

OLYDEFENCE Product Evidence Checklist

Public checklist for product identity, configuration, quality, traceability and lifecycle review

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Classification	Public / business review	Document owner	OLYDEFENCE — OLYVENRA LTD
Language	English	Status	Publication-ready / controlled master
Purpose			
Use this checklist to assemble evidence references. The exact package is risk-based and contract-specific; do not transmit controlled material until a secure route is approved.			

Organisation and authority

Legal registration and registered address; ownership and authorised signatory; licence/registration or public mandate; manufacturer appointment and authorised territory; buyer/end-user authority where applicable.

Product identity and configuration

Legal manufacturer and production site; generic designation; exact model/configuration, part/catalogue number and revision; country of origin; serial/lot schema; software/firmware or accessory baseline where relevant.

Technical and conformity evidence

Current specification; exact standard and edition; test report with issuer/accreditation and tested configuration; certificate/approval with scope and status; labels, markings, instructions and limitations; material/environmental evidence where applicable.

Quality and acceptance

Quality-system scope; configuration/change control; authorised substitution; counterfeit avoidance; inspection and acceptance plan; certificate/release record; nonconformance, deviation/concession and corrective-action process.

Traceability and controlled transfer

Party and beneficial-owner review; destination/end-user/end-use documentation; classification and licence responsibility; sanctions/embargo screen; re-transfer controls; serial/lot custody and post-delivery records.

Packaging, logistics and storage

Unit of issue; packaging, marking and preservation; dangerous-goods description and packing instruction; transport mode and permits; storage conditions and access control; shelf/service life, inspection and stock rotation.

Lifecycle and support

Warranty; spares/consumables; training audience and competency; maintenance authority, tooling and calibration; technical publications; incident/complaint/recall process; obsolescence; return, destruction or disposal.

Category-specific additions

- Ammunition: lot, production/acceptance date, remaining life, surveillance, dangerous-goods and disposal records.
- Ballistic/respiratory protection: threat/hazard, exact complete configuration, conditioning, fit, limitations, care and service life.
- Tactical clothing/footwear: user/climate/task, fabric/construction, size curve, protective performance, laundering/care and replenishment.
- First-aid/trauma kits: approved contents, trained-user protocol, legal manufacturer/device status, lot/sterility/expiry, storage and replenishment; medicines excluded from public scope.

Do not send classified, export-controlled, operationally sensitive or personal medical information through the public form. Product-specific technical data, pricing, stock, licences, end-user records and contracts are shared only after due diligence through an approved channel.

Portfolio governance: configuration, origin, availability, evidence, licensing, end user, end use and support scope are validated for each authorised requirement before supplier engagement or technical release.