

# EEG Patterns in Epilepsy Syndromes

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# Virtual EEG course 2022

18<sup>TH</sup> SEPTEMBER



สมาคมโรคสมองชักแห่งประเทศไทย

TIME	SCHEDULE	SPEAKERS
08.20-08.30 u.	Opening remarks	พลตรี นพ.ชาครินทร์ ณ บางช้าง
08.30-09.00 u.	Normal awake EEG	อ.นพ.ศรित्रาวุธ วงษ์เวียงจันทร์
09.00-09.30 u.	Normal sleep EEG	อ.พญ.ชูปณี สมบูรณ์
09.30-10.00 u.	Normal variants	ผศ.นพ.อิริวัฒน์ สุนทรพันธ์
10.00-10.30 u.	How to localize waveforms on the EEG?	รศ.พญ.กนกวรรณ บุญญพิสิฏฐ์
10.30-11.00 u.	Artifacts	อ.นพ.กุลเสถียร ศักดิ์พิชัยสกุล
11.00-11.30 u.	Non-epileptiform abnormalities	อ.พญ.สุดา จิรสกุลเดช
11.30-12.30 u.	Lunch	
12.30-13.00 u.	Epileptiform abnormalities	ผศ.พญ.ณัฐรุจ วิศวกรรมดิษฐกุล
13.00-13.30 u.	Interictal and ictal patterns in generalized seizures	รศ.นพ.ชัยยศ คงคศิริสม
13.30-14.00 u.	Interictal and ictal patterns in focal seizures	พ.อ.หญิง ผศ.ภริณี สุวรรณภักดี
14.00-14.30 u.	EEG patterns in epilepsy syndromes	ผศ.พญ.กมลวรรณ กตัญญูวงศ์
14.30-15.00 u.	EEG patterns in encephalopathy and coma	ผศ.นพ.อภิสิทธิ์ บุญเกิด
15.00-15.30 u.	EEG patterns in status epilepticus	อ.นพ.พิพาทร ตุ่มนาค
15.30-16.00 u.	How to write the EEG report?	ผศ.นพ.ชูศักดิ์ ลิ้มภัย
16.00-16.30 u.	Post-test	ผศ.พญ.กมลวรรณ กตัญญูวงศ์

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# Outline

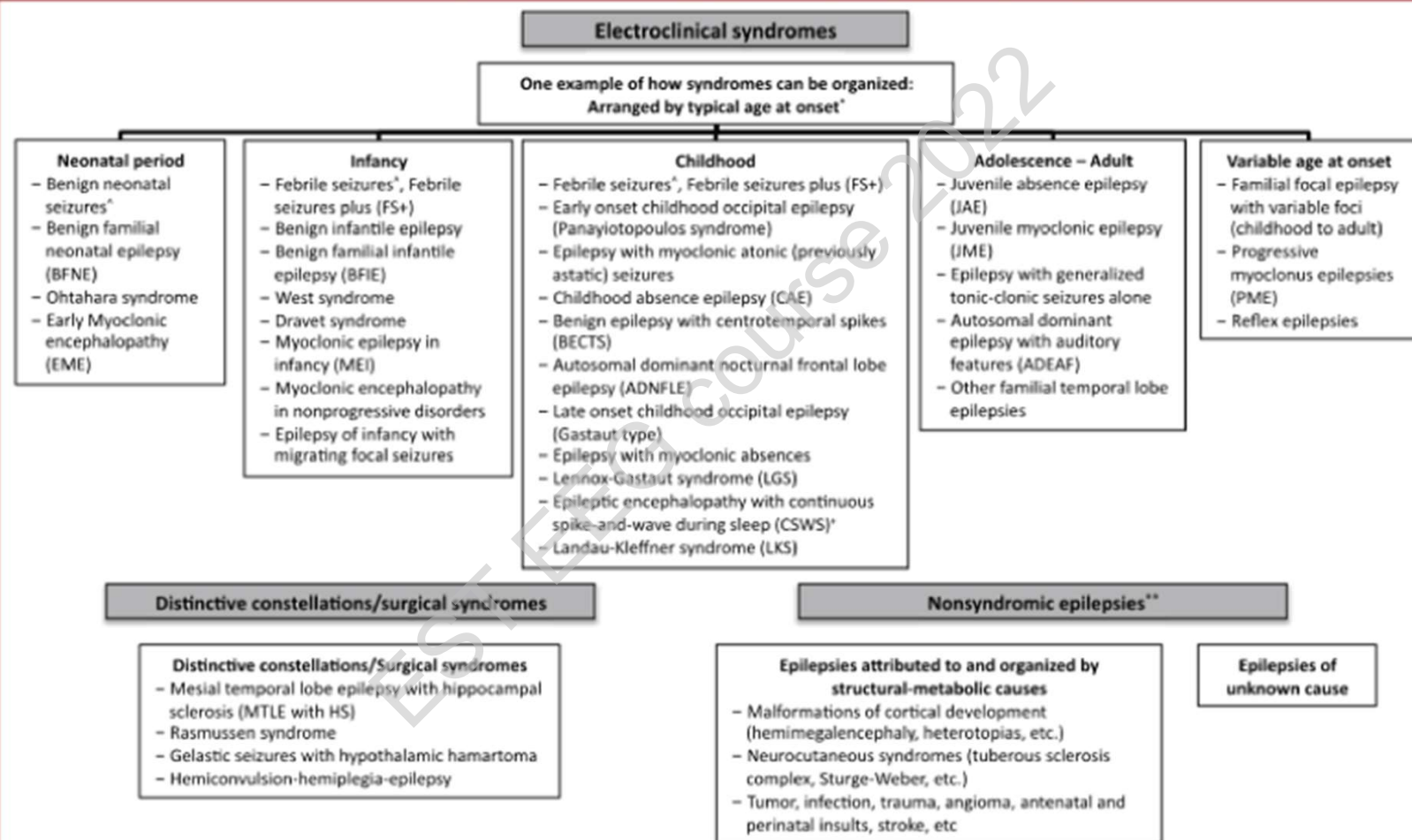
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- Important EEG information/pattern in some epilepsy syndromes

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# ILAE Proposal for Revised Terminology for Organization of Seizures and Epilepsies 2010

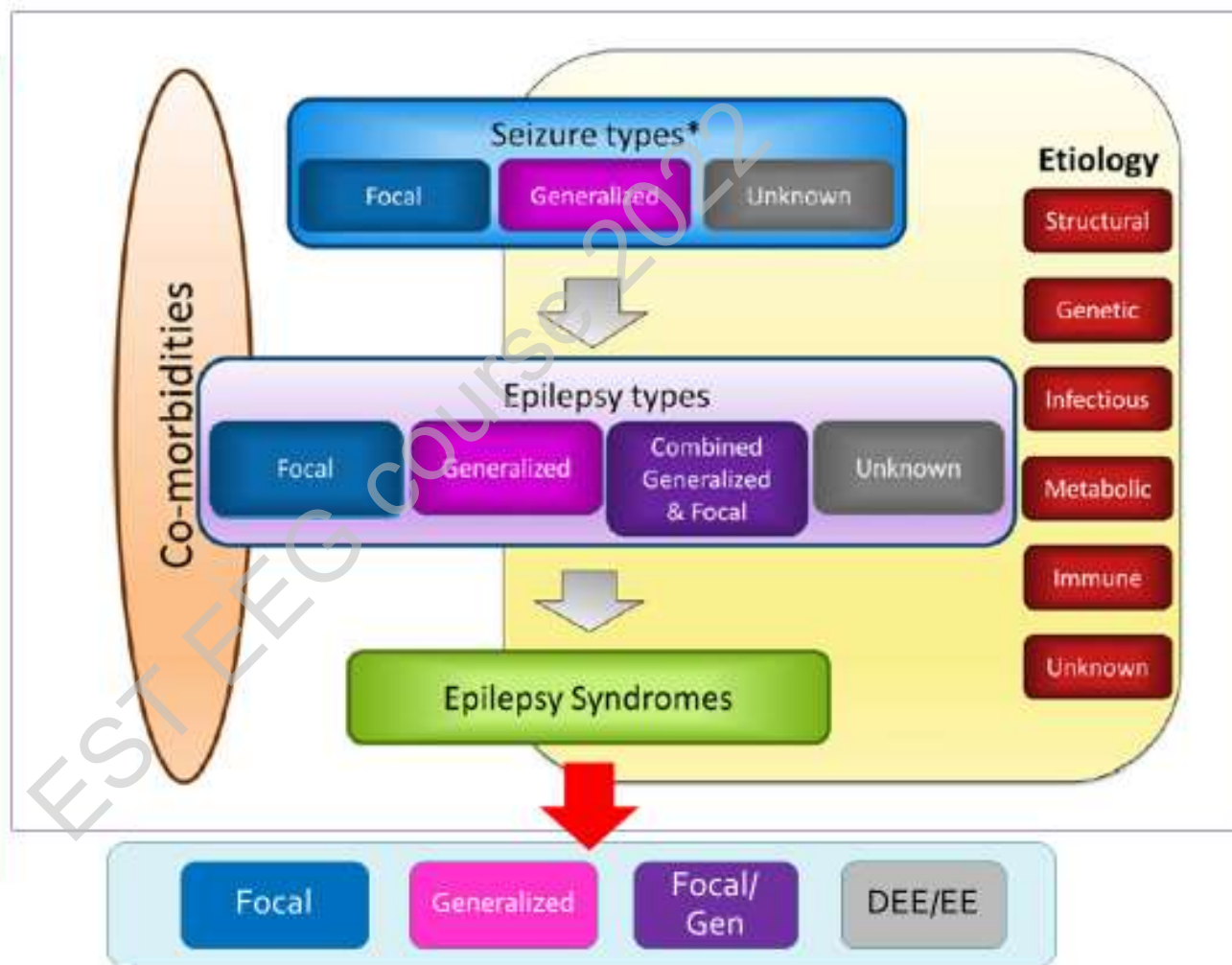
## Electroclinical Syndromes and Other Epilepsies Grouped by Specificity of Diagnosis



# Epilepsy Classification 2017

+

## Epilepsy syndromes 2022



## ILAE classification and definition of epilepsy syndromes with onset in neonates and infants: Position statement by the ILAE Task Force on Nosology and Definitions

Sameer M. Zuberi<sup>1</sup> | Elaine Wirrell<sup>2</sup> | Elissa Yozawitz<sup>3</sup> | Jo M. Wilmschurst<sup>4</sup> | Nicola Specchio<sup>5</sup> | Kate Riney<sup>6,7</sup> | Ronit Pressler<sup>8,9</sup> | Stephane Auvin<sup>10</sup> | Pauline Samia<sup>11</sup> | Edouard Hirsch<sup>12</sup> | Santiago Galicchio<sup>13</sup> | Chahnez Triki<sup>14</sup> | O. Carter Snead<sup>15</sup> | Samuel Wiebe<sup>16</sup> | J. Helen Cross<sup>17,18</sup> | Paolo Tinuper<sup>19,20</sup> | Ingrid E. Scheffer<sup>21</sup> | Emilio Perucca<sup>22,23</sup> | Solomon L. Moshé<sup>24,25,26</sup> | Rima Nabbout<sup>27</sup>

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DOI: 10.1111/epl.17236

## ILAE definition of the Idiopathic Generalized Epilepsy Syndromes: Position statement by the ILAE Task Force on Nosology and Definitions

Edouard Hirsch<sup>1</sup> | Jacqueline French<sup>2</sup> | Ingrid E. Scheffer<sup>3</sup> | Alicia Bogacz<sup>4</sup> | Taoufik Alsaadi<sup>5</sup> | Michael R. Sperling<sup>6</sup> | Fatema Abdulla<sup>7</sup> | Sameer M. Zuberi<sup>8</sup> | Eugen Trinka<sup>9,10</sup> | Nicola Specchio<sup>11</sup> | Ernest Somerville<sup>12</sup> | Pauline Samia<sup>13</sup> | Kate Riney<sup>14,15</sup> | Rima Nabbout<sup>16</sup> | Satish Jain<sup>17</sup> | Jo M. Wilmschurst<sup>18</sup> | Stephane Auvin<sup>19,20</sup> | Samuel Wiebe<sup>21</sup> | Emilio Perucca<sup>22,23</sup> | Solomon L. Moshé<sup>24</sup> | Paolo Tinuper<sup>25,26</sup> | Elaine C. Wirrell<sup>27</sup>

Received: 24 April 2021 | Revised: 16 March 2022 | Accepted: 17 March 2022

DOI: 10.1111/epl.17241

## International League Against Epilepsy classification and definition of epilepsy syndromes with onset in childhood: Position paper by the ILAE Task Force on Nosology and Definitions

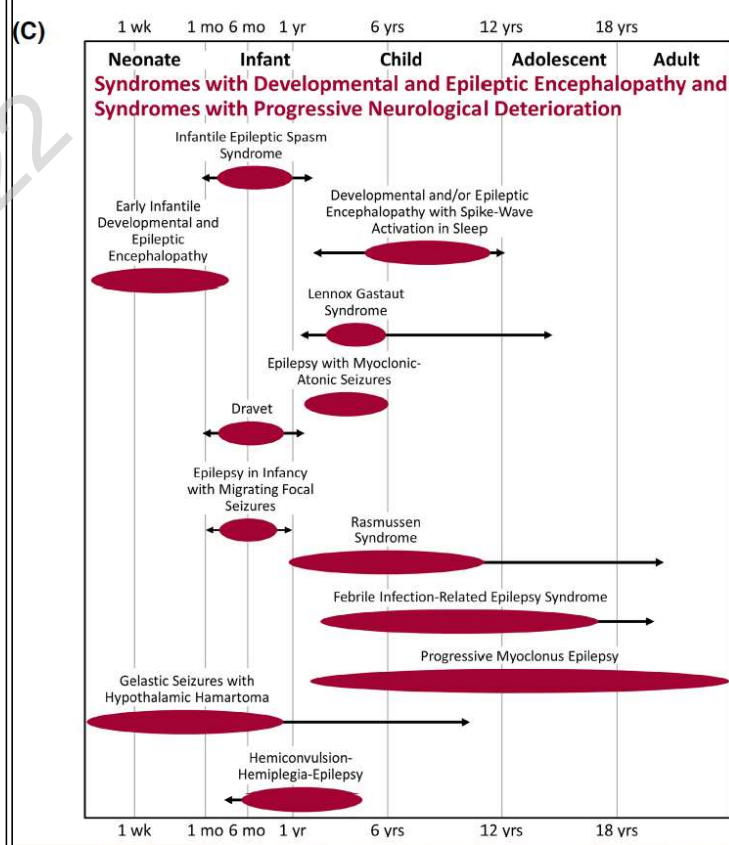
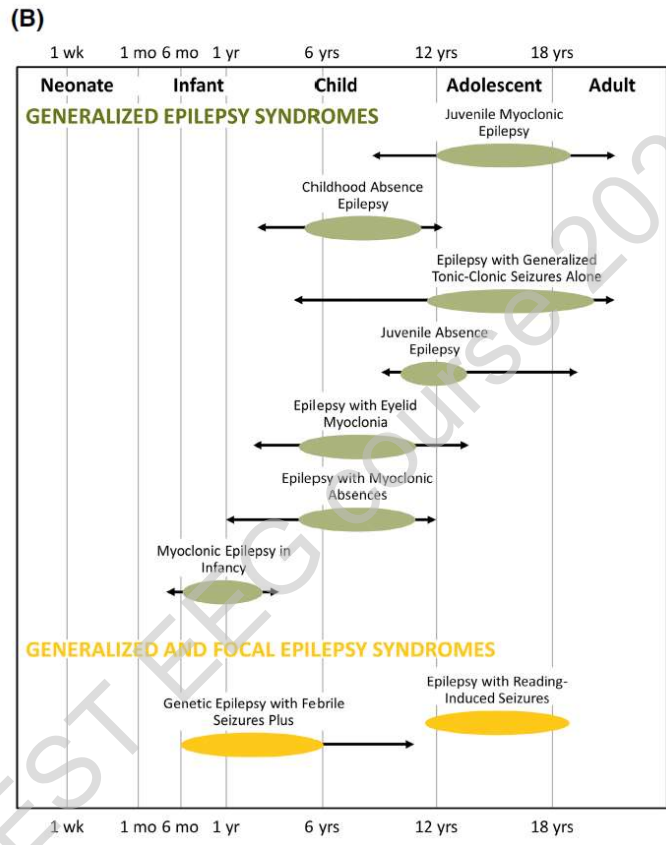
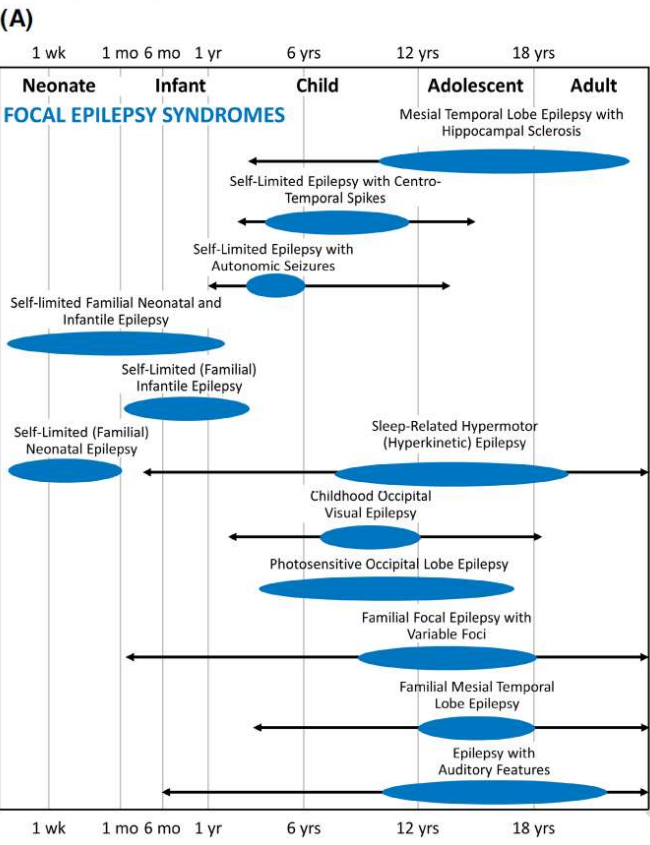
Nicola Specchio<sup>1</sup> | Elaine C. Wirrell<sup>2</sup> | Ingrid E. Scheffer<sup>3</sup> | Rima Nabbout<sup>4</sup> | Kate Riney<sup>5,6</sup> | Pauline Samia<sup>7</sup> | Marilisa Guerreiro<sup>8</sup> | Sam Gwer<sup>9</sup> | Sameer M. Zuberi<sup>10</sup> | Jo M. Wilmschurst<sup>11</sup> | Elissa Yozawitz<sup>12</sup> | Ronit Pressler<sup>13</sup> | Edouard Hirsch<sup>14</sup> | Sam Wiebe<sup>15</sup> | Helen J. Cross<sup>16</sup> | Emilio Perucca<sup>17,18</sup> | Solomon L. Moshé<sup>19</sup> | Paolo Tinuper<sup>20,21</sup> | Stéphane Auvin<sup>22</sup>

Received: 23 April 2021 | Revised: 12 March 2022 | Accepted: 16 March 2022

DOI: 10.1111/epl.17240

## International League Against Epilepsy classification and definition of epilepsy syndromes with onset at a variable age: position statement by the ILAE Task Force on Nosology and Definitions

Kate Riney<sup>1,2</sup> | Alicia Bogacz<sup>3</sup> | Ernest Somerville<sup>4,5</sup> | Edouard Hirsch<sup>6,7,8</sup> | Rima Nabbout<sup>9,10,11</sup> | Ingrid E. Scheffer<sup>12</sup> | Sameer M. Zuberi<sup>11,13,14</sup> | Taoufik Alsaadi<sup>15</sup> | Satish Jain<sup>16</sup> | Jacqueline French<sup>17</sup> | Nicola Specchio<sup>18</sup> | Eugen Trinka<sup>19,20,21</sup> | Samuel Wiebe<sup>22</sup> | Stéphane Auvin<sup>23,24,25</sup> | Leonor Cabral-Lim<sup>26</sup> | Ansuya Naidoo<sup>27,28</sup> | Emilio Perucca<sup>29,30</sup> | Solomon L. Moshé<sup>31,32</sup> | Elaine C. Wirrell<sup>33</sup> | Paolo Tinuper<sup>34,35</sup>





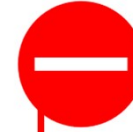
## Mandatory

- Criteria that must be present to diagnose the syndrome



## Alerts

- Absent criteria in vast majority of patients but rarely can be seen
- Rethink the diagnosis
- R/O other conditions



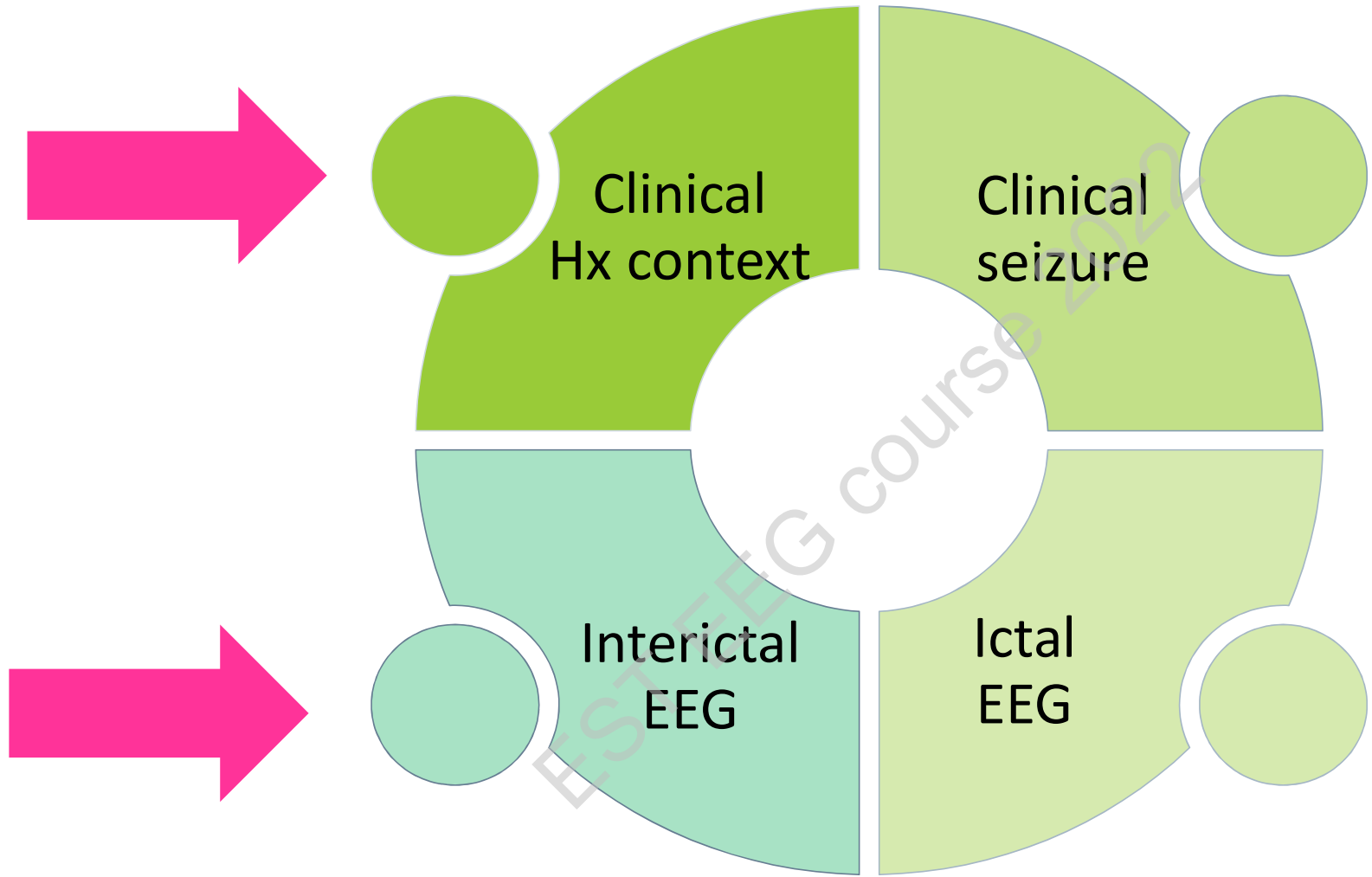
## Exclusionary

- Criteria that must be absent to diagnose the syndrome



# Syndrome Core Diagnostic Criteria

	Mandatory	Alert	Exclusionary
Seizure type			
<b>EEG</b> ( <i>interictal</i> )	<input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
Age at onset			
Development at onset			
Neurological exam			
Are MRI or <b>ictal EEG</b> required For Dx			
Other studies-genetic			
Syndrome without laboratory confirmation			



# EEG Diagnostic Criteria in epileptic syndrome


	Mandatory	Alert	Exclusionary
<b>EEG</b> ( <i>interictal</i> )			
<b>Ictal EEG</b> required For Dx			

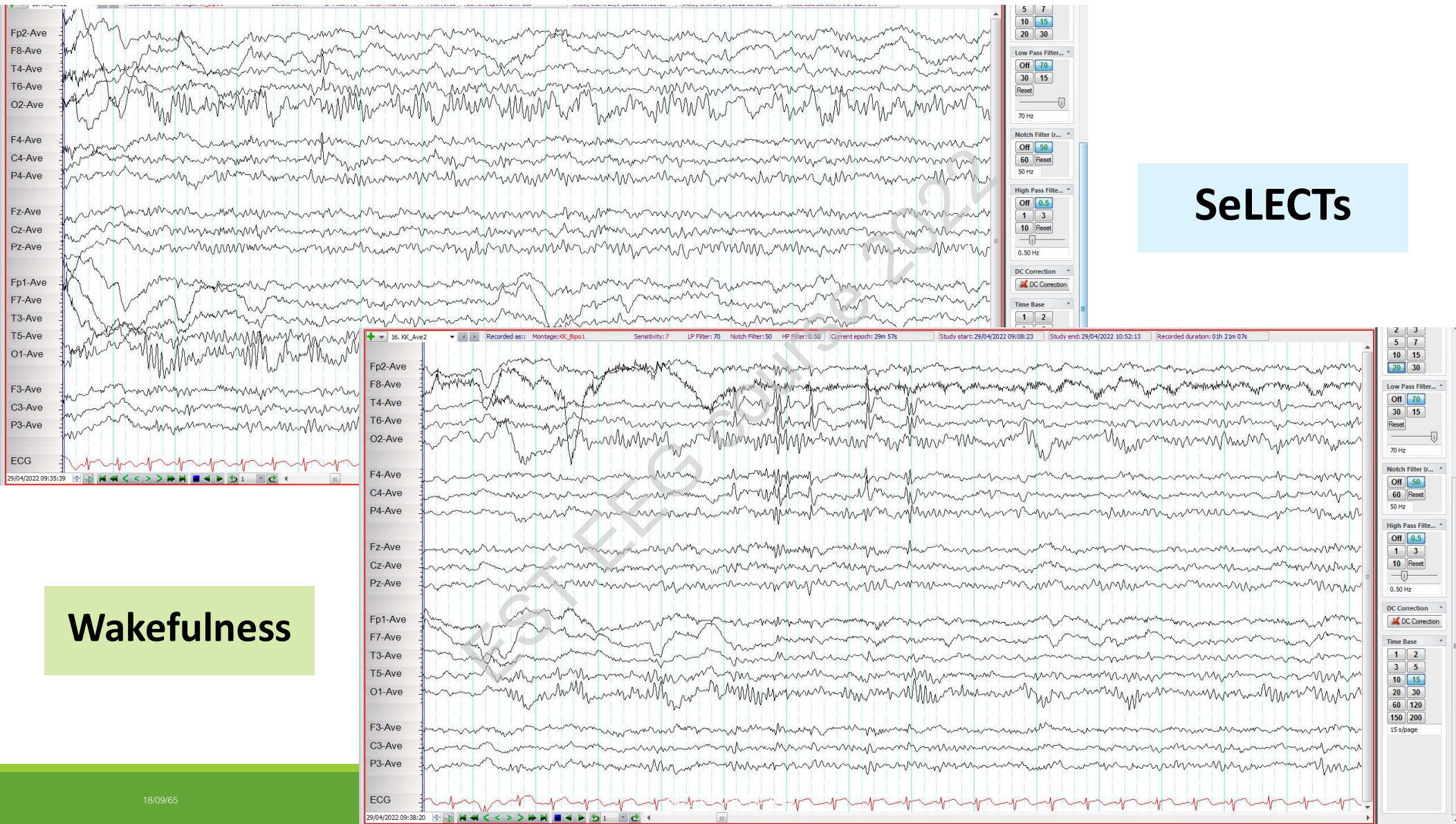
# Self-limited epilepsy with centrotemporal spikes (SeLECTs)

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- Previous : BRE-Benign Rolandic Epilepsy  
: BECTS-Benign epilepsy with centrotemporal spikes

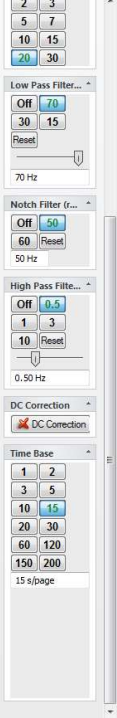
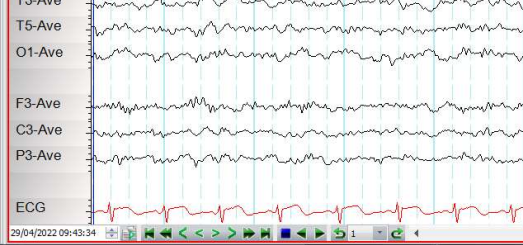
# SeLECTs EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
EEG ( <i>interictal</i> )	<p>High amplitude C-T biphasic epileptiform abnormality</p>	<ul style="list-style-type: none"> <li>Sustained focal slowing not limited to postictal</li> <li>Persistently unilat C-T SW on serial EEG</li> <li>Lack of sleep activation of CT SW</li> </ul>	
<b>Ictal EEG</b> required For Dx	<p><b>No need for Ictal EEG</b></p> 		





**SeLECTs**



**Sleep**

# SeLECTs



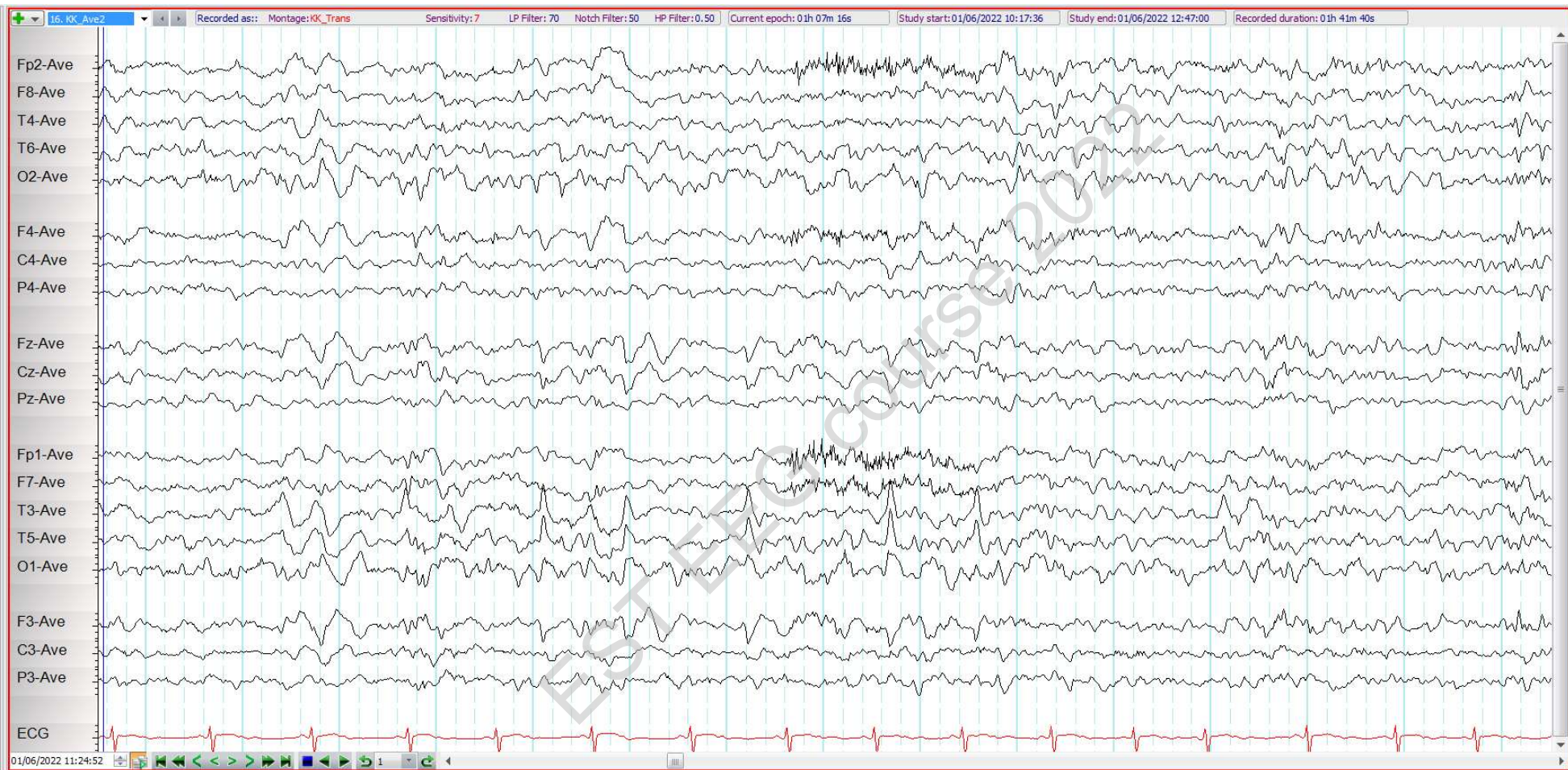


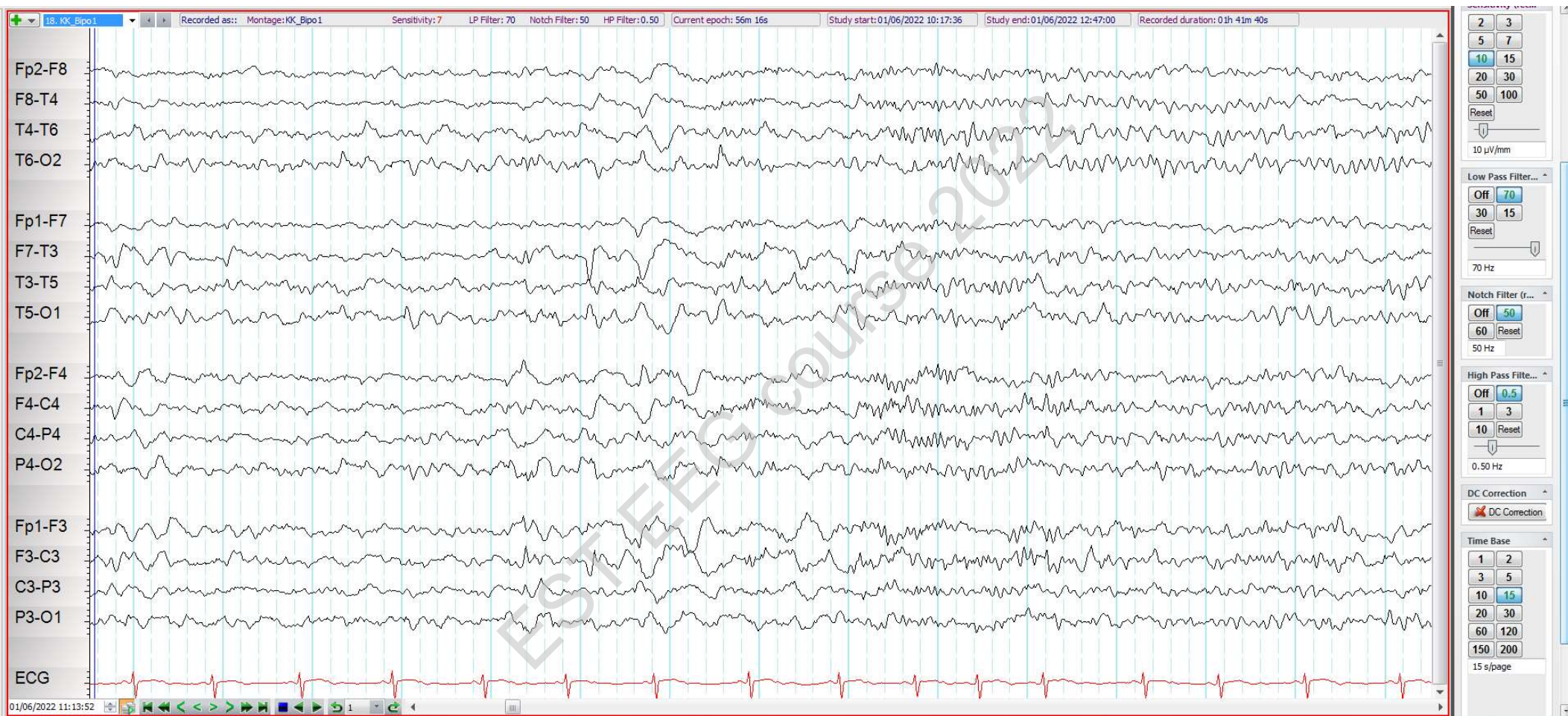
# SeLECTs exclusion

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- R/O by atypical history context, EEG exclusion, imaging
- Imaging: R/O structural lesion, FCD etc
- Example of the 3 previous pages is EEG of patient who has left hemispheric encephalomalacia







# Sleep-related hypermotor (hyperkinetic) epilepsy (SHE)

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# SHE EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
<b>EEG</b> ( <i>interictal</i> )		<ul style="list-style-type: none"><li>Fq epileptiform d/c outside F area</li></ul>	
<b>Ictal EEG</b> required For Dx	Not required		

# GGE

## Genetic Generalized Epilepsies

### Idiopathic Generalized Epilepsies

Childhood  
Absence  
Epilepsy  
*CAE*

Juvenile  
Absence  
Epilepsy  
*JAE*

Epilepsy with  
Generalized  
Tonic-Clonic  
Seizures Alone  
*GTCA*

Juvenile  
Myoclonic  
Epilepsy  
*JME*

### Epileptic Encephalopathy

Epilepsy with  
Myoclonic-Atonic Seizures  
*EMaTS*

### Developmental and Epileptic Encephalopathy

Epilepsy with  
Eyelid Myoclonia  
*EEM*

Epilepsy with  
Myoclonic Absences  
*EMA*

Myoclonic Epilepsy  
in Infancy  
*MEI*

### Developmental Encephalopathy

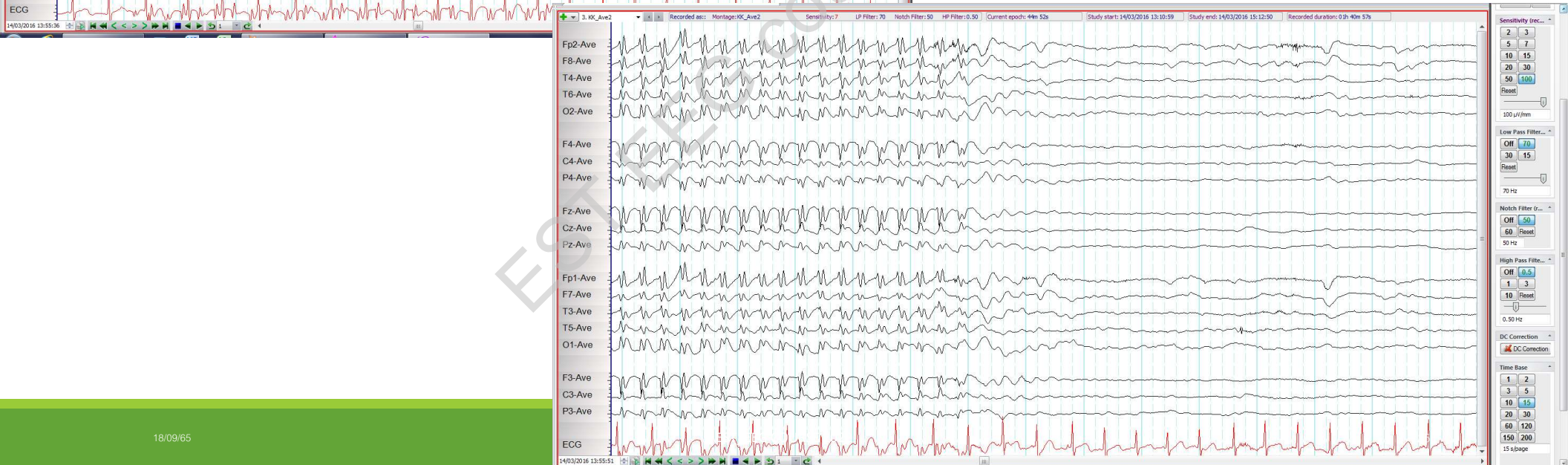
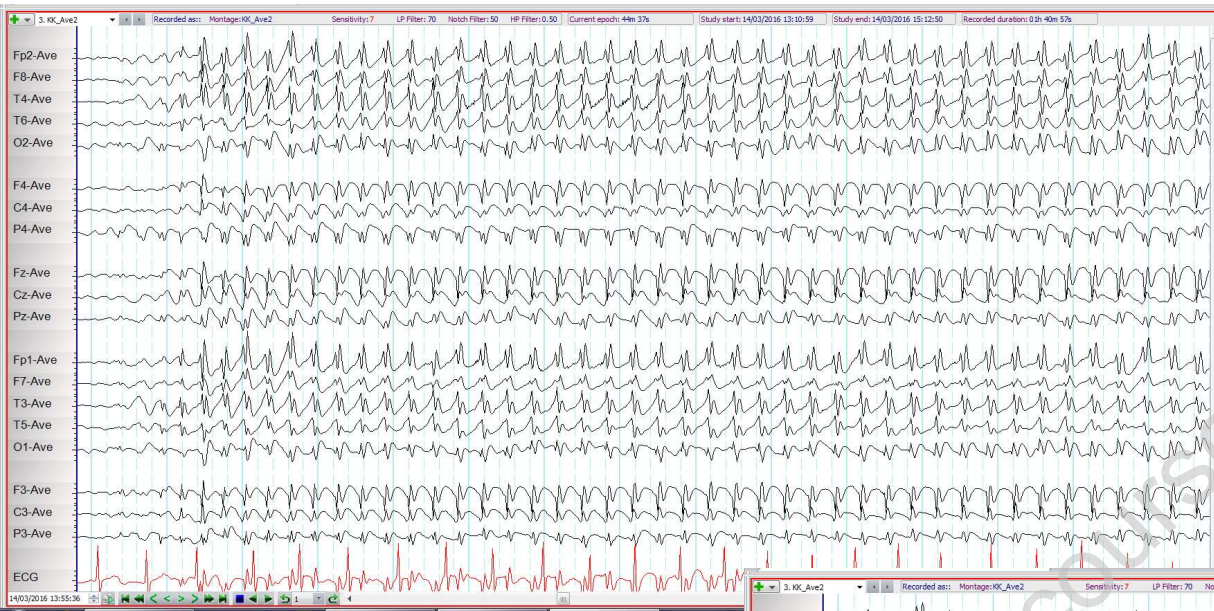
# CAE EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
<b>EEG</b> ( <i>interictal</i> )	Paroxysmal 3 (2.5-4) Hz gen S/W	<ul style="list-style-type: none"> <li>• Unilat d/c</li> <li>• No 3 Hz S/W d/c during HV in untreated pt</li> <li>• Persistent slowing B/G</li> <li>• Typical staring without EEG correlate</li> </ul>	Diffuse B/G slowing
<b>Background</b>	OIRDA 21%		< 2, > 13 yrs
<b>Ictal EEG</b> required For Dx	<b>No need</b> , but untreated pt will have a recorded absence sz on routine EEG.		





# Untreated CAE



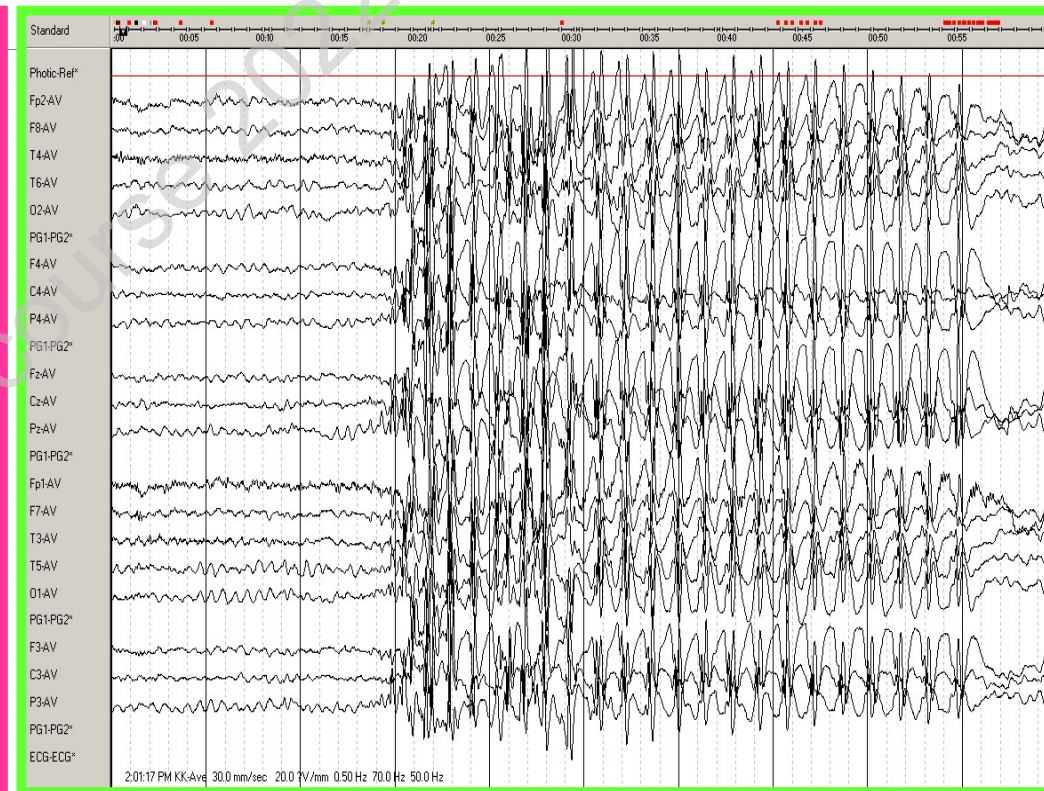
# JAE EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
<b>EEG</b> ( <i>interictal</i> )	Paroxysmal <b>3 -5.5</b> Hz gen S/W	<ul style="list-style-type: none"> <li>No 3-5.5 Hz S/W d/c during HV in untreated pt</li> <li>Persistent slowing B/G without sedative drug</li> </ul>	<ul style="list-style-type: none"> <li>Unilat focal d/c</li> <li>Diffuse b/g slowing</li> <li>Typical spell with no EEG correlate</li> </ul>
<b>Ictal EEG</b> required For Dx	<b>No need</b> , but untreated pt will have a recorded absence sz on routine EEG.		< 8, > 20 yrs



# Ictal EEG CAE vs. Ictal EEG JAE

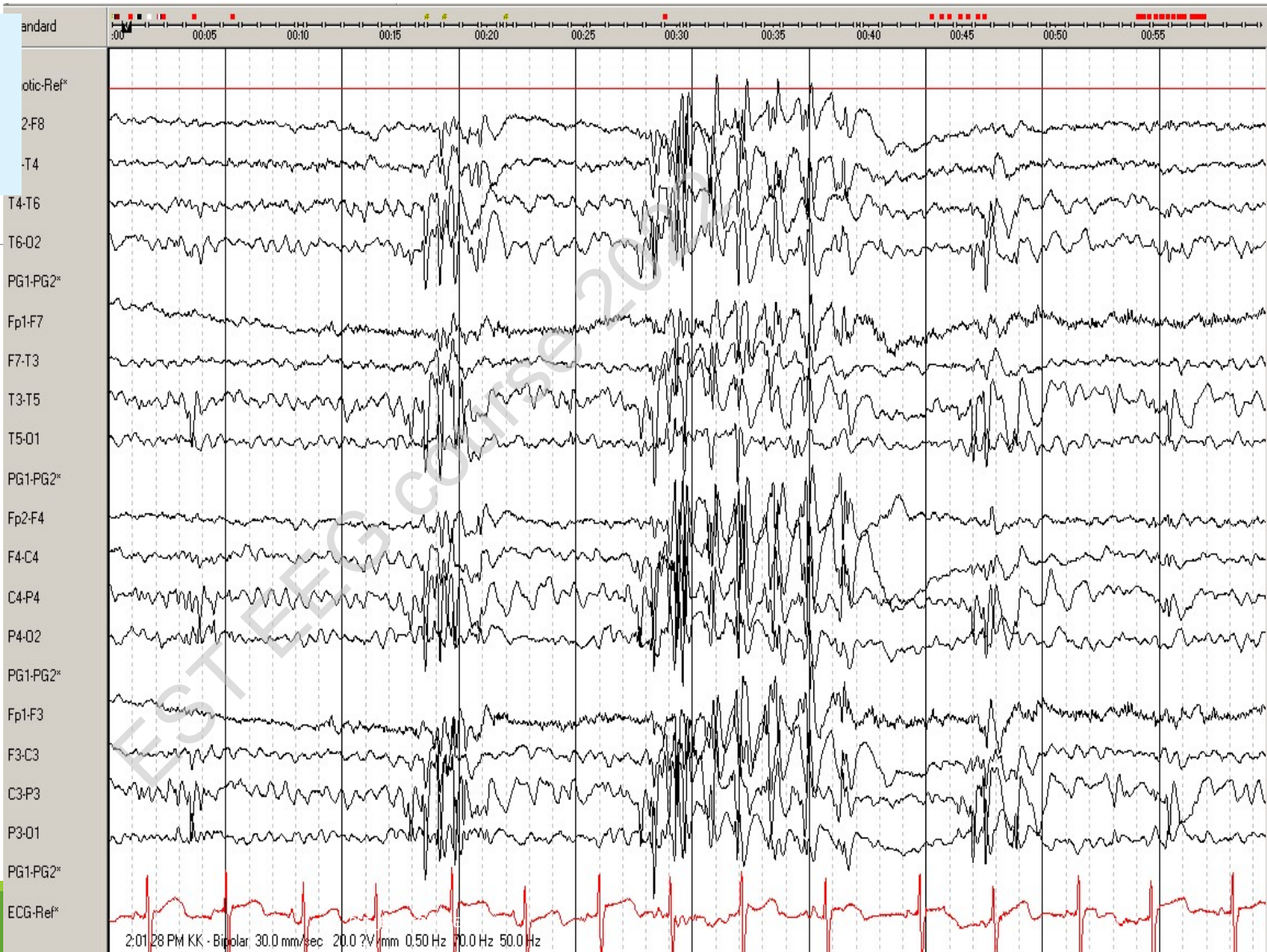
*More disorganized d/c in JAE*



# JME EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
<b>EEG</b> ( <i>interictal</i> ) <ul style="list-style-type: none"> <li>• 3 -5.5 Hz gen S/W</li> <li>• GPSW</li> </ul>			<ul style="list-style-type: none"> <li>• Gen slow SW &lt; 2.5 Hz</li> <li>• Unilat focal d/c</li> <li>• Focal slowing</li> <li>• Diffuse b/g slow that is not postictal</li> <li>• Myoclonic sz but no GPSW</li> </ul>
<b>Ictal EEG</b> required For Dx EST EEG course 2022	Not required		< 8, > 40 yrs

# JME in untreated pt



# EEG in treated JME

Normal

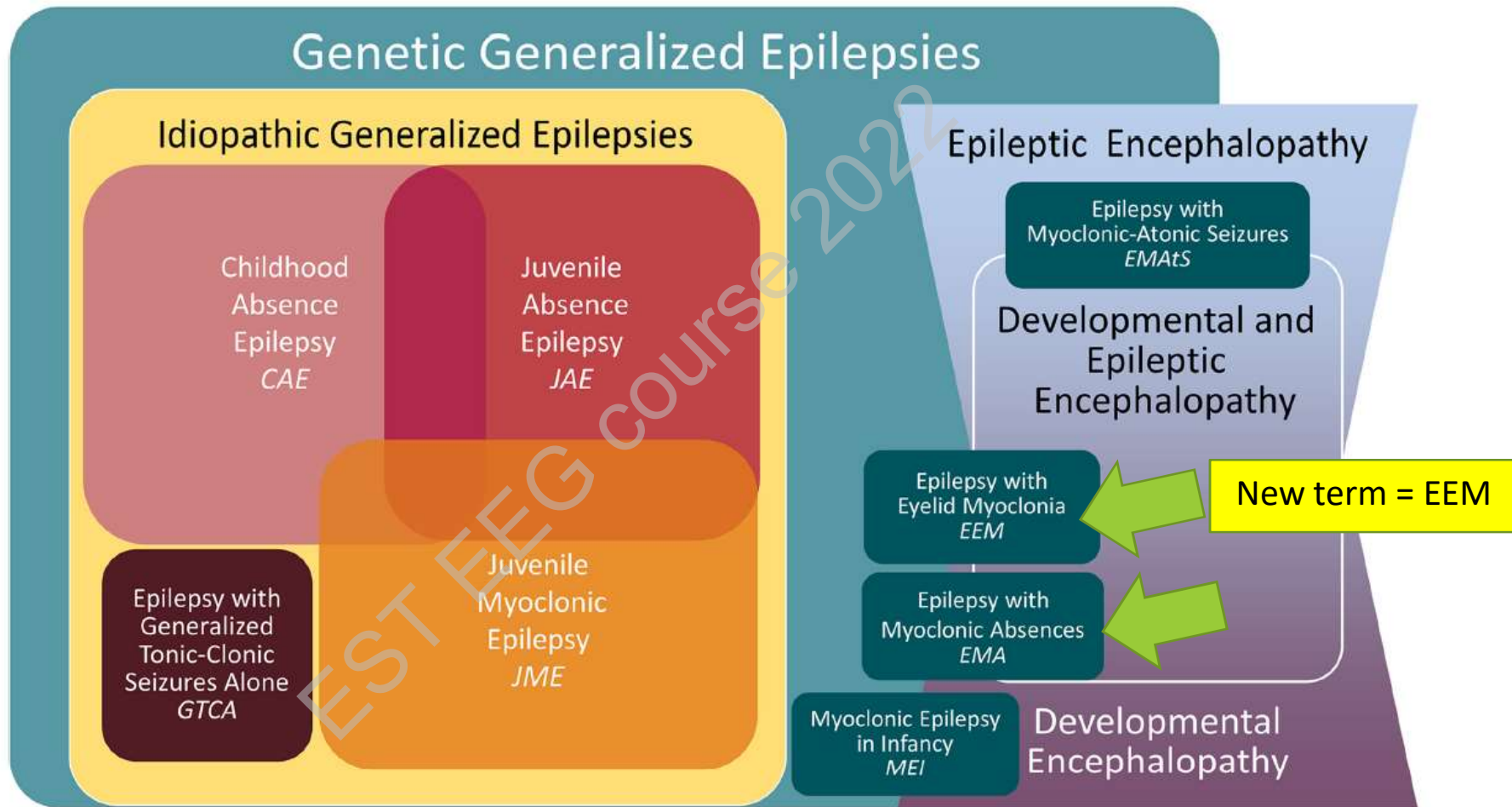


# EEG in treated JME

Isolated/subtle abnormal



# GGE



18/09/2022 Old term: Eyelid Myoclonia with Absence (EMA) but now EMA = epilepsy w myoclonic absence



# EEM EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
<b>EEG</b> ( <i>interictal</i> ) Eye closure and intermittent PS produces 3-6 Hz GPW, gen polyspikes			<ul style="list-style-type: none"> <li>• Gen slow SW &lt; 2.5 Hz</li> <li>• Unilat focal d/c</li> <li>• Focal slowing</li> <li>• Diffuse b/g slow that is not postictal</li> <li>• Lack of EEG correlate w typical event</li> </ul>
<b>Ictal EEG</b> required For Dx	Not required		< 2, > 14 yrs


# Infantile epileptic spasms syndrome (IESS)

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- Previous: West syndrome

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# IESS EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
<b>EEG</b> ( <i>interictal</i> )	<b>Interictal</b> <ul style="list-style-type: none"> <li>Hypsarrhythmia</li> <li>Multifocal/focal d/c (might be seen early after spasms)</li> </ul>	<b>Interictal</b> <ul style="list-style-type: none"> <li>Normal EEG</li> <li>SB pattern</li> </ul>	<b>Ictal</b> Normal EEG during clinical spasms
<b>Ictal EEG</b> required For Dx	<u>No</u> need for Ictal EEG. <u>If</u> no hypsarrhythmia or no epileptiform discharge – <b>need Ictal EEG.</b> 		

# Hypsarrhythmia pattern

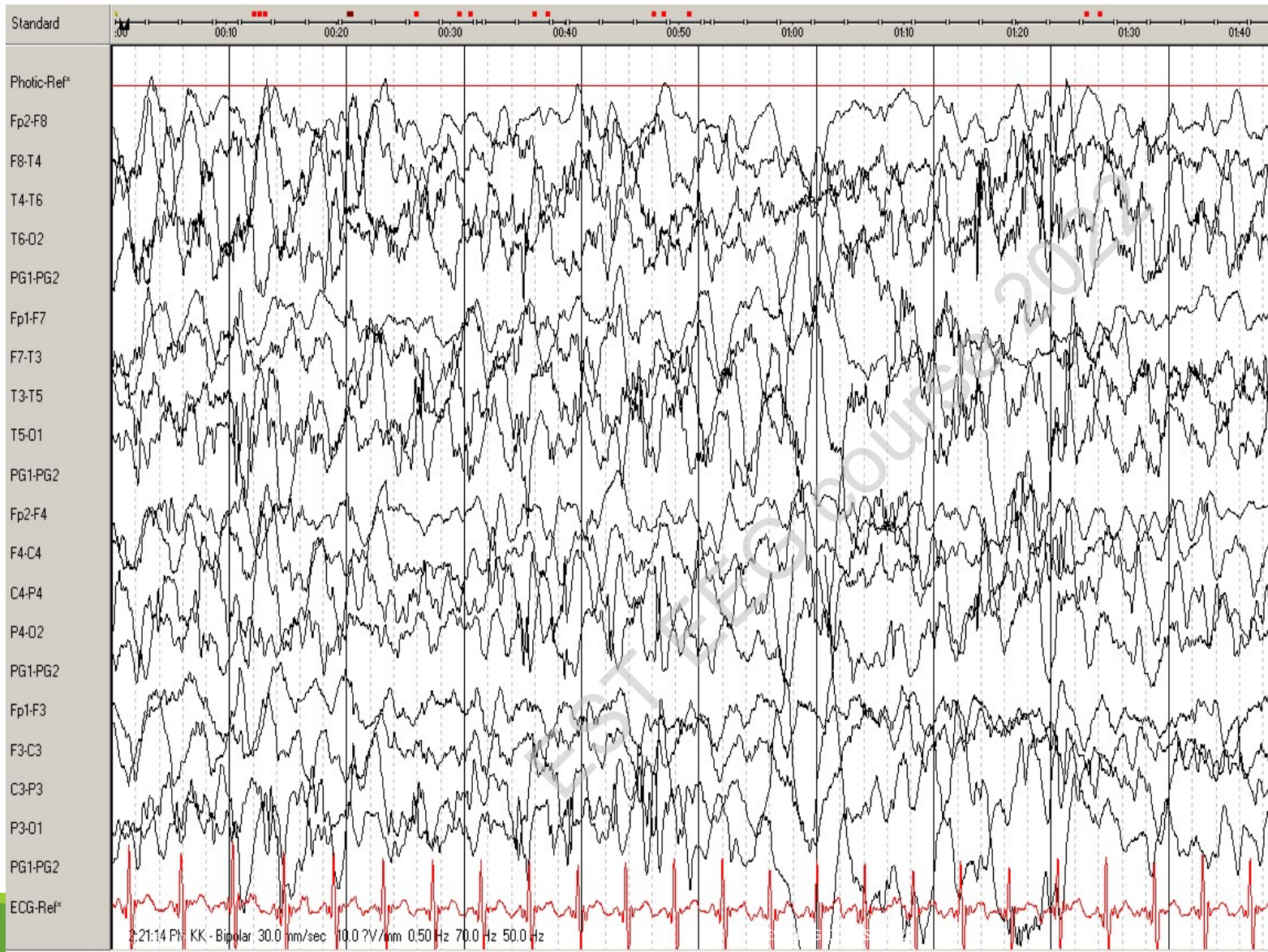
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- Abnormal interictal high amplitude waves and a background of irregular spikes
- Wakefulness: continuous high amplitude (> 200 Hz), gen polymorphic slowing
  - : no organized background
  - : multifocal spikes

# Modified Hypsarrhythmia

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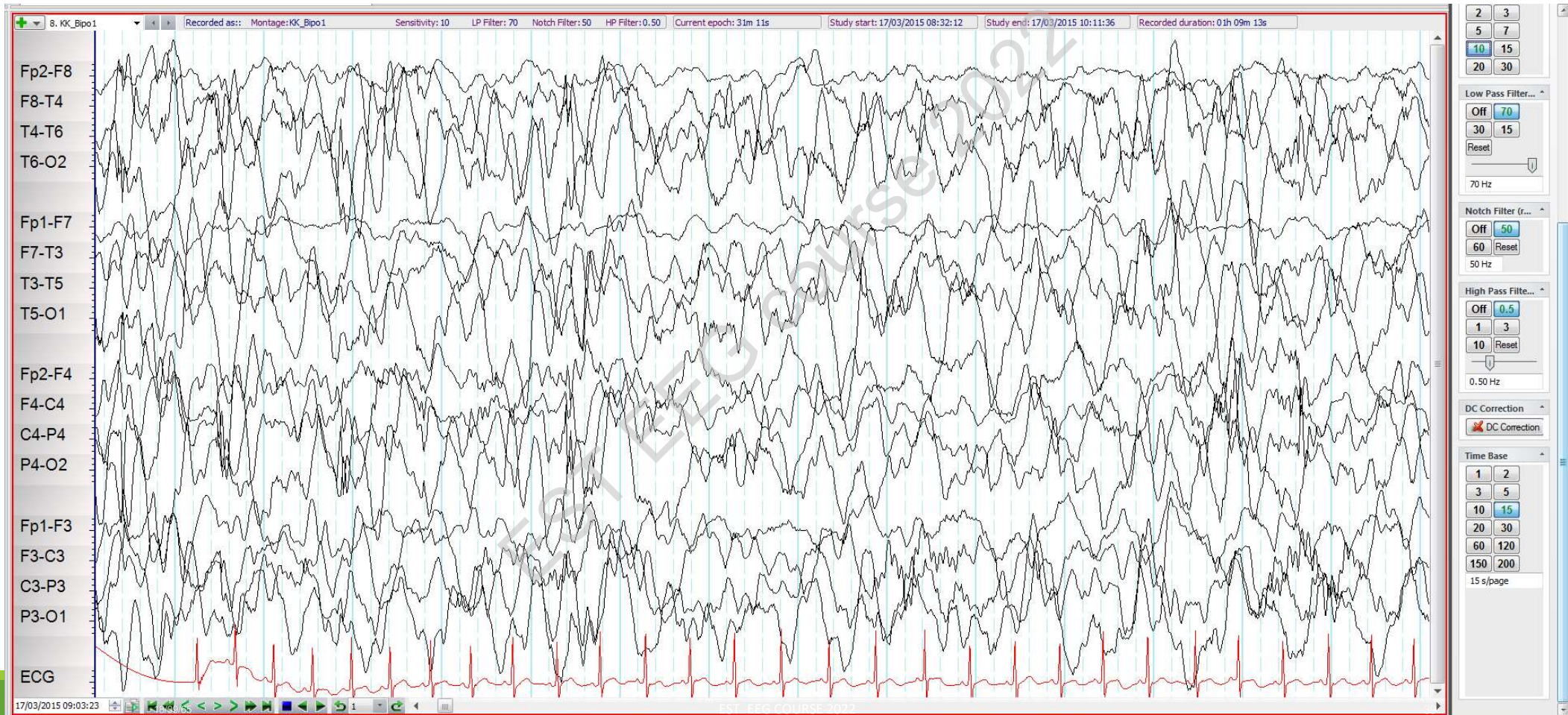
- Hypsarrhythmia with interhemispheric synchrony
- Hypsarrhythmia with voltage asymmetries
- Hypsarrhythmia with focus of abnormal discharges
- Hypsarrhythmia with episodes of attenuation
- Hypsarrhythmia consisting primarily of high voltage slow activity with few sharp waves or spikes



IESS  
or  
West

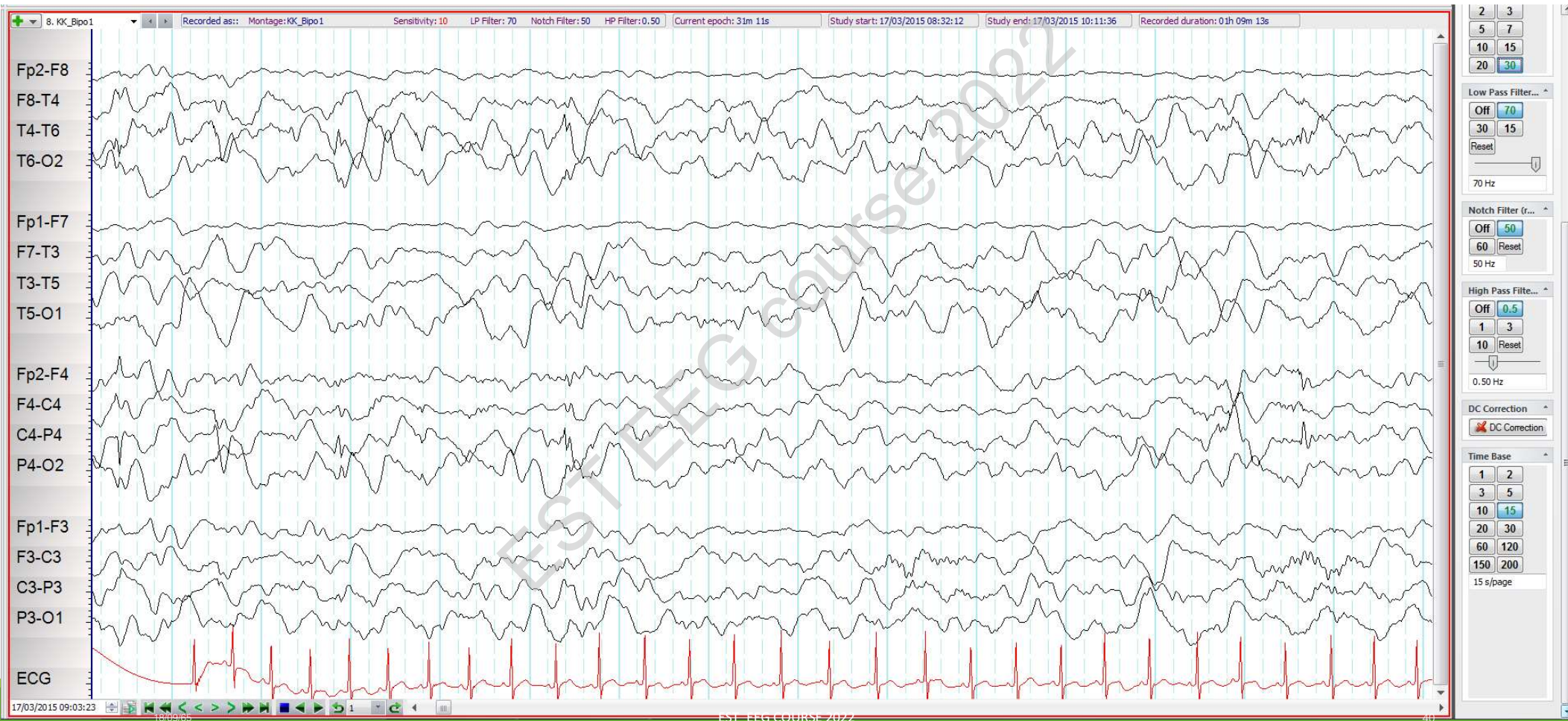
10  $\mu\text{v}/\text{mm}$

# Hypsarrhythmia-IESS



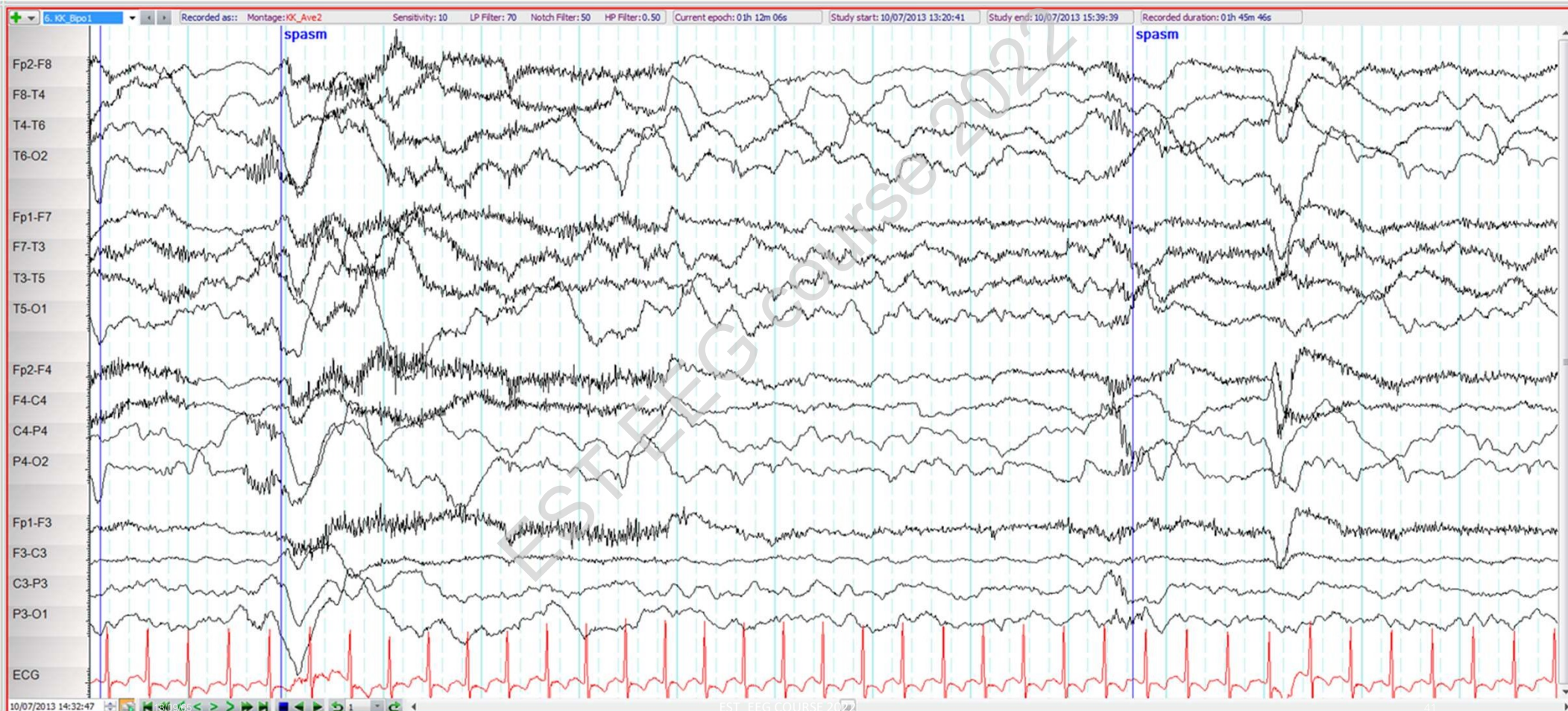
30 uv/mm

# Modified hypsarrhythmia-IESS





# Ictal EEG in IESS

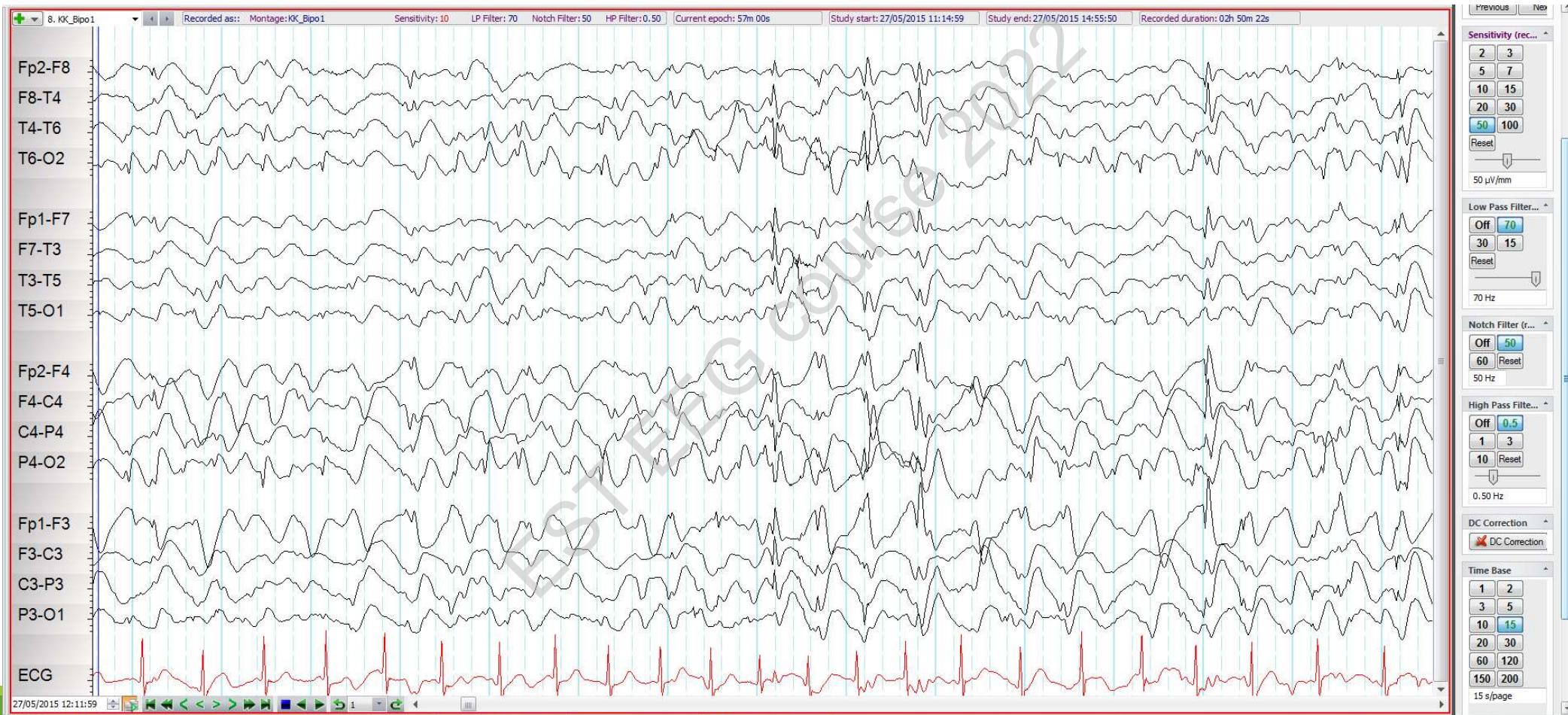


# LGS EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
<b>EEG</b> ( <i>interictal</i> )	<b>Interictal</b> <ul style="list-style-type: none"> <li>Gen slow &lt; 2.5 Hz SW complexes</li> <li>Gen PFA in sleep</li> </ul>	PPR at low fq (suggest CLN2)	Persistent focal abnormal without gen slow SW
<b>Ictal EEG</b> required For Dx	<u>Not</u> required. But if strongly consider myoclonic- atonic seizure syndrome.		



# LGS: EEG-bipolar montage



# LGS: EEG-average montage



# DEE-SWAS, EE-SWAS

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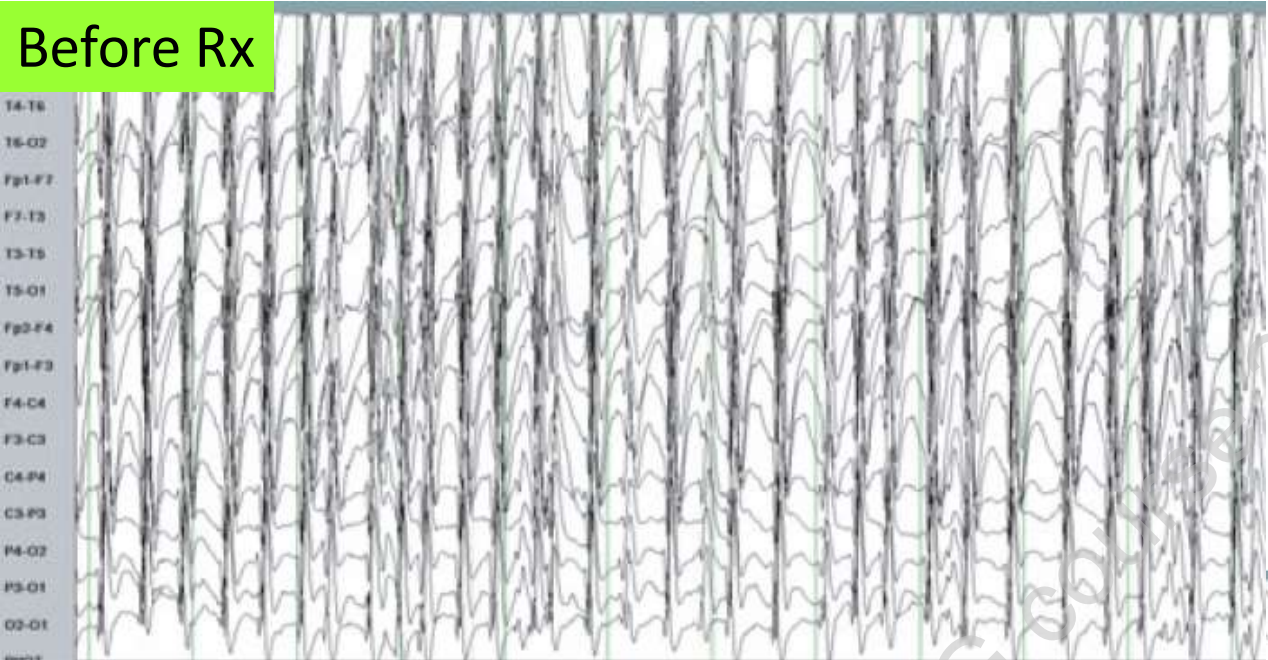
- New term: **D**evelopmental and/or **E**pileptic encephalopathy with **S**pike-wave **A**ctivation in **S**leep
- Previous term: CSWS/ESES

# DEE/EE-SWAS EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
<b>EEG</b> ( <i>interictal</i> ) <ul style="list-style-type: none"> <li>• Slow (1.5-2 Hz) spike and wave in NREM sleep</li> <li>• Abnormality activated in sleep</li> </ul>		<ul style="list-style-type: none"> <li>• Gen PFA in sleep (LGS)</li> <li>• Gen slow s/w in both awake and asleep stage (LGS)</li> </ul>	<b>Epileptic spasm</b>
<b>Ictal EEG</b> required For Dx	<u>Not</u> required		
<b>Sleep EEG</b>	Needed		

Before Rx

CSWS



After Rx

# Epilepsy with Myoclonic-atonic seizure

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# Epilepsy w myoclonic-atonic seizure EEG Diagnostic Criteria

	Mandatory	Alert	Exclusionary
<b>EEG (interictal)</b> <ul style="list-style-type: none"> <li>• Gen 2-6 Hz spike and wave</li> <li>• Polyspike/wave</li> </ul>		<ul style="list-style-type: none"> <li>• Gen PFA in sleep</li> <li>• Gen slow S/W</li> <li>• PPR at low Fq</li> </ul>	<ul style="list-style-type: none"> <li>• Hypsarrhythmia</li> <li>• Persistent focal abnormality</li> </ul>
<b>Ictal EEG</b> required For Dx	<u>Not</u> required. But if suspicious of LGS, epileptic spasms Ictal EEG should be recorded.		

# EEG: Epilepsy w myoclonic-atonic seizures



# Summary: Pattern that we have learned

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- SeLECTs
- SHE
- CAE/JAE/JME
- EEM
- IESS
- LGS
- DEE/EE-SWAS
- Epilepsy with myoclonic atonic

# Syndrome Core Diagnostic Criteria

	Mandatory	Alert	Exclusionary
Seizure type			
<b>EEG</b> ( <i>interictal</i> )	<input type="text"/>	<input type="text"/>	<input type="text"/>
Age at onset			
Development at onset			
Neurological exam			
Are MRI or <b>Ictal EEG</b> required For Dx			
Other studies-genetic			
Syndrome without laboratory confirmation			

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Thank you for your time

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