

Aims. To determine the prevalence of emotionally unstable personality disorder (EUPD) attending a community mental health team (CMHT) in a major Irish city

To describe the current psychiatric care afforded to this cohort of service user

Method. Clinical chart review of all 328 patients attending a CMHT outpatient in an urban setting was carried out. Patients diagnosed with EUPD or displayed features of EUPD were identified. Data on the various interventions offered to this cohort of service users were collected and compared against current guidelines.

Result. Out of the 328 patients actively attending the service, almost 17% ($n = 55$) were diagnosed with EUPD and further 6% ($n = 19$) were found to display features of EUPD such as emotional dysregulation, self-harming behaviour and cognitive distortions. Comorbid psychiatric disorder such as mood or anxiety spectrum disorder was diagnosed in 23% ($n = 17$) of this cohort. Meanwhile, 8% ($n = 6$) was diagnosed with addiction disorders and 5% ($n = 4$) was diagnosed with a comorbid personality disorder. A significant proportion of 77% ($n = 57$) were prescribed psychotropic medication with 51% ($n = 29$) being on more than one psychotropic medication. Antidepressants, antipsychotics and hypnotics were the three most common medications prescribed at the rate of 89% ($n = 51$), 30% ($n = 17$) and 28% ($n = 16$) respectively. A majority of 66% ($n = 49$) were offered intervention from a multi-disciplinary team (MDT) member with 47% ($n = 23$) being offered more than one type of intervention. Referrals to community mental health nurses and psychology service were the two most common interventions offered with a referral rate of 59% ($n = 29$) and 55% ($n = 27$) respectively. 28% ($n = 21$) of service users with EUPD or EUPD traits has had at least one hospital admission while attending the CMHT and 46% ($n = 34$) have been admitted to the day hospital at least once.

Conclusion. The prevalence of EUPD in our outpatient sample corresponds with findings in previous studies. Standard psychiatric care is the most common option available to the majority of general adult patients with EUPD in Ireland due to the lack of any national treatment programme and scarce availability of specialised therapeutic approaches such as dialectical behavioural therapy within community mental health teams. Our CMHT will attempt to integrate mentalization-based treatment into our outpatient management of EUPD patients taking into account current clinical guidelines for management of EUPD and resources needed for training and delivering the intervention.

Evaluating appropriate use of nicotine replacement therapy on acute adult psychiatric units and adverse events related to smoking bans on wards

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Aims. To assess if patients are offered appropriate and adequate NRT (nicotine replacement therapy) upon admission to an acute adult inpatient ward.

To evaluate the number of adverse events occurring which are related to mandatory temporary abstinence from cigarette smoking on inpatient wards.

Background. Not all patients are able or willing to quit smoking when admitted to secondary care. In line with NICE Guidance, psychiatric inpatients should not be permitted to smoke inside the hospital building or outside on hospital grounds. NRT is

the most widely used smoking cessation aid. It aims to temporarily replace the nicotine from cigarettes to reduce motivation to smoke and nicotine withdrawal symptoms. NRT should be offered to patients who need support with nicotine withdrawal during an inpatient stay. Trust guidelines state nursing staff are able to administer NRT to patients on admission without prescription, reducing cravings and withdrawal symptoms such as agitation and anxiety which can lead to adverse events such as aggressive behaviour. Cravings may result in the patients self-discharging or absconding from the ward to smoke.

Method. A retrospective review of electronic records was conducted of all inpatients admitted to two acute adult units over a three month period (65 patients). Patients were identified as smokers or non-smokers. Search words used included: 'smoking', 'NRT', 'nicotine', 'cig' to search for relevant entries. Data collected included whether NRT was offered and given by nursing staff on admission and adverse events related to smoking.

Result. Data from 65 patient admissions were reviewed (31 males, 34 females, mean age 37 years). 37 (57%) patients were identified as smokers. NRT was offered and accepted by 17% of patients on admission and not recorded in 77% of admissions. NRT was declined when offered by 3% of patients.

Adverse events related to smoking occurred in 38% of 'smoking' patients. 40% of these adverse events occurred in first 72 hours of admission. Adverse events include verbal conflicts, physical aggression towards nursing staff and smoking in patient areas.

Conclusion. The majority of patients were not offered NRT on admission or this was not accurately documented in clinical notes. When offered NRT, a large proportion of patients accepted it demonstrating its acceptability amongst this patient group. There is a high rate of adverse events related to smoking on wards. More accurate documentation is required to ensure NRT is being sufficiently offered to patients to reduce possible withdrawal symptoms and adverse events.

Evaluation of paediatric liaison psychiatry services in England 2015-2019

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Aims. Liaison psychiatry services (LPSs) provide psychiatric care to general medical patients. This paper aims to evaluate LPS provision for children and young people in England.

Method. The annual Liaison Psychiatry Surveys of England (LPSEs) included questions on paediatric services from 2015 (LPSE-2). Questions were developed in consultation with NHS England and the Liaison Faculty of the Royal College of Psychiatrists. We analysed data from LPSE-2 and three subsequent surveys.

LPSs were systematically identified by contacting all acute hospitals with Type 1 emergency departments listed by NHS England. All identified LPSs were emailed a copy of the questionnaire, with follow-up emails and telephone contact for non-responders. Responses by email, post or telephone were accepted.

Result. The number of acute hospitals with access to paediatric LPSs increased from 29 (16%) in 2015 to 46 (27%) in 2019; all of these hospitals had access to adult LPSs. The number of paediatric LPSs with at least 11 full time equivalent (FTE) mental

health practitioners (MHPs) has increased from 6% to 24% and from none to 16% with 13 FTE or more MHPs. For both LPSE-4 and LPSE-5, there were only two acute hospitals where both 8 FTE MHPs and 1.5 FTE consultants were present. For LPSE-4, only one site met the Core 24 criteria (for adults - there are no criteria for paediatric LPSs) of 11 FTE MHPs and 1.5 FTE consultants, and for LPSE-5, both these sites exceeded them. Other paediatric services did not meet the adult core 24 criteria for a LPS.

Acute hospitals with access to 24/7 paediatric LPSs increased from 12% to 19% between LPSE-4 and LPSE-5. In LPSE-5 68% of paediatric LPS worked to a one-hour response time target to the ED. This is an increase from 42% (14/33) in LPSE-4.

Conclusion. There are still far fewer paediatric than adult LPSs, but the provision of paediatric LPSs improved from 2015 to 2019, with more services, more staffing, and faster response times. Services need to continue to improve as few services match the adult core 24 criteria for an LPS.

A question of information mismatch in the SPC and PIL on the effect of ADHD stimulant medications on tourette's syndrome

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Aims. To assess the quality of information provided by pharmaceutical companies to patients and doctors regarding the impact of stimulant medications indicated for the treatment of Attention Deficit Hyperactive Disorder (ADHD) on Tourette's syndrome (TS) and tics in children and its implication on treatment.

Background. It is estimated that between 35% to 90% of TS patients also have ADHD. However, there remains a pervasive belief that the use of stimulants to treat ADHD symptoms in children with comorbid tic disorders is contraindicated because of concerns about possible tic exacerbation. Recent studies have disproved this, which is reflected in United Kingdom (UK) and European ADHD and TS guidelines. Pharmaceutical companies are legally required to provide a Summary of Product Characteristic (SPC) and Patient Information Leaflet (PIL) for each medicine as it is an integral part of the marketing authorisation approval. The SPC contains vital information for the usage and prescription of a drug for use by healthcare professionals. The PIL included in the medication packaging is a patient-friendly version of the SPC.

Method. The available stimulant medications licenced for use in paediatric patients with ADHD in the UK were identified through the Medicines & Healthcare products regulatory Agency (MHRA) website. The SPC and PIL were then accessed from the Electronic Medicines Compendium (EMC) website. Those not on the site were obtained directly from the marketing authorisation holder. Any direct mention of tics or Tourette's in the contraindication, warning and caution, or side effect section were documented. The information was then tabulated and compared.

Result. Of the three stimulant drug types, 17 variations are currently available for use in the UK. There were inconsistencies found between the SPC and PIL in reference to the impact of these drugs on tics and TS in all 17 licenced medication. Most discrepancy was found in regard to TS as a side effect (16/17) and

also tics (15/17). TS is also listed as a contraindication in the SPC and PIL for all available variety of Dexamphetamine class drugs. This is inconsistent with current clinical evidence and guidelines.

Conclusion. The disparities in information regarding the impact of stimulant medications on tics and TS can have wide ranging effects. Outcomes could include poor patient adherence, or prevention of initiation of potentially beneficial treatment. It would benefit to standardize the information between these two documents to minimize inconsistencies in understanding between doctor and patient.

An audit into the physical health monitoring of patients who are prescribed antipsychotics in HMP Birmingham

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Aims. To assess the compliance of physical health monitoring with NICE and Maudsley prescribing guidelines for those patients prescribed antipsychotics in HMP Birmingham. To assess secondary objectives including who prescribed the antipsychotics (GP vs psychiatrist), the indication and diagnosis they are prescribed for (licensed or otherwise) and which antipsychotics were usually prescribed.

Background. Patients with psychosis or schizophrenia have a reduced life expectancy of 15-20 years when compared to the general population. The physical health effects of the medication prescribed for these conditions play a large role in this. Physical health monitoring and appropriate intervention is vital to reduce the discrepancy in life expectancy and improve the quality of life of these patients.

Method. Notes of 105 patients in total at HMP Birmingham were reviewed to assess whether the primary outcomes of weight, waist circumference, physical observations, blood tests, medical systems review and education/lifestyle advice were done at the correct times. Secondary objectives of which antipsychotics were prescribed, the profession of the prescriber and the indication for the medications (or diagnosis) were also audited.

Result. Antipsychotics were initiated by both GP's and psychiatrists. Appropriately, there were no prescriptions for clozapine. Olanzapine and quetiapine were the most common antipsychotics prescribed. Not all medications were prescribed for licensed indications and some lacked documentation of both a mental health diagnosis and indications in terms of symptoms. Average BMI of patients was overweight, with BMI ranging as high as 45. The pre-prescription, 12 weekly and annual physical health checks had poor compliance. Those that were completed in line with NICE and Maudsley guidelines were done so by coincidence at the time of diabetic reviews.

Conclusion. The physical health monitoring of patients on antipsychotics in HMP Birmingham is not currently compliant with clinical guidelines. There needs to be improved systems in place for the monitoring of physical health both before prescriptions are initiated and after at the NICE recommended intervals. Amongst other actions, improved computer reminders and training of existing and new team members will be done. The monitoring requirements will be re-audited in 6 months following immediate implementation of the recommendations outlined below.