Performance, Quality & Safety

World Health Organization

WHO IMD-PQS Annual Review 2026

PRODUCT RESELLER - FORM A

(COMPANY REVIEW FORM)

*** One Form A only per applicant/PQ Holder ***

SECTION 1a. CHAIN OF CUSTODY — MANDATORY

Original product manufacturer company name:
Reseller company name (current applicant / PQ Holder):

SECTION 1b. RESELLER DETAILS — MANDATORY

Registered Company Name:		
License Issuing Country:		
Manufacturing site postal address: Road name & number: Complement: Postal code:	City:	Country:
Manufacturing site web address:		
Manufacturing site contact number Country code:	Area code:	Telephone:

Contact person 1 name:		
Contact person email address:		@
Contact person name contact number: Country code:	Area code:	Telephone:

Contact person 2 name:		
Contact person email address:		@
Contact person name contact number Country code:	Area code:	Telephone:

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SECTION 1c. PRODUCT MANUFACTURER DETAILS MANDATORY

!! USE THIS SECTION TO PROVIDE INFORMATION ABOUT THE ORIGINAL PRODUCT MANUFACTURER !!

Registered Company Name (MANUFACTURER):
License Issuing Country:
Manufacturing site postal address: Road name & number: Complement: Postal code: City: Country:
Manufacturing site web address:
Manufacturing site contact number Country code: Area code: Telephone:

Contact person 1 name:
Contact person email address: @
Contact person name contact number: Country code: Area code: Telephone:

Contact person 2 name:
Contact person email address: @
Contact person name contact number Country code: Area code: Telephone:

SECTION 2a. LICENSING - RESELLER LICENCES MANDATORY

IMPORTANT: Only a valid **Business Registration** document is acceptable as licence.

Translations - Any original document in languages other than English must be accompanied by an official notarized English translation.

INSTRUCTIONS: Complete the information in Table 2 below.

One (1) certified copy (.pdf) of each licence must also be submitted.

Resellers that do not provide valid licences will not be considered.

The submission of invalid licences will result in the automatic suspension of all of the manufacturer's products until valid licenses are provided.

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Table 2a. – Mandatory **RESELLER** Licence

		Formal document no.	Issue date (DD-MMM-YY)	Expiry date (DD-MMM-YY)
MANDATORY:	**Company license** or **Registration document**			
Optional:	Manufacturing license			
Optional:	Other:……..			

SECTION 2b. ORIGINAL MANUFACTURER LICENCES — MANDATORY

!! USE THIS SECTION TO PROVIDE INFORMATION ABOUT THE ORIGINAL PRODUCT MANUFACTURER !!

IMPORTANT: Only a valid **Business Registration** document is acceptable as licence.

Translations - Any original document in languages other than English must be accompanied by an official notarized English translation.

INSTRUCTIONS: Complete the information in Table 2 below.

One (1) certified copy (.pdf) of each licence must also be submitted.

Manufacturers that do not provide valid licences will not be considered.

The submission of invalid licences will result in the automatic suspension of all of the manufacturer's products until valid licenses are provided.

Table 2b. – Mandatory **MANUFACTURER** Licence

		Formal document no.	Issue date (DD-MMM-YY)	Expiry date (DD-MMM-YY)
MANDATORY:	**Company license** or **Registration document**			
Optional:	Manufacturing license			
Optional:	Other:……..			

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SECTION 3a. CERTIFICATION - RESELLER CERTIFICATES MANDATORY

IMPORTANT:

Certificates expiring before July 14th 2026 must be accompanied by a copy of a renewal request letter**.** This is a **mandatory** requirement.

A website link to a certification authentication page is mandatory for certification verification. **PLEASE VERFIY THAT WEBLINK FUNCTIONS.**

INSTRUCTIONS:

Complete the information in Table 2 below.

One (1) certified copy (.pdf) of each certificate is required**.**

Translations - Any original document in languages other than English must be accompanied by an official notarized English translation.

Resellers that do not provide valid certificates will not be considered. Email confirmation of a pending certificate validation will not be accepted.

The submission of invalid certificates will result in the automatic suspension of all manufacturer products until valid certificates are provided.

All certification bodies should be ILAC accredited.

Table 3a. – Mandatory Certification

Requirements per IMD-PQS category	Certificate	Certificate accreditation year (e.g. 2015)	Certification authority (certified by)	Website link for certification verification *MANDATORY*	Expiry date (DD-MMM-YY)
ALL CATEGORIES Except E008 & E013	**ISO 9001:2015***				
E001 E003 E004	**ISO 14001:2015***				
E008 E013	**ISO 13485:2016***				
Optional	**93/42/EEC**				
Other approval marks (e.g. CE, UL, FDA or EMA):					

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SECTION 3b. ORIGINAL MANUFACTURER CERTIFICATES MANDATORY

!! USE THIS SECTION TO PROVIDE INFORMATION ABOUT THE ORIGINAL PRODUCT MANUFACTURER !!

IMPORTANT:

Certificates expiring before July 14th 2026 must be accompanied by a copy of a renewal request letter**.** This is a **mandatory** requirement.

A website link to a certification authentication page is mandatory for certification verification. **PLEASE VERFIY THAT WEBLINK FUNCTIONS.**

INSTRUCTIONS:

Complete the information in Table 2 below.

One (1) certified copy (.pdf) of each certificate is required**.**

Translations - Any original document in languages other than English must be accompanied by an official notarized English translation.

Manufacturers for whom valid certificates are not provided will not be considered. Email confirmation of a pending certificate validation will not be accepted.

The submission of invalid certificates will result in the automatic suspension of all manufacturer products until valid certificates are provided.

All certification bodies should be ILAC accredited.

Table 3b. – Mandatory **MANUFACTURER** Certification

Requirements per IMD-PQS category	Certificate	Certificate accreditation year (e.g. 2015)	Certification authority (certified by)	Website link for certification verification MANDATORY	Expiry date (DD-MMM-YY)
ALL CATEGORIES Except E008 & E013	**ISO 9001:2015***				
E001 E003 E004	**ISO 14001:2015***				
E008 E013	**ISO 13485:2016***				
Optional	**93/42/EEC**				
Other approval marks (e.g. CE, UL, FDA or EMA):					

SECTION 4. CHANGES TO MANUFACTURING INFORMATION MANDATORY

INSTRUCTIONS: If no manufacturing changes have occurred, or are expected to occur, between April 2025 and March 2026, please tick **NO**.

If changes **HAVE** occurred, resellers must submit documentation verifying the amended information.

Table 4 – Changes to manufacturing information

		Please provide details of changes:
Are there any changes to the manufacturer's **company name** and/or **status**?	YES ☐ NO ☐	
Are there any changes to the **manufacturing site**?	YES ☐ NO ☐	