

Patient-Focused Drug Development and Patient-Centered Clinical Care: *Working Together*

A Case Study in Multiple Sclerosis



Agenda and Workshop Panelists

- Introduction of Panelists and Workshop Topics

 Robert McBurney, PhD, Accelerated Cure Project & iConquerMS™ PPRN
- Experience of MS and Treatment
 Fodorica Palzani, Dr. (Law), Italian MS Social
 - Federica Balzani, Dr. (Law), Italian MS Society
- Patient-Centered Clinical Care and Patient-Reported Outcomes
 - Giampaolo Brichetto, MD PhD, Clinical Director Rehabilitation Center, Italian MS Society
- Patient-Focused Drug Development
 - Schiffon Wong, MPH, Exec. Dir., Global Evidence & Value Development, EMD Serono
- Audience Participation: Polls, Q&A and Contributions



Who is Participating in the Workshop? - Poll

RAISE YOUR HAND

- Patient or patient representative
- Healthcare practitioner
- Med. specialty society representative
- Pharmaceutical company personnel
- Device company personnel
- Diagnostic company personnel
- Service/Analytical company personnel

- Academic/Institutional researcher
- Representative of regulators
- HTA Agency representative
- Insurance Co. representative
- Government payers personnel
- ISPOR representative/staff
- Media representative
- Other



4

Who has Experience in....?

RAISE YOUR HAND

- Patient-Focused Product (Drug, Device, Dx) Development
- Patient-Centered Clinical Care
- Patient-Reported Outcomes
- Clinician-Reported Outcomes
- Performance Outcomes
- Shared Treatment Decision-Making
- Clinical Decision Support Tools



5

Workshop Topics

Robert McBurney



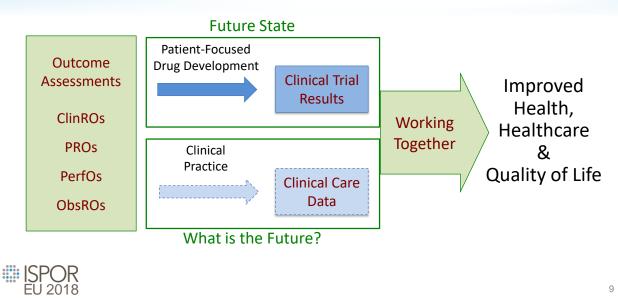
Topics to be Addressed in the Workshop

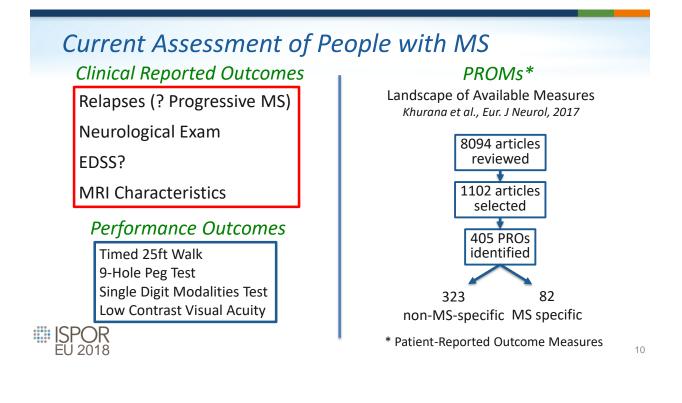
- What symptoms, etc., affect or matter most to people with MS
- Does clinical care or drug development address the symptoms, etc., that matter most to people affected by MS?
- What can be done/learnt in the clinic to enable patient-centered care?
- How can patient-focused drug development be achieved?
- How patient-focused drug development can work with patient-centered clinical care to improve outcomes for people with MS?

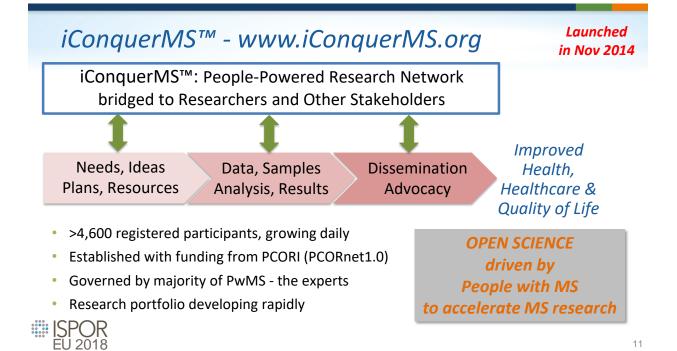
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Impact Areas for Outcome Assessments







What Affects iConquerMS™ Participants Most?

Rank Order	Neuro-QoL Domain (5-point Likert scale questions) Scored: 1[worst], 2, 3, 4 , 5[best]	Average Score (N = ~1,400)
1	Fatigue	2.89
2	Satisfaction with Social Roles and Activities	3.09
3	Sleep Disturbance	3.59
4	Positive Affect and Well Being	3.59
5	Ability to Participate in Social Roles and Activities	3.60
6	Anxiety	3.67
7	Cognitive Function	3.71
8	Emotional and Behavioral Dyscontrol	3.92
9	Lower Extremity Functional Mobility	3.93
10	Depression	4.11
11	Stigma	4.17
12	Communication	4.41
13	Upper Extremity Function Fine Motor ADL	4.54

All respondents as of May 2016

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The other symptoms and quality of life domains have <u>rarely</u> been available in MS clinical trial results and submissions

Mobility is captured in the regulatory outcome measure Expanded Disability Status Scale (EDSS)¹

1. ClinRO

13

iConquerMS[™] *is Collaborating in 2 Related Areas*





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Experience of MS and Treatment

Federica Balzani



Experience of MS and Treatment



From Injective Therapy to Oral Therapy

Symptom Domains that Matter to People with MS

✓ Fatigue
✓ Bladder dysfunction
✓ Walking and mobility
✓ Anxiety
✓ Depression
✓ Visual problems
✓ Hearing problems
✓ Pain
✓ Cognitive losses
✓ Ataxia and tremor



- \checkmark Balance and falls
- √Vertigo/dizziness
- √Temperature sensitivity
- √Sleep disturbances
- √Speech difficulties
- √Swallowing difficulties
- √Emotionalism
- ✓ Spasticity and spasm
- \checkmark Contracture at joints

17

Design with The End in Mind



Patient-Reported Outcomes: Design with the End in Mind MYTH: PROs are by definition patient-centered

REALITY: Not necessarily. Many PROs were developed with insufficient understanding about what matters to patients and inadequate partnership with patients in measurement development.

MYTH: If patients are reporting on an outcome, it matters to them.

REALITY: Many patients can be compliant with filling out surveys regardless of whether the questions resonate with what is important to them. PROMs can have adequate psychometric properties....yet still not capture outcomes that matter to patients.



MILKEN INSTITUTE

M multi-act

MULTI-ACT and the Science of Patient Input

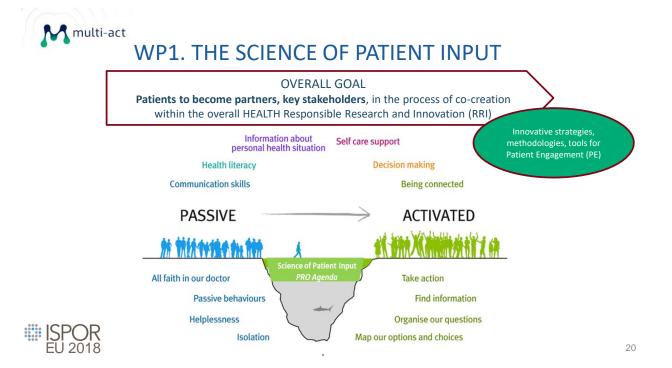
A Collective Research Impact Framework and multi-variate models to foster the true engagement of actors and stakeholders in Health Research and Innovation



Paola Zaratin, MULTI-ACT Project Coordinator on behalf of MULTI-ACT Consortium

> Multi-Act This project has received funding from the European Union's Horizon 2020 Research and Innovation Programme under the Grant Agreement No. 787570





Patient-Centered Clinical Care and Outcomes Assessments

Giampaolo Brichetto



MS Functional Domains & PROs in a Clinical Setting

Measuring MS is a hard challenge mainly due to the variety of functional problems that may affect individual patients

A universally accepted measurement instrument that is precise, reliable, easy to administer, captures the key neurological domains affected by MS, is sensitive at various levels of disability and that accurately reflects neurological and neuropsychological disability is lacking





MS Functional Domains & PROs in a Clinical Setting

Italian MS Society Foundation in 2013 promoted and funded the initiative:



"A new functional **PRO**file to **MO**nitor the **PRO**gression of disability in **M**ultiple **S**clerosis" (**PROMOPRO-MS**).



PROMOPRO-MS and Clinical Care



There is consensus in the clinical and scientific community that not only the long-established **Expanded Disability Status Scale (EDSS)**¹ (Kurtzke, 1983), but also the **Multiple Sclerosis Functional Composite (MSFC)**² (Fischer et al., 1999) **are inadequate to capture the change of the patients' clinical condition** (Cohen et al., 2012). 1. ClinRO 2. PerfO

PROMOPRO-MS: Key Questions

What is a set of outcomes related to the functional domains that really matter to People with MS? mobility, upper limb dexterity, fatigue, cognitive performance, emotional status, bladder continence, quality of life

How do we validate a "functional profile" of MS based on meaningful variables and measures?





PROMOPRO-MS and Clinical Care

The study is continuously collecting in a **real-life setting**, every four months, data from PwMS* followed at the Rehabilitation Centers of the Italian MS Society Foundation

- ClinRO EDSS Expanded Disability Status Scale
- ClinRO FIM[™] Functional Independence Measure
 - PRO Abilhand (perceived manual ability in daily life)
 - PRO OAB-Q Overactive Bladder Questionnaire
 - PRO M-FIS Modified-Fatigue Impact Scale
- PerfO **SDMT** Symbol Digit Modality Test
- PRO MoCA Questionnaire
- PerfO PASAT Paced Auditory Serial Addition Test
- PRO HADS Hospital Anxiety and Depression Scale
- PRO LSI Life Satisfaction Index



*No inclusion/exclusion criteria were considered apart from MS diagnosis as inclusion criteria



PROMOPRO-MS and Clinical Care

The proposed set of outcomes is currently used to measure disease progression and the efficacy of treatments (drugs and/or rehabilitation) in a real-life clinical setting.

27

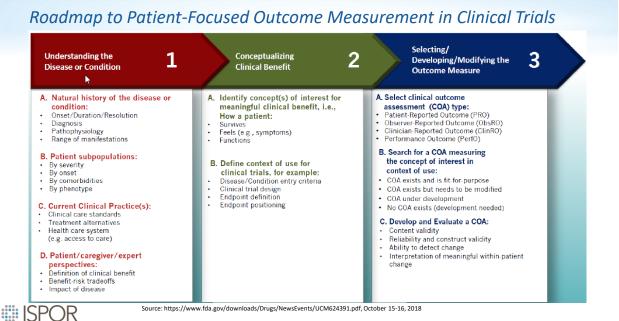
Patient-Focused Drug Development

Schiffon Wong

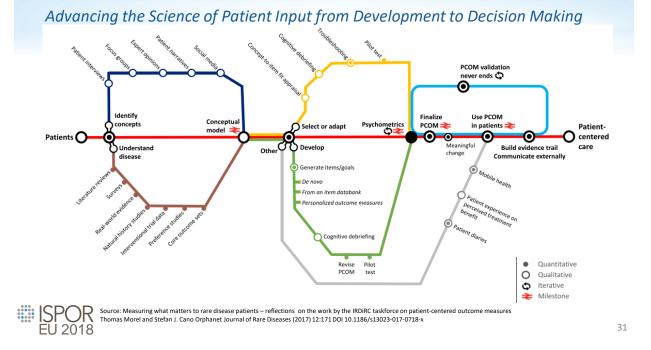




Meaningful, systematic consultation, involvement, and partnership with patients and their caregivers, will enhance the value of innovative medical products and improve the integration of therapeutics into the context of patients' lives



30



Transformative Collaboration: ACP/iConquerMS™ & EMD Serono

Patient feedback on	Patient feedback on	Establishing criteria for	Generate additional
design of PRO endpoints*	results of PRO measures	interpreting PRO	evidence/insights on MS
in MS clinical trials	validation studies	endpoints	disease burden
 Is there suitable coverage of domains most important to patients? Is the timing of PRO assessments consistent with patient experience? Identification of relevant aspects of patient experience to be assessed outside RCTs 	 Feedback on whether PRO measures reflect the patients' voice. Adding the patients' voice to the way validation study results are reported and communicated 	 Patient insights on clinical significance criteria for several PRO measures Conceptualizing the magnitude of benefit patients would consider relevant for the investigational therapeutic 	✓ Co-generate research questions, which will be investigated with the iConquerMS [™] patient- powered research network using mixed methods

*As a component of a mixed methods research program to support the development of PROs EU 2018

Practical Considerations from Conceptualization to Execution



Select program & identify priorities

Search & identify partner Consider complementary interests, resources, & capabilities

Joint brainstorming on potential scope See "eye-to-eye"

mutual benefits

Joint development of detailed scope Align on objectives, expectations, & define

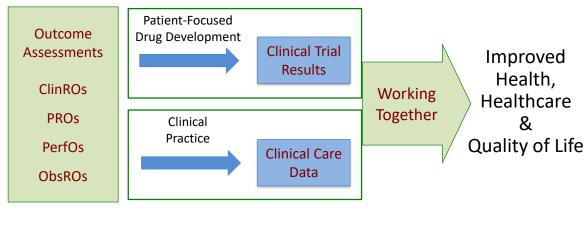


Audience Participation: Q&A and Contributions

Robert McBurney, Discussion Facilitator



How can Stakeholders make this a Reality?



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Workshop Questions

- From your stakeholder perspective, what are top priorities, or low-hanging fruit, for bringing together patient-centered clinical care and patient-focused product development?
 - Patient / Patient Representative
 - Product developer
 - Regulator
 - Healthcare technology assessor
 - Payer / Government
 - HEOR researcher
 - Medical specialty professional society
 - ISPOR



36

Workshop Questions

- What challenges are currently facing stakeholders in engaging with patients in patient-centered clinical care or patient-focused product development?
- What needs to be done to address the current challenges ?
- What opportunities do you see for translating the patient perspectives and insights captured during drug R&D into patient-centric care in clinical practice?



37

Workshop 14

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Thank You

