

Defence Supplier Registration Form

Risk-based qualification of manufacturers, authorised suppliers, integrators and service organisations

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

Purpose

Use this form for initial qualification or periodic renewal of a proposed OLYDEFENCE supplier. Registration does not constitute approval, representation, exclusivity or a purchase commitment. Review depth is proportionate to product, jurisdiction, destination, role and information sensitivity.

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.

[Read the OLYVENRA Privacy Notice](#)

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.

Legal identity, ownership and authorised role

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	
Ultimate beneficial owners, ownership percentages and government ownership/control	
Directors, officers and authorised signatories	
Registered, manufacturing, storage and service sites	

Product scope, provenance and controlled-goods authority

Information requested	Response / evidence reference
Manufacturer, authorised supplier, integrator, maintainer or service role	
Current licences, registrations, OEM appointments and approved territories	
Controlled and non-controlled product/service families offered	

Information requested	Response / evidence reference
NCAGE/CAGE, NSN or codification capability, where contractually applicable	
Legal manufacturer, production site and authorised supply route	
Critical subcontractors, service providers and flow-down controls	

Quality, evidence and configuration control

Information requested	Response / evidence reference
Quality-management system scope, certificates and issuing bodies	
AQAP or customer-specific quality requirements, where contractually applicable	
Test laboratories, accreditation scope and current report controls	
Configuration baseline, drawing/specification control and change notification	
Provenance, counterfeit avoidance, authorised-channel and suspect-item controls	
Incoming, in-process, final inspection and release authority	
Nonconformance, concession/deviation and corrective-action process	

Traceability, nonconformance and lifecycle support

Information requested	Response / evidence reference
Serial, lot or batch identification and traceability model	
Serial/lot records, retention period and customer traceability access	
Complaint, incident, field action, recall and customer-notification process	
Shelf/service-life, storage, inspection and stock-rotation controls	
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	
Spares, repair, calibration, technical publications and obsolescence capability	
Product, professional and other relevant insurance evidence	

Integrity, sanctions, human rights and information security

Information requested	Response / evidence reference
Applicable sanctions, embargo and restricted-party review status	
Anti-bribery, gifts, facilitation-payment and third-party controls	
Agents, brokers, commissions, success fees and payment recipients	
Sanctions, debarment, export-control breaches, investigations or enforcement history	
Actual or potential conflicts of interest and government relationships	
Human-rights/IHL due-diligence policy and escalation process	
Forced labour, child labour, modern slavery and responsible-minerals controls	
Information-security framework, incidents and controlled-technical-data protections	
Financial standing, insolvency and going-concern matters	

Supporting evidence checklist

Information requested	Response / evidence reference
Registration extract, tax registration and registered-address evidence	
Ownership chart and beneficial-owner identity evidence	
Current licences, registrations, authorisations and appointment letters	
Certificates, audit summary, accreditation and product test evidence	
Sample serial/lot genealogy, certificate and release record	
Anti-bribery, sanctions, human-rights/IHL, supply-chain and information-security policies	
Two relevant institutional customer or professional references	

Declaration and authorised sign-off

Declaration and authorised sign-off

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	

Declaration and authorised sign-off

Date