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1 July 2021



RE Official Information Act request CDHB 10622

I refer to your email dated 1 June 2021 requesting the following information under the Official Information Act from Canterbury DHB regarding children in seclusion in mental health facilities. Specifically:

1. Number of seclusion incidents each year since 2018 until today for patients aged under 12.

There are no seclusion incidents recorded for this age group during the date range of your request.

2. Number of seclusion incidents each year since 2018 until today for patients aged 13-18.

Please refer to **Table one** (below).

Table one: Seclusion events 2018 to 2021 year to date for patients aged 13-18

Years	Seclusion events
2018	6
2019	19
2020	27
2021	13

Please note We have included 18-year olds, so some of the events were for people under adult services.

3. How long were patients in seclusion over that period and in what conditions? Please provide a breakdown of average, minimum and maximum seclusion duration for each year (2018, 2019, 2020 and 2021 to date if possible) for patients aged under 12, and for patients aged 13-18.

Please refer to **Table two** (overleaf).

Table two: Seclusion hours for patients aged 13-18 for years 2018-2021 year to date.

Years	Sum of Duration Hours	Minimum of Duration Hours	Maximum of Duration Hours	Average of Duration Hours
2018	52.44	0.2	16.08	8.74
2019	276.77	1.17	26.67	14.57
2020	220.19	0	48	8.16
2021	181.79	1.67	43.67	13.98

Notes: There are no seclusion incidents recorded for patients under the age of 12 years during the date range of your request. We have included 18-year olds, so some of the events were for people under adult services.

As per the Health and Disability services standards attached as **Appendix 1**, Child and Family Inpatient Service uses seclusion for safety reasons only and seclusion only occurs in an approved and designated seclusion room.

Designated seclusion rooms are audited annually by the Chief of Psychiatry and the Director of Nursing. The audit tool used for this is attached as **Appendix 2**. A person can only be placed in seclusion in accordance with the provisions outlined in s7 of the Mental Health (Compulsory Assessment and Treatment) Act 1992.

4. Number of patients put in seclusion each year since 2018 with their age, or age range if possible, and how many times they were placed in seclusion.

Please refer to **Table three** (below).

Table three: Number of patients aged 13-18 put in seclusion for years 2018 -2021 year to date*

Years	Distinct Count of HCU
2018	4
2019	6
2020	10
2021	3

^{*}Number of events per person has not been provided on the basis of privacy due to low numbers. Declined pursuant to section 9(2)(a) of the Official Information Act.

Notes:

- There are no seclusion incidents recorded for patients under the age of 12 years during the date range of your request.
- We have included 18-year olds, so some of the events were for people under adult services.

I trust that this satisfies your interest in this matter.

You may, under section 28(3) of the Official Information Act, seek a review of our decision to withhold information by the Ombudsman. Information about how to make a complaint is available at www.ombudsman.parliament.nz; or Freephone 0800 802 602.

Please note that this response, or an edited version of this response, may be published on the Canterbury DHB website after your receipt of this response.

Yours sincerely

Tracey Maisey

Executive Director

Planning, Funding & Decision Support

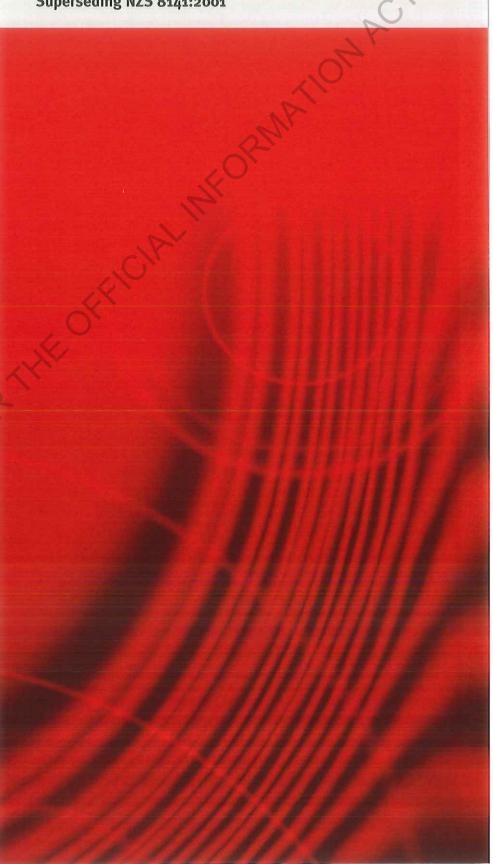


New Zealand Standard

APPENDIX 1

Health and Disability Services (Restraint Minimisation and Safe Practice) Standards -Seclusion

Superseding NZS 8141:2001



RELEASED UNDER THE OFFICIAL INFORMATION ACT

New Zealand Standard

HEALTH AND DISABILITY SERVICES (RESTRAINT MINIMISATION AND SAFE PRACTICE) STANDARDS

2.3: SECLUSION
NOHO MÖWAI

RELEASED UNDER THE OFFICIAL INFORMATION ACT

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AFELERAGED UNDER THE OFFICIAL INFORMATION ACT

RELEASED UNDER THE OFFICIAL INFORMATION ACT

FOREWORD

The main intent of NZS 8134.2 is to reduce the use of restraint in all its forms and to encourage the use of least restrictive practices. It is crucial that providers recognise which interventions constitute restraint and how to ensure that, when practised, restraint occurs in a safe and respectful manner.

Restraint should be perceived in the wider context of risk management. Restraint is a serious intervention that requires clinical rationale and oversight. It is not a treatment in itself but is one of a number of strategies used by service providers to limit or eliminate a clinical risk. Restraint should only be used in the context of ensuring, maintaining, or enhancing the safety of the consumer, service providers, or others. All restraint policies, procedures, practices, and training should be firmly grounded in this context.

This Standard covers all forms of restraint and supersedes NZS 8141:2001.

NZS 8134.2 *Health and disability services (restraint minimisation and safe practice) Standards* includes referenced and related documents and guidelines, guidance on restraint minimisation and safe practice, and on the application of the Standard, along with the following Standards:

- (a) NZS 8134.2.1 Restraint minimisation
- (b) NZS 8134.2.2 Safe restraint practice
- (c) NZS 8134.2.3 Seclusion.

Each is to be read in conjunction with NZS 8134.0 *Health and disability services (general) Standard*, as this contains the definitions and audit framework information applicable across the health and disability suite

ETHICAL AND LEGAL CONSIDERATIONS

Practice is guided by ethical principles that include acting for the consumer's good (beneficence), avoiding harm to the consumer (non-maleficence), avoiding harm to self and others, and respecting the dignity of the consumer and the consumer's human rights.

The Standard should be implemented in ways that respect these and other ethical principles and at all times promote the interests, safety, and well-being of all involved.

Any unauthorised restriction on a consumer's freedom of movement could be seen as unlawful. Organisations should develop clear policies and procedures to guide service providers in the implementation of the Standard, and seek legal advice if necessary.

Seclusion and restraint shall not be used by providers for punitive reasons.

MEDICATION

The term chemical restraint is often used to mean that rather than using physical methods to restrain a consumer at risk of harm to themselves or others, various medicines are used to ensure compliance and to render the person incapable of resistance. Use of medication as a form of 'chemical restraint' is in breach of this Standard.

All medicines should be prescribed and used for valid therapeutic indications. Appropriate health professional advice is important to ensure that the relevant intervention is appropriately used for therapeutic purposes only.

G 3

Seclusion is a form of restraint that can only be used within the Mental Health (Compulsory Assessment and Treatment) [MH (CA and T)] Act 1992, and the Intellectual Disability (Compulsory Care and Rehabilitation [ID(CCR)] Act 2003.

Seclusion can be legally implemented subject to the conditions specified in the MH (CA and T) Act. The legal basis of seclusion for consumers under MH (CA and T) Act is set out in s71 of the Act and s148 of ID(CCR) Act. Seclusion should be used for as short a time as possible and is best conceived as a safety mechanism rather than a therapeutic intervention or treatment. The decision to seclude should be an uncommon event, used as a final alternative and subject to strict review. The information in NZS 8134.2.3 is provided with the expectation that although seclusion is legal, services will be proactive in reducing and minimising/avoiding its use.

Seclusion should not occur as part of a routine admission procedure or for punitive reasons. The MH (CA and T) Act requires that, except in an emergency, seclusion shall be used only with the authority of the responsible clinician. If not involved in the immediate decision, the responsible clinician must be informed of the seclusion as soon as possible, at least at the start of the next working day, and should review the decision. A doctor must assess the secluded consumer as soon as possible; this should be within two hours. The specificity of the assessment shall be appropriate to the level of risk and likelihood of harm occurring to the consumer. Wherever practicable, the two clinicians involved should be the consumer's own nurse and doctor.

The ID (CCR) Act requires that, except in an emergency, seclusion may only be used with the authority of a care manager who must ensure the care recipient is not secluded for longer than is necessary to achieve the purpose of secluding the care recipient.

- **G 3.1.1** Procedural guidelines for the use of seclusion can be found on http://www.moh.govt.nz.
- **G 3.1.2** Seclusion should only be used to prevent violent behaviour compromising safety.
- G 3.1.4 Seclusion should only be used with great caution, and with intensive monitoring in the following circumstances:
 - (a) Where the consumer has had escalating requirements for medication and there is:
 - (i) Evidence of serious recent side effects
 - (ii) Likelihood of serious side effects;
 - (b) Physical deterioration;
 - (c) Where the consumer is in need of intensive assessment and/or observation, especially where there is a history suggestive of trauma, ingestion of unknown drugs/substances, or organic diagnosis.
- **G 3.2.4** In addition to the environment being safe, wherever possible:
 - (a) Doors should open outwards;
 - (b) While in seclusion consumers should be able to wear their own clothing and retain some of their personal possessions if their safety is not compromised.
 - (c) Items are provided if required by the consumer, if their safety is not compromised.
 - (d) A means of orientation (time, date, news, and other information) is provided;
 - (e) The service provider facilitates timely access to washing, showering, and toilet facilities in or adjacent to the area;
 - (f) There is access to a safe external area to assist with reintegration.

SECLUSION NOHO MŌWAI

Outcome 3 Consumers receive services in the least restrictive manner.

Seclusion shall only be used by services with approved seclusion facilities. All use of seclusion shall comply with NZS 8134.2.2 and NZS 8134.2.3.

SAFE SECLUSION USE WHAKAMAHI MÖWAI HAUMARU

Criteria

	Standard 3.1	Servic	es demonstrate that all use of seclusion is for safety reasons only.
	Criteria	The cri	teria required to achieve this outcome shall include the organisation ensuring:
		3.1.1	The service has policies and procedures on seclusion that meet the requirements contained in 'Seclusion under the Mental Health (Compulsory Assessment and Treatment Act 1992' (MoH).
MHA* & ID**		3.1.2	Consumers are subject to the use of seclusion when there is an assessed risk to the safety of the consumer, to other consumers, service providers, or others.
es II)^^ 	3.1.3	There exists a legal basis for each episode of seclusion.
		3.1.4	Any factors that may require caution must be assessed for each episode.

therapeutic relationships is considered and documented.

The likely impact the use of seclusion will have on the consumer's recovery and

APPROVED SECLUSION ROOMS RUMA MÕWAI KUA WHAKAAFA

3.1.5

Seclusion only occurs in an approved and designated seclusion room. Standard 3.2

	3.2.1 The seclusion room provides adequate lighting, room temperature, and ventilation.
	3.2.2 The seclusion room allows the observation of the consumer and allows the
	consumer to see the head and shoulders of the service provider.

The seclusion room provides a means for the consumer to effectively call for attention.

The criteria required to achieve this outcome shall include the organisation ensuring:

The seclusion room contains only furniture and fittings chosen to avoid the potential for harm.

applies to mental health and addiction services only

^{**} applies to intellectual disability services only

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Approval of Minister of Health received on 25 September 2008. Approved by the Standards Council on 25 September 2008 to be a New Zealand Standard pursuant to the provisions of section 10 of the Standards Act 1988.

JUNDER THE OFFICIAL INFORMATION ACT

First published: 8 October 2008

The following SNZ references relate to this Standard:

Project No. P 8134 Draft for comment No. DZ 8134 Typeset by: Standards New Zealand Printed by: The Colour Guy







Seclusion Room / Area Audit Tool Audit date: Unit: Criteria Rm. No. Rm. No. Rm. No. Rm. No. **Comments and recommendations** Adequate light, heat, ventilation Means of easy observation, that also allows the consumer to see the head and shoulders of the observer Means for a secluded consumer to call for attention Fittings recessed to avoid potential for harm Doors open outwards flush with walls and the environment should be pleasant and minimally stimulating Assistance be given to providing a means of orientation (time, date, news and other information)





	Seclusion Room / Area Audit Tool					
Un	it:					Audit date:
	Criteria	Rm. No.	Rm. No.	Rm. No.	Rm. No.	Comments and recommendations
7	There is access to washing or showering facilities in or adjacent to the area				MION	
8	There is access to an equally safe area to assist with reintegration			CICIA		
Additional comments and recommendations:						
Ар	Approved for use Date:					
Re	Repairs completed Confirmation date:					
Sig	Signature Director of Area Mental Health Services:					Date:
Signature Director of Nursing:					Date:	