

## **CLIENT SITUATION**

A top-20 global biopharmaceutical company was developing two unrelated assets requiring companion diagnostics (CDx) and was seeking insight into the eligible populations and access barriers in different markets.

## PRMA CONSULTING SOLUTION

## Targeted research provided insight into the HTA issues to consider

- Building on previous extensive research on CDx-drug pairs, we reviewed and summarized country-specific regulatory and P&R processes and data requirements for CDx market access.
- A range of CDx-drug pairs relevant to the client's products were carefully selected.
  Case studies were developed to investigate HTA issues, the role of clinical guidelines, CDx pricing, and CDx adoption processes.
- Evidence requirements to support market access for both the drugs and the CDx were identified.

## Critical insights supported the client's market access strategy

- From this, the client had a concise description of the processes and challenges in developing CDx-drug pairs, including critical insight from key internal stakeholders in each scope market. This allowed them to:
  - optimize and rationalize budget allocation
  - change project prioritization
  - develop recommendations and key questions for affiliate teams.
- A robust framework was developed that could be applied to future CDx-drug pairs.
- The client gained a clear understanding of how CDx could support their assets.



Our cross-functional team found the evaluation very useful. I want to thank everyone for delivering this.

Director, Global Health Economics



