

# OUTCOMES AND THEIR PREDICTORS IN SCHIZOPHRENIA – SYSTEMATIC REVIEWS

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- The aim is to present key results of recent systematic reviews and meta-analyses on outcomes of schizophrenia.
- The included reviews focus on proportion of recovery in schizophrenia and how other clinical and functional outcomes are predicted by family history of psychosis, onset age, and duration of untreated psychosis (DUP).

Table 1

Study	Recovery% (95% CI)
Hong Kong (incidence cohort, WHO)	0.0 (0.0, 0.4)
Johanson 1958	1.2 (1.0, 1.4)
Bond 1921	2.1 (1.7, 2.5)
Suvisaari et al. 2009	3.1 (2.7, 3.5)
Auslander and Jeste 2004	3.2 (3.0, 3.4)
Wolter et al. 2010	3.9 (3.5, 4.3)
Dublin (DOSMed incidence cohort)	5.0 (4.0, 6.0)
Nagasaki (DOSMed incidence cohort)	5.8 (5.2, 6.5)
Rupp and Fletcher 1940	6.4 (6.2, 6.6)
Möller et al. 2011	7.3 (6.8, 7.8)
Gottlieb 1940	8.0 (7.5, 8.6)
DeLisi et al. 1998	8.0 (7.2, 8.8)
Lambert et al. 2009	8.1 (8.0, 8.2)
Obembe et al. 1995	8.7 (7.5, 9.9)
Nyman and Jonsson 1983	9.0 (8.4, 9.6)
Mannheim (RAPyD incidence cohort)	9.1 (8.2, 10.0)
Kaleda 2009	9.7 (9.3, 10.1)
Lauronen et al. 2005	9.9 (9.2, 10.6)
Sofia (RAPyD incidence cohort)	10.0 (9.0, 11.0)
Huber et al. 1980	10.0 (9.7, 10.3)
Muller et al. 1951a	12.0 (11.3, 12.7)
Beijing (prevalence cohort, WHO)	12.1 (11.2, 13.0)
Prague (DOSMed incidence cohort)	12.5 (11.4, 13.6)
Malamud and Render 1939	13.0 (12.5, 13.5)
Harris et al. 1956	13.0 (12.4, 13.6)
Harrow et al. 1978	13.9 (13.1, 14.7)
Langfeldt 1937	16.0 (15.2, 16.8)
Bland and Orn 1978	16.3 (15.1, 17.5)
Robinson et al. 2004	16.4 (15.5, 17.4)
Qureshi et al. 1987	16.7 (15.5, 17.9)
Achte 1967b	16.8 (16.0, 17.6)
Henisz 1966	16.9 (16.3, 17.5)
Achte 1967a	17.7 (16.9, 18.5)
Angst and Preisig 1995	18.4 (17.4, 19.1)
Wootton et al. 1935	18.9 (18.0, 19.8)
Henry et al. 2010	19.0 (18.5, 19.5)
Modestin et al. 2003	19.7 (19.1, 20.3)
Muller et al. 1951b	20.0 (19.1, 20.9)
Nottingham (DOSMed incidence cohort)	20.4 (19.2, 21.6)
Vazquez-Barquero et al. 1999	23.0 (21.9, 24.1)
Rennie 1939	25.2 (24.5, 25.9)
Ciampi 1980	26.6 (26.0, 27.2)
Holmboe and Astrup 1957	29.0 (28.3, 29.7)
Cali (IPSS prevalence cohort)	31.8 (30.1, 33.5)
Ogawa et al. 1987	32.4 (31.1, 33.7)
Moscow (DOSMed incidence cohort)	32.4 (30.6, 34.2)
Chennai (incidence cohort, WHO)	36.4 (35.1, 37.8)
Chandigarh, urban (DOSMed incidence cohort)	37.0 (34.7, 39.3)
Agra (IPSS prevalence cohort)	51.9 (49.2, 54.6)
Rajotte and Denber 1963	58.0 (55.9, 60.1)
Overall	16.4 (14.5, 18.2)

## RECOVERY

Recovery was defined as improvements in both clinical and social domains with a two-year good outcome for at least one of the domains. In all studies, a follow-up of at least two years was required (Jääskeläinen et al. 2013).

We identified 50 studies with data suitable for inclusion. The median proportion (25%–75% quantiles) of patients who met our recovery criteria was 13.5% (8.1%–20.0%) (Table 1).

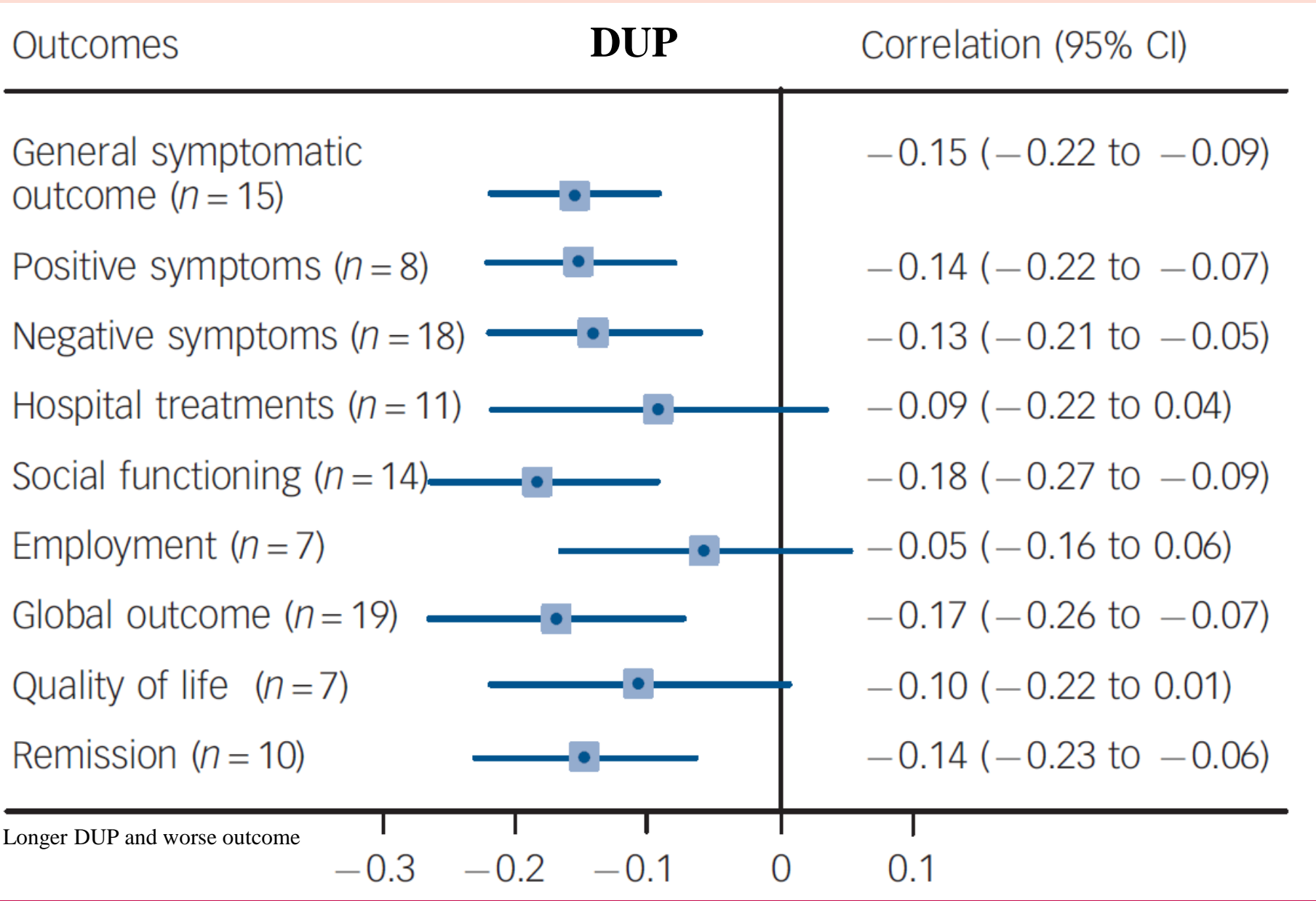
Recovery proportions did not differ by gender. Proportion of recovered cases had not increased in recent decades. Countries with poorer economic status had higher recovery proportions. See Table 2.

## PREDICTORS OF CLINICAL AND FUNCTIONAL OUTCOMES

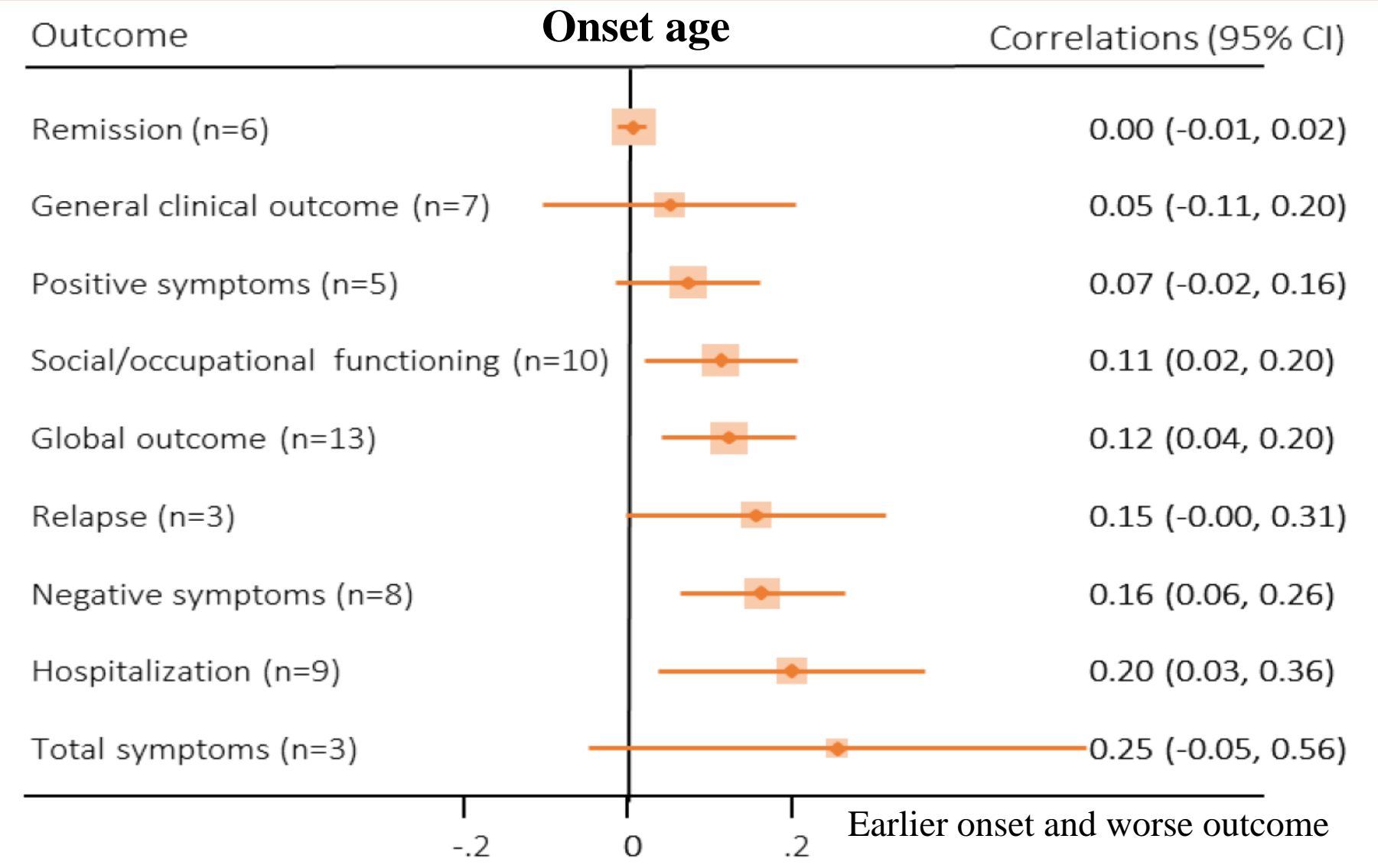
14 studies investigated associations between family history of psychosis and outcomes. Family history of psychosis was associated with poor occupational ( $r=0.17$ ) and global ( $r=0.13$ ) outcome (Käkälä et al. 2014).



33 studies investigated associations between DUP and outcomes. DUP associated with poor general symptomatic outcome, more severe positive and negative symptoms, lower likelihood of remission, poor social functioning, and poor global outcome (correlations 0.13–0.18) (Penttilä et al. 2014).



81 studies investigated associations between onset age and outcomes. Earlier onset age associated with more hospitalizations, negative symptoms, relapses, worse social/occupational functioning, and poorer global outcome (correlations 0.11–0.17) (Immonen et al. In press).



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## CONCLUSIONS

- Despite major changes in treatment options in recent decades, the proportion of recovered cases has not increased. Current outcomes are unsatisfactory and their exact mechanisms, trajectories, and predictors are partly unknown.
- Family history of psychosis has a relatively small but statistically significant negative effect on the occupational and global outcome.
- Longer duration of untreated psychosis was associated with poorer outcomes.
- Earlier age at onset has a small, but significant negative impact on some of the outcomes.
- Studied predictors, family history, onset age and duration of untreated psychosis, may correlate with each other and further studies are needed to study their interactions.

