Positive and negative symptom predominance in the screening period in acute schizophrenia trials: An exploratory analysis

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SUBMISSION DETAILS

What is the Methodological Question Being Addressed? In the current analyses we wanted to assess symptom predominance changes (changes in severity of positive and negative symptoms) in acute schizophrenia clinical trials and their impact on last visit change in the blinded data.

Introduction We have previously reported that on average 38% of subjects entering acute schizophrenia clinical trials have negative symptoms more severe than positive symptoms (Daniel and Kott; 2017). We have as well observed that the positive vs. negative symptom predominance may change between screening and baseline. In the current analysis we further explored this phenomenon of symptom predominance changes in screening period.

Methods Screening and baseline data were pooled from 15 acute schizophrenia clinical trials. The severity of positive and negative symptoms was derived from the PANSS positive and negative subscales. Symptom predominance was determined based on the difference between positive and negative symptoms as positive predominance (positive symptoms more severe than negative), negative predominance and no predominance. Lastly we assessed whether symptom predominance changed between screening and baseline and if so, in which direction. The impact of change in predominance between screening and baseline was then assessed in the blinded data on the last visit PANSS change from baseline, correcting for study and baseline severity using regression analysis.

Results The dataset consisted of 4,667 pairs of screening and baseline ratings and 4,144 last visit assessments. 35% of screening visits and 36% of baseline visits had negative symptoms more severe than positive, in 7% negative and positive symptoms were equally severe at both screening and baseline. Symptom predominance between screening and baseline changed in 9.3% of cases, in 4.8% from negative to positive and in 4.5% from positive to negative. No effect on last visit change from baseline was observed.

Conclusion In the current dataset negative symptom predominance occurred in around 35% of screening or baseline visits. Changes in symptom predominance in the screening period occurred in over 9% of data and were equally likely to be from negative to positive or from positive to negative symptoms. In the blinded data the changes seem to have no effect on end of treatment PANSS change, but more meaningful differences might be seen in unblinded datasets. We plan to reassess the impact on symptom predominance change on unblinded datasets as these become available.

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Keywords

Keywords	
Positive symptoms	
Negative symptoms	
Symptom predominance	
Acute schizophrenia	
Clinical Trial	

Guidelines I have read and understand the Poster Guidelines

Disclosures if applicable All authors are full time employees of Signant Health. The poster was financially supported by Signant Health.

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