Submission form: Building Amendment Bill proposals for regulations for Building Product Information Requirements, the modular component manufacturer certification scheme, and the product certification scheme

The Ministry of Business, Innovation and Employment (MBIE) would like your feedback on proposals for regulations for Building Product Information Requirements, the modular component manufacturer certification scheme, and the product certification scheme (CodeMark). Please provide your feedback by **5pm, on 11 June 2021.**

When completing this submission form, please provide comments and reasons explaining your choices. Your feedback provides valuable information and informs decisions about the proposals.

We appreciate your time and effort taken to respond to this consultation.

Instructions

To make a submission you will need to:

- 1. Fill out your name, email address, phone number and organisation.
- 2. Fill out your responses to the discussion document questions. You can answer any or all of these questions in the <u>discussion document</u>. Where possible, please provide us with evidence to support your views. Examples can include references to independent research or facts and figures.
- **3.** If your submission has any confidential information:
 - i. Please state this in the email accompanying your submission, and set out clearly which parts you consider should be withheld and the grounds under the Official Information Act 1982 (Official Information Act) that you believe apply. MBIE will take such objections into account and will consult with submitters when responding to requests under the Official Information Act.
 - ii. Indicate this on the front of your submission (e.g. the first page header may state "In Confidence"). Any confidential information should be clearly marked within the text of your submission (preferably as Microsoft Word comments).
 - iii. Note that submissions are subject to the Official Information Act and may, therefore, be released in part or full. The Privacy Act 1993 also applies.

How to submit this form

4. Submit your feedback:

- i. As a Microsoft Word document by email to building@mbie.govt.nz with subject line: Consultation: Building Amendment Bill proposals for regulations
- ii. By mailing your submission to:

Consultation: Building Amendment Bill proposals for regulations Building System Performance Building, Resources and Markets Ministry of Business, Innovation and Employment PO Box 1473

Wellington 6140 New Zealand

Submitter information

MBIE would appreciate if you would provide some information about yourself. If you choose to provide information in the section below it will be used to help MBIE understand the impact of our proposals on different occupational groups. Any information you provide will be stored securely.

Your name, email address, phone number and organisation

Name:		Teena Hale Pennington		
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Phor	ne number:	027 527 5273		
Orga	nisation:	Te Kāhui Whaihanga New Zealand Institute of Architects		
	name or oth that MBIE m MBIE may up www.mbie.s	Act 1993 applies to submissions. Please tick the box if you do <u>not</u> wish your er personal information to be included in any information about submissions ay publish. pload submissions or a summary of submissions received to MBIE's website at govt.nz. If you do <u>not</u> want your submission or a summary of your submission to our website, please tick the box and type an explanation below:		
I do n	ot want my s	ubmission placed on MBIE's website because [insert reasoning here]		
Please check if your submission contains confidential information				
I would like my submission (or identifiable parts of my submission) to be kept confidential and have stated my reasons and ground under section 9 of the Official Information Act the believe apply, for consideration by MBIE.				

Supply chain responsibilities to meet Building Product Information Requirements

1.	Do you think the clear?	Do you think the split of responsibilities across the supply chain for information requirements is clear?					
	☐ Yes	☑ Yes, with changes	□ No	\square Not sure/No preference			
	Please explain yo	our views.					
	(along with supp		on standards) to ensure co	e supplying that information mpliance with Building Code.			
	require • relation accurace	ements/standards. Who has	responsibility? Is it equally irements and 'reach' offshore	etation of information to NZ y shared — joint responsibility? ore around ensuring information and updated?			
2.	, ,	• •	•	should be responsible for ler to comply with information			
		\square I agree in part	☐ No, I don't agree	☐ Not sure/no preference			
	Please explain yo	Please explain your views.					
		Accurate and evidenced information must be available for a product or system at its source, that means with the NZ manufacturer or the importer/distributor.					
3.	-	th the proposal that distri s they supply comply with		uld be responsible for ensuring nts?			
		\square I agree in part	☐ No, I don't agree	\square Not sure/no preference			
	Please explain yo	our views.					
	distributors and able to substant	retailers). All parties within	the system must act ethic	ss of role in supply chain (e.g., ally, responsibility and must be or quality assurance systems			
4.	requirements on	th MBIE's assessment of t n (1) manufacturers and in think the proposals will ha	nporters, and (2) distribu	itors and retailers? If not, what			
	Manufacturers a ⊠ Yes, I agree	nd importers: □ I agree in part	☐ No, I don't agree	☐ Not sure/no preference			
	Distributors and ⊠ Yes, I agree	retailers: ☐ I agree in part	☐ No, I don't agree	☐ Not sure/no preference			

Is there anything you would like to tell us about the reason(s) for your choice?

Any participant (e.g., business) in the supply chain has clear responsibilities and obligations to ensure building products/systems they manufacture/supply have met minimum information requirements and it is accurate. The discussion document is unfortunately silent on the issue of 'risk and liability' attributable to manufacturers and suppliers, outside of the modular components and MCM. Te Kāhui Whaihanga considers it important to acknowledge and protect 'downstream users of information' (e.g., architects and engineers) from code non-compliance to the extent that the non-compliance is attributable to a reasonable reliance on what turns out to be incorrect or incomplete product information or certification (whether wilful or careless) provided by a product manufacturer or supplier.

The supplied information from manufacturers/distributors would benefit from being accessible from an independent public e-portal. This ensures an ongoing archive of information and assist with sector/user trust and confidence of the repository of information.

Product information needs to be current (a HSWA requirement) and there needs to be version control so that product information available at the date the design services were provided forms the base for accountabilities. Such a system could ultimately be integrated with online consenting, which would provide efficient lodgement, review, and system assurances. Any system needs to be respected and developed with industry users if it is to meet professional's needs.

Content of information to be provided about building products

5.	Does the minimum set of information required for all building products look reasonable? If not, what information requirements should be added or removed?							
	☐ Yes	\square Yes, with changes	⊠ No	☐ Not sure/No preference				
	information Safety at W	a) <u>Health and Safety in Design</u> - There will be some overlap between the proposed building product information under the Building Act and the information requirements of PCBUs under the Health and Safety at Work Act 2015 (HSWA) in relation to structures at workplaces and 'designers' as upstream PCBUs and the associated duties associated with health and safety by design.						
	b) A structure as defined under that Act includes components and parts of structures. From a d perspective, the building product information supplied by manufacturers and suppliers should edesigners to comply with the information requirements of the HSWA and the additional duties in designers as upstream PCBUs.							
	c) <u>Scope and limitations on use</u> – there should be a minimum expectation of providing clear information and references to the relevant building code clauses and standards and limitations of unwhich are evidenced by testing.							
		<mark>ty periods</mark> - The information should or warranty with a clear claims pro		periods and be backed by a consistent				
		contact – there should be a contact and/or clarifications.	ct number and ema	il provided to ensure enquires can be				
	for climate	change programme and the MBIE information should be targeted aro	procurement guide	mitments included within the Building (climate change, recently added), ad embodied carbon, waste, carbon				
6.	Do you agree with the proposal that manufacturers and importers must make claims about how their building product meets relevant Building Code clauses?							
	⊠ Yes, I agr	ee 🔲 I agree in part	☐ No, I don't ag	ree ☐ Not sure/no preference				
	Is there anything you would like to tell us about the reason(s) for your choice?							
	This is essential information that should be made available and has been contributor to time/cost delays in the current system. Manufacturers and importers must demonstrate compliance and not just make a claim. Too often information is provided to 'market' a project/system rather than demonstrate compliance.							
	Compliance can be demonstrated by verification methods or providing suitable testing data to be verified by independent bodies. The certification and quality management system need to be reviewed/certified at appropriate intervals determined by a risk assessment. Unfortunately, there are too many examples of a "certified system" like Harditex, Monotex, VentClad to name a few that pass tests in a lab to receive a BRANZ appraisal and, they are not constructed onsite the same way.							
	accurately i	It will also be important to attribute the information to a point in time reference that can be sourced accurately into the future. Again, important technical product information, including testing data have been removed from online sources and paper-based catalogues, even though they need to be relied upon for consenting, warranty, and liability periods.						

7. What challenges would manufacturers and importers face in making claims about how the building product meets relevant Building Code clauses?

One of the key challenges will be accessing available knowledge and expertise regarding codifying the product/system to the NZ building code clauses. Building consenting functions around the country often struggle to retain and attract staff and alternative career pathways for this knowledge may be more attractive in a product role.

Clear guidance will be required for manufacturers and importers on the content of claims particularly regarding referencing – testing, supporting evidence, verifications, etc.

Manufacturers / importers have a duty to understand how their products meet relevant Building Code clauses and provide the relevant information to ensure the necessary performance standard is achieved. It is critical that information includes relevant Code Clauses for stated scope / limitations of use. (Proposal 3).

Manufacturers/importers who do not have the systems to provide this information must be prevented from supplying into the NZ market. The risk and liability otherwise transfer to downstream users of the information – architects, designers, and consent authorities.

8.	, 0		'	orters to use the compliance ompliance with the Building
	⊠ Yes, I agree	☐ I agree in part	□ No, I don't agree	☐ Not sure/no preference

This is a critical aspect of the system and provision of this information must be mandatory with no exceptions.

9. What other requirements or guidance would you recommend to ensure the information provided is relevant and accurate?

Some parameters or attributes that must be provided with the information:

- date of statement/performance claims

Please explain your views.

- a transparent version control process of information updates/changes.
- use of third-party verification and/or regular re-verification for critical products/systems.

Further work is needed on the information and evidence required in support of product substitutions.

Supply chain data and information standards

10.	the requirement t	o make evidenced claim	the likely impacts on manuns about the Building Code nk the proposals will have		
	⊠ Yes, I agree	\square I agree in part	☐ No, I don't agree	☐ Not sure/no preference	
	Is there anything y	ou would like to tell us	about the reason(s) for yo	our choice?	
	There will also be a positive impact for those manufacturers and importers who invest in quality information to the market.				

11.	. Do you agree that all information requirements should be met prior to supply of a building product and that information be kept up to date with the latest version of that product? If not, what other requirements do you think would be reasonable?					
	⊠ Yes, I agree	\square I agree in part	☐ No, I don't agree	☐ Not sure/no preference		
	Is there anything y	ou would like to tell us	about the reason(s) for yo	our choice?		
	It is vital that product information is time referenced, accessible historically, kept current and updated when there are any changes made to the content or manufacturing process of a building product and/or in response to regulatory/code changes.					
	be evidenced to m	neet its stated performand	not be released to market use and product information is /substituted for earlier versions.			
12.	Do you agree that the supply chain a		oe provided in structured (data and accessible across		
		\square I agree in part	☐ No, I don't agree	☐ Not sure/no preference		
	Please explain you	r views.				
	adapt and require		structure is essential – and the added (e.g., safety in design in the sign in	nere needs to be flexibility to 2015, operational carbon		
13.	Do you think it is r be made available		l information to be disclos	ed about building products to		
		\square I agree in part	☐ No, I don't agree	☐ Not sure/no preference		
	Is there anything y	ou would like to tell us	about the reason(s) for yo	our choice?		
	Absolutely.					
14.	-	the proposal for all bui mation provided online		nique identifiable code that		
	☐ Yes, I agree	⊠ I agree in part	☐ No, I don't agree	☐ Not sure/no preference		
	Is there anything y	ou would like to tell us	about the reason(s) for yo	our choice?		
	Clearly an easier of	oncept to administer for p	oroducts – more challenging	for systems.		

Transition period

15. Do you agree with proposal for an 18 month transition period after building product information requirement regulations are made before they come into force? If not, what would be a reasonable timeframe?

☐ Yes, I agree	☑ I agree in part	☐ No, I don't agree	☐ Not sure/no preference
transition (say 12	months). The intent for i		rry performance with a shorter on has been well signalled to the ow.

Modular component manufacturer certification scheme

Prescribing the kinds of building products that would be 'modular components' and scopes of certification

such as open frames and trusses, enclosed panels/units, volumetric structures, and whole buildings as 'modular components'?							
	☐ Yes, I agree	\square I agree in part	☐ No, I don't agree	\square Not sure/no preference			
	Please explain yo	our views.					
	[insert response	here]					
2.		ufacturer certifiction boo	efit in developing a syster dies describe the scope o				
	[insert response	here]					
3.	Which, if any, of system do you pr		n which to base the propo	osed scope of certification			
	\square Option 1	\square Option 2	☐ Option 3	\square Not sure/no preference			
	Please explain yo	Please explain your views.					
	[insert response	here]					
	odular compo gistration	nent manufacturei	certification body	accreditation and			
4.	Do you think the proposed regulatory settings provide confidence in the certification bodies that would be accredited and registered within the modular component manufacturer certification scheme?						
	-	tory settings to be accred	dited:				
	☐ Yes	☐ Yes, with changes	□ No	☐ Not sure/No preference			
	Proposed regulat ☐ Yes	tory settings to be registed \Box Yes, with changes	ered: No	☐ Not sure/No preference			
	Please explain your views.						
	[insert response	here]					
5.	•	How do you think the proposed regulatory settings for certification bodies might affect their uptake of the modular component manufacturer certification scheme?					
	[insert response	here]					

Modular component manufacturer certification scheme

Modular component manufacturer certification and registration

6.	Do you think the proposed regulatory settings provide confidence in the modular component manufacturers that would be certified and registered within the scheme?						
	Proposed regula ☐ Yes	tory settings to be certified:	□ No	☐ Not sure/No preference			
	Proposed regula						
	☐ Yes	\square Yes, with changes	□ No	\square Not sure/No preference			
	Please explain yo	our views.					
	[insert response here]						
7.	•	e proposed regulatory settings to resumer protection?	for modular compon	ent manufacturers provide			
	☐ Yes	\square Yes, with changes	□ No	☐ Not sure/No preference			
	Please explain your views.						
	[insert response	e here]					
8.	How might the proposed regulatory settings for modular component manufacturers have different impacts for different kinds of manufacturers that may wish to participate in the scheme?						
	[insert response	here]					
9.	To what extent do you think modular component manufacturers will benefit from the proposed regulatory settings, and what costs do you think they might face when trying to meet the proposed settings?						
	[insert response	here]					

certificates might have?

[insert response here]

Modular component manufacturer certification scheme

Audits within the modular component manufacturer scheme 10. Do you agree with the proposal that auditing parties will use a prescribed risk assessment to decide the frequency and type of audits they will use for those being audited? ☐ Yes, I agree ☐ I agree in part ☐ No, I don't agree ☐ Not sure/no preference Please explain your views. [insert response here] 11. What costs do you think the proposed audit requirements might have for modular component manufacturers, given that the fees for audits would be set through contract between the manufacturer and its modular component manufacturer certification body? [insert response here] 12. Do you agree with modular component manufacturer certification bodies and modular component manufacturers having three months to make changes outlined in an audit report following an audit? Please explain your views. ☐ Yes, I agree ☐ I agree in part ☐ No, I don't agree ☐ Not sure/no preference Please explain your views. [insert response here] Modular component manufacturer's certificates 13. Do you support manufacturers being responsible for transportation, storage and assembly of modular components that they manufacture within the modular component manufacturer certification scheme? What impacts might this have on manufacturers? ☐ Yes ☐ Yes, with changes ☐ No ☐ Not sure/No preference [insert response here] 14. To what extent do you think the information that is proposed to be required on manufacturer's certificates will provide clarity for different parties within the modular component manufacturer certification scheme? [insert response here] 15. What costs do you anticipate that providing the proposed information on manufacturer's

Product certification scheme

Implement registration requirements for product certification bodies

1.	Do you consider that the proposed fit and proper test and notification requirements would be effective criteria to establish if a product certification body should operate in the scheme?						
	⊠ Yes	☐ Yes, with changes	□ No	☐ Not sure/No preference			
	Please explain your views.						
	[insert response	here]					
2.		· · ·	escribe an adequate mean his stage? Please explain y	•			
	☐ Yes, I agree	☑ I agree in part	☐ No, I don't agree	☐ Not sure/no preference			
	Please explain yo	our views.					
	A standard 'base	eline' technical competency	evaluation should be include	ed in the PCB application process.			
3.	•	that MBIE has proposed t roduct certification body	the right requirements for registration?	what must go on an			
	⊠ Yes	\square Yes, with changes	□ No	\square Not sure/No preference			
	Is there anything you would like to tell us about the reason(s) for your choice?						
	[insert response here]						
lm	plement regis	stration requiremen	its for certificates				
4.	Do you agree with the MBIE's assessment that the proposals for certificate information will improve the usability of product certificates?						
	⊠ Yes, I agree	\square I agree in part	☐ No, I don't agree	☐ Not sure/no preference			
	Is there anything you would like to tell us about the reason(s) for your choice?						
	[insert response	here]					
5.	Are there any gaps or issues with current certificates that MBIE have missed that should be addressed by changes to Regulation 14 or Schedule 2?						
	[insert response	here]					

Product certification scheme

Improve scheme requirements for product certification body accreditation

6.	Do you consider that the product certification body accreditation proposals will improve the alignment of scheme documents?						
	oximes Yes $oximes$ Yes, with changes $oximes$ No $oximes$ Not sure/No preference						
	Please explain yo	our views.					
	[insert response	here]					
7.	•	there will be any co oposals? If so, what a	•	sues with the pr	oduct certification body		
	⊠ Yes	□ No	☐ Not sur	e/No preference	2		
	Is there anything you would like to tell us about the reason(s) for your choice?						
	Potential for increased costs to manufacturer who has to pay for PCB fees. Likelihood is that these will be passed through the supply chain and ultimately the Client.						
8.	What further clarification related to the proposal to require product certification bodies to only accept test reports from competent testing facilities may be required?						
	Strengthening of CodeMark will require certainty that PCB's will only accept test reports from competent testing facilities certified as meeting International Standards.						
9.	Do you agree wi	th proposal 8 to revo	oke existing	Regulation 7A?			
	 ⊠ Yes, I agree Please explain years 	\square I agree in parour views.	t 🗆 I	No, I don't agree	☐ Not sure/no preference		
	[insert response	here]					

Product certification scheme

Strengthen requirements for product certification body audits and reviews of certificates

10.	reasonable? If not, what requirements should be amended, added or removed?					
	⊠ Yes	\square Yes, with chang	es	□ No	☐ Not sure/No preference	
	Please explain yo	our views.				
11.	What cost impacts do you consider the product certification body audit proposals will have? Will costs change compared to the current requirements?					
	There may be some increase to cover certification requirements.					
12.	Is three years th	e correct minimum f	frequency fo	or certification rev	view?	
	☐ Yes	⊠ No	☐ Not sur	e/No preference		
	Please explain your views.					
	Certification rev		cted followir	g a risk-based appı	roach, with three years being the	

Regulated fees for the modular component manufacturer certification scheme and the product certification scheme

Regulated fees for the modular component manufacturer certification scheme and the product certification scheme

Registration fees for modular component manufacturer certification scheme

1.	Do you agree with MBIE's estimated cost drivers for modular component manufacturer certification body and modular component manufacturer registration?					
	☐ Yes, I agree ☐ I agree in part ☐ No, I don't agree ☐ Not sure/no preference ☐ Please explain your views.					
	[insert response here]					
2.	To what extent might the prescribed registration fees create a barrier to entry and ongoing participation in the scheme?					
	[insert response here]					
Accreditation and audit fees for modular component manufacturer certification scheme						
	Do you agree with MBIE's assumption that the fee structure and level for assessing modular component manufacturer certification body accreditation is comparable to that for assessing building consent authority accreditation?					
	☐ Yes, I agree ☐ I agree in part ☐ No, I don't agree ☐ Not sure/no preference ☐ Please explain your views.					
	[insert response here]					
	Do you agree with MBIE's proposed fee structure for modular component manufacturer certification body accreditation and audits?					
	☐ Yes, I agree ☐ I agree in part ☐ No, I don't agree ☐ Not sure/no preference					
	Is there anything you would like to tell us about the reason(s) for your choice?					
	[insert response here]					
	To what extent might the prescribed audit fees create a barrier to entry and ongoing participation in the scheme?					
	[insert response here]					

Regulated fees for the modular component manufacturer certification scheme and the product certification scheme

Registration fees for product certification scheme

6.	Do you agree with MBIE's assessment of the options for structuring registration fees for product certification bodies and certificates? Please explain your views.				
	☐ Yes, I agree	\square I agree in part	t □ No, I don't agree	☐ Not sure/no preference	
	Please explain your views.				
	[insert response	here]			
	Do you consider that the proposed fees for registration of product certification bodies and certificates are set at the right level? Please explain your views.				
	☐ Yes	\square Yes, with change	es 🗆 No	\square Not sure/No preference	
	Please explain yo	ur views.			
	[insert response	here]			
Ac	creditation an	d audit fees for	r product certification so	cheme	
	Would the proposed fees for product certification body accreditation and audits of product certification bodies create any practical issues? If so, what would the issues be?				
	☐ Yes	⊠ No	\square Not sure/No preference		
	Is there anything	you would like to te	ell us about the reason(s) for yo	our choice?	
	Do you consider that the proposed fees for product certification body accreditation and audits of product certification bodies are set at the right level?				
	⊠ Yes	□ No	☐ Not sure/No preference		
	Please explain yo	ur views.			
Exp	pected impact	ts			
	Will the prescribed fees have a significant impact on the costs of participating in the schemes?				
	□ Yes	⊠ No	☐ Not sure/No preference		
	Is there anything	you would like to te	ell us about the reason(s) for yo	our choice?	
11.	Do vou have anv	other comments on	the proposals?		
	2 ,22 2		· · · · · · · · · · · · · · · · · · ·		