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Critical condition: Alzheimer's and identity in Carla Subirana's Nedar (2008)

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This article reads Carla Subirana's documentary film Nedar against a background of panic in the mainstream media (in Spain as in the United States and the United Kingdom) about Alzheimer's disease and the apocalyptic demography presupposed by an aging population. It considers the history of Alzheimer's disease and diagnosis to propose that while technologies for the medical representation of the disease are becoming globalised, cultural responses to experiences of the disease are more distinctive. The article queries the use of illness as metaphor in Nedar and suggests that Alzheimer's, dementia, and amnesia are used to portray the irretrievable loss of information about personal and collective experience in Civil War and post-war Spain. The article details the film's juxtaposition of illness and archives within an exploration of the debate articulated by Susan Sontag and Nancy Scheper-Hughes over the appropriate use of disease metaphors: Alzheimer's complicates this debate because patients are sometimes unable to tell their own stories. The representation of amnesic disease is further complicated in a Catalan and Spanish context by a history of politically sanctioned forgetting and by the culturally specific value of memory. The article concludes by indicating that Subirana succeeds in using the connotative value of illness metaphors whilst also reaching an accommodation with her relatives' illnesses which differs from the practice of forgetting memory advocated by some of those who work with Alzheimer's patients in the English speaking world. It is also argued that unlike many other narratives of Alzheimer's disease, Nedar avoids recourse to the genre of horror, an eventuality which is in part facilitated by the film's origins in an investigative memorial project.

Keywords: Carla Subirana; Nedar; aging; Alzheimer's disease; historical memory; illness as metaphor

Introduction: Alzheimer's, apocalyptic demography, and the hypercognitive

Alzheimer's disease is indisputably a condition which demands critical attention in the sense that it has emerged recently as one of the diseases which, in the

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developed world, is especially prominent in the public awareness of incurable pathologies. Representations of Alzheimer's, such as Richard Eyre's *Iris* in 2001, a film based on a series of pathographic memoirs written by John Bayley, and campaigning by celebrity patients like Terry Pratchett, have further added to the profile of the disease in the United Kingdom. In the United States, similarly, *The Alzheimer Project* (Hoffman 2009), a series of television documentaries shown through HBO in 2009 and fronted by Maria Shriver, has further heightened the profile of Alzheimer's within the mainstream media. Using fear as a nosological index, each instalment of *The Alzheimer Project* began with a graphic which informed the audience that Alzheimer's is the second most feared disease in America after cancer (see Figure 1). The HBO website which accompanied the series tells readers that

One of the most devastating forms of memory loss is Alzheimer's disease, an irreversible and progressive brain disorder that slowly destroys memory and thinking skills. Today, Alzheimer's is *the second most-feared illness* in America, following cancer, and may affect as many as five million Americans. ('About the Project' 2009, emphasis added)

The appeal to fearsomeness invites viewers to recognise the impact, seriousness, and epidemic qualities of a condition with a spectrum of symptoms, both medical and physical, which evolve over years and may be different for each individual; the protean nature of Alzheimer's challenges the simplistic disease-of-the-week format often adopted by television in its portrayal of illness and this was reflected in the fact that *The Alzheimer Project* ran over four episodes and was supported by additional online content and supplementary materials.

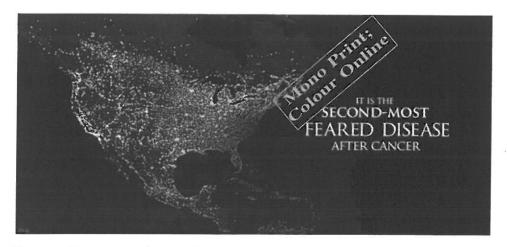


Figure 1. Screen capture from opening sequence of The Alzheimer Project.

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As scholars of medical anthropology like Ann Robertson have noted, the representation of Alzheimer's as a dread condition usually goes hand in hand with a discourse of apocalyptic demography so that the threat posed by the disease is exponentially magnified by the greying of society in developed nations:

The prevailing belief is that an increasing aging population means increasing demands on the resources of society, including health care resources [...]. According to this scenario people will live longer but sicker [...]. This greater morbidity of increasing numbers of elders, so the argument goes, will drive up health care costs. This 'bankruptcy hypothesis of aging' in which 'oncoming hordes of elderly' [...] deplete national health care budgets constitutes 'apocalyptic demography'. (1991, 135)

Using diagnostic terminology from medicine, we could say that Alzheimer's discourse grafts together as co-morbidities the perceived threat of an aging society and the perceived fearsomeness of dementia. Robertson argues that the expansion of Alzheimer's diagnosis is a response to what she terms the biomedicalization of old age, an interpretive shift which allocates to a pathology contained within an individual body the wider problems and demands posed by an aging population: 'By framing the "problems of aging" and, therefore, the solutions as biomedical, this view [...] ignores nonmedical issues – such as poverty, isolation, the loss of role and status – and thus, [it] effectively depoliticizes the problems of aging' (1991, 136). The aging of society in developed countries has coincided with the apogee of what Stephen Post calls a hypercognitive culture, a coincidence which means that 'nothing is as fearful as AD because it violates the spirit (geist) of self-control, independence, economic productivity, and cognitive enhancement that defines our dominant image of human fulfilment' (2000, 245).

When we turn to the United States National Institute on Aging's (NIA) preamble to its work on Alzheimer's, Robertson's contention about the comorbidity of apocalyptic demography and dementia is graphically borne out. The NIA contextualises the social impact of Alzheimer's by saying:

According to recent estimates, as many as 2.4 million to 5.1 million Americans have AD. Unless the disease can be effectively treated or prevented, the number of people with AD will increase significantly if current population trends continue. That's because the risk of AD increases with age, and the U.S. population is aging. The number of people aged 65 and older is expected to grow from 39 million in 2008 to 72 million in 2030, and the number of people with AD doubles for every 5-year interval beyond age 65 [...]. In the years to come, AD is expected to pose physical and emotional challenges for more and more families and other caregivers, in addition to those with the disease. The growing number of people with AD and the costs associated with the disease also will put a heavy economic burden on society. (NIA 2010)

It is noteworthy here that the elderly are doubly stigmatised. Transgressive in the first place by dint of fitting uneasily into a work based ethic where non-

productivity meets with suspicion and disdain, senior citizens now pose an even greater threat by virtue of their ever increasing vulnerability to an economically and socially draining dementia. As Elizabeth Herskovits observes: 'The Alzheimer's Disease construct has contributed to a monsterizing of senility such that the present day lived experience of senility and aging has become more unpleasant and horrific than it was previously' (1995, 153).

Just as many of the mechanical technologies for diagnosing Alzheimer's have been transplanted wholesale from the English speaking world to Spain, so too, it seems, has the discourse of apocalyptic demography in which the future of cerebral grey matter and society's senior or grey sector is couched. For example, the Fundación Alzheimer España (FAE) replicated the rhetorical device which conjoins an explosion in numbers of elderly people with a newly dreadful disease:

Afecta al 5-7% de las personas de más de sesenta y cinco años. Cerca de 650.000 personas están afectadas en España y se manifiestan más de 100.000 nuevos enfermos al año. Si se estima el número medio de miembros de una familia a 4, son más de dos millones las personas que ven su vida trastornada por la enfermedad. Es la causa de invalidez, dependencia y mortalidad más frecuente en los mayores. Teniendo en cuenta el envejecimiento de la población y el futuro incremento de personas mayores de 80 años, se prevé que el número de enfermos se duplique en 2020 y triplique en 2050. (FAE 2010b)

To represent the disease and its social consequences graphically, the Fundación uses the image of an iceberg, the massive threat which comes from nowhere and which without sufficient vigilance will shipwreck the unwary national vessel. This is not the first time that a national health campaign has used an iceberg to represent the threat posed by a disease. In the United Kingdom, a government-sponsored awareness campaign used a similar graphic to depict the threat of AIDS in 1986. In Spain the FAE not only builds Alzheimer's up into an apocalypse of pathological demography but also emphasises that among European countries Spain is likely to be the most severely affected because of the combination of its low birth rate and rapidly aging population (FAE 2010a).

As in the United States and the United Kingdom, there has also been in Spain an efflorescence of lay representations of the disease and two of the most striking recent films with an Alzheimer's disease theme have been made by Catalan directors, namely Carla Subirana with her first feature-length documentary, *Nedar*, and Albert Solé with *Bucarest: la memòria perduda*, both released in 2008. Although both of these films are autobiographical and documentary, there is a precursor to their combination of pathography and family history in David Trueba's 2003 film adaptation of Javier Cercas's *Soldados de Salamina* (2001).

In Trueba's film, Lola Cercas talks to her elderly and infirm father during a visit to his nursing home about the work she has been doing to complete a journalistic assignment on the Spanish Civil War. Looking totally distracted, indifferent, and confused, her father's response is very simple as well as devastating. He asks her:



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'¿Qué guerra?'. Although Soldados de Salamina is not the focus of this article, I include a detail from this scene of memory transmission — or interruption — between generations because my point is that Nedar (and Bucarest: la memòria perduda) can be seen as an exploratory expansion of this moment in Trueba's film where national and social amnesia are fused with the inexorable erasure of memory implied by a construction of old age centred around dementia, something which is arguably an offshoot of the increasing awareness of the extent to which Alzheimer's is affecting the older age groups in Spanish society. And what I propose in the more detailed attention to Nedar, in particular, which follows, is that this equation of the personal and the collective memories produces a nosology of Alzheimer's disease which is specific to Spanish and Catalan discourse.

Representing Alzheimer's and the economy of memory in Spain and Catalonia

I want to suggest that in *Nedar* Alzheimer's disease is not only represented within the discourse of a condition which is critical *for* Spain's social economy, but that it is furthermore part of a discourse which is critical *of* that social economy. Medical anthropologists and cultural historians like Anne Davis Basting (1998, 2009) and Janelle S. Taylor (2008) have pointed to the role of Alzheimer's disease and old age in performing cultural work, and my contention in this article is that Subirana steps away from the more clichéd representations of Alzheimer's disease to uncover some of the functions performed by Alzheimer's, and by people diagnosed with Alzheimer's (PDWAD), in an economy of recall and oblivion. Where does it leave the person diagnosed with Alzheimer's when his or her brain is reified as a fragile store of forensic evidence? Can a metaphorical deployment of illness which has power and usefulness for raising social consciousness of historical memory loss work without prejudice to the individual patient?

Alzheimer's sits at the threshold between mind and body, between the neurological and the psychological, between procedural and propositional memory, between autonomy and dependency views of self and personhood; and, as a closer reading of *Nedar* suggests, this disease whose resurgence in the 1970s was hatched in international symposia and conferences sits in a Spanish context, furthermore, on the threshold between informally sanctioned collective neglect and legislatively binding care for the past.²

Although Alzheimer's takes its name from Alois Alzheimer's description of a link between neurofibrillary plaques and pre-senile dementia, the initial impetus for construction of an Alzheimer's diagnostic practice in the first decade of the twentieth century came from other clinicians who wanted to counter psychoanalysis by proving an organic basis for aberrant behaviour. From the outset, then, the disease diagnosis has constructed a patient centred, individualised, and contained account of malaise in contradistinction to an

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interpersonal or systemic reading of dementia.³ Drawing on Karen Lyman (1989), Robertson suggests that

The ready acceptance of the clinical construct of AD can be attributed to its efficacy in creating order out of chaos both for clinicians and families; a 'disease' construct serves to normalize and render manageable the 'disturbing' and 'disruptive' behavior of the PDWAD. The 'biomedical model of dementia', ultimately fails to take into account any notion of the extent to which AD may be socially produced and reproduced. (1991, 142)

The metaphorical values afforded Alzheimer's disease in recent pathographical films made by directors from Catalonia are thus engaged with a form of narrative which grafts as metaphor for a collective amnesia an illness defined in part by a history which resists inter-subjective meaning. Subirana's act of de-personalising the illness and of reading it across a social history is in itself a political gesture. But, at the same time, we must ask what happens to the illness in the midst of these various projects for using illness to achieve semantic engineering. In order to approach these questions it is helpful briefly to step back and to consider the genesis of Subirana's Alzheimer documentary.

Investigating memory

Some of the footage in Nedar was shot as much as eight years before the film was released. The director took pictures of her grandmother to practise for her degree and, as she has said in interviews, did not set out either to make a film about the Civil War or about Alzheimer's (Semana Internacional de Cine 2009). The film was to chronicle an investigation into the missing data in her family's biography and she did not plan to feature in it herself. The focal point would be Juan Arroniz, Subirana's late maternal grandfather, and the project was intended to retrieve some historical information about a man who had disappeared not only physically but who had also been written out of the family's history just as the political elements with which he may have been linked were written out of the country's history by the victors in the Civil War. Among the few facts Subirana knows about her grandfather before her investigation is that he was executed in 1940 at Camp de la Bota. Her grandmother has spoken very little of him and the pared down family of three women seems to have reached within the domestic sphere an agreement not to mention him, echoing in this silence the wider pacto de olvido which had, supposedly, expedited Spain's transition to democracy. The film which was eventually to become Nedar started out as a project meant to document and to unravel this structure of silence. Subirana says in her narration towards the beginning:

A casa era molt natural no parlar sobre el meu avi; potser l'absència del meu pare pesava massa així que mai em vaig preocupar per aquesta altra absència. L'únic que sabia era que l'havien afusellat el 1940, com a tants d'altres. Volia esbrinar qui era aquell home i per què va morir afusellat.

In the course of this project to retrieve memory, however, both of the director's two immediate family members are diagnosed with Alzheimer's, a condition which erodes precisely the resource around which the film - as it had been originally conceived - was hinged. Subirana's grandmother, Leonor, is in her 80s when the filmmaker embarks on the project and very soon after is diagnosed with Alzheimer's. Close on the heels of Leonor's death, Subirana's mother, Ana, who is in her 60s, is also diagnosed with pre-senile dementia. Thus, as the filmmaking project was unfolding, decisions about what to forget and what to remember shifted from being entirely elective to being determined in part by biology and the steady progress of a disease. In the course of the documentary we see that for Leonor and Ana – Subirana's two most immediate sources of information – the withholding of information (or simply the preference not to remember certain elements of family history) is no longer a matter of habit or of expedience: their memories are being shredded by the implacable progress of a dementia which is indifferent to changes in the political landscape. As in Bucarest: la memòria perduda, Nedar's account of the decimation of the individual capacity for memory dramatises the loss of historical memory. It does this in the framework of a documentary which had initially been conceived as a narrative whose piecing together of the remaining fragments of information about little known family members would contribute to the retrieval of a bigger picture of a political and national history, one eroded by neglect and the deliberate decisions to forget taken in the 1980s. In the project as originally conceived, specific memories of individuals would have contributed towards the building of a more inclusive collective memory; in the project as it was reshaped by Alzheimer's, the incomplete or partial collective memory corresponds not only to the specific memories of individuals which can no longer be salvaged, but also to the loss of the ability to remember per se.

Archiving the mind

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300 301 At a point in the narrative where it becomes clear that Leonor's dementia has advanced to a point where it is divesting her not only of her memory but also of her individuality, Subirana observes: 'A mesura que el meu avi es dibuixava en la meva imaginació, la meva àvia anava desapareixent'. The recovery of memory and cognitive impairment are here explicitly allied, and, arguably, the threat to the project of retrieving a record of the past from oblivion becomes part of what Nancy Scheper-Hughes calls 'the disease double': 'Societal and cultural responses to disease create a second illness in addition to the original affliction [...] the layers of stigma, rejection, fear, and exclusion that attach to particularly dreaded diseases' (1986, 137). The editing of the film reinforces the metaphorical equation which conjoins the individual memories of Subirana's relatives with archival documentation and other information. For example, in one of the last scenes shot before Leonor's diagnosis, Subirana says, in reference to her grandmother's confused and uninformative answers to her questions about her grandfather:

Aquell dia em vaig adonar que era com si ella em digués: 'A mi no m'ho preguntis. Si vols saber, ara et toca a tu'. Poc després, la meva àvia es va posar malalta. L'informe mèdic deia: 'Presenta un dèficit cognitiu greu amb síndrome afàsica, apràxica, anòsica, amnèsica important'. La veritat és que no hi entenia res.

Visually, the film marks this turn in the narrative by juxtaposing a scene of Carla comforting her grandmother and stroking her temples with another scene of the stacks at the Arxiu Nacional de Catalunya (Figure 2). The implication is clearly of parity between an individual's brain as a storehouse of information and an archive as a documentary extrapolation synthesising the sum of such individualised recollection. It is almost as if Alzheimer's provided the solution to the question Cercas's metafiction posed about the facticity of history: seen through the discourse of Alzheimer's, history is neither fact nor fiction but an organic corporeal network of synaptic activity. The privileging by Alzheimer's of a concept of the brain as a mechanical electrostatic computer of flesh lends itself well to this equation, and here Leonor's illness is inscribed by the narrative not so much within an apocalypse of demographic massification as one of cultural death and extinction, a trope which echoes the title of Carles Casajuana's (2009) recent novel, L'últim home que parlava català. Each member of the community who succumbs to dementia is like another folder in the archive of national and cultural identity being destroyed, and since dementia threatens precisely autonomy, recognition, sovereignty, and independence, the diagnosis also lends itself to being a metaphor for a stymied and frustrated self governance of a community of individuals.

Since there are several examples in *Nedar* of elision between representations of people with Alzheimer's and historical documentation, the deployment of this metaphor is clearly intentional. For example, Subirana edits together two scenes which both involve the consultation of a graphic illustration from a test



Figure 2. Juxtaposition of two screen captures from *Nedar* illustrating shift from Leonor as patient with memory loss to interior of stacks as collective memory.

determining the identity status of one of her relatives. The first of these two scenes sewn together illustrates Ana undergoing the MRI scan which allows doctors to confirm her mother's diagnosis with Alzheimer's from the visual representation of her brain tissue which it produces. The second scene illustrates Subirana herself consulting her grandfather's criminal record which includes a microfiche reproduction of his fingerprints. The image we see on the screen shifts from a brain with signs of the amyloid plaques and taut angles characteristic of Alzheimer's to a document bearing a fingerprint of the patient's father, a man she never knew (Figure 3). Just as her grandmother disappeared when her grandfather was forming some kind of shape in her mind, Carla's mother's brain is being eroded in the same frame in which her father has left a digital trace of presence. The paternal stories which for so long remained unknown seem to fill the gaps being left by the onset of the disease, as if the recovery of historical memory were both threatened by and a compensatory symptom of the loss of individual memory.

The memory loss suffered by Ana and Leonor unfolds against the background of the introduction of legislative obligation to remember. The 'Ley por la que se reconocen y amplían derechos y se establecen medidas en favor de quienes padecieron persecución o violencia durante la Guerra Civil y la Dictadura', better known as the 'Ley de memoria histórica', came into effect in 2007 and indeed Memorial Democràtic, which supported the production of *Nedar*, is governed by a statutory obligation to recover and disseminate facts and information pertaining to the time between the Civil War and the early 1980s.⁴

Almost coetaneous with the 'Ley de memoria histórica' another law which has received less attention was passed in Spain in 2006, the 'Ley de promoción de la autonomía personal y atención a las personas en situación de dependencia y a las familias', better known as the 'Ley de dependencia'. I would argue that *Nedar* can be seen not only against the backdrop of the legislative obligation to remember, but also through the prism of the other law which enshrines the obligation to take

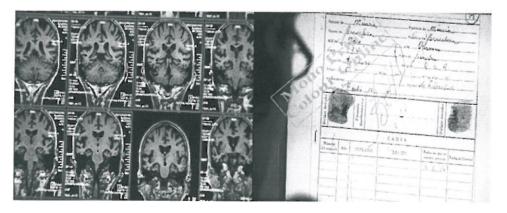


Figure 3. Juxtaposition of two screen captures from *Nedar*: MRI images of Ana's brain (left) and a document from Juan Arroniz's criminal record illustrating his fingerprints (right).



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care. Taking care of memory becomes for the director a way to take care of her mother and her grandmother, and inasmuch as Alzheimer's affects not only the patient but the family, the effort to recover what her relatives did not know or could not know becomes part of the therapeutic response to the disease. Unlike some of the sons and daughters in *The Alzheimer Project* who are depicted by the series as being primarily concerned with their parents' individual suffering, Subirana does not focus exclusively on loss but also on what remains to be discovered. And yet, as illustrated by the editing alongside each other of scenes of personal loss and the documentary repositories of historical record, the individuated pathographies of Subirana's relatives are nevertheless stitched into a narrative which also addresses the loss of memory on a larger scale.

Alzheimer's and PDWAD as metaphor

Is the appropriation of the suffering of people diagnosed with Alzheimer's disease as a metaphor which articulates a debate about the recovery of historical memory a violent or insensitive act? Could it be held that when someone is suffering from memory loss it is prejudicial to him or her to be burdened furthermore with the weight of remembering for an entire nation or people? Some consideration of the rather polarised views on the metaphorical exploitation of illness can help to answer this question. While Susan Sontag () was opposed to the use of illness as metaphor, others, such as Nancy Scheper-Hughes, have critiqued this gesture as one which 'would have us - patients all on one level - retreat into the safe haven of radical materialism offered in scientific biomedicine' (1986, 138). Scheper-Hughes goes on to suggest that Sontag's opposition to metaphorical value for pathology produces 'the reification of illness (as disease alone) [and] is a form of self alienation, of false consciousness' (1986, 138). The question is further complexified by the fact that Alzheimer's strikes precisely at the ability to self narrate so that patients become what some theorists of life-writing have described as unautobiographical subjects.⁵ One symptom of the disease is indeed anosognosia, an effect whose name is a lexical compound which combines disease and knowledge only to negate both with its initial alpha privative; at a certain stage in the disease some persons diagnosed with Alzheimer's no longer know that they are ill and nor can they quantify how ill they are. Alexithymia. another symptom, curtails the ability to describe or interpret feelings, emotional or physical. We can only speak of a disease affecting someone with anosognosia, then, if we understand the ailment to be one that is recognised intersubjectively, and since the patient cannot report his or her experience of the disease, it can only be known by proxy. And how else can we represent someone else's suffering, which we can never know existentially, without recourse to metaphor? The same inability to describe illness suffered by an Alzheimer's patient creates the obligation on the part of anyone involved in advocacy for him or her to use metaphor if that person is to try to convey the patient's condition without the

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benefit of knowledge which is impossible even for the patient him or herself to access. As Scheper-Hughes writes:

We are without a language with which to address mind-body-society interactions and so are left hanging in mid air, suspended in hyphens that testify to the radical disconnectedness of our thoughts. We resort to such fragmented concepts as the biosocial, the psycho-somatic, the psycho-social, the somata-social as a feeble way of expressing the complex and myriad ways that our minds speak to us through our bodies, and the ways in which society is inscribed on the expectant canvas for our flesh and bones, blood and guts. (1986, 137)

Rather than seeing Subirana's yoking of illness to historical memory (and the threat of its erasure) only as a political act in reference to the stymied recovery of national memory, then, we could see in it an attempt to use the hyphenation made possible by film editing to negotiate the social alexia which otherwise resists making the connections between mind, body, and society necessary for a non-privatised understanding of illness.

Whereas *Bucarest: la memòria perduda* uses the image of a maze to describe Albert Solé's exploration of his father's Alzheimer's and of his country's amnesia, Subirana's film returns at crucial points in its narrative to the image of the swimming pool. Between interviews and between discoveries about her grandfather, the camera follows her swimming lengths underwater, as if her body were tracing links between the nodes in the story through the discourse of medicine and politically constrained knowledge in which we are all awash. The first scene of the director diving into the pool is sewn between opening titles partly comprised of an ultrasound scan and other footage of Leonor pottering about with some plants: at one end of the pool is the scan of a foetus, at the other are images of a grandmother, and the narrator swims between the two the length of the film and vectors the coordinates of a disease which, in the film's portrait, exists socially. In an interview with José David Cáceres Tapia, Subirana spoke at length about her use of swimming and of the pool as metaphors in the film:

El agua es la memoria. Nadar para reencontrarse con ella, un zambullido en busca de unas respuestas imposibles e intangibles... El agua como el primer líquido presente en nuestras vidas, el amniótico, al que vuelvo para reconciliarme con lo más primigenio, con lo más esencial. Es un punto de partida del presente, una necesidad física que se traduce en un medio para simbolizar todo un viaje interior hacia un pasado tan escurridizo como una pastilla de jabón [...]. El agua es también un espacio reservado para el propio espectador, donde inevitablemente su historia se filtra en todas las lagunas que el guión plantea. Un terreno donde la particularidad de la historia se puede convertir en un lugar común. (Cáccres Tapia 2009)

This conception of activity in the water is unlike that seen in Richard Eyre's film *Iris*, for example, where swimming is equally prevalent. As in the book by John Bayley (1998) on which the Iris Murdoch biopic was based, *Iris* uses bathing scenes to represent what her husband perceived to be Murdoch's slide into

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depersonalisation. Despite the different function, it is hard to believe, however, that Subirana did not intend to reference an important precursor in the cinematic representation of Alzheimer's disease.

'No vull l'oblit': the value of scar tissue

Pia Kontos (2006) and Anne Davis Basting (2009) have proposed that we should forget memory if we are to value the lives of people diagnosed with advanced Alzheimer's outside an epistemological framework in which selfhood is synonymous with a fully functioning cognitive ability. Kontos uses Maurice Merleau-Ponty's notion of an immanent corporeal selfhood derived from the body's intuitive forward facing direction, and Bourdieu's notion of shared selfhood in a communal habitus, to find identity among a community of Jewish Alzheimer's patients brought together in a residential home. The context allows her to read even gestures and unformed words or lists of syllables as meaningful (Kontos 2006, 195-217). Similarly, in Forget Memory Davis Basting (2009) urges carers of the elderly and of people diagnosed with Alzheimer's to concentrate on who the patient is, not on who he or she was or on what he or she once knew. At first sight, these kinds of convalescent responses to the depersonalising and dehumanising effects of a diagnosis based on loss of cognition seem to be foreclosed by Subirana's yoking of personal recollection to historical memory and yet, if we listen attentively to how she describes her relationship with her grandmother, even before Leonor became cognitively impaired, she suggests they could relate to each other outside the boundaries of sense and abstraction. The director revisits the cinema where her grandmother worked as a cleaner for decades and tells us over a scene depicting the cloakroom where she spent many hours together with her grandparent: 'En aquell lavabo fèiem juntes la travessa i encara que cap de les dues no sabia res de futbol. ens fèiem companyia' (emphasis added). This observation echoes Subirana's gloss on the doctors' jargon-filled report describing her grandmother's condition: 'La veritat és que no hi entenia res'. It is out of this 'res' that Nedar strives to make something and such details point to a rapport which already sidesteps the construction of identity through the assertion of cognitive ability and may in part explain why Subirana avoids some of the traps Janelle S. Taylor identifies in her model of common narratives of Alzheimer's:

When it comes to speaking about dementia, horror seems to be the default genre. A person you love, and to whom you are bound by unbreakable ties, turns out to be someone you do not know at all, who does not 'care' about you and may even seek to harm you; this is the classic Gothic plot [...] or, alternatively, a person dies but their body lives on: this is the basic zombie story. (2008, 321)

Nedar's incorporation of a parallel film within a film – black and white sequences where actors play out a *noir* reconstruction of Juan Arroniz's clandestine activities and the hold up of a shoe shop for which he was arrested and executed – inscribes

the framing story within the genre of detective fiction or investigative journalism, rather than gothic horror or the zombie genre (see Figure 4). The metaphorical reading of Ana's and Leonor's illnesses, then, redirects the narrative towards a register which pulls pathography away from the monsterizing of senility. By affording her immediate family members a role in the recovery of historical memory, Subirana also interrupts the eventuality proposed by Paul John Eakin whereby Alzheimer's patients become 'subjects outside discourse, subjects culturally uninscribed' (as quoted in Roy 2009, 42). Nedar's metaphorical framework can also be seen as consistent with the way in which the Catalonian government has chosen to commemorate cultural landmarks. For example, a plaque at the former site of Camp de la Bota, where Ana walks her dog, is inscribed with the text of 'Aniversari', a poemby the doctor and poet Marius Torres which, in the context of a memorial addressed to the public, can be understood as an exhortation to the community to forge a kind of peace not dependent on oblivion in which wounds - mental as well as physical - should in some sense be valued as a record of the past:

Que en els meus anys la joia recomenci sense esborrar cap cicatriu de l'esperit. Oh pare de la nit, del mar i del silenci, jo vull la pau – però no vull l'oblit.

Subirana's deployment of illness as metaphor, then, entails both benefit and risk, and the film's negotiation between these two possible outcomes suggests that at stake there is a different conception of self – less anchored to a paradigm which values self-sufficiency (and thus hypercognitive autonomy) above all else – than the one presupposed by Anglophone narrators of Alzheimer's for whom finding a



Figure 4. A screen shot from one of the *noir* sequences in *Nedar* where actors perform Carla Subirana's reconstruction of the events which led to her grandfather's arrest.

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way to appreciate the life of a person diagnosed with the disease means circumventing established notions of personhood altogether. Taylor claims that

because a person's sense of self is grounded in his or her membership in a cultural group, when the political system in which they live fails to recognise the cultural identity of the group to which they belong this causes real harm to individuals. (2008, 325)

Since Subirana is working within a Catalonian and Spanish cultural discourse where this is well understood, she starts from a premise of selfhood which is already distinct from that which underpins many critiques of Alzheimer's discourse and hence the metaphorical value ascribed by her film to dementia functions bilaterally: an amnesic society is pathologised for forgetting its history and for forgetting its elders as much as individual patients are lumbered with the representative weight of a collective cognitive impairment.

Alzheimer's and cultural texts

The work of Janice Graham (2006) on Alzheimer's disease as a cultural text and her deconstructing of the cultural precepts which underpin dementia diagnosis are also germane to my reading of *Nedar*. Reference to Graham's contribution to the literature follows appositely from the previous discussion of Subirana's adaptation of a one-size-fits-all diagnostic practice to the specificities of a history particular to a given family and a national community struggling against cultural amnesia. According to Graham,

[t]he fervent activity directed at international standardization in [dementia] case ascertainment and assessment of severity is an attempt to gain [...] common language, a psychiatric Esperanto [that] constitutes no more than a necessary prerequisite for a rationally based nosology. (2006, 88–89)

Graham sees the dissemination of Alzheimer's diagnostic practice as an element in the flattening out and homogenisation of medical cultures, and by extension of cultures more generally. She proposes that 'the hard facts of neurological signs have gained diagnostic supremacy over that of the social affect and general well being of the person' (2006, 91), and draws on Julian Leff's discussion of an epistemological approach to psychiatry, one that

generates a view of psychiatry as a cultural product of Western societies in the same way as are the diatonic scale and Coca-Cola. These products can be exported all over the world and may be appreciated by the local populace, but are no more valid than indigenous music or beverages. (Leff 1990, 305)

Since it includes a number of fly on the wall scenes where Ana's submission to cognitive and photographic tests is frankly portrayed, *Nedar* allows us to see the collision of a homogenising diagnostic model – in this case a neurological one – with aspects of cultural specificity.

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Some viewers - Corbella (2011), for example - have read scenes where Ana is asked to draw a clock face or to recite a list of words as incidental occurrences within the film, and these scenes are furthermore recreations of internationally standardised tests for cognitive impairment, such as the 3MS (Modified Mini Mental State), the Seven Minute Neurocognitive Screening Battery, and the CDT. As usual, medicine resorts to acronyms and abbreviations to imply a level of complexity which is not necessarily to be found on closer inspection. For example, the CDT is simply the clock drawing test, where Alzheimer's and dementia patients around the world are asked to draw a clock face showing eleven minutes past ten: failure in the test indicates loss of temporal and spatial awareness. It is this test which Subirana recreates for her mother at home. In a hospital setting. Ana is asked by a doctor to repeat a list of words – as many as she can remember. There is a correct way to answer wrongly: in other documentaries about Alzheimer's (The Alzheimer Project, for example) we see patients who can only remember the last word in the list, or who cannot remember anything, or who cannot remember the question. Here, however, Ana's wrong answers are also wrong in the wrong way. To questions seeking information about the names of her parents she answers with impressions about the Civil War and this list of words speaks not only to a deficit of short term memory but also to a surfeit of connotative memory. The word list contains the noun 'presó' and this sets Ana off on a train of thought about the injustices suffered by her father, his imprisonment, and his execution. If on the one hand we might have reservations about Subirana's recourse to a homogenising discourse of dementia diagnosis as a way of recovering or of spotlighting the loss of culturally specific memory, on the other hand the disjuncture between the standardisation of the tests and the peculiarity of some of the results in fact resists the evacuation of local specificity against which scholars such as Graham caution.

As Alzheimer's pursues its course through her family, Subirana gradually uncovers more information about her grandfather. Joaquim Jordà – the director of *Mones com la Becky* (1999), to which *Nedar* has been likened (see Martí Freixas 2008) – insists that if the police report contends that Arroniz shouted '[1]os rojos también tenemos derecho a la vida' as he held up a shoe shop, he must surely have been politically active. Subirana finds her grandfather's sister and learns that he was indeed active within an anarchist-syndicalist cell. Her great aunt goes regularly to a local pharmacy to pick up her medication at the same address which formerly housed the offices of the group to which her late brother belonged, a coincidence which seems not only to concretise the displacement of radicalism by Foucauldian biopower but also further to authorise the film's knitting together of biomedicine and cultural memory.

Conclusion: forgetting memory?

As Subirana reaches some sketchy conclusions about the circumstances leading to her grandfather's arrest and his subsequent disappearance from her family's history, her mother's diagnosis becomes clear and the editing of the film again signals resonances between an oppressive political apparatus and the narrow confines of medical discourse. Subirana situates immediately before the scenes of her mother's MRI scans a portrait of Ana at the monument to Camp de la Bota, as if the executions continued and as if the sentence against her mother's cognitive person were a further form of assassination and cultural erasure. Arguably, this is more deleterious to the Alzheimer's patient than other uses of metaphor in the film; it is certainly harder to mitigate. But the juxtaposition of the two forms of representation also recalls Georges Canguilhem's observation that 'the science of life no longer resembles a portrait of life' (1994, 317). The pictures at the memorial could also be seen as memorialising the living person before the dehumanising work of the machines and test batteries begins. Robertson points out that

[w]ith the growth in medical technology, such as the imaging technology used in the investigation of AD, the panoptic gaze focuses on ever more detailed examinations of the experience of aging, including brain function. This kind of technology, combined with the technology of cognitive and psychological testing has been partly responsible for the recent focus on AD. (1991, 137)

Nedar brings to the surface the cradle-to-grave function of such invasive imaging in shaping and demarcating an individual's progress through life, and perhaps we can also ask if it and other films like it are not implicated in the visual reification of Alzheimer's disease. How far is *Nedar* a critique of techniques of visualising medical epistemology and how far does it contribute to the legitimisation of those same technologies?

While I have only been able here to sketch out an answer to this question and to those I posed earlier, an approach to *Nedar* informed by the debate over the exploitation of disease as metaphor does indicate with some degree of conclusiveness that Spain's forgetting of some of its past complicates the forgetting of memory proposed by some of those working with Alzheimer's as a form of reconciliation with the effects of the disease. Inasmuch as Spain's history of forgetting pathologises erasure, this makes it harder in Spain to accept on an individual level the forgetfulness of amnesia and thus Spanish and Catalan politics have become imbricated in the aetiology of Spanish and Catalan presentations of Alzheimer's disease. Yet at the same time as Subirana activates the powerful illness metaphors of Alzheimer's disease to dramatise collective historical losses, she also looks for family attachments and intimacies which escape the cognitive functions impaired by dementia and thus does not entirely turn over her family members' identities to a larger memorial project. To put it another way, whilst extrapolating from her immediate family members' pathographies a pathological

historical lacuna Subirana is also able to create a more intimate account of her maternal family which finds biography beyond and between pathography. As a discussion of the film alongside the debate within medical anthropology over the value of illness metaphors has also illustrated Subirana – to an extent – cannot in any case represent Alzheimer's nor the people closest to her diagnosed with Alzheimer's without recourse to metaphor since an effect of the illness is to take away from a subject precisely the ability to narrate her own life. Given that *Nedar* does not have recourse to metaphors of horror in reaching an accommodation with this metaphorical imperative (as many accounts of the disease in the literature in English do), it has been my contention in this article, furthermore, that close study of the film demonstrates that while diagnostic practices for dementia illnesses may tend towards standardisation, the creative responses which they provoke do not. There are culturally significant differences between the paradigms adopted by narrators of Alzheimer's across languages and cultures.

Nedar's exploration of the interstices between a globalising discourse of an Alzheimer's apocalypse and the personal and more culturally specific iterations of disease experience makes manifest that traces of identity remain even where it has been bulldozed by political tyranny or where it is being overwritten by the plaques and tangles which impair cognition.

Notes

- 1. Nicolas Roeg's 'Iceberg' was one of five television spots broadcast in the UK in 1986 designed to shock people into practising safer sex. '[F]eaturing falling icebergs inscribed with the word "Aids". This was public information at its most sensational. Expensive special effects and high-concept production design brought public information filmmaking into the realm of state-of-the-art corporate advertising' (McGahan 2003-10). John Hurt narrated the films and during the course of 'Iceberg' he warns viewers: 'You can't always tell if someone is infected and unless we're all a lot more careful, the people who have died so far will be just the tip of the iceberg'. The 'Iceberg' and 'Tombstone' films can be viewed within the BFI's screenonline archive.
- 2. Robert Katzman and Katherine L. Bick (2000) chronicle the resurgence of interest in Alzheimer's disease in the 1960s and 1970s, and note the rise in the number of scholarly papers on the disease following the first modern symposium on it in 1969 and the roll out in the same decade of electron microscope technology which allowed the study of brain pathology at the ultrastructural level. While only 10 articles were published on Alzheimer's disease in 1966, the number rose to 2,372 in 1996 (2000, 110). Herskovits (1995, 150) sustains that 'one can persuasively argue that Alzheimer's recent emergence on the biomedical terrain is an artifact arising from local sociocultural and political dynamics'.
- 3. Emil Kraepelin was the first scientist to use the term Alzheimer's disease, in 1910. Konrad Maurer, Stephan Volk, and Hector Gerbaldo (2000) consider several possibilities for Kraepelin's hasty creation of the eponym (which attached the disease to Kraepelin's Munich Nervenklink) and find plausible among them his 'wish to show the superiority of his school over psychoanalytical theories and to show (vis-à-vis Freud) that some mental disorders were organically based' (2000, 26).
- 4. 'L'article 54 de l'Estatut d'autonomia de Catalunya estableix l'obligació de la Generalitat i els altres poders públics de vetllar pel coneixement i el manteniment de la memòria històrica de Catalunya com a patrimoni col·lectiu que testimonia la resistència i la lluita pels drets i les

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- Ilibertats democràtiques, i disposa que amb aquesta finalitat han d'adoptar les iniciatives institucionals necessàries per al reconeixement i la rehabilitació de tots els ciutadans que han patit persecució com a conseqüència de la defensa de la democràcia i l'autogovern de Catalunya' (see Department d'Interior, 2008).
- 5. Wendy Roy refers to the work of Sidonie Smith and Paul John Eakin in this regard. She cites Smith's inclusion of Alzheimer's among a list of other conditions which consign those who suffer from them to 'unautobiographical' lives which can only be known thanks to the representation performed by others. She also cites Eakin's suggestion that memory loss disabilities may prevent the performance of any kind of self-narration (Roy 2009, 41-45).
- 6. Subirana has suggested in an interview that the scenes shot at the swimming pool are the ones which allow the strands of the film's narratives to agglutinate (Surt TV, 2009). This idea is also discussed by Prout (2011, 101–104).
- 7. Michael Philpot (2004) assesses the CDT's usefulness as a stand alone test and also provides an overview of some of the key literature relevant to this form of discursive test.

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