ADULT WELLNESS / SEXUAL HEALTH MANUFACTURING

A practical capability statement for overseas buyers.

WODE Biotechnology Co., Ltd. supports brands, distributors and qualified sourcing teams with OEM/ODM development, private-label packaging, selected finished-product wholesale, bulk supply and reference-sample development. This pack summarizes what WODE can discuss, what must be confirmed per project, and where buyers can review the supporting sourcing resources.

LEGAL COMPANY	FOUNDED	BUSINESS CONTACT
WODE Biotechnology Co., Ltd.	12 October 2006	info@wodebiotech.com

PRODUCT CAPABILITY MAP

Four current commercial product lines.

The following portfolio is the approved public scope. Exact formula, claims, packaging, MOQ, documentation and destination-market feasibility are confirmed against each project.

01 Personal lubricants and selected concentrates

Water-based and gel formats, selected hybrid or silicone directions, functional variants, powder lubricant concentrate and external-use glow-effect concepts. Routes may include custom formulation, private label, bulk finished formula, selected concentrate and finished goods.

wodebiotech.com/en/private-label-water-based-lubricant-manufacturer/

02 Pre-intimacy cleansing and rinse systems

Intimate wash, ready-to-use rinse formats and portable powder-to-rinse development platforms. Applicator, water source, dissolution, compatibility and destination-market wording require project-specific confirmation.

wodebiotech.com/en/intimate-wash-care-oem/

03 Men's external-use intimate care

External-use essences, delay-care sprays, soothing gels and single-, three- or ten-wipe private-label formats. Product names, ingredients, directions and claims are reviewed for the destination market.

wodebiotech.com/en/delay-spray-wipes-oem/

04 NEWSCEN rapid-test distributor projects

Qualified B2B review for NEWSCEN HIV (1+2) and Treponema Pallidum antibody rapid-test enquiries. WODE is the manufacturer and NEWSCEN is a WODE-manufactured diagnostic product line. Distribution is subject to current document and destination-market review.

wodebiotech.com/en/newscen-distributor-document-review/

AVAILABLE SUPPLY ROUTES

COMMERCIAL ROUTE	WHAT IT MEANS	KEY CONFIRMATION
OEM / ODM and private label	Formula direction, sampling, packaging and finished production under the buyer's brand.	Formula, pack, quantity, market, claims and documents.
Bulk or semi-finished supply	Bulk finished formula or selected concentrate for qualified buyer-managed filling.	Buyer filling controls, container compatibility and shipment route.
Finished-product wholesale	Selected brand-neutral or ready product formats for approved commercial projects.	Available format, pack, label responsibility and destination market.
Reference sample development	Lawful reference-sample review, measurable target definition, prototype work and pilot preparation.	Reference rights, target attributes, acceptance criteria and scale-up scope.

PROJECT CONTROL

From buyer brief to a reviewable quotation.

A useful first enquiry does not need a complete technical specification. It should establish the commercial and product boundaries that control feasibility, sampling and quotation.

01	**Brief**	Target market, product family, supply route, intended use and expected quantity.
02	**Formula direction**	Existing family, measurable reference attributes or lawful sample for review.
03	**Sample**	Prototype direction, feedback criteria, target packaging and evaluation plan.
04	**Production review**	Quantity, component lead time, records, testing and commercial terms.
05	**Delivery scope**	Final product, bulk supply or pilot batch, destination and buyer responsibilities.

WHAT TO INCLUDE IN AN RFQ

PRODUCT Family, texture, intended application and preferred direction.	**SUPPLY MODEL** Private label, finished wholesale, bulk finished formula, selected concentrate or pilot batch.	**PACKAGING** Bottle, tube, sachet, pouch or buyer-supplied component.
QUANTITY Forecast, launch quantity and repeat-order expectation.	**MARKET** Destination country, language and known compliance expectations.	**TIMING** Target sample date, launch window and required delivery route.

DOCUMENT SCOPE

Documents are reviewed against the selected product, assessed sample, batch, package and shipment. A report for one lubricant sample must not be presented as universal approval for every formulation or destination. WODE's public buyer guide describes a 2026 GHS Rev.10 MSDS and sea-transport classification report for one assessed lubricant sample; applicable records for a new project are confirmed during review.

BUYER RESOURCES

Use one evidence-based route for the next discussion.

The WODE buyer resource centre is available in 14 languages. It covers project selection, low-MOQ planning, packaging, quotation inputs, bulk versus finished supply, documentation scope and the route from a reference sample to pilot-batch review.

Buyer resource centre	wodebiotech.com/en/oem-buyer-resources/
Product capability portfolio	wodebiotech.com/en/product-portfolio/
OEM / ODM manufacturing	wodebiotech.com/en/oem-odm-manufacturing/
RFQ checklist	wodebiotech.com/en/lubricant-oem-rfq-checklist/
MSDS and transport document guide	wodebiotech.com/en/personal-lubricant-msds-transport-documentation/
Packaging and MOQ guide	wodebiotech.com/en/private-label-packaging-moq-guide/
Reference sample to pilot batch	wodebiotech.com/en/reference-sample-to-pilot-batch/
Request for quotation	wodebiotech.com/en/contact-rfq/

START A QUALIFIED PROJECT DISCUSSION

Tell us the product family, market, supply route and expected quantity.

info@wodebiotech.com

https://wodebiotech.com/en/contact-rfq/

COMPANY FACTS

Legal company: WODE Biotechnology Co., Ltd.
Established: 12 October 2006
Address: Unit D, Building A, Nankai Industrial Park, Ya'an Road, Nankai District, Tianjin, China
Manufacturer relationship: WODE is the manufacturer. NEWSCEN is a WODE-manufactured diagnostic product line.

This capability statement is a sourcing aid, not a universal quotation, certification or market authorization. Final feasibility, specification, claims, MOQ, lead time, documentation and destination-market requirements are confirmed in writing for the selected project.