

Controlled Buyer Requirement Form

Institutional defence, security, protective-equipment and field-support requirements

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Classification	Controlled business form	Document owner	OLYDEFENCE — OLYVENRA LTD
Language	English	Status	Publication-ready / controlled master

Purpose

Use this controlled form to define a lawful, non-sensitive institutional requirement. OLYDEFENCE reviews authority, parties, final destination, end user, end use, classification, licensing, sanctions, anti-diversion controls, product evidence, logistics and lifecycle support before any commercial progression.

Completion guidance

Complete every applicable field. Use generic, non-sensitive descriptions and cite evidence by document title, issuer, number, revision and date. Mark Not applicable with a reason. OLYDEFENCE may request clarification, enhanced due diligence or a secure document-room exchange.

Privacy and secure document handling

Do not send classified information, export-controlled technical data, weapon configuration details, personal identity documents, medical records or credentials by ordinary email or this public form. Provide only references until an approved secure channel and authorised recipients are confirmed.

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Controlled-supply and secure-information boundary

This form is not an offer, product approval, licence, end-use authorisation, legal advice or commitment to transact. Controlled items proceed only through competent authorities and lawful licensed routes. Public first-aid kit scope excludes medicines; any medicine request must use a separate licensed channel where lawful.

Organisation identity and authority

Information requested	Response / evidence reference
Full legal name and trading name	
Registration number, country and registered address	
Corporate website and verifiable public profile	
Authorised contact: name, title, department, email and telephone	
Public mandate, procurement authority or basis to represent the organisation	
Organisation type and role in the proposed transaction	

Requirement and procurement route

Information requested	Response / evidence reference
Tender, framework, project or opportunity reference	
Procurement route and contracting authority	
Non-sensitive requirement summary and operational environment	
Product family or service category	
Indicative quantity, units of issue and forecast	

Information requested	Response / evidence reference
Required delivery, implementation or readiness date	
Indicative budget/currency, if disclosable	

Parties, destination, end user and end use

Information requested	Response / evidence reference
Buyer and contracting entity	
Consignee, importer and delivery location	
Final end-user organisation and unit, where lawfully disclosable	
Final destination country and declared location	
Documented intended end use and authorised user population	
Custody, accountability, secure storage and issue controls	
Re-transfer, re-export, resale or onward-disclosure restrictions	
Agents, brokers, advisers, freight parties and other intermediaries	

Classification, licensing and responsible transfer

Information requested	Response / evidence reference
Proposed military, dual-use, firearms, PPE, medical-device, dangerous-goods or other classification	
Applicable control-list, customs/tariff or dangerous-goods references	
Export, import, transit, brokering or other licence owner and status	
Applicable sanctions, embargo and restricted-party review status	
Human-rights and international humanitarian law risk considerations	
Diversion indicators, enhanced controls and escalation route	

Product identity, configuration and evidence

Information requested	Response / evidence reference
Legal manufacturer, production site and authorised supply route	
Generic designation, model/configuration, part or catalogue number	
Mandatory interfaces, environmental conditions and performance criteria	
Current specification, test report, certificate, label and instruction references	
Serial, lot or batch identification and traceability model	

Information requested	Response / evidence reference
Approved-equivalent and substitution/change-control rules	
Inspection, test, sampling and acceptance plan	
Customer-controlled compatibility or integration requirement	

Ammunition-specific through-life controls, if applicable

Information requested	Response / evidence reference
Generic ammunition family, authorised platform/interface and training/service status	
Manufacturer, designation, lot/batch, production date and proof/acceptance records	
Shelf life, surveillance/inspection status and remaining-life requirement	
Dangerous-goods description, packaging instruction and transport/storage conditions	
Accounting, stock rotation, incident, recovery and disposal responsibilities	

Protective equipment and medical-kit controls, if applicable

Information requested	Response / evidence reference
Defined threat/protection level and exact standard/test edition	
Model/configuration-specific laboratory, conformity and conditioning evidence	
User population, sizing, ergonomics, coverage, mass and care requirements	
First-aid/trauma kit type, approved contents list and trained-user protocol	
Medical-item manufacturer, device status, lot, sterility, expiry and replenishment	
Confirmation that medicines are excluded or routed separately through licensed channels	

Logistics, storage, security and lifecycle

Information requested	Response / evidence reference
Incoterm, delivery point, schedule and receiving requirements	
Packaging, marking, labelling, preservation and unit-of-issue requirements	
Transport mode, dangerous-goods responsibility, permits and security plan	
Storage environment, access control, inventory and condition monitoring	
Authorised training audience, scope and competency evidence	
Maintenance authority, inspection, calibration, tooling and return-to-service records	

Information requested	Response / evidence reference
Spares, consumables, warranty, service levels and obsolescence plan	
Return, demilitarisation, destruction, recycling or disposal responsibilities	

Declaration and authorised sign-off

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I confirm that the information supplied is complete and accurate to the best of my knowledge, that I am authorised to submit it, and that the organisation will promptly disclose material changes. I understand that submission does not create approval, appointment, confidentiality, availability, exclusivity, licence eligibility or an obligation to transact.

Information requested	Response / evidence reference
Authorised declarant: full name	
Position and organisation	
Basis of authority to submit this form	
Signature	
Date	