GILEAD | Medical Information

Hepcludex® (bulevirtide-gmod) Pregnancy and Breastfeeding

This document is in response to your request for information regarding the use of Hepcludex® (bulevirtide-gmod [BLV]) for the treatment of chronic HDV infection in pregnant or breastfeeding women.

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The full indication, important safety information, and boxed warnings are available at: www.gilead.com/-/media/files/pdfs/medicines/hdv/hepcludex/hepcludex_pi.

Product Labeling[1]

Indications and Usage

BLV is indicated for the treatment of chronic HDV infection in adults without cirrhosis or with compensated cirrhosis.

This indication is approved under accelerated approval based on a decrease in HDV RNA and ALT normalization. An improvement in disease-related clinical outcomes has not been established. Continued approval for this indication may be contingent upon verification and description of clinical benefit in a confirmatory trial(s).

Dosage and Administration

Recommended dosage in adults

The recommended dosage in adults is BLV 8.5 mg once daily administered by subcutaneous injection.

Use in Specific Populations

Pregnancy

Risk summary

There are insufficient human data on the use of BLV during pregnancy to inform a drug-associated risk of birth defects and miscarriage. The estimated background risk of major birth defects and miscarriage for the indicated population is unknown. In the US general population, the background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2% to 4% and 15% to 20%, respectively.

In nonclinical reproductive toxicity studies, BLV demonstrated no adverse effect on embryofetal development when administered to pregnant rats and rabbits at systemic

exposures (area under the concentration-time curve) 4- and 37-fold relative to exposure in humans at the recommended human dose (RHD).

Data: animal data

BLV was administered via subcutaneous injection to pregnant rats and rabbits (2.5 mg/kg/day) on Gestation Days 6 through 17 and 6 through 20, respectively, and also to rats from Gestation Day 6 to Lactation/Postpartum Day 20. There were no adverse effects on embryofetal development in rats and rabbits. During organogenesis, exposure in rats and rabbits was 4 and 37 times higher, respectively, than the exposure in humans at the RHD. In a pre/postnatal development study in rats, BLV (2.5 mg/kg/day) was administered via subcutaneous injection from Gestation Day 6 to Lactation Day 21. No effects were observed in the offspring at maternal exposures 3 times the exposure at the RHD.

Lactation

Risk summary

It is not known whether BLV is present in human breast milk, affects human milk production, or has effects on the breastfed infant. In nonclinical pre- and postnatal developmental rat studies, BLV was not measured in the plasma of pups or in the milk of nursing animals. However, due to its high protein binding, liver tropism, and high specificity for sodium taurocholate co-transporting polypeptide, BLV is not likely to be secreted in milk. The developmental and health benefits of breastfeeding should be considered along with the mother's clinical need for BLV and any potential adverse effects on the breastfed child from BLV or from the underlying maternal condition.

Clinical Studies

Clinical trials in adults with chronic HDV infection without cirrhosis or with compensated cirrhosis

Trial MYR301

The efficacy of BLV once daily in the treatment of adults with chronic HDV infection without cirrhosis or with compensated cirrhosis is based on data through Week 144 from a multicenter, randomized, open-label, parallel-arm phase 3 trial, Trial MYR301 (NCT03852719), in which 100 participants received BLV 8.5 mg once daily. The MYR301 protocol specified the BLV dose as 10 mg; however, a dose recovery study later showed that the delivered dose was 8.5 mg.

Clinical Data on BLV Use During Pregnancy and Breastfeeding

In a phase 3 study (MYR301) and phase 2/2b clinical studies (MYR202, MYR203, and MYR204), pregnant or breastfeeding female patients were excluded.[2-5]

Literature Search

Additionally, a literature search was conducted in Ovid MEDLINE and Embase databases for studies published up to August 11, 2025, using the search terms Hepcludex, bulevirtide,

pregnancy, breastfeeding, lactation, HDV, and related search terms. No relevant citations were found.

References

1. Enclosed. Gilead Sciences Inc. HEPCLUDEX® (bulevirtide) injection, for subcutaneous use. US Prescribing Information. Foster City, CA.
2. Wedemeyer H, Schoneweis K, Bogomolov P, et al. Safety and efficacy of bulevirtide in combination with tenofovir disoproxil fumarate in patients with hepatitis B virus and hepatitis D virus coinfection (MYR202): a multicentre, randomised, parallel-group, open-label, phase 2 trial. *Lancet Infect Dis.* 2022.
3. Asselah T, Chulanov V, Lampertico P, et al. Bulevirtide Combined with Pegylated Interferon for Chronic Hepatitis D [Supplementary Appendix]. *N Engl J Med.* 2024;391(2):133-143.
4. Wedemeyer H, Aleman S, Brunetto MR, et al. A Phase 3, Randomized Trial of Bulevirtide in Chronic Hepatitis D.[Protocol]. *N Engl J Med.* 2023;389(1):22-32.
5. Lampertico P, Bogomolov PO, Chulanov V, et al. Phase 2 Randomised Study of Bulevirtide as Monotherapy or Combined With Peg-IFNalpha-2a as Treatment for Chronic Hepatitis Delta [Supplement]. *Liver Int.* 2025;45(2):e70008.

Product Label

For the full indication, important safety information, and boxed warning(s), please refer to the Hepcludex US Prescribing Information available at: www.gilead.com/-/media/files/pdfs/medicines/hdv/hepcludex/hepcludex_pi.

Follow-Up

For any additional questions, please contact Gilead Medical Information at:

☎1-866-MEDI-GSI (1-866-633-4474) or www.askgileadmedical.com

Adverse Event Reporting

Please report all adverse events to:

Gilead Global Patient Safety ☎ 1-800-445-3235, option 3 or www.gilead.com/utility/contact/report-an-adverse-event

FDA MedWatch Program by ☎ 1-800-FDA-1088 or ✉MedWatch, FDA, 5600 Fishers Ln, Rockville, MD 20852 or www.accessdata.fda.gov/scripts/medwatch

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