The Olmstead Decision Has Been Misinterpreted

“The Olmstead decision, which interprets the Americans with Disabilities Act, is so clear that persistent misinterpretation by federal agencies can only be described as purposeful.”

Introduction

VOR is a national nonprofit advocacy organization that has for 32 years supported the right of individuals with intellectual and developmental disabilities (I/DD) to receive services and supports according to their individual choice and need, regardless of setting.

We take this opportunity – the 25th Anniversary of the ADA - to set the record straight about what the landmark Olmstead decision actually says about the ADA’s “integration regulation.”

July 26, 2015 marks the 25th Anniversary of the Americans with Disabilities Act (ADA). Olmstead v. L.C., 527 U.S. 581 (1999) is the landmark Supreme Court decision interpreting the ADA’s “integration regulation,” which states:

“a public entity must administer services, programs, and activities in the most integrated, least restrictive setting appropriate to the needs of qualified individuals with disabilities.” [28 C.F.R. § 35.130(d)].

The Olmstead Court considered whether the ADA’s prohibition of discrimination by a public entity required “placement of persons with mental disabilities in community settings rather than in institutions.” (Olmstead at 587)

The Court’s answer: A “qualified yes.” (Id., emphasis added)

Since 1999, however, federal agencies and some advocacy organizations have lost sight of the individual when considering the rights established by the Olmstead decision. A thoughtful decision which balanced individual need, choice, and the state’s resources against a statute that also required consideration of individual needs, has been discarded for an interpretation that the ADA, as interpreted by Olmstead, allows only community placement.

In sharp contrast, both the Olmstead decision and the ADA eschew absolutes. Neither support only community care or only institutional care:

[O]ne conclusion seems plain: some disabled individuals can benefit from community placement, and some may not. While all disabled are covered by the ADA, different remedies are recognized by the statute for different degrees of disability.” [Brief of VOR et al., as Amici Curiae, in Olmstead v. L.C., at 6 (February 3, 1999)]

The *Olmstead* decision and the law are so clear that persistent misinterpretation by federal agencies can only be described as purposeful.

**The History of the *Olmstead* Decision: Lois Curtis, Elaine Wilson, and VOR**

In 1995, two women, Lois Curtis and Elaine Wilson, courageously pursued their right under the ADA and its integration regulation to receive person-centered services and supports in the most integrated setting appropriate to their individualized needs. Their lawsuit against the State of Georgia, *Olmstead v. L.C.*, made it all the way to the Supreme Court, and in 1999, culminated into what is now recognized as the landmark decision interpreting the ADA’s Title II “integration regulation.”

VOR and 141 of its affiliates and supporting organizations stood by Lois and Elaine, filing an *Amicus Curiae* brief in support of their right to receive community-based supports, while also arguing that the ADA does not mandate a singular placement option – the community – for all citizens with disabilities. Instead, VOR argued that the ADA’s protections extended to individuals with “degrees of disability” who are entitled to receive services, supports and accommodations according to what is most “appropriate to the needs of the individual,” which may sometimes be an institution.

Quoting VOR’s brief, the Court agreed:

“Each disabled person is entitled to treatment in the most integrated setting possible for that person — recognizing on a case-by-case basis, that setting may be an institution.” *Olmstead v. L.C.*, 527 U.S. 581, 605 (1999) (*quoting* Brief of VOR et al., as *Amici Curiae* at 11).

The Court was clear that the ADA’s integration regulation does not mandate that people with disabilities be forced from facility-based care (“institutions”) when doing so was not appropriate to individual needs or contrary to individual choice.

The Court found that Lois and Elaine had suffered discrimination under the ADA, pointing to Lois and Elaine’s choices, their treating professionals opinion that for them community placement was appropriate factored directly into the Court’s holding, and the fact that the State had agreed to provide such services. Accordingly, the Court held community placement is only required (not always required), and institutionalization is only “unjustified,” when -

“[a] the State’s treatment professionals have determined that community placement is appropriate, [b] the transfer from institutional care to a less restrictive setting is not opposed by the affected individual, and [c] the placement can be reasonably accommodated, taking into account the resources available to the State and the needs of others with mental disabilities. *Olmstead*, 527 U.S. at 587 (*emphasis added*).

The Court expressly recognized an ongoing need for facility-based care (“institutional” care) for some individuals with disabilities, stating –

“We emphasize that nothing in the ADA or its implementing regulations condones termination of institutional settings for persons unable to handle or benefit from community settings...Nor is there any federal requirement that community-based treatment be imposed on patients who do not desire it.” Id. at 601-602.
Purposeful Misinterpretation

*Olmstead* is often incorrectly referred to as a community-only / deinstitutionalization mandate by disability advocates. Federal agencies, such as the Department of Justice, the Administration on Intellectual and Developmental Disabilities, the Centers for Medicare & Medicaid Services, and the National Council on Disability perpetuate this misinterpretation of the *Olmstead* decision and pursue forced deinstitutionalization.

The National Council on Disability, an agency within the U.S. Department of Health and Human Services, has published “Deinstitutionalization: Unfinished business” which encourages the closure and displacement of individuals with profound disabilities from homes of four or more residents, without regard for individual choice and need and contrary to *Olmstead*.

The Administration on Intellectual and Developmental Disabilities, within HHS, has likewise denigrated a proven system of facility-based care through lobbying, class action lawsuits and other tactics in support of forced deinstitutionalization, aimed at evicted the most vulnerable people I/DD from their facility homes, under the guise of “integration” for all, contrary to *Olmstead*.

The Centers for Medicare and Medicaid Services finalized a new regulation that defines “community” settings for the purpose of receiving Medicaid Home and Community-Based Services (HCBS) funding. Complying with the rule is proving costly for the states, resulting in states cutting services and adding to existing gaps left either by historic deinstitutionalization or the closing of congregate settings to comply with the new Rule. As a result, the whole system of care is being undermined in the name of inclusion and integration, contrary to *Olmstead*.

The U.S. Department of Justice’s Civil Rights Division is charged with enforcing “federal statutes prohibiting discrimination on the basis of race, color, sex, disability, religion, familial status and national origin,” including the ADA. Its technical assistance website on *Olmstead* is titled:

*“Olmstead: Community Integration for Everyone”* ²

These are not mere words. The Department of Justice’s (DOJ’s) *Olmstead* enforcement actions bear out an interpretation of *Olmstead* that mandates “community integration for everyone” by pursuing forced, without any regard to the choice or needs of affected individuals, contrary to *Olmstead*. Consider U.S. v. Arkansas³ in which Federal Judge J. Leon Holmes in the first paragraph of his decision dismissing DOJ’s claims wrote:

Most lawsuits are brought by persons who believe that their rights have been violated. Not this one. The Civil Rights Division of the Department of Justice brings this action on behalf of the United States of America against the State of Arkansas and four state officials in their official capacities alleging that practices at Conway Human Development Center violate the rights of its residents guaranteed by the Fourteenth Amendment, the Americans with Disabilities Act, and the Individuals with Disabilities Education Act. All or nearly all of those residents have parents or guardians who have the power to assert the legal rights of their children or wards. Those parents and guardians, so far as the record shows, oppose the claims of the United States. Thus, the United States is in the odd position of asserting that certain persons’ rights have been and are being violated while those persons—through their parents and guardians—disagree. U.S. v. Arkansas, 4:09-cv-33 ( 2011, E.D. Ark.) (emphasis added).

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Over 150 media reports in more than 30 states in the past two decades reveal systemic concerns in small settings for people with intellectual and developmental disabilities (I/DD), including deaths, abuse, neglect, isolation, and financial malfeasance.⁴

Across the country, thousands of fragile citizens have needlessly died in community settings as advocates aggressively pursued community integration without regard to the preparedness of these settings to receive fragile individuals, even while proponents began to acknowledge the predictable problems. As noted Samuel Bagenstos, former Principal Deputy Assistant Attorney General in DOJ’s Civil Rights Division and a key litigator in deinstitutionalization cases –

“It should not be surprising that the coalition of deinstitutionalization advocates and fiscal conservatives largely achieved their goal of closing and downsizing institutions and that deinstitutionalization advocates were less successful in achieving their goal of developing community services.”[“The Past and Future of Deinstitutionalization Litigation,” 34 Cardoza L. Rev. 1, 21 (2012)].

More recently dramatic increases in mortality of individuals with I/DD in community settings has been widely publicized in Georgia and Virginia, two states which are in the process of implementing DOJ Olmstead settlements. In Georgia, there have been nearly 500 deaths in both 2013 and 2014 of individuals with I/DD in community settings, including 62 individuals who transferred from facility-based care per settlement requirements. In Virginia, the rate of mortality of those individuals with I/DD who transferred from facility care per settlement terms is double that of facility-based individuals.

**Conclusion**

Like the integration regulation itself, the Supreme Court’s decision requires consideration of an individual’s needs, while also holding that individual choice also matters.

However, don’t take our word for it. Read the Olmstead decision.⁵

There are tragic outcomes that lie in the wake of forced deinstitutionalization, which since 1999 has been improperly pursued in Olmstead’s name.

Lost in pursuit of the “integration” ideal is concern for the individual. Person-centered planning, which is held up as the ideal by the government, advocates, and nonprofit organizations alike, is short-changed by system change advocacy to eliminate specialized care options for those who need it, which is exactly what the Olmstead court cautioned against. Instead, Olmstead and the ADA reinforce the right of choice, requiring a level of integration that meets the needs of the individual. This right of choice is preserved only if we maintain a full range of options to meet the full range of needs for a diverse population.

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