Thank you for allowing me to testify.

I have also submitted a written statement.

My name is Thaddeus Pope

of the Health Law Institute
and a Professor
at Mitchell Hamline
School of Law.

I specialize in legal and ethical issues in end-of-life medical treatment.

I have published

over 100 articles in this area, including in the New England Journal of Medicine, JAMA,

Chest,

and the New York Times.

The bill you are considering has been extensively tested.

A basically identical law has been in effect in Oregon since 1997 in Washington since 2008 and in Vermont since 2013

It becomes effective in California in June 2016

That is 30 years of combined experience.

There is no better or more relevant track record

on which to evaluate the bill before you.

There are an overwhelming number of safeguards in this bill that control

access to aid in dying medication

The patient must be an adult.

Must be a resident of Minnesota.

The patient must have an incurable and irreversible illness anticipated to cause death within six months

Must have decision making capacity.

This must be confirmed by both an attending and a consulting physician

that the patient is suffering from impaired judgment, the must refer the patient for mental health counseling

The attending physician must fully inform the patient of her alternative options

The bill requires two signed written requests

They must be witnessed

They must be made 15 days apart

They must be made by the patient herself

This cannot be done through an advance directive, health care agent, surrogate, or guardian.

Finally, once the patient obtains the aid in dying medication, she must ingest it herself.

That is why this is "aid" in dying not "euthanasia."

These are the **same** safeguards in Oregon, Washington, Vermont, and California.

Multiple independent studies
have uniformly concluded
that they are effective
and that there has been no abuse.

The health authorities in Oregon and Washington have collected decades of data

(1) Very few patients use the law.

Last year, 132 Oregonians ingested aid in dying prescriptions.

That is less than **one-half of one percent** of the Oregonians who die each year.

(2) The demographics of that narrow population.

show that aid in dying is not being foisted onto minorities or the vulnerable.

Instead,

it is overwhelmingly used by educated, insured, white cancer patients.

99%	Health insurance
97%	White
90%	Age > 65
72%	College educated

And these patients did not use aid in dying As an alternative to hospice

Over 92% used it with hospice.

This is why one-third of patients who get aid in dying prescriptions never ingest the medication

The bill provides **opt-outs**to accommodate physicians
who have an objection

But this does not mean that participation
Will be concentrated
in just a few physicians

The 218 prescriptions written in Oregon in 2015 were written by 106 different physicians.

Opponents point to cases in **Belgium**As evidence of a **slippery slope**.

Some Belgians have obtained aid in dying even though they are **not** terminally ill.

But this argument is misplaced.

Assisted death was already prevalent before the practice was legalized there.

Belgium never "slipped" from being less permissive to being more permissive of aid in dying.

Belgium was always more permissive.

There is no slipping in the United States either

No U.S. state has ever enacted legislation with different, fewer, or weaker safeguards.

There is no evidence

that physicians have failed to comply with safeguards.

not a single criminal case.

not a single health licensing board action.

Neither has Disability Rights Oregon, the state's Protection and Advocacy System, received any complaint of exploitation or coercion of any individual with disabilities

More than two decades ago,
the Minnesota Legislature
confirmed the right
of Minnesotans
to refuse life-sustaining treatment.

chronically and critically ill patients across the state

by withholding or withdrawing dialysis, mechanical ventilation, and other interventions.

But some terminally ill patients are not dependent on any such technologies.

This bill gives these patients

The same freedom

To control the manner of their death

Thank you