Jessica Watson, CCRC

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Research Affiliations

Offices

Preferred Research Partners, Inc.
11219 Financial Centre Parkway, Suite 320
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Preferred Research Partners Arkansas Center for Sleep Medicine
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Education:
Arkansas State University-Beebe
Pulaski Technical College
Associate of Applied Science in Business Technology with emphasis in Legal Assistant
Graduation Date: May 2010
3.65 GPA

Work Experience:
Preferred Research Partners, Inc., Little Rock, AR
August 2012 - Present
Clinical Research Coordinator
Position involves conducting study visits & ensuring patient safety, AE/SAE reporting, facilitating informed consent process, collection of IRB correspondence, patient recruitment, CRF and source documentation completion, laboratory collection and processing.

Licensure/Certifications:

- October 2014: Certified Clinical Research Coordinator
- BLS Provider 2024
- FibroScan certified

Research Experience:

- 2012: UCB, Restless Legs Syndrome in Adolescents – SP1004
- 2012: Bristol Myers Squibb, Type II Diabetes & Hypertension – MB102073
- 2012: Synergy Pharmaceuticals, Chronic Idiopathic Constipation – SP304-20210
- 2012: UCB, Restless Legs Syndrome in Adolescents – SP1005

- 2012: Bristol Myers Squibb, Type II Diabetes & Hypertension – MB102077
- 2012: Forest Research Institute, Hypertension – NAC-MD-01
- 2012: Ardea, Gout – RDEA594-301
- 2012: Orexigen, Obesity – NB-CVOT
- 2012: Novartis, High Triglycerides – CLCQ908C2201
- 2012: Furiex Pharmaceuticals, Diarrhea Predominant Irritable Bowel Syndrome – 27018966IBS3001
- 2012: Lexicon Pharmaceuticals, Diarrhea Predominant Irritable Bowel Syndrome – LX1033-1.201
- 2012: Tioga Pharmaceuticals, Diarrhea Predominant Irritable Bowel Syndrome – ASMP3001
- 2012: Shionogi, Inc., Opioid Induced Constipation – 1107V9221
- 2012: Tsumura Pharmaceuticals, Crohn's Disease – TU100P2T2
- 2012: Coronado Biosciences, Crohn's Disease – CNDO-201-003
- 2012: Shire, GERD – SPD557-206
- 2012: UCB, Healthy Adult, Phase I – SP1031
- 2012: UCB, Restless Legs Syndrome – RL0003
- 2012: GSK, Restless Legs Syndrome – RXP114025
- 2013: Ardea BioSciences, Gout – RDEA594-306
- 2013: Dr. Reddy's Laboratories, Chronic Idiopathic Constipation – DRL-USG01-L/2012
- 2013: Purdue Pharma L.P., Chronic Low Back Pain & Opioid-Induced Constipation – ONU3705
- 2013: Cubist Pharmaceuticals, Opioid-induced Constipation – 5945-SOIC-12-05
- 2013: Cubist Pharmaceuticals, Opioid-induced Constipation – 5945-OIC-12-04
- 2013: Ferring International Pharmascience Center US, Inc., Chronic Idiopathic Constipation – 000080
- 2013: Ferring International Pharmascience Center US, Inc., Chronic Idiopathic Constipation – 000081
- 2013: Synergy Pharmaceuticals, Inc., Chronic Idiopathic Constipation (OLE CIC) – SP304203-01
- 2013: Shionogi, Inc., Opioid Induced Constipation – 1315V9232
- 2013: Menarini, Diarrhea-Predominant Irritable Bowel Syndrome – NAK-07
- 2013: Synergy Pharmaceuticals, Inc., Chronic Idiopathic Constipation (CIC3) – SP304203-00
- 2014: Evoke, Diabetic Gastroparesis – METO-IN-003
- 2014: Evoke, Diabetic Gastroparesis – METO-IN-004
- 2014: Eisai, Long-Term incidence of MACE events & conversion to Type II Diabetes Mellitus in Obese & Overweight Subjects with Cardiovascular Disease or Multiple CV Risk Factors – APD356-G000-401
- 2014: A Phase 3, Randomized, Double Blind, Multicenter, Placebo Controlled Study to Evaluate the Efficacy and Safety of Febuxostat 40 mg XR, 80 mg XR, 40 mg IR and 80 mg IR in Subjects With Gout – Takeda Development Center Americas, Inc. – FEB-XR_301

- 2014: A Double-Blind, Placebo-Controlled, Parallel-Group, Multicenter, Multiregional, One Year Study to Assess the Efficacy and Safety of Twice Daily Oral Rifaximin Delayed Release Tablets for Induction of Clinical Remission with Endoscopic Response at 16 Weeks followed by Clinical and Endoscopic Remission at 52 Weeks in Subjects with Active Moderate Crohn's Disease – Salix Pharmaceuticals, Inc. – RECD3125
- 2014: A Randomized Double-Blind, Placebo Controlled, Flexible and Fixed Dose, Parallel Group Study of Extended-Release Lorazepam (EDG004) for the Treatment of Generalized Anxiety Disorder (GAD) – Edgemont Pharmaceuticals – EDG004-003
- 2014: A Randomized, Double-Blind, Placebo-Controlled, Crossover Study To Assess The Effects Of TC-6499 on Gastric Emptying Time In Diabetic Subjects with Gastroparesis – Targacept – TC-6499-12-CLP-005
- 2014: Xenon, RLS Genetic Biomarker – XEN-RLS
- 2015: Rhythm Pharmaceuticals, Diabetic Gastroparesis – RM-131-009
- 2015: Jazz Pharmaceuticals, Narcolepsy & Excessive Daytime Sleepiness – 14-002
- 2015: Jazz Pharmaceuticals, Obstructive Sleep Apnea & Excessive Daytime Sleepiness – 14-003
- 2015: Jazz Pharmaceuticals, Long-term extension study for Obstructive Sleep Apnea or Narcolepsy with Excessive Daytime Sleepiness – 14-005
- 2015: Neurocrine Biosciences, Inc., Tardive Dyskinesia – NBI-98854-1304 (KINECT3)
- 2015: Theravance, Diabetic or Idiopathic Gastroparesis – 0099
- 2015: Synergy Pharmaceuticals, Inc., Irritable Bowel Syndrome-Constipation – SP304203-04
- 2015: Forest Research Institute, Inc., Irritable Bowel Syndrome-Constipation &/or Chronic Idiopathic Constipation – LIN-MD-10
- 2015: Janssen Research & Development, Treatment-resistant Depression – ESKETINTRD3001 (TRANSFORM-1)
- 2015: Janssen Research & Development, Treatment-resistant Depression – ESKETINTRD3003 (SUSTAIN-1)
- 2015: Neurocrine Biosciences, Inc., Tardive Dyskinesia – NBI-98854-1402 (KINECT4)
- 2015: Naurex Inc., Major Depressive Disorder – NRX1074-C-202
- 2015: Gilead, Ulcerative Colitis – GS-US-326-1100
- 2015: Sunovion Pharmaceuticals, Adult Attention Deficit Hyperactivity Disorder (ADHD) – SEP360-301
- 2015: Ardelyx – IBS-C – TEN-01-301
- 2016: Ferring International Pharmascience Center US, Inc., Ulcerative Colitis – 000174
- 2016: Ferring International Pharmascience Center US, Inc., Ulcerative Colitis – 000175
- 2016: Conatus Pharmaceuticals, Non-alcoholic Steatohepatitis (NASH) – IDN-6556-12

- 2016: Synergy Pharmaceuticals, Irritable Bowel Syndrome with Constipation (IBS-C) – SP304203-06
- 2016: Janssen Research & Development, Treatment-resistant Depression – ESKETINTRD3008
- 2016: Eisai Pharmaceuticals, Elderly Insomnia – E2006-G000-304
- 2016: Luitpold Pharmaceuticals, Inc., Restless Legs Syndrome – 1VIT14037
- 2016: Braintree Laboratories, Chronic Constipation – BLI400-302
- 2016: Braintree Laboratories, Opioid Induced Constipation – BLI801-203
- 2016: Applied Proteomics – Colorectal Cancer Screening – API-CP001
- 2017: Takeda Pharmaceuticals – Diabetic or Idiopathic Gastroparesis – TAK906-1002
- 2017: Conatus Pharmaceuticals – Non-Alcoholic Cirrhosis – IDN-6556-17
- 2017: Shire Pharmaceuticals – Preschooler ADHD – SPD489-347
- 2017: Shire Pharmaceuticals – Preschooler ADHD – SPD489-348
- 2017: Allergan – Diabetic Gastroparesis – RLM-MD-01
- 2017: Braintree Laboratories, Bowel Prep for Colonoscopy – BLI4700-302
- 2017: Tonix Pharmaceuticals, Military-Related PTSD – TNX-CY-P301
- 2017: Vanda Pharmaceuticals, Inc. – Diabetic or Idiopathic Gastroparesis – VP-VLY-686-2301
- 2017: Janssen – Major Depressive Disorder – CNTO136MDD2001
- 2017: Allergan – Pediatric Major Depressive Disorder – LVM-MD-11
- 2017: Allergan – Diabetic Gastroparesis – RLM-MD-04
- 2018: Allergan – Diabetic Gastroparesis – RLM-MD-03
- 2018: Tonix Pharmaceuticals, Military-Related PTSD – TNX-CY-P303
- 2018: Janssen Research & Development, Insomnia – 42847922ISM2005
- 2018: Shire Pharmaceuticals – Preschooler ADHD PK – SHP465-112
- 2018: Shire Pharmaceuticals – Adolescent ADHD – SHP465-308
- 2018: Shire Pharmaceuticals – Adolescent ADHD – SHP465-309
- 2018: AbbVie – Testosterone Replacement Therapy for Hypogonadism – M16-100
- 2018: Otsuka Pharmaceuticals – Post Traumatic Stress Disorder – 331-201-00061
- 2018: Idorsia Pharmaceuticals – Insomnia – ID-078A301
- 2018: Idorsia Pharmaceuticals – Insomnia – ID-078A303
- 2018: Janssen Research & Development – Major Depressive Disorder – 67953964MDD2001
- 2018: Neos Therapeutics – Attention Deficit Disorder in Children 4-6 y/o – NT0202.1009
- 2018: Neos Therapeutics – Attention Deficit Disorder in Children 4-6 y/o – NT0202.1010
- 2018: Tonix Pharmaceuticals - Military-Related PTSD – TNX-CY-P306
- 2018: Allergan – Diabetic Gastroparesis – 3071-305-020
- 2018: Ironwood Pharmaceuticals – GERD – C3718-302
- 2019: Tonix Pharmaceuticals - PTSD – TNX-CY-P302
- 2019: Apnimed – Obstructive Sleep Apnea – APN-002
- 2019: Aptinyx – Post-Traumatic Stress Disorder – NYX-783-2004

- 2019: Axsome Pharmaceuticals – Treatment-Resistant Depression – AXS05-301
- 2019: Biohaven Pharmaceuticals – Obsessive Compulsive Disorder – BHV4157-202
- 2019: Otsuka Pharmaceuticals – Post Traumatic Stress Disorder – 331-201-00071
- 2019: Takeda – Obstructive Sleep Apnea – Phase 1 – TAK-925-2001
- 2019: LG Chem – Gout with Hyperuricemia – LG-GDCL002
- 2019: Applied Therapeutics - Diabetic Cardiomyopathy / Stage B Heart Failure at High Risk of Progression to Overt Heart Failure (Stage C Heart Failure) – AT-001-2001
- 2019: Adare Pharmaceuticals – Eosionophilic Erosive Esophagitis – SP-1011-003
- 2019: Allergan - Liver Fibrosis in Adult Subjects with Nonalcoholic Steatohepatitis – 3152-301-002
- 2019: Allergan – Constipation in Children 6-17 y/o – LIN-MD-64
- 2019: Allergan – Pediatric Participants with Functional Constipation (FC) or Irritable Bowel Syndrome with Constipation (IBS-C) – LIN-MD-66
- 2019: Allergan - Pediatric Participants (Age 2 to 5 Years) with Functional Constipation – LIN-MD-67
- 2019: Braintree Laboratories - Bowel Prep for Colonoscopy – BLI4900-302
- 2019: Phathom Pharmaceuticals – Erosive Esophagitis – EE-301
- 2019: Phathom Pharmaceuticals – H. Pylori – HP-301
- 2019: Vanda Pharmaceuticals – Diabetic or Idiopathic Gastroparesis – VP-VLY-686-3301
- 2019: Vibrant Ltd – Chronic Idiopathic Constipation – V270
- 2020: Takeda – Idiopathic Hypersomnia – Phase 1 – TAK-925-2002
- 2020: Neurogastrx – Diabetic or Idiopathic Gastroparesis – NG101-201
- 2020: Allergan – Major Depressive Disorder – 2006-308-008
- 2020: Otsuka – Pediatric Attention Deficit Hyperactivity Disorder – 405-201-00010
- 2020: Apnimed – Obstructive Sleep Apnea – APN-006
- 2020: Janssen – Treatment-resistant Depression – 54135419TRD4005
- 2020: Janssen – Major Depression & Insomnia – 42847922MDD3002
- 2020: Novavax – COVID-19 Vaccine – 2019nCoV-301
- 2020: Supernus – ADHD in Preschoolers – 812P401
- 2020: GlaxoSmithKline – Meningococcal B Vaccine - 205416
- 2021: Otsuka Pharmaceuticals – ADHD in Children – 405-201-00046
- 2021: Apnimed – Obstructive Sleep Apnea – APC-003
- 2021: Apnimed – Obstructive Sleep Apnea – APC-003-OLE
- 2021: Moderna – CMV Vaccine – mRNA-1647-P301
- 2021: Phathom Pharmaceuticals – Non-erosive Reflux Disease – NERD-201
- 2021: Acerus Pharmaceuticals – Ambulatory BP monitoring in Hypogonadism – NAT-2020-01
- 2021: Purdue Pharma LP – ADHD in Preschoolers & Children – ADA4004
- 2021: BlackThorn – MDD with Anhedonia &/or GAD – K2-MDD-201
- 2021: Relmada – Major Depressive Disorder – REL-1017-302

- 2021: Relmada – Major Depressive Disorder – REL-1017-310
- 2021: AbbVie – Pediatric Migraine - 3110-305-002
- 2021: AbbVie – Pediatric Migraine – 3110-306-002
- 2021: Otsuka Pharmaceuticals – ADHD in Children & Adolescents – 405-201-00016
- 2021: Otsuka Pharmaceuticals – ADHD in Children & Adolescents – 405-201-00017
- 2021: Otsuka Pharmaceuticals – ADHD in Children & Adolescents – 405-201-00021
- 2021: Revolo Pharmaceuticals – Eosinophilic Esophagitis – RVLO 121-04
- 2021: Eliem Therapeutics – MDD in Perimenopausal Females – ETX-020155-202
- 2022: Sommetrics – Sleep Apnea – SOM-029
- 2022: Mind Med – Generalized Anxiety Disorder – MMED008
- 2022: Merck – Treatment Resistant Depression – MK1942-006
- 2022: 89 Bio – NASH – BIO89-100-122
- 2022: Phathom Pharmaceuticals – Non-erosive Reflux Disease – NERD-301
- 2022: Alto Neurosciences – PTSD &/or MDD – ALTO-100-001
- 2022: Braintree Laboratories – Erosive Esophagitis – BLI5100-301
- 2022: Braintree Laboratories – Non-erosive Reflux Disease – BLI5100-302
- 2022: Ellodi Pharmaceuticals, L.P. – Eosinophilic Esophagitis – SP-1011-005
- 2022: Hepion Pharmaceuticals. – Non-alcoholic Steatohepatitis – HEPA-CRV431-207
- 2022: Idorsia Pharmaceuticals, Inc. – Pediatric Insomnia – ID-078A205
- 2022: Janssen – Bipolar Depression – 55308942BIP2001
- 2022: Merck – HPV Vaccine & mRNA COVID-19 Vaccine in 9-11 y/o – V503-076
- 2022: Novavax – Pediatric COVID-19 Vaccine – 2019nCoV-503
- 2022: Perception Neuroscience, Inc. – Treatment Resistant Depression – PCN-101-21
- 2022: Phathom Pharmaceuticals – Adolescents with Symptomatic GERD – VPED-102
- 2022: Supernus Pharmaceuticals – Child & Adolescent ADHD – 812P412
- 2023: Alto Neurosciences – MDD – ALTO-100-201
- 2023: AbbVie – Pediatric Episodic Migraine – M21-199
- 2023: AbbVie – Pediatric Episodic Migraine – M21-201
- 2023: Corium – Pediatric ADHD in ages 4-12 y/o – KP415.P01
- 2023: Corium – Pediatric ADHD – KP415.P02
- 2023: CinDome – Adult Diabetic Gastroparesis – CIN-102-123
- 2023: Tonix – Adult Major Depressive Disorder – TNX-TI-M201
- 2023: Tonix- Adult Chronic Migraine – TNX-OX-CM201
- 2023: Braintree Laboratories – Gastroesophageal Reflux Disease – BLI5100-303

- 2023: Phathom Pharmaceuticals – Adolescents with Symptomatic GERD – VPED-103
- 2023: Shionogi – COVID-19 & prevention in household contacts – 226T1331
- 2023: Apnimed – Sleep Apnea – APC-APN-304
- 2023: Lilly – Diabetic Peripheral Neuropathy Pain – J2P-MD-LXBD
- 2023: Supernus – Pre-schooler ADHD – 812P310
- 2023: Supernus / Navitor – Treatment Resistant Major Depressive Disorder – NAV-17A-007
- 2023: Bayer – Sleep Disturbances Associated with Menopause – 22423 (NIRVANA)
- 2023: Usona – Major Depressive Disorder – PSIL301
- 2024: Axsome – Major Depressive Disorder – SOL-MDD-301
- 2024: Axsome – Excessive Daytime Sleepiness with Shift Work Disorder – SOL-SWD-301
- 2024: Axsome – Major Depressive Disorder in Children & Adolescents – AXS-05-MDD-402
- 2024: Axsome – Major Depressive Disorder in Children & Adolescents (extension) – AXS-05-MDD-403
- 2024: AbbVie – Menstrual Migraine Prevention – M23-714
- 2024: Apnimed – Obstructive Sleep Apnea – APC-APN-305
- 2024: MindMed – Generalized Anxiety Disorder – MM120-300/301
- 2024: Apnimed – Obstructive Sleep Apnea – APC-APN-306
- 2024: Vistagen – Social Anxiety Disorder – PH94B-CL043
- 2024: Vanda – Diabetic & Idiopathic Gastroparesis – VP-VLY-686-3303
- 2024: Phathom – Pediatric GERD – VPED-104
- 2024: Alkermes – Narcolepsy – ALKS2680-202
- 2024: Neumora – Major Depressive Disorder – NMRA-335140-301
- 2024: Neumora – Major Depressive Disorder – NMRA-335140-501
- 2024: AbbVie – Migraine Prevention in Adults – M24-873
- 2024: Lilly – Chronic Weight Management & Obstructive Sleep Apnea – J1I-MC-GZBK/GSA2
- 2025: Alto Neuroscience – Bipolar Depression – ALTP-100-211
- 2025: Cindome Pharma, Inc. - A Randomized, Double-Blind, Placebo-Controlled, Multicenter Study to Evaluate the Efficacy and Safety of CIN-102 (Deudomperidone) in Adult Subjects With Idiopathic Gastroparesis – CIN-102-124
- 2025: Phathom Pharmaceuticals, Inc. - A Phase 2, Randomized, Double-Blind, Multi-Center, 3-Part Study in Adult and Adolescent Subjects with Eosinophilic Esophagitis (EoE) to Evaluate the Safety and Efficacy of Vonoprazan 10 mg and 20 mg Compared to Placebo After 12 Weeks and to Evaluate the Safety and Efficacy of Vonoprazan 10 mg and 20 mg Up to 52 Weeks – EoE-201

- 2025: Eli Lilly & Company - An Adaptive, Dose-Ranging, Phase 2 Study of Eltrekibart and Mirikizumab for the Treatment of Adult Patients with Moderately to Severely Active Ulcerative Colitis – I7J-MC-DSAG
- 2025: Sanofi-Aventis - An induction study to investigate the efficacy and safety of duvakitug in participants with moderately to severely active Ulcerative Colitis – EFC18325
- 2025: Sanofi-Aventis - A multicenter, multinational, randomized, double-blind, placebo-controlled Phase 3, induction study to evaluate the efficacy and safety of duvakitug in participants with moderately to severely active Crohn's Disease – EFC18326
- 2025: Sanofi-Aventis - A multicenter, multinational, randomized, double-blind, placebo-controlled Phase 3, maintenance study to evaluate the efficacy and safety of duvakitug in participants with moderately to severely active Crohn's disease – EFC18327
- 2025: Sanofi-Aventis - A Multicenter, Multinational, Randomized, Double-blind, Placebo-Controlled, Phase 3 Maintenance Study to Evaluate the Efficacy and Safety of Duvakitug in Participants with Moderately to Severely Active Ulcerative Colitis – EFC18359
- 2025: Mirador Therapeutics, Inc. - A Phase 2, Multi-Center, Platform Study to Evaluate the Safety, Efficacy, Pharmacokinetics, and Pharmacodynamics with Multiple Therapies in Participants with Active Crohn's Disease or Active Ulcerative Colitis (ASCEND-IBD – MT-100-201/PROBEUC/PROBECD
- 2025: Orion Corporation - EFFICACY AND TOLERABILITY OF TASIPIMIDINE AFTER 3 REPEATED BED-TIME DOSES IN PATIENTS WITH INSOMNIA DISORDER WITH A 4-WEEK EXTENSION PART – 3110012
- 2025: VistaGen Therapeutics, Inc. - A US, Phase 3, Multicenter, Randomized, Double-blind, Placebo-controlled Clinical Trial of Fasedienol Nasal Spray for the Acute Treatment of Anxiety Induced by a Public Speaking Challenge in Adult Subjects with Social Anxiety Disorder, with an Open-Label Extension (PALISADE-3) - PH94B-CL042
- 2025: Mind Medicine, Inc. – Major Depressive Disorder (MDD) – MM123-310 (EMERGE)
- 2025: Atai Pharmaceuticals – Treatment Resistant Depression (TRD) – VLS-01-203
- 2025: AbbVie – Pediatric Chronic Migraine ages 12-17 y/o – M23-712
- 2025: Alkermes – Idiopathic Hypersomnia – ALKS2680-203
- 2025: Alkermes – Narcolepsy – ALKS2680-301
- 2025: Mineralys Therapeutics, Inc. – Moderate to Severe Sleep Apnea (OSA) – MLS-101-207
- 2025: Transcend Therapeutics – Post-Traumatic Stress Disorder (PTSD) - TSND201-PTSD-301
- 2025: Axsome Therapeutics – Fibromyalgia – AXS-14-FM-301

Training Documentation:

- June 2024: IATA training
- October 2023: Stericycle IATA Training
- August 2023: CITI GCP training

- October 2021: IATA training
- June 2021; CITI GCP training
- September 2014: IATA training
- July 2014: National Institutes of Health web-based training on “Protecting Human Research Participants”
- March 2014: Stericycle Biohazardous Waste Training
- March 2014: Stericycle DOT Hazardous Materials Training
- October 2013: Stericycle Blood Bourne Pathogen Training
- September 2012: National Institutes of Health web-based training on “Protecting Human Research Participants”
- September 2012: IATA training
- September 2012: GCP Manual/Code of Federal Regulations Review

  Phlebotomy Training
  Informed Consent Training
  Vital Signs Training
  EKG Training
  Coordinator Training
  Job Shadowing
  Blood Borne Pathogen Training
  CRF Training
  SOP Review
  Lab Processing Training
  HIPAA Training