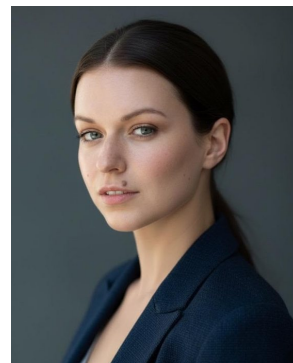


Your licensed EU wholesale partner.

GDP-certified pharmaceutical wholesaler in Vilnius, Lithuania, licensed since 2 October 2026. Cold chain, immunologicals and blood-derived products permitted.

Justina Krukonytė-Racz – Director, licensed pharmacist. Former Responsible Person for wholesale distribution at Tamro-Phoenix, Lithuania. Board member, Lithuanian Pharmacists' Association.



Licence – public record extract (VVKT, Lithuania)

Licence holder	Clusi Pharma, UAB. Company code 307404023
Licence	Wholesale Distribution Authorisation No. 0985, issued 2 October 2026, valid
Permitted activities	Procurement, supply, export
Permitted products	With an EEA marketing authorisation; without one, for export or named-patient supply
Special categories	Cold chain, immunologicals, blood-derived products
Operating site	Savanorių pr. 178, Vilnius, Lithuania (EU, Schengen, Eurozone)
Public record	vapris.vvkt.lt/vvkt-web/public/subsidiaries/0985/5724

What we will do

- **GDP wholesale partner.** Once your product is authorised and released in the EU, it moves under our wholesale licence, GDP system and Responsible Person – procurement, storage and supply across the EU, and export.
- **Coordinate the import side.** Non-EU product needs an import authorisation holder (MIA) and Qualified Person batch release. We coordinate MIA and QP partners so the chain is compliant end to end.
- **Party to your transaction, for a fee.** Buyer, supplier or exporter, whichever role the deal needs.
- **Facilitate.** Introductions, structure and documentation, from CPHI to product on a shelf in Europe.
- **Cold chain.** Low-temperature handling is on the licence. Vaccines, biologics, insulin.

What we will not do

- **We won't fund your project.** No capital into registrations, licensing, stock or expenses.
- **We won't act as your marketing authorisation holder.** A wholesale licence moves authorised products; it doesn't authorise them.
- **We won't import or release batches ourselves.** Non-EU products require an import authorisation and QP batch release; we coordinate with partners who provide this.
- **We won't cut corners on compliance.** Every supplier, customer and shipment is qualified under GDP.
- **We won't say yes to everything.** If your product can't move through our authorisation, you'll hear it in the first conversation.

How it works

1. **First conversation.** You tell us the product, its regulatory status and the markets you need. We tell you plainly whether it can move through our authorisation.
2. **Qualification.** Supplier and customer qualification, quality agreement, and the import route (MIA and QP) agreed with partners where the product comes from outside the EU.
3. **Product moves.** Procurement, GDP storage and supply across the EU, or export, under our licence and Responsible Person.

Let's continue the conversation. Most partners reach me on WhatsApp.

WhatsApp / call +370 655 06475 · justina@clusipharma.com · clusipharma.com/cphi

Also open to discussing acquisition of a licensed, operating EU wholesale entity.

