



Healing today. Curing tomorrow.

SANUWAVE Health, Inc.
Barry Jenkins, CFO
Bernie Laurel, VP of Sales and Marketing
678-578-0103
investorrelations@sanuwave.com

Lippert/Heilshorn & Associates
Anne Marie Fields (Investors)
212-838-3777
afields@lhai.com
Mackenzie Mills (Media)
212-838-3777
mmills@lhai.com

FOR IMMEDIATE RELEASE

**SANUWAVE TO PRESENT AT THE RODMAN & RENSHAW
13th ANNUAL GLOBAL INVESTMENT CONFERENCE**

ALPHARETTA, GA, September 7, 2011 – SANUWAVE Health, Inc. (OTCBB: SNWV), an emerging medical technology company focused on the development and commercialization of noninvasive, biological response activating devices in regenerative medicine, today announced that the Company will participate in the Rodman & Renshaw 13th Annual Global Investment Conference taking place September 12-13, 2011 at the Waldorf Astoria Hotel in New York City.

Christopher M. Cashman, President and CEO of SANUWAVE, will present a corporate overview and update on Monday, September 12, 2011 at 2:00 p.m. Eastern time.

The presentation will be webcast live at <http://www.wsw.com/webcast/rrshq20/snwv> and also at <http://www.sanuwave.com/investors/investorevents.html>, where it will be archived for 90 days.

Company management will be available for one-on-one meetings with investors participating in the Rodman & Renshaw Global Investment Conference. For those who would like to schedule an appointment with SANUWAVE's management, please contact Anne Marie Fields, Lippert/Heilshorn & Associates, Inc., at 212-838-3777 or at afields@lhai.com or contact your Rodman & Renshaw representative.

About SANUWAVE Health, Inc.

SANUWAVE Health, Inc. (www.sanuwave.com) is an emerging regenerative medicine company focused on the development and commercialization of noninvasive, biological response activating devices for the repair and regeneration of tissue, musculoskeletal and vascular structures. SANUWAVE's portfolio of products and product candidates activate biologic signaling and angiogenic responses, including new vascularization and microcirculatory improvement, helping to restore the body's normal healing processes and regeneration. SANUWAVE intends to apply its PACE[®] technology in wound healing, orthopedic/spine, plastic/cosmetic and cardiac conditions. Its lead product candidate for the global wound care market, dermaPACE[®], is CE marked for treatment of the skin and subcutaneous soft tissue and recently completed its highly positive pivotal Phase III, Investigational Device Exemption (IDE) clinical trial in the U.S. for the treatment of diabetic foot ulcers. SANUWAVE researches, designs, manufactures, markets and services its products worldwide, and believes it has demonstrated that its technology is safe and effective in stimulating healing in chronic conditions of the foot (plantar fasciitis) and the elbow (lateral epicondylitis) through its U.S. Class III PMA approved Ossatron[®] device, as well as stimulating bone and chronic tendonitis regeneration in

the musculoskeletal environment through the utilization of its Ossatron, Evotron™ and orthoPACE® devices in Europe.

Forward-Looking Statements

This press release may contain “forward-looking statements” within the meaning of the Private Securities Litigation Reform Act of 1995, such as statements relating to financial results and plans for future business development activities, and are thus prospective. Forward-looking statements include all statements that are not statements of historical fact regarding intent, belief or current expectations of the Company, its directors or its officers. Investors are cautioned that any such forward-looking statements are not guarantees of future performance and involve risks and uncertainties, many of which are beyond the Company’s ability to control. Actual results may differ materially from those projected in the forward-looking statements. Among the key risks, assumptions and factors that may affect operating results, performance and financial condition are risks associated with the marketing of the Company’s product candidates and products, unproven pre-clinical and clinical development activities, regulatory oversight, the Company’s ability to manage its capital resource issues, competition, and the other factors discussed in detail in the Company’s periodic filings with the Securities and Exchange Commission. The Company undertakes no obligation to update any forward-looking statement.
