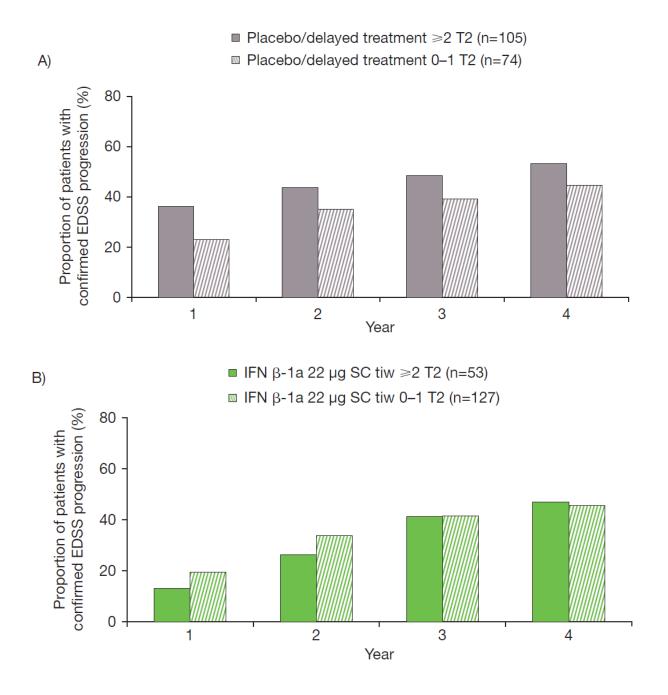
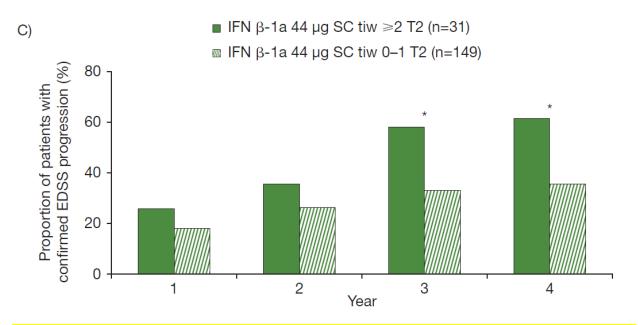
Additional file 6





Supplementary Fig. 5 Proportion progressed at each year by ≥2 versus 0–1 active T2 lesions at

12 months.

(a) Placebo/delayed treatment, ≥ 2 versus 0–1 T2 lesions at 12 months; (b) IFN β -1a 22 µg SC tiw, ≥ 2 versus 0–1 T2 lesions at 12 months; (c) IFN β -1a 44 µg SC tiw, ≥ 2 versus 0–1 T2 lesions at 12 months. *p*-values indicate differences between patients with differing lesion loads at 12 months within the treatment group. No statistically significant differences were seen in the placebo/delayed treatment or IFN β -1a 22 µg SC tiw groups. Values were calculated with a logistic-regression model with predictor (≥ 2 vs. 0–1 T2 lesions) as a fixed effect; number of relapses within the previous 2 years, age, baseline EDSS score, and baseline burden of disease were independent variables, and *p*-values were calculated for the predictive effect of T2 lesion subgroups.

**p*<0.05.

EDSS: Expanded Disability Status Scale; IFN β-1a: interferon beta-1a; SC: subcutaneously; tiw: three times weekly.