

Vaping goods

Pharmacy self-assessment tool

Compliance with storage and supply requirements

Product compliance and storage

- ☐ Are all vaping goods sold listed on the [TGA's list of notified vapes](#)?
- ☐ Are all vaping goods stored behind the counter and not available for self-selection by consumers?
- ☐ There are no advertisements for vaping goods displayed inside or outside the pharmacy.

Supply pathways and regulatory requirements

- ☐ Are vaping goods with a nicotine concentration greater than 20mg/mL only supplied on prescription?
- ☐ Are all vaping goods supplied via an established pathway i.e. TGA's [AP scheme](#) or [SAS](#)?
- ☐ Are processes in place to ensure Schedule 3 supply requirements are met, including:
 - ☐ confirming the vaping goods is for smoking cessation or management of nicotine dependence
 - ☐ confirming the patient is aged 18 years or older
 - ☐ sighting patient identity and age
 - ☐ obtaining informed consent
 - ☐ submitting an SAS Category C notification within 28 days of supply
- ☐ Are there processes in place to ensure Schedule 3 supply is restricted to:
 - ☐ no more than one month's supply at a time
 - ☐ no more than one supply per patient per month
- ☐ Are there processes in place to ensure that patients are informed about:
 - ☐ alternative smoking cessation supports and therapy
 - ☐ appropriate dose and frequency of use as per patient age, weight and severity of condition
 - ☐ titration plan and expected duration of therapy
 - ☐ potential drug interactions with current medicines
 - ☐ contact details of smoking cessation support services
 - ☐ the unapproved status of vaping goods (i.e., not TGA assessed for safety, quality or efficacy).

Safety and reporting

- ☐ Are there processes in place to ensure adverse effects and faulty vaping goods are reported to the TGA and sponsor of the vaping good.

For more information, visit the [Vaping Goods](#) webpage or submit an [online enquiry](#).