RESPONSE FORM

Codex Committee on Food Labelling Electronic Working Group on Allergen Labelling

Second Consultation Paper October 2022

Request for comments on Part 1 – Revision of the GSLPF

The EWG are asked to provide comments on the proposed draft revisions to the GSLPF as discussed in Part 1 of the consultation paper. This includes providing comments on the proposed definitions at Section 2 of Part 1.

Comments on Part 1:

Allergy & Anaphylaxis Australia (A&AA) agree with the amendments made to the draft revisions to the GSLPF. The amendments are consistent with the feedback that we have previously provided.

We have one comment to make with regard to the definition of food allergy below.

Food allergy is defined as an adverse health effect arising from a specific immune-mediated response that occurs reproducibly on oral exposure to a given food, which may or may not be mediated by food-specific immunoglobulin class E (IgE) antibodies.

It's almost always oral exposure, however cooking of fish can aerolise protein, which can trigger anaphylaxis. People can also have mild to moderate allergic reactions to touch.

Request for comments on Part 2 – Proposed draft PAL guidelines

Please provide responses to the following questions about the PAL guidelines.

Question 1: Do you support the Chair's proposal for the title, purpose, and scope?			
Do you support the Chair's proposal for th	le lille, purpose, and scope?		
Yes ⊠	No □		
Please provide reasons for your answer: A&AA agree that the title should be 'Guidelir Labelling' and it is not necessary to include "A and industry understand the term, Precautiona believe that to ensure PAL is valuable to cons titled, "Requirements for the use of Precaution	Advisory" in the title. As outlined, consumers ry Allergen Labelling (PAL). However, we umers, it should be mandated and therefore be		
A&AA generally agree that the purpose is "To approaches to the effective use of PAL for conabout the risk from the unintentional presence during the production of food," however we be should replace just "cross contact". This uses community. In addition, our understanding is to plate', which would encompass growing, metransporting of raw ingredients. A&AA agree with the scope, "These guideline unintended presence of allergens caused by created the production of the purpose is "To approache the purpose i	mmunicating to consumers with food allergy of allergens in food due to cross-contact relieve that cross contact/cross contamination language that is better understood by the that "production of food" means from 'paddock nanufacturing as well as the storage and respectively. The production of the storage and respectively.		
Question 2: Do you support the Chair's proposal for the definitions of 'allergen' and 'precautionary allergen labelling'?			
Yes ⊠	No □		
Please provide reasons for your answer: A&AA agree that the definition of allergen should be, "food and ingredients listed in sections 4.1.2.4 and where applicable 4.1.2.5 of the General Standard for the Labelling of Prepackaged foods.			
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Question 3: Do you support the inclusion of the new revised principle 4.1?		
Yes ⊠	No □	
Please provide reasons for your answer: A&AA support that the use PAL be restricted presence of an allergen(s) cannot be controlled to absolve food business operators from using	d. It should not be used as a blanket statement	
Question 4: Do you support the revised wording of pri	nciple 4.2?	
Yes ⊠	No □	
of a risk assessment which shall include, but it We believe that risk assessment should be evi	ion to use PAL should be based on the findings as not limited to, quantitative risk assessment. dence/science based, otherwise it is subjective be not accurately convey information required	
Question 5: Do you support the including in section 4. reference doses (RfD) based on ED05 as	3.1 of the draft guideline the established recommended by the Expert Committee?	
Yes ⊠	No □	
agreed by expert committee but suggest that to data may become available. This is likely to o	DD05). We accept that they are based on levels hey do need review at points in time when more occur with the high number of oral food olled settings. In addition, we are moving to the	

more information on people that are having anaphylaxis. This will allow us to so monitor this

space once we move to ED05.

Question 6: Do you support principle 4.4 which allows national authorities to determine reference doses for the regional list of allergens in section 4.2.1.5 of the proposed revised GSLPF?		
Yes ⊠	No □	
	nust include clinical immunology/allergy	
Question 7: Do you support including location and form presentation of PAL?	nat aspects in the principles for the	
Yes ⊠	No □	
Please provide reasons for your answer: The presentation of PAL needs to be simple, of statements should be clearly distinguishable of visualised near the list of ingredients and any simple.	n packet and need to be able to easily be	

Question 8:

Which option for a single statement for PAL do you prefer and why?

Option 1 – 'Not suitable for people with a x allergy' or 'Not suitable for x allergy'

Option 2 – 'May contain x'

Option 3 – 'May be present: x'

Option 1 X□	Option 2	Option 3	Other □		
Please provide reasons for your answer. If answering 'Other', please describe your proposed option and explain why you support this. A&AA prefer option 1 as we believe "not suitable for people with a X allergy" it is the simplest message for all and communicates our desire for people with food allergy to avoid those products which have been assessed as being 'higher risk". If we move to ED05 we need to move to a statement that people need to easily recognise. We believe that Option 1 could become the 'new' trusted PAL single statement used as we transition to ED05.					
Question 9: Do you support the inclusion of a principle on the need to indicate on the label (e.g. through the use of a symbol) that a risk assessment has been undertaken?					
		I	monan.		
Yes ⊠		No □			
Please provide reasons for your answer: A&AA believe that a symbol should be included on the label if a risk assessment has been done as otherwise there is no way to communicate to consumers that the product has undergone a risk assessment. We believe that this would instil some confidence in consumers, knowing that the product has undergone risk assessment. Currently packaged food without a PAL may be more of a risk than a product with a PAL and consumers have no way of knowing this.					
		Question 10: Do you agree with the proposed principle on education to ensure understanding about PAL by consumers, health care providers and food business operators?			
Do you agree with th					
Do you agree with th					

Question 11:

education messaging is consistent.

Do you support the Chairs' proposal to incorporate the draft PAL guideline as an annex to the GSLPF?

consumers, health care providers and food business operators. This is important if

PAL is going to be meaningful, accurate and effective. It is important that the

Yes ⊠	No □		
Please provide reasons for your answer: A&AA support PAL as an annex rather than as a stand-alone guidance as PAL needs to be consistent with GSLPF and as they both relate to allergen labelling.			