

Health Tech Company – PPT Slides and IG Posts

Client provided **white papers** and **technical slides**, which I reformatted into **less-technical slides** and **Instagram posts**. The sales team used the new slides for physician presentations, leaving the white papers behind for further details.

Following are three samples for: **pulmonary arterial hypertension, GLP-1 inhibitors, and COPD**.

Each sample is three pages: the **technical slide** that was provided to me, the **less-technical** slide I created, and the **Instagram posts** that I created.

(I've redacted the company name, drug names, and some other sensitive text.)



3 High

Comparative effectiveness of endothelin receptor antagonists (ERA) in Japanese patients with pulmonary arterial hypertension: a real-world administrative claims database study

Situation

██████████ is the latest ERA approved for PAH. There was no evidence on comparative effectiveness of three ERAs in real world setting post approval, particularly in Asian countries. Evidence was needed to evaluate the effectiveness of these ERAs on persistence before and after ██████████ approval in Japan (2015).

Solution

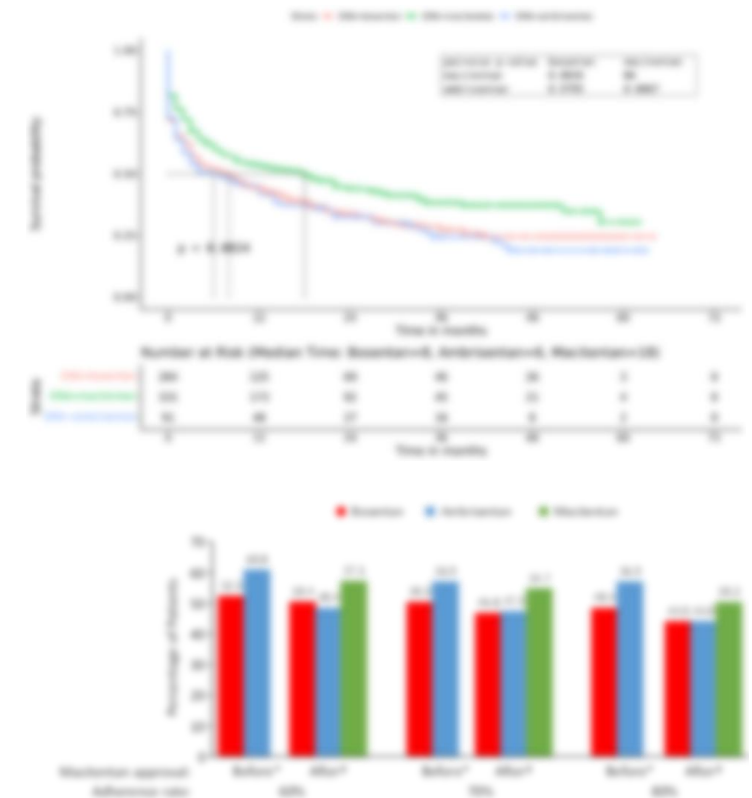
Generate evidence to understand the effectiveness of recent treatments using real-world data by conducting a retrospective observational study with an aim to increase awareness amongst health care providers to improve patient outcomes. Persistence to all three ERAs before and after launch of latest ERA was analysed.

~~Use of RWD~~ Combine with 'Solution'

██████████ used real-world data from the Japanese Medical Data Vision administrative claims database between April 2008 and November 2020. Persistence was defined as the time between the start of the index ERA and treatment discontinuation, censoring, or death. Propensity score (PS) adjustment was applied to minimize confounding effects among treatment groups.

Outcome

Real-world data for Japanese patients with PAH showed that persistence was ██████████ as long for ██████████, after its approval, versus ██████████ and ██████████.



*The study was also replicated in Australia (PBS10) and US (DRG) showing similar results. Both have been presented in Pulmonary Hypertension Society Australia & New Zealand (PHSANZ) 11th Annual Scientific Meeting and in European Society of Cardiology (ESC) 2022 (in the respective countries).

Generating evidence from real-world data to evaluate persistence to therapy for pulmonary arterial hypertension



Challenge

The Customer leveraged [redacted] patient-centric intelligence to evaluate persistence of its treatment compared to two others, using real-world data.



Results

Using real-world data from an administrative claims database, [redacted] found that persistence was [redacted] for the Customer's product, as compared to the other two.



Goals

The Customer was looking to address:

- Comparative effectiveness
- Assess persistence before and after Customer's treatment approval
- Increase awareness among healthcare providers



Impact

This allowed the Customer to:

- Provide key differentiation for its brand
- Influence clinicians toward their product
- **Increase prescriptions and reduce risk**



Patient-centric Intelligence in Action

Evaluating persistence to therapy from claims database

- Top 5 Pharma Client



Challenge

There was previously no evidence of comparative effectiveness of three therapies in real-world settings, post-approval.



Objective

The customer needed an assessment of persistence pre- and post-approval, and to evaluate comparative effectiveness.



Results

██████████ found that persistence was twice as long for the customer's product, as compared to the other two.



Impact

The customer was able to provide key differentiation for its brand, influence clinicians toward its product, increase prescriptions and reduce risk.

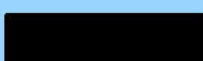


10+
Years in Business

20+
Therapeutic Areas

100+
Pharma Customers

100+
Clinical Indications





Analysis of off-label utilization of GLP-1 inhibitors

Situation

- GLP-1 have long been used (not under PBS restriction) for weight-loss.
- PBS hypothesized that patients may have been prescribed GLP-1s with PBS support outside PBS restriction.
- In preparation for the DUSC review, research was required to understand the breakdown of patients potentially using GLP-1 within/outside PBS restriction.

Solution

- designed the solution and method to examine:
- Criteria of in/outside PBS usage based on patients initiating on GLP-1s.
 - Determine concomitant use of medications (by class) with GLP-1.

Use of RWD ~~Combine with 'Solution'~~

Retrospective observational study using real-world data was conducted using the PBS 10% dataset

Outcome

At end of each year there are renegotiations between sponsors' and the government to determine and optimise reimbursement criteria and status in PBS. The Drug Utilisation Sub Committee (DUSC) would determine whether patients were utilising drugs in-line with its' set regulations. The results were used by the sponsor to predict the DUSC review assisting them to renegotiate ahead of time their drugs reimbursement criteria and status.

Breakdown of unique patients who initiate GLP-1 within and outside the PBS restriction from 2017 to 2022

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- 3.4. Quantifying patients co-prescribed with SGLT-2 and/or DPP-4

3.1. Quantifying patients prescribed with SGLT-2 and/or DPP-4

In the first stage, we identified the number of patients who initiated GLP-1 within and outside PBS restriction. We then analysed the number of patients who initiated GLP-1 with SGLT-2 and/or DPP-4. We then analysed the number of patients who initiated GLP-1 with SGLT-2 and/or DPP-4 and who were also prescribed with SGLT-2 and/or DPP-4.

Category	2017	2018	2019	2020	2021	2022
GLP-1 with SGLT-2 and/or DPP-4	1,234	1,567	1,890	2,123	2,456	2,789
GLP-1 without SGLT-2 and/or DPP-4	3,456	3,789	4,123	4,456	4,789	5,123
Total	4,690	5,356	6,013	6,579	7,245	7,912

3.2. Quantifying patient intention to reformatin and/or SU

We analysed the number of patients who initiated GLP-1 with SGLT-2 and/or DPP-4 and who were also prescribed with SGLT-2 and/or DPP-4. We then analysed the number of patients who initiated GLP-1 with SGLT-2 and/or DPP-4 and who were also prescribed with SGLT-2 and/or DPP-4.

Category	2017	2018	2019	2020	2021	2022
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Total	4,690	5,356	6,013	6,579	7,245	7,912

3.3. Quantifying patients in different claim history categories

We analysed the number of patients who initiated GLP-1 with SGLT-2 and/or DPP-4 and who were also prescribed with SGLT-2 and/or DPP-4. We then analysed the number of patients who initiated GLP-1 with SGLT-2 and/or DPP-4 and who were also prescribed with SGLT-2 and/or DPP-4.

Category	2017	2018	2019	2020	2021	2022
GLP-1 with SGLT-2 and/or DPP-4	1,234	1,567	1,890	2,123	2,456	2,789
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Total	4,690	5,356	6,013	6,579	7,245	7,912

3.4. Quantifying patients co-prescribed with SGLT-2 and/or DPP-4

We analysed the number of patients who initiated GLP-1 with SGLT-2 and/or DPP-4 and who were also prescribed with SGLT-2 and/or DPP-4. We then analysed the number of patients who initiated GLP-1 with SGLT-2 and/or DPP-4 and who were also prescribed with SGLT-2 and/or DPP-4.

Category	2017	2018	2019	2020	2021	2022
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Total	4,690	5,356	6,013	6,579	7,245	7,912

Analyzing off-label utilization of diabetes drugs within and outside a government subsidy program's criteria



Challenge

The Customer leveraged [redacted] patient-centric intelligence to determine usage of drugs in advance of a subcommittee review for actual usage and financial cost.



Results

Using real-world data from the program, [redacted]'s study provided a factual resource for the Customer to predict the subcommittee's review findings.



Goals

The Customer was looking to address:

- Understanding usage in/outside program's criteria for patients initiating in the drug class.
- Determining concomitant use of other drug classes with this one.



Impact

This allowed the Customer to:

- Be prepared to react to the subcommittee's findings expeditiously
- Submit data-based arguments to renegotiate reimbursement criteria and status



██████████
Patient-centric Intelligence in Action

Analyzing off-label utilization of drugs for government subsidy program

- Top 5 Pharma Client



Challenge

An upcoming subcommittee review of a drug's off-label use could potentially affect the drug's future reimbursement criteria and status.



Objective

The customer engaged ██████████ to address usage in/outside the program's criteria and to determine concomitant use of other drug classes with the customer's.



Results

██████████ study provided a factual resource for the customer to predict the subcommittee's review findings.



Impact

The customer was able to respond to the subcommittee's report expeditiously and submit data-based arguments to renegotiate reimbursement criteria and status.

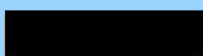


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COPD Analysis - Triple Therapy sizing analysis for PBAC submission

Situation

Pharmaceutical Benefits Advisory Committee (PBAC) is the expert advisory body that advises the Minister about which treatments should be made available on the Pharmaceutical Benefits Scheme (PBS). In preparation for PBAC submission the client undertook a market analysis of patients on COPD treatment.

Solution

defined the patient cohort of interest and algorithms to determine episodes of care, regimens of treatment, conversion to scripts and their equivalent beneficiary type. Post defining these segments the following could be examined:

- The number of patients on COPD and its growth rate
- The scripts dispensed over time (for COPD) and their beneficiary type

Use of RWD - Combine with 'Solution'

A retrospective cohort analysis was conducted using sample of prescriptions in Pharmaceutical Benefit Schedule (PBS) data from 2006-January through to 2018-March identifying scripts for all scripts.

Outcome

The study identified patients in COPD, the annual patient numbers and growth rates in Australia.

Project Description

Data Set Description

Defining COPD patients

Results 1: Number of Patients on Triple Therapy and Annual Growth Rate

Results 2: Number of Scripts by Beneficiary Type

Results 3: Number of Prescriptions by Beneficiary Type



Analyzing the COPD market to inform listing submission for government reimbursement



Challenge

The Customer leveraged [redacted]'s patient-centric intelligence to analyze complex data that involved combination drugs, inferred conclusions, overlapping conditions, and unclear treatment pathways.



Results

By defining the patient cohort and pertinent algorithms and applying inference and therapeutic knowledge, [redacted] identified triple-therapy patients, annual patient numbers and growth rates.



Goals

The Customer was looking to address:

- The number of patients on triple therapy
- Scripts dispensed over time
- Scripts' beneficiary types



Impact

This allowed the Customer to:

- Present a submission to apply for their drug to be reimbursed.
- Save time in the submission process and reduce the risk of not earning reimbursement status.



██████████
Patient-centric Intelligence in Action

Analyzing a therapeutic market to inform submission for reimbursement

- Top 5 Pharma Client



Challenge

The condition in question was complex to analyze because it involved combination drugs, inferred conclusions, overlapping conditions, and unclear treatment pathways.



Objective

The customer sought analysis for the number of patients on triple therapy, scripts dispensed over time, and scripts' beneficiary types.



Results

██████████ identified triple-therapy patients, annual patient numbers and growth rates.



Impact

██████████ analysis saved the customer time in the submission process and reduced the risk of an unfavorable reimbursement ruling.

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