Chapter One—Introduction and Context

Overview

Though this project follows many artistic and conceptual avenues, can be characterized by the exploration of three questions: **how has disability been presented in art and media?** What are we losing if those with disabilities are “othered” and excluded? **How can disability be ethically portrayed in art?** As a creative thesis, the exploration is structured in two parts, a series of six portraits and a written component. The aim of this project is to highlight the humanity of disability, to show people with a disabilities as humans with agency, and to discuss what has been missed and what will be lost if social spaces, specifically artistic ones, are closed off not only to ethical representation of those with disabilities but to their actual presences.

In order to achieve these goals, I was highly intentional with the selection of the individuals portrayed in this project. Thus, I chose three close friends, each with a different disability. Their engagement and participation in the process was a vital aspect of the agency I hoped to highlight, so each individual was interviewed and was highly involved throughout the project. Building upon our relationship, our interview, and our collaborative approach, I painted both a large-scale portrait and a smaller, detailed portrait of a defining quality of the sitter. Though these are not self-portraits, I hope that I have captured my subjects as authentically and genuinely as possible.

The written portion of this project is supplemental to the artistic work described above and is broken into four chapters. This chapter serves as an introduction to the project and its context, including an overview of disability studies in art historical discourse, its relationship with disability, and the stereotypes it enforced. The second
chapter offers an insight into the artists that have questioned the notion of the “Other” in their work, many of them specifically challenging the stereotypical historical presentation of those with disabilities. These artists—such as Frida Kahlo, Mary Duffy, Marc Quinn, Richard Avedon, and many others—have been crucial to my research and inspiring to my work. The third chapter builds upon the foundation that those artists laid, tracing what I hope this project adds to the conversation and the exploration of disability’s role in art. As it focuses on agency and humanity, it is the conceptual and philosophical backing of the painting component. The fourth and final chapter is strictly reserved for the creative process. It discusses my growth as a painter and portraitist as it culminates with this series, and it documents the creative process I embarked upon with the subjects of the portraits.

Context

Though disability is ever-present throughout history as a quality of the human condition, it has been underrepresented, misconstrued, and rejected. Specifically in art history, “people with disabilities have often served as visual and cultural objects, rather than as active participants in and creators of culture and media.”¹ Reduced to nothing aside from their disability, they are depicted by artists and interpreted by historians as an othered group. Amanda Cachia articulates this process:

There is an internalized, almost unconscious assumption of able-bodiedness in art theory and praxis—if the assumption becomes “disrupted” by nonnormative corporeal forms, then these forms have historically been rejected and marked as pathological, diseased, and “other.”²


Thus, even when disability is displayed in artwork, it is pushed out of the scope of the normative, human experience. Simultaneously in art history, artists who were disabled themselves were not provided the opportunity to engage with their disability in their art. While Claude Monet was visually impaired later in his life and Henri de Toulouse-Lautrec was short-statured due to a congenital bone disorder, “there are no written accounts on the lived experiences of disability for each of these prominent artists, as their disabilities were perceived as irrelevant and invisible.”

Survey of Art History

In order to provide the necessary context of disability in art history, I turn to Professor Keri Watson’s cohesive chronological survey of the ways in which disability has been artistically portrayed in the Western tradition and how those portrayals led to harmful stereotypes of people with disabilities. For brevity, I will condense her survey into two major themes: disability in religion and disability as a novelty. Both highlight how those with disabilities have been simultaneously excluded from being active participants in their culture and stereotypical objects of unwarranted attention. This paradox of exclusion and inclusion permeates disability history and is an important theme that will be drawn upon throughout the course of this project.

Beginning with the fall of Hellenism and the rise of Christianity, disability transitioned from a societal burden to a sickness requiring supernatural healing. Works such as Niccolò Vicentino’s *Christ Healing the Lepers* (ca. 1540-1550) and El Greco’s

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3 Ibid., p. 165.

*Christ Healing the Blind* (1570) reflect the negative attitude towards disability. While this characterization of disability was born from the Christian commission to care for the downtrodden, it has led to the common religious stereotype that disabilities were a form of divine punishment, that they “were cursed by Satan, and that their disability was a result of sin.”  

Both of these paintings are excellent examples of how such a stereotype strips those with disabilities to cast them instead as othered beings, so lowly that only an act of divine will could restore them to their rightful place. As the message of Christianity spread, so did the power it bestowed upon its followers. By the middle of the 17th century, apostles and other Christians can be observed healing the disabled in art history as well.

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This shift in thought concerning disability is evident in writings from prominent church leaders of the 15th and 16th centuries. Not only did Martin Luther categorize those with cognitive disorders as “filled with Satan,” he even “advised that children with severe mental retardation be drowned, because they lacked souls.”⁶ John Calvin also claimed that predestination did not hold a place for those with disabilities.⁷ As the Church climbed as a world power, such religious attitudes towards disability deeply entrenched the stigma of disability and led to the characterization of certain individuals as the Other.

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It is from this religious stereotype of disability that the medical classification of disability arises; no longer was this stereotype confined to the religious, but it spread to the secular realm, as doctors and scientists searched for “cures” to disability.\textsuperscript{8} Eugenics, Social Darwinism, and the "survival of the fittest” became the general attitudes and proposed solutions to disability, further entrenching the identity of disability as subhuman and extra-societal, as the Other.\textsuperscript{9} Eventually, both the medical and religious stereotypes enabled media to portray those with disabilities as a thing to be feared. Villains in fiction consistently exhibited physical deformities and disabilities—characters such as Quasimodo, Captain Hook, Dr. Strangelove, and Frankenstein all encourage the trope that disabilities are evil and nonhuman.

\textsuperscript{8} M. Miles, “Martin Luther and Childhood Disability in 16th Century Germany: What Did He Write? What Did He Say?”

Reinforced by eugenics and propaganda, the evil stereotype of those with disabilities enabled historical figures like Hitler to push extensive social exclusion and even genocide of those with disabilities.¹⁰ Negative stereotypes such as these, however, did not always culminate in such exclusive efforts; interestingly, an immense level of fascination with disability coexisted with such disdainful attitudes, causing those with disabilities to also be viewed as a novelty.

Falling back to the 15th century, Watson traces the growth of the view of disability as a novelty based on the development of the so-called “Freakshow.”¹¹ Undesirable members of society, especially those with physical and intellectual disabilities, were paraded and exhibited around the world for profits, often abandoned afterwards. Hieronymus Bosch’s Ship of Fools (ca. 1490-1500), depicting a group of societal outsiders, has ties to this exploitation of those with disabilities. The characters in the painting, all deemed societal burdens of the time, are portrayed as “deranged, frivolous, and oblivious inhabitants who are seemingly ignorant of their course.”¹² Because neurodiversity was little understood in the renaissance era, often mental illness and intellectual disability were lumped into the category of lunacy. In his translation of Michel Foucault’s Madness & Civilization: A History of Insanity in the Age of Reason, Jose Barchilon comments on the stigma displayed in Bosch’s Ship of Fools:

Renaissance men developed a delightful, yet horrible way of dealing with their mad denizens: they were put on a ship and entrusted to mariners because folly, water, and sea, as everyone then “knew,” had an affinity for each other. Thus, “Ships of Fools” crisscrossed the sea and canals of Europe with their comic and

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¹¹ Ibid.

¹² Ibid.
pathetic cargo of souls. Some of them found pleasure and even a cure in the changing surroundings, in the isolation of being cast off, while others withdrew further, became worse, or died alone and away from their families. The cities and villages which had thus rid themselves of their crazed and crazy, could now take pleasure in watching the exciting sideshow when a ship full of foreign lunatics would dock at their harbors.¹³

![Figure 7. Hieronymus Bosch, Ship of Fools, ca. 1490-1500, oil on panel.](image)

Such fascination with disability, Watson explains, led to the upper and aristocratic classes to take individuals with disabilities on as “pets.”¹⁴ Such treatment of disability is depicted in paintings of jesters in royal courts. Typically, jesters with dwarfism were illustrated as subservient and subhuman pets to their masters, like in Gaspar de

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Crayer’s *Portrait of Philip IV of Spain with His Court Dwarf* (ca. 1632) and Giacomo Vighi’s *Charles Emmanuel I of Savoy as a child accompanied by a dwarf* (ca. 1573). In these paintings, the artists not only placed the jesters in physically lower positions in the painting—like below and behind their master—but they even portrayed emotions of subservience, adoration, and contentment on their faces, reinforcing their social position as well. Diego Velázquez’s *Portrait of Sebastián de Morra* (ca. 1645), more recently reidentified as *The Buffoon El Primo*, depicts on the other hand a more formal and respectful approach to dwarf portraiture that differed from others of its time. One of the few that focuses solely on the jester, the sitter feels solemn and even distrustful. Even so, in Velázquez’s portrait, the jester is granted much more respect than many other court jesters of the time.


With the rise of the entertainment industry, individuals such as P.T. Barnum exploited and profited off of individuals with disabilities by marketing them as “living
curiosities” in museums and “freak shows.” Opening his American Museum in 1841, P.T. Barnum catalogs several individuals with disabilities as “profitable attractions” in his *An Illustrated Catalogue And Guide Book To Barnum’s American Museum*. He first names ‘General Tom Thumb,’ a world-renowned individual with dwarfism, along with his wife and other ‘dwarfs’—‘Commodore Nutt’ and ‘Miss Minnie’—as the first curiosities of his museum. Their names alone exhibit the ridicule and belittlement these individuals were subjected to because of their disabilities. Among his other attractions, Barnum included a family of giants, siblings with albinism, and individuals with microcephaly in the guide to his collection of oddities.

Photographers and artists document the rise of such establishments and their attractions, giving a glimpse into the ways in which disability was exploited for profit. What scholars have deemed “Freak Show Portraiture” features photographs of individuals with disabilities, especially those who gained renown because of their extraordinary bodies. Such portraiture of the nineteenth century—*carte-de-visites* and cabinet-card depictions of Freak Show performers—falls into the stylistic traditions of Victorian portraiture, specifically bourgeois self-portraiture. While many of these individuals capitalized off of the public’s interest in their disability and lived comfortable, successful lives, they were often known for their disability alone. In many of the

16 Ibid., 106
17 Ibid., 107.
18 Ibid., 107-113.
photographs and portraits of individuals with disabilities during this period, their name, agency, individuality, and often humanity, are stripped away.

Before an analysis of this photographic industry, it is necessary to offer a definition, clarification, and context for the terms “freak” and “Freak Show.”

Guy C. M. Kirkwood, in his analysis of Freak Show photography, does so in a concise statement:

By “Freak Shows” I refer to the practices of displaying human difference for profit, particularly popular in America from the mid-nineteenth century until around the 1920s and 1930s. However, the display of “extraordinary bodies” was as much a performance involving stylized modes of presentation as a bodily “condition.” It involved portraying oneself or being portrayed—in a certain manner or pose, drawing on historically and culturally contingent conceptualizations of what was considered “freakish” or “normal,” both of which were infused with fluid and contested elements of race, gender, ablebodiedness, and sexuality.

Freak show photography, Kirkwood explains, was no longer the scientific documentation that was used in eugenics to “type” those outside of the norm—whether in categories of race, gender, disability, etc.—but had shifted towards an industry that capitalized on disability:

Freakishness was marketed and packaged and, like the fables of abundance, this commodification involved more than simply meeting the forces of supply and demand—it involved engaging with dominant ways of being in the world, contested identities, and shifting worldviews.

With the rise of this commodification of disability, individuals with disabilities were documented as novelties. Work from Charles Eisenmann and Diane Arbus offers a specific insight into the attitude surrounding disability during this period.

20 Ibid.
21 Ibid.
22 Ibid., 6.
After the success of P.T. Barnum’s American Museum, similar establishments displaying bodily abnormalities and curiosities became a trend in the entertainment industry. Photographers from across the world flocked to places like the Bowery area in New York City because of its abundance of Freak Shows. Charles Eisenmann, “a German immigrant who came to America in 1868, proved highly successful in this trade for over a decade and a half, and left behind more photographic portraits of extraordinarily bodied performers than anyone else.” At the time, small, pocket-sized photographs—called “carte-de-visites” or “cabinet cards”—were immensely popular, and were collected, becoming “a common feature in Victorian homes.” Eisenmann capitalized upon this interest, focusing his work on capturing “Freaks” in small portraits.

It is important to acknowledge that Charles Eisenmann did expand the restricted perception of individuals with disabilities by the ways in which he had his subjects sit and pose. He employed more bourgeois poses and styles, granting much more respect to individuals with disabilities. Still, however, Eisenmann along with other Freak Show photographers greatly profited off of capturing disability as a novelty.

Kirkwood comments on Eisenmann’s method of posing his subjects:

Extraordinary bodies could be portrayed in cabinet cards not simply as an example of a particular typology of bodily difference, but as part of a performance that emphasized the abundance of embodied possibilities, thus both stabilizing and undercutting dominant cultural narratives.

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23 Ibid., 10.
24 Ibid., 10.
25 Ibid., 10.
26 Ibid., 12.
27 Ibid., 12.
These posed photographs were an extension of the performers’ acts at their Freak Shows. While individuals with disabilities were granted freedom of expression in this style of portraiture, it also enabled photographers to capitalize off of their bodily differentness, their Otherness. No matter how pompously dressed or stylishly posed, disability was the central focus of Eisenann’s photography of individuals with disabilities.


Diane Arbus, a twentieth-century photographer, is also recognized as working within the category of Freak Show photography. Though she lived much later than Eisenmann, she is acknowledged to have treated her subjects, especially those with disabilities, arguably worse than those of Eisenmann’s age. She had an odd fascination with the outsiders of society—she often focused on populations that differed from the norm in ability, sexuality, and ethnicity. Germaine Greer, who posed for Arbus in the 1970s, comments from personal experience on the photographer and her treatment of her subjects:
Though formally Arbus is within the tradition of freak photography, there is an important difference between her and her predecessors like the Eisenmann Studio or Obermann and Kern. Their subjects had names, lots of them, stage names and real names. The giants, dwarves, midgets, conjoined twins, bird-girls, bearded women and dog-faced boys whose photographs appear on thousands of postcards were all professionals. Often the notes on the postcards spoke of them as well-educated and happily married. Arbus's nameless subjects are denied such confederacy and performativity. She often uses the devices of the older tradition in her treatment of otherwise unremarkable subjects. Her famous Identical Twins, Roselle, NJ of 1967 are posed as if they were joined at the shoulder and hip and had only three arms between them. She reduced her subjects to generic phenomena by the names she chose for them: Jewish Couple, Puerto Rican Housewife, Albino Sword-Swallow. My ordeal resulted in a picture called Feminist In Hotel Room. No permission for the reproduction of what is an undeniably bad picture was ever requested.

Greer goes on:

The language Arbus uses about her photographic practice is revealing: “Freaks was a thing I photographed a lot. It was one of the first things I photographed and it had a terrific kind of excitement for me. I just used to adore them. I still do adore some of them.” “Freaks” (a word 21st-century sensibility finds hard to use) is “a thing”, a medium for her use that Arbus finds quite distinct from herself.

Greer pinpoints a crucial component of Arbus’ attitude towards “Freaks.” She reduced them to their Otherness, documenting their difference or disability but neglecting to even record their name or allow them to approve any of the photos. While some praise Arbus for including disabled, queer, and other diverse populations in her work, the attitude she displays above is not inclusive by any means. Arbus viewed her subjects—her “Freaks” —as specimens different from herself. This insensitive fascination with disability especially in art is a reflection of the treatment of disability as a novelty and commodity that has permeated entertainment culture since the late eighteenth century. It is artists such as Diane Arbus that have prolonged such negative treatment of disability: “Arbus is


29 Ibid.
not an artist who makes us see the world anew; she embeds us in our own limitations, our lack of empathy, our kneejerk reactions, our incuriosity and lack of concern. Hers is a world without horizons where there is no escape from self."\(^{30}\)


\(^{30}\) Ibid.
Chapter Two—Inspiration and research

“The second chapter offers an insight into the artists that questioning the notion of the Other in their work, many of them specifically challenging the stereotypical historical presentation of those with disabilities. These artists—such as Frida Kahlo, Mary Duffy, Marc Quinn, Richard Avedon, and many others—have been crucial to my research and inspiring to my work.”

Despite the negative stereotypes of disability—such as the views of disability in religion or in the entertainment industry—permeating art history, there are several artists whose work pushes against those norms. While some of these artists specifically addressed the topic of disability and others did not, each of these artists challenged the stigma of Otherness held by contemporary society with their work in portraiture, and their art has been vital to my research and composition of this project. This chapter serves to introduce the artists that proved especially influential in my research of disability in art. First, I will introduce them and their work, and then I will illustrate the ways in which they redefined the art of portraiture and influenced this project.

It would be beneficial to begin with a discussion of artists who challenged and redefined portraiture, though not specifically addressing disability. In their photography, artists such as Richard Avedon, Nan Goldin, Amy Sherald, and Kehinde Wiley have challenged the narrative in portraiture and the stereotypes that surrounded those who had been misrepresented and excluded from art. As this project is a work of portraiture, these artists have have been indispensable for both my creative development and for the evolution of this project.

Richard Avedon, born in 1923, is arguably one of the most influential photographers of the 20th century. He first gained fame in the fashion industry, after he
revolutionized the field by “elevating fashion photography to an art form.”

When Mitchell Wilder, the director of the Amon Carter Museum of American Art in Fort Worth, saw Avedon’s work in 1978, “he gave the photographer free license to photograph his view of the American West.” From 1979 to 1984, Avedon toured through “13 states and 189 towns from Texas to Idaho, conducting 752 sittings and exposing 17,000 sheets of film” in his exposition of the people of the American West.

Visiting ranches, rodeos, truck stops, oil fields, and slaughterhouses, Avedon focused on the true, rural populations:

Rather than playing to the western myths of grandeur and space, he sought out people whose appearance and life circumstances were the antithesis of mythical images of the ruggedly handsome cowboy, beautiful pioneer wife, dashing outdoor adventurer, or industry mogul. The subjects he chose for his portraits were more ordinary people, coping with personal cycles of boom and bust.

The photographer did not seek to glamorize or romanticize the people and lifestyle he captured, but “brought their human frailties to the forefront.” His portraits were “dramatically large” and black and white against a white backdrop, removing the subject from “any reference to place.”

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32 Ibid.

33 Ibid.

34 Ibid.

35 Ibid.

36 Ibid.
Both conceptual and physical qualities of Avedon’s “In the American West” have influenced this project. First, his careful aim to neither glorify his subjects’ way of life nor differentiate his subjects from the rest of humanity. Avedon sought to capture people as their authentic selves, yet he simultaneously removed them from their background, forcing his viewers to be challenged by the stereotypes and stigma that they held for the subjects and their lifestyles. This aspect of Avedon’s work has been one that I have
sought to emulate in my own. I photographed my subjects in their home, amidst their
daily life, but I utilized a neutral background, to draw focus to the subject alone. While
Avedon’s sitters often have a poised and posed composure, I sought to capture the
sitters in their most genuine and natural state.

An artist who specifically focused on capturing her subjects as their natural,
unposed selves, unlike her contemporary Richard Avedon, is Nan Goldin. Born in 1953,
Goldin experienced a traumatic childhood and left home in her early teens. Her older
sister Barbara, whom Goldin was very close with, took her own life when Goldin was
eleven.37 The secrecy and avoidance with which her parents dealt with Barbara’s death
led Goldin to seek intimacy in her photos. After leaving home, Goldin was taken in by a
community of drag queens in Boston—she recognized them as her new family and
wanted to document their lives together, thus her early work as a portraitist revolved
around that world. It included intimate snapshots into the lives of the Boston queer
community, the AIDs epidemic, and the sexual lives of Goldin and her friends.

Though many aspects of Goldin and her work influenced my project, one specific
aspect of her creative process had a critical effect. She viewed her work as an ongoing,
collaborative, creative process with the people around her. Unlike artists like Diane
Arbus or even Richard Avedon, Goldin stated that she never sought to restrict her
subjects—her friends, her family—to a single image:

I don’t believe in the decisive moment. I’m interested in the cumulative images,
and how they affect each other, the relationships between them. There is so
much more said than by a single image.38

www.moma.org/artists/7532.

Like Goldin, I hope to offer more than a singular image of the subjects of my paintings, thus I portrayed each individual in a two-part series. The first is a formal portrait, similar to the dramatically oversized and stark portraits of Avedon’s “In the American West.” The second part, however, is much more like Goldin’s concept of using multiple portraits to convey an ongoing narrative of an individual’s life. Instructed by our interview and guided by our relationship, I painted each of these smaller portraits with the subject’s favorite qualities about themselves in mind, offering an intimate snapshot into an individual’s life, personality, and humanity. Goldin achieves a high level of intimacy in her work. Similarly in my project, I aimed to encapsulate and capture more than just the outward appearance of my subjects.


The next artist that greatly influenced this project is the contemporary painter, Amy Sherald. In 2016, Sherald was the first woman in twelve years to win the Smithsonian National Portrait Gallery’s Outwit Boochever Portrait Competition with her
Miss Everything (Unsuppressed Deliverance).\textsuperscript{39} Soon after, Sherald was selected to paint First Lady Michelle Obama for her official portrait, bringing so much "unimagined public attention," that the National Portrait Gallery had to relocate the piece to a larger room.\textsuperscript{40}

Her portraits—large-scale pieces with hyper-realistic or even surreal features—deal with how Black people exist in the world, forcing viewers to “confront the psychological effects of stereotypical imagery on African-American subjects."\textsuperscript{41} Sherald displays her subjects with an air of quiet grace and poise—a sense of balance and peace pervades her work. Often juxtaposing a vague, yet stereotype-inciting title with a portrait of a Black person, Sherald forces her viewers to confront the subconscious stereotypes that they hold of the individual depicted in the portrait.

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\textsuperscript{40} Ibid.

\textsuperscript{41} Ibid.
Though there are many admirable qualities in Sherald’s paintings, I found the way in which she paints each of the individuals in her portraits to be the most influential. The poise, gracefulness, and power Sherald instilled in every painting, enabling her to challenge her viewers and their beliefs, are qualities I sought to portray in each of my portraits.

The next artist who proved especially significant in my research is Kehinde Wiley. Though the focus of Wiley’s work is thematically similar to Amy Sherald, his method is different. Through his portraiture, Wiley has sought to insert African American men and women into art, specifically in positions of power. He uses classical portraits as inspiration to create images of Black men and women—often utilizing elements of pop culture and bright patterns to ensure that he is not just pasting Black people into a historically whitewashed art scene, but that he is showcasing the Black experience, narrative, and culture.
Wiley’s work has been influential in my research in two specific ways: his artistic techniques and conceptual motives. While he often maintains traditional painting methods, Wiley creates artworks that push the boundaries of modern portraiture through his use of color, pattern, and pop culture references. In my project, I strove to do something similar: I wanted to maintain many of the formal qualities of traditional portraiture—like realistic proportions and color—but with a modern twist. I utilized neutral backdrops and modern objects—like tattoos or toys—in my portraits. Secondly, the way he paints a historically disenfranchised group in a position of power, agency, and beauty is a quality I sought to emulate in my project. Though I did not take specific classical portraits and reimagine them representing a marginalized group as Wiley does, I did call on classical portraiture in technique and seek to portray each of the individuals I painted with the same respect that Wiley often inscribes in his portraits.

Though the artists discussed greatly aided my development as a portraitist and the evolution of this project, the following individuals, who specifically dealt with disability in their work, have been the most vital: Frida Kahlo, Mary Duffy, and Alison Lapper. Of course, there are numerous artists, especially recently, that have created work relating to disability, but these artists, their work, and their creative process have been the main focus of my research. Not only did they challenge the stereotypes surrounding disability, they specifically used disability—the experience in a disabled body—as a narrative and expression in their work.

Frida Kahlo, the first and arguably most well-known artist who influenced this project, created much work that pertained to her experience as a disabled woman. After encountering polio as a young child and surviving a traumatic bus accident as a teenager, Kahlo’s life was riddled with complications and procedures. Throughout her life, she used painting as a mode to process and invite others into her daily experience in her own body. While Kahlo’s portraits are often complex, she depicts her own lived experiences—her pain, sorrows, joys, memories, and disability—as an embodied narrative. She does not isolate her disability as some other experience but includes it as an expression of her humanity. As Mine Inahara has argued, “constructed from a bricolage of her lived experiences,” Kahlo’s paintings are not focused only upon her disability, but rather on wholistic self-portraiture. “She continues, her work can be seen as a creative inquiry into her life, death, pain, disability, and suffering, in which she is communicating the nature of living in/through a body such as hers. This inquiry is made

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43 Ibid., 220.
through the mediation of art and thereby her experience of living is transformed through her art.”

Kahlo’s conceptualization of holistic representation has been priceless in this project. Similar to the inspiration I gathered from the manner in which Nan Goldin crafted an ongoing narrative through multiple images, I hope to emulate Kahlo and her approach to embodied portraiture. As I paint individuals with disabilities, I neither want to extract nor elevate their disability from their other qualities—doing so would not offer a holistic representation of who they are—nor do I wish to diminish the reality that their lived experience includes their disability. Through the two portraits of each individual, I hope I offer a glimpse into who they are as human beings.


44 Ibid., 224.
Mary Duffy and Alison Lapper—the next two artists that influenced my study of disability in art—have both created work that is an extension of Kahlo’s holistic and embodied portraiture. However, these two artists used their own bodies as a medium in performance art. Mary Duffy, born in 1961 without arms, is an expressionist painter and performance artist. I have followed her work as a painter for years, but have only recently come upon her performance art in researching this project.

Beginning in the late 1980s, Duffy utilized performance art to express her experience as a disabled person. Reminding her audience of famous statues from antiquity, Tobin Siebers has argued that “Duffy presents herself to the audience fully nude or draped, while reciting statements challenging the vision of her as defective and claiming her place alongside the Venus as a disabled beauty.” As she chants the hateful things said to her on account of her disability, viewers are forced to reevaluate their standard of beauty and its connection with disability. Documented via photographs, these performances take on a form of a narrative of Duffy and her experience as a disabled person.

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woman with a disability.

Figure 39. Mary Duffy, *Cutting the Ties that Bind (Heroes)*, 1987, photograph.

Duffy reflects on her series, *Cutting the Ties that Bind*:

By confronting people with my naked body, with its softness, its roundness and its threat I wanted to take control, redress the balance in which media representations of disabled women are usually tragic, always pathetic. I wanted to hold up a mirror to all those people who had stripped me bare previously...the general public with naked stares, and more especially the medical profession [that caused those stares in the first place].  

She goes on to address her performance work as a whole:

I wanted to create new images. Images that would celebrate, challenge and change. My body became a kind if metaphor. I worked through photography. I wanted to confront and contradict my own image of myself. I wanted to create a mirror image, creating illusion of sameness and subtly introduce the notion of difference. Maybe we are not all the same, maybe difference could be welcomed

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and is worth celebrating. The question I was asking myself and other people was, can you understand and accept my pride in differentness, and really wanting to celebrate and share it? My work is an exploration of difference and all that it implies—deviance, death, deformity, distrust, dismay, and also of delight, wonder, variety, inventiveness, creativity and uniqueness. It is a statement about and with my body. A body which does not have quite all the pieces to fit the human jigsaw, and is therefore not always considered whole. It is about looking and seeing new ways, new images where there has never been an image before without an able-bodied cultural prejudice. It is about finding a new identity for myself as a woman with a disability and my right to define that identity and its politics for myself.47

Similarly to Mary Duffy, Alison Lapper also calls upon classical art and reimagines it with her own body. Born without arms and shortened legs, Lapper utilizes the similarities between her body and statues such as Venus de Milo and Winged Victory. As Siebers explains, “like Duffy, Lapper engages in a deliberate recuperation of her own image as belonging among the most celebrated and valued representations of female beauty in the history of art.”48


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Both Duffy and Lapper’s form of self-portraiture challenge aesthetic norms in art and society. Viewers are forced to confront the presupposed disconnect between the disabled bodies they are looking at and the works of art that those bodies resemble—thus, viewers must reimagine their standards for beauty and extinguish their standards of able-bodiedness and retroactively reassess art history. Siebers, in his analysis of disability aesthetics, references this retroactive process:

The images that Duffy and Lapper make of themselves are seen as beautiful because they recall so powerfully the vision of beauty affirmed in the history of art by the Venus de Milo. But these images also change retroactively the perception of the Venus, for her beauty now incorporates necessarily the presence of disability. We cannot see Duffy and Lapper without seeing the Venus, and we cannot see the Venus without seeing Duffy and Lapper.49

While my paintings are not self-representation and are therefore nowhere near the intimate, embodied, and holistic representations that Lapper and Duffy offer in their self-portraits, I have been inspired by their references to classical art. In style, I have attempted to emulate qualities—such as the softness and absence of brushstrokes—that classical portraits often carry. In some of my smaller portraits—like Faith’s Arms and Zack and Woody—I have been able to subtly reference classical representations of hands. Similar to Duffy and Lapper, my goal with these subtly stylistic elements in my paintings is to force my viewers to retroactively reassess art history and the aesthetic standards it enforces. I hope my portraits remind viewers of classical portraiture in form and technique, yet I hope they are simultaneously challenged—like Lapper and Duffy’s audiences—to confront the disparity between the people portrayed in my project and the history of portraiture, the able-bodied norms it enforces, and its underrepresentation and misrepresentation of people with disabilities.

49 Ibid.
Finally, the last artist, and the one whose work bears the most stylistic and philosophical similarities to this project, is Chris Rush. The writer and illustrator is alive but has very little digital footprint or social media presence. I stumbled across one of his paintings that was briefly mentioned in Rosemarie Garland-Thompson’s *Staring: How We Look*, and was shocked at the similarity between our work. Unfortunately, Garland-Thompson’s virtual source for his work no longer exists, and as far as I can find, Rush has published no statements or comments on his work. I have attempted to contact him via social media but to no avail. Thus, the best context I can provide is how Garland-Thompson discusses his work in her book, and how I believe his portraits relate to my paintings. Of Rush and his portrait *Swim II* (figure 42) Garland-Thompson comments:

Rush gets between his subjects and our discomfort by softening their differences with the medium of conte crayon and posing them with great dignity. Rush’s pictures navigate between us and them, attending carefully to the visual relationship by gratifying our “deep curiosity” while at the same time inviting “empathy” and “sensitivity. . . One of Rush’s most arresting drawings presents a young woman in the regal profile pose we know from the familiar commemorative portraits of the Italian Renaissance. Her likeness emerges from the sharp line her stately features form against the background; her nose and chin lift imperially; her eyes gaze impassively down on the world beneath her. Her head is turbaned with a richly colored and ornately patterned aristocratic headdress, and her shoulders reveal a simple but elegant gown. On first glance, she looks like a modern Florentine lady. On second glance, however, we recognize a face we have never seen in a portrait. We see the distinct features of a person with Down syndrome, her hair wrapped in a bright beach towel, her face in a faraway reverie, and a simple heart tattooed on her shoulder below her bathing suit strap. The portrait invites us to stare, engrossed perhaps less with the “strangeness” of this woman’s disability and more with the strangeness of witnessing such dignity in a face that marks a life we have learned to imagine as unlivable and unworthy, as the kind of person we routinely detect in advance through medical technology and eliminate from our human community.50

Though the concepts of staring and portraiture will be addressed in the following chapter, it is important to highlight the similarities between this project and Rush’s portrait. Unfortunately, I have been unable to ascertain if he has completed any other portraits similar to *Swim II*, or if it is the only one. Nevertheless, the dignity and regality embedded in Rush’s portrait are qualities I aim to emulate in my own.

Figure 42. Chris Rush, *Swim II*. Conte crayon on paper. Portraits are life size in scale. No date.

Though not all directly pertain to disability, each of these artists and their work have challenged, provoked, and reimagined aesthetic standards of beauty in their representational work. They have each, in their own ways, deeply informed my research and paintings and have given me the foundation to expand the conversation of disabled representation in portraiture.
Chapter Three

Behind the Portraits

Despite the work of the artists discussed in the previous chapter, disability is still, as a whole, othered by society. Whether consciously or subconsciously, the general, able-bodied population does not consistently consider disability to be a quality of humanity. Thus, disabled individuals are pushed to the outskirts of society, severely underrepresented in art, media, and everyday life.

In her book *Staring: How We Look*, Rosmarie Garland-Thompson expounds upon this topic. She explains that disabled individuals continue to be othered on account of staring in American culture and how it has led to the simultaneous fascination and disgust with anything that differs from normalcy. Though she offers an extensive analysis of staring and its effect on all who are othered from society, the analysis she offers specifically pertaining to disability is essential to the discussion of the representation of individuals with disabilities. She categorizes the different types of staring and the subsequent relationships between the staree and starer—the starer, most often those that meet all the standards of society’s definition of normalcy, and the staree, those who do not qualify as “normal,” often people within the disabled community.

Though Garland-Thompson addresses several negative forms of staring in her book, her analysis of “domination staring” is most applicable here. She explains:

In humans, a stare can also communicate social status, conferring subordination on a staree and ascendancy on a starer. . . But human dominance staring is usually more complex than chest pounding. Often staring as a manifestation of dominance veils aggression with a restraint enabled by the hierarchy the staring
enacts. In other words, a harsh stare can do the work of a foot on the neck because the subordinate accedes to the system of domination that is in place.\(^{51}\)

The “system of domination”—in this case ableism and stigma surrounding disability—is what empowers the starer. As we have seen in the previous two chapters, negative staring can have dominance over the starees. Freak shows and their photographers capitalized off of able-bodied society’s fascination and the system of domination in place to exploit those with disabilities. In such a system, even well-meaning artists—such as Diane Arbus, who is often praised as being a hero of diversity—exploit and enable negative staring of individuals with disabilities, thus perpetuating the problem.

Along with negative staring, however, exists a paradox. Garland-Thompson calls it ‘looking away’:

Consequently, our eager stares often quickly shift to uncomfortable looking away. Our ocular id, in other words, jerks our eyes toward a stimulating sight and our ocular super-ego guiltily retracts them. We may withdraw a stare in simple deference to propriety or parental prohibition. Charges of rudeness further encourage us to cut and run. Sometimes, however, truncated stares come from our distress at witnessing fellow humans so unusual that we cannot accord them a look of acknowledgment. To be suddenly confronted with a person extraordinary enough to provoke our most baroque stares withers our ready curiosity and we turn away, snuffing out the possibility for mutual recognition. If the knowledge that staring delivers is unbearable, the expected elasticity of human connection that mutual looking offers becomes brittle. When we suddenly find ourselves face to face with some momento mori or our most dreaded fate—we look away.\(^{52}\)

Looking away is equally as harmful—if not more so—as negative staring. Rather than objectifying or capitalizing, however, looking away denies one's disability and humanity. She continues, “looking away is an active denial of acknowledgment rather than the


\(^{52}\) Ibid., 79.
tacit tipping of one’s hat to ordinary fellow citizens expressed in simply not noticing one another. Looking away is for [an individual with a disability] a deliberate obliteration of [their] personhood.\textsuperscript{53} Thus, in social settings, those within the disabled community are underrepresented and often excluded.

For the past several decades, however, there has been an increase in the discussion of disability and its relation to society and humanity. Questions pertaining to disability have been raised in academic works across many different disciplines: what does it mean to be human? What qualities of human life should be valued? How do we, as a society, ethically respond to difference?\textsuperscript{54} “Much of the new work,” Catherine Kudlick explains in her article, “Disability History: Why We Need the ‘Other’,” “springs from disability studies, an interdisciplinary field dating from the mid-1980s that invites scholars to think about disability not as an isolated, individual medical pathology but instead as a key defining social category on par with race, class, and gender.”\textsuperscript{55} In her book, \textit{Claiming Disability: Knowledge and Identity}, Simi Linton offers another helpful definition of disability studies:

Disability studies takes for its subject matter not simply the variations that exist in human behavior, appearance, functioning, sensory acuity and cognitive processing, but, more crucially, the meaning we make of those variations. . . It is an interdisciplinary field based on a sociopolitical analysis of disability and informed both by the knowledge base and methodologies used in the traditional liberal arts, and by conceptualizations and approaches developed in areas of new scholarship.\textsuperscript{56}

\textsuperscript{53} Ibid., 83.

\textsuperscript{54} Catherine J. Kudlick, “Disability History: Why We Need Another ‘Other.’” \textit{The American Historical Review} 108, no. 3 (2003): 764.

\textsuperscript{55} Ibid.

Along with the rise in the discussion of disability and all it entails, there has also been an explosion of the discussion of disability in political and advocative fields. Individuals with disabilities have forced the able-bodied community to acknowledge their existence. Though there is still much progress to be made, there have been many legal and societal advances toward accessibility for and inclusion of the disabled community in American society over the past several decades.

Yet, for some reason, art—its history, study, and practice—has not kept pace with society’s growth in disability studies. Despite “art education’s longstanding social justice theory and advocacy. . . disability—among the most widely and severely oppressed human conditions—has been excluded from that tradition. Despite [the call] for a socio-political orientation of disability two decades ago, and the flourishing of disability theory and activism that supports [that call], the field continues to resemble orthodox special education discourses that largely ignore the first-hand perspectives of disabled students, teachers, researches, artists, and others.”

Disabled individuals have been and continue to be excluded from artistic spaces. Thus, art—its theory, history, and practice—has failed to properly and ethically include and represent individuals with disabilities. In some ways, art history has corresponded with Garland-Thompson’s analysis and utilized the culture of staring to capitalize off of those with disabilities: photographers such as Charles Eisenmann and Diane Arbus are prime examples. Yet at the same time, art history— with its underrepresentation and exclusion of those with disabilities—also falls under Garland-Thompson’s categorization of looking away from those with disabilities, denying their humanity. Disabled artists “continue to be wary of

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entering, lest they be cast as overcoming savants,” and able-bodied artists either fail to
include people with disabilities in their artwork or fail to do so ethically.\footnote{Petra Kuppers, “Diversity: Disablility.” Art Journal 75, no. 1 (2016): 97.}

The problem, then, is this: we, as an able-bodied society, have simultaneously
ogled at and capitalized off of individuals with disabilities yet failed to acknowledge their
humanity alongside their disability; we need to look at individuals with disabilities—to
include them in our social spaces, to acknowledge their humanity— but, with such a
negative history surrounding our treatment of the disabled community, we, as an able-
bodied society, do not know how to do so ethically. Portraiture, if done rightly, provides a
space for ethical looking at individuals with disabilities. It enables an able-bodied
audience to encounter disability, unencumbered by the historical sense of ‘otherness,’
and forces them to reconcile their stereotypical views of disability with the human being
looking back at them from the canvas. The philosophical notions behind ethical
portraiture—specifically of disabled individuals—can be lengthy. To be both concise and
practical, I will discuss the physical steps I took as an able-bodied artist to ensure
ethical portraiture of individuals with disabilities.

In painting a portrait of anyone, there is a moral component that is not present in
self-portraiture. Not only must the artist be conscious of the great honor it is to represent
someone else to the world, but they must also be conscious of the history and
stereotypes of the individual they are painting. As an able-bodied individual, I do not
know or pretend to know what it is to have a disability. Though I hold deep love and
respect for the friends that I have painted in this project, I recognize that they are not
only different from me in ability, but in the way that they experience the world and
subsequently the way in which the world experiences them. With the project, I sincerely
hope it is clear that my goal is not to speak for the individuals that I am painting nor to
speak to the disabled experience. I recognize that as an able-bodied person, my art will
never add what a disabled artist’s work will to that conversation. Rather, my goal with
this project is to create a bridge for my viewers—a space that closes the gap between
the simultaneous staring and looking away, the unwarranted attention and exclusion,
and the rejection and fascination with individuals with disabilities. Garland-Thompson
explains of portraits of individuals with disabilities:

They grant us more than permission to stare; they use the clout of high art to
transform our staring from a breach of etiquette or an offensive intrusion into an
act of appreciation. These portraits enable visual pilgrimages of deliberate
contemplation that might be scuttled in a face-to-face encounter on the street.
The invitation to look that a portrait offers precludes our skittish staring and
instead allows us to look deep and long into these unfamiliar faces made
strangely familiar.59

Not only do I hope this project offers a space where people with disabilities can be
viewed with respect, dignity, and love by able-bodied viewers, but I also hope that it
challenges able-bodied stereotypes and standards that have been upheld in art and
society. Garland-Thompson goes on in a quote that is most essential to the heart of this
project:

Conferring dignity on people whose differences draw stares is the challenge to
which these portraits of disabled people rise. These portraits intervene between
starees and starers to offer respectful, even beautiful, pictures of people we
have not learned to look at in this way. They revalue devalued people, the kinds
of people most of us have only glimpsed in institutions or in medical pictures with
black boxes over the eyes. This anonymity that medical photographs impose on
a staree also prevents the person pictured from staring back at the viewer. . .

Rush’s portraits rework the way we usually stare, however. They keep us looking rather than looking away.\textsuperscript{60}

Because I was immensely conscious of the negative history of misrepresentation and underrepresentation of individuals with disabilities, I chose individuals with disabilities whom I already knew and had developed a relationship with. I felt that, out of everyone I could have painted, I knew these three individuals best and could therefore render the most holistic portrait of them. From the beginning of the project, I knew that I wanted the subjects of the paintings heavily involved in the creative process. Thus, I interviewed each individual, picked the reference photo with them and their family, and painted the qualities, interests, and traits that they wished me to portray. Though the details of each relationship, interview, and creative process will be discussed in the following chapter, I will offer a brief overview as it pertains to the present discussion.

For the first portrait, I painted Faith. In knowing her for over a year when the project started, I knew she loved tattoos, but it was after interviewing her that I earned that her tattoos were a way in which she displayed her personality. Faith’s arms, embroidered with colorful references to her favorite books, quotes, songs, and more, are the subject of my second portrait. The second portrait-sitter is Zack, whom I had also known for over a year at the beginning of this project. Through our friendship, I noticed that Zack always carried a Woody doll with him, but it was through our interview that I learned that Woody is the only toy that Zack has ever played with. Woody has become a part of Zack’s family and a friend to Zack’s friends. Though tattered and stained, Woody is an integral part of Zack’s life and personality and is the focus of my fourth image. My third subject is Kevin. I have been friends with Kevin the longest of all,\textsuperscript{60} Ibid.
almost four years. Though I knew Kevin loved disc golf (I have been beaten by him several times), it was through the interview that I learned how much it means to him. His lucky disc is one that he made a hole-in-one within the Redwood forest, and it is the central point of the sixth painting.

The next practical step I took to ensure ethical portraiture of disability is the style and technique with which I painted the portraits. I called upon classical portraiture—its softness, realism, and color—to paint these six portraits. Conscious of the history of exploitive and ‘Freak Show’ portraiture, I sought to capture each individual as unposed as possible. For each, I interviewed and photographed them in their own home, to encourage them to feel comfortable, in control, and unposed.

Though it cannot supplement face-to-face interaction between able-bodied and disabled individuals, portraiture provides a space for able-bodied people to ethically gaze at people with disabilities, and it opens the door to destigmatizing disability. With this project, I hope to have created such a space.
Chapter Four

The Portraits

The purpose of this chapter is to introduce the individuals portrayed in this portrait series, thus I will discuss our friendship, our interview, and how these culminated in my portraits of them. I want to be explicitly clear that I recognize I am relaying the experiences of another individual, who has experiences much different than mine. The experiences of Zack, Faith, and Kevin—their lives, personalities, interests, and disabilities—are not my own. I recognize with great honor and care the privilege it is to share another person’s story. I also recognize that I will not attempt to completely understand Zack, Faith, and Kevin’s experiences. Rather, I hope to share and portray—through our friendship and experience together—each person as authentically as I have come to know them. In the interviews, I asked about each person’s favorite thing about themselves, what they would people to know about them when they see the paintings. It is crucial to note here the importance of the role of a caregiver in communicating. Zack, who is limited in his verbal communication, is aided by his mother, who knows him better than anyone, in answering the questions.

I first came to know Zack a little over a year ago, as he began working at the local coffee shop Bitty and Beau’s in Waco where a close friend of mine was the manager, but Zack and I also participated in YoungLife together at Baylor. Through these, I came to know Zack, his sense of humor, his love for his toy Woody, his competitive spirit, and so much more. Zack has Down syndrome, so I cannot speak to his experience as an individual with a disability, but I can attest to his genuine friendship that he has offered to me.
As I sat with Zack and his mom at their kitchen table in September for our interview, Zack welcomed me as if I was one of his family. After I explained the project, we talked about his favorite drinks to make at Bitty and Beau's, what he planned to do for the rest of the day, and finally about the paintings. When I asked Zack about his favorite thing about himself, he and his mom communicated that Zack's ability to love and welcome all people is one of his best traits. He also communicated that Woody—his battered Toy Story doll—was one of his favorite things about himself.

After some thought, I realized how much those two aspects of Zack went hand-in-hand. Anyone who has come to know Zack, has come to know Woody, who travels with him everywhere. To Zack’s friends, Woody has become an integral part in their relationship with Zack. When we would play games at YoungLife events, Woody would compete alongside Zack, sometimes even competing in the games as an individual player—one of Zack’s staple comedic bits was whenever we played freeze dance, he would try to blame Woody for his own moves after the music had stopped. To all of Zack’s friends, Woody was associated with Zack, his personality, his joy, his acceptance, his love, and his laughter. Thus, Zack, his mom, and I felt that Woody was one of the defining qualities of Zack and deserved to be present in his portrait.

As we stepped on the front porch to take some reference photos, Zack began laughing at something his mom was saying, so I quickly pulled out my phone to capture the moment. The photo we chose shows Zack (figure 43), giggling so much his eyes are barely open. I then began taking some photos of Zack holding Woody. Again, Zack was focusing on his mom, absentmindedly fiddling with Woody as they conversed. The result was a renaissance-esque photo—the way that Zack’s fingers softly cradled Woody
remind me of hands depicted in Da Vinci or Michelangelo’s artworks (figure 44).

Figures 43 and 44. Reference photos of Zack.

Like Zack, I came to know Faith through her job at Bitty and Beau’s. The summer before the opening of the coffee shop, I volunteered at the hiring fair, where I met Faith for the first time. Despite the stress of a job interview and the hectic environment, Faith introduced herself to me and spoke with me for several minutes. She was kind, personable, and excited to be present. Faith has cerebral palsy and has lived her life with a wheelchair as an extension of her body. Thus, I do not claim to understand her experience or speak for her, I merely hope to share the glimpse of Faith that I have gained through our relationship.

Ever since our first meeting, Faith has exhibited nothing but kindness and friendliness towards me. As our friendship grew from our first meeting in the summer of 2021, I have come to know Faith, her love of music, her creativity, her sense of justice,
and her venerability. Because Faith is extremely open with her story, her struggles, and her pain, she has been able to uplift and encourage others.

In April, I sat with Faith in her bedroom for our interview. After catching up and after discussing the scope of this project, we began talking about Faith and what she loved about herself. Among many things, her favorite quality about herself is her tattoos. For the past several years, Faith has utilized her arms as a space to express her personality and share bits of her story. Studded with song lyrics, verses, plants, animals, and much more, Faith’s two arms have become a canvas. As we sat in her room, she pointed to several, not only explaining what each meant to her, but what she was feeling at the time she got them. She explained that many of her tattoos are reminders of her strength, of the trials and pain that she has pushed through. Though I had known Faith for almost a year at the time, it was one of our most intimate conversations. We both knew that Faith’s tattoos must be included in her portrait.

As Faith was telling me about her tattoos, I asked if it was alright if I started taking photos while we talked. Among the many I took, the final reference photo displayed many of her favorite and most colorful tattoos, but most notably, it exhibits her namesake piece, “keep the faith” (figure 45). After Faith went through each of her tattoos, she sat for the portrait. Natural and authentic, Faith laughed in one of the first photos I took. That one, we decided, would be the final reference for the portrait (figure 46).
Of all the three people painted, I have known Kevin the longest and therefore know him the best. During my first year of college, I met Kevin at a Best Buddies party, a student-lead organization that seeks to build friendships between college students and those within the disability community. It was a Halloween party, and Kevin—well over six feet—was dressed in a perfect replica of a WWII general. When we met, he gave me one of his tight hugs, and from then on we were friends. Kevin has an intellectual disability, so while I will not attempt to articulate his experience as a disabled individual, I will attest to the friendship that he has shared with me and do my best to share the person I know Kevin to be.

Over the course of my four years at Baylor, my friendship with Kevin has continued to grow. Despite the fact that we had to meet on zoom over the course of the pandemic,
it was Kevin who was the first to join and the last to leave every Zoom call. Aside from Best Buddies, we have played many rounds of disc golf and putt-putt, two of Kevin’s favorite activities. Since I met Kevin for the first time in the fall of 2019, I have heard from him every single week. He has never failed to reach out to see how I have been and what I have been doing. With each call, he consistently signs off with the question, “when are we hanging out next?” I have been blessed by Kevin’s exuberant joy, his deep love for his friends, and his unending care for those around him.

As I sat with Kevin and his mom in their home, we discussed the project and what it would look like. Then, I asked Kevin what his favorite thing about himself was, what he wanted people to know about him when they saw his portrait. Immediately, Kevin got up and went into his room. His mom and I followed and watched as he lugged his large disc golf backpack out of his closet. He placed it on his bed, unzipped it, and decisively slid out his favorite one, a peach disk with a cheetah across the front. It seemed clear to me and Kevin’s mom that he wanted viewers to know about his disc golf. After snapping some photos of Kevin holding his disc, he pulled out his phone and showed me a video of him making a hole-in-one in the redwood forest with that very disc, his best shot to date. As Kevin showed me the video, he was bursting with excitement.

Similar to the relationship between Zack and his doll Woody, Kevin and his disc are a recognizable pair. Any of Kevin’s friends know that he is a very skilled disc golfer—many of them know from personally losing to Kevin’s far throws and hole-in-ones. They also know the joy that disc golf brings him. As an opportunity to be around people, disc golf has enabled Kevin to grow even closer with his friends. Even though I am terrible at disc golf, Kevin sees our matches as an opportunity both to deepen our
friendship. As we walk around the course, he will ask me about my friends, tell me about his family, talk about the trips he is excited about, and ask me about music that I like. One of Kevin’s discs is a reminder of the intentional friendship that he displays to all around him. Being friends with Kevin is refreshing—the humor, genuine interest, and kindness that he has exhibited toward me throughout our friendship have made him one of my most cherished friendships from college. After discussing it with him and his mom, I decided that Kevin and his disc will provide an authentic narrative of the Kevin I have been so honored to know.

Kevin and I stepped out onto his front porch to take some photos, and I realized I had to stand up on my tiptoes to achieve the right angle because Kevin is much taller than I am. He posed for a few with a soft smile, but soon laughed, showing an open-mouthed grin. After sending a few to his mom, we decided on the one that captured his laugh (figure 47). Then, Kevin posed with his disc, his hands instinctively holding it about the edges, turning the cheetah-clad face towards me (figure 48).

Figures 47 and 48. Reference photos of Kevin.
Though I sought uniformity in each of the six portraits in color and style, I was careful to still exhibit each individual as distinctive. Their personalities, interests, hobbies, along with their disabilities have enabled them to offer genuine friendship to those around them, myself included. I have been deeply honored to not only know each person but to be entrusted to share a glimpse of who they are.
Figure 49. Natalie Stitt, Zack, 36 by 48 inches, oil on canvas.
Figure 50. Natalie Stitt, *Zack and Woody*, 22 by 28 inches, oil on canvas.
Figure 51. Natalie Stitt, *Faith*, 36 by 48 inches, oil on canvas.
Figure 52. Natalie Stitt, *Faith and Her Tattoos*, 22 by 28 inches, oil on canvas.
Figure 53. Natalie Stitt, *Kevin*, 36 by 48 inches, oil on canvas.
Figure 54. Natalie Stitt, *Kevin and His Disc*, 22 by 28 inches, oil on canvas.