

MEDIA UPDATE

Novartis data highlight benefit of early treatment initiation in patients with secondary progressive multiple sclerosis (SPMS)

- *Novartis presented a total of 34 abstracts at the Congress of the European Academy of Neurology (EAN), emphasizing its strong multiple sclerosis (MS) portfolio with 20 abstracts*
- *Long-term efficacy data from EXPAND, previously presented at the American Association of Neurology Congress, show patients with SPMS continuously treated with Mayzent® (siponimod) experienced a sustained effect in delaying disability for up to 5 years, demonstrating the advantages of early treatment¹*
- *These data also show that for every 2 years of treatment with Mayzent patients can achieve 1 year of delay of progression and cognitive decline²*
- *EXPAND data demonstrated Mayzent consistently slowed cortical grey matter and thalamic atrophy across all SPMS patient subgroups, including those with less active disease and higher disability³*

Basel, May 28, 2020 — Novartis announced today that Mayzent data from the Phase III EXPAND trial were published in the *European Journal of Neurology* after the 6th EAN, held virtually due to COVID-19. These data, which included separate post hoc analyses from the Phase III EXPAND trial, continue to build on existing clinical evidence that Mayzent has significant impact on reducing the risk of disease progression, including physical disability and cognitive decline for patients with SPMS⁴.

“We are pleased to highlight data demonstrating that Mayzent helps slow disability progression and declining cognitive function,” said Norman Putzki, M.D., Global Head of Development Neuroscience. “For people living with MS, it’s essential to get ahead of disease progression and treat early so that they can maintain as much independence as possible in the future. We are dedicated to reimagining MS treatment and bringing hope to people with progressive diseases like MS.”

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generally be identified by words such as “potential,” “can,” “will,” “plan,” “may,” “could,” “would,” “expect,” “anticipate,” “seek,” “look forward,” “believe,” “committed,” “investigational,” “pipeline,” “launch,” or similar terms, or by express or implied discussions regarding potential marketing approvals, new indications or labeling for the investigational or approved products described in this media update, or regarding potential future revenues from such products. You should not place undue reliance on these statements. Such forward-looking statements are based on our current beliefs and expectations regarding future events, and are subject to significant known and unknown risks and uncertainties. Should one or more of these risks or uncertainties materialize, or should underlying assumptions prove incorrect, actual results may vary materially from those set forth in the forward-looking statements. There can be no guarantee that the investigational or approved products described in this media update will be submitted or approved for sale or for any additional indications or labeling in any market, or at any particular time. Nor can there be any guarantee that such products will be commercially successful in the future. In particular, our expectations regarding such products could be affected by, among other things, the uncertainties inherent in research and development, including clinical trial results and additional analysis of existing clinical data; regulatory actions or delays or government regulation generally; global trends toward health care cost containment, including government, payor and general public pricing and reimbursement pressures and requirements for increased pricing transparency; our ability to obtain or maintain proprietary intellectual property protection; the particular prescribing preferences of physicians and patients; general political, economic and business conditions, including the effects of and efforts to mitigate pandemic diseases such as COVID-19; safety, quality, data integrity or manufacturing issues; potential or actual data security and data privacy breaches, or disruptions of our information technology systems, and other risks and factors referred to in Novartis AG’s current Form 20-F on file with the US Securities and Exchange Commission. Novartis is providing the information in this media update as of this date and does not undertake any obligation to update any forward-looking statements contained in this media update as a result of new information, future events or otherwise.

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References

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3. Fox R, et al. Effect of Siponimod on Grey Matter Atrophy in Patients with Secondary Progressive Multiple Sclerosis: Subgroup Analyses from the EXPAND Study. *Eur J Neurol*. 2020;27(1).
4. Kappos L, Cree B, Fox R, et al. Siponimod versus placebo in secondary progressive multiple sclerosis (EXPAND): a double-blind, randomized, phase 3 study. *Lancet*. Published online March 22, 2018. [http://dx.doi.org/10.1016/S0140-6736\(18\)30475-6](http://dx.doi.org/10.1016/S0140-6736(18)30475-6).

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