The Socioeconomic Impact of Disability Progression in Multiple Sclerosis: A Retrospective Cohort Study of the German **NeuroTransData Registry**

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BACKGROUND

- Disability progression on the Expanded Disability Status Scale (EDSS) is a common outcome measure of disease progression in clinical studies of multiple sclerosis (MS)^{1,2}
- The socioeconomic burden of MS goes far beyond the costs arising from the healthcare setting³

METHODS

Evaluation

NTD MS registry

- Registry network comprises 66 neurology and psychiatry clinics in Germany
- Database maintained since 2008
- Includes 22,000 PwMS with an average observation period of 5 years Routinely captures clinical and socioeconomic data, as well as patient-reported outcomes All PwMS, regardless of disease-modifying therapy (DMT) treatment and MS subtype, were eligible for inclusion Patients provided informed consent and explicitly agreed to secondary use of their data

Evaluation of HCRU and costs (Euro 2019)

- Full covariate information available for 7,286 patients
- Analyses conducted from the societal perspective with costs derived from public sources
- Stratified Health care resource utilisation (HCRU) by EDSS including:

NeuroTransData (NTD) MS registry contains real-world data for patients with MS (PwMS) living in Germany⁴

• Disability progression leads to decreased quality of life (QoL) and increased socioeconomic costs⁵

Statistical analysis

- Multivariate linear mixed regression modelling evaluated the associations between EDSS and 1) Qol 2) Cost
- Results were adjusted for confounders
- NTD treating center included as a random effect

- **Direct medical costs** (inpatient care, day admissions, tests, medication, DMTs)
- **Direct non-medical costs** (investments, community services, informal care)
- Indirect costs (short- and long-term employment) absences, pension status)

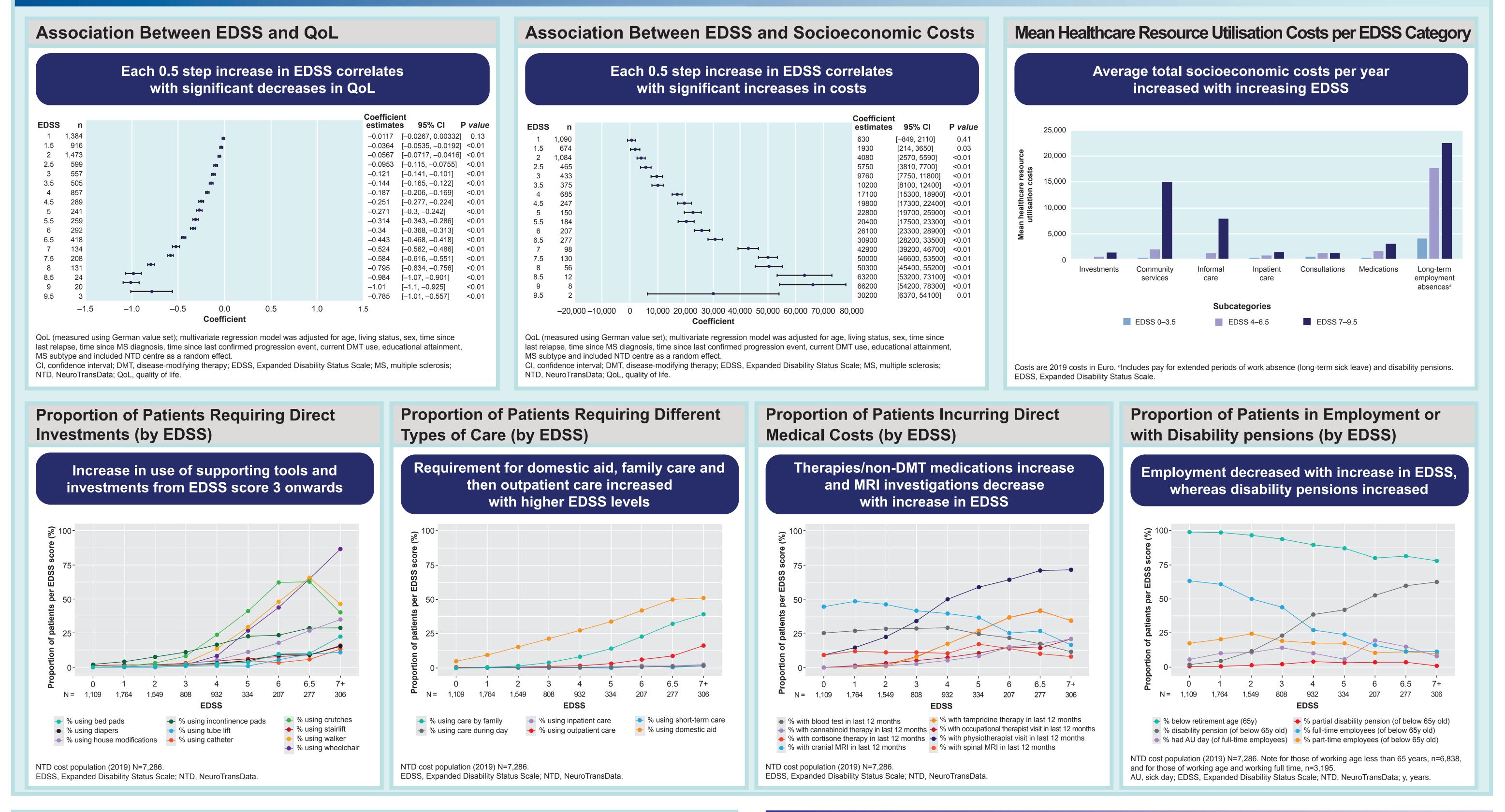
Evaluation of QoL

- Full covariate information available for 9,821 patients
- Utilities estimated from the EQ-5D-5L values recorded in the NTD database using the German value set published in Ludwig et al. (2018)⁶



The aim of this study was to evaluate the impact of disease progression on societal costs and QoL using data from the German NTD MS registry

RESULTS



STRENGTHS & LIMITATIONS

CONCLUSIONS

- Data on the full range of medical and societal costs on the impact of MS disease progression are not systematically captured during routine clinical practice within the NTD registry
- With regard to historical data, to ensure generalisability of findings regarding HCRU and costs, this analysis has been limited to a single recent year (2019) while for utility analyses, it is not expected that temporal trends exist in the relation between QoL and EDSS and thus data from 2009–2019 are used
- Informal care was recorded as binary response ('yes'/'no') without hours of care being recorded, and this could have led to inaccurate assignment of costs. Also, costs for investments were not directly recorded and had to be estimated based on assigning unit costs per investment type
- A strength of our study is the representative sampling of PwMS patients overcoming limitations of prior studies relying on survey-based recruitment, which may have led to exclusion of healthier patients at low EDSS and sicker patients at high EDSS
- Sampling differences may have led to observation of lower costs at lower EDSS and lower QoL at higher EDSS, compared with a previous survey-based study⁷



Each 0.5 step increase in EDSS score correlates with significant increases in socioeconomic costs and decreases in QoL

- Within a clinically representative MS population, increases in EDSS scores lead to decreases in QoL and increases in socioeconomic costs
- These findings highlight the socioeconomic burden of disease progression in MS
- Such findings highlight the potential value of halting or delaying disease progression through the early use of high efficacy therapies

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DISCLOSURES

Paul Dillon is an employee of F. Hoffmann-La Roche Ltd, Basel, Switzerland and has share/ownership of F. Hoffmann-La Roche Ltd. **Yanic Heer** is an employee of PricewaterhouseCoopers (PwC), Zurich, Switzerland. Eleni Karamasioti is an employee of PricewaterhouseCoopers (PwC), Zurich, Switzerland. Erwan Muros-Le Rouzic is an employee of and shareholder in F. Hoffmann-La Roche Ltd, Basel, Switzerland. Giuseppe Marcelli is an employee of F. Hoffmann-La Roche Ltd, Basel, Switzerland. Danilo Di Maio is an employee of F. Hoffmann-La Roche Ltd, Basel, Switzerland. Stefan Braune received honoraria from Kassenärztliche Vereinigung Bayerns and health maintenance organisations for patient care; and from Biogen, Merck, NeuroTransData Novartis and Roche for consulting, project management, clinical studies and lectures; he also received honoraria and expense compensation as a board member of NeuroTransData. Gisela Kobelt is President of EHE International GmbH. Jürgen Wasem is Professor for Health Services Management at University Duisburg-Essen, North Rhine-Westphalia, Germany. He received a honorarium for consulting study concept and quality assurance of data calculations

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