

INDICATIONS

PERCLOT Absorbable Hemostatic Powder is indicated in surgical procedures (except neurological and ophthalmic) as an adjunctive hemostatic device to assist when control of suture line bleeding or capillary, venous, and arteriolar bleeding by pressure, ligature, and other conventional procedures are ineffective or impractical.

IMPORTANT RISK INFORMATION

Do not inject or place **PERCLOT** into blood vessels such as artery or vein as potential for embolization and death may exist.

Do not use **PERCLOT** for treatment of severe or extreme bleeding.

Do not inject into bladder or ureteral lumen.

Single use only. Do not re-use. Do not re-sterilize. Re-use or reprocessing of a single use device may lead to contamination and compromised device function or structural integrity.

Safety and efficacy of **PERCLOT** have not been clinically evaluated in children (less than 21 years old) and pregnant or lactating women.

PERCLOT should be used with caution in the presence of infection or in contaminated areas of the body. If signs of infection or process develop where **PERCLOT** has been applied, re-operation may be necessary in order to allow drainage.

Safety and efficacy of **PERCLOT** in neurological and ophthalmic procedures have not been established.

Safety and efficacy of **PERCLOT** have not been clinically evaluated for use in controlling post-partum bleeding or menorrhagia.

Once hemostasis is achieved, excess **PERCLOT** should be removed from the site of application by irrigation and aspiration particularly when used in the pericardial cavity and around foramina of bone, orifices of bony canals, the spinal cord, and/or the optic nerve and chiasm. **PERCLOT** achieves its maximum swelling within 10 minutes when exposed to blood or other fluids. Dry, white **PERCLOT** should be removed. The possibility of the product interfering with normal function and/or causing compression of surrounding tissues due to swelling is reduced by removal of excess dry material.

The effect of this product on patients with known sensitivity to starch or starch-derived materials has not been studied.

The efficacy of **PERCLOT** in achieving hemostasis in cortical bone and spinal bleeding has not been studied in randomized clinical trials.

Blood vessels, suture line gaps, and large needle holes with a diameter of ≥ 2 mm must be ligated prior to **PERCLOT** application.

Do not apply more than 50g of **PERCLOT** in diabetic patients as it has been calculated that amounts in excess of 50g could affect the glucose load.

As with other hemostatic agents, do not apply **PERCLOT** to sites where there is negative peripheral venous pressure as material may be drawn into the vascular system potentially resulting in life-threatening thromboembolic events.

Bx Only. For sale and proper use please refer to full device instructions for Use for Contraindications, Warnings and Precautions.

ORDERING INFORMATION

PRODUCT	ORDER NUMBER	QUANTITY
PERCLOT 3g - 8cm Standard Tip	PCUS003	5 EA / 1 BX
PERCLOT 20cm Extender Tip	PCET000	5 EA / 1 BX (2 Applicators / Pouch)
PERCLOT 38cm Laparoscopic Tip	PCET008	10 EA / BX

For questions or ordering information, please contact your Baxter representative or call 888-229-0001.
advancedsurgery.baxter.com

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Baxter Elastometric Guide

Michael Hoelscher

A red circular graphic with a gradient, appearing as a stylized arrow or a partial circle, located to the right of the author's name.

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Table of Contents **Baxter Elastometric Guide**

1. Understanding the eBook Baxter Elastometric Guide
 - The Rise of Digital Reading Baxter Elastometric Guide
 - Advantages of eBooks Over Traditional Books
2. Identifying Baxter Elastometric Guide
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Baxter Elastometric Guide
 - User-Friendly Interface
4. Exploring eBook Recommendations from Baxter Elastometric Guide
 - Personalized Recommendations
 - Baxter Elastometric Guide User Reviews and Ratings

- Baxter Elastometric Guide and Bestseller Lists
- 5. Accessing Baxter Elastometric Guide Free and Paid eBooks
 - Baxter Elastometric Guide Public Domain eBooks
 - Baxter Elastometric Guide eBook Subscription Services
 - Baxter Elastometric Guide Budget-Friendly Options
- 6. Navigating Baxter Elastometric Guide eBook Formats
 - ePub, PDF, MOBI, and More
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- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Baxter Elastometric Guide
 - Highlighting and Note-Taking Baxter Elastometric Guide
 - Interactive Elements Baxter Elastometric Guide
- 8. Staying Engaged with Baxter Elastometric Guide
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Baxter Elastometric Guide
- 9. Balancing eBooks and Physical Books Baxter Elastometric Guide
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Baxter Elastometric Guide
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Baxter Elastometric Guide
 - Setting Reading Goals Baxter Elastometric Guide
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Baxter Elastometric Guide
 - Fact-Checking eBook Content of Baxter Elastometric Guide
 - Distinguishing Credible Sources

13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
14. Embracing eBook Trends
 - Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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