

# Early diagnosis of brain tumours in childhood

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# Some statistics

- ❖ One quarter of all childhood cancers occur in the brain
- ❖ Brain tumours kill more children and young people than leukaemia
- ❖ It takes longer for brain tumours to be diagnosed in the UK than in many other countries
- ❖ Early detection of brain tumours can improve outcomes
  - saving lives and reducing long-term disability

# Challenges in diagnosing childhood brain tumours

- ❖ Relatively rare
  - GPs will typically see only one, maybe two, in their whole career
- ❖ Initial symptoms mimic common, less serious illnesses
- ❖ Varied presentation
- ❖ Fluctuate in severity
- ❖ Differ according to tumour location and developmental stage of the child

# The presenting features of brain tumours: a review of 200 Southampton cases

*Wilne SH, Ferris R, Nathwani\* A, Kennedy CR.*

*Archives of Disease in Childhood, 2006, 91:502-06.*

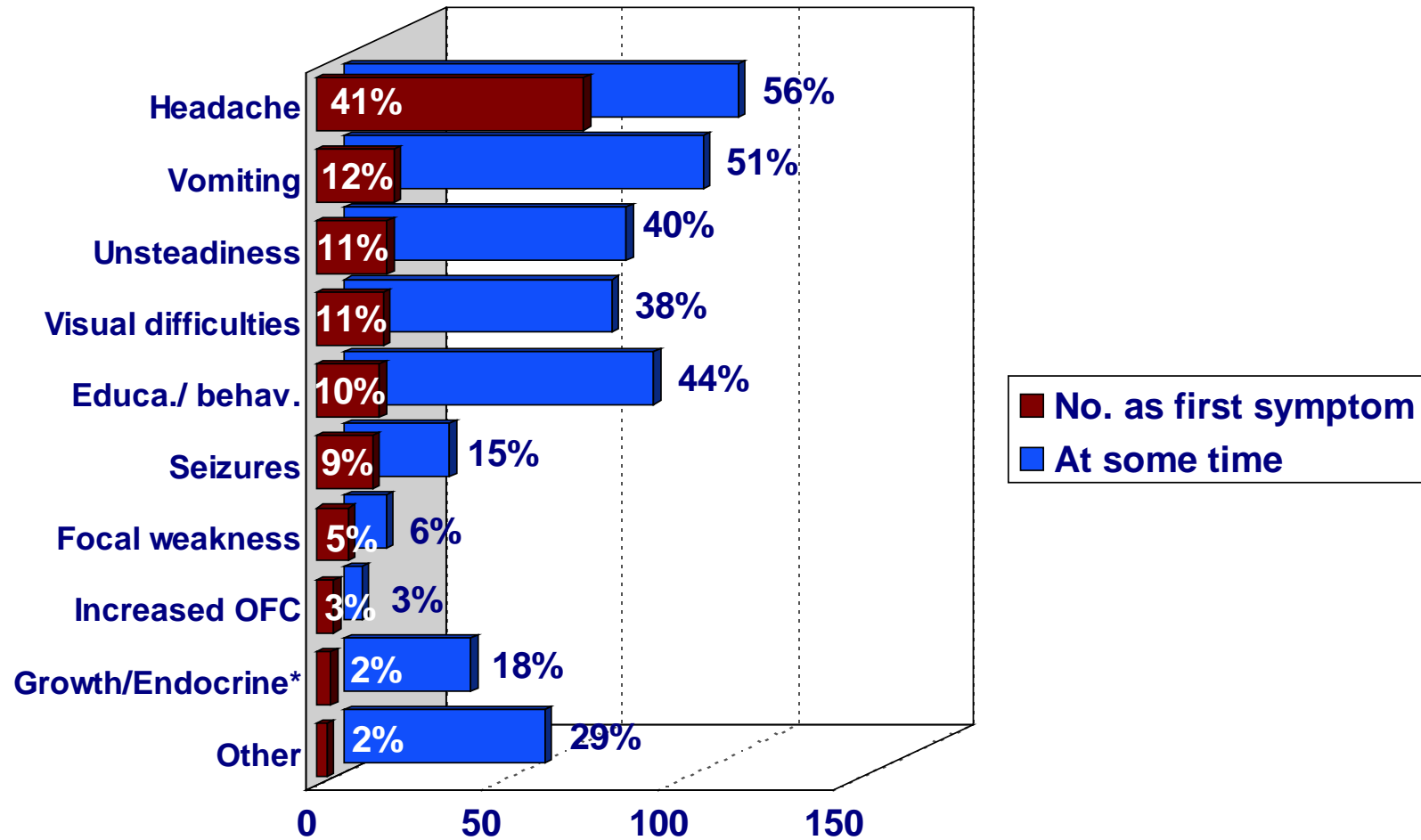
# Methods

- Consecutive admissions to tertiary referral centre in UK
- Retrospective analysis of hospital notes
- Discharge diagnosis of CNS tumour
- >14 year period (Jan 1987 to June 2001)
- 9 symptom complexes

# Results

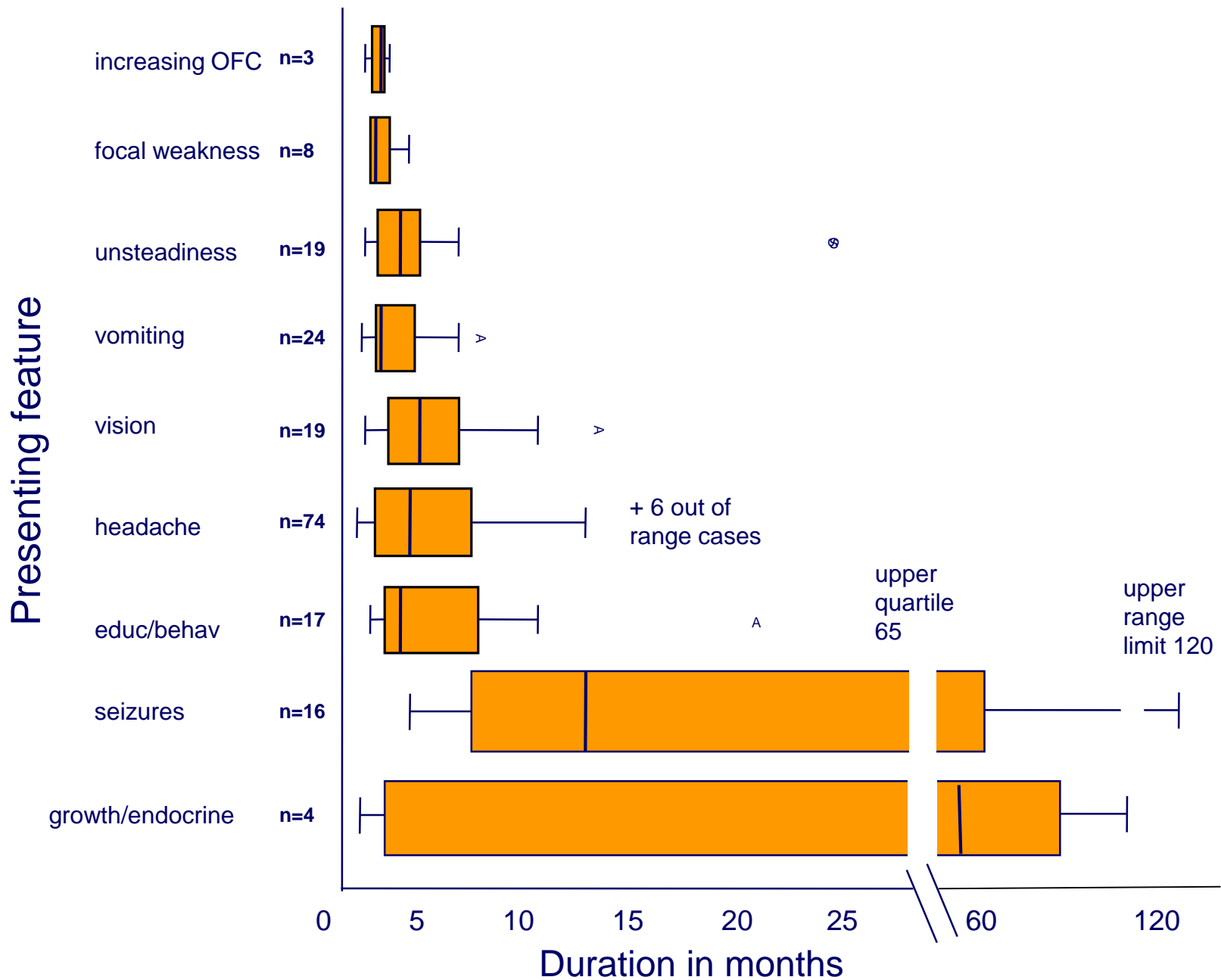
- 204 sets of notes reviewed
- 200 eligible patients
- Age range 15wks – 17yr (mean 7.4yrs)
- Gender 86 F: 114 M (3:4)
- Brain tumour + other diagnosis – 8%
  - 6 NF1, 1 NF2, 1 JCA, 2 TS, 1 DMD, 3 shunts, 2 SLD
- Hydrocephalus at presentation – 43%

# Relative frequency of symptoms in 200 children with brain tumours



\* Includes symptoms of weight loss

Duration of symptoms (months)

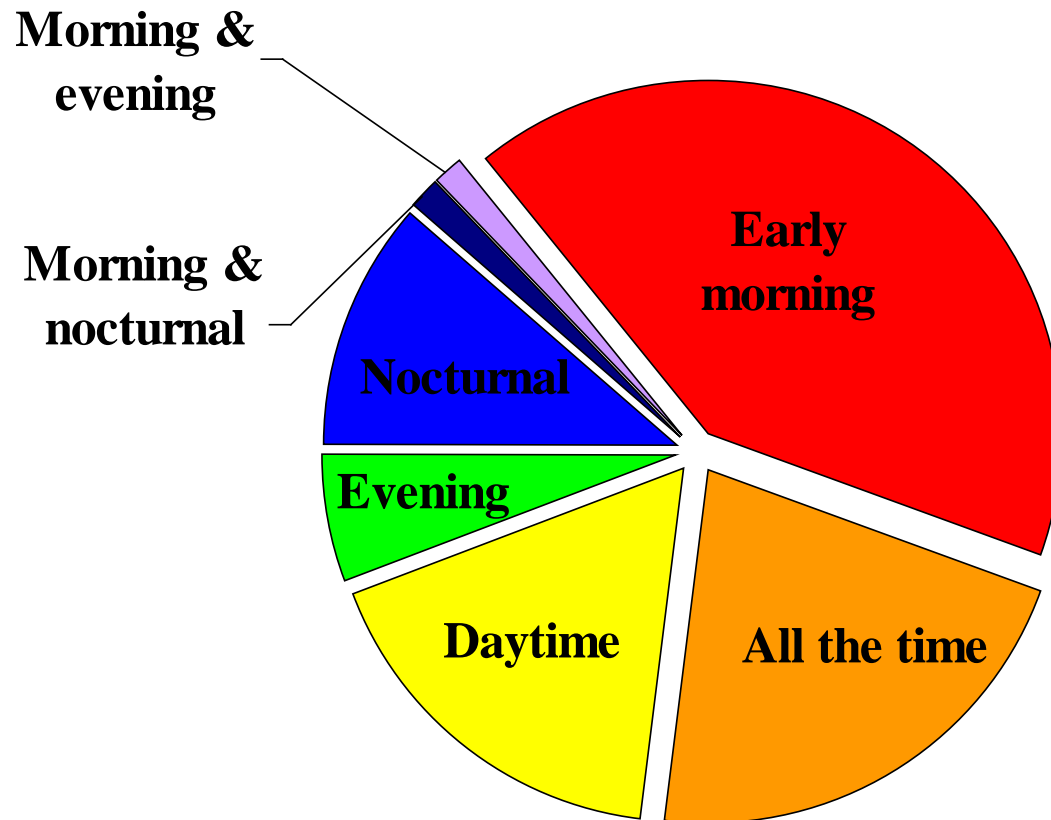




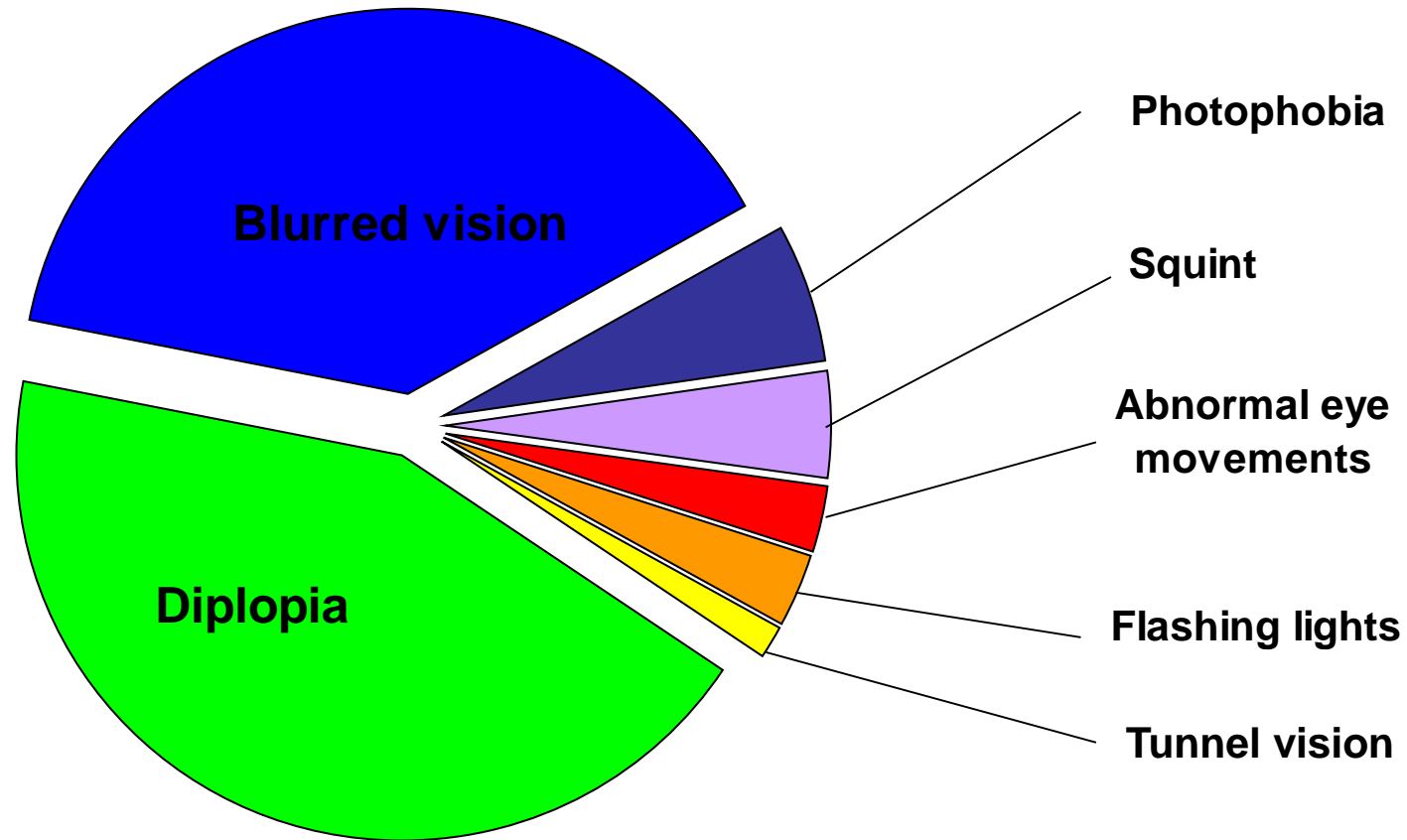
# Factors associated with a median symptom interval of less than the median duration.

Variable	n	Odds Ratio of less than average symptom interval	95% CIs for Odds Ratio	P value
<b>Univariate analysis</b>				
Infratentorial location	94	2.33	1.24 to 4.37	0.009
Posterior fossa tumour	94	3.67	1.39 to 9.71	0.009
High grade tumour	73	5.13	2.62 to 10.00	<0.001
Age less than or equal to 3 years	45	3.94	1.58 to 9.8	0.003
<b>Multivariate analysis</b>				
High grade tumour	73	4.83	2.43 to 9.62	<0.001
Age less than or equal to 3 years	45	3.46	1.32 to 9.09	0.012

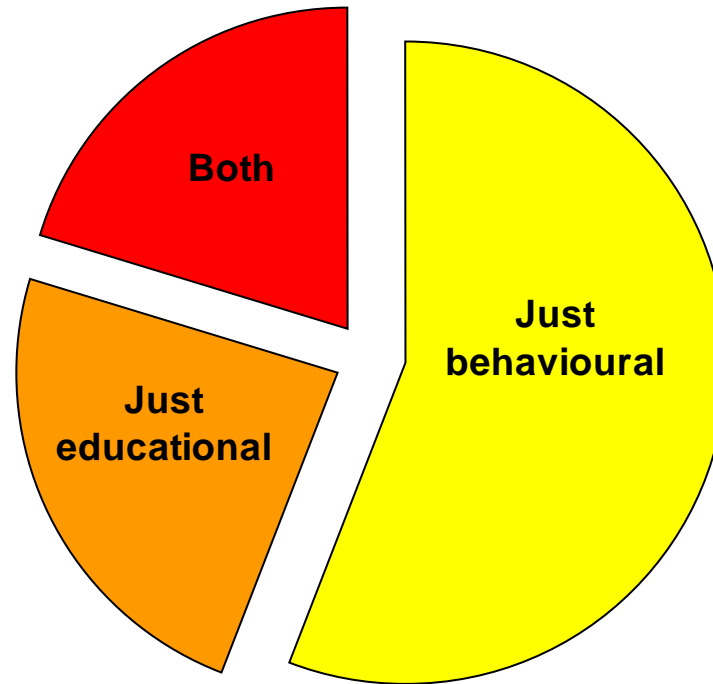
# Diurnal pattern of headache ( n = 70 recorded )



# Visual disturbance (n = 69 / 76 recorded)

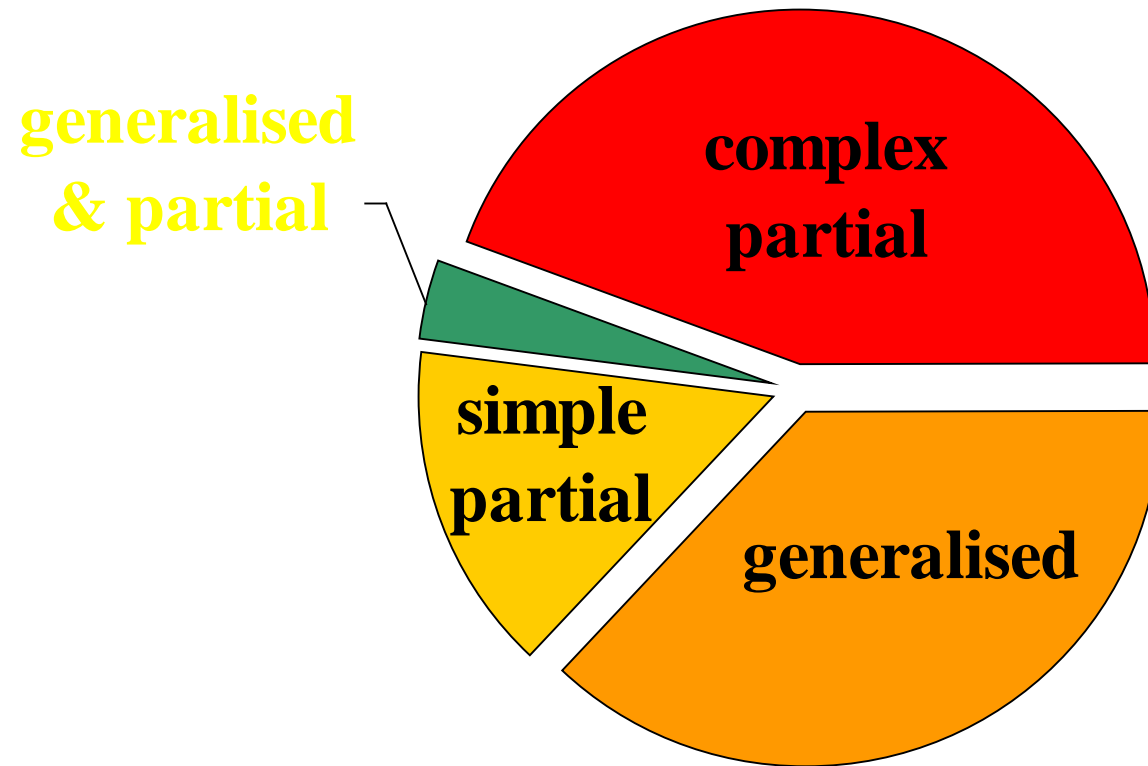


# Behavioural / Educational Problems ( n = 79 recorded )

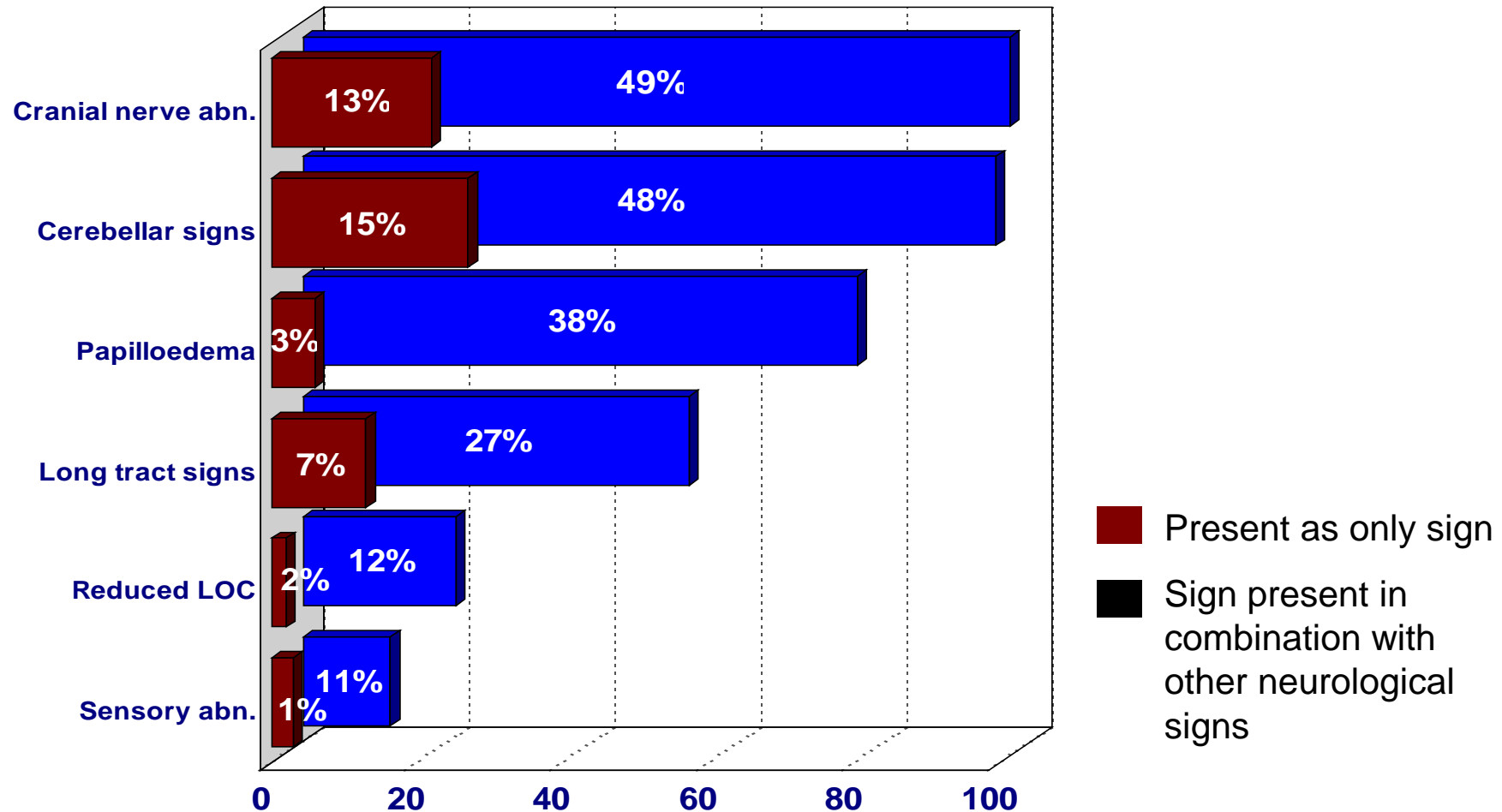


# Seizures

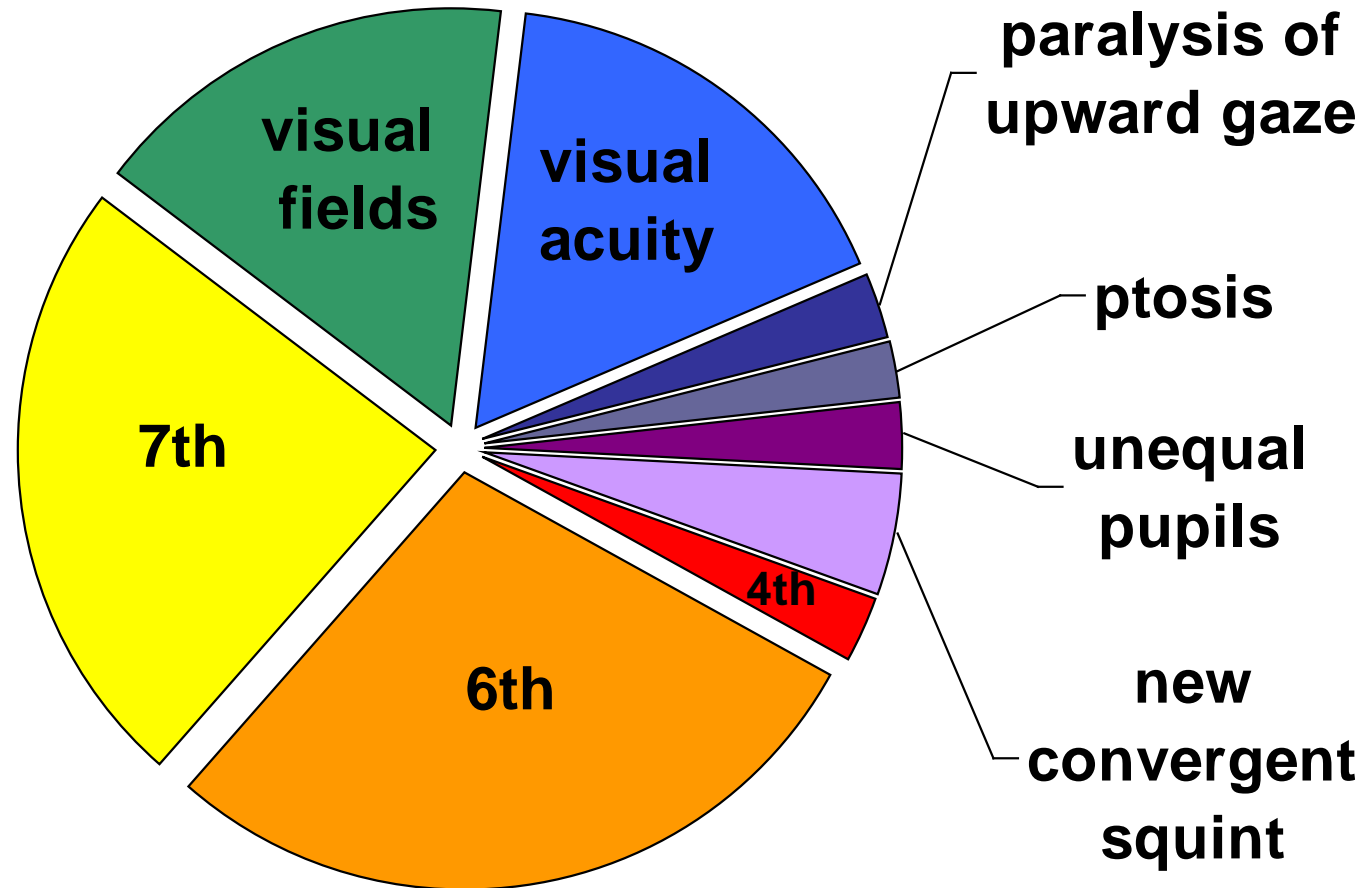
( n = 27 recorded )



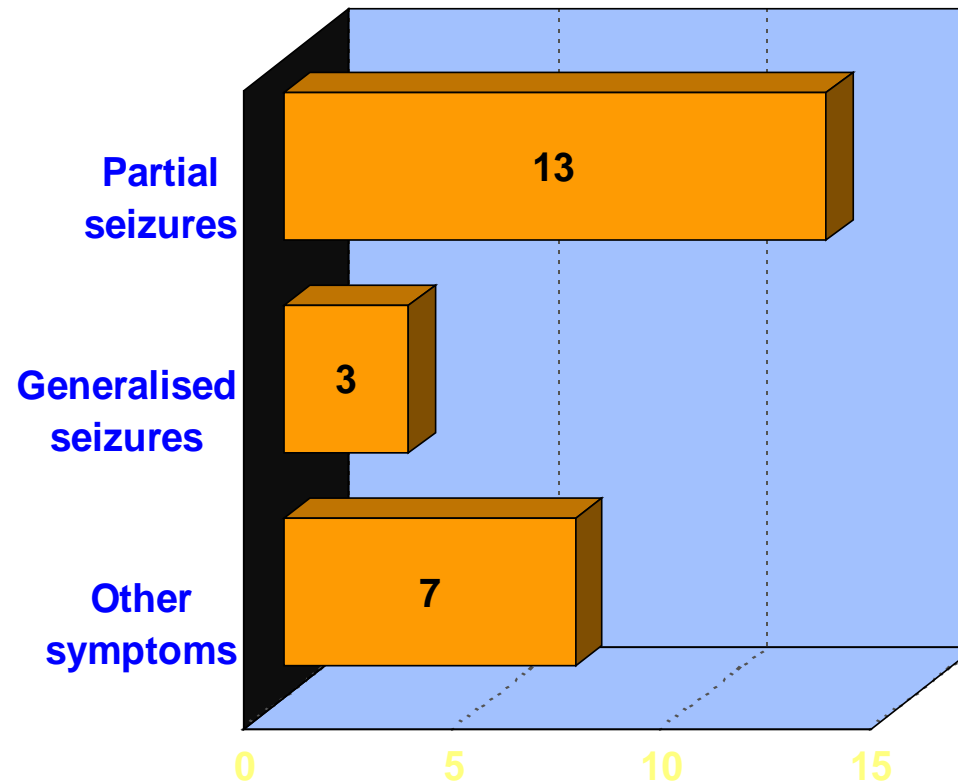
# Abnormal neurological signs n = 175 (88%)



# Cranial nerve involvement (n = 31/97 recorded) - Piechart



23 children (12%) with no abnormal neurological signs



‘MRI is particularly important in those who develop epilepsy before the age of two years...or who have any suggestion of a focal onset on history, examination or EEG (unless clear evidence of benign focal epilepsy).’ NICE guidelines on management of epilepsy , 2nd consultation, April 2004



# Conclusions

- Enquire about not only vomiting and unsteadiness but also visual, educ/behav, endocrine problems
- Examine not only motor system but also cranial nerves (esp fundi, visual acuity & fields, eye movements), growth and head size
- Always consider cranial imaging for partial Sz or gen Sz with either additional Sx or persistent focal EEG

# Summary

- Headaches
  - affect more than half, diurnal in 2/3
  - always assoc with other symptoms esp vomiting (>75%)
- Visual, educ/behav Sx, unsteadiness each affect c. 40%
- Endocrine, educ/behav, visual Sx, H/A or Sz  $\Rightarrow$  longer history
- Abnormal neurol signs present in 90%
  - including 50% with cranial nerve, 50% with crblrr signs
  - 20% had papilloedema and/or cranial nerve signs alone
- Neither abnormal neurol signs nor seizures in v few (3.5%)

# References

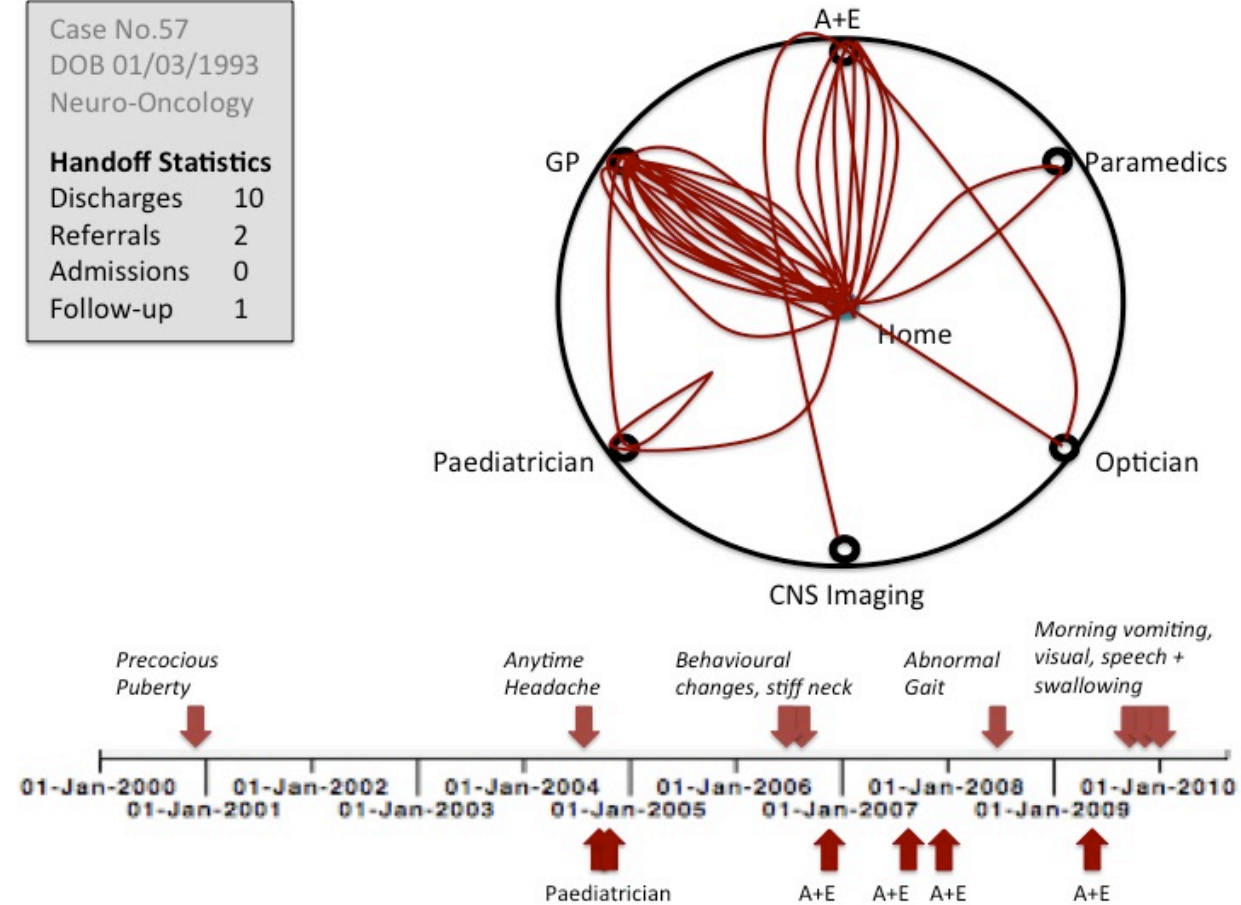
- **The presenting features of brain tumours: a review of 200 cases.** Wilne SH, Ferris R, Nathwani A, Kennedy CR.  
*Arch Dis Child*, 2006, 91:502-06.
- **The presentation of childhood central nervous system tumours: a systematic review and meta-analysis.** Wilne S, Collier J, Kennedy C, Koller K, Grundy R, Walker D.  
*The Lancet Oncology*, 2007, 8: 685-695

# Many children present with signs and symptoms associated with a brain tumour, but are **not immediately diagnosed**

Brain tumours mimic very common childhood illnesses, complicating diagnosis

Consequently children often have **multiple contacts** with health professionals before diagnosis

Case No.57	
DOB 01/03/1993	
Neuro-Oncology	
<b>Handoff Statistics</b>	
Discharges	10
Referrals	2
Admissions	0
Follow-up	1



# Headsmart Project

- ❖ Research project (2002 onwards)
  - ❖ pathways and time taken to diagnosis for children and young adults with a brain tumour
- ❖ Led by Professor David Walker
  - ❖ Professor of Paediatric Oncology,  
Brain Tumour Research Centre, Nottingham (CBTRC)
  - ❖ acts as Clinical Lead on HeadSmart
- ❖ Findings published in *Lancet Oncology*
  - ❖ a systematic review and meta-analysis of the literature on childhood brain tumour presentation – 2007



# HeadSmart Partnership & Campaign Created



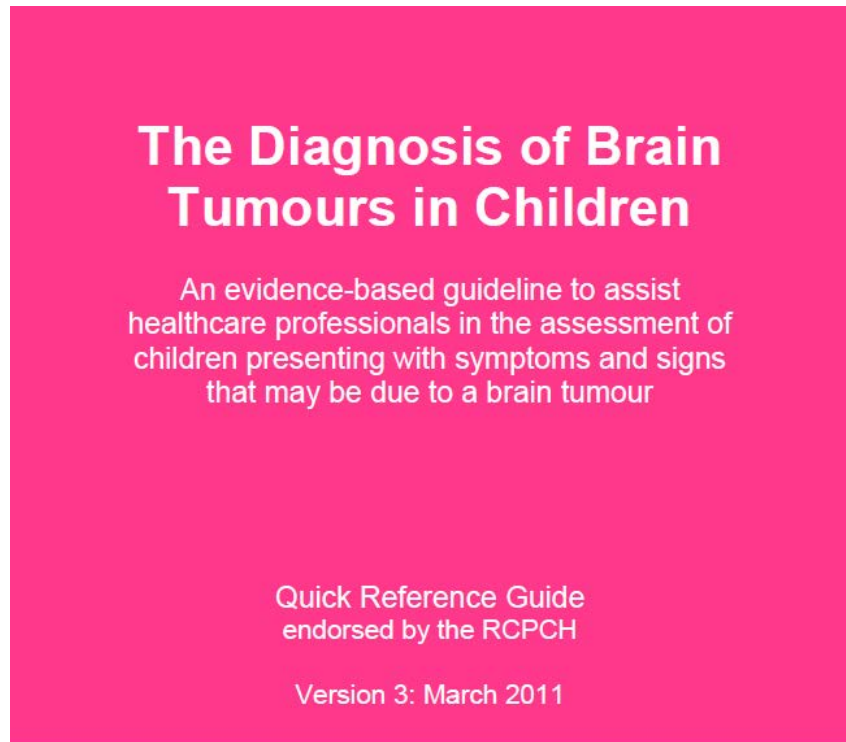
- ❖ Four organisations unite to undertake HeadSmart campaign - 2008



- ❖ Campaign launched - June 2011
- ❖ Working throughout with others



# Guideline Development



- ❖ CBTRC literature review results used in development of **Diagnosis of Brain Tumours in Children guideline** – 2007
- ❖ Supports health professionals
- ❖ Identification of symptoms/ signs of brain tumours
- ❖ Indications for imaging
- ❖ Referral times for imaging
- ❖ Endorsed by RCPCH in 2008
- ❖ Published in ADC in 2010
- ❖ NHS Evidence Accreditation received in 2011
- ❖ Used as basis for development of HeadSmart materials



# Initial Dissemination - **HeadSmart** Launch



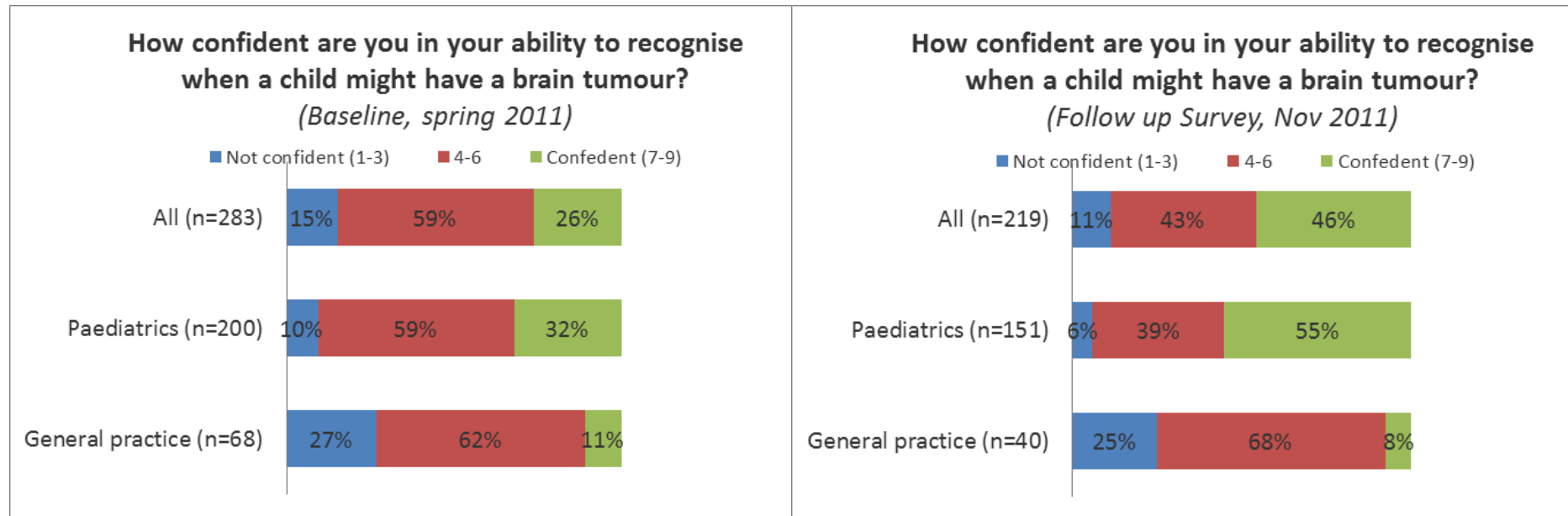
- ❖ Information pack sent to:-
  - Every GP surgery in East Midlands
  - Clinical champions
  - Campaign supporters (through SDBTT)
- ❖ Information pack included:-
  - Explanation of HeadSmart (leaflet)
  - Posters
  - Symptom cards
- ❖ Media campaign (national and regional)



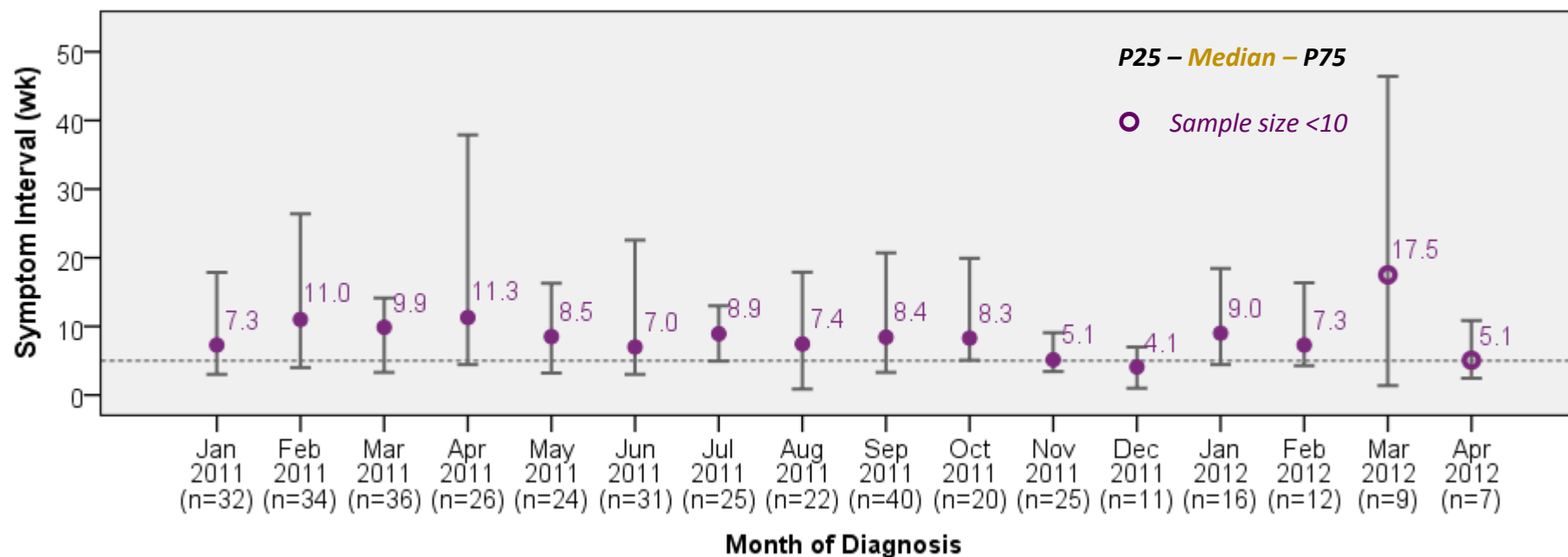


# Evaluating HeadSmart

- ❖ Professional and public awareness surveys
  - ❖ 11% of the public aware of HeadSmart
  - ❖ Health professionals more confident



## Symptom Interval by Month of Diagnosis (weeks)



### Median Symptom Interval

<b>All patients (n=376)*</b>	<b>8.1 weeks (min 0, MAX 398.1)</b>	<b>1.9 months (min 0, MAX 91.6)</b>
Pre-launch (n=155)	9.3 weeks (min 0, MAX 398.1)	2.1 months (min 0, MAX 91.6)
Post-launch (n=219)	7.5 weeks (min 0, MAX 363.6)	1.7 months (min 0, MAX 83.6)

\* Two patients with date of diagnosis missing, therefore numbers don't add up.

# Resources specifically for health professionals

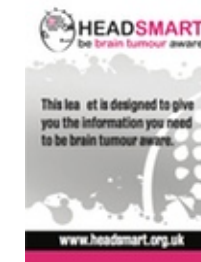
## ❖ 'Diagnosis of Brain Tumours in Children' guideline

- NHS Evidence Accredited, RCPCH endorsed, best practice guideline

<http://www.rcpch.ac.uk/dren%20Guideline>

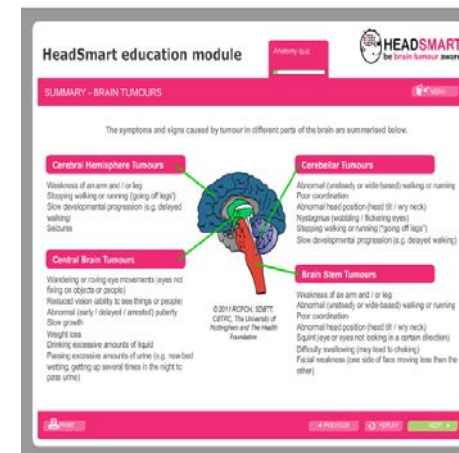


## ❖ Detailed symptoms leaflet



# Resources specifically for health professionals

- ❖ Online interactive Education Module  
[www.headsmart.org.uk/edu/launch.html](http://www.headsmart.org.uk/edu/launch.html)



“Interesting, informative and definitely needed. I would like to forward this to all the paediatricians and GPs I know”

*(Consultant Paediatrician specialising in Paediatric Oncology)*