

Five-Year Follow-up of a Randomized Multicenter Trial of Intensive Early Intervention vs Standard Treatment for Patients With a First Episode of Psychotic Illness

The OPUS Trial

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Context: Intensive early treatment for first-episode psychosis has been shown to be effective. It is unknown if the positive effects are sustained for 5 years.

Objective: To determine the long-term effects of an intensive early-intervention program (OPUS) for first-episode psychotic patients.

Design: Single-blinded, randomized, controlled clinical trial of 2 years of an intensive early-intervention program vs standard treatment. Follow-up periods were 2 and 5 years.

Setting: Copenhagen Hospital Corporation and Psychiatric Hospital, Aarhus, Denmark.

Patients: A total of 547 patients with a first episode of psychosis. Of these, 369 patients were participating in a 2-year follow-up, and 301 were participating in a 5-year follow-up. A total of 547 patients were followed for 5 years.

Interventions: Two years of an intensive early-intervention program vs standard treatment. The intensive early-intervention treatment consisted of assertive community treatment, family involvement, and social skills training. Standard treatment offered contact with a community mental health center.

Main Outcome Measures: Psychotic and negative symptoms were recorded. Secondary outcome measures were use of services and social functioning.

Results: Analysis was based on the principles of intention-to-treat. Assessment was blinded for previous treatment allocation. At the 5-year follow-up, the effect of treatment seen after 2 years (psychotic dimension odds ratio [OR], -0.32 ; 95% confidence interval [CI], -0.58 to -0.06 ; $P = .02$; negative dimension OR, -0.45 ; 95% CI, -0.67 to -0.22 ; $P = .001$) had equalized between the treatment groups. A significantly smaller percentage of patients from the experimental group were living in supported housing (4% vs 10%, respectively; OR, 2.3; 95% CI, 1.1-4.8; $P = .02$) and were hospitalized fewer days (mean, 149 vs 193 days; mean difference, 44 days; 95% CI, 0.15-88.12; $P = .05$) during the 5-year period.

Conclusions: The intensive early-intervention program improved clinical outcome after 2 years, but the effects were not sustainable up to 5 years later. Secondary outcome measures showed differences in the proportion of patients living in supported housing and days in hospital at the 5-year follow-up in favor of the intensive early-intervention program.

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PSYCHOSIS IS RELATED TO GREAT stress and is a burden for society; the personal consequences of relapse and the fear of future relapse are often overwhelming for patients.¹ Birchwood and colleagues² have hypothesized that there exists a critical period just after the debut of psychosis in which a window of opportunity exists to intervene and improve the long-term course of illness.

Previous studies³⁻⁶ have shown that it is possible to intervene during the early course

of illness to secure a better outcome, but there is no evidence that indicates how long early interventions need to be active to prevent relapse. Rosenheck et al⁷ have shown that for a large population of homeless people with severe mental disorders, the advantages of assertive community treatment (ACT) can be sustained after transfer to other services if clinicians are allowed the flexibility to decide at which stage a patient is ready for such a transfer.

This study (the OPUS trial) is the largest randomized clinical trial comparing the

intensive early-intervention program (OPUS) with standard treatment (community mental health centers) for patients experiencing their first episode of psychosis⁸ and the first to report outcome after 5 years of follow-up. The intensive early-intervention program consisted of ACT, psychoeducational family treatment, and social skills training. The experimental treatment was carried out for 2 years.

The intensive early-intervention program has shown significant positive effects on psychotic and negative symptoms, secondary substance abuse, treatment adherence, success with lower dosages of antipsychotic medication, and a higher satisfaction with treatment after 2 years of treatment.⁴ It is unknown if these positive effects are sustainable after the experimental treatment ends and patients are referred to standard treatment according to their needs.

The 3 core elements in the intensive early-intervention program aim to provide the patient with the skills to cope with the illness, manage their own medication, and reduce stress. If patients have learned to use these skills independently or with support from their families, they will theoretically help them to continue medication and buffer stress during the course of illness and when the experimental treatment is no longer active. The positive effects of ACT,^{9,10} psychoeducational family intervention,¹¹⁻¹³ and social skills training¹⁴⁻¹⁷ for patients with psychosis are well documented separately during the time when treatment is active, but follow-up studies indicate that the effects disappear when treatment ends.^{18,19}

This 5-year follow-up assesses the patients 3 years after the transition to standard treatment and offers the opportunity to investigate whether the intensive early-intervention treatment is able to sustain its positive effects.

Two hypotheses were tested; the first was linked to the primary outcome measure, and the second to the secondary outcome measure. (1) Patients who were allocated to the intensive early-intervention program for 2 years and then transferred to standard treatment have a better clinical outcome, measured by the level of psychotic and negative symptoms, global functioning, substance abuse, depression, and suicidal behavior, compared with patients allocated to standard treatment throughout the first 5 years of treatment. (2) Patients who were allocated to the intensive early-intervention program for 2 years and then transferred to standard treatment have a better social outcome compared with patients allocated to standard treatment; social outcome is measured as more patients living independently, fewer hospitalizations (less use of supported housing and days in hospital), and more patients with a competitive job or studying.

METHODS

PARTICIPANTS

A total of 547 patients were included in the trial during the period from January 1998 until December 2000. Patients were included from both inpatient and outpatient mental health services in Copenhagen and Aarhus. At the time of inclusion, patients were aged between 18 and 45 years and came in contact with mental health services for the first time with a diagnosis within the schizophrenia spectrum (measured using the F2-category codes

Table 1. Sociodemographic and Clinical Characteristics of 547 First-Episode Psychotic Patients at Entry Into the Trial of the Intensive Early-Intervention Program vs Standard Treatment

Characteristics	No. (%)	
	Intensive Early-Intervention Program (n=275)	Standard Treatment (n=272)
Sociodemographic		
Male	159 (58)	164 (60)
Mean (SD) age, y	26.6 (6.4)	26.6 (6.3)
Married	16 (6)	14 (5)
Being a parent	42 (15)	37 (14)
Completed high school	98 (36)	83 (31)
Education		
None	163 (60)	156 (59)
Being educated	38 (14)	31 (12)
Short education, skilled	54 (20)	52 (20)
Longer education	17 (6)	24 (9)
Living conditions		
Living alone vs with partner or child	208 (76)	213 (80)
Living with parents	49 (18)	41 (15)
Living in supervised setting	1 (0)	2 (1)
Homeless	14 (5)	10 (4)
Inpatient at randomization	117 (43)	127 (47)
Clinical		
Median duration of untreated psychosis, wk ^a	46	53
Diagnosis		
Schizophrenia	185 (67)	177 (65)
Schizotypal disorder	42 (15)	37 (14)
Delusional disorder	12 (4)	13 (5)
Brief psychosis	19 (7)	26 (10)
Schizoaffective disorder	10 (4)	15 (5)
Unspecified nonorganic psychosis	7 (2)	4 (1)
Psychopathology scores^b		
Psychotic dimension	2.8 (1.4)	2.6 (1.4)
Negative dimension	2.2 (1.2)	2.2 (1.2)
Disorganized dimension	1.0 (0.9)	1.0 (1.0)
Substance abuse		
Diagnosis of substance abuse	73 (27)	73 (27)
Suicidal ideation and behavior		
Suicide attempt ever	82 (32)	77 (31)
Suicide attempt during last year	48 (21)	52 (23)
Suicidal thoughts during last year	124 (58)	136 (64)
Social functioning		
Mean (SD) GAF symptoms	32.7 (10.3)	34.4 (11.0)
Mean (SD) GAF functioning	41.6 (13.6)	41.0 (13.1)

Abbreviation: GAF, global assessment of functioning (scale).

^an=429; was not assessed for patients who were diagnosed with schizotypal disorder and schizophrenia simplex.

^bScores of assessment scales with values ranging from 0 to 5.

of the *International Statistical Classification of Diseases, 10th Revision [ICD-10]*²⁰, and none of them had received antipsychotic medication for more than 12 continuous weeks. Patients were followed and reassessed after 2 and 5 years.

The basic characteristics of the cohort are shown in **Table 1**. Representativity was good. A total of 90% of the patients in Aarhus and 63% in Copenhagen were registered in the psychiatric case registers as having had their first contact with psychiatric services in the same period, and they were diagnosed within the same diagnostic spectrum. The official registers revealed that patients who were not included were significantly

older and fewer of the patients had a diagnosis of schizophrenia (66% in the trial compared with 42% in the register). Of the eligible patients, only 5% refused to participate.

RANDOMIZATION

Patients were centrally randomized to the intensive early-intervention program or standard treatment. In Copenhagen, randomization was carried out through centralized telephone randomization at the Copenhagen Trial Unit. The allocation sequence was a computer-generated ratio of 1:1 in blocks of 6, and stratified for each of 5 centers. In Aarhus, the researchers contacted a secretary by telephone when they had finished the entry assessment of each patient. The secretary then drew 1 lot from among 5 red and 5 white lots out of a black box. When the block of 10 was used, the lots were redrawn. Block sizes were unknown to the investigators.

TREATMENTS

The trial was pragmatic, comparing the intensive early-intervention program, which was defined by a set of protocols, with treatment as usual.

INTENSIVE EARLY-INTERVENTION PROGRAM

The intensive early-intervention program consisted of 3 core elements: ACT,^{8,9} family treatment,^{12,21} and social skills training.^{15,22} Two multidisciplinary teams were established and trained to provide the intensive early-intervention program for 2 years. The caseload ratio was 1 researcher for every 10 patients. The patients were designated a primary staff member who was responsible for maintaining contact and coordinating the treatment within the team and across social services and other involved institutions. Patients were visited at their homes or other places in the community, or seen at their primary team member's office, according to the patients' preferences. During hospitalization, responsibility for the patient was transferred to the hospital, but the primary staff member maintained contact with the patient at least weekly. Office hours were Monday to Friday, from 8 AM to 5 PM. Outside of office hours, patients could leave messages on the staff's cell phone and be assured that the team would respond the next morning. A crisis plan was developed and trained for with each patient. The aim of the intensive early-intervention program was to offer an individual plan of treatment for each patient. If patients were reluctant to be treated, the team stayed in contact with the patient and tried to find a common focus for collaboration to thereby motivate the patient to continue treatment.

The team always tried to get in contact with at least 1 family member and motivate the family to participate in a psychoeducational group. Family treatment followed the McFarlane manual for psychoeducational treatment²¹ for multiple family groups and included 18 months of treatment for 1.5 hours every second week in a multiple-family group with 2 therapists and 4 to 6 patients with their families. The focus was on problem solving and development of skills to cope with the illness.

Patients with impaired social skills were offered social skills training focusing on medication, coping with symptoms, conversation, and problem-solving skills in a group with a maximum of 6 patients and 2 therapists.

The fidelity of the treatment program, measured with the index of fidelity of assertive community treatment,²³ was 70% in Copenhagen and Aarhus. The factors responsible for the reduced fidelity were time-limited treatment, 24-hour coverage in other settings, and about 2 contacts weekly with each patient, the patient's family, and collaborating partners.

The intensive early-intervention program was phase specific, meaning that the primary team member carefully assessed when patients were ready for a specific treatment modality.

STANDARD TREATMENT

Standard treatment usually consisted of offering the patient treatment at a community mental health center. Each patient was in contact with a physician, a community mental health nurse, and in some cases, a social worker. Home visits were possible, but office visits were the general rule. A staff member's caseload in the community mental health centers varied between 20 and 30 patients. Outside of office hours, patients could refer themselves to the psychiatric emergency department. Such psychosocial treatments as supported counseling, psychoeducation, and family contact were provided infrequently and in a less intensive and unsystematic way, and only in a minority of cases. For more details see Petersen et al.⁴

TRANSITION FROM INTENSIVE EARLY-INTERVENTION PROGRAM TO STANDARD TREATMENT

After 2 years of the intensive early-intervention program, patients from the intervention group were transferred to standard treatment. For a few patients, this consisted of a transfer to only their general practitioner. The transition to standard treatment was carried out as gradually and gently as possible, but naturally the break in the relationship with the contact person in the intensive early-intervention program could cause feelings of loss for the patient. Introduction to standard treatment was a high priority, and the transition period could last up to 2 months.

ANTIPSYCHOTIC MEDICATION

Patients in both treatment groups were offered antipsychotic drugs according to guidelines from the Danish Psychiatric Society, which recommend a low-dose strategy for patients with a first episode of psychotic illness and the use of second-generation antipsychotic drugs as a first choice.

ASSESSMENTS

Independent investigators conducted the 5-year follow-up interviews. All investigators were blind to previous treatment allocation. For practical reasons, independent investigators at the 2-year follow-up could not be blinded.⁴

At entry, the 2-year follow-up, and the 5-year follow-up, information on the following topics was collected: (1) main diagnosis and substance abuse, based on the Schedule for Clinical Assessment in Neuropsychiatry ([SCAN] version 2.0 in 1998 and SCAN 2.1 since 1999)²⁴; (2) symptoms according to the Scale for Assessment of Psychotic Symptoms (SAPS) and Scale for Assessment of Negative Symptoms (SANS)²⁵ (data are analyzed according to the 3 dimensions: psychotic, negative, and disorganized, with values ranging from 0-5)²⁶; (3) sociodemographic factors²⁷; (4) course of illness with Life Chart Schedule²⁸; (5) global assessment of functioning and symptoms (GAF)²⁹; (6) duration of untreated psychosis, assessed at entry to the trial with the Interview for Retrospective Assessment of Onset of Schizophrenia³⁰; and (7) suicidal behavior, measured by self-reporting of suicide attempts and suicidal ideation.³¹

The algorithms from the SCAN interview were used to investigate whether patients fulfilled the general criteria for depression according to the ICD-10.

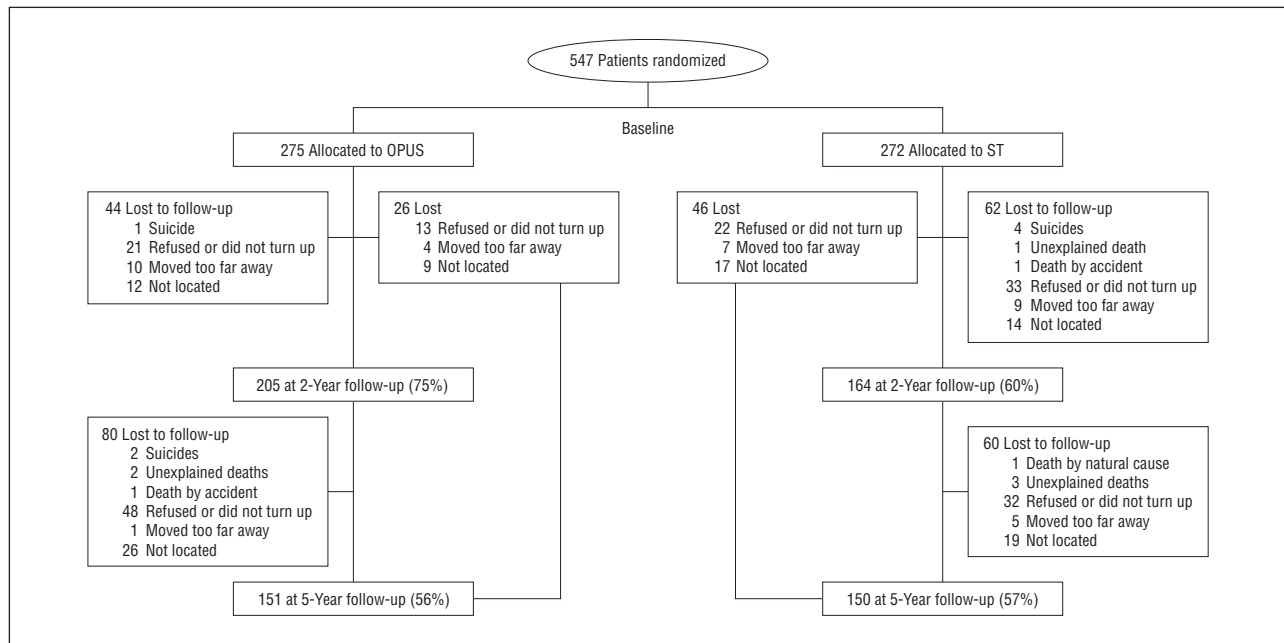


Figure 1. Flowchart of patients through study. OPUS indicates intensive early-intervention program; ST, standard treatment.

OTHER DATA SOURCES

Using the unique Danish official registers, it is possible to accomplish a complete follow-up of all patients regarding a range of relevant process and outcome measures. Information about days spent in hospital, emergency department visits, outpatient contacts, housing situation, and vocational situation were collected from the registers for all patients (100% follow-up rate) included in the trial, except those who had died or emigrated. A list of all supported housing institutions was made manually, so that all institutions were captured. A great effort was made to ensure that all institutions used are supported housing facilities for patients with mental health problems. Most of the institutions are staffed 24 hours daily.

The following information was gathered: (1) In the Danish Civil Registration System,³² all persons alive and residing in Denmark are registered and assigned a 10-digit personal identification number. The register contains continuously updated information. (2) Information about days in hospital, emergency department contacts, and outpatient contacts was collected from the Danish Psychiatric Central Register.³³ (3) A database with addresses for all supported housing facilities in all counties and municipalities was combined with address information in the Civil Status Register, thereby providing information about independent living and supported housing.³² (4) Employment, family situation, sick leave, and early-age pension were extracted from the Integrated Database for Labour Market Research.²⁷ (5) Mortality and cause of death were drawn from the Cause of Death Register.³⁴

INTERRATER RELIABILITY

All investigators were trained in conducting the SCAN interview at the World Health Organization collaborating center and trained in the SCAN, SAPS, and SANS with live interviews. Twenty five SCAN interviews were conducted. During the 5-year follow-up period, all raters from the 2-year follow-up and the 5-year follow-up did reliability interviews with SANS and SAPS. The intraclass correlation coefficient was 0.90 for the negative dimension and 0.92 for the psychotic dimension; both are classified as very good agreement.³⁵

BLINDING

Raters at the 5-year follow-up were blinded to patients' previous treatment allocation. After each interview, raters made a guess as to which treatment they believed the patient had most likely received. The reliability between the guessed and true treatment allocation were measured as a κ coefficient, 0.23, indicating poor reliability, which means that a fair level of blinding was obtained.³⁶

OUTCOME MEASURES

The primary outcome measures were psychotic and negative symptoms (SANS and SAPS) and social functioning (functional GAF). Secondary outcomes included secondary diagnosis of substance abuse, medication and use of services, depressive symptoms, suicidal behavior, housing situation, and vocational situation.

STATISTICAL ANALYSIS

Because of the attrition from the follow-up interviews (**Figure 1**), the influence of missing data on the 2- and 5-year outcome measures had to be considered. Hence, data from the SANS, SAPS, and GAF were subjected to further analysis using a mixed-model analysis with a repeated-measurements model with unstructured variance matrix (**Table 2**). This approach assumed that the distribution of missing data could be estimated from the information from previous interviews. The condition for using this method is the assumption that data were missing at random when taking into consideration the information extracted from entry and 2-year follow-up interviews.³⁷⁻³⁹ The following covariates were entered in the repeated measurements model: treatment, substance abuse at entry, sex, and age. The values from entry for the respective outcome measures (SAPS, SANS, and GAF) were included automatically, because they are included in the model and no treatment effect was allowed for at entry. Alternative approaches to managing the skewed attrition are sensitivity analyses and multiple imputations. According to Little et al,³⁷ multiple im-

Table 2. Clinical Outcome of Patients With a First Episode of Psychotic Illness Who Participated in the Intensive Early-Intervention Program or Standard Treatment^a

	Mean (SD)							
	2-Year Follow-up (n=369)				5-Year Follow-up (n=301)			
	OPUS (n=205)	ST (n=164)	Estimated Mean Difference (95% CI)	P Value of Difference	OPUS (n=151)	ST (n=150)	Estimated Mean Difference (95% CI)	P Value of Difference
Psychotic dimension	1.06 (1.26)	1.27 (1.40)	-0.32 (-0.58 to -0.06)	.02	1.41 (1.62)	1.31 (1.60)	0.04 (-0.3 to 0.39)	.83
Negative dimension	1.41 (1.15)	1.82 (1.23)	-0.45 (-0.67 to -0.22)	< .001	1.73 (1.29)	1.82 (1.46)	-0.05 (-0.34 to 0.24)	.73
Disorganized dimension	0.37 (0.56)	0.50 (0.73)	-0.12 (-0.25 to -0.00)	.06	0.42 (0.75)	0.47 (0.76)	-0.14 (-0.27 to 0.06)	.22
GAF symptoms	51.18 (15.01)	48.67 (15.92)	2.45 (-0.32 to 5.22)	.08	53.46 (16.64)	53.78 (17.79)	-0.16 (-3.97 to 3.37)	.96
GAF function	55.16 (15.15)	51.13 (15.92)	3.12 (0.37 to 5.88)	.03	55.36 (17.28)	54.16 (18.41)	1.34 (-2.65 to 5.34)	.51

Abbreviations: CI, confidence interval; GAF, global assessment of functioning (scale); OPUS, intensive early-intervention program; ST, standard treatment.

^aEstimated mean differences are based on a repeated measurement model in unstructured variance matrix. Treatment site, sex, substance abuse, and age were included as covariates.

putations are not necessary when data has already been entered in a repeated measurement model. Sensitivity analyses were carried out.

Odds ratios for treatment effects were calculated using logistic regression analysis, and mean differences were estimated through analysis of variance for continuous variables.

In accordance with the intention-to-treat principle, all patients were analyzed in the treatment groups to which they were randomly allocated, regardless of whether they had completely followed the scheduled design. All statistical analyses were done with the Statistical Package for the Social Sciences 11.0 (SPSS, Inc, Chicago, Illinois).

POWER CALCULATION

It was expected that the mean (SD) reduction in psychotic symptoms measured by SAPS would be 1 (1.3) point for the patients allocated to standard treatment. At a minimum, it should be possible to detect a 50% greater reduction in psychotic symptoms in the experimental group at the .05 level of significance and with power of 0.9. Using the Pocock formula,⁴⁰ 142 patients would be required for each study group being followed.

RESULTS

ATTRITION FROM STUDY

The flowchart (Figure 1) shows attrition from the study after 2 and 5 years. After 2 years, the attrition was skewed; 75% of the patients from the intensive early-intervention program attended the follow-up interview, as did 60% of the standard group. Furthermore, analysis of attrition at the 2-year follow-up revealed that patients from Copenhagen, patients who had not completed high school, and those with substance abuse at entry were more likely to not attend the 2-year follow-up.

Regarding the 5-year follow-up, no significant differences were found between entry and 1-year follow-up measures regarding sex, educational level, treatment site, duration of untreated psychosis, and psychotic and negative symptoms between those patients who attended the 5-year follow-up and those who did not. Nor were differences found between the whole group and each treat-

ment group when analyzed, except that patients not attending the 5-year follow-up were slightly older.

SENSITIVITY ANALYSIS OF PSYCHOTIC SYMPTOMS, NEGATIVE SYMPTOMS, AND SUBSTANCE ABUSE

Owing to the large attrition from the study (Figure 1), 2 assumptions were tested regarding patients who did not participate in the 5-year follow-up interview. The first assumption was that the nonparticipants' level of psychotic and negative symptoms and substance abuse would be the same as the last observation; therefore, the nonparticipants' entry values and 2-year values (if available) for the psychotic and negative dimensions and substance abuse were carried forward to the 5-year follow-up. The second assumption was that nonparticipants had experienced a total remission of psychotic and negative symptoms and substance abuse. On this basis, their values at the 5-year follow-up were set at 0. Data were analyzed using both assumptions, but it was found that neither of the 2 possibilities showed any significant differences between the 2 groups regarding level of symptoms or substance abuse.

Although no selection bias could be detected, it is still necessary to be cautious about possible bias with a follow-up rate of only 57%.

PRIMARY OUTCOME MEASURES

When analyzing clinical data with a repeated measurement model (Table 2), no significant difference between the 2 treatment groups was observed at the 5-year follow-up (psychotic dimension odds ratio [OR], 1.41 vs 1.31; mean difference, 0.04; 95% confidence interval [CI], -0.3 to 0.39, $P = .83$ and negative dimension OR, 1.73 vs 1.82; mean difference, -0.05; 95% CI, -0.34 to 0.24; $P = .73$). In other words, the psychotic effect seen after 2 years of treatment (psychotic dimension OR, 1.06 vs 1.27; mean difference, -0.32; 95% CI, -0.58 to -0.06; $P = .02$ and negative dimension OR, 1.41 vs 1.82; mean difference, -0.45; 95% CI, -0.67 to -0.22; $P < .001$) had disappeared during the 3 years that passed between the

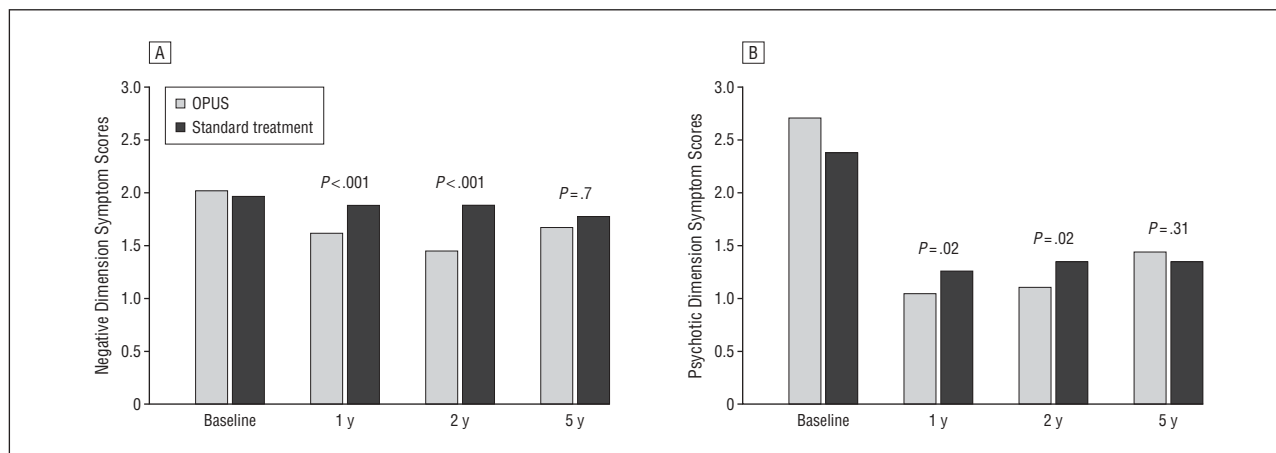


Figure 2. Mean symptom values for patients in the intensive early-intervention program (OPUS) vs standard treatment, according to the Scale for Assessment of Psychotic Symptoms and Scale for Assessment of Negative Symptoms²⁵ at baseline, 2-year follow-up, and 5-year follow-up for the negative (A) and psychotic (B) dimensions. Values range from 0 to 5.

2- and 5-year follow-ups, when the experimental treatment was no longer active. This development is illustrated in **Figure 2**.

Functional GAF also demonstrated an effect of the intensive early-intervention program after 2 years of treatment (55.16 vs 51.13; mean difference, 3.12; 95% CI, 0.37-5.88; $P = .03$), but this difference was also not present after 5 years (55.36 vs 54.16; mean difference, 1.34; 95% CI, -2.65 to 5.34; $P = .51$).

SECONDARY OUTCOME MEASURES

Depression, Substance Abuse, and Suicidal Behavior

The intensive early-intervention program significantly reduced substance abuse at the 2-year follow-up, but not at the 5-year follow-up (**Table 3**). No effect was found of the intensive early-intervention program on depression and suicidal behavior after 2 or 5 years.

Antipsychotic Medication

The proportion of patients receiving first- or second-generation antipsychotic medication was not significantly different in the 2 treatment groups after 2 or 5 years. Patients in the intensive early-intervention program received significantly lower doses of second-generation antipsychotic medication after 2 years, but not after 5 years (Table 3).

Use of Services

When analyzing data from the registers, it was found that patients who had participated in the intensive early-intervention program during the first 2 years spent significantly fewer days in hospital than patients who had received standard treatment (mean, 96 vs 123 days; mean difference, 27.4 days; 95% CI, 0.57-54.32; $P = .05$) (**Table 4**). From 2 to 5 years, there was no significant difference in mean days in hospital between patients in the 2 groups (58 vs 71 days; mean difference, 13.1 days;

95% CI, -12.5 to 38.7; $P = .31$). Because days in hospital decreased in this period, lack of power makes it impossible to say that the difference is not coincidental; hence, the result may be a type II error. Investigating the entire period of 5 years, patients in the intensive early-intervention program had 20% fewer mean days in hospital than patients in standard treatment (149 vs 193 days; mean difference, 44 days; 95% CI, 0.15-88.12; $P = .05$) (not shown).

Social Outcomes

All data on social outcome are derived from the registers that made it possible to follow up all patients except those who had died (Table 4). During the 5 years, 16 patients had died. A total of 7 patients died of suicide, 3 of unexplained causes, 1 by accident, and 1 of natural causes. Standard mortality rate was 11 compared with the general population in Copenhagen and Aarhus Counties, aged 18 to 45 years.

The proportion of patients living in supported housing did not differ between treatment groups after 2 years of treatment, but at the 5-year follow-up, only 4% of the patients in the experimental group were living in supported housing, whereas 10% from the standard treatment group were (OR, 2.3; 95% CI, 1.1-4.8; $P = .02$). The mean (SD) days spent in supported housing for the patients from the experimental group in the time period from 2 to 5 years was 57 (213) vs 102 (282) in the standard treatment group (mean difference, 45.1 days; 95% CI, 0.31-89.9; $P = .05$). The range of days spent in supported housing from entry to the 5-year follow-up was 5 to 1094 days for the experimental group and 27 to 1094 days for the standard group (not shown).

When analyzing the proportion of patients working or being educated, it was found that after 5 years, 61% of patients from the intensive early-intervention program and 59% from the standard treatment group were not working or studying. In this age range in the general population, 20% are not working or studying.²⁷ There were no significant differences between the 2 treatment groups.

Table 3. Substance Abuse, Use of Antipsychotic Drugs, and Suicidal Ideation and Behavior of Patients With a First Episode of Psychotic Illness Who Participated in the Intensive Early-Intervention Program or Standard Treatment

	No. (%)									
	2-Year Follow-up (n=436) ^a					5-Year Follow-up (n=301)				
	OPUS (n=243)	ST (n=193)	Difference in Percentages (95% CI)	OR (95% CI) / Parameter Estimate (95% CI)	P Value	OPUS (n=151)	ST (n=150)	Difference in Percentages (95% CI)	OR (95% CI) / Parameter Estimate (95% CI)	P Value
Substance abuse and suicidal behavior and ideation										
Diagnosis of substance abuse ^b	43 (17)	40 (21)	4 (-3 to 11)	0.5 (0.3 to 1.0)	.04	33 (22)	36 (24)	2 (-0.076 to 0.12)	0.8 (0.52 to 1.53)	.68
Diagnosis of depression ^b	37 (15)	35 (18)	3 (-4 to 11)	0.8 (0.5 to 1.5)	.47	37 (25)	29 (19)	-6 (-0.04 to 0.14)	0.8 (0.44 to 1.3)	.27
Attempted suicide during follow-up ^b	20 (8)	20 (10)	2 (-3 to 7)	0.8 (0.4 to 1.7)	.51	13 (9)	14 (9)	0 (-0.06 to 0.07)	0.9 (0.4 to 2.1)	.86
Suicidal thoughts during last 2 y ^b	113 (56)	86 (53)	3 (-0.1 to 0.07)	0.9 (0.7 to 1.6)	.62	83 (55)	93 (62)	7 (-0.03 to 0.19)	0.7 (0.4 to 1.1)	.15
Antipsychotic drug use										
Any antipsychotic drugs at follow-up date ^b	146 (60)	107 (55)	-5 (-14 to 4)	1.2 (0.9 to 1.8)	.23	98 (65)	98 (65)	0 (-0.1 to 0.11)	1.0 (0.6 to 1.6)	.93
First-generation drugs only ^b	25 (17)	21 (20)	3 (-4 to 10)	0.9 (0.5 to 1.4)	.51	32 (32)	22 (22)	-10 (-0.02 to 0.2)	0.6 (0.35 to 1.16)	.11
Second-generation drugs only ^b	72 (49)	45 (42)	-7 (-16 to 2)	1.3 (0.9 to 1.9)	.15	85 (87)	88 (90)	3 (-0.2 to 0.06)	1.3 (0.5 to 3.2)	.50
Mean (SD) equivalents of haloperidol, mg ^c										
First- or second-generation drugs ^c	4.3 (2.8)	5.3 (3.4)		-0.7 (-1.4 to 5.6)	.07	2.8 (3.2)	2.3 (2.0)		-0.48 (-1.24 to 0.28)	.21
First-generation drugs only ^c	3.3 (2.7)	3.0 (3.4)		0.29 (-1.01 to 1.6)	.66	1.9 (4.6)	1.6 (2.0)		-0.3 (-2.41 to 1.78)	.76
Second-generation drugs only ^c	4.0 (2.4)	4.9 (2.9)		-0.91 (-1.6 to -0.2)	.01	2.5 (2.1)	2.1 (1.8)		-0.35 (-0.93 to 0.22)	.22

Abbreviations: CI, confidence interval; OPUS, intensive early-intervention program; OR, odds ratio; ST, standard treatment.

^aResults from 2-year follow-up are presented in a previous article.⁴

^bOdds ratios and *P* values based on logistic regression analyses.

^cIn calculating equivalence, 100 mg of chlorpromazine was estimated to be equivalent to 2 mg of haloperidol. *P* values are based on analysis of variance.

REMISSION

The Life Chart Schedule²⁸ offered an opportunity to analyze the course of illness during the previous 2 years up to the 5-year follow-up (**Table 5**). Results showed that there were no significant differences between the 2 groups as to whether the course had been continuous or episodic, or if the patient had not been psychotic at all during the past 2 years.

COMMENT

The results of this large randomized controlled trial disprove our first hypothesis; 2 years in the intensive early-intervention program showed no effect on the clinical outcome at the 5-year follow-up, nor on psychotic or negative symptoms, global functioning, substance abuse, depression, or suicidal behavior. The positive effects on psychotic and negative symptoms and global functioning seen after 2 years of treatment were, in other words, not sustainable after the experimental treatment ended.

The second hypothesis regarding social outcome was partly confirmed; the results demonstrated that patients from the experimental group were living more independently (less use of supported housing), and spent significantly fewer days in hospital during the 5-year period. These results indicate that, to some extent, patients

from the experimental treatment group fared better with regard to adapting to normal life outside institutions. It is debatable how much attention should be paid to the statistically significant result regarding independent living. On one hand, data on supported housing are a secondary outcome measure, but are very robust, reliable, and valid, especially considering that the number of days spent in supported housing after 2 and 5 years also differ significantly. These results offer unique insight into the patients' social abilities and capability to live alone without support, and are the best proxy measure for social functioning in our study. Further, Marshall et al⁹ showed that ACT had an effect on adherence, days in hospital, employment, and housing situation, meaning that even though housing situation is a secondary outcome measure, it is highly relevant.

On the other hand, if a Bonferroni correction had been applied to the figures, the significant difference between the 2 groups regarding independent living would have disappeared. According to Schulz et al,⁴¹ caution should be exercised when there are more than 15 end points and only 1 turns out to be statistically significant. Because we have fewer end points, a Bonferroni correction of the data was not chosen.

The intensive early-intervention program added substantial cost to treatment, but this was counterbalanced by the reduced cost of other health services during this

Table 4. Use of Health Services and Social Outcome for Patients With a First Episode of Psychotic Illness Who Participated in the Intensive Early-Intervention Program or Standard Treatment^a

	Percentage of Patients (n=547)									
	2-Year Follow-up					5-Year Follow-up				
	OPUS	ST	Difference in Percentages (95% CI)	OR (95% CI)/ Parameter Estimate (95% CI)	P Value	OPUS	ST	Difference in Percentages (95% CI)	OR (95% CI)/ Parameter Estimate (95% CI)	P Value
Use of services										
Mean (SD) No. of days in hospital ^b	96 (146.7)	123 (170.6)		27.4 (0.57 to 54.32)	.05	58 (145.1)	71 (154.9)		13.1 (-12.5 to 38.7)	.31
Median No. of days in hospital ^c	25	52			.04	0	0			
Not hospitalized ^d	89 (32)	73 (27)	-5 (-0.1 to 0.02)	1.3 (0.88 to 1.84)	.20	157 (57)	148 (54)	-3 (-0.1 to 0.06)	1.0 (0.74 to 1.52)	.66
No outpatient contacts last year ^d	20 (7)	85 (31)	24 (17 to 31)	0.2 (0.1 to 0.3)	<.001	127 (46)	133 (48)	4 (-0.04 to 0.13)	0.8 (0.6 to 1.2)	.41
Mean (SD) use of emergency department, No. ^b	1.43 (2.6)	1.71 (4.0)		0.28 (-0.28 to 0.87)	.33	1.9 (4.6)	2.2 (5.9)		0.3 (-0.6 to 1.2)	.51
Social outcome										
Not living independently ^d	19 (7)	18 (7)	0 (-0.04 to 0.04)	1.0 (0.5 to 1.9)	.99	11 (4)	26 (10)	6 (-0.09 to -0.006)	2.3 (1.1 to 4.8)	.02
Mean (SD) No. of days in protected homes ^b	30 (105)	35 (122)		5.2 (-24.6 to 14.3)	.59	57 (213)	102 (282)		45.1 (0.31 to 89.9)	.05
Not working or being educated ^d	142 (52)	151 (56)	4 (-0.02 to 0.14)	0.8 (0.5 to 1.1)	.20	159 (57)	148 (54)	-3 (-0.1 to 0.06)	1.1 (0.8 to 1.6)	.57
Mean (SD) No. of days absent from work owing to illness ^b	75 (125)	80 (125)		-5.1 (-19.1 to 28.7)	.69	45 (115)	40 (95)		-5.9 (-26.1 to 15.29)	.58
Mean (SD) No. of days on early pension ^b	58 (163)	75 (186)		16.9 (-15.9 to 49.1)	.31	382 (437)	430 (457)		48.3 (-37.2 to 133.8)	.26
Living alone or alone with children ^{d,e}	243 (88)	219 (81)	-7 (-0.1 to -0.00)	1.6 (0.9 to 2.7)	.05	226 (82)	206 (75)	-7 (-0.1 to 0.01)	1.4 (0.8 to 2.2)	.16

Abbreviations: CI, confidence interval; OPUS, intensive early-treatment program; OR, odds ratio; ST, standard treatment.

^aTwo-year figures show the period from entry to 2-year follow-up; 5-year figures show the period from 2-year follow-up to 5-year follow-up. Data are from registers with 547 patients followed up after both 2 and 5 years. Patients who had died are not included in analysis.

^bP values based on analysis of variance.

^cP values based on Mann-Whitney U test for independent sample.

^dOdds ratios and P values based on logistic regression analyses.

^eLiving alone or alone with children mean that the patient is either living totally by himself or herself or that he or she is a single mother or father raising his or her children alone, with no partner in the household.

intervention, especially the savings on supported housing after 5 years.

The external validity of the trial was high owing to good representativity.

Dropout analyses showed that patients not participating in the 2-year follow-up⁴ had a poorer prognosis, but the same analysis of patients participating in the 5-year follow-up revealed that it was not a select group of patients who were participating. Even though it is not possible to detect a possible attrition bias at the 5-year follow-up, it cannot be refuted that the large attrition at the 5-year follow-up may have biased the results.

Power calculations showed that 142 patients were required in each arm, and this requirement was fulfilled; hence, it is not likely that the 5-year follow-up results are subject to a type II error, but a 50% reduction in symptoms is possibly a very optimistic expected difference.

In the 5-year follow-up, performance bias may have influenced the results, owing to the fact that the patients from the experimental group, unlike the patients from the standard group, experienced a shift in treatment continuity when they were transferred to standard treatment after 2 years.

The raters at the 5-year follow-up were blinded to previous treatment allocation, and data from the registers offered 100% follow-up of patients, thus minimizing the risk of our results being biased. On the other hand, assessment at the 2-year follow-up may have been associated with a biased rating owing to the lack of blinding; therefore, these results may have been subject to detection bias.

Our findings are in accordance with findings from other studies regarding outcome after 2 years of early intensive treatment,^{3,5,6} but these studies do not investigate the effect of the experimental treatments for an extended time period.

In strict terms, it must be concluded that the benefits of the intensive early-intervention program after 2 years

Table 5. Remission and Relapse During Last 2 Years Before 5-Year Follow-up^a

	5-Year Follow-up, No. (%)		
	Intensive Early-Intervention Program (n=151)	Standard Treatment (n=150)	Differences in Percentages (95% CI)
Episodic course of illness ^b	21 (14)	19 (13)	-2 (-0.06 to 0.1)
Continuous course of illness ^c	67 (45)	65 (44)	-2 (-0.12 to 0.1)
Not psychotic ^d	62 (41)	64 (43)	2 (-0.13 to 0.09)

Abbreviation: CI, confidence interval.

^aBased on Life Chart Schedule.²⁸

^bEpisodic characterized as no psychotic episode lasting longer than 6 months.

^cContinuous characterized as no remission lasting longer than 6 months.

^dNot psychotic meaning no psychotic episode during last 2 years.

were not sustainable, and no basic changes in illness were seen after 5 years from the start of the program.

Our results give rise to questions about how long early-intervention services should be offered to patients to maintain good clinical and social outcomes. Second, this trial pinpoints the intrinsic problem of early-intervention services, namely how to make the transition to normal life as gentle as possible for those patients who no longer need treatment, or who need a less intensive treatment program, while at the same time maintaining continuous treatment for those who develop a chronic course of illness.

There is a need to continuously investigate whether the short-term (12-24 months)³⁻⁶ outcome achieved with specialized early-intervention services can be sustained for a longer time period. There is also a need to investigate the consequences of a prolonged treatment program that lasts for the entire 5 years, which is hypothesized to cover the critical period.² More randomized controlled trials are also needed that try to determine which specific elements from specialized early intervention, if not all, need to be offered for an extended time period. It would also be of great interest for future research to carry out analysis of the mediating factors of the different treatment modalities to determine whether some subgroups of the sample might be faring differently from others. This would give the field of research an indication of whether the experimental treatment is particularly beneficial for some groups of patients compared with other groups.

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