. The Pharmacy Board of Australia is inviting feedback on its draft revised Guidelines for compounding of medicines (the draft revised guidelines). Optional questions have been provided below and you may wish to address some or all of these in your response.

Please note this survey contains the same questions as the response template (Word document). Please choose only ONE method of responding to avoid duplicating your submission.

. Please provide your details below

Name	
Organisation name (if applicable)	The Tasmanian Pharmacy Authority
Contact email	

. Published submissions will include the names (if provided) of the individuals and/or the organisations that made the submission unless confidentiality is requested.

Do you want your responses to be published after public consultation?

- Yes
- O No
- . When providing feedback, please include the relevant guideline number/section that your feedback refers to.
- . The revised compounding guidelines include additional content on medicine supply pathways to consider before deciding if it is appropriate to compound a medicine (Guideline 1 When to compound medicines)
- Q1. 1. Is the new content on medicine supply pathways clear and helpful? Why or why not?

The content is concise, providing a clear decision-making pathway for pharmacists to consider prior to compounding a medicine, including non-prescription medicines. It will provide regulatory bodies with a clear framework for auditing/inspection or investigation into a pharmacists' practice is required

. When providing feedback, please include the relevant guideline number/section that your feedback refers to.

active ingredients and their strengths, and all inactive ingredients) must be provided to the patient when requested (Guideline 13 Supporting informed patient choice). Providing patients with information about the ingredients in their compounded medicine will support patient choice and safer patient outcomes.
Q2. 2. Do you agree that the formula for their compounded medicine must be provided when requested by the patient? Why or why not?
The formula should be provided to the patient, by knowing all the ingredients contained in the compound medicine they can then make an informed decision to take it or not.
. When providing feedback, please include the relevant guideline number/section that your feedback refers to.
. The revised compounding guidelines include content that is specific to medicines compounded for animal patients.
Q3. 3. Is the new content that is specific to medicines for animal patients clear and helpful? Why or why not?
Yes, provides differentiation for pharmacists when considering compounding veterinary medicines compared to human medicines.
. When providing feedback, please include the relevant guideline number/section that your feedback
Q4. 4. Is there any content that needs to be changed, added or deleted in the revised guidelines? If so, please provide your suggestions and reasons.
Q4. 4. Is there any content that needs to be changed, added or deleted in the revised guidelines? If so,
Q4. 4. Is there any content that needs to be changed, added or deleted in the revised guidelines? If so, please provide your suggestions and reasons.
Q4. 4. Is there any content that needs to be changed, added or deleted in the revised guidelines? If so, please provide your suggestions and reasons.  No
Q4. 4. Is there any content that needs to be changed, added or deleted in the revised guidelines? If so, please provide your suggestions and reasons.  No  Q5. 5. Is the language of the revised guidelines clear and is the structure helpful? Why or why not?  Clear and concise  Q6. 6. Please provide any other feedback about the revised guidelines.
Q4. 4. Is there any content that needs to be changed, added or deleted in the revised guidelines? If so, please provide your suggestions and reasons.  No  Q5. 5. Is the language of the revised guidelines clear and is the structure helpful? Why or why not?  Clear and concise