



25 November 2022

**Shortage of LONITEN minoxidil 10mg tablets and alternative supply arrangement under Section 19A of the Therapeutic Goods Act**

Dear Healthcare Professional,

This communication is sent by ORSPEC Pharma to notify your organisation that due to the shortage of **LONITEN minoxidil 10mg tablets (AUST R 12309)**. ORSEPC Pharma has arranged the supply of an alternative product on a temporary basis.

**Minoxidil 10mg tablets (Roma Pharmaceuticals)**, are NOT registered in Australia and supply is granted under an exemption granted by the Therapeutic Goods Administration (TGA) under Section 19A of the Therapeutic Goods Act, 1989 until **30 August 2023**.

**Minoxidil 10mg tablets (Roma Pharmaceuticals)**, are approved for use under Section 19A for the following indications:

*As adjunctive therapy in adults with severe refractory hypertension which has failed to respond to extensive multiple therapy.*

*When used in combination with an accompanying diuretic and beta-blocker, minoxidil MINOXIDIL has been shown to reverse encephalopathy and retinopathy in severe hypertensives.*

The s19A approved Uk product is identical in active ingredient and strength to the Australian registered product. The two products differ in their pack sizes. The differences are noted below:.

	<b>ARTG product</b> LONITEN minoxidil 10mg tablets (AUST R 12309).	<b>S19A product</b> Minoxidil 10mg tablets (Roma Pharmaceuticals)
<b>Nature and contents of container</b>	The tablets are supplied in HDPE <b>bottles of 100.</b>	Minoxidil Tablets are packed in opaque PVC/PVDC/ALU <b>blisters.</b> Each blister contains 10 tablets. <b>Pack size: 60 tablets.</b>

**Minoxidil 10mg tablets (Roma Pharmaceuticals)**, are registered in the United Kingdom and are packaged in English language. For dosing and administration information, please refer to the Australian Product Information for LONITEN® (MINOXIDIL) tablets available at <https://www.ebs.tga.gov.au/>

### **Adverse Event Reporting**

Reporting any suspected adverse event is important for the continued monitoring of the safety of all medicines. Any adverse events which are experienced with **Minoxidil 10mg tablets (Roma Pharmaceuticals)**, should be reported by healthcare professionals and patients to ORSPEC Pharma on 024 339 4239 or email at [sas@orspecpharma.com](mailto:sas@orspecpharma.com). Alternatively, this information can be reported to the TGA at <https://www.tga.gov.au/reporting-problems>

Please forward this information to relevant staff members in your organisation.

For further information, please contact ORSPEC Pharma on 024 339 4239 or email [sas@orspecpharma.com](mailto:sas@orspecpharma.com).

Yours sincerely,

A handwritten signature in black ink, appearing to read "Deon Scheepers".

Deon Scheepers  
Managing Director  
ORSPEC Pharma