

PRESCRIBING FOCUS

August 2024

Supply Issues with Pancreatic Enzyme Replacement Therapy (PERT) including Creon® and Nutrizym® 22 Capsules

As detailed in the National Patient Safety Alert ([NatPSA/2024/007/DHSC](#)) issued on 24th May 2024 and the [Specialist Pharmacy Services Medicines \(SPS\) Supply Tool](#) (registration required), **the resupply date for Nutrizym 22® gastro-resistant capsules is mid-August 2024 and for Creon® 25,000 and 10,000 capsules is 2026**. The NatPSA requests that only **one month's supply** is issued at a time to help stock distribution. The DHSC had also issued [Serious Shortage Protocols](#) for Creon 25,000 and 10,000 capsules restricting the quantity that can be provided to patients to one month's supply.

A joint societies' [Position Statement](#) includes some useful clinical advice regarding these stock shortages. Of interest within this document is the **equivalence table** on page 7 and **content of imported pancreatic products** in Appendix 4 of the Clinicians' Position Statement. Patients with Pancreatic Exocrine Insufficiency prescribed PERT are likely to need a change in their repeat prescriptions due to the changing availability of products. Unfortunately, this may need to be altered on each prescription depending on product availability. SPS has also produced a [conversion tool](#) to help identify equivalent and alternative products for patients currently taking any UK licensed PERT. There is also advice on the [Community Pharmacy England website](#). This helpfully **includes links** to the websites of the specialist importers that have **confirmed they can source unlicensed preparations**.

The ICB would support practices prescribing unlicensed imports where licensed alternatives are unavailable, in line with the NatPSA/SPS advice. However, they will need to [liaise with the prospective dispenser](#) to confirm which products they can source. Where a prescriber wishes to prescribe a specially manufactured or imported product, an **FP10 paper prescription** will need to be issued including the words "**Specified imported product (Special Order)**". Practices are requested **not to refer** patients to secondary care to obtain supplies unless they usually obtain prescriptions from the hospital.

Medicines Safety

Avoiding look-alike, sound-alike (LASA) errors through safer prescribing. (PrescQIPP Bulletin 349)

PrescQIPP has produced [this bulletin](#) (May 2024) to provide "best practice" support on minimising the risk of look-alike, sound-alike (LASA) prescribing errors occurring due to similarities between medicines, which can result in patient harm.

LASA errors can occur at any stage of medication use or processes: prescribing, transcribing or documenting, ordering, dispensing, administering and monitoring. The bulletin focuses on LASA errors at the point of prescribing.

The PrescQIPP website (registration required for login) provides a wide range of information including evidence-based clinical guidance.
See: <https://www.prescqipp.info/>

For further information, speak to your ICB MO pharmacist/pharmacy technician.

Did You Know?....

Ivermectin.

[Coventry and Warwickshire APC](#) have designated the recently licensed **Ivermectin 3mg tablets** as specialist advised for 2nd line treatment of scabies if treatment with permethrin 5% cream has failed.

Topiramate and High Potency Topical Steroids.

June's [MHRA Drug Safety Update](#) announced the introduction of a new Pregnancy Prevention Programme (PPP) for **topiramate**. **Topiramate is now contraindicated in pregnancy and in women of childbearing potential unless the conditions of the PPP are fulfilled.**

May's [MHRA Drug Safety Update](#) reminded healthcare professionals of the possibility of severe side effects from the prolonged use of **high potency topical steroids**.

Tirzepatide (Mounjaro®)

[Coventry and Warwickshire APC](#) have designated Tirzepatide (Mounjaro®) as **specialist advised** for the treatment of **Type 2 diabetes in adults**. Prescribing tirzepatide solely for **weight loss** (in the absence of type 2 diabetes) is **not clinically supported or funded by C&W ICB** until [NICE](#) have evaluated its use for this indication. Tirzepatide is administered by **subcutaneous injection once a week**. The Mounjaro® KwikPen® contains **4 doses (one pen = 4 weeks supply)**.

All strengths of Mounjaro® are currently available: 2.5 mg/0.6ml, 5mg/0.6ml, 7.5 mg/0.6ml, 10 mg/0.6ml, 12.5 mg/0.6ml, 15 mg/0.6ml as Mounjaro KwikPens® solution for injection 2.4ml pre-filled pens.