Predictors of Psoriasis Relapse After Guselkumab Withdrawal: An Observational **Analysis of Chinese Patients**

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Background



Guselkumab, a fully human monoclonal antibody that inhibits the biological activity of interleukin-23,1 has received conditional approval in China for the treatment of moderate-to-severe plaque psoriasis based on results from global studies^{2,3}



A post-approval commitment (PAC) study (NCT04914429) previously demonstrated the efficacy of guselkumab in Chinese patients with moderate-to-severe psoriasis,⁴ and was conducted to understand maintenance of response after guselkumab withdrawal and time to psoriasis relapse in a real-world setting



The primary results on maintenance of response following withdrawal of guselkumab are shown in a concurrent presentation

Objective



Here we explore potential predictors of psoriasis relapse after withdrawal of guselkumab treatment in Chinese patients with moderate-tosevere plaque psoriasis

Methods Study design

• This post-PAC, prospective, observational real-world study included Chinese adult patients with moderate-to-severe plaque psoriasis who achieved a Psoriasis Area and Severity Index (PASI) 75 response at the time of receiving the last scheduled dose of guselkumab (Week 44) in the preceding PAC study (index date; Figure 1) and subsequently withdrew from guselkumab treatment

Secondary endpoints

- also assessed
- Associations between patients' demographic and clinical features with clinical outcomes of relapse or non-relapse were explored using Cox proportional hazards models to estimate hazard ratios (HRs) and 95% confidence intervals (CIs)

Key Takeaways

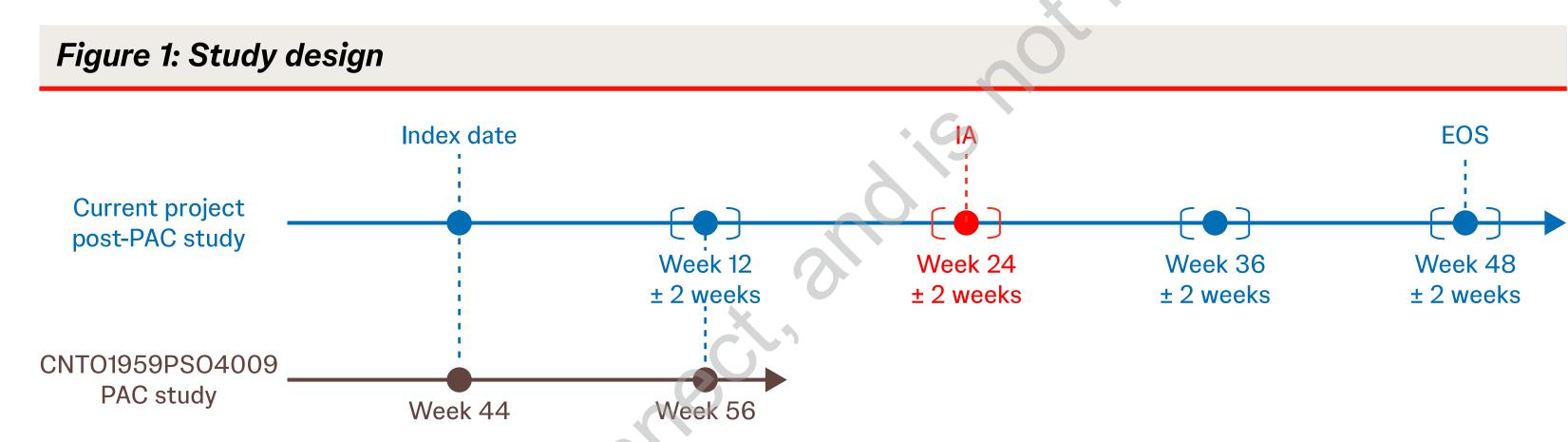


Among Chinese patients with moderate-to-severe plaque psoriasis who discontinued guselkumab treatment, overweight status (compared with normal body mass index) and history of systemic treatment (compared with no prior systemic treatment) were significantly associated with a higher risk of relapse



Changes in DLQI score over time, indicating worsened quality of life among patients who relapse after treatment withdrawal, may signal the need for resuming treatment, and highlight the importance of monitoring DLQI

- We present sociodemographic and clinical characteristics of patients in the relapse (i.e., those starting a systemic treatment for psoriasis after guselkumab withdrawal) and non-relapse groups
- Change in Dermatology Life Quality Index (DLQI) score during the study follow-up period was



Results

Demographic characteristics at the index date

- Of 243 eligible patients included in this post-PAC study, 103 (42.4%) experienced a relapse of psoriasis
- Among patients in the relapse group (**Table 1**):
 - mean (standard deviation [SD]) age was 41.7 (13.0) years
 - 87.4% were male
 - 15.5% had a family history of psoriasis
 - 68.0% previously received systemic treatment; 5.8% previously received a biologic
- Among patients in the non-relapse group (n = 140) (**Table 1**):
 - mean (SD) age was 40.8 (11.8) years
 - 74.3% were male
 - 11.4% had a family history of psoriasis
 - 55.0% previously received systemic treatment; 4.3% previously received a biologic

DLQI scores over time

- At the index date, DLQI scores were available for 80 (77.7%) patients in the relapse group and 107 (76.4%) patients in the non-relapse group
- Mean DLQI scores worsened from the index date to Week 48 in the post-PAC study

EOS, end of study; IA, interim analysis; PAC, post-approval commitment

- In the relapse group:
 - mean DLQI (SD) score at the index date was 1.9 (3.11)
 - mean DLQI (SD) change from the index date was (Figure 2)
 - 0.9 (3.94) at Week 12 8.1 (7.17) at Week 48
- In the non-relapse group:
 - mean DLQI (SD) score at the index date was 3.0 (4.93)
- mean DLQI (SD) change from the index date was (Figure 2)
 - 0.1 (4.89) at Week 12 6.6 (7.70) at Week 48

Effects of demographic and clinical characteristics on psoriasis relapse

- Multivariate Cox regression analysis (**Figure 3**) showed:
 - an increased risk of relapse for patients with overweight status compared with those with normal body mass index (HR 1.813 [95% CI 1.021-3.217]; P = 0.042)
 - a numerically greater, but nonsignificant, risk for patients with obese status compared with those with normal body mass index (HR 1.811 [95% CI 0.991–3.311]; P = 0.054) an increased risk of relapse for patients who received previous systemic treatment compared with those who received no prior systemic treatment (HR 1.915 [95% CI 1.163-3.154]; P = 0.011)
 - a numerically greater, but nonsignificant, risk of relapse for patients with a history of past biologic treatment compared with bio-naive patients (HR 2.443 [95% CI 0.977–6.111]; P = 0.056)

Table 1: Demographic characteristics at the index date, by relapse status

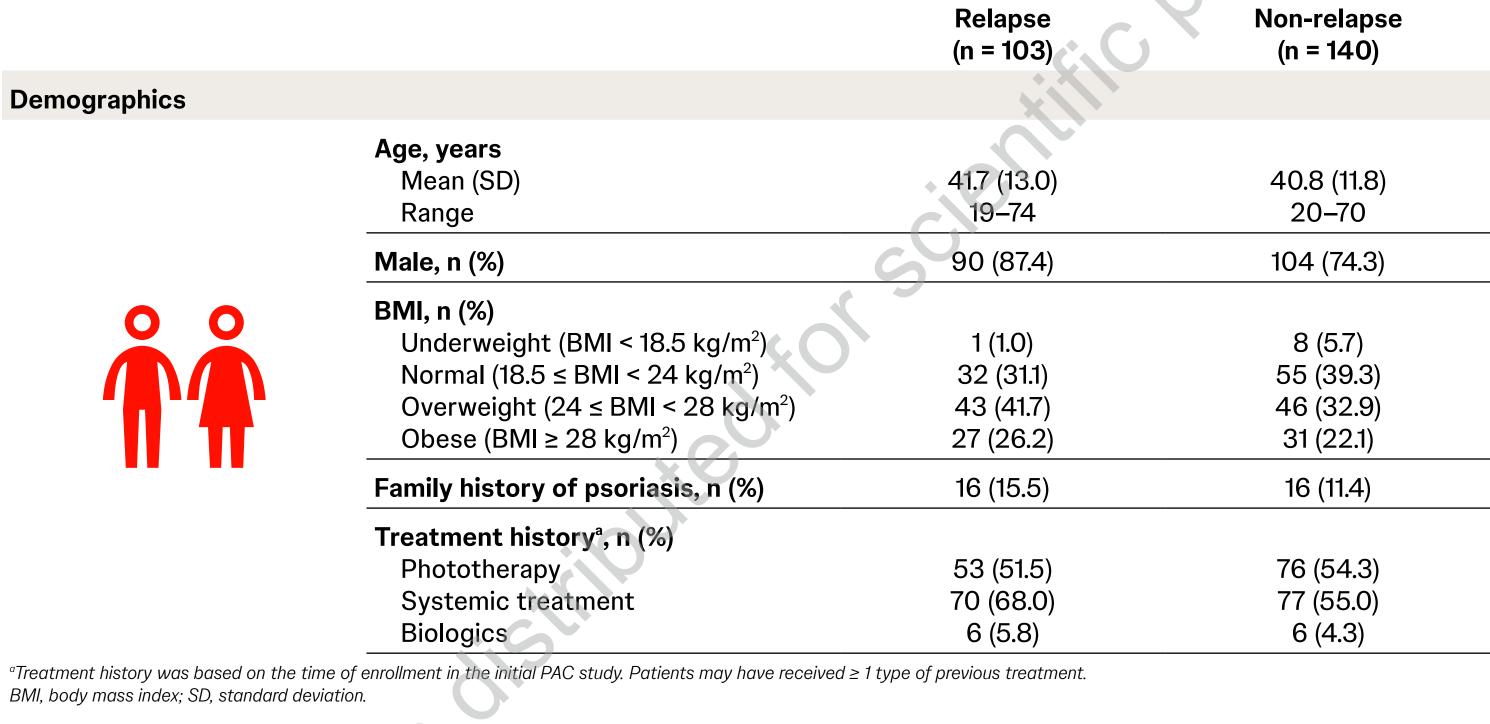
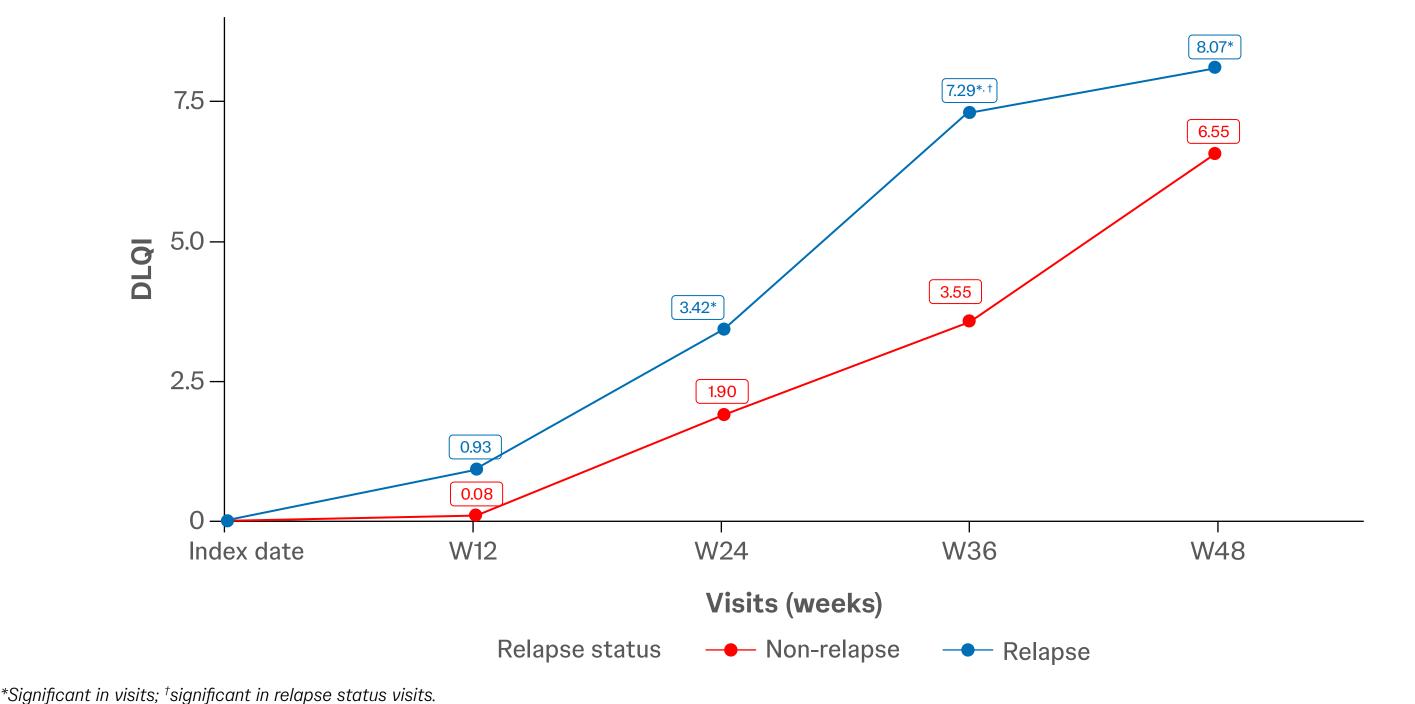
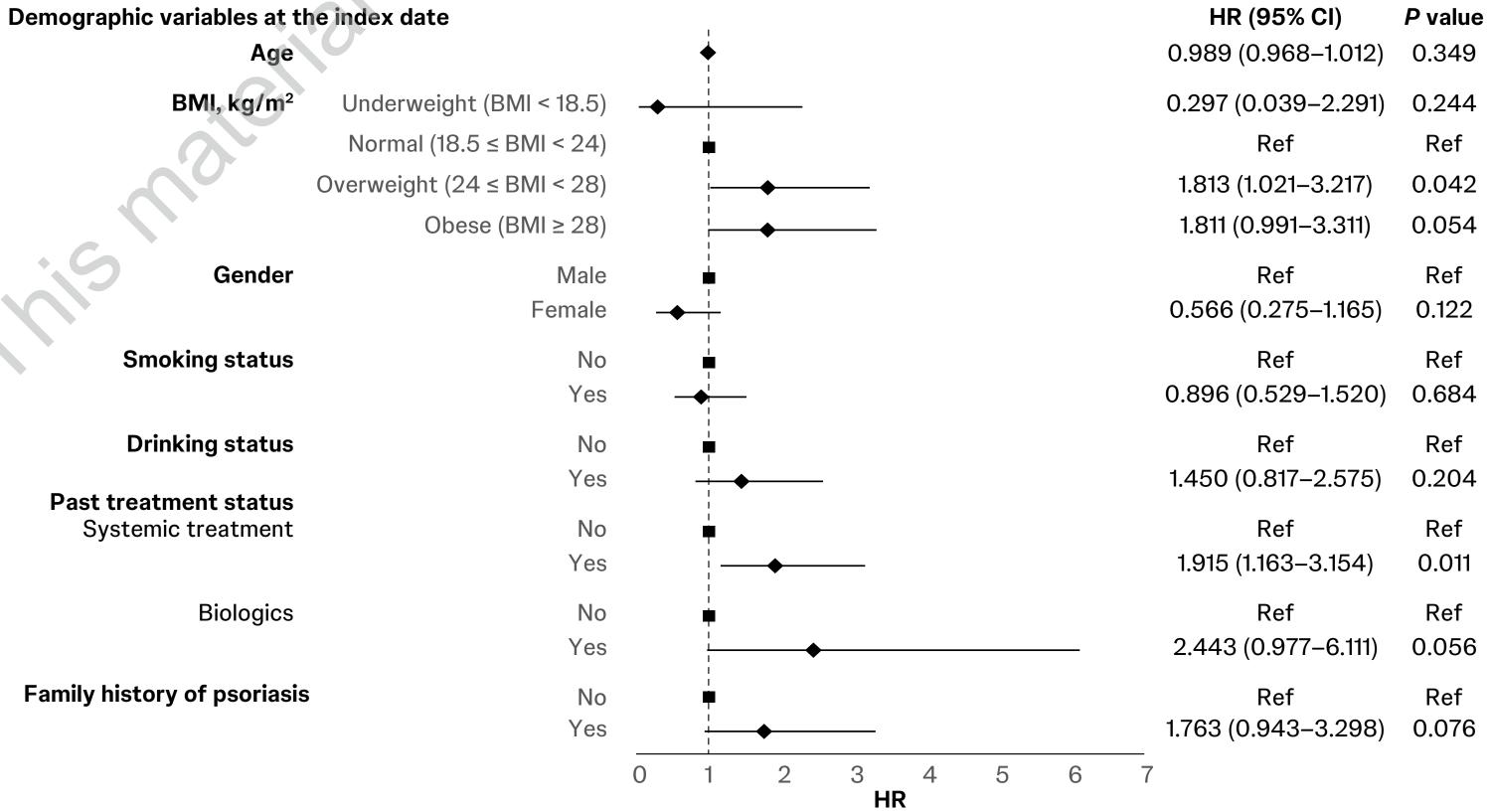


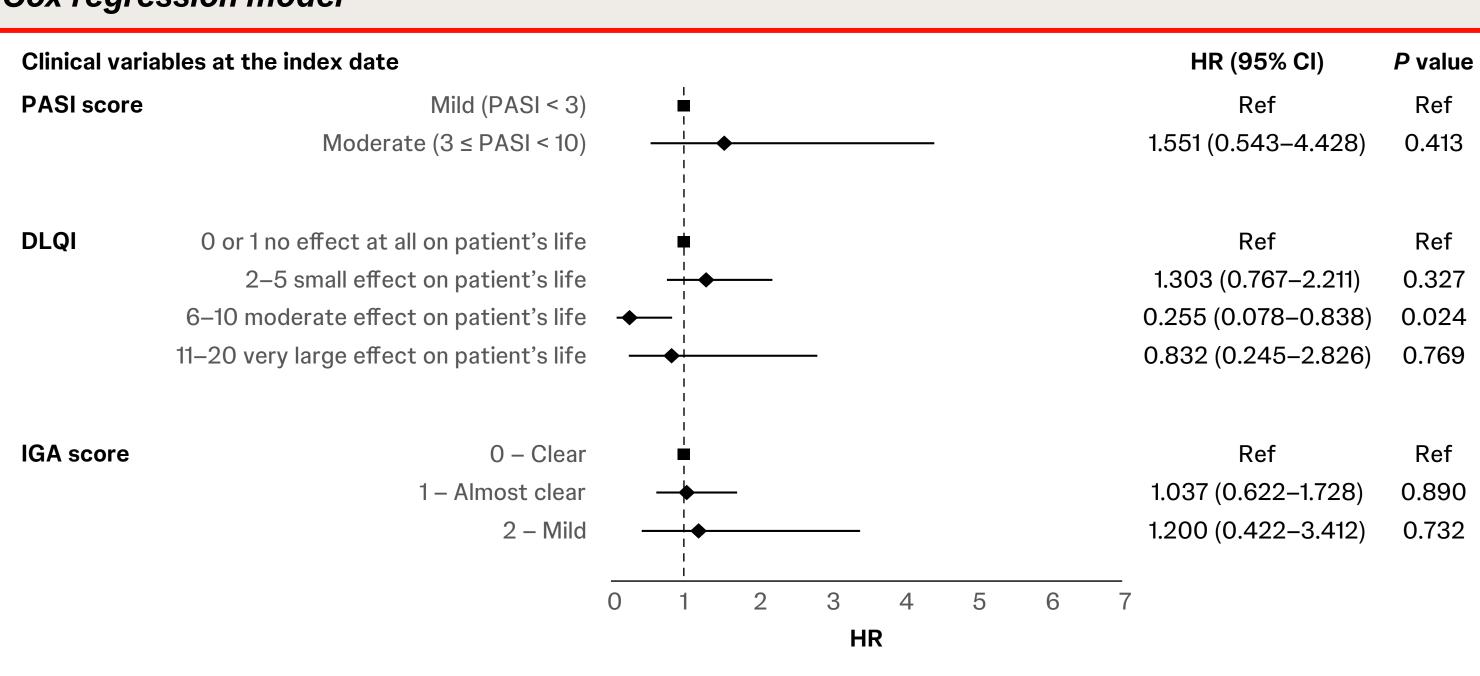
Figure 2: Mean DLQI score over time, by relapse status



DLQI, Dermatology Life Quality Index; W, Week.

Figure 3: Effects of demographic and clinical characteristics on psoriasis relapse – multivariate Cox regression model





BMI, body mass index; CI, confidence interval; DLQI, Dermatology Life Quality Index; HR, hazard ratio; IGA, Investigator Global Assessment; PASI, Psoriasis Area and Severity Index.