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What is that's going on here? A multidimensional time concept is foundational to framing for decision making in situations of uncertainty

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Abstract

STEM disciplines are the dominant culture in K-12 education. With its study of organs and diseases that afflict patients' bodies, Western evidence-based medicine is seen and understood in the modern cultural paradigm as a science and as the practice in which a subject, the doctor, acts on an object; the patient's body—a dominant culture in the patient's journey. However, with the continually evolving high-technological and medical knowledge, life-saving therapeutic options are life-changing. They can range from changes in the diet, requiring structural and cultural changes in family life, to changes related to the experiences of learning to live tethered to a machine that is partly inside and partly outside one's body or with somebody else's heart. In this article, we show how competing needs to personalize care for the patient as a person forcefully emerge in response to evidence-based medicine's global cultural dominance. We highlight two fundamental issues emerging in decision-making processes: (1) Framing evidence-based knowledge, uncertainties of the course of the disease and options, and (2) working with different, equally important, and often at odds *conceptions of time* in the care for the Other. Through the longitudinal analysis of moment-to-moment interactions in high-tech medicine encounters of a patient, his family, and the team caring for them, we show how *framing* and *different conceptions of* time emerge as issues, are profoundly interconnected, and are addressed by participants to care for a patient confronting existential decisions.

Keywords Care \cdot Framing \cdot Multidimensions of time \cdot Professional vision \cdot Shared decision making

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Abstract in Italian

La cultura scientifica ha un ruolo dominante nell'istruzione scolastica negli USA. Con lo studio degli organi e delle malattie che affliggono il corpo dei pazienti, la medicina occidentale, basata sulle prove di efficacia clinica, è vista e intesa nel paradigma culturale moderno sia come scienza sia come attività pratica in cui un soggetto, il medico, agisce sull'oggetto, il corpo del paziente – la medicina è la cultura dominante del percorso che deve intraprendere il paziente.

Tuttavia, con le conoscenze mediche e tecnologiche in continua evoluzione, le moderne opzioni terapeutiche salvavita stanno modificando l'esistenza dei pazienti. Le terapie possono variare dal mutamento della dieta, il che richiede trasformazioni strutturali e culturali all'interno della vita familiare, a cambiamenti connessi alla necessità di imparare a convivere con una macchina che è in parte interna e in parte esterna al proprio corpo o con un cuore o un altro organo estraneo.

In questo articolo mostriamo come le esigenze di personalizzare l'assistenza al paziente in quanto persona emergano con forza in contrapposizione al predominio culturale dominante della medicina basata sulle prove di efficacia clinica. Evidenziamo qui due questioni fondamentali che emergono nei processi decisionali: 1) Inquadrare (framing) la conoscenza basata sulle prove di efficacia clinica, le incertezze del decorso della malattia e delle opzioni terapeutiche e 2) lavorare con concezioni di tempo diverse ma ugualmente importanti, e spesso in contrapposizione quando si cura l'Altro.

Attraverso l'analisi longitudinale delle interazioni momento-per-momento durante gli incontri di medicina high-tech tra un paziente, la sua famiglia e l'equipe medica, mostriamo come il framing e le diverse concezioni di tempo emergono come problemi, sono profondamente interconnessi e sono affrontati da coloro che curano il paziente che deve prendere decisioni esistenziali.

How to care for a person catapulted into a world that is unfamiliar, where the person confronts choices that infringe on the lived experiences and where the person's needs and queries are often in sharp contrast to those acknowledged by the dominant culture?

Educators continually confront this question, albeit in the specificity of each situation and community, when meeting first-generation students attending college, immigrant and migrant youth (Suárez-Orozco, 2004), and youth who do not experience familial and/ or cultural continuity (Miller et al., 2015). In science education, Glen Aikenhead (1996) depicts the powerful image of students who need to "border-cross" from their lived experiences into the subculture of science and science teaching to be able to learn science. In all the STEM disciplines, there is a long-lasting discussion on how to best implement what Gloria Ladson-Billings defines as culturally relevant pedagogy (1995) that is responsive to different ways of knowing (Gutiérrez & Rogoff, 2003), inhabiting the world (Elmesky & Seiler, 2007) with religious beliefs (Roth, 1997), lived experience, cultural norms and values (Meyer & Crawford, 2011) that are in contrast to those accepted in the scientific paradigms and classroom context (Mavuru & Ramnarain, 2020). *Cultural Studies of Science Education* has been examining these issues since its foundation in 2006 by Kenneth Tobin and Wolff-Michael Roth.

In this paper, we discuss how similar issues are tackled by medical practitioners in the practice of high-tech medicine of Advanced Heart Failure (AdHF), when patients with malfunctioning hearts are confronted with the decisions to accept to live with the heart from another human being (heart transplantation) and/or with wearable mechanical circulatory support devices (e.g., a Total Artificial Heart, BiVAD) that can be at odds with the person's sense of what is acceptable. We do so by analyzing recorded data from moment-to-moment interactions and showing how tensions between the dominant Western medicine cultural paradigm (Cunningham & Andrews, 1997) with its continually evolving high-technological and medical knowledge and the need to personalize care for the patient, as a person, are always at play and are continually emerging and in need to be resolved in the interactions of the AdHF medical team and patients and their families.

To better make sense of the context of AdHF, we start by taking the reader in the situation experienced by a patient, Mr. Spencer. We then develop our questions from this scenario.

Mr. Spencer sees his wife's face coming slowly into focus. Her voice, first a distant sound, becomes closer and louder as she yells, "Don't you leave me! Keep your eyes open, STAY AWAKE HONEY! DON'T YOU LEAVE ME!".

Mr. Spencer is being resuscitated. He is in the Coronary Care Unit (CCU), diagnosed with a rare, aggressive and often fatal autoimmune disease, Giant Cell Myocarditis, that drove him into cardiac arrest, again. Being in CCU is a constant reminder of this experience and of the difficult choices, patients and families have to make. As Mr. Spencer recounts:

Mr. Spencer. Segment 1_ Every patient in the CCU unit has serious heart issues, so I was not the only person going Code Blue¹ from time to time. The alarm and announcement occurred multiple times each day. "Code Blue! Code Blue! [...] Code Blue! Code Blue! [...]." We would hear people running down the hall. One night it was in the room next door. For nearly an hour we listened to the elevated voices calling out directives and continually re-announcing vital numbers. The frantic activity sent furniture or other heavy items crashing into the wall. The alarm and flashing lights continued for the duration. I grappled aloud with Tracy on how far do we go as Christians to scratch and claw at hanging on to this life when we know that this world is not our home. It weighed heavy on her when I expressed that I didn't want to be shocked and pounded and pumped incessantly like that.

(¹Code blue is an emergency announcement over a hospital's public address system requiring a team of healthcare providers to rush to the specific location and begin immediate resuscitative efforts of a patient in cardiopulmonary arrest.)

Western medicine with its study of organs and diseases that afflict patients' bodies is seen and understood in the modern cultural paradigm as a science and also as the practice in which a subject, the doctor, acts on the object, the patient's body (Timmermans & Almeling, 2009). In this paradigm, issues of objectivity (Daston & Galison, 2009) and objectification for the subject to understand, study and control things and other beings (Haraway, 1988) are considered important and necessary (Harding, 2015) but are incompatible with Mr. Spencer's agentic sense of being a person, not an object, and his aversion for overtreatment and potential futility (Clark, 2007) of high-tech medicine.

How to address Mr. Spencer's reluctance toward high-tech medicine treatments perceived as medical futility by him but not by his family? Mr. Spencer's Giant Cell Myocarditis disease developed suddenly. One day he was decorating the Christmas tree with his family, the next he was acutely falling ill and progressing toward AdHF. As per Guidelines Directed Medical Therapy for Giant Cell Myocarditis, Mr. Spencer has been treated with high doses of immunosuppressant medication. At the same time, in agreement with Mr. Spencer, the AdHF team has initiated the evaluation process for heart transplantation as a backup plan. Later, Mr. Spencer will recount about these days:

Mr. Spencer. Segment 2. It is interesting how one gradually begins to accept things he would have never accepted before. For years I had reflected on the passing of my grandparents and others and was fairly adamant that I never wanted to be on life support. I certainly would not have been amenable to the idea of a heart transplant. When it was my time to go, I just wanted to be let go. Of course, I always figured that would come in my 80's or 90's, not at the fresh age of 50.

Mr. Spencer has to yet come to terms with the idea of living with the heart from somebody else. In sociocultural terms as Margrit Shildrick (2011) puts it, "the heart stands in for a range of inherently human attributes [...] that are at the core of selfhood" (p. 234), which for Mr. Spencer includes a spiritual sense of self. As Federica Raia and Mario Deng (2015) discuss, the possibility of accepting the idea of heart transplantation becomes possible when one is compelled to uncover one's body as made of substitutable parts. In this process, "my heart" then transforms into a malfunctioning or broken "tool" that cannot support my life. Learning to accept "things [he] would have never accepted before" is now becoming part of Mr. Spencer's life experience.

How to frame the possibilities, the uncertainties, and course of the disease in high-tech AdHF in a meaningful context for Mr. Spencer confronting existential decisions?

Ethnographic work on caring for AdHF patients points to the ideological contradictions emerging from the competing needs to—on the one hand—objectify bodies and organs and—on the other hand—personalize care for the patient as a person (Sharp, 1995). Competing needs to personalize care for the patient as a person (Sharp, 2006) requires a multidimensional understanding of time (Raia, 2020). Indeed, a cohesive sense of a person's *past* and *future* possibilities, are both necessary to make sense of who we are and our actions in the *present* making the existential time nonlinear (Heidegger, 1962). However, within the modern cultural paradigm of science, and therefore in western medicine, time is one of the fundamental quantities, measured by a clock (clock-time) and is a linear progression from the past to the present into the future.

How to resolve the competing needs of the complementary concepts of clock-time and existential time to safely and meaningfully care for Mr. Spencer?

To address these questions, we highlight two issues: (1) *framing* the evidence-based knowledge of AdHF, uncertainties of the course of the disease and options for a meaning-ful decision-making process facing the patient, Mr. Spencer, and (2) Working with different, equally important and often at odds *conceptions of time* in the care for the Other. We discuss them below, reviewing some of the literature from different disciplinary perspectives that help us frame ours. After describing the research model, we present sequences of

talk-in-interaction during two medical encounters between Mr. Spencer, his family, and the AdHF team. We ground our understanding in the study of how the issues of *framing* and different *dimensions of time* manifest in the moment-to-moment interactions and show that they are deeply interconnected.

Framing

The concept of framing, in its original formulation by Gregory Bateson (1972), serves to define context, organize perceptions of behavior, and shape how something makes sense to participants. Developing his concept of frame analysis, Erving Goffman (1974) suggests that people use frames to answer the basic question of "what is it that's going on here?" Since then, the concept has been utilized in different domains of social interactions.

In the learning sciences, situations in which professionals need to attend to the other's learning, such as in the interactions between teacher and students, framing is understood as the response to student thinking, as an attunement to, respect for, and engagement with intellectual resources students bring to bear in disciplinary contexts. In medical decision-making literature, framing is understood in a context of decision making, as organizing the possibilities of treatments and the choices a patient faces (Wirtz et al., 2006). It has been studied in primary care (e.g., Elwyn et al., 2012) and in oncology (Gattellari et al., 2002). These studies show that framing is often neglected or given too little time by practitioners in their interactions with patients. We tentatively adopt both definitions to study how framing is done in the moment-to-moment interactions, in which making-sense of a disease and therapeutic options require learning different medical concepts and a different language that is specific to the medical disciplinary context.

Dimensions of time

Wolff-Michael Roth, Kenneth Tobin and Stephen Ritchie (2008) show how classroom teaching and learning is mediated by different dimensions of time. Specifically, they identify not only misalignments between the temporally linear organization of the school day, and the way students and teachers experience and organize their activities, but that "rosters, deadlines and calendars, reporting periods, announcements on the PA by the principal, pre-lab activities and after-school activities, and being on time and being late create "temporal orientation, timing, pace, cycles, and rhythms" (p. 136). This time dimension, we call organizational, governs teachers' and students' experiences and possibility of success.

As we discussed above, clock-time is one of the fundamental quantities in science, measured by a clock with a linear progression from the past to the present into the future. As Raia (2020) discusses, the linearity of clock-time implies that the past has causal influence over the action of the present and the present over future action.

Existential time emerges from an individual's lived experience. In sociocultural theory, existential temporality has been explored in part through Lev Vygotsky's theory of play (Vygotsky, 1978) and Michael Cole's (2007) notion of prolepsis.

As Vygotsky discusses: "If human activity were limited to reproduction of the old, then the human being would be a creature oriented only to the past and would only be able to adapt to the future to the extent that it reproduced the past. It is precisely human creative activity that makes the human being a creature oriented toward the future, creating the future and thus altering his own present" (Vygotsky, 2004, p. 9). The child confronting the existential demands of becoming is always in dialogue with her future: "*play gives a child a new form of desires*. It teaches her to desire by relating her desires to a fictitious "I," to her role in the game and its rules. In this way, a child's greatest achievements are possible in play, achievements that tomorrow will become her basic level of real action and moral-ity" (Vygotsky, 1978, p. 100).

In Cole's notion of prolepsis, a child is put always in dialogue with her future while confronting the existential demands of becoming, to make sense of how parents' expected goals for the future of their child constrain and guide their child's upbringing actions in the present. The nonlinearity of the process is introduced by the parents' culturally mediated, imagined future for their child. Both approaches assume cultural continuity (Cole, 2007, p. 239), while Raia's Relational Ontology (2020) focuses on caring-for-the-Other, catapulted into a world that is unfamiliar and threatening, does not assume cultural continuity. Indeed, she shows that to care for another, a professional needs to take into account not only a projection into future possibilities but also a past, no matter how disconnected it feels to the current situation. Building on Martin Heidegger's work, she shows that both are necessary because the *past* modulates the person's sensibility to attune to what is relevant to this person, and the *future* modulates the sense of possibilities of being this person. Indeed, a person's past and future possibilities together are necessary to make sense of the present, by organizing how things show up in the world as mattering in the present, and what it makes sense to do for this person. Specifically, Raia shows how the physician, in dialogic interactions, builds with and for the patient an existential temporal horizon: the physician recruits *past* events in the patient's life as salient for both the patient and physician and, after grounding the patient in a common relevant past, projects the patient into possibilities of being this person (*future*), he then returns to present actions in which the patient can inhabit a meaningful and shared *present* with the physician.

Within the Relational Ontology framework, Mr. Spencer's cohesive sense of the existential horizon is fragmented, when confronting death as the end of being in the world, and facing the world of AdHF, unfamiliar and disconnected from his existential *past* and projection into *future* possibilities of existing in the world. Based on this, we understand Mr. Spencer's sense of meaningful engagement in activities with others in the current situation (*present*) developing as an existential crisis.

As discussed above, the possibility of accepting the idea of heart transplantation can be possible in situations of existential crises in AdHF (Raia and Deng, 2015). That is, when Mr. Spencer is compelled to uncover his body as made of substitutable parts, his heart transforms into a malfunctioning or broken "tool" that cannot support his life. Following Timothy Koschmann et al. (1998) who, building on Martin Heidegger, Aleksei Leont'ev and John Dewey, posit that a breakdown of fluent pragmatic doing is necessary to learn, Mr. Spencer's acceptance of "things [he] would have never accepted before" can be seen as a process of learning within the AdHF practice.

Research model

This work is part of an ongoing ethnographic and participatory research project studying the practices of teaching, learning and patient care in the high-tech medical practices of an AdHF program in a large US university hospital (Raia and Deng, 2015) with more than 500 h of recorded medical encounters between healthcare professionals (n > 25) and patients (n > 125) (IRB#11–003,590).

The research model developed to study this practice (Raia, 2018) proceeds iteratively in three stages of data collection and analysis:

Stage 1: Encounter Recordings of AdHF medical encounters are longitudinally audio/ videotaped for a period of 1–4 years.

Stage 2: Cogenerative dialoguing (cogen) (Roth and Tobin, 2004). Weekly video-/ audiotaped sessions with participating healthcare practitioners (co-authors here) whose interactions were recorded in Stage 1. Together participants and researchers review participants' taped interactions to make sense of their practices and discuss the emerging elements and themes. We report here parts of these weekly audio/video reviewing sessions (cogen session) in making sense of the data.

Stage 3: An ethnomethodological informed microethnographic (Erickson, 1996) analysis of the practice-recordings identifies the resources used by participants to organize their conduct and reciprocal accountability. Events identified by researchers and participants in Stage 2 are transcribed utilizing some of the transcription symbols (Sacks et al., 1974) reported below.

Although our analysis is based on collections encompassing multiple encounters from the data corpus, here we present a "single-case analysis" and report data from medical encounters of one of the 125 recorded patients, Mr. Spencer (pseudonym). This choice, as also Karen Lutfey and Douglas Maynard (1998, p. 323) discuss in their work on medical encounters, is consistent with Emanuel Schegloff's organization of analysis in which a range of phenomena from a larger corpus of "talk-in-interaction are brought to bear on the analytic explication of a single fragment of talk" (1987, p. 101).

Transcription system

[Left square bracket, on two successive lines with utterance by different speakers marks the point at which the talk above is overlapped by the other talk a line below
=	Equal signs in pairs indicates that there is no discernable silence between the end of the first and the start of the next utterance, the first is ' <i>latched</i> ' to the following
ord-word	A dash between words marks an acceleration of talk
(0.5)	Number in parentheses indicate silence in seconds
:	Colons indicates that the sound that immediately precedes the colon has been sensibly prolonged or stretched
word	Bold and italic indicates some kind of stress or emphasis, which maybe signaled by a change in pitch and/or amplitude
Word	Capital letters indicate raised pitch or volume
(())	Double parentheses enclose comment by the transcriber
? ,	Intonation: Punctuation symbols are used to mark intonational changes rather than as grammatical symbols: A period indicates a falling contour A question mark indicates a rising contour A comma indicates a falling-rising contour.
(hh)	indicates breathiness rather than laughter in the midst of a word; in the cases included it is near sobbing.

Moment-to-moment interactions

December 29th

The full case study of the data from our corpus is reported in Raia et al. (2021). Here we report two sets of data from two medical encounters with Mr. Spencer, bringing a new analysis of the data showing how issues of *framing* and different *dimensions of time* manifest in the moment-to-moment interactions and are deeply interconnected in the care for the Other. The first set shows how framing emerges in talk-in-interactions and is complexified by the needs of taking into account different dimensions of time. The second set shows the encounter with different healthcare professionals and how Mr. Spencer confronts choices, as shown in Fig. 1, that infringe on his lived experiences, needs and queries that are in sharp contrast to those acknowledged by the medical dominant culture, and how the participants resolve these tensions.

Heart transplant waiting time is not an hour, a day, a week, a month

This encounter introduces us to Dr. D, the AdHF attending, a cardiologist specialized in heart transplantation and mechanical assist device implantation who is conducting his

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Fig. 1 Mr. Spencer's timeline. Mr. Spencer falls ill after Thanksgiving. In mid-December, a biopsy of the heart confirms Giant Cell myocarditis. Despite immunosuppression, Mr. Spencer's heart function worsens, and the team recommends evaluation for heart transplantation (HTx). At the end of December, the AdHF team recommends *BiVAD implantation as a bridge* to HTx. On New Year's Day, the team recommends *ECMO as a short-term bridge to a long-term BVAD bridge* to HTx. Thick dashed lines indicate the encounters we report here. Dr. D is a cardiologist specialized in AdHF, Dr. S is an AdHF surgeon

hospital rounds in the CCU accompanied by two trainees, the general cardiologist (GC) fellow and the AdHF fellow. They enter Mr. Spencer's room. It's December 29, and Mr. Spencer had suffered multiple episodes of arrhythmia. Mr. Spencer sits in his hospital bed, next to him stands his wife and CCU Nurse RN N. Dr. D discusses the recovery trajectory of Mr. Spencer's heart based on the results from the latest ultrasound rounding with him listen attentively:

1	Dr. D:	when uhm we were in with the team uhm
2		we uhm uh were talking about
3		from the update ultrasound (0.7)
4		it is not surp <i>risi</i> ng that things aren't uhm (0.8)
5		in-quotation-mark
6		just turning around (0.8)
7		it's uhm from the overall condition
8		something that uhm (1.0) is u::h (1.2)
9		gonna-be? (0.4) ch a: lleng in g

There is no possibility of recovering Mr. Spencer's heart function. As the ultrasound results (line 3) show, things are not turning around (lines 4–6); the situation is challenging (line 9). Dr. D talks slowly with short acceleration (line 5), interspersed with hesitations ("uhm" lines 1, 2, 4, 7, and 8) suggestive of the talk's delicate nature (Silverman & Peräkylä, 1990). Dr. D's slow but inexorable pace, the hesitations, and the number and the length of the notable silences (>0.4 s) places a significant emphasis on the coming words. Prominence is given to the utterances "not surprising" (line 4), emphasizing the syllables.

The pause after a fast uttered "gonna-be" (line 9) produces an effect of suspension marking, even more, the importance of what is uttered afterward: "cha:llenging" is followed by another pause, this time, marking the gravity of the situation: the possibility of recovering Mr. Spencer's heart function is vanishing.

10	Dr. D:	we will be uhm (1.4) intensifying
11		the immunosuppressive therapy (0.8) uhm
12		and uhm (0.6) we will continue to (0.9)
13		closely monitor? (1.2)
14		uhm (1.4) but it's uhm
15		(0.5) ((swallows deeply)) (0.4)
16		even in the b: <u>e</u> st (.) possible scen <u>a</u> rio
17		(1.2) that we
18		as we said yesterday hum
19		are uhm observing since you ca:me
20		here and we started the therapy? (0.6)
21		it's going to be some ((clears his voice))
22		first of all long term immunosuppression
23		which is the $least$ (1.0) of a problem
24		but there is still (0.6) even at best possible immunosuppression
25		l:ong term uhmm some limitations in (.) good outcomes
26		in other words
27		this transplantation option
28		just n <u>ee</u> ds (0.4) to be recommended
29		as a (.) close standby backup option (1.8)

Dr. D continues slowly to move toward the challenges ahead. Tracing back to the familiar step of immunosuppressive therapy to Mr. Spencer, Dr. D moves toward the challenges without mentioning them yet (lines 14–16). He moves in little increments, going back to familiar issues and then forthwith hesitations (e.g., "uhm" line 14). In silence or by clearing his voice (e.g., line 21), he pauses, pre-annunciating the importance of what is uttered afterward. What comes modifies what has been just uttered; the familiar immunosuppressive therapy, becomes long-term, the long-term therapy then becomes "the least of a problem" (line 23).

Lines 1–25 cumulatively point to Dr. D's involvement in the situation that is not just related to a decision-making process (Skelton et al., 2002) regarding what to do next with these new results or to minimize resistance to reach consensus (Robertson et al., 2011). Dr. D's talk (line 1–29) displays an emotional stance (M. Goodwin & Goodwin, 2000) pointing to a participation in the disappointment that the option of transplantation is now closer (line 29) on the horizon with Mr. Spencer's heart function worsening, and Mr. Spencer's hope to retain his native heart is vanishing.

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30	Dr. D:	the (.) problem being
31		$\underline{\mathrm{i} \mathrm{f}}$ for some reason we feel that things aren't heading the right direction
32		from that time point <u>on</u> to
33		transplantation
34		is not (1.0) an <i>hour</i> period
35		it's not (.) a <i>day</i> period
36		it's not necessarily a <i>week</i> period
37		it's not necessarily a <i>month</i> period
38		the median waiting time for heart transplantation
39		in this-region-of-the-United-States is three (.) point (.) zero (.) months
40		which i:s [hospital name]
41		which is the shortest in the country? for (.) population reasons (0.9)
42		and that is why then
43		there may be this necessity of an additional bridge to transplantation
44		if that medication isn't stable enough
45		(1.0) talking about these pumps (1.0)

Dr. D is at a very delicate point in the conversation. Introducing heart transplantation as a very close option brings other complications. With no heart function recovery, the option of heart transplantation includes the possibility of a bridge to heart transplantation via mechanical assist devices. With Mr. Spencer's worsening condition, the immunosuppressive therapy can no longer sustain his heart. The waiting period for heart transplantation can vary in time (lines 31-39). Dr. D slowly proceeds toward the challenges ahead. Being approved for heart transplantation requires waiting. The more hesitant talk we have on lines 10-25 for Mr. Spencer's vanishing hope of recovering gives way to a reciting voice (lines 34–37). The increments of clock-time attempt to measure a definite, yet unknown amount of time. The listener is taken into a rising and falling rhythm repeated each line 34–37 to admit the only certainty. There will be a waiting time. This waiting period is a liminal space for patients and their families, who often report that they are caught between hope for an offer of a donor heart, guilt knowing that the heart offer requires someone else's death and another family's tragedy and an ongoing concern about their current, fragile, status. Clock-time enters this space in a strident juxtaposition to existential time.

Framing in practice

The new development in Mr. Spencer's condition requires thinking more purposefully on how to reframe the scenario of a bridge to transplantation (line 43) by machine implantation, "these pumps" (line 45), i.e., the BiVAD, now looming on the horizon. It is an "open heart surgery" creating considerable anxiety in patients not only because of the procedure's complexity and potentially serious complications but also for fear that during the months of recovery their condition could worsen; they would die attached to a machine that is half inside and half outside their bodies before having a heart transplantation.

46	Mr. Spencer:	that makes (.) that makes (.) perfect sense to me
47		(5.0)
48		you're basically saying
49		go ahead and put me on the list (1.8)
50		and then (2.0) ((clears his voice))
51		the only question would be ((clears his voice))
52		I:f things came up earlier (1.2)
53		and we weren't really sure whether this heart (0.4)
54		was going (0.5) to make it or not
55		we have to make a split second decision=
56	Dr. D:	=that's br ill ia:nt thought (0.6)

For a person witnessing the dialogue, Dr. D's utterances such as "that is why then" (line 42) and "talking about these pumps" (line 45) would require an elaboration to be understood. Equally difficult to comprehend are the first lines 1–10 and all the references to the "list" to which Mr. Spencer also refers (line 49). Mr. Spencer's utterances, such as "if things come up earlier" (line 52), would require some elaboration for a casual witness to make sense of them; elaboration that we are providing based on ethnographic work. However, it makes perfect sense to both Dr. D and Mr. Spencer, who have the sense of "what is that's going on here" (Goffman, 1974). Dr. D has discussed the need for *framing* during the first visit as fundamental for the patient to learn to contextualize therapeutic options, uncertainties, and potential challenges that could present themselves in the patient's illness and life journey (Raia et al., 2021).

With the disease's erratic behavior, there is now a need to discuss, within the frame already in place, the possibility of a bridge to transplantation with machine implantations as a viable option. Dr. D does not interject during the five-second silence (line 47) at the end of Mr. Spencer's statement of agreement (line 46) nor during Mr. Spencer's numerous pauses. This muted request serves as an open invitation for Mr. Spencer to continue to talk after having introduced the transplantation option and is particularly noticeable because the silences on lines 49 and 52, like the one on line 47, are at the end of a finite statement and could signal to Dr. D that he should continue talking. An agreement as to what makes "perfect sense" is not enough to grasp "what's going on here," in Mr. Spencer's life. Deciding to accept heart transplantation is an existential choice, and the gravity of what has been said needs to be fully grasped.

Dr. D needs to care for Mr. Spencer (Raia, 2020), who is existentially shaken by the now very real possibility of losing his heart. What needs to be fully existentially grasped must start from what Mr. Spencer will say next.

"And then" (line 50) ... Mr. Spencer's doubt emerges. The two-second silence and the clearing of his voice emphasize what is uttered after "the question" remaining, Mr. Spencer clears his voice again. If the heart offer comes up too early (line 52), there is still some chance that his heart could recover (53–54). A "split-second decision" on a life-changing option must be made (line 55) to accept or not to accept the heart offer.

Nuancing the discussion on making decisions within a much more complex context of uncertainty is a "brilliant thought" (line 56). But, what about the bridge to transplantation through machine implantation? Dr. D follows Mr. Spencer's lead. First to be discussed is "the split-second decision" uncertainty, a heart offer could come up "earlier" (line 52) when Mr. Spencer's heart function could still be sufficiently stable (lines 52–53).

January 1st

Difficult decisions

Mr. Spencer's condition is deteriorating fast. A few days later, on New Year's Day, Mr. Spencer is facing the dreaded option of a bridge to transplantation—the implantation of machines (HVAD-BiVAD) surgery. Today he meets for the first time his AdHF surgeon, Dr. S. Together they discuss a new heart-ware HVAD-BiVAD that Mr. Spencer prefers over other types of BiVAD, the sequence of surgeries, and the machine implantation first and after recovery, heart transplantation. We enter in Mr. Spencer's CCU room when the AdHF team members are assembled around the bed: Dr. D, Dr. S, CCU nurse RN, AdHF Fellow, and Mr. Spencer's family: wife, brother, and father, standing beside. Dr. D is talking when Mr. Spencer shows irregular heartbeat. The situation is becoming urgent:

1	Dr. D:	There is one one onh uhh
2		ad thought that I want to share
3		let's a <i>ssu</i> me next twenty four hours
4		you are on the highest urgency waiting list
5		and a ssu me that the arrhythmias are stable uh
6		BEE::P BEE:::P ((beeping sound from heart monitor for sudden arrhythmias))
7		which uh may not be the case
8		Uhm then we can wait for the heart
9		uhm uh for these next short time periods days=
10	Dr. S:	=did you just feel something just now
11	Mr. Spencer:	I did a little like a little bump

Dr. D is delineating the scenarios again. If arrhythmias remained stable (line 5), Dr. D was reasoning, but the loud piercing sound of the heart monitoring system alerted the room to the onset of new episodes of arrhythmias (ventricular tachycardia) (line 6). Dr. D repairs the sentence on line 5 with "which uh may not be the case" (line 7) and treads carefully not to exclude the possibility of waiting for a few days to see if a heart offer arrives while preparing for machine implantation (lines 12–15). A few days, a series of 24-h periods or the liminal waiting period for Mr. Spencer. Clock-time again enters this space in a strident juxtaposition to existential time.

Dr. S, who was inspecting the monitor latches into Dr. D's talk (line 10). Did Mr. Spencer feel the ventricular tachycardia (v-tach)? (line 10). He did; "a little bump" (line 11). The alarms marks Mr. Spencer's v-tach, his heart at 190 beats/minute; clock-time or existential threat for Mr. Spencer? There is a good chance that this unstable v-tach

will degenerate into the potentially lethal ventricular fibrillation (v-fib). The team needs to consider an additional emergency but temporary step: have Mr. Spencer on an artificial heart and lung machine (ECMO). Dr. D resumes his talk:

12	Dr. D:	yeah and if it's not stable and uhm
13		we are preparing for the mechanical support uhm
14		the preparation for the HVAD the uhm
15		implantable device just takes a bit=
16	Dr. S:	=a little bit
17	Dr. D:	and=
18	Dr. S:	=lem me ask you one thing uhm
19		if you do need need to go on ECMO though
20		just to get to the a safe place to get you to the VAD
21		would you have any issue with that
22		cause that's not a
23		that's something we need to address right now

Dr. D's talk is in stark contrast to the loud beeping sound of the alarm; he proceeds slowly, as if the situation, turning into a possible emergency, has no effect on his talk. He starts taking Mr. Spencer through the steps toward the prospect of needing an ECMO, pointing to the time it takes to prepare for BiVAD implantation surgery, a process of framing. It "just takes a bit" (line 15), but Dr. S intervenes (lines 16-18). As if in a relay race, he picks up the "bit" from Dr. D and runs with it. First with a calm but inadequately casual tone ("would you have any issue with that" line 21), not to alarm Mr. Spencer, as Dr. S discusses in cogen. Dr. S presents the prospect of going on ECMO a step "just" to get Mr. Spencer to a safe place (line 20); as if a simple decision on a preference and not on a significant component of life support can be made by Mr. Spencer. Then, with a sharp acceleration, Dr. S asks for an immediate verbal consent ("right now," line 23). The situation is escalating. Everybody can feel it in the room with the incessant beeping from the heart monitor. In cogen, both Dr. D and Dr. S recount their concern about Mr. Spencer's aversion to excessive medical interventions, and the impossibility of coming to terms with and agreeing to an ECMO if he coded because of v-fib.

Dr. S: you're constantly surveying a situation. At the beginning of this conversation, Mr. Spencer was discussing with me, and his vitals were stable. But with the occurrence of these events, you realize that the situation is changing. Mr. Spencer may arrest. You feel the urgency in my tone." Mr. Spencer could code, leaving the us physicians with only a slim opportunity to save his life.

A similar perception and understanding of the situation was expressed by Dr. D and the CCU Nurse, who in cogen reflects:

I felt he [Mr. Spencer] was, he was going to die. I really felt it because his v-tach. His v-tach was very fast, and his hemodynamics were deteriorating.

Dr. S's recognition of the urgency in his tone indicates what, in that situation, is salient to him to highlight and how he interprets it. His emotional engagement on what matters to him is consistent with Raia's (2020) definition of *existential time*. CCU Nurse N brings together the fear of having her patient die (what matters to her, Raia, 2020) with a clock-time analysis monitoring the irregular heartbeat of Mr. Spencer, the *bit bit bit* rhythm measured in clock-time.

As we have shown in the beginning, Mr. Spencer is incessantly against an additional intervention—"bridge" procedure; he probes deeper:

35	Mr. Spencer:	=three three procedures to get to a heart transplant
36	Dr. S:	It is it's a it's a bridge to a bridge so to speak

In the event that ECMO is necessary, Mr. Spencer counts the three procedures, ECMO, VAD implantation and heart transplantation, whereas Dr. S reformulates them, framing them as one continuum: a bridge-to a bridge-to heart transplantation, and at the same time, caring for Mr. Spencer, orienting him toward a definitive therapy, heart transplantation.

37	Dr. S	it's it's
38		but if you if something happens to you and
39		your heart stops? and people are doing CPR?
40		one of the options to try to get you out of it?
41		chemically and with chest compressions? uh and
42		hope that you come back? and
43		the other option-if those don't and
44		then withdraw if things are not going the way that
45		we want them to:?
46	Mr. Spencer:	o:r
47	Dr. S:	or we put you on ECMO and
48		that gives you a little bit of uh of t ime (.)

The situation can escalate, Dr. S continues: if Mr. Spencer's heart stops and he needs to be resuscitated one option is to try CPR (measured in compression per minute, clock-time), heart synchronization with defibrillation (i.e., electric shocks) and/or by chemical resuscitation. But then stop (withdraw, line 44) if there is no response. Or? Asks Mr.

Spencer. "Or we put you on ECMO." That gives Mr. Spencer a bit more time. A few days at most that Mr. Spencer would remember as an eternity.

62	Mr. Spencer:	no that's that's <i>three</i> (1.0) open heart (.) surgeries=
63	Dr. S:	=it's-not-open-heart
64		it goes through your groin (0.8)
65		we do it through the groin uhm
66	Mr. Spencer:	the pump hangs out of my leg
67	Dr. S:	the the <i>cannu</i> las hang out
68		the pumps-at-the-bedside are much like your uh
69		looks something you know
70		like something like this size right here it it
71		I'm not so sure on ECMO you would even be aware it's there (.)

Mr. Spencer has often discussed his reservation with excessive intervention to save his life, as he did with his wife when he overheard the electrical shocks, and the incessant pounding to resuscitate a patient next door to him in CCU "how far do we go as Christians to scratch and claw at hanging on to this life when we know that this world is not our home." The ECMO implantation is for him an extra procedure, to get to transplantation, and one too many.

As if to plug gushing water, Dr. S rushes to utter the words ("=it's-not-open-heart" (line 63)) and amends Mr. Spencer' understanding that ECMO is another open heart surgery (line 62). But the pumps would be hanging out of his body, Mr. Spencer retorts. The cannulas, rectifies Dr. S, will be visible hanging out. Rushing the words "pumps-at-the-bedside," Dr. S moves to show how small the pumps are to revise Mr. Spencer's perception. But maybe more is needed to convince Mr. Spencer that undergoing the ECMO procedure is essential to have a longer meaningful life with his family. Dr. S, as he commented in cogen, needs 'a nuclear option,' "think about your family *too*: when you make a decision" (line 78).

72	Dr. S	<i>okay</i> okay
73		let me let me just give you a suggestion cause uh
74		i'm a father and a husband <i>too</i> uhm (.)
75		it would
76		it could get you to the point
77		where you could get the VAD to the transplant (3.0)
78		and think of your family too: when you make this decision
79	Wife	ple:ase (6.0)
80	Dr. D:	maybe uh you uh
81		also want to talk about this uh for a [moment
82	Brother:	[((inaudible)) of us too (0.6)
83		please agree to it (2.0)

Dr. D supports Dr. S' move and calls on the family to speak more. The conversation continues with Dr. S clarifying to Mr. Spencer that the reason why he wants him to accept the possibility of an ECMO procedure is not because Mr. Spencer is unstable but on the account of the more complex preparation and the time required (up to 24 h) to initiate the HVAD BiVAD implantations surgery.

Different dimensions of time are at play here. The altered beating of the heart (v-tach) measured in beats per minutes (clock-time) changes the mood in the room—all fear for Mr. Spencer's life including Mr. Spencer—and marks a change in the emotional response made visible by Dr. S acceleration of speech and rush into completion of consent.

In cogen, the two physicians comment further:

Dr. S: In the back of my mind, I'm thinking this is a big undertaking. It could go awry. This could be terrible, it's the first one, and I'm advocating for something which I'd never done before. You know, we're going to do the best we can, which is bringing in the support of people who've seen it done before and can guide us and proctor us through this process. [...] As a surgeon, you have to figure out all this; and have to figure out how to portray confidence, but not cockiness, being upfront and forthright with the patient; confidence with full disclosure.

Dr. D: [...] At this part of the encounter, I feel like walking on eggs, you know—the two of us [Dr. D and Dr. S] think it should be done [agree to a possible ECMO implantation] and need to communicate it in a way that he [Mr. Spencer] doesn't shut off feeling he is pushed into authorizing what he may think are futile medical procedures. [..] My task is to deliver the patient to the surgical team in a meaningful way, that is, with an authentic decision made. Then I go home and have a glass of wine with my wife. Yes, it is really like the metaphor of the relay we discussed. Dr. S takes up from here. He calls home and says I'm not gonna be home tonight because I'm gonna spend the next eight hours in the operating room. So that's exactly how it feels.

Different time dimensions are also evident in the discussion in the cogen session. While Dr. D will measure time in existential time with his wife, Dr. S will not go home; he will spend eight hours in the operating room. Dr. S expresses also what Roth et al. (2008) describe as time imposed by institutional demands (we call *organizational time*). Implanting the new type of BiVAD will require Dr. S to assemble a larger team to include personnel from the company that makes the HVAD BiVAD, to follow the FDA guidelines ensuring maximum patient safety. He needs time to organize the team for the surgery and make sure he has the consent from Mr. Spencer to proceed. Dr. S will need to take these actions as soon as he walks out of Mr. Spencer's room so that the surgery can be done by the evening.

For Dr. S caring for Mr. Spencer and "returning him to his family" is what matters. In Raia's Relational Ontology, the *past* and *future* possibilities of being organize how things show up in the world as mattering in the present. Dr. S' description expresses an existential temporality as also reported to Mr. Spencer in the encounter:

95	Dr. S:	these are big undertakings I
96		you know I have to assemble not just our team but
97		I have to assemble the ou tside team
98		that comes in too that helps manage these devices-it-is
99		and as I <i>sai</i> d
100		the right sided device is not something that
101		we have a lot of experience with ((ECMO team enters))

Unexpected disruptions

Unexpectedly for Dr.S, Dr. D, Mr. Spencer and his family, the ECMO team enters the room (line 101). Its entrance could not have happened at a worse time, because, as shown on lines 101–105, not only does it interrupt Dr. S's explanation, but also, because the ECMO team is usually called when a patient has coded already and either had previously consented to an ECMO procedure or the consent process did not take place yet because the patient is incapable of consenting. This could suggest to Mr. Spencer that Dr. S and Dr. D are not taking into account his strong reservation against ECMO implantation. But Mr. Spencer has not coded and the doctors did not call the ECMO team.

With the entrance of the ECMO team, the speaker's orientation also changes from speaking *to* Mr. Spencer, into speaking to the ECMO team *about* Mr. Spencer. (e.g., Mr. Spencer "is talking to us," Dr. S insists, lines 107, 113, 120). In this situation, all participants have access to the conversation with the ECMO team. To show this change, we underlined the ECMO team members and those speaking to them (second columns in the transcript).

102 103	Dr. S	because it just hasn't been used for that ((ECMO team talks to CCU Nurse N in the back ground)) [it's it's like using uh
104	ECMO Team:	[((inaudible))
105		[it's it's it's
106	ECMO Team:	[we need shock him out of it
107	<u>Dr. S:</u>	he's TAlki ng to us ((turns to ECMO team))
108	ECMO M1:	just a second let us take a look at here
109	<u>ECMO M2:</u>	I think we should have art-line
110	<u>ECMO M3:</u>	уеа
111	ECMO M2:	just so you we know
112	ECMO M1:	yeah
113	<u>Dr. S:</u>	but he's TALK ing to us r eal ly
114	ECMO M2:	I just gonna ATP him
115		((inaudible))
116		is this the most recent pressure for the cycle
117	<u>RN. N:</u>	I just did one
118		I'll do another one
119	<u>Dr. S:</u>	Look at the heart wave form he clearly perfuses
120		he is talking
121		he said some sentences with this pressure
122	ECM0 M2:	I don't think you wanna (.) leave him in this
123	Mr. Spencer:	hrh hrh ((clears his throat))
124	ECMO M3:	Did it just start now
125	AdHF Fellow:	yeah while we were talking about it
126	Dr. D:	Uh uh
127	ECMO M3:	Are you guys scaring him?
128	ECMO M2:	eh eh
129	<u>Dr. S:</u>	No he had the
130	Dr. D:	I think we are scaring you
131	Mr. Spencer:	no:hh ((crying))

Upon entry into Mr. Spencer's room, the ECMO team confirms the emergency character of the situation wanting to shock Mr. Spencer to restore his heart rhythm (line 106), place an arterial line (art-line line 109) and initiate an anti-tachycardia-pacing (ATP-line, line 114). At the same time, the ECMO team needs to acknowledge that Dr. S is correct, something must be stable and not emergency-like requiring the three actions (lines 106, 109, and 114) immediately. Dr. S repeats three times that Mr. Spencer " is *TALKing* to us" (line 107) "*but* he's *TALK*ing to us r*eally*" (line 113), "he is talking" (line 120), indeed Mr. Spencer did not code! "Look at the heart wave" urges Dr. S to the ECMO team interpreting it as the sign of good perfusion (line 119). Yes, the waveform (the heart rhythm) trumps Mr. Spencer talking to the team in the interpretation of Mr. Spencer's ability to be present and capable of decision making.

The ECMO team continues talking with Dr. S and CCU Nurse N. In the meantime, Dr. D talks to Mr. Spencer and his family. Here, the pronoun "we" changes from "we", the hospitals teams, "are scaring you," Mr. Spencer (line 130) into indicating "we" Mr. Spencer *and* the AdHF team, "we have a decision in place" (line 133):

132	Dr. D:	I would think th	at would be pa	rt of the situation		
133		the important thing is				
134		we have decisions in place				
135		on the listing no	w that'll be for	malized		
136		number two				
137		we have a decis	ion in place of	biventricular assist device		
138		that should be if	f possible HVAI) BiVAD		
139		what we're righ	t now discussir	ig is		
140		the t i:me to asse	emble everythir	ng will take so:me		
141		and if we don't	have the <i>time</i>	uh		
142		we have right ne	ow available a	PVAD BiVAD but		
143		those hang out o	of the body:			
144		you have a state	ment in place	hat you don't want tha:t		
145		we have				
146		if more urgently	bridging to eit	her of these two		
147		a VA-ECMO requ	ired currently	a t hou ght that uh		
148		you would not v	vant that			
149		uhm this is by t	he way			
150		the VA-ECMO is	not the type of	assist heart pump surgery		
151		that of size of a	uhm mechanica	al support or heart transplant		
152		it's mo:re as we	call it a big ca	theter based uh approach		
153		So it's this uh				
154		it doesn't have t	he same quality	y and		
155		it's only for a fe	w da:ys to uh l	nave that longer term pump in place		
156		that would be a	n important in	my opinion		
157		a very favorable decision to make to uh uh to accept that				
158		but uhm just from my perspective				
159		it's not the same like three uh stage to heart transplant				
160		it's um [VA-ECM <i>[B</i>	10 you know <i>E::::</i>			
		BE:::::	BE::::	BE::::		

From line 134 to 142, Dr. D uses the pronoun "we" to include Mr. Spencer, his family and the AdHF team to summarize all the important decisions in place (i.e., lines 134, 137, 139, 141, 142). First, to accept heart transplantation ("being on the list" line 135). "Number two" (line 136), the decision to have a bridge to transplantation via biventricular assist device. Rather than having visual access to the blood rotating into the pumps, which Mr. Spencer would have if he had chosen the PVAD BiVAD implantation, Mr. Spencer prefers the HVAD BiVAD because the actual pumps are implanted inside the body (lines 137–141). At lines 144 and 148, Dr. D isolates Mr. Spencer's preference (you). The first time, line 144, points to the choice between PVAD BiVAD and HVAD BiVAD, the PVAD BiVAD requiring less preparation and organization and therefore less time during which Mr. Spencer can develop more arrhythmias. The second time (line 148) is the discussion about a possible ECMO procedure. In highlighting Mr. Spencer's preference "you" Dr. D voices to Mr. Spencer that his preferences are acknowledged, however, having defined the second "currently a thought" (line 147) Mr. Spencer's rejection of a possible ECMO procedure is not yet accepted as the final decision by the doctor.

Dr. D continues. He supports Dr. S's description, downsizing the ECMO implantation in comparison with HVAD-BiVAD implantation and heart transplantation. He defines it as a catheter procedure (149–154), and therefore, as Dr. S has done, *frames* Mr. Spencer's understanding of a three-stage procedure rather than two (BiVAD implantation with possible ECMO as a bridge to transplantation) or even one a bridge to a bridge to transplantation. By highlighting Mr. Spencer's preferences first and then at lines 155–156 and 157 stating his opinion, Dr. D points to the role distribution, a patient who has the right of making a final decision in consultation with a professional who needs to state his/her professional advice and therefore defining this process as a shared decision-making process. Dr. D expresses his assessment of the acceptance of the possibility of ECMO implantations as "a very favorable decision" (line 157).

After Dr. D's words, the intensity of the situation seems to have lost the turbulence generated by the sudden entry of the ECMO team, however as a coda, the loud sound of a beeper suddenly erupts, the ECMO team enters the conversation with Mr. Spencer putting electrodes on his chest and side (lines 160–162), the defibrillator's mechanical voice speaks the automated words: backup without signs of compression backup without signs of compression, bit bit bit the heart monitor marks the accelerated heart beats generated by a new v-tach episode.

161	ECMO M2:	we are puttin backup withou	g them as a ba at signs of com	ck-up now? pression
162		We are not pl	anning to give	you shock?
		backup withou	ut signs of com	pression
163	Dr. D:	(hhH)HHH ((deep loud sighs))
164	CCU Nurse N:	just putting th	nem on	
		bit bit bit	bit bit bit	bit bit bit
165	Dr. D:	(hhH)HHH sł	nare your thou	ghts anytime (hh)hh
		bit bit bit	bit bit bit	bit bit bit
166		I now want u	s to (.) work th	rough this you know
167	Mr. Spencer:	I don't want ((hh) <i>((as if a</i>	bout to cry))
168		stuff hanging	out uh	
		bit bit bit	bit bit bit	bit bit bit
169		except for an	electrical cord	or
		bit bit bit	bit bit bit	bit bit bit
170		that that's wh	ny that's accept	cable
		bit bit bit	bit bit bit	bit bit bit
171		it would let m	ne go h om e (0	4) for a while
		bit bit bit	bit bit bit	bit bit bit
172	Dr. D:	ye s bit bit bit	bit bit bit	bit bit bit
173	Mr. Spencer:	till we had a	transplant and	have a somewhat normal <i>life</i> =
		bit bit bit	bit bit bit	bit bit bit
174	Dr. D:	=the HVAD Bi	VAD <i>absolutel</i>	<i>y=</i>
		bit bit bit	bit bit bit	bit bit bit

bit bit bit bit bit bit bit bit bit. The 3 bits per second sound of ventricular tachycardia (v-tach) now is ongoing and clearly audible via the audio signal of the heart rhythm motor. The situation can turn into an emergency if the v-tach does not dissipate quickly as it did before. Dr. D's sighs (lines 163 and 165) are deep, as if conveying the gravity of what

is happening. Dr. D asks Mr. Spencer to share his thoughts, "I now want us to (.) work through this you know" (line 166) as, he had share in cogen "to deliver to the surgical team the patient in a meaningful way that is with an authentic decision made." Together they review the decisions in place once again: Mr. Spencer does not want to see the pumps pumping his blood outside his body; the electrical cord will connect the pumps inside his body to the batteries and computers (lines 168–169). He wants to be able to go home (line 171) and recover there with a somewhat normal life (line 173). Dr. D immediately translates Mr. Spencer's decision into the available choice: the HVAD BiVAD, emphasizing they are in agreement (*yes*, line 172; *absolutely*, line 174).

A faster pace, a reorganization of communication spaces

Dr. S joins Dr. D and Mr. Spencer. Dr. S intervenes (line 176), overlapping first with Mr. Spencer (lines 176) and then with Dr. D (line omitted) to describe the ECMO. However, Mr. Spencer is not interested in what the device is. He is interested in how the device impacts his short- and long-term life:

189	Mr. Spencer:	i'm laying flat on my back the whole time?
190	Dr. S:	you cannot walk with them
191		you're absolutely right
192		you lay on your back uhm

∅	Springer	

222

Dr. D:

193	Dr. S	MY FEELING is [that	
194	Dr. D:	[but it's [only also for a very few [days that	
195	Dr. S:	[yes [a day or two	
196	Dr. D	that you know	
197		this is not a (0.4) living on the VA-ECMO	
198	Dr. S:	no:	
199	Dr. D:	this [is a (.) an <i>acut</i> e support to	
200	Dr. S:	[no no right	
201	Dr. D:	a (.) decision making (.) [to do the uhm	
202	Dr. S:	[at most it's two to [three days	
203	Dr. D:	[a long term	
204		[assist pump	
205	Dr. S:	[while we get your	
206	Dr. D:	[a-few-days	
207	Dr. S:	[your situation [clarified and	
208	Dr. D:	[no-more-than that	
209	Dr. S:	make sure your brain	
210		is everything is okay after you went on it and	
211		then and then	
210 211 212		then and then we would go but it's not a long term uh=	

FRONT STAGE

BACKSTAGE

213	Dr. D:	=it's very similar conceptually to the uhm	Dr. S :	(turns to (talks to ECMO team and AdHF
214		impella that you had in Santa Monica		Fellow))
215		very very similar on the you know		((not audible))
216		it's a <i>sho</i> rt <i>te</i> rm support until a emh		
217 218		next step is taken for a few days		or redline him to the OR do we have everything in place for the VA:D
219		Yea::h		
220			AdHF F	I mean it would be okey
221	Dr. S:	Do we have all the administra	ative stuff in	n place?

Yes yes that's in place

Both Dr. D and Dr. S act immediately as they are used to in their respective practices, taking charge in an emergency. Their overlaps show their moves. Dr. S suspends answering Mr. Spencer's question and with a loud voice is shifting into a higher gear, moves fast into convincing Mr. Spencer of the benefit of the ECMO. Dr. D does the same. The apparent competition for the lead fast resolves in organized teamwork: Dr. D talks to Mr. Spencer and his family in what Goffman (1974) defined as a frontstage, while Dr. S organizes the plan in backstage, where Mr. Spencer and his family have no auditory access, a common division of space in medical care (Raia & Smith, 2020). A sudden change of a situation, from urgent to emergency, requires multiple rapid adjustments.

223		du:::m
224	ECMO:	I don't want to leave him in this?
225		We want want to ATP him our of it
226	CCU Nurse:	Would you like it in V1
		BE::: BE:::: BE:::: bit bit bit bit bit bit bit bit bit
227	Mr. Spencer:	I haven't changed my mind
		bit bit bit bit bit bit bit bit
228	ECMO :	Ok starting ATP guys
		bit bit bit du:m bit bit bit du:m bit bit bit du:m bit bit bit du:m du:m DU:::M DU:::M DU:::M DU:::M DU:::M DU:::M DU:::M DU:::M DU:::M DU:::M DU:::M DU:::M Hu:r Hu:r Hu: Hu: BE::: BE:::
229	Dr. D:	would you like a (.) moment with wife and dad without us being here
230	Mr. Spencer:	Uhm (10.0) ((the reast of the team talk to ecmo team in the background))
231		I don't I don't necessarily want you guys to leave
232		cause you're watching over my stability right now
233	Dr. D:	yes that's right
234	Mr. Spencer:	so what
235	Dr. D:	or uhm maybe we could just uh
236		can you both maybe also just share your thoughts ((addresses family))
237		come here yeah ((team talking in the background))
238		we can we can use this moment to to uh
239		respecting whatever you want to be done ((to Mr. Spencer))
240		to go through this right now
241		we have we have <i>all the</i> ti:me
242		you know?

The ECMO team charges the machine (line 223) to deliver electrical pacing impulses that are 10 bits/minute faster than Mr. Spencer's v-tach heart rate (Anti-Tachycardia-Pacing, ATP, line 225). The emergency situation does not change Mr. Spencer's mind about the ECMO procedure (line 227). The ECMO team proceeds to ATP (line 228). The v-tach stops. For now.

Dr. D asks Mr. Spencer if he would like a moment alone with his family (line 230). Mr. Spencer's wife and brother have already voiced their hope that Mr. Spencer would accept the ECMO procedure if necessary (lines 79 and 83). As Dr. D reflects in cogen, alone with the members of his family, Mr. Spencer would feel in a safe and trusted existential space where he does not feel pressured by the presence of and intervention by the AdHF team. But Mr. Spencer is scared to be left without medical supervision, watching over his stability (lines 231–232) and Dr. D repairs his action by inviting the family, who had retreated from the bed during the work of the ECMO team, to come back closer to Mr. Spencer, share their thoughts with the promise of respecting Mr. Spencer

preferences and supporting him in a shared decision-making process; "we have all the time, you know?".

Mr. Spencer's wife commented on that moment as follows:

He [Mr. Spencer] didn't want the the mechanical heart support [ECMO] and the family all decided to pray around him. Dr. D put his hand on [Mr. Spencer] 's arm and he stood there with us and he just waited for us to have that time together praying. His presence was just there the whole time. Very, very supportive. And I think that the decision had a lot to do with it.

Contra-punctual tempo

In cogen session, Dr. D comments:

Listening to this sentence, "we have all the time" I feel myself in that situation, trying to create an atmosphere where Mr. Spencer feels in a space that is not rushed, not driven, not motivated by the fear of dying and the associated anxiety in the team; the urgency, the indicators like the alarms and the behavior of people talking about the illnesses as opposed to talking about it with him. So I'm seeing myself actively creating a contra-punctual atmosphere; is like Joan Sebastian Bach's composition.

Dr. D contrasts clock-time with existential time as elements that are in contrast but all part of one composition. Dr. D continues:

It is interesting to see myself working on this. Just take what we just listen to, those five sentences [lines 136-241]. The last one of these is "we have all the time" [line 241], which is absurd; we have no time because right now he is dying. The sentence before is "respecting whatever you want" [line 239], and the sentence before this one is to his father, brother, and wife to stand by next to him. They were basically paralyzed and not interacting with him. So, I'm trying to ease them into saying something to this patient who is at risk of dying right now. The sentence before this is "share your thoughts" [line 236]. It's the first of the five. I went backward. "Share your thoughts," it is so important for me to listen to this; my voice is a contrapunctual to everything else that's going on. It's the opposite of the high pitch of all the alarms and alarmed voices. I am trying to own the space with the patient that allows for some existential and spiritual connections and against the gradient of the emergency.

The Resident doctor participating in cogen responds to Dr. D:

I totally agree. In this moment, Dr. S and the ECMO team, like everyone else, are moving in one direction: start ECMO! Save the patient! Like: we don't have ANY time. Dr. D and the patient are taking a moment, a different direction. The sentence 'we have all the time' was so funny to me; my heart rate started speeding up; even just listening to the situation, I felt like, there is no time! We have no time! Dr. D takes the power because he's making time, as if there was time, 'we are creating time' on the basis of the patient's values, dying versus getting a procedure against his will.

You [Dr. D] are balancing that all. I think it speaks to how well you [Dr. D] understand the patient, you know, and I also think the patient signals to you that he understands how seriously ill he is, and you listen. A lot of docs would think that the patient doesn't get it, you know because he is blocking a potentially lifesaving procedure. But Mr. Spencer understands; he says I don't think you should leave the room because you're watching over me because I'm unstable. So, he signals to you that he gets it, you get it. And the patient has the power to stop [the procedure]. So, both you and the patient are having the power in this moment because every other physician and nurse in that room are like, 'Go! Go! Go!' preparing for that moment, and the patient is saying: 'I'm not sure yet.'

Saying 'we have all the time' honestly almost makes you laugh in a way because it's so ... In a doctor-centered perspective, it's so crazy to say that because we don't have time. In the doctor's typical perception, it's like: we have to save this person's life, and we don't have that much time. The way we cope with that stress is to do something, you know, and they're all moving towards that, and it takes a lot to stop that momentum, yeah, and create that space. So, it is in contrast to everyone else's pace in the room; and also like me as a listener, because even me as a listener, I'm like, oh my God, he's going to code right there if we don't start ECMO. I know listening to this, I, I can't imagine doing what you did in this moment because I still have, I just feel the adrenaline. I think that would be really hard to do in that moment for me. I think it's very impressive.

More than an epistemologically defined professional vision

The above discussion among the healthcare professionals in cogen unpacks for us different stages of training (Resident, Fellow, and Attending) *highlight* the situation as relevant, pausing the recording to comment on this moment. They all *code* the situation as very urgent with the possibility of degenerating into an emergency, expressing a similar professional vision as previously defined by Charles Goodwin (1994). However, the emotional responses are different. While the Resident and the Cardiology Fellow, feeling the adrenaline rising, are drawn into the room's emergency mood, the AdHF Attending, Dr. D, acts differently. The Resident describes Dr. D's counter-punctual move to the emergency mood as "stopping that momentum" that was drawing all, including the Resident, into an emergency mood. Both the Resident and Fellow also recognize Dr. D's act as something that they have not yet mastered and a "very impressive" skill that helps the patient partake in a *shared decision*-making process.

When the emergency situation unfolds, both Dr. D and Dr. S act immediately as they are used to in their respective practices, taking charge in an emergency. They similarly highlight and code a change in the situation from urgent to emergency. What the Resident describes as the immediate stress reaction to a possible emergency "is do something," translates for the Attendings into immediately taking the lead of the situation. In this boundary space, the sudden change of a situation, from urgent to emergency, requires multiple rapid adjustments which resolve the apparent competition and moves the focus from the individual professional's capacity and skills to a teamwork context: Dr. D talks to Mr. Spencer and his family, while Dr. S organizes the plan. Within the larger background context of caring for Mr. Spencer, the local organization of the two professionals' responses is organized not only by what they highlight and how they code the situation at hand but, as Raia et al. (2021) discuss, also according to the diversified professional goals and anticipated actions. Dr. D's goal, as he has expressed above "is to deliver to the surgical team the patient in a meaningful way that is with an authentic decision made." Dr. S's goal is to consent Mr. Spencer and to immediately start the preparation for surgery, making sure that the procedure is safe and has a successful outcome.

The fast reorganization and division of backstage and frontstage activities shows how well Dr. S and Dr. D know how to rely on each other and support their respective professional goals and needs, which combined provide the care Mr. Spencer needs. Dr. S and Dr. D have been working closely together for some years, with the shared background of caring for the patient, they continuously monitor each other's differential actions, making sure they contribute to the complementary final goal of care. It is this cooperation that allows Dr. D to take a counter-punctual stream of actions that ensure that Mr. Spencer can have a safe space where to share his thoughts and pray with his family in a decisive moment of his care. It is based on this that Dr. S, as he also discussed in cogen, could, a few moments later, leave Mr. Spencer's CCU room to come back after Mr. Spencer and his family under the supervision of Dr. D had time to talk and pray together.

Framing-for and-with the other

Dr. D's particular counter-punctual move can only be successfully accomplished because in the past days Dr. D, Mr. Spencer and his family have built trustful relations through the process of *framing-for and -with the Other*. Dr. D, Mr. Spencer and his family together have confronted numerous uncertainties. Jürgen Kasper et al. (2008) identify multiple sources of uncertainty pertaining to: diagnoses, prognoses, and treatment of disease; patient's preferred degree of involvement, issues of patient social integration, and mastering of life requirements; and physician's competence and trustworthiness—which includes how information about risk is provided and how the patient deciphers it. Throughout the excerpts of two days, we have seen how Dr. D framed-for and-with Mr. Spencer these uncertainties, centering them around Mr. Spencer's expectations and existential queries. By analyzing talk-in-interaction, we showed how Mr. Spencer and Dr. D implicitly refer to previous acts of framing in their conversation. However, Dr. D did not only attend to Mr. Spencer's thinking or to his intellectual understanding of options and procedures but is *caring* for Mr. Spencer (Raia, 2020) as a person who is existentially shaken by the very real possibility of losing his heart, losing his way of life and facing death. Dr. D's acts of care can be accomplished by the doctor's talk, punctuated by silences and repetitions, creating an existential space in which Mr. Spencer's doubts on new uncertainties and complex options can emerge and be discussed and framed for and with Mr. Spencer in the moment. On New Year's Day, new uncertainties emerge: having to possibly receive an emergency bridge to a bridge to transplantation (ECMO), and meeting his surgeon for the first time and a need to assess the physician's competence and trustworthiness (Kasper et al., 2008). During the escalation from urgency to emergency situation, Dr. D creates again a space for Mr. Spencer's needs where there is "all the time" for Mr. Spencer to be able to pray with his family. Indeed, Dr. D creating a space for Mr. Spencer and his family to be able to pray, and partaking in it as a professional, identifies Mr. Spencer's spiritual life as salient. It is an existential *past* modulating Mr. Spencer's sonsibility to attune to what is relevant to him, and a *future* sense of possibilities of being himself so that Mr. Spencer can inhabit a meaningful and shared *present* with the physician to face the decision to accept an ECMO.

To care-for-the-other, catapulted into a world that is unfamiliar, where the person confronts choices that infringe on the lived experiences and where the person's needs and queries are often in sharp contrast to those acknowledged by the dominant culture required all participants to admit a complementarity of existential, organizational and clock-time in medical practice to help the person, Mr. Spencer, to be able to make sense of "what is that's going on here" and support more complex professional visions of different team members.

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