



2024–2025 Pediatrics Review Syllabus

Behavioral Medicine & Substance Use Disorder:

Page 98, Autism Spectrum Disorder (ASD)

<i>Text currently reads:</i>	<i>Text should read:</i>
<p>Autism Spectrum Disorder (ASD)</p> <ul style="list-style-type: none">• Affects 1:36 children<ul style="list-style-type: none">– Males ~ 4× more likely to be affected than females<ul style="list-style-type: none">• May be underdiagnosed in females– Increased incidence in Black and Hispanic children– Rate in siblings: 10–20%  <ul style="list-style-type: none">• ~ 50% of affected children have associated intellectual disability• More common with certain genetic conditions<ul style="list-style-type: none">– Tuberous sclerosis– Fragile X syndrome– Angelman syndrome– Rett syndrome– Noonan syndrome– Trisomy 2– Neurofibromatosis 1– CHARGE syndrome– DiGeorge syndrome– Untreated phenylketonuria	<p>Autism Spectrum Disorder (ASD)</p> <ul style="list-style-type: none">• Affects 1:36 children<ul style="list-style-type: none">– Males ~ 4× more likely to be affected than females<ul style="list-style-type: none">• May be underdiagnosed in females– Increased incidence in Black and Hispanic children– Rate in siblings: 10–20%  <ul style="list-style-type: none">• ~ 50% of affected children have associated intellectual disability• More common with certain genetic conditions<ul style="list-style-type: none">– Tuberous sclerosis– Fragile X syndrome– Angelman syndrome– Rett syndrome– Noonan syndrome– Trisomy 21– Neurofibromatosis 1– CHARGE syndrome– DiGeorge syndrome– Untreated phenylketonuria

Metabolic Disorders:**Page 400, Audience Response Answers and Explanatory Information**

<i>Text currently reads:</i>	<i>Text should read:</i>
<p>AR 5 E. Scheduled oral cornstarch supplementation</p> <p>Answer: E. Scheduled oral cornstarch supplementation</p> <ul style="list-style-type: none">• This is a classic presentation for a GSD; the lab values are consistent with the elevated lipids and lactate, as well as the enlarged liver• The problem is an inability to break down glycogen to glucose; so, provide the body with a slow-releasing form of glycogen via cornstarch• Fatty acids and protein supplementation will not address the need for glycogen; glucagon will force the body to release what glycogen it can, but then will get low glucose again• Repeat IV dextrose does not address the need for ability to live independently and may force the body to store more glycogen that it is not using	<p>AR 5 E. Scheduled oral cornstarch supplementation</p> <p>Answer: E. Scheduled oral cornstarch supplementation</p> <ul style="list-style-type: none">• This is a classic presentation for a GSD; the lab values are consistent with the elevated lipids and lactate, as well as the enlarged liver• The problem is an inability to break down glycogen to glucose; so, provide the body with a slow-releasing form of glucose via cornstarch• Fatty acids and protein supplementation will not address the need for glycogen; glucagon will force the body to release what glycogen it can, but then will get low glucose again• Repeat IV dextrose does not address the need for ability to live independently and may force the body to store more glycogen that it is not using