# Why FDA-Registered Class II Colon Hydrotherapy Equipment Matters

When choosing a Colon Hydrotherapy provider, the equipment used is just as important as the training and experience of the therapist.

In the United States, Colon Hydrotherapy equipment should be **FDA-registered as a Class II Medical Device**. This classification means the manufacturer must meet specific standards for safety, performance, sanitation, and quality control.

Why is this important?

✔ **Patient Safety** – FDA-registered systems are designed with features intended to help provide a safe and comfortable treatment experience.

✔ **Sanitation Standards** – Modern Class II devices utilize disposable components and closed water pathways designed to reduce contamination risks between clients.

✔ **Backflow Prevention** – Properly engineered systems include safeguards that help prevent contaminated water from flowing backward into the clean water supply, an important feature for protecting both clients and facilities.

✔ **Quality Manufacturing** – Equipment manufactured in the USA under FDA regulations must comply with quality system requirements and ongoing oversight that help ensure consistency and reliability.

As consumers become more health-conscious, it is wise to ask your provider what equipment they use and whether it is FDA-registered for use in the United States.

At Living Water Rejuvenation Centers, we believe your health deserves the highest standards of safety, sanitation, and professionalism. That's why we proudly use **FDA-registered Class II Colon Hydrotherapy equipment manufactured in the USA**, providing our clients with confidence as they pursue their wellness journey.

Your health is worth asking questions—and choosing quality matters.