## Syndromes by Age of Onset: (check all that apply)

Neonatal Period Table

| Neonatal Period | Present? |
| --- | --- |
| Benign familial neonatal epilepsy (BFNE) | No  Possible  Probable  Definite  Unknown  N/A |
| Early myoclonic encephalopathy (EME) | No  Possible  Probable  Definite  Unknown  N/A |
| Ohtahara syndrome | No  Possible  Probable  Definite  Unknown  N/A |

Infancy Table

| Infancy | Present? |
| --- | --- |
| Epilepsy of infancy with migrating focal seizures | No  Possible  Probable  Definite  Unknown  N/A |
| West syndromes | No  Possible  Probable  Definite  Unknown  N/A |
| Myoclonic epilepsy in infancy (MEI) | No  Possible  Probable  Definite  Unknown  N/A |
| Benign infantile epilepsy | No  Possible  Probable  Definite  Unknown  N/A |
| Benign familial infantile epilepsy | No  Possible  Probable  Definite  Unknown  N/A |
| Dravet syndrome | No  Possible  Probable  Definite  Unknown  N/A |
| Myoclonic encephalopathy in nonprogressive disorders | No  Possible  Probable  Definite  Unknown  N/A |

Childhood Table

| Childhood | Present? |
| --- | --- |
| Febrile seizures plus (FS+; can start in infancy) | No  Possible  Probable  Definite  Unknown  N/A |
| Early onset benign childhood occipital epilepsy (Panayiotopoulos type) | No  Possible  Probable  Definite  Unknown  N/A |
| Epilepsy with myoclonic atonic seizures | No  Possible  Probable  Definite  Unknown  N/A |
| Benign childhood epilepsy with centrotemporal spikes (BCECTS) | No  Possible  Probable  Definite  Unknown  N/A |
| Autosomal-dominant nocturnal frontal lobe epilepsy (ADNFLE) | No  Possible  Probable  Definite  Unknown  N/A |
| Late onset childhood occipital epilepsy (Gastaut type) | No  Possible  Probable  Definite  Unknown  N/A |
| Epilepsy with myoclonic absences | No  Possible  Probable  Definite  Unknown  N/A |
| Lennox-Gastaut syndrome | No  Possible  Probable  Definite  Unknown  N/A |
| Epileptic encephalopathy with continuous spike-and-wave during sleep (CSWS) | No  Possible  Probable  Definite  Unknown  N/A |
| Landau-Kleffner syndrome (LKS) | No  Possible  Probable  Definite  Unknown  N/A |
| Childhood absence epilepsy (CAE) | No  Possible  Probable  Definite  Unknown  N/A |

Adolescence – Adult Table

| Adolescence – Adult | Present? |
| --- | --- |
| Juvenile absence epilepsy (JAE) | No  Possible  Probable  Definite  Unknown  N/A |
| Other familial temporal lobe epilepsies | No  Possible  Probable  Definite  Unknown  N/A |
| Juvenile myoclonic epilepsy (JME) | No  Possible  Probable  Definite  Unknown  N/A |
| Progressive myoclonus epilepsies (PME) | No  Possible  Probable  Definite  Unknown  N/A |
| Autosomal Dominant Epilepsy with Auditory Features (ADEAF) | No  Possible  Probable  Definite  Unknown  N/A |
| Epilepsy with generalized tonic-clonic seizures alone | No  Possible  Probable  Definite  Unknown  N/A |

Less Specific Age Relationship Table

| Less Specific Age Relationship | Present? |
| --- | --- |
| Familial focal epilepsy with variable foci (childhood to adult) | No  Possible  Probable  Definite  Unknown  N/A |
| Reflex epilepsies | No  Possible  Probable  Definite  Unknown  N/A |

Distinctive Constellations Table

| Distinctive Constellations | Present? |
| --- | --- |
| Mesial temporal lobe epilepsy | No  Possible  Probable  Definite  Unknown  N/A |
| Rasmussen syndrome | No  Possible  Probable  Definite  Unknown  N/A |
| Gelastic seizures with hypothalamic hamartoma | No  Possible  Probable  Definite  Unknown  N/A |

Other Localization Related Epilepsies Table

| Other Localization Related Epilepsies | Present? |
| --- | --- |
| Temporal lobe epilepsy | No  Possible  Probable  Definite  Unknown  N/A |
| Frontal lobe epilepsy | No  Possible  Probable  Definite  Unknown  N/A |
| Occipital lobe epilepsy | No  Possible  Probable  Definite  Unknown  N/A |
| Parietal lobe epilepsy | No  Possible  Probable  Definite  Unknown  N/A |
| Focal epilepsy (specific localization unknown) | No  Possible  Probable  Definite  Unknown  N/A |

Other Table

| Other | Present? |
| --- | --- |
| The epilepsy does not fit into one of these specific electro-clinical or distinctive constellations categories | No  Possible  Probable  Definite  Unknown  N/A |

## Additional Information

If two or more epilepsy syndromes were selected as present, rate the confidence level that these are distinct syndromes:

No Confidence

Possible

Probable

Definite

Unknown

N/A

## GENERAL INSTRUCTIONS

Based on the current International League Against Epilepsy (ILAE) guidelines, this CRF Module is recommended to classify syndromes for all epilepsy studies. Only one syndrome should be checked for a given time point, however it is possible to have had a syndrome in infancy that develops into another syndrome during childhood. Therefore, if the form is used more than once during follow-up, the possible evolution of syndromes can be codified.

The following definitions should be used when completing this form:

* Not Present/None = The summary of evidence suggests no possibility
* Possible = The summary of evidence suggests less than 50% confidence level
* Probable = The summary of evidence suggests greater than 50% confidence level
* Definite = The summary of evidence suggests 100% confidence level
* Unknown = The summary of evidence is not sufficient to support a finding
* N/A = Not Applicable; to be used at the discretion of the Principal Investigator based on study design

## REFERENCES

Berg AT, Berkovic SF, Brodie MJ, Buchhalter J, Cross JH, van Emde Boas W, Engel J, French J, Glauser TA, Mathern GW, Moshé SL, Nordli D, Plouin P, Scheffer IE. Revised terminology and concepts for organization of seizures and epilepsies: report of the ILAE Commission on Classification and Terminology, 2005-2009. Epilepsia. 2010 Apr;51(4):676-85. Epub 2010 Feb 26.

Freely available online at: [Classifications and Terminology Report](http://www.ilae.org/Visitors/Centre/ctf/ctfoverview.cfm)