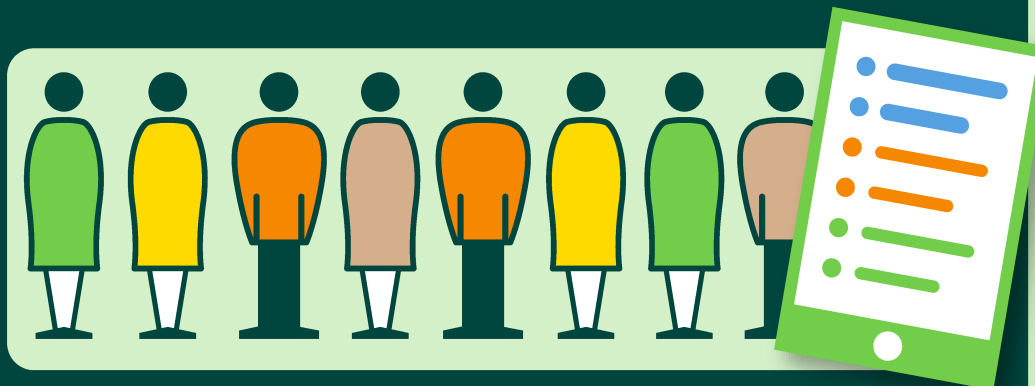


The impact of filgotinib on pain control in the Phase 3 FINCH studies

A post hoc analysis of the **FINCH 1, 2 & 3** studies was performed to evaluate the effects of filgotinib on pain in patients with RA

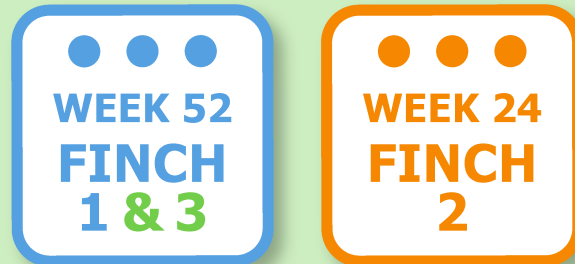


Pain was assessed on a 100 mm visual analogue scale (VAS)



Key findings

Filgotinib reduced pain from as early as **week 2** and responses were sustained throughout the studies up to:



In FINCH 1

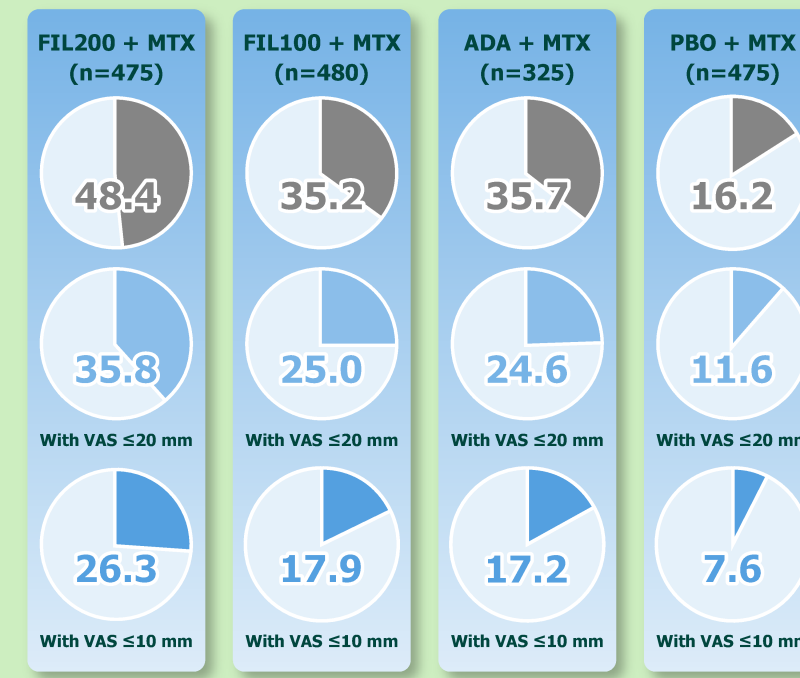
a greater proportion of patients treated with **filgotinib 200 mg** vs **adalimumab** achieved VAS pain ≤ 10 or ≤ 20 mm



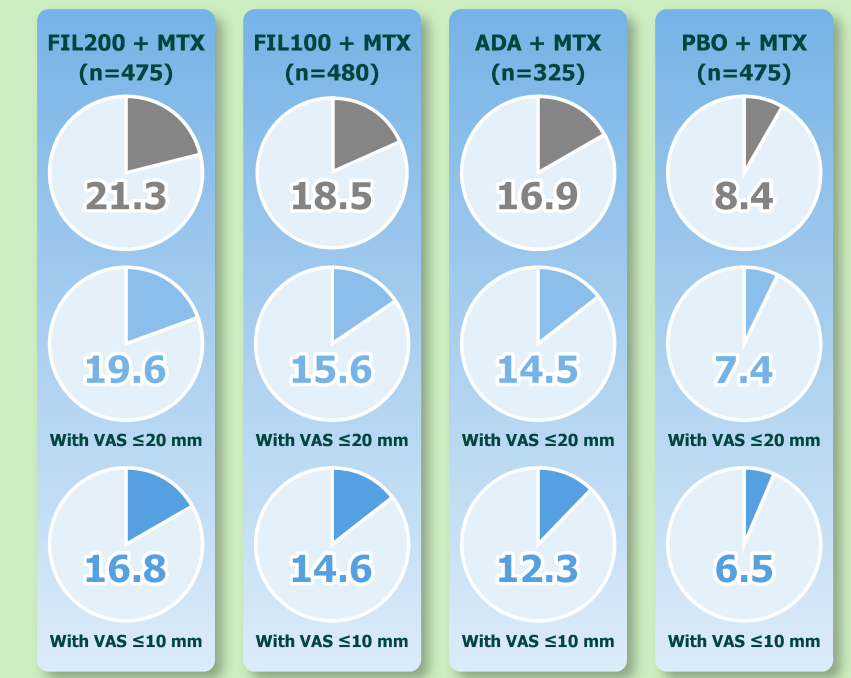
in addition to DAS28-CRP or CDAI remission at **WEEK 24**

FINCH 1

DAS28-CRP Remission (% of patients)



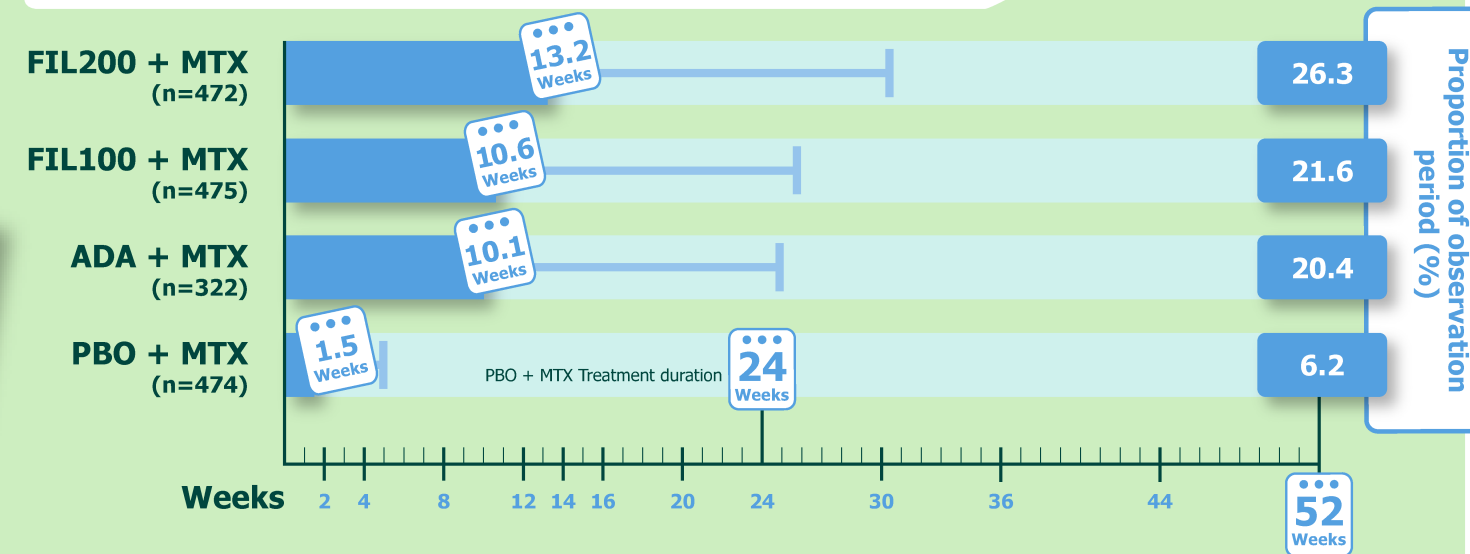
CDAI Remission (% of patients)



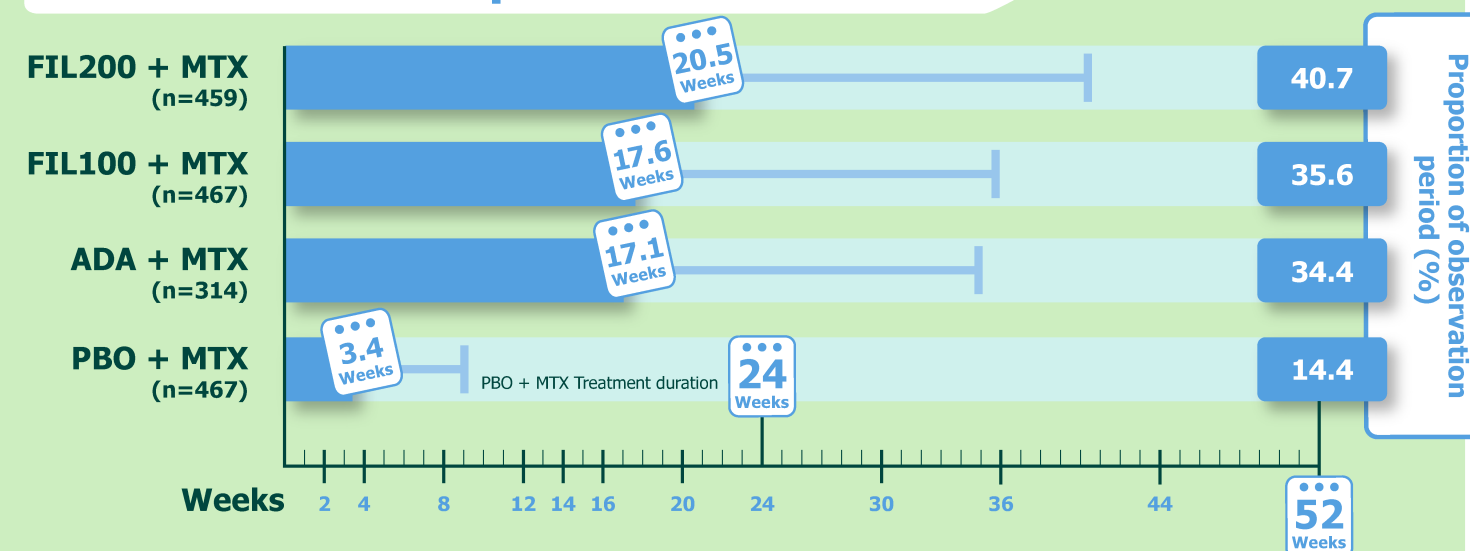
In FINCH 1

the time during which VAS pain was ≤ 10 or ≤ 20 mm was longest with **filgotinib 200 mg** and comparable between **adalimumab** and **filgotinib 100 mg**

Duration of VAS pain score ≤ 10 mm



Duration of VAS pain score ≤ 20 mm

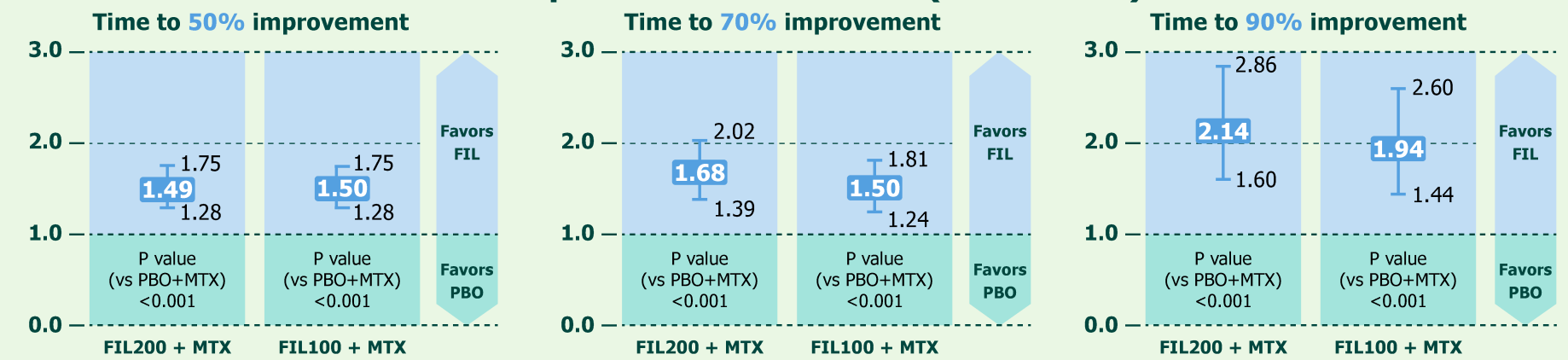


Reductions in pain in FINCH 1

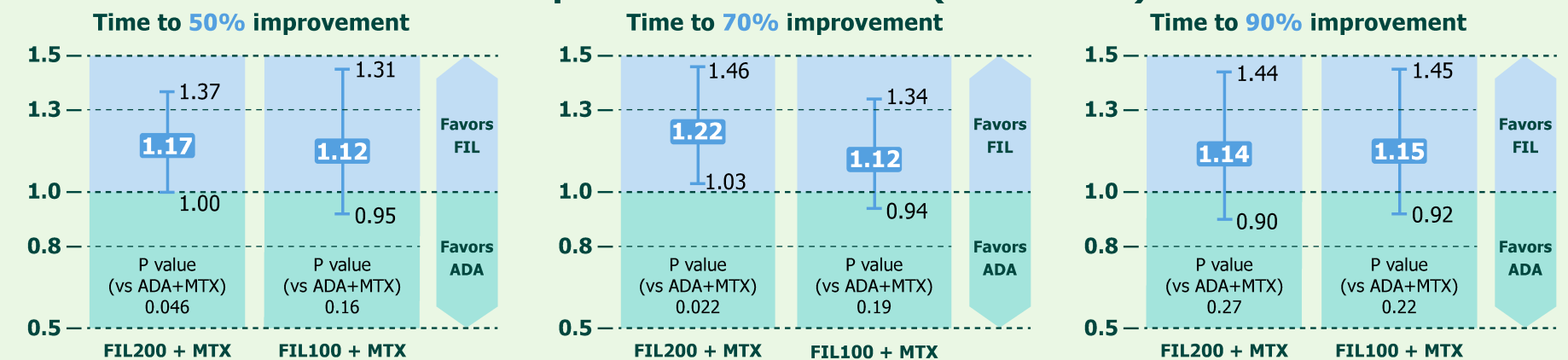
of **30%, 50%, 70% & 90%**

These were all reached earlier with either **filgotinib 200 mg** or **100 mg** than with placebo and were reached earlier with **filgotinib 200 mg** for 30%, 50%, and 70%, but not 90%, than with **adalimumab**

HR comparison of FIL vs PBO (HR 95% CI)



HR comparison of FIL vs ADA (HR 95% CI)



Similar findings were reported for filgotinib in

FINCH 2 & 3