

Epilepsy					
i. Age of seizure onset:	days / weeks / months / years ( <u>circle</u> )				
ii. Age of peak seizure frequency:	:days / weeks / months / years ( <u>circle</u> )				
iii. Seizures ongoing?	S $\square$ No - on / off treatment? ( <u>circle</u> )				
iv. Number of seizure types $\ \Box$ 1	□ 2 □ 3 □ >3				
a. Seizure type 1					
i. Age onset:	days / weeks / months / years ( <u>circle</u> )				
ii. Seizure type:					
☐ focal - evolving to bilateral convulsion ☐ atonic ☐ clonic ☐ epileptic spasms ☐ unclassified	aired awareness □ hemiclonic: R / L / Both ( <u>circle</u> ) on □ tonic □ tonic □ myoclonic □ myoclonic-atonic				
Seizure description:					
iii. Seizure duration - average	sec / min / hourssec / min / hours				
iv. Approximate total number of seizur	es:				
v. Frequency of seizures ☐ hourly	☐ daily ☐ weekly ☐ monthly ☐ yearly ☐ other				
If other, please specify:					
vi. Seizure type ongoing? ☐ yes	☐ no - age of offset days / weeks / months / years				
vii. Is there a video? ☐ yes	□ no				
b. Seizure type 2					
i. Age onset	days / weeks / months / years (circle)				



Tura e di sa				
ii. Seizure type				
$\square$ absence - typical $\square$ atypical absence $\square$ myoclonic absence $\square$ absence with eyelid myoclonia				
☐ focal – aware ☐ focal - impaired awareness ☐ hemiclonic: R / L / Both (circle)				
$\square$ focal - evolving to bilateral convulsion $\square$ tonic-clonic $\square$ tonic				
□ atonic □ clonic □ myoclonic □ myoclonic-atonic				
☐ epileptic spasms ☐ unclassified				
Seizure description				
iii. Seizure duration - average sec / min / hours maximum sec / min / hours				
iv. Approximate total number of seizures				
v. Frequency of seizures $\square$ hourly $\square$ daily $\square$ weekly $\square$ monthly $\square$ yearly $\square$ other				
If other, please specify:				
vi. Seizure type ongoing? ☐ yes ☐ no - age of offset days / weeks / months / years				
vii. Is there a video? ☐ yes ☐ no				
c. Seizure type 3				
i. Age onset days / weeks / months / years (circle)				
ii. Seizure type				
□ absence - typical □ atypical absence □ myoclonic absence □ absence with eyelid myoclonia				
☐ focal – aware ☐ focal - impaired awareness ☐ hemiclonic: R / L / Both (circle) ☐ focal - evolving to bilateral convulsion ☐ tonic ☐ tonic				
☐ focal - evolving to bilateral convulsion ☐ tonic ☐ tonic ☐ myoclonic ☐ myoclonic-atonic				
□ epileptic spasms □ unclassified				
Seizure description				
iii. Seizure duration - average sec / min / hours maximum sec / min / hours				
iv. Approximate total number of seizures				
v. Frequency of seizures  hourly daily weekly monthly yearly other				
v. Frequency of seizures $\square$ hourly $\square$ daily $\square$ weekly $\square$ monthly $\square$ yearly $\square$ other If other, please specify:				



vii. Is there a video? ☐ yes ☐ no				
d. Seizure type >3				
If seizures >3:				
Age onset, seizure type, description, duration, number				
of seizures, frequency, ongoing?				
Electroencephalogram				
i. EEG at epilepsy onset $\qed$ yes $\qed$ no				
ii. EEG at epilepsy evolution □ yes □ no				
EEG at epilepsy onset				
i. Age of EEG: days / weeks / months / years ( <u>circle</u> )				
ii. Interictal background				
☐ normal ☐ background abnormality – focal / generalised ( <u>circle</u> )				
□ burst-suppression □ hypsarrhythmia / modified hypsarrhythmia				
iii. Interictal epileptiform activity				
$\square$ none $\square$ unifocal $\square$ multifocal $\square$ generalised $\square$ both focal and generalised				
Location if focal:				
Type if generalised: ☐ generalised spike-wave >3.5Hz ☐ generalised spike-wave 2.5-3.5Hz				
$\Box$ generalised spike-wave <2.5Hz $\Box$ generalised paroxysmal fast				
$\square$ generalised elecotrodecrement $\square$ other				
If other, please specify:				
iv. Proportion of EEG during which epileptiform activity is seen (awake or asleep)				
□ 0% □ 0-25% □ 26-50% □ 51-75% □ 76-100%				
v. Seizures recorded?				



vi. Seizure type and location				
Seizure type and location				
Seizure type and location				
Seizure type and location_				
EEG at epilepsy evolution				
i. Evolution:				
ii. Age of EEGdays / weeks / months / years				
iii. Interictal epileptiform activity				
□ none □ unifocal □ multifocal □ generalised □ both focal and generalised				
Location if focal:				
Type if generalised: ☐ generalised spike-wave >3.5Hz ☐ generalised spike-wave 2.5-3.5Hz				
☐ generalised spike-wave <2.5Hz ☐ generalised paroxysmal fast				
☐ generalised elecotrodecrement ☐ other				
If other, please specify:				
iv. Proportion of EEG during which epileptiform activity is seen (awake or asleep)				
$\square$ 0% $\square$ 0-25% $\square$ 26-50% $\square$ 51-75% $\square$ 76-100%				
v. Seizures recorded?				



vi. Seizure type and location				
Seizure type and location				
Seizure type and location				
Syndromes				
a. Epileptic syndrome at presentation				
<ul> <li>□ Dravet syndrome</li> <li>□ Early myoclonic encephalopathy</li> <li>□ Epilepsy of infancy with migrating focal seizures</li> <li>□ Landau-Kleffner syndrome</li> <li>□ Ohtahara syndrome</li> <li>□ unclassified</li> <li>□ other classified</li> <li>■ b. Epileptic syndrome at evolution</li> </ul>	☐ Early onset epileptic encephalopathy (onset <3m) ☐ Epilepsy with myoclonic-atonic seizures ☐ Febrile infection-related epilepsy syndrome ☐ Lennox-Gastaut syndrome ☐ West syndrome			
	□ Forth, anget arillagtic an earth play other (200)			
<ul><li>□ Dravet syndrome</li><li>□ Early myoclonic encephalopathy</li></ul>	<ul><li>☐ Early onset epileptic encephalopathy (onset &lt;3m)</li><li>☐ Epilepsy with myoclonic-atonic seizures</li></ul>			
☐ Epilepsy of infancy with migrating focal seizures	☐ Febrile infection-related epilepsy syndrome			
$\square$ Landau-Kleffner syndrome	$\square$ Lennox-Gastaut syndrome			
☐ Ohtahara syndrome	☐ West syndrome			
unclassified				
□ other classified				
Age at evolutiondays / weeks / months	; / years ( <u>circle</u> )			
Other epilepsy features				
i. Convulsive status epilepticus $\ \square$ no $\ \square$ yes	s - number of episodes			
ii. Non-convulsive status epilepticus ☐ no	☐ yes - number of episodes			
iii. Febrile seizures □ no	☐ yes - agesdays / months / years			
iv. Seizure triggers (does it trigger a specific seizure type	2)			



v. Seizure free period >6 months?	□ no	□ yes - age	days / weeks / months / years				
Dura	ation of seizure fre	eedom	days / weeks / months / years				
Treatment whilst seizure free							
Treatment							
i. Current							
ii. Previous							
iii. Non-pharmacologic treatments:							
$\square$ none $\square$ ketogenic diet	□ vagus nerv	re stimulation	other – specify:				
☐ epilepsy surgery: resesction – hem corpus callosotomy / other – specify:	•		section – lobular / resection – tubular ,				
Details (start and end date):							
Beneficial treatment:							
Exacerbating treatment:							
Development							
i. Was development ever normal?	□ no	□ yes					
ii. Development plateaued	□ no	$\square$ yes - age and to	rigger				
Details							
iii. Development regressed	□ no	$\square$ yes - age and t	rigger				
Details							
iv. Developmental outcome $\qed$ n	ormal						
☐ isolated delay - domain							
v. Outcome severity							
☐mild ☐moderate	□severe	$\square$ profound					
vi. Developmental milestones: age sat	walke	ıd					
single words							
		vords together					
vii. Visually attentive	□ no	□ yes					
viii. Hearing	□ normal	□ impaired - sens	sorineural / conductive ( <u>circle</u> )				
ix. Best gross motor skill:  ☐ none ☐ head control ☐ rolling ☐ sitting ☐ walking ☐ running							



x. Best language skill:					
□ none □ cooing □ complex babble □ < 5 words					
$\square$ <20 words $\square$ < 50 words $\square$ <100 words $\square$ 2-words together					
□ short phrases (<6 words) □ sentences					
xi. Autism spectrum disorder $\Box$ no $\Box$ yes					
xii. Psychiatric / behavioural problems					
Other neurologic features					
i. Movement disorder $\ \square$ no $\ \square$ yes - specify type and if persistent / paroxymal					
Type of movement disorder:					
ii. Tone abnormality □ no □ yes - type and distribution					
Type of abnormality:					
Other:					
iii. Head size					
□ normal □ microcephalic – congenital □ microcephalic – acquired □ macrocephalic – congenital □ congenital □ macrocephalic – acquired					
Other: Non-neurologic features					
i. Dysmorphic features $\Box$ no $\Box$ yes - specify					
ii. Neurocutaneous features					
iii. Growth abnormality □ no □ yes - small / large for age (circle)					
iv. Other abnormalities $\square$ no $\square$ yes - specify					
v. Enteral feeding required $\Box$ no $\Box$ yes – specify					
□ nasogastric tube □ nasojejunal tube □ percutaneous endoscopic gastrostomy					
□ percutaneous endoscopic jejunostomy □ other − specify					
Detectable out endoscopic jejunoscomy otner – specify					



vi. Age at which enteral feeding was used:		months / years ( <u>circle</u> )			
Perinatal					
i. Morphologic / growth abnormality	□ no	□ yes -	- specify below		
ii. Amniotic fluid abnormality	□ no	□ yes -	- polyhydramnios ,	/ oligohydramnios ( <u>circle</u> )	
iii. Other problems in pregnancy	□ no	□ yes ·	☐ yes - specify below		
(eg illness, bleeding, maternal medicati	ons)				
iv. Gestation		weeks			
v. Resuscitation required	□ no	□ yes -	- specify below		
vi. Other problems in neonatal period	□ no	□ yes -	- specify below		
Results of investigations					
i. MRI-brain performed		□ no	□ yes -	age and result	
ii. CT-brain performed (if no MRI)		□ no	□ yes -	age and result	
iii. Chromosomal microarray		□ normal	☐ abnormal	$\square$ not assessed	
Result:					
iv. Other genetic testing performed		□ no	□ yes -	details below	
Type, result, laboratory location, date_					
v. Other abnormal results		□ no	□ yes -	details below	
Type, result, laboratory location, date_					