RCPCH Epilepsy Quality Improvement Programme August 2021 – April 2022

Improving first year of care pathways

ICB West Yorkshire & Harrogate Healthcare partnership

RCPCH Epilepsy Quality Improvement Programme project team:

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Project aim:

Finalise and make progress on a WYH CYPF Epilepsy Group first year of care pathway. Establish a standardised first year of care pathway for patients with epilepsy diagnosis and improving patient information on the process, (subprojects include the areas of transition, reducing waiting times and improving referral pathways for first referral).

Background

- West Yorkshire and Harrogate Health and Care Partnership Children, Young People, and Families (WYH CYPF) Epilepsy Group is an Integrated Care Board (ICB) in England. It brings together all health and care organisations in six places within the Bradford District and Craven Health Care, Calderdale, Harrogate, Kirklees, Leeds and Wakefield. It is not a new organisation, but a new way of working to meet the diverse needs of our 2.7 million people living in the area, of whom 23% (570,000) are children and young people. Building on local plans in our 6 Places, they have a shared commitment and a simple goal; the aim is for everyone in West Yorkshire and Harrogate to have a great start in life and the support they need to stay healthy and live longer.)
- The WYH CYPF Epilepsy Group recognised that there currently is not a first year of care for CYP with epilepsy in the 6 acute trusts or outside of WYH that fully captures the above. As a result, it was agreed to develop the first ICB level WYH CYPF First Year of Care pathway for epilepsy.



Area of focus

- 1. Mapping how we are currently performing against each of the components that make up the pathway.
- 2. Decide whether each Trust focuses on a different component of the pathway and shares learning or pick an area where each of the six Trusts is weakest and focus on improving standards in that area before moving on to another part of the pathway.
- 3. Engagement with CYP and families on the first year of care pathway.

Participant Trusts within the ICS and team projects:

Project aim	Team working on which project aim
Establish a standardised referral pathway to reduce waiting times for	Airedale NHS Foundation Trust
first seizure patients seen via GP and A&E.	
To provide an equitable transition service by using group transition	Bradford Teaching Hospitals
information sessions to young people with epilepsy aged 14, 15 and 16	
years	
To review appointment process and reduce waiting times of over 6	Leeds Teaching Hospital NHS Trust
months and triage overdue follow-up appointments	
To develop a standardised referral system for paediatric seizure, with	Mid Yorkshire Hospitals NHS
the aim of reducing wait times to within national guidance.	
Non-participant	Harrogate and District NHS Foundation Trust
Non-participant	Calderdale & Huddersfield NHS Foundation
	Trust

Project Outcomes	Team working on which project aim
Overall patients seen within 2 weeks of referral (epilepsy and non-epilepsy):	Airedale NHS Foundation Trust
46.34%. They have achieved their aim and are looking into other areas of	
improving their service.	
While participating in the programme, the team received eight responses from	Bradford Teaching Hospitals
patients regarding their ideas on how to improve their transition service.	
Unfortunately, due to significant capacity issues the team was unable to act on	
the views collected from their patients. Young people with epilepsy will in the	
future be provided with explanations of the actions taken by the service	
regarding what they are trying to achieve and enable their understanding.	
Reduced patient consultant waiting list from a backlog of over 100 patient	Leeds Teaching Hospital NHS Trust
appointments, focusing on those with confirmed epilepsy or strongly	
suspicious of epilepsy.	
• Established a reliable list of confirmed epilepsy patients awaiting review that	
is easily accessible by the lead consultant and ESN.	
The internal system for patient access to epilepsy nurse support has been	
demonstrated to be robust and effective.	Nial Vaulahina I I agaitala NII IC
Developed a system to efficiently improve the pathway of new patient referrals received from A&E and the ward to ensure essential details were	Mid Yorkshire Hospitals NHS
included with the referral to allow for appropriate triage within paediatric	
epilepsy clinics.	
 Integrated a new process into an existing Trust guideline for new children 	
presenting in ED and inpatient wards with suspected epilepsy.	
All documents accessible on the internal intranet have clear and consistent	WDODOU TOUT
messaging.	RCPCH EQIP Epilepsy quality improvement programme

Lessons learnt

Communication processes	 Many ICBs face the challenge of engaging all individual Trusts within their networks and understanding the needs of their service and the patients and families that use these services. Having a communications plan in place will help to identify the gaps in communication and information that are lacking within areas and will help to refer to and tailor information and communicate with multiple Trust level teams. Not all NHS staff are aware of who is their ICB lead, and the WYHHP is fortunate that is not the case. Like the WYHHP, ICBs should implement quarterly meetings with individual Trusts to help provide support and learning across their regents and begin to tackle variation by encouraging support for each other and sharing best practices.
Team structures and dynamics	To increase team building and support, ICBs should form an executive team involving the ICB Clinical Lead and ICB Project Manager with individual Trust level teams that at the very least include a clinical lead, ESN, and administrator. • During the course of the EQIP, WYHHP lost their project management support, which made it increasingly challenging to obtain support for the ICB lead. Therefore, it is important that all ICBs have a project manager or coordinator in place to work alongside the ICB lead.



Lessons learnt

Challenges and complexities	 There were many challenges experienced within an ICB. ICB leads have the task of engaging with Trust level service teams that are different in population sizes and demographics. Services that are under-resourced and experience other unique challenges experienced by each service team within their region. This means ICBs will need strong leadership and process mapping tools when navigating these challenges to begin the complex task of reducing variation.
Measuring/Monitoring variation.	 All Trust service level teams within the ICB's should be a part of a national audit or implement processes for measuring and monitoring areas of improvement, working together on their chosen quality improvement topic to share best practices and overcome challenges collectively to begin to reduce variation. Action planning with clear measures that provide baseline local data are good ways to start problem-solving in areas of concern.

Visual presentation of team project intervention

Team video presentation



Project aim

Establish a standardised referral pathway to reduce waiting times for first seizure patients seen via GP and A&E.

Background

Nice guidelines published first in 2004 recommend that all children who have had a first non-febrile seizure should be seen as soon as possible (within 2 weeks) by a paediatric consultant with expertise in epilepsy. This standard is not always achievable. Airedale has had an established paediatric epilepsy clinic since 2005, with an average first referral waiting time of around 10–12 weeks.

Area of focus

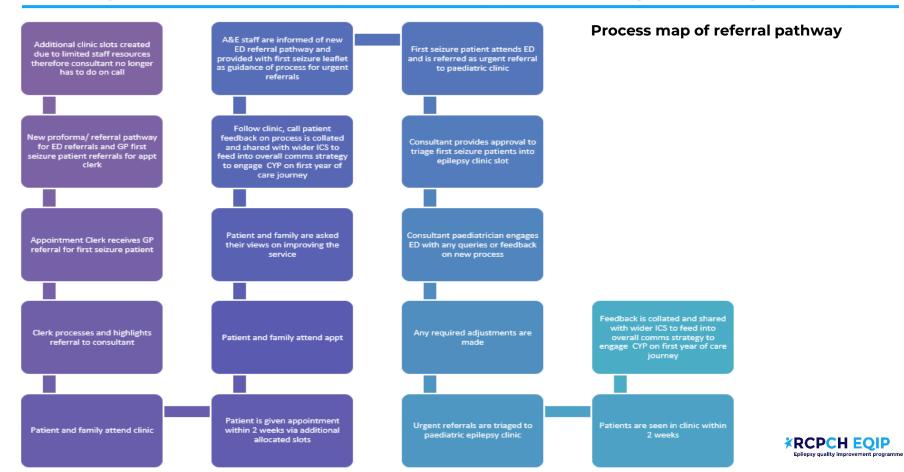
To see at least 50% of patients within 2 weeks after referral.



Changes

- Focusing on two areas of improvement, reducing waiting times and creating a public-facing engagement document that explains the pathway of what patients and families should expect during their first year of care.
- Engaged A&E doctors and junior doctors regarding allocating all new seizure referrals to the team and shared the RCPCH leaflet on epilepsy first seizure.
- Engaged with appointment clerk to set proforma for alerting service of patients with first seizures.
- New patient referrals are seen within 8–4 weeks, but the consultant is temporarily no longer on on-call shifts; therefore, the plan is to see more patients by triaging them with the aim of seeing them in the clinic within 4–2 weeks.
- NICE guidance on two-week referrals is quite challenging; therefore, the project aim has changed slightly for at least 50% to be seen within 4 weeks.
- One appointment slot was kept free for patients referred from A&E/admissions/GP if required to be seen within 2 weeks.
- Monitoring progress via local data on the number of increased patients seen and the reduced waiting time.





Results

- Total patient referred to Epilepsy clinic: 40 (GP, A+E, Hospital admissions)
- From October 2021 to February 2022
- Diagnosed with Epilepsy 5/40 (12.5%)
- Waiting time range: 1-8 weeks. Mean 3.3 weeks
- Waiting time after new measures: 3.3 weeks.
- Waiting time last year 2020: 4.3 weeks
- Waiting time for Epilepsy patients: 1-2 weeks
- Waiting time for non-Epilepsy patients mean: 3.4 weeks (37 % seen within 2 weeks)

Challenges

- During the participation of the programme, this service experienced a lack of ESN support due to a long-term illness, leaving the consultant solely running the service.
- Reducing variation for first seizure referral waiting times from 8 to 2 weeks.





Local data was captured on the number of increased patients seen and the reduced waiting time from 2004 to 2021.



Audit results cohort 3/4 - Professional Input – Airedale NHS Foundation Trust

NICE recommends that children and young people presenting with suspected seizure are seen by a specialist in the diagnosis and management of epilepsies within 2 weeks of presentation (Quality statement 1).

Airedale paediatric epilepsy service have continuously been able to provide CYP with access to a paediatric consultant with 'expertise in epilepsy, offering a defined consultant or associate specialist led epilepsy clinic seeing patients at a secondary level.

The percentage of CYP with input from a Paediatrician with expertise increased from 85% 2021 to 100% in 2022, achieving above regional and national averages.

Percentage of CYP with input from:	2019	2020	2021	2022	2021 – YPEN	2021 – England & Wales	2022 – YPEN	2022 – England & Wales
Paediatrician with expertise OR paediatric neurologist (PI.1)	90%	95%	95%	100%	82%	88%	95%	91%
Paediatrician with expertise	90%	95%	85%	100%	90%	85%	96%	89%
Paediatric neurologist	40%	33%	50%	30%	24%	25%	11%	20%
Epilepsy specialist nurse	100%	86%	90%	95%	83%	80%	96%	80%



Audit results cohort 3/4 - Time since first referral – Airedale NHS Foundation Trust

The percentage of CYP that were seen by a paediatrician with expertise in epilepsy within 2 weeks of first referral increased to 45% in 2021, above regional and national averages. In 2022, 25% of CYP were seen within 2 weeks of first referral achieving above national average. The percentage of CYP that were seen by a paediatrician with expertise in epilepsy at 16+ weeks was 10% in 2021 and 5% in 2022.

Time since first referral to first paediatric assessment	2019	2020	2021	2022	2021 – YPEN	2021 – England & Wales	2022 – YPEN	2022 – England & Wales
0 – 2 weeks	50%	38%	45%	25%	19%	23%	25%	21%
2 – 4 weeks	10%	33%	10%	25%	14%	12%	17 %	11%
4 – 6 weeks	10%	5%	15%	15%	11%	9%	12%	11%
6 – 8 weeks	10%	5%	0%	10%	6%	7 %	12%	8%
8 – 10 weeks	0%	5%	5%	5%	6%	5%	11%	7 %
10 – 12 weeks	0%	0%	0%	10%	4%	3%	5%	5%
12 – 14 weeks	0%	5%	0%	0%	4%	3%	2%	3%
14 – 16 weeks	0%	0%	0%	5%	2%	2%	1%	2%
16 + weeks	0%	5%	10%	5%	20%	13%	9%	13%



Outcomes

- Overall, the service was able to increase the number of patients seen within 2 weeks of referral (epilepsy and non-epilepsy) by 46.34%.
- The service has established improved relationships and engagement with ED staff.
- Increased effective engagement with patients and families when obtaining their views on continuous improvement and how to maintain a high standard of care.
- National audit results show that the percentage of CYP with input from a paediatrician with expertise increased from 85% in 2021 to 100% in 2022, achieving above regional and national averages.
- The percentage of CYP that was seen by a paediatrician with expertise in epilepsy within 2 weeks of the first referral increased to 45% in 2021, above regional and national averages. In 2022, 25% of CYP were seen within 2 weeks of the first referral achieving above the national average.

Lessons learnt

- Collaboration and engagement with other departments (A+E, appointment clerks) have had a positive impact.
- Clear referral guidance and an efficient triaging system helped reduce waiting time.
- Providing RCPCH leaflets after children and young people have experienced a first seizure helped families with their concerns and provided increased knowledge.



Bradford NHS Foundation Trust – ICS West Yorkshire & Harrogate Healthcare partnership (transition)

Project aim

To provide an equitable transition service by using group transition information sessions to young people with epilepsy aged 14, 15 and 16 years.

Background

Previously, the transition information session provided by children's ESN visited young people with epilepsy at home and went through the 'Ready, Steady, Go' questionnaire to guide the discussion, which became ineffective. An internal audit was completed in January 2020, which showed that there were 84 young people with epilepsy aged 14, 15, and 16 years old on the service caseload. Only 12% of patients had received some transition input from the children's ESN. This was due to the service being 50% understaffed (RCN guidelines). Additionally, ESNs were providing group information sessions to their patients, which proved an efficient and effective method of promoting patient education.

Area of focus

Develop a transition information session with the aim of promoting empowerment and understanding of epilepsy to improve the quality of life for young people with epilepsy through education on the psychosocial aspects of living with epilepsy. Promote skills in communication, decision-making, and assertiveness to enhance a sense of control and independence.

Bradford NHS Foundation Trust – ICS West Yorkshire & Harrogate Healthcare partnership (transition)

Changes

- Used an internal database to identify young people with epilepsy aged 14–16 years of age.
- Received eight responses collated by the children's ESN via phone using a proforma to obtain their views on how they want the transition information session to be delivered.
- The team will engage the young people with epilepsy when they attend a clinic appointment to obtain their views.

Ensure we have the following place: Liaise with Use resources colleagues about consent Use feedback such as how the responses from risk assessment Youth Link Invite patients on Run session and patients to Scotland schedule 2 workers a suitable capture ideas resources and review what their session with the (safequarding date/time and feedback needs and ideas keep updated on feedback focus and tech issues) older ones 16+ to set up the how to escalate session well responded issues raised (especially if out of hours)



Bradford NHS Foundation Trust – ICS West Yorkshire & Harrogate Healthcare partnership (Transition)

Results

Responses from YPWE aged 14, 15, and 16 years old

75% would attend group transition information session

Of which 12.5% would attend virtual group transition information session

12.5% would prefer one to one session from ESN

12.5% would prefer online eLearning session

40% would attend group transition information session during school hours

60% would attend group transition information session out of school hours

Of which 20% would attend at the weekend

50% would not like their parent/carer present at transition information session

37.5% would prefer their parent/carer present at transition information session

12.5% don't know if they'd like their parent/carer present at transition information session

Challenges

- The team experienced time restraints due to being understaffed, and with no additional resources available, the team was unable to complete their EQIP project.
- If the data showed that preference was for one-to-one sessions, this would be very challenging for the service to take action.

Bradford NHS Foundation Trust – ICS West Yorkshire & Harrogate Healthcare partnership (transition)

Outcomes

- While participating in the programme, the team received eight responses from patients regarding their ideas on how to improve their transition service.
- Young people with epilepsy will in the future be provided with explanations of the actions taken by the service regarding what they are trying to achieve and enable their understanding.
- Beyond the EQIP, the service team plans to:
 - Continue planning group transition information sessions with patients,
 - Share the results of the data with patients and their families.
 - Distribute a letter of thanks for providing their views.

Lessons learnt

- Young people with epilepsy will need an explanation of what the service is trying to achieve to enable their understanding.
- Young people with epilepsy will need time to speak to the service team, either on the phone or in person in the clinic.
- The service will consider the views of their patients concerning how they want the transition information session to be delivered.



Project aim

To review appointment process and reduce waiting times of over six months and triage overdue review/follow-up appointments.

Background

Due to limited capacity within the team due to reduced staffing and long-term sickness, referrals for new patients have increased to over 16 weeks, and overdue patient follow-up appointments with the consultant waiting lists have increased to patients waiting for over six months. The team chose to first focus on reducing follow-up patient lists while mapping a new referral process for first seizure patients, reducing waiting times from over 16 weeks.

Area of focus

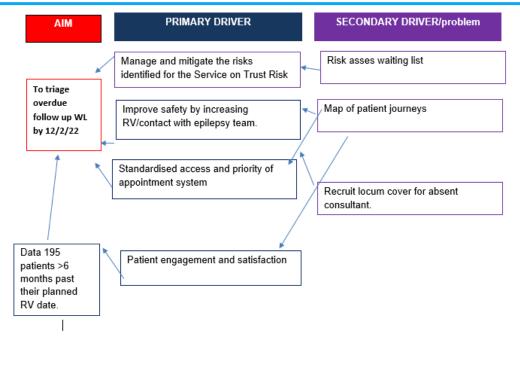
Review of follow-up patient appointment pathways and quality of service provided to patients on waiting lists with the aim of reducing waiting lists.



Changes

- The clinical lead has returned from long-term sickness and plans to review the follow-up patient appointment pathway, supporting the team.
- The team has uncovered that the electronic system WinDip, used within the Trust to process GP referrals, is inaccurately sending referrals to neurology instead of the paediatric epilepsy service team.
- The team had initial discussions regarding the waiting lists to uncover an outstanding 184 follow-up patients that are on average six months overdue.
- The team went through and reviewed the status of each patient on the list and triaged patients that needed to be seen as a matter of urgency.
- The team sent each patient a letter to apologise for the long waiting times and request that families contact the service to inform them of their immediate needs.





Driver diagram of project plan

CHANGE IDEAS

Create whole patient list on PPM for 2nd consultant.

Create list for urgent RVs to be shared with rest of epilepsy team.

Create standard letter template to send to all patients to apologise for delay, explain team cover issue, that patient records are being reviewed and prioritised in order of urgency.

Ask patients to contact ESN via email in usual way if have urgent seizure related issue they have been waiting to discuss in clinic.

Ask patients to contact secretary if they feel they need different review timing.

Ask patients to contact secretary to identify anything they needed whilst waiting for their review (other than a DR review) so we can try to address that.

Identify any patients who could be transferred out of epilepsy clinic to general clinic.

Test and implement SOP for speciality patient WL triage.



Results

Of the 184 patients on the review list:

- 53 had required review or medication advice.
- All had accessed the support offered via epilepsy nurses.
- 97 patients in total reviewed out of 184.
- Nurse clinic 4 reviews none identified they experienced any difficulties whilst waiting for overdue appointment review.
- Feedback: (suboptimal approach) questions were asked directly, either by a consultant or an epilepsy nurse, to the parent/carer rather than the child, but there was clear evidence that all patients had been contacted by their epilepsy service for support when needed.

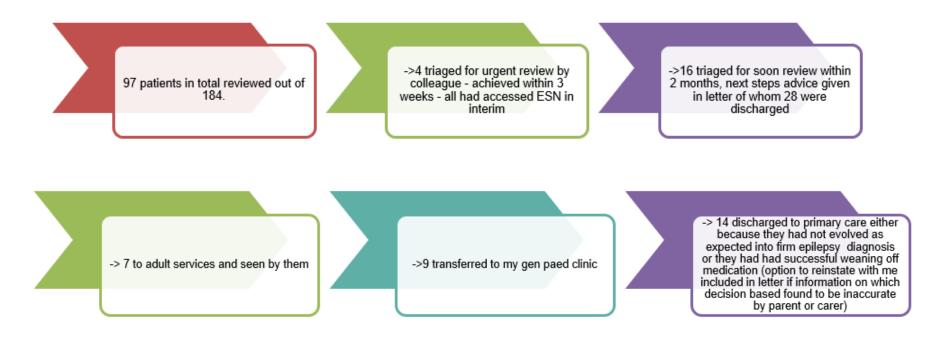


Challenges

- The epilepsy nurse response time had increased from 1 week to 3 weeks for non-urgent responses. This is a direct result of the increased volume of absences and reduced capacity within the team.
- The service had been resilient in terms of patient safety.
- Colleague morale, workload, and resilience are likely to have been adversely impacted.
- The part-time consultant within the team carried the larger epilepsy patient workload. This meant a disproportionate deficit of epilepsy coverage was needed during the period of absence, which impacted the service hugely. There was a failure to recruit epilepsy consultant cover despite two recruitment advertisements circulated within the Trust.



Results





Outcomes

- Through the mapping and review of follow-up patient lists, the service did not experience any serious incidents in terms of patient safety issues.
- Reduced patient follow-up waiting list focusing on those with confirmed epilepsy or strong suspicion of epilepsy.
- Established a reliable list of confirmed epilepsy patients easily accessible by the lead consultant and ESN.
- The internal system for patient access to epilepsy nurse support has been demonstrated to be robust and effective.
- The maximum wait period for a medical review was 18 months. An example of a patient being supported revealed having four ESN reviews with agreed actions and advice provided.
- There is potential for more nurses and non-consultant grade colleagues to increase the number of patients that can be seen and reviewed with specialist epilepsy consultant oversight.
- Beyond the EQIP, the service team plans to:
 - Complete the review of the rest of the epilepsy waiting list.
 - Explore better ways of obtaining patient feedback about the service in the affected time frame,
 - · Explore the possibility of increased colleague sessions,
 - Explore the possibility of a dedicated registrar epilepsy clinic (historically never possible because of recurrent issues at rota level).

Lessons learnt

- · Individual patients had information reflected back to them.
- Patient feedback revealed that they felt they had undergone a review of their circumstances, which equipped the ESN
 or colleague performing the review. This review demonstrated that patients were able to follow the advice given to
 them by the epilepsy nurses before the next appointment date and received it appropriately, which helps to keep
 them safe.

Visual presentation of team project intervention

Team poster



Project aim

To develop a standardised referral system for paediatric seizure, with the aim of reducing wait times within national guidance.

Background

Nice guidelines published first in 2004 recommend that all children who have had a first non-febrile seizure should be seen as soon as possible (within 2 weeks) by a paediatric consultant with expertise in epilepsy. Problems faced at the Mid Yorkshire paediatric epilepsy service include:

- The referral process isn't standardised.
- Families receive little information about what will happen next and what to do if an event happens again.
- Many children are seen in the first seizure clinic when, in fact, they haven't had a seizure.
- Most referrals do not contain sufficient detail to allow appropriate triage to decide if the first seizure clinic is the most appropriate clinic for the child.
- · Lack of knowledge of how to make an appropriate referral.

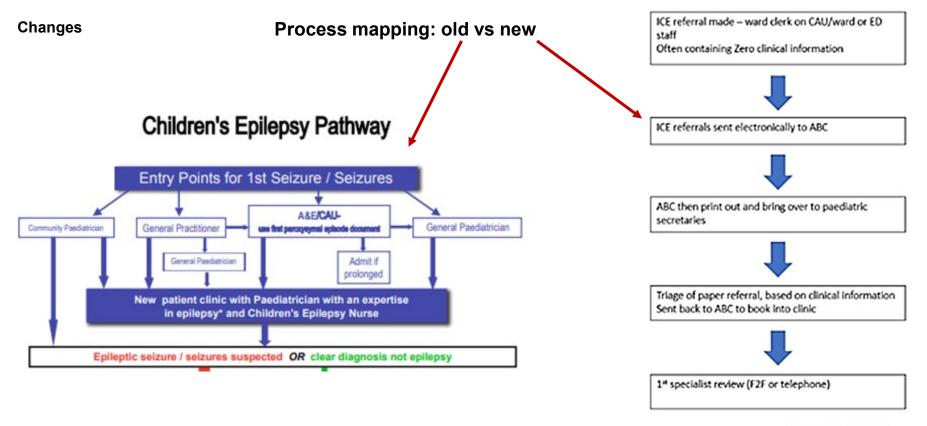
Area of focus

The team's aim is to target the new referrals received from A&E and the inpatient ward in order to make the process more efficient by getting better-quality referrals.

Changes

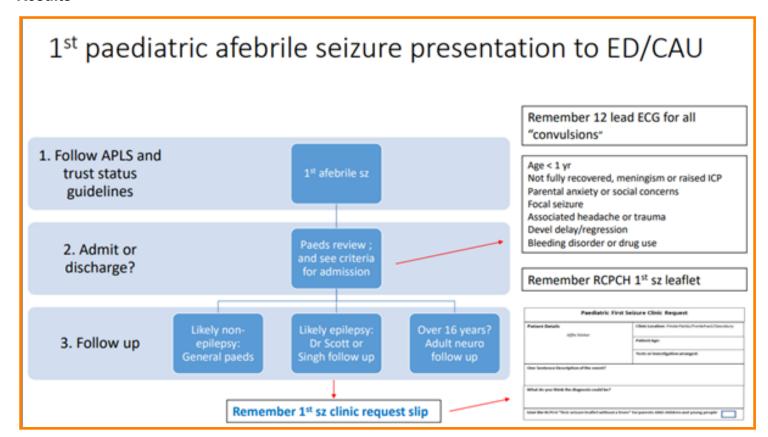
- The team is creating a poster that will be hung in the assessment unit and on the ward on both sites so that staff know what they're meant to be doing.
- The team has created a slip for ED staff to fill in when they enter the referral details on the Trust system called ICE.
- The slip has minimal information. Staff fill in two questions as well as some basic biographic details and include a reminder of the contact details of the paediatric epilepsy service teams to share with patients and their families.
- The team has been engaging with juniors at handover on the wards so that they are aware of the change in process and able to support them with any queries.
- The team will begin evaluating the effectiveness of the process and fine-tuning any details once they begin receiving real-world data.
- The team has sent the paperwork out to the paediatric consultants and middle grade consultants.





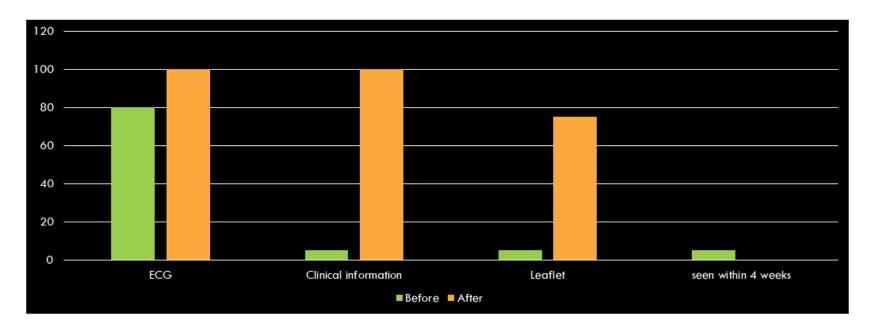


Results





Results



Challenges

Compliance from the team using different methods for referring patients.



Outcomes

- Developed a system in conjunction with current IT systems. After using paper methods to improve, there is a need for electronic referral.
- Educated staff on the use of the slip and encouraged use of the form.
- Incorporating information from paper form into electronic form or adapting adult form
- Integrated a new process into an existing Trust guideline for new children presenting with query epilepsy.
- All documents accessible on the internal intranet have clear and consistent messaging.

Lessons learnt

- It's important to create 'buy-in' from peers and other health professionals when establishing a new process.
- · Testing small changes at a time prevents feeling overwhelmed.
- Collating feedback that feeds into improvement and then measuring the changes is important when abandoning or implementing new processes.
- Ensure you are clear on your purpose and that your colleagues understand the purpose of change to help sustain it.
- · Working well within a support team is beneficial.
- There are many complexities that exist when working within an ICS that need to be reviewed.





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