

PEMBROLIZUMAB AND ATEZOLIZUMAB AS POSSIBLE **EQUIVALENT FIRST-LINE THERAPEUTIC ALTERNATIVES IN PD-L1-EXPRESSING TRIPLE-NEGATIVE BREAST CANCER**



M.D RIVAS RODRÍGUEZ, A.GIL GARCÍA, A. ROJAS ALBARRÁN, M. GRAGERA GÓMEZ, H. VÁZQUEZ VELÁZQUEZ, J.F RANGEL MAYORAL University Hospital Complex of Badajoz, Hospital Pharmacy, Badajoz, Spain Marilolirr.2612@gmail.com

BACKGROUND

Recent studies have established the influence of the immune system on disease progression in triple negative breast cancer (TNBC) patients.

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To determine if pembrolizumab and atezolizumab can be considered equivalent first-line

therapeutic alternatives (ATE) by using a common comparator, for patients with locally

recurrent unresectable or metastatic unresectable TNBC in adults whose tumors express PD-L1

and who have not received prior chemotherapy

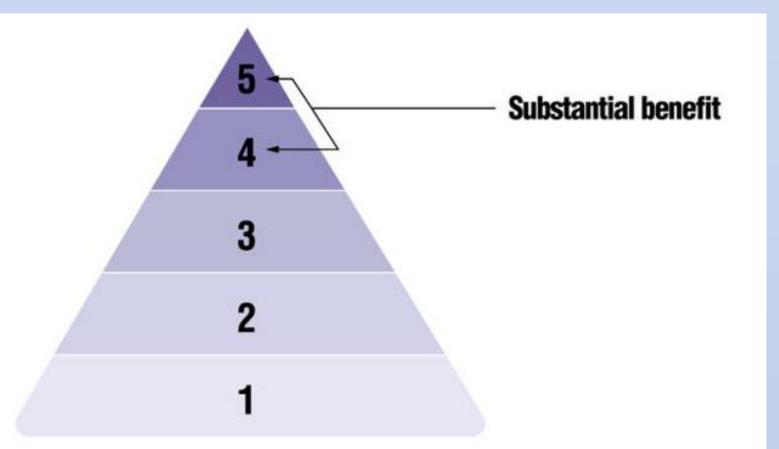
MATERIALS AND METHODS

Bibliographic search Phase III randomised clinical trials of first-line treatments for TNBC.

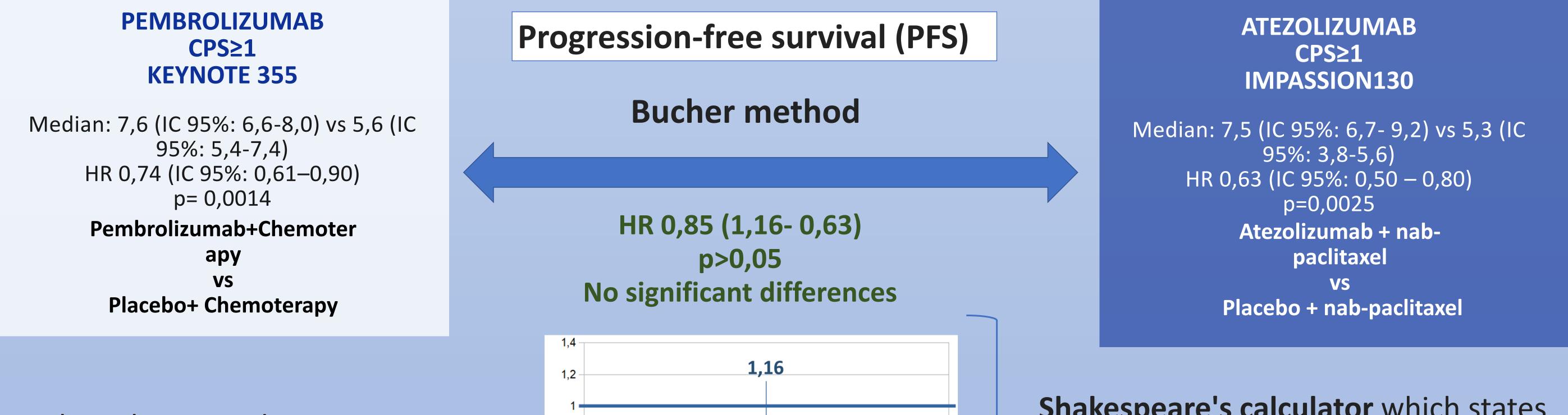
- The indirect comparison was performed with the **Bucher method**.
- The variable selected to determine clinical equivalence was progressionfree survival (PFS).
- The maximum acceptable difference as a clinical non-inferiority standard Delta (D), and its inverse were set at 0.65 and 1.54, respectively. They were established by ESMO-Magnitude of Clinical Benefit Scale.

0,8

0,6

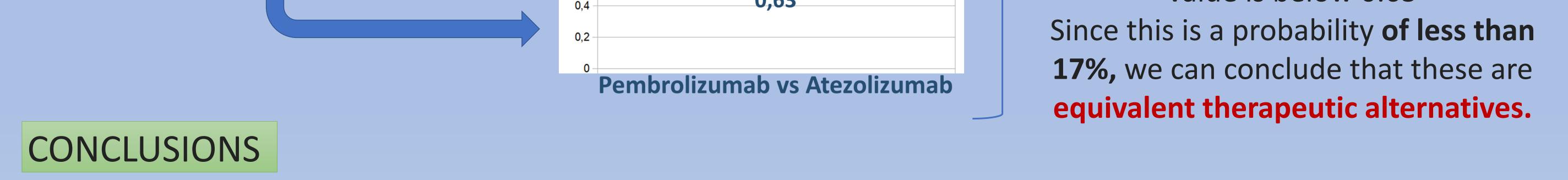






Delta value according to ESMO guidelines 0,65

Shakespeare's calculator which states that there is a **4.25%** probability that the value is below 0.65



0,63

Pembrolizumab and atezolizumab could be considered ATE, however, recent studies such as the Impassion 131 bring a great deal of uncertainty to this determination.

