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## FENhance 1 & 2 Phase III Studies in RMS

## FENhance 1 & 2 [NCT04586010, NCT04586023] **Patient population** • RMS diagnosis, aged 18-55 years • Disease duration <10 years from symptom onset • EDSS score of 0.0-5.5 at screening Double-blind treatment (1:1 randomization) **Treatment Active Comparator** Fenebrutinib Teriflunomide Primary endpoint: annualized relapse rate Non-MRI assessments Q12Wk week BL 48 **12** 24 36 96 Patients discontinuing treatment enter safety follow-up cCDP12<sup>†</sup> MRI OLE ≈96 weeks (optional)

<sup>†12-</sup>week composite confirmed disability progression (cCDP12) is a composite outcome measure defined as the first occurence of: a CDP-EDSS confirmed at 12 weeks OR a sustained increase of at least 20% from baseline in 9-HPT score. CDP-EDSS is defined as a sustained increase of ≥1.0 points from baseline scores ≤5.5 (or ≥0.5 points from baseline EDSS scores ≥6.0).



