



CURRICULUM VITAE

SIGNATURE:		DATE:	1/5/15
NAME: TITLE:	Jeremy B. McCandless, MD Principal Investigator	DATE UPDATED:	January 2015

RESEARCH SITE ADDRESS & PHONE:

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EDUCATION/TRAINING:

EDUCATION/TRAINING:	DEGREE <i>(if applicable)</i>	YEAR(s)	FIELD OF STUDY
University of Utah Hospitals and Clinics Salt Lake City, UT	Fellowship	2010	Adult Reconstruction
University of Utah and Hospitals and Clinics Salt Lake City, UT	Resident	2009	Orthopedic Surgery
University Hospitals of Cleveland, Cleveland, OH	Internship	2005	General surgery and Orthopedics
Case Western Reserve University College of Medicine, Cleveland, OH	M.D.	2004	Medicine
University of Cincinnati McMicken College of Arts and Sciences Cincinnati, OH	B.S.	1999	Biology

BOARD CERTIFIED:

BOARD CERTIFIED/ELIGIBLE	YEAR(s)	SPECIALTY
American Board	2012	Orthopedic Surgery
American Board	2009	Orthopedic Surgery

POSITIONS AND EMPLOYMENT:

2014-Present	Principal Investigator, TriWest Research Associates, LLC, El Cajon, CA
2014-Present	Private Practice, Jeremy McCandless, MD, El Cajon, CA
2010-2014	Private Practice, Hofmann Arthritis Institute, Salt Lake City, UT
2009-2010	Attending Physician, George E. Whalen VAMC, Salt Lake City, UT

PUBLICATIONS:

Hofmann, AA, McCandless J. "Posterior Cruciate Sacrificing Total Knee Arthroplasty"
Insall and Scott, Surgery of the Knee 5th Edition

Hofmann, AA, McCandless J. "Imageless Computer Navigation in TKA: The Simpler Wave of the Future" Insall and Scott, Surgery of the Knee 5th Edition

Hofmann, AA, McCandless J. "Highly Crosslinked Polyethylene in Total Knee Arthroplasty" Orthopedic Knowledge Update: Hip and Knee Reconstruction, American Academy of Orthopedic Surgeons

CLINICAL RESEARCH EXPERIENCE:

2009-Present, Principal Investigator
Outcome of Total Knee Arthroplasty in Obese versus Non-obese patients with Unresurfaced Patellas

2008-2009, Co-Principal Investigator
Computer Navigation of Knee Flexion: The Patellar Thickness Effect

2007-Present, Principal Investigator
Regional Osteoporosis and Muscle Mass Loss after Lower Extremity Trauma

Janssen: A Multicenter, Parallel-group Study of Long-term Safety and Efficacy of XXXX (sirukumab) for Rheumatoid Arthritis in Subjects Completing Treatment in Studies CNTO136ARA3002 (SIRROUND-D) and CNTO136ARA3003 (SIRROUND-T)

Santarus: A Randomized, Double-blind, Placebo-Controlled, Multiple Ascending Dose Study to Evaluate the Safety, Pharmacokinetics, Pharmacodynamics, and Efficacy of Escalating Doses of XXXX in Patients with Active Rheumatoid Arthritis with Inadequate Response to Disease-Modifying Anti-rheumatic Drug(s)

HGS: A 5-Year Prospective Observational Registry to Assess Adverse Events of Interest and Effectiveness in Adults with Active, Autoantibody-Positive Systemic Erythematosus Treated with or without BENLYSTA™ (belimumab) in Routine Clinical Practice

Teva Branded Pharmaceutical Products R&D, Inc.: A Phase 2, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Safety and Efficacy of Topically Applied XXXX (4% and 8% w/w Ointment) in Patients with Primary Osteoarthritis Affecting a Single Knee

Amgen Inc.: An Open-Label, Single-Arm Extension Study to Evaluate the Long-Term Safety And Efficacy Of XXXX In Subjects With Moderate To Severe Rheumatoid Arthritis

Flexion: A Double-Blind, Randomized, Parallel Group, Dose-Ranging Study to Assess the Safety and Efficacy of XXXX for the Treatment of Pain in Patients with Osteoarthritis of the Knee

Fidia Farmaceutici S.p.A.: A multi-center, parallel, double-blind, randomized, placebo-controlled study to evaluate the safety and effectiveness of XXXX, a new viscoelastic hydrogel, for the treatment of osteoarthritis of the knee

Novartis: A 56-week, randomized, multi-center, double-blind, placebo-controlled, phase IIa/IIb study to evaluate the safety and efficacy of XXXX on total lean body mass and physical performance in patients after surgical treatment of hip fracture

Takeda Development Center Americas, Inc. (TDC): A Phase 3, Randomized, Double Blind, Multicenter, Placebo Controlled Study to Evaluate the Efficacy and Safety of XXXX 40 mg XR, 80 mg XR, 40 mg IR and 80 mg IR in Subjects With Gout

Amgen: A Prospective, Observational study for the Psychometric Evaluation of Novel Migraine-Related Functional Impact Instrument in Subjects with Episodic and Chronic Migraine

Genentech: A Randomized, Double-Blind Trial assessing the Impact of Methotrexate Discontinuation on the Efficacy of Subcutaneous XXXX with Methotrexate Therapy

Liventa: Participate in a Clinical Research Registry AmnioClear™ LCT Knee Registry

Synergy: A Randomized, 12-Week, Double-Blind, Placebo-Controlled Study to Assess the Safety and Efficacy of Plecanatide (3.0 and 6.0 mg) in Patients with Chronic Idiopathic Constipation