

OrthoPulse™ Ultra

Industry Leading Innovation and Performance
Combined with Unmatched Clinical and Economic Value
for Acute and Chronic Musculoskeletal Pain

Extracorporeal Pulse Activation Technology (EPAT®) System

Powered by Storz Medical

Regenerative Medicine Redefined





OrthoPulse™ Ultra – The Next EPAT® Generation

The appeal of the new OrthoPulse™ Ultra acoustic pressure wave device lies in its inspiring design, the “Active-tip-control” hand-piece display and, the CuraMedix web site for expanding knowledge, education and awareness.

With more than 15,000 devices sold, the EPAT® line powered by Storz Medical is the top-selling acoustic pressure wave technology in the world. Compact dimensions, high quality and reliability, low maintenance costs, the silent “Air Power” drive and wide range of use, make the OrthoPulse™ an acoustic pressure wave icon in treatment of musculoskeletal pain.

EPAT® Technology Systems Non-invasive musculoskeletal treatment systems are comprised of a control unit, specialized hand-pieces and transmitters. EPAT® treatment systems can be utilized in the private practice, clinic, ASC and/or hospital environments.

Applications

Generally, acute or chronic musculoskeletal pain and/or pain that significantly impairs mobility or quality of life including:

- Foot and ankle
- Lower extremity
- Upper extremity
- Back and chest
- Shoulder
- Neck
- Myofascial trigger points



The Design

The pioneering Single Frame Casing and innovative three-dimensional design of the OrthoPulse™ Ultra combines unmatched performance, reliability and value.

The Hand-Piece

System functionality and Individual Parameter Settings (IPS-Controls) have been integrated into the new “Active-Tip-Control” display on the hand-piece make treating patients easier.

The CuraMedix Web Site

Access to the latest regenerative medicine treatment information including research, articles, physician and patient testimonials, and more.

Patient Care Redefined

Innovative Technologies for Advanced Healing

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Important Information for U.S. Customers:

Certain devices and references made herein to specific indications of use may have not received clearance or approval by the United States Food and Drug Administration. Practitioners in the United States should first consult with their local CuraMedix representative in order to ascertain product availability and specific labeling claims. Federal (USA) law restricts certain devices referenced herein to sale, distribution, and use by, or on the order of a physician, dentist, veterinarian, or other practitioner licensed by the law of the State in which she/he practices to use or order the use of the device.