Establishing Criteria for Pediatric Epilepsy Surgery Centers Levels of Care: Report from the ILAE Pediatric Epilepsy Surgery Task Force


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Glossary of Terms

CT computed axial tomography
MRI magnetic resonance imaging
fMRI functional magnetic resonance imaging
DTI diffusion tensor imaging
EEG electroencephalogram
vEEG video electroencephalogram
PET positron emission tomography
SPECT single photon emission computed tomography
MEG magnetoencephalogram
MSI magnetic source imaging
TMS transcranial magnetic stimulation
VNS vagal nerve stimulator
DBS deep brain stimulation
RNS responsive nerve stimulation
ICU intensive care unit
PICU pediatric intensive care unit
T Tesla
AVM arterio-venous malformations
WHO World Health Organization

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Pre-surgical evaluation and surgery in the pediatric age group are unique in challenges related to caring for the very young, range of etiologies, choice of appropriate investigations, and surgical procedures. Accepted standards which define the criteria for levels of pre-surgical evaluation and epilepsy surgery care do not exist. Through a modified Delphi Process involving 61 centers with experience in pediatric epilepsy surgery across 20 countries, including low-middle to high income countries, we established consensus, for two levels of care. Levels were based on age, etiology, complexity of pre-surgical evaluation and surgical procedure. Competencies were assigned to the levels of care relating to personnel, technology, and facilities. Criteria were established when consensus was reached (≥75% agreement). Level 1 care consists of children age nine years and older, with discrete lesions including hippocampal sclerosis, undergoing lobectomy or lesionectomy, preferably on the cerebral convexity and not close to eloquent cortex, by a team including a pediatric epileptologist, pediatric neurosurgeon, and pediatric neuroradiologist with access to vEEG and 1.5T MRI. Level 2 care, also encompassing Level 1 care, occurs across the age span and range of etiologies (including tuberous sclerosis complex, Sturge-Weber syndrome, hypothalamic hamartoma) associated with MRI lesions that may be ill-defined, multilobar, hemispheric or multifocal, and includes children with normal MRI or foci in/abutting eloquent cortex. Available Level 2 technologies includes 3T MRI, other advanced
MR technology including fMRI and DTI (tractography), PET and/or SPECT, source localization with EEG or MEG, and the ability to perform intra- or extra-operative invasive monitoring and functional mapping, by a large multi-disciplinary team, with pediatric expertise in epilepsy, neurophysiology, neuroradiology, epilepsy neurosurgery, neuropsychology, anesthesia, neurocritical care, psychiatry, and nursing. Levels of care will improve safety and outcomes for pediatric epilepsy surgery and provide standards for personnel and technology to achieve these levels.

**Key Points Box:**

1) Levels of pediatric epilepsy surgery care established by a modified Delphi method involving multidisciplinary epilepsy professionals from across the globe

2) Two levels of care established based on clinical complexities and institutional competencies

3) Standardized criteria designed to elevate care and improve safety of all infants and children with epilepsy undergoing epilepsy surgery

**Introduction**

The specific skills, procedures, and resources required for pre-surgical evaluation and epilepsy surgery in children have steadily gained acknowledgement over the past 30 years. Several efforts of the International League Against Epilepsy (ILAE) Pediatric Epilepsy Surgery Task Force have delineated the status of pediatric epilepsy surgery in an effort to gain visibility and standardize care. The number of pediatric epilepsy surgical procedures has increased over the last decade, in contrast to adults. Nevertheless, there is an increasing recognition that epilepsy surgery in children is under-utilized. Historically, epilepsy surgery was regarded as a treatment of last resort whereas more recently surgery is considered standard of care in appropriately selected patients, being superior to treatment with a third medication. There is also evidence that surgery may mitigate the detrimental effects of ongoing seizures and medications, leading to improved cognition or prevention of further cognitive decline.
as well as reduced morbidity and mortality. Furthermore, there is evidence that delays in epilepsy surgery increase the risks of not achieving seizure control, poor cognitive outcomes, and death.\textsuperscript{13, 14, 15}

The first ILAE Epilepsy Surgery Task Force document\textsuperscript{16} acknowledged the unique characteristics of early age epilepsies and candidacy for surgery. Among these are the issue of brain development, the existence of age-specific epileptic syndromes, the prospect of plasticity with recovery of function, the pathological substrates and etiologies specific to the pediatric focal epilepsies, and the various surgical procedures performed. These features justify dedicated resources for specialty pediatric epilepsy surgery centers highlighting the importance of having pre-surgical evaluation and neurosurgery teams with specialized skills and settings - infants and children are not small adults. A set of standards were established for physicians detailing when to refer children with drug-resistant epilepsy to specialists. In the second document the range of the pathological substrates that occur in pediatric populations and types of surgical procedures was affirmed in a worldwide survey of 20 pediatric epilepsy surgery centers.\textsuperscript{17} A third document\textsuperscript{18} provided a range of views and consensus on the use of diagnostic testing for a range of pediatric entities determined by etiology/pathology. These recommendations were based on a literature review and consensus discussion, admittedly in the context of limited high-level evidence. Included was a recommendation for an epilepsy surgery evaluation pathway, priorities for diagnostic testing, and implementation of resection strategies including indications for invasive monitoring. A fourth document\textsuperscript{19} surveyed pediatric epilepsy neurosurgeons on current practice and surgical approaches, in the context of varying pathology and procedures, including technical aspects of operating room practice. This was followed by a survey of surgical practice patterns in children with epileptogenic foci near or in eloquent cortex along with recommendations for minimizing and reporting neurological deficits.\textsuperscript{20}

These documents emphasize the special and particular needs of pediatric epilepsy surgical care. Epilepsy surgery is evolving in its standards for evaluation, consideration, and execution. The number of centers that perform epilepsy surgery in children has increased, yet the resources, expertise, candidacy, and practice patterns vary considerably; adult skills are not transferable to children. Many worldwide regions do not have designated epilepsy centers for
surgical evaluation and may be unaware of the technology and skills required. The identification and consideration of a child as a surgical candidate (“presurgical care”), in addition to the selection of appropriate diagnostic tools and performance of surgery, includes knowledge of pediatric epilepsy, neuroanatomy, neurophysiology, and of the numerous technical approaches. For these reasons there remain gaps in the care of children with drug-resistant epilepsy who are potential candidates for epilepsy surgery. Here we seek to address these gaps by setting standards and resources necessary to achieve the aims delineated as levels of care. These efforts are not only necessary to set standards for basic and advanced epilepsy surgery care, but importantly serve as a resource for centers to achieve the technology, personnel, skills, and facilities to be a pediatric epilepsy and epilepsy surgery center. The manuscript also contributes to the Domain 4 (competencies 4.1 to 4.7) of the ILAE curriculum.21

Several countries have put forward varying forms of guidelines or standards for epilepsy surgery care (Chile, Brazil, Germany/Austria/Switzerland, England, China, United States; Table 1; See Supplementary Materials I for details). Some of these are legislated, some by self-authorized oversight bodies, while others are consensus statements. They define single level or at most two levels of surgical care. When two levels are proposed the first level is more basic (lesionectomy, lobectomy) and the second level more complex care or procedures (e.g. hemispherectomy, eloquent cortex). The first level typically includes CT, MRI (≤ 1.5T), and EEG; the second level typically includes higher field MRI (3T), DTI, PET, SPECT, and invasive monitoring of any kind. Few distinguish children as a separate entity. The English national Childhood Epilepsy Surgical Services (CESS) designate care of those younger than five years to certified pediatric centers only, and the Chileans place infants in Level Two centers but without other comment of pediatric expertise or designation.22 The American National Association of Epilepsy Centers (NAEC) offer a pediatric designation, but unlike the English, the American criteria do not address etiology, diagnostic complexity, or specify surgical procedures.23 Only the English and the American criteria include a system of auditing. The European Reference Network for Rare and Complex Epilepsies and the Chinese designate volume of evaluation and surgical procedures.24, 25

Previous national efforts share some common elements, others hold unique elements, but all have limitations. They do, however, form the basis of ILAE efforts going forward to achieve
worldwide consensus for these criteria, based on expert opinion and experience in a large group of professionals from different backgrounds. Here we delineate the elements of care and expertise necessary (e.g. personnel, technology, facilities) to perform pediatric epilepsy surgery according to different levels of care and to be considered a pediatric epilepsy surgery center. We also presume each center holds the commensurate skills and resources necessary to provide presurgical care for children with epilepsy.

Methods

A modified Delphi method was used\textsuperscript{26, 27, 28} (See Figure 1) to achieve consensus on aspects and elements of care. Items to be rated were generated, based on a literature review, by a core team (See Supplementary Materials II) according to complexity of care and competencies as elements for pediatric epilepsy surgery care (see below). Round I consisted of rating, on a seven-point scale, each item (134 Items, See Supplementary Materials III). The presumption was that two or three levels (basic, middle, advanced) of care would become evident from the responses from members of the ILAE pediatric surgery task force members (29 participants). However, the results demonstrated an absence of consensus on any items.

It was therefore decided at the Task Force meeting in Barcelona 2017, convened to review and discuss survey results, to establish two levels of care and have up or down votes on elements of care required for each of these levels (Round II). Round II was further divided into two tiers of voting based on Complexity, Round IIA; and Competencies, Round IIB. Specifically, the two levels were established first by considering Complexity of care (age, etiology, complexity of pre-surgical evaluation, type of surgery) to decide what patients would be assigned to Level 1 (basic epilepsy surgery) or Level 2 (complex epilepsy surgery). Once levels were determined in the tier one survey (Round IIA), we then assigned Competencies to each of the two levels by vote in a second-tier vote (Round IIB) that included items related to the required technology (derived from commonly used and new technology advocated for epilepsy surgery evaluation and care), resources, personnel, and facilities for each level. For some items, access to resources not otherwise on site (i.e. rehabilitation, MEG, neuropathology) were deemed...
sufficient. Competencies were also considered for a series of particular clinical circumstances, such as infants, operating in or near eloquent cortex, and multifocal epilepsy.

The Round II questions were drafted, discussed, revised, and finalized by members of the core team. The methodological approach for Round IIA and then Round IIB were similar. The Round II questions were circulated to the wider participant group (n=80 members; Table 2 for participating members/sites). All voting was anonymous, was performed on an electronic platform entered directly into a RED Cap database, included feedback with questions, comments, and provided participants the opportunity to identify gaps in Round II items. Unclear items were clarified, and when necessary removed or substituted, and new items added. Consensus was defined as 75% agreement or more on a given item; items with 60-74% agreement were included in a subsequent round.

For round IIA, participants were asked, for each item, to select the type of center that was appropriate for each item (Level 1, level 2 or neither; See Supplementary Materials III). Initial voting, IIAi, involved 51 items. A second round of voting (n=28 items) ensued (IIAii) and included clarifying questions. In the second round (IIAii) results of the first round (IIAi) of voting were provided and members asked if they agreed with the majority, and if not to provide reasons for disagreement. A final, third round (IIAiii), occurred for a minority of items (n=1 item). Some items did not achieve consensus. After each round, results were tabulated and circulated to participants.

For round IIB, participants were asked, for each item, to select aspects of technology, personnel, resources/facilities, and experience that was appropriate (Level 1 or level 2 care, or neither; See Supplementary Materials III). Initial voting, IIBi, involved 159 items. A second round of voting (n=53 items) ensued (IIBii) and included clarifying questions. In the second round (IIBii) results of the first round (IIBi) of voting were provided and members asked if they agreed with the majority, and if not to provide reasons for disagreement. A final, third round (IIBiii), occurred for a minority of items (n=5 items). Some items did not achieve consensus. After each round, results were tabulated and circulated to participants.
In the principal round of voting (Round II) we aimed to assure greater distribution across the globe as well as economic diversity. Eighty centers were invited from the following sources: participants in the pediatric neurosurgical survey\textsuperscript{19}, sites identified as having epilepsy surgery centers for children from ILAE chapter lists, NAEC centers (for the USA), the European Reference Network for Rare and Complex Epilepsies, and ILAE pediatric epilepsy surgery task force members. Sixty-one centers participated in the principal (Round II) survey for Complexities and subsequently Competencies. Participating centers and collaborators are listed in Table 2 (See figure 2, Map). Results are descriptive based on 75% agreement. It is presumed a case conference will occur with members of the specialty team to review history, clinical characteristics, results of diagnostic studies, and to agree on surgical options and approaches.\textsuperscript{18}

Results

Participating Sites

Sixty-one centers out of 80 invited participated in the principal (Round II) survey. These centers represented 20 countries (Figure 2); 76% came from WHO high income economies, 17% upper middle income economies, and 7% lower middle income economies. Of those answering surveys for their centers, 65% were neurologists, 27% neurosurgeons and 8% from other disciplines. Twenty-nine sites also conducted epilepsy surgery on adult patients. The number of first time epilepsy resections per year on patients 18 and younger (excludes VNS, or prior procedures for invasive monitoring) ranged from 2 to 165, with a mean of 35, and a median of 28 (Figure 3).

Level 1 Pediatric Epilepsy Surgery Center Criteria (see Table II)

Complexities

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Age: Children nine years and older, the same as the earlier neurosurgical consensus survey.\textsuperscript{19}

Etiology: Children with single and discrete lesions on MRI, including tumors, well-defined focal cortical dysplasias, cavernomas, hippocampal sclerosis (single pathology), and cysticercosis.

Location/procedures: Lesionectomy, temporal lobectomy, amygdalo-hippocampectomy, resections on the cerebral convexity away from eloquent cortex, and placement of VNS.

Competencies

Personnel: Pediatric neurologist, at least one pediatric epileptologist (who is also a pediatric neurologist), a pediatric neurosurgeon, a pediatric neuroradiologist, an anesthesiologist skilled with children, and a social worker.

Technology: Should include the ability to perform routine EEG and prolonged video-EEG monitoring. Standard imaging includes CT and 1.5 T MRI using a dedicated epilepsy protocol.\textsuperscript{29,30}

Resources/Facilities: Include designated pediatric beds which may reside within an adult hospital but they must be clearly delineated as pediatric care, and a pediatric intensive care unit (PICU).

Experience: Evaluating and treating over 100 pediatric patients per year with epilepsy (median number subsequently accepted in second round vote). Minimum number of pediatric epilepsy surgeries, excluding VNS, is 15 per year.

Level 2 Pediatric Epilepsy Surgery Center Criteria

Complexities
All Level 1 ages, etiologies, and surgical procedures are included in Level 2 care.

*Age:* children eight years and younger are restricted to level 2 centers.

*Etiology/Lesion characteristics:* Hippocampal sclerosis with dual pathology, arteriovenous malformations, encephalomalacia due to ischemic or other injury, hemispheric malformations of cortical development, post infectious/inflammatory conditions and Rasmussen’s encephalitis, tuberous sclerosis complex, Sturge-Weber syndrome, and hypothalamic hamartoma are Level 2, as are MRI-negative patients and those with poorly demarcated or multiple lesions.

*Location/procedures:* In addition to temporal lobar and discrete convexity lesions, Level 2 locations include extra-temporal lobar, multilobar, hemispheric, multifocal, interhemispheric, and subcortical regions. Procedures may involve eloquent cortex, insula or basal regions. This indicates a range of procedures including corticectomy, lesionectomy, lobectomy, amygdalo-hippocampectomy, multi-lobar resection, hemispheric resections and disconnections, and resections in deep, interhemispheric and insular regions and eloquent cortex. Additional stimulation procedures include deep brain stimulation and responsive nerve stimulation, in addition to vagal nerve stimulation. Invasive intracranial monitoring including subdural and stereo EEG monitoring are Level 2 (see below).

Clinical characteristics of children undergoing epilepsy surgery only in level 2 centers include children with developmental regression/epileptic encephalopathy, specific developmental or medical syndromes, and genetic abnormalities. Comorbid, complicated medical problems (not further defined) and ongoing status epilepticus are also restricted to Level 2 centers.

**Competencies**

*Personnel:* Includes a child neurologist, a minimum of two pediatric epileptologists, a pediatric neurophysiologist (who may also be a pediatric epileptologist), a pediatric epilepsy neurosurgeon preferably trained in functional/stereotactic epilepsy surgery, a pediatric neuro-radiologist, a
pediatric neuropsychologist, a child psychiatrist, a pediatric anesthesiologist, a neuropathologist, a social worker, pediatric neurology nursing, a nutritionist/dietician, and a program coordinator.

**Technology:** Includes routine EEG, long-term video-EEG, CT, 3T MRI, functional MRI, DTI (white matter tractography), angiography, functional radio-ligand imaging (PET and/or SPECT), source localization capability (but not specified between high-density EEG or MEG, or fMRI/EEG). MRI follows a dedicated epilepsy protocol.\textsuperscript{29, 30} Unique to Level 2 is the capacity for invasive EEG recording (intraoperative ECoG, subdural EEG, stereo EEG) and cortical stimulation mapping (intra- or extra-operative). Neuro-navigation, a widely available MRI based frameless system for intraoperative localization of brain structures and lesions before a resection is performed, is deemed standard of care.

**Resources/Facilities:** Dedicated pediatric services that may be linked to an adult facility or be free standing, a dedicated pediatric epilepsy monitoring unit (EMU) (NOT on adult floor/service), a dedicated PICU and access to rehabilitation facilities.

**Experience:** Evaluating and treating a minimum of 225 children with epilepsy a year in clinic (median number subsequently accepted in a second round vote) and conducting a minimum number of 20 epilepsy surgeries per year, excluding VNS. Twenty resections was the median number of epilepsy surgical procedures per year recommended from the initial vote. In a second vote the group then voted to accept 20 as the minimum number of definitive epilepsy surgical procedures per year.

**Other Level 2 Considerations.**

While there was consensus not to split Level 2 further, there was near unanimous (95%) or unanimous view that a subset of complexities were sufficiently unique to warrant specific competencies, thus perhaps suggesting a more specialized subset of level 2: There was a unanimous view that children younger than six months with multifocal epilepsy/lesions, a seizure focus in or near eloquent cortex, those with normal MRI or an indiscrete lesion were particular to specialized pediatric care. There was near unanimous (>95%) view for children less...
than 24 months; children with tuberous sclerosis complex, hypothalamic hamartoma, insula, multi-lobar (but not hemispheric) epilepsy/lesions belonged in a Level 2 setting.

A series of eight clinical circumstances that represent particular challenges were presented to ascertain consensus on additional competencies beyond those agreed upon for Level 2: 1) Epilepsy surgery in infants six months and younger, 2) children younger than 24 months, 3) operating near or in eloquent cortex, 4) operating on poorly demarcated lesions, 5) muti-lobar resection (without clear MRI abnormalities), 6) operating in insular cortex, 7) children with tuberous sclerosis, and 8) hypothalamic hamartoma. Consensus was that a pediatric neuroanesthesiologist be available when operating on infants younger than 6 months of age. There are commonalities for those with normal MRI, ill-defined lesions, and multi-lobar abnormalities where two functional imaging modalities (PET, SPECT (as opposed to one or the other in Level 2). Insular surgery benefits from access to ictal/interictal SPECT. A minimally-invasive method of some form was deemed necessary for hypothalamic hamartoma surgery. fMRI is deemed standard for normal MRI and operating near or in eloquent cortex.

*No Consensus*

No agreement was achieved regarding corpus callosotomy being designated level 1 or 2. Similarly, level designation for the presence of psychiatric comorbidity or intellectual disability was not established.

*Not Elements of Standard Care*

It is also notable that some technologies, especially newer technology, and facilities were not yet deemed essential or standard of care, including MEG, Transcranial Magnetic Stimulation (TMS), High Frequency Ultrasound (HiFUS), intraoperative MRI, minimally invasive surgery (of some form), and a dedicated Neuro PICU.
Discussion

We present recommendations for two levels of pediatric epilepsy surgery care, Level 1 basic care and Level 2 advanced care, founded on the unique features of patient complexity including age, etiology, and location/procedure and center competencies including personnel, technology, facilities, and case volume. Children age 8 years and below should receive care at Level 2 Centers. The goal of establishing criteria is to elevate care, and improve safety of all infants and children with epilepsy undergoing epilepsy surgery, and allows centers to petition for resources, technology, and personnel.

Our levels of care and delineation of technology and personnel match previous guidelines and standards. Ours differ by bringing several elements together, determined by consensus through a Delphi process of successive and tiered votes, among more than 60 international pediatric epilepsy centers, and incorporate pediatric specific skills. The choice of age eight years as a major threshold of complexity (Level 2) matches the views from the prior neurosurgical survey. There is strong agreement that the younger the patient, the more essential it is that the child be cared for by a pediatric epilepsy specialty unit. This proposed view is concordant, with some modification, to the Chilean and English regulations. There was (near) unanimous view that the youngest (<24 months of age, especially < 6 months) require advanced and skilled teams for pre-surgical and surgical care. Beyond age, commonalities for Level 2 include an array of advanced structural and functional imaging technologies, the capacity for intracranial EEG monitoring and cortical stimulation, and pediatric specialty personnel.

Other guidelines generally do not comment on experience, either in patients seen or in children operated upon. There are some data to suggest that one needs to operate (resections/disconnections) on 15 patients with epilepsy per year to maintain skills as outcomes; adverse perioperative events are greater when volume falls below 15. For these reasons 15 is considered the minimum number of resections for a Level 1 center. For Level 2 centers, 20 is deemed the minimum but further study may be needed to ascertain numbers needed to establish and maintain advanced operative skills. We chose as another minimum the number of children...
with epilepsy evaluated each year, viewing the number of children evaluated for epilepsy surgery to be more open to variable interpretation. Previous work stressed the importance of case conferences for any level of care.\textsuperscript{18}

There were some areas where consensus was not reached; by default they are assumed to fall within Level 2 but not excluded from level 1. The introduction of technology is a major part of epilepsy surgery evaluation. A challenge with diagnostic and therapeutic technology is that high-quality studies to evaluate their utility are often lacking to achieve class 1 or class 2 evidence status, especially in children.\textsuperscript{32} New technology is often expensive with a poorly defined utility, but often assumed added diagnostic value. Thus, MEG is often valued but not deemed standard, similarly intra-operative MRI, and TMS are relatively new technologies that are used by some centers but not commonly accepted in practice. 3T MRI appears to have been accepted as standard practice, and may have helped change practice. It is unclear what the role 7T MRI or post processing image algorithms will assume in the future, as well as technology yet to be developed.\textsuperscript{33, 34, 35}

There will be exceptions and differences of view with implementing these standards. For example, cavernomas may be multifocal whereas a younger child may have a clearly distinct lesion outside of eloquent cortex. Cysticercosis is viewed as a distinct lesion but those hailing from areas where cysticercosis is endemic thought it a Level 2 and not a Level 1 entity, based on complex subtleties of the entity and decision making as to whether surgery is required. Several participants commented that additional skills may be required to ascertain whether hippocampal sclerosis is isolated or accompanied by dual pathology. While the treatment of AVMs was considered by many as standard neurosurgical care, when accompanied by epilepsy it was deemed specialty care for level 2 centers. Similarly, although VNS placement can be performed at Level 1 centers, excluding candidacy for resective surgery is important and may require diagnostic competencies available at level 2 centers.

There are limitations to this effort. The survey was conducted mainly by experienced epilepsy center directors, or senior members of established epilepsy programs, who may carry a bias for particular technologies, etiologies, or procedures. While we have identified the needs for
pediatric epilepsy surgery, the evaluation cannot take place without a setting for pediatric epilepsy care. This is implied in the personnel, technology, and patient volumes presented but not a primary purpose of this endeavor. Comment was not made on epilepsy genetics, increasingly central to pediatric neurology, let alone epilepsy care, including the consideration of surgical candidacy. The ability to care for children with genetic epilepsies is noted in complexities for Level 2 but specific expertise in epilepsy genetics was not considered. We did not address range of medical complexities that may be present. We have also not surveyed the specifics of what technology is needed in the operating room other than neuro-navigation systems; many of these needs, however, are covered in the previous neurosurgical survey.

Nor have we defined what is considered an expert and the pathway of training to achieve expertise. A previous ILAE Pediatric Epilepsy Surgery Task Force survey (unpublished observations) noted wide ranging experience in training of pediatric neurologists, epileptologists, and neurophysiologists, and more variability in the training of pediatric epilepsy neurosurgeons. In some countries, epileptologists are also trained as electrophysiologists, in others the two are distinct disciplines. A future task will be to define the training and experience necessary to be an expert.

Finally, resource limited economies are less well represented in this survey. Yet, expertise and technology are critical for specialty care and it is hoped that by delineating these features it will be easier to petition for the requisite training and technology necessary to improve and standardize care in these settings.

We proposed criteria for epilepsy centers so that programs will have the equipment, personnel, skills, and volume to optimize outcomes and elevate quality of care. We do not aim to deny care to children in resource-limited regions of the world where Level 1 centers do not exist. Such children may be offered epilepsy surgery at facilities without all the corresponding competencies, resources, and experience outlined above, but they would not be deemed to be epilepsy centers. Rather the long-term aim is to build the necessary resources and experience outlined here, step by step. While we have stated the requirements for Level 2 care, that does not preclude Level 1 centers from incorporating Level 2 technology and personnel. For example, 3T
MRI, or some form of functional imaging, strengthens the armamentarium at that institution and may help identify children with a previously unrecognized focal and operable abnormalities.

**Conclusion**

We identified two levels of care based on complexities and competencies, with volume of patient care experience, and minimum number of annual and epilepsy surgery resections. The basic Level 1 center cares for children 9 years and older, with a single defined lesion, employing a pediatric epileptologist and a general pediatric neurosurgeon, utilizing video-EEG, and 1.5T MRI. The advanced Level 2 center cares for children of all ages, in a pediatric setting, with a team of skilled pediatric clinicians trained in neurology, epilepsy, neurophysiology, neuroimaging, and pediatric epilepsy surgery. Level 2 centers utilize an array of advanced diagnostic technology including 3T MRI, fMRI, PET or SPECT; have invasive EEG monitoring capacity; conduct surgery in those with normal MRI, involvement near and in eloquent cortex, and deep lesions; and, provide specialty service on the in-patient floor, PICU, and rehabilitation services. An argument can be made for additional skills and technology in those younger than 24 months, and those with multi-lobar or deep lesions. Future efforts should include defining criteria for specialty training. The goal of establishing criteria is to elevate care, and improve safety of all infants and children with epilepsy undergoing epilepsy surgery and allows centers to petition for resources, technology, and personnel.

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Ethical Publication Statement.
We confirm that we have read the Journal’s position on issues involved in ethical publication and affirm that this report is consistent with those guidelines.

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**Figure 1. Delphi Process Flow Chart:** Through a modified Delphi Process we established consensus, for two levels of care. Round I consisted of rating, on a seven-point scale, each item. Round II was further divided into two tiers of voting based on Complexities, Round IIA; and Competencies, Round IIB. A second round of voting ensued (IIAii) and included clarifying questions. A final, third round (IIAiii), occurred for a minority of items. Initial voting, IIBi
occurred and then a second round of voting ensued (IIBii) and included clarifying questions. A final, third round (IIBiii), occurred for a minority of items.

**Figure 2. Map of Survey Participants:** Participation involved 61 centers with experience in pediatric epilepsy surgery across 20 countries, including low-middle through to high income countries.

**Figure 3. Surgical Resections Per Year by Sites:** The mean number of surgical resections per year was 35 and the median was 28.

<table>
<thead>
<tr>
<th>Feature</th>
<th>USA NAEC</th>
<th>Brazil</th>
<th>Chile</th>
<th>Europe ERN</th>
<th>England CESS</th>
<th>German Speaking</th>
<th>China</th>
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**Table 1: Worldwide Epilepsy Surgery Criteria**

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*NAEC: National Association of Epilepsy Centers*

*ERN: European Reference Network*

*CESS: Children’s Epilepsy Surgery Service*

*DGFE: Deutsche Gesellschaft für Erziehungswissenschaft (German Society for Epileptology)*

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Table 2: Survey Participants and Sites

USA (18)

<table>
<thead>
<tr>
<th>Physician</th>
<th>Institution</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>William Davis Gaillard</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chima Oluigbo</td>
<td>Children’s National Medical Center</td>
<td>Washington, DC</td>
</tr>
<tr>
<td>Prasanna Jayakar</td>
<td>Nicklaus Children’s Hospital</td>
<td>Miami, FL</td>
</tr>
<tr>
<td>Mark Libenson</td>
<td>Boston Children’s Hospital</td>
<td>Boston, MA</td>
</tr>
<tr>
<td>Ajay Gupta</td>
<td>Cleveland Clinic</td>
<td>Cleveland, OH</td>
</tr>
<tr>
<td>Jack Kerrigan</td>
<td>Phoenix Children’s Hospital</td>
<td>Phoenix, AZ</td>
</tr>
<tr>
<td>Susan Arnold</td>
<td>Dallas University of Texas Southwestern</td>
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<tr>
<td>Gary Mathern</td>
<td>University of California Los Angeles</td>
<td>Los Angeles, CA</td>
</tr>
<tr>
<td>Jeff Ojeman</td>
<td>Seattle Children’s Hospital, University of Washington</td>
<td>Seattle, WA</td>
</tr>
<tr>
<td>Shlomo Shinnar</td>
<td>Montifiore Hospital, Einstein University</td>
<td>New York, NY</td>
</tr>
<tr>
<td>Sudha Kessler</td>
<td>Children’s Hospital of Philadelphia</td>
<td>Philadelphia, PA</td>
</tr>
<tr>
<td>Jason Doescher</td>
<td>Minnesota Epilepsy Group</td>
<td>Minneapolis, MN</td>
</tr>
<tr>
<td>Elaine Wirrell</td>
<td>Mayo Clinic</td>
<td>Rochester, MN</td>
</tr>
<tr>
<td>Adam Hartman</td>
<td>Johns Hopkins Hospital</td>
<td>Baltimore, MD</td>
</tr>
<tr>
<td>Doug Nordli</td>
<td>University of Chicago</td>
<td>Chicago, IL</td>
</tr>
<tr>
<td>Jim Whelass</td>
<td>Memphis Children’s Hospital</td>
<td>Memphis, TN</td>
</tr>
<tr>
<td>Michael Handler</td>
<td>Colorado Children’s Hospital</td>
<td>Denver, CO</td>
</tr>
<tr>
<td>Howard Weiner</td>
<td>Texas Children’s Hospital, Baylor University</td>
<td>Houston, TX</td>
</tr>
<tr>
<td>Nathalie Jette</td>
<td>Icahn School of Medicine at Mount Sinai</td>
<td>New York, NY</td>
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Canada (2)

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<thead>
<tr>
<th>Physician</th>
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<tbody>
<tr>
<td>Elizabeth Donner</td>
<td>Hospital for Sick Children, University of Toronto</td>
<td>Toronto, ON</td>
</tr>
<tr>
<td>Mary Connolly</td>
<td>University of British Columbia</td>
<td>Vancouver, BC</td>
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Europe (20)

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<tr>
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<th>Location</th>
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<tr>
<td>J Helen Cross, Martin Tisdall</td>
<td>Great Ormond Street Hospital for Sick Children, University College London</td>
<td>London, United Kingdom</td>
</tr>
<tr>
<td>Timothy Martland</td>
<td>Central Manchester University Hospitals</td>
<td>Manchester, United Kingdom</td>
</tr>
<tr>
<td>Ailsa McLellan</td>
<td>The Royal Hospital for Sick Children</td>
<td>Edinburgh, United Kingdom</td>
</tr>
<tr>
<td>Tilman Polster</td>
<td>Mara Hospital, Bethel Epilepsy Centre</td>
<td>Bielefeld, Germany</td>
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<tr>
<td>J (Hans) Holthausen</td>
<td>Schon Kliniken</td>
<td>Vogtareuth, Germany</td>
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<tr>
<td>Julia Jacobs</td>
<td>University Medical Center Freiburg</td>
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<tr>
<td>Kees P Braun</td>
<td>University Medical Center Utrecht</td>
<td>Utrecht, Netherlands</td>
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<tr>
<td>Bertil Rydenhag</td>
<td>University of Gothenburg</td>
<td>Gothenburg, Sweden</td>
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<tr>
<td>Liisa Metsahonkala</td>
<td>Helsinki University Hospital</td>
<td>Helsinki, Finland</td>
</tr>
<tr>
<td>Laura Tassi</td>
<td>Claudio Munari Epilepsy Surgery Center, Ospedale Niguada</td>
<td>Milan, Italy</td>
</tr>
<tr>
<td>Luca Del Palma</td>
<td>Sapienza University</td>
<td>Rome, Italy</td>
</tr>
<tr>
<td>Carmen Barba</td>
<td>University of Florence</td>
<td>Florence, Italy</td>
</tr>
<tr>
<td>Alexis Arzimanoglou</td>
<td>University Hospitals of Lyon and Lyon Neuroscience Research Centre</td>
<td>Lyon, France</td>
</tr>
<tr>
<td>Mathilde Chipaux</td>
<td>Rothschild Foundation Hospital</td>
<td>Paris, France</td>
</tr>
<tr>
<td>Didier Scavarda and Agnes Trebuen</td>
<td>Aix-Marseille University / La Timone Hospital in Marseille</td>
<td>Marseille, France</td>
</tr>
<tr>
<td>Victoria San Antonio, Alexis Arzimanoglou</td>
<td>Barcelona Children's Hospital</td>
<td>Barcelona, Spain</td>
</tr>
<tr>
<td>Pavel Krsek</td>
<td>Motol University Hospital, Prague</td>
<td>Prague, Czech</td>
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<table>
<thead>
<tr>
<th>Physician</th>
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<tr>
<td>Petia Dimova</td>
<td>St. Ivan Rilski University Hospital, Sofia, Bulgaria</td>
<td>Sofia, Bulgaria</td>
</tr>
<tr>
<td>Sergiusz Józwiak</td>
<td>Medical University of Warsaw</td>
<td>Warsaw, Poland</td>
</tr>
<tr>
<td>Philippe Kahane</td>
<td>Grenoble-Alpes University and Hospital</td>
<td>Grenoble, France</td>
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**South/Central America (4)**

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<tr>
<td>Arthur Cukiert</td>
<td>San Paolo Epilepsy Clinic</td>
<td>San Paolo, Brazil</td>
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<tr>
<td>Hugo Pomata</td>
<td>Comprehensive Epilepsy Centre, Institute for Neurological Research, Buenos Aries</td>
<td>Buenos Aries, Argentina</td>
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<tr>
<td>Mario A. Alonso-Vanegas</td>
<td>National Institute of Neurology and Neurosurgery</td>
<td>Mexico City, Mexico</td>
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<tr>
<td>Helio Machado</td>
<td>University of Ribeirao Preto</td>
<td>San Paolo, Brazil</td>
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**Africa (1)**

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<th>Location</th>
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<tbody>
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<td>Jo Wilmshurst</td>
<td>University of Cape Town</td>
<td>Cape Town, South Africa</td>
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**Australia (3)**

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<td>Deepak Gill</td>
<td>The Children's Hospital at Westmead</td>
<td>Sydney, Australia</td>
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<tr>
<td>Simon Harvey/Wirginia Maixner</td>
<td>The Royal Children's Hospital</td>
<td>Melbourne, Australia</td>
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<tr>
<td>Stephen Malone</td>
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**Asia (13)**

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<td>Sarat Chandra</td>
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<td>Gleneagles Global Hospitals</td>
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<td>Xian Lun Zhu</td>
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<td>Lixin Cai</td>
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<td>Beijing China</td>
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<td>Jianxiang Liao</td>
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<td>Japan</td>
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<td>Piradee Suwanpakdee</td>
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**Table III. Levels of Care**

**Level 1.**

**Complexities**

**Age:**

Children aged 9 years and older
Etiology:
Tumors
Focal cortical dysplasia (discrete)
Cavernoma
Hippocampal sclerosis (isolated in medial TLE)
Cysticercosis
Lesion characteristics: well demarcated

Procedures:
Lesionectomy
Temporal lobectomy
Amygdala-Hippocampectomy
Convexity away from eloquent cortex
VNS

Competencies

Personnel:
Child neurologist
Pediatric epileptologist (1)
Pediatric neurosurgeon
Pediatric neuroradiologist
Anesthesiologist skilled with children
Social worker

Equipment:
Routine EEG
Prolonged video EEG
CT
MRI 1.5T

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Facilities:
Designated Pediatric Beds (may be within an adult hospital),
Pediatric ICU

Experience:
Evaluate minimum of 100 epilepsy patients/year
Minimum 15 resections/year

Level 2 (includes all Level 1)

Complexities

Age:
Children aged 8 years and younger

Etiology/Lesion characteristics:
Hippocampal sclerosis associated with dual pathology
AVM
Encephalomalacia (which may reflect prior ischemic or other injury)
Post inflammatory (including post infectious)
Rasmussen’s encephalitis
Hemispheric malformation of cortical development
Tuberous sclerosis complex
Sturge Weber
Hypothalamic hamartoma
Poorly-demarcated lesions (unclear margins/borders)
MRI negative

Location/procedures:
Extra-temporal lobar
Clinical Characteristics of children undergoing epilepsy surgery:

Encephalopathy
Developmental regression/epileptic encephalopathy
Associated developmental or medical syndrome
Genetic abnormalities
Complicated medical problems/(including those medically ill)
Ongoing status epilepticus

Personnel:
Child neurologist,
Pediatric epileptologist (minimum 2)
Pediatric neurophysiologist
Pediatric (epilepsy) neurosurgeon (minimum 1)

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previously trained in functional/stereotactic epilepsy neurosurgeon
Pediatric neuropsychologist
Child psychiatrist
Pediatric neuroradiologist
Pediatric anesthesiologist
Epilepsy neuropathologist
Social worker
Pediatric neurology nursing
Nutritionist/dietician
Coordinator

Technology:
Routine EEG
Long-term video-EEG
CT
MRI 3T
Functional MRI
White Matter Tractography
Angiography
PET and/or SPECT
Some form of source localization capability (not specified between 3D EEG, MEG, or fMRI/EEG)
Cortical stimulation mapping (intra or extra-operative)
Invasive recording capability (intraoperative ECoG, subdural EEG, stereo-EEG)
Neuro-navigation

Resources/ Facilities:
Certified EMU (pediatric) (NOT on adult floor/service)
Dedicated pediatric services
PICU (NOT general ICU)
Rehabilitation facilities

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Experience:

Evaluate minimum of 225 epilepsy patients/year
Minimum 20 resections/year
Operations/Sites/Year

Number of Pediatric Surgery Operations per Year

Number of Sites

1-10 11-20 21-30 31-40 41-50 51-60 61-70 71-80 81-90 91-100 100+

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