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CURRICULUM VITAE

SIGNATURE:		**DATE:**	08-27-2019
NAME: **TITLE:**	Patrick C. Astourian, PA-C Sub Investigator	**DATE UPDATED:**	Aug 2019

RESEARCH SITE ADDRESS & PHONE:

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EDUCATION/TRAINING:

INSTITUTION AND LOCATION	DEGREE *(if applicable)*	YEAR(s)	FIELD OF STUDY
San Diego State University San Diego, California	B.S.	1998	Microbiology
Samuel Merritt College Oakland, California	M.S.	2002	Physician Assistant

BOARD CERTIFIED:

BOARD CERTIFIED/ELIGIBLE	YEAR(s)	SPECIALTY
Board Certified	2002	Physician Assistant

POSITIONS AND EMPLOYMENT:

2019-Present	Sub-Investigator Investigator, ACRC Studies, Poway, CA
2014-Present	Sub-Investigator Investigator, BioSolutions Clinical Research Center, La Mesa, CA
2011-Present	Physician Assistant, Grossmont Orthopedic Medical Group, La Mesa, CA
2008-2011	Physician Assistant, William Eves M.D, Chula Vista, CA
2004-2008	Physician Assistant, San Diego Arthritis Medical Clinic, , San Diego, CA
2003-2004	Physician Assistant, US NAVY, San Diego, CA
2002-2003	Physician Assistant, De Anza Urgent Care/Family Practice, Calexico, CA

PUBLICATIONS:

Ultrasound guided musculoskeletal injections, Patrick Astourian MS PA-C Issues and Answers in Treating Osteoarthritis
Date: 03-04-2006
Issue Number: 2

Regional Rheumatic Pain Syndrome: What You Should Know
Abstract/Deck: Offering a unique case study approach, this author discusses various forms of regional rheumatic pain syndrome and provides key treatment insights.
Author: By Patrick Astourian, MS, PA-C Date: 09-01-2006
Issue Number: 5

CLINICAL RESEARCH EXPERIENCE:

Sub-Investigator

Genentech Rheumatoid Arthritis 2004- 2008 (Efficacy Assessor)
Hoffmann La-Roche Rheumatoid Arthritis anti-TNF failure 2005-2008 (Efficacy Assessor) Genentech Rheumatoid Arthritis MTX naive 2005-2008 (Efficacy Assessor)
Genentech Rheumatoid Arthritis anti-TNF naive 2005-2008 (Efficacy Assessor) Celltech Rheumatoid Arthritis 2005- 2008 (Efficacy Assessor)
Scios-Rheumatoid Arthritis 2004- 2006 (Efficacy Assessor)
Hoffmann La-Roche Rheumatoid Arthritis anti-TNF nai've2005-2008 (Efficacy Assessor) GSK- Opioid induced bowel dysfunction 2005-2008
Cephalon- Chronic Low Back Pain 2005- 2008 Cephalon- Neuropathic Pain 2005- 2008
Bristol Meyers Squibb- Rheumatoid Arthritis 2005- 2008 Pfzier-Rheumatoid Arthritis 2005- 2008
Pfzier-Osteoarthritis 2005- 2008
GSK- Rheumatoid Arthritis 2005- 2008
Abbott- Osteoarthritis 2005-2008
Forrest- Fibromyalgia 2005- 2008
Zars- Osteoarthritis 2004- 2006

Liventa Bioscience ArnnioClear™ LCT Knee Registry.

Celution Prepared Adipose Derived Regenerative Cells in the Treatment of Osteoarthritis of the Knee: A Double-blind, Placebo Controlled, Multi-center Safety and Feasibility Study.

Centrexion a phase 2 study to evaluate the analgesic efficacy of two dose levels of CNTX-XXXX, compared to placebo at 4 weeks, when administered as a single (AI) injection to the index knee in patients with knee osteoarthritis with pain score 5-9.

Viking a Phase 2a Study ambulato1y and recovery from subjects l" hip fracture 2 to 7 weeks post injury.

Pfizer a phase 2a study a monoclonal antibody that binds to and inhibits the actious of nerve growth factor in patient with knee Osteoarthritis.

Axsome a phase 3 study to assess the Efficacy and Safety ofXXXX-02 administered orally to subjects with knee osteoarthritis associated with bone marrow lesions

Lannett a phase 3 investigation of topical application of Cocaine HCL 4% solution on safety and efficacy and Cocaine HCL I 0% solution on safety in local (topical) anesthesia for diagnostic procedures and surgeries on or through the accessible mucous membranes of the nasal cavities

Abbvie Phase 2a Study evaluating the safety, efficacy and Pharmacodynamic effects of XXX-XXX in patients with knee osteoarthritis
Pfizer Pharmaceuticals: A Phase ill. Multi-Center, Randomized, Double blind, Controlled study of the long-term analgesic efficacy of XX.XX alone or in combination with non-steroidal anti-inflammatory drugs (NSAIDS) versus NSAIDS alone in patients with Osteoarthritis of the knee or hip
Samumed Phase 2 Study 24 week, multicenter, randomized double blind placebo-controlled study Evaluating the safety and efficacy of XXXXX for the treatment of moderately to severely symptomatic knee osteoa1thritis

Regeneron Phase 3 Randomized double-blind, multi-dose, Placebo and Naproxen controlled study to evaluate the efficacy and safety of Fasinumab in patients with pain due to osteoarthritis of the hip or knee

Centrexion Phase 3, Randomized, Double-blind, Placebo-controlled, Single, lnjection,52-Week Study to Evaluate the Efficacy and Safety of an Intra-articular Injection of CNTX-4975-05 in Subjects with Chronic, Moderate-to-severe Osteoarthritis Knee Pain

A Randomized, Double-blind, Parallel Group, Multicenter Study to Compare the Pharmacokinetics, Phermacodynamics, Safety, and Efficacy of SAITIOI versus MabThera® versus Rituxan® in Patients with Rheumatoid Arthritis (RA).

A Phase 3, Multicenter, Observational Long-term Study Evaluating the Safety, Tolerability, and Efficacy of Treatment of SMXXXXX or Placebo Previously Injected in the Target Knee Joint of Subjects with Moderately to Severely Symptomatic Osteoarthritis

A Phase III, Multicenter, Randomized, Double-Blind Clinical Trial to Assess the Efficacy and Safety of Ciprofloxacin 0.3% plus Fluocinolone acetonide 0.025% Otic Solution Compared to Ciprofloxacin 0.3% Otic solution and to Fluocinolone acetonide 0.025% Otic Solution in the Treatment of Acute Otitis Externa (AOE)

A Phase 2 Randomized, Double-Blind, Placebo-Controlled Study to Evaluate Efficacy and Safety of BMS-XXXXX in Subjects with Systemic Lupus Erythematosus

Phase 2A, FXOOl6-XX-XXX, A Randomized, open-label, study comparing the systemic exposure to Triamcinolone Acetonide following a single Intra-articular injection of Extended release FXXXX, or immediate release TAcs (Triamcinilone Acetonide Suspension) in patients with osteoarthrosis of the shoulder (Glenohumeral) or hip