

CURRICULUM VITAE
OF
JEANNE M. CARROLL, RN, BA, CCRC

BUSINESS ADDRESS

Parkview Research Center
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Fort Wayne, Indiana 46845

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EDUCATION

1992 – 1995 Associate of Science, Nursing
Lutheran College, Fort Wayne, IN

1985 – 1989 Bachelor of Arts, Psychology
Huntington College, Huntington, IN

1981 – 1985 Diploma
Snider High School, Fort Wayne, IN

CURRENT POSITION

04/2019 – Present Research Project Lead
Parkview Research Center

PROFESSIONAL EXPERIENCE

05/2018 – 03/2019 Neurosciences Service Line Manager
Parkview Health

10/2016 – 05/2018 Neurosciences Service Line Clinical Manager
Parkview Health

02/2015 – 10/2016 Stroke & Clinical Research Coordinator
Parkview Health

10/2009 – 01/2013 Program Coordinator
StrokeCareNow Network

06/2002 – 02/2015 Clinical Research Coordinator
Parkview Research Center

07/2001 – 04/2002 Clinical Research Coordinator
Northeast Indiana Research, LLC

08/1999 – 2003 Staff Nurse, Emergency Department
Parkview Hospital Randallia

04/1997 – 08/1999 Staff Nurse, Surgical/Trauma ICU
Parkview Hospital Randallia

02/1996 – 04/1997 Staff Nurse, ICU/Medical/Surgical/Telemetry
Caylor-Nickel Medical Center

01/1991 – 12/1995 Social Worker
Big Brothers/Big Sisters of Greater Fort Wayne

PROFESSIONAL LICENSURE

Indiana State Board of Nursing, Registered Nurse License (#28131480A)

AFFILIATIONS

Parkview Regional Medical Center	11109 Parkview Plaza Drive Fort Wayne, Indiana 46845
Parkview Hospital Randallia	2200 Randallia Drive Fort Wayne, Indiana 46805
Parkview Physicians Group	10501 Corporate Drive Fort Wayne, Indiana 46845
Parkview Mirro Center for Research & Innovation	10622 Parkview Plaza Drive Fort Wayne, Indiana 46845
Parkview Research Center Clinical Trials Office	3942 New Vision Drive Fort Wayne, Indiana 46845

CERTIFICATIONS

2019 – Present QPR Suicide Prevention Gatekeeper Training/Certification

2006 – Present Certified Clinical Research Coordinator

PUBLICATIONS, ABSTRACTS & PRESENTATIONS

- 2023 Drouin M, Flanagan M, Carroll J, Kerrigan C, Henry H, Toscos T. Piloting a Peer Support Program for Patients Who Screen Positive for Intimate Partner Violence, Suicidal Ideation, and Depression. *Healthcare*. 2023; 11(17):2422. <https://doi.org/10.3390/healthcare11172422>
- 2023 Li, J., Bohn, C., Todd, N., Pater, J., Carroll, J., Henriksen, B., Chang, F.L. (April 2023). Peripheral Neuropathy in Long-COVID Patients: Demographic Distribution and Risk Factors. American Association of Neurology annual meeting.
- 2022 Pater, J., Chang, F.L., Carroll, J., Stienecker, R.S., Toscos, T., Guha, S. (November 2022). Charting the Changing Nature of Post-COVID Symptoms: Initial Findings from a Longitudinal Study. Presented at the American Medical Informatics Association Annual Meeting. Washington D.C.
- 2022 Bohn, C., Li, J., Todd, N., Pater, J., Carroll, J., Henriksen, B., and Chang, F.L. (August 2022). Predicting Cognitive Decline in Long-COVID Patients: A Demographic and Comorbid Analysis Using BrainCheck Cognitive Assessment. Presented at the 2022 IU Student Research Fellowship (SERF) Research Symposium. Fort Wayne, IN.
- 2022 Li, J., Bohn, C., Todd, N., Pater, J., Carroll, J., Henriksen, B., and Chang, F.L. (August 2022). Peripheral Neuropathy in Long-COVID Patients: Demographic Distribution and Medical Risk Factors. Presented at the 2022 IU Student Research Fellowship (SERF) Research Symposium. Fort Wayne, IN.
- 2019 Fawver, J., Flanagan, M., Carroll, J., Mirro, M. (2019) The association between COMT genotype and bupropion treatment response in outpatients with major depressive disorder. Poster presentation, Psych Congress Conference
- 2013 Hatzigeorgiou, M., Rao, P., Litchin, N., Khatri, R., Carroll, J., and Chang, F. (2013) The Feasibility of iPad Technology for Remote Stroke Assessment when in Moving Ambulance. Poster presentation, Midwest Alliance for Health Education (MAHE) Conference
- 2012 Binz, S., Khatri, R., Carroll, J., Prusakov, P., and Chang, F. (2012) The Feasibility of iPad Technology for Remote Acute Ischemic Stroke Assessment. Poster presentation, Midwest Alliance for Health Education (MAHE) Conference

CLINICAL RESEARCH EXPERIENCE

- | | | |
|------|---------------------------|----------------------------|
| 2015 | PATH Hypothermia Registry | University of Pennsylvania |
| | MaRISS Stroke Study | University of Miami |

	CREST 2	NIH Sponsored
2014	SOCRATES Minor Stroke/TIA Study	Astra-Zeneca
	MAGNIFY Spinal Cord Injury Study	Acordia Therapeutics, Inc.
2013	SILVER AMI: Risk Stratification in Older Persons with Acute Myocardial Infraction	NIH Sponsored
	PRISMS Stroke Study	Genentech
	SWIFT PRIME Ischemic Stroke Study	Covidien
	EQUALITY On the Road Telestroke Study	Parkview Research Center/ Ft. Wayne Neurological Center
	ATACH II ICH Study	NIH Sponsored
	CPT-MD-32 Bacteremia Study	Cerexa
2012	EQUALITY Telestroke Study	Parkview Research Center/ Ft. Wayne Neurological Center
2011	OSPREY SFA Stent Study	Terumo Medical Corporation
2011	SuperNOVA SFA Stent Study	Boston Scientific Corporation
	ATACH-II ICH Stroke Study	NIH Sponsored
2010	POINT TIA/Minor Stroke Study Pfizer Ischemic Stroke Study	NIH Sponsored Pfizer
	NEST-3 Ischemic Stroke Study	PhotoThera
	ICTUS 2/3 Ischemic Stroke Study	NIH Sponsored
	The Celestial Study	Biotronik, Inc.
	The IMPACT Study	Biotronik, Inc.
	S-ICD System Clinical Investigation	Cameron Health
	The Quick Flex Study	St. Jude Medical
2009	The SJ4 Study	St. Jude Medical

	PICS Study	Penumbra, Inc.
	ORION Iliac Stent Study	Boston Scientific Corporation
2008	RESOLUTE US Coronary Stent Trial VIRGO AMI Registry	Medtronic, Inc. NIH Sponsored
	Tracer Thrombin Receptor Antagonist	Schering-Plough
	MERCI Registry	Concentric Medical, Inc.
2007	NEST-2 Ischemic Stroke	PhotoThera
	ANCROD Ischemic Stroke	Neurobiological Technologies, Inc.
	MERCK Ischemic Stroke	MERCK & Co., Inc.
	CHOICE Carotid Stent Trial	Abbott Vascular Devices
	PROTECT Carotid Stent Trial	Abbott Vascular Devices
2006	EXACT Carotid Stent Trial	Abbott Vascular Devices
2005	FAST Hemorrhagic Stroke	Novo Nordisk
2005	DIAS 2 Ischemic Stroke	Forest Laboratories
	ACT I – Carotid Revascularization Endarterectomy vs. Stenting Trial	Abbott Vascular Devices
	TRIUMPH Cardiogenic Shock ETC-588-007 Reverse Lipid Transport	Arginox Pharmaceuticals, Inc. Esperion Therapeutics
2004	CHANT Hemorrhagic Stroke	Astra-Zeneca
	SAINT II Ischemic Stroke	Astra-Zeneca
	CREST-Carotid Revascularization Endarterectomy vs. Stenting Trial	NINDS/University of Medicine and Dentistry of New Jersey
2003	Pfizer A1611005 Ischemic Stroke	Pfizer
	RReact Ono Ischemic Stroke	Ono Pharmaceutical Co., Ltd.
	COMPASS-HF – Chronicle offers Management to Patients with Advanced Signed and Symptoms of Heart Failure	Medtronic, Inc.

2002	Informatics/CPOE	Northeast Indiana Innovation Center
	Security Carotid Stenting Trial	Abbott Vascular Devices/Perclose
	Carotid Stent IDE Trial	Parkview Research Center/Parkview Hospital/ Fort Wayne Cardiology
	Randomized, Double-Blind, Placebo-Controlled study of XXXX Provided to Patients with Erectile Dysfunction	Eli Lilly
	Randomized, Placebo-controlled, Double-Blind, Parallel Design Phase 3 Trial of the Efficacy and Safety of XXXX in Male Patients with Erectile Dysfunction	NexMed
2002	A Multicenter, Fixed Dose Study with a Double-Blind, Randomized, Placebo-Controlled, Parallel Group Phase and An Open Label Phase, to Investigate the Time-to-Onset of Activity of XXX	Pfizer
	A Randomized, Placebo-controlled, Double-Blind, Crossover Design Phase 2 Study of the Efficacy and Safety of XXXX in Patients with Erectile Dysfunction using Penile Rigidity and Tumescence Monitoring	NexMed
	Randomized, Double-Blind, Crossover Study to Investigate the Time-To-Onset of a Single Oral Dose Of 20 mg XXXX	Bayer
	Two Phase, Double-Blinded, Randomized, Parallel Group Design, Multicenter Study of XXX 0.4 mg Versus Placebo In Male Patients with Acute Urinary Retention Related to Benign Prostatic Hyperplasia	Boehringer Ingelheim

2001-2002	Randomized, Placebo Controlled, Double-Blind, Crossover Design Pilot Trial of the Efficacy and Safety of XXXX for Women in Women with Female Sexual Arousal Disorder	Qualilife
2001	Preliminary Efficacy Study in Pre-menopausal Women with Normal or Impaired Sexual Function Due to Acquired Arousal And/or Orgasm Disorder Comparing XXXX 1.0 mg To Placebo: Double-Blind with 8-Week home Treatment	Pharmacia

Jeanne M. Carroll, RN, BA, CCRC

Date