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COST-EFFECTIVENESS ANALYSIS OF UPADACITINIB AS A TREATMENT OPTION FOR PATIENTS WITH RHEUMATOID ARTHRITIS IN THE KINGDOM OF SAUDI ARABIA

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OBJECTIVE

To evaluate cost-effectiveness of upadacitinib (ts-DMARD) as 1st-line treatment (1L) versus current treatment pathway among patients with RA in the KSA, who had inadequate response to prior conventional-DMARDs and/or biologic-DMARDs from the societal (includes indirect costs) perspective

CONCLUSIONS



Upadacitinib as 1st line treatment for the management of patients with moderate-to-severe RA projects improved health outcomes at lower budget over 10-year time horizon compared to the current treatment pathway.



Upadacitinib may bring significant reduction in healthcareresources utilization in KSA, majorly due to reduced cost of drug-administration, monitoring, hospitalization, surgical-cost, and indirect-costs (productivity loss).



Although the current cost effectiveness analysis doesn't estimate cost savings with adalimumab-originator as 2nd-line treatment for patients with moderate RA, it is perceived that this will not offset the use of adalimumab-originator over adalimumab-biosimilar.

Zeyad AlZahrani speaker and advisory honoraria from Pfizer, AbbVie, Janssen and Roche, Ibrahim Alhomood: speaker and advisory honorarium from Amgen Pfizer, Lilly, GSK, AbbVie, lanssen and Roche, Hana Al Abdulkarim, Hind Almodaimegh aila Abu Esba, Suzan Attar, Waleed Husain, Bedor Al Omari have nothing to disclose, Omneya Mohamed and Yuvraj Sharma are full-time employees of IQVIAAG, Mai Alsaqa'aby is a full-time employee of IQVIA Solutions Saudi Arabia, Ahmed Roshdy and Tharwat Hamad are full-time employee at AbbVie Biopharmaceuticals GmbH and hold company's shares and Ali Anwar is a full-time employee of AbbVie Biopharmaceuticals GmbH and may hold company shares.

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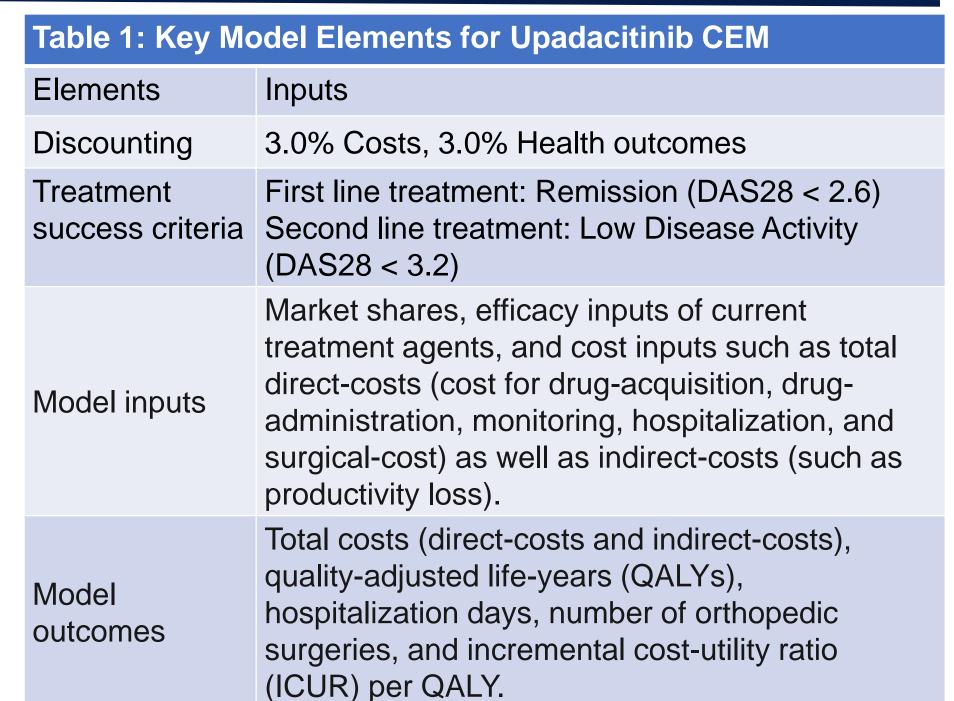
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INTRODUCTION

- Rheumatoid Arthritis (RA) is a chronic inflammatory disease Over a 10-year time horizon in a cohort size of 100 characterized by symmetrical inflammation of synovial joints with higher comorbidity and mortality rates. 1,2
- In the Kingdom of Saudi Arabia (KSA), the prevalence of RA is estimated to be 2.2 per thousand people.³
- The aim of RA management is to achieve remission or reduction of disease activity.^{4,5}
- More recently, the targeted synthetic disease modifying antirheumatic drugs (ts-DMARDs) such as upadacitinib have emerged as an alternative advanced treatment option in RA. Upadacitinib has demonstrated significantly higher rates of remission and low disease activity in all its pivotal trials. 6
- However, the advantages of low disease activity, which might Inputs were retrieved from literature and/or obtained seem self-evident to the rheumatologists at large, are yet to be perceived and still evolving among other stakeholders.^{7,8}

METHODS

- patients, Cost-Effectiveness Model (CEM) analyzed current KSA market for two scenarios:
- Scenario 1: Current treatment pathway (1L: adalimumab-originator/biosimilar, 2L: other biologic-DMARDs/tofacitinib) Versus New treatment pathway (1L: upadacitinib, 2L: adalimumab-biosimilar)
- Scenario 2: Current treatment pathway (1L: adalimumab-originator/biosimilar, 2L: other biologic-DMARDs/tofacitinib) Versus Alternate treatment pathway upadacitinib, adalimumab-originator)
- from interviews with key experts.

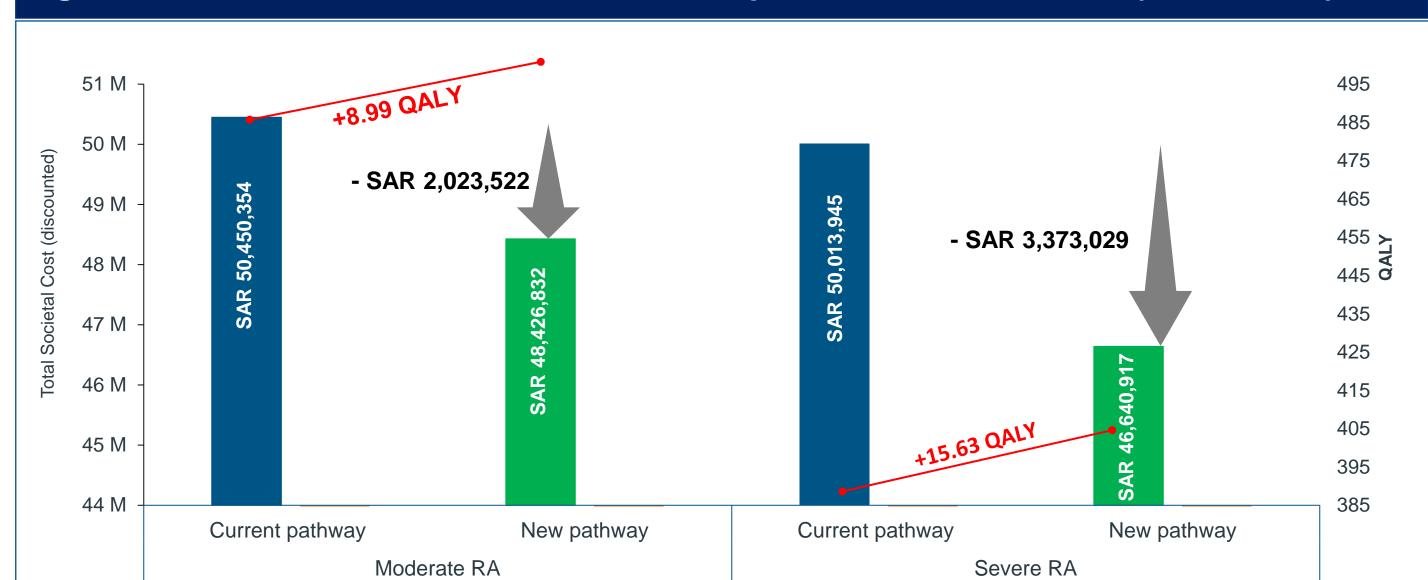


RESULTS

Scenario 1: Current Treatment Pathway vs. New Treatment Pathway

- New treatment pathway among moderate and severe RA patients leads to higher QALYs gain (+8.99 and +15.63) at lower societal-cost (cost difference: -SAR 2,023,522 and -SAR 3,373,029) (Figures 2, Table 2).
- Thus, as 1L, upadacitinib projects 'dominant' ICUR per QALY over current treatment pathway (Table 2).
- Also, new treatment pathway projects reduced hospitalization days (-14.83 and -11.41) and reduced number of orthopedic surgeries (-8.36 and -6.54) among moderate and severe RA patients, respectively (Table 2).

Figure 2: Total Societal Cost and QALY with Upadacitinib - Scenario 1 (Discounted)



QALY: Quality-adjusted Life-year; RA: Rheumatoid Arthritis; SAR: Saudi Riyal					
Table 2: Health Outcomes with Upadacitinib - Scenario 1 (Discounted)					
Health outcomes	Current Treatment Pathway	New Treatment Pathway	Incremental		
Moderate RA					
QALYs	479.04	488.03	+8.99		
Hospitalization Days	239.73	224.90	-14.83		
No. of Orthopaedic surgeries	122.65	114.29	-8.36		
	ICUR per QALY		Dominant		
Severe RA					
QALYs	386.77	402.40	+15.63		
Hospitalization Days	217.78	206.37	-11.41		
No. of Orthopaedic surgeries	111.11	104.57	-6.54		
ICUR per QALY			Dominant		

Scenario 2: Current Treatment Pathway vs. Alternate Treatment Pathway

- Alternate treatment pathway projects 'dominant' ICUR per QALY for patient with severe RA (QALY gain: +15.63, societal cost difference: -SAR 164,536) (Figure 3, Table 3).
- However, for moderate RA, it is associated with additional societal cost of SAR 1,255,696 for improved QALY (+8.99) over current treatment pathway (ICUR per QALY: SAR 139,742) (Figure 3, Table 3).
- Overall, the alternate treatment pathway projects reduced hospitalization days (-14.83 and -11.41) and reduced number of orthopedic surgeries (-8.36 and -6.54) among moderate and severe RA patients, respectively (Table 3).

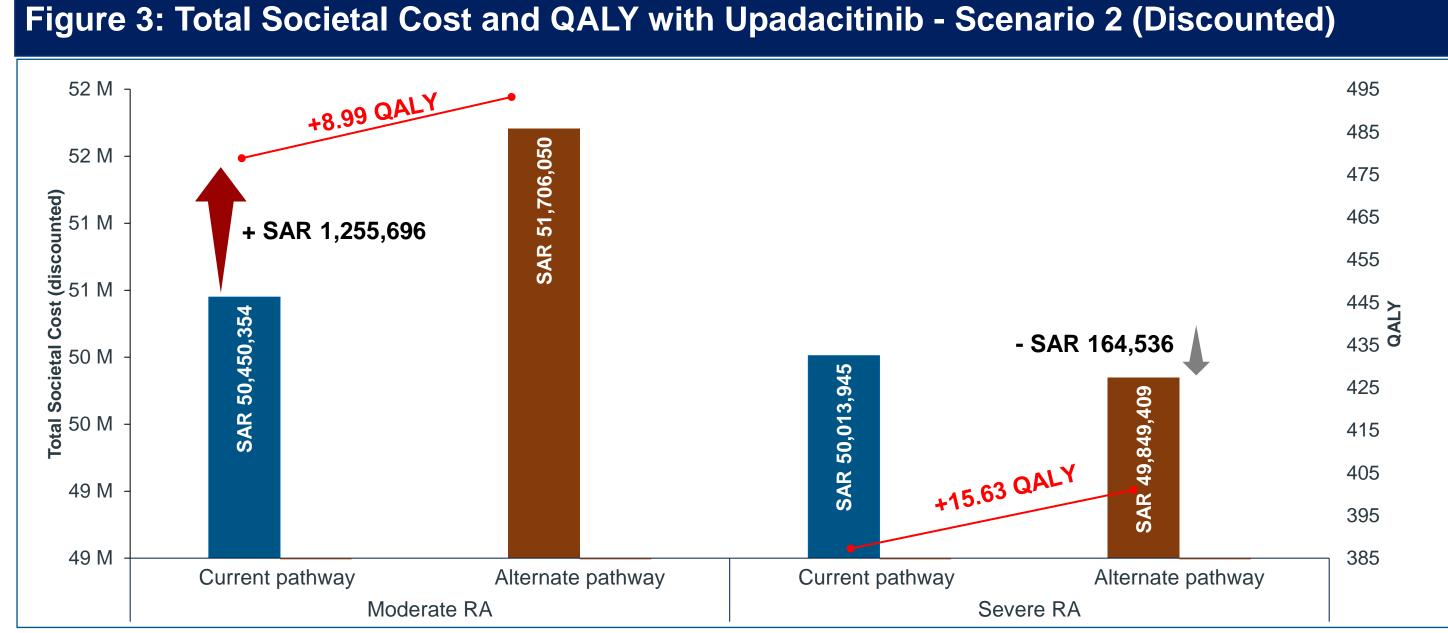


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ICUR: Incremental cost-utility ratio; QALY: Quality-adjusted Life-year; RA: Rheumatoid Arthritis

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