



ARTHREX ACP® AUTOLOGOUS CONDITIONED PLASMA INTRA-ARTICULAR (IA) INJECTIONS FOR KNEE OSTEOARTHRITIS (OA) CLINICAL TRIAL

Dr. Cole, Dr. Yanke and Dr. Weber
are currently enrolling patients in the Arthrex ACP®
Autologous Conditioned Plasma Clinical Trial at
Midwest Orthopaedics at Rush.

The purpose of this clinical trial is to evaluate the effectiveness of ACP IA injections using Arthrex's ACP Double Syringe System compared to Normal Saline IA injections. This treatment was designed to investigate the safety and effectiveness of ACP in treating pain in subjects with primary OA of the knee. The Arthrex ACP system allows rapid and efficient concentration of platelets and growth factors from autologous blood for use at the treatment site. If successful, possible benefits may include significant decrease in knee pain, improvement in knee function and range of motion.

Participants in the trial are randomized to receive either the ACP treatment or the Normal Saline in a 2:1 ratio. The study will follow participants for 12 months post treatment to assess functional outcomes, need for treatment revisions and adverse events.

Basic Inclusion Criteria:

- Between the ages of 18 and 70
- Has diagnosis of knee osteoarthritis

**REFER A
PATIENT!**

**For more information or to refer a patient
for enrollment in this trial, contact:**

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Further information regarding this trial can be found

at www.clinicaltrials.gov and

www.ACPclinicaltrial.com



For more information
on the trial please visit:
www.ACPClinicalTrial.com