



WHO IMD-PQS Annual Review 2026 PRODUCT MANUFACTURER – DECLARATION

*** Submit ONE declaration only per applicant/PQ Holder ***

complete and that the information provided in Forms A and B is accurate, correct and complete and that the documents submitted along with those forms are genuine. I undertake to inform the IMD-PQS Secretariat in writing of any changes to the information already provided and to update the information on these forms if requested to do so by the IMD-PQS Secretariat.

I also hereby confirm that the each of the following mandatory and supplemental documents are included in the submission, as required:

	PRODUCT MANUFACTURER – CHECKLIST	/
MANDATORY:	Form A – one form completed (in word.doc format)	
MANDATORY:	Form B – one form completed for each product (in word.doc format)	
MANDATORY:	Company licence	
If required	Notarised translations of licences that are not in English or French	
MANDATORY:	All relevant ISO certifications (See Form B)	
If required	Notarised translations of certificates that are not in English or French	
MANDATORY:	Copy of a the "Product Data Sheet" for each product	
If required	A hand-annotated <u>"Product Data Sheet"</u> indicating any changes required to administrative or technical product information.	

CONTINUES BELOW →

¹ https://apps.who.int/immunization standards/vaccine quality/pqs catalogue/

² https://apps.who.int/immunization_standards/vaccine_quality/pgs_catalogue/





CONFIRMATION OF PRODUCT WITHDRAWALS IN 2026			
Do you wish to withdraw any of your currently-prequalified products as a part of this 2026 Annual Review?	Yes No		
If you answered "Yes", please list the IMD-PQS code of each product you wish to withdraw as a part of this 2026 Annual Review:			
E00 /			
E00 /			
E00 /			
(Add lines as required)			
SIGNATURE REQUIRED:			
Authorised signature:			
For and on behalf of (Company name):			
Date:			
IMPORTANT: SUBMIT ONLY <u>ONE</u> DECLARATION FORM PER PREQUALIFICATION HOLDER.			