

## Footnotes for the Infographic

## Antipsychotics: Benefits, Risks and Limitations

## from Onward Mental Health

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- [1] Citrome L et al, Schizophrenia, Clinical Antipsychotic Trials of Intervention Effectiveness (CATIE) and number needed to treat: how can CATIE inform clinicians, Int J Clin Pract. 2006, PMID: 16893436, https://goo.gl/iQXmVa. Lieberman J et al, Effectiveness of Antipsychotic Drugs in Patients with Chronic Schizophrenia, N Engl J Med. 2005, PMID: 16172203, https://goo.gl/hQeWv5
- [2] Leucht S et al, Sixty Years of Placebo-Controlled Antipsychotic Drug Trials in Acute Schizophrenia: Systematic Review, Bayesian Meta-Analysis, and Meta-Regression of Efficacy Predictors, 2017, Amer Jof Psychiatry, <a href="https://goo.gl/bndxBq">https://goo.gl/bndxBq</a>. Note: At least a "minimal" response [we call it "minimal benefit"] occurred in 51% of the antipsychotic group versus 30% in the placebo group, and 23% versus 14% had a "good" response [we call it "substantial benefit"]. 23% 14% = 9% see substantial benefit attributable to ("due to") antipsychotics, so 100% 9% = 91% do NOT see "substantial benefit" "due to" antipsychotics. 51%-30% = 21% see "minimal benefit" attributable to ("due to") antipsychotics, so 100%-21% = 79% do NOT see "minimal benefit" "due to" antipsychotics.
- [3] Ucok, Sexual dysfunction in patients with schizophrenia on antipsychotic medication, Eur Psych, 2007, <a href="MID:17344032"><u>PMID: 17344032</u></a>. Young SL et al, "First do no harm." A systematic review of the prevalence and management of antipsychotic adverse effects, <a href="PMID:25516373">PMID: 25516373</a>, <a href="https://goo.gl/on3k62">https://goo.gl/on3k62</a>.
- [4] Miller D, Extrapyramidal side-effects of antipsychotics in a randomised trial, Br J Psychiatry. 2008, PMC2801816. "...Table 1 probability of having a parkinsonism event within 1 year for people with no parkinsonism at baseline with adjustment for baseline covariates shows 37%–44% for the four second-generation antipsychotics and 37% for perphenazine". Note: we have used the midpoint percentage of 40% in the infographic. The data used is the large CATIE study from footnote #1.
- [5] Fusar-Poli P et al, Progressive brain changes in schizophrenia related to antipsychotic treatment? A meta-analysis of longitudinal MRI studies, Neurosci Biobehav Rev. 2013, <a href="PMCID: PMC3964856">PMCID: PMC3964856</a>.
- [6] Waddington JL, Mortality in schizophrenia. Antipsychotic polypharmacy and absence of adjunctive anticholinergics over the course of a 10-year prospective study, Br J Psychiatry 1998, PMID: 9926037. Joukamaa M et al, Schizophrenia, neuroleptic medication and mortality. Br J Psychiatry, 2006, PMID: 16449697. Ito H et al, Polypharmacy and excessive dosing: psychiatrists' perceptions of antipsychotic drug prescription. Br J Psychiatry. 2005, PMID: 16135861.
- [7] Rajkumar, AP et al, Endogenous and antipsychotic-related risks for diabetes mellitus in young people with schizophrenia: a Danish population-based cohort study, Am J Psychiatry. 2017, PMID: 28103712.
- [8] Xiang Y et al, Almost All Antipsychotics Result in Weight Gain: A Meta-Analysis, 2014, PMCID: PMC3998960;
- [9] Harrow M et al, A 20-Year multi-followup longitudinal study assessing whether antipsychotic medications contribute to work functioning in schizophrenia, 2017, Psychiatry Research, PMID: 28651219.
- [10] Harrow M, Do all schizophrenia patients need antipsychotic treatment continuously throughout their lifetime? A 20-year longitudinal study, Psychological Medicine, 2012, PMID: 22340278, https://goo.gl/HwUOj8; Wunderink et al, Recovery in remitted first-episode psychosis at 7 years of follow-up of an early dose reduction/discontinuation

or maintenance treatment strategy: long-term follow-up of a 2-year randomized clinical trial, JAMA Psychiatry. 2013, <a href="MID:23824214">PMID: 23824214</a>.

[11] Ray et al, Atypical Antipsychotic Drugs and the Risk of Sudden Cardiac Death, NE J Med 2009, PMCID: PMC2713724.