

Technology in Parkinson disease: Challenges and Opportunities

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ABSTRACT

In both daily practice and clinical trials, it remains difficult to reliably evaluate patients' functioning in everyday life, over long periods of time, and using methods sensitive to relevant milestones. Examples include fluctuating events (e.g., response to medication), rare incidents (e.g., falls), and non-motor complications (e.g., orthostatic hypotension). The miniaturization, sophistication, proliferation, and accessibility of sensing technologies are now enabling the acquisition of data suitable to capture more and previously inaccessible phenomena in Parkinson disease (PD). However, a gap remains between the amount of information gathered using these technologies and the corresponding insights into disease complexity that should be gained to satisfy diagnostic and therapeutic needs. Challenges include the variety of non-compatible technology platforms, the feasibility of wide-scale and long-term deployment of sensor technology (in particular among vulnerable elderly patients), and the gap between the "big data" acquired with sensitive measurement technologies and their limited clinical application. Major opportunities could be realized if new technologies are developed as part of open-source and/or open-hardware platforms enabling multi-channel data capture, sensitive to the broad range of motor and non-motor problems that characterize PD, and adaptable into self-adjusting, individualized treatment delivery systems. The International Parkinson and Movement Disorders Society Task Force on Technology is entrusted to convene engineers, clinical researchers, and patients to promote the development of integrated measurement and closed-loop therapeutic systems with high patient adherence that also serve to: 1) encourage the adoption of clinico-pathophysiologic phenotyping and early detection of critical disease milestones; 2) enhance tailoring of symptomatic

therapy; 3) improve subgroup targeting of patients for future testing of disease modifying treatments; and 4) identify objective biomarkers to improve longitudinal tracking of impairments in clinical care and research. This article summarizes the work carried out by the Task Force toward identifying challenges and opportunities in the development of technologies with potential for improving the clinical management and quality of life of individuals with PD.

INTRODUCTION

During the last decade, a multitude of technology-based objective measures of parkinsonian impairments have been developed, bringing with them the promise of substantially changing the diagnostic, monitoring and therapeutic landscape in Parkinson disease (PD). Sensors, mobile communications, cloud computing, advanced analytics, and the Internet of Things (wireless connectivity of all electronic devices),^{1, 2} are among the innovations that have the potential to transform healthcare and our approach to patients with chronic, complex and fluctuating disorders. With the abundance of new technologies, their growing power and versatility, and the smart algorithms upon which they increasingly rely, our field needs to ponder the basic questions of *why* we should even consider adding alternative measures, *what clinical needs* should be addressed, *what instruments* should be used, *how to deploy* new technologies with minimal burden, disruption and cost and maximal compliance, and whether they *replace or complement* existing resources. Unfortunately, technology developers appear to be operating in competing “islands of expertise” whose focus may be redundant, thus increasing the risk of duplicating rather than extending progress while potentially making their technologies incompatible with those of other developers. Also, not all technologies are primarily driven by burning questions from within the clinical field, sometimes creating technical solutions that – clever as they may be – remain in search of a clinical indication.

In the absence of well-established and validated biomarkers for diagnosis or disease progression, PD remains a clinically defined disease. Today, clinical scales and traditional patient-reported outcomes continue to be the primary assessment tools or

endpoints in both clinical care and research in PD. However, there is growing awareness that technology-based objective measures (TOMs) may improve the sensitivity, accuracy, reproducibility and feasibility of objectively capturing the full complexity and diversity of changes in motor and non-motor behaviors.³⁻⁷ Examples include the difficulty to reliably evaluate fluctuating events (e.g., the variable response to medication), to capture rare incidents (e.g., falls, or freezing of gait) or to assess behaviors that, by definition, take place over long periods of time outside the clinical examination room (e.g. physical activities in everyday life). The ability to remotely capture behavioral data and use it to optimize treatment strategies has the potential to finally “close the loop” and address critical knowledge gaps.⁸ Despite multiple ongoing projects by stakeholders in academia and industry, it remains challenging to determine what initiatives are ready to be scaled up, and what type of deployment would ensure the highest yield in the future.

This review summarizes the deliberations of the International Parkinson and Movement Disorders Society Task Force on Technology. As a first step, we centered on assessing the landscape of wearable devices and other technologies for individualized assessments, as well as the therapeutic and scientific uncertainties they stand to fill, rather than on assessing the clinimetric properties of any of the growing list of measurement technologies that have become available in the last 10 years. For the latter purpose, a number of reviews have been recently published.⁹⁻¹¹ The Task Force is entrusted with bringing together experts from the device and biopharmaceutical industry, clinical researchers, engineers, and patients to brainstorm on needed developments to advance PD research and care. We aim at appraising the extent to which technology and data analysis

in general, and TOMs in particular, can bring robust granularity to the clinical complexity of PD in order to facilitate clinico-pathophysiologic phenotyping, the detection of prodromal symptoms, the improvement of diagnostic accuracy, progression monitoring, and to begin the process of integrating technology-based diagnostics and actionable pharmacological and non-pharmacological therapeutics for clinical applications.

DEFINITIONS AND OBJECTIVES

TOMs are the outcome of device-based instrumented clinical tests conducted by clinicians in standardized environments to objectively *measure specific behaviors*, or self-administered by patients to *detect and monitor* impairments in specific or overall function in everyday life. Initially, TOMs targeted motor phenomena, such as gait or balance, and were gathered in specialized movement laboratories.¹² More recently, they have been extended to devices worn by the patient (i.e., wearable sensors and systems) in the clinic and –for remote monitoring– in the home or community settings (**Figure 1**).¹³⁻¹⁵ The goal of wearable technologies is to maximize the “ecological” validity as well as the temporal and spatial resolution of capturing motor and non-motor phenomena that are naturally expected to change over time. As such, wearable technology may provide a more “realistic” portrayal of behaviors of interest in clinical and research settings.¹⁶ In addition, in the research arena, increasing the number of data points through greater temporal and spatial resolution of a targeted behavior is expected to reduce the sample size required to evaluate the effect of therapeutic interventions assuming the ratio of signal to noise can be maintained or improved.¹⁷

Important goals of TOMs are to provide objective parameters in the detection and monitoring of motor and non-motor functions, thereby enhancing the quality of treatment delivery and allowing for personalized care (**Table 1**). Currently available wearable technologies (inertial sensors, surface electromyography [EMG]) are – with variable success – capable of capturing the number and intensity of multiple activities, such as the frequency and amplitude of movements during the day and while asleep, the frequency with which tremor and dyskinesia appear and disappear during the day, and the fluctuations in the severity of gait and balance impairments.¹⁸⁻²² The use of consumer wearable technologies in medicine is becoming increasingly more common. For instance, in the field of sleep medicine, the use of actigraphy for sleep monitoring is already starting to supplant more traditional methods like polysomnography due to its validity, lower cost, and ability to evaluate individuals in their homes over a longer period of time.²³ Advanced wearable technologies can also precisely monitor skin conductance, respiratory rate, blood pressure, oximetry and provide surface EMG, electrocardiography (ECG) and electroencephalography (EEG) tracings.²⁴⁻²⁹ Furthermore, the ability to collect TOMs using smart devices (mobile phones, tablets, and smart watches) provides additional opportunities to collect and analyze numerous clinically-relevant parameters (e.g., posture, balance, dexterity, voice and speech patterns, facial expression, eye tracking, medication and exercise compliance and adherence) and develop communication portals to improve patient engagement and self-management.

Caveat emptor: Why measure at all? The often-implied assumption that the sole existence of a PD symptom justifies its measurement and that all PD-related phenomena

should be measured must be dispelled. A measure is justified if it enhances our understanding of a complex disease or aids in testing or delivering a therapy. The use of measurements to improve therapy is filled with rich examples from other branches of medicine (e.g., glucose monitoring for insulin pumps, cardiac defibrillators). It should be remembered that every qualitative clinical assessment is a form of measurement and that the use of quantitative measures carries potential for improving the decision-making process as to the need and dose of therapy. Implicit however is that what is being measured represents a therapeutic target and hence the measurement must be relevant to the treatment question (**Table 1**).

The case for multi-domain, multi-sensor, integrated technology. PD is characterized by considerable inter- and intra-subject clinical variability in clinical symptoms. What matters most for one patient in the motor sphere may not be as important for another (e.g., tremor vs. freezing of gait vs. sleep disturbance) given the different levels of functional disability.³⁰ Fluctuations in daily functioning in some patients may only include non-motor phenomena (e.g., fatigue or anxiety).³¹ Even if we were to accurately measure the most overt deficit, most patients display a repertoire of motor and non-motor endpoints that vary within and between days, with varying impact on their quality of life.³² Thus, a multi-domain, multi-sensor, smart technology is needed to determine the source of all relevant changes, identify individualized disease fingerprints, and develop truly personalized therapeutic approaches.

CHALLENGES

The need for monitoring non-motor symptoms. The development of wearable systems to monitor individuals with PD has been focused heavily on motor aspects of the disease (e.g., tremor, bradykinesia, gait impairment, and dyskinesia)³³⁻³⁵ that are also, albeit with lower sensitivity and specificity, evaluated by clinical scales. Despite recent advances in the quantification of motor symptoms such as tremor, these endpoints often bear only modest quantitative agreement with measures of quality of life.^{36, 37} Indeed, patient priorities and sources of disability often arise from non-motor deficits (e.g., depression, anxiety, fatigue, orthostatic hypotension, sleep disturbance). Unfortunately, relatively few studies have thus far focused on capturing the fluctuations of these complex disease manifestations, marked by high variability within and between days.^{38, 39} The development of TOMs for non-motor endpoints has relied upon labor intensive or computerized, laboratory-based measurements (e.g., cognitive function, heart-rate variability, blood pressure changes, or sleep).^{40, 41} There is an urgent need for developing unobtrusive systems to monitor non-motor endpoints in the home and community settings.

Limitations of sensors used to monitor motor symptoms. Biomechanical sensors such as accelerometers, gyroscopes, and magnetometers are well suited for the detection of tremor, bradykinesia, gait impairment, and motor complications, such as dyskinesia. However, data collected in the home and community settings using these sensors do not always provide sufficient information to achieve a reliable clinical assessment of motor symptoms. For instance, it is difficult to infer from the sensor data alone if slowness of movement (as detected using biomechanical sensors) can be used as a proxy of

bradykinesia, or is the result of fatigue or other factors related to the context in which a motor task is performed (e.g., slow walking due to fear of falling). Also, the resolution of biomechanical sensors is quite restricted to the anatomical area on which they are applied, which may yield low quantitative agreement with the wider range of motor disability, quality of life, and other measurable patient-relevant endpoints.^{36, 37}

Discrepancy between clinical needs and research. Endpoints that may be ideally suited for a clinical study may not necessarily be relevant or applicable in clinical care. The relevance of specific TOMs to assessing the impact of parkinsonian symptoms on patients' quality of life may be difficult to evaluate. For instance, fluctuations in motor symptoms and complications such as dyskinesia may have a complex quantitative relationship with measures of disability.⁴² In addition, the measurement target and timeline of data capture differ depending on the goals of a study. For example, an instrumented test that captures finger tapping over several hours may suffice to track the immediate response to a dopaminergic therapy. However, monitoring disease progression over time involves more complex targets and longer data collection, such as physical activity levels, gait speed, rate of falls, and a variety of periodic or continuously gathered measures of motor, cognitive, or other non-motor functions. In many cases, accuracy and reliability of these TOMs may not yet be sufficient to justify their deployment in Phase III clinical trials.

Lack of compatibility among wearable systems. Most wearable systems developed to monitor individuals with PD are not compatible with one another. As a result, it may be

cumbersome or impossible to combine data gathered by TOMs developed by different manufacturers. This makes it difficult to guide behavioral changes or therapeutic interventions. Furthermore, devices developed by different manufacturers for the same purpose may not always yield the same result, raising questions about the validity of the mathematical algorithms that govern the data processing. Few currently available systems gather synchronized data from multiple body segments before transfer to a computer for whole-body analysis,⁴³ in a way that is fully compatible with the simultaneous use of platforms developed by different manufacturers

Limitations of available analytical methods. Despite our ability to collect and store extremely large datasets of TOMs, our ability to algorithmically analyze and synthetically display clinically and disease-relevant information to physicians and patients, remains limited. Here, clinical expertise is needed, for instance, to eliminate the “clinical noise” in the data analytical efforts. In addition, technical expertise is needed so that the field can take advantage of the tremendous advances that have been achieved over the past two decades in research areas such as signal processing and machine learning. Data analysis techniques that leverage advances in these research areas are important to achieving clinically meaningful TOMs.

Practical limitations in user engagement. Software and hardware components of wearable systems are often not as user friendly or compelling to adopt as they should be.⁴⁴ Currently, patient and caregiver engagement with wearable and mobile technology is modest, as shown by a recent study demonstrating that 32% of users stop using

wearables after 6 months, and 50% after just over a year.⁴⁵ Similarly, there is a high dropout rate amongst smartphone apps users: 26% of apps are used only once and 74% of apps are not used more than 10 times.^{45, 46} Lack of motivation to use wearables/self-monitoring systems should not be underestimated, particularly in the absence of meaningful feedback provided to their users. Preliminary evidence suggests that patient empowerment and their inclusion as active players in the development of research activities may favorably impact compliance.⁴⁷ Research is needed to determine the characteristics of wearable systems for long-term monitoring of motor and non-motor symptoms that would be acceptable to patients. In particular, we need to ascertain the number of sensors needed to accurately monitor PD symptoms without negatively affecting compliance in a clinical context.

OPPORTUNITIES

Wearable systems provide the opportunity to measure and monitor the individual variability of motor and non-motor phenomena, minimize rater bias, and increase sensitivity to subclinical but possibly relevant physiologic changes (**Table 2**).^{4, 10, 48, 49}

Standard measurement platform. Several companies have tested or are in the process of assessing a variety of methods to probe individual motor and non-motor constructs. To avoid duplication of investments and efforts, an opportunity exists to identify the technologies and approaches with most versatility, greatest ease of deployment, least patient and physician encumbrance, and lowest cost. It should no longer be a question of whether a given motor phenomenon can be measured in yet a different manner (which it

can), but on how to choose a standard platform of TOMs behind which developers and end-users can coalesce. Efforts towards standardization – guided by the MDS Task Force on Technology but endorsed or sanctioned by regulatory agencies such as the Food and Drug Administration (FDA) and the European Medicines Agency (EMA) – will greatly facilitate technology adoption and the integration of different systems.

Multi-domain measurements. With different types of disabilities comes the need for tailored measurement approaches to support the design of individualized interventions. Tremor measurement may be completely irrelevant to individuals with an akinetic or postural-instability gait-disorder (PIGD) phenotype of PD. Continuous step monitoring to capture freezing of gait episodes would be futile in patients without gait impairment. Systems that are designed for multi-domain data capture could provide researchers and clinicians with the flexibility of choosing the sensors to monitor individuals with different phenotypes of PD. In exploratory studies, an approach based on multi-domain data capture would increase the likelihood of finding relevant changes in one of the many “channels” of the system. This approach would also provide the opportunity for assessing correlated effects of symptoms of interest across other domains.

Better phenotyping and subtyping. Tremor and tremorless (akinetic/PIGD) variants of PD are grossly defined clinical phenotypes, based on mainly observational evidence, with substantial heterogeneity.⁵⁰ Besides these clinical phenotypes, there likely exist several disease subtypes defined by autonomic, cognitive or other domains of disability that could be captured by multichannel systems.⁵¹ In addition, it is conceivable that the

greater resolution of TOMS may detect novel phenomena that could serve to more sensitively stratify certain PD subtypes and serve as (or assist in the development of) biomarkers of disease progression.

Precision medicine. By identifying areas of dysfunction and their relationship with therapy, TOMs can be used to provide customized feedback to individual patients and possibly stratify criteria that “predict” the responsiveness to distinct treatment paradigms, in a way that is similar to how consumer-based wearable devices already measure level of activity, sleep disturbances, etc. Smart algorithms could be developed to generate specific recommendations that would be made available directly to patients and clinicians to motivate changes in treatment and lifestyle-related behaviors, tailored to each person’s specific individual needs and disabilities. This approach would provide value for end-users (both patients and their care team) and thereby improve adherence.

Closed-loop (feedback) systems. Data collected using wearable sensors could be used to trigger device-based interventions. Much as ECG sensing is used in cardiac defibrillators to trigger the delivery of stimulation pulses, data collected from sensors positioned on the limbs and trunk could be used to predict, for instance, the onset of a freezing episode. The system may detect an increase in cadence with a corresponding decrease in step length or a change in frequency of the lower leg oscillations.³⁴ The detection of such motor behaviors could trigger a device designed to deliver proprioceptive cues that could lead to a change in postural control and stepping pattern, that in turn could prevent a fall.^{43, 52, 53}

Real-time symptom tracking. TOMs could offer real-time, continuously captured, rater-independent data, in contrast with clinical assessments that rely upon subjective information gathered during sporadic, in-clinic evaluations.⁵⁴ Continuous monitoring of parkinsonian symptoms could replace diary-based recordings of fluctuations and be used to track periods of OFF, OFF with dyskinesia (not currently captured using the Hauser diary), ON, and ON with dyskinesia over the time span of several days.

The promise of remote monitoring. TOMs based on the use of wearable systems could improve healthcare delivery by providing assessment data when patients are not in the clinic. This possibility is particularly relevant for individuals with PD who live in areas with limited access to care. TOMs could provide ecologically valid data to help clinicians monitor responses to therapy and individualize management in order to optimize outcomes.⁴⁹ Remote monitoring also offers the opportunity for healthcare cost reduction.⁸

Better monitoring, better patient engagement, better outcomes. Innovation in sensor and communication technologies alongside mobile connectivity has enabled a process of “medical democratization”.¹ The creation or support of TOMs for remote, continuous monitoring provides an opportunity for healthcare providers to scale and extend services offered to patients in order to better manage their health.⁵⁵ TOMs can capture meaningful aspects of function that improve personalized patient care through an intuitive, interconnected, and energy-efficient interface.^{56, 57}

POTENTIAL PITFALLS IN DEVELOPING TOMs

Clinimetric validation pitfall. A number of studies have been focused on developing methods to derive TOMs that parallel the clinical assessment scales commonly utilized in clinic. These are important but potentially misguided efforts toward validating new TOMs at the clinic or home settings by attempting to force a simple quantitative agreement with widely used, previously validated subjective or semi-objective rating scales or questionnaires. However, it could be argued that, in theory, a “perfect” objective measurement should have a *complicated* quantitative match with an “imperfect” subjective one. This is because a clinical rater integrates many sources of information in order to produce a subjective score, including prior experience and expectations. So there is no *a priori* reason to believe that assessments performed by clinical raters would lead to a simple quantitative agreement with data features derived from sensor data. Indeed, the relationship between TOMs and subjective clinical scores may be highly complex and extremely difficult to ascertain in practice.

To the extent that we are seeking more sensitive and “ecologically valid” technologies, TOMs may agree only loosely with clinical scales. An important aim of TOMs is to improve upon, rather than act as surrogates of, previously developed clinical scales. As these are developed, clinicians and regulatory agencies will need to consider that a new TOM which appears to provide clinically relevant measures of movement characteristics, but which does not correlate with the motor section of the Unified Parkinson Disease Rating Scale [UPDRS], for example, could be accepted as valid on the basis of its own merits if it can accurately represent patient-relevant endpoints. Engineers and clinicians

alike should be reminded to “think outside the box” and use the power of the technology to develop new scoring paradigms rather than solely generate sensor-based versions of existing clinical scores.

Ecological validity pitfall. Efforts in ensuring validity, or the degree to which we are truly measuring what we intend to measure, increase in complexity with proliferating technologies, evolving in different platforms, and on different targets with unclear ecological and clinical relevance. To this end, before developing new TOMs, researchers would need to determine which constructs generated by routine clinical observations, and standardized by clinical scales and medical devices, are truly relevant to patients within such domains and are meaningful contributors to the performance of activities of daily living (ADLs). For example, if “dyskinesia” is not relevant as a construct to patients and is not a significant contributor to the performance of ADLs in their ecological environment, do we invest in maintaining its primacy in future technologies? Efforts to ensure system interoperability and to build open data repositories will help distinguishing relevant from futile TOMs.

“Big data” pitfall. It has been demonstrated that an abundance of behavioral data *can* be captured from individuals and populations using largely unstructured, “crowd-sourced” efforts. These data may differentiate populations of loosely defined “Parkinson’s disease” (on the basis of generic measures of movement abnormalities) from “normal” subjects. However, although these measures can provide valuable background information at the population and community level, they cannot substitute for a careful neurological

examination, deep clinical phenotyping, and assessment via laboratory studies. At best, they complement but do not replace the phenomenological and pathophysiological granularity required for PD subtyping, much less predict the response of an individual to treatment. Ultimately, the reproducibility and responsiveness of individually selected TOMs confirmed beyond small pilot studies and accounting for contextual information and confounders, should prevail over simply obtaining a large body of population-level data.

PREPARING FOR TOMS IN CLINICAL CARE AND THE RESEARCH SETTING

Wearable systems that are used to gather TOMs in the home and community settings could generate real-time, accurate, sensitive, and rich datasets including contextual information and data such as the time of medication and food intake. While TOMs are typically derived from wearable sensors, contextual information is captured using companion applications (e.g., mobile apps and web-based applications). Wearable systems that are used to gather TOMs also provide an opportunity for multidirectional interactions among investigators/clinicians and patients/caregivers at a reasonable cost. In the clinical care setting, the use of wearable systems to generate TOMs could decrease the need for outpatient visits while maintaining high-quality care and high patient engagement. Likewise, the integration of TOMs with virtual-visit interfaces has the potential to greatly improve the accuracy and value of telemedicine visits. In the clinical research setting, the use of wearable systems could enhance protocol adherence and patient compliance, leading to fewer missing data points. Also, an appropriate choice of TOMs could lead to prospectively collecting data with high signal-to-noise ratio (with

regard to the effect size of interest) hence reducing the required sample size and the resulting study costs. The composite of TOMs, companion applications designed to gather contextual information and pharmacogenomics could enable precision medicine interventions.⁵⁸

Integrating technologies. The development of new TOMs is currently advancing in isolated silos rather than as part of concerted actions aimed to implement open platforms. The development of open platforms would be highly desirable in the context of obtaining comprehensive information on patients and populations of interest. An open platform may not yet be fostered by the brave new world of health-driven technologies. However, there are reasons for optimism. While in the traditional medical device market, short-term financial forces drive the creation of proprietary measuring instruments at the expense of multi-channel, interconnected systems, consumer-driven market forces are pushing heavily in the opposite direction, i.e. towards the development of open technology platforms. As consumer technologies evolve to achieve the clinimetric sophistication required for application in the clinical management of individuals with PD, the move towards shared, interoperable software and hardware for applications in research and clinical practice is also emerging.

Smart delivery of treatment. Justification for the development and adoption of TOMs is strongest when presented in the context of improving the clinical management of individuals with PD. TOMs can be used as part of closed-loop systems designed to assist in the controlled delivery of medications. The development of such systems requires

manufacturing high-performance, energy-efficient and energy-harvesting sensors and storage modules.⁵⁹ In this context, the development of nanomembranes and stretchable electronics on a polymeric substrate for intimate mechanical contact with soft tissue has been proposed.⁶⁰ A critical unmet need is the ability to connect multi-sensor diagnostics to self-guided therapies in a closed-loop system. In the field of neuromodulation, deep brain stimulation (DBS) electrodes may also act as sensors, capable of recording local field potentials (for the presence or absence of beta-band oscillations) to automatically program the amplitude and frequency of stimulation, thus effectively closing the loop between measuring and treating.⁶¹

Objective measures of specific movement impairments can also be used to tailor therapy. Proof of concept systems have been developed to predict the onset of pathological tremor using surface electromyographic (sEMG) and acceleration data, which could inform the design of the next generation of non-invasive closed-loop predictive ON-OFF controllers for DBS.^{62,63} In the realm of physical rehabilitation, subtle asymmetries in gait, limitations in joint range of motion, or excessive postural sway indicating poor balance may be difficult to observe clinically but can be addressed by rehabilitation specialists when identified using TOMs. Through TOMs, therapists could personalize the therapy prescribed to each individual.⁴³ A simple clinical measure such as the time needed to walk a specified distance does not provide the therapist with an understanding of the spatial and temporal gait performance or the musculoskeletal and dynamic balance characteristics that cause poor mobility. These factors could be captured using TOMs and hence guide the choice of appropriate therapeutic approaches. Longitudinal monitoring of

TOMs also has the potential for identifying small improvements or declines related to the intervention or the progression of the disease that could lead to changes in the prescribed rehabilitation intervention.

Regulatory needs and commercialization. We anticipate that TOMs will eventually be routinely used in both clinical practice and research settings. Despite the promise of greater sensitivity and the presumed accuracy of collected data, regulatory validation of TOMs as efficacy and safety measures will require dedicated studies. The path to marketing for TOMs appears long and risky considering the short lifecycle of technological innovation and the costs associated with their development. Unlike drug development, where there is substantial precedent and a regimented path for marketing authorization, commercialization, and license protection, the path for TOMs and digital health solutions remains to be defined. It is critical that key stakeholders share the costs and financial rewards of technology development, implementation, and maintenance in order to accelerate and preserve innovation and growth. Despite opportunities to meet all stakeholder needs, the business model for development and deployment of TOMs in healthcare remains to be determined. Currently, healthcare payers show little incentive to financially reimburse TOMs, despite the promise for healthcare cost reduction, population management and delivery on high quality, efficient care. The lack of incentive may be driven by initial expenditures and the complexity of a rapidly evolving, but not yet fully integrated, technology market. Providers are also reluctant to fully adopt TOMs despite early evidence that they can improve patient outcomes and lead to overall improved care and patient satisfaction. Clinicians may not yet view TOMs as an

opportunity to support clinical decision-making and increase productivity. Patients are also reluctant to pay out of pocket despite opportunities for improved access to better care and better outcomes. Building a solid ‘business case’ – including properly designed cost-effectiveness studies – is much needed. A value-based care approach could be an attractive solution, where deployment of TOMs is funded as part of an integrated care solution where providers are rewarded for good outcomes per invested dollar, and where the decision to engage TOMs is left to the providers and patients. Finally, in addition to funding agencies, regulatory bodies such as the FDA and the EMA should increase efforts toward establishing programs that encourage the adoption of standards aimed to assure interoperability of wearable systems and the development of open data repositories.

Integration into medical care and reimbursement. Payers do not yet provide reimbursement for medical services provided by TOMs and companion apps. This limits the rate of innovation and the opportunities for integration of TOMs into medical care. Establishing reimbursement mechanisms will require demonstration that, along with the enhancements in diagnostics and therapeutics, TOMs can be integrated in quality control concepts, help reduce costs, and improve patients’ quality of life while guarding against privacy concerns. Quantifying clinical benefits of interventions using TOMs is anticipated to become increasingly important in healthcare as the allocation of resources is expected to be tied to objective outcome measures.

CONCLUSIONS AND NEXT STEPS

Despite challenges, the continuous improvements in technological sophistication, versatility, and wearability of sensors have created opportunities to collect disease-relevant data using targets consequential to patients and sensitive to PD-specific symptoms and milestones. In order to translate these opportunities into enhanced care, better self-management options for PD patients, and overall improved health care outcomes, technologies will need to be 1) developed as open platforms and integrated with electronic medical record systems, 2) suitable for the acquisition of data that captures motor and non-motor phenomena, and 3) integrated in treatment delivery systems. The International Parkinson and Movement Disorders Society Task Force on Technology aims at reversing the current model of simply adapting available technologies to meet patient management and research endpoints. As such, the Task Force will assist in improving the academic and regulatory environments for technology developers by encouraging the sanctioning of open standard platforms for technology-based measurements and treatments by, e.g., the FDA and EMA. This collaborative endeavor will materialize in the development of integrated, multi-channel, and in many instances closed-loop feedback systems that can achieve more sophisticated clinico-pathophysiologic characterization, better informed tailoring of symptomatic therapy, greater patient engagement and self-assessment, and better subgroup targeting of patients for testing of future disease-modifying treatments.

Authors' roles

1. Research project: A. Conception, B. Organization, C. Execution
2. Manuscript Preparation: A. Writing of the first draft, B. Review and Critique

AJE: 1A, 1B, 1C, 2B

PB: 1B, 1C, 2C

FN, WM, JMD, JK, BME, AM, FH, AEL, RR, JG, AN, MH, MAL, IL, TS, RD, MAB,

KK, AK, CG, J-FD, GM, LK, JMH: 1C, 2B

BRB: 1B, 1C, 2B

SP: 1A, 1B, 2B

Figure Legend

Figure 1. Schematic representation of a subject undergoing monitoring in the home setting using wearable and ambient sensors. The technology shown includes a wireless unit strapped around the wrist, band-aid like sensors attached to the lower limbs, a wearable camera worn as a pendant, a smart watch, and a mobile phone clipped on the belt used as gateway to relay the data to the cloud to assess specific functions (using its embedded sensors) as well as to communicate with the subject (using customized apps). Ambient sensors and computer technologies are utilized in the home settings to gather additional information or replace wearable sensors when wearable sensors cannot be used. The integration of wearable technology with smart devices enables remote monitoring of patients with PD and real-time feedback to clinicians, family/caregivers, and the patients themselves.

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Prof. Bloem currently serves as Associate Editor for the Journal of Parkinson's disease, serves as editorial board member of Physiotherapy Canada, has received honoraria from serving on the scientific advisory board for Danone and Zambon, has received fees for speaking at conferences from AbbVie and Teva, and received research support from the Netherlands Organization for Scientific Research, the Michael J Fox Foundation, the Prinses Beatrix Foundation, the Stichting Parkinson Fonds, the National Parkinson Foundation, the Hersenstichting Nederland, and the Parkinson Vereniging.

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