# An Economic Model Assessing Costs, Relapses, and Antipsychotic Adherence of Switching Medicaid Beneficiaries With Schizophrenia From Oral Antipsychotics to Once-Monthly, Once-Every-Three-Months, and Once-Every-Six-Months Paliperidone Palmitate

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## Background

- Schizophrenia is associated with substantial clinical and economic burden in the United States (US), with an estimated prevalence ranging from 1.46% to 4.23% among Medicaid beneficiaries across 7 states and total excess costs for patients with schizophrenia, relative to those without, of more than $340 billion in the US in 2019[1-3]
- The large clinical and economic burden of schizophrenia highlights the importance of long-term compliance to antipsychotic (AP) treatment to improve symptoms and reduce the likelihood of relapse[4]
- Long-acting injectable (LAI) APs, including paliperidone palmitate, have been shown to be associated with improved AP adherence and clinical outcomes compared to oral APs (OAPs), while remaining cost-neutral[5-8]
- In an evolving LAI treatment space with emergent real-world data, there is a need to better understand the impact of paliperidone palmitate treatment pathways on economic and clinical outcomes among a population of Medicaid beneficiaries with schizophrenia

## Objective

- To evaluate the plan-level impact on healthcare costs, relapse rates, readmission rates, and average AP adherence of switching Medicaid beneficiaries with schizophrenia from OAPs to once-monthly paliperidone palmitate (PP1M) alone, and with subsequent transitions to once-every-three-months (PP3M) and once-every-six-months paliperidone palmitate (PP6M)

## Methods

### Model framework/structure

- A Markov model with twelve 3-month cycles over a 36-month time frame was developed from a Medicaid payer perspective (**Figure 1**)
- Three cohorts of adults with schizophrenia were included: those treated with once- or twice-daily OAPs, those switching to PP1M only, and those switching to PP1M with subsequent on-label transitions to PP3M and PP6M
- For the switching cohorts, 20% of the population was switched to PP1M, based on real-world evidence of LAI utilization rates among Medicaid beneficiaries[9]
- The model evaluated and compared total plan-level healthcare costs (based on costs of AP treatment and relapse, reported in 2025 US dollars [USD]), relapse rates, readmission rates, and average AP adherence among the three cohorts of patients

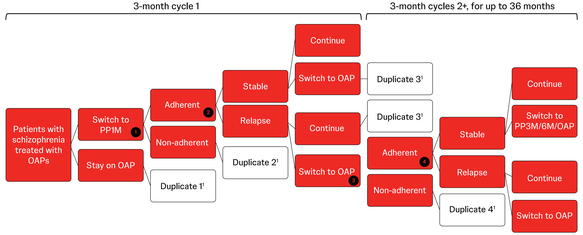
### Model assumptions

- At each 3-month cycle, patients may or may not be adherent to AP treatment, may or may not experience a relapse, and could continue receiving the same AP treatment or switch treatment
- The risk of relapse differed according to treatment and adherence status, and medication switch rates differed according to relapse and adherence status; patients could have experienced multiple relapses and treatment switches (up to 12) over the 36-month model time horizon
- Patients transitioning from PP1M to PP3M/PP6M had to do so on-label (i.e., after two 3-month cycles of PP1M, at the correct corresponding dose strength, with no evidence of relapse during the current cycle)[10,11]
- Patients on PP3M/PP6M could switch to another paliperidone palmitate formulation, or an OAP, while patients on an OAP could only switch to another OAP
- Patients switching to PP1M were assumed to be part of the PP1M treatment arm regardless of whether they switched to an OAP over the course of the model

### Model inputs

- All model inputs including population inputs, rates of relapse, readmission, adherence, and treatment switches, as well as costs of treatment and relapse, were literature-based[10,12-18]
- Considering both statutory and inflation discounts, a 70% discount was used for PP1M and PP3M treatment costs; for PP6M, given that it is a newer product, a 60% discount was applied, in line with prior literature[12,19]

**Figure 1: Model structure**




OAP = oral antipsychotic; PP1M = once-monthly paliperidone palmitate; PP3M = once-every-three-months paliperidone palmitate; PP6M = once-every-six-months paliperidone palmitate.
†At this point the branch is being duplicated from the corresponding number. Duplication does not include any switches to PP3M and PP6M from the OAP cohort.

## Results

- In a hypothetical health plan of 1 million Medicaid beneficiaries, an estimated 13,419 adult members with schizophrenia treated with OAPs incurred plan-level costs of $1,253.5 million, based on 33,671 relapses, 7,778 readmissions, and average AP adherence of 69.5% over the 36-month model time horizon (**Table 1**)

**Table 1: Plan-level costs, relapses, readmissions, and average AP adherence associated with switching 20% of Medicaid beneficiaries with schizophrenia from OAPs to PP1M, with subsequent transitions to PP3M and PP6M**

| | Patients on any OAPs [A] | Patients switching to PP1M only [B] | Difference [B – A] | Patients switching to PP1M, PP3M, and PP6M [C] | Difference [C – A] |
|---|---|---|---|---|---|
| **Plan-level costs (2025 USD)** | | | | | |
| Year 1 | $417,824,890 | $379,828,955 | –$37,995,934 | $378,754,322 | –$39,070,567 |
| Net cost per patient switched per year | - | - | –$14,158 | - | –$14,558 |
| Year 2 | $417,824,890 | $381,940,519 | –$35,884,370 | $377,788,663 | –$40,036,226 |
| Net cost per patient switched per year | - | - | –$13,371 | - | –$14,918 |
| Year 3 | $417,824,890 | $385,536,042 | –$32,288,848 | $380,857,921 | –$36,966,969 |
| Net cost per patient switched per year | - | - | –$12,031 | - | –$13,774 |
| **Total** | **$1,253,474,669** | **$1,147,305,517** | **–$106,169,152** | **$1,137,400,906** | **–$116,073,763** |
| Net cost per patient switched over 3 years | - | - | –$39,559 | - | –$43,250 |
| **Relapses** | | | | | |
| Year 1 | 11,224 | 9,614 | –1,610 | 9,545 | –1,678 |
| Year 2 | 11,224 | 9,782 | –1,442 | 9,517 | –1,706 |
| Year 3 | 11,224 | 9,926 | –1,297 | 9,609 | –1,614 |
| **Total** | **33,671** | **29,322** | **–4,349** | **28,672** | **–4,999** |
| **Quality measures** | | | | | |
| **Readmissions** | | | | | |
| Year 1 | 2,593 | 2,221 | –372 | 2,205 | –388 |
| Year 2 | 2,593 | 2,260 | –333 | 2,198 | –394 |
| Year 3 | 2,593 | 2,293 | –300 | 2,220 | –373 |
| **Total** | **7,778** | **6,773** | **–1,005** | **6,623** | **–1,155** |
| **Average AP adherence**[1] | | | | | |
| Year 1 | 69.5% | 78.9% | 9.4% | 82.5% | 13.0% |
| Year 2 | 69.5% | 76.3% | 6.8% | 86.8% | 17.3% |
| Year 3 | 69.5% | 75.6% | 6.1% | 86.8% | 17.3% |
| **Average** | **69.5%** | **76.9%** | **7.4%** | **85.4%** | **15.9%** |

AP = antipsychotic; OAP = oral antipsychotic; PP1M = once-monthly paliperidone palmitate; PP3M = once-every-three-months paliperidone palmitate; PP6M = once-every-six-months paliperidone palmitate; USD = United States Dollars.
[1]Average AP adherence was reported among patients who switched and was compared between the cohort of patients remaining on OAPs relative to the 20% of patients who switched to PP1M, as well as the 20% of patients with subsequent on-label transitions to PP3M and PP6M.

### Plan-level costs

- Switching 20% (n=2,684) of patients from OAPs to PP1M only resulted in plan-level cost savings of $106.2 million (**Figure 2**)
- Incorporating subsequent transitions to PP3M and PP6M led to additional plan-level savings of $9.9 million relative to patients treated with PP1M only, representing $116.1 million in plan-level cost savings relative to patients treated with OAPs

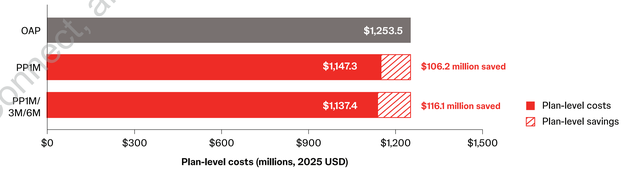
### Relapses and Readmissions

- Switching 20% of patients from OAPs to PP1M only avoided 4,349 relapses, with subsequent transitions to PP3M and PP6M avoiding an additional 650 relapses (**Figure 3**)
- Similarly, switching patients from OAPs to PP1M avoided 1,005 readmissions, and subsequent transitions to PP3M and PP6M additionally avoided 150 readmissions

### Adherence

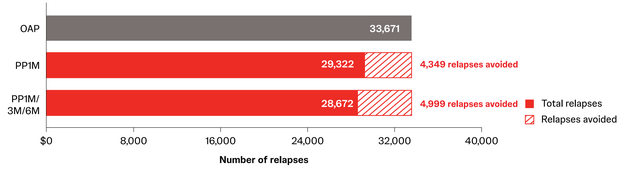
- Among the 20% of patients who switched from OAPs to PP1M only, switching to PP1M resulted in a 7.4 percentage point increase in average adherence to any AP, while incorporating transitions to PP3M and PP6M resulted in an additional 8.5 percentage point increase in average AP adherence (**Figure 4**)

**Figure 2: Total plan-level costs and cost savings associated with switching 20% of Medicaid beneficiaries with schizophrenia from OAPs to PP1M, with subsequent transitions to PP3M and PP6M**



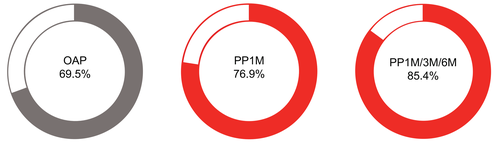

OAP = oral antipsychotic; PP1M = once-monthly paliperidone palmitate; PP3M = once-every-three-months paliperidone palmitate; PP6M = once-every-six-months paliperidone palmitate; USD = United States Dollars.

**Figure 3: Total relapses and relapses avoided associated with switching 20% of Medicaid beneficiaries with schizophrenia from OAPs to PP1M, with subsequent transitions to PP3M and PP6M**




OAP = oral antipsychotic; PP1M = once-monthly paliperidone palmitate; PP3M = once-every-three-months paliperidone palmitate; PP6M = once-every-six-months paliperidone palmitate.

**Figure 4: Average AP adherence associated with switching 20% of Medicaid beneficiaries with schizophrenia from OAPs to PP1M, with subsequent transitions to PP3M and PP6M**




AP = antipsychotic; OAP = oral antipsychotic; PP1M = once-monthly paliperidone palmitate; PP3M = once-every-three-months paliperidone palmitate; PP6M = once-every-six-months paliperidone palmitate.

## Limitations

- Results reported are specific to patients with schizophrenia covered by Medicaid and may not be generalizable to patients with other types of insurance or those who are uninsured
- Furthermore, as the estimated number of adult Medicaid beneficiaries with schizophrenia in a 1 million member plan is based on single-state prevalence data, plan-level cost savings and relapses avoided may vary if national prevalence estimates differ
- Model inputs were based on data available in published literature and may not fully represent outcomes observed in clinical practice; analyses based on real-world data are warranted to confirm the findings of this study
- Given that PP6M is a newer product, the treatment discount was based on assumptions, and real-world rebates may vary widely

## Conclusions

- Based on the findings of this 36-month Markov model, switching patients from OAPs to PP1M is associated with notable plan-level cost savings driven by avoided relapses and increased adherence to APs
- These results highlight the economic benefit of paliperidone palmitate LAIs, with longer-acting formulations PP3M and PP6M leading to additional incremental cost savings, further reductions in relapse rates, and continued AP adherence improvements
- This study offers important insights that should be taken into account in real-world clinical practice to inform treatment decisions

**Disclosures**
CB and YD are employees of Johnson & Johnson. LM, BL, LD, and DP are employees of Analysis Group, ULC., a consulting company that has provided paid consulting services to Johnson & Johnson which funded the development and conduct of this study.

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