

## Review

# Video-Based Analyses of Parkinson's Disease Severity: A Brief Review

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Accepted 10 February 2021

Pre-press 3 March 2021

**Abstract.** Remote and objective assessment of the motor symptoms of Parkinson's disease is an area of great interest particularly since the COVID-19 crisis emerged. In this paper, we focus on a) the challenges of assessing motor severity via videos and b) the use of emerging video-based Artificial Intelligence (AI)/Machine Learning techniques to quantitate human movement and its potential utility in assessing motor severity in patients with Parkinson's disease. While we conclude that video-based assessment may be an accessible and useful way of monitoring motor severity of Parkinson's disease, the potential of video-based AI to diagnose and quantify disease severity in the clinical context is dependent on research with large, diverse samples, and further validation using carefully considered performance standards.

**Keywords:** Parkinson's disease, video, artificial intelligence, machine learning

## INTRODUCTION

The objective assessment of motor severity in Parkinson's disease (PD) is a major priority not only for the clinical follow up of individual patients and their objective response to drug changes, but also in the evaluation of experimental approaches in clinical trials. Since the COVID-19 crisis emerged, routine in-person assessment of PD severity has become impractical or undesirable in many cases; many older patients diagnosed with PD are considered part of the 'at risk group' and have been formally advised to shield [1]. This has resulted in remote/video approaches being encouraged to support patients suf-

fering from chronic illnesses such as PD [2–4] as well as in current clinical trials of PD [5].

Video assessments can also facilitate the evaluation of patients in the absence of their regular dopaminergic medication, which can be useful in the assessment process for treatments such as deep brain stimulation (DBS) surgery, where an "off" and "on" medication assessment is required as standard practice. Avoiding the need to travel to a clinic or hospital in the "off medication" state can be far more comfortable for patients, may reduce the duration of the time they spend in a suboptimal state, and reduce expenses due to travel costs and parking fees.

Alongside this, there have been a number of attempts to develop the automated rating of PD severity using video-recordings and Artificial Intelligence (AI)/Machine Learning techniques. This may provide support to clinicians when identifying and diagnosing

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disease. Given this growing interest in using a) video based/ remote assessments to assess PD severity, and b) AI rating of PD, this review aims to discuss the major issues that must be recognised as part of the potential role of video assessment and analysis in future management of PD patients, in the context of the growing use of Digital Health Technologies in PD.

## CLINICAL TOOLS

The most widely used tool for the assessment of PD is the Movement Disorder Society Unified Parkinson's Disease Rating Scale (MDS UPDRS). Part 3 of the MDS-UPDRS is the gold standard assessment tool for measuring motor signs of PD [6, 7]. Other validated scales exist for the evaluation of dyskinesia and tremor, but for the purpose of this review, discussion will be limited to the MDS UPDRS part 3, although the same principles may apply to the other scales.

## LITERATURE SEARCH

We searched PUBMED, Web of Science and SCOPUS i) from 2000 to 2020 using "Video assessment", "Parkinson's disease", "Telemedicine", "Telehealth" and "MDS-UPDRS" and ii) from 2016 to 2020 using "Video assessment", "Artificial Intelligence", "Machine Learning", "Automated" and "Parkinson's disease" "Motor symptoms" as key words. Reference lists from the identified articles were cross-checked to identify any other potentially eligible studies.

## STUDY SELECTION

We included observational and experimental studies conducted in PD patients in which i) items of MDS-UPDRS part 3 measured by a clinician through videos, were utilised as at least one of the outcome measures and ii) studies that used motor symptoms of PD measured by a machine learning algorithm relying on video-assessments. We excluded reviews and studies written in languages other than English. All retrieved abstracts were independently screened. The full texts of potentially relevant articles were retrieved for further assessment and were included if they met the above criteria above.

## CAN PD MOTOR DISABILITY BE ADEQUATELY CAPTURED ON VIDEO?

### *Direct comparisons of video-based and live evaluations of the MDS-UPDRS*

Before considering whether computer vision or machine learning techniques can improve upon traditional human rating of PD patients, it is important to consider the initial impact of using video rather than live face to face examination. Video-based administration of the MDS-UPDRS motor section has been successfully explored as a way to measure motor function of patients with PD [8–14], even among older individuals with PD who have substantial disability [15]. Studies directly comparing video-based and face-to-face scores of the MDS-UPDRS have shown moderate-good agreement, with intraclass correlation coefficients (ICC) ranging from 0.53–0.78 [9, 12, 15, 16].

Closer scrutiny of the studies comparing video-based and face-to-face analyses of PD symptoms reveals poor agreement however for specific elements of the MDS-UPDRS. Whilst studies have shown good agreement between live and video evaluations for scores of postural stability and gait [17], the same investigators have shown poor agreement for items of bradykinesia [10, 17] and tremor [17]. It is thought that because the tasks measuring bradykinesia involve rhythmic and continuous movements, technical difficulties such as poor internet connection, time lags and motion blur could affect accurate scoring [13]. In addition, multiple elements are included in rating bradykinesia, which may add greater complexity to assessing these tasks compared with rating more uniform manifestations such as gait and balance [18, 19]. There are also reports of difficulty rating rest tremor via videos due to the manner of rating tremor amplitude in centimetres, which may not be easily discerned [13].

However, a recent study comparing video and in-person assessments of upper limb function, which utilised a series of standardized measures including motor speed and tremor in 21 patients with PD, found good agreement across all measures, with ICC ranging from 0.75–0.99 [20]. It should be noted here that participants had access to high speed Internet and measures were completed in real time via Skype. Data from the PREDICT-PD study corroborates this view, demonstrating that subtle subclinical signs of PD can still be discerned from video assessments [21]. Whilst much of the research discussed

140 above demonstrates that video assessment can serve  
141 the clinical management of PD [8–14], a recent  
142 analysis of the STEADY-PD trial has shown that  
143 virtual visits involving video-based assessments of  
144 motor symptoms, is both feasible and comparable  
145 to in-person assessments [15], which suggests that  
146 video-assessments may also be of use in clinical  
147 trials. The current research into video assessments  
148 of PD motor symptoms has shown that video-based  
149 assessments can be carried out in patients own homes  
150 [12, 13, 16, 22] as well as in clinic [9, 17]. Method-  
151 ological issues with conducting video assessments  
152 in these respective environments are described in  
153 Table 1 which also outlines some of the other chal-  
154 lenges associated with video assessment of PD motor  
155 severity.

156 It should be noted here that the majority of  
157 studies were video telehealth consultations, where  
158 video-based MDS-UPDRS part 3 assessments were  
159 carried out as a secondary interest [8–16]. Stud-  
160 ies conducting formal video-based MDS-UPDRS  
161 motor assessments measured only some items of the  
162 MDS-UPDRS [20] or did not compare scores with in-  
163 person assessments [21]. No study to the best of our  
164 knowledge has validated video-assessment against  
165 in-person assessment of the MDS-UPDRS.

### 166 *Incomplete assessment*

167 During video administration of the MDS-UPDRS,  
168 there is an inability to perform parts of the motor  
169 exam such as assessment of rigidity and postural sta-  
170 bility, which require a hands-on assessment. Asking  
171 an untrained carer or family member to perform a Pull  
172 test to assess postural stability may lead to falling and  
173 injury. Many studies investigating video versus live  
174 administration of the MDS-UPDRS therefore omit  
175 these items from the assessment [9, 10, 12, 15].

176 The inability to assess patient rigidity may rep-  
177 resent a major limitation for video assessment  
178 especially among individuals in whom this is a major  
179 feature. More importantly, the restricted examination  
180 of patients (when limited to the MDS-UPDRS), may  
181 not detect the presence of co-morbid signs contrib-  
182 uting to a patients' disability. An example may be a  
183 patient with progressively worsening balance due to  
184 cervical myelopathy or sensory neuropathy, which  
185 may only be evident following examination of ten-  
186 don reflexes or distal sensory examination. While  
187 the MDS-UPDRS score is designed to be used with-  
188 out direct interpretation whether a change in score  
189 is due to PD progression or not, day to day clinical

190 evaluation of patients needs to consider whether other  
191 explanations may exist for a change in PD severity  
192 suggested by MDS-UPDRS part 3 scoring.

193 Despite these concerns, there are data to show that  
194 a modified MDS-UPDRS, in which elements that  
195 require a physical exam, including postural stabil-  
196 ity and rigidity, are excluded from rating, can remain  
197 a reliable and valid assessment of motor function.  
198 In a secondary analysis of the CALM-PD clinical  
199 trial, which compared a modified MDS-UPDRS to  
200 the standard motor UPDRS (including all items),  
201 found that the modified versus standard UPDRS  
202 was cross-sectionally ( $ICC \geq 0.92$ ) and longitudi-  
203 nally ( $ICC \geq 0.92$ ) reliable and valid [25].

204 Nonetheless, the use of objective measures such  
205 as wearable sensors may be used in conjunction  
206 with video assessment, to partially compensate for  
207 the missing data from the MDS-UPDRS scores for  
208 these items. A recent study with 32 patients with  
209 PD found that patient-worn wearable sensors com-  
210 bined with machine learning techniques were able to  
211 accurately predict clinician-assigned MDS-UPDRS  
212 scores for rigidity in 85.4% of cases [26]. Likewise,  
213 wireless accelerometers have been shown to suc-  
214 cessfully detect postural instability in patients with  
215 PD, produce scores that correlate with scores from  
216 gold standard assessments, and detect slight postu-  
217 ral abnormalities in early PD [26–32]. Consequently,  
218 video-based analyses in conjunction with the support  
219 of wearable sensors may give us the means to create  
220 an ecologically valid clinical picture remotely. Whilst  
221 a discussion of wearable sensors is beyond the scope  
222 of this article, wearable sensors have a large role in  
223 Digital Health Technologies and will be covered in a  
224 separate article in this issue.

### 225 **COMPUTER VISION VIDEO ANALYSIS**

226 Despite the challenges outlined, the recording of  
227 movement using video opens the possibility of using  
228 AI/Machine Learning techniques to quantitate human  
229 movement, which may be potentially useful in the  
230 diagnosis of movement disorders such as PD, and  
231 their longitudinal assessment. This has the theoret-  
232 ical advantage of improved objectivity and access and  
233 therefore improved signal to noise ratio in compar-  
234 ison to clinician ratings, inevitably subject to fatigue  
235 and intra- and inter-rater variability.

236 Computer vision defines humans as articulated  
237 objects with parts moving according to these artic-  
238 ulation points. Detecting human poses from a  
239 single viewpoint presents many challenges given the

Table 1  
Identified challenges and potential solutions associated with rating motor symptoms of Parkinson's disease via video

	Challenges	Potential Solutions
Video quality	In a recently completed study using state-of-the-art tablet technology in which the MDS-UPDRS was conducted via Facetime on 10 participants with PD, neurologists still reported difficulty in detecting quick involuntary movements and small amplitude tremor due to the quality and resolution of videos [12].	Wifi/ 5G
Patient Burden	<p>More elderly people are the most infrequent users of technology and the internet [23] and therefore, patients may have issues accessing technology devices and the internet in their own homes, particularly if they have cognitive impairment [24].</p> <p>The majority of the current research comparing video-based and face-to-face analyses of PD, focuses on samples that are relatively younger, highly educated, familiar with the Internet and present with milder symptoms of PD [8–10, 12, 13, 16, 17].</p>	<p>Secure web based uploads. High resolution video cameras Assistance for remote monitoring of more disabled patients with PD is essential. Moreover, these latter individuals represent the group most at risk of poor outcomes if exposed to infections such as COVID19.</p> <p>Replicate studies in the broader population living with PD.</p>
Inadequate patient visualisation	<p>Studies conducting video-assessments of motor symptoms in patient's own homes have presented challenges with the environment such as space constraints [13, 22]. This may make it difficult to visualise the patient's entire body, which is required for full assessment of the MDS-UPDRS [13] and their gait.</p> <p>Some studies are carried out at designated facilities with nurses on hand to assist with video set-up and in-person administration of the MDS-UPDRS [9, 17], which may not be conflated with findings from video assessments carried out in the home environment, which is unlikely to have the presence of a qualified clinician. Other studies that demonstrate the MDS-UPDRS conducted via videoconference in patient's own homes [12, 13, 16, 22] is of course of greater relevance in the context of patients potentially shielding from COVID-19. On one hand, conducting an assessment in the patient's natural environment may provide ecological validity of the clinical picture. However, longitudinal comparisons of scores may need to consider the context in which the video examination was performed.</p>	<p>Ensure adequate camera position and request (if possible) a family member to support the set up of video equipment.</p>
Inconsistent Video setting		<p>Compromise may be necessary on occasion according to disease stage/ purpose of evaluation.</p>

## Short-term evidence

The majority of research considering video-based assessment of PD is limited by short-term studies. Concrete conclusions cannot be made about the long-term use of videos to analyse PD symptoms, nor the validity of video-based methods in replacing face-to-face assessment.

By contrast, one longitudinal study that compared in-person assessments with virtual visits in 195 patients with PD found no difference between groups in MDS-UPDRS score changes as well as no difference between groups in standardized measures of quality of life over 12 months [8]. This suggests that video-based analyses of PD can be used over a medium term period with low risk to patient's clinical outcomes or quality of care.

Future longitudinal research may give insight into the efficacy of longer-term video-based assessments of motor symptoms with built in mechanisms to trigger face to face clinical examinations when necessary.

complexity of human structure. Put simplistically, computer vision uses low level features such as edges, shapes, colour, texture, and combines these with higher level features such as context and motion, prior models of human body parts and enhanced deep learning algorithms to assemble a human body model from a two dimensional image (2D) [33].

Several companies and academic research labs are attempting to develop machine-learning algorithms to aid in the measurement of PD severity. Strategic differences exist between them, either to provide an AI estimate of the modified MDS-UPDRS, to provide an AI estimate of sub-items of the MDS-UPDRS, or to provide an AI estimate of movement fluidity independently of the items incorporated in the MDS-UPDRS. All of these commercial ventures have challenges to overcome. The success of the pose estimation depends on many factors, such as the whole body being captured in the image as well as many additional issues relating to the lighting level, the background, the possible presence of other people, all of which represent major challenges for computer vision to recognise the human pose on a 2D image. Additionally, there are challenges involved for participants unable to follow the instructions appropriately leading to incomplete or inconsistent data.

In a recent study combining video-based analyses with machine learning techniques, severe motion blur and fluency issues with videos made it difficult for the AI system to score aspects of bradykinesia in 60 patients with PD [34]. Rating items of bradykinesia using the MDS-UPDRS requires scoring the fluency and quality of movement, and likewise characterising small amplitude tremor relies on discernment. Work utilising computer vision-based methods should therefore consider that accurate scoring of these items via video relies on the quality of video. Table 2 summarises the progress made by difference commercial approaches to video analysis of PD. Discussion of the machine-learning techniques utilised in the examples presented in Table 2 are beyond the scope of this review, please see [35] for further reading.

Beyond the demonstration that technical difficulties can be overcome, the interface has to be acceptable to the users (both patients and clinicians) and the data needs to be safely and securely stored (with informed consent) while meeting data protection requirements. Ultimately the analysis pipeline should be as automated as possible while still needing quality control checks to ensure a patient has engaged properly with the appropriate movement item. See

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Table 2  
Commercial approaches to video-analyses of Parkinson's disease

Company	MDS-UPDRS task measured	Description and method	Examples and further reading
Parkinson's Analysis with Remote Tasks (PARK)	Measures some items of the UPDRS to derive a composite score that indicates disease severity. This includes: facial expression, speech, finger taps, hand movements, pronation/supination, postural tremor, speech.	An online framework (available at parktest.net) that uses deep learning which allows patients to perform this abbreviated version of the MDS UPDRS score using a computer from anywhere in the world [36].	The framework can automatically discriminate between PD vs. non-PD with 82.5% accuracy [37]. They additionally inform subtle differences in facial expressions between PD and non-PD individuals that are possible for machines to quantify, invisible to a human eye, but The use of facial information to diagnose and quantify PD is acknowledged as a separate field in the assessment of PD and is beyond the scope of this review. <a href="https://machinemedicine.com/">https://machinemedicine.com/</a>
Machine Medicine Technologies	Measures all items of UPDRS to derive a composite score that indicates disease severity	Incorporates the traditional clinician rated MDS UPDRS onto a smartphone/tablet capable of recording segments of video footage which can immediately be quantified, using novel AI algorithms, according to item e.g. Right hand finger taps, with the aim of deriving a modified AI UPDRS score which they have termed "Kelvin UPDRS" (See Fig. 1a).	
Tencent Medopad	Diagnoses and quantifies the severity of PD using far fewer items than the traditional UPDRS.	Uses an AI system through a smartphone app that assesses a patient performing hand movements by identifying frequency and amplitude of movement to generate a score for the patient, which determines the severity of their PD.	<a href="https://www.tencent.com/en-us/articles/2200927.html">https://www.tencent.com/en-us/articles/2200927.html</a>
OpenPose	This includes: upper and lower limb movements. Quantifies gait not based on the MDS-UPDRS	Developed a deep learning-based system capable of discriminating, parkinsonian from normal gait, based on estimating cadence of periodic gait steps from sequential gait features via 2D videos taken on any accessible device	Estimated cadence of gait from the sequential gait features using the short-time pitch detection approach and combined machine learning algorithms and found discrimination performance for detecting mild PD gait from healthy controls was 0.754– 0.957 (AUC). Furthermore, when comparing gait sequences before versus after DBS treatment had a discrimination performance of 0.980 (AUC) [38]. OpenPose was able to estimate gait parameters relative to gold standard measures in healthy adults with a high degree of accuracy showing strong Pearson and intra-class correlation coefficients (0.671– 0.964) [39].

Table 2  
(Continued)

Company	MDS-UPDRS task measured	Description and method	Examples and further reading
Convolutional Pose Machines	Quantifies toe tapping and leg agility items of the UPDRS and derives dyskinesia severity by quantifying dysfunction during a communicating and drinking task according to the Unified Dyskinesia Rating Scale (UDysRS).	A deep learning based system that analyses 2D videos to a). automatically quantify parkinsonism according to toe tapping and leg agility items of the UPDRS and b) detect and estimate the severity of levodopa induced dyskinesia according to communication and drinking tasks measured by the Unified Dyskinesia Rating Scale (UDysRS).	Convolutional Pose Machines have been used to identify features of movement trajectories (e.g., kinematic, frequency) and these features were used to train random forests to score severity of parkinsonism and levo-dopa induced dyskinesia (LID). Found AUC for detecting LID was 0.930, severity estimation: $r = 0.661$ . For parkinsonism, leg agility was better for severity estimation ( $r = 0.618$ ) while toe tapping was better for detection (AUC = 0.773) [40].
Deep Lab Cut	Quantifies finger taps based on the MDS-UPDRS.	A system using transfer learning with deep neural networks that incorporates UPDRS measures of bradykinesia, which can track finger tapping in standard smartphone video recordings and automatically quantify speed, amplitude and rhythm of movements.	Computer measures derived from coordinates produced by DeepLabCut video tracking correlated highly with clinical ratings of bradykinesia (Spearman correlation coefficients: $-0.56$ speed, $0.61$ amplitude, $-0.50$ rhythm for MDS-UPDRS, $0.69$ combined for MDS-UPDRS) [41].
Microsoft Kinect	Identifies gait features associated with clinical scores of UPDRS gait measures.	A motion sensing input device to record standard colour videos and provide coordinates of 25 body joints in 3D space. Uses an inbuilt depth sensor, containing a monochrome Complementary Metal Oxide Semiconductor (CMOS) sensor and infrared projector that creates 3D images, combined with a measure that uses infrared light to capture 3D movement patterns.	Videos of walking bouts of natural gait were collected using a Microsoft Kinect sensor and onboard color camera which were processed to extract 3D and 2D gait features. Extracted features were put into subsequent regression models to identify features which correlate with clinical scores. It was found that measures of gait extracted from videos were significantly associated with UPDRS-gait clinical measures with regression models achieving accuracies of 61.4% and 62.1% for 2D and 3D features, respectively [42].
		Utilised clinically by incorporating regression models for gait assessment in neurological diseases.	

292 Figs. 1a and 1b that depict two example processes of  
293 using automated video assessment tools.

294 The development of such tools and applications to  
295 an illness such as PD requires extensive model opti-  
296 misation using data from large numbers of individuals  
297 and then complex validation with careful considera-  
298 tion of the gold standard against which the tool should  
299 be validated, given that our human clinical skills are  
300 intrinsically flawed, and patient performance varies

according to fatigue, medication and time of day. In  
addition, most machine learning techniques such as  
those described above, are supervised, for example,  
an AI model dedicated to scoring a video accord-  
ing to the MDS-UPDRS is trained using clinician's  
scores of MDS-UPDRS [34]. Therefore, at best, the  
accuracy of machine learning techniques will be as  
good as the clinician assessment of MDS- UPDRS,  
which already presents issues with inter/intra rater

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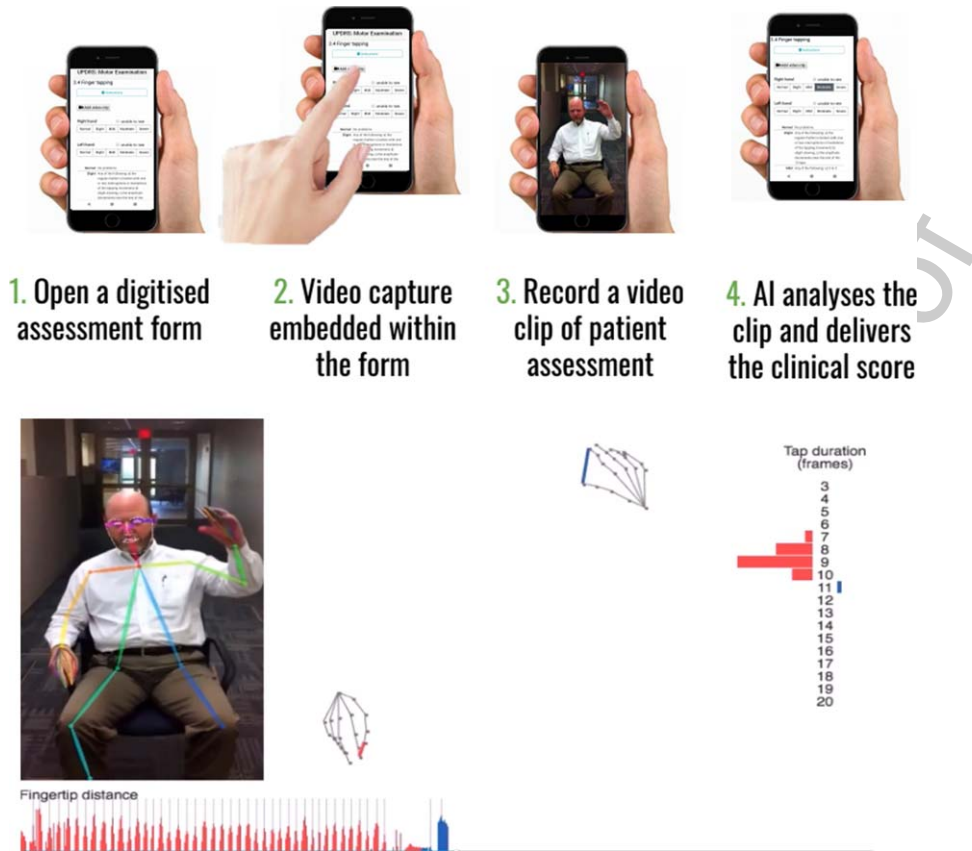


Fig. 1a. Example of Kelvin, a platform that allows the user to record 2D videos of patients with PD on any accessible device, and an inbuilt AI system will analyse the clip and denote scores according to items of the UPDRS. Reproduced with permission.



Fig. 1b. Example of Park, a platform that allows the user to perform the UPDRS score at home. Reproduced with permission.

310 variability, thus there is a need to demonstrate greater  
 311 inter/intra assessment reliability using AI tools which  
 312 would further the argument that AI can provide truly  
 313 objective ratings. In addition, a deep learning method  
 314 (utilised by multiple companies in Table 2), works by  
 315 training a model using example data that is inputted,  
 316 to be able to subsequently identify PD from novel data  
 317 and this heavily relies on large amounts of high quality  
 318 labelled data, to ensure that the model achieves  
 319 state of the art accuracy. One caveat with the current

research into automated video-based assessments is  
 that often the models are trained using small samples  
 of cognitively intact, predominantly white partici-  
 pants, that are relatively younger, and present milder  
 symptoms of disease (Hoehn and Yahr stage 2) [34,  
 37, 38, 41], which may bias the automated assessment  
 framework and thus findings cannot necessarily be  
 generalised to the wider population of people living  
 with PD. One study has demonstrated that it is possi-  
 ble to apply automated video-based assessment to an

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330 older population with lower cognitive status [42], but  
331 this is yet to be demonstrated in the older population  
332 of people with PD, who have a more severe disease  
333 status.

334 So far, recent research developing computer vision-  
335 based methods for AI analysis have shown  
336 success in scoring bradykinesia, gait and facial  
337 expressions, as well as AI scores showing good correlation  
338 with clinician ratings among these studies  
339 [34, 38–44]. At present, the application of AI to diagnosis  
340 of movement disorder patients will likely only  
341 ever be used in conjunction with human expert movement  
342 disorders clinicians, since rigorous validation  
343 of AI technology and subsequent regulatory approval  
344 would be required. Nevertheless, with computer  
345 vision-based analysis of PD there appears to be potential  
346 for repeated, longitudinal data collection, with  
347 internal consistency.

## 348 FUTURE DIRECTIONS

349 In the future, in addition to video-based assessment  
350 to measure MDS-UPDRS-part 3 scores, we  
351 may even see more applications of passive sensing  
352 towards early diagnosis and possible referral for PD.  
353 For example, with users' permission, any video feeds,  
354 e.g., even from videoconferencing sessions could be  
355 analysed for subtle variations of micro expressions  
356 over time as an early detector of PD. The ethical  
357 implications of this are of course potentially myriad.  
358 While uncertainty around video quality and the noise  
359 in the data diminishes the human performance on  
360 measuring MDS-UPDRS score, with recent advances  
361 in AI [45], it is possible to reduce the video bandwidth  
362 usage to one-tenth, resulting in high quality video  
363 despite having low-bandwidth. While many  
364 irregularities in the data may appear noise to humans;  
365 for an AI, these are patterns. With enough data, the  
366 noise can be modelled and successfully decoupled.  
367 In addition, future work could consider comparing  
368 machine-generated scores with clinician assessments  
369 of motor symptoms in blinded OFF and ON medication  
370 conditions to assess the AI ability to detect  
371 different clinical status in the same patient, which  
372 would provide detail into the sensitivity of AI as well  
373 as further validate machine-learning techniques for  
374 clinical purposes. Furthermore, large datasets representing  
375 a wider population across race, gender, geography and  
376 socio-economic boundaries would be key in order to  
377 facilitate an equitable machine-learning outcome.

## CONCLUSION

378  
379 In summary, the move towards remote video measurement  
380 of PD severity has been greatly accelerated by the  
381 COVID19 pandemic. As such, it is vital that clinicians  
382 and researchers devise a valid and safe way to continue  
383 support and monitoring of patients with PD without  
384 exposure to infectious risks, while also without losing  
385 important details currently captured in face-to-face  
386 assessments. Whilst video-based assessment of the  
387 MDS-UPDRS presents some challenges, it is likely that  
388 remote video capture is an accessible means for  
389 neurologists to continually monitor and support  
390 patients living with PD. However, this process must  
391 also consider the circumstances in which a face-to-face  
392 consultation should be triggered, for example, to  
393 evaluate the emergence of atypical features of  
394 parkinsonism or other causes for deterioration in  
395 the clinical signs. At present, whilst the growing  
396 use of Digital Health Technologies is full of promise  
397 for supporting chronic neurological conditions such  
398 as Parkinson's disease, it seems that using automated  
399 video assessments for diagnostic purposes and to  
400 accurately quantify disease severity depends on  
401 research with large, diverse samples and further  
402 validation, in order to best represent and thus be  
403 useful for the 10 million people worldwide with  
404 Parkinson's disease at present.

## CONFLICT OF INTEREST

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406 TF has received grant funding from Cure Parkinson's  
407 Trust, National Institute for Health Research, John  
408 Black Charitable Foundation, Michael J Fox Foundation,  
409 Van Andel Research Institute, Defeat MSA. TF has  
410 received funding from Innovate UK to collaborate in  
411 the assessment of Kelvin-UPDRS with Machine  
412 Medicine Technologies. He has received honoraria  
413 for talks sponsored by Bial, Profile Pharma, Boston  
414 Scientific and has served on Advisory Boards for  
415 Pepton pharmaceuticals, Handl therapeutics, Living  
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