## 1

#### Abled subjectivity is tied up in a two-tiered affective response that explains disabled life – primary pity which reflects disability upon the ego threatening its ability status, which invokes secondary pity to overcorrect for the shattered-ego necessitating disabled death.

Mollow 15 The Disability Drive by Anna Mollow A dissertation submitted in partial satisfaction of the requirements for the degree of Doctor of Philosophy in English in the Graduate Division of the University of California, Berkeley Committee in charge: Professor Kent Puckett, Chair Professor Celeste G. Langan Professor Melinda Y. Chen Spring 2015 //ACCS JM

Much as the differentiation between the inseparable processes of primary and secondary narcissism rests on a distinction between building up and breaking down the ego, a similar heuristic distinction gives structure to my concepts of primary and secondary pity. To be clear, pity and narcissism are not the same thing: if narcissism can be understood as love of the self, pity involves a complex affective reaction to the suffering of someone else. Primary pity entails a response to the image of another person succumbing to what I have termed the “tragedy of disability.”121 Primary pity arises when one witnesses a fall of the self, a collapse of the ego; such falling is at once painful and pleasurable to observe. In other words, primary pity could be described as a vicarious experience of the tragedy of disability. A great deal of the pain and pleasure of primary pity center on questions about what, or who, this fallen self is. When most people think about pity, we refer to an affect in which, to adopt Edelman’s phrase, we purport to “feel for the other.” But as with primary narcissism, in which the self has not yet been constituted, and therefore cannot be said to enter into intersubjective relations with an “other,” primary pity entails a mixing up of self and other such that the ego, in becoming permeable to pain that may properly belong to “someone else,” is profoundly threatened in its integrity. Primary pity is that intense pain-pleasure complex that is provoked by the image of a suffering other who, it seems momentarily, both is and is not one’s self. This affective response can feel unbearable, as seen in Siebers’s formulation: one “cannot bear to look...but also cannot bear not to look.” Primary pity is difficult to bear because it involves a drive toward disability (one cannot bear not to look), which menaces the ego’s investments in health, pleasure, and control—because to contemplate another person’s suffering is to confront the question, “Could this happen to me?” Such a prospect, although frightening, may also be compelling; in this way, primary pity replicates the self-rupturing aspects of sexuality. Indeed, the unbearability of primary pity reflects its coextensiveness with sexuality. Sex, or the Unbearable, a book coauthored by Edelman and by Lauren Berlant, argues that sex “unleashes unbearable contradictions that we nonetheless struggle to bear” (back cover). This claim accords with Freud’s account of sexuality as a “pleasurable” “unpleasure” that the ego can never fully master or control (Three 49,75). As Leo Bersani puts it in his reading of Freud, “the pleasurable unpleasurable tension of sexual enjoyment occurs when the body’s ‘normal’ range of sensation is exceeded, and when the organization of the self is momentarily disturbed”; thus, “sexuality would be that which is intolerable to the structured self” (Freudian 38). Primary pity is also intolerable to the structured self, because it entails a fascination with the fantasy of a self in a state of disintegration or disablement. Secondary pity is something else, although it cannot wholly be differentiated from primary pity. Secondary pity attempts to heal primary pity’s self-rupturing effects by converting primary pity into a feeling that is bearable. As with secondary narcissism, secondary pity involves both an attempt to get back to that ego-shattering state of painfully pleasurable primary pity, and at the same time to defend against that threat to the ego by aggrandizing oneself at someone else’s expense. Secondary pity refers to all those ego-bolstering behaviors that most people think of when they talk about pity. Disabled people are all too familiar with these behaviors: the saccharin sympathy, the telethon rituals of “conspicuous contribution,” the insistence that “they” (i.e., nondisabled people) could never endure such suffering. More commonly known in our culture simply as “pity,” secondary pity encompasses our culture’s most clichéd reactions to disability: charity, tears, and calls for a cure. Correlatives of these commonplace manifestations of secondary pity are the obligatory claims that disabled people’s suffering is “inspiring.” Indeed, the speed with which conventional cultural representations of disability segue from overt expressions of pity to celebrations of “the triumph of the human spirit” highlights the ways in which secondary pity, as a defense against primary pity’s incursions, reinforces the ego’s fantasy of sovereignty. Secondary pity, in other words, can be seen as a variation of secondary narcissism: these affects enlarge the ego of the pitier or the narcissist at the expense of someone else. But primary pity is not the same as either primary narcissism, secondary narcissism, or secondary pity. Unlike primary narcissism, a feeling that emerges out of a relation to the world in which notions of “self” and “other” do not obtain, primary pity does depend upon the constructs of self and other, although these constructions are unstable and are continually threatening to come undone. Primary pity can thus be envisioned as a threshold category occupying a liminal position between the total denial of the other that is inherent to primary narcissism and the rigid structure of (superior) self and (inferior) other that constitutes secondary narcissism and secondary pity. My concept of primary versus secondary pity also differs from Freud’s primary- secondary narcissism distinction at the level of genealogy. Like Freud’s account of primary and secondary narcissisms, my model of primary and secondary pities involves a temporal transition; but whereas Freud imagines the movement from primary to secondary narcissism as a passage from an earlier to a later stage of an individual’s development, the temporal shift from primary to secondary pity happens much more quickly than this. It happens in an instant: that moment in which we feel primary pity and then, almost before we can blink, deny that we feel or have felt it. The denial is understandable: who wants to admit that one gets pleasure from the sight of another person’s suffering—or, to make matters worse, that this pleasure derives in part from the specter of disability’s transferability, the possibility that this suffering could be—and, fantasmatically, perhaps already is—an image of one’s own self undone?

#### The 1AC’s belief of a better future becomes complicit in the logic of rehabilitative futurism, which is threatened by the Disabled Child.

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“Let us begin our reexamination of Tiny Tim with a discussion of No Future, a text in which Tiny Tim takes a prominent position. No Future is a text with a target: the book takes aim at “the Child whose innocence solicits our defense,” a trope that Edelman names as the emblem of an ideology that he terms “reproductive futurism” (2). According to Edelman, commonplace cultural invocations of the figure of the Child (“not to be confused with the lived experiences of any historical children”) uphold “the absolute privilege of heteronormativity” (11, 2). Defying pronatalist social imperatives, Edelman names queerness as “the side of those not fighting for the children‟” (3) and urges queers to accept the culture’s projection of the death drive onto us by saying explicitly what Law and the Pope and the whole of the Symbolic order for which they stand hear anyway in each and every expression or manifestation of queer sexuality: Fuck the social order and the Child in whose name we’re collectively terrorized; fuck Annie; fuck the waif from Les Mis; fuck the poor, innocent kid on the Net; fuck Laws both with capital ls and with small; fuck the whole network of Symbolic relations and the future that serves as its prop. (No Future 29) Elsewhere, I have argued that No Future’s impassioned polemic is one that disability studies might take to heart. Indeed, the figure that Edelman calls “the disciplinary image of the ‘innocent’ Child” is inextricable not only from queerness but also from disability (19). For example, the Child is the centerpiece of the telethon, a ritual display of pity that demeans disabled people. When Jerry Lewis counters disability activists’ objections to his assertion that a disabled person is “half a person,” he insists that he is only fighting for the Children: “Please, I’m begging for survival. I want my kids alive,” he implores (in Johnson, Too Late 53, 58). If the Child makes an excellent alibi for ableism, perhaps this is because, as Edelman points out, the idea of not fighting for this figure is unthinkable. Thus, when Harriet McBryde Johnson hands out leaflets protesting the Muscular Dystrophy Association, a confused passerby cannot make sense of what her protest is about. “You’re against Jerry Lewis!” he exclaims (61). The passerby’s surprise is likely informed by a logic similar to that which, in Edelman’s analysis, undergirds the use of the word “choice” by advocates of legal abortion: “Who would, after all, come out for abortion or stand against reproduction, against futurity, and so against life?” (16). Similarly, why would anyone come out for disability, and so against the Child who, without a cure, might never walk, might never lead a normal life, might not even have a future at all? The logic of the telethon, in other words, relies on an ideology that might be defined as “rehabilitative futurism,” a term that I coin to overlap and intersect with Edelman’s notion of “reproductive futurism.” If, as Edelman maintains, the future is envisaged in terms of a fantasmatic “Child,” then the survival of this future-figured-as-Child is threatened by both queerness and disability. Futurity is habitually imagined in terms that fantasize the eradication of disability: a recovery of a “crippled” or “hobbled” economy, a cure for society’s ills, an end to suffering and disease. Eugenic ideologies are also grounded in both reproductive and rehabilitative futurism: procreation by the fit and elimination of the disabled, eugenicists promised, would bring forth a better future.” (68-69)

#### The alternative is to disable the figure of the “human” – instead of seeing disability as a redeemable position within civil society, the alternative weaponizes disability’s structural position against the human. If we win their starting point is ableist they cannot weigh the consequences of it.

Mollow 12 [Mollow, Anna. "Is Sex Disability? Queer Theory and the Disability Drive." Sex and Disability, by R. McRuer and A. Mollow, Durham, Duke UP, 2012, pp. 306-10. http://www.sfu.ca/~baw2/GSWS826/Mollow.pdf] //ACCS JM

But here a problem emerges: as we have seen, the writing by Bersani and Edelman that I have examined forwards powerful arguments against the project of becoming human. Urging queers to embrace the “inhumanity of the sinthomosexual,” Edelman observes that the liberal goal of expanding the category of “human” to encompass those presently excluded from it will not “stop the cultural production of figures” made to embody the inhumanity of the death drive (No Future 107). What would it mean for disability theory to embrace disabled people’s figuration as inhuman? As we contemplate this possibility, a moment from How I Became a Human Being may give us pause. After a presentation by the physicist Stephen Hawking at the U.C. Berkeley campus, O’Brien posed the following question: “Doctor Hawking, what can you say to all the disabled people who are stuck in nursing homes or living with their parents or in some other untenable situation and who feel that their life is over, that they have no future?” (Human Being 230). A response that might be derived from Edelman’s book—that there is, and can be, no future, since the future, by definition, can only ever be a fantasy (“always / A day / Away,” in Annie’s paean to “Tomorrow”)—hardly seems more adequate than Hawking’s reply: “All I can say is that one must do the best one can in the situation in which one finds oneself ” (No Future 30; Human Being 231). In light of O’Brien’s question, Edelman’s embrace of the death drive, or Bersani’s celebration of what he calls “the breakdown of the human itself in sexual intensities,” can easily appear as irresponsible theoretical indulgences (“Rectum” 29). Indeed, the word “irresponsible” is one that Bersani himself uses when he reflects, at a distance of thirteen years, on “Is the Rectum a Grave?”: “Much of this now seems to me a rather facile, even irresponsible celebration of ‘self- defeat.’ Masochism is not a viable alternative to mastery, either practically or theoretically” (“Sociality” 110). This remark highlights important shifts and ambivalences in Bersani’s thinking over the course of his career, which may serve as an entry into the question of the status of the human in disability theory. Bersani and Edelman are often cited, as if in the same breath, as proponents of an “antisocial” or “antirelational” “thesis” in queer theory, in opposition to which some critics of their work, such as Muñoz, have defined their own projects as “utopian.”17 But Bersani’s work, rather than conforming to either side of a utopian/antirelational binary, often reveals an interest in thinking in both of these ways at once. For example, writing of passages in his book, Homos (published in 1995) that are frequently cited as the origin of the “antirelational thesis,” Bersani describes the “performance of antirelationality” that he celebrates in Jean Genet’s Funeral Rites as a “utopic form of revolt” (“Sociality” 103; emphasis added). This joining of the utopian and the antirelational corresponds to what Bersani describes, in an essay published in 2004, as a central concern throughout his career: “a dialogue (both conciliatory and antagonistic) between” Foucault and Freud (“Fr- oucault” 133). In this essay and other recent writings, Bersani moves away from the “Freudian” and toward the “Foucauldian.” Worrying that the psychoanalytic (or antirelational) side of this paradigm may be politically irresponsible (insofar as its insistence on the intractability of the death drive seems “resistant to any social transformations whatsoever”), Bersani has become increasingly interested in the creation of what, invoking a phrase of Foucault’s, he calls “new relational modes” (“Fr- oucault” 134). Interestingly, this “admittedly utopic” project often employs a rhetoric of futurism, both reproductive and rehabilitative (Bersani, “Fr- oucault” 134). For example, in a reading of Plato’s Symposium, Bersani approvingly observes that “the goal of a love relation with Socrates” is “the bringing to term of the other’s pregnancy of soul” (“Sociality” 110; 117).18 Not only a pregnancy but perhaps also a rehabilitation of the soul is at stake at moments in which a utopian impulse is evident in Bersani’s work—as when, for example, he speaks of effecting “a curative collapse of social difference,” or of enabling a future enjoyment of “as yet unarticulated pleasures” that have thus far been “suppressed and crippled” (Homos 177; “Fr- oucault” 137; emphasis added). If, as these examples suggest, Edelman is correct in asserting that we cannot think of the future without reference to the Child—and if I am right in suggesting that the overlapping ideology of rehabilitative futurism is equally pervasive and insidious—then how should disability theory answer O’Brien’s question? The disability rights movement, of course, has already provided compelling responses: protestations against the injustice of institutionalization, critiques of the nursing home lobby, and advocacy for attendant programs. Theoretically, it could be said that the goal of de- institutionalization is merely a liberal one, as it aims only to include disabled people within the social fabric. Yet in this instance (and many similar ones), an imperfect politics clearly seems better than no politics at all. But what is the role of disability theory in relation to this politics? Is it, as Paul Longmore described disability studies in 2003, to serve as the “academic counterpart to disability rights advocacy” (Burned 2)? Or should disability theory conceive of itself as sometimes in tension with this movement (as queer theory often is in relation to the mainstream lGbt movement)? Insofar as it has acted as a “counterpart” to the disability rights movement, disability studies has made crucial contributions to what might be called a humanizing enterprise. It has offered, for example, myriad analyses of the reasons for our society’s willingness—its desperation, even—to dehumanize and exclude disabled people, even to the point of locking them up. But when sex enters the picture, things get complicated. Consider, for example, the following remark, made by a doctor to a group of patients at one of O’Brien’s rehabilitation hospitals: “You may think you’ll never have sex again, but remember . . . some people do become people again” (Human Being 80). The doctor’s comment points to a paradox that inheres in any conversation about sex and disability: disabled people, it is implied here, are less than fully human because they are presumed not to “have sex”—but sex, psychoanalysis shows us, is radically dehumanizing, effecting a “shattering” of “the structured self” rather than its entrenchment in personhood or identity. This paradox is at the root of the double bind I discussed in the introduction to this chapter, in which disability simultaneously figures sexual excess and sexual lack: disabled people are regarded as sexually deficient and therefore not fully human, but at the same time, disabled people register as less than human because disability is the ubiquitous figure for a dehumanizing, identity- disintegrating force that resembles sex. If, as the second half of this paradoxical construction suggests, assertions of humanity are in necessary conflict with expressions of sexuality, then perhaps disability theory should, rather than seeking to humanize the disabled (insisting that disabled people be treated “as human beings”), instead ask how disability might threaten to undo, or disable, the category of the human. It might do so in part by attending to the insights Bersani’s and Edelman’s readings of psychoanalytic theory yield, according to which sex, far from enabling us to “become people,” ruptures the self and dehumanizes us all. But what, then, would become of disability politics? Critics of No Future— despite Edelman’s insistence that its argument pertains to “figurality,” not to “being or becoming” the death drive—tend to read the book as advocating, on a literal level, the abandonment of hope and political goals (No Future 17; 25).19 As noted earlier, however, it is “politics as we know it” that Edelman refuses, and even this refusal does not mean that queers should stop insisting on “our equal right to the social order’s prerogatives” (No Future 3; 29; emphasis added). Edelman further clarifies this point in his essay “Ever After”: “Without for a moment denying the importance that distinguishes many [political] projects, I want to insist on the need for an ongoing counterproject as well: a project that’s willing to forgo the privilege of social recognition” (473; emphasis added). Such a counterproject—one that can be read as possibly opposing the humanizing impulse behind O’Brien’s narration of How I Became a Human Being—may take shape in some of O’Brien’s own poetry. While the title of his autobiography speaks of becoming human, his unpublished poem “Femininity” disrupts this trajectory. O’Brien writes of lying: Naked on the gurney in the hospital corridor, surrounded by nurses, tall, young, proud of their beauty, admiring my skinny cripple body. “You’re so thin, you should’ve been a girl.” “I wish my eyelashes were as long as yours.” “Such pretty eyes.” I thought or think I thought or wish I’d said, “But your bodies work. Get scissors, cut my cock and balls off. Make me a girl, without anaesthesia, make me a girl, make me a girl.”20 Much of the unnerving intensity of these lines derives from what, invoking Bersani, we might refer to as their embrace of “the suicidal ecstasy of being a woman” (or a girl, or queer, or disabled); from their rejection, that is, of the ideology of rehabilitative futurism, and from their refusal to engage in a “redemptive reinvention” of sex or disability. O’Brien’s speaker does not plead with the nurses who admire his “skinny cripple body” to “cure me” or “make me walk again.” Nor does he attempt to redefine his body (which does not “work”) as merely a manifestation of human variation. Suffering and lack, rather than being dissociated from disability, are amplified and eroticized: “cut my cock and balls off . . . without anaesthesia,” the speaker implores, the repetition of his plea (“make me a girl, / make me a girl”) evoking the repetitiveness of a drive. “Femininity” can indeed be read as an instantiation of the disability drive: disability in this poem, like “the rectum” in Bersani’s essay, “is the grave in which the masculine [and nondisabled] ideal of proud subjectivity is buried.” It will of course be tempting to evade this “nightmare of ontological obscenity” (“Rectum” 29), this fantasy of unbecoming human.21 But the dehumanizing double binds that so persistently structure cultural representations of sex and disability suggest that such evasions may be futile. Intrinsically obscene, yet inherently asexual: rather than attempting to assume a different position within this impossible paradigm, disability theory should perhaps underscore its pervasiveness as evidence of a disability drive; as a sign, that is, that our culture’s desexualization of disabled people functions to defend against a deeply rooted but seldom acknowledged awareness that all sex is incurably, and perhaps desirably, disabled.

#### You should adopt epistemological disablement over certainty. Their attempt to construct a universal guide to action ignores the ways in which uncertainty, confusion, and the overall drive determine us. This means their framework should be ep suspect because it is contingent on a complete knowledge of the self which the disability drive denies.

**Mollow 15** The Disability Drive by Anna Mollow 2015 // UTDD

“How, then, might we begin to acknowledge our own determination by the drive? Any knowing of the drivethat we might hope to achieve must, on account of the structural barriers that render the drive unthinkable, be an effort characterized by failure and incompletion—that is, we might say, by epistemological disablement**.** The term “epistemological disablement” will appear frequently in this dissertation, as I will argue that coming into close proximity with the disability drive produces states of cognitive and affective uncertainty, confusion, and incapacity that are akin to disability. In the works that I shall analyze, epistemological disablement will often be performed on a textual level, as theorists and narrators seem to lose control of what they want to say about disability. These moments of epistemological disablement are often disavowed by theorists and narrators and are instead projected onto disabled people. When this happens, disabled people’s impairments are depicted as the result of an insufficiency of self-knowledge that is assumed not to determine nondisabled subjects. I will challenge these characterizations of disabled people not only by arguing for the value of “cripistemologies” (that is, ways of knowing that arise from disabled people‟s lived experiences) but also by using drive theory to undermine belief in the possibility of a transparent and wholly knowable self, whether disabled or nondisabled.18 My two-pronged approach to the issue of epistemological disablement may seem to present a paradox: on the one hand, I am asserting that disabled people’s lived experiences generate important knowledge about disability; yet at the same time I am seeking to destabilize the very notion of self-knowledge. Let me be clear, then, that in undertaking this double endeavor I do not forward all-or-nothing claims either “for” or “against” the possibilities of self- knowledge. I will not assert that people cannot ever know anything reliable about themselves, but I will also not suggest that truth claims derived from personal knowledge about disability are infallible. Instead, this dissertation highlights the limits of complete self-knowledge for nondisabled and disabled subjects alike, while at the same time interrogating the social dynamics that give rise to imbalances in the distribution of epistemological authority to particular subjects on the basis of their perceived status as disabled or nondisabled.” (4)

#### Communicative spheres always zone out disability – breaking down notions of progress is necessary in the face of social death. Thus, the role of the judge is to question ideological optimism in the classroom, and the ballot is to vote for the debater who best disrupts notions of progress within civil society.

Selck 16 [Selck, Michael L. "Crip Pessimism: The Language of Dis/ability and the Culture that Isn't." (Jan 2016) // WHSRS and Lex VM]

Despite the fact that a large basis of American culture is founded on ability, dis/ability rarely enters the dominant public communication sphere. The unpleasant and visceral questions that accompany communication about dis/ability have been strategically re-zoned and relocated like so many dis/abled patients, veterans, and transients. Yet, when conversation about dis/ability does seem to permeate the ideological walls of ability the messages are inspirationally distorted and optimistic. My time researching dis/ability in academia found that the conversation there mimicked the exploitive inspirational humaninterest trope found in cinema and journalism. To break the optimistic silence I set out with a performance art piece titled Under The Mantle to advance a theme of crip-pessimism, which intended to raise the stakes of contemporary dis/ability research. The beginning of this essay takes the time to detail the vast theoretical backgrounds of critical disability theory and philosophical pessimism. In the following section I reviewed intercultural communication literature for dis/ability because much of the theory literature I drew from existed outside the communication studies discipline. The evidenced lack of intercultural dis/ability artifacts up against a dis/ability centric performance art project necessitated an interdisciplinary multi-method framework. In that framework I demonstrate how autoethnography is significant to dis/ability studies because it illuminates even the most mundane able-bodied norms. In the final sections I offer a textual description of the performance and hone in on three explicit arguments that augment traditional thinking about dis/ability and communication. The trouble I encountered with dis/ability research in communication studies has to do with the way American culture understands offensive communication. Political correctness as a disciplining communication concept dictates what terms are socially acceptable at a given time. Political correctness underscores how many communication studies programs operate within the rubric of conflict (Wilderson, 2010). The thinking that suggests simply avoiding offensive terms will diminish oppression is within the rubric of conflict because it understands the oppression as materially reconcilable. What crippessimism does, and what UTM performed, is skepticism that speaking inspirationally and avoiding speaking offensively about dis/ability would end disablism. Instead I argued that what dis/ability represents is an antagonism, it is an oppression so much more foundational to the core of American values that linguistic reforms would not even scratch the surface. The significance of antagonism is that it raises the stakes of dis/ability research. The end goal of research should not be to service the meta-theoretical assumptions of the paradigm (Kuhn, 1962), because consequently the researcher never stops to ask if the assumptions of the paradigm are ethical, valid, or effective. Crippessimism is a call for some demolition and redistribution of communicative identity paradigms. If the radical promise of our theories is nothing more than a call for social stability then they are complicit in the neoliberal eugenic project. We need to theorize so that there is nothing already ‘given’ or taken for granted. Often in those moments, like the moments of so many textbooks, the underlying optimism goes completely unquestioned. Crip-pessimism as a theme is characterized by negotiating debates surrounding the efficacy of identity politics. Arguments that fit within the theme ask why the disabled should abandon their bodies in the political sphere. Social death has already occurred, the dis/abled are being rendered culturally unintelligible and physically fungible. So what we need when we are having discussions about how to progress is a theory that breaks down the notion of progress. The recognition and need for a theory like this comes about when we ask central dis/ability questions like: ‘when did eugenics end?’ and ‘where is disability in U.S. society before and after the passage of the Americans with Disabilities Act?’ and ‘globally has the Convention on the Rights of Persons with Disabilities reconciled the antagonism of disablism?’. These are the questions that I want to end on and encourage communication and dis/ability scholars alike to take up. As scholars and mass media engines continue to project dis/ability within the rubric of conflict our collective reliance on capitalism and neoliberalism grow deeper. It is my hope at the end of this project that my voice both in performing and in writing encourages more scholarship detailing the omnipresence of disablism in American culture. Under The Mantle is a reminder to me that all representations of dis/ability have consequences and in many cases all we need to witness those consequences is a slight perspectival shift.

# Accessibility

## 1

#### Abled subjectivity is tied up in a two-tiered affective response that explains disabled life – primary pity which reflects disability upon the ego threatening its ability status, which invokes secondary pity to overcorrect for the shattered-ego necessitating disabled death.

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pity involves a reaction to the “tragedy of disability.” Primary pity is unbearable to contemplate another person’s suffering is to question, “Could this happen to me?” Secondary pity attempts to heal the ego at someone else’s expense and calls for a cure. we feel primary pity and then deny that we have felt it.

#### The 1AC’s belief of a better future becomes complicit in the logic of rehabilitative futurism, which is threatened by the Disabled Child.

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the image of the Child” is inextricable from disability the Child is a display of pity that demeans disabled people. the Child makes an excellent alibi for ableism because not fighting for is unthinkable. The logic relies on “rehabilitative futurism,” Futurity is imagined that the eradication of disability would bring a better future.

#### The alternative is to disable the figure of the “human” – instead of seeing disability as a redeemable position within civil society, the alternative weaponizes disability’s structural position against the human. If we win their starting point is ableist they cannot weigh the consequences of it.

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expanding the category of “human” to encompass those excluded will not “stop the inhumanity of the drive rather than conforming to a utopian/antirelational binary think both at once disability studies has offered reasons for society’s desperation to dehumanize and exclude disabled people the double bind simultaneously figures excess and lack disability theory should rather than humanize the disabled instead disable the human. abandon politics Suffering and lack, rather than dissociated from disability, are amplified It will be tempting to evade unbecoming human. But the dehumanizing double bind suggest that evasions may be futile. rather than to assume a different position within this impossible paradigm, disability theory should underscore its pervasiveness as evidence of a disability drive

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Any knowing of the drive must be an effort characterized by failure and incompletion—that is epistemological disablement coming into proximity with the disability drive produces states of affective uncertainty These moments are often disavowed by theorists and are instead projected onto disabled people disabled people are depicted as the result of an insufficiency of self-knowledge drive theory undermine belief in the possibility of a transparent and whol self this highlights the limits of complete self-knowledge for subjects while interrogating social dynamics that give rise to imbalances in epistemological authority

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dis/ability rarely enters the communication sphere. The questions that accompany dis/ability have been relocated like many patients Political correctness as communication underscores how programs operate understands the oppression as materially reconcilable dis/ability is an antagonism linguistic reforms would not scratch the surface Social death has occurred we need discussions about how to break down progress