

PRALUENT: A POWERFUL* PCSK9i THAT TICKS ALL THE BOXES¹⁻⁴



- ✓ **Intensive, fast, and sustained LDL-C reduction^{1,2*}**
- ✓ **The only LLT beyond statins associated with a reduction in all-cause mortality^{2,†}**
- ✓ **Demonstrated to get patients to LDL-C goal^{3†}**
- ✓ **Established, long-term safety profile¹**
- ✓ **Unique[‡] once-monthly pen for your patients requiring >60% LDL-C reduction^{2,4}**

*62.7% LDL-C reduction compared to placebo at 4 months in ODYSSEY OUTCOMES trial.¹ †Based on a summary of ten phase 3 trials (five placebo-controlled, five ezetimibe-controlled) in high and very high-CV-risk patients.² ‡In patients following an ACS event in the ODYSSEY OUTCOMES trial, sustained at 54.7% relative to placebo.¹

ACS = acute coronary syndrome; **CV** = cardiovascular; **LDL-C** = low-density lipoprotein cholesterol.

References:

1. Schwartz GG, Steg PG, Szarek M, et al. Alirocumab and cardiovascular outcomes after acute coronary syndrome. *N Engl J Med.* 2018;379(22):2097-2107.
2. PRALUENT (alirocumab) Summary of Product Characteristics.

To report any side effect(s): Saudi Arabia: The National Pharmacovigilance and Drug Safety Centre (NPC): SFDA call center: 19999, E-mail: npc.drug@sfga.gov.sa , Website: <https://ade.sfga.gov.sa/>

Full Prescribing Information is available upon request:

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