

**ROMA**

17-18 marzo 2026

**NEURO**Young<sup>5<sup>th</sup> edition</sup>  
next generation in neurologia

**Intelligenza Artificiale:  
dalla diagnosi al monitoraggio clinico**

Lazzaro di Biase, MD, PhD

Neurology Unit, Campus Bio-Medico University Hospital

Brain Innovations Lab

[l.dibiase@policlinicocampus.it](mailto:l.dibiase@policlinicocampus.it)

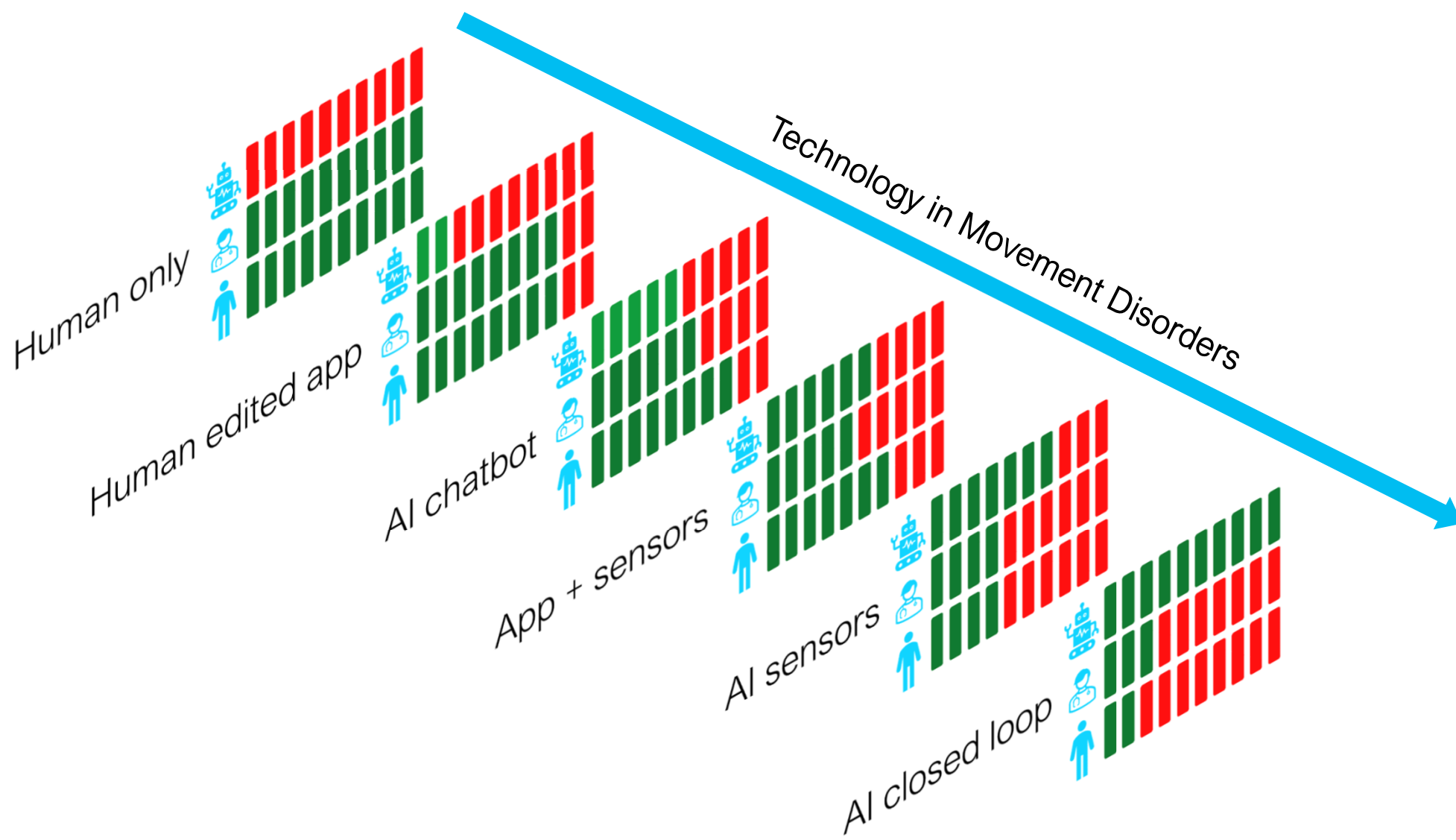


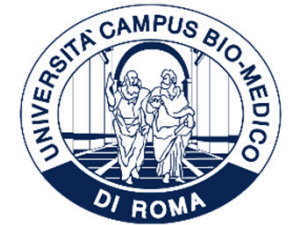
# Disclosures

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L. di Biase has received a speaker honoraria from Bial, consultant honoraria from Abbvie, Bial and Boston Scientific and research funding from Zambon, is the scientific director and one of the shareholders of Brain Innovations Srl, a University spinoff of Campus Bio-Medico University of Rome.

L. di Biase has received support to attend national and international conferences from Abbvie, Bial, Zambon, Boston Scientific.





# Research vs clinical practice

## RESEARCH

## CLINICAL PRACTICE

Accuracy

High

High

Precision

High

High

Cost/benefit

from Low to High

Low

Invasiveness

from Low to High

Low

User experience

from Easy to Complex

Easy

Ethics committee approval

Needed

Not needed

Med. Dev. Regulatory approval

Partial

Needed (FDA, CE mark...)

Lazzaro di Biase, MD, PhD

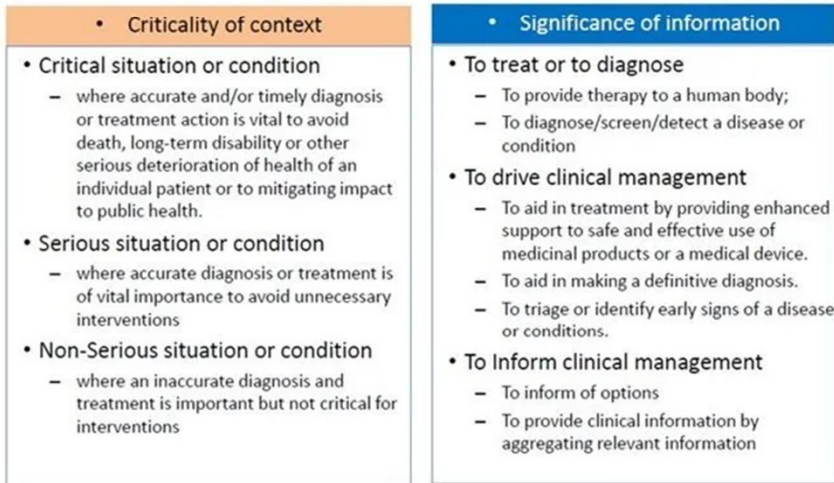
[l.dibiase@policlinicocampus.it](mailto:l.dibiase@policlinicocampus.it)

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**Software as a Medical Device: "software intended to be used for one or more medical purposes that perform these purposes without being part of a hardware medical device."**

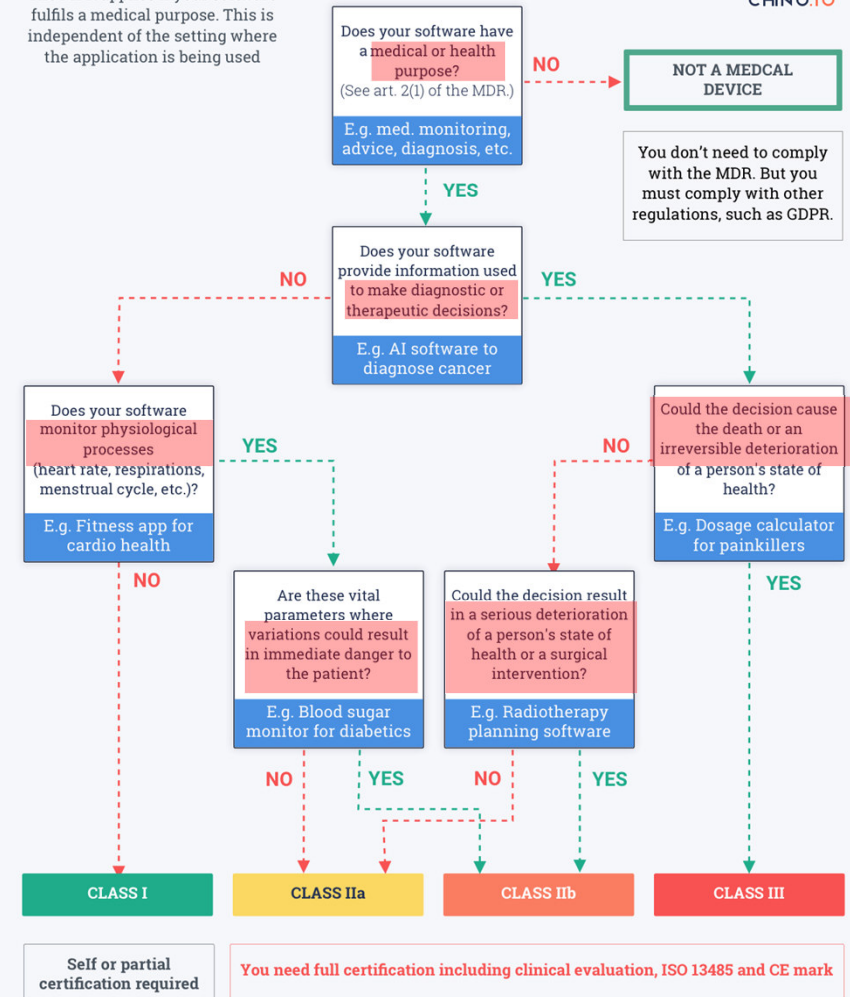
## Clearly defining Intended use of SaMD



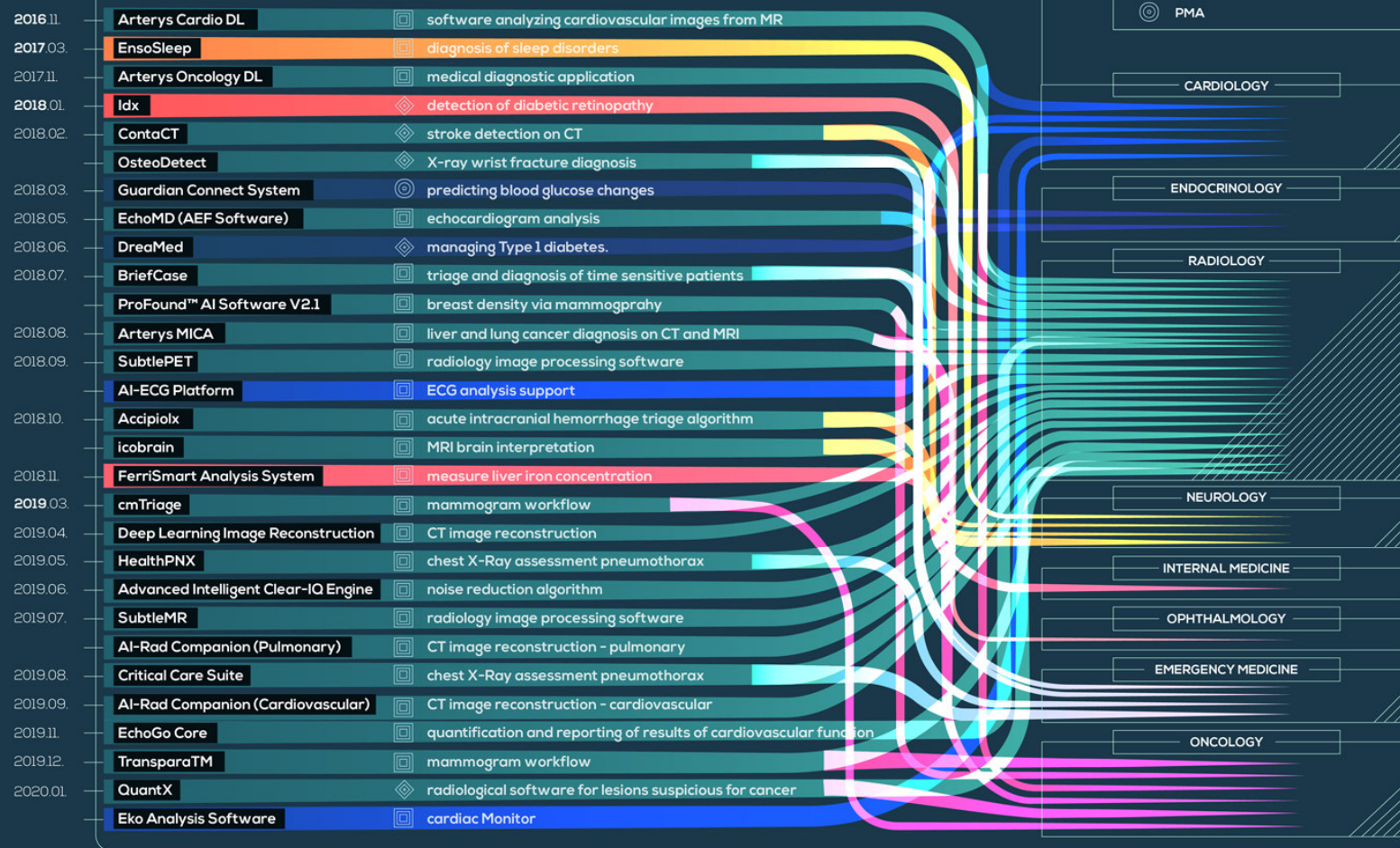
State of Healthcare Situation or Condition	Significance of Information Provided by SaMD to Healthcare Decision		
	Treat or Diagnose	Drive Clinical Management	Inform Clinical Management
Critical	IV	III	II
Serious	III	II	I
Non-Serious	II	I	I

## What MDR class is my software?

The MDR applies if your software fulfils a medical purpose. This is independent of the setting where the application is being used



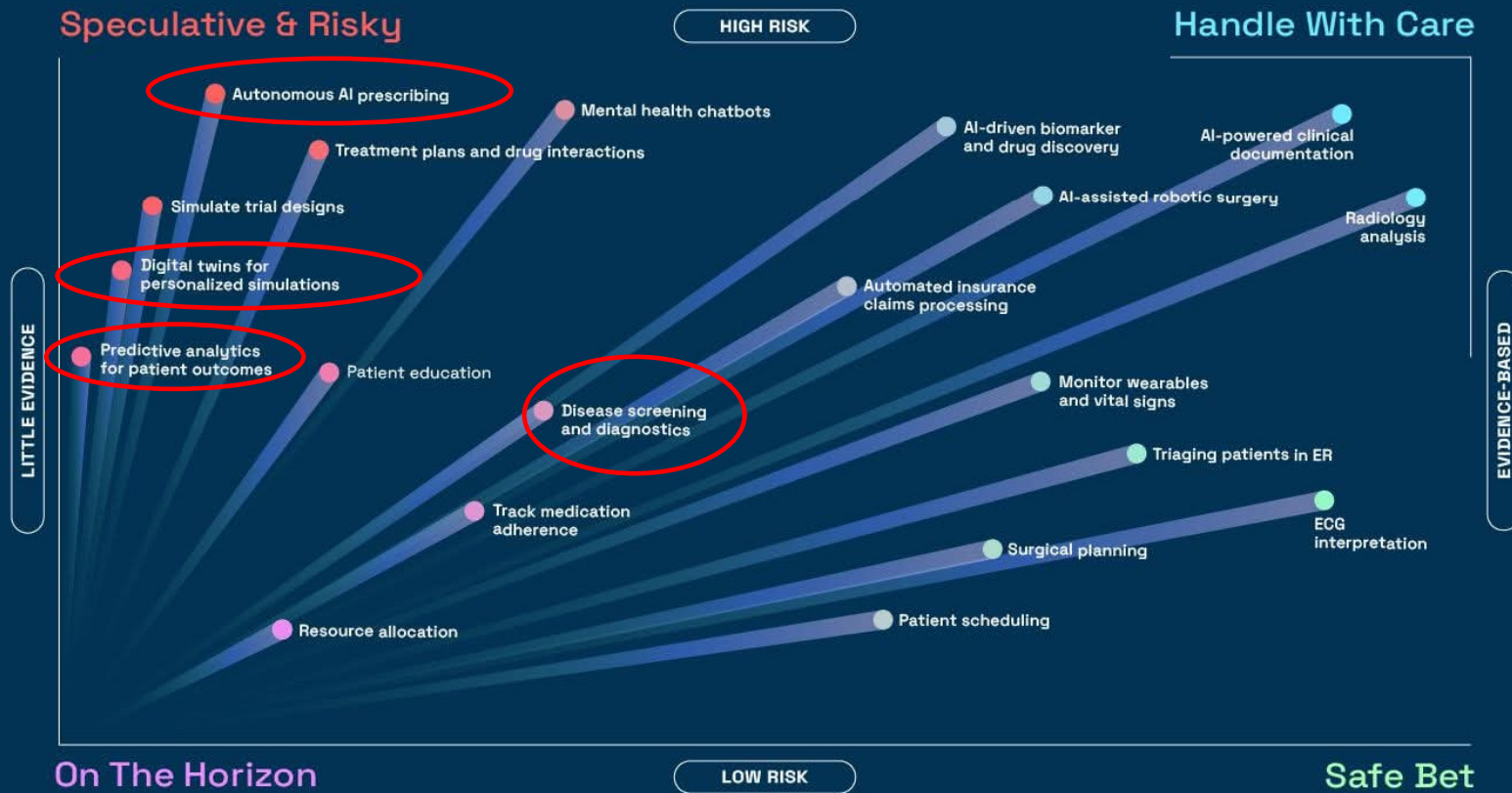
## FDA APPROVALS FOR ARTIFICIAL INTELLIGENCE-BASED DEVICES IN MEDICINE



Benjamins, S., Dhunoo, P. & Meskó, B. The state of artificial intelligence-based FDA-approved medical devices and algorithms: an online database. *npj Digit. Med.* **3**, 118 (2020).

# Navigating AI Use Cases in Healthcare

FROM HYPE TO EVIDENCE,  
FROM SPECULATION TO SAFE BETS



# Perceptual abilities =

Detection  
(information gathering) + Cognition  
(information processing)

Humans

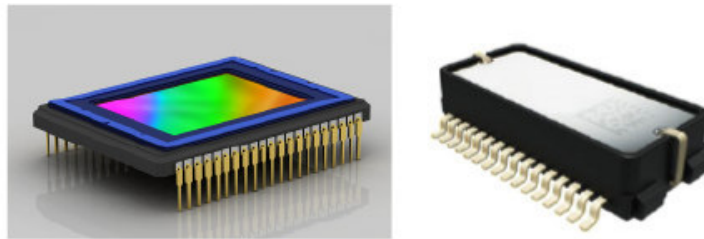


Sensory receptors



Brain

Machines



Sensors



AI and  
other information systems

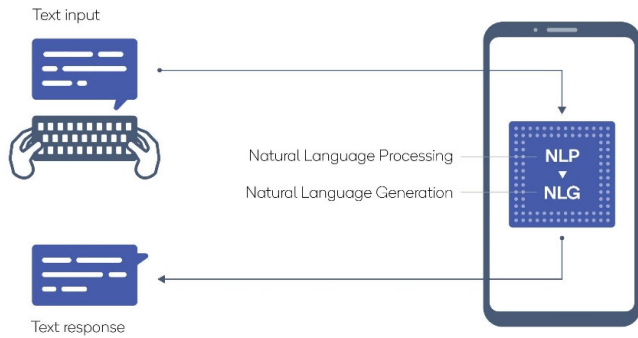
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TELE CONSULT

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# TELECONSULT: CHATBOT

## Text-based personal assistant



**GIORNATA NAZIONALE PER IL PARKINSON**

**26.11.2016**  
ore 9.00 - 14.00  
aula P12|Policlinico

**PROGRAMMA**  
Seminaro "Malattia di Parkinson: ultime novità diagnostiche e terapeutiche"

**09.00 - 09.45**  
I sintomi motori e non motori della Malattia di Parkinson  
**Dott. Lazzaro di Biase**

**09.45 - 10.30**  
Terapie della fase avanzata della Malattia di Parkinson  
**Dott. Lazzaro di Biase**

**10.30 - 11.15**  
Asili non farmacologici per la Malattia di Parkinson  
**Dott. Alessandro Di Santo**

**11.15 - 12.00**  
Nuovi approcci diagnostici e terapeutici e nuove tecnologie  
**Dott. Lazzaro di Biase**

**12.00 - 13.00**  
Il Neurologo risponde: l'Università Campus Bio-Medico di Roma in collaborazione con IBM presenta un assistente virtuale per i medici ed i caregiver.  
**Dott. Enrico Cereda, AD IBM Italia**

**13.00 - 14.00**  
Discussione finale e domande dei partecipanti

**ICBEBIONE**  
Obbligatorio fino ad esaurimento posti. Modalità di pagamento link: [www.policlinicoambrosiano.it/parkinsonday](http://www.policlinicoambrosiano.it/parkinsonday) oppure attraverso il QR code

**Policlinico Università Campus Bio-Medico**  
Via Ardeatina 296 - 00185 Roma

**GNP2016**

**Eventi**  
**Segreteria Neurologia**  
Al Policlinico  
Numero: 06.851221 - 06.85122100  
[secreto@paul.unroma3.it](mailto:secreto@paul.unroma3.it)  
**Dott. Lazzaro di Biase**  
06.791034  
[liber@unroma3.it](mailto:liber@unroma3.it)

**IP**

Il Neurologo risponde UNIVERSITÀ CAMPUS BIO-MEDICO DI ROMA

Ciao, sono Watson e posso darti alcune informazioni sul morbo di Parkinson.

IL NEUROLOGO RISPONDE  
Watson può aiutarti a conoscere la Malattia di Parkinson.  
Puoi chiedere informazioni sul morbo, sulle sue cause vere o presunte, sui suoi sintomi, sulla terapie e sugli aspetti che impattano la qualità di vita del malato.  
Inizia fargli domande così come faresti con il tuo medico.

Fai una domanda a Watson...



# Parkinson's disease

## MISDIAGNOSIS

One diagnosis out  
three is wrong



# 800k

YEARLY MISDIAGNOSIS

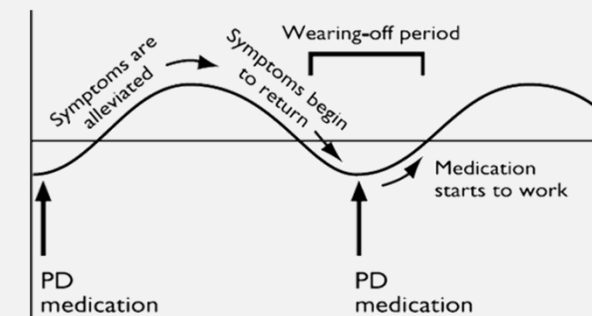
## SYMPTOMS MONITORING

No objective disease  
biomarkers



## THERAPY MANAGEMENT

Costant fluctuation around  
ideal motor status



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# Parkinson's disease

## MISDIAGNOSIS

One diagnosis out  
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# 800k

YEARLY MISDIAGNOSIS

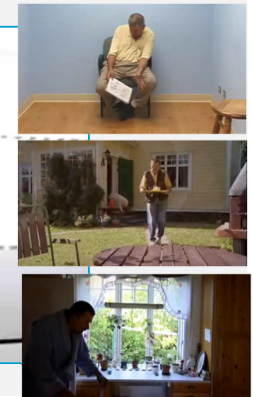
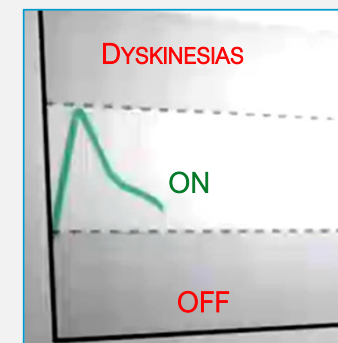
## SYMPTOMS MONITORING

No objective disease  
biomarkers



## THERAPY MANAGEMENT

Costant fluctuation around  
ideal motor status



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TELE DIAGNOSIS

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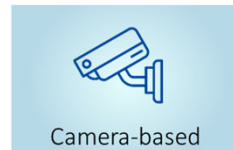
TELE MONITORING

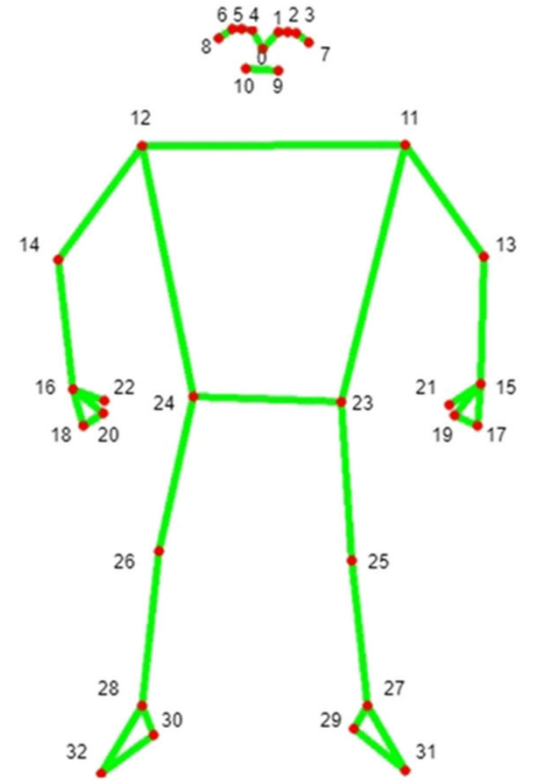
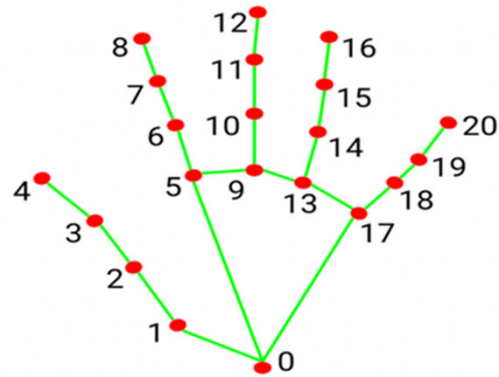
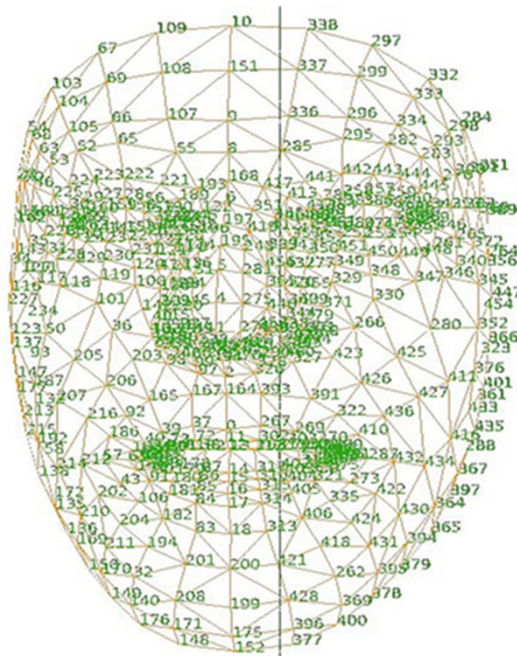
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# Computer Vision

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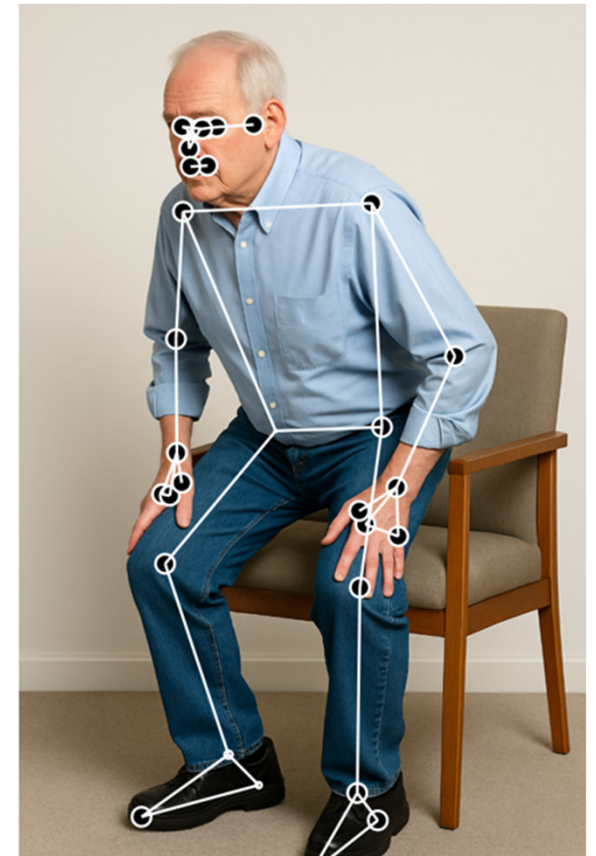




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AI  
Computer vision for movement analysis

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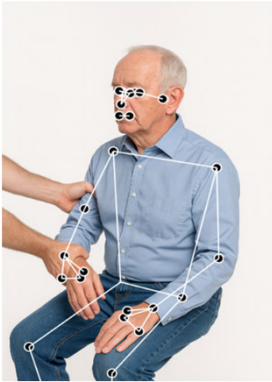


# MDS-UPDRS

**FACE EXPRESSIONS**



**RIGIDITY**



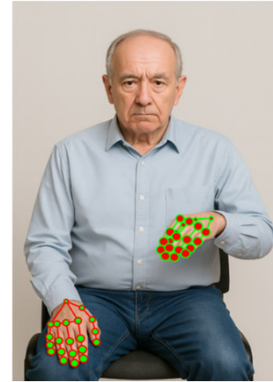
**FINGER TAPPING**



**HAND MOVEMENTS**



**PRONOSUPINATION**



**TOE TAPPING**



**LEG AGILITY**



**ARISING FROM  
A CHAIR**



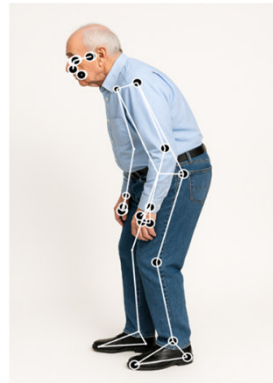
**GAIT**



**POSTURAL STABILITY**



**POSTURE**



**POSTURAL TREMOR**



**KINETIC TREMOR**



**REST TREMOR**



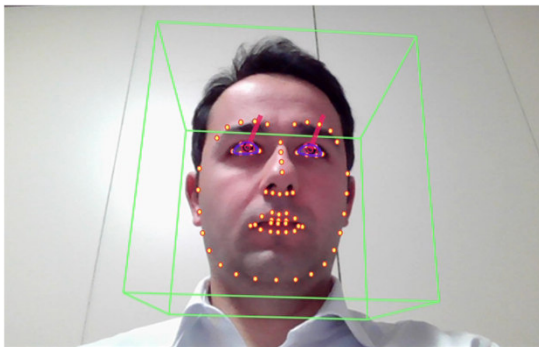
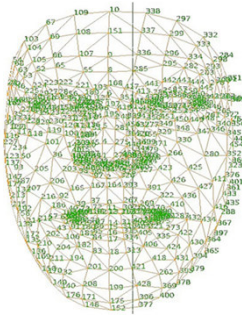
# Televisit



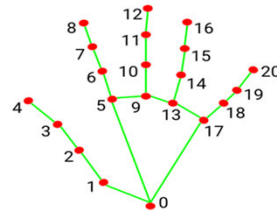
## AI

### Computer vision for movement analysis

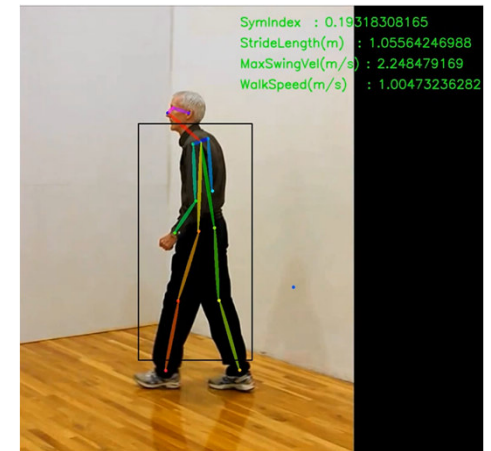
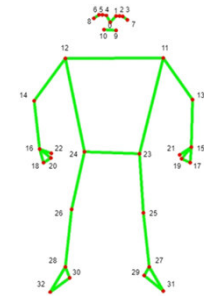
#### Face



#### Hand



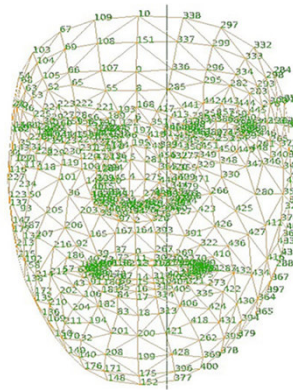
#### Full body



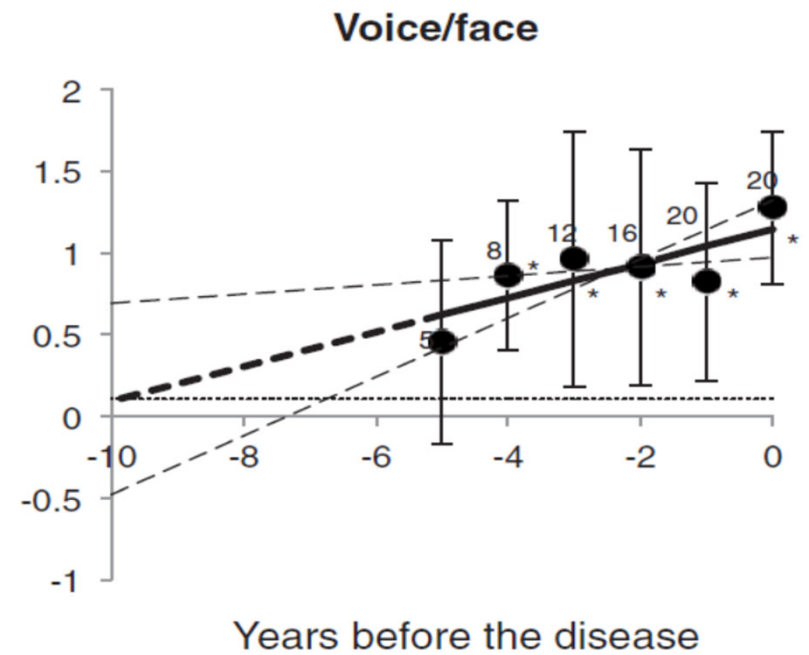
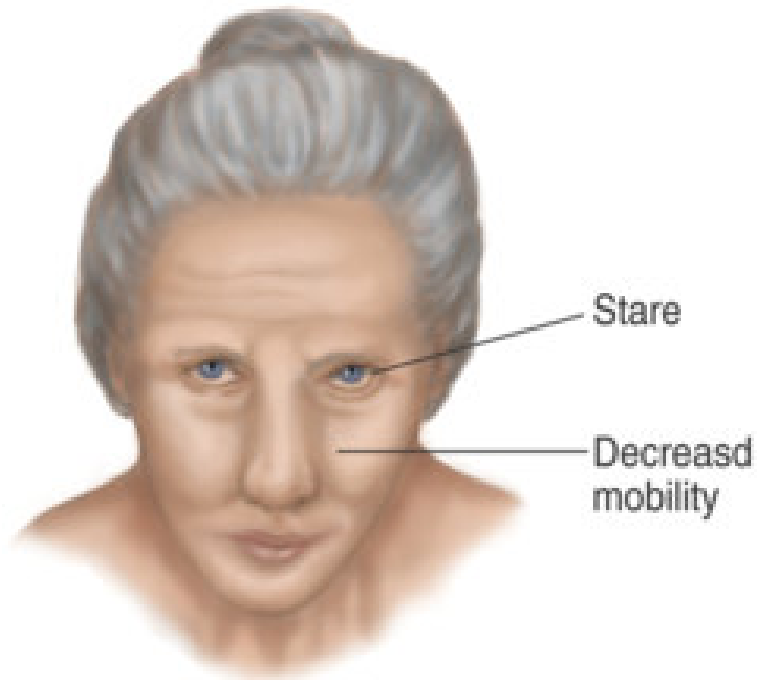
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# FACE Analysis

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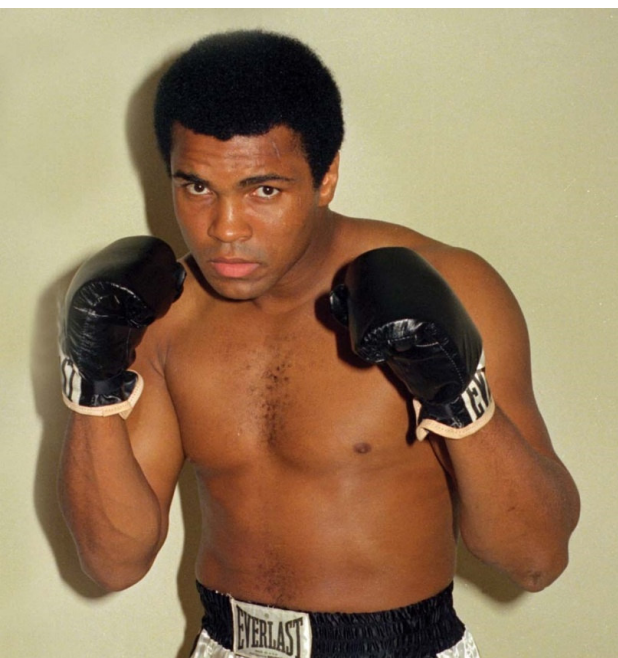


# Hypomimia

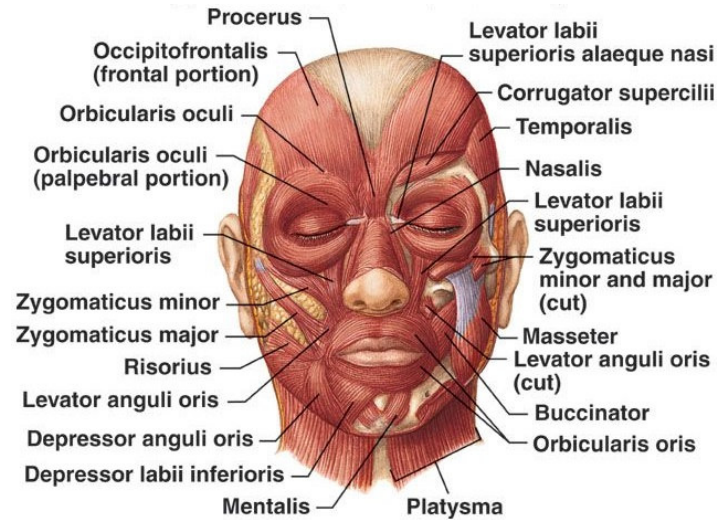
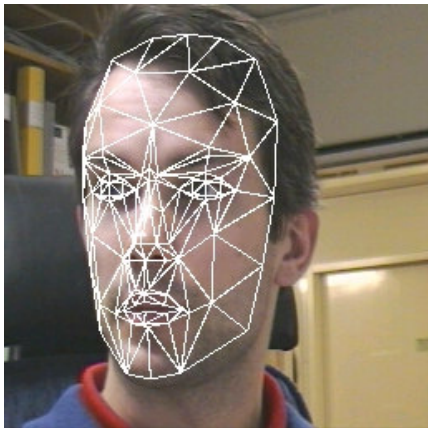


(Postuma et al., 2012)

# Hypomimia

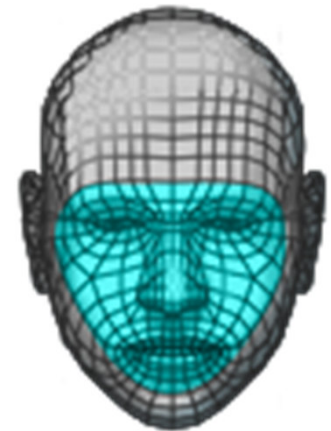


# Quantitative Facial Expression Analysis



0101101

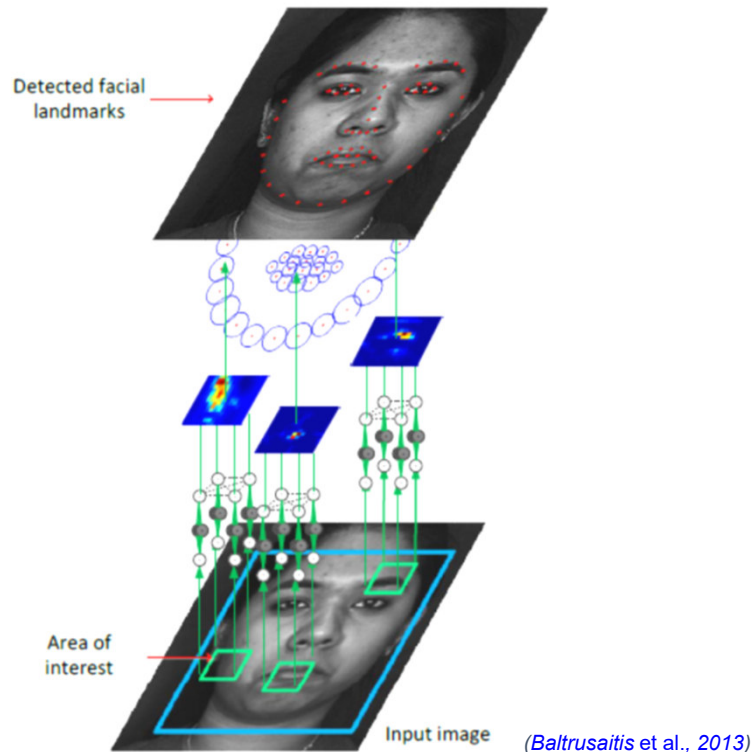
Digital data



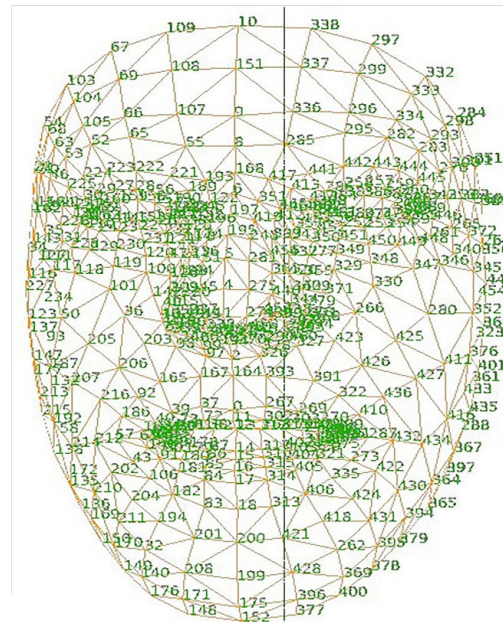


Camera-based

# Facial Action Coding System (FACS)



## FACE LANDMARKS



## EYE LANDMARKS



(Amos et al., 2016)

Lazzaro di Biase, MD, PhD

[lazzaro.dibiase@braininnovations.eu](mailto:lazzaro.dibiase@braininnovations.eu)

Brain Innovations

# Face analysis



Camera-based

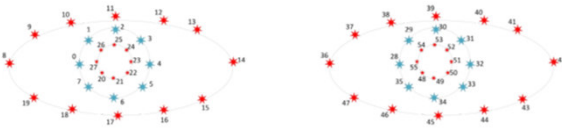
FPS: 23 Confidence: 97%

Action Units	
Presence	Intensity
AU01 - Inner Brow raiser	AU01 - Inner Brow raiser
AU02 - Outer Brow raiser	AU02 - Outer Brow raiser
AU04 - Brow lowerer	AU04 - Brow lowerer
AU05 - Upper lid raiser	AU05 - Upper lid raiser
AU06 - Cheek raiser	AU06 - Cheek raiser
AU07 - Lid tightener	AU07 - Lid tightener
AU09 - Nose wrinkler	AU09 - Nose wrinkler
AU10 - Upper lip raiser	AU10 - Upper lip raiser
AU12 - Lip corner puller (smile)	AU12 - Lip corner puller (smile)
AU14 - Dimpler	AU14 - Dimpler
AU15 - Lip corner depressor	AU15 - Lip corner depressor
AU17 - Chin Raiser	AU17 - Chin Raiser
AU20 - Lip Stretcher	AU20 - Lip Stretcher
AU23 - Lip tightener	AU23 - Lip tightener
AU25 - Lips part	AU25 - Lips part
AU26 - Jaw drop	AU26 - Jaw drop
AU28 - Lip suck	AU28 - Lip suck
AU45 - Blink	AU45 - Blink

Head Rotation			
Rotation	Turn: -3°	Up/down: -19°	Tilt: 1°
Location	X: -1 mm	Y: 21 mm	Z: 443 mm

# Parkinson's disease hypomimia predictor (PHP)

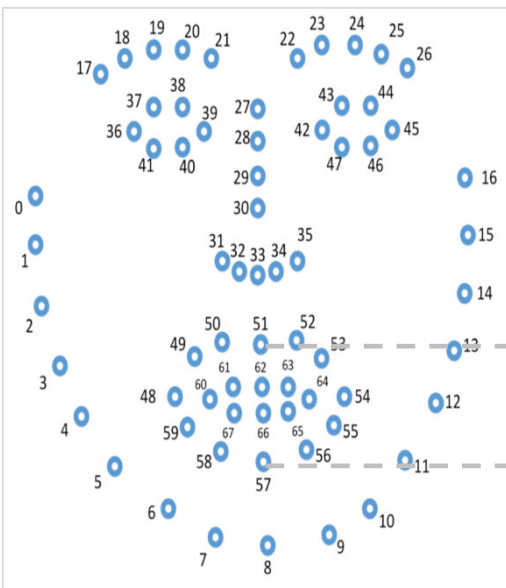
## EYE LANDMARKS



### Blinking intensity

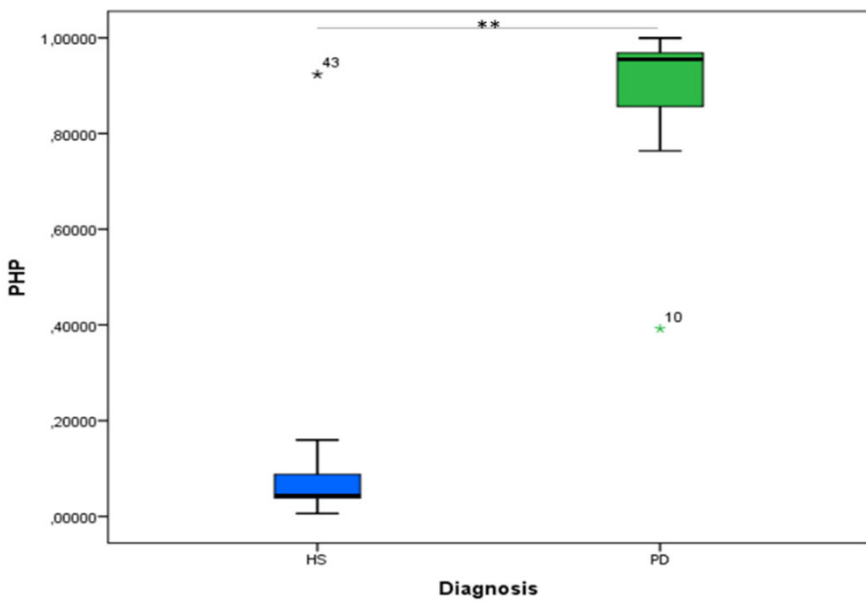
max.AU45.r: Maximum value AU45 (blink: relaxation of *levator palpebrae* and contraction of *orbicularis oculi, pars palpebralis*)

## FACE LANDMARKS

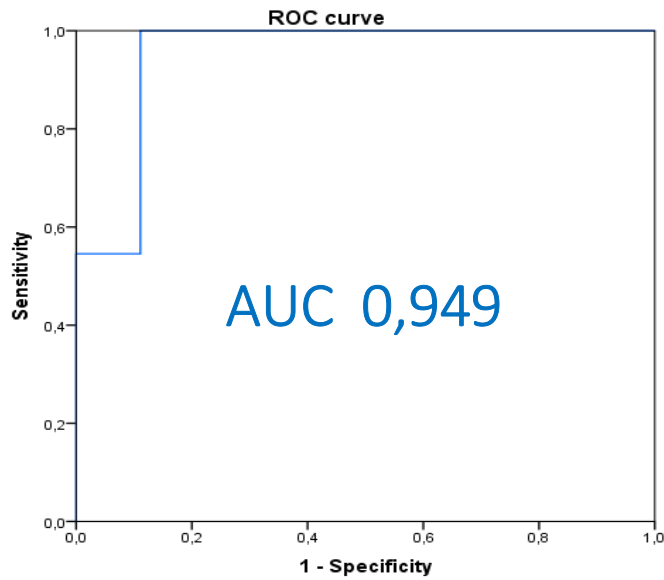


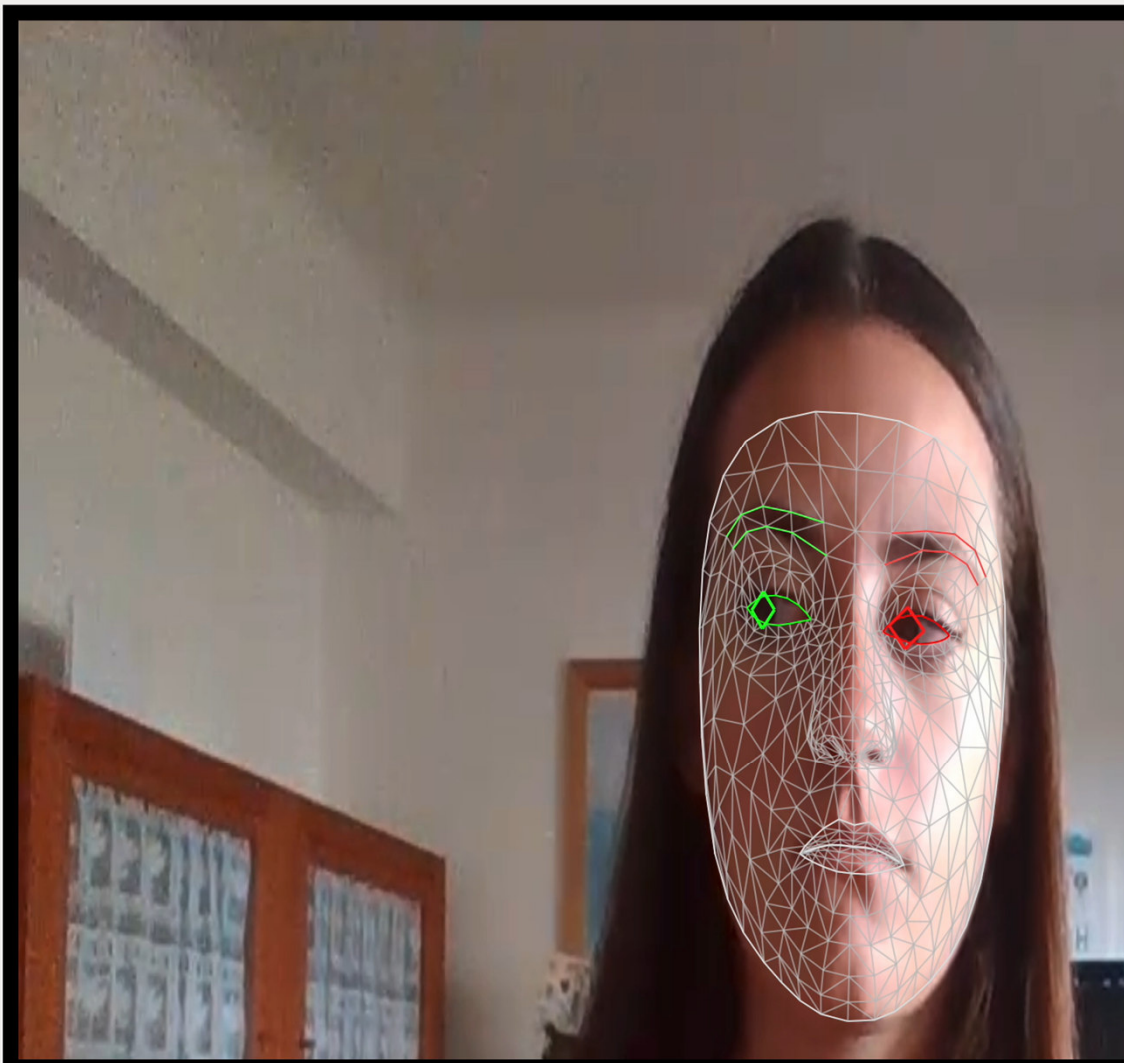
### Lips distance

max.lips.2D.1: Maximum value between face landmarks 51 and 57



Legend= \*\*: Mann-Whitney U test  $p < 0,01$





## Analisi

STOP

Seleziona una feature da analizzare

Seleziona... ▾

1,0  
0,8  
0,6  
0,4  
0,2  
0

## Movement Analysis

Patients  ▾

### Models

- Face Model
- Pose Model
- Hand Model

Selected Model: face

Disable analysis

× Close

# VIDEO ANALYSIS

## EYE BLINKING

---

ON

Number of blinking:0  
Frequency: 0.0 Hz

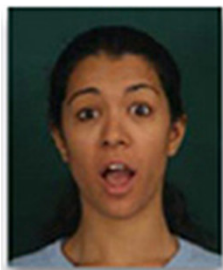


OFF

Number of blinking:0  
Frequency: 0.0 Hz



# UNIVERSAL FACIAL EXPRESSIONS



Surprise



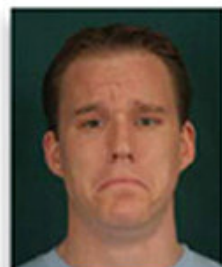
Happiness



Contempt



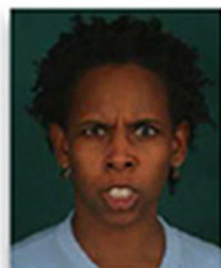
Fear



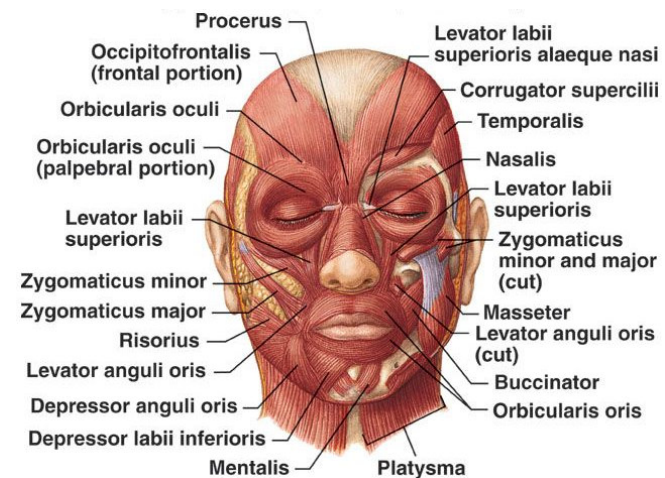
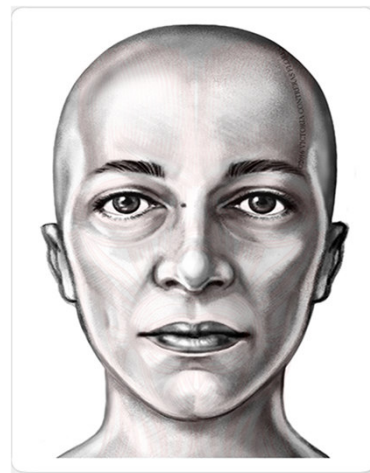
Sadness



Disgust



Anger



# UNIVERSAL FACIAL EXPRESSIONS



Surprise



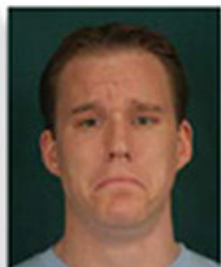
Happiness



Contempt



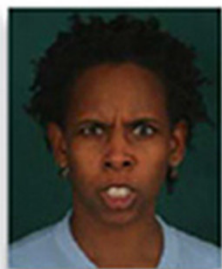
Fear



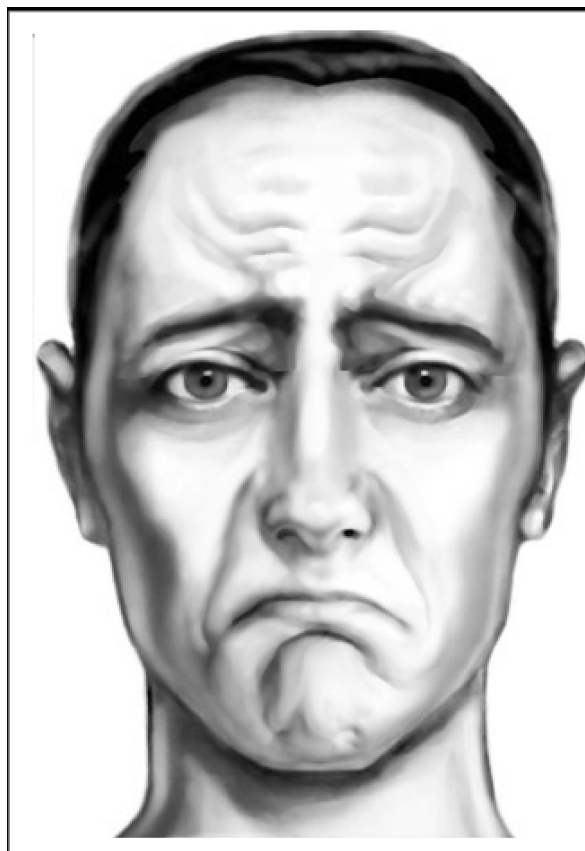
Sadness



Disgust



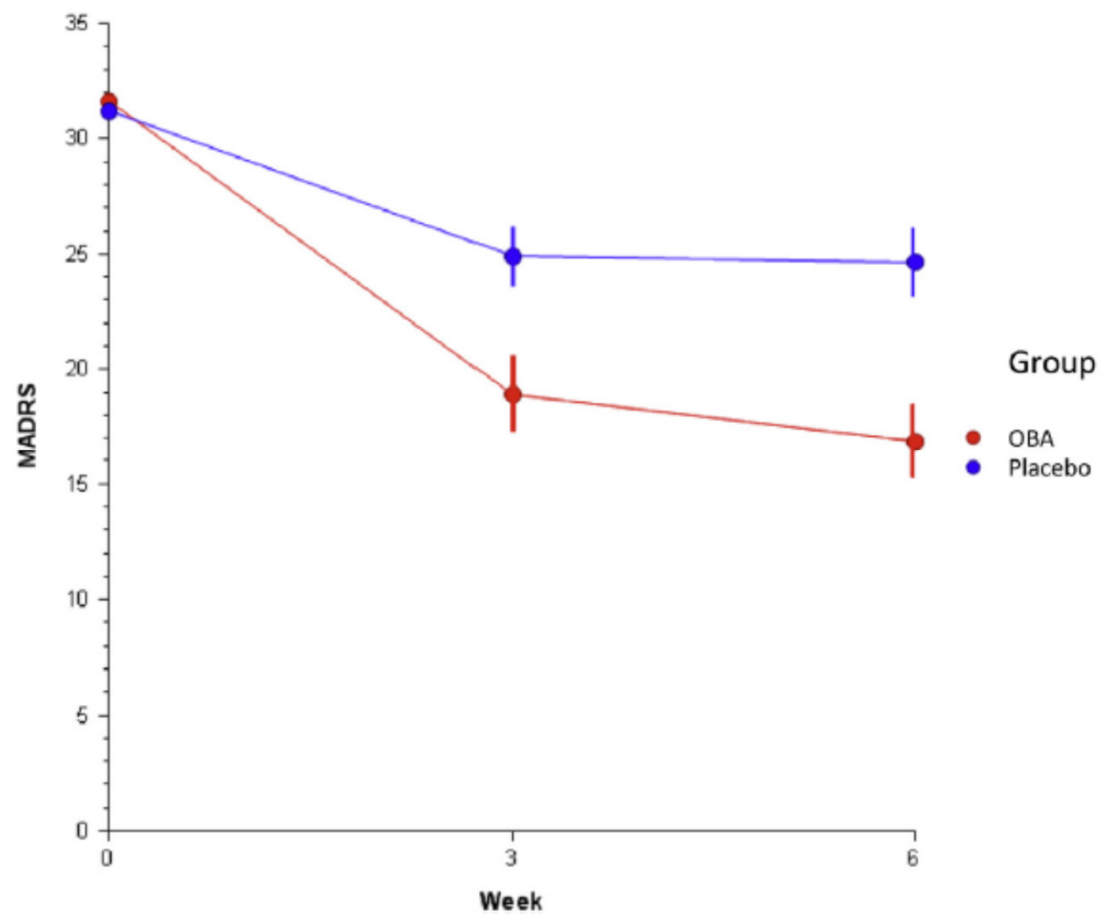
Anger



<b>Frontalis</b>	<b>Eyebrows</b>	elevation	↑
<b>Corrugator</b>		oblique Mov.	↗ ↘
OrbicularisOculi		oblique Mov.	↗ ↘
Procerus	<b>Eyelids</b>	depression	↕
NasalisTransversa		elevation	↑
<b>NasalisAlaris</b>		direction of the look	↗ ↘
Nose Expander		depression	↕
OrbicularisOris	<b>Wings of Nose</b>	elevation	↑
Caninus		dilatation	↔
QuadratusLabiiSup.		<b>depression</b>	↓
OwnElevator	<b>Corners of mouth</b>	elevation	↑
Zygomaticus Min.		upward retraction	↕
Zygomaticus Maj.		retraction	↔
Risorius		<b>depression</b>	↓ ↘ ↙
Buccinatorius	<b>Upper Lip</b>	elevation	↑
<b>QuadratusLabii Inf.</b>	<b>Lower Lip</b>	eversion	↕
<b>Triangularis</b>		depress.+ retracc.	↔
<b>Mentalis</b>	<b>Lips</b>	compression	⊗
MASTICATORS		pursing	⊖
<b>Platysma</b>		protusion	⊕
		Parting	⊖
		separa.+ later. mov	↕
		separa.+later. mov. 2	↔
	<b>Maxilar</b>	depression	↓
	<b>Neck</b>		



## DEPRESSION



Finzi E, Rosenthal NE. Treatment of depression with onabotulinumtoxinA: a randomized, double-blind, placebo controlled trial. J Psychiatr Res. 2014

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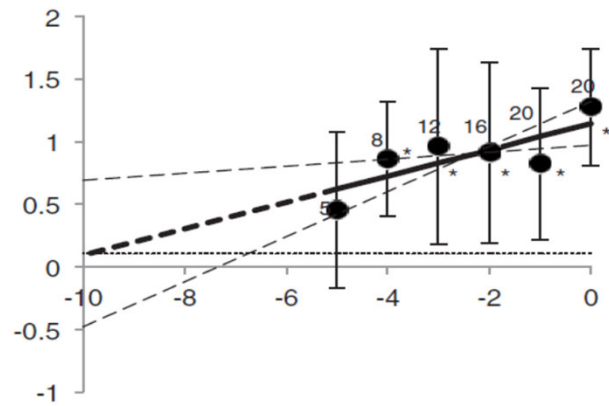
# VOICE Analysis

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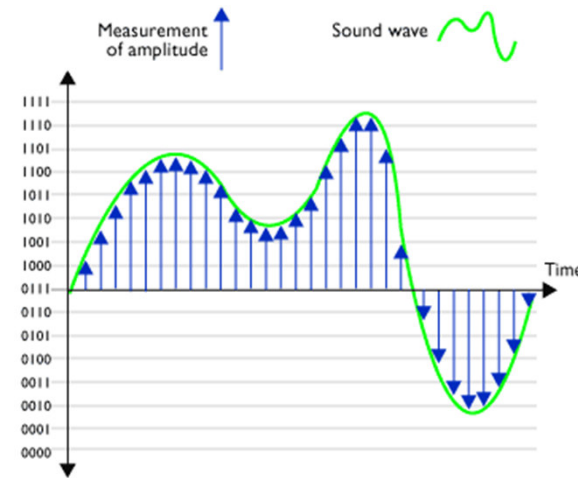
# Quantitative Speech Analysis

Voice/face



Years before the disease

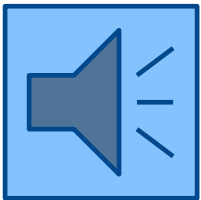
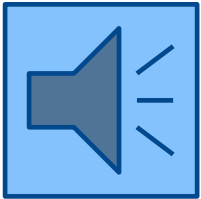
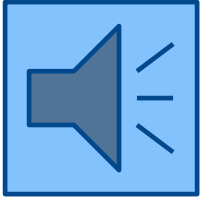
*(Postuma et al., 2012)*



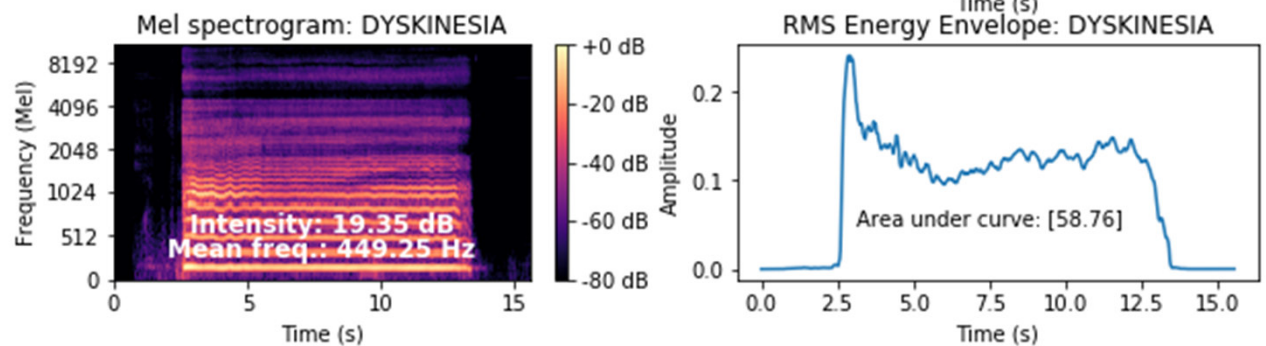
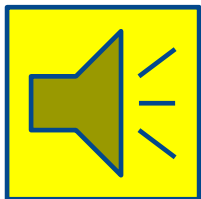
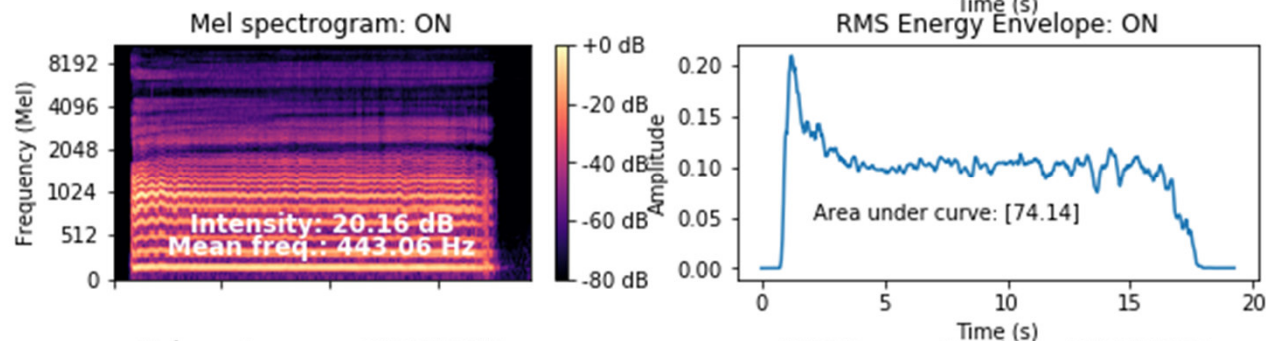
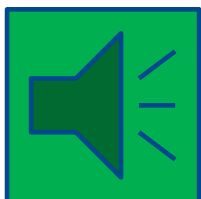
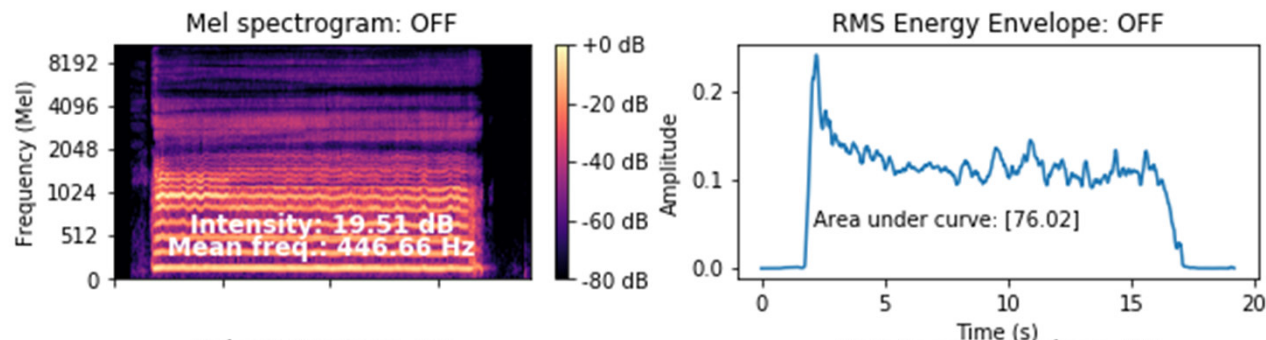
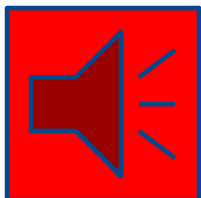
0101101  
Digital data

OBJECTIVE  
NUMERICAL  
OUTPUT

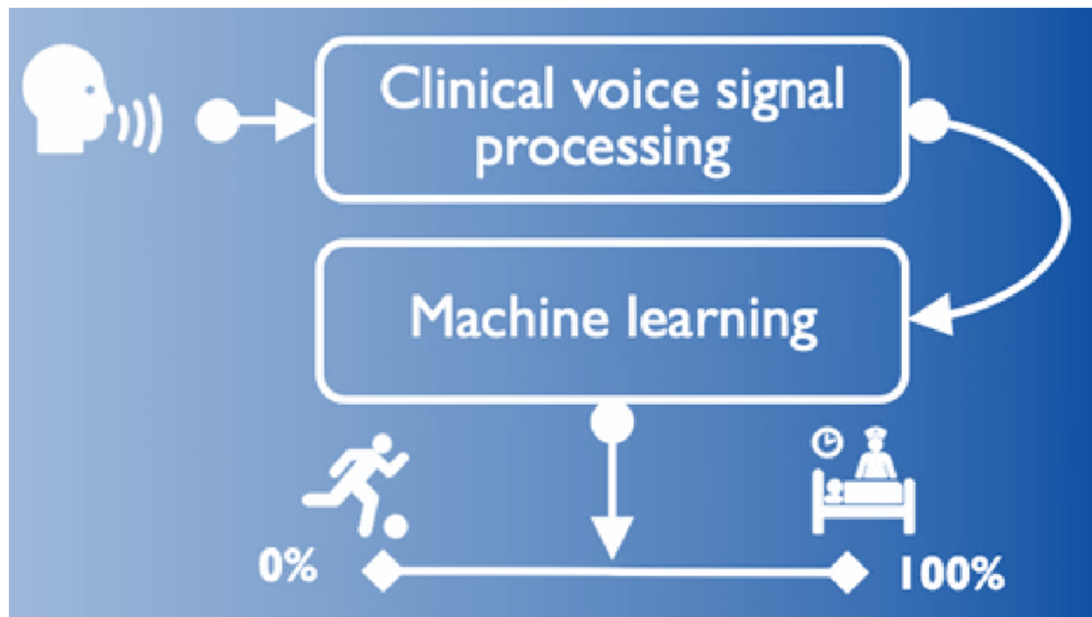
# Audio analysis



# Audio analysis



# Quantitative Speech Analysis

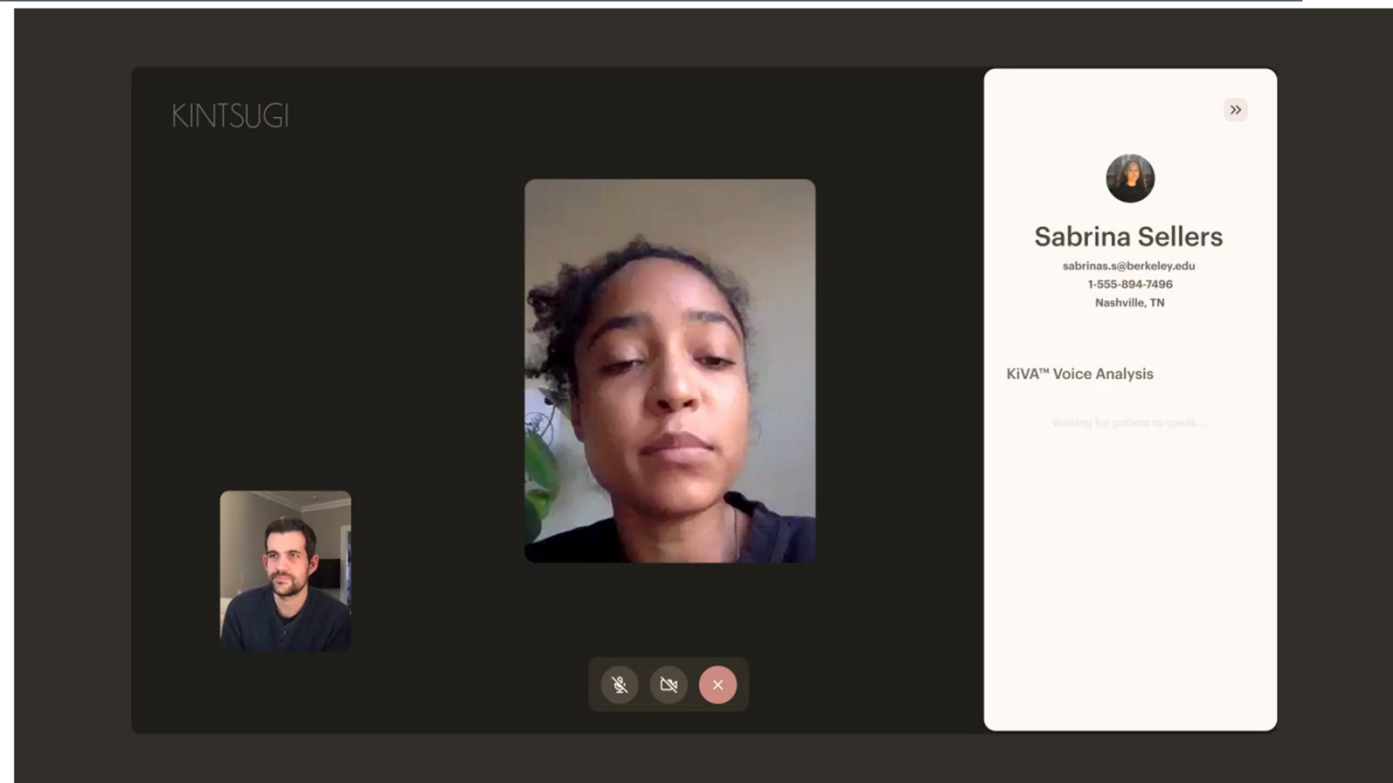
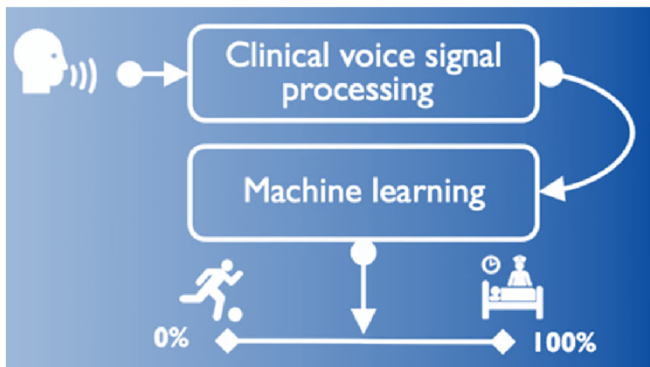


MACHINE VS CLINICIANS  
2 UPDRS points difference

Tsanas A, J R Soc Interface. 2011 Jun  
6;8(59):842-55. doi: 10.1098/rsif.2010.0456.

# Quantitative Speech Analysis

Voice biomarkers for Anxiety and Depression

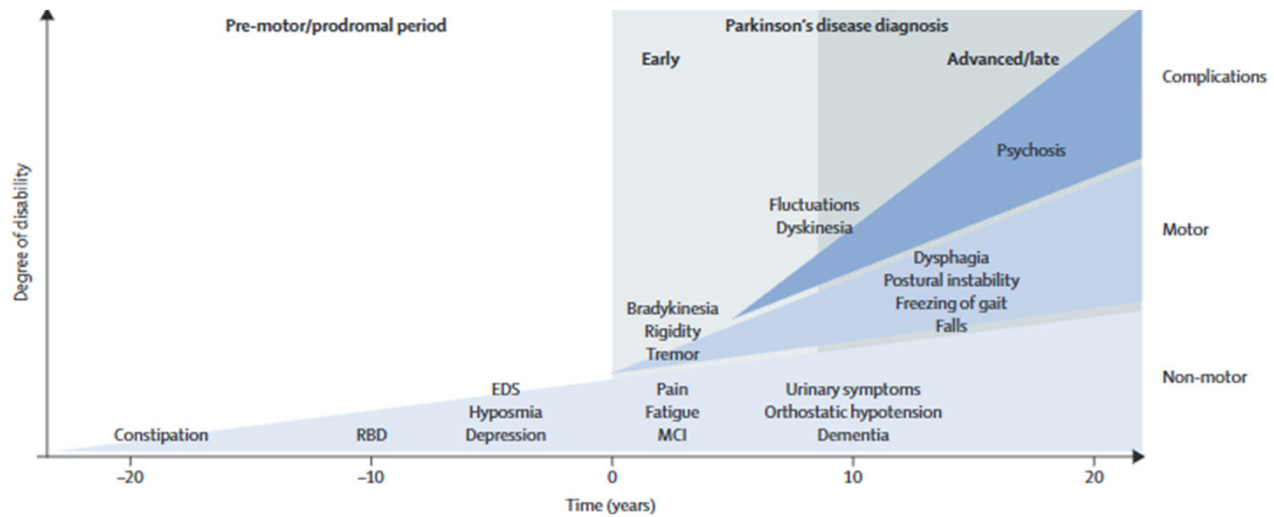


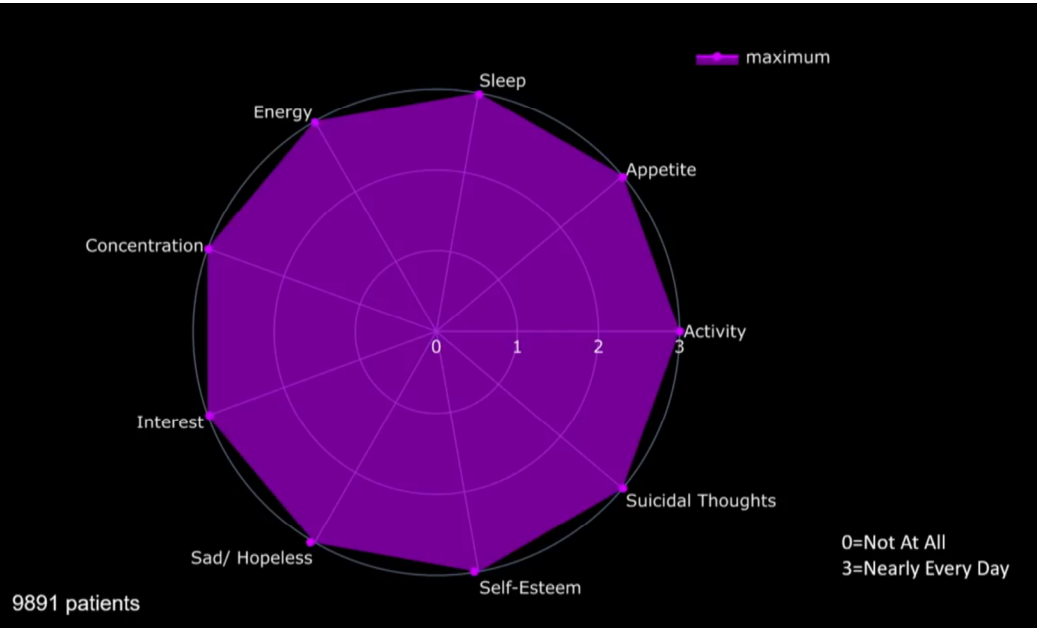
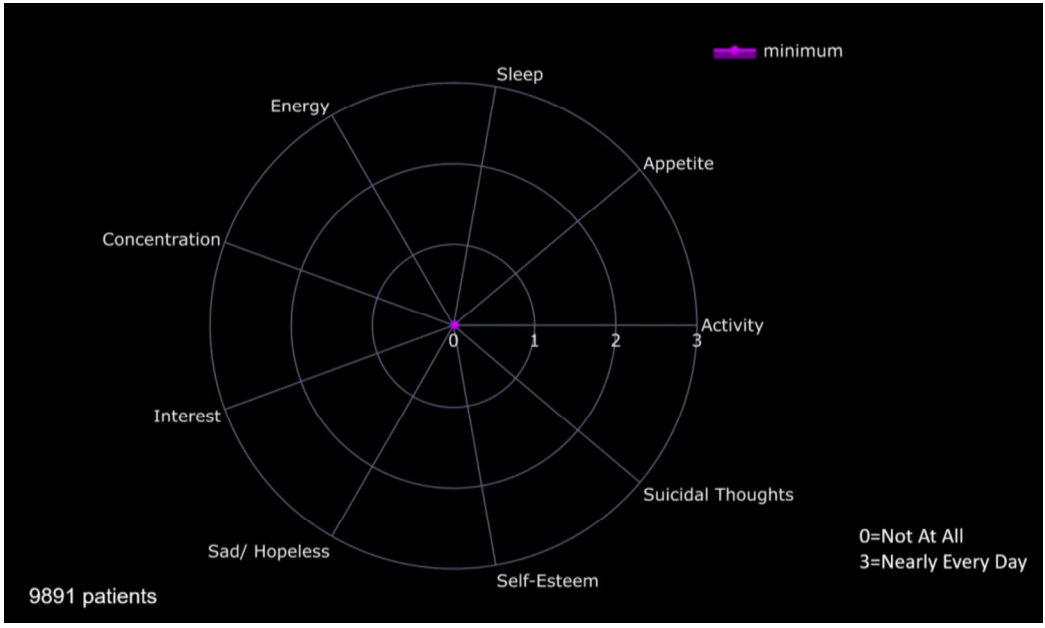
The screenshot shows the Kintsugi telehealth interface. At the top left, the "KINTSUGI" logo is visible. The main area contains two video feeds: a smaller one in the bottom left showing a male provider and a larger one in the center showing a female patient. To the right of the video feeds is a white sidebar containing the patient's profile information: a circular profile picture, the name "Sabrina Sellers", the email "sabras.s@berkeley.edu", the phone number "1-555-894-7496", and the location "Nashville, TN". Below this information, it says "KIVA™ Voice Analysis" and "Waiting for patient to speak...". At the bottom of the video feeds, there are three control buttons: a microphone icon, a video camera icon, and a red "X" icon.

---

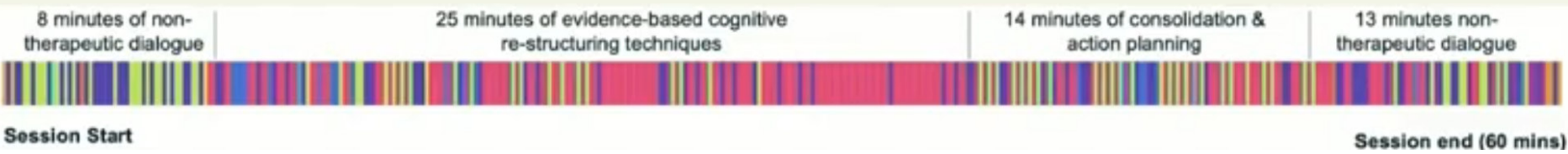
# DEPRESSION

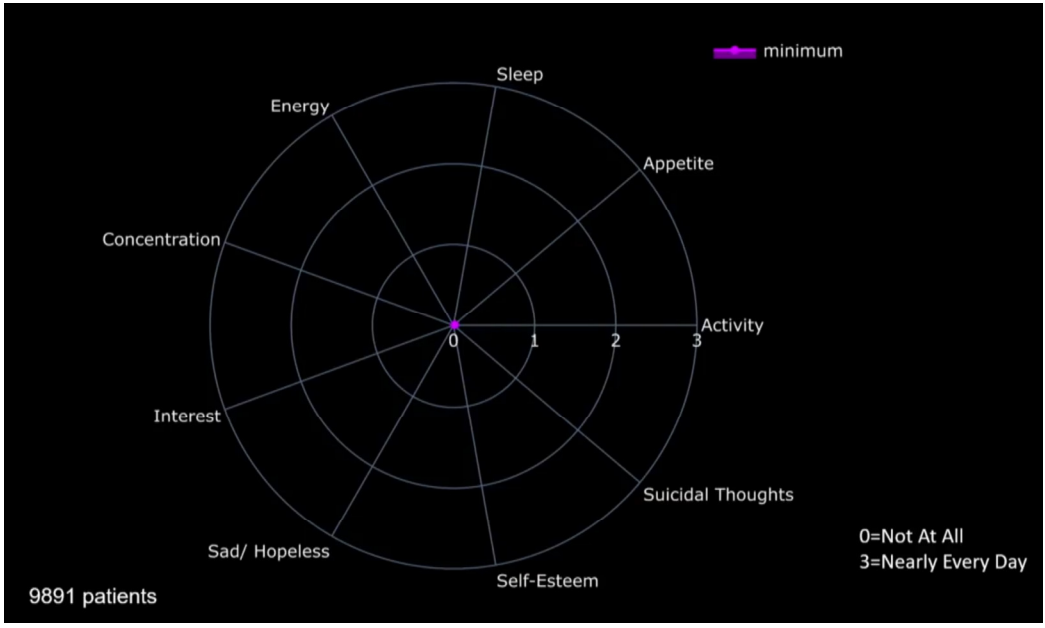
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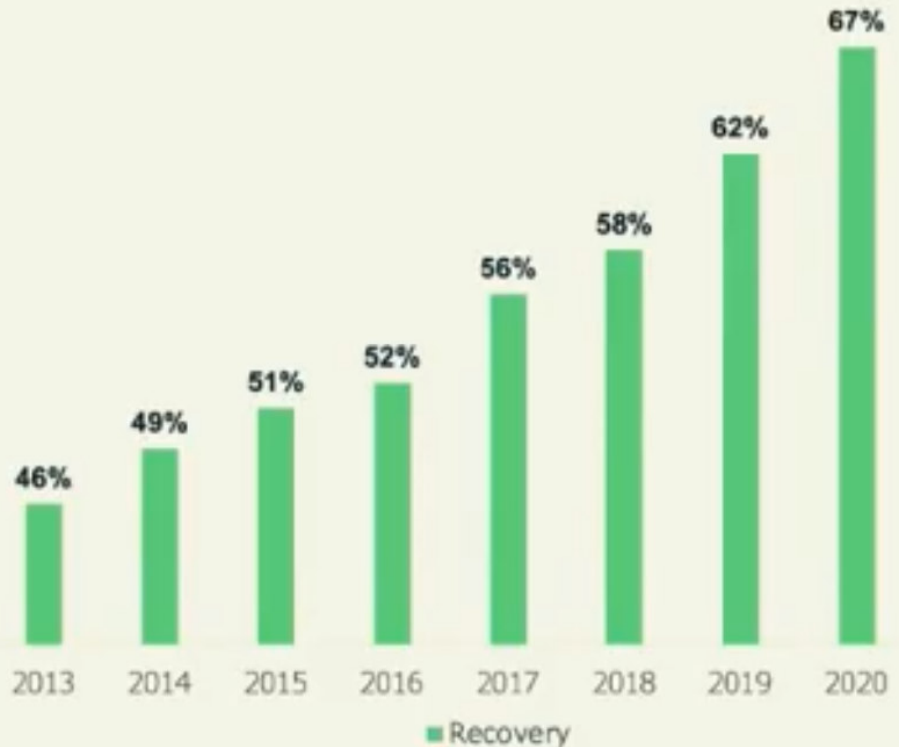
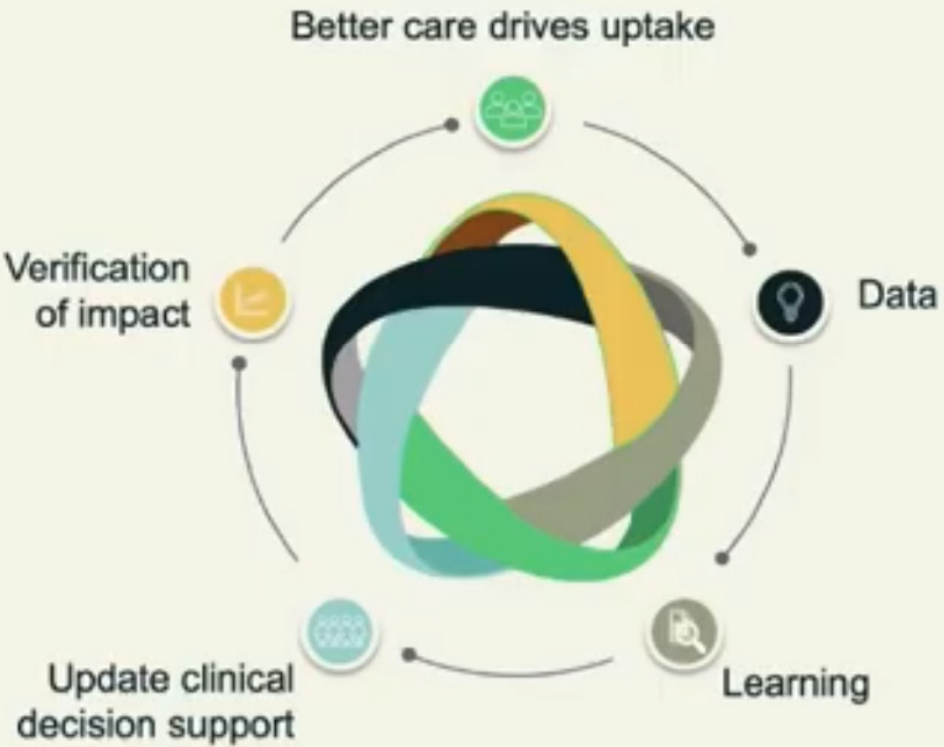


# Telehealthcare Platforms Can be Used to Collect Decision-Action-Outcomes Data at Scale





# Data-enabled care models can improve outcomes in short cycles



Clinical Recovery Rates 2013-2020 (IESO Benchmark Cohort Depressive Episode and Generalised Anxiety Disorder)



eightbillionminds™



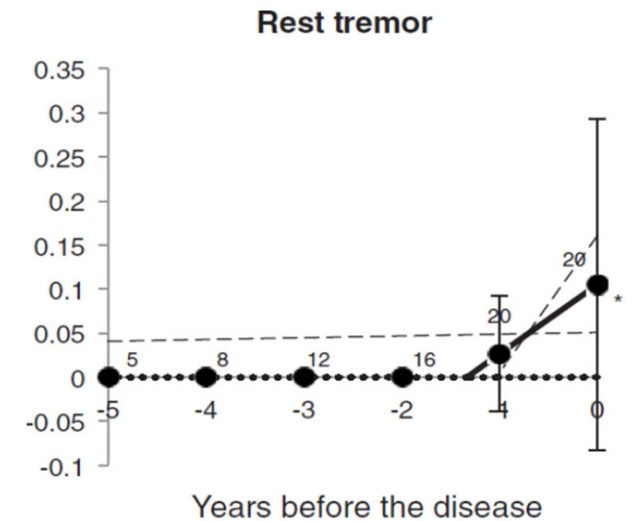
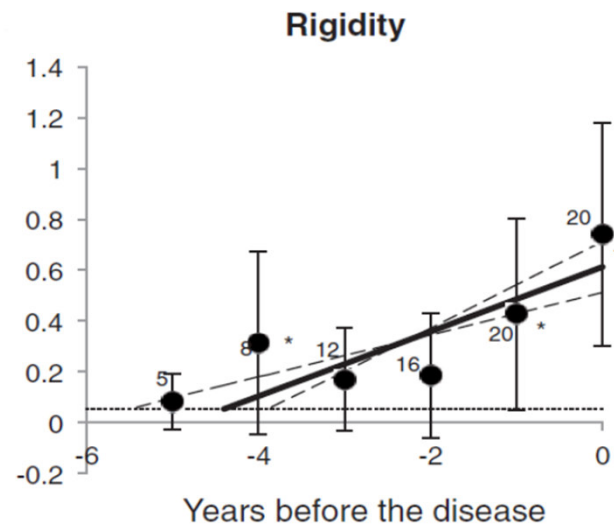
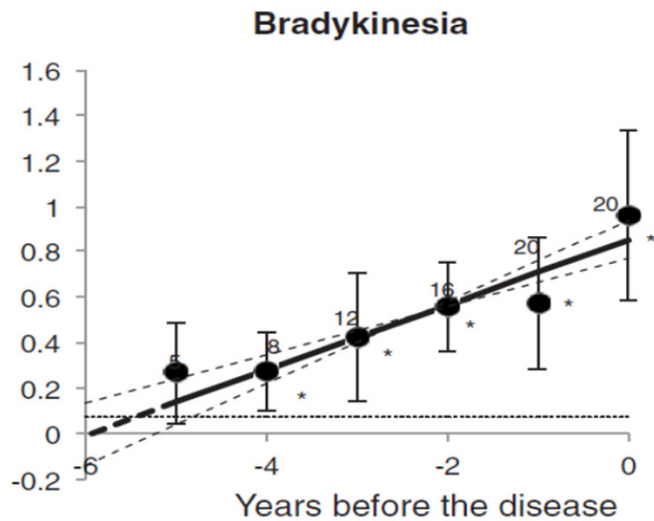
---

# MOVEMENT Analysis

---



# Early diagnosis



## FINGER TAPPING

## HAND MOVEMENTS

## PRONATIO-SUPINATION

## TOE TAPPING

## LEG AGILITY

0: Normal: No problem.

1: Slight:

Any of the following: a) the regular rhythm is broken with one or two interruptions or hesitations of the movement; b) slight slowing; c) the amplitude decrements near the end of the task.

2: Mild:

Any of the following: a) 3 to 5 interruptions during the movements; b) mild slowing; c) the amplitude decrements midway in the task.

3: Moderate:

Any of the following: a) more than 5 interruptions during the movement or at least one longer arrest (freeze) in ongoing movement; b) moderate slowing; c) the amplitude decrements starting after the 1st open-and-close sequence.

4: Severe:

Cannot or can only barely perform the task because of slowing, interruptions or decrements.

## QUANTITATIVE ANALYSIS

VARIABILITY OF FREQUENCY

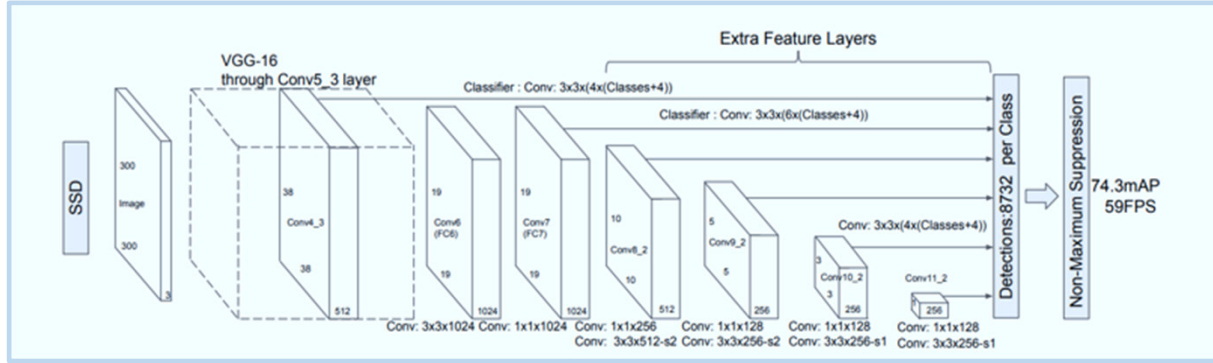
AMPLITUDE (M)

VELOCITY (M/S)

# VIDEO



# Deep learning Landmarks detection

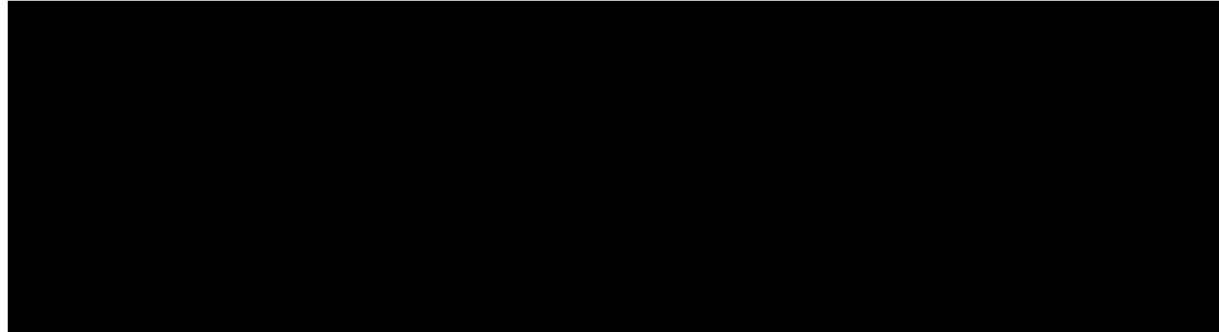


Camera-based

## VIDEO

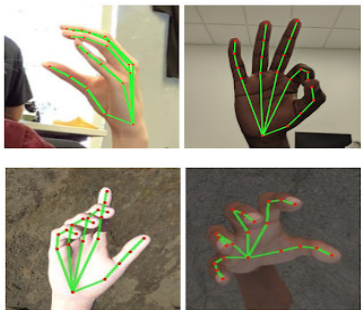
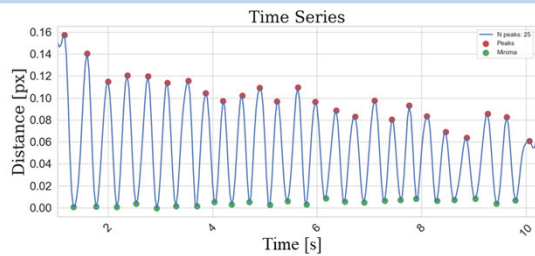


## Deep learning Landmarks detection

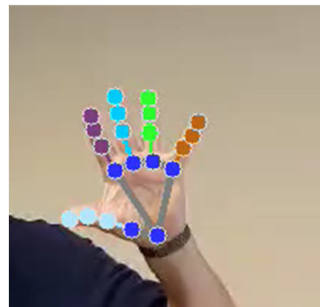


Camera-based

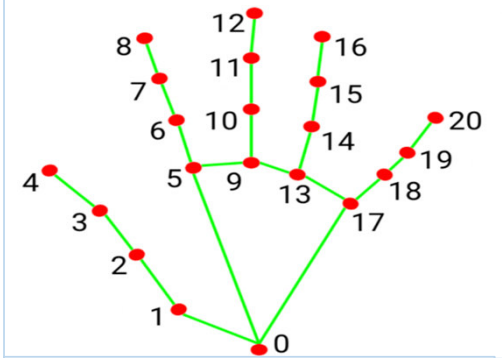
## Kinematic analysis and Pose detection



## VIDEO with landmarks



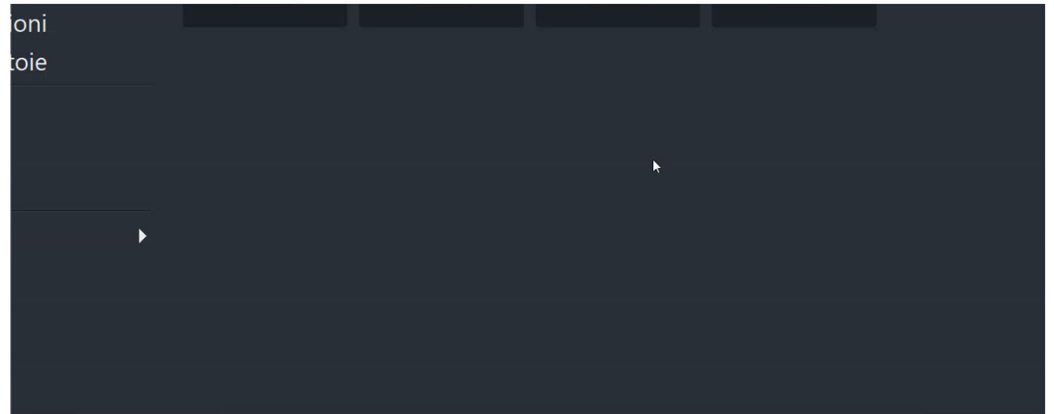
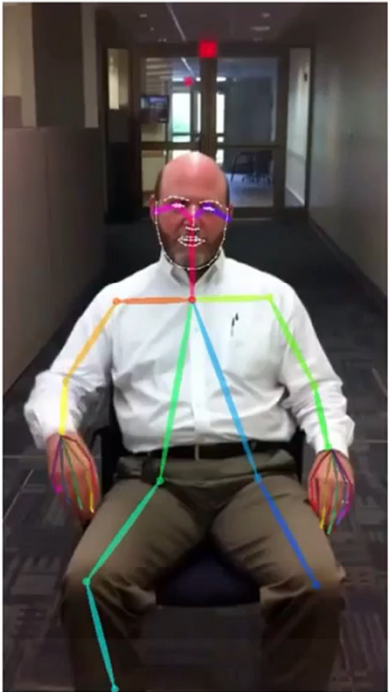
## Hand Landmarks



- |                       |                       |
|-----------------------|-----------------------|
| 0. WRIST              | 11. MIDDLE_FINGER_DIP |
| 1. THUMB_CMC          | 12. MIDDLE_FINGER_TIP |
| 2. THUMB_MCP          | 13. RING_FINGER_MCP   |
| 3. THUMB_IP           | 14. RING_FINGER_PIP   |
| 4. THUMB_TIP          | 15. RING_FINGER_DIP   |
| 5. INDEX_FINGER_MCP   | 16. RING_FINGER_TIP   |
| 6. INDEX_FINGER_PIP   | 17. PINKY_MCP         |
| 7. INDEX_FINGER_DIP   | 18. PINKY_PIP         |
| 8. INDEX_FINGER_TIP   | 19. PINKY_DIP         |
| 9. MIDDLE_FINGER_MCP  | 20. PINKY_TIP         |
| 10. MIDDLE_FINGER_PIP |                       |



Camera-based



Lazzaro di Biase, MD, PhD

[lazzaro.dibiase@braininnovations.eu](mailto:lazzaro.dibiase@braininnovations.eu)

Brain Innovations



## Analisi Blinking

STOP

Seleziona una feature da analizzare

✓ Seleziona...

Finger Tapping

Hand Open-Close

1,0  
0,8  
0,6  
0,4  
0,2  
0

## Movement Analysis

Patients

### Models

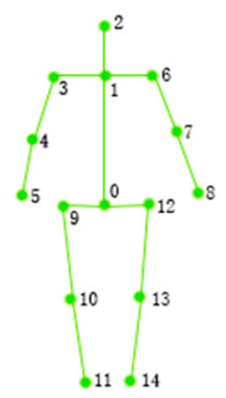
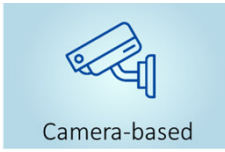
- Face Model
- Pose Model
- Hand Model

Selected Model: hand

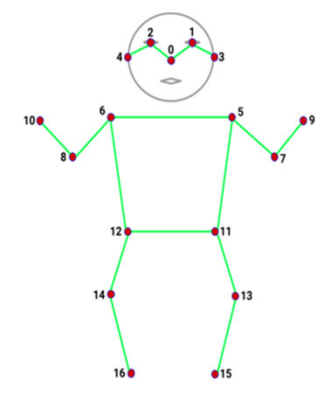
Disable analysis

× Close

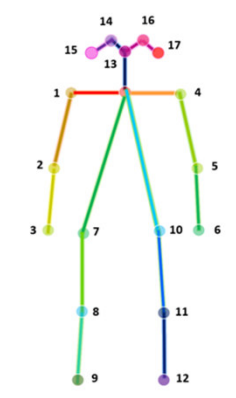
# Full body analysis



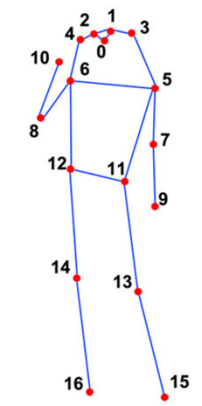
**Openpose**  
15 pt



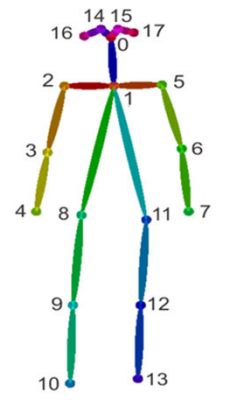
**COCO / MoveNet / PoseNet  
/ Yolo / Effcient-pose**  
17 pt



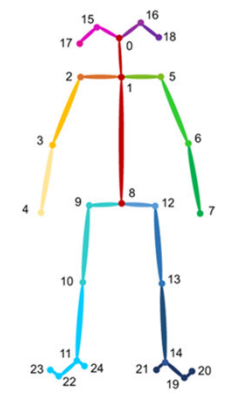
**Alphapose**  
17 pt



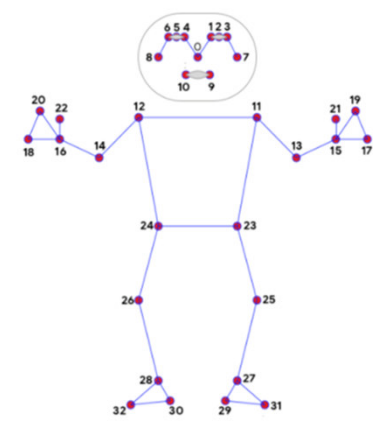
**HRNET**  
17 pt



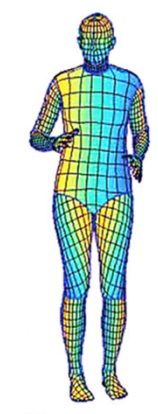
**Openpose**  
18 pt



**Openpose**  
25 pt



**Mediapipe Balzepose**  
33 pt



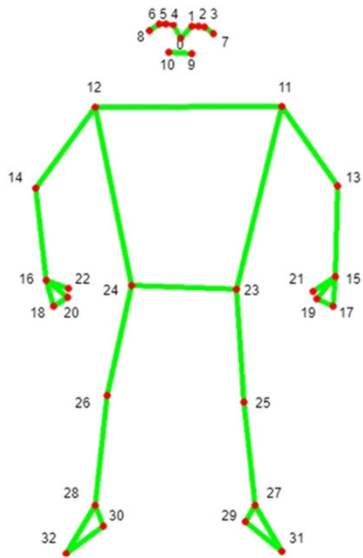
**Densepose**  
3D

di Biase, L., et al. (2025). *Sensors*.  
<https://doi.org/10.3390/s25206373>

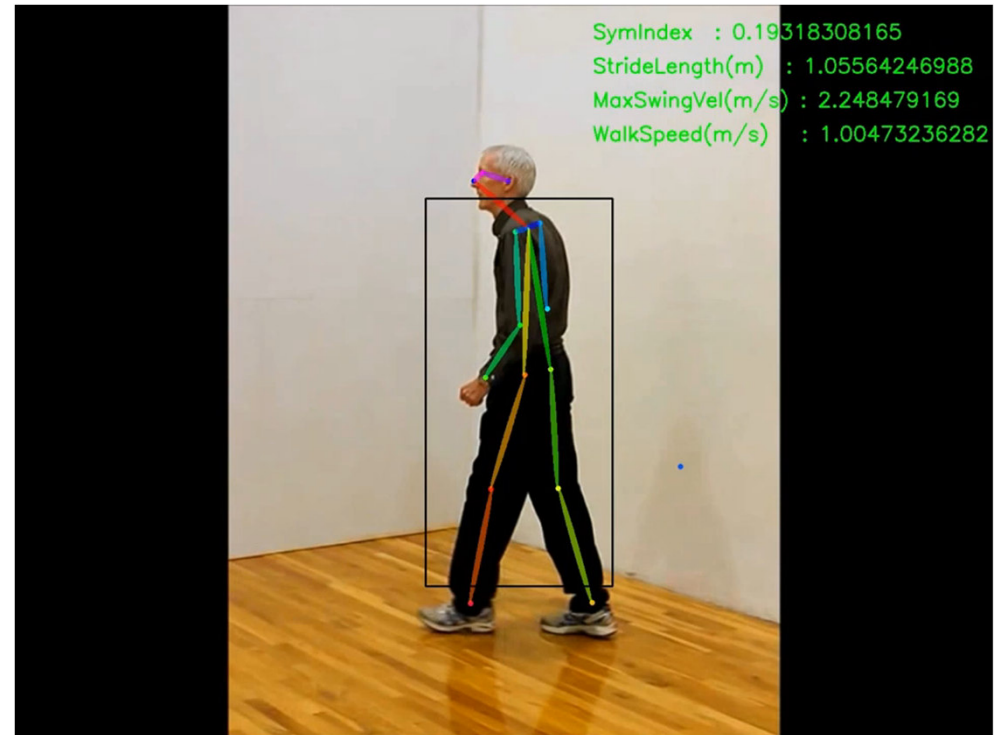


Camera-based

# Full body analysis



- |                    |                      |
|--------------------|----------------------|
| 0. nose            | 17. left pinky       |
| 1. left eye inner  | 18. right pinky      |
| 2. left eye        | 19. left index       |
| 3. left eye outer  | 20. right index      |
| 4. right eye inner | 21. left thumb       |
| 5. right eye       | 22. right thumb      |
| 6. right eye outer | 23. left hip         |
| 7. left ear        | 24. right hip        |
| 8. right ear       | 25. left knee        |
| 9. mouth left      | 26. right knee       |
| 10. mouth right    | 27. left ankle       |
| 11. left shoulder  | 28. right ankle      |
| 12. right shoulder | 29. left heel        |
| 13. left elbow     | 30. right heel       |
| 14. right elbow    | 31. left foot index  |
| 15. left wrist     | 32. right foot index |
| 16. right wrist    |                      |



Lazzaro di Biase, MD, PhD

[lazzaro.dibiase@braininnovations.eu](mailto:lazzaro.dibiase@braininnovations.eu)

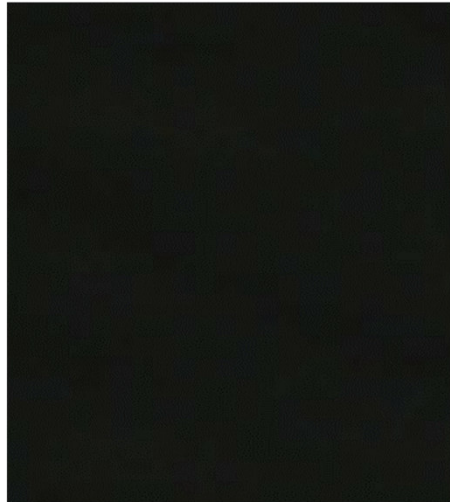
Brain Innovations

# Bradykinesia

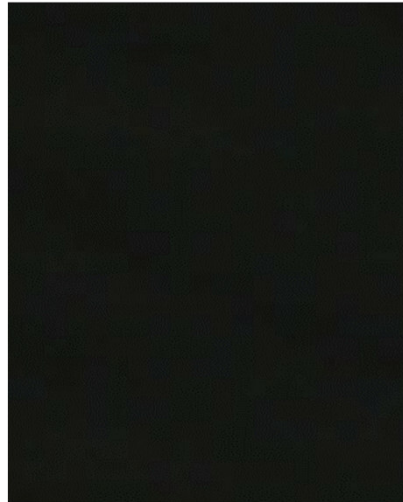
Slowness of initiation of voluntary movements

with progressive reduction in speed and amplitude of repetitive action

- 0: Normal: No problems.
- 1: Slight: Any of the following: a) the regular rhythm is broken with one or two interruptions or hesitations of the tapping movement; b) slight slowing; c) the amplitude decrements near the end of the 10 taps.
- 2: Mild: Any of the following: a) 3 to 5 interruptions during tapping; b) mild slowing; c) the amplitude decrements midway in the 10-tap sequence.
- 3: Moderate: Any of the following: a) more than 5 interruptions during tapping or at least one longer arrest (freeze) in ongoing movement; b) moderate slowing; c) the amplitude decrements starting after the 1st tap.
- 4: Severe: Cannot or can only barely perform the task because of slowing, interruptions or decrements.



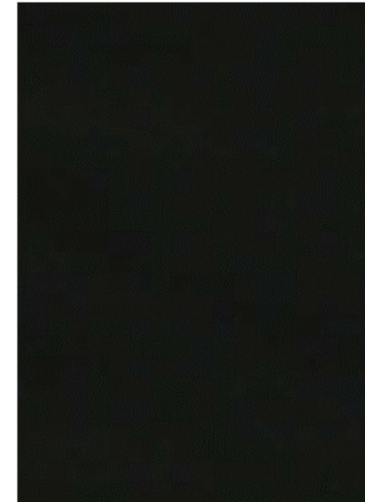
Normal



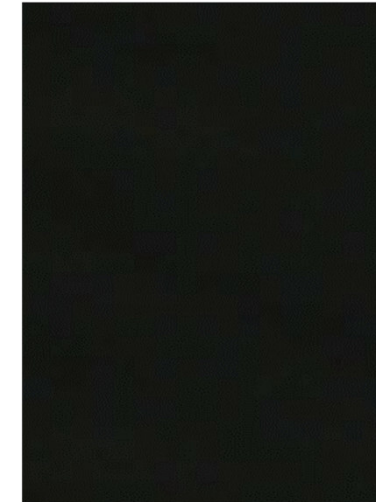
Slight Bradykinesia



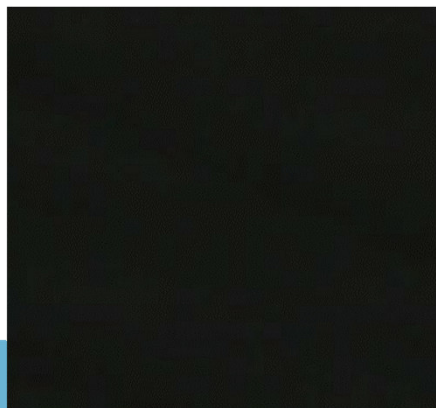
Mild Bradykinesia



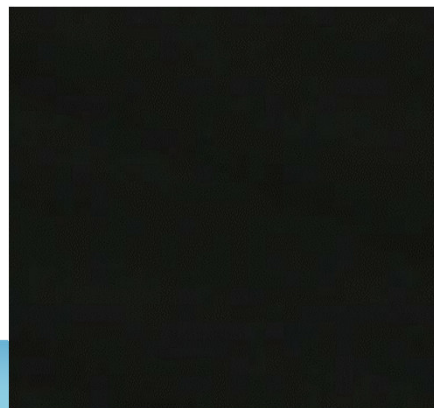
Moderate Bradykinesia



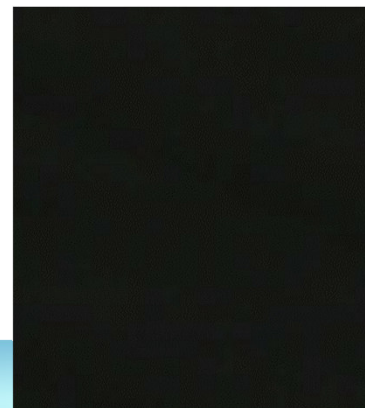
Severe Bradykinesia



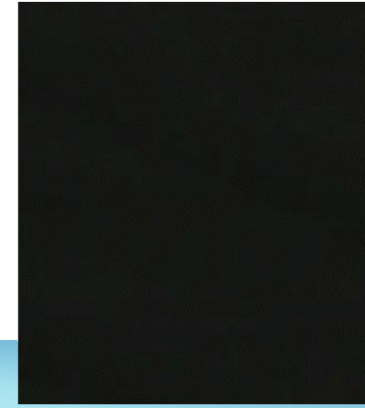
Normal



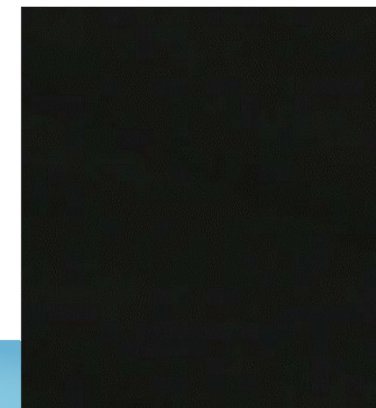
Slight Bradykinesia



Mild Bradykinesia



Moderate Bradykinesia



Severe Bradykinesia

# Bradykinesia

- **Slowness** of initiation of voluntary movements
- with **progressive reduction** in **speed** and **amplitude** of repetitive action

- 0: Normal: No problems.
- 1: Slight: Any of the following: a) the regular rhythm is broken with one or two interruptions or hesitations of the tapping movement; b) slight slowing; c) the amplitude decrements near the end of the 10 taps.
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- 4: Severe: Cannot or can only barely perform the task because of slowing, interruptions or decrements.

Normal



Slight Bradykinesia



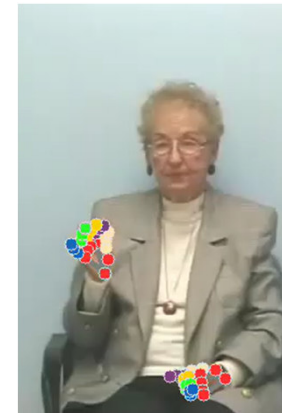
Mild Bradykinesia



Moderate Bradykinesia



Severe Bradykinesia



a: 0,15 px  
v: 0,91 px/s

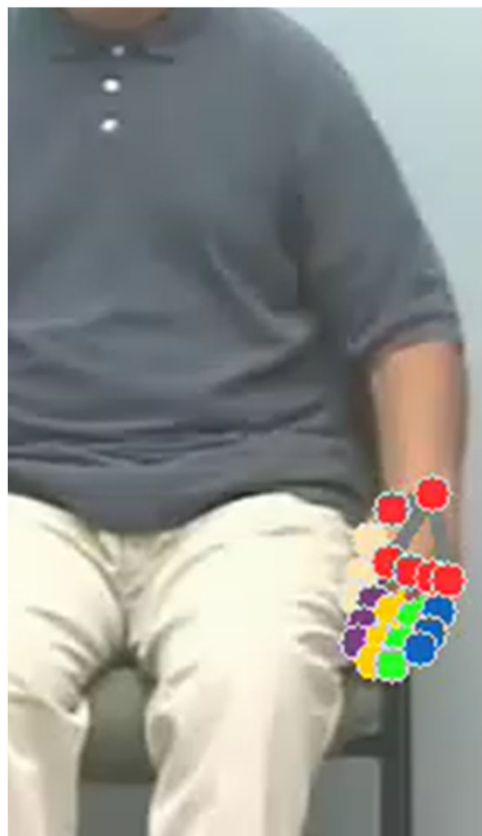
a: 0,07 px  
v: 0,44 px/s

a: 0,05 px  
v: 0,14 px/s

a: 0,01 px  
v: 0,13 px/s

a: 0,02 px  
v: 0,08 px/s

# Rest Tremor



f: 6,1 Hz



f: 2,4 Hz

# VIDEO ANALYSIS

## FINGER TAPPING

---

**ON**

Number of touch:0  
Frequency: 0.0 Hz



**OFF**

Number of touch:0  
Frequency: 0.0 Hz

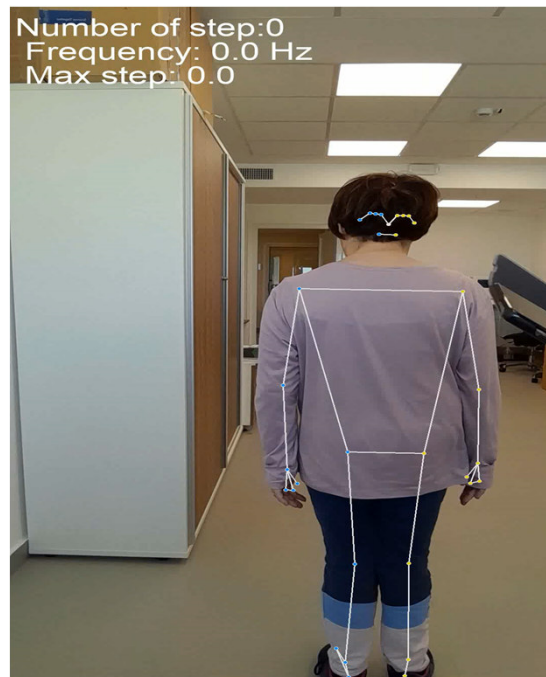


# VIDEO ANALYSIS

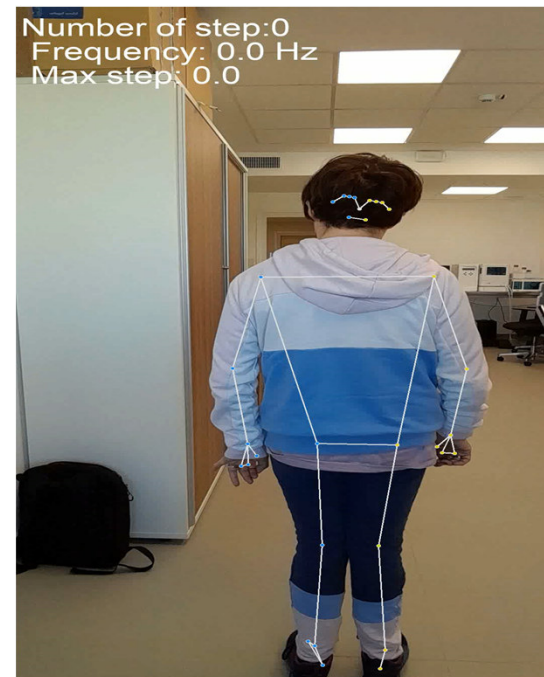
GAIT

---

**ON**  
**mild dyskinesias**



**OFF**



# Caso clinico: caratteristiche generali del paziente

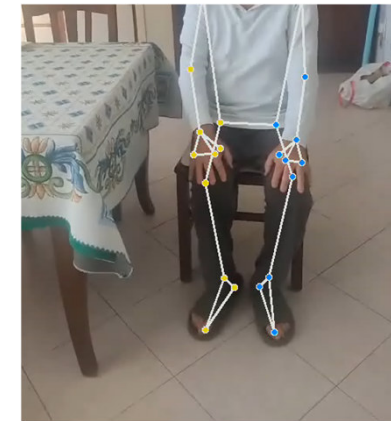
## Overview dei paziente

- Età al momento dell'impianto: 68 anni
- Sesso: M
- Situazione familiare: caregiver (moglie), figlia
- Status lavorativo: pensionato

## Caratteristiche cliniche

- Età alla diagnosi: 59 anni
- Durata di malattia: 9 anni
- Anno di comparsa delle fluttuazioni motorie: 65 anni
- Off: 3-4 ore, frequenti cadute, Discinesie 2-3 ore
- Presenza di sintomi non motori: iposmia, stipsi, RBD, sindrome ansioso-depressiva, difficoltà prassico costruttiva con profilo cognitivo globale conservato, allucinazioni e shopping compulsivo
- Sonno: insonnia intermedia
- Presenza di acinesia al risveglio e notturna
- Comorbidità: deficit ipofisario (apoplezia ipofisaria)

OFF



f 0,54 Hz

OFF



f 0,44 Hz

# Caso clinico: caratteristiche generali del paziente

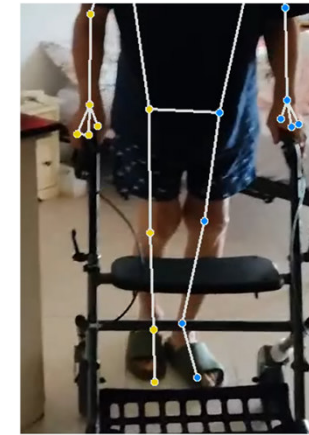
## Overview dei paziente

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- Sonno: insonnia intermedia
- Presenza di acinesia al risveglio e notturna
- Comorbidità: deficit ipofisario (apoplezia ipofisaria)

OFF



vel 0,64 px/s

ON



vel 1,69 px/s

# Caso clinico: caratteristiche generali del paziente

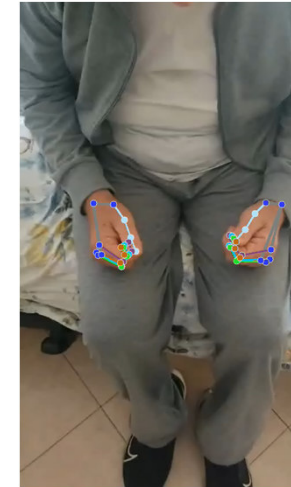
## Overview dei paziente

- Età al momento dell'impianto: 68 anni
- Sesso: M
- Situazione familiare: caregiver (moglie), figlia
- Status lavorativo: pensionato

## Caratteristiche cliniche

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- Sonno: insonnia intermedia
- Presenza di acinesia al risveglio e notturna
- Comorbidità: deficit ipofisario (apoplessia ipofisaria)

OFF



f 0,1 Hz

ON



f 0,8 Hz

# Caso Clinico: implementazione della terapia

## Descrizione dell'implementazione

- **Modalità di implementazione:** ambulatorio
- **Durata dell'implementazione:** 1h (due visite di 30 min separate da 3 h)
- **LEDD pre impianto** 825 mg
- **Terapia infusionale sottocutanea con FOSCARBIDOPA/FOSLEVODOPA 120/2400 mg:**

LE <sub>24</sub> (valore di LE derivante da tutti i medicinali orali contenenti levodopa assunti in un periodo di veglia di 16 ore), (mg)	Velocità di infusione oraria iniziale raccomandata per Duodopa soluzione per infusione sottocutanea (ml/h) somministrato nel corso delle 24 ore
<400	0,15
400-499	0,15-0,17
500-599	0,17-0,20
600-699	0,20-0,24
700-799	0,24-0,27
800-899	0,27-0,30
900-999	0,30-0,34
1000-1099	0,34-0,37

- **Dose base** 0,25 ml/h
- **Dose alta** 0,27 ml/h
- **Dose notturna** 0,22 ml/h
- **Dose Extra** 0,20 ml
- **Dose di carico** 0,6 ml
- **Ago** 9 mm

- **Terapie associate:** Safinamide 100 mg
- **Formazione al paziente/caregiver:** in sede ed a casa
- **Efficacia clinica:** buona
- **Off:** 1 ora (trattato con dose extra), regressione delle cadute, Discinesie: lievi; risoluzione di acinesia notturna e al risveglio
- **Soddisfazione del paziente e del caregiver:** ottima

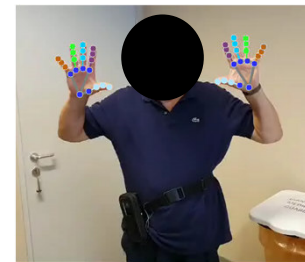
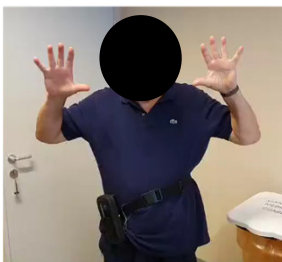
ON



f 1 Hz

$$\frac{\text{ON}}{\text{OFF}} = \times 2$$

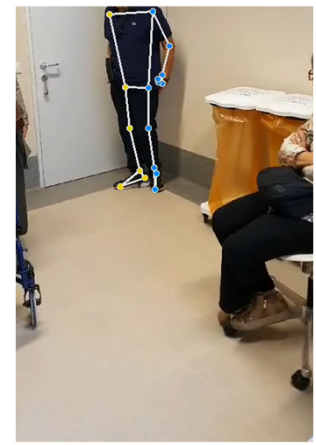
ON



f 0,85 Hz

$$\frac{\text{ON}}{\text{OFF}} = \times 8,5$$

ON



vel 1,5 px/s

$$\frac{\text{ON}}{\text{OFF}} = \times 2,3$$

# Parkinson's disease telemanagement



OFF

vel 0,64 px/s

**Ldopa Orale**  
825 mg (6 dosi ogni 3 h)



ON

vel 1,5 px/s

**SC FosLDopa**  
Dose base 0,25 ml/h 16h/die  
Dose notturna 0,22 ml/h 8h/die  
Dose alta 0,27 ml/h  
Dose Extra 0,20 ml



ON

vel 1,5 px/s

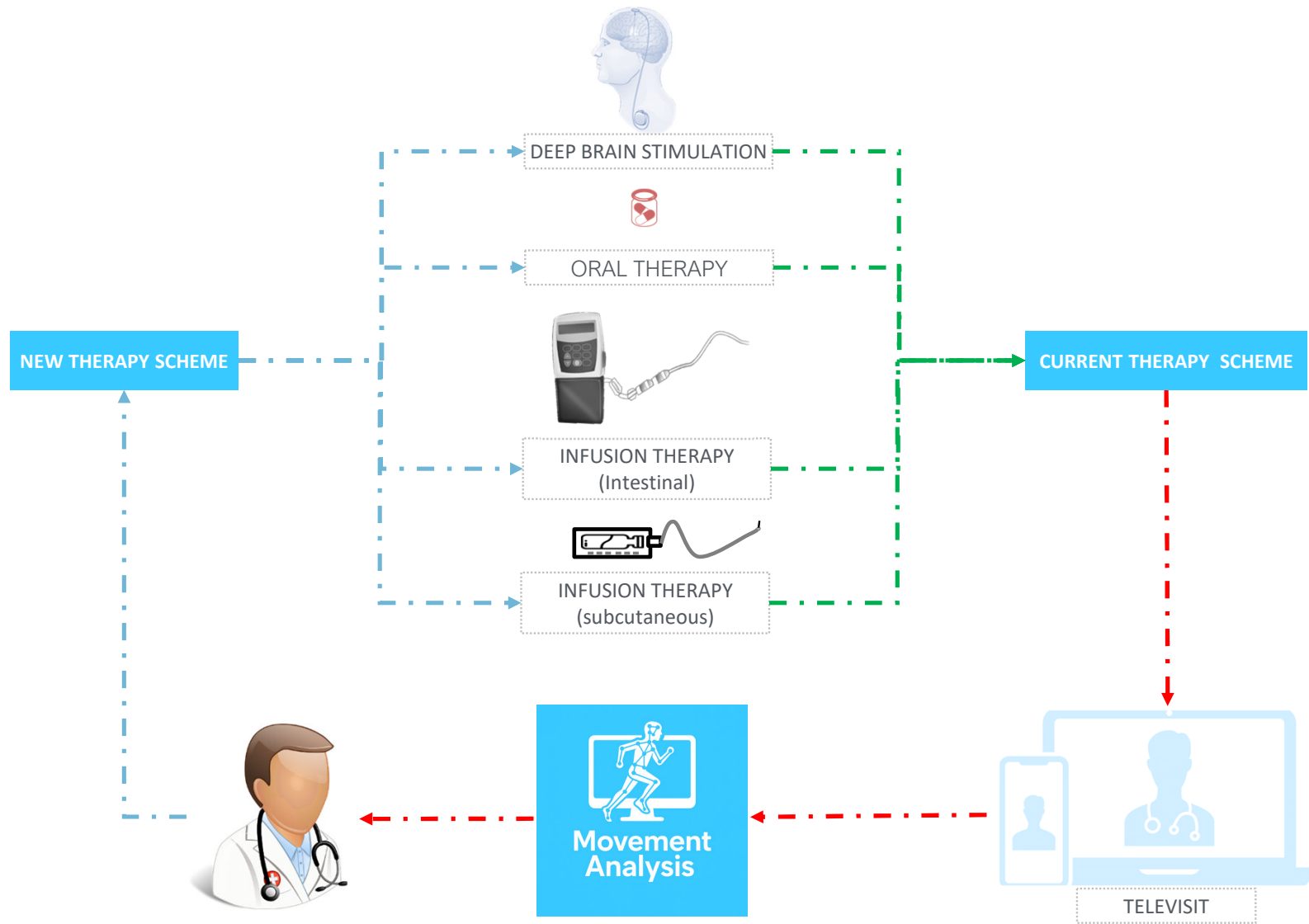
**SC FosLDopa**  
Dose alta 0,27 ml/h 16h/die  
Dose notturna 0,22 ml/h 8h/die  
Dose base 0,25 ml/h  
Dose Extra 0,20 ml



ON

vel 1,7 px/s

**SC FosLDopa**  
Dose alta 0,30 ml/h 16h/die  
Dose notturna 0,25 ml/h 8h/die  
Dose base 0,28 ml/h  
Dose Extra 0,3 ml



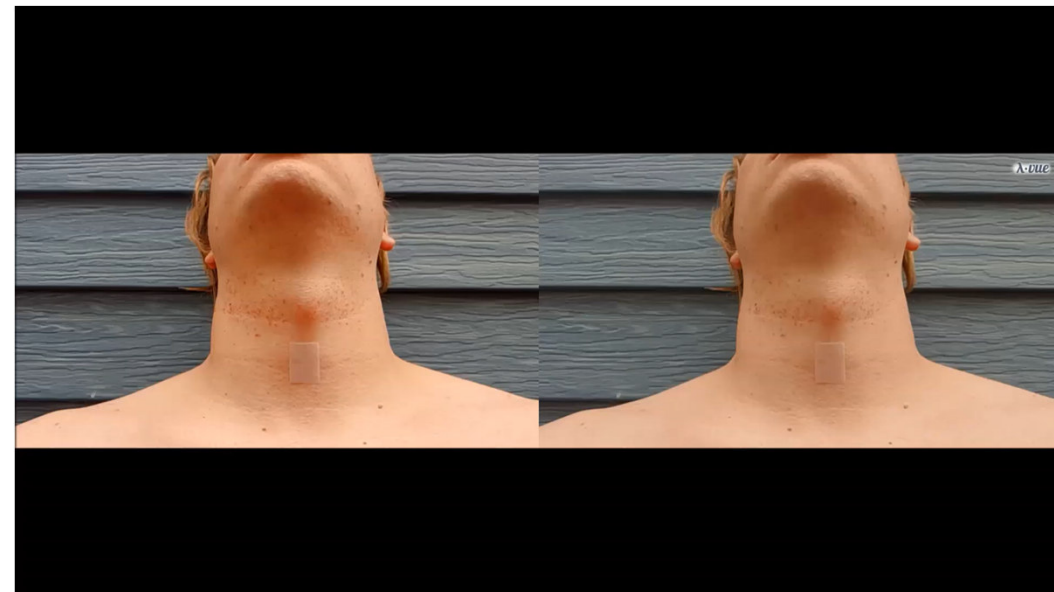
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# VIDEO Magnification

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# Video magnification

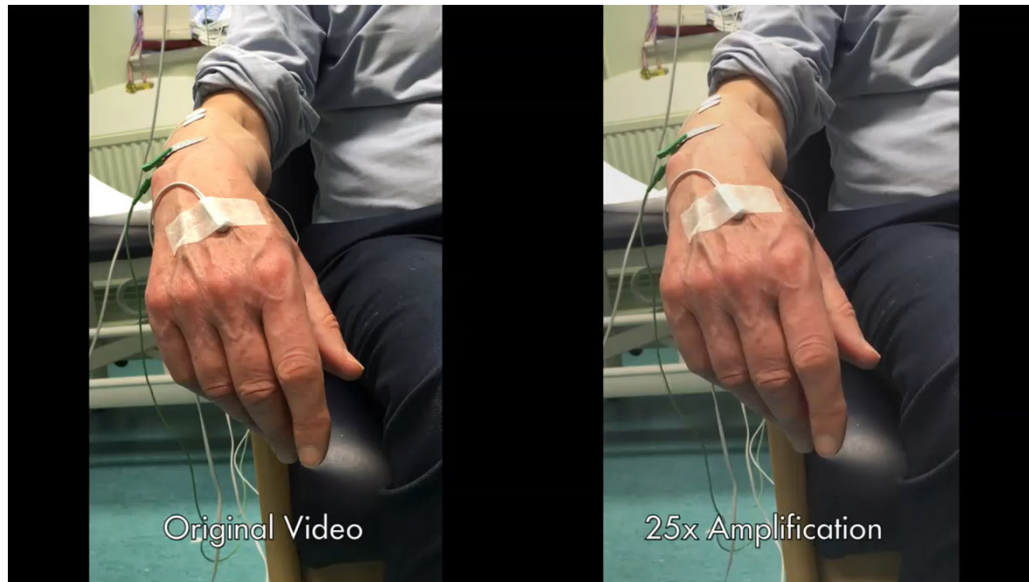


Lauridsen H, et al. Extracting physiological information in experimental biology via Eulerian video magnification. *BMC Biol.* 2019 Dec 12;17(1):103. doi: 10.1186/s12915-019-0716-7. PMID: 31831016; PMCID: PMC6907275.

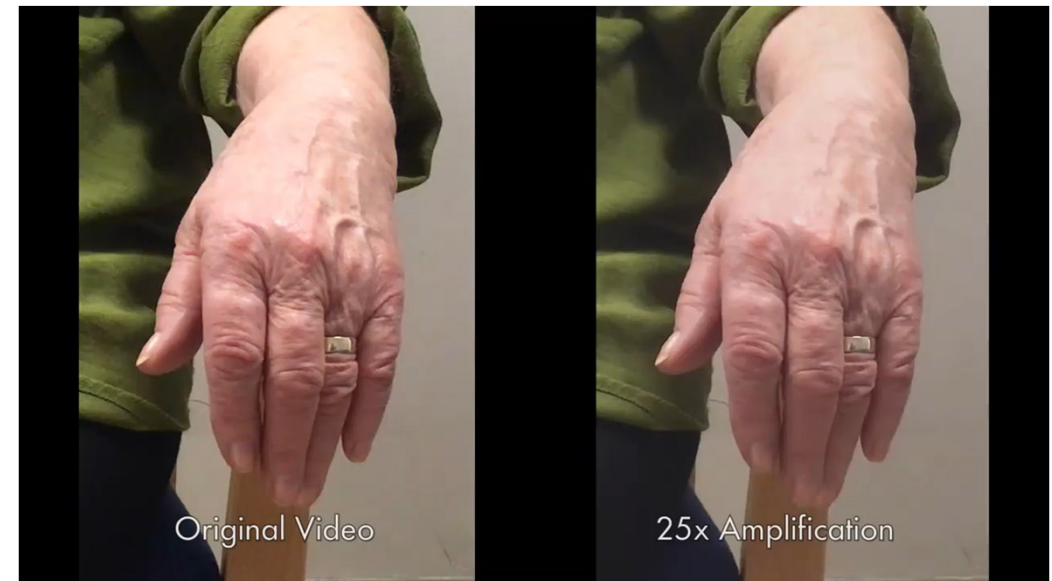
# Video magnification

---

## Parkinson's disease

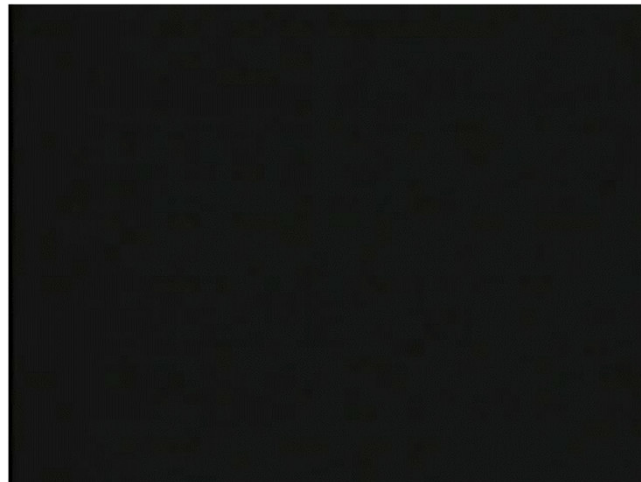


## Healthy subject



Williams S, et al. Seeing the unseen: Could Eulerian video magnification aid clinician detection of subclinical Parkinson's tremor? *J Clin Neurosci*. 2020 Nov;81:101-104. doi: 10.1016/j.jocn.2020.09.046. Epub 2020 Oct 2. PMID: 33222895.

# Rigidity



- 0: Normal: No rigidity.
- 1: Slight: Rigidity only detected with activation maneuver.
- 2: Mild: Rigidity detected without the activation maneuver, but full range of motion is easily achieved.
- 3: Moderate: Rigidity detected without the activation maneuver; full range of motion is achieved with effort.
- 4: Severe: Rigidity detected without the activation maneuver and full range of motion not achieved.

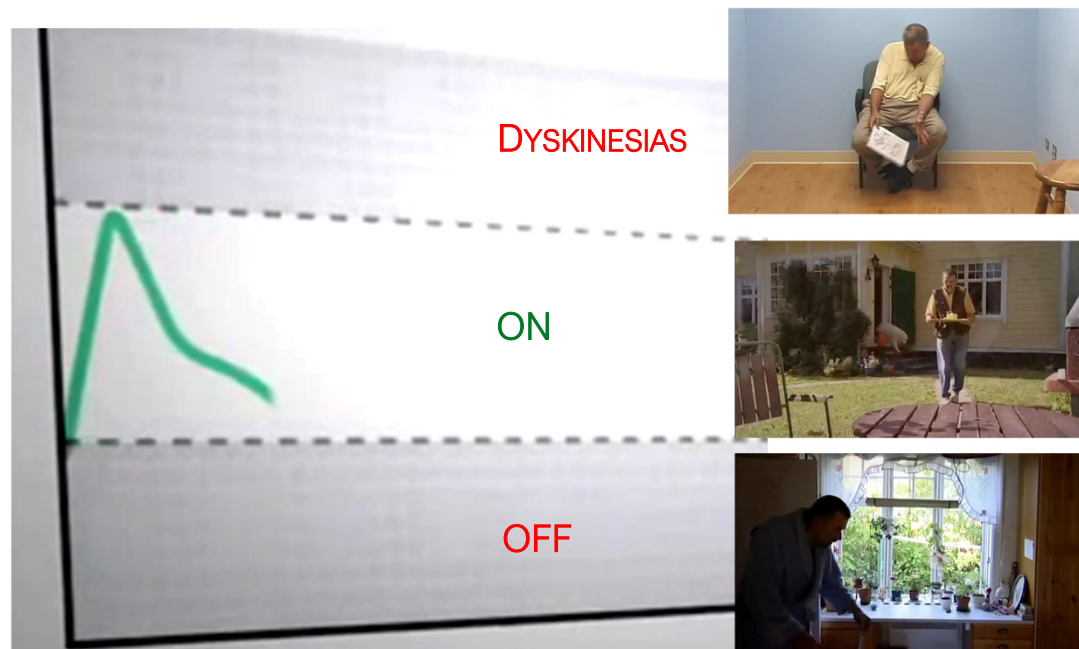
# Rigidity



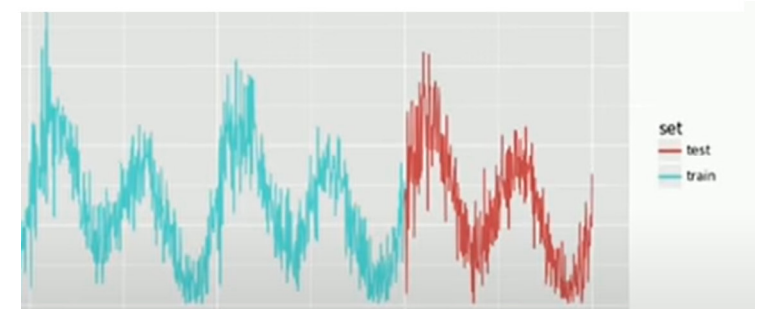
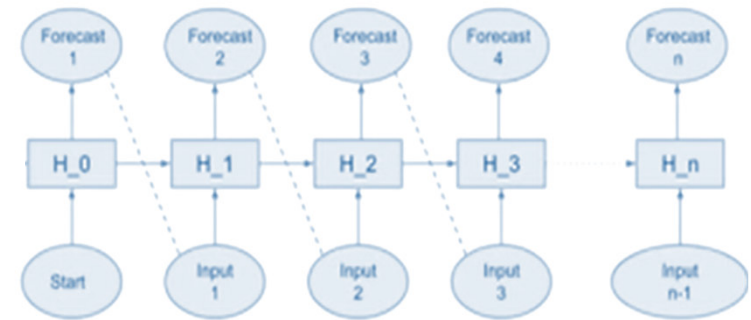
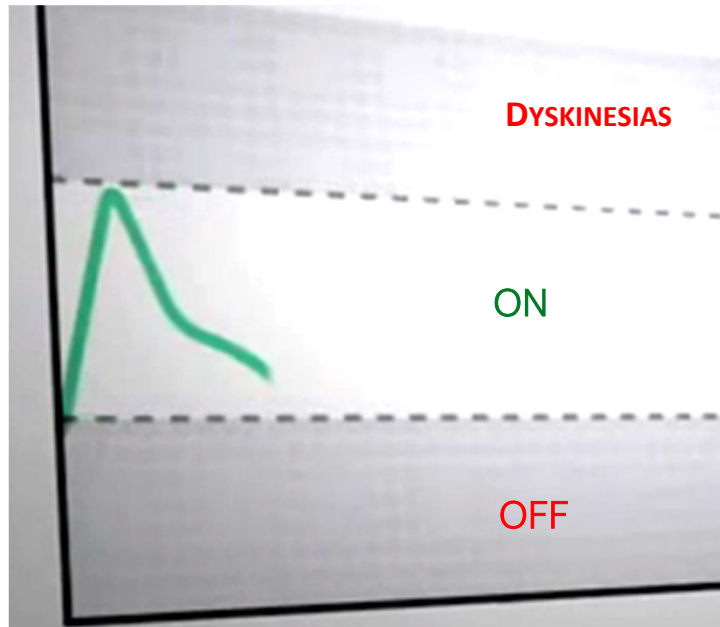
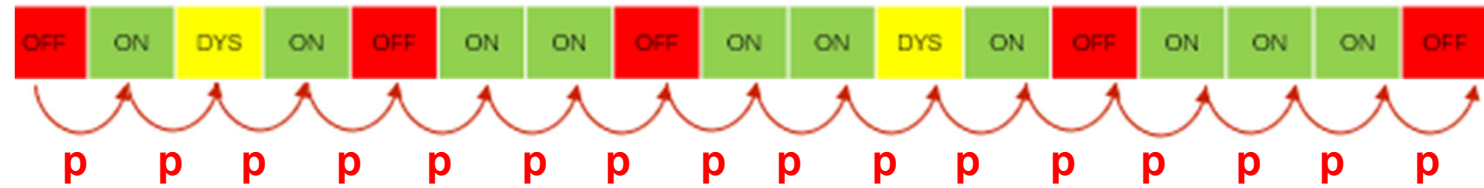
Slow motion 0.5x

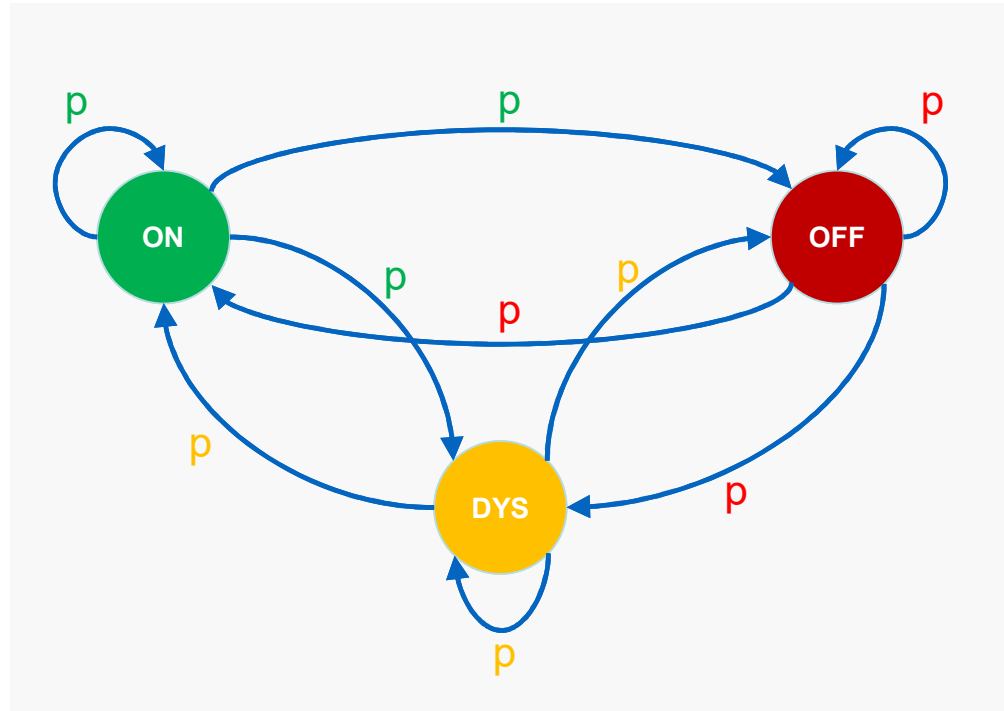
# Motor Fluctuations

Costant fluctuation around ideal motor status

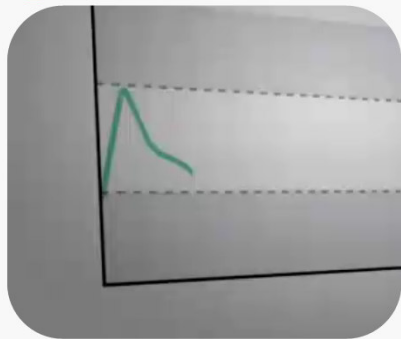


# Therapy management

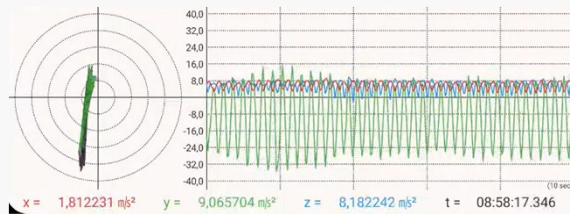




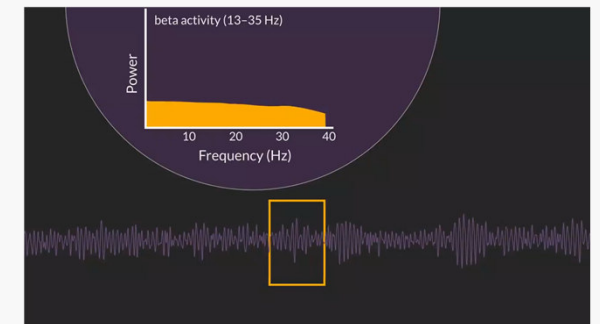
### BIOCHEMICAL MONITORING



### MOVEMENT MONITORING

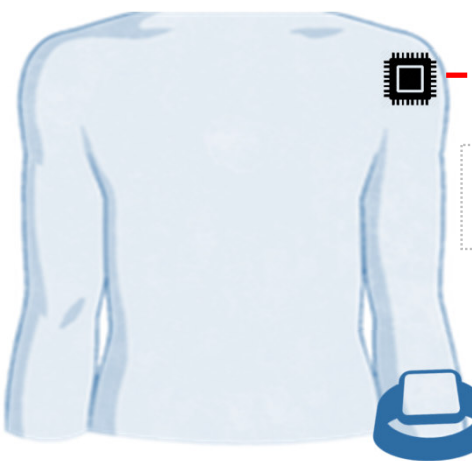


### BRAIN ACTIVITY MONITORING





LFP  
MONITORING



BIOCHEMICAL  
MONITORING



KINEMATIC  
MONITORING

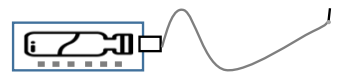
DEEP BRAIN STIMULATION



ORAL THERAPY



INFUSION THERAPY  
(Intestinal)



INFUSION THERAPY  
(subcutaneous)

NEW THERAPY SCHEME

CURRENT THERAPY SCHEME

SENSING SYSTEM

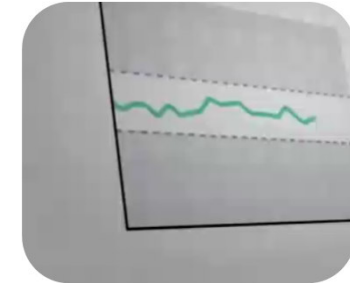
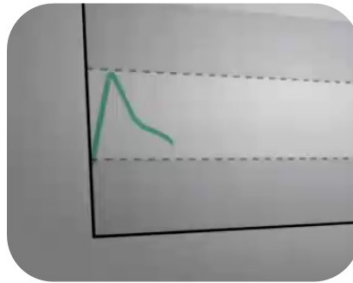
ADAPTIVE CLOSED-LOOP  
motor function stabilization algorithm

di Biase, Lazzaro. (2021). Adaptive closed-loop therapy system for Parkinson's disease. <https://doi.org/10.5281/zenodo.4767082>

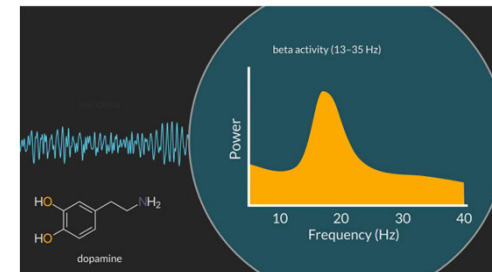
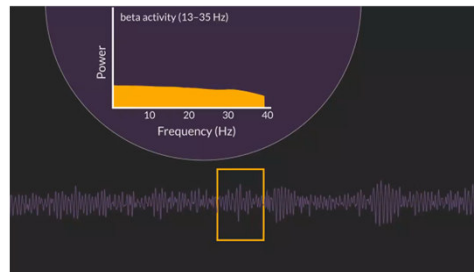
# Therapy management solutions



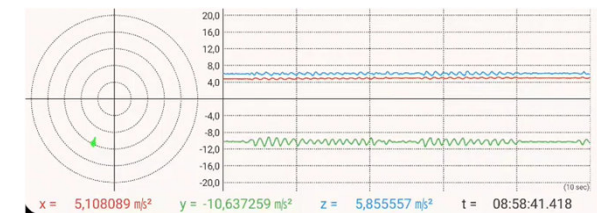
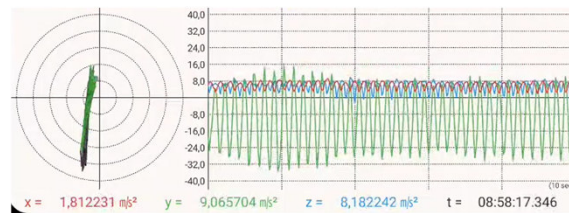
BIOCHEMICAL  
MONITORING



BRAIN ACTIVITY  
MONITORING



MOVEMENT MONITORING



di Biase L, et al. Adaptive, personalized closed-loop therapy for Parkinson's disease: biochemical, neurophysiological, and wearable sensing systems. *Expert Rev Neurother*. 2021 Dec;21(12):1371-1388. doi: 10.1080/14737175.2021.2000392. Epub 2021 Nov 17.

Lazzaro di Biase, MD, PhD

[l.dibiase@policlinicocampus.it](mailto:l.dibiase@policlinicocampus.it)

Neurology Unit, Campus Bio-Medico University Hospital

Brain Innovations Lab

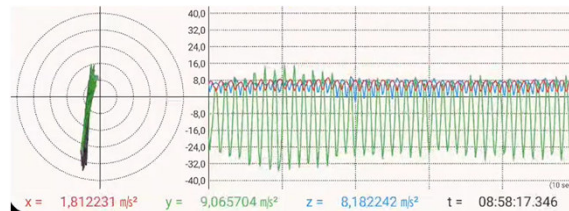
---

# Movement monitoring

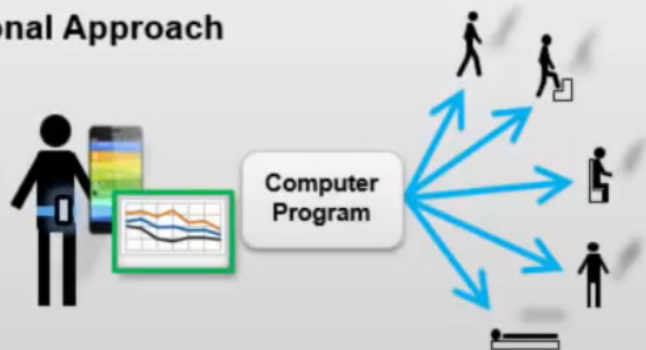
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Wearable IMU



## Traditional Approach



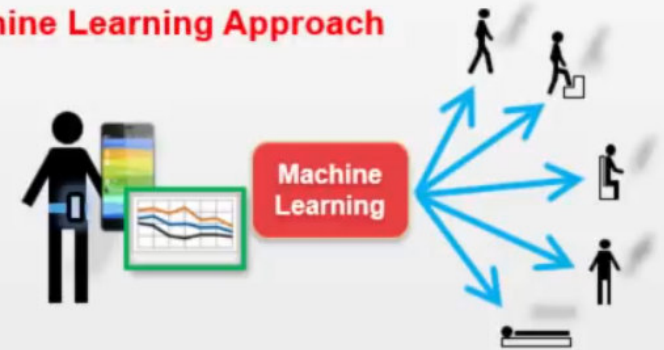
### Hand Written Program

If  $X_{acc} > 0.5$   
then "SITTING"  
If  $Y_{acc} < 4$  and  $Z_{acc} > 5$   
then "STANDING"  
...

### Formula or Equation

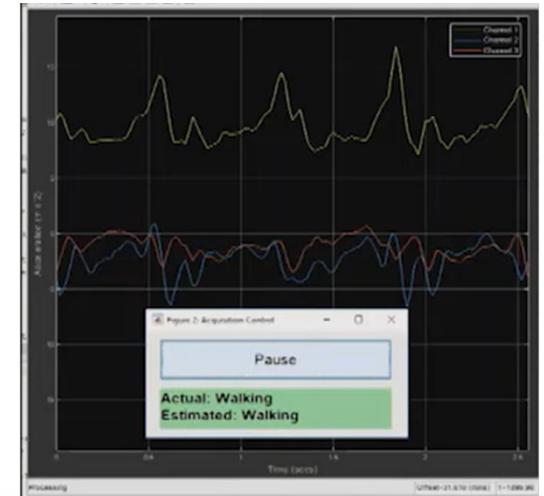
$$Y_{activity} = \beta_1 X_{acc} + \beta_2 Y_{acc} + \beta_3 Z_{acc} + \dots$$

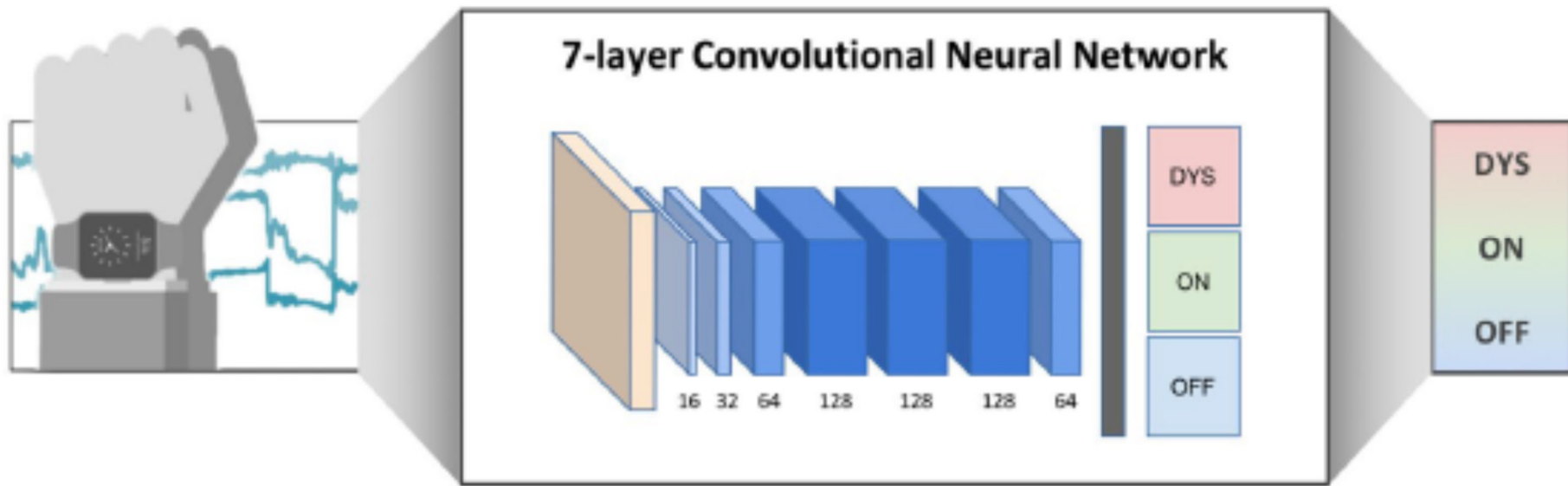
## Machine Learning Approach



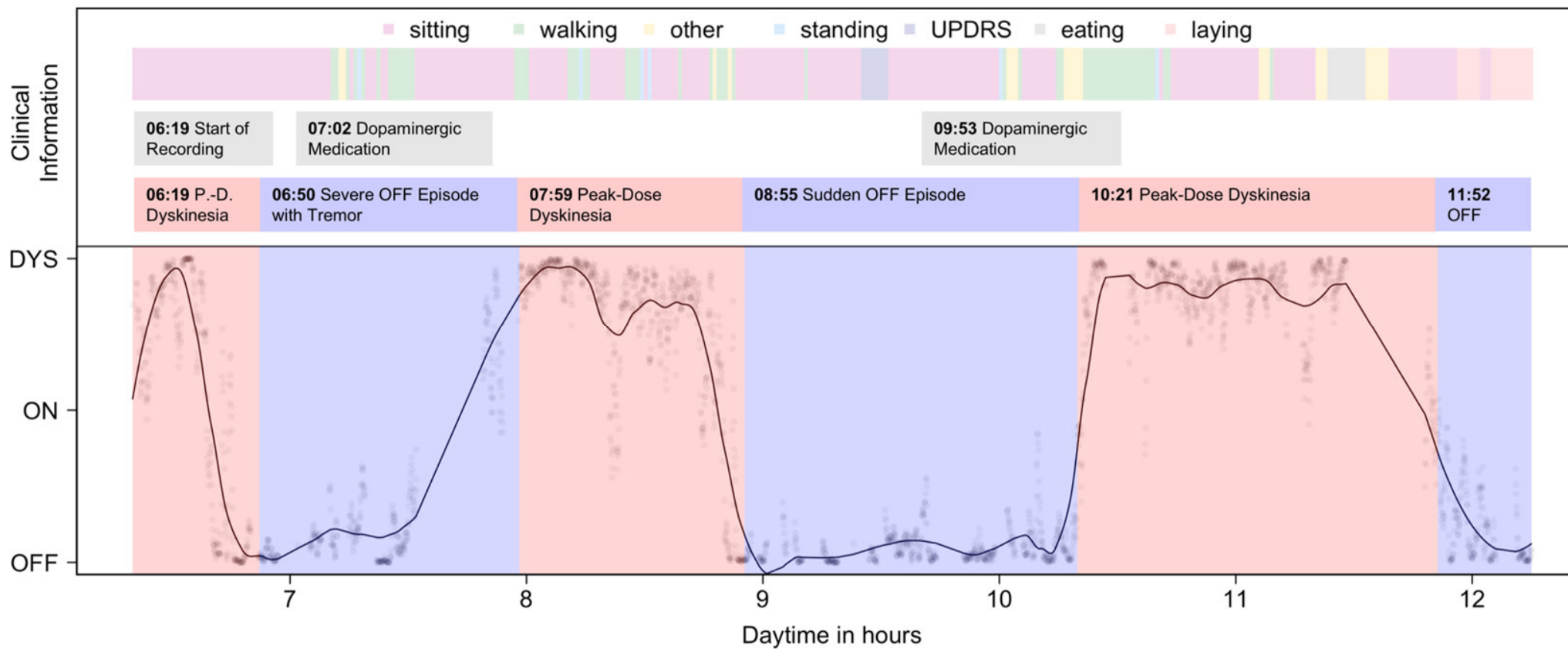
*model*: Inputs → Outputs

*model* = < Machine Learning Algorithm > (sensor\_data, activity)





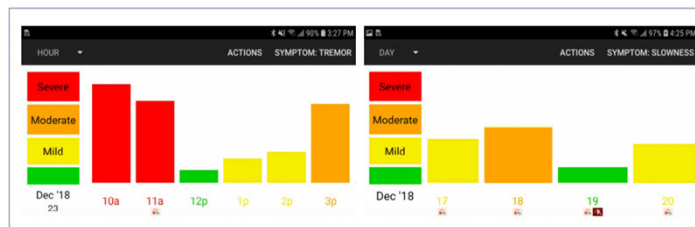
Pfister - High-Resolution Motor State Detection in Parkinson's Disease Using Convolutional Neural Networks



Pfister - High-Resolution Motor State Detection in Parkinson's Disease Using Convolutional Neural Networks



Objective Motor Assessment Using A Smartwatch



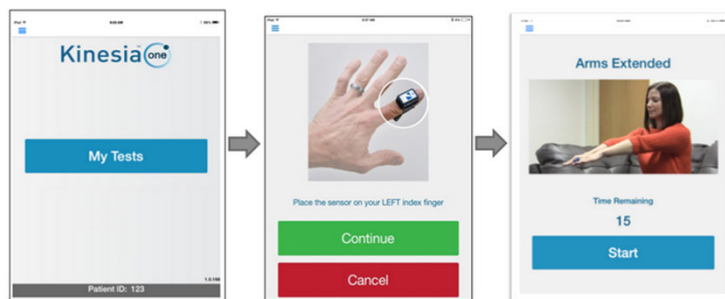
DEVELOPMENT STAGE



CERTIFICATIONS



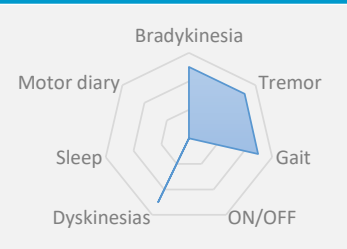
Objective, Task-Based Motor Assessments Using Sensor & App



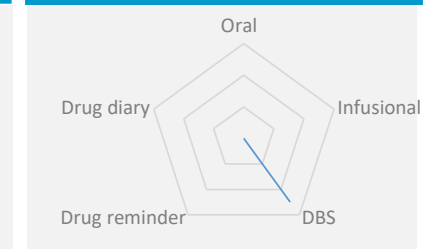
DIAGNOSIS



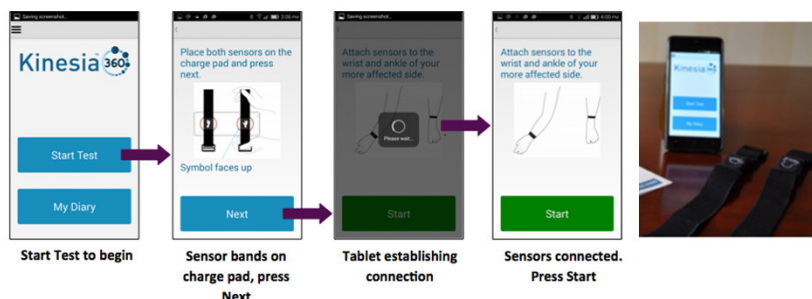
SYMPTOMS MONITORING



THERAPY MANAGEMENT

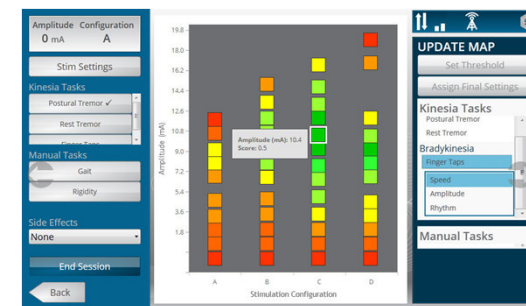
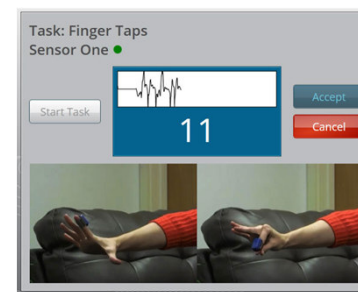


Objective, Continuous Symptom Monitoring Using Wearables & App



Visualize Symptom Response To Deep Brain Stimulation

The Kinesia ProView kit includes a carrying case, tablet PC with integrated broadband, and charge pad with wireless sensors to capture motor symptom severity.



# Task-Based Motor Assessments



Kinesia<sup>™</sup>one

Time	Rest Tremor	Postural Tremor	Finger Taps Speed	Finger Taps Amplitude	Finger Taps Rhythm	Dyskinesia
7:01 AM	4.0	3.5	2.5	2.4	2.2	0.0
7:02 AM	SINEMET (100mg)					
7:32 AM	3.4	3.3	1.7	1.4	1.0	0.0
8:01 AM	3.0	3.0	1.8	1.8	1.2	0.0
8:34 AM	2.9	2.8	1.3	1.2	1.0	0.0
9:00 AM	2.8	2.4	1.2	1.1	1.2	0.0
9:23 AM	2.8	2.6	1.0	1.0	1.0	0.0
10:00 AM	2.6	2.8	1.0	1.0	1.0	0.0
10:33 AM	3.2	3.3	1.5	1.9	1.5	0.0
11:01 AM	3.5	3.5	2.3	2.2	2.0	0.0
11:30 AM	3.7	3.8	2.0	2.0	1.8	0.0
12:00 PM	SINEMET (100mg)					
12:01 PM	3.3	3.8	2.6	2.7	2.0	0.0
12:32 PM	3.2	3.4	1.8	1.9	2.0	0.0
1:08 PM	2.6	3.1	2.0	1.4	1.8	0.0
1:28 PM	2.6	2.9	1.5	1.2	1.7	0.0
2:00 PM	2.7	2.7	1.3	1.0	1.5	0.0
2:32 PM	2.9	2.6	1.0	1.2	1.7	0.0
3:00 PM	3.0	2.9	1.1	1.5	1.3	0.0
3:29 PM	3.3	3.1	1.4	1.7	1.7	0.0
4:02 PM	3.8	3.6	1.6	1.8	1.8	0.0
4:30 PM	3.9	3.8	1.9	1.9	2.0	0.0
5:01 PM	3.9	3.9	2.5	2.4	2.0	0.0
5:15 PM	SINEMET (100mg)					
5:29 PM	3.5	3.6	2.1	2.2	2.0	0.0
6:02 PM	3.3	3.5	2.0	2.1	1.6	0.0
6:30 PM	3.0	2.9	1.9	2.0	1.5	0.0
7:00 PM	2.8	2.5	1.5	1.8	1.3	0.0
7:33 PM	2.6	2.6	1.2	1.5	1.1	0.0
8:04 PM	2.6	2.6	1.0	1.4	0.9	0.0
8:30 PM	2.9	2.8	1.2	1.5	1.1	0.0
9:02 PM	3.3	3.2	1.3	1.6	1.4	0.0
9:33 PM	3.5	3.6	1.6	1.8	1.8	0.0
10:00 PM	3.8	3.9	2.0	1.9	2.1	0.0
Mean	3.2	3.2	1.6	1.7	1.6	0.0
Fluctuation	0.4	0.5	0.5	0.4	0.4	0.0

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 Neurology Unit, Campus Bio-Medico University Hospital  
 Brain Innovations Lab

# Task-Based Motor Assessments



Kinesia<sup>™</sup>one

Time	Rest Tremor	Postural Tremor	Finger Taps Speed	Finger Taps Amplitude	Finger Taps Rhythm	Dyskinesia
7:01 AM	4.0	3.5	2.5	2.4	2.2	0.0
7:02 AM	SINEMET (100mg)					
7:32 AM	3.4	3.3	1.7	1.4	1.0	0.0
8:01 AM	3.0	3.0	1.8	1.8	1.2	0.0
8:34 AM	2.9	2.8	1.3	1.2	1.0	0.0
9:00 AM	2.8	2.4	1.2	1.1	1.2	0.0
9:23 AM	2.8	2.6	1.0	1.0	1.0	0.0
10:00 AM	2.6	2.8	1.0	1.0	1.0	0.0
10:33 AM	3.2	3.3	1.5	1.9	1.5	0.0
11:01 AM	3.5	3.5	2.3	2.2	2.0	0.0
11:30 AM	3.7	3.8	2.0	2.0	1.8	0.0
12:00 PM	SINEMET (100mg)					
12:01 PM	3.3	3.8	2.6	2.7	2.0	0.0
12:32 PM	3.2	3.4	1.8	1.9	2.0	0.0
1:08 PM	2.6	3.1	2.0	1.4	1.8	0.0
1:28 PM	2.6	2.9	1.5	1.2	1.7	0.0
2:00 PM	2.7	2.7	1.3	1.0	1.5	0.0
2:32 PM	2.9	2.6	1.0	1.2	1.7	0.0
3:00 PM	3.0	2.9	1.1	1.5	1.3	0.0
3:29 PM	3.3	3.1	1.4	1.7	1.7	0.0
4:02 PM	3.8	3.6	1.6	1.8	1.8	0.0
4:30 PM	3.9	3.8	1.9	1.9	2.0	0.0
5:01 PM	3.9	3.9	2.5	2.4	2.0	0.0
5:15 PM	SINEMET (100mg)					
5:29 PM	3.5	3.6	2.1	2.2	2.0	0.0
6:02 PM	3.3	3.5	2.0	2.1	1.6	0.0
6:30 PM	3.0	2.9	1.9	2.0	1.5	0.0
7:00 PM	2.8	2.5	1.5	1.8	1.3	0.0
7:33 PM	2.6	2.6	1.2	1.5	1.1	0.0
8:04 PM	2.6	2.6	1.0	1.4	0.9	0.0
8:30 PM	2.9	2.8	1.2	1.5	1.1	0.0
9:02 PM	3.3	3.2	1.3	1.6	1.4	0.0
9:33 PM	3.5	3.6	1.6	1.8	1.8	0.0
10:00 PM	3.8	3.9	2.0	1.9	2.1	0.0
Mean	3.2	3.2	1.6	1.7	1.6	0.0
Fluctuation	0.4	0.5	0.5	0.4	0.4	0.0
Time	Rest Tremor	Postural Tremor	Finger Taps Speed	Finger Taps Amplitude	Finger Taps Rhythm	Dyskinesia
6:55 AM	3.9	3.4	2.6	2.5	2.3	0.0
6:57 AM	SINEMET (300mg)					
7:28 AM	2.5	3.0	1.7	1.4	1.0	0.0
7:59 AM	0.5	1.9	1.8	1.5	1.2	1.3
8:30 AM	0.3	0.9	0.3	0.5	1.0	2.9
9:05 AM	0.1	0.5	0.2	0.2	1.2	3.5
9:33 AM	0.3	0.4	0.0	0.0	1.0	3.8
10:02 AM	0.5	0.1	0.5	0.3	1.0	3.7
10:31 AM	1.5	2.0	1.0	0.5	1.5	2.9
10:58 AM	3.0	3.1	2.3	2.2	2.0	0.0
11:35 AM	3.5	3.4	2.0	2.0	1.8	0.0
11:50 PM	SINEMET (300mg)					
11:56 PM	1.1	2.7	2.3	2.2	2.0	0.0
12:30 PM	0.2	2.0	1.8	1.9	2.0	3.0
1:04 PM	0.1	1.4	2.0	1.4	1.8	3.3
1:38 PM	0.0	1.1	0.8	0.9	1.7	3.5
2:02 PM	0.0	1.0	0.6	1.0	1.5	3.6
2:30 PM	0.2	1.0	1.0	1.2	1.7	2.4
3:07 PM	0.4	0.7	1.1	1.5	1.3	1.1
3:33 PM	0.5	1.3	1.4	1.7	1.7	0.0
4:03 PM	2.6	1.5	1.6	1.8	1.8	0.0
4:28 PM	3.5	2.0	1.9	1.9	2.0	0.0
5:00 PM	3.8	2.2	2.1	2.1	2.0	0.0
5:05 PM	SINEMET (300mg)					
5:39 PM	3.5	2.2	2.1	2.2	2.0	0.0
6:03 PM	2.3	2.0	2.0	2.1	1.6	0.0
6:29 PM	1.7	1.3	1.9	2.0	1.5	0.5
7:05 PM	0.8	1.1	1.5	1.8	1.3	1.0
7:36 PM	0.6	0.8	1.2	1.5	1.1	2.3
8:01 PM	0.3	0.6	1.0	1.4	0.9	3.8
8:28 PM	0.2	1.0	1.2	1.5	1.1	3.7
9:00 PM	0.3	1.1	1.3	1.6	1.4	1.3
9:34 PM	0.3	2.0	1.6	1.8	1.8	0.5
9:59 PM	2.8	2.3	2.0	1.9	2.1	0.0
Mean	1.3	1.6	1.4	1.5	1.6	1.6
Fluctuation	1.3	0.9	0.7	0.6	0.4	1.5

→  
Increase dose by 200mg, Dose interval unchanged

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# Task-Based Motor Assessments



Kinesia<sup>™</sup>one

Time	Rest Tremor	Postural Tremor	Finger Taps Speed	Finger Taps Amplitude	Finger Taps Rhythm	Dyskinesia
7:01 AM	4.0	3.5	2.5	2.4	2.2	0.0
7:02 AM	SINEMET (100mg)					
7:32 AM	3.4	3.3	1.7	1.4	1.0	0.0
8:01 AM	3.0	3.0	1.8	1.8	1.2	0.0
8:34 AM	2.9	2.8	1.3	1.2	1.0	0.0
9:00 AM	2.8	2.4	1.2	1.1	1.2	0.0
9:23 AM	2.8	2.6	1.0	1.0	1.0	0.0
10:00 AM	2.6	2.8	1.0	1.0	1.0	0.0
10:33 AM	3.2	3.3	1.5	1.9	1.5	0.0
11:01 AM	3.5	3.5	2.3	2.2	2.0	0.0
11:30 AM	3.7	3.8	2.0	2.0	1.8	0.0
12:00 PM	SINEMET (100mg)					
12:01 PM	3.3	3.8	2.6	2.7	2.0	0.0
12:32 PM	3.2	3.4	1.8	1.9	2.0	0.0
1:08 PM	2.6	3.1	2.0	1.4	1.8	0.0
1:28 PM	2.6	2.9	1.5	1.2	1.7	0.0
2:00 PM	2.7	2.7	1.3	1.0	1.5	0.0
2:32 PM	2.9	2.6	1.0	1.2	1.7	0.0
3:00 PM	3.0	2.9	1.1	1.5	1.3	0.0
3:29 PM	3.3	3.1	1.4	1.7	1.7	0.0
4:02 PM	3.8	3.6	1.6	1.8	1.8	0.0
4:30 PM	3.9	3.8	1.9	1.9	2.0	0.0
5:01 PM	3.9	3.9	2.5	2.4	2.0	0.0
5:15 PM	SINEMET (100mg)					
5:29 PM	3.5	3.6	2.1	2.2	2.0	0.0
6:02 PM	3.3	3.5	2.0	2.1	1.6	0.0
6:30 PM	3.0	2.9	1.9	2.0	1.5	0.0
7:00 PM	2.8	2.5	1.5	1.8	1.3	0.0
7:33 PM	2.6	2.6	1.2	1.5	1.1	0.0
8:04 PM	2.6	2.6	1.0	1.4	0.9	0.0
8:30 PM	2.9	2.8	1.2	1.5	1.1	0.0
9:02 PM	3.3	3.2	1.3	1.6	1.4	0.0
9:33 PM	3.5	3.6	1.6	1.8	1.8	0.0
10:00 PM	3.8	3.9	2.0	1.9	2.1	0.0
Mean	3.2	3.2	1.6	1.7	1.6	0.0
Fluctuation	0.4	0.5	0.5	0.4	0.4	0.0
Increase dose by 200mg, Dose interval unchanged						
Time	Rest Tremor	Postural Tremor	Finger Taps Speed	Finger Taps Amplitude	Finger Taps Rhythm	Dyskinesia
6:55 AM	3.9	3.4	2.6	2.5	2.3	0.0
6:57 AM	SINEMET (300mg)					
7:28 AM	2.5	3.0	1.7	1.4	1.0	0.0
7:59 AM	0.5	1.9	1.8	1.5	1.2	1.3
8:30 AM	0.3	0.9	0.3	0.5	1.0	2.9
9:05 AM	0.1	0.5	0.2	0.2	1.2	3.5
9:33 AM	0.3	0.4	0.0	0.0	1.0	3.8
10:02 AM	0.5	0.1	0.5	0.3	1.0	3.7
10:31 AM	1.5	2.0	1.0	0.5	1.5	2.9
10:58 AM	3.0	3.1	2.3	2.2	2.0	0.0
11:35 AM	3.5	3.4	2.0	2.0	1.8	0.0
11:50 PM	SINEMET (300mg)					
11:56 PM	1.1	2.7	2.3	2.2	2.0	0.0
12:30 PM	0.2	2.0	1.8	1.9	2.0	3.0
1:04 PM	0.1	1.4	2.0	1.4	1.8	3.3
1:38 PM	0.0	1.1	0.8	0.9	1.7	3.5
2:02 PM	0.0	1.0	0.6	1.0	1.5	3.6
2:30 PM	0.2	1.0	1.0	1.2	1.7	2.4
3:07 PM	0.4	0.7	1.1	1.5	1.3	1.1
3:33 PM	0.5	1.3	1.4	1.7	1.7	0.0
4:03 PM	2.6	1.5	1.6	1.8	1.8	0.0
4:28 PM	3.5	2.0	1.9	1.9	2.0	0.0
5:00 PM	3.8	2.2	2.1	2.1	2.0	0.0
5:05 PM	SINEMET (300mg)					
5:39 PM	3.5	2.2	2.1	2.2	2.0	0.0
6:03 PM	2.3	2.0	2.0	2.1	1.6	0.0
6:29 PM	1.7	1.3	1.9	2.0	1.5	0.5
7:05 PM	0.8	1.1	1.5	1.8	1.3	1.0
7:36 PM	0.6	0.8	1.2	1.5	1.1	2.3
8:01 PM	0.3	0.6	1.0	1.4	0.9	3.8
8:28 PM	0.2	1.0	1.2	1.5	1.1	3.7
9:00 PM	0.3	1.1	1.3	1.6	1.4	1.3
9:34 PM	0.3	2.0	1.6	1.8	1.8	0.5
9:59 PM	2.8	2.3	2.0	1.9	2.1	0.0
Mean	1.3	1.6	1.4	1.5	1.6	1.6
Fluctuation	1.3	0.9	0.7	0.6	0.4	1.5
Decrease dose by 100mg, Decrease dose interval by 2 hours						
Time	Rest Tremor	Postural Tremor	Finger Taps Speed	Finger Taps Amplitude	Finger Taps Rhythm	Dyskinesia
7:00 AM	3.5	3.2	2.7	2.5	2.4	0.0
7:01 AM	SINEMET (200mg)					
7:31 AM	2.0	2.1	1.9	2.1	2.2	0.0
8:00 AM	0.6	0.7	0.3	0.5	1.0	0.0
8:33 AM	0.3	0.5	0.2	0.2	1.2	0.0
8:59 AM	0.2	0.2	0.0	0.0	1.0	0.0
9:22 AM	0.2	0.0	0.5	0.3	1.0	0.0
9:59 AM	1.1	1.5	1.0	0.5	1.5	0.0
10:32 AM	SINEMET (200mg)					
11:00 AM	1.2	1.3	1.5	1.4	1.5	0.0
11:29 AM	0.3	0.3	0.5	0.6	2.1	0.0
11:59 PM	0.2	0.2	0.3	0.3	1.0	0.0
12:00 PM	0.1	0.0	0.4	0.1	2.3	0.0
12:31 PM	0.2	0.6	0.6	0.1	2.1	0.0
1:07 PM	1.2	1.6	1.7	1.6	1.7	0.0
1:27 PM	SINEMET (200mg)					
1:59 PM	1.0	0.8	1.0	0.9	1.0	0.0
2:31 PM	0.3	0.7	0.3	0.8	0.9	0.0
2:59 PM	0.2	0.5	0.2	0.5	0.9	0.0
3:28 PM	0.0	0.3	0.2	0.8	0.9	0.0
4:01 PM	0.5	0.8	0.9	1.6	1.7	0.0
4:29 PM	1.3	1.7	1.6	2.1	2.1	0.0
5:00 PM	SINEMET (200mg)					
5:14 PM	1.0	1.5	1.0	0.9	1.0	0.0
5:28 PM	0.3	0.6	0.3	0.8	2.4	0.0
6:01 PM	0.2	0.3	0.2	0.5	2.0	0.0
6:29 PM	0.0	0.0	0.2	0.8	1.7	0.0
6:59 PM	0.5	0.2	0.9	1.6	1.2	0.0
7:32 PM	1.3	0.9	1.6	2.1	1.0	0.0
8:03 PM	SINEMET (200mg)					
8:29 PM	0.8	0.6	0.5	0.7	0.5	0.0
9:01 PM	0.0	0.2	0.2	1.1	0.9	0.0
9:32 PM	0.0	0.1	0.9	1.6	1.3	0.0
9:55 PM	0.5	0.6	1.9	2.0	1.9	0.0
Mean	0.7	0.8	0.8	1.0	1.5	0.0
Fluctuation	0.7	0.7	0.7	0.7	0.5	0.0

Lazzaro di Biase, MD, PhD

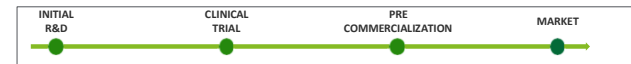
[l.dibiase@policlinicocampus.it](mailto:l.dibiase@policlinicocampus.it)

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DEVELOPMENT STAGE



CERTIFICATIONS



DIAGNOSIS	SYMPTOMS MONITORING	THERAPY MANAGEMENT
✗	<p>Bradykinesia</p> <p>Motor diary</p> <p>Sleep</p> <p>Dyskinesias</p> <p>ON/OFF</p> <p>Tremor</p> <p>Gait</p>	<p>Oral</p> <p>Drug diary</p> <p>Drug reminder</p> <p>Infusional</p> <p>DBS</p>

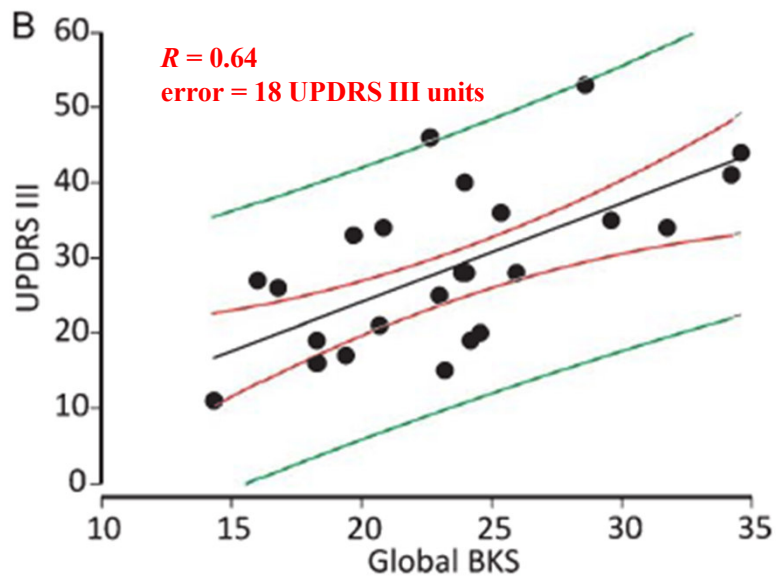
## PKG® Remote Monitoring Process



### Bradykinesia score (BKS):

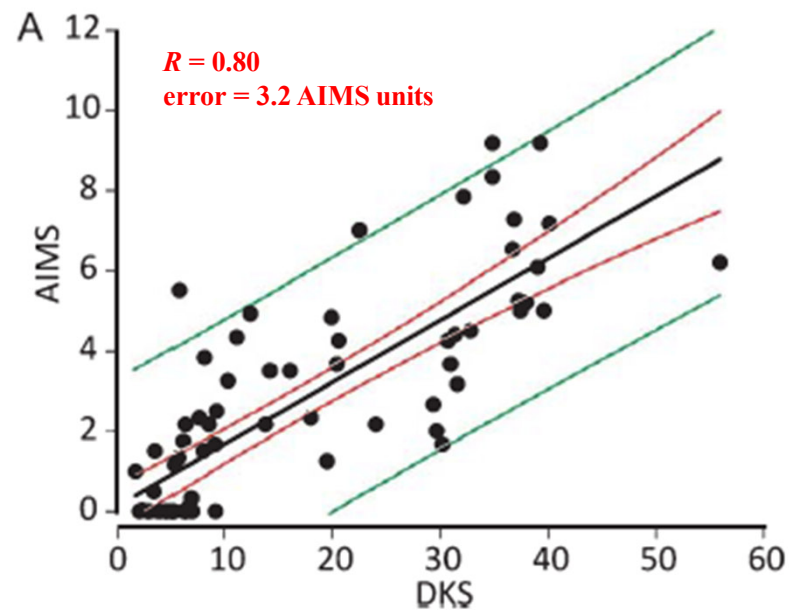
- **Lower** Mean Spectral Power (MSP) within 0.2 and 4 Hz,
- **Lower** Peak acceleration
- **Longer** Time without movement.

Bradykinetic subjects move with lower accelerations, amplitude, energy, and with longer intervals between movements than normal subjects.



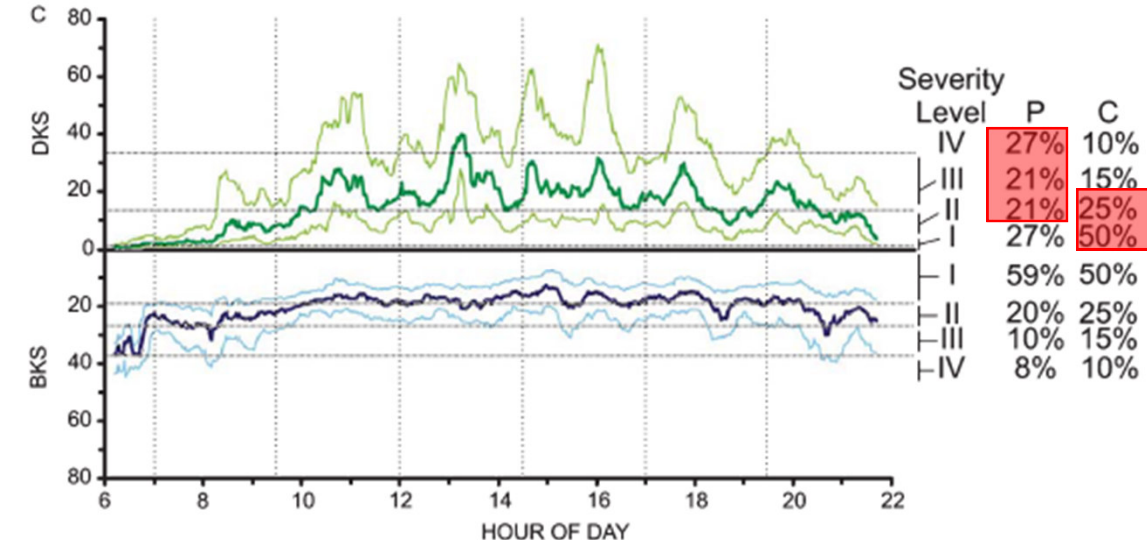
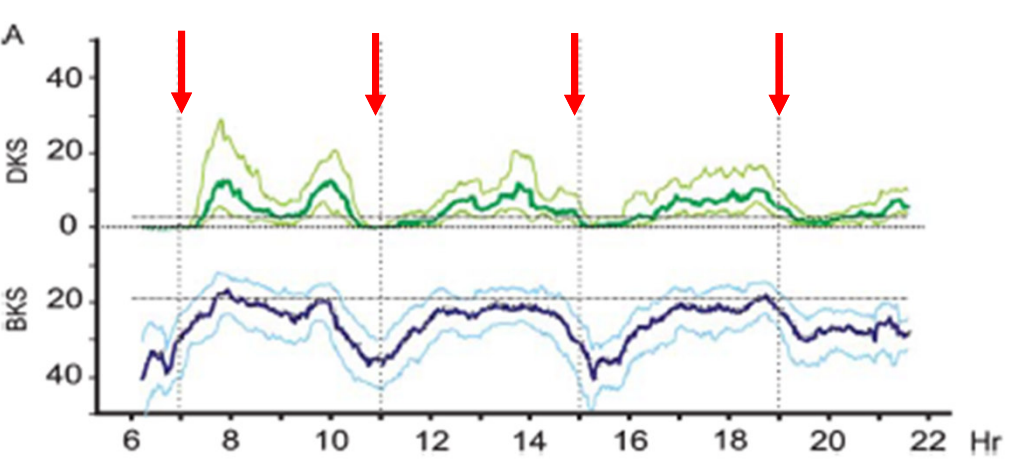
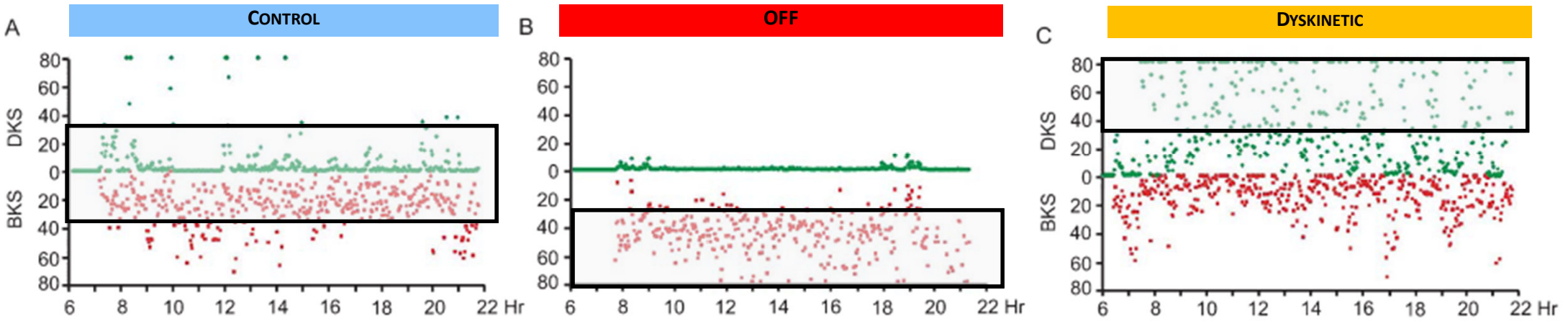
### Dyskinesia score (DKS):

- **Larger** Mean Spectral Power (MSP)
- **Greater** Acceleration than the mean.
- **Shorter** Time without movement.



Griffiths RI, et al. J Parkinsons Dis. 2012;2(1):47-55. doi: 10.3233/JPD-2012-11071. PMID: 23939408.

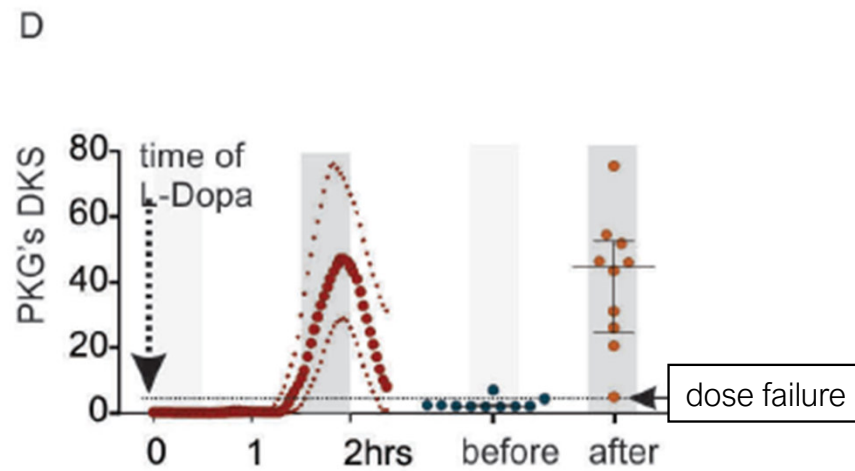
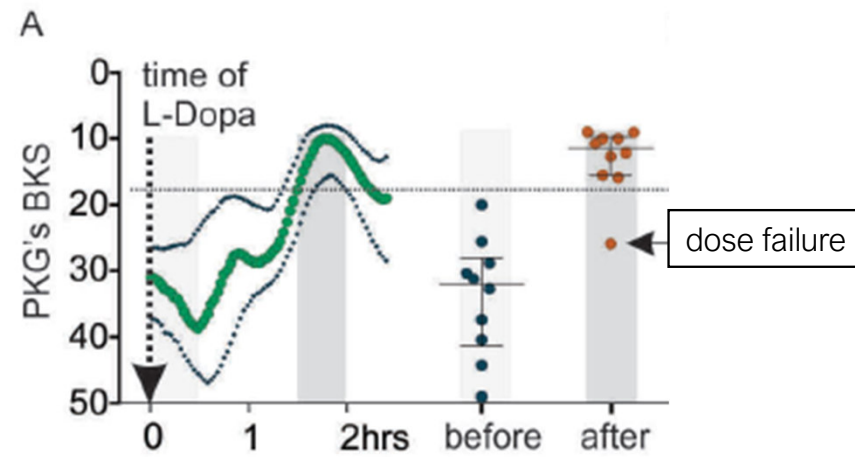
# BRADYKINESIA AND DYSKINESIA



Griffiths RI, et al. J Parkinsons Dis. 2012;2(1):47-55. doi: 10.3233/JPD-2012-11071. PMID: 23939408.

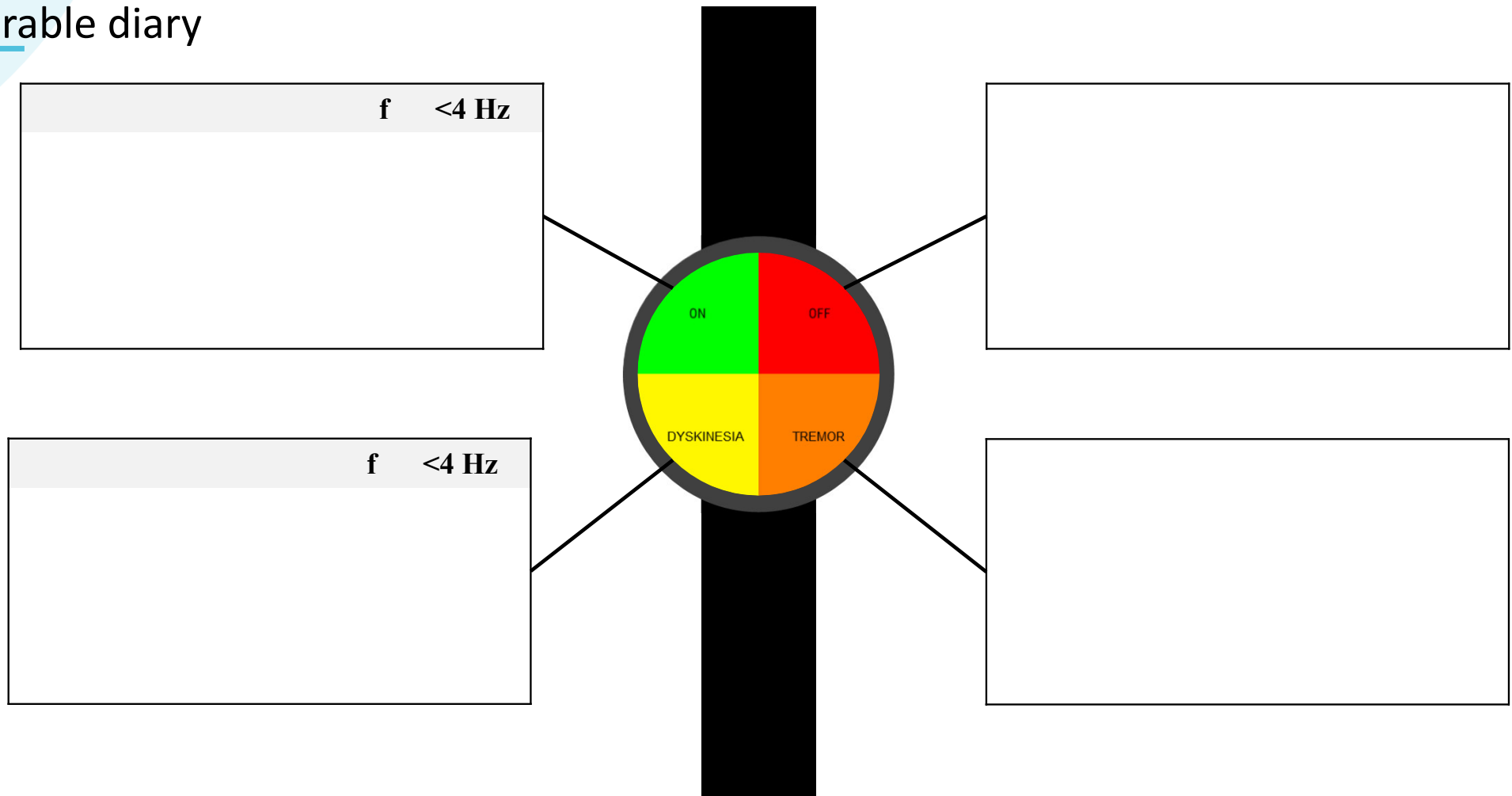
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# BRADYKINESIA AND DYSKINESIA

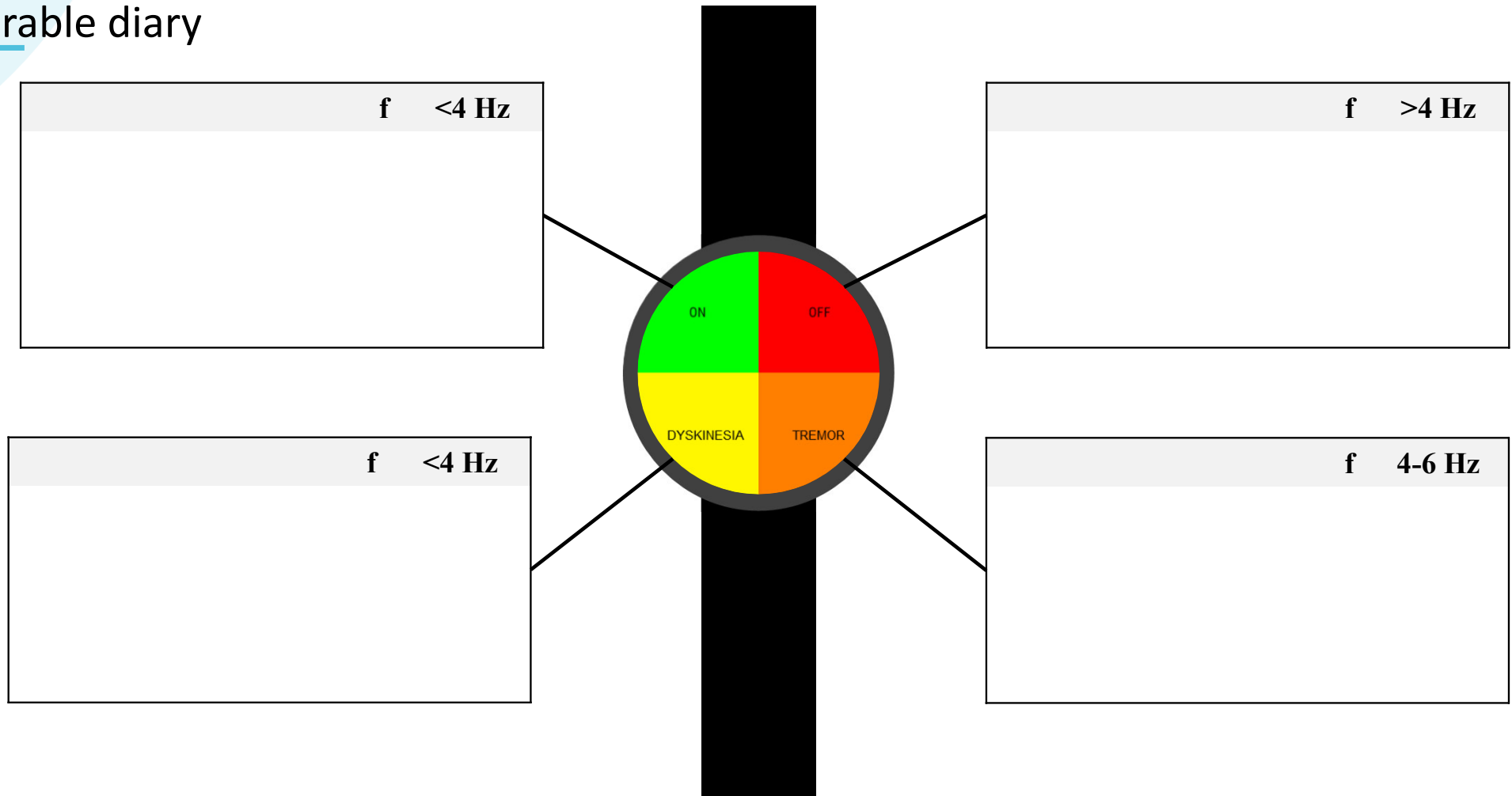


Horne, M., et al. (2016). In *CNS* (Vol. 2, No. 1, pp. 15-22).

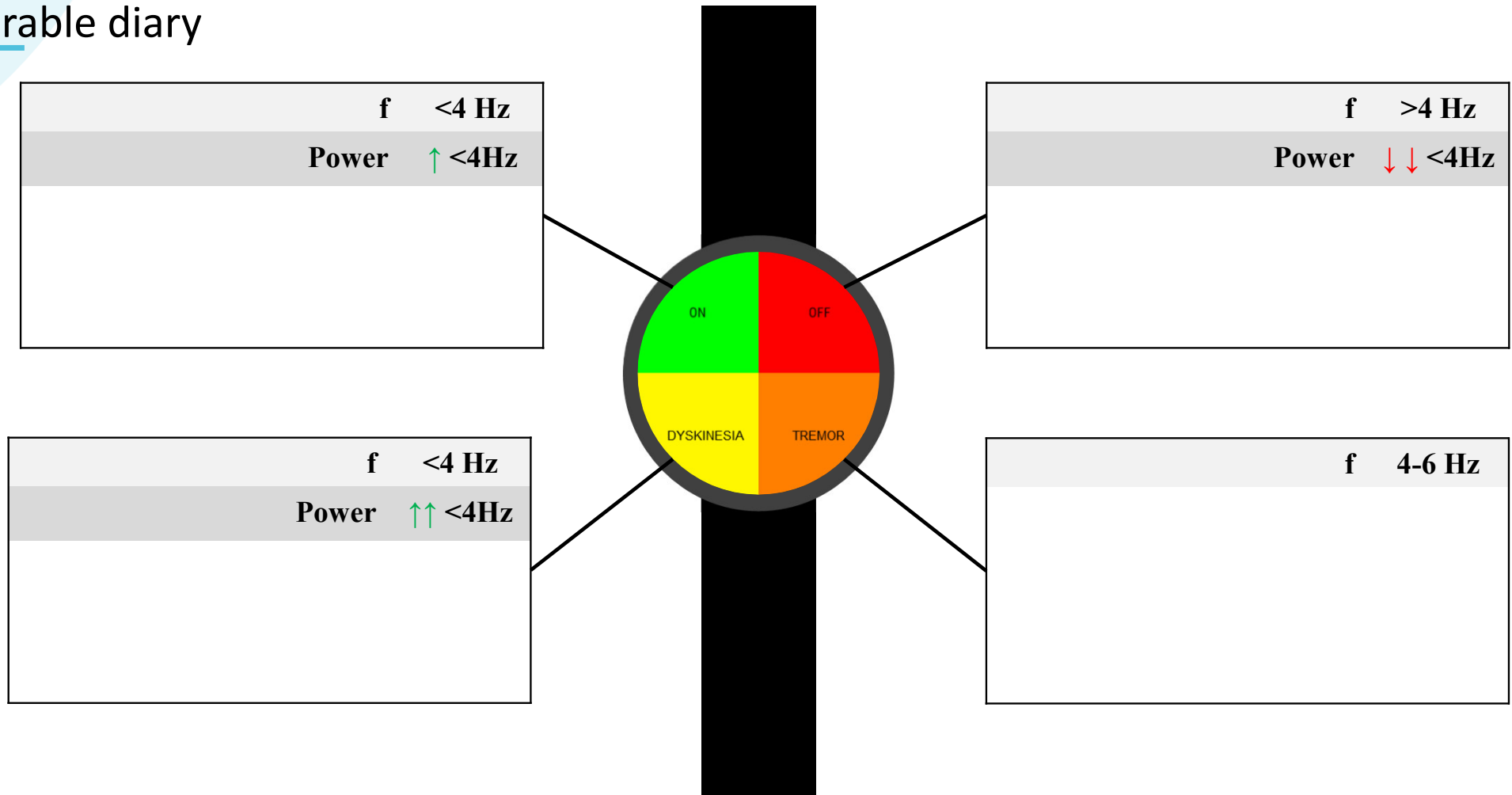
## Wearable diary



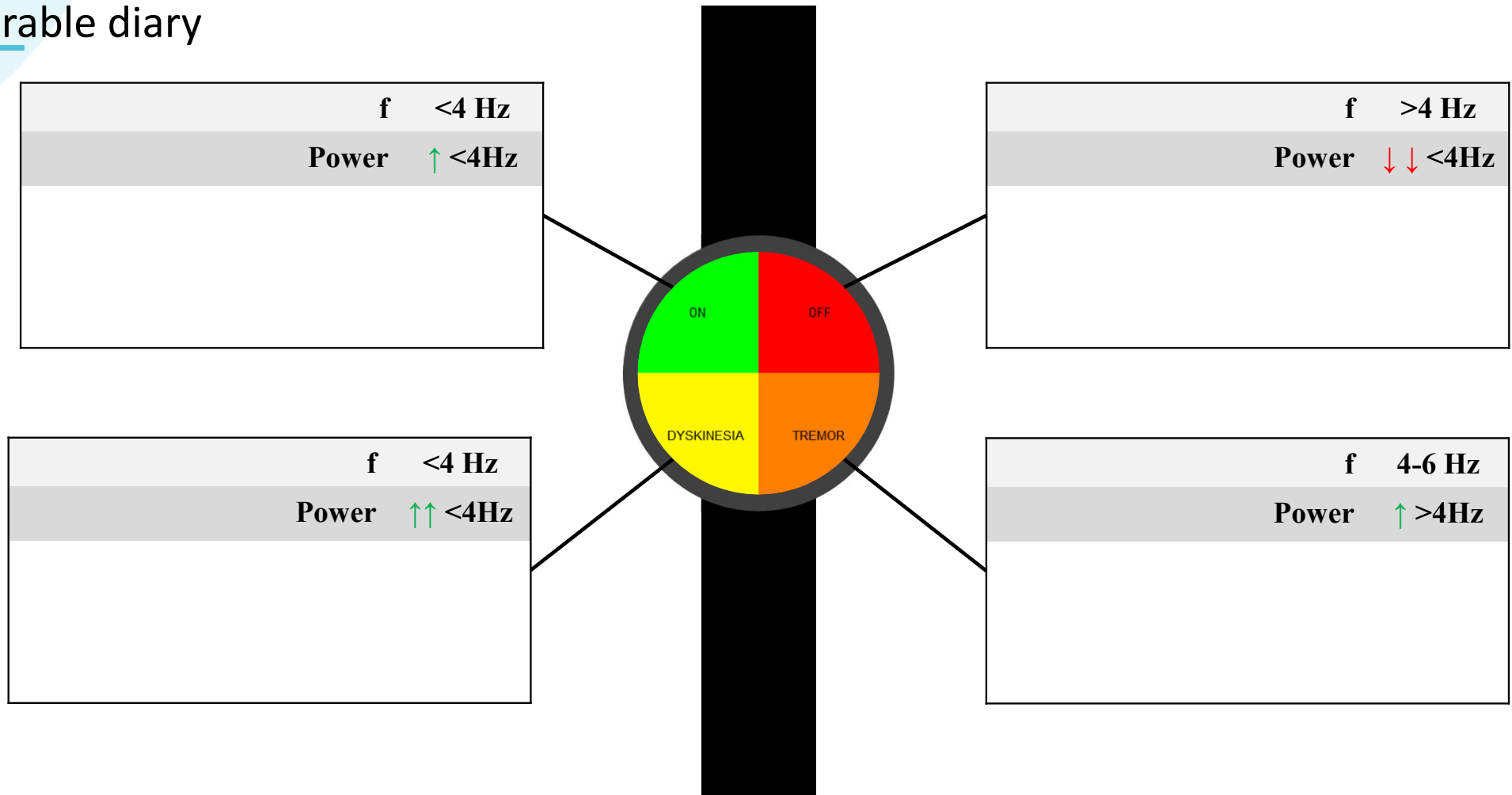
## Wearable diary



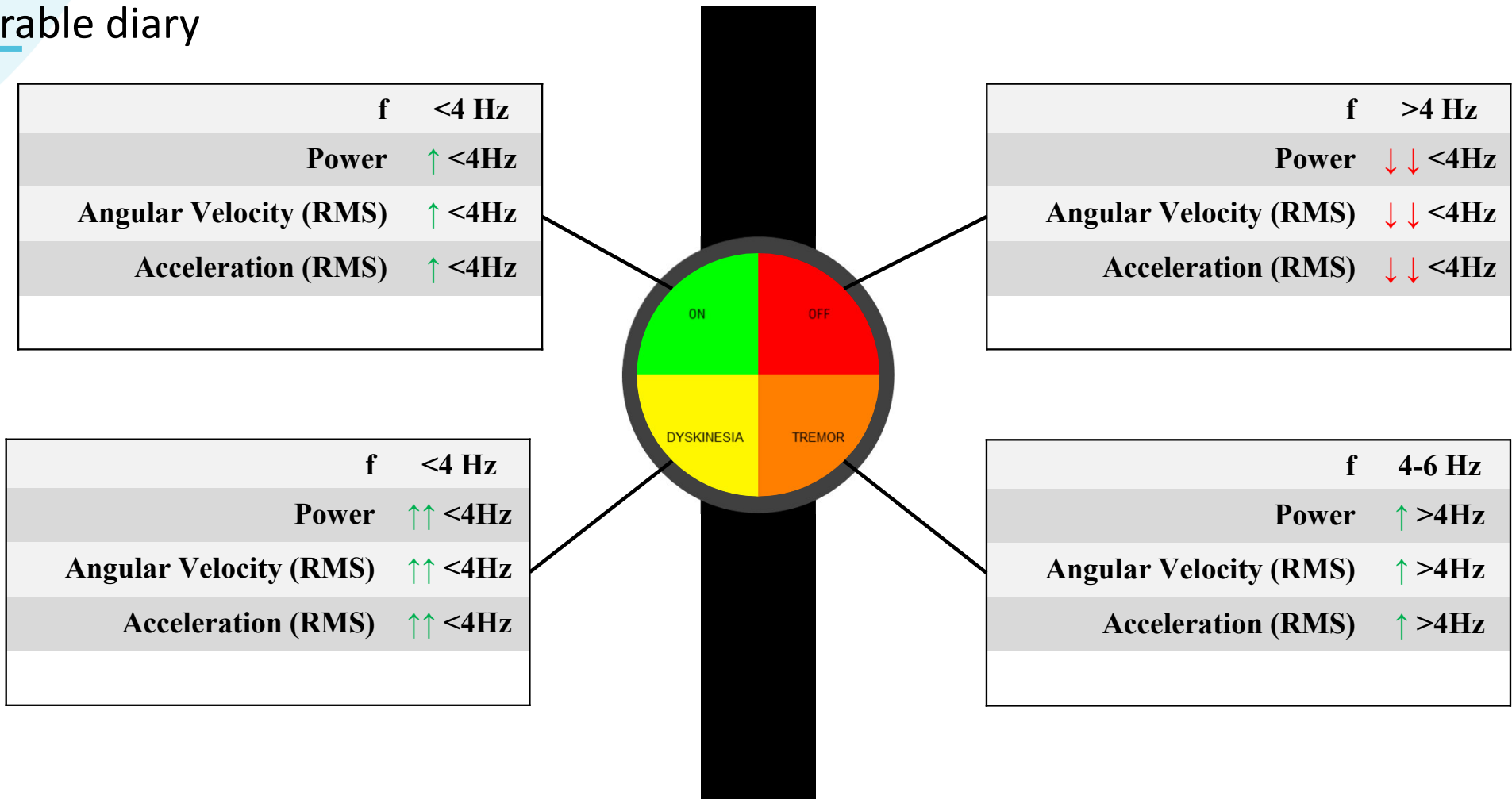
## Wearable diary



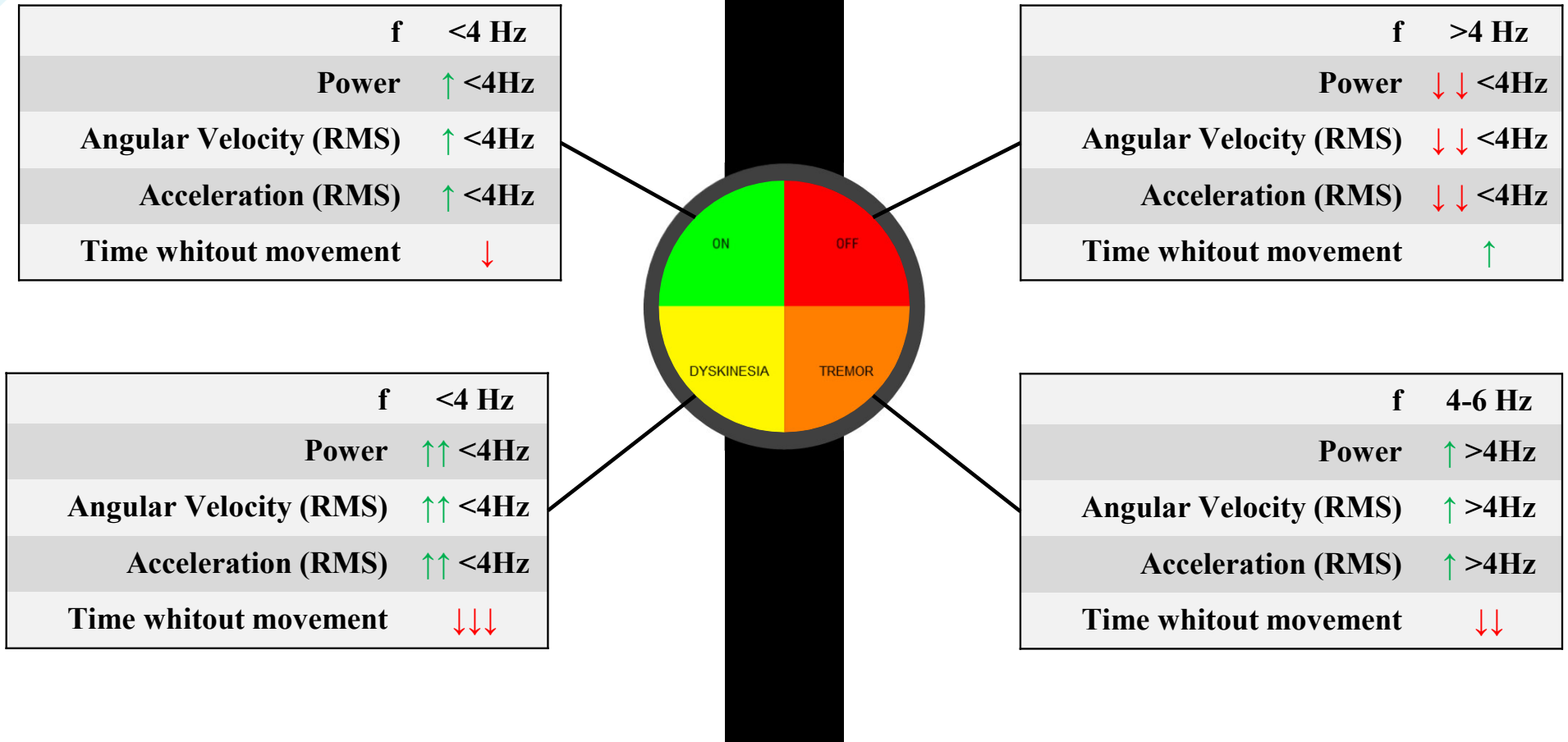
## Wearable diary



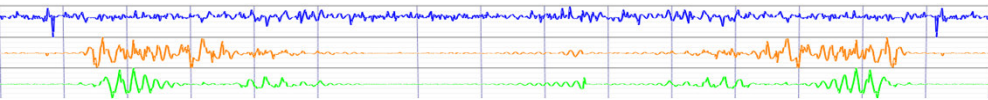
# Wearable diary



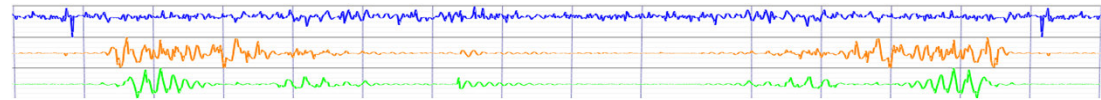
# Wearable diary

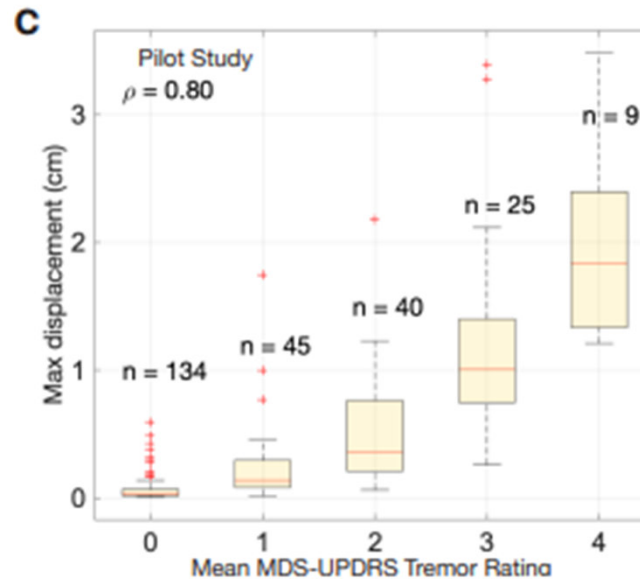
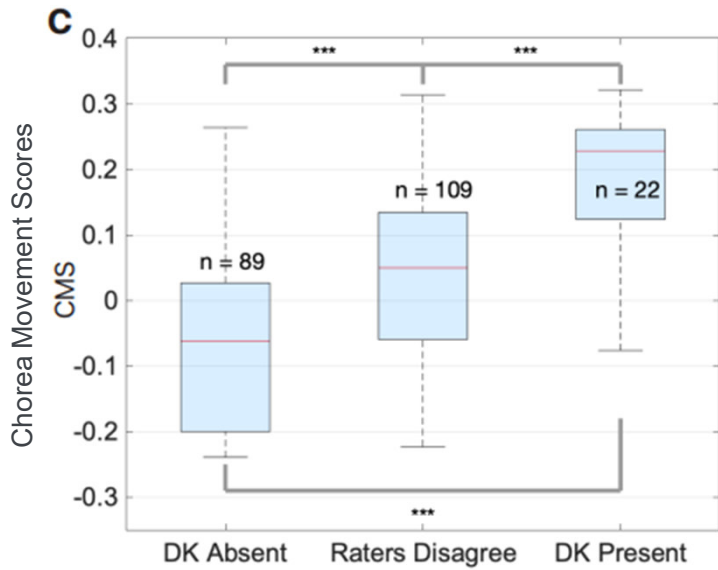


# Bradykinesia



# Dyskinesia





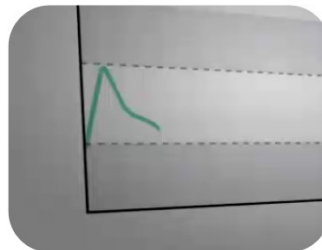
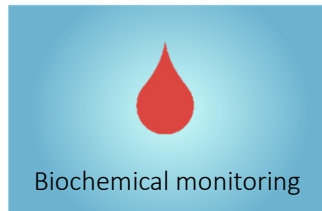
Activity	% Tremor	% Chorea
Walking	2%	1%
Brushing Teeth (Manual)	8%	4%
Riding Bus	2%	13%
Knitting	0.3%	14%
Cycling	0.4%	31%
Musical Instrument	2%	61%









Powers R, et al. Smartwatch inertial sensors continuously monitor real-world motor fluctuations in Parkinson's disease. *Sci Transl Med.* 2021. doi: 10.1126/scitranslmed.abd7865.

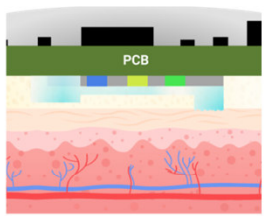
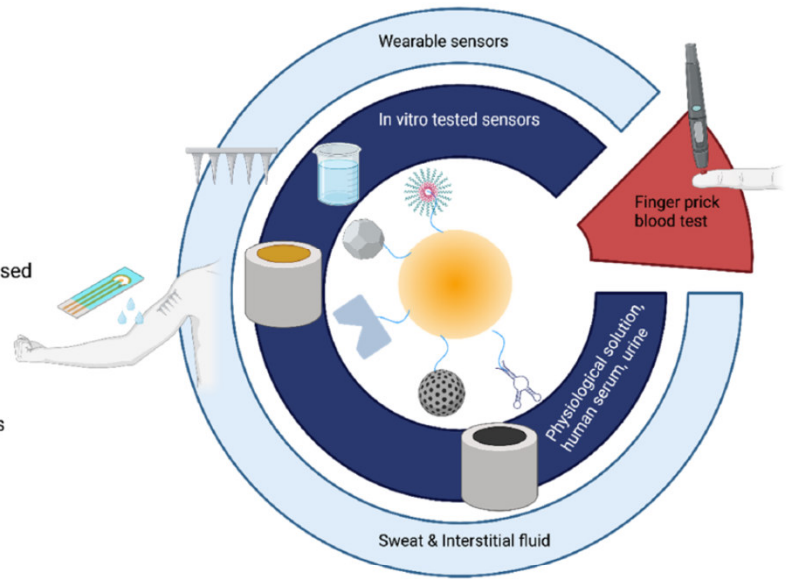
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# Biochemical monitoring

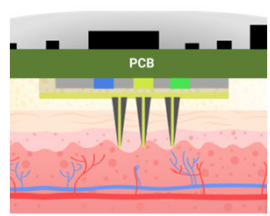
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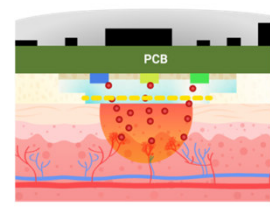
-  Aptamer-based
  -  Polymer-based
  -  Metal-based
  -  Enzyme-based
  -  Carbon Nanomaterial-based
- 
-  Wearable sensors
  -  In vitro sensors
  -  Point of Care



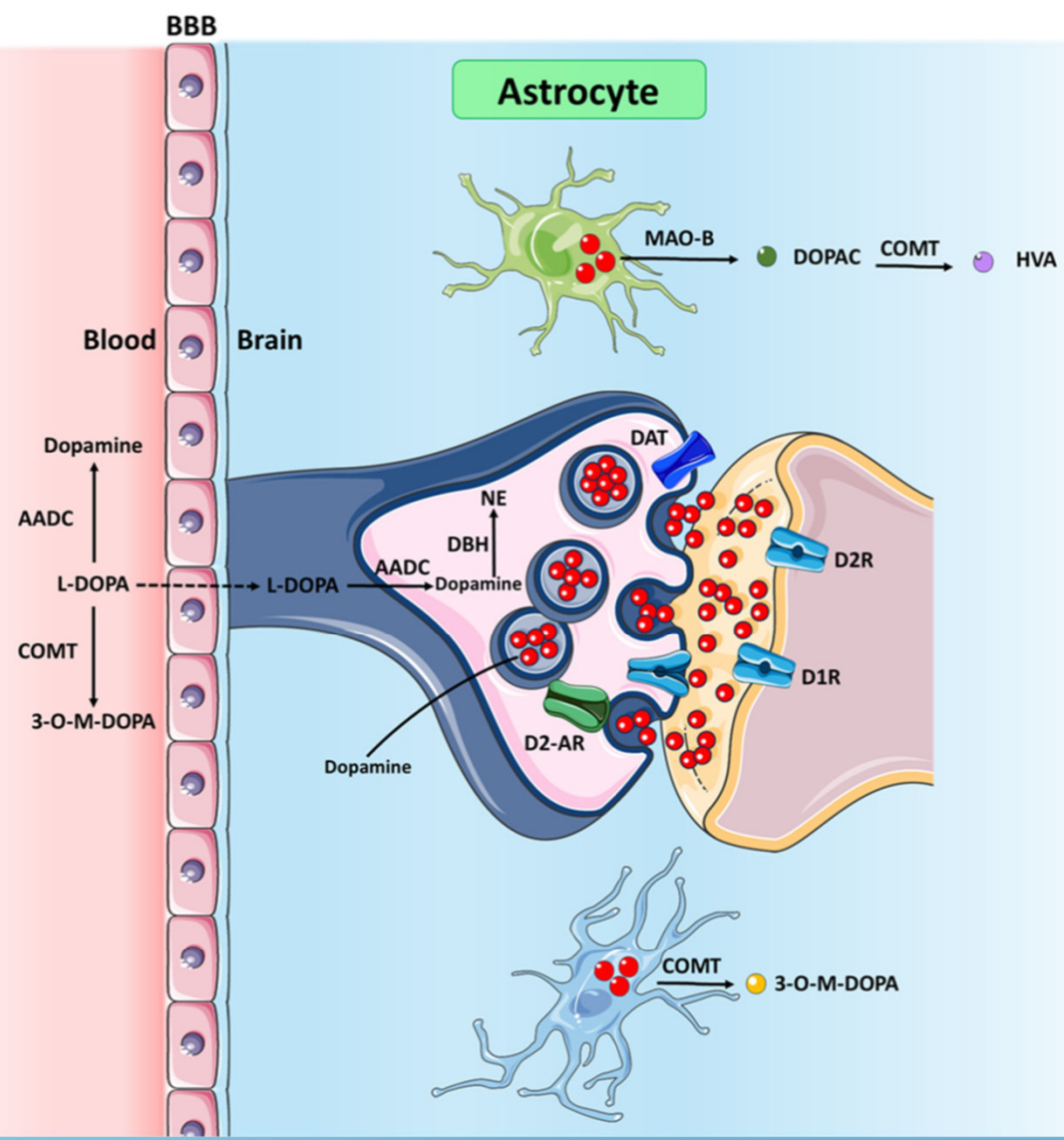
Sweat



ISF

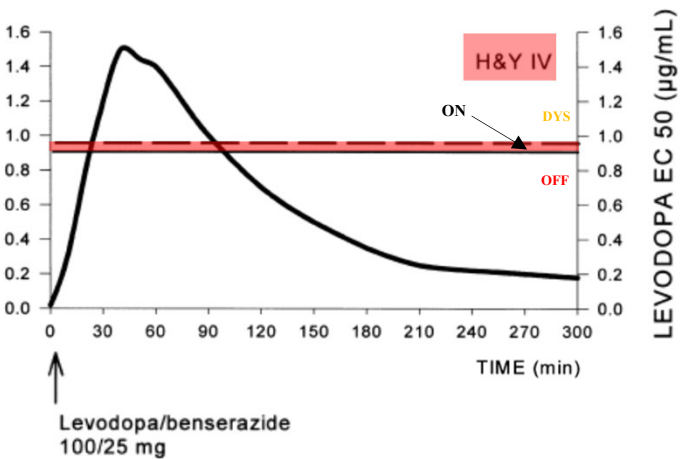
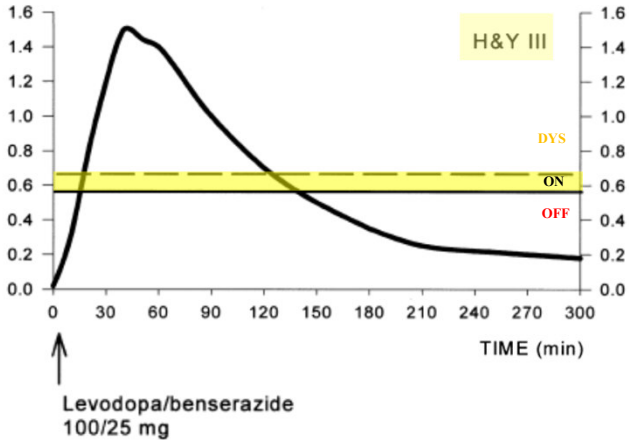
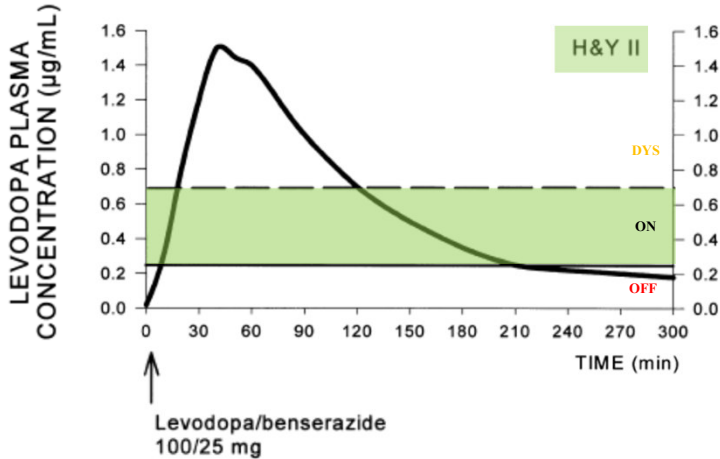


Blood



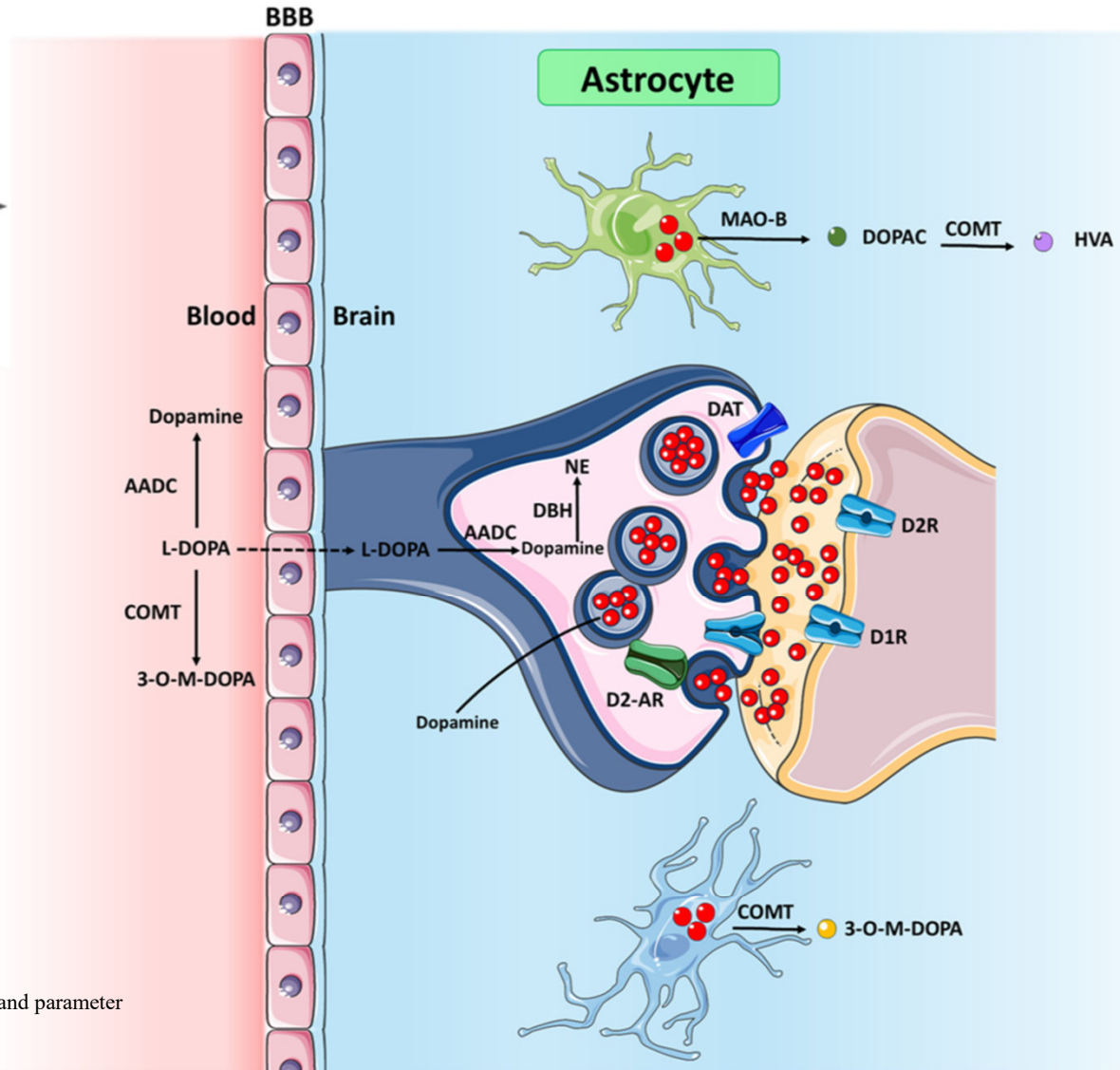
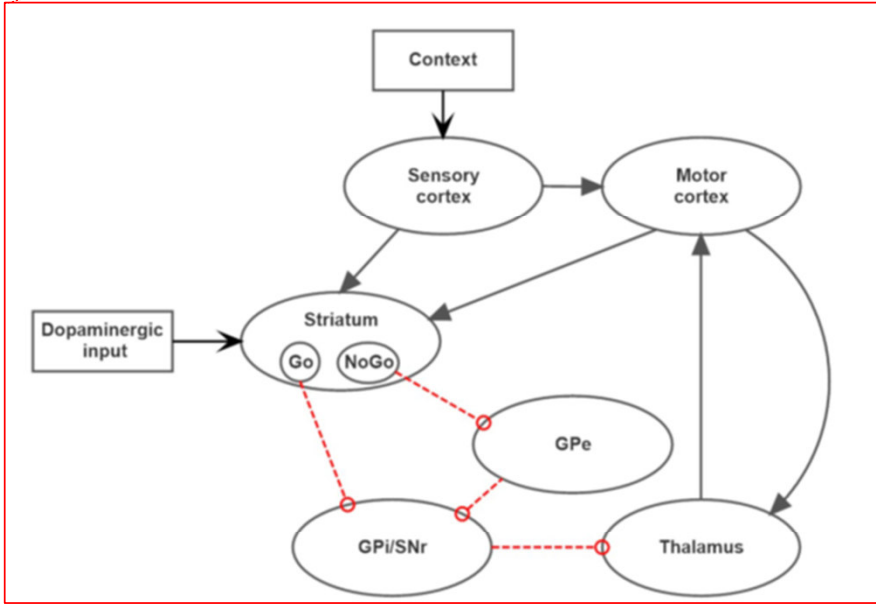
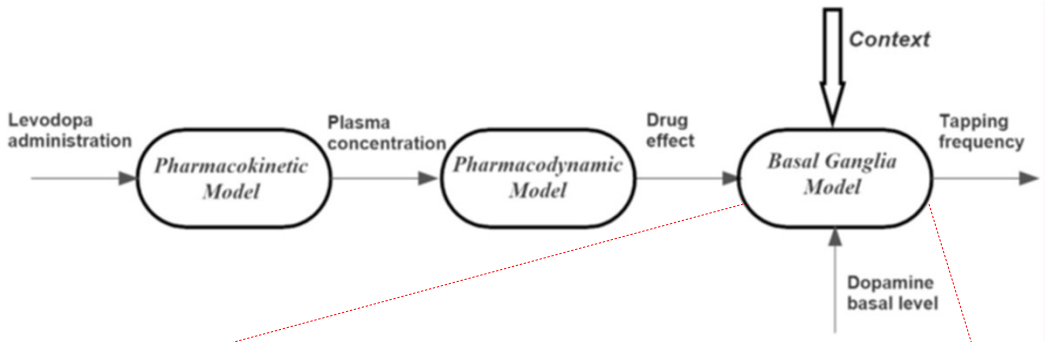
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# Pharmacokinetics and Pharmacodynamics



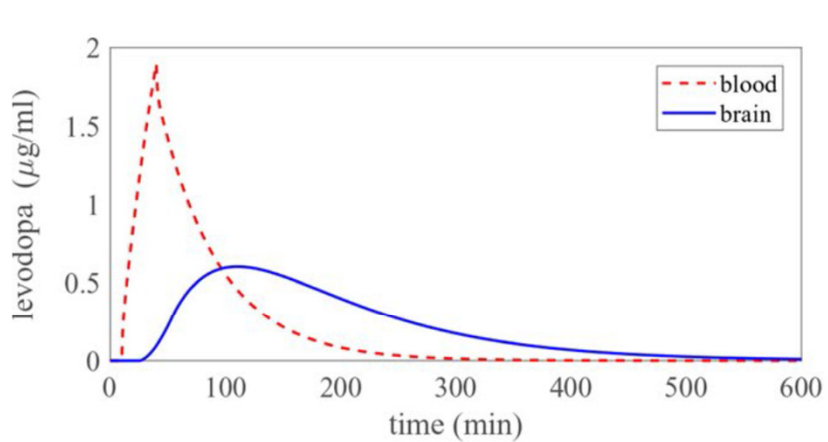
Contin M, et al. Ther Drug Monit. 2001 Dec;23(6):621-9. doi: 10.1097/00007691-200112000-00005. PMID: 11802094.

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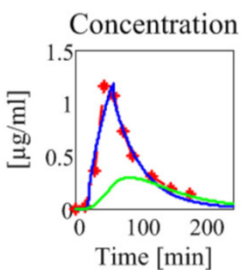


Ursino M, Magosso E, Lopane G, Calandra-Buonaura G, Cortelli P, Contin M (2020) Mathematical modeling and parameter estimation of levodopa motor response in patients with parkinson disease. PLoS ONE 15(3): e0229729 <https://doi.org/10.1371/journal.pone.0229729>

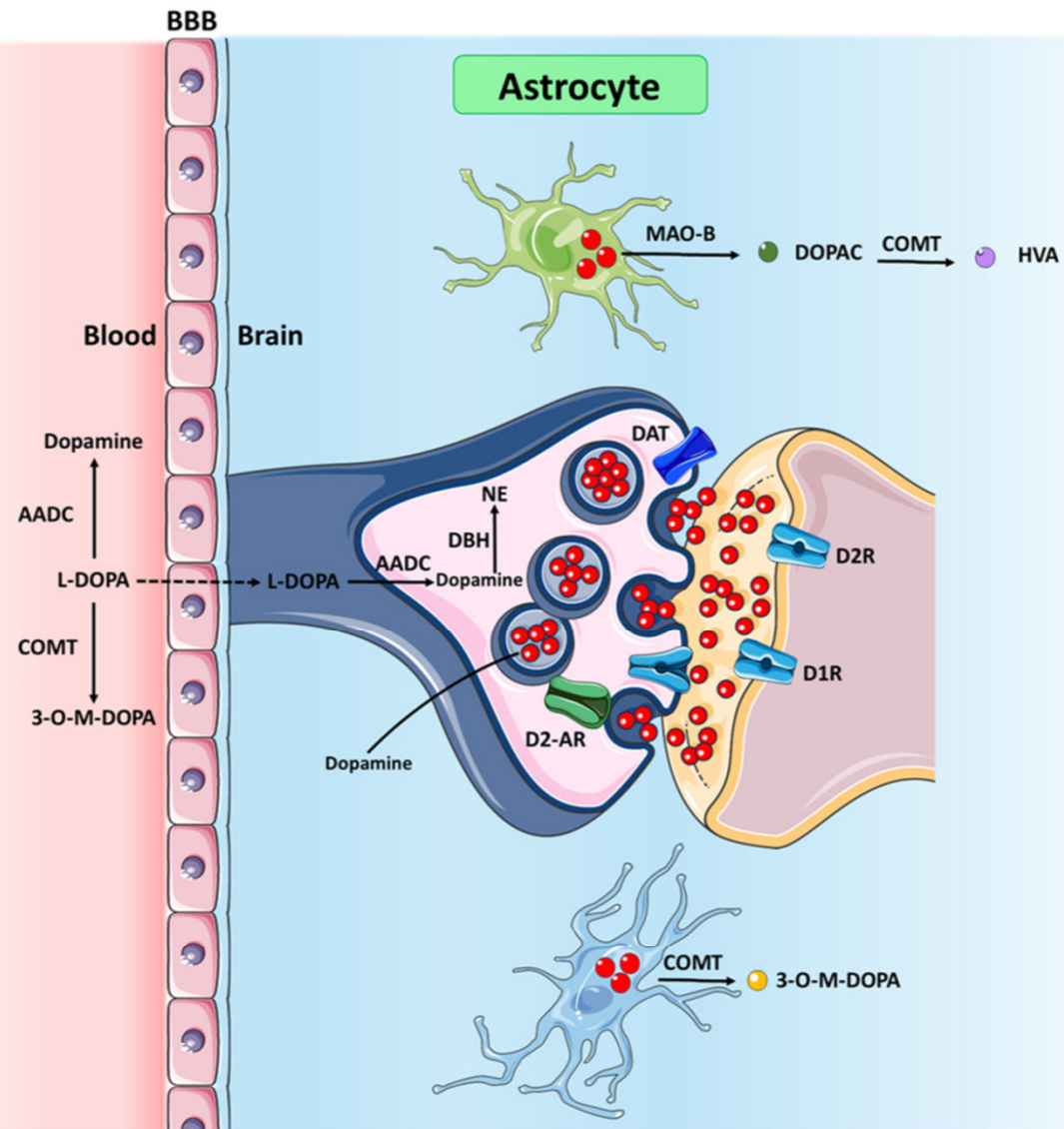
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Ursino M, Baston C. Aberrant learning in Parkinson's disease: A neurocomputational study on bradykinesia. *Eur J Neurosci.* 2018;47:1563– 1582. <https://doi.org/10.1111/ejn.13960>



Baston, C., et al. (2016). *Frontiers in human neuroscience*, 10, 280.

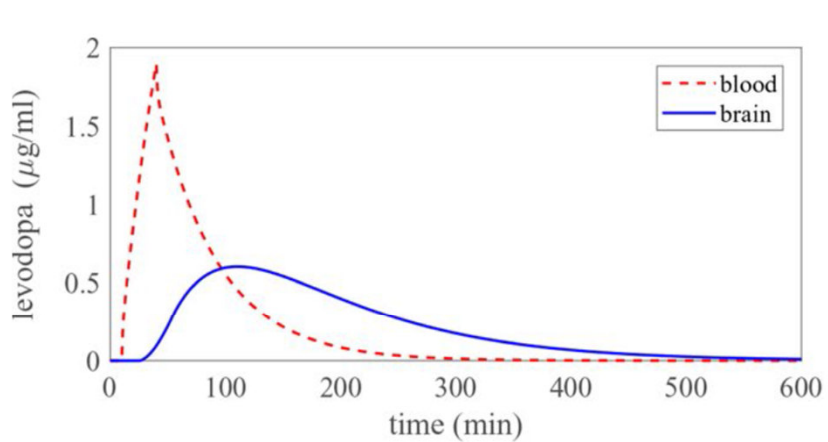


Lazzaro di Biase, MD, PhD

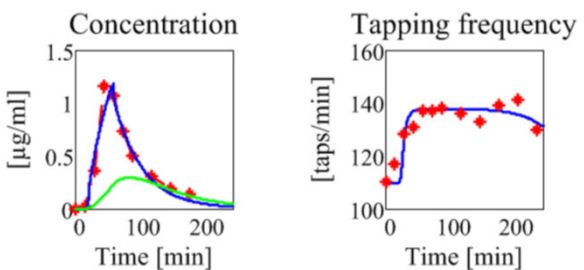
[l.dibiase@policlinicocampus.it](mailto:l.dibiase@policlinicocampus.it)

Neurology Unit, Campus Bio-Medico University Hospital

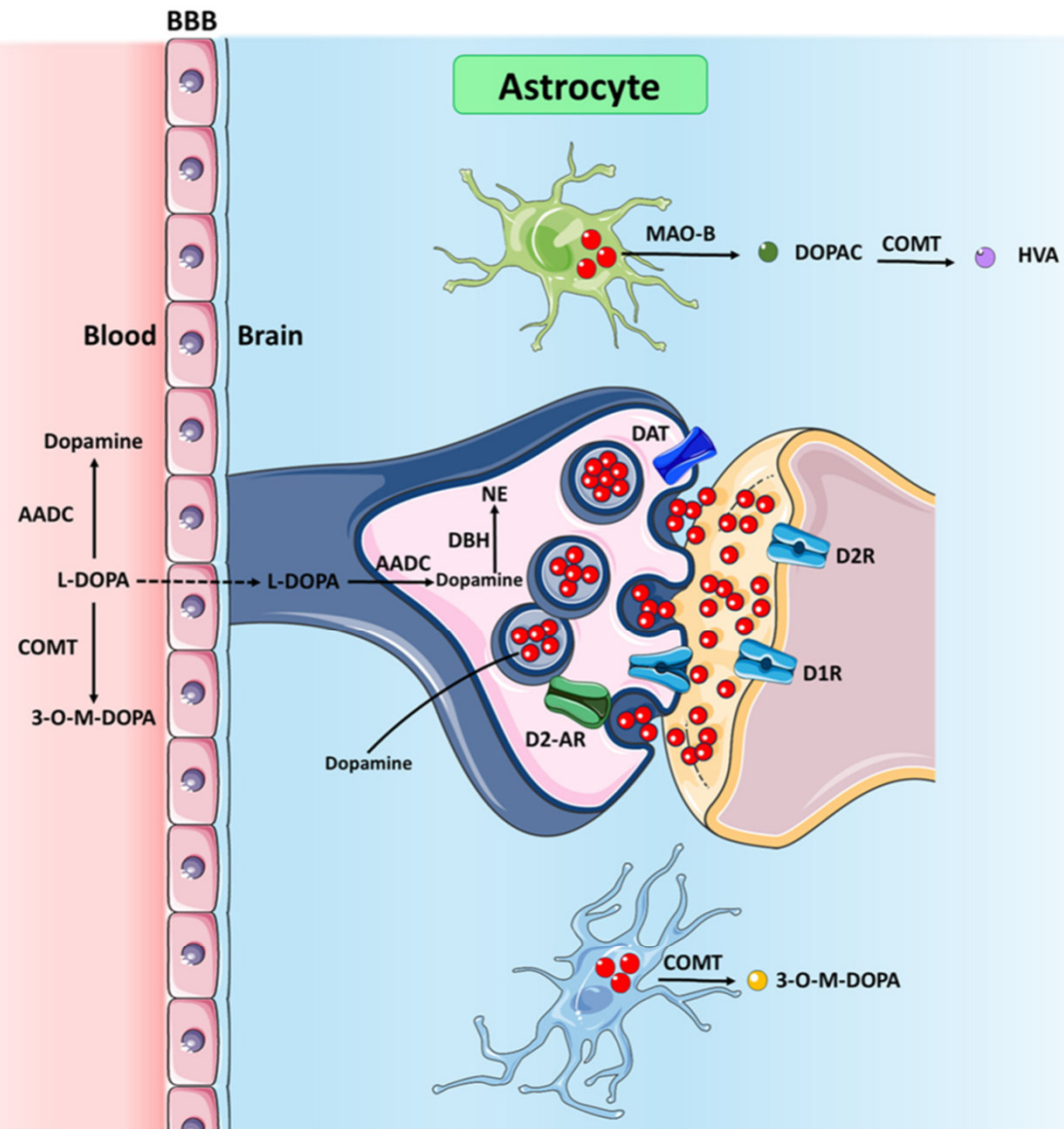
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Ursino M, Baston C. Aberrant learning in Parkinson's disease: A neurocomputational study on bradykinesia. *Eur J Neurosci.* 2018;47:1563– 1582. <https://doi.org/10.1111/ejn.13960>



Baston, C., et al. (2016). *Frontiers in human neuroscience*, 10, 280.

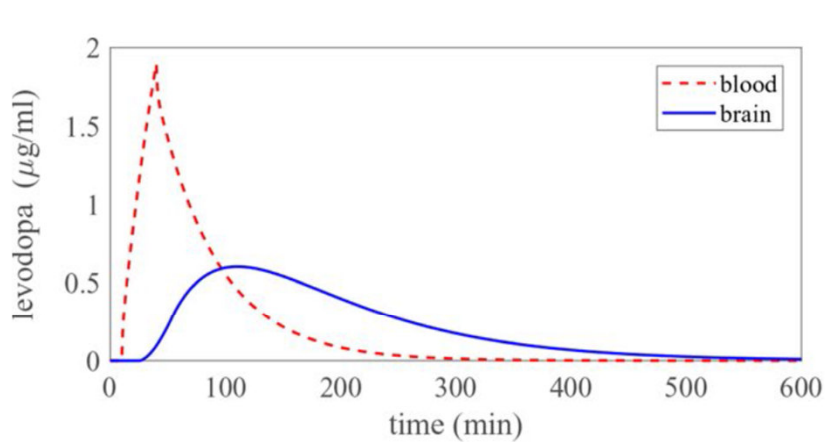


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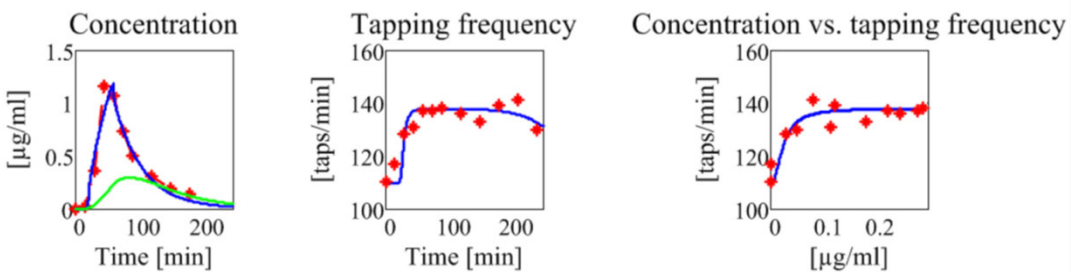
[l.dibiase@policlinicocampus.it](mailto:l.dibiase@policlinicocampus.it)

Neurology Unit, Campus Bio-Medico University Hospital

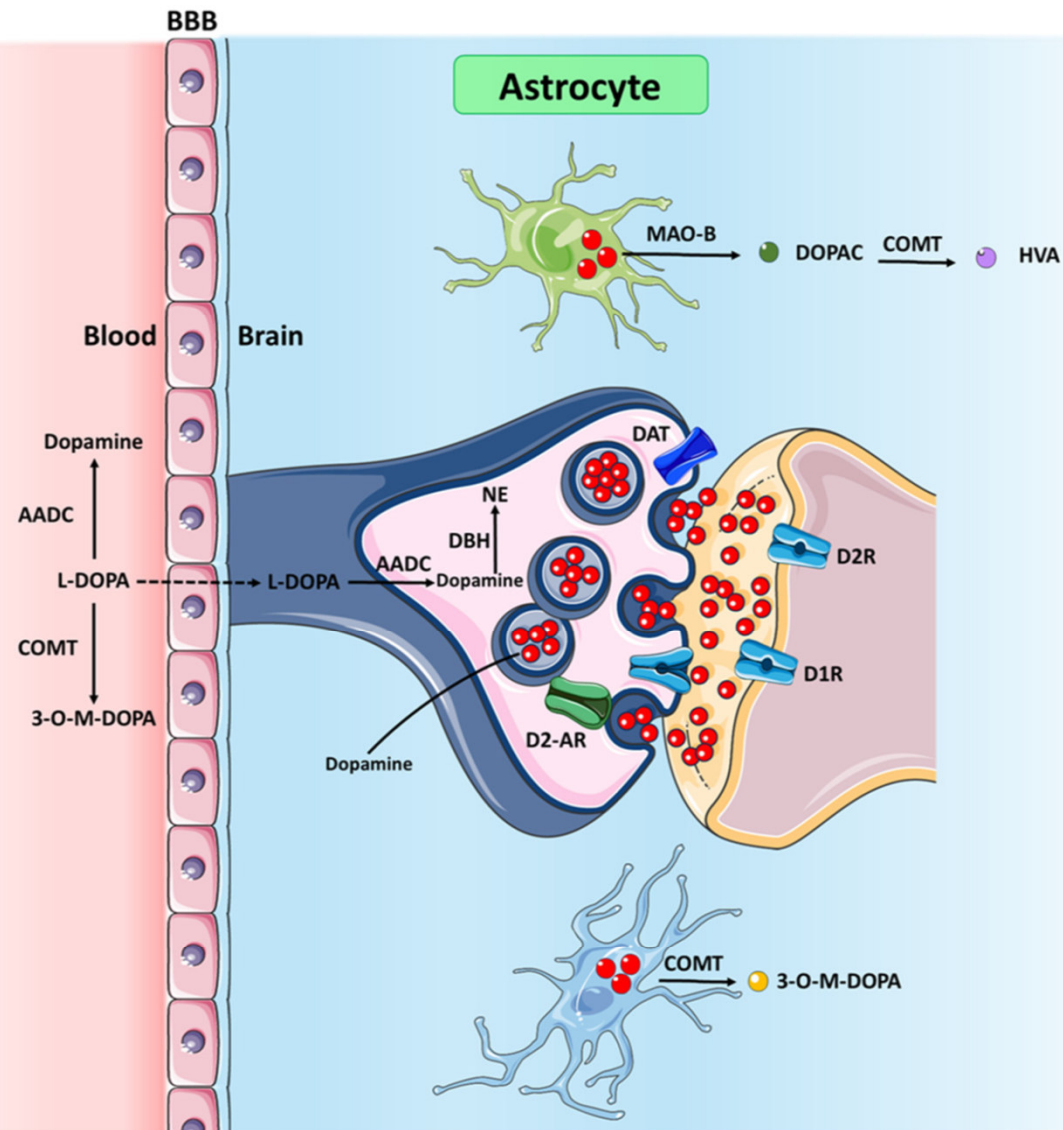
Brain Innovations Lab



Ursino M, Baston C. Aberrant learning in Parkinson's disease: A neurocomputational study on bradykinesia. *Eur J Neurosci.* 2018;47:1563–1582. <https://doi.org/10.1111/ejn.13960>



Baston, C., et al. (2016). *Frontiers in human neuroscience*, 10, 280.



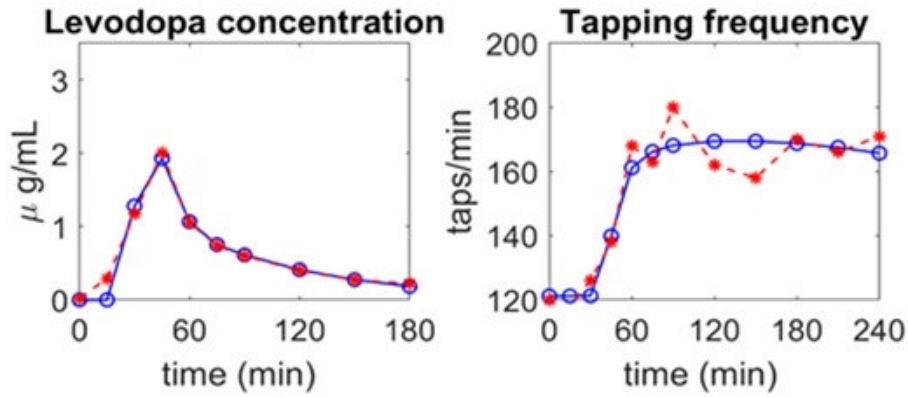
Lazzaro di Biase, MD, PhD

[l.dibiase@policlinicocampus.it](mailto:l.dibiase@policlinicocampus.it)

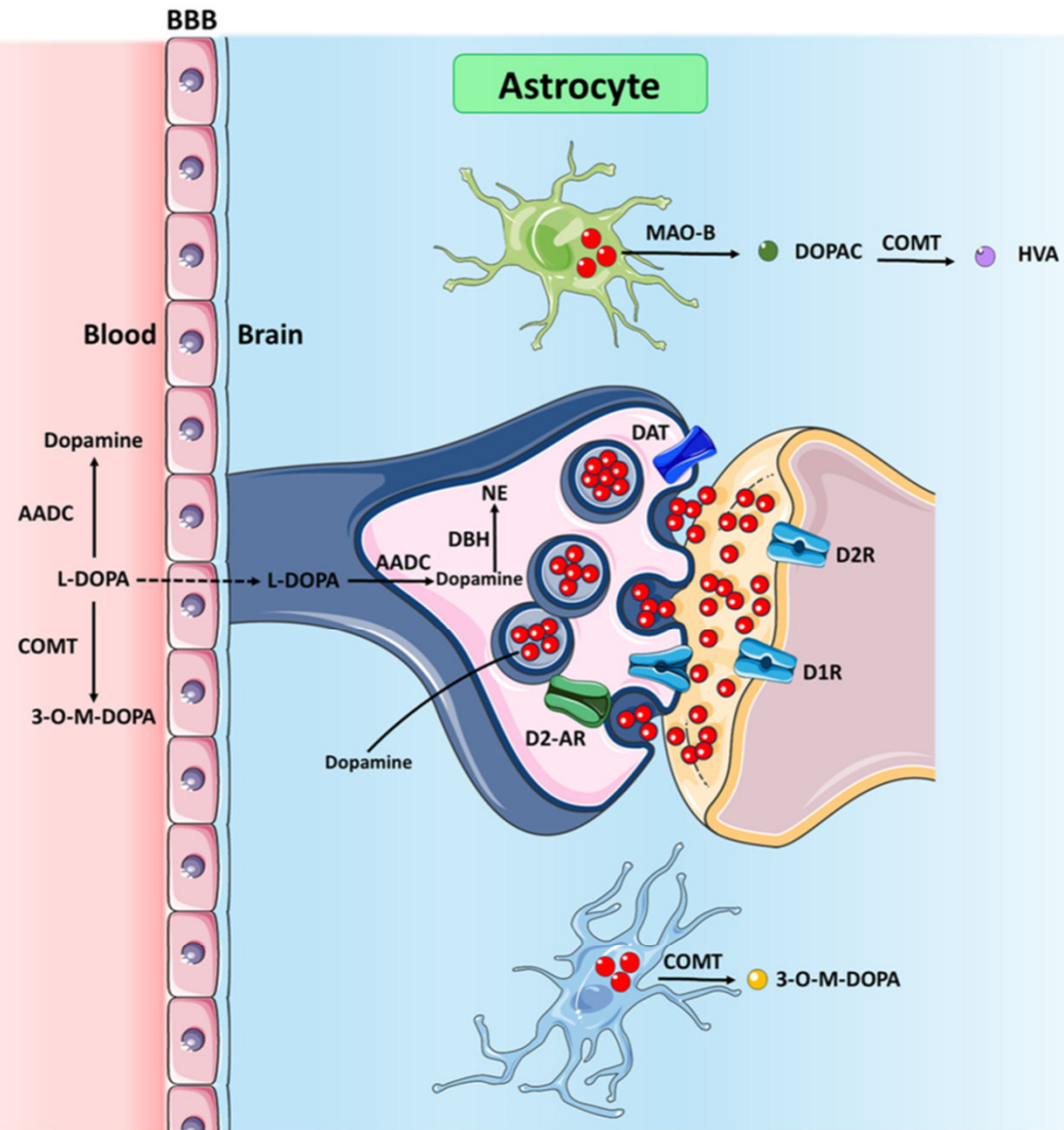
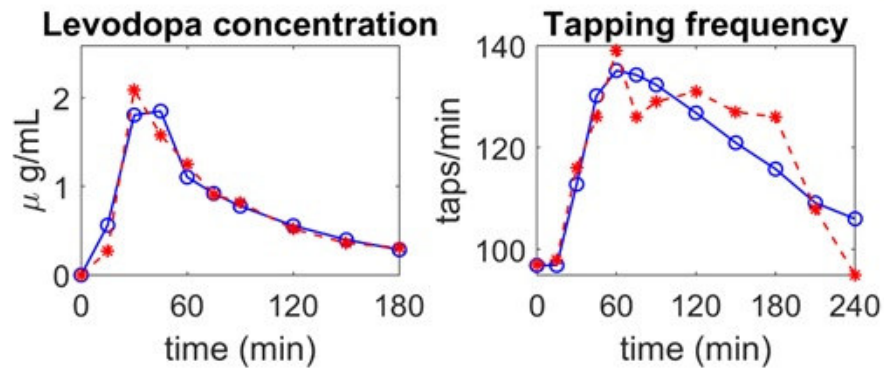
Neurology Unit, Campus Bio-Medico University Hospital

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### STABLE PD



### FLUCTUATING PD

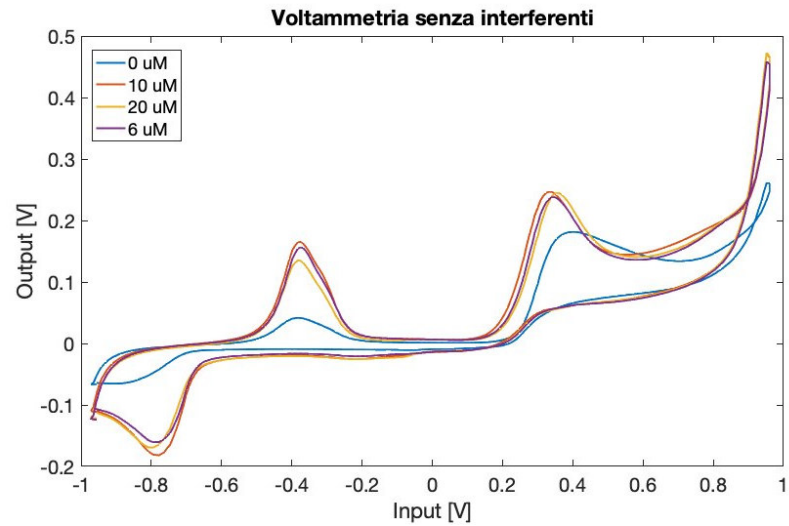


Ursino M, et al. PloS one, 15(3), e0229729 (2020).

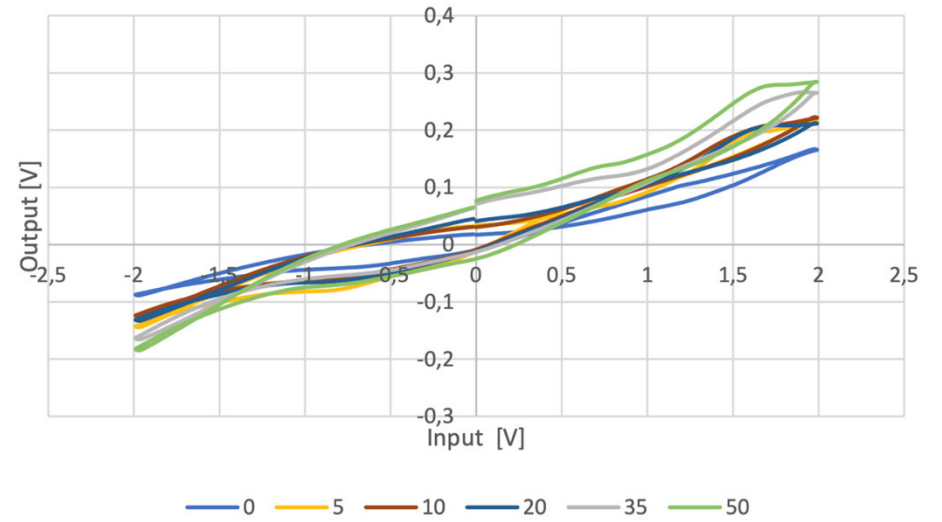
di Biase L., et al. (2021). Expert Review of Neurotherapeutics, DOI: 10.1080/14737175.2021.2000392

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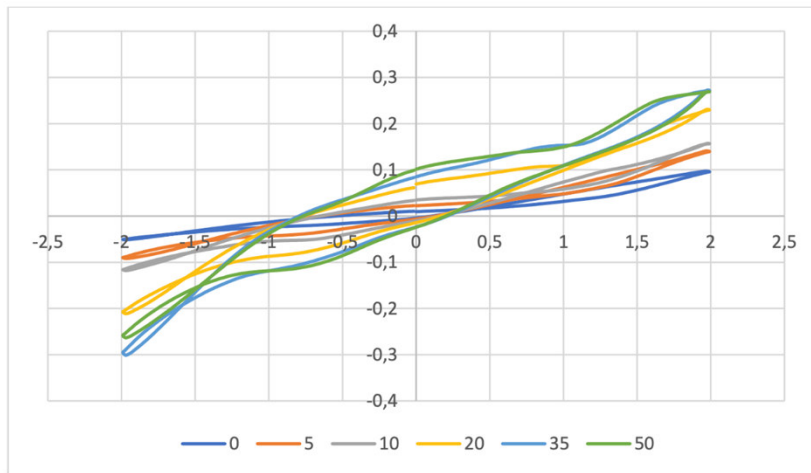
**L-DOPA:** 0,6,10,20  $\mu\text{M}$   
Prediction Model  $\rightarrow$  **6.93  $\mu\text{M}$**



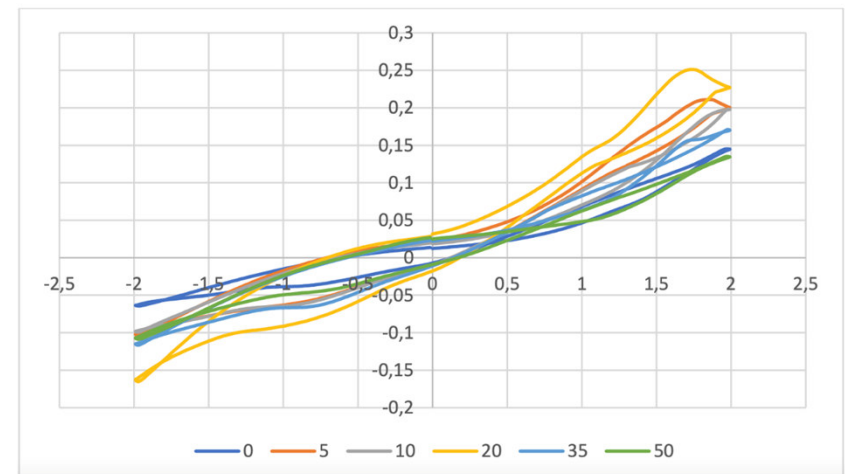
**DOPAC:** 0,5,10,20,35,50  $\mu\text{M}$   
Prediction Model  $\rightarrow$  **1.93  $\mu\text{M}$**



**3-MethoxyTyramine (3MT):** 0,5,10,20,35,50  $\mu\text{M}$   
Prediction Model  $\rightarrow$  **2.84  $\mu\text{M}$**



**3-O-Methyldopa (3-OMD):** 0,5,10,20,35,50  $\mu\text{M}$   
Prediction Model  $\rightarrow$  **5.37  $\mu\text{M}$**

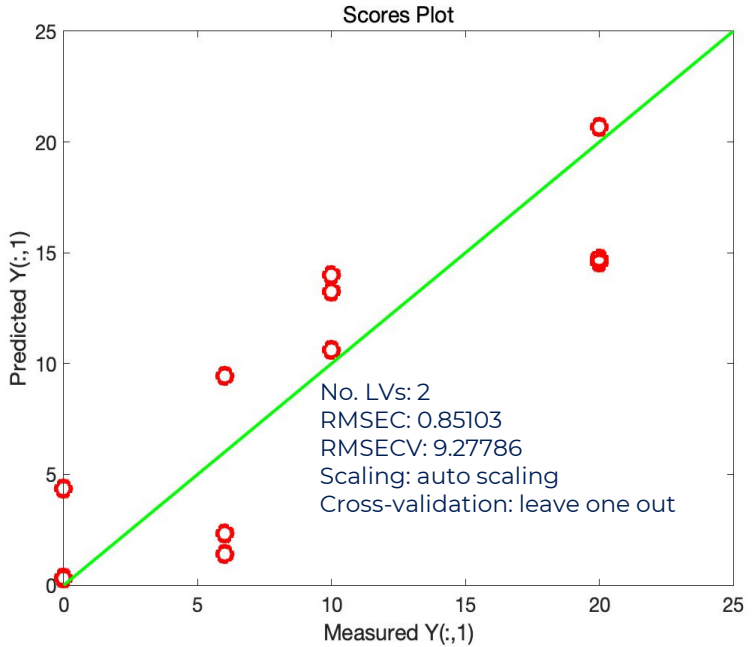
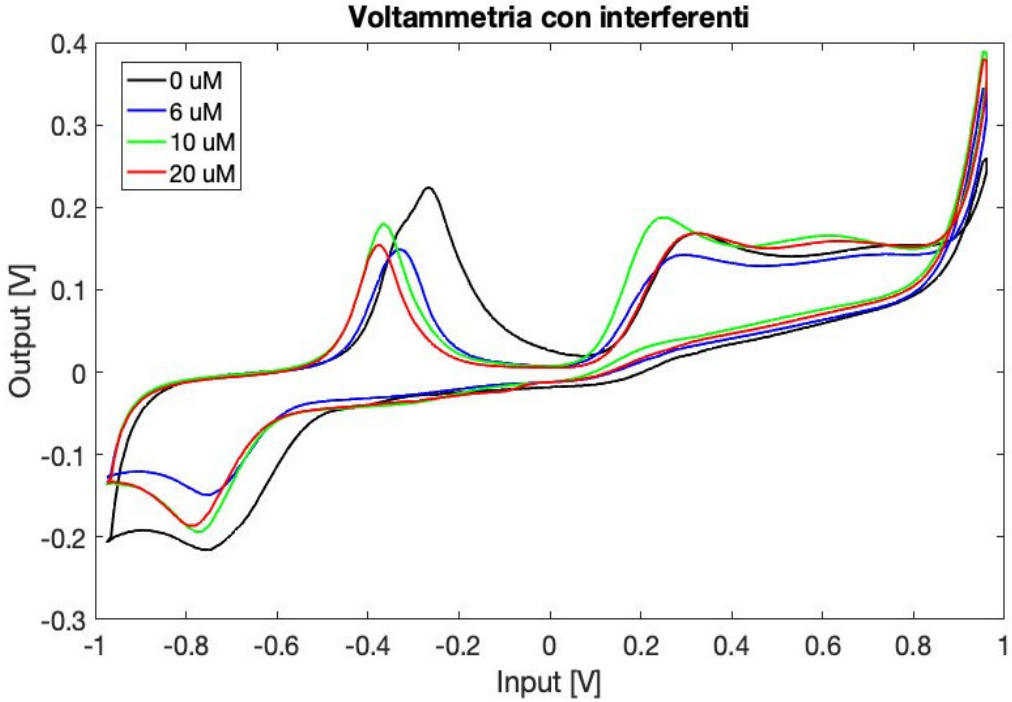


**L-DOPA:** 0,6,10,20  $\mu\text{M}$

with the presence of **interferents** :

- **Lactic acid (20 mM)**
- **Ascorbic acid (50  $\mu\text{M}$ )**
- **Urea (10 mM).**

Prediction Model  $\rightarrow$  **9.27  $\mu\text{M}$**



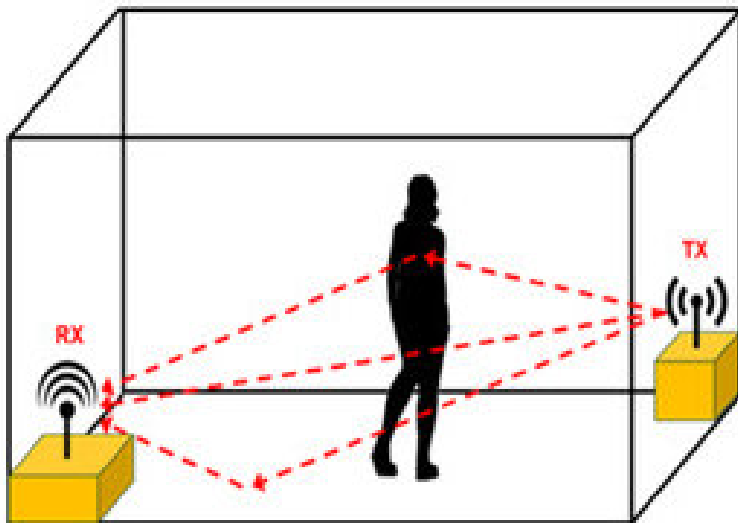
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# IR MONITORING

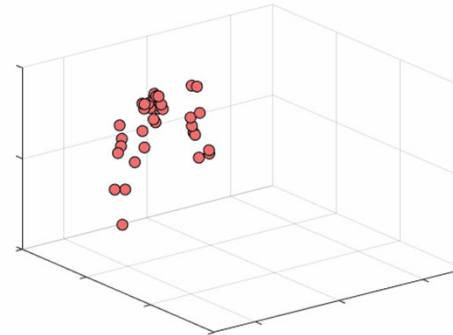
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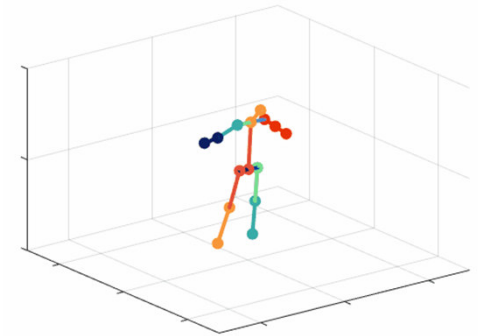
Radar



Radar Point Cloud:



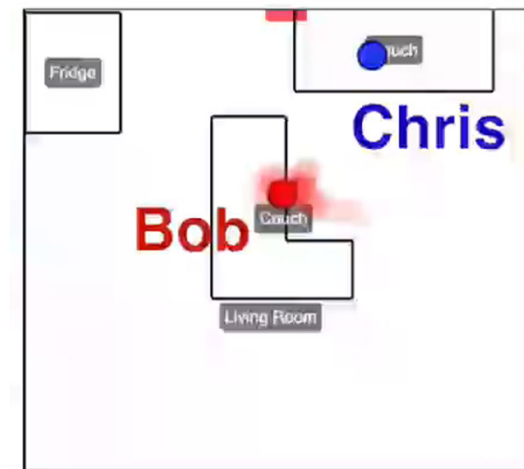
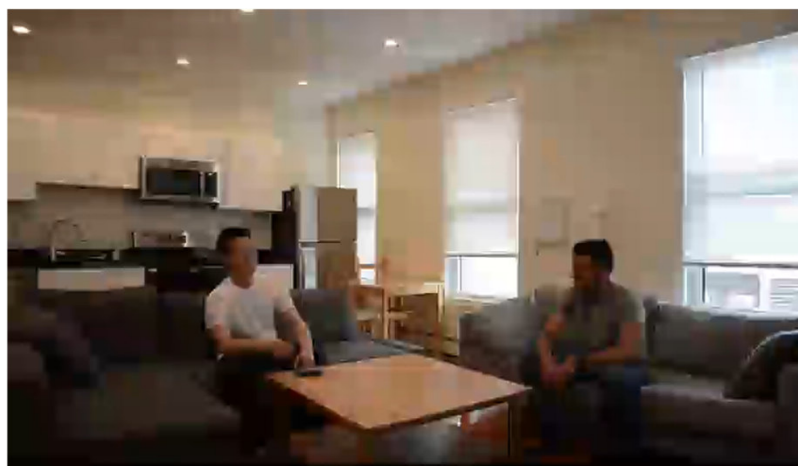
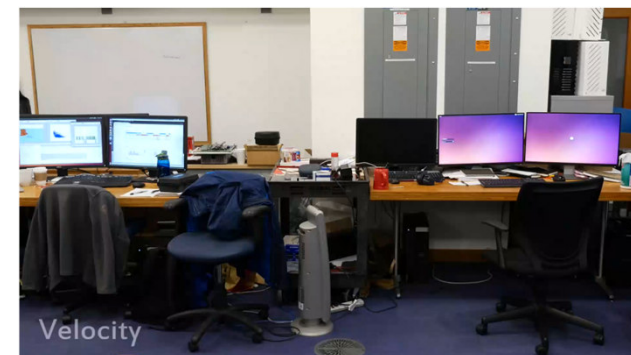
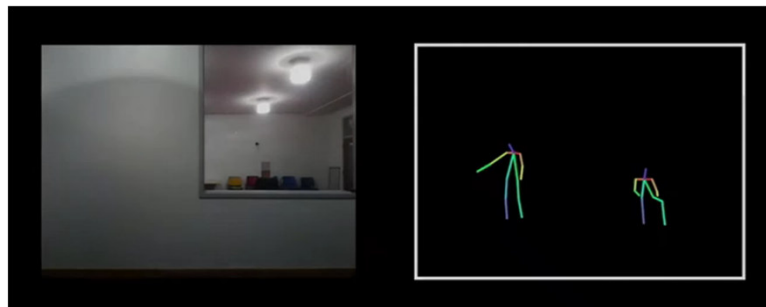
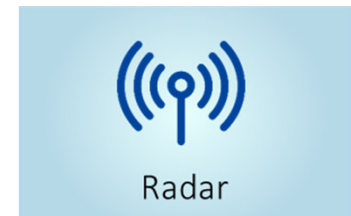
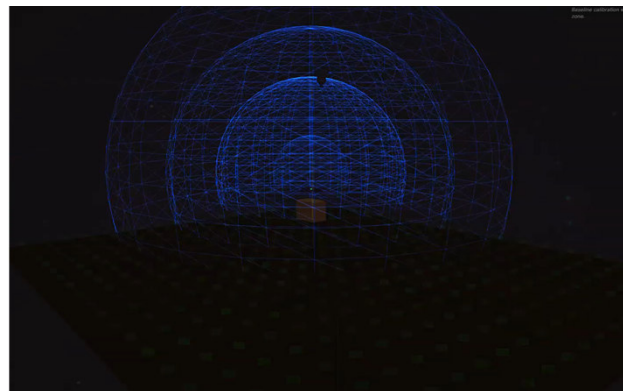
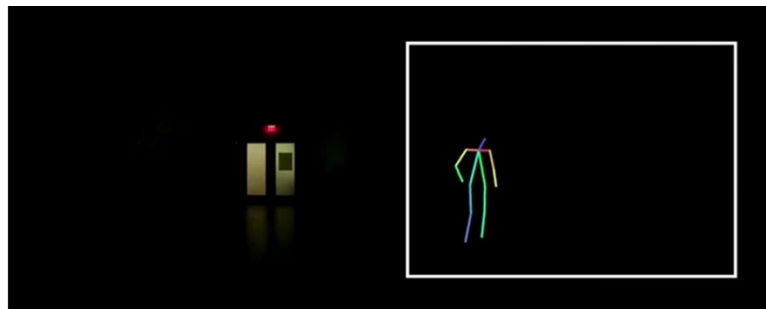
Radar Estimation:

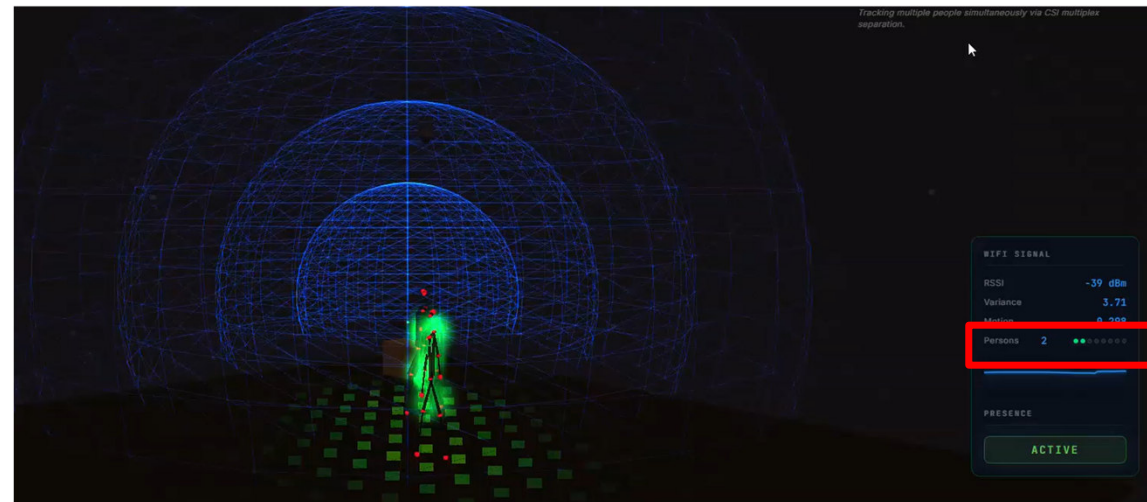
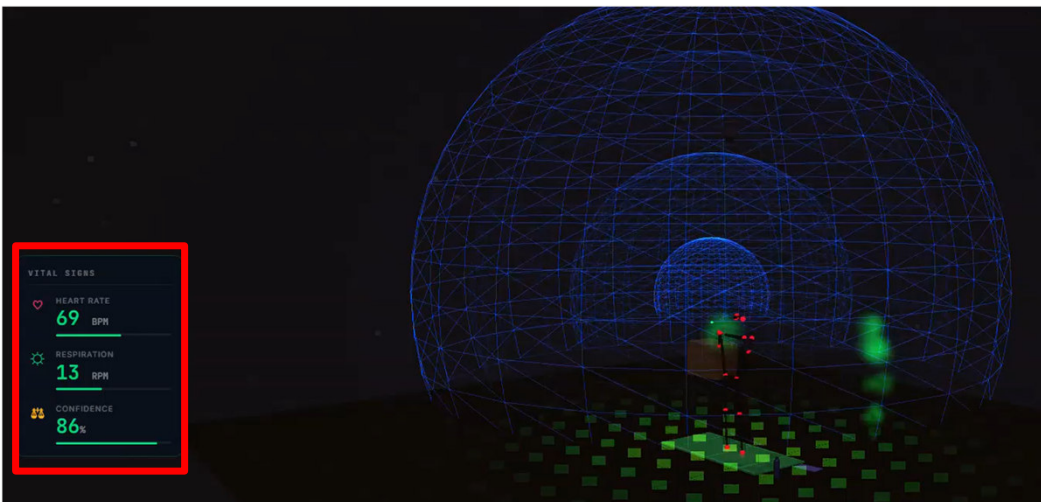
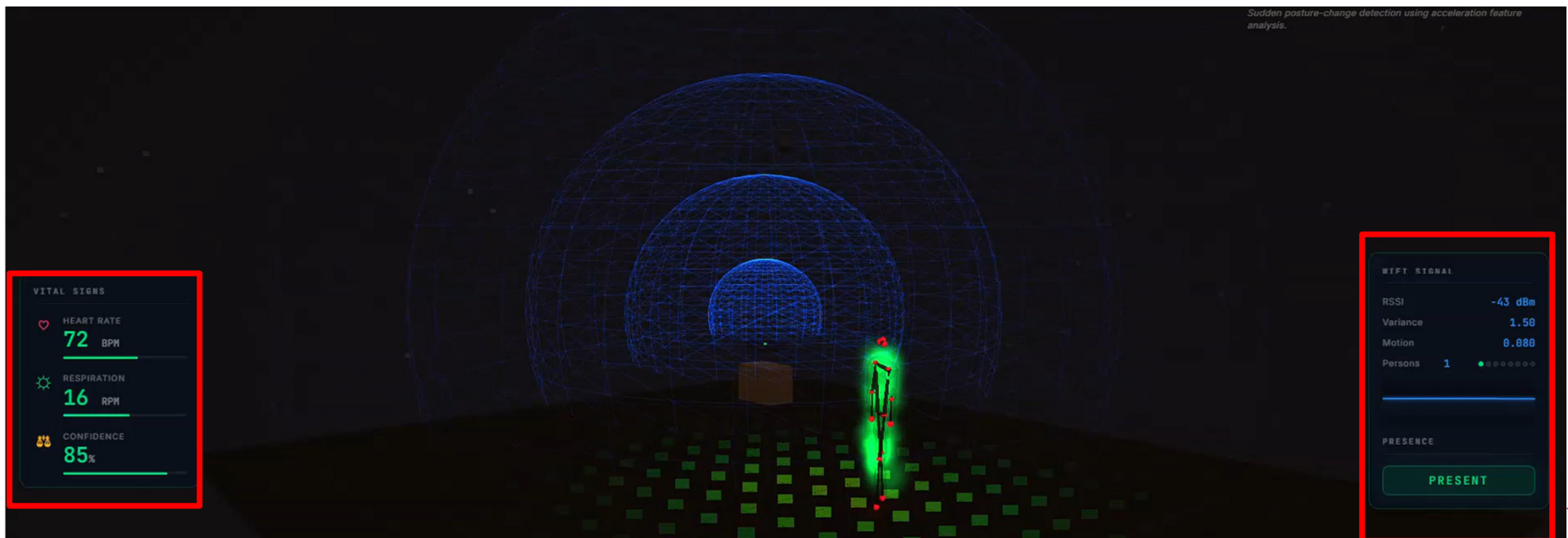


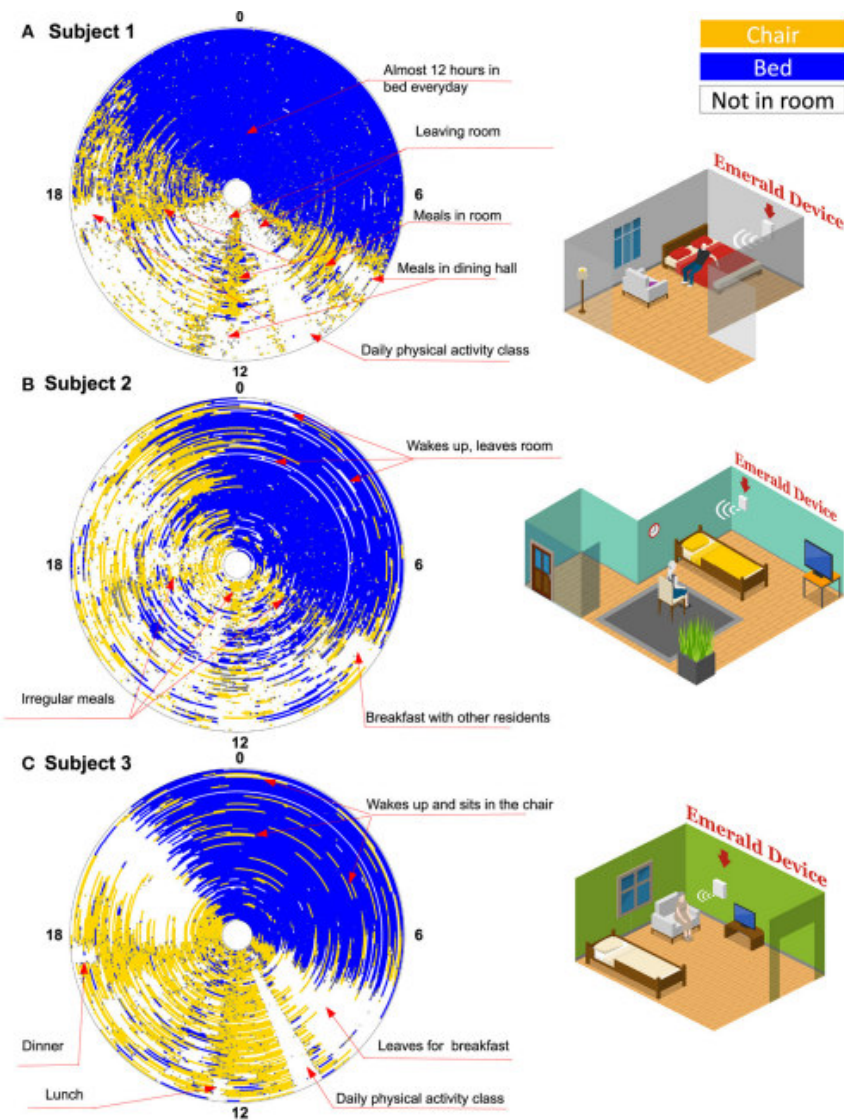
di Biase, L., et al. (2022). Markerless Radio Frequency Indoor Monitoring for Telemedicine: Gait Analysis, Indoor Positioning, Fall Detection, Tremor Analysis, Vital Signs and Sleep Monitoring. *Sensors*, 22(21), 8486. <https://doi.org/10.3390/s22218486>

An, S., Li, Y., & Ogras, U. (2022). mri: Multi-modal 3d human pose estimation dataset using mmwave, rgb-d, and inertial sensors. *Advances in neural information processing systems*, 35, 27414-27426.

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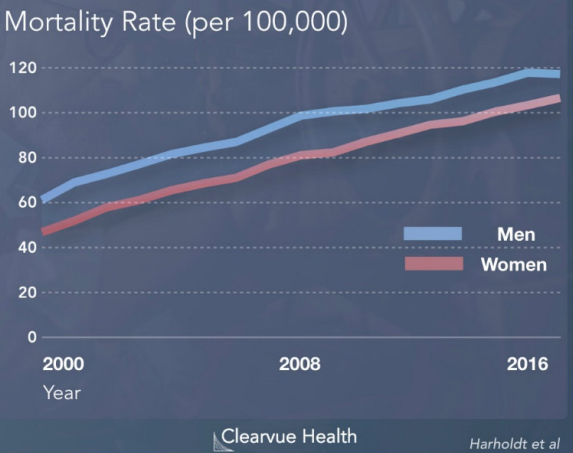




Zhang, G., et al., *Contactless In-Home Monitoring of the Long-Term Respiratory and Behavioral Phenotypes in Older Adults With COVID-19: A Case Series*. *Front Psychiatry*, 2021. **12**: p. 754169.

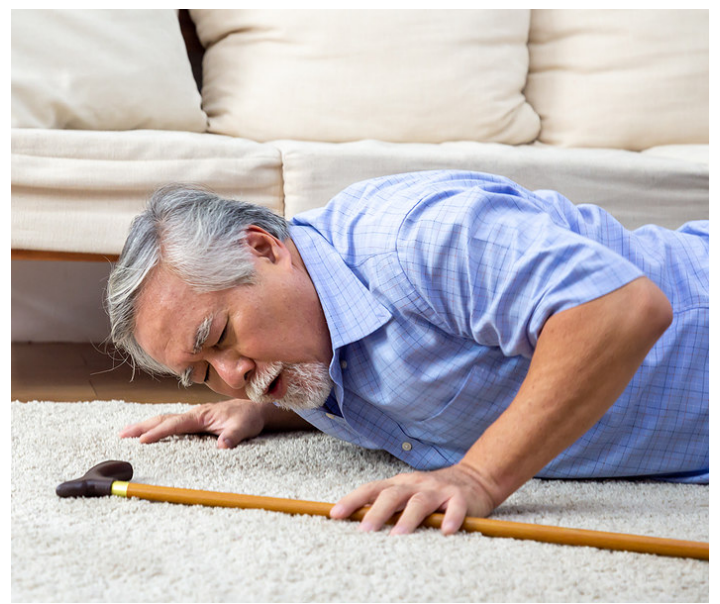
# Mortality Rate from Falls

Falls are a leading cause of death for the elderly. More elderly men and women are dying from falls every year.



# Low Impact Fall: Mortality

Even low impact falls at ground level can cause serious injury and mortality in the elderly.

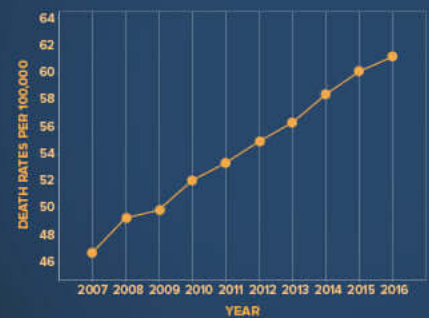


# Falls: Mortality Rate by Age

Fall mortality increases exponentially with age

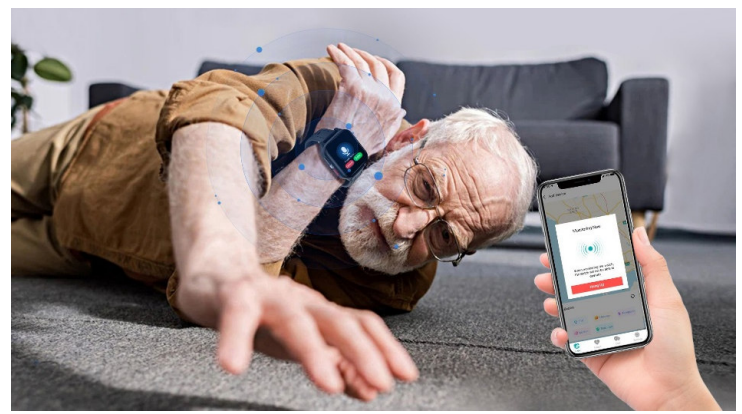


# Fall Death Rates in the U.S. INCREASED 30% FROM 2007 TO 2016 FOR OLDER ADULTS



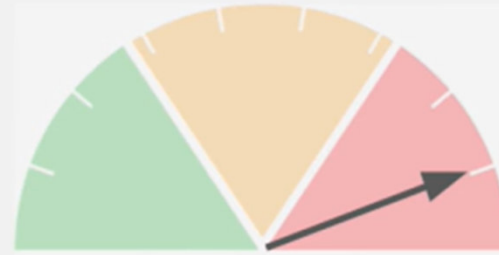
If rates continue to rise, we can anticipate **7 FALL DEATHS EVERY HOUR BY 2030**

Learn more at [www.cdc.gov/HomeandRecreationalSafety](http://www.cdc.gov/HomeandRecreationalSafety).

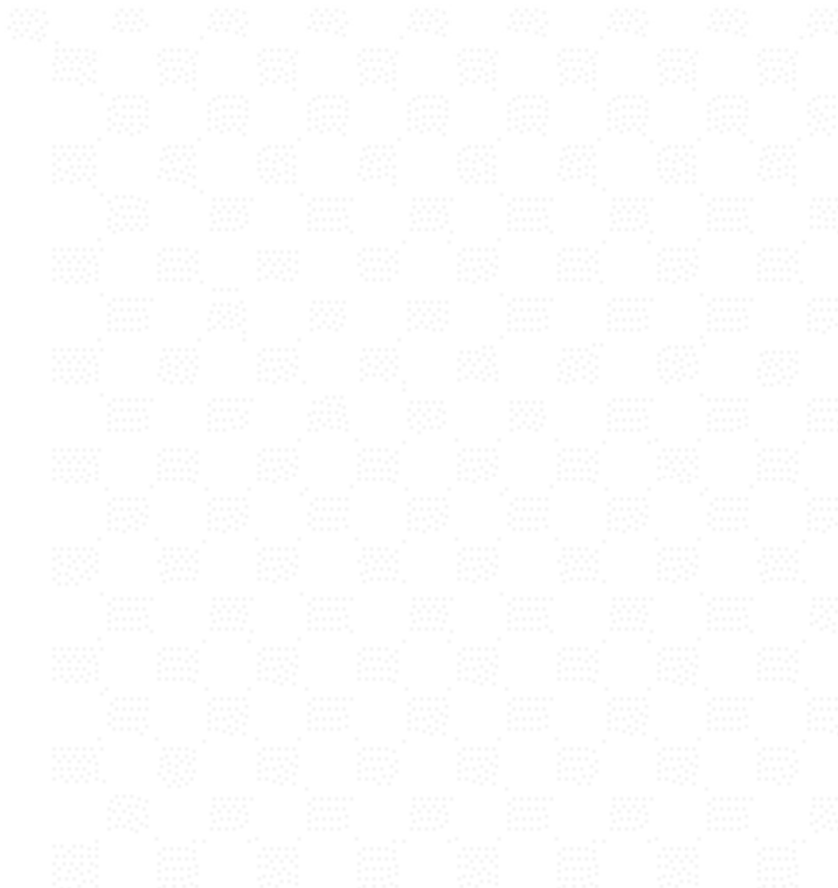




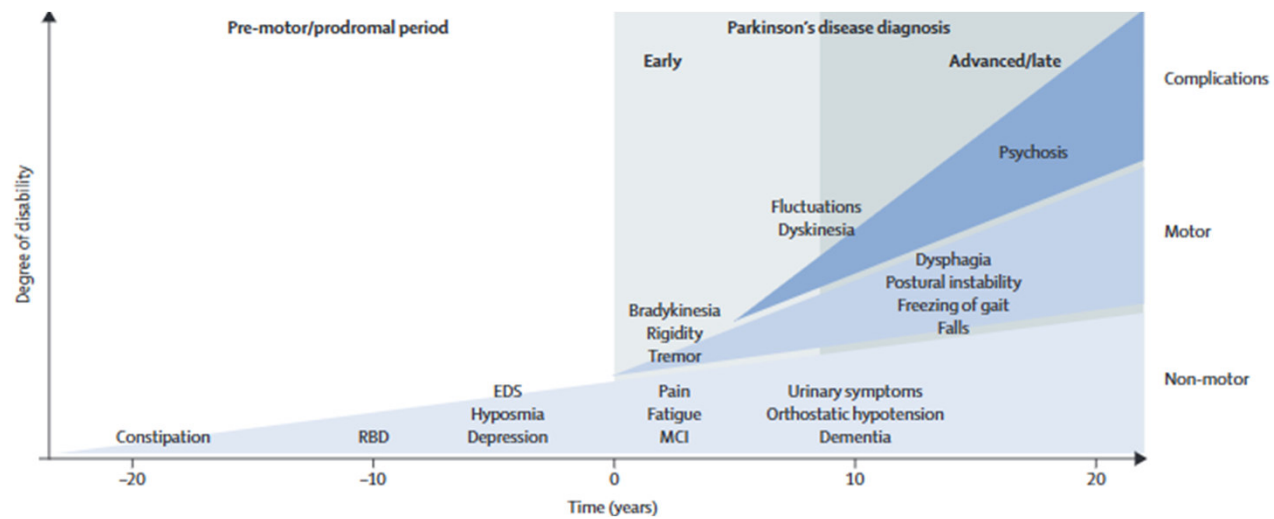
Name : James Smith  
PN : 07865467



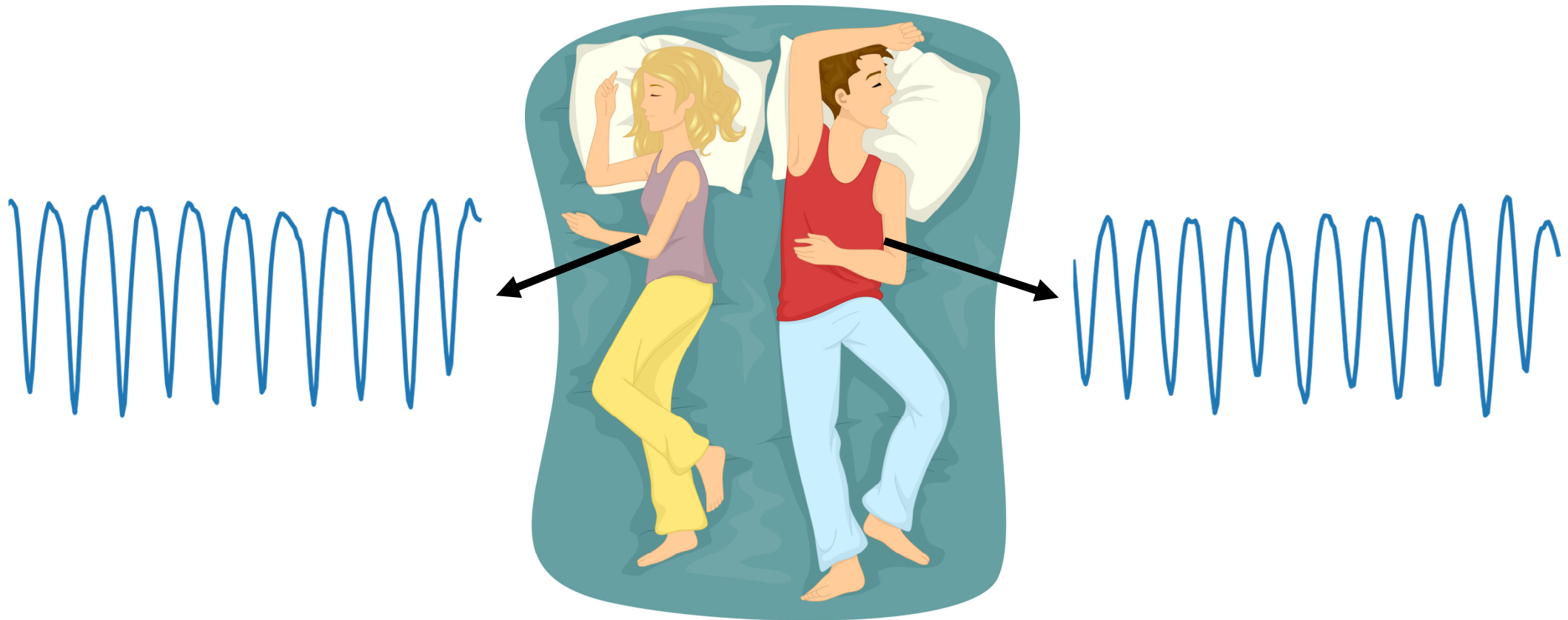
Total Falls Risk

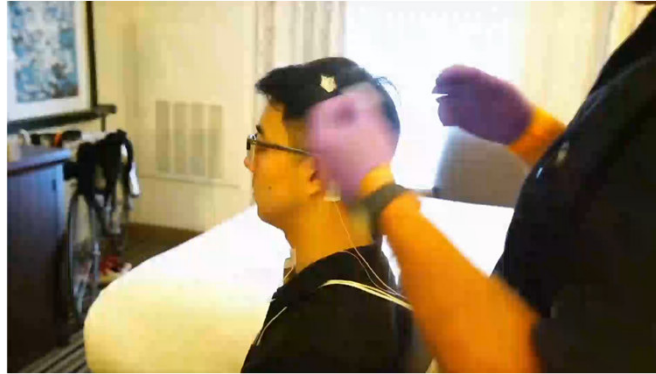
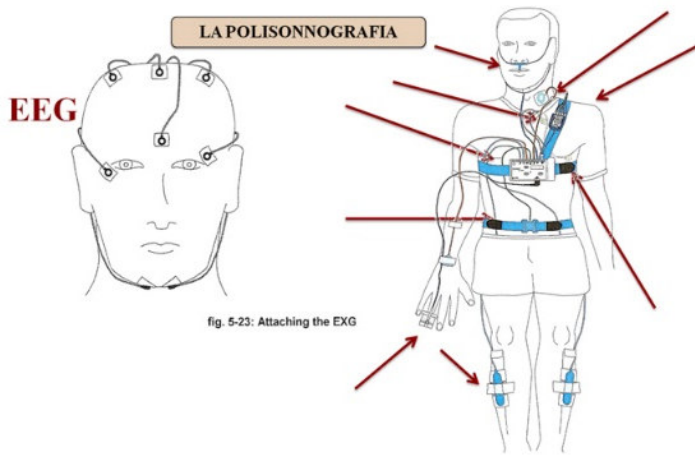


# SLEEP MONITORING

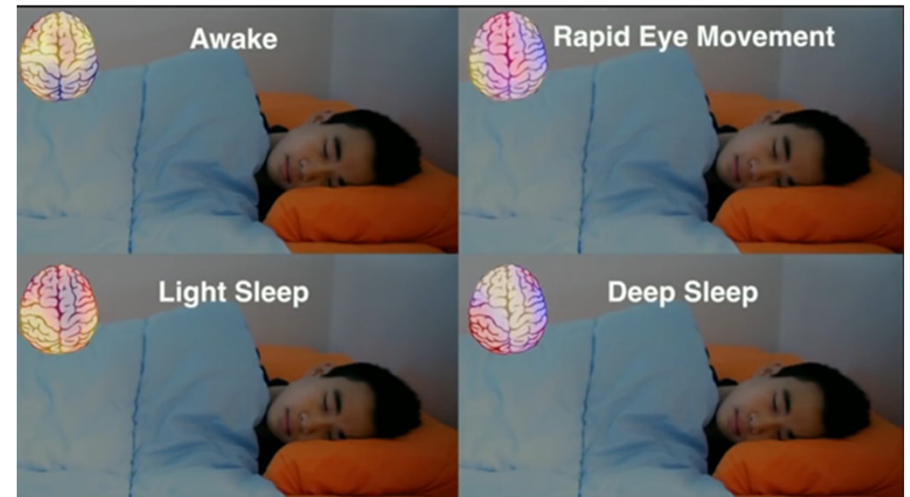
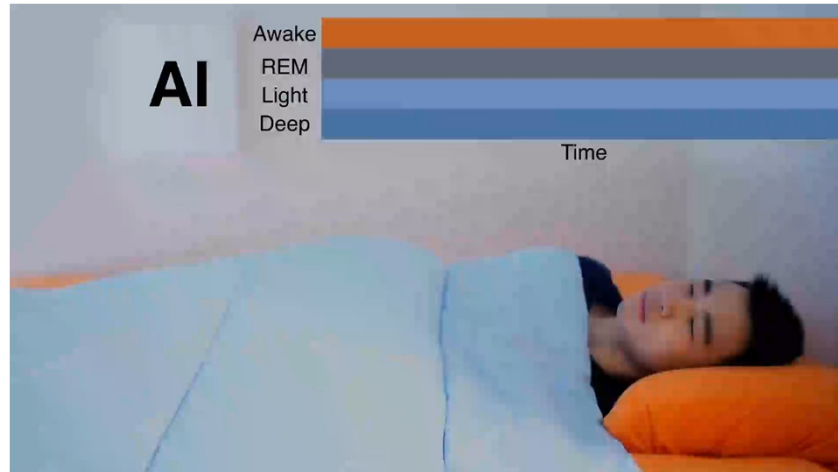
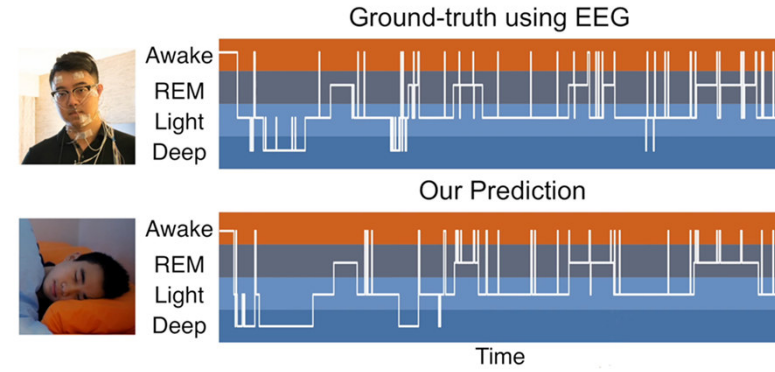


# Non-Contact Sleep Analysis





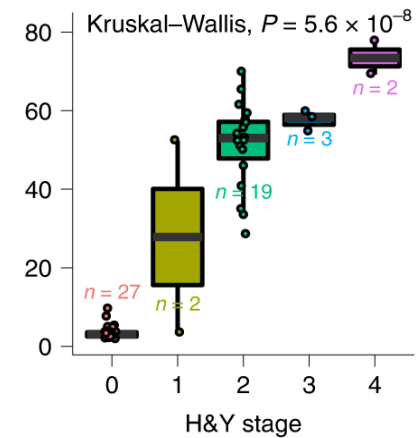
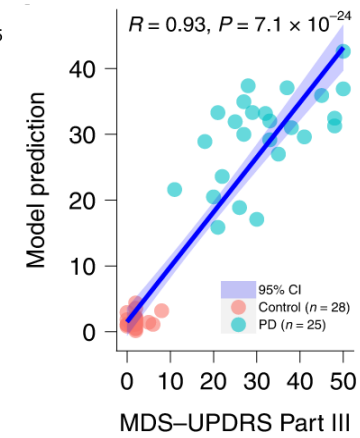
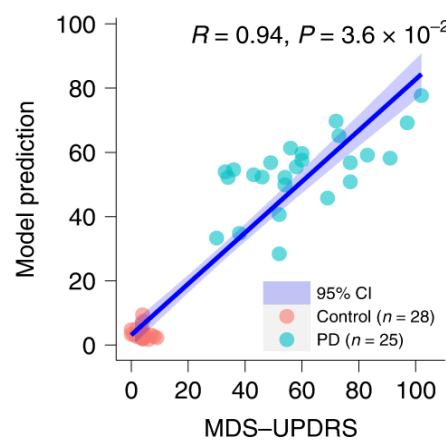
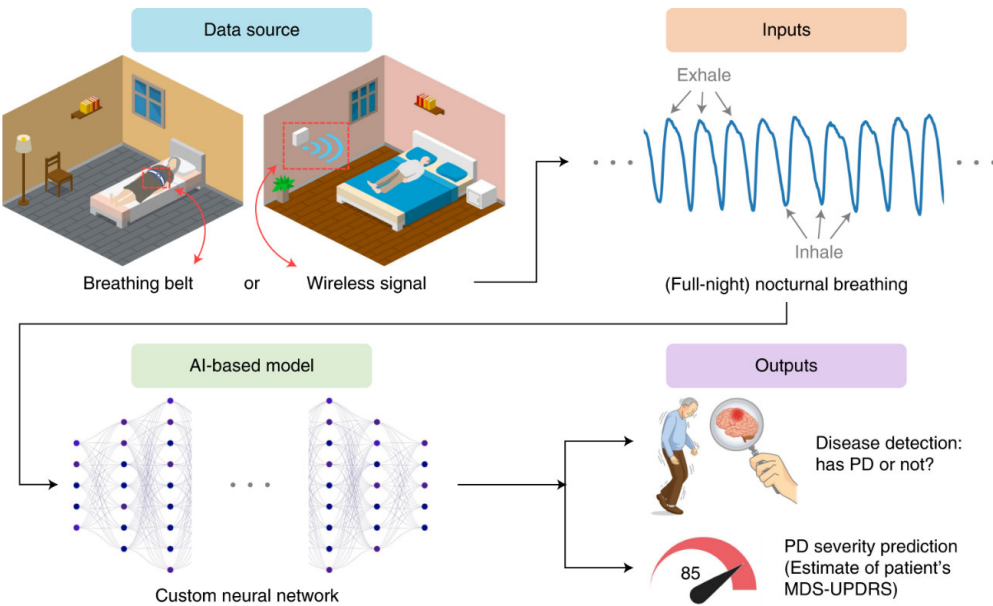
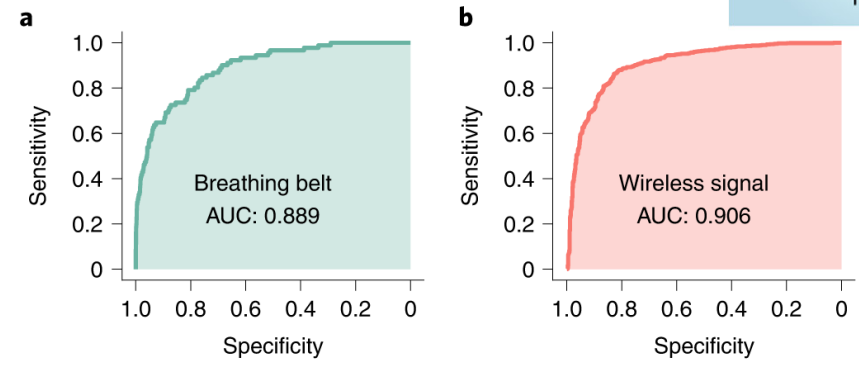
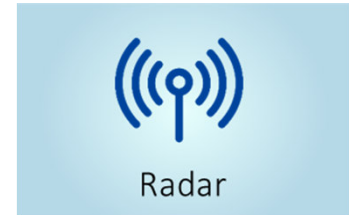
Representative Example Acc = 80%





**OPEN**  
**Artificial intelligence-enabled detection and assessment of Parkinson's disease using nocturnal breathing signals**

Yuzhe Yang<sup>1</sup>, Yuan Yuan<sup>1</sup>, Guo Zhang<sup>1</sup>, Hao Wang<sup>1,2</sup>, Ying-Cong Chen<sup>1</sup>, Yingcheng Liu<sup>1</sup>, Christopher G. Taroll<sup>3,4</sup>, Daniel Crepeau<sup>5</sup>, Jan Bukartyk<sup>6</sup>, Mithri R. Junna<sup>7</sup>, Aleksandar Videnovic<sup>8</sup>, Terry D. Ellis<sup>9</sup>, Melissa C. Lipford<sup>7</sup>, Ray Dorsey<sup>3,4</sup> and Dina Katabi<sup>1,10</sup>



Yang, Y., et al (2022). Artificial intelligence-enabled detection and assessment of Parkinson's disease using nocturnal breathing signals. Nature Medicine, 28(10), 2207–2215. <https://doi.org/10.1038/s41591-022-01932-x>

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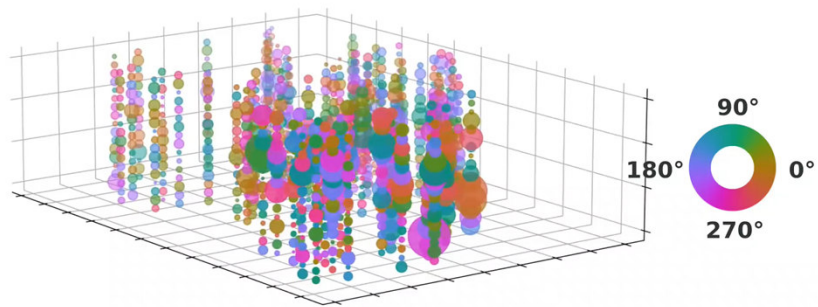
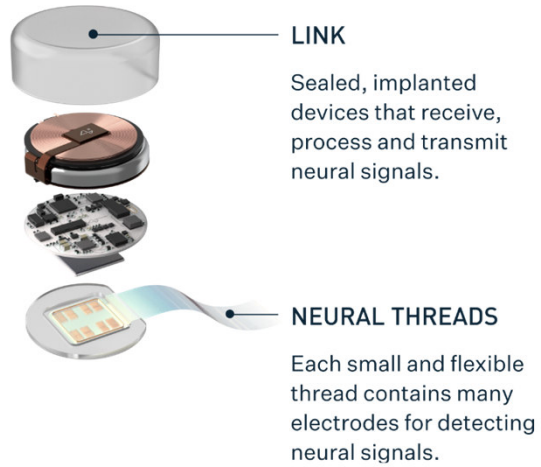
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# Brain Computer Interfaces

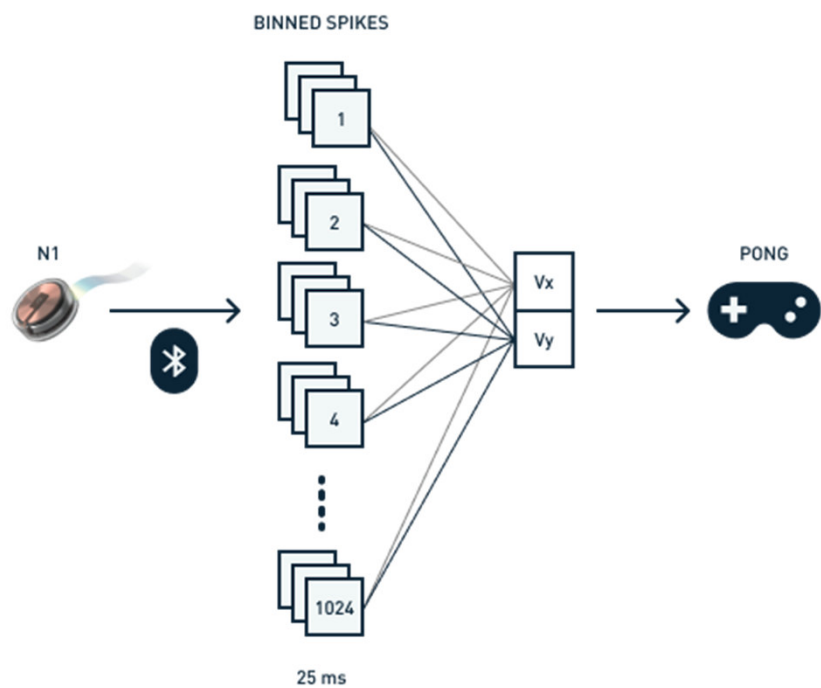
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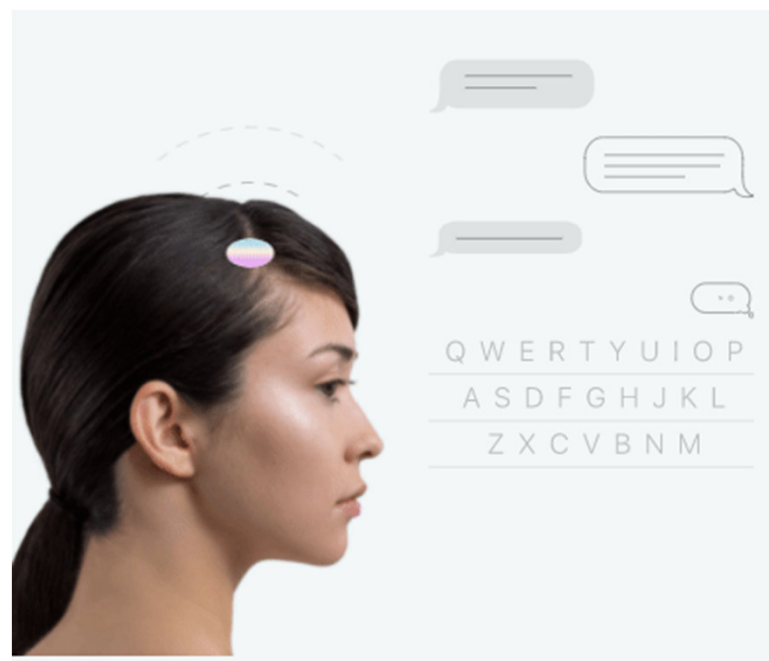
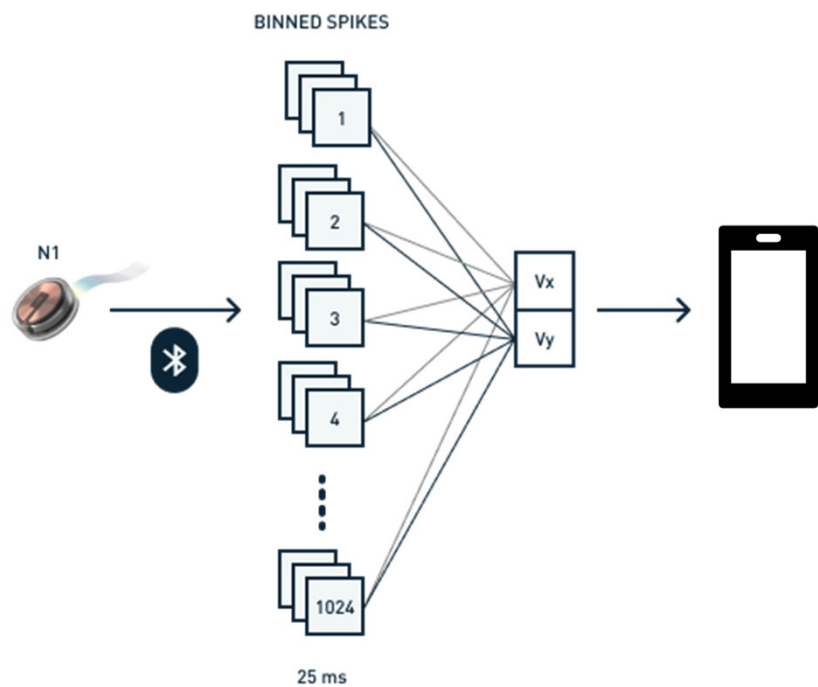
# BCI MOTOR CORTEX - NEURALINK

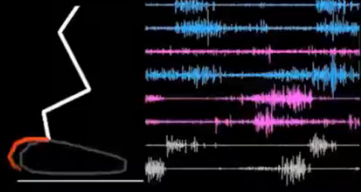


# BCI MOTOR CORTEX - NEURALINK

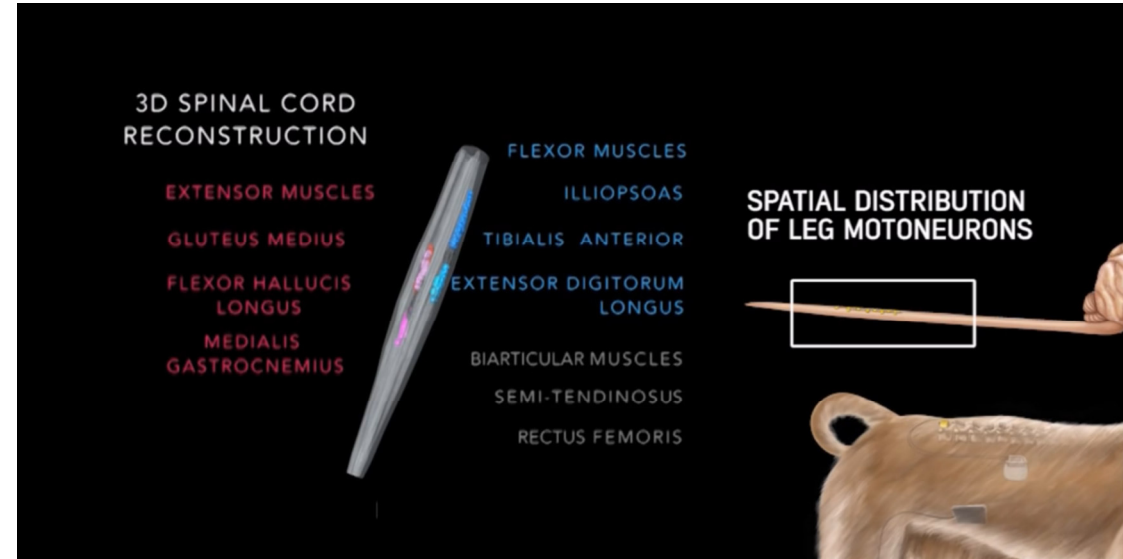
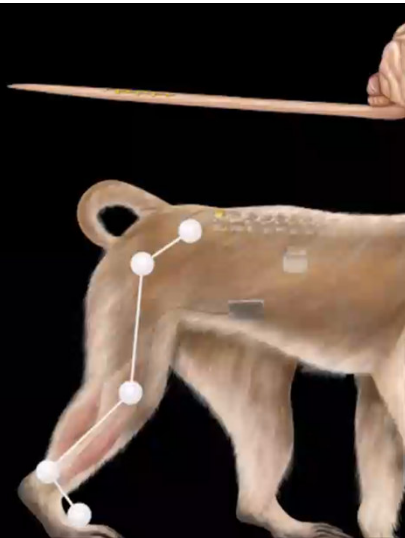


# BCI MOTOR CORTEX - NEURALINK

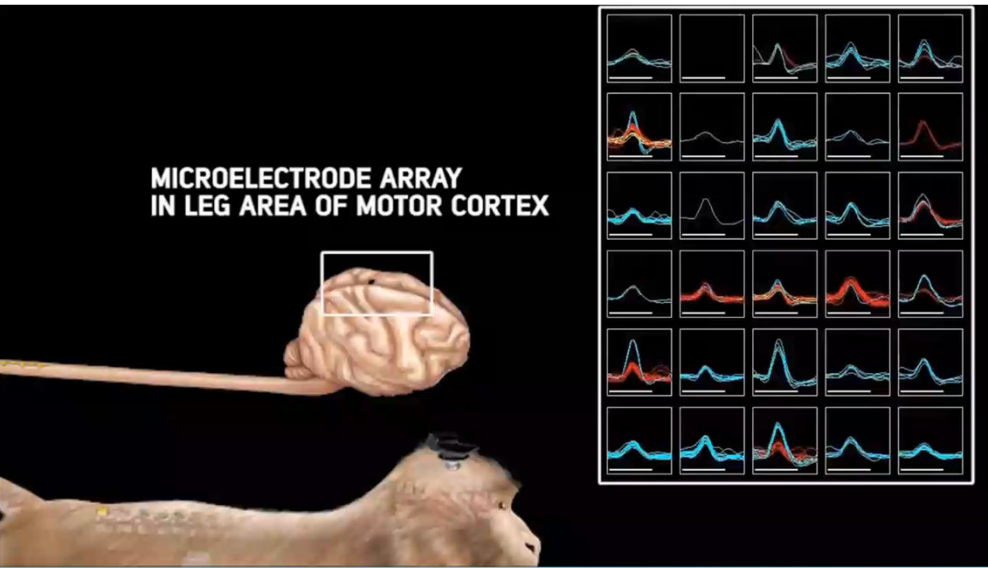




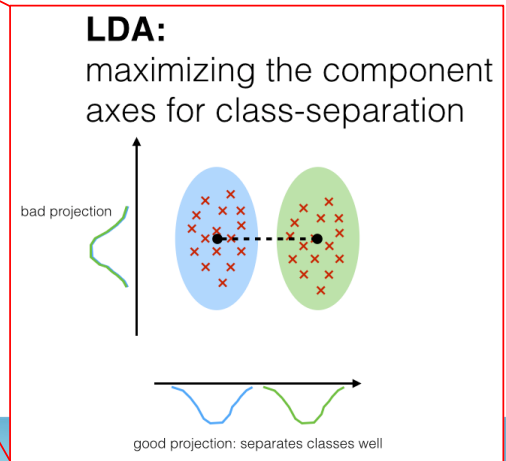
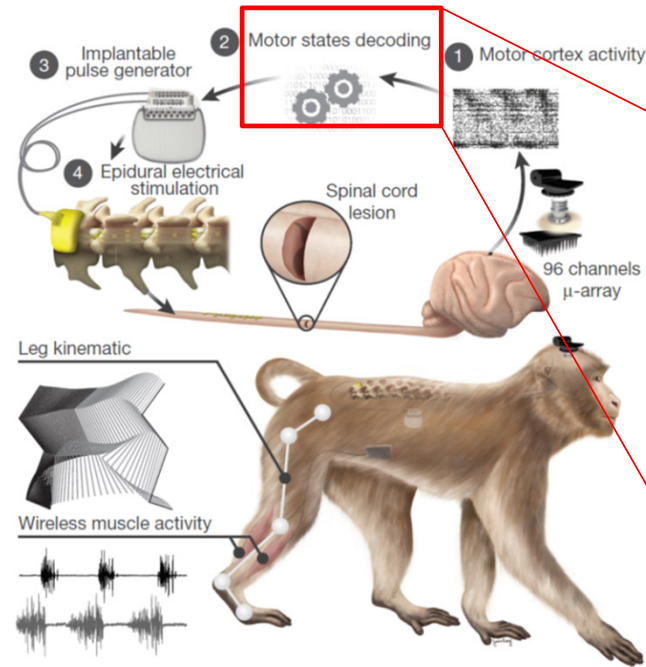
**LEG KINEMATICS AND MUSCLE ACTIVITY DURING GAIT**



Capogrosso, M., et al. (2016). *Nature*, 539(7628), 284-288.



**MICROELECTRODE ARRAY IN LEG AREA OF MOTOR CORTEX**

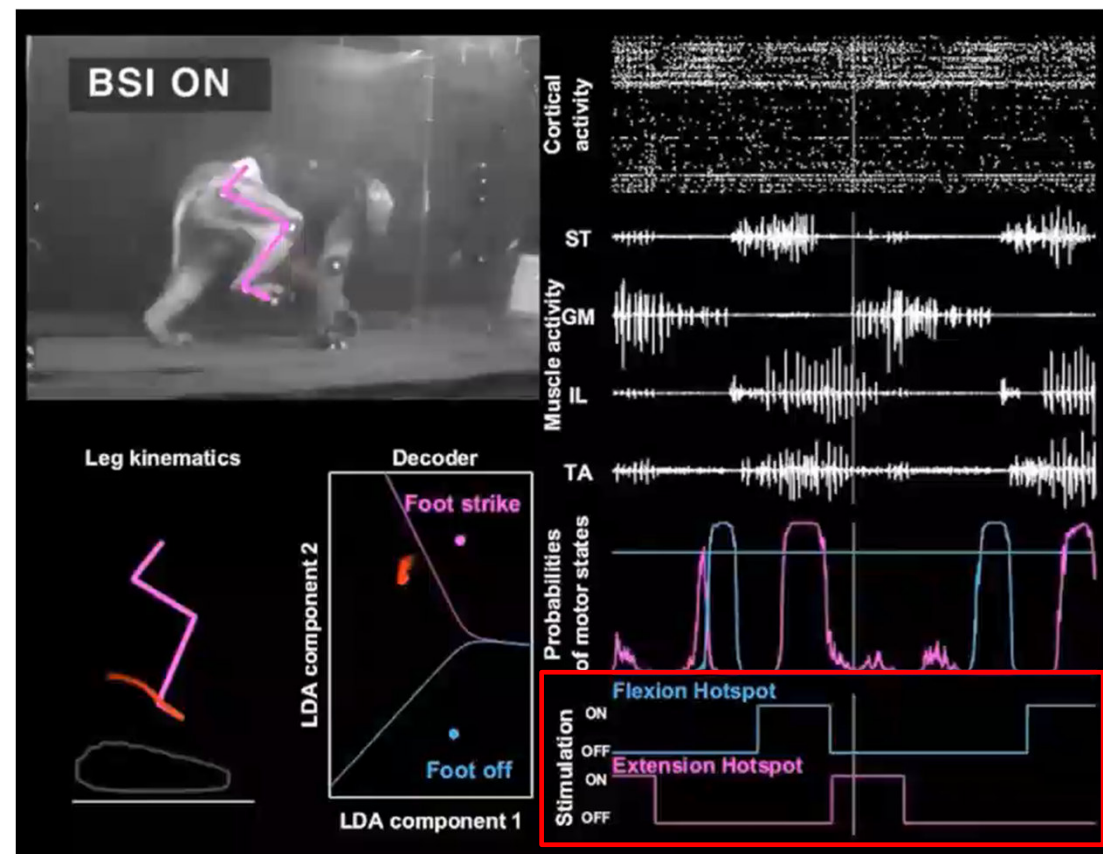
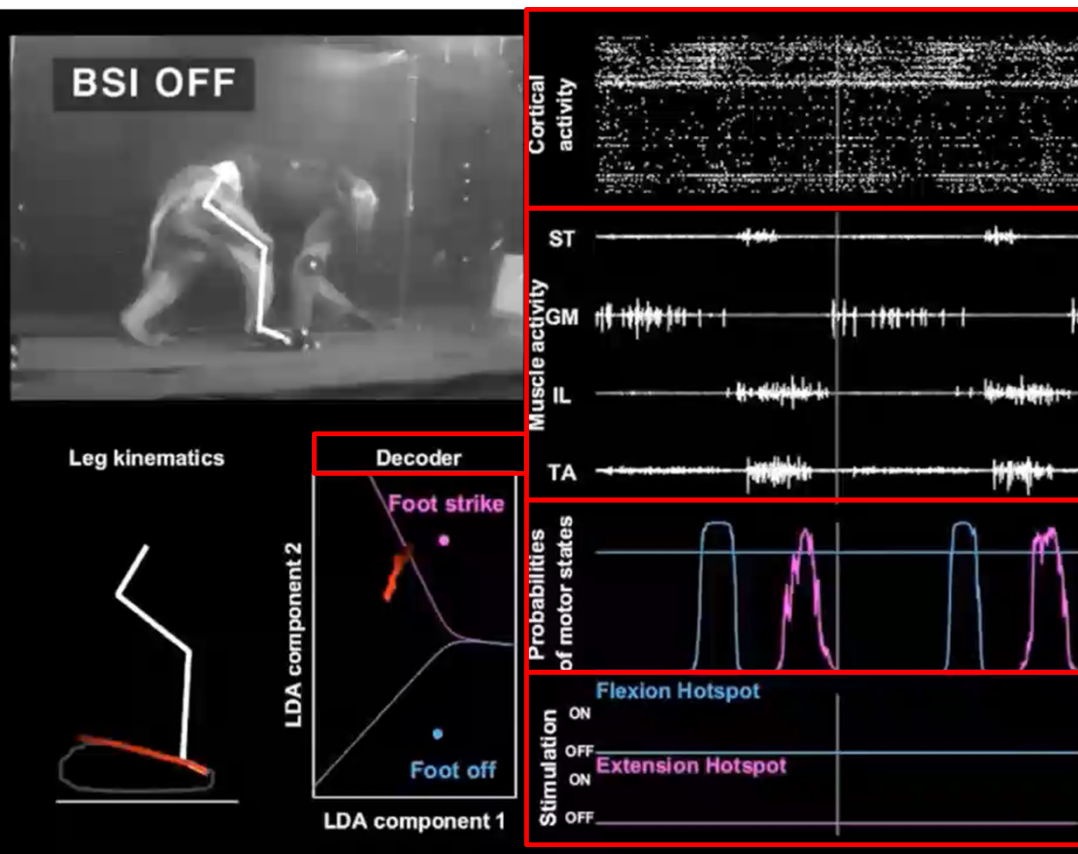


**LDA:**  
maximizing the component axes for class-separation

bad projection

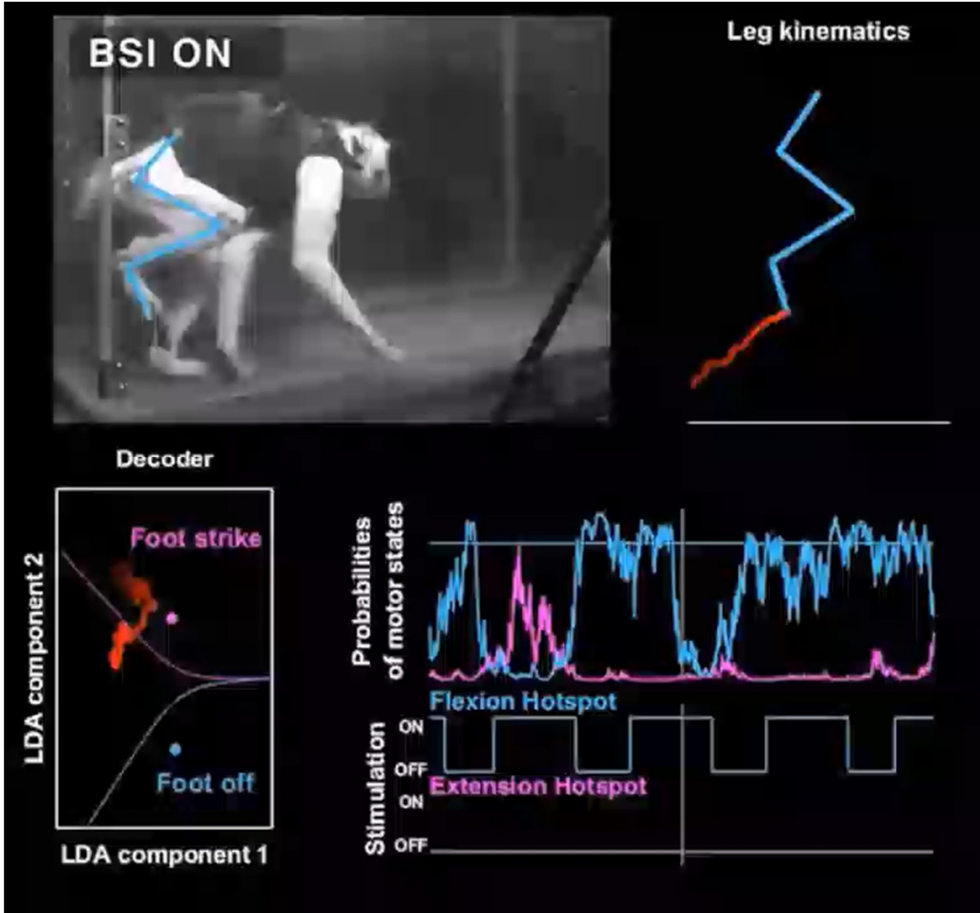
good projection: separates classes well

# BRAIN-SPINE INTERFACE: INTACT MONKEY

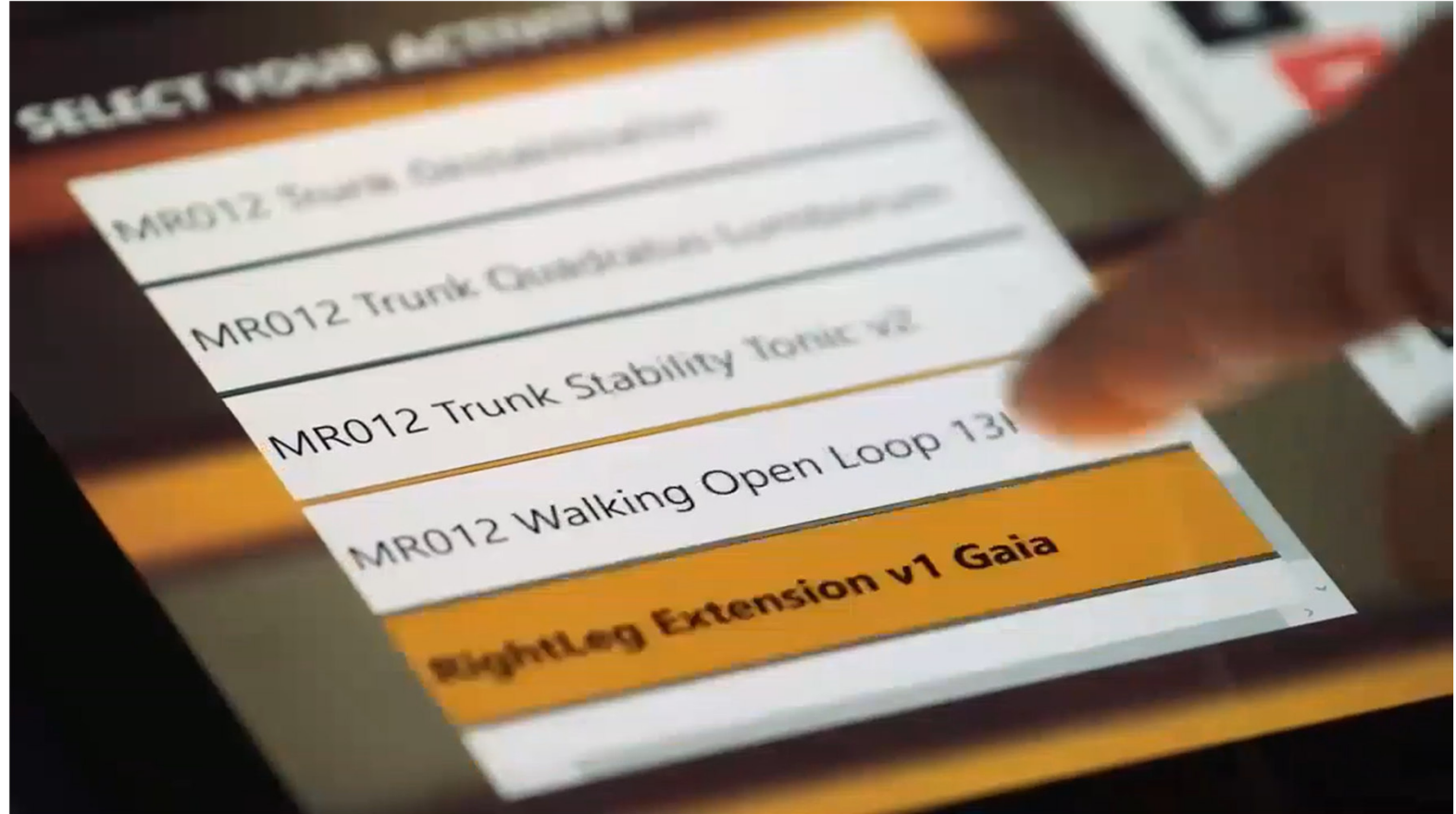


Capogrosso, M., et al. (2016). A brain–spine interface alleviating gait deficits after spinal cord injury in primates. *Nature*, 539(7628), 284–288.

# BRAIN-SPINE INTERFACE: POST SPINAL CORD LESION



Capogrosso, M., et al. (2016). A brain–spine interface alleviating gait deficits after spinal cord injury in primates. *Nature*, 539(7628), 284-288.

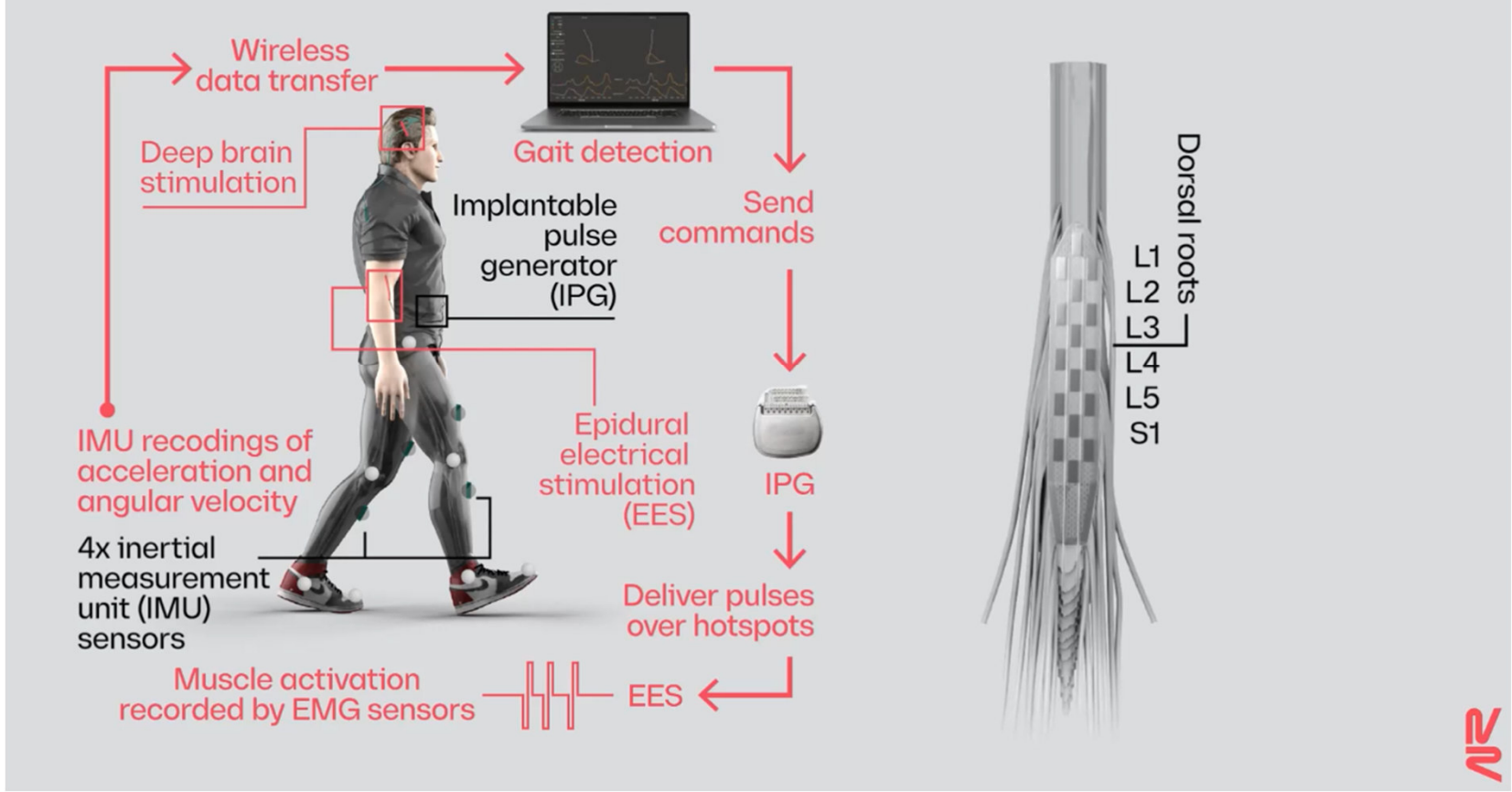


A brain-controlled spinal cord neuroprosthesis  
that alleviates locomotor deficits due to  
Parkinson's disease




Milekovic, T., et al. *Nat Med* **29**, 2854–2865 (2023).  
<https://doi.org/10.1038/s41591-023-02584-1>

# KINEMATICALLY-CONTROLLED SPINAL CORD NEUROPROSTHESIS




Spinal Cord Stimulation  
OFF

A brain-controlled spinal cord neuroprosthesis that alleviates locomotor deficits due to Parkinson's disease




A brain-controlled spinal cord neuroprosthesis that alleviates locomotor deficits due to Parkinson's disease




Spinal Cord Stimulation  
ON

A brain-controlled spinal cord neuroprosthesis that alleviates locomotor deficits due to Parkinson's disease



A brain-controlled spinal cord neuroprosthesis that alleviates locomotor deficits due to Parkinson's disease



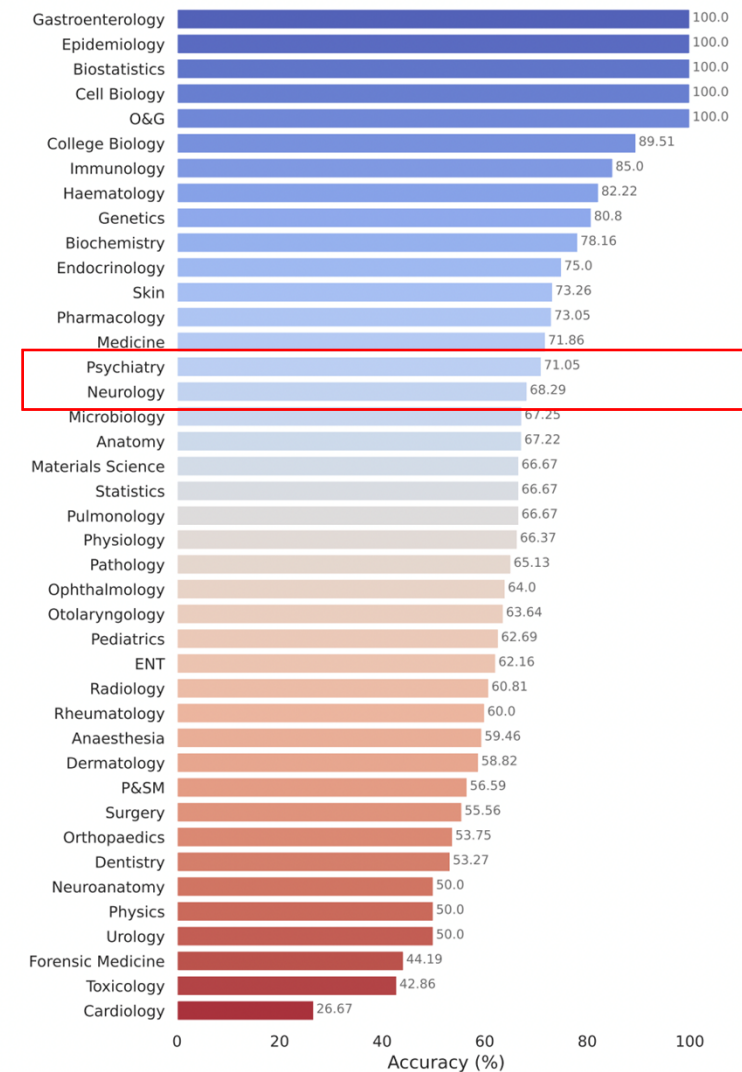
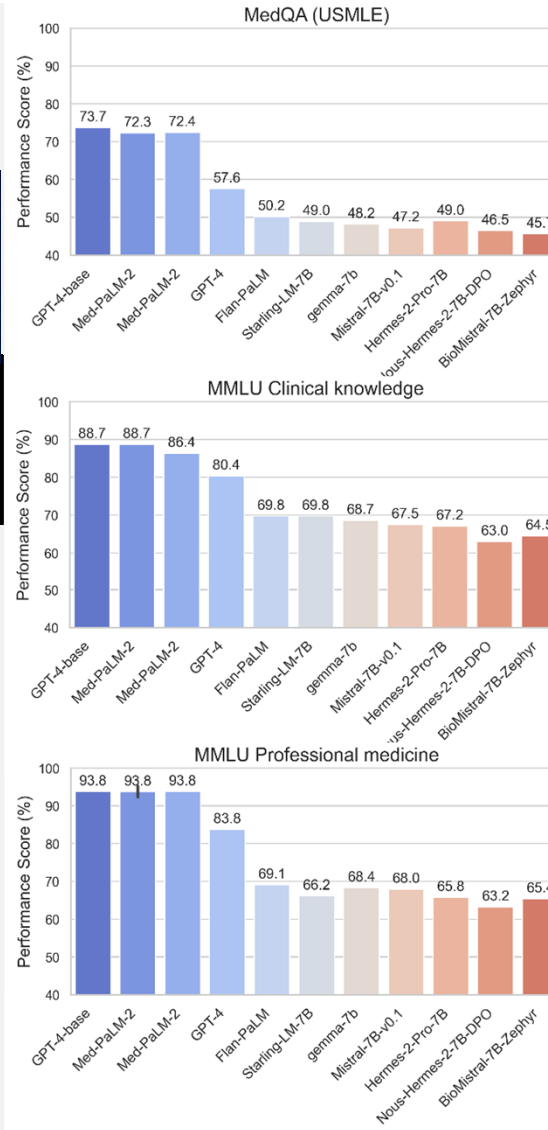
Milekovic, T., et al. *Nat Med* **29**, 2854–2865 (2023).  
<https://doi.org/10.1038/s41591-023-02584-1>

# Healthy Technology Act 2025 US Leads the AI Powered Prescription Revolution

IN THE HOUSE OF REPRESENTATIVES

JANUARY 7, 2025

Mr. SCHWEIKERT introduced the following bill; which was referred to the Committee on Energy and Commerce



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 Neurology Unit, Campus Bio-Medico University Hospital  
 Brain Innovations Lab

# WHY THE HUMAN BRAIN IS STILL SMARTER—AND MORE AMAZING—THAN AI



**12W**



**2,7 billion W**