LAUNCH PLANNING & 360° REIMBURSEMENT
Build strategic imperatives for clinical and commercial planning that address access, utilization, and reimbursement potential based on the identification of hurdles and competitive landscape.

- Understand implications of varying launch and approval dates for new indications with specific regard to price and utilization.

- Assess context for formulary review with respect to unmet needs, current environment, competitive landscape, etc.

- Understand current and future payer landscape, likely formulary access process, and access hurdles from the perspective of P&R decision-makers from key plans.

Environmental Assessment & Scenario Playbooks

- Presents the current clinical and reimbursement landscape including:
  - Treatment approaches
  - Treatment satisfaction
  - Unmet needs
  - Develop launch scenarios for antimicrobial

Decision-Maker Interviews

- Payer Panel consisting of top-tier decision-makers who lead and set policy for formulary access and price negotiation

Market-Specific Strategic Implications

- Offers insight on key implications and strategic recommendations to optimize P&R positioning and market access

<table>
<thead>
<tr>
<th>Conclusions</th>
</tr>
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<tbody>
<tr>
<td>1. In either scenario, market access is not hindered due to perceived value for NP</td>
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<td>2. Establish NP benchmark price without the NP indication except in Italy</td>
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<td>3. Launch with cSSSI indication and robust NP data file in market 1, market 2; wait for NP indication in country x</td>
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Establish NP benchmark price without the NP indication except in Italy.
Launch with cSSSI indication and robust NP data file in market 1, market 2; wait for NP indication in country x.
Quantitative Launch Pricing Research through different price finding methodologies seeking to identify price response and profit functions to determine the profit-maximizing price

**Attributes Validation**
- Through qualitative discussions with payers and KOLs, identified the attributes, levels, and potential pricing points that have impact on physician's prescribing behavior

**Quantitative Target Profile Substantiation**
- Discrete choice modeling was applied to best understand prescribing decision-making based on behavioral response (rather than preference judgment)

**Market Simulation and Pricing Strategy**
- For each scenario, unique demand curves were generated and incorporated into a market simulation model
- The simulator allows the client full insight into the influence of specific product attributes on price and demand

**Client Need**

Understand the impact of a competitor entering in the market at 50% off the current price

Gauge the potential share of various efficacy, toxicity, dosing frequency, and cost per cycle performance scenarios associated with the licensing of an oncologic

Derive value-based price optimization curve and price elasticity from qualitative and quantitative inputs

<table>
<thead>
<tr>
<th>Treatment of Condition</th>
<th>Attribute A (Portability)</th>
<th>Level 1: Portable</th>
<th>Level 2: Not portable</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Attribute B (Length of Administration)</td>
<td>Level 1: 5 minutes</td>
<td>Level 2: 20 minutes</td>
</tr>
<tr>
<td></td>
<td>Attribute C (FDA Approval)</td>
<td>Level 1: Approved for children 1 to 4</td>
<td>Level 2: Not Approved for children 1 to 4</td>
</tr>
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</table>
Understanding the attributes and nuances of the product which attain favorable versus unfavorable reimbursement status across various markets and across key health plans.

Identify the therapeutic areas where products launched with a non-reimbursement strategy are most likely to succeed.

Evaluate the implications of identified product characteristics/factors on the uptake of non-reimbursed products in comparison to reimbursed competitors.

Provide a strategic decision-making tool to help the client to assess future reimbursement strategies for its portfolio.

Product Evaluation:
- Identify and define values for the factors that are most relevant to succeed with a non-reimbursement strategy.
- Conduct historic product performance analysis to identify the negative and positive factors to influence the uptake of a non-reimbursed product.
- Perform correlation analysis to clearly delineate the factors most important when determining a successful non-reimbursement strategy.

Opportunity Assessment Matrix:
- Applying therapeutic area and macroenvironmental relevant factors and build an opportunity matrix to identify which therapeutic areas are most suitable for products launching with a non-reimbursement strategy in each country.

Interactive Assessment Tools:
- Strategic decision-making tools are provided to clients to perform future assessments for their pipeline when considering the launch with a non-reimbursement strategy in each individual plan analyzed.
A small-sized biopharmaceutical company with a focus on antivirals for immunocompromised patients approached the Certara US team. They had developed a Phase II pipeline agent for treatment of adenovirus in allogeneic stem cell transplant recipients. However, their product development strategy faced a major challenge: there currently is a major gap in multicenter studies of incidence and management of adenovirus infection in allogeneic hematopoietic cell transplant (allo-HCT) recipients.

The project demonstrates Certara’s ability to take a customized approach on highly complex path-to-market challenges, and leverage strong client relationships to tailor content that creates unique value and positively impacts external stakeholders. The client chose Certara given our extensive expertise in rigorous primary research processes (e.g. physician procedures, recruitment, contracting, fees, IRB process) and our ability to navigate a complex US system with an advanced approach to capture physician insight. The results of the research led to a transformation in the client’s value proposition, but also provided their key stakeholders (transplant physicians) with helpful insights about patient management. For example, many of them thanked the Certara teams they engaged with for the opportunity to compare how their center performs on a national level. Our engagement with this client led to a multi-year partnership involving US and ex-US markets.
A large pharma company approached Certara to develop a plan to derive evidence-based strategies to proactively ensure franchise success through an evolving market.

**Key Objective**

Achieve franchise profitability goals through 2015 as the market and client's franchise face challenges from new competitors, patent expirations, and changes in care delivery models.

**Certara Methodology**

- Comprehensive secondary research and gap analysis informed the research plan.
- Primary research of internal stakeholders (n=18) and payer customers (n=22) provided intelligence for an updated market access landscape.
- Using additional internal expertise collected through an account team survey (n=62) and war game exercises (n=40), a strategic market access plan was developed.

**Results & Deliverables**

- Recommendations in the Strategic Market Access Plan have been executed.
- Client’s principal product has retained market leadership in most plans and geographies through the launch of a successor compound.
A client requested evidence-based guidance of the current landscape and target product profile to identify the value drivers and advance the asset in product development.

1. DESCRIPTION
   A client requested evidence-based guidance of the current landscape and target product profile to identify the value drivers and advance the asset in product development.

2. KEY OBJECTIVE
   - Understand current patient journey and treatment landscape
   - Review process of novel therapies within the neonatal intensive care unit (NICU) and the billing and reimbursement process in the NICU
   - Assess the clinical research designs of a novel treatment of an orphan disease to assist in guiding the commercial development and clinical teams

3. Certara METHODOLOGY
   - In-depth interviews with 20 respondents in the US including pediatric ophthalmologists, retinal specialists, NICU medical directors, NICU nurse managers, NICU pharmacists, hospital coding/reimbursement experts, and payers

4. RESULTS
   - Description of types of NICUs in US, key decision makers, adoption of new treatments and their fit into current processes
   - Map of reimbursement process within the hospital and NICU by payer type
   - Patient journey including patient characteristics, screening, treatment and outcomes of this rare disease
   - Comprehensive product assessment by respondents and expected utilization by HCPs
   - Development of a cost model and price recommendation
   - Findings were used by the client to plan the Phase III clinical trial
1. **DESCRIPTION**

A large pharma company requested a plan to derive evidence-based strategies to proactively ensure franchise success through an evolving market.

2. **KEY OBJECTIVE**

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4. **RESULTS & DELIVERABLES**

- Recommendations in the Strategic Market Access Plan have been executed.
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Many of the recommendations in the Strategic Market Access Plan have been executed:

**Actionable strategic recommendations from the Strat-MAP**

- Overarching Market Access Strategic Initiative #1: Execute A Product Re-Vitalization
- Overarching Market Access Strategic Initiative #2: Engineer as a therapeutic area solutions company, capitalizing on the emerging market access environment

- Communicate Product Value Drivers
- Integrate Resources to Gain Synergies
- Innovate in Contracting
- Execute and Competition
- Optimize Market Access for Therapeutic Area Assets
- Gain Experience in Physician Risk Models
- Embrace a Corporate Culture and Stay with it
- Invest in Value-added Programs adding Value to Brands
Impactful science from bench to market
WE ARE CERTARA

90% of all novel drugs approved by the U.S. FDA in the past three years were supported by Certara software or services

1,700 companies, academic institutions, global non-profits, and leading regulatory agencies in 60 countries partner with Certara

850+ Employees including 250+ PhD, PharmD, and MD consultants

110+ Global regulatory submissions written and approved in the last three years from a team of 250+ Regulatory science consultants

Certara software is used by major regulatory agencies and considered a “gold standard” by the U.S. FDA
MEET OUR SENIOR US TEAM

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