End of Life Choice Bill

Member's Bill

Explanatory note

General policy statement

Purpose

This Bill gives people with a terminal illness or a grievous and irremediable medical condition the option of requesting assisted dying.

The motivation for this Bill is compassion. It allows people who so choose, and are eligible under this Bill, to end their lives in peace and dignity, surrounded by loved ones

The Bill carefully defines those eligible for assisted dying, details a comprehensive set of provisions to ensure this is a free choice, made without coercion, and outlines a stringent series of steps to ensure the person is mentally capable of understanding the nature and consequences of assisted dying.

Background

Bills relating to assisted dying have been debated twice before by the New Zealand Parliament. The first time was in 1995, when Michael Laws' Death with Dignity Bill was defeated 61/29 at first reading. The second time was in 2003, when Peter Brown's Death with Dignity Bill was defeated 60/58 at first reading.

Since that time, evidence and developments have established that there are serious problems with the current state of the law in New Zealand that will be ongoing without a legislative solution. Extensive evidence and analysis considered in the case of *Seales v Attorney-General* [2015] NZHC 1239 demonstrated that, without a change in the law, some people in New Zealand are suffering unbearably at the end of the lives and are taking their lives earlier than they would if assisted dying were legally available to them. There was broad consensus that palliative care cannot alleviate all suffering, including suffering that is unbearable for a person.

Continuation of the status quo cannot be an option when the risks can be managed and the law targeted to the small but significant group of competent adults who are not vulnerable and who wish to die without unbearable suffering and pain. Analysis of overseas jurisdictions where assisted dying is permitted demonstrates that concerns, including concerns about abuse of the vulnerable, have not materialised and that risks can be properly managed through appropriate legislative safeguards. In countries where assisted dying is permitted, medical practitioners and organisations have adapted well, developing guidelines that inform ethical and medical practice.

A change in the law will bring benefits beyond alleviating unbearable suffering and avoiding people being at risk of premature death. The evidence considered in *Seales* and overseas studies show that, when assisted dying is permitted, the quality and uptake of palliative care increases and the doctor-patient relationship is positively enhanced.

In other cases in New Zealand, the courts are treating the family members who have assisted their loved ones to die at their request with increasing leniency and compassion.

Evidence of this includes a case in 2008 in which a sentence of 6 months community detention plus 150 hours community work was given to a man who assisted his mother to die, when his mother had been suffering from the final stages of terminal stomach cancer. Another case in 2012, of a man who whose wife had been suffering from primary progressive multiple sclerosis (PPMS), resulted in a discharge without conviction for his assisting his wife to die. The public reaction to such trials has been overwhelmingly compassionate and understanding.

These cases highlight the very difficult position family members can be placed in when assisted dying is not legally available for their loved ones. It also demonstrates further issues with the current state of our law, under which it is becoming permissible, in effect, for family members to assist loved ones to take their own lives. This is clearly less ideal, less clear, and considerably more risky than a regulated process in which medical practitioners can, in limited circumstances, assist those who are suffering.

The state of the law in New Zealand is increasingly out of step with public opinion and with developments overseas. The *Seales* case has promoted nationwide debate around assisted dying and has demonstrated evidence of strong public support for legalising it. There will be those who remain opposed to assisted dying, including those who are opposed because of their personal views and religious or cultural beliefs. This Bill ensures that medical practitioners who are opposed to the practice are under no obligation to advise on or provide assisted dying.

This Bill provides a legal pathway that prevents medical practitioners from being charged with an offence when the request for assisted dying comes from the expressed will of a person who is suffering unbearably.

Under this Bill, it remains a criminal offence to assist a person to die except by an action undertaken by a medical practitioner in the very limited circumstances pre-

scribed. This Bill will provide the option of relief to a small but significant number of people who suffer unendurably during the terminal days or weeks of a difficult illness, despite the best that palliative care can offer, or to those who suffer unendurably as a result of a grievous and irremediable medical condition, despite the best available medical care.

Main Provisions

The Bill defines a person eligible for assisted dying as someone who:

- is aged 18 years or over
- has New Zealand citizenship or is a permanent resident
- suffers from a terminal illness likely to end their life within 6 months or has a grievous and irremediable medical condition
- is in an advanced state of irreversible decline in capability
- experiences unbearable suffering that cannot be relieved in a manner that he or she considers tolerable
- has the ability to understand the nature and consequences of assisted dying.

The Director-General of Health will establish a group of medical practitioners, serviced by the Ministry of Health, known as the Support and Consultation for End of Life in New Zealand (SCENZ) Group. This group will maintain a list of medical practitioners, specialists in mental health, and pharmacists willing to participate in assisted dying. The group will be responsible for allocating replacement and independent medical practitioners, thereby ensuring that the attending medical practitioner does not choose the replacement or independent practitioner. The group will write standards of care, provide advice on medical and legal procedures, and provide practitioners practical assistance if requested.

A Registrar (assisted dying) will be nominated by the Director-General of Health and will have a number of responsibilities under this Bill. These responsibilities include maintaining a register of forms lodged with the Registrar as this Bill requires, cosigning prescriptions, establishing a procedure to deal with complaints, and reporting to the Minister of Health and End of Life Review Committee as required.

No person is obligated to take a role under this Bill, although medical practitioners who conscientiously object must refer people to the SCENZ Group.

Process

A legal process will be created for a person who asks to receive assisted dying. Two medical practitioners will have to be satisfied that the person meets the criteria required.

The attending medical practitioner must provide the person with the prognosis for the terminal illness or grievous and irremediable medical condition and inform the person of the irreversible nature and anticipated impacts of assisted dying. The medical practitioner must also encourage the person to talk about his or her choice with family, friends, and counsellors, ensure that the person has had the opportunity to talk to his

or her choice of people, and do his or her best to ensure that the person has chosen assisted dying free of any pressure or coercion. Should the person wish to proceed with his or her choice, the medical practitioner must provide the prescribed form to the person.

The medical practitioner must then contact the SCENZ Group, which will refer the person requesting assisted dying to a second medical practitioner, who must be independent of the person and of the initial medical practitioner. The second practitioner must read the person's files and examine the person and decide whether the person is a person who is eligible for assisted dying.

After the reading of the person's files and the examination of the person have occurred, should either medical practitioner be uncertain about whether the person requesting assisted dying is competent, the two medical practitioners must jointly refer the person to a specialist with a relevant scope in mental health. The specialist must read the person's files, examine the person, and provide his or her opinion about whether the person requesting assisted dying is competent.

Once this process has occurred, the person must be informed about whether he or she is eligible for assisted dying. Should a person be told he or she is not eligible, the reason for this must be explained.

At each step of this process, a prescribed form – the signed request form, the opinion of the attending medical practitioner, the opinion of the independent medical practitioner, and when required the opinion of the specialist – must be sent independently to the Registrar, so that the Registrar has a complete record of the decisions taken and opinions formed, regardless of whether the request has been approved or declined.

In the case that the person is found to be a person eligible for assisted dying, the medical practitioner must discuss with the person the likely timing and make provisional arrangements to be available. When the person wishes to exercise the option of assisted dying, he or she must tell the medical practitioner, who must inform the person of the medication options available, ask the person to choose one of them and to choose a suitable time for the administration of the prescribed medication, and ensure that the person knows that he or she can change his or her mind at any time. At least 48 hours prior to the administration of the prescribed medication, the practitioner must inform the registrar, who will check that all safeguards have been correctly followed and will then co-sign the prescription for the medication.

At the chosen time of the administration, the medical practitioner must confirm that the person wishes to receive the medication. Should the person wish to receive the medication, the medical practitioner must administer it and either be available to the person until death occurs or arrange for another medical practitioner to be available to the person until death occurs. The medical practitioner must then report the death to the registrar within 14 working days.

A review of the operation of this Act must start three years after commencement and be completed within six months. The report must be presented to Parliament. Thereafter, a review must be conducted every five years.

Clause by clause analysis

Clause 1 states the title.

Clause 2 describes the commencement date.

Clause 3 defines terms.

Clause 4 contains the definition of person who is eligible for assisted dying.

Clause 5 provides that the Act binds the Crown.

Clauses 6 to 18 deal with assisted dying.

Clause 19 to 22 set out accountability mechanisms.

Clauses 23 to 28 cover a range of related matters.

David Seymour

End of Life Choice Bill

Member's Bill

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The	Parliament of New Zealand enacts as follows:
1	Title
	This Act is the End of Life Choice Act 2017.
2	Commencement
	This Act comes into force 6 months after the date on which it receives the Royal assent.
	Part 1
	Preliminary provisions
3	Interpretation
	In this Act, unless the context requires another meaning,—
	assisted dying means the administration by a medical practitioner of a lethal dose of medication to a person to relieve his or her suffering by hastening death
	attending medical practitioner means a person's medical practitioner
	competent means having the ability described in section 4(f)
	conscientious objection means an objection to doing anything authorised or required by this Act
	Director-General means the Director-General of Health
	health practitioner has the meaning given to it by section 5 of the Health Practitioners Competence Assurance Act 2003
	independent medical practitioner means a medical practitioner who is independent of an attending medical practitioner and the person
	medical practitioner means a health practitioner who—

(a)	_	gistered with the Medical Council of New Zealand as a practitioner e profession of medicine or is deemed to be so registered; and	
(b)	holds	a current practising certificate	
		eans the Minister of the Crown who is responsible for the adminisis Act—	5
(a)	unde	the authority of a warrant; or	
(b)	unde	the authority of the Prime Minister	
mini	stry m	eans the Ministry of Health	
perso tion		o is eligible for assisted dying has the meaning given to it in sec-	10
phar	macist	means a health practitioner who—	
(a)	-	gistered with the Pharmacy Council as a practitioner of the profes- of pharmacy or is deemed to be so registered; and	
(b)	holds	a current practising certificate	
psycl psycl		t means a medical practitioner whose scope of practice includes	15
psycl	nologis	st means a health practitioner who—	
(a)	_	sistered with the Psychologists Board as a practitioner of the profes- of psychology or is deemed to be so registered; and	
(b)	holds	a current practising certificate	20
regis	trar m	eans the registrar (assisted dying) nominated under section 21	
revie	w com	mittee means the committee established under section 20	
SCE	NZ me	ans Support and Consultation for End of Life in New Zealand	
SCE	NZ Gr	oup means the body established under section 19	
speci	alist n	neans a psychiatrist or a psychologist.	25
	_	f person who is eligible for assisted dying	
In thi	s Act,	person who is eligible for assisted dying means a person who—	
(a)	is age	ed 18 years or over; and	
(b)	is—		
	(i)	a person who has New Zealand citizenship as provided in the Citizenship Act 1977; or	30
	(ii)	a permanent resident as defined in section 4 of the Immigration Act 2009; and	
(c)	suffe	rs from—	
	(i)	a terminal illness that is likely to end his or her life within 6 months; or	35

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		(ii)	a grievous and irremediable medical condition; and					
	(d)	is in	an advanced state of irreversible decline in capability; and					
	(e)	(e) experiences unbearable suffering that cannot be relieved in a manner that he or she considers tolerable; and						
	(f)	has t	he ability to understand—	5				
		(i)	the nature of assisted dying; and					
		(ii)	the consequences for him or her of assisted dying.					
5	Act	binds 1	the Crown					
	This	Act bi	nds the Crown.					
			Part 2	10				
			Assisted dying					
6	Con	scienti	ous objection					
(1)			oes not require a person to do anything to which the person has a bus objection.					
(2)	Sub	sectio	on (1)—	15				
	(a)		ies despite any legal obligation to which the person is subject, how- the obligation arises; and					
	(b)	does	not apply to the requirement in section 7(2) .					
7	Effe	ct of c	onscientious objection					
(1)	This	section	n applies when—	20				
	(a)	-	rson tells the attending medical practitioner under section 8(1) that person wishes to have the option of receiving assisted dying; and					
	(b)	the a	ttending medical practitioner has a conscientious objection.					
(2)	The	attendi	ng medical practitioner must tell the person that—					
	(a)	the n	nedical practitioner has a conscientious objection; and	25				
	(b)		person may ask the SCENZ Group for the name and contact details replacement medical practitioner.					
(3)	in th	is Act	on chooses to have the replacement medical practitioner, references to the attending medical practitioner mean the person's replacement actitioner, except in subsection (2) and section 8(1) .	30				
8	Req	uest m	ade					
(1)			who wishes to have the option of receiving assisted dying must tell ng medical practitioner of his or her wish.					
(2)	The	attendi	ng medical practitioner must—					

the prognosis for the terminal illness or grievous and irremediable

give the person the following information:

(a)

(i)

		. ,	medical condition; and	
		(ii)	the irreversible nature of assisted dying; and	
		(iii)	the anticipated impacts of assisted dying; and	5
	(b)		with the person about his or her wish at intervals determined by the ress of his or her terminal illness or medical condition; and	
	(c)	ensur care;	re that the person understands his or her other options for end of life and	
	(d)		re that the person knows that he or she can change his or her mind at time; and	10
	(e)		urage the person to talk about his or her wish with others such as ly, friends, and counsellors; and	
	(f)	ensur one;	re that the person knows that he or she is not obliged to talk to any- and	15
	(g)		re that the person has had the opportunity to talk about his or her with those whom he or she chooses; and	
	(h)		is or her best to ensure that the person expresses his or her wish free pressure from any other person by—	
		(i)	talking with other health practitioners who are in regular contact with the person; and	20
		(ii)	talking with members of the person's family approved by the person; and	
	(i)	sisted	plete the first part of the prescribed form requesting the option of asd dying by recording the actions he or she took to comply with agraphs (a) to (h).	25
9	Requ	iest co	onfirmed	
(1)	This	section	n applies after section 8 is complied with.	
(2)		-	on wishes to proceed, the attending medical practitioner must give the prescribed form requesting the option of assisted dying.	30
(3)	The j	person	must—	
	(a)	sign a	and date the second part of the form; or	
	(b)	-	resent when the second part of the form is signed and dated as deed in subsection (4) .	
(4)			part of the form may be signed and dated by a person other than the shom it relates if—	35
	(a)	the p	erson to whom it relates cannot write for any reason; and	

	(b)	the p	person to whom it relates requests the other person to sign and date ad		
	(c)		person who signs and dates the part notes on it that he or she did so the person to whom the form relates; and		
	(d)	the p	erson who signs and dates the part is not—	5	
		(i)	a health practitioner caring for the person to whom the part relates; or		
		(ii)	a person who knows that he or she stands to benefit from the death of the person to whom the part relates; or		
		(iii)	a person aged under 18 years; or	10	
		(iv)	a person with a mental disability.		
(5)	The	attendi	ng medical practitioner must—		
	(a)	be pr	resent when—		
		(i)	subsection (3)(a) is complied with; or		
		(ii)	subsections (3)(b) and (4) are complied with; and	15	
	(b)	colle	ect the form; and		
	(c)	send	the completed form to the registrar.		
10	First	opini	on reached		
(1)	This	section	n applies after section 9 is complied with.		
(2)	The	attendi	ng medical practitioner must reach the opinion that—	20	
	(a)	the p	erson is a person who is eligible for assisted dying; or		
	(b)	the p	erson is not a person who is eligible for assisted dying; or		
	(c)	-	person would be a person who is eligible for assisted dying if the on's competence were established as described in section 12 .		
(3)	The	attendi	ng medical practitioner must—	25	
	(a)	comp	plete a prescribed form recording his or her opinion; and		
	(b)	send	the completed form to the registrar.		
11	Seco	nd opi	inion reached		
(1)			section applies if the attending medical practitioner reaches the opinion ribed in section 10(2)(a) or (c) .		
(2)	The	attendi	ng medical practitioner must—		
	(a)		he SCENZ Group for the name and contact details of an independ- nedical practitioner; and		
	(b)		he independent medical practitioner for his or her opinion on whether person is a person who is eligible for assisted dying.	35	
(3)	The	indepe	ndent medical practitioner must—		

	(a)	read	the person's files; and				
	(b)	exan	nine the person; and				
	(c)	reacl	n the opinion that—				
		(i)	the person is a person who is eligible for assisted dying; or				
		(ii)	the person is not a person who is eligible for assisted dying; or	5			
		(iii)	the person would be a person who is eligible for assisted dying if the person's competence were established as described in section 12 .				
(4)	The	The independent medical practitioner must—					
	(a)	comp	plete a prescribed form recording his or her opinion; and	10			
	(b)	send	the completed form to the registrar; and				
	(c)	send	a copy of the completed form to the attending medical practitioner.				
12	Thir	d opin	ion reached, if necessary				
(1)	This	section	n applies if—				
	(a)	the f	ollowing situation exists:	15			
		(i)	the attending medical practitioner reaches the opinion described in section 10(2)(a) ; and				
		(ii)	the independent medical practitioner reaches the opinion described in section 11(3)(c)(iii) ; or				
	(b)	the f	ollowing situation exists:	20			
		(i)	the attending medical practitioner reaches the opinion described in section 10(2)(c) ; and				
		(ii)	the independent medical practitioner reaches the opinion described in section 11(3)(c)(i) ; or				
	(c)	the f	ollowing situation exists:	25			
		(i)	the attending medical practitioner reaches the opinion described in section 10(2)(c) ; and				
		(ii)	the independent medical practitioner reaches the opinion described in section 11(3)(c)(iii) .				
(2)	The	medica	al practitioners must jointly—	30			
	(a)	ask the SCENZ Group for the name and contact details of a specialist; and					
	(b)	ask t	he specialist for his or her opinion on whether the person is compe-				
(3)	The	special	ist must—	35			
	(a)	read	the person's files; and				
	(b)	exan	nine the person; and				

	(c)	reach	the opinion that—			
		(i)	the person is competent; or			
		(ii)	the person is not competent.			
(4)	The	special	ist must—			
	(a)	comp	plete a prescribed form recording his or her opinion; and	5		
	(b)	send	the completed form to the registrar; and			
	(c)	send	a copy of the completed form to—			
		(i)	the attending medical practitioner; and			
		(ii)	the independent medical practitioner.			
13	Nega	ative d	ecision made on request	10		
(1)			n (2) applies if the attending medical practitioner reaches the opined in section 10(2)(b) .			
(2)		attendi o the p	ng medical practitioner must explain the reasons for his or her opinerson.			
(3)	Sub	sectio	n (4) applies if—	15		
	(a)	the independent medical practitioner reaches the opinion described in section 11(3)(c)(ii) ; or				
	(b)	the fo	ollowing situation exists:			
		(i)	a specialist is asked for his or her opinion under section 12(2)(b) ; and	20		
		(ii)	the specialist reaches the opinion described in section 12(3)(c)(ii).			
(4)	tend	the pe	ndent medical practitioner or the specialist, as appropriate, must at- rson with the attending medical practitioner to explain the reasons er opinion to the person.	25		
(5)	The	attendi	ng medical practitioner must—			
	(a)	_	plete a prescribed form recording the actions taken to comply with section (2) or (4); and			
	(b)	send	the completed form to the registrar.			
14	Posi	tive de	cision made on request	30		
(1)	This	section	n applies if—			
	(a)	the fo	ollowing situation exists:			
		(i)	the attending medical practitioner reaches the opinion described in section 10(2)(a) ; and			
		(ii)	the independent medical practitioner reaches the opinion described in section 11(3)(c)(i) ; or	35		

	(b)	the fo	ollowing situation exists:				
		(i)	a specialist is asked for his or her opinion under section 12(2)(b); and				
		(ii)	the specialist reaches the opinion described in section 12(3)(c)(i).	5			
(2)	The	attendi	ng medical practitioner must—				
	(a)	tell thing; a	he person that the person is a person who is eligible for assisted dy- and				
	(b)		ass with the person the progress of the person's terminal illness or yous and irremediable medical condition; and	10			
	(c)	discu	ss with the person the likely timing of the assisted dying; and				
	(d)		e provisional arrangements to be available to administer the medica- at the time indicated.				
15	Med	icatior	n chosen				
(1)	This	section	n applies after section 14 is complied with.	15			
(2)		-	the person wishes to exercise the option of receiving assisted dying, he must tell the attending medical practitioner.				
(3)	The	The attending medical practitioner must—					
	(a)		he person about the following methods for the administration of a l dose of medication:	20			
		(i)	ingestion, triggered by the person:				
		(ii)	intravenous delivery, triggered by the person:				
		(iii)	ingestion through a tube:				
		(iv)	injection; and				
	(b)	ask t	he person to choose one of the methods; and	25			
	(c)		he person to choose the time at which he or she wishes the medica- to be administered; and				
	(d)	ensur any t	re that the person knows that he or she can change his or her mind at ime.				
(4)			hours before the chosen time of administration, the attending med- oner must—	30			
	(a)	write	the appropriate prescription for the person; and				
	(b)	advis	se the registrar of the method and time chosen; and				
	(c)	provi	ide the registrar with the prescription.				
(5)		registra	ar must check that the process in sections 8 to 14 has been com-	35			

(6)		e registrar is satisfied that the process in sections 8 to 14 has been com- l with, the registrar must—					
	(a)	co-sign the prescription for the person; and					
	(b)	provide the co-signed prescription to the attending medical practitioner.					
16	Med	ication administered	5				
(1)	This	section applies after section 15 is complied with.					
(2)		At the chosen time of administration, the attending medical practitioner must ask the person if he or she chooses to receive the medication.					
(3)		e person chooses not to receive the medication, the attending medical practier must—	10				
	(a)	remove the medication from the room; and					
	(b)	return the medication to the pharmacist who dispensed it; and					
	(c)	complete a prescribed form recording the actions taken to comply with paragraphs (a) and (b); and					
	(d)	send the completed form to the registrar.	15				
(4)		e person chooses to receive the medication, the attending medical practi- er must administer it by—					
	(a)	providing it to the person, for the methods described in section 15(3)(a)(i) and (ii) ; or					
	(b)	providing it, for the methods described in section 15(3)(a)(iii) and (iv) .	20				
(5)	The	attending medical practitioner must—					
	(a)	be available to the person until the person dies; or					
	(b)	arrange for another medical practitioner to be available to the person until the person dies.	25				
(6)		he purposes of subsection (5) , the medical practitioner is available to the on if the medical practitioner—					
	(a)	is in the same room as the person; or					
	(b)	is not in same room as the person but is in close proximity to the person.					
17	Deat	th reported	30				
(1)	med regis	Within 14 working days of a person dying as a result of the administration of medication under section 16 , the attending medical practitioner must send the registrar a report in the prescribed form containing the information described in subsection (2) .					
(2)	The	information is—	35				
	(a)	the attending medical practitioner's name; and					
	(b)	the person's name; and					

	(c)	the person's last known address; and				
	(d)	the fact that the person died; and				
	(e)	a description of how the attending medical practitioner complied with section 14(2) ; and				
	(f)	which of the methods described in section 15(3)(a) was used; and	5			
	(g)	a description of the administration of the medication; and				
	(h)	whether any problem arose in the administration of the medication and, if so, how it was dealt with; and				
	(i)	the place where the person died; and				
	(j)	the date and time when the person died; and	10			
	(k)	the name of the medical practitioner who was available to the person until the person died; and				
	(1)	the names of any other health practitioners who were present when the person died.				
(3)	The	registrar must send the report to the review committee.	15			
18	Unu	sed medication returned				
(1)	Subsection (2) or (3) applies if—					
	(a)	a prescription is written under section 15(4)(a); and				
	(b)	the medication is not dispensed before the person for whom the prescription was written dies.	20			
(2)		e attending medical practitioner holds the prescription when the person he or she must—				
	(a)	destroy it; and				
	(b)	complete a prescribed form recording the action taken to comply with paragraph (a) ; and	25			
	(c)	send the completed form to the registrar.				
(3)	If the	e registrar holds the prescription when the person dies, he or she must—				
	(a)	destroy it; and				
	(b)	complete a prescribed form recording the action taken to comply with paragraph (a) .	30			
(4)	Sub	section (5) applies if—				
	(a)	a prescription is written under section 15(4)(a); and				
	(b)	the medication is dispensed but not used before the person for whom the prescription was written dies.				
(5)	The	attending medical practitioner must—	35			
	(a)	return the medication to the pharmacist who dispensed it; and				

(b)	complete a prescribed form recor	ding the action	taken to	comply	with
	paragraph (a); and				

(c) send the completed form to the registrar.

Part 3 Accountability

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19 SCENZ Group

- (1) The Director-General must establish the SCENZ Group by appointing to it the number of medical practitioners that the Director-General considers appropriate.
- (2) The functions of the SCENZ Group are—

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- (a) to make and maintain a list of medical practitioners who are willing to act for the purposes of this Act as—
 - (i) replacement medical practitioners:
 - (ii) independent medical practitioners:
- (b) to provide a name and contact details from the list, when this Act requires the use of a replacement medical practitioner or independent medical practitioner, in such a way as to ensure that the attending medical practitioner does not choose the replacement medical practitioner or independent medical practitioner:
- (c) to make and maintain a list of health practitioners who are willing to act 20 for the purposes of this Act as specialists:
- (d) to provide a name and contact details from the list, when this Act requires the use of a specialist, in such a way as to ensure that neither the attending medical practitioner nor the independent medical practitioner chooses the specialist:

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- (e) to make and maintain a list of pharmacists who are willing to dispense medication for the purposes of **section 16**:
- (f) to provide a name and contact details from the list when **section 16** is to be applied:
- (g) in relation to the administration of medication under **section 16**,— 30
 - (i) to prepare standards of care; and
 - (ii) to advise on the required medical and legal procedures; and
 - (iii) to provide practical assistance, if assistance is requested.
- (3) The ministry must service the SCENZ Group.

20 Review committee

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(1) The minister must appoint an end of life review committee consisting of—

	(a)	a medical ethicist; and					
	(b)	a medical practitioner who practises in the area of end of life care; and					
	(c)	another medical practitioner.					
(2)	The	The review committee has the following functions:					
	(a)	to consider reports sent to it under section 17(3); and	5				
	(b)	to report to the registrar about its satisfaction or otherwise with the cases reported; and					
	(c)	to recommend actions that the registrar may take to follow up cases with which the review committee was not satisfied.					
21	Regi	istrar (assisted dying)	10				
(1)		The Director-General must nominate an employee of the ministry as the registrar (assisted dying).					
(2)	The	registrar must establish and maintain a register recording the following:					
	(a)	prescribed forms held by the registrar; and					
	(b)	the review committee's reports to the registrar; and	15				
	(c)	the registrar's reports to the minister.					
(3)	The	The registrar must consult the Privacy Commissioner—					
	(a)	before establishing the register; and					
	(b)	at regular intervals while maintaining the register.					
(4)		The registrar must establish and maintain a procedure to deal with complaints 2 about breaches of this Act.					
(5)	The registrar must report to the minister by the end of 30 June each year on the following matters for the year:						
	(a)	the total number of deaths occurring under section 16 :					
	(b)	the total broken down into deaths occurring through each of the methods described in section 15(3)(a) :	25				
	(c)	the number of complaints received about breaches of this Act:					
	(d)	how the complaints were dealt with:					
	(e)	any other matter relating to the operation of this Act that the registrar thinks appropriate.	30				
(6)		The registrar must perform any other functions that this Act requires the registrar to perform.					
22	Revi	Review of operation of Act					
(1)	Three years after the commencement of this Act, the ministry must start a review of the operation of this Act and must complete it within 6 months of starting it.						

(2)

(2)		Every 5 years after the date of the last review, the ministry must start another review of the Act and must complete it within 6 months of starting it.				
(3)	Every review must consider whether any amendments to this Act are necessary or desirable.					
(4)	Ever	y review must be the subject of a report to the minister.	5			
(5)						
		Part 4				
		Related matters				
23	Reg	ulations prescribing forms	10			
		Governor-General may, by Order in Council, make regulations prescribing as for the purposes of this Act.				
24	Oth	er rights and duties not affected				
(1)	Noth	ning in this Act affects a person's rights to—				
	(a)	refuse to receive nutrition:	15			
	(b)	refuse to receive hydration:				
	(c)	refuse to receive life-sustaining medical treatment.				
(2)		ring in this Act affects a medical practitioner's duty to alleviate suffering in rdance with standard medical practice.				
25	Effe	ct of death under this Act	20			
		erson who dies as a result of the provision of assisted dying is taken for all oses to have died as if assisted dying had not been provided.				
26	Imn	nunity in civil or criminal proceedings				
	omis	erson is immune from liability in civil or criminal proceedings for acts or sions in good faith and without negligence in providing or intending to ide assisted dying.	25			
27	Offe	nces				
(1)	A pe	erson commits an offence who—				
	(a)	wilfully fails to comply with a requirement in this Act; or				
	(b)	completes or partially completes a prescribed form for a person without the person's consent; or	3(
	(c)	alters or destroys a completed or partially completed prescribed form without the consent of the person who completed or partially completed it.				
(2)	The	person is liable on conviction to either of both of—	35			

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- (a) a term of imprisonment not exceeding 3 months:
- (b) a fine not exceeding \$10,000.

Amendments to Births, Deaths, Marriages, and Relationships Registration (Prescribed Information) Regulations 1995

- (1) This section amends the Births, Deaths, Marriages, and Relationships Registration (Prescribed Information) Regulations 1995.
- (2) Replace regulation 7(1)(a)(xiii) with:
 - (xiii) the cause or causes of the person's death, subject to **subpara-graph (xiiia)**:
 - (xiiia) in respect of a person who died as a result of the provision of assisted dying under the End of Life Choice Act **2017**, the cause or causes of death as if assisted dying had not been provided:
 - (xiiib) in respect of a person who died as a result of the provision of assisted dying under the End of Life Choice Act **2017**, the fact that the person died as a result of the provision of assisted dying under the End of Life Choice Act **2017**:
 - (xiiic) the interval between onset of the cause of death and death, in respect of each cause of death, subject to **subparagraph** (xiiid):
 - (xiiid) in respect of a person who died as a result of the provision of assisted dying under the End of Life Choice Act **2017**, the interval between onset of the cause of death and death by assisted dying, in respect of each cause of death:

Wellington, New Zealand:

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