

Association of the Novel Clinical Trial Endpoint Multicomponent Improvement (MCI) with Event-Free Survival in Patients Living with Pulmonary Arterial Hypertension in the US

Tracey Weiss¹; Dominik Lautsch¹; Dena R. Ramey¹; Bennett Lane^{2,3}; Vladimir Valtchinov^{2,4}; Vallerie McLaughlin⁵

¹Merck & Co., Inc., Rahway, NJ, USA; ²TriAxia Health, Boston, MA, USA; ³University of Cincinnati, Cincinnati, OH, USA; ⁴Brigham and Women's Hospital, Boston, MA, USA; ⁵University of Michigan Health, Ann Arbor, Ann Arbor, MI, USA

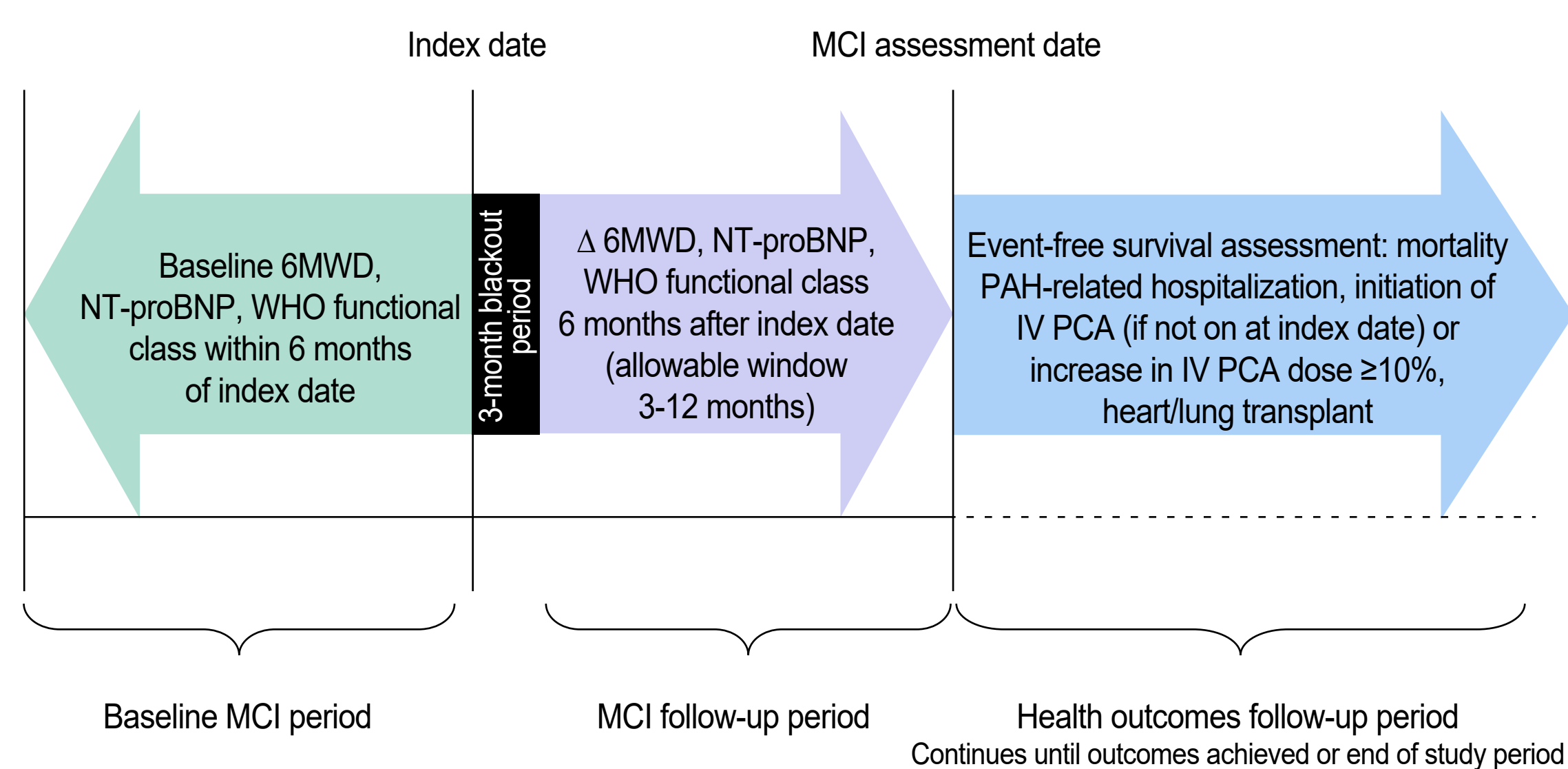
Purpose

- Pulmonary arterial hypertension (PAH) is a rare, progressive disorder in which remodeling of the pulmonary vasculature leads to increased pulmonary arterial pressure (PAP), eventually leading to right ventricular failure and death
- Three noninvasive measures – 6-minute walk distance (6MWD), World Health Organization functional class (WHO FC), and N-terminal pro-hormone B-type natriuretic peptide (NT-proBNP) – are used to assess clinical and risk status
- Multicomponent improvement (MCI) is a novel endpoint in the phase 3 trial for activin signaling inhibitor sotatercept. It is defined by attaining all 3 criteria of:
 - 6MWD increase of >30 m
 - WHO FC improvement or maintenance of WHO FC I or II
 - NT-proBNP reduction of ≥30% or NT-proBNP <300 pg/mL
- We conducted an exploratory evaluation of the association between MCI and event-free survival in real-world (RW) patients living with PAH in the US

Methods

- A retrospective cohort study was conducted in the TriAxia Health dataset, comprising electronic medical record data for 3,180 patients living with PAH from 4 academic medical centers (Massachusetts General Brigham, Stanford University, University of Pittsburgh Medical Center, University of Arizona)
- As shown in **Figure 1**, 3 study time periods were defined:
 - Baseline MCI period: 6-month window when all 3 baseline MCI components were collected, ending with observation of third MCI component at the “index date”
 - MCI follow-up period: 3-12 months after index date, ending with observation of third follow-up MCI component at “MCI assessment date”
 - Health outcomes follow-up period: Continues from MCI assessment date until outcome achieved or end of study period

Figure 1. Schematic of study time periods



- PAH was identified 1 of 3 ways:
 - Formal Dx by academic pulmonologist or cardiologist diagnosis **or**
 - Diagnostic right-heart catheterization (RHC) results consistent with PAH (mPAP >20 mmHg, PAWP ≤15 mmHg, and PVR >2WU). (Note: Criteria can be met by a single RHC or by data from multiple RHCs obtained within a 90-day period) **or**
 - Coding-based (ICD-9/ICD-10) diagnosis of PAH (I27.0, I27.20, I27.21, I27.81, I27.83, I27.89, I27.9) with no recorded codes for causes of other PH groups (I27.22, I27.23, I27.24, I27.29)
- All PAH patients aged ≥18 years receiving PAH therapy and with baseline 6MWD, WHO FC, and NT-proBNP and follow-up NT-proBNP and 6MWD (3-12 months later) were included
 - Per standard clinical practice, WHO FC was assumed unchanged at follow-up if not otherwise noted
- MCI achievement was evaluated at follow-up
- The proportion of patients with event-free survival was stratified by MCI achievement
- A time-to-event analysis for event-free survival over 10 years using Cox proportional hazards model with log-rank test was performed
 - Events were PAH-related hospitalization, initiation or titration of prostacyclin analogue, or heart/lung transplantation. R version 4.4.2 survival package performed computations

Disclosures

Funding for this research was provided by Merck Sharp & Dohme LLC, a subsidiary of Merck & Co., Inc., Rahway, NJ, USA (MSD). TW, DL, and DR are employees of MSD and are Merck & Co., Inc., Rahway, NJ, USA shareholders. BL & VV are employees of TriAxia Health, who was paid to perform this analysis by MSD. VM has received consulting fees from MSD.

Results

- The analysis included 116 patients. Mean (±SD) follow-up was 4.8 ± 3.3 years. Key demographic and clinical characteristics of the study cohort are in **Tables 1 and 2**

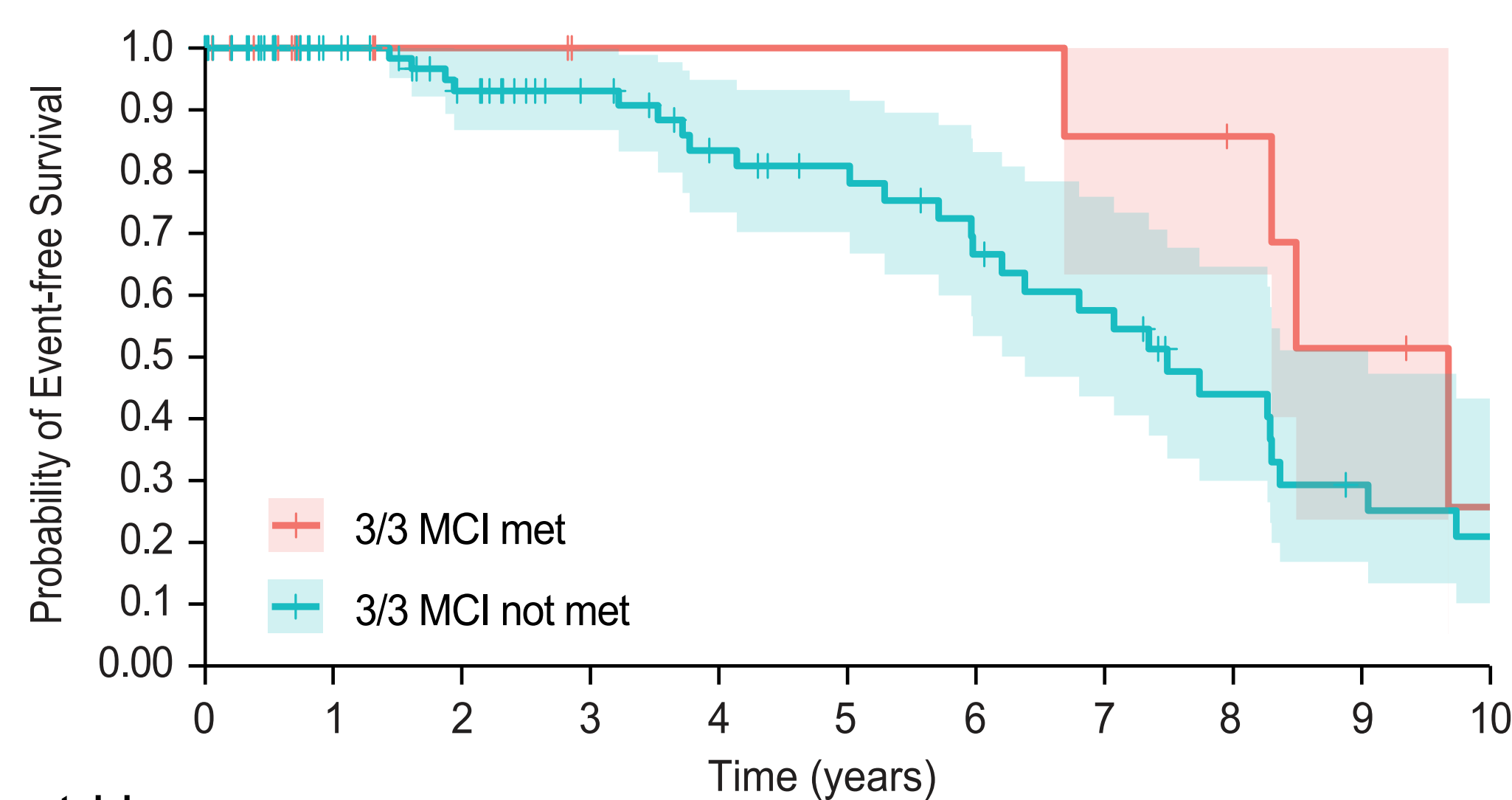
Table 1. Study population characteristics

Patient characteristics	N=116	%	Median (IQR)
Age at index date	–	–	54.0 (23.5)
Sex	Female	69.0%	
BMI	–	–	27.7 (9.9)
Timing of diagnosis	Newly dx (<6 mon of index date)	65	56.0%
	Prevalent	51	44.0%
PAH subtype	IPAH+HPH+DTI	65	56.0%
	PAH-CTD	26	22.4%
	PAH-OTHER	25	21.6%

Table 2. Study population clinical assessments

Clinical Assessments	N=116	%	Median (IQR)
Baseline			
PAH treatment at baseline	Monotherapy	39	33.6%
	Dual	22	19.0%
	Triple	1	0.9%
	None recorded	54	46.6%
6MWD	–	–	346.2 (188.1)
NT-proBNP	–	–	686.0 (1657.5)
WHO FC	I	13	11.2%
	II	43	37.1%
	III	54	46.6%
	IV	6	5.2%
Follow-up			
PAH treatment at follow-up	Monotherapy	48	41.4%
	Dual	49	42.2%
	Triple	4	3.4%
	None recorded	15	12.9%
6MWD	–	–	397.2 (188.3)
NT-proBNP	–	–	452.0 (1133.5)
WHO FC	I	16	13.8%
	II	48	41.4%
	III	48	41.4%
	IV	4	3.4%

Figure 2. Time-to-event analysis Kaplan-Meier curve for event-free survival



- 19 patients (16.4%) achieved MCI
- Over 10 years, 31/96 patients not achieving MCI (32.3%) and 4/19 patients achieving MCI (21.1%) experienced an event (HR 0.343 [95%CI, 0.118-0.996])
- This result is borderline significant with a log-rank $P = 0.049$. Due to small sample size, the results must be interpreted with caution (**Figure 2**)

Limitations

- Real-world data from US academic centers is not representative of all US PAH patients
- No data collected prospectively; analysis limited to data recorded during clinical practice

Conclusion

- Achievement of MCI was associated with 10-year event-free survival in a real-world US cohort of PAH patients

Access Slides



<https://bit.ly/3R44bZj>

Copies of this presentation obtained through QR (Quick Response) codes are for personal use only and may not be reproduced without permission of the authors.

Access Poster



<https://bit.ly/4aPC6Pi>