



# Bexicaserin (LP352) Open-Label Extension (OLE) Interim Analysis

JUNE 10, 2024

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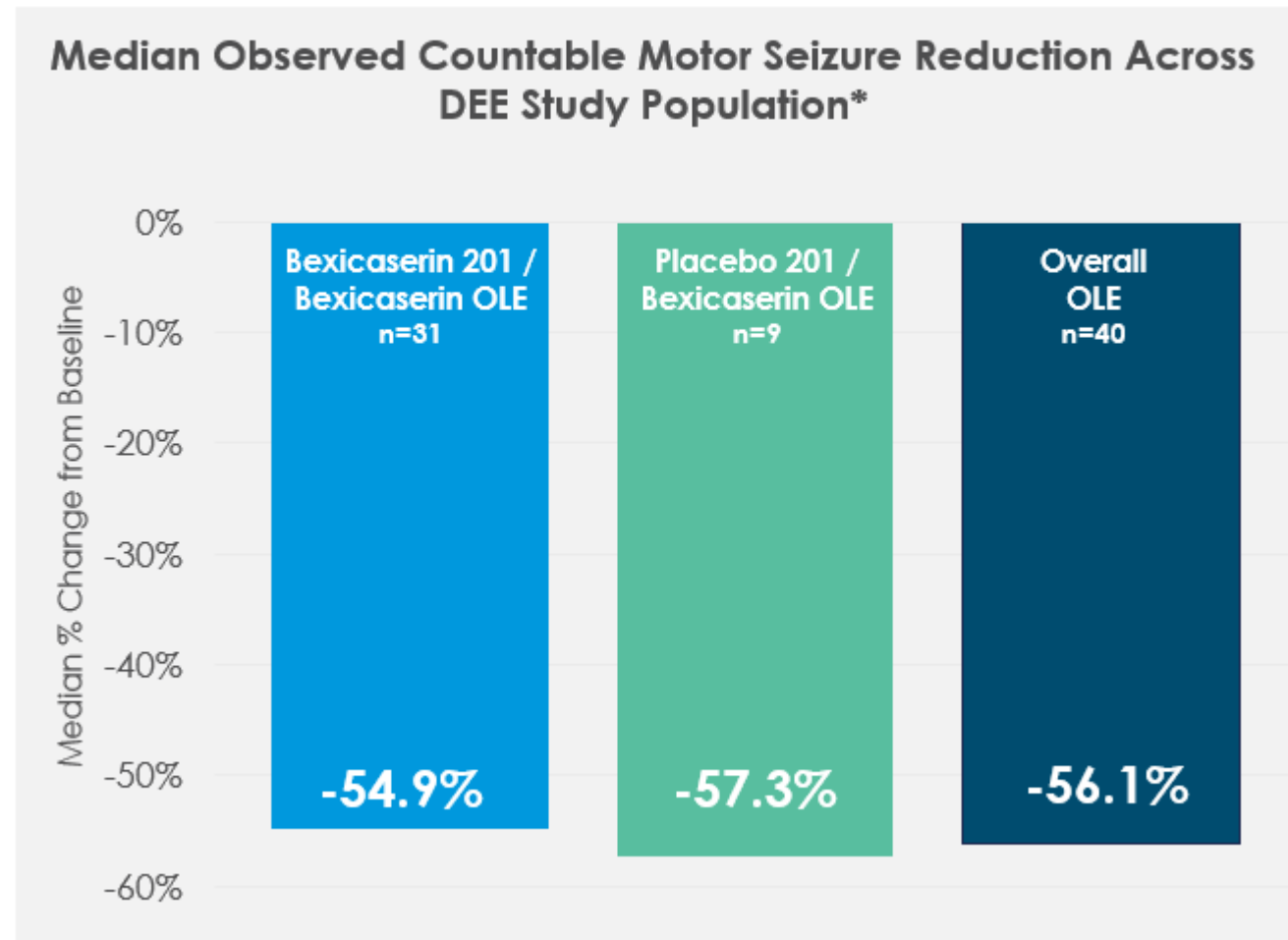
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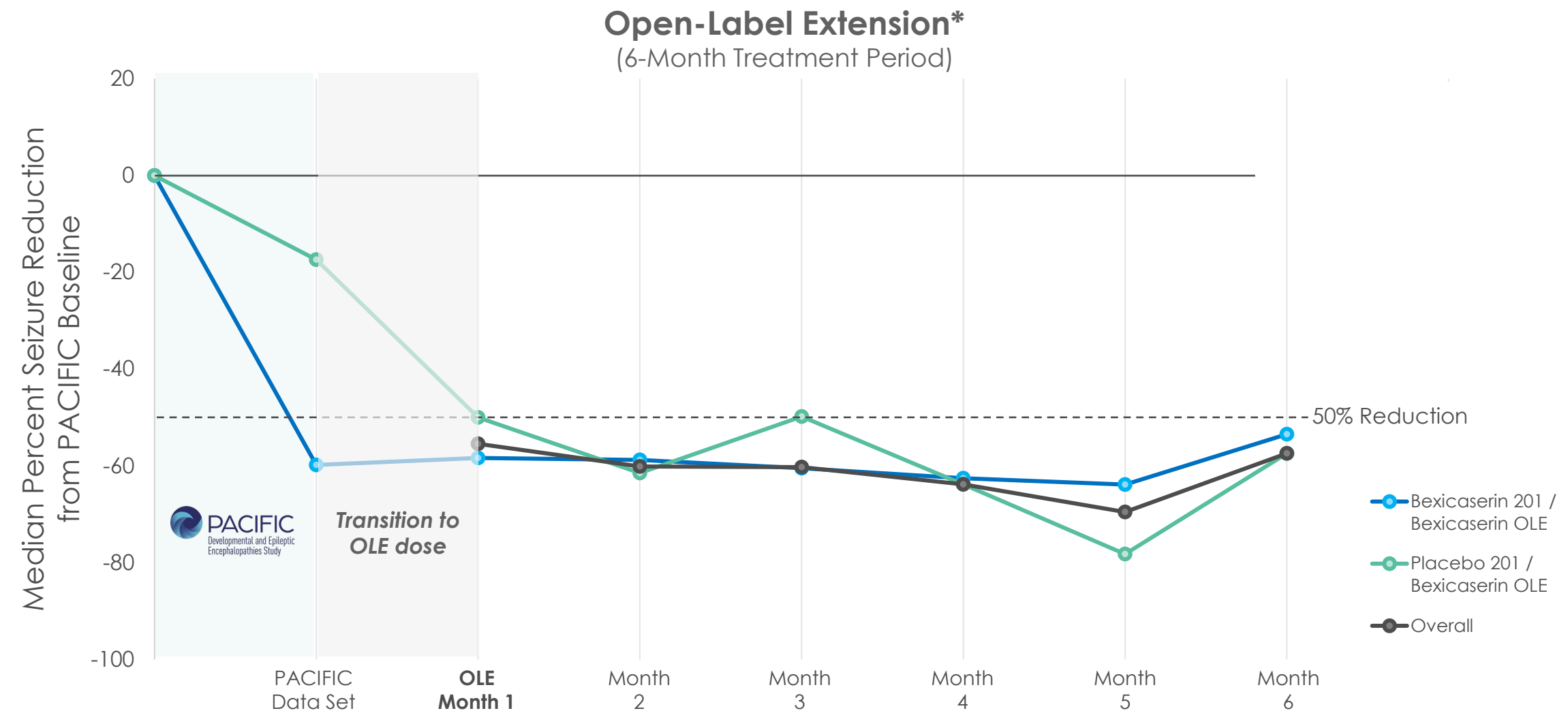


# Summary: Interim Analysis from Bexicaserin (LP352) OLE Study

- **100% of PACIFIC completers continued into OLE** (95.1% in OLE at 6-months)
  - PACIFIC completers n=41 (DS=3, LGS=20, DEE Other=18)
- **Sustained response** over an approximate 6-month treatment period
- **Favorable safety and tolerability** results observed
- **PACIFIC (201 Study) Placebo participants:**
  - **All successfully titrated up** and entered maintenance phase of the OLE
  - **Motor seizure reduction** consistent with bexicaserin efficacy observed in PACIFIC
- **Global Phase 3 Program on track** to initiate later this year



# Bexicaserin (LP352) Median Observed Countable Motor Seizure Reduction in OLE



# Summary: Bexicaserin (LP352) Safety and Tolerability in the OLE Interim Analysis

	Overall (N = 41)
Parameter	n (%)
Safety Set	41 (100)
Full Analysis Set	40 (97.6)

- SAEs in the overall group were comprised of pneumonia, pneumonia bacterial, change in seizure presentation, seizure, and agitation
- One participant discontinued from the study due to the adverse event of lethargy (2.4%) during the titration period
- One participant discontinued from the study by the withdrawal of consent (2.4%) during the first month of maintenance
- **Favorable safety and tolerability results observed**

	Overall (N = 41)
Preferred Term*	n (%)
Upper respiratory tract infections	5 (12.2)
COVID-19	3 (7.3)
Pneumonia	3 (7.3)
Sinusitis	3 (7.3)
Seizure	3 (7.3)
Decreased appetite	3 (7.3)

