# **Cost-Effectiveness of Dupilumab in the Treatment of Moderate-to-Severe Prurigo Nodularis in France**

Allali, N.<sup>a</sup>, Vataire, A.<sup>a</sup>, Sutour, L.<sup>b</sup>, Bahloul, D.<sup>a</sup>, Payan, M.<sup>c</sup>, Thomas, RB.<sup>d</sup>, Kuznik A.<sup>d</sup>, Sambuc, C.<sup>e</sup>, Tavi, J.<sup>a</sup>

<sup>a</sup>Sanofi, Gentilly, France; <sup>b</sup>Sanofi, Cambridge, MA, USA; <sup>c</sup>Sanofi, Netherlands; <sup>d</sup>Regeneron, Sleepy Hollow, NY, USA, <sup>e</sup>VYOO Agency, Paris, France



### INTRODUCTION



Dupilumab is the first biologic approved to treat adults with moderate-to-severe prurigo nodularis (MS-PN) who are uncontrolled with topicals. PN is a subtype of chronic pruritus, characterized by an intense itch and pruriginous lesions.



Cost-effectiveness of dupilumab in MS-PN was assessed by the Economic Commission (CEESP) of the French Haute Autorité de Santé (HAS).

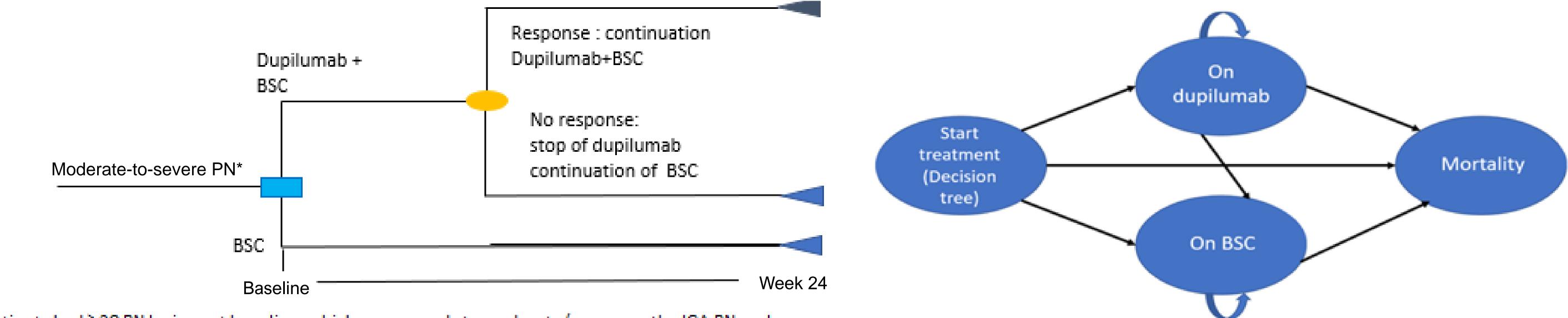
#### OBJECTIVE

The objective of the presented study was to perform a cost-effectiveness analysis of dupilumab to treat MS-PN versus best supportive care (BSC), from a French societal National Health perspective.

The presented approach highlights the value of dupilumab based on effectiveness related to response-year (ry) gained.

## **METHODS**

The 24-Week decision tree followed by a lifetime horizon Markov model structure is showed below:



\*Patients had ≥ 20 PN lesions at baseline, which corresponds to moderate/severe on the IGA-PN scale.

The response to treatment was defined as a minimum improvement of 4 points on the worst-itch numerical rating scale. Costs and healthcare resource utilization were informed by a French patients' survey with PN, with a 2.5% discount rate for costs over a 20-year time horizon. Cost distribution among health states assumed that treatment responders would utilize healthcare resources similarly to patients with mild PN, and non-responders to patients with MS-PN. Sensitivity analysis explored different scenarios, including change in response rate in both treatment arms, cost of disease management, and treatment discontinuation rate.

The model structure was validated by the French Health Authority (HAS).

RESULTS									
	Dupilumab +				Incremental			ICER	
Costs (€) - discounted	138,356					103,043			
Life years in response - discounted	7,844		2,848		4,996			20,623	
			-10 0	000€ -	-6 000 €	-2 000 €	2 000 €	6 000 €	10 000 €
In the base case analysis, over a 20-year time horizon, the treatment with dupilumab versus BSC resulted in a gain of 5		Respor	nse Rate - BSC						
incremental response-years and an incremental cost of		Response Rate - Dupilumab							
€103,043; yielding an ICER (Incremental Cos Ratio) of 20,623 €/ry.	st-effectiveness	Cost - PN non-responder							
	All	All-cause discontinuation - Dupilumab							
The one-way sensitivity analysis shown in the diagram describes the responder rates in each		Cost	- PN responder						
arm as the main drivers of the model.		Lower bound Higher bound							

# CONCLUSIONS

Dupilumab showed a significantly greater response-year gain in patients with MS-PN versus BSC. The ICER/ry in MS-PN was comparable to atopic dermatitis, another dermatologic indication of dupilumab reimbursed in France, demonstrating a similar cost-effectiveness of dupilumab in PN.

This analysis based on cost per response-year is limited by the comparison and the interpretation challenges versus strategies in other therapeutic areas as opposed to the traditional cost per QALY analysis.

#### **CONFLICTS OF INTEREST**

Noémie ALLALI, Jules TAVI, Anne-Lise VATAIRE, Lisa SUTOUR, Donia BALHOUL, Marine PAYAN are/were Sanofi employees and may hold stock and/or stock options in the company. Ryan THOMAS, Andreas KUZNIK are Regeneron employees and may hold stock and/or stock options in the company. Clea SAMBUS is a Vyoo Agency employee and may hold stock and/or stock options in the company.

This study was funded by Sanofi and Regeneron.

**ISPOR Europe 2024 | Barcelona, Spain | 17-20 November**