Valproate, also known as valproic acid (brand names include Epilim and Depakote) is an effective medication used to treat epilepsy¹ and bipolar disorder.² Although unlicensed for treatment of other conditions in the UK, we are aware of ‘off-label’ use for migraine or chronic pain.³

In girls and women of childbearing potential, valproate should be initiated and supervised by a specialist and only when other medications have not been tolerated or have been found to be ineffective.

Unborn babies exposed to valproate during pregnancy are at very high risk (30-40 in every 100)⁴,⁵,⁶,⁷ of neurodevelopment disability - such as lower intelligence and autistic spectrum disorders, and also at risk (10 in every 100) of other birth defects.⁸ This has been increasingly recognised and reflected in strengthened regulatory guidance issued in 2014.⁹ In 2015 the Medicines and Healthcare products Regulatory Agency (MHRA) published the valproate toolkit, providing a set of resources for patients, GPs, pharmacists and specialists.³ This was added to in February 2016⁹ and April 2017 www.gov.uk/government/publications/toolkit-on-the-risks-of-valproate-medicines-in-female-patients.¹¹ These resources emphasise the need to avoid the use of valproate in girls and women of childbearing potential; warn women of the very high risks to the unborn child of valproate in pregnancy; and emphasise the need for effective contraception planning and specialist oversight of changes to medication when planning a pregnancy, as abrupt changes to medication can be harmful.

The MHRA resources have had widespread dissemination. This has resulted in a change of clinical practice in some organisations but evidence suggests a further concerted effort is needed to ensure professionals are informing all girls and women of childbearing age. This evidence includes:

• a survey of women in April 2016 that found of those taking valproate (n=624), 20% were not aware of any of the risks of valproate in pregnancy and <20% had received any of the educational materials¹²
• a National Reporting and Learning System (NRLS) search for incidents involving valproate and reported since January 2015 identified 13 reports that indicated valproate had been prescribed, including two that specifically reported no discussion of the risks in pregnancy had occurred. For example: “Patient … on valproate. No discussion in notes about information or risks given to young female patient taking valproate.”

The actions in this alert ask all organisations to undertake systematic identification of girls and women who are taking valproate, and ensure the MHRA resources are used to support them to make informed choices.

### Actions

**Who:** GP practices, community pharmacies,* acute trusts, mental health and learning disabilities trusts, specialist trusts and all other organisations providing NHS funded-care where valproate is prescribed or dispensed

**When:** To begin as soon as possible and be completed by 6 October 2017

1. Identify how the resources signposted in this alert can be used to support fully informed decisions on the use of valproate by girls and women of childbearing age.
2. Develop an action plan to ensure all girls and women of or nearing childbearing age taking valproate are systematically identified so that all relevant resources can be used to plan their care.
3. Ensure relevant resources are embedded in clinical practice for current and future patients by revising local training, procedures and protocols.
4. By circulating this Alert or through local alternatives (such as newsletters and local awareness campaigns) ensure staff are aware of the MHRA resources and understand their role in local plans to identify all girls and women of childbearing age taking valproate.

*Community pharmacies should deliver all actions that are within their remit, but systematic identification will typically need to be undertaken by the organisation prescribing valproate.

### Sharing resources and examples of work

If there are any resources or examples of work developed in relation to this alert you think would be useful to others, please share them with us by emailing patientsafety.enquiries@nhs.net

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**Patient Safety improvement.nhs.uk/resources/patient-safety-alerts**

See page 2 for references, stakeholder engagement and advice on who to direct this alert to.
Technical notes

Patient safety incident reporting
National Reporting and Learning System (NRLS) searches for incident dates between 1 January 2015 and 31 December 2016 exported to the NRLS on or before 27 February 2017. Extraction used drug and brand names and misspells of valproate, valproic, Depakote, Convulex, Epilim, Episenta, Epival. Three searches were conducted; on incidents reported as death and severe harm for all settings and specialties; on no harm, low harm and moderate harm incidents in obstetric specialties; and on no harm, low harm and moderate harm incidents outside obstetric specialties where the medication keywords occurred alongside keywords related to pregnancy or contraception. These searches identified 15 relevant incidents (nine where there was the potential for pregnancy, and six where pregnancy occurred). Valproate was actually prescribed in 13 of the 15 incidents reported, and two of those reports noted that no contraceptive advice was given.

References

Stakeholder engagement
- National Patient Safety Response Advisory Panel (for a list of members and organisations represented on the panel, see improvement.nhs.uk/resources/patient-safety-alerts/)

Advice for Central Alerting System officers and risk managers
This alert asks for a systematic approach to contacting all affected patients, and therefore needs co-ordinated implementation rather than separate action by individual teams or departments. We recommend that acute or specialist trusts seek advice from their clinical director for neurology, clinical director for paediatrics and medication safety officer who will be able to identify who to direct this alert to. We recommend that mental health and learning disability trusts seek advice from their medical director and medication safety officer who will be able to identify who to direct this alert to. We recommend that GP practices and community pharmacies consider who would be the most appropriate person to co-ordinate local action before wider dissemination of the alert.