

Switching from inotersen to eplontersen in patients with hereditary transthyretin-mediated amyloidosis with polyneuropathy: Analysis from NEURO-TTRansform

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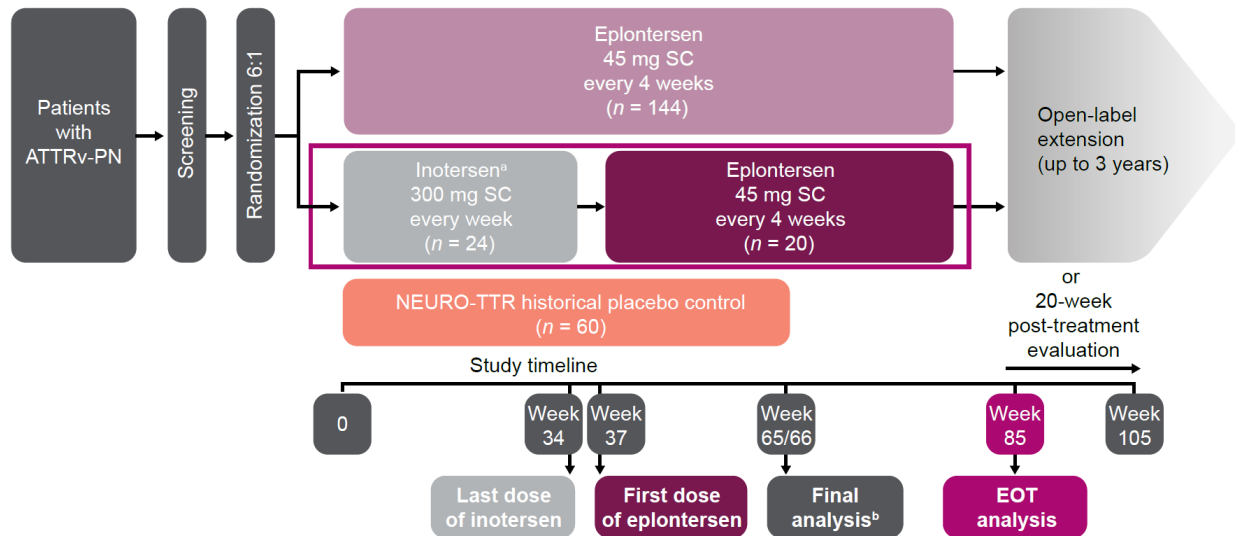
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Supplementary materials

Supplementary Fig. 1. NEURO-TTRansform study design

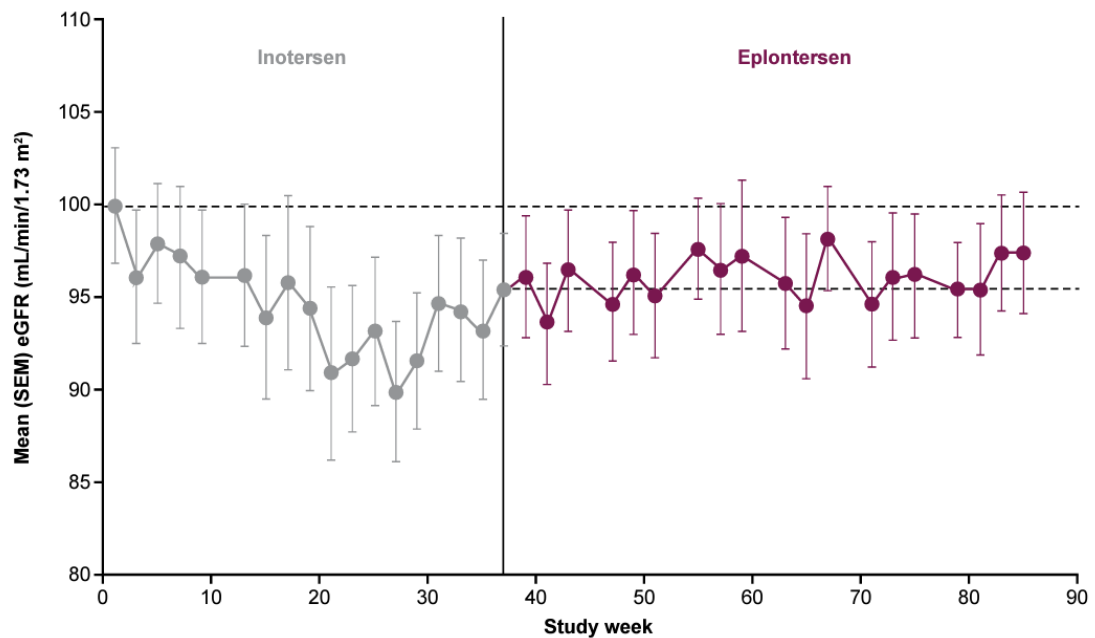


^aInotersen reference group was intended to confirm sufficiently comparable disease progression and treatment response patterns between NEURO-TTR (NCT01737398) and NEURO-TTRansform.

^bFinal analysis refers to final historical placebo-controlled efficacy analysis.

ATTRv-PN, hereditary transthyretin amyloidosis with polyneuropathy; EOT, end-of-treatment; SC, subcutaneous.

Supplementary Fig. 2. eGFR in patients randomized to inotersen who then switched to eplontersen



eGFR, estimated glomerular filtration rate; SEM, standard error of the mean.