The Society for Clinical Child and Adolescent Psychology (SCCAP): Initiative for Dissemination of Evidence-based Treatments for Childhood and Adolescent Mental Health Problems

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Center for Children and Families

Workshop Psychotherapy for Children with Bipolar Disorder

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Center for Children and Families

Just Because I'm Bipolar 14 Year Old 8th Grader Former MF-PEP Study Participant Just because I'm bipolar I'm not a freak I'm not weird I just want to be noticed Just because I'm bipolar I still have feelings I still have emotions I just have trouble expressing them Just because I'm bipolar I can still be trusted I can still be reliable Just because I'm bipolar—I'm still a normal kid

Conflict of Interest/Funding Dr. Fristad receives royalties from CFPSI: MF-PEP and IF-PEP Workbooks Guilford Press: Raising a Moody Child: How to Cope with Depression and Bipolar Disorder Psychotherapy for Children with Bipolar and Depressive Disorders ■ APPI: Clinical Manual for Management of Bipolar Disorder in Children and Adolescents Children's Interview for Psychiatric Syndromes (ChIPS)

Workshop Goals Participants will learn: Beside the How BPSD presents in children Biological interventions Psychosocial treatment for BPSD Multi-family psychoeducational psychotherapy (MF-PEP) Individual-family psychoeducational psychotherapy (IF-PEP)

Part 1 of 6

Childhood Bipolar Disorder—On the Rise? Lofthouse & Fristad, 2004, Clinical Child & Family Psychology Review

Literature review—174 articles/chapters **26 before 1980 36** during the 1980s 66 during the 1990s **46 from 2000-2002** Amazon search—18 books **15** from 2000 to 2003 ■ Websites—5 since 1999 Time—cover article, Aug 19, 2002

2005 Google Internet Search Leffler & Fristad (2006)

Topic	Number
childhood mood disorders	517,000
adolescent mood disorders	577,000
childhood depression	3,100,000
adolescent depression	3,630,000
childhood bipolar disorder	483,000
adolescent bipolar disorder	757,000
childhood mania	248,000
adolescent mania	645,000

National Trends in the Outpatient Diagnosis and Treatment of Bipolar Disorder in Youth *Moreno, et al, Arch Gen Psychiatry.* 2007; 64:1032-1039

- Compared outpt visits in 1994-1995 and 2002-2003 for individuals aged 0 to 19 years vs those aged 20 years or older diagnosed with bipolar disorder
- Demographic, clinical, and treatment characteristics were compared from '99-'03
- Patient visits (n=962) were tallied from the Nat'l Ambulatory Medical Care Survey
- Estimated annual number of office-based visits with a diagnosis of bipolar disorder
 - Youth: 0.0025% (94-95), 1% (02-03)
 Adults: 0.9% (94-95) 1.7% (02-03)

Under- or Over-diagnosis?

"It is possible that pediatric bipolar disorder, previously underdiagnosed, is now being appropriately recognized at earlier ages. The median age at onset of bipolar disorder has been located between ages 19 and 23 years, indicating that in 50% of patients, the illness starts at a younger age. Long delays in treatment seeking have been previously documented when the onset occurs in childhood or in adolescence, perhaps owing to problems with clinical recognition.

Under- or Over-diagnosis?

In recent years, there has been an increase in academic attention devoted to pediatric bipolar disorder... In addition, childhood bipolar disorder has been regularly featured in the popular press. These developments may have raised clinical and public awareness and promoted appropriate treatment seeking and clinical recognition of the condition at earlier ages"

Additional Thoughts

If 50% of patients have their onset prior to age 19, the # of patients should be roughly half the adult figures (ie, 0.45% [94-95], 8.5% [02-03]). This suggests a 180-fold underutilization of visits in 94-95

Any good pendulum swings before settling in the middle. Overdiagnosis typically follows underdiagnosis. It is critical that clinicians learn to accurately diagnose so they can settle in the middle.

What Makes Diagnosing Mood Disorders Tricky with Children?

- What's the mood disorder and what are the child's traits?
- How do I tell the ordinary ups and downs apart from the "clinical" ups and downs?
- What's a "normal" reaction to a bad event (eg, divorce) and what's not?
 If Mom/Dad has a mood disorder, "whose illness is it?"

Rule #1: Take a video, not a snapshot

Assess behaviors

- over time
- in context

TAKE YOUR TIME!

If truly BPSD, this is *probably* a lifetime diagnosis and is *not* to be considered "casually"—the stakes are FAR too high

Longitudinal review of symptoms

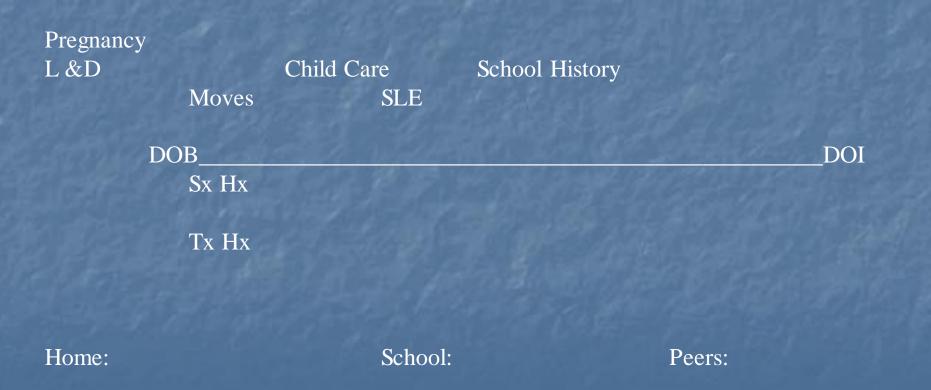
Determine symptom onset, offset, duration in relation to:

- Social history
- Treatment history
- Functioning at home, school and w/ peers

Lifeline—Document

Above line: pregnancy, labor and delivery, age in yrs, calendar yrs, moves, life stressors, child care arrangements, school placement

Below line: physical health (onset, offset) & treatment, mental health (onset, offset, mood & co-morbid diagnosis) & treatment, current functioning (home, school, peers)



How to Diagnose BPSD

Medical history

Allergies, asthma, chronic illnesses, staring spells, injuries (especially head trauma)
 Treatments for the above

7-10 day washout for prescription drugs

- 2-3 week washout for steroids, fluoxetine
- Review previous lab findings, brain imaging
- Drug screen if ANY suspicion of illicit use

Rule #2: Decide what's in and what's out

 Comorbid conditions (what's in)

 Differential diagnosis (what's out) Children's Interview for Psychiatric Syndromes Weller, Weller, Rooney & Fristad, 1999, American Psychiatric Press, Inc. (www.appi.org)

Does NOT assess

personality

- cognitive capacity (mental retardation, learning disabilities)
- tic/Tourette, autistic spectrum, panic disorders
 Does NOT replace a clinician
- DOES provide a standardized initial assessment procedure for clinical and research purposes

Brief

- Thorough coverage--20 disorders & stressors
- Psychometrics available for 6-18 year olds
- Easy to administer, score, interpret
- Storage requirements are minimal

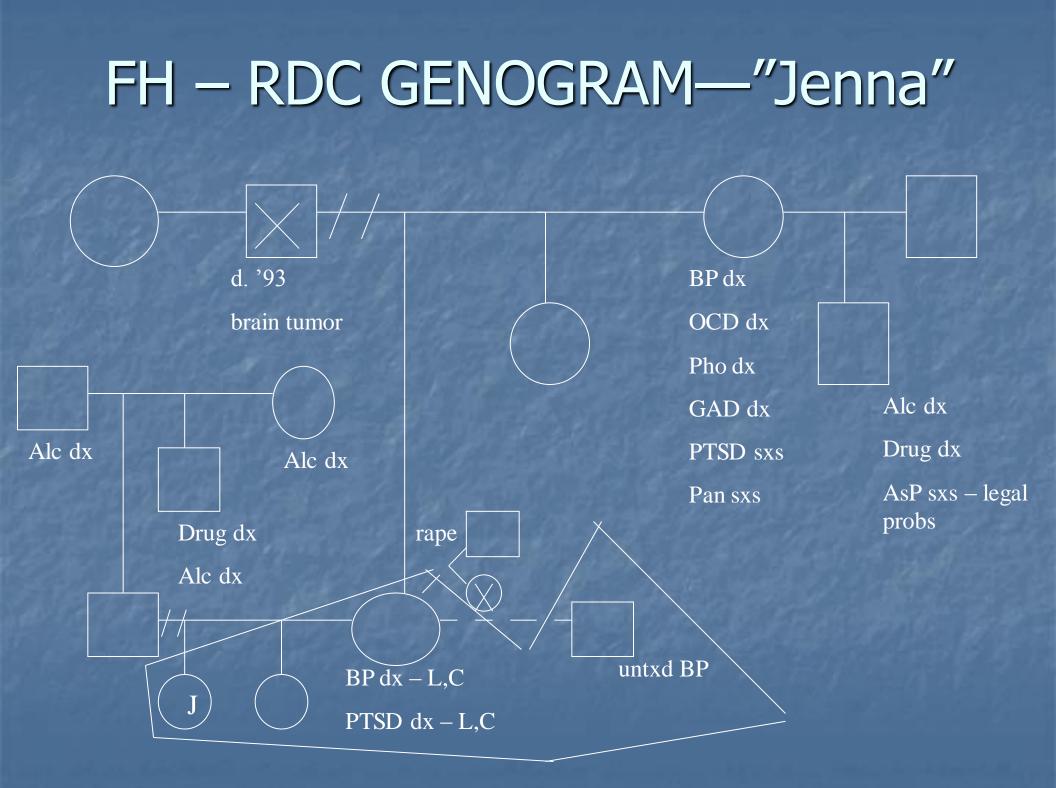
Rule #3: We diagnose children, not families BUT, families give us good clues and therapeutic material

Why Do People Get Mood Disorders?

Part of the story is genetics...

1 in 3 adopted persons *with* bipolar disorder have biological parents with mood disorders (compared to 1 in 50 adopted persons *without* bipolar disorder)
If 1 parent has a mood disorder, 27% offspring + If 2 parents have a mood disorder, 74% offspring +
If one twin has a mood disorder--

The Other Twin	Identical	Non-identical
Depression	54%	19%
Bipolar Disorder	67-79%	15-20%



Rule #4: Measure progress

Paper-and-Pencil Measures

Youngstrom EA, Findling RL, Youngstrom JK & Calabrese JR (2005). Toward an evidence-based assessment of pediatric bipolar disorder. J Clinical Child and Adolescent Psychology, 34, 433-448.

 Parent report: best source of information
 Child report: doesn't provide incremental data
 Teacher report: not validated

____feeling today?

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How is ____

How to Diagnose (Cont'd)

Obtain information from AT MINIMUM--Parent and child Parents: Ask about current, best, worst Children: Ask about current, try for best, worst Adolescents: Use your judgment 1/3 of symptoms reported by child only "classic" manic symptoms frequently reported by child only *Tillman et al, 2004* Use "tie breaker" strategy if one says YES and the other says NO (ie, observation, other parent, other source) PREFERABLY—Primary care physician, teacher, other clinicians

FIND Criteria: Treatment Guidelines for Children and Adolescents with Bipolar Disorder:
 Kowatch, Fristad, Birmaher, Wagner, Findling, Hellander & the Child Psychiatric Workgroup on Bipolar Disorder, J Am Acad Child Adol Psychiatry, 2005
 Frequency: Symptoms occur most days of

- the week
- Intensity: Symptoms cause
 - extreme disturbance in one domain
 - Moderate disturbance in two or more domains
- <u>Number</u>: Symptoms occur 3-4 or more times/day
- <u>D</u>uration: Symptoms occur > 4 hours/day (total)

Tracking Mood Changes: Depressive Spectrum Disorders

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Normal Major Depressive Disorder Dysthymic Disorder

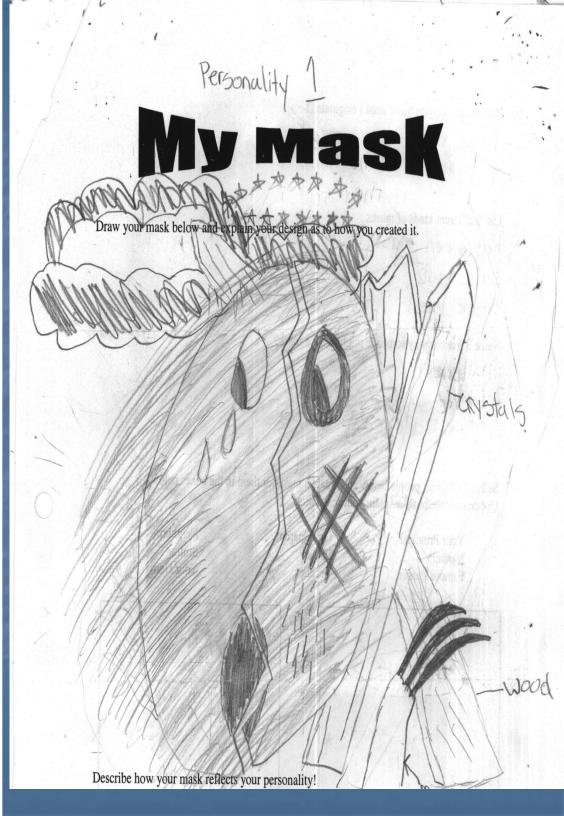
Tracking Mood Changes: Bipolar Spectrum Disorders

Bipolar I Bipolar II Cyclothymia

Tracking Mood Changes: Rapid Cycling/Mixed States

Describe how your mask reflects your personality!

The black eye represents all the mental punches I get from people. The stars circling above my head represents how I'm confused. My searching eyes represent that I'm trying to find the right path, but I'm lost so I can't find it. My broken face represents that I'm a broken person. The hair across my face shows that I can be whipped around easily. The tears are of loneliness. The crystals= dreamer/pretender. Claw and hand = outcast.



Defining Mood Disorders

- Symptoms cause distress &/or interfere with family, school, friends or work
- Symptoms are NOT because of other drugs or illness
- Symptoms do NOT directly follow the loss of a loved one
- Symptoms occur at the same time and for an extended period of time

Defining the Conditions: MDD Major Depressive Disorder

Need 1 or both of these: Impaired mood Sad/anxious Irritable/angry loss of interest Complaints of boredom Previously fun activities aren't fun anymore

Need 3-4 of these (5 total):

- impaired sleep
- impaired appetite
- poor concentration
- fatigue
- restlessness/lethargy
- worthlessness/guilt
- suicidal/morbid ideation

Symptoms last
 2 weeks

When Will My Child Get Better...? The MDD Picture Birmaher et al, 96
Single episode length: 7-9 months
90% get well by 1.5-2 years
6-10% stay impaired

Recurrence
 40%, 2 yrs
 70% 5 yrs

Defining the Conditions: Dysthymia

The "low grade fever" of mental health!

Particularly hard to diagnose

This can and should be treated

Dysthymic Disorder (DD) MOOD (lasting 1 year) sad irritable Two or more of: Impaired appetite Impaired sleep Fatigue Low self-esteem Impaired concentration/thinking Hopeless feelings

When Will My Child Get Better? The DD Picture Kovacs et al, 94
Single untreated episode: 4 years
MDD episode usually comes 2-3 years after DD onset

Can lead to:
Bipolar disorder: 13%
Substance abuse: 15%

Defining the Conditions: Seasonal Affective Disorder (SAD) Most common: fall/winter: "hibernating" depression--increased sleep and appetite, carbohydrate craving, decreased activity spring/summer: nondepressed or manic

Defining the Conditions: Psychotic Symptoms

Some children with mood disorders experience: hallucinations hearing voices seeing things sometimes-- smelling or feeling delusions special messages special powers other unusual thoughts/ideas These occur when mood symptoms are severe They go away when the mood disorder is treated This is NOT schizophrenia!

Defining the Conditions: Suicidal Risks

Time

During/right after inpatient treatment During a crisis Following suicide of a close friend/relative +/- life events Warning signs Talking about death/suicide Saying good-byes, making wills, giving away belongings

Other factors:

- depressed, hopeless
- drug/alcohol use
- impulsive/angry
- physical/sexual abuse
- runaway
- past attempt
- self-destructive
- perfectionistic
- ACCESS TO GUNS!

Who is at Risk for Bipolar Disorder? Birmaher et al, 96

About 1/4 - 1/2 of depressed children develop bipolar disorder within 2-5 yrs

Risk factors include:
 symptoms of psychomotor retardation or psychosis
 + family history-- bipolar disorder
 ++ family history--mood disorder

Medication induced hypomania

Defining the Conditions: Mania (Bipolar Disorder) MOOD (1 week—this differs for children) elevated expansive irritable 3 (4 if irritable mood) of: grandiosity decreased need for sleep increased talking racing thoughts distractible increased activity/agitation foolish/reckless behavior

Defining the Conditions: Hypomania

MOOD changes (4-7 days)
Associated symptoms (same as mania)
Functioning clearly "out of character"
Altered mood & behavior noted by others
Symptoms not severe enough to be called MANIA

Bipolar Spectrum Diagnoses

Bipolar Disorder I (BP-I): M + DBipolar Disorder II (BP-II): m + D Cyclothymia: m + dBipolar Disorder NOS (BP-NOS) Define why One symptom short? Duration insufficient? Episodes not clearly defined? Informants sketchy, need to observe before finalizing diagnosis?

Defining the Conditions: Cyclothymia

Less severe highs and lows than bipolar disorder
Causes disruption
Can be tricky to diagnose
Decide when/how to treat

Part 2 of 6

Differential Diagnosis & Comorbidities

Health Conditions that Mimic BPD Temporal lobe epilepsy Hyperthyroidism Closed or open head injury Multiple sclerosis Systemic lupus erythematosus (SLE) Alcohol related neurodevelopmental disorder Wilson's disease

Medications that May Increase Cycling Abouesh et al, 2002, J Clin Psychopharmacol ANY biological intervention for depression Tricyclic antidepressants Serotonin specific reuptake inhibitors Serotonin and norepinephrine reuptake inhibitors Light box Aminophylline Corticosteroids Sympathomimetic amines (eg, pseudoephedrine) Antibiotics (eq, clarithromycin, erythromycin, amoxicillin) Illicit drugs

BPD vs ADHD: Symptoms that Overlap Geller et al. (2002)

Symptoms	EOBD	ADHD
Irritability	98%	72%
↑ Speech	97%	81%
Distractibility	93%	96%
1 Energy	100%	95%

BPD vs ADHD: Symptoms that Differ Geller et al. (2002)

Symptom	EOBD	ADHD
Elated Mood	89%	13%
Grandiosity	86%	5%
↓ Sleep	40%	6%
Flight of ideas	71%	10%
Hypersexuality	43%	6%
Suicidality	25%	0%
Psychosis	60%	0%

Is it BPD or A/PTSD?

Symptoms of BPD and A/PTSD overlap Symptoms can also co-occur Post 2006-% of adults w/ BPD who experienced abuse/neglect in childhood 52%--childhood onset 34%--adolescent onset 21%--early adult onset (19-29 yrs) 20%--late adult onset (30+ yrs)

Posttraumatic Stress Disorder

Traumatic event occurred

- The <u>child's reaction</u> involved intense fear, helplessness, or horror that might appear as disorganized or agitated behavior, with <u>new</u> and <u>persistent</u> examples of:
- Reexperiencing the trauma-- > 1 of:
 - recurrent and intrusive distressing recollections (can be displayed as repetitive play)
 - recurrent distressing dreams (these may be frightening but
 - w/o recognizable content)
 - acting or feeling like the traumatic event is recurring (this can include a sense of reliving the experience, illusions, hallucinations, and dissociative flashbacks)
 - intense psychological distress at exposure to internal or external reminders of the trauma
 - physiological reactivity on exposure to internal or external reminders

Posttraumatic Stress Disorder (continued) <u>Avoiding</u> reminders of the trauma /emotional numbing-->3 of: avoiding thoughts, feelings, or conversations associated with the trauma avoiding activities, places, or people that remind one of the trauma forgetting an important aspect of the trauma Iosing interest in activities feeling disconnected from others restricted range of affect (e.g., unable to have loving feelings) expecting a shortened life (e.g., does not expect to have a career, marriage, children, or a normal life span) *Increased arousal*- > 2 of: difficulty falling or staying asleep irritability or outbursts of anger difficulty concentrating/hypervigilance exaggerated startle response

Acute Stress Disorder

- Traumatic event occurred
- Child experienced an *intense emotional response*
- Reexperiencing, avoidance, and arousal ~PTSD
- During or after the trauma, <u>dissociative symptoms</u> occur-- > 3 of:
 - feeling numb, detached, or devoid of emotions
 Feeling "in a daze" (unaware of surroundings
 - Derealization
 - Depersonalization
 - dissociative amnesia (i.e., can't recall an important aspect of the trauma)
- Lasts > 2 days and < 4 wks, occurs within month of event</p>

Is it Mania or Something Else?

Euphoria

Normal: Special occasions, transitory

- Drug-induced disinhibition: Steroids, illicit drugs
- Carefully examine contextual cues to determine +/-

Irritability

- Ubiquitous: MDD, DD, ODD, PDD, Anxiety disorders, ADHD, schizophrenia
- Medication side-effects: stimulant wear off, SSRI adverse event
- Normal: hungry, hot, tired children

Is it Mania or Something Else? Grandiosity: *True talent:* check it out Peers unavailable: fantasy play may persist can the child distinguish fantasy from reality? Normal: understand the child's age developmental context persistence effects on behavior (eg, playing Superman vs) jumping out of the window because you are Superman)

Is it Mania or Something Else? Decreased Need for Sleep NOT the same as decreased sleep! Ruminations of a depressed, anxious child Poor sleep hygiene Excessive environmental stimuli Excitatory medications Results in fatigue the next day Full of energy Common time to get in trouble (eg, sexual content on TV)

Is it Mania or Something Else? Pressured Speech

Affective arousal: excited, nervous or angry children may speak quickly
 ADHD: often chronic "motor mouths"

Racing Thoughts

Young/Low IQ/Language Disorder: Get an "interpreter"!

Is it Mania or Something Else?

Distractibility

ADHD: Establish a baseline. There needs to be a *change* from baseline for children with ADHD to count as a symptom of mania (not just when medications are wearing off) *Depression:* impaired concentration common *Anxiety:* preoccupation common *Learning disabilities:* distracted while doing schoolwork

Is it Mania or Something Else? Increased Goal Directed Activity Psychomotor agitation: common and nonspecific • *Gifted children:* may be highly productive work tends to be focused and accomplishments accrue Depressed/anxious/traumatized children: may be agitated or demonstrate "nervous habits"

Is it Mania or Something Else?

Excessive Involvement in Pleasurable/Risky Activities

Sexual abuse: sexual acting out often anxious/compulsive in nature

Hypersexuality: often has erotic, pleasureseeking quality, excessive, violates social norms (OK in private between consenting adults, NOT OK in public with a child initiating unwanted behavior toward adult)

Is it Mania or Something Else?

Psychosis

Perceptual distortions: falling asleep (hypnogogic)/waking up (hypnopompic)—see or hear things

Suicidality
 Not pathognomonic
 Critical to assess

Co-occurring Disorders

Behavior Disorders

 attention deficit hyperactivity disorder (ADHD), oppositional-defiant, conduct, tic/Tourette, substance use/abuse

Anxiety Disorders

separation anxiety, generalized anxiety, phobias, acute/post-traumatic stress, obsessive-compulsive, social phobia, panic disorder

Eating Disorders

anorexia, bulimia, obesity

Learning Disorders

- reading, writing, math, language
- Developmental Disorders
 - Autism, Asperger's, PDD-NOS

Treating Co-occurring Disorders Some co-occurring disorders will be successfully treated IN PART by mood disorder treatments many anxiety disorders some behavior disorders some eating disorders Most will require some other specialized intervention other medications/psychotherapy school intervention learning to cope with symptoms of those disorders

Workshop Goals Participants will learn: How BPSD presents in children Biological interventions Psychosocial treatment for BPSD Multi-family psychoeducational psychotherapy (MF-PEP) Individual-family psychoeducational psychotherapy (IF-PEP)

Types of Treatment Biological Social Medications school-based Lights interventions **ECT** Home-based interventions Nutritional Psychological Respite Out-of-home Individual Therapy placement Family Therapy Parent Guidance Group Therapy

Medication Issues--#1 Medications do not cure a mood disorder, but they help *manage* it They can STOP or LESSEN current symptom severity They can PREVENT or DECREASE impairment from future episodes Medications DO NOT: solve every problem For medications to do their best, it is ESSENTIAL that families are an *active partner* in treatment!

Medication Issues--#2 Partial responses are common Don't despair! May need to change Dose Time medicine is taken Type of medicine Full response Is NOT necessarily a reason to discontinue medication May be useful to prevent future episodes Plan with your doctor when/how to stop medicine

If/when you stop medicine
Do so *with input from your doctor*Stopping abruptly may lead to unpleasant side effects

Mood stabilizers are NOT addicting!

Ask for fact sheets about medications Know common & serious side-effects (side) effects often decrease as the body adjusts to new medicine) Know what to do if a serious side-effect occurs Know how to monitor the medication ALWAYS tell both the primary care doctor and psychiatrist about ALL medications *(including*) herbal remedies) being taken

Timetable

Many medicines take several weeks to work BE PATIENT, DON'T GIVE UP TOO SOON! "Physical" symptoms may improve before mood and thinking does Make a plan (& a backup plan!) to remember to take medicine eg, pill holder, morning ritual... Know what to do about missed doses (check w/ your doctor)

Plan ahead to get refills on time

Why Blood Levels?

- Some medicines need to be monitored by blood tests
- Same dose can produce different levels in different people
- Need this to stay safe
- How to do?
 - Get levels
 - 12 HRS AFTER THE LAST DOSEBEFORE THE NEXT DOSE
 - Postpone the morning dose until AFTER the blood draw

Medication Update Kowatch & Post, Bipolar Network News '08, 12(2): 1-4 Much progress in the past 3 years 11 RCTs of substantial size Unequivocal evidence for atypicals Topirimate (Topomax): 4 – RCTS in adults; + on some secondary outcomes in youth; wt loss

Mood Stabilizers

Kowatch & Post, Bipolar Network News '08, 12(2): 1-4

Drug	Evidence	Problems	Tolerability	Utility
Valproate (Depakote)	(++)	Weight gain, polycystic ovarian syndrome	B+	B+/A-
Lithium	++ FDA	Weight, GI, thyroid, acne, requires blood levels	B	B+
Carbama- zepine	(++) (Tegretol)	Rash, rare bone marrow suppression	C+	В
Oxcarba- zepine	(+/-) (Trileptal)	Hyponatremia, sedation	B	B-
Lamotrigine (Lamictal)	(0)	1:2500, severe rash	В	D

() Ambiguous data; ++ substantial evidence/multiple series; +/- minimal evidence; 0 no effect; FDA=approved or approval pending

Atypical Antipsychotics Kowatch & Post, Bipolar Network News '08, 12(2): 1-4

Drug	Evidence	Problems	Tolerability	Utility
Aripiprazole (Abilify)	+++ FDA	Akathisia, minimal wt gain, GI upset	A-	А
Clozapine (Clozaril)	+++	Seizures, drooling, wt gain, requires white blood cell monitoring	С	С
Olanzapine (Zyprexa)	+++ FDA	Weight gain +++, sedation	B-	B-
Quetiapine (Seroquel)	+++ FDA	Weight gain ++, sedation	В	А-
Risperidone (Risperal)	+++ FDA	Prolactin increases, weight gain ++	В	В
Ziprasidone (Geodon)	+++ FDA	Minimal wt gain, increases QTc (not clinically problematic)	A-	A

+++ strong evidence, placebo controlled RCT; FDA=approved or approval pending

Medication Algorithm Kowatch & Post, Bipolar Network News '08, 12(2): 1-4		
STEP	MEDICATION	
1	AA or MS If poor tolerability, switch within drug class	
2	Add other drug class (AA or MS)	
3	Add lithium	
4	Add combination within class	
5	Add an AA	
6	Switch the AA	
7	Switch MS to carbamazepine	
AA=atypical antipsychotic: MS=mood stabilizer Add low dose stimulant		

AA=atypical antipsychotic; MS=mood stabilizer. Add low dose stimulant (amphetamine/methylphenidate) for residual ADHD; if +, continue; lf -, switch type of stimulant

Treating Comorbid Disorders Kowatch et al, 2005, JAACAP

Guidelines primarily anecdotal Stabilize the mania FIRST Review what problems remain Treat sequentially Use psychosocial treatments when Treatments are evidence-based Families are willing Trained therapists are available

Treating Comorbid ADHD Kowatch et al, 2005, JAACAP

Occurs in 70-90% of children and 30-40% of adolescents with BPD Medications—first-line Low and slow Psychosocial parent training school consultation

Treating Comorbid ODD-CD Kowatch et al, 2005, JAACAP

Medications: those for mood may help for behavior disorders

Psychosocial: parent training

Environmental: may need residential placements, foster care, etc.

Treating Comorbid Anxiety Kowatch et al, 2005, JAACAP **Psychosocial: USE FIRST!** - CBT Medications: SSRIs: be cautious Buspirone (Buspar): not effective Benzodiazepines: limited data, \uparrow abuse potential, cognitive side-effects; might want to use short-term until other meds

begin to work

Treating Comorbid Substance Abuse Kowatch et al, 2005, JAACAP

TREAT BOTH IMMEDIATELY Wilens et al, 1999, JAACAP Medications Lithium Geller et al, 1998, JAACAP Psychotherapy Family therapy Latimer et al, 2003, Drug & Alcohol Dependence; Liddle & Dakof, 1995, NIDA Research Monograph

Treating Other Comorbid Conditions Kowatch et al, 2005, JAACAP Pervasive developmental disorder PDD program Mental retardation Ditto Seizures and/or migraines Use dual-purpose medications (eg, DVP, CBZ, OXC) Premenstrual dysphoric disorder SSRIs AFTER stabilization of mood

Maintenance Treatment Kowatch et al, 2005, JAACAP Agents that get you well keep you well Discontinuation leads to relapse 37.5% vs 92.3% relapse in those who stayed on vs went off meds, Strober et al, 1990 Work toward monotherapy Try reductions in summer/over breaks Environment should be stable Go slow Stay on maintenance dose 18 months poststabilization Use cost-benefit analyses to decide

Managing Side-Effects

Dizzy: Stand up slowly Dry mouth: Drink water Use sugarless gum/candy **Constipation:** Eat high fiber diet Drink 6-8 glasses of water/day Persistent nausea: Take medicine with meals or in divided doses

Managing Side-Effects

Increased thirst/urination:

Drink 6-8 glasses of liquid/day

- Avoid high calorie beverages!
- Make school plan to use the bathroom more frequently

Tremor:

Take with meals or in divided doses

Avoid caffeine

Excessive weight gain:

- Balanced diet & regular exercise
- Avoid drastic diets &/or diet pills

Managing Side-Effects

Skin Sensitivity Use sunscreen Wear protective clothes Avoid sunlight/sunlamps **Impaired Sleep** Have routine sleep habits Don't let the weekend disrupt this by > 1 hour No exercise/caffeine in late evening Wake at regular time EVEN IF TIRED! Don't nap during the day

Part 3 of 6

Treatment: Light Therapy Swedo et al, 1997, JAACAP, 36(6): 816-821 N=28, 7-17 yrs (Boston, Washington, DC) Double-blind, placebo-controlled crossover design 1 hr bright-light tx+2 hours dawn stimulation Total depression, atypical depression, typical depression scores improved with phototherapy 78% parents, 80% children reported phototx was "best" condition **CLINICAL GUIDELINES** Move to a warmer, sunnier climate OR

 Sit for 20-40 minutes, 1-2X/day in front of a special lamp ECT: Electroconvulsive Therapy *Walter & Rey, 6/97* Epidemiologic study (NSW) of all adolescent ECT from '90-'96

Results from 42 pts mimic adult lit

- half had marked/total improvement
- pts w/ mood disorders derived the most benefit

- side effects were transient and minor

 comorbid personality disorder predicted poorer response

propofol was associated w/ shorter seizures

Electroconvulsive Therapy (ECT) Practice Parameters, J Am Acad Child Adol Psychiatry, 2004;43(12):1521–1539

Treatment of choice for adults w/: pregnancy, catatonia, neuroleptic malignant syndrome, other medical conditions that contraindicate standard medication regimens
 Limited case studies indicate ECT is beneficial for youth

w/:

Mania, rapid cycling, depressed phase--BPD

Requires two board certified child and adolescent psychiatrists to independently review records and recommend ECT

Uncommonly used, has potential for adverse events but often provides significant symptomatic relief

Nutritional Interventions

Omega-3 Fatty Acids

Multi-nutrient complex

Changes in Dietary Ω 3 Scheffer, 2008, in Geller & Del Bello "Treatment of Bipolar Disorder in Children and Adolescents" • Hunter-gatherers: $\Omega 3: \Omega 6$, 1:1 Modern ratio: 1:100 or greater <10% adult Americans obtain</p> sufficient Ω 3 in their diets Rates are worse for children

Ω3 Treatment: Depression in Children

Nemets et al, '06, Amer J Psychiatr, 163(6): 1098-1100
28 children aged 6 to 12 with MDD, 16-week RCT

N=13, Ω3 (2:1 EPA: DHA, 380-400 mg EPA and 180-200 DHA); n=15, placebo (pbo)

8 children dropped out and were deleted from analyses (5, pbo; 3, Ω3)

No clinically significant side effects reported

 Ω 3 Treatment: **Depression in Children** Nemets et al, '06, Amer J Psychiatr, 163(6): 1098-1100 >50% ↓ in depressive sxs: 7/10 Ω3 vs 0/10 pbo Remission: $4/10 \Omega 3 vs 0/10 pbo$ Most response noted at Wk 16 was achieved by Wk 12 Findings support Ω3 as potentially beneficial for childhood depression. Risk: benefit ratio may be superior to that of traditional anti-depressants

Ω3 Treatment: Bipolar Disorder in Children Wozniak et al, '07, European Neuropsychopharmacology, 17, 440-447

- 8-wk open-label trial of Ω3 fatty acids (7:1 EPA:DHA)
- Monotherapy (only exception: stimulants)
- 20 outpatients aged 6-17 with bipolar disorder
- 1.3 to 4.3g of Ω 3 daily, 85% taking ≥2.0 g/day
- 16 participants completed the entire trial
 Drop-outs were d/t lack of efficacy, not side effects

Ω3 Treatment of Bipolar Disorder in Children Wozniak et al, '07, European Neuropsychopharmacology, 17, 440-447

 \square Ω 3 caused few, mild side effects, usually GI

Manic symptoms: ↓ by 30% in 50%; ↓ by 50% in 35%

Participants on $\geq 2.0g \ \Omega 3$ improved more than those on < 2.0g

Depressive symptoms: 40% rated much/very much improved

Nutrition & Mental Health

Iron	Produces ATP-brain energy, maintains oxygen levels in brain, binding of serotonin & dopamine in frontal cortex, neurotransmitter production	
Copper	Neurotransmitter production, ratio to copper important-metabolic functioning	
Zinc	Found in glial cells & neurons, protein synthesis & regulation of gene expression, involved in >200 enzymatic reactions-many in brain, central role in metabolism	
B1	Neuronal health, deficiencies impact cardiovascular & PNS- Wernicke's encephalopathy	
B6	Deficiencies may decrease serotonin & GABA in brain	
B12	Essential to brain & CNS functioning maintains myelin sheaths, synthesis of monoamine neurotransmitters, alters folate levels	
D	Deficiencies related to depression & abnormal brain development	
Е	Anti-oxidant, protects cell membranes potentially effecting brain functioning	
Folate	Neurotransmitter synthesis, relationship btw tryptophan & serotonin, deficiencies related to dementia & mood d/o	
(Hutto, 1997; Kaplan et al., 2007; McCarty, 2000; Meador et al., 1993; Takeda, 2001; Velez-Pardo et al.,		

(Hutto, 1997; Kaplan et al., 2007; McCarty, 2000; Meador et al., 1993; Takeda, 2001; Velez-Pardo et al., 1995)

Multi-Ingredient Supplements Kaplan et al, 2007, Psych Bulletin, 133(5), 747-760

Past nutritional research focused on individual vitamins and minerals

To understand results of these studies, need to know potential mechanisms underlying the positive effects of nutrients on mood.

Multi-Ingredient Supplements: **Theorized Mechanisms of Action** Kaplan et al, 2007, Psych Bulletin, 133(5), 747-760 Mood dysregulation may result from innate metabolism malfunctions that ultimately affect brain functioning Mood instability may result from deficiencies in methylation of molecules responsible for completing DNA transcription, switching on genes, regulating protein generation, activating enzymes, and synthesizing neurotransmitters Nutrition deficiencies may alter gene expression and lead to mood instability Unstable mood may result from long-latency effects of nutrient deficiencies that alter brain development directly or by way of dysfunctional nutrient absorption

Impact of Diet & Nutrition on Brain Function Benton, Eur J Nutr 47(3):25-37, 2008

Diet and nutrition affect the structure and functioning of the brain
 In adulthood, the brain contributes 20% of the human basal metabolic rate

In neonates, this number is as high as 44%

Nutrition deficiencies, particularly in prenatal and neonatal stages, may contribute to long-term physical and behavioral detriments

Vitamin-Mineral Treatment of Anti-social Behavior in Children

Schoenthaler and Bier (2000) Journal of Alternative & Complementary Medicine, 6(1), 7-18

468 6--12 yr olds from two working class, primarily Hispanic elementary schools in the Southwestern US RCT— 1/2 placebo, 1/2 daily vitamin-mineral supplementation at 50% of the US recommended daily allowance (RDA) for 4

months Outcome variable: disciplinary infraction

Outcome variable: disciplinary infractions

Vitamin-Mineral Treatment of Anti-social Behavior in Children Schoenthaler and Bier (2000) Journal of Alternative & Complementary Medicine, 6(1), 7-18

Vitamin-Mineral group had, compared to the placebo group

47% lower mean rating of antisocial behavior (1.0 vs. 1.9 disciplinary actions)

 lower ratings for every type of recorded infraction: threats/fighting, vandalism, being disrespectful, disorderly conduct, assault/battery, defiance, obscenities, refusal to work or serve, endangering others, and nonspecified offenses

 Ω & Vitamin-Mineral Treatment of **Antisocial Behavior** Gesch et al 2002 British Journal of Psychiatry. 181: 22-28 Double-blind clinical trial—231 prisoners 142 days on recommended daily doses of two supplements one capsule of Forceval, a 25-ingredient vitamin supplement four capsules of Efamol Marine, containing 1260mg linoleic acid, 160mg gamma linolenic acid, 80mg EPA and 44mg DHA or placebo (identically appearing oil-based gelatin) capsules)

Ω3 & Vitamin-Mineral Treatment of Antisocial Behavior

Gesch et al 2002 British Journal of Psychiatry. 181: 22-28

Neither group reported notable side effects

 Supplement group experienced a 26% decrease in overall infringements resulting in disciplinary reports compared to placebo group (p<.03)

EMPower (EMP+) www.truehope.com

Contains 36 ingredients
Vitamins:

A, C, D, E, B1, B2, B3, B5, B6, B9, B12, H (Biotin)
Minerals:

 Calcium, Phosphorus, Magnesium, Potassium, Iodine, Zinc, Selenium, Copper, Manganese, Chromium, Molybdenum, Iron, Nickel
 Other:

 dl-Phenylalanine, Glutamine, Citrus Bioflavonoids, Grape Seed, Choline, Inositol, Ginkgo Biloba, Methionine, Germanium, Boron, Vanadium

EMP: Animal Data Halliwell & Kolb (2003) Society for Neuroscience Abstracts, 29, 459.11., 29, 459.411

- Newborn rats received frontal or posterior parietal lesions on Day 3
- Subsequently fed either normal rat chow or rat chow enhanced with a rodent-appropriate dose of EMP+
- At Day 60, supplemented animals, compared to unsupplemented rats
 - exhibited deficit reversals (eg, performance on spatial learning tasks)
 - had significant regrowth of cortical tissue
 - were significantly calmer

EMP: Open Label Adult BPD Study

Kaplan et al, '01, J Clin Psychiatr, 62(12): 936-944

- 11 adults aged 19-46 yrs diagnosed w/ BPD and resistant to conventional treatments
- Treatment occurred for 6-21 months with no medication restrictions (i.e., participants could continue use of concurrent psychiatric medications under the supervision of their psychiatrist)
- Outcome variables:
 - Hamilton Depression Rating Scale (HAM-D: Hamilton, 1960)
 - Young Mania Rating Scale (YMRS: Young et al., 1978)
 - Brief Psychiatric Rating Scale (BPRS; Overall & Gorham, 1962)

EMP: Open Label Adult BPD Study Kaplan et al, '01, J Clin Psychiatr, 62(12): 936-944 Results indicated **55-66%** reduction in HAM-D, YMRS and BPRS scores 50% decrease in need for psychotropic medications only one mild side effect reported, infrequent/transitory nausea, which occurred most commonly when participants took their supplement without food Conclusion: more studies are needed

EMP: Child Bipolar Study *Popper, 2001, <u>J Clin Psychiatr</u>, 62(12): 933-935*10-year old boy with bipolar disorder ABACB design

- A: 4 mos, daily severe temper tantrums for multiple hrs
- **B**:
 - 2 days after full EMP dose, behavior improved significantly
 - by 5 days, tantrums/irritability ceased
- A: After 14 days, EMP was discontinued
 - Within 2 days, tantrums resumed
- C: different supplement used, resulting in (per parent and teacher reports) 60% of the benefit
 B: EMP resumed, tantrums/irritability resolved

EMP: Open Label Bipolar Study Popper, 2001, J Clin Psychiatr, 62(12): 933-935

22 patients with bipolar disorder 10 adults, 9 adolescents and 3 children Mild side effects (eg, headache) common ■ 19/22 (86%) responded positively 11/15 patients (73%) previously on psychiatric medications remained stable without resumption of these medications at both 6-month and 9-month follow-up assessments

EMP: Open-Label ABAB Trial Kaplan et al, 2002, J Ch Adol Psychopharm, 12(3): 205-219 2 boys w/ irritability, mood lability, explosive rage 8-year old: atypical obsessive-compulsive disorder (OCD) and ADHD Course of treatment: ABAB A: Baseline/EMP withdrawal: consistent explosive rage, irritability, obsessions-guns B: On EMP: these behaviors almost completely dissipated. Obsessive thoughts ceased, frequency and duration of temper outbursts \downarrow significantly, mood fluctuations minimized. After \geq 2 years of treatment, the boy remained well and free of side-effects while taking 25% of his initial dose of EMP

EMP: Child Case Series Kaplan et al, 2004, J Child Adol Psychopharm, 14(1), 115-122

N=11 children aged 8—15, all with mood and/or behavioral problems

Intake diagnoses: n=3, BPD (1+anxiety, 1+behavior); n=2, Asperger's+anxiety; n=5, ADHD+ anxiety +/or behavior disorders; and n=1, Praeder-Willi syndrome, ODD + anxiety

9/11 (82%) completed the open-label trial

EMP: Child Case Series Kaplan et al, 2004, J Child Adol Psychopharm, 14(1), 115-122

- Intent-to-treat analyses indicated significantly ↓ scores from baseline to final visit on Youth Outcome Questionnaire (YOQ: p<.001) and Young Mania Rating Scale (YMRS: p<.01)</p>
- For the 9 completers, improvement was significant on 7/8 (88%) Child Behavior Checklist scales, YOQ and YMRS (p values ranged from <.05 to <.001)
 Conclusion: need PCTs
- Conclusion: need RCTs

OSU Case Study: EMP+ Frazier, Fristad & Arnold, 2009, J Ch Adol Psychopharm, <u>19(4)</u>: 453-460.

Boy diagnosed at age <u>5 yrs, 11 mos</u> with BP-NOS, rule-out generalized anxiety disorder (GAD)

severe mood cycling, sadness, irritability, selfharming behaviors, sleep disturbance, severe tantrums, elevated mood, poor peer relations, low frustration tolerance, flight of ideas, aggressive behavior, hyperactivity and impulsive negative behaviors

OSU Case Study: EMP+

By <u>age 8</u>: diagnoses evolved into BP-I, enuresis, GAD, learning disorder-not otherwise specified (LD-NOS), communication disorder-not otherwise specified

impairing anxiety and worsening mood symptom intensity and cycling, increased destructive behavior, transient suicidal ideation, and increased global impairment

OSU Case Study: EMP+

By <u>ages 10-11</u>: developed psychotic features (auditory hallucinations), obsessions and compulsions

increasingly disrespectful and aggressive behaviors. Intrusive, command hallucinations when anxious, told him to act on his compulsions, do things he did not want to do, and threatened him, $\leq \sim 100$ X/wk telling him, "If you don't do this you'll surely die." and "Don't listén to them [referring to parents and other adults]". In 2006, the voices told him he would die on the day John Glenn dies. Symptoms became impairing to the point that he was removed from his private school and was home schooled.

OSU Case Study: Conventional Treatment

Medication+psychoeducational psychotherapy Many medication trials d/t intolerable side effects +/or inadequate treatment response lithium citrate, risperidone, lithium carbonate, clonidine, flax seed, desmopressin, omega-3 fatty acids, trazodone, gabapentin, valproic acid, propranolol, quetiapine, aripiprazole, lorazepam, and lamotrigine No medication combination maintained symptom amelioration and adequate global functioning over an extended period of time

2008	1/21/08	1/22/08	1/29/08	1/31/08	2/1/08
Begin EMPower Plus (EMP+)	EMP+ 5 pills/day Lithobid Lamictal	EMP+ 10 pills/day, Lithobid Lamictal	EMP+ 15 pills/day, cut ¼ Lithobid & ¼ Lamictal	EMP+, cut Lithobid & Lamictal by ½	EMP+, cut Lithobid & Lamictal by 1⁄2
Taper off all psycho- tropic medica- tions between 12/07- 2/08			slight headache, improve- ment in bowels	irritability after taking regular meds, "downer" after lunch, dizziness & extreme tiredness before bed	slight headache in AM- gone by lunch

2/2/08	2/3/08	2/4/08
EMP+, cut Lithobid & Lamictal by ½	EMP+, cut Lithobid & Lamictal by 1⁄4	EMP+, cut Lithobid & Lamictal by ¼ (no regular meds in AM)
slight irritability in early afternoon, very tired in PM after receiving regular meds	"felt, looked, & acted great", no irritability, no social issues tired & dizzy after receiving regular meds in PM	

2/10/08	3/6/08	5/2/08
EMP+	EMP+ only	EMP+ only
only		antest sugar set
	focused & efficient in school, good peer relationships, calm & playful, sleeping well, regular BMs, irritable & angry if pushed around 3-4PM followed by genuine apology, "mouthy" at times, decreased fidgeting, some compulsions but are ignored, child feels "natural"	good peer interactions, enjoys school, active in sports, all symptoms stable no longer has dry skin

Additional Follow-Up

- May, '08:
 EMP+ dose from 15 to 18 capsules/day d/t
 irritability, ? mood cycling, and "odd" behavior
 - Mother and boy reported this relieved symptoms
- Summer, '08: ↓ EMP+ to 15 capsules/day
- Fall, '08: Started regular public school (for the first time), plays on a school soccer team, maintains friendships, parents report improvements in his functioning while taking EMP+ are greater than those he has made in the past on medications
- His only additional intervention has been drinking whey protein mixed in milk when very active. During sports seasons, he takes ½ cup of whey protein mixed with milk in the morning, which according to his mother, helps keep him "clear, slowed down, peaceful, settled and happy"
 <u>Summer, '09:</u> Continues to remain stable, a "normal" teenager <u>Summer, '11</u>: Needed to ↑ dose

Open Label Pilot Exploration of a Nutritional Supplement for Childhood Mood Dysregulation *Frazier, Fristad, Arnold, in press, J Comp Alt Med*

- N=10, 2-month open label trial of EMP+
- Explore feasibility, provide preliminary efficacy data
- Participants had to remain off all psychotropics for 11 wks (3 wks prior to EMP+ and 8 wks of EMP+)
 Any other treatment could continue
 Seven assessments conducted

 interviews with parent and child
 brief physical exam
 blood draws at the second and last visit

Participants

■ 10 children age 6-12; *M* = 8.9, *SD* = 2.0 60% male ■ 90% White, 10% Hispanic ■ IQ estimate: *M* = 98.7, *SD* = 9.5 Income: \$20,000 to >\$100,000 ■ 10% BP-I, 30% BP-NOS, 60% sub BP-NOS Sub BP-NOS: 20% MDD & 10% Dysthymia 100% comorbid behavior d/o, 60% comorbid anxiety, 10% enuresis 70% completed full open-label study

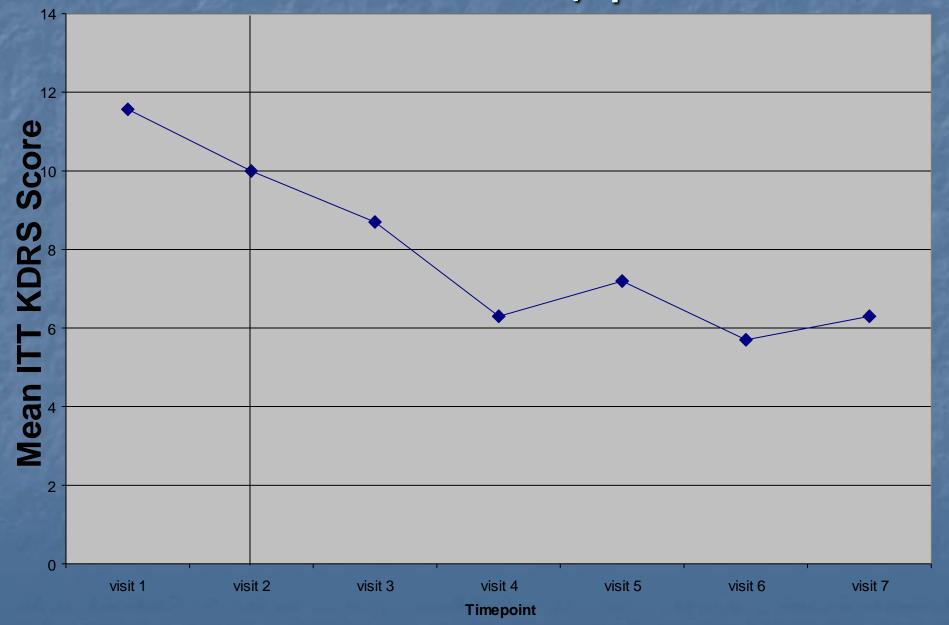


1 capsule tid, increased up to a max of 5 capsules tid • Average 46.4 days on EMP+ (SD = 23.3, median = 55)No concomitant psychotropics N=3 on psychotropic medications at enrollment; completed \geq 3-week washout \square N=1 n-3 + melatonin n-=1 n-3 fatty acids; Focalin 10mg added after Visit 5 N=10 continued pre-existing psychosocial services outside of study

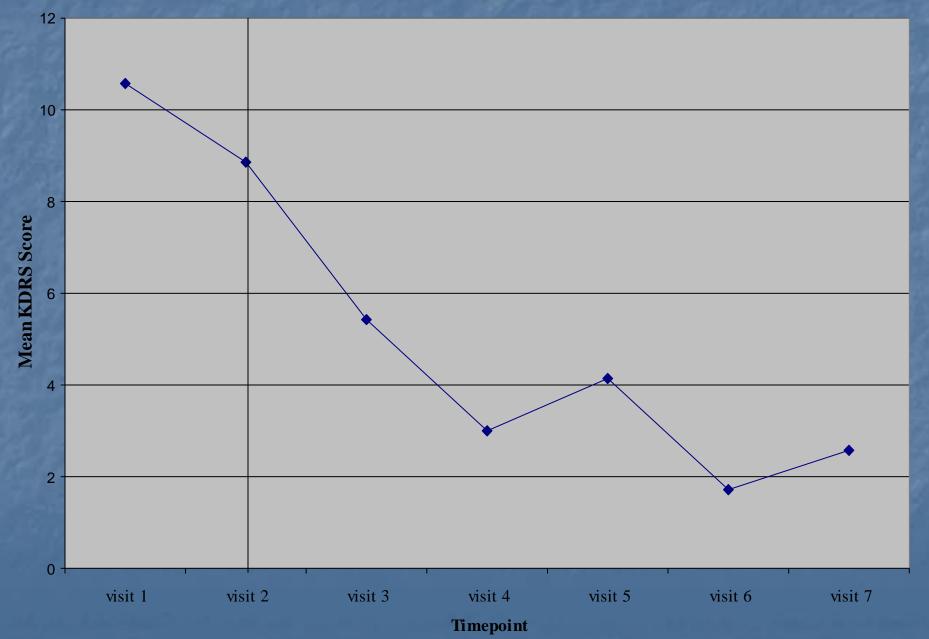
Side Effects

4 mild nausea w/o food 1x, 1 vomited 1x
3 mild initial insomnia 1x
1 mild ↑ appetite 1x
1 nocturnal enuresis 1x

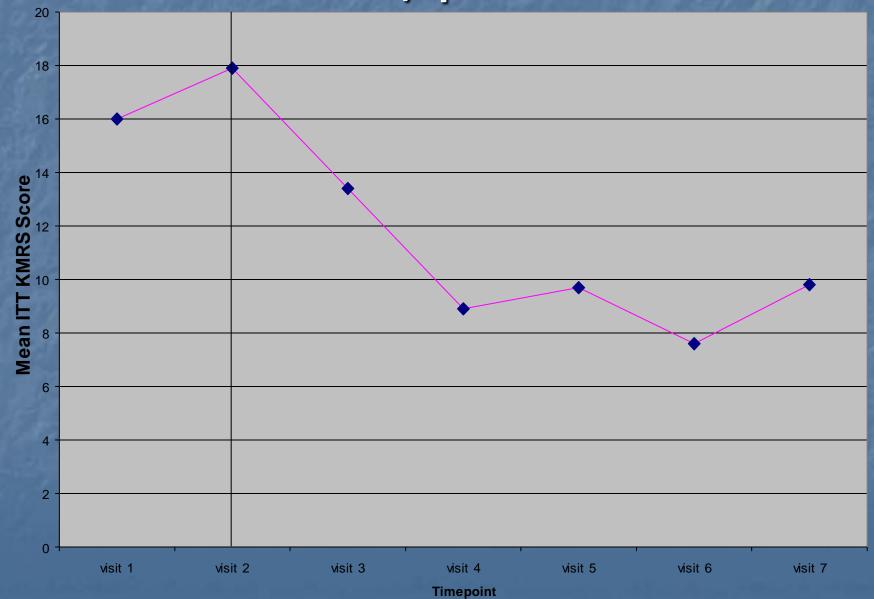
ITT (N=10) Depression Scores Over Time, p<.06



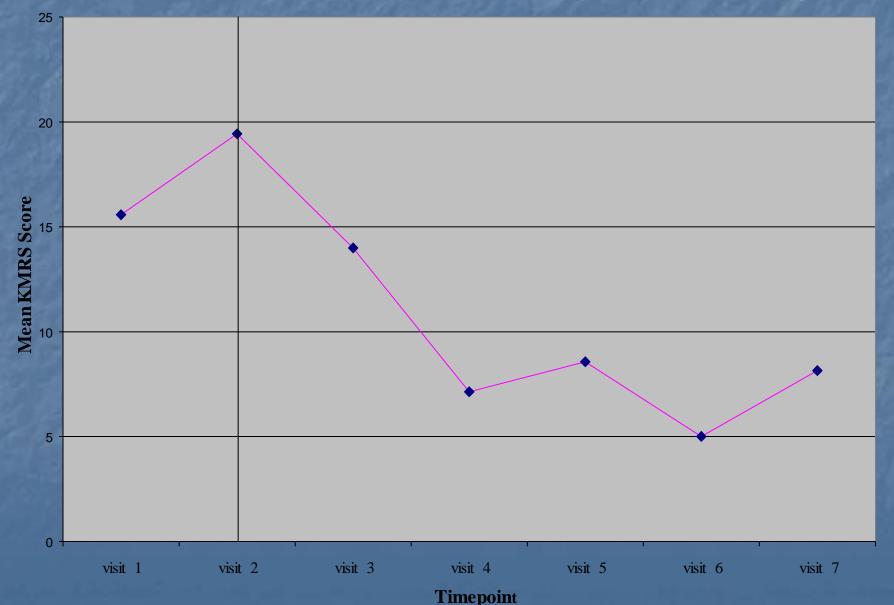
Mean Depression Scores Over Time (n=7) Study Completers, p<.05



ITT (n=10) Mania Scores Over Time, p<.01



Mean Mania Scores Over Time (n=7) Study Completers, p<.05



Discussion-Medication Compliance

3 participants dropped out due to tolerability/palatability/compliance Of the 7 completers, all maintain high compliance All maintained at least 93% 2 maintained 100% Those who fit swallowing inclusion criterion displayed excellent treatment adherence

Treatment Response

Suggest depression and mania symptoms over time
 Open-label design, interpret w/ caution

- Placebo-effects, 18.7%-33.6% in review of adult BPD medication trials (Smith, Cornelius, Warnock, Tacchi & Taylor, 2007)
- Anecdotal follow-up
 - 4 continued on EMP+ post study completion
 - 2 were considering cost, unknown
 - 1 began mood stabilizer
 - 1 began antidepressant
 - 2 unknown

 Future, more scientifically rigorous placebo-controlled trials appear warranted

Workshop Goals Participants will learn: How BPSD presents in children Biological interventions Psychosocial treatment for BPSD Multi-family psychoeducational psychotherapy (MF-PEP) Individual-family psychoeducational psychotherapy (IF-PEP)

Psychotherapy for Children with Bipolar and Depressive Disorders

> Mary A. Fristad Jill S. Goldberg Arnold Jarrod M. Leffler

The OSU Psychoeducational Psychotherapy (PEP) Program

Orientation

- Nonblaming/growth-oriented
- Biopsychosocial—uses systems and cognitive-behavioral techniques
- Education + Support + Skill Building → Better Understanding → Better Treatment + Less Family Conflict → Better Outcome

Three formats

- Multi-family psychoeducational psychotherapy (MF-PEP)
- Individual family psychoeducational psychotherapy (IF-PEP)
- workshops

How to Conceptualize **Family-Based Intervention** Historically, families Have been blamed Have not gotten useful information/support/skill building This can result in families being "skittish" or "defensive" about family-based intervention

Goals of Psychoeducation

Teach parents and children about The child's illness & its treatment Provide support Peers ("I'm not the only one") Professionals - understand the disorder Build skills problem-solving communication symptom management

Psychoeducation: Treatment Goal

If you give a man a fish, he will eat for a day. If you teach a man to fish, he will eat for a lifetime.



Pro's and Con's of MF-PEP +Peer support- for children and parents +In-vivo social skills training +Parents and children learn from each other -Can't tailor make treatment for specific needs of child/parent -Need to consider attend to group dynamics when selecting members -Schedule harder for families to accommodate

Therapist (3) Requirements: MF-PEP

Parent & Lead Child Therapist: be familiar with Mood disorders Children/families Group therapy Child Co-Therapist: be familiar with Behavior management Excellent role for trainee

Additional Features of MF-PEP Point system—accumulate at Each session-for completed projects, participation Review game (cash in after playing) Immediate reinforcers (eg, Starbursts) News of the Week Games (social skills, thematically linked) Think about group composition Don't leave any member "stranded" SES, gender, symptom severity Therapist workbook includes: Keep in Mind, Leader Tip, Examples, Games, Posters+ prompts Child Workbook page prompts

Pro's and Con's of IF-PEP

+Can flexibly administer treatment modules
 +Includes Healthy Habits, siblings, school professional contact

- +In-the-bank sessions allow for repeating and/or augmenting of treatment
- +Easier to schedule for families
- -Don't meet others struggling with unique issues of mood disorder
- -Social skills training with peers not available in session

Therapist (1) Requirements: IF-PEP

Be familiar with
 Mood disorders
 Children and families

Resources

OATS-Bipolar, 2011-2014, NIMH R34

- OATS=Omega3 and Therapy Study
- Kayden Healy, <u>614-293-4908</u>
- N=60
- 12 week trial
- 8-14 years
- BP-NOS, cyclothymic disorder
- No meds/psychotherapy in previous month except stable stimulants, sleeping aids

	Omega3	Placebo	TOTAL
IF-PEP	15	15	30
Active Monitoring	15	15	30
TOTAL	30	30	60

Books for Children

Brandon & the Bipolar Bear -- *T. Anglada* My Bipolar, Roller Coaster, Feelings Book & Workbook—*B. Hebert*

The Storm in My Brain -- Child & Adolescent Bipolar Foundation (CABF): 1-847-256-8525, www.bpkids.org

Kid Power Tactics for Dealing with Depression
 -- N. & S. Dubuque

Matt, The Moody Hermit Crab -- C. McGee

Anger Mountain—*B. Hebert*

Books for Adolescents

- When Nothing Matters Anymore: A Survival Guide for Depressed Teens -- *B. Cobain* Recovering from Depression: A Workbook for Teens -- *M. E. Copeland & S. Copans* Conquering the Beast Within: How I Fought
- Depression & Won...& How You Can, Too -- *C. Irwin*

Mind Race: A Firsthand Account of One Teenager's Experience with Bipolar Disorder – *P.E. Jamieson & M.A. Rynn*

Children's Literature

The Phoenix Dance Dia Calhoun, award winning author Farrar, Straus & Giroux, NY, 2005

 Based on the Grimms' Twelve Dancing Princesses

 Explores the experience of bipolar disorder in an adolescent girl

Books for Parents

Raising a Moody Child: How to Cope with Depression and Bipolar Disorder -- M.A. Fristad & J.S. Goldberg-Arnold New Hope for Children & Teens with Bipolar Disorder—*B. Birmaher* The Childhood Bipolar Disorder Answer Book— T. Anglada & S.M. Hakala The Bipolar Child -- D. & J. Papalos A Parent's Survival Guide to Childhood Depression -- S. Dubuque

Books for Adults

- Out of the Darkened Room: Protecting the Children and Strengthening the Family When a Parent is Depressed --*W. Beardslee*
- Living Without Depression & Manic Depression -- M. E. Copeland
- An Unquiet Mind -- K. Redfield Jamison
- Thoughts & Feelings: Taking Control of Your Moods & Your Life -- M. McKay, M. Davis & P. Fannin
- The Bipolar Survival Guide: What You and Your Family Need to Know --D.J. Miklowitz
- Winter Blues: Seasonal Affective Disorder- What it is and How to Overcome it -- N.E. Rosenthal

More Books to Read

General Parenting

- How to Talk So Kids Will Listen & Listen So Kids Will Talk --Faber & Mazlish
- The Explosive Child -- R. Greene
- The Optimistic Child -- *M. Seligman*

Sibling Issues

- Siblings Without Rivalry -- A. Faber & E. Mazlish
- Turbo Max: A Story For Siblings of Bipolar Children -- T. Anglada

Understanding Psychiatric Disorders

- It's Nobody's Fault -- H. Koplewicz
- Understanding Psychiatric Medications
 - Straight Talk About Psychiatric Medications for Kids --- T. Wilens

Miscellaneous

- I Am Not Sick, I Don't Need a Help! -- X. Amador & A.L. Johanson
- The Thyroid Sourcebook -- M.S. Rosenthal

Educational Websites

Information re: BPD for Parents, Children and Educators

- <u>www.bpchildren.com</u>
- <u>www.schoolbehavior.com</u>
- <u>www.bpkids.org</u>
- www.josselyn.org/Store.htm

Special Education Advocacy -- <u>www.wrightslaw.com</u>

 National Association of Therapeutic Schools and Programs—www.natsap.org

Internet Special Education Resources (ISER)

<u>www.iser.com/index.shtml</u>

Groups/Websites – Adults, Families & Children

National Alliance on Mental Illness (NAMI) 1-800-950-6264 <u>www.nami.org</u> Mental Health America (NMHA) 1-703-684-7722 www.nmha.org Depressive & Bipolar Support Alliance (DBSA) 1-800-826-3632 www.dbsalliance.org Families for Depression Awareness (FFDA) 1-718-890-0220 www.familyaware.org Child & Adolescent Bipolar Foundation (CABF) 1-847-492-8519, <u>www.bpkids.org</u> Juvenile Bipolar Research Foundation (JBRF) 1-866-333-5273, www.bpchildresearch.org BP Children <u>1-732-909-9050 (fax) www.bpchildren.com</u>

Additional Resources

Light Therapy: Center for Environmental Therapeutics www.cet.org Nutritional Intervention: EMpower Plus 1-888-878-3467 <u>www.truehope.com</u> Omega-Brite 1-800 383 2030 www.omegabrite.com Evidence-Based Treatments: www.effectivechildtherapy.com PEP Workbooks www.moodychildtherapy.com

Note that Parts 4-6 follow the PEP manuals.

PEP & MF-PEP Resources

Moody Child HOW TO COPE WITH DEPRESSION AND BIPOLAR DISORDER

Mary A. Fristad, PhD and Jill S. Goldberg-Arnold, PhD

Raising a

Books & DVD for parents or therapists order from www.amazon.com

beyond the

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Dr. Fristal has been the principal or co-principal invostigator on to two doors factoria, tatia, and local grants. Bearthy, the wasmented on the state of the state minimum state of the sta

Professed by Video Propulsion, Inc. Bit minutes © 2007, Mary A. Frankel, Ps.D., ABPP and Video Propulsion, Inc. All Rights Reserved. Bipolar Children and Ther Fam

beyond the book bipolar children and their families an indepth interview with Mary A. Fristad, Ph.D., ABPP



bipolar children and their families an in-depth interview with Mary A. Fristad, Ph.D., ABPF



Treatment Manual—2011, Guilford Press

Clinical Manual for

Children and

Adolescents

Robert A. Kowatch, M.D. Mary A. Fristad, Ph.D., ABPP Robert L. Findling, M.D. Robert M. Post, M.D.

Management of **Bipolar Disorder in**

Home Study Course— *for professionals* Taped 2 day seminar by Dr. Fristad 6 or 12 hours Continuing Education credit \$95 for CD or cassette \$65 for test scoring/reporting <u>www.jkseminars.com</u>

<u>www.moodyennetherapy.com</u>

Child, Parent & Child Therapist MF-PEP Workbooks

Child & Parent PEP Workbooks

Thank You, The End

mary.fristad@osumc.edu

Clinic #: 614-293-9600

I670 Upham Drive Suite 460G Columbus, OH 43210-1250 For more information, please go to the main website and browse for more videos on this topic or check out our additional resources.

Additional Resources

Online resources:

1. Society of Clinical Child and Adolescent Psychology website: <u>http://effectivechildtherapy.com</u>

2. Psychoeducational Psychotherapy Workbooks: <u>www.moodychildtherapy.com</u>

Books:

1. Fristad, M.A., Goldberg Arnold, J.S. & Leffler, J. (2011). *Psychotherapy for Children with Bipolar and Depressive Disorders*. New York: Guilford Press.

Selected Peer-reviewed Journal Articles:

1. Fristad, M.A., Verducci, J.S., Walters, K. & Young, M.E. (2009). The impact of multi-family psychoeducational psychotherapy in treating children aged 8-12 with mood disorders. *Archives of General Psychiatry*. *66*(9);1013-1021.

2. Kaplan, B.J., Crawford, S.G., Field, C, J., & Simpson, J.S. (2007). Vitamins, minerals, and mood. *Psychological Bulletin*, 133(5) 747-760.

3. Kowatch, R.A., Fristad, M.A., Birmaher, B., Wagner, K.D., Findling, R.L., Hellander, M., & the Child Psychiatric Workgroup on Bipolar Disorder (2005). Treatment guidelines for children and adolescents with bipolar disorder. *Journal of the American Academy of Child and Adolescent Psychiatry* 44, 213-235.

4. Lofthouse, N. and Fristad, M. (2004) Psychosocial interventions for children with bipolar disorder. *Clinical Child and Family Psychology Review*, 7, 71–88.

5. Youngstrom, E.A., Findling, R. L., Youngstrom, J. K., & Calabrese, J. R. (2005). Toward an evidence-based assessment of pediatric bipolar disorder. *Journal of Clinical Child & Adolescent Psychology*, 34, 433-448.







Center for Children and Families