

Testimony of Diane Coleman, JD, President and CEO of Not Dead Yet Opposing Maryland HB 933/SB 845 End-of-Life Option Act March 7, 2023



I am submitting testimony as a person with an advanced neuromuscular condition. I'm also the founder and President of Not Dead Yet, a national disability rights group that opposes legalization of assisted suicide. I have personal experience with the uncertainty of terminal predictions by doctors, as well as the risks of assisted suicide laws for people who use breathing support or otherwise depend on ongoing treatments for serious chronic conditions. I will also discuss the data from Oregon, which substantiates our concerns.

My Personal Story of Mistaken Prognosis

At the age of six I was diagnosed as having muscular dystrophy and my parents were told that I would die by the age of 12. A few years later I was re-diagnosed with spinal muscular atrophy (SMA), a progressive neuromuscular condition which has a longer lifespan. A few years ago, new DNA tests resulted in another revised diagnosis, another neuromuscular condition. Regardless of the exact diagnosis, since age eleven, I have used a motorized wheelchair.

I am now age 68. About 20 years ago I started using breathing support at night. Over the years, the pressures required to sustain my breathing have increased. About seven years ago, I went into respiratory failure and I now use what is called a non-invasive ventilator over 22 hours a day. If I did not use this support, I would go into respiratory failure and die in a fairly short time.

Throughout my adult life, I have worked full time, first as an attorney and then directing nonprofit disability related organizations. Since 2012, working from home, I have continued to run Not Dead Yet, which has four staff, a few contractors and numerous volunteers across the country. I have spoken at conferences, guest lectured at universities, published articles, submitted testimony in legislatures, and provided the day-to-day management an organization requires. I'm not saying this

to be "inspirational", but to make it clear that people like me can and do work, and/or do lots of other worthwhile and fun things and should not be written off.

How Assisted Suicide Laws Endanger People Like Me

As a severely disabled person who depends on life-sustaining treatment, I would be able to qualify for assisted suicide at any time if I lived where assisted suicide is legal. As an Oregon official has clarified in writing, any person who becomes terminal because they do not receive treatment, for any reason, would qualify for assisted suicide under an Oregon type law. If the reason that I could not get treatment were an inability to afford insurance co-pays, I would be eligible for assisted suicide. For over a year, my breathing support had a \$500 per month co-pay which I had to pay out of pocket till my employer changed health plans.

If I became despondent, for example if I lost my husband or my job, and decided that I wanted to die, I would not be given the same suicide prevention as a nondisabled and healthy person who despaired over divorce or job loss. Where assisted suicide is legal, I would be treated completely differently due to my condition. By denying equal suicide prevention and other supports to people deemed "terminal", assisted suicide laws are inherently discriminatory against older, ill and disabled people. The discrimination of state licensed health professionals denying equal suicide prevention and instead providing suicide assistance is a fundamental violation of the Americans with Disabilities Act.

In these days of private managed care companies taking over Medicare and Medicaid, where providing expensive care has a tendency to reduce profits, we should at least question whether there is an inherent conflict of interest in having healthcare providers administrate a state sanctioned assisted suicide program.

Anyone could ask for assisted suicide, but doctors are the gatekeepers with the power to decide who's eligible. Since early 2020, the COVID pandemic has revealed that people with disabilities have been denied treatment for the virus due to their disabilities and pervasive healthcare provider biases about our "quality of life." According to a Harvard researcher, "In our survey of 714 practicing US physicians nationwide, 82.4 percent reported that people with significant disability have worse quality of life than nondisabled people. Only 40.7 percent of physicians were very confident about their ability to provide the same quality of care to patients with disability"² This is only the most recent academic confirmation of the real-life experiences shared by many disabled people. This is among the many factors that lead the National Council on Disability in 2019 to issue formal report entitled "The Danger of Assisted Suicide Laws."³

What I've Learned From the Oregon Assisted Suicide Data

Oregon's assisted suicide law, the oldest in the U.S., is held up as the model for other states. I've examined the Oregon state assisted suicide reports from a disability rights perspective and want to share with you what I've learned.

One of the most frequently repeated claims by proponents of assisted suicide laws is that there is "no evidence or data" to support any claim that these laws are subject to abuse, and that there has not been "a single documented case of abuse or misuse" in the 24 reported years. These claims are demonstrably false.

Regarding documented cases, please refer to a description of individual cases and source materials compiled by the Disability Rights Education and Defense Fund entitled Oregon and Washington State Abuses and Complications. For an in-depth analysis of several cases by Drs. Herbert Hendin and Kathleen Foley, please read Physician-Assisted Suicide in Oregon: A Medical Perspective. 5

The focus of the discussion below is the <u>Oregon Public Health Division data</u>.⁶ These reports are based on forms filed with the state by the physicians who prescribe lethal doses and the pharmacies that dispense the drugs. As the early state reports admitted:

"As best we could determine, all participating physicians complied with the provisions of the Act. . . . <u>Under reporting and noncompliance is thus difficult to assess because of possible repercussions for noncompliant physicians reporting to the division</u>." (Emphasis added.)

Further emphasizing the serious limits on state oversight under the assisted suicide law, Oregon authorities also issued a release in 2005 clarifying that they have No authority to investigate Death with Dignity case.

Nevertheless, despite these apparently intended limitations in data gathering, the Oregon state reports substantiate some of the problems and concerns raised by opponents of assisted suicide bills, including the Maryland bill.

Non-Terminal Disabled Individuals Are Receiving Lethal Prescriptions In Oregon

The Oregon Public Health Division assisted suicide reports show that nonterminal people received lethal prescriptions every year except the first.

The prescribing physicians' reports to the state include the time between the request for assisted suicide and death for each person. Prior to the 2020 report, the online state reports did not reveal how *many* people outlived the 6-month or 180-day terminal prognosis. Instead, the reports gave that year's median and range of the number of days between the request for a lethal prescription and

death. This was on page 13 of the <u>2019 annual report</u>. In 2019, at least one person lived 1503 days, the longest reported duration between the request for assisted suicide and death so far. In every year except the first year, the reported upper range has been significantly longer than 180 days.

The 2020 report revisits this survival issue and states that 4 percent of individuals outlived their 6-month prognosis⁹ and 3.8% in 2021.¹⁰ This does not take into account the individuals who took the drugs quickly but may have survived if they had waited longer.

The definition of "terminal" in the Oregon statute only requires that the doctor (or in some bills non-physician providers) predict that the person will die within six months. There is no requirement that the doctor consider the likely impact of medical treatment in terms of survival. Unfortunately, while short-term terminal predictions of some conditions, such as some cancers, are fairly well established, this is far less true six months out, as the bill provides, rather than one or two months before death, and is even less true for other diseases.

In addition, it should be noted that the attending physician who determines terminal status and prescribes lethal drugs is not required to be an expert in the disease condition involved, nor is there any information about physician specialties in the state reports.

Furthermore, as noted above, many conditions will or may become terminal if certain medications or routine treatments are discontinued – e.g. insulin, blood thinners, pacemaker, CPAP, etc. Any person who becomes terminal because they do not receive treatment for any reason, such as lack of insurance coverage, or costly deductibles and co-pays, or government budget cuts in Medicare and Medicaid, would qualify for assisted suicide under an Oregon type law.

Oregon reports that non-cancer conditions found eligible for assisted suicide have grown over the years to include: neurological disease, respiratory disease, heart/circulatory disease, infectious disease, gastrointestinal disease, "endocrine/metabolic disease (e.g. diabetes)", infectious disease and, in the category labeled "other", anorexia, arthritis, arteritis, blood disease, hernia, complications from a fall, kidney failure, medical care complications, musculoskeletal system disorders, sclerosis, and stenosis (2021 Report, pages 12 & 14). The addition of **anorexia** to Oregon's and Colorado's reports as a condition some physicians have viewed as warranting assisted suicide demonstrates how empty the law's purported "safeguards" are.

The Only Certifiers of Non-Coercion And Capability Need Not Know the Person

Four people are required to certify that the person is not being coerced to sign the assisted suicide request form, and appears capable: the prescribing doctor, second-opinion doctor, and two witnesses.

In most cases over the years, the prescribing doctor is a doctor referred by assisted suicide proponent organizations. (See, M. Golden, Why Assisted Suicide Must Not Be Legalized, 11 section on "Doctor Shopping" and related citations). The Oregon state reports say that the median duration of the physician patient relationship was 5 weeks in 2021, and 11 weeks over all years (2021 Report, page 13). Thus, lack of coercion is not usually determined by a physician with a longstanding relationship with the patient. This is significant in light of well-documented elder abuse-identification and reporting problems among professionals in a society where an estimated one in ten elders is abused, mostly by family and caregivers. (Lachs, et al., New England Journal of Medicine, Elder Abuse. 12)

The witnesses on the Oregon <u>request form¹³</u> need not know the person either. In Oregon, only one of them may be an heir, but neither of the witnesses need to actually know the person (the Oregon forms say that if the person is not known to the witness, then the witness can confirm identity by checking the person's ID).

In addition, the definition of "capacity to make medical decisions" in the Maryland bill allows third parties to communicate orally for the person to the doctors and witnesses, providing for the patient to be "communicating through a person familiar with a patient's manner of communicating." This is especially dangerous for people with speech impairments, such as from a stroke, brain injury or other neurological disability.

So neither doctors nor witnesses need know the person well enough to certify that they are not being coerced. What might be an example of coercion? Many elder and disabled people would say the threat of being put in a nursing home.

No Evidence of Consent or Self-Administration At Time of Death

In less than half the reported cases, the Oregon Public Health Division reports also state that no health care provider was present at the time of ingestion of the lethal drugs or that it is "unknown" whether a provider was present (2021 Report, page 13). Without requiring an independent witness, there is no way to confirm whether the lethal dose was self-administered and consensual. The initial request for assisted suicide does not prove that the person wanted to go through with it, as shown by the reported decision by many not to do so.

Therefore, although "self-administration" is touted as one of the key "safeguards", in about half the cases, there is no evidence of consent or self-administration at the time of ingestion of the lethal drugs. If the drugs were, in some cases, administered by others without consent, no one would know. The request form

constitutes a virtual blanket of legal immunity covering all participants in the process based on a mere claim of "good faith."

Essentially, proponents of legalizing assisted suicide have taken what is fundamentally a third-party legal immunity statute and marketed it deceptively as a personal rights statute. This deception relies, in part, on the common misconception that suicide is illegal, which it is not.

Pain Is Not the Issue, Unaddressed Disability Concerns Are

The top five reasons doctors give for their patients' assisted suicide requests over all reported years are not pain or fear of future pain, but psycho-social issues that are well understood by the disability community: "loss of autonomy" (91%), "less able to engage in activities" (90%), "loss of dignity" (73%), "burden on others" (48%) and "losing control of bodily functions" (44%) (2021 Report, page 13).

These reasons for requesting assisted suicide pertain to disability and indicate that over 90% of the reported individuals, possibly as many as 100%, are disabled at the time of their assisted suicide request.

Three of these reasons (loss of autonomy, loss of dignity, feelings of being a burden) could be addressed by consumer-directed in-home long-term care services, but no disclosures about or provision of such services is required. Some of the reported reasons are clearly psycho-social and could be addressed by disability-competent professional and peer counselors, but this is not required either. Moreover, less than 4% of patients who request assisted suicide were referred for a psychiatric or psychological evaluation (and only 2 patients in 2021), despite studies showing the prevalence of depression in such patients. Even if a bill requires such a referral, it's only purpose is to determine whether the person has "impaired judgment."

Basically, the law operates as though the person's reasons don't matter, and nothing need be done to address them.

Conclusion

The Oregon assisted suicide data demonstrates that people who were not actually terminal received lethal prescriptions in all 24 reported years except the first, and that there is little or no substantive protection against coercion and abuse. Examples of abuses have surfaced through mainstream media and professional journals, but not through the superficial and unmonitored physician self-reporting system used by the state. Moreover, reasons for requesting assisted suicide that sound like a "cry for help" with disability-related concerns are apparently ignored. Thus, even the paltry data collected by the state substantiates serious problems with the implementation of assisted suicide laws.

For all these reasons, I urge you to vote no on the Maryland assisted suicide bill HB 933/SB845. The dangers of mistakes, coercion and abuse are simply too high.

https://drive.google.com/file/d/1xOZfLFrvuQcazZfFudEncpzp2b18NrUo/view

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https://www.oregon.gov/oha/PH/PROVIDERPARTNERRESOURCES/EVALUATION RESEARCH/DEATHWITHDIGNITYACT/Documents/year23.pdf (page 11)

https://www.oregon.gov/oha/PH/PROVIDERPARTNERRESOURCES/EVALUATION RESEARCH/DEATHWITHDIGNITYACT/Documents/year24.pdf (page 5)

¹ https://www.washingtontimes.com/news/2018/jan/11/diabetics-eligible-physician-assisted-suicide-oreg/;

² Lisa I. lezzoni, et al., Physicians' Perceptions Of People With Disability And Their Health Care, https://www.healthaffairs.org/doi/10.1377/hlthaff.2020.01452

³ https://ncd.gov/sites/default/files/NCD Assisted Suicide Report 508.pdf

⁴ https://dredf.org/wp-content/uploads/2015/04/Revised-OR-WA-Abuses.pdf

⁵ https://dredf.org/wp-content/uploads/2012/08/Hendin-Foley-Michigan-Law-Review.pdf

⁶https://www.oregon.gov/oha/PH/PROVIDERPARTNERRESOURCES/EVALUATION RESEARCH/DEATHWITHDIGNITYACT/Pages/ar-index.aspx

⁷ https://dredf.org/wp-content/uploads/2012/08/Oregon-DHS.pdf

⁸https://www.oregon.gov/oha/PH/PROVIDERPARTNERRESOURCES/EVALUATION RESEARCH/DEATHWITHDIGNITYACT/Documents/year22.pdf

¹¹ https://dredf.org/public-policy/assisted-suicide/why-assisted-suicide-must-not-be-legalized/

http://www.nejm.org/doi/full/10.1056/NEJMra1404688; See also, "New Report on Elder Abuse in Australia: Implications for Legalising Euthanasia" (Jan. 2022) https://www.australiancarealliance.org.au/new_report_on_elder_abuse_in_australia_implications_for_legalising_euthanasia_

¹³http://www.oregon.gov/oha/PH/PROVIDERPARTNERRESOURCES/EVALUAT IONRESEARCH/DEATHWITHDIGNITYACT/Documents/pt-req.pdf