

WHO Prequalification Programme
WHO PUBLIC ASSESSMENT REPORT (WHOPAR)

[RH067 trade name]*

Desogestrel/ethinylestradiol 150 micrograms/30 micrograms tablets

[RH067 trade name], manufactured at Laboratorios Leon Farma SA, Villaquilambre, Leon 2400824193, Spain, was included in the WHO list of prequalified medicinal products for contraception for women on 27 October 2020.

[RH067 trade name] is indicated as an oral combined hormonal contraceptive (CHC) medicine for women. Detailed information on the use of this product is described in the summary of product characteristics (SmPC), which can be found in this WHOPAR.

The active pharmaceutical ingredients of [RH067 trade name] are ethinylestradiol and desogestrel, two types of female sex hormones, oestrogen and progestogen.

The efficacy and safety of desogestrel and ethinylestradiol are well established based on extensive clinical experience in female contraception

For details on the uses of this product and for side effects and warnings, see Part 4 of this WHOPAR (summary of product characteristics).

On the basis of data submitted and public information on the use of desogestrel and ethinylestradiol for contraception, the team of assessors advised that [RH067 trade name] is of acceptable quality, efficacy and safety to allow inclusion of [RH067 trade name] in the list of prequalified medicinal products.

Summary of prequalification status for [RH067 trade name]:

The table shows the status of relevant completed activities, including the dates of WHO internal quality assurance.

* Trade names are not prequalified by WHO. This is the national medicines regulatory authority's responsibility.

Initial acceptance	Date	Outcome
Status on PQ list	27 October 2020	listed
Pharmaceutical quality	21 October 2020	MR
Bioequivalence	22 October 2020	MR
Safety, efficacy	NA	NA
GMP (re-)inspection		
API	NA	NA
FPP	08 March 2018	MR*
GCP/GLP (re-)inspection	08 August 2019	MR*
API: active pharmaceutical ingredient FPP: finished pharmaceutical product GCP: good clinical practice [quality standard] GLP: good laboratory practice [quality standard]	GMP: good manufacturing practice [quality standard] MR: meets requirements MR*: desk review (based on recent inspection reports) NA: not applicable, not available PQ: prequalification	