




Primary endpoints: OR at Day 28, including CR and PR; DOR at Day 56; cumulative incidence of cGvHD at 1 year; OS at 1 year; cumulative incidence of FFS at 1 year; median OS and FFS

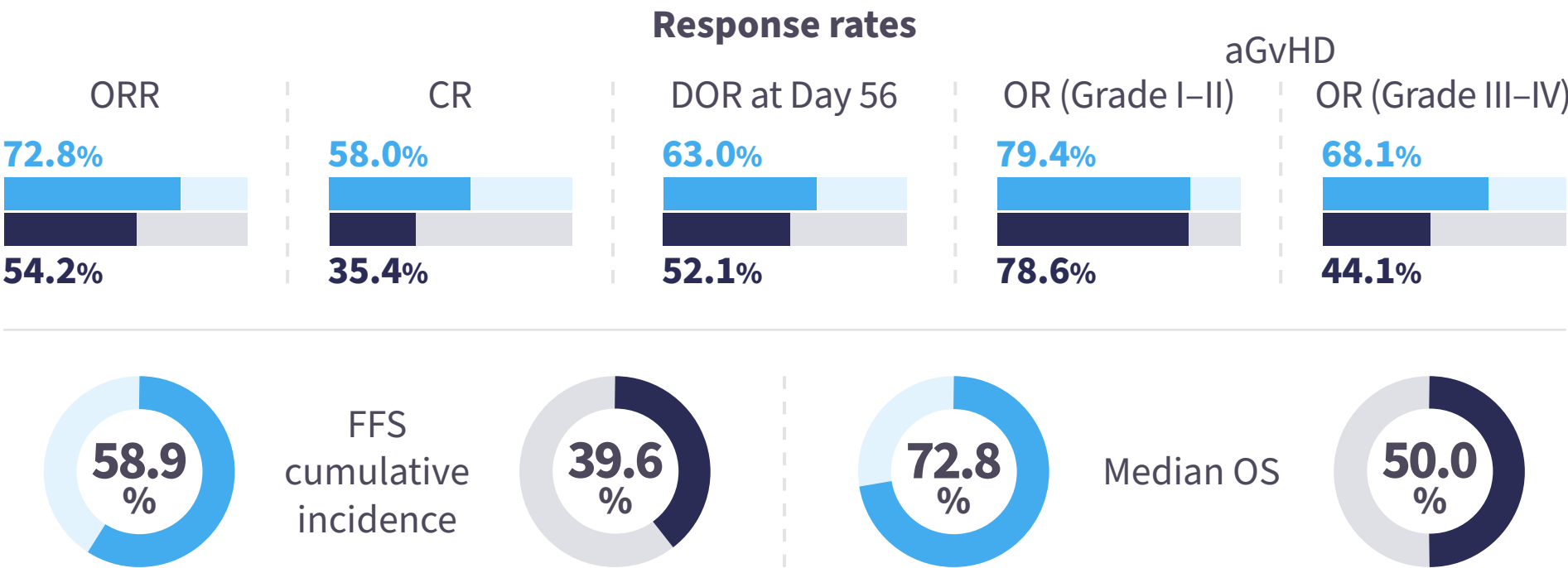
Eligibility: Adult patients with SR-aGvHD

TRIAL DESIGN

Drug 1  Ruxolitinib	Drug 2  Ruxolitinib	Drug 3  Basiliximab
5 mg/bid (if platelet count is $20 \times 10^9/L$)	10 mg/bid (If platelet count is $> 30 \times 10^9/L$)	20 mg once Days 1, 4, 8, 15, and 21.
for 3 years	for 0–3 years	for 21 days

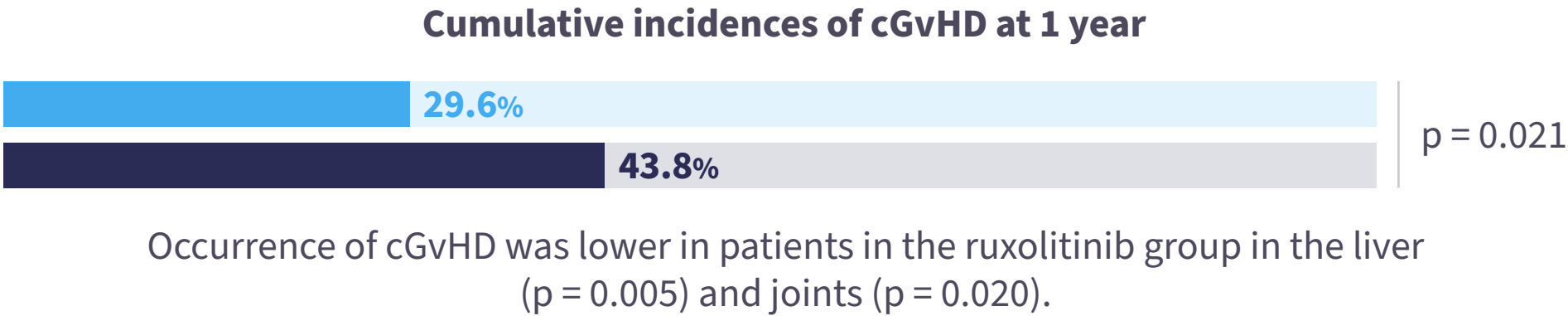
Ruxolitinib
Basiliximab

EFFICACY



Ruxolitinib
Basiliximab

SAFETY



Ruxolitinib showed favorable results in the treatment of SR-aGvHD and reduced the incidence of cGvHD compared with basiliximab.

Abbreviations: aGvHD, acute graft-versus-host disease; bid, twice a day; cGvHD, chronic GvHD; CR, complete response; DOR, duration of response; FFS, failure-free survival; OR, overall response; OS, overall survival; PR, partial response; SR-aGvHD, steroid-refractory aGvHD.

Lui, et al. *Ann Hematol.* 2023;102(10):2865-2877. DOI: 10.1007/s00277-023-05361-9