Date: 12 Nov. 2026

CRITICAL RECALL

Chuan Xin Lian Jie Du Fang a.k.a. Andrographis Antitox Formula (CM105)

All batches within expiry

ARTG: AUST L 99762 Chuan Xin Lian Jie Du Fang a.k.a. Andrographis Antitox Formula (CM105)

TGA Ref: RC-2026-RN-00931-1

Dear Sun Herbal Distributor & Customer,

Sun Herbal, following agreement with the Therapeutic Goods Administration (TGA), is conducting a recall of all batches that are still within expiry for the above listed medicine Chuan Xin Lian Jie Du Fang a.k.a. Andrographis Antitox Formula (CM105). The product has been supplied to you/your organisation.

WHAT IS THE PROBLEM?	• Since 2019, listed medicines containing *Andrographis paniculata* (Andrographis) for supply in Australia have been required to carry a label warning about the risk of allergic reactions including severe reactions such as anaphylaxis. • However, despite these warnings, the TGA has continued to receive a sustained high number of reports of anaphylaxis associated with Andrographis-containing listed medicines, which are considered low-risk medicines. • Andrographis has now been removed from the *Therapeutic Goods (Permissible Ingredients) Determination (No. 2) 2026*. This means the ingredient is no longer available for use in new listed (low risk) medicines for supply in Australia. • Further information on this decision is available at Andrographis removed from listed medicines framework because of allergy risk | Therapeutic Goods Administration (TGA)
WHAT IS THE HAZARD/RISK?	• Anaphylaxis is the most severe type of allergic reaction and is a medical emergency. • It requires immediate treatment with adrenaline (epinephrine). Delayed treatment can be life-threatening. • Andrographis-related anaphylaxis reactions are unpredictable and can happen the first time someone takes it, or after taking it previously without any issue.
WHAT SHOULD YOU DO?	**Distributors/retailers** • Inspect your stock **immediately** and quarantine all stock of Chuan Xin Lian Jie Du Fang a.k.a. Andrographis Antitox Formula (CM105) on hand to prevent further use or distribution.

	• Complete the attached Customer Response form immediately even if you do not have any affected stock and return it to info@sunherbal.com.au so we can reconcile this recall. • Return affected stock on hand with the completed response form to the following address: Unit 5, 25 Garema Circuit Kingsgrove NSW 2208 • Following receipt of returned stock and a completed response form we will contact you to arrange a refund/credit for the returned stock. • Ensure relevant staff members are informed of this recall. • If you have supplied or transferred any of the above product to another facility or organisation, provide that facility with a copy of this letter immediately. • Place this letter in a prominent position for at least one month. • Retailers are recommended to place the 'Public Notice' attached to this letter in their storefront to alert consumers about the recall. **Consumers** • Consumers are advised that if they have the above product, to not use it. • Consumers can return the product to the place of purchase. • If someone is suspected of having anaphylaxis, call for emergency medical help immediately by dialling '000'.
CONTACT INFORMATION	For questions regarding this letter, please contact: Keith Chan (02) 9759 2927 / info@sunherbal.com.au

Thank you for your assistance in helping us to manage this recall.



Keith Chan

Managing Director