

February 21, 2016





www.theautismdoctor.com



# Low Dose Naltrexone & Autism Spectrum Disorder



# Low Dose Naltrexone & Autism Spectrum Disorder

In her 2006 report, Dr. Jaquelyn McCandless concluded, "As an effective, nontoxic, non-addicting, and inexpensive behavioral and immunomodulating intervention, LDN is joining our biomedical arsenal to help more and more children recover from autism as well as helping anyone with autoimmune diseases and cancer."

Background - describe ASD and Epidemic Proportions

Diagnosis - new DSM 5.0 criteria

Treatment - Traditional therapies Medical therapies Biomedical therapies

LDN protocols

**Recent Results** 

Conclusions



# An Alternative Biomedical Approach to Autism





# An Alternative Biomedical Approach to Autism





# An Alternative Biomedical Approach to Autism



![](_page_6_Figure_1.jpeg)

![](_page_7_Figure_1.jpeg)

![](_page_8_Figure_0.jpeg)

# Autism **Spectrum** Disorder Implementation of DSM 5.0

![](_page_9_Picture_1.jpeg)

![](_page_9_Picture_2.jpeg)

AMERICAN PSYCHIATRIC ASSOCIATION

![](_page_9_Picture_4.jpeg)

DSM IV to DSM 5.0:

![](_page_10_Figure_2.jpeg)

![](_page_10_Picture_3.jpeg)

#### DSM-5 AUTISM SPECTRUM DISORDER

persistent social communication & social interaction

deficits in social-emotional reciprocity

deficits in nonverbal communicative behaviors used for social interaction

deficits in developing, maintaining and understanding relationships

![](_page_11_Picture_6.jpeg)

![](_page_11_Picture_7.jpeg)

restricted & repetitive behavior patterns

stereotyped or repetitive motor movements

insistence on sameness or inflexible adherence to routines highly restricted, fixated interests OR

hyper or hyper reactivity to sensory input

unusual interest in sensory aspects of the environment

©TheAutismDoctor.com

![](_page_11_Picture_14.jpeg)

Genetic – Fragile X, Rett's S., Trisomy 21?, Copy number variations

Gastro-intestinal

![](_page_12_Picture_3.jpeg)

- Most common in practice
  - If a person acts like they have 'ants in their pants', sometimes they actually do!
- Immunologic Asthma, eczema, food sensitivities, frequent ear infections, ?vaccination Microbiome alterations
- Syndromes Down's S., Metabolic disturbances, Unclassified
- Birth complications Cerebral palsy, 'Mental Retardation', "Premies'
- Metabolic & Nutritional Vitamin deficiency, Lipid abnormalities, Picky eaters, MTHFR

Boy vs. Girl

![](_page_12_Picture_11.jpeg)

![](_page_12_Picture_12.jpeg)

### Autism Spectrum Disorders Medical Workup

H&P

Specialized Audiology MRI EEG

Chromosomes

CBC

**Metal Levels** 

**Thyroid Screen** 

Appropriate additional baseline information

![](_page_13_Picture_8.jpeg)

### Autism Spectrum Disorders Medical Workup

#### G-I Health –

Comprehensive evaluation Toilet training can be much more effective when gut health is restored.

#### Food Allergies -

The presence of immunoglobulins indicates inflammation results. Reducing overall energy-depleting reactions for use elsewhere

#### Vitamin levels –

Low levels in most of our patients D3 especially important in immune health

#### **Comprehensive metabolic profile**

Baseline, drug tolerance and interactions

#### Lipid panel

High – nutrition? Low – eye contact, immune health

![](_page_14_Picture_11.jpeg)

### Autism Spectrum Disorders Available Treatments

#### Conventional

Anxiety

Abilify Risperdal Intuniv

Stimulant

Methamphetamine Ritalin Strattera

SSRI

Zoloft Prozac

Anti-Seizure Trileptal Depakote

![](_page_15_Picture_9.jpeg)

### Autism Spectrum Disorders Available Treatments

Alternative

Homeopathic Naturopathic Chelation Cranial-sacral NAET MMS Helminth HBOT Stem cell therapies

Biomedical Gut health Nutritional optimization Immune regulation

Traditional OT, PT, S&L ABA

![](_page_16_Picture_5.jpeg)

Treatment – depends on underlying conditions(s), age and severity of:

#### **Behavioral difficulties** –

Especially aggression, self-injurious, destructive

#### Speech apraxia –

No conventional treatment exists

![](_page_17_Picture_6.jpeg)

#### B-endorphin disregulation in autistic and self-injurious behavior: A neurodevelopmental hypothesis

Curt A. Sandman Ph.D.\*

Article first published online: 12 OCT 2004

DOI: 10.1002/syn.890020304

Copyright © 1988 Alan R. Liss, Inc.

![](_page_18_Picture_6.jpeg)

Synapse Volume 2, Issue 3, pages 193–199, 1988

![](_page_18_Picture_8.jpeg)

- B-endorphin disregulation in autistic and self-injurious behavior: A neurodevelopmental hypothesis. Curt A. Sandman Ph.D.\*, et. al. 1988
- Certain peptides influence neurodevelopmental processes.
- Earlier studies indicated that some of those compounds improved behavioral efficiency in retarded individuals.
- Recent studies have shown that opiate blockers reduce treatment-resistant selfinjurious behaviors.
- Prenatal protein dysregulation, addiction to endogenous opiates and elevated pain threshold have been proposed to account for this behavior.

![](_page_19_Picture_6.jpeg)

- B-endorphin disregulation in autistic and self-injurious behavior: A neurodevelopmental hypothesis. Curt A. Sandman Ph.D.\*, et. al. 1988
- In study one, four SIB patients were given 0, 25, 50 or 100 mg of naltrexone on separate weeks in a double blind, Latin square design. A specific dose dependent reduction in SIB was observed in three patients.
- In study two, plasma b-endorphin was measured in 40 patients with SIB, a related behavior, stereotypy (ST) or controls. SIB and ST patients had higher levels of endorphin than controls.
- These data added new support for the role of b-endorphin in a treatmentresistant patient group.

![](_page_20_Picture_5.jpeg)

![](_page_21_Picture_1.jpeg)

Journal of the American Academy of Child & Adolescent Psychiatry

Volume 28, Issue 2, March 1989, Pages 200-206

![](_page_21_Picture_4.jpeg)

Articles

#### Naltrexone in Autistic Children: An Acute Open Dose Range Tolerance Trial

MAGDA CAMPBELL, M.D., JOHN E. OVERALL, Ph.D., ARTHUR M. SMALL, M.D., MAE S. SOKOL, M.D., ELIZABETH KAY SPENCER, M.D., PHILLIP ADAMS, Ph.D., RODGER L. FOLTZ, Ph.D., KIM M. MONTI, B.S., RICHARD PERRY, M.D., MITCHELL NOBLER, M.D., EUGENE ROBERTS, Ph.D.

![](_page_21_Picture_8.jpeg)

- Naltrexone in Autistic Children: An Acute Open Dose Range Tolerance Trial. Magda Campbell, M.D., et. al. 1989
- The safety and efficacy of naltrexone was explored in an open acute dose range tolerance trial in 10 hospitalized autistic children, ages 3.42 to 6.50 years (mean, 5.04).
- Naltrexone was given in ascending doses: 0.5, 1.0, and 2.0 mg/kg/day.
- Behavioral side effects were observed as early as ½ hour after dosing.
- Ratings on the Children's Psychiatric Rating Scale showed that withdrawal was reduced across all three dose levels
- 0.5 mg/kg/day dose resulted in increased verbal production
- 2.0 mg/kg/day dose resulted in reduction of stereotypies.

![](_page_22_Picture_8.jpeg)

Naltrexone in Autistic Children: An Acute Open Dose Range Tolerance Trial. Magda Campbell, M.D., et. al. 1989

Mild sedation of brief duration was the only side effect.

Electrocardiogram, liver function tests, and all other laboratory studies remained unchanged throughout the study.

These preliminary findings require replication in a larger sample of patients under double-blind and placebo controlled condition.

![](_page_23_Picture_5.jpeg)

Naltrexone in autistic children: A double-blind and placebo-controlled study. Campbell, Magda; et.al. Psychopharmacology Bulletin, Vol 26(1), **1990**, 130-135.

18 autistic children, aged 3.08–7.99 yrs.

Randomly assigned to NAL or placebo and received daily doses over a period of 21 days.

NAL was superior to placebo according to blind clinical global consensus ratings.

Other behavioral rating measures, such as the Children's Psychiatric Rating Scale (W. Guy, 1976) did not confirm this result.

There was only a suggestion that NAL reduced fidgety and hyperactive behavior and tended to alleviate overall symptomatology in older children.

NAL did not appear to affect discrimination learning.

![](_page_24_Picture_8.jpeg)

# Naltrexone and other potential new pharmacological treatments of autism.

Panksepp, Jaak; Lensing, Patrick; Leboyer, Marion; Bouvard, Manuel P. Brain Dysfunction, Vol 4(6), Nov-Dec **1991**, 281-300.

![](_page_25_Picture_3.jpeg)

Naltrexone and other potential new pharmacological treatments of autism. Panksepp, Jaak; Lensing, Patrick; Leboyer, Marion; Bouvard, Manuel P. Brain Dysfunction, Vol 4(6), Nov-Dec 1991, 281-300.

Summarizes results from a series of open and doubleblind trials that have yielded positive therapeutic effects with low doses of naltrexone, including: reductions in autistic stereotypes, aggressiveness, and self-injurious behaviors, and the production of heightened prosocial emotional attitudes that are accompanied by increased smiling, eye contact, attention, and attempts to communicate.

![](_page_26_Picture_3.jpeg)

### Naltrexone and other potential new pharmacological treatments of autism.

Panksepp, Jaak; Lensing, Patrick; Leboyer, Marion; Bouvard, Manuel P. Brain Dysfunction, Vol 4(6), Nov-Dec **1991**, 281-300.

The positive behavioral change seems to be enhanced by social support, and how such features of therapeutic situations can be maximized to optimize clinical benefits... is discussed.

![](_page_27_Picture_4.jpeg)

![](_page_28_Picture_1.jpeg)

Journal of the American Academy of Child & Adolescent Psychiatry

Volume 32, Issue 6, November 1993, Pages 1283-1291

![](_page_28_Picture_4.jpeg)

Case Study

#### Naltrexone in Autistic Children: Behavioral Symptoms and Attentional Learning

MAGDA CAMPBELL, M.D. &, LOWELL T. ANDERSON, PH.D., ARTHUR M. SMALL, M.D., PHILLIP ADAMS, PH.D., NILDA M. GONZALEZ, M.D., MONTIQUE ERNST, M.D., PH.D.

![](_page_28_Picture_8.jpeg)

Naltrexone in Autistic Children: Behavioral Symptoms and Attentional Learning. Magda Campbell MD, et. al. 1993

#### Objective

To assess critically the short-term efficacy and safety of naltrexone in autistic children and its effects on discrimination learning in the laboratory.

#### Method

Forty-one children, all inpatients, ages 2.9 to 7.8 years, completed the study. Naltrexone reduced hyperactivity and had no effect on discrimination learning in the laboratory.

There was a suggestion that it had a beneficial effect on decreasing selfinjurious behavior.

Untoward effects were mild and transient.

![](_page_29_Picture_8.jpeg)

Naltrexone in Autistic Children: Behavioral Symptoms and Attentional Learning. Magda Campbell MD, et. al. 1993

#### Conclusion

In the present study, naltrexone significantly reduced only hyperactivity, and no serious untoward effects were observed.

The effectiveness of naltrexone in the treatment of autism and self-injurious behavior requires additional assessment in a sample of children with moderate to severe self-injurious behavior.

![](_page_30_Picture_5.jpeg)

![](_page_31_Picture_1.jpeg)

Journal of the American Academy of Child & Adolescent Psychiatry

Volume 34, Issue 2, February 1995, Pages 223-231

![](_page_31_Picture_4.jpeg)

#### ARTICLE

#### Naltrexone in Young Autistic Children: A Double-Blind, Placebo-Controlled Crossover Study

Barbara K. Kolmen, M.D. Å, Heidi M. Feldman, M.D. Ph.D., Benjamin L. Handen, Ph.D., Janine E. Janosky, Ph.D.

From the Child Development Unit, Children's Hospital of Pittsburgh, Departments of Pediatrics (Drs. Kolmen, Feldman, and Handen), Psychiatry (Dr. Handen), and Clinical Epidemiology (Dr. Janosky), University of Pittsburgh School of Medicine.

![](_page_31_Picture_9.jpeg)

Naltrexone in Young Autistic Children: A Double-Blind, Placebo-Controlled Crossover Study Barbara K. Kolmen, M.D., et. al. 1995

#### Objective

This study evaluated the efficacy and safety of naltrexone an opiate blocker, in the treatment of autism.

#### Method

Thirteen children with autistic disorder, aged 3.4 to 8.3 years (mean 5.4)

Naltrexone, 1.0 mg/kg, was given daily in a randomized, double-blind, placebocontrolled crossover design.

Dependent measures included parent and teacher Clinical Global Impressions (CGI), Conners Rating Scales, and Naltrexone Side-Effects (SE) Rating Scale; laboratory CGI, movement actometer readings, and a 10-second interval recording system analysis of on-task, communication initiations, disruptive behavior, and self-stimulation.

Naltrexone in Young Autistic Children: A Double-Blind, Placebo-Controlled Crossover Study Barbara K. Kolmen, M.D., et. al. 1995

#### Results

Eight of 13 subjects improved in two or more settings.

Changes in parent measures... and Teacher CGI achieved statistical significance.

Teacher SE-Restlessness and initiation of communication in the clinic showed a trend toward improvement.

Actometer readings improved in two children who were very active at baseline. Adverse side effects were behavioral, mild, and transient.

Administering the bitter tablet was a challenge.

#### Conclusion

Naltrexone offers promise as an agent for modest improvement of behavior and social communication in young children with autism.

Parent and teacher measures can be useful in outpatient trials to evaluate change.

![](_page_33_Picture_11.jpeg)

![](_page_34_Picture_1.jpeg)

#### **Psychiatry Research**

Volume 58, Issue 3, 16 October 1995, Pages 191-201

![](_page_34_Picture_4.jpeg)

Article

#### Low-dose naltrexone effects on plasma chemistries and clinical symptoms in autism: a double-blind, placebo-controlled study

Manuel P. Bouvard<sup>a</sup>, Marion Leboyer<sup>b</sup>, Jean-Marie Launay<sup>c</sup>, Christophe Recasens<sup>a</sup>, Marie-Hélène Plumet<sup>a</sup>, Delphine Waller-Perotte<sup>a</sup>, François Tabuteau<sup>c</sup>, Dominique Bondoux<sup>c</sup>, Michel Dugas<sup>a</sup>, Patrick Lensing<sup>d</sup>, Jaak Panksepp <sup>1</sup>/<sub>e</sub>, <sup>w</sup>

![](_page_34_Picture_8.jpeg)

Low-dose naltrexone effects on plasma chemistries and clinical symptoms in autism: a double-blind, placebo-controlled study. Manuel P. Bouvard<sup>a</sup>, et. al. 1995

#### Abstract

Month-long naltrexone (NTX) treatment at a daily oral dose of 0.5 mg/kg/day vs. Placebo

Modest clinical benefits were achieved with both... marginally better overall results NTX.

Degree of improvement appeared to be related to plasma chemical profiles.

Massively elevated levels of  $\beta$ -endorphin were observed in all children.

70% of the children exhibited abnormally low levels of adrenocorticotropic hormone, elevated norepinephrine (60%) arginine-vasopressin (50%) and serotonin (20%).

![](_page_35_Picture_8.jpeg)

Low-dose naltrexone effects on plasma chemistries and clinical symptoms in autism: a double-blind, placebo-controlled study. Manuel P. Bouvard<sup>a</sup>, et. al. 1995

The best clinical responders exhibited the clearest normalization of the elevated plasma chemistries, especially in C-terminal- $\beta$ -endorphin and serotonin.

The results suggest that NTX only benefits a subgroup of autistic children, who may be identified by the presence of certain plasma abnormalities.

![](_page_36_Picture_4.jpeg)

![](_page_37_Picture_1.jpeg)

#### **Psychiatry Research**

Volume 58, Issue 3, 16 October 1995, Pages 203-215

![](_page_37_Picture_4.jpeg)

Article

#### Placebo-controlled acute dosage naltrexone study in young autistic children

Sophie H.N. Willemsen-Swinkels, Jan K. Buitelaar 📥 , Florence G. Weijnen, Herman van Engeland

![](_page_37_Picture_8.jpeg)

Placebo-controlled acute dosage naltrexone study in young autistic children Sophie H.N., et. al. 1995

Double-blind, placebo-controlled, crossover trial 23 autistic children were treated with a single 40-mg dose naltrexone Naltrexone treatment failed to produce significant changes in social behavior, reduced irritability and target scores on behavior checklists.

The playroom data indicated that naltrexone significantly affected indices of activity and attention.

![](_page_38_Picture_4.jpeg)

Ann. Ist. Super. Sanità, vol. 32, n. 3 (1996), pp. 351-359

#### Opioid-immune interactions in autism: behavioural and immunological assessment during a double-blind treatment with naltrexone

Renato SCIFO (a), Matteo CIONI (b), Alfredo NICOLOSI (c), Nunzio BATTICANE (b), Cataldo TIROLO (b), Nuccio TESTA (b), Maria C. QUATTROPANI (d), Maria C. MORALE (b) Francesco GALLO (e) and Bianca MARCHETTI (e)

![](_page_39_Picture_4.jpeg)

- Opioid-immune interactions in autism: behavioural and immunological assessment during a double-blind treatment with naltrexone. Scifo R1, et. al. 1996
- 12 patients, 7 to 15 years
- Double-blind crossover study with NAL at the doses of 0.5, 1.0 and 1.5 mg/kg q48 hours.
- Responders 7/12 displayed "... a significant reduction of the autistic symptomatology."
- The behavioural improvement was accompanied by alterations in the distribution of the major lymphocyte subsets, with a significant increase of the T-helper-inducers (CD4+CD8-) and a significant reduction of the T-cytotoxic-suppressor (CD4-CD8+) resulting in a normalization of the CD4/CD8 ratio.
- Changes in natural killer cells and activity were inversely related to plasma betaendorphin levels.

![](_page_40_Picture_7.jpeg)

Opioid-immune interactions in autism: behavioural and immunological assessment during a double-blind treatment with naltrexone. Scifo R1, et. al. 1996

It is suggested that the mechanisms underlying opioid-immune interactions are altered in this population of autistic children and that an immunological screening may have prognostic value for the pharmacological therapy with opiate antagonists.

![](_page_41_Picture_3.jpeg)

![](_page_42_Picture_1.jpeg)

**Biological Psychiatry** 

Volume 39, Issue 12, 15 June 1996, Pages 1023-1031

![](_page_42_Picture_4.jpeg)

#### The effects of chronic naltrexone treatment in young autistic children: A double-blind placebo-controlled crossover study

Sophie H.N. Willemsen-Swinkels, Jan K. Buitelaar 📥 , Herman van Engeland

![](_page_42_Picture_7.jpeg)

The effects of chronic naltrexone treatment in young autistic children: A doubleblind placebo-controlled crossover study. Sophie H.N., et. al. 1996

23 autistic children, aged 3–7 years Mean daily dosage of 1 mg/kg naltrexone for 4 weeks.

On average, parents' checklists and playroom data could not differentiate between naltrexone treatment and placebo treatment; however, teachers significantly favored naltrexone treatment.

They reported a decrease in hyperactivity and irritability. No effects of naltrexone on social and stereotypic behavior could be demonstrated.

![](_page_43_Picture_5.jpeg)

![](_page_44_Picture_1.jpeg)

Journal of the American Academy of Child & Adolescent Psychiatry

Volume 36, Issue 11, November 1997, Pages 1570-1578

![](_page_44_Picture_4.jpeg)

#### ARTICLES

#### Naltrexone in Young Autistic Children: Replication Study and Learning Measures

Barbara K. Kolmen, M.D. 📥 , Heidi M. Feldman, M.D., Ph.D., Benjamin L. Handen, Ph.D., Janine E. Janosky, Ph.D.

![](_page_44_Picture_9.jpeg)

Naltrexone in Young Autistic Children: Replication Study and Learning Measures Barbara K. Kolmen, M.D, et. al. 1997

This study expanded upon previous work on naltrexone efficacy and safety in young autistic children and assessed performance on learning measures.

#### Method

Eleven children with autistic disorder, aged 3.0 to 8.3 years Naltrexone, 1.0 mg/kg, was given daily in a randomized, double-blind, crossover design.

#### Results

...The combined study sample showed improvement on all parent measures and on Teacher CGI and SE-Restlessness compared with baseline and placebo. Eleven of the 24 children improved in two or more settings. Scores on learning measures did not change across conditions.

#### Conclusions

Naltrexone was associated with modest improvement of behavior in 11 of 24 children but learning did not improve.

![](_page_46_Picture_1.jpeg)

Journal of the American Academy of Child & Adolescent Psychiatry

Volume 38, Issue 5, May 1999, Pages 587-593

![](_page_46_Picture_4.jpeg)

Special Section

#### Naltrexone and Communication Skills in Young Children With Autism

HEIDI M. FELDMAN, M.D., PH.D. 📥 , BARBARA K. KOLMEN, M.D., ALDA MARIA GONZAGA, B.A.

![](_page_46_Picture_8.jpeg)

**The effects of chronic naltrexone treatment in young autistic children**: A doubleblind Naltrexone and Communication Skills in Young Children With Autism H.M. Feldman, M.D., et. al. **1999** 

#### **Objectives**

To evaluate the effect of naltrexone on communication skills of young children with autism.

#### Method

Twenty-four children with autism, 3.0 to 8.3 years old (mean 5.1) - randomized, double-blind, placebo-controlled, crossover trial.

Naltrexone, 1.0 mg/kg, or placebo was administered daily for 2 weeks. Communication was evaluated from videotaped samples of seminaturalistic parent-child interaction.

![](_page_47_Picture_7.jpeg)

The effects of chronic naltrexone treatment in young autistic children: A double-blind Naltrexone and Communication Skills in Young Children With Autism H.M. Feldman, M.D., et. al. **1999** 

#### Results

No differences were found between the naltrexone and placebo conditions in any of the measures of children or parents' communication.

Significant correlations were found between the child's number of words and developmental quotient (Spearman p = 0.58, p = .003) and between the child's and parent's number of words (p = 0.55, p = .005).

#### Conclusions

In this short-term study, the medication did not lead to improvement in communication, a core deficit of autism.

![](_page_48_Picture_7.jpeg)

#### Self-injurious behavior and the efficacy of naltrexone treatment: A quantitative synthesis

Frank J. Symons<sup>†,\*</sup>, Andrea Thompson and Michael C. Rodriguez

Article first published online: 20 DEC 2004 DOI: 10.1002/mrdd.20031

Copyright @ 2004 Wiley-Liss, Inc.

![](_page_49_Figure_5.jpeg)

#### Mental Retardation and Developmental Disabilities Research Reviews

Special Issue: Treatment Efficacy Volume 10, Issue 3, pages 193–200, August 2004

![](_page_49_Picture_8.jpeg)

Self-injurious behavior and the efficacy of naltrexone treatment: A quantitative synthesis. Frank J. Symons<sup>†,\*</sup>, et. al. 2004

People with mental retardation, autism, and related developmental disabilities who self-injure are treated with a wide array of behavioral techniques and psychotropic medications.

Despite numerous reports documenting short-term and some long-term changes in self-injury associated with the opiate antagonist naltrexone hydrochloride, no quantitative review of its efficacy has been reported.

A quantitative synthesis of the peer-reviewed published literature from 1983 to 2003 documenting the use of naltrexone for the treatment of self-injurious behavior (SIB). Individual-level results were analyzed given subject and study characteristics.

![](_page_50_Picture_5.jpeg)

- Self-injurious behavior and the efficacy of naltrexone treatment: A quantitative synthesis. Frank J. Symons<sup>†,\*</sup>, et. al. 2004
- A sample of 27 research articles involving 86 subjects with self-injury was reviewed.
- Eighty percent of subjects were reported to improve relative to baseline (i.e., SIB reduced) during naltrexone administration and 47% of subjects SIB was reduced by 50% or greater.
- Males were more likely than females to respond.
- No significant relations were found between treatment outcomes and autism status or form of self-injury.
- Results are discussed with respect to future efficacy work related to study outcomes and the pharmacological treatment of self-injury.

![](_page_51_Picture_7.jpeg)

Efficacy and Safety of Naltrexone Use in Pediatric Patients with Autistic Disorder. G M ElChaar, et. al. Ann Pharmacother 2006

**OBJECTIVE:** To review the efficacy and safety of naltrexone in pediatric patients with autistic disorder (AD).

DATA SOURCES: Literature search

Three case reports, 8 case series, and 14 clinical studies were identified as pertinent.

#### DATA:

Naltrexone has been used most commonly at doses ranging from 0.5 to 2 mg/kg/day Found to be predominantly effective in decreasing self-injurious behavior.

![](_page_52_Picture_7.jpeg)

Efficacy and Safety of Naltrexone Use in Pediatric Patients with Autistic Disorder. G M ElChaar, et. al. Ann Pharmacother 2006

#### DATA:

Naltrexone may also attenuate hyperactivity, agitation, irritability, temper tantrums, social withdrawal, and stereotyped behaviors. Patients may also exhibit improved attention and eye contact. Transient sedation was the most commonly reported adverse event.

#### **CONCLUSIONS:**

A child affected by AD may benefit from a trial of naltrexone therapy, particularly if the child exhibits self-injurious behavior and other attempted therapies have failed. Serious adverse effects have not been reported in short-term studies.

![](_page_53_Picture_6.jpeg)

#### Low-Dose Naltrexone (LDN) for Mood Regulation and Immunomodulation in ASD

#### Jaquelyn McCandless, M.D., April 2006

I completed a preliminary eight-week informal study on 15 of my autism patients May-June 2005 applying 3mg of LDN transdermally between 9 and 12 p.m.

Eight of the 15 children in this study had positive responses, with five of these eight having results considered quite phenomenal according to their parents.

The primary positive responses are in the area of mood regulation, cognition, language, and socialization.

Two small children responded better when changed to 1-1/2mg dosing.

#### Low-Dose Naltrexone (LDN) for Mood Regulation and Immunomodulation in ASD

Jaquelyn McCandless, M.D., April 2006

No allergic reactions were noted, and the primary negative side effect was insomnia and earlier awakening, usually fairly short-lived.

As an effective, non-toxic, non-addicting, and inexpensive behavioral and immunomodulating intervention, LDN is joining our biomedical arsenal to help more and more children recover from autism as well as helping anyone with autoimmune diseases and cancer.

As an FDA approved medication, it must be prescribed and must also be compounded for the tiny dosing required.

#### Child Development Center of America, PA

During 2014, 53 patients chosen out of 393 total visits Received 3mg/1cc as cream after 9PM

<b>Response to LDN Treatment</b>	Number of Patients (out of 53)
"Significant Improvement"	5
"Satisfactory Results"	19
"No Improvement/Unsatisfactory Results"	16
"No Follow-Up Information Received"	4
"LDN Prescribed but Usage Not Initiated"	9
T-11-10 1 Descentes to LDN Treatment has ACD Detionst	

Table 10.1 Responses to LDN Treatment by ASD Patients.

![](_page_56_Picture_5.jpeg)

#### **Jacob's Story**

![](_page_57_Picture_2.jpeg)

- Success depends on:
- Accurate diagnosis
- Choose candidates most likely to succeed
- Address gut and metabolic issues first
- Address other co-morbidities as they exist or arise

![](_page_58_Picture_6.jpeg)

#### Form

Liquid?

Pill?

Cream?

#### **Expectations**

First few days Few weeks Long term

#### Side effects

5% stop 25% Renew 70% Not enough/ no effect

![](_page_59_Picture_9.jpeg)

#### **Treatment Works**

Journal of Autism and Developmental Disorders December 2014, Volume 44, Issue 12, pp 2981-2995

Date: 12 Sep 2014

#### Autism Treatment in the First Year of Life: A Pilot Study of Infant Start, a Parent-Implemented Intervention for Symptomatic Infants

S. J. Rogers, L. Vismara, A. L. Wagner, C. McCormick, G. Young, S. Ozonoff

![](_page_60_Picture_5.jpeg)

### **Treatment Works**

12-week, low-intensity treatment with seven symptomatic infants ages 7–15 months

Parents mastered the intervention and maintained skills after treatment ended

Four comparison groups were matched from a study of infant siblings

The treated group of infants was significantly more symptomatic than most of the comparison groups at 9 months of age but was significantly less symptomatic than the two most affected groups between 18 and 36 months.

At 36 months, the treated group had much lower rates of both ASD and DQs under 70 than a similarly symptomatic group who did not enroll in the treatment study.

![](_page_61_Picture_6.jpeg)

#### Conclusions

![](_page_62_Picture_1.jpeg)

#### **LDN – Autism - 2016**

![](_page_62_Picture_3.jpeg)

Complex, multi-system condition leading to behavioral and developmental signs and symptoms.

**Epidemic proportions** 

Treatable condition – combination of therapies and medical intervention at earliest stages provides best chance for recovery

![](_page_63_Picture_4.jpeg)

### **LDN Conclusions -2016**

Continuum of brain, gut, inflammation protocols

Anecdotal evidence - Small sample size, and inconsistent evaluative methods

Few therapeutic breakthroughs Learning more about genetics and susceptibility Learning more about environmental effects

Given risks, in highest responder population, Naltrexone worth a try

Continuing problems Education Older patients

![](_page_64_Picture_6.jpeg)

![](_page_65_Picture_0.jpeg)

![](_page_65_Picture_1.jpeg)

#### TheAutismDoctor.com Brian D. Udell MD

#### www.theautismdoctor.com

![](_page_65_Picture_4.jpeg)