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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 (OEM) company name:

Reseller company name (PQ holder):

SECTION 1b. RESELLER DETAILS

MANDATORY

!! USE THIS SECTION TO PROVIDE INFORMATION ABOUT THE RESELLER COMPANY / PQ HOLDER !!

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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CONTINUES ON NEXT PAGE →



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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. 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 VERIFY 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*				



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Optional	93/42/EEC				
Other approval marks (e.g. CE, UL, FDA or EMA):					

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 VERIFY 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*				



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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

		<i>Please provide details of changes:</i>
Have there been any changes to the company name and/or status in the last 12 months?	YES <input type="checkbox"/> NO <input type="checkbox"/>	
Have there been any changes to the product manufacturing site address in the last 12 months?	YES <input type="checkbox"/> NO <input type="checkbox"/>	