

**CURRICULUM VITAE
FOR**

Ronald L. Baird, DO

3364 North 500 East

Provo, UT 84604

Phone: (801) 995-2634

dr.ronald.baird@gmail.com

EDUCATION AND CERTIFICATIONS

- 1998 Board Certification in Family Practice Medicine
American Board of Family Medicine
- 1998 Family Practice Residency
Utah Valley Family Practice Residency,
IHC Hospital, Provo, Utah
- 1996 General Rotating Internship
Park Lane Hospital, Kansas City, MO
- 1995 Doctor of Osteopathic Medicine
University of Health Science College of Osteopathic Medicine
Kansas city, MO
- 1990 Bachelors of Science, Biochemistry
Brigham Young University, Provo, Utah

EMPLOYMENT HISTORY:

- 2020 – Present **Olympus Healing Center - Medical Director**
2870 E 3300 S
Salt Lake City, UT
- 2016 – Present **Chateau Recovery - Medical Director**
375 Rainbow Ln
Midway, UT
- 2017 – 2019 **Cirque Lodge Recovery - Medical Director**
RR 3 Box A-10
Sundance, UT

EMPLOYMENT HISTORY (continued):

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|-------------------|---|
| 2017 – 2020 | Copper Sage Recovery - Medical Director 14014 S 2200 W Bluffdale, UT |
| 2016 – 2019 | Pinnacle Recovery - Medical Director 6196 South Holladay Blvd Holladay, UT |
| 2016 – 2018 | Daylight Recovery - Medical Director 672 East Union Square Sandy, UT |
| 2013 – 2017 | True North Treatment Center - Medical Director 234 N Orem Blvd Orem, UT |
| 2008 - 2010 | Clinical Investigator Aspen Clinical Research, LLC Orem, UT |
| 1998 – 2010 | President/Owner – Private Practice Canyonview Family Practice Provo, UT |
| 1997 – 2010 | Part-time Emergency Room Coverage Orem Community Hospital Orem, UT |
| Jul '98 – Nov '98 | Full-time Faculty Member Utah Valley Family Practice Residency |
| 1997 - 1998 | Weekend Emergency Room Coverage Sanpete Valley and Delta Hospitals |
| 1992 – 1993 | Teaching Assistant University of Health Sciences, Physiology Dept. |
| 1992 – 1993 | Laboratory Teaching Assistant University of Health Sciences, Osteopathic Manipulation Dept. |
| 1992 (summer) | Teaching Assistant University of Health Sciences, Anatomy Dept. |
| 1990 – 1991 | Laboratory Technician/Biochemist Murdock Health Care, Quality Assurance Dept. |

PROFESSIONAL LICENSE:

322409-1204 State of Utah Medical License
Active License
Expires 05/31/2022

PROFESSIONAL MEMBERSHIPS

- American Academy of Family Practice
- American College of Family Practitioners
- Utah Medical Association

PROFESSIONAL AFFILIATIONS:

- Utah Valley Regional Medical Center, Provo, UT
- Orem Community Hospital, Orem, UT
- Timpanogos Regional Hospital, Orem, UT

REFERENCES:

- References available upon request:

CLINICAL RESEARCH TRAINING:

THERAPEUTIC AREAS OF RESEARCH EXPERIENCE

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|----------------------|----------------------------------|
| ADME | Phase I, II clinical trials |
| Alzheimer's | Phase I, II clinical trials |
| Cardiology | Phase II, III clinical trials |
| Diabetes | Phase II, III clinical trials |
| Hypercholesterolemia | Phase III clinical trials |
| Hypertension | Phase III clinical trials |
| Acute / Chronic Pain | Phase I, II, III clinical trials |

DETAILED CLINICAL TRIAL EXPERIENCE:

“A Phase 2a, Randomized, Blinded, Placebo and Active controlled, 2-Period Crossover Study to Assess the Analgesic Efficacy, Safety, and Tolerability of XXXXXXXXXX in subjects with Post Herpetic Neuralgia (PHN)”

“A Multi-Center, Double Blind, Placebo-Controlled, Study to evaluate the Safety and Efficacy of Oral XXXXXXXXXX in Subjects with Euvolemic Hyponatremia”

“A Double Blind, Efficacy and Safety of Duloxetine versus Placebo in the Treatment of Children and Adolescents with Major Depressive Disorder”

“A Phase I, Multi-Centered, Randomized, Placebo-Controlled, Double Blind Study to assess the Safety, Pharmacokinetics, Pharmacodynamics, and Immunogenicity of Intravenous XXXXXXXXXX administered in a Single Dose, Dose Escalation stage followed by a Multi-Dose. Parallel Treatment stage in patients with Mild to Moderate Alzheimers”

“A Phase IV, Double-Blind, Multi-Center, Placebo-Controlled, Randomized Withdrawal, Safety and Efficacy Study of XXXXXX in Adults age 18-55 with Attention Deficit Hyperactive Disorder (ADHD)”

“A Randomized, Double-Blind, Placebo Controlled, Dose ranging study of the Safety and Efficacy of XXXXXXXXXX in Subjects with Typee II Diabetes Mellitus on Stable Metformin Therapy”

“A Multi-center, Randomized, Controlled Study to Investigate the Safety and Tolerability of XXXXXXXXXX vs. Standard Medical Care in Treating Iron Deficiency Anemia”

“A Multi-center, Randomized, Controlled Study to Investigate the Safety and Tolerability of A Single Dose of XXXXXXXXXX vs. Standard Medical Care in Treating Iron Deficiency Anemia in Subjects Who are Not Dialysis Dependent”

“A Multi-center, Randomized, Active Controlled Study to Investigate the Efficacy and Safety of XXXXXXXXXX in Patients with Iron Deficiency Anemia (IDA)”

“A 24-Week, Multicenter, Double-Blind, Randomized, Parallel-Group, Dose Ranging Study of the Efficacy and Safety of oral doses of XXXXXXXXXX 5, 10, and 30 mg and placebo on top of an established treatment regimen of either olanzapine, risperidone/paliperidone, quetiapine or aripirazole monotherapy in the treatment of Cognitive Impairment in Schizophrenia.”

“A Randomized, Double-Blind, Active-Controlled Crossover Study to Evaluate the Efficacy and Safety of XXXXXXXXXX Compared With Immediate-Release Oxycodone for the Management of Breakthrough Pain in Opioid Tolerant Patients With Chronic Pain, Followed by a 12-Week Open-Label Extension to Evaluate the Impact of XXXXXXXXXX on Patient Outcomes”

“A Randomized, Double-Blind, Placebo-Controlled, Dose-Ranging, Exploratory, 28-Day Study to Examine the Effects of XXXXXXXXXX on Blood Pressure and Glucose Tolerance in Patients With Mild to Moderate Hypertension and Impaired Glucose Tolerance”

“The effect of insulin detemir in combination with XXXXXXXXXX and metformin compared to XXXXXXXXXX and metformin in subjects with type 2 diabetes. A 26-week, randomised, open-label, parallel-group, multicentre, multinational trial with a 26-week extension”

“A Randomized, Double-blind, Parallel-Group, Multicenter Study to Evaluate the Retention Rate, Efficacy, Safety, and Tolerability of XXXXXXXXXX, Topiramate and Levetiracetam as Adjunctive Therapy in Subjects with Partial Onset Seizures”

“Multicenter, Double-Blind, Randomized, Placebo-Controlled, Three-Arm, Parallel Group Study to Evaluate the Efficacy and Safety of XXXXXXXXX as Adjunctive Therapy in Subjects with Refractory Partial Seizures due to Epilepsy on up to Two Concomitant Antiepileptic Medications”

“Multicenter, Open-Label Extension Study to Evaluate the Long-Term Safety and Tolerability of XXXXXXXXX as Adjunctive Therapy in Subjects with Refractory Partial Epilepsy on up to Two Concomitant Antiepileptic Medications”

“A Randomized, Double-Blind, Placebo- and Active-Controlled, Parallel-Group, Multicenter Study to Determine the Efficacy and Safety of XXXXXXXXX When Used in Combination With Metformin Compared With Metformin Plus Sitagliptin, Metformin Plus Glimepiride, and Metformin Plus Placebo in Subjects With Type 2 Diabetes Mellitus”

“A Randomized, Open-Label, Parallel-Group, Multicenter Study to Determine the Efficacy and Long-Term Safety of XXXXXXXXX Compared With Insulin in Subjects With Type 2 Diabetes Mellitus”