

FOR IMMEDIATE RELEASE

Contact:
Paul Byerley
Chief Executive Officer
+1 320.259.4321
pbyerley@artossinc.com

Artoss Announces Completion of \$1 Million Funding Round

Financing allows expanded U.S. commercial and clinical presence

St. Cloud, MN, September 24, 2018 / -- Artoss, Inc. (St. Cloud, MN), a medical device company that is pioneering the use of advanced bone graft substitutes to treat multiple orthopaedic indications, announced the successful completion of its recent funding round of Unsecured Convertible Notes. The round, which began in 2017 with a target of raising \$500,000, was oversubscribed to twice that amount, representing a total of \$1.0 million.

James J. Cassidy, Ph.D., Managing Director of Artoss, Inc., the exclusive North American distributors for NanoBone[®] Bone Graft in orthopaedic surgery, said, "We have been overwhelmed by the interest shown in our second round of angel funding. We are especially grateful to those investors who participated in our 2016 funding round and chose to increase their investment in this round as well as to our new investors for their support of Artoss."

Paul Byerley, Managing Director of Artoss, Inc. said, "We are very pleased with our commercial success to-date and appreciate the support of our investors. Surgeon feedback on the clinical performance of NanoBone SBX Putty and QD remains extremely positive. We look forward to expanding our distribution network to put our superior bone grafting solution in the hands of more surgeons."

NanoBone has consistent and cost-effective outcomes in bone grafting in multiple orthopaedic indications. NanoBone achieves these outcomes with no preparation required, excellent handling for surgeons, creation of an osteogenic matrix within 14 days, and clinical and radiographic healing in as little as 8 weeks. The dramatic growth in NanoBone procedures further secures its growing market position as a leader in the bone repair space. There is growing acceptance by surgeons that not all bone graft substitutes are the same, given the rapid osteogenic matrix formation of NanoBone within 14 days.

###

About NanoBone

NanoBone was first cleared by the Food and Drug Administration (FDA) in 2015 after having been successfully used in Europe for ten years. Clinical publications have identified bone graft substitutes as an increasingly important component of many orthopaedic

procedures. Consistent healing in orthopaedic procedures is a significant unmet clinical need and one where NanoBone may provide an effective option.

NanoBone is a commercially available product in the U.S. intended to fill bony voids or gaps of the skeletal system (i.e., extremities, posterolateral spine and pelvis). These osseous defects are surgically created or the result of traumatic injury to the bone and are not intrinsic to the stability of the bony structure. NanoBone[®] SBX Putty resorbs and is replaced with bone during the healing process.

###

About Artoss, Inc.

Artoss, Inc. (St. Cloud, MN) is a medical device company that is a leading distributor of bioidentical bone graft substitutes to treat multiple orthopaedic indications and is dedicated to the development of tools and products for effectively treating a variety of orthopaedic conditions requiring surgical intervention. The company markets, NanoBone Bone Graft Substitute, which is gives clinical outcomes comparable to autograft with less postoperative pain and complications. Artoss, Inc. has an experienced management team with extensive experience in orthopaedic medical devices.