

September 2, 2025

Dear Hand & Stone Network,

We've got exciting news! Dermalogica's new **Pro Pen Microneedling System** has officially received **FDA 510(k) clearance** and will debut in early 2026 as a **Class II medical device** — this milestone reflects Dermalogica's long-standing commitment to advancing professional treatments through both **innovation and education**.

Dermalogica's **Pro Pen NanoInfusion Systems** (current and upcoming) and their accessories will continue to be classified as **cosmetic devices**. These are designed to gently exfoliate, smooth the skin, and boost product absorption — and that purpose will stay the same moving forward. Because the **Pro Pen NanoInfusion System** is intended for esthetic use and is non-invasive, it is exempt from the FDA's medical device pre-market requirements.

If you would like to learn more about the new Pro Pen Microneedling System or have any additional questions, please reach out to Danielle Guinn, Senior Sales Manager, National Accounts or Lorilee Williams, Senior Sales Director, Corporate Accounts.

Best,



Tim Bunch
General Manager, North America