

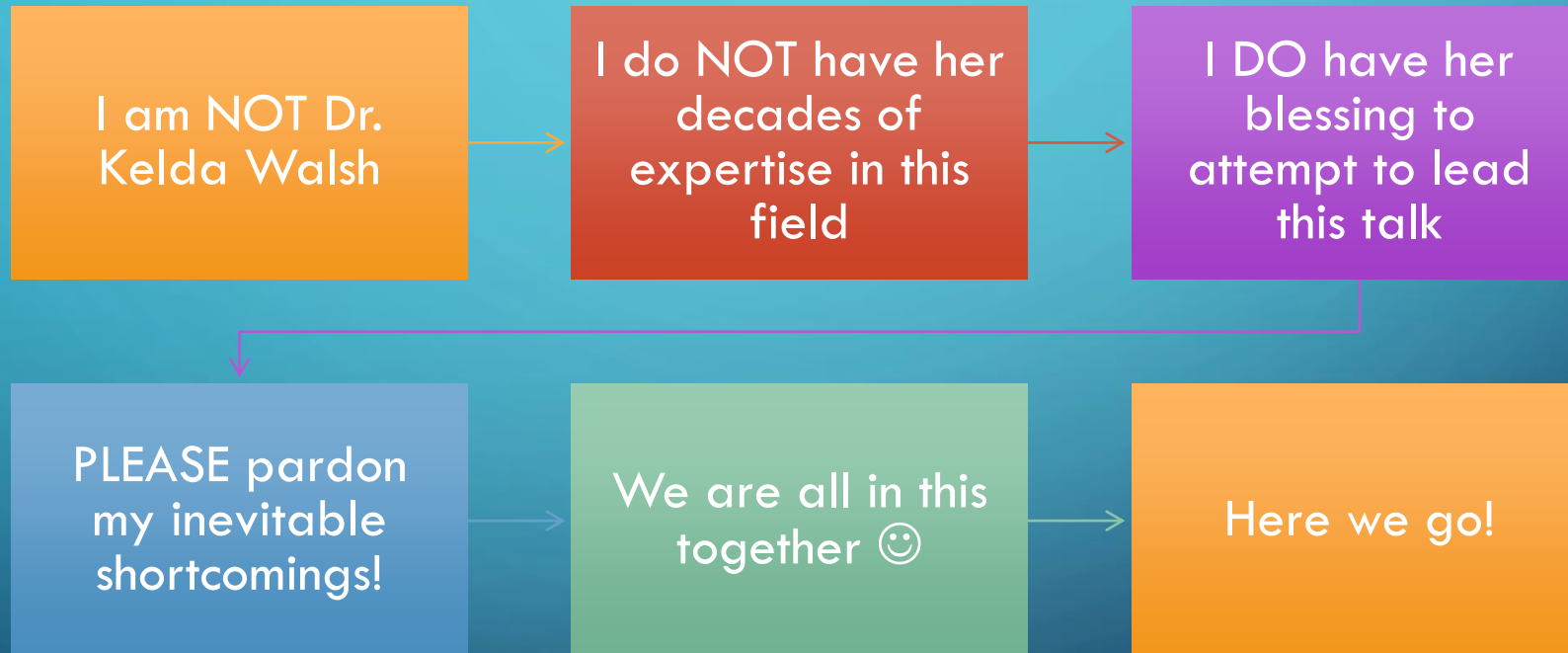
MEDICATION MANAGEMENT OF PEDIATRIC OCD AND ANXIETY DISORDERS

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CLINICS

HI! I'M CLAIRE.

I'M A 5TH YEAR POST GRADUATE TRAINEE, NEAR THE END OF MY CHILD & ADOLESCENT PSYCHIATRY FELLOWSHIP



COVID-19 RELATED ANXIETY: RESOURCES

Mental Health America

[Mhanational.org/covid19](https://www.mhanational.org/covid19)

- Resources on coping with COVID anxiety for children, parents and adults
- Activity book “**Coping after a disaster**” for 3-10 year olds
 - (identifies many kinds of disasters, including health disasters)
- Emphasis: keeping normal schedule, emphasizing adults are working to keep children safe, identifies helping adults.
- Minimize exposure to media; share limited information with children.



FDA- APPROVED ANXIETY MEDICATIONS

For pediatric obsessive-compulsive disorder (OCD)

- Clomipramine (Anafranil - CMI; 10 and older)
- Fluvoxamine (Luvox - FLV; 8 and older)
- Fluoxetine (Prozac - FLX; 7 and older)
- Sertraline (Zoloft - SER; 6 and older)

For generalized anxiety disorder (GAD)

- Duloxetine (Cymbalta, 7 and older)

For non-OCD pediatric “anxiety”

- Chlordiazepoxide (Tranxene)
- Hydroxyzine (Vistaril, Atarax)



OCD

FIRST STUDIED

MOST DATA



ASSESSING OCD SEVERITY

Identify the current obsessions & compulsions

Assess severity

- Level of distress
 - Both patient & family
 - Assess level of family accommodation
- Frequency & total time spent on obsessions or compulsions. Is it increasing?
 - “How long can you go without worrying about OCD symptoms or doing rituals?”
- Assess ability to resist or delay
- Assess avoidance

WHAT IS THE MEDICATION RESPONSE?



- In pediatric OCD studies of serotonin reuptake inhibitors baseline OCD scores were reduced by 30-38% in responders
 - Studies included clomipramine plus the SSRIs
 - Child Yale-Brown Obsessive-Compulsive Scale (CY-BOCS) is the gold standard measurement in studies
- Patients often remain symptomatic despite adequate medication doses
- Patients need therapy

MED ALGORITHM FOR OCD

Exposure and ritual prevention therapy
(no meds if mild to moderate)

Fluoxetine or sertraline
(start right away if symptoms are severe)

If first choice fails, use the other one
(30% better? *Think twice* before switching)

If second choice fails, assess therapy.
Is the child getting real exposure therapy?

Consider psychiatry referral for
clomipramine or antipsychotic augmentation



WHY ISN'T --- ON THE ALGORITHM?

Fluvoxamine

- It wears off really quickly. More SRI discontinuation symptoms if doses missed (clomipramine has this issue too).
- It does have FDA approval, best option if you need an SSRI when sertraline/fluoxetine aren't working.

Paroxetine

- **ARGH!** Don't use it! Just don't!

Bupirone

- Data for OCD is poor.
- Data for children for any purpose is even poorer.

Escitalopram

- Approved for pediatric depression and adult depression/GAD.
- Not a lot of data for pediatric anxiety.

WHAT THE BIG
PEDIATRIC OCD
STUDY TELLS US:

PEDIATRIC OCD
TREATMENT
STUDY
(POTS)



12-week, 97 completers

7- to 17-year-olds without Tourette disorder



Compared efficacy of SER, CBT, COMB, PBO

Time to respond, predictors of response, relapse rate



Mean peak dose: 133 in COMB, 170 in SER only



Very precise adherence to therapy manual



Multiple iterations (POTS Jr (therapy only), etc.)

POTS OUTCOME

Remission (CY-BOCS \leq 10)

- COMB (54%) > E/RP (39%) > SER (21%) > PBO (4%)
- SER effect size 0.66
- 2 children with over-activity
- No suicidal ideation, hypomania, mania

Comorbid tics reduced likelihood that SER would work, although E/RP was still effective

- COMB > E/RP > SER = PBO
- *Therapy is EXTRA important for patients with OCD + tics*



OTHER OUTCOMES


Poorer Outcomes:

- More externalizing symptoms
- High family accommodation of OCD
- More aggression/ODD (to medication-only treatment)
- More severely ill kids were less likely to go into remission

With a family history of OCD, all treatment conditions were equally effective

- (i.e. E/RP didn't seem to work as well; were parents having trouble coaching?)

OCD psychoed (not E/RP) was not highly effective in follow-up study



NON-OCD ANXIETY DISORDERS

“TRIAD” ANXIETY DISORDERS:
SEPARATION ANXIETY DISORDER (SEP),
SOCIAL ANXIETY DISORDER (SOC),
GENERALIZED ANXIETY DISORDER (GAD)

MED ALGORITHM FOR ANXIETY DISORDERS

Anxiety management therapy focused on
EXPOSURE
(no meds if mild to moderate)

Fluoxetine or sertraline
(start right away if symptoms are severe)

If first choice fails, use the other one

If second choice fails, assess therapy.
Is the child getting real exposure therapy?

Consider psychiatry referral, duloxetine

CHILD- ADOLESCENT ANXIETY MULTIMODAL STUDY

(CAMS)



442 completers 7- to 17-year-olds, mean age 10.7



SEP, SOC, and/or GAD (78% had GAD)



SER, CBT, PBO, COMB, x 12 weeks




CBT: “Coping Cat” based



Mean final dose:

133.7+/-59.8 mg in COMB, 146+/-60.8 in SER groups



CAMS RESULTS

COMB > SER = CBT > PBO

Effect sizes: $0.86 > 0.45 = 0.31$ (statistically insignificant difference)

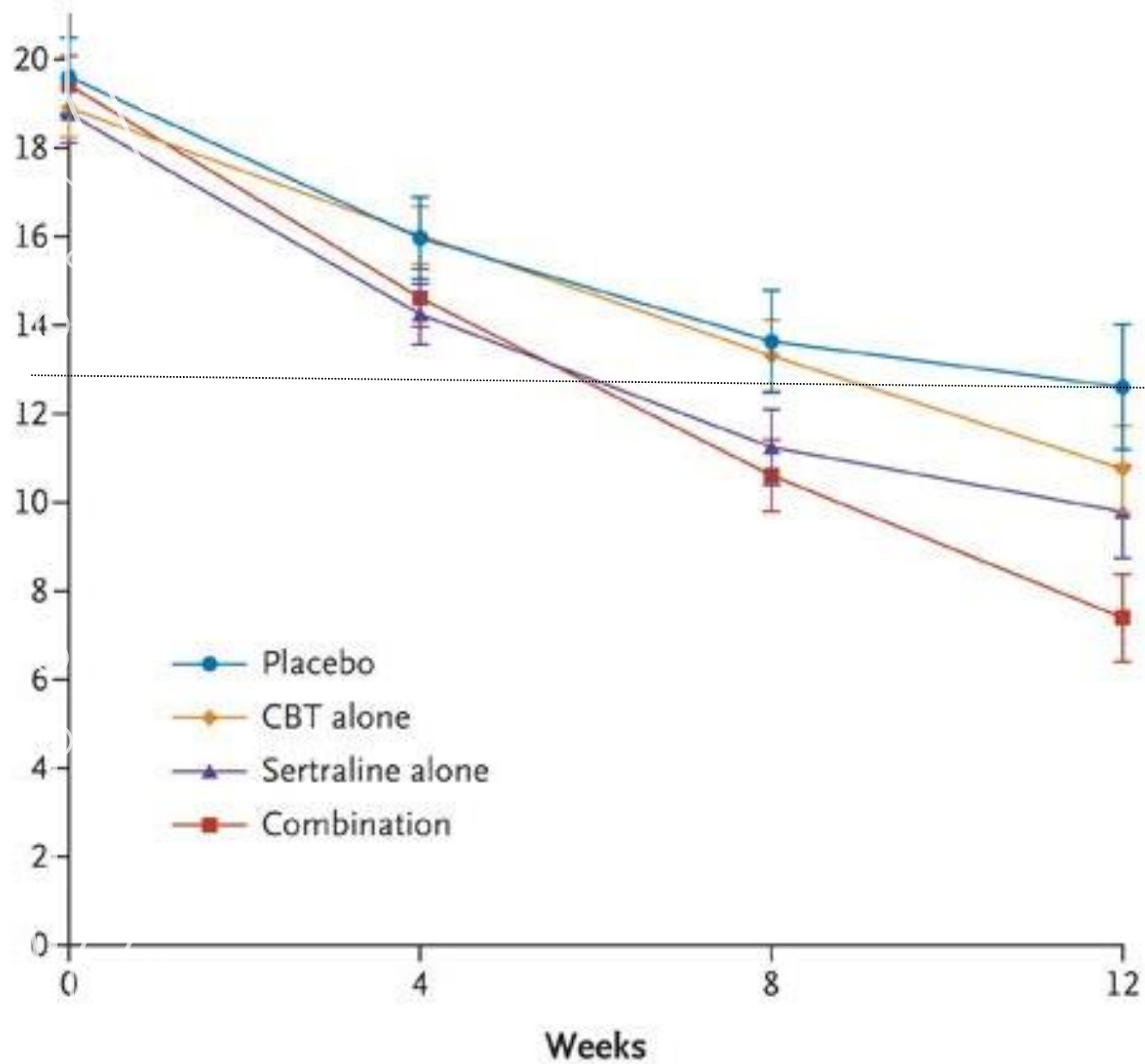
NNT: $1.7 > 3.2 = 2.8$

1 episode of non-suicidal self-harm and homicidal ideation

PARS (score >13 c/w mod anxiety disorder) (next slide)

(PARS = Pediatric Anxiety Rating Scale,
gold standard developed for studies/research, not usually clinically)

Expected Mean Score



PARS SCORE > 13
CONSISTENT WITH
MODERATE ANXIETY
DISORDER

(WALKUP JT ET AL., 2008)

CHILD/
ADOLESCENT
ANXIETY
MULTIMODAL
EXTENDED
LONG-TERM
STUDY

(CAMELS)
2014



**CAMS follow-up a mean of 6 years post study.
288/488 pts completed follow-up, Avg age 17**



**CAMS treatment condition not associated with remission,
anxiety severity, or global function at follow-up**



**CAMS initial responders much more likely to be in remission at
follow-up (OR 1.83)**



Remission rate at follow-up:

- 48.8% (combination)
- 51.9% (medication)
- 45.8% (CBT)



CAMELS REMISSION PREDICTORS

Male gender

Higher family functioning at
baseline

- Clear rules, more trust
- Higher-quality interactions by parent report



MEDICATION ADVERSE EFFECTS

SSRIS

CLOMIPRAMINE (CMI)

DULOXETINE (DUL)

SSRI ADVERSE EFFECTS

Sedation, insomnia, GI distress, rare diarrhea, tremor (especially sertraline)

Suicidal thinking likely more common in depressed teens than anxious children

Activation in some, particularly young children

Reduced platelet aggregation via 5-HT_{2A} platelet receptors

More nosebleeds, bruises, rare GI bleed

Recommend patients be cautious with NSAIDs and ASA

CMI ADVERSE EFFECTS

A tricyclic (SER and NE reuptake inhibitor)

Primarily metabolized by CYP 2D6
(FLV increases level)

High blood levels are toxic

Prolongs QTc → torsades des pointes

Anticholinergic
(dry mouth, constipation, urinary retention)

Antihistaminergic
(weight gain, somnolence)

CMI MONITORING



BASELINE EKG

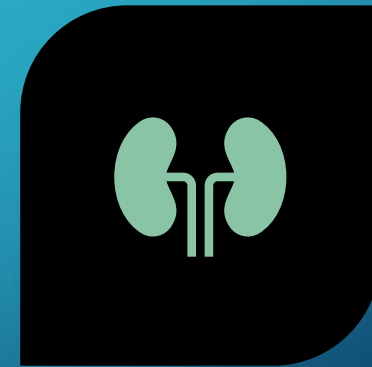
KEEP QTC BELOW 440 MSEC

AT LEAST ANNUAL EKGs



STARTING DOSE 1
MG/KG/DAY OR LESS

MAX 3 MG/KG/DAY



FOLLOW BLOOD LEVELS
FOR TOXICITY

DUL ADVERSE EFFECTS

SNRI (like venlafaxine)

More noradrenergic at high doses

More serotonergic at low doses

Headache
Abdominal pain, constipation
Somnolence
Loss of appetite
Sweating
Perhaps mild increase in BP

Typically started in 20-30 mg range
Peak dose usually 60-90



THAT'S ALL FOR NOW! BE SAFE OUT THERE!
HAPPY ST. PATRICK'S DAY
FROM OUR SOCIAL DISTANCING CENTER TO YOURS!