# HI, I'M DIANA CAITO

## HEMOPHILIA COMMUNITY CLINICAL EDUCATION MANAGER

HEMLIBRA®
emicizumab-kxwh
injection for subcutaneous use | 150 mg/mL

**"I'm a nurse who is dedicated and passionate about providing support and education to the hemophila community."**

## A LITTLE BIT ABOUT ME:

- I am born and raised in IN
- I am the youngest of 7 children
- I love to spend time with my family and friends traveling
- I have been working with the bleeding disorders community since 2017

As a Hemophilia Community Clinical Education Manager, I'm so proud to be part of an amazing team of nurses that provides support and resources for the hemophilia community. I'd love to connect with you and answer any questions you may have about HEMLIBRA.

## Let's get connected today!


**Directly:**

caito.diana@gene.com

(317) 703-4864 (call or text)

**Online:**

hemlibra.com/signup


**It's important to know that Clinical Education Managers do not provide medical advice. Please consult your doctor as needed.**

### What is HEMLIBRA?

HEMLIBRA is a prescription medicine used for routine prophylaxis to prevent or reduce the frequency of bleeding episodes in adults and children, ages newborn and older, with hemophilia A with or without factor VIII inhibitors.

### What is the most important information I should know about HEMLIBRA?

**HEMLIBRA increases the potential for your blood to clot. People who use activated prothrombin complex concentrate (aPCC; Feiba®) to treat breakthrough bleeds while taking HEMLIBRA may be at risk of serious side effects related to blood clots.**

**These serious side effects include:**

- **Thrombotic microangiopathy (TMA),** a condition involving blood clots and injury to small blood vessels that may cause harm to your kidneys, brain, and other organs
- **Blood clots (thrombotic events),** which may form in blood vessels in your arm, leg, lung, or head

Please see Important Safety Information, including **Serious Side Effects**, as well as the HEMLIBRA full Prescribing Information and Medication Guide.

## What is the most important information I should know about HEMLIBRA? (cont'd)

**Carefully follow your healthcare provider's instructions regarding when to use an on-demand bypassing agent or factor VIII, and the dose and schedule to use for breakthrough bleed treatment. If aPCC (Feiba®) is needed, talk to your healthcare provider in case you feel you need more than 100 U/kg of aPCC (Feiba®) total.**

**The most common side effects of HEMLIBRA include:** injection site reactions (redness, tenderness, warmth, or itching at the site of injection), headache, and joint pain. These are not all of the possible side effects of HEMLIBRA. You can speak with your healthcare provider for more information.

## What else should I know about HEMLIBRA?

**See the detailed "Instructions for Use" that comes with your HEMLIBRA for information on how to prepare and inject a dose of HEMLIBRA, and how to properly throw away (dispose of) used needles and syringes.**

- **Stop taking your prophylactic bypassing therapy the day before you start HEMLIBRA**
- **You may continue taking your prophylactic factor VIII for the first week of HEMLIBRA**

HEMLIBRA may interfere with laboratory tests that measure how well your blood is clotting and create an inaccurate result. Speak with your healthcare provider about how this may affect your care.

These are not all of the possible side effects of HEMLIBRA. Speak to your healthcare provider for medical advice about side effects.

Side effects may be reported to the FDA at (800) FDA-1088 or www.fda.gov/medwatch. You may also report side effects to Genentech at (888) 835-2555.

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