



## Confirmation of WHO Active Pharmaceutical Ingredient Prequalification (CPQ)

<b>Date:</b>	12 July 2017
<b>WHO prequalification number:</b>	WHOAPI-234
<b>Active pharmaceutical ingredient (API):</b>	Sulfadoxine
<b>API specification number:</b>	FPS/130-03, Version 03
<b>Re-test Period:</b>	36 months
<b>Storage conditions</b>	Do not store above 30°C, protect from moisture, protect from light

### API Manufacturers:

Anuh Pharma Ltd  
Manufacturing Block AB-3  
E-17/3&E17/4 M.I.D.C, Boisar  
Tarapur, Taluka –Palghar, Dist: Thane-401 506  
Maharashtra state  
India

### API Intermediate manufacturers: *(in addition to the API manufacturers above)*

Not applicable.

This is to confirm that Sulfadoxine, manufactured by Anuh Pharma Ltd, has been prequalified by the World Health Organization (WHO). Further information on the API prequalification procedure can be located on the Prequalification Team - Medicines Assessment web page:

[http://www.who.int/prequal/info\\_applicants/API\\_info\\_applicants.htm](http://www.who.int/prequal/info_applicants/API_info_applicants.htm).

API prequalification provides an assurance that the supplied API is of good quality. The comprehensive evaluation procedure has two components: assessment of the API master file (APIMF) to verify compliance with WHO norms and standards, and assessment of the sites of API manufacture to verify compliance with WHO GMP requirements.

The decision to prequalify Sulfadoxine, manufactured by Anuh Pharma Ltd, is particular to the specific details assessed during evaluation, such as sites of manufacture, method of manufacture, control of the API and retest period.

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