Weight Gain and Metabolic Syndrome in Children Exposed to Second-Generation Antipsychotic Medications



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BACKGROUND

- Second-generation antipsychotic medications (SGA) are associated with adverse metabolic effects
- National guidelines recommend screening after SGA initiation
- Guideline adherence within pediatric primary care is uncertain

OBJECTIVES

- Determine the rate of screening and detection rates of metabolic syndrome after SGA initiation
- Compare changes in weight among SGA exposed children unexposed controls

DATA SOURCE & METHODS

Data Source

- Electronic health record network of networks or super-network (CER^2)
- Two-hundred twenty-two U.S. pediatric primary care practic

Study Population & Exposure Status

- Children ages 3-18 years with minimum 180 days observati time between 2000 and 2016
- SGA exposure was determined by the presence of at least one SGA prescription

Primary Outcomes

- Metabolic syndrome, based on modified WHO definition
 - Obesity
 - 2. Hypertension
 - Dyslipidemia 3.
 - Elevated glucose level/Diabetes Mellitus

Analysis

- Rates of metabolic syndrome screening and detection were summarized using descriptive statistics
- Mean change (\pm SD) from baseline weight at 3, 6 & 12 months after SGA initiation were compared to propensityscore matched unexposed children
- Separate analyses were performed by age at initiation (<11y vs ≥11y)



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	RESULTS	: SGA-EXPOSED	
	Table 1: Demographics		
ion tain		Eligible subjects w (N= 12,9	
	Age at index SGA		
	3 – 5 years	884 (7	
	6 – 12 years	6297 (4	
	13 – 18 years	5730 (4	
lO	Male gender	8158 (6	
	Race		
	African-American	3980 (3	
	American Indian/Alaska native	35 (0.3	
/ork	Asian/Pacific Islander	98 (0.8	
	Caucasian	6427 (5	
ces	Mixed	174 (1	
	Unknown	2197 (1	
_	Ethnicity		
tion	Hispanic	1047 (8	
	Non-Hispanic	10,573 (8	
	Unknown	1291 (1	

RESULTS: METABOLIC SYNDROME SCREENING AND DETECTION



tested (360/2145 [17%], p<0.001)

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COHORT DESCRIPTION			
vith ≥1 SGA Rx 911) 7%)	 Baseline MH Diagnoses ADD 43% Non-OCD Anxiety disorder 23% Depression 15% Psychotic Disorders 13% PDD/Autism 12% 		
49%)	Concurrent Medications	Baselin	
14%) 33%)	 Stimulants 41% Antidepressants 32% 	Weight	
,0,10)	 Anti-anxiety 4% 	3 mon	
81%)		6 mon	
3%)	Index SGA and Treatment	12 mc	
3%)	Patterns	FEMAL	
50%)	 Index SGA Risperidone 50% 		
%)	 Aripiprazole 30% 		
7%)	 Quetiapine 14% 		
	Other SGA 6%	Baselin	
8%)	 67% children had ≥1 SGA Rx Modion SCA duration was 1.1 	Weight	
82%)	vears (IQR $0.38 - 2.31$)	3 mon	
0%)		6 mon	

Table 2: Metabolic Changes after SGA Initiation*

	Age at index SGA	
	Age <11 yrs	Age ≥11 yrs
	N = 5,216	N = 7,695
ty	974 (27%)	1175 (31%)
tension	612 (18%)	543 (15%)
mal cholesterol	86 (11%)	190 (18%)
mal glucose testing	82 (10%)	162 (12%)
iteria for metabolic	19 (5%)	45 (10%)
ome** (3 out of 4)		

* Follow-up time was within 3 years, or 6 months following last SGAM prescription (whichever comes earlier)

** Based on the World Health Organization (WHO) criteria for metabolic syndrome; percentages reflect those with follow-up measurements only

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RESULTS: WEIGHT CHANGE

Table 3: Short-Term Weight Change Age at index SGA Age <11 years Age ≥11 years ≥1 SGA Rx No SGA ≥1 SGA Rx No SGA N= 1,556** N= 5,598 N= 1,707** N= 6,317 ne weight (kg) 62.7 ± 21.3 62.7 ± 20.5 30.9 ± 10.6 | 28.6 ± 10.9 change at: 1.5 ± 2.0 0.9 ± 1.3 2.6 ± 3.7 1.1 ± 2.6 nths 2.4 ± 2.8 2.7 ± 3.5 nths 1.8 ± 1.7 3.9 ± 5.0 7.2 ± 7.2 $\textbf{5.4} \pm \textbf{5.0}$ 4.6 ± 3.9 3.7 ± 2.5 onths .ES Age at index SGA Age <11 years Age ≥11 years ≥1 SGA Rx ≥1 SGA Rx Unexposed Unexposed N= 1,796 N= 542** N= 1,181** N= 5,041 31.5 ± 11.8 ne weight (kg) 29.8 ± 12.1 64.6 ± 20.1 60.6 ± 17.9 change at: 1.8 ± 3.5 0.7 ± 2.6 1.9 ± 2.2 1.0 ± 1.4 nths $\textbf{3.1} \pm \textbf{2.8}$ 2.1 ± 2.0 2.6 ± 5.9 1.4 ± 3.4 nths 4.2 ± 7.5 2.9 ± 4.6 5.4 ± 4.7 4.4 ± 2.9 12 months

*Mean values and ± standard deviations are shown

**Only subjects with baseline weight measured on the day of SGAM initiation are included

CONCLUSION

• Screening for metabolic syndrome in pediatric primary care after SGA initiation was low, despite accelerated weight gains

• Prospective studies are needed to better delineate metabolic changes and identify targets for invention including greater adherence to screening recommendations.

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