** of the following criteria?
  - The patient has shown improvement while on therapy
  - The patient had a trial of or contraindication to ONE preferred agent: Dupixent (dupilumab), Rinvoq (upadacitinib)
  - Cibinqo will NOT be used concurrently with other systemic biologics (e.g., Adbry [tralokinumab-ldrm], Dupixent [dupilumab]) for atopic dermatitis or other JAK inhibitors (e.g., Xeljanz [tofacitinib], topical Opzelura [ruxolitinib]) for any indication

If yes, **approve for 12 months by HICL with a quantity limit of #1 per day.**

If no, do not approve.

**RENEWAL 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 **ABROCITINIB (Cibinqo)** requires the following rule(s) be met for renewal:

- A. You have refractory, moderate to severe atopic dermatitis (a type of skin condition)
- B. You have shown improvement while on therapy
- C. You had a trial of or contraindication (harmful for) to ONE preferred medication: Dupixent (dupilumab), Rinvoq (upadacitinib)
- D. You will NOT use Cibinqo concurrently (at the same time) with other systemic biologics (such as Adbry [tralokinumab-ldrm], Dupixent [dupilumab]) for atopic dermatitis or other Janus kinase (JAK) inhibitors (such as Xeljanz [tofacitinib], topical Opzelura [ruxolitinib]) for any indication

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 Cibinqo.

**REFERENCES**

- Cibinqo [Prescribing Information]. New York, NY: Pfizer Labs; February 2023.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A

Commercial Effective: 10/01/23

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P&T Approval: 07/23

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