

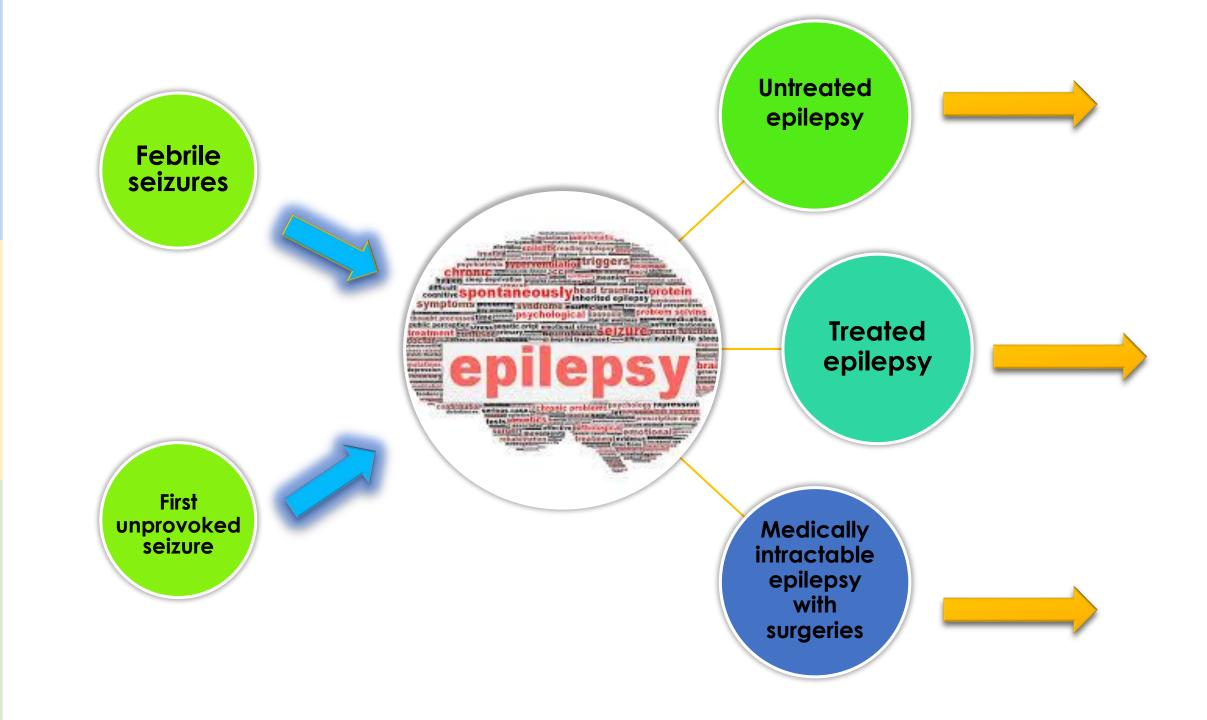


Pillar of the Kingdom



# Natural course and prognosis of epilepsy

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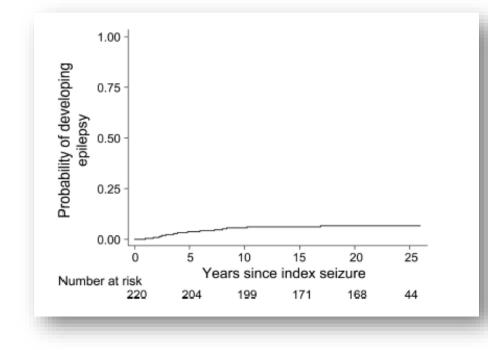
# Natural course and prognosis of febrile seizures

#### Long-term prognosis of febrile seizures

- Population-based studies: risk of later developing epilepsy between 2 and 7% (depending on duration of follow up)
   Chungath M and Shorvon S; Nat Clin Pract Neurol 2008
- 1980s to 2012, with a mean follow-up of 21.6 years
   6% of the children developed subsequent epilepsy (compared with a population risk of 1.4%)
  - age-specific incidence risk of developing epilepsy is almost 10 times (SIR = 9.7, 95% CI 5.7-16.4)

Neligan A et.al; Neurology 2012

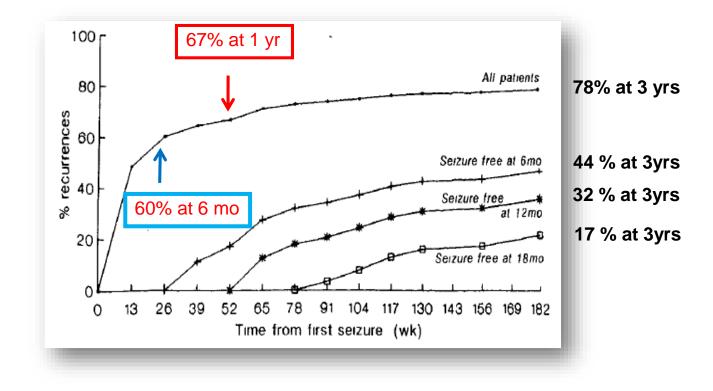
- 12-yr F/U: 1/3 simple febrile convulsions
- Risk factors: 1) number of FC: The more febrile convulsions that occurred, the more likely was subsequent epilepsy 2) complex FC



# Natural course and prognosis of first unprovoked seizure

#### Early prognosis: Risk of recurrence after first unprovoked seizure

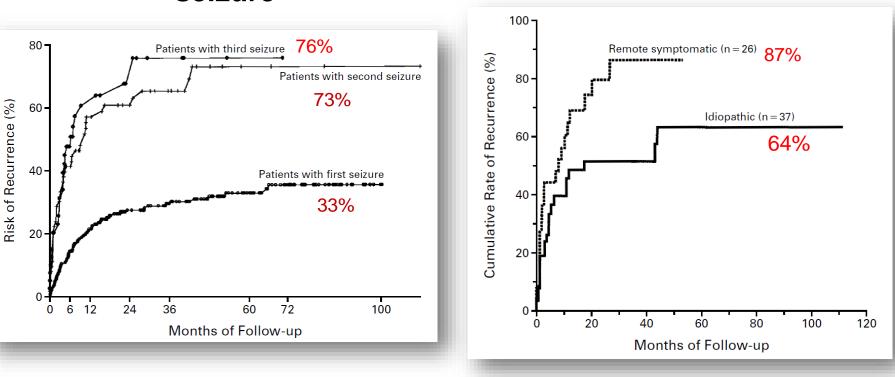
Pooled estimate of 2 year recurrence risk = 42% (30-50%)



Berg AT and Shinnar S Neurology 1991 Hart YM et.al; The Lancet 1990

#### **Risk of recurrence**

#### After first, second and third seizure



**Etiology** 

Hauser WA et.al; NEJM 1998

#### What are the predictors of recurrence?

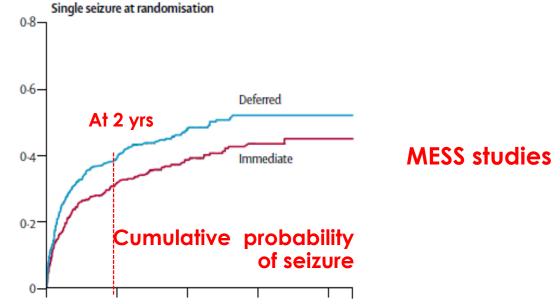
	Predictor	Pooled RR of recurrence	Pooled risk of 2 year recurrence (%)
Abnormal neurological status and abnormal EEG	Abnormal neurological status <u>Normal EEG</u> Epileptiform abnormalities in EEG Non-epileptiform abnormalities in EEG Aetiology and EEG combined		57 27 58 37
	ldiopathic + normal EEG Idiopathic + abnormal EEG Remote symptomatic + normal EEG Remote symptomatic + abnormal EEG	1.9 1.4	24 48 48 65

EEG, electroencephalogram; RR, relative risk.

Camfield P and Camfield C Epilepsia 2000

### Immediate vs deferred treatment after a first unprovoked seizure

Marson A et.al The Lancet 2005



2 yr-remission: 69% vs 61%

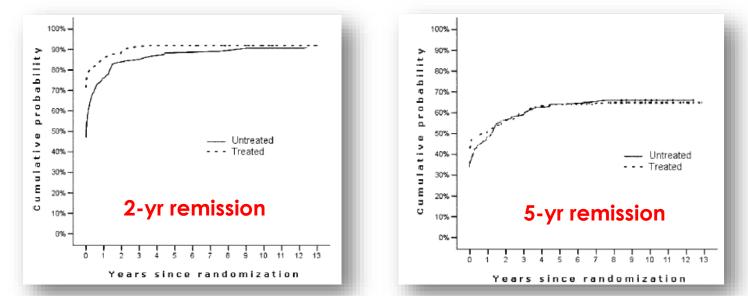
At 3 yrs: 1-3 yrs sz remission: 74% vs 71%

At 5 yrs: 3-5 yr sz remission: 76% vs 77%

"Immediate antiepileptic drug treatment reduces the occurrence of seizures in the next 1–2 years, but does not affect long-term remission in individuals with single or infrequent seizures"

### Immediate vs deferred treatment after a first unprovoked seizure

FIR.S.T. (First Seizure Trial) studies



 Treatment of the first seizure increased the probability of a 2-year remission in the first 3 years; however, the difference disappeared after a longer period of follow-up

(only patients with GTCs were included)

Maurizio AL et.al Neurology

#### **Prediction of risk of seizure recurrence**

	Prognostic Index	
Starting value		
One seizure prior to presentation	0	
Two or three seizures prior to presentation	1	
Four or more seizures prior to presentation	2	
A dd If present		
Neurological disorder or deficit, learning disability, or developmental delay	1	MESS studies
Abnormal EEG	1	
Risk classification group for seizure recurrence*	Final score	
Low risk	0	
Medium risk	1	
High risk	2-4	
"See table 3 for probabilities of no seizure recurrence at specific time points for each o		
Table 4: Prognostic index, with integer values		

"There is little benefit to immediate treatment in patients at low risk of seizure recurrence, but potentially worthwhile benefits are seen in those at medium and high risk"

Kim LG et.al The Lancet 2006

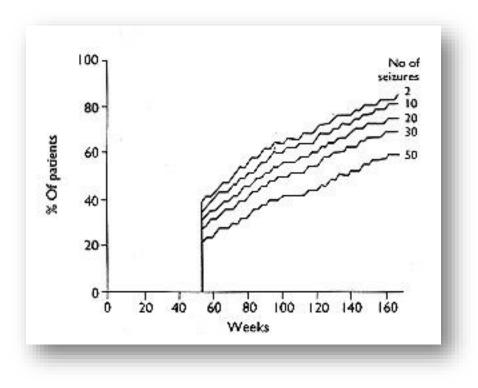
# Natural course and prognosis of untreated epilepsies

#### Introduction

- The course of disorders from onset to resolution, without interventions (Last, 1988)
- Evidence-based treatments with proven efficacy alter the natural course of disorders
- Prospective studies in untreated patients are thus not possible.

## Effect of duration and number of seizure prior to treatment

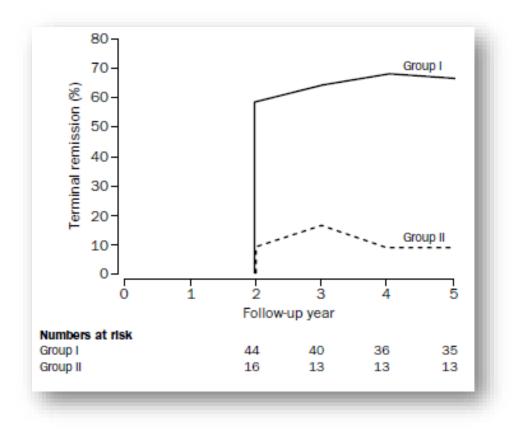
Gower's observation and Reynolds EH studies pointed out that the longer the history of epilepsy the worse the longer term prognosis



Prospective study: 241 adults with newly diagnosed epilepsy treated with one drug

> Gowers WR 1881 Reynolds EH; BMJ 1995 Reynolds EH et.al; Epilepsia 1989

## Effect of number of seizure prior to treatment



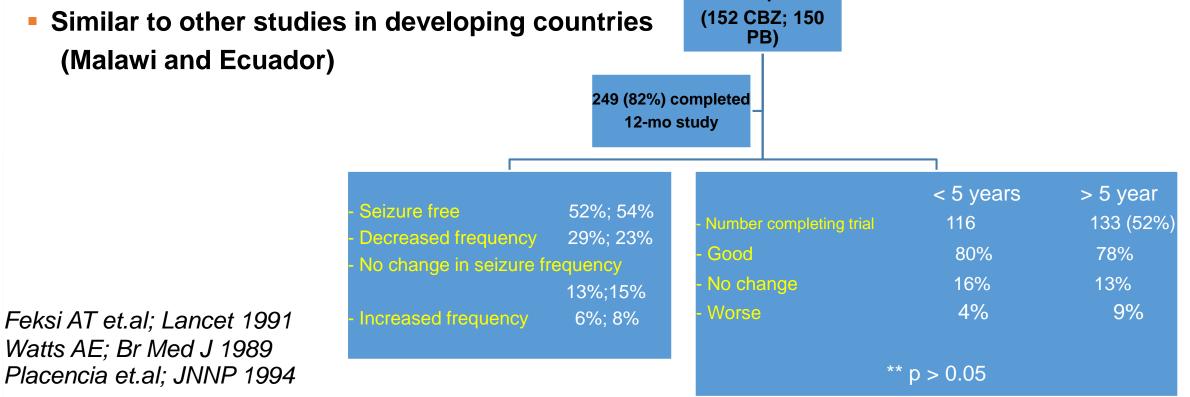
**135 adult patients with partial seizure or GTCs** 

- Treated with either PB or PHT
- Primary outcome: 2year seizure freedom

Group I: good compliance coupled with lifetime total of ≤ 30 GTCs Group II: poor compliance and lifetime total ≥ 30 GTCs

## The effects of AEDs on long-lasting untreated epilepsy

- A study in Kenya (Lancet, 1991) : a finding that does not support the suggestion that the disorder becomes intractable if not treated early.
- Neither length of history of epilepsy nor number of seizures before treatment influenced effect of therapy
   302 pts



# Natural course and prognosis of treated epilepsies

#### **"2001"**

TABLE 2. SUCCESS OF ANTIEPILEPTIC-DRUGREGIMENS IN 470 PATIENTS WITH PREVIOUSLYUNTREATED EPILEPSY.

VARIABLE	No. (%)
Response to first drug	222 (47)
Seizure-free during continued therapy	207 (44)
with first drug	
Remained seizure-free after discontinuation of first drug	15 (3)
Response to second drug	61 (13)
Seizure-free during monotherapy with second drug	41 (9)
Remained seizure-free after discontinuation of second drug	20 (4)
Response to third drug or multiple drugs	18 (4)
Seizure-free during monotherapy with third drug	6 (1)
Seizure-free during therapy with two drugs	12 (3)
Total	301 (64)

 470 patients with newly-diagnosed epilepsy

#### Seizure-free for at least 1 year

- 1<sup>st</sup> drug: 47%
- 2<sup>nd</sup> drug mono: 13%
- 3<sup>rd</sup> drug mono: 1%
- Two drugs: 3%

Medically controlled: 64% Medically refractory: 36%

#### **Effectiveness of AEDs**

#### **"2012"**

Drug regimens	No. of patients	Seizure-free on monotherapy	Selzure-free on combination	Total no. seizure-free	% of cohort seizure-free	% Selzure-free on regimen
First	1,098	543	0	543	49.5	49.5
Second	398	101	45	146	13.3	36.7
Third	168	26	15	41	3.7	24.4
Fourth	68	6	5	11	1.0	16.2
Flfth	32	1	з	4	0.4	12.5
Sixth	16	1	1	2	0.2	12.5
Seventh	9	1	1	2	0.2	22.2
Eighth	3	0	0	0	0.0	0.0
Ninth	2	0	0	0	0.0	0.0

**Medically controlled: 68%** 

**Medically refractory: 32%** 

Despite the introduction of more than 15 new AEDs (since 1985), there is limited evidence endorsing improved outcomes in the common adult epilepsies over the past 30 years

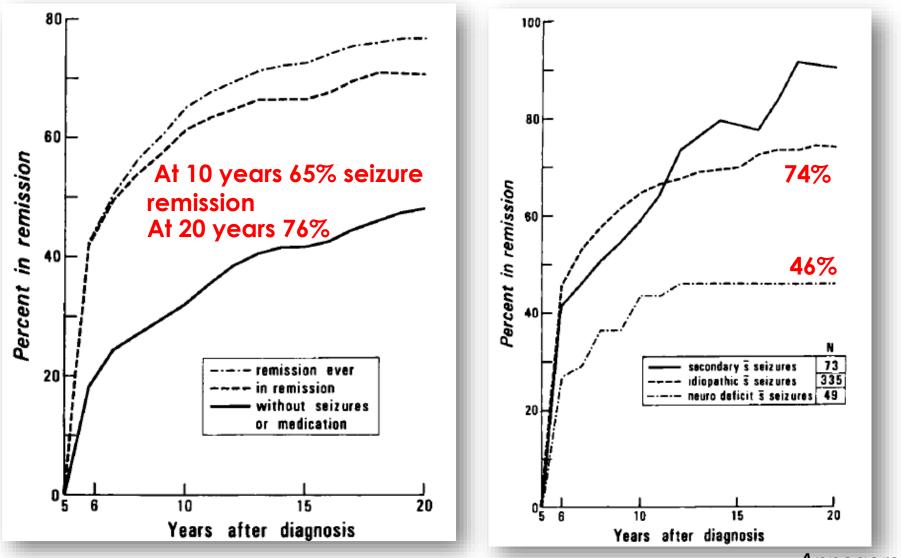
Based on two large studies by the same group, there has been slightly increased rate of seizure freedom from 64% to 68%

Loscher W. and Schmidt D; Epilepsia 2011

#### Late prognosis

- Based on 3 longitudinal community-based studies with long-term follow up
  - Mayo Clinic Record linkage study (US)
  - Tonbridge study (UK)
  - Turku study (Finland) (childhood-onset epilepsy)

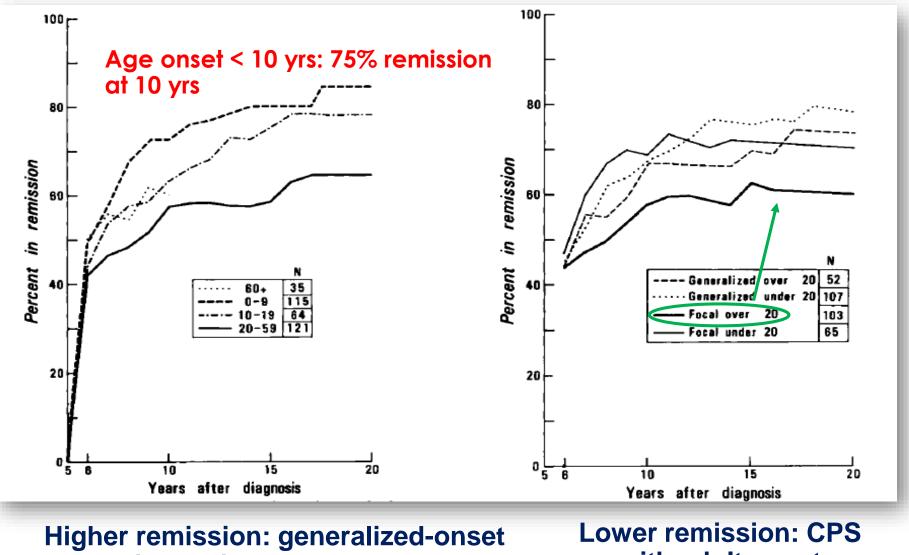
#### Mayo clinic study (20 years follow up)



475 pts
followed at
least 5 yrs
141 pts
20 yrs

Annegers et.al; Epilepsia 1979

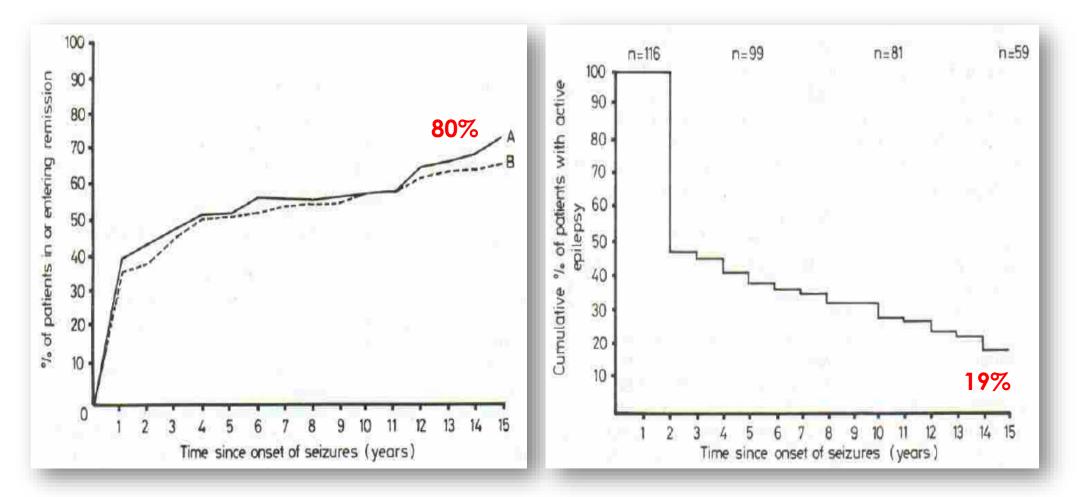
#### Mayo clinic study



seizure diagnosed < 10 yrs

with adult onset

#### Tonbridge study (15 years follow up)



About one fifth (20%) of the patients continued to have seizure (chronic epilepsy)

Goodridge and Shorvon; BMJ 1983

#### **Tonbridge study**

• At 5 years after the first seizure

- of those whose epilepsy was still active, only 21% achieved subsequent terminal remission as compared with 96% of those who were already in remission

" the longer seizure continues to occur, the lower the probability for subsequent remission"

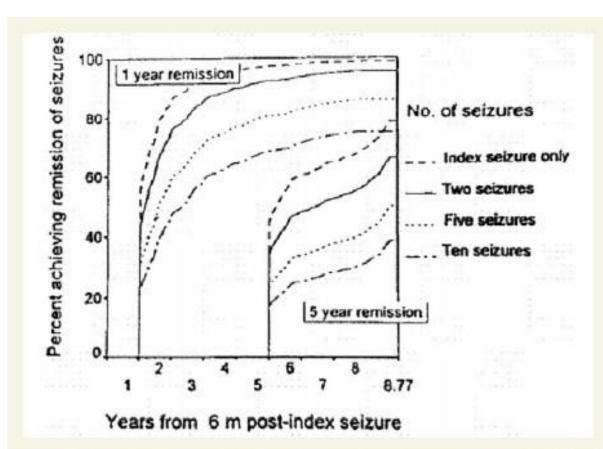
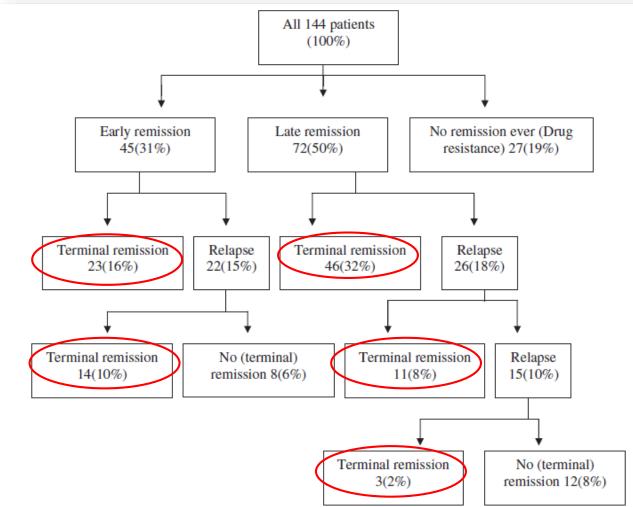


Figure 2 NGPSE: the influence of seizure density on long-term remission. The percentage of patients achieving remission in those who had experienced one (dashed line), two (solid line), five (dotted line), or 10 (dashed and dotted line) seizures in the period from the index seizure to 6 months.

The more number of seizure in the 6 months after the first seizure, the lesser is the chance of long-term remission

### Turku study (37 years follow up)

#### Seizure before the age of 16 years



67% achieved terminal remission (5year seizure freedom at the end of follow up)

- 19% drug resistant
- 19% entered terminal remission after a relapse

Sillanpaa M and Schmidt D; Brain 2006

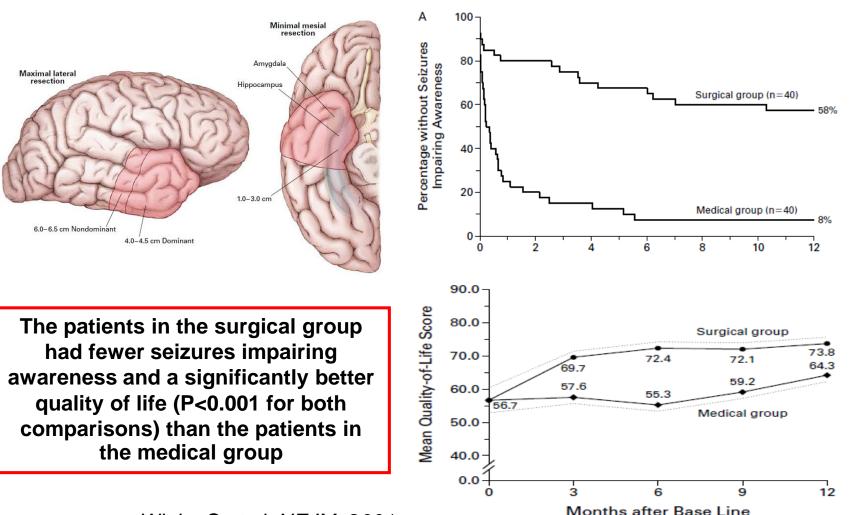
#### Conclusion

• 2/3 of the patients (58-65%) achieved 5-year cumulative terminal remission at 7-10 years follow up

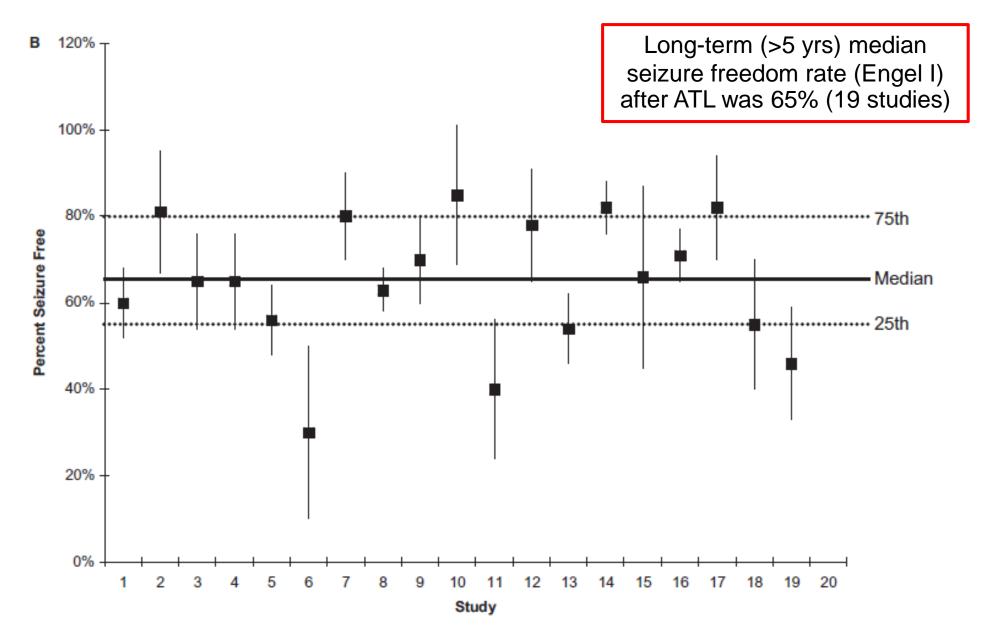
- 3/4 of the patients (67-78%) with childhood-onset epilepsy achieved 3-5 year remission at 12-37 years follow up
- Neurological deficits, age onset, seizure type, number of early seizure influence on long-term remission rate

### Natural course and prognosis of medically intractable epilepsy with surgeries

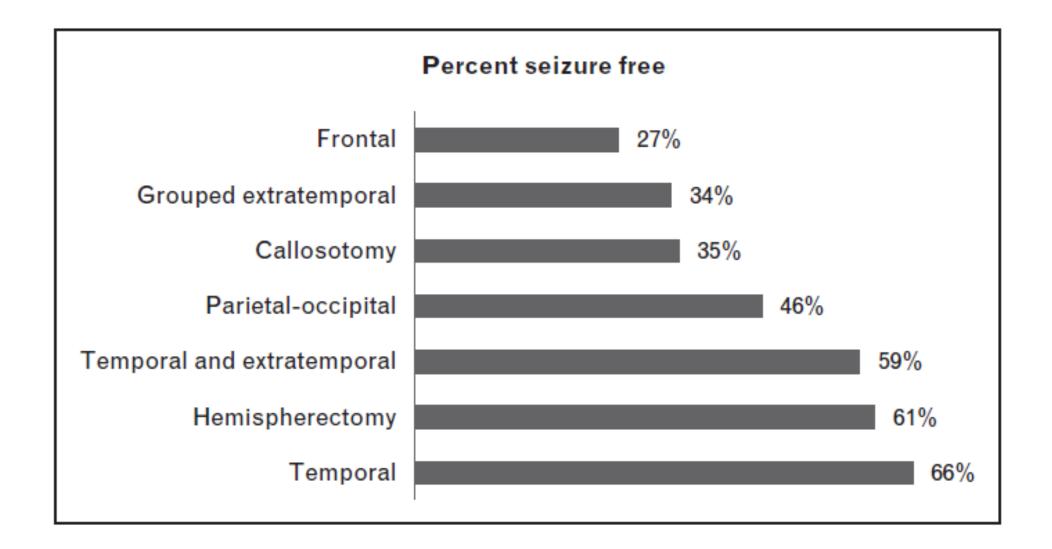
#### **RCT of surgery for temporal lobe surgery**



Wiebe S et.al; NEJM 2001

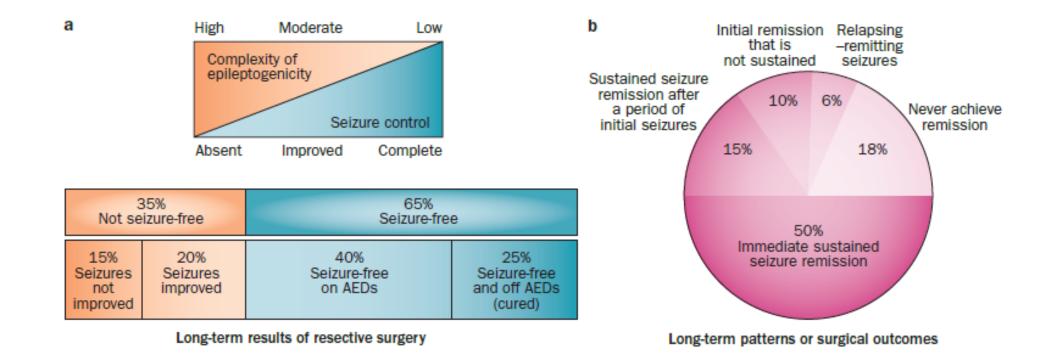


Tellez-Zenteno JF et.al; Brain 2005



Tellez-Zenteno JF et.al; Brain 2005 Weibe S and Jette N; Curr Opin Neurol 2012

## The outcome of surgical treatment in patients with epilepsy



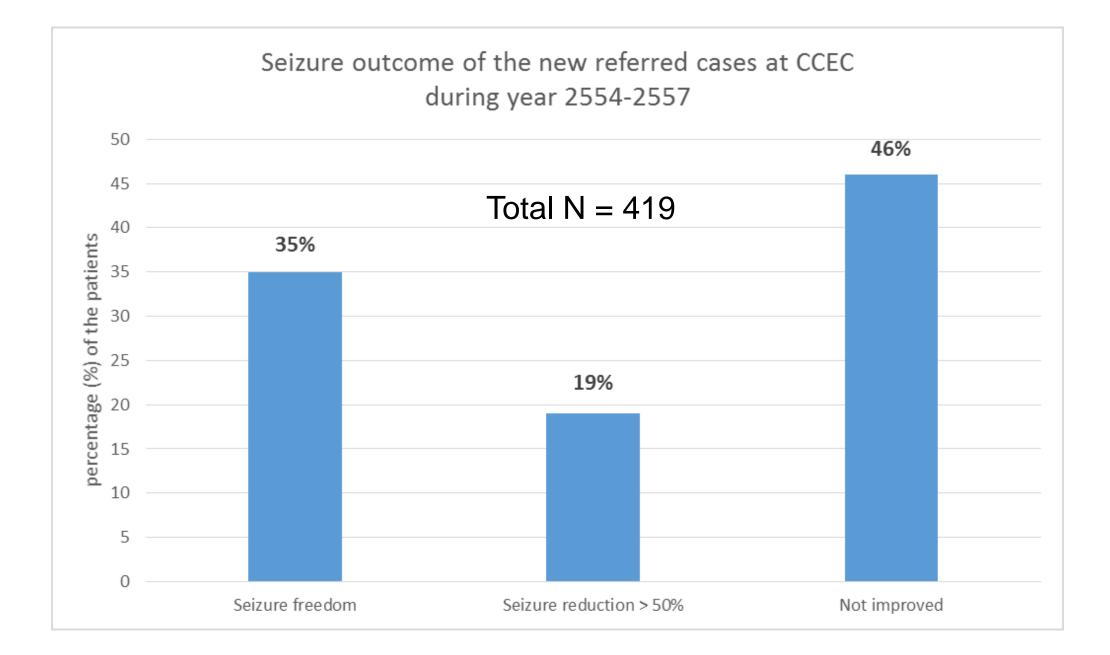
Wiebe S and Jette N; Nat Rev Neurol 2012

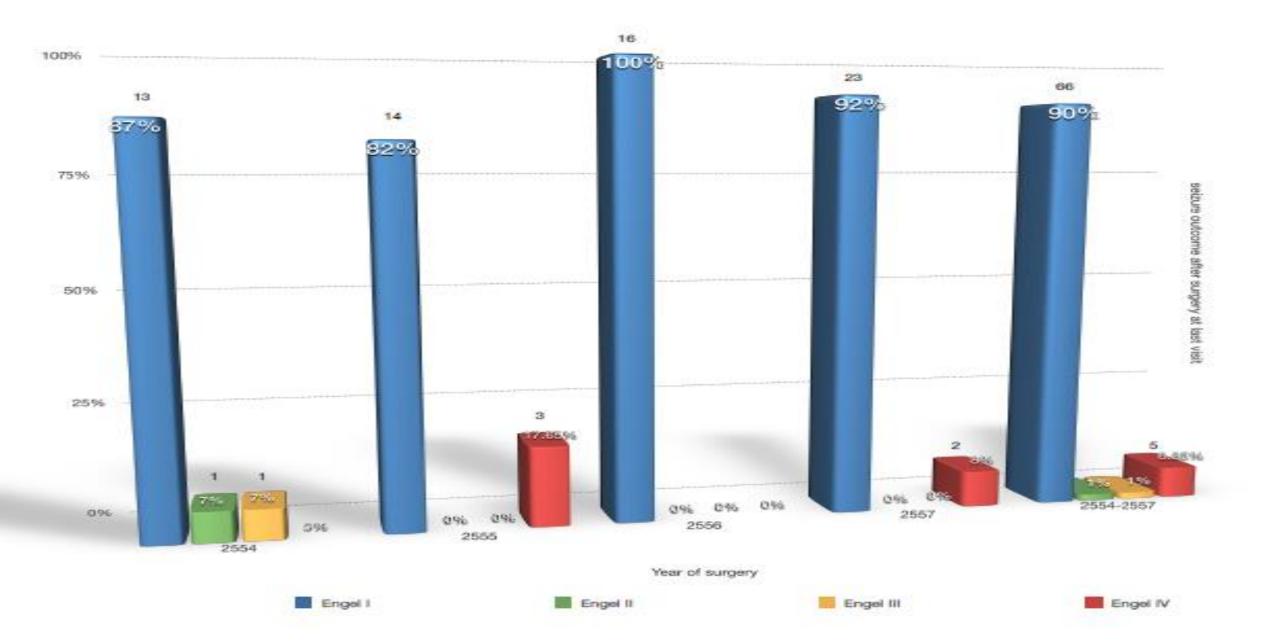
#### **Referred cases at CCEC**

Continuing seizures due to inadequate AEDs adjustment

**Medically intractable TLE** 

Difficult surgical cases requiring extensive investigations





## Long-term courses of Juvenile myoclonic epilepsy (JME)

#### JME

- Original report on JME, published in 1957, Janz and Christian highlighted the rarity of spontaneous remission, as well as the high risk of relapse after AED discontinuation
- Further studies with follow-up periods not longer than 5 years confirmed the initial observation
- Based on 3 recent population-based studies with long-term follow up varying from 25 to 63 years after epilepsy onset
  - 24, 31, and 66 patients respectively (2009, 2012, 2013)
  - 78%, 67.7%, and 59.1% achieved seizure remission

- 25%, 28.6%, and 28.2% were seizure-free off medication for at least the last 5 years

Camfield CS and Camfield PR; Neurology 2009 Geithner J et.al; Epilepsia 2012 Senf P et.al; Neurology 2013

#### **Predictors for seizure recurrence**

- Additional absence seizure at onset of JME
- GTCs preceded by bilateral myoclonic seizures
- A long duration of epilepsy with unsuccessful treatment
- AED polytherapy
- Occurrence of photoparoxysmal responses





### Thank you for your attention