

Evaluation of Concurrent Medications Pre- and Post Initiation of Long-Acting Injectable Antipsychotic Therapy

Lindsay Pokallus PharmD

Lehigh Valley Health Network, Lindsay.Pokallus@lvhn.org


Laurence Karper MD

Lehigh Valley Health Network, Laurence.Karper@lvhn.org

Jessica Price PharmD, BCPS

Lehigh Valley Health Network, Jessica_B.Price@lvhn.org

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EVALUATION OF CONCURRENT MEDICATIONS PRE- AND POST INITIATION OF LONG-ACTING INJECTABLE ANTIPSYCHOTIC THERAPY

Lindsay Pokallus, PharmD; Laurence Karper, MD; Jessica Price, PharmD, BCPS • LEHIGH VALLEY HEALTH NETWORK • ALLENTOWN AND BETHLEHEM, PA

PURPOSE

Retrospectively evaluate concurrent psychotropic medications in schizophrenic patients prior to- and during long-acting injectable (LAI) antipsychotic therapy

BACKGROUND

- **Approximately one percent (1%) of the adult population (2.4 million) has schizophrenia¹**
- **Schizophrenic patients oftentimes have multiple comorbid mental and substance-abuse issues, and other health-related comorbidities²**
- **Medication compliance is of immense importance in these patients^{2,3,4}**
 - Non-adherence or partial medication compliance occurs in >50% of schizophrenic patients⁵
- **Primary goals of therapy: minimize or eliminate the symptoms of schizophrenia, maximize the patients' quality of life and social functioning, maintain a level of mental stability⁶**
- **Meta-analysis reviewed 613 outpatient 'mirror-image' studies⁷**
 - Assessed evidence of efficacy for oral versus LAI antipsychotic medications⁷
 - Hospitalization days for patients taking LAI antipsychotic medications versus oral preparations: 17,860 versus 75,492 days⁷
 - Relapse rates for patients using LAI antipsychotics compared to oral preparations: 30% versus 47.1% patients, statistically significant⁷

- **Olfson et al. found that patients receiving LAI preparations compared with oral antipsychotics continued to require multiple concurrent oral psychotropics⁸**
 - Antidepressants, anxiolytics/hypnotics, mood stabilizers, and oral antipsychotics⁸
- **Shi et al. evaluated characteristics of patients taking LAI antipsychotics versus oral antipsychotics⁹**
 - Two-thirds of LAI antipsychotic group concurrently received more psychotropic medications compared to those on oral antipsychotics only⁹
 - LAI antipsychotic group tended to be more difficult-to-treat, non-compliant and refractory to traditional oral antipsychotics⁹
- **If LAI antipsychotics can decrease the number of concurrent psychotropic medications, this may increase patient compliance**
- **No trials directly assess the number of concurrent psychotropic medications prior to- and during LAI antipsychotic therapy**

STUDY DESIGN

- **Retrospective chart review**
- **Inclusion criteria:**
 - Patients prescribed LAI antipsychotic (fluphenazine decanoate, haloperidol decanoate, paliperidone palmitate, and long-acting risperidone) for a minimum period of 180 days for schizophrenia
 - Patients who received LAI antipsychotic between January 2004 and July 2010
- **The primary outcome of the study will be the number of concurrent psychotropic medications prior to LAI antipsychotic therapy, and during LAI antipsychotic treatment**
- **The secondary outcome will be patient relapse rates prior to LAI antipsychotic therapy, and during LAI antipsychotic treatment**

METHODS

- **Chart review evaluating number of concurrent psychotropic medications and relapse rates prior to- and during LAI antipsychotic treatment**
 - Number of concurrent psychotropic medications will be tallied for 180 days prior to- and 180 days during LAI antipsychotic treatment
 - Relapse rate will be examined, as indicated by number of emergency room visits, hospitalizations and admissions for a schizophrenic episode
 - Psychotropic medications: any medication used to treat symptoms of a mental disorder, including schizophrenia, depression, bipolar disorder, anxiety disorders, and attention deficit-hyperactivity disorder (ADHD)
- **Other patient data to be collected includes:**
 - Age, gender, weight, renal function
 - Patient comorbidities, age and date of schizophrenia diagnosis, previously documented failed medications
 - Dates and durations of admissions for schizophrenia exacerbations
 - Patient Diagnostic and Statistical Manual of Mental Disorders (DSM-IV-TR) information for diagnosis, Positive and Negative Syndrome Scale (PANSS) symptoms
 - Factors that may have contributed to acute episodes or mental/mood status changes (including acute illness, trauma, illicit drug use)
- **Following data collection, the number of concurrent psychotropic medications and relapse rates will be compared**

DISCLOSURE

Authors of the presentation have the following to disclose concerning possible financial or personal relationships with commercial entities that may have a direct or indirect interest in the subject matter of this presentation.

- Lindsay Pokallus – nothing to disclose
- Laurence Karper – nothing to disclose
- Jessica Price – nothing to disclose

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