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Predictors for Work Participation in Individuals with an Autism Spectrum Disorder: A Systematic Review

Anja Holwerda · Jac J. L. van der Klink · Johan W. Groothoff · Sandra Brouwer

Abstract  Introduction Research shows that only about 25% of people with autism are employed. Method We conducted a systematic review on factors facilitating or hindering work participation of people with autism in longitudinal studies. An extensive search in biomedical and psychological databases yielded 204 articles and 18 satisfied all inclusion criteria. We assessed the methodological quality of included studies using an established criteria list. Results Seventeen factors were identified and categorized as disease-related factors, personal factors or external factors. Limited cognitive ability was the only significant predictor consistently found for work outcome. Functional independence and institutionalization were both reported by one study to be significantly related to work outcome. Inconsistent findings or non significant findings were reported for the other fourteen factors. Conclusion These findings emphasize the need for more high quality cohort studies focussing on work participation as the main outcome among people with Autism.

Keywords  Autism · Work participation · Predictors

Introduction

Work participation is considered as an increasingly important health outcome [1]. On the individual level it contributes to health and welfare [2]. On the societal level demographic pressure due to ageing and shrinking populations make a broad participation more and more imperative. At the same time participation in work by vulnerable groups is complicated by increasing demands in work. Young disabled people willing to enter the workforce experience barriers in acquiring and retaining work. Despite the relevance and although the impact of autism on social outcomes has been described in quite a few studies in the existing literature [3–7], the body of knowledge regarding factors facilitating or hindering work participation of people with autism is limited.

Autism

Autism, a life-long lasting developmental disability, affects social functioning, behavior, learning and cognition [8, 9]. According to Kobayashi et al. [3] three in four individuals with autism also have intellectual disabilities. Autism spectrum disorders (ASD) seem to be more prevalent in boys than girls [10].

Autism and Work Participation

Adults with autism have typically not been considered suitable candidates for employment in the work force [8, 11, 12]. Especially the social deficits typical for most
people with autism hinder their integration in the work force [4, 13]. Research shows that only about 25% of people with autism are employed. These are mostly the more high-functioning individuals. Unemployment rates for individuals with ASD as well as mental retardation are especially high [14]. They are mostly in sheltered employment, if employed at all [15].

Notwithstanding these limitations, there are several opportunities for work for these young disabled people, like regular work (including supported employment), sheltered employment, daytime activity or voluntary work [8]. People with autism can benefit from employment socially as well as personally [11]. Employment can provide a daytime structure that is helpful for this group as well as social contacts that are otherwise difficult for them to maintain [8, 11, 16]. Having a job also may facilitate their self-confidence, self-worth, independence and autonomy [11, 17].

Factors Associated with Autism and Work Participation

The available reviews describing autism and social outcome suggest that the majority of individuals with autism is unable to lead an independent life, including employment [13, 18–20]. Most individuals suffer (severe) persisting impairments in communication and social life limiting their independence and social functioning considerable, especially as demands on social adaptation and functioning increase with age [7, 13, 19]. Howlin [6] suggests that, as far as high functioning individuals are concerned, employment levels may be more dependent on the area individuals live in and the available support services than on any other factors. Also access to supported employment programs for this group may increase chances to find and retain appropriate jobs [6, 21]. IQ, communicative speech at 5–6 years of age, the level of mental retardation, and other comorbidity are mentioned as important predictors for outcome in individuals with ASD [18–20, 22]. Of those with a comorbid intellectual disability (IQ scores <50) few are capable of employment. Outcome for individuals with an IQ between 50 and 70 is more variable, but not much better. Outcome for individuals with an IQ of 70 or more seemed to be more promising but also more difficult to predict [6, 18–20, 22]. Besides deficits in cognitive and social functioning, limited independent performance and high dependence on caregiver support are considered important contributors to restricted outcome for individuals with ASD [9, 13].

To our knowledge, the literature on factors associated with work participation in adulthood for people with ASD has not yet been reviewed systematically. In a recent review factors influencing the work participation of young disabled starters entering the labor market were identified [23]. They found that male gender, higher education, high psychosocial level of functioning, low depression and high dispositional optimism were promoting factors in relation to employment. However, in their review no studies on autism were included.

The aim of this review is to systematically investigate the prognostic factors facilitating or hindering work participation of people with ASD.

Methods

We started conducting a systematic review of the scientific literature on prognostic factors related to work participation of people with ASD. However, we found only one study focusing on factors in relation to work outcome as a primary outcome measure; in most studies work outcome measures were incorporated in an overall social outcome. Therefore, we decided to include also studies looking at overall social outcome, incorporating employment, which provided valuable information about work outcome as well. Studies on overall social outcome including employment, not reporting on work outcome specifically, were not included. The International Classification of Functioning was selected as an underlying framework because it takes the multidimensional nature of work participation into account and provides a broad view on predictors [1].

The first (AH) and second reviewer (SB) discussed search strategy, criteria for selecting studies, quality assessment and data extraction to reach consensus. In case of disagreement the third reviewer (JvdK) made the final decision.

Literature Search

An extensive search in biomedical and psychological databases was performed (PubMed, PsycINFO, Embase, Cinahl, ERIC, SocINDEX) to find relevant articles, using MeSH terms, subheadings and free text words. Original studies (in English, Dutch, German and French) were identified that were published till June 2011. Only longitudinal studies were included to be able to distinguish predictors of work outcome. The search strategy consisted of an autism component and a work-related outcome measure.

In Table 1 the search terms are presented.

To select relevant studies for this review, the following eligibility criteria were defined: (1) Studies reported on factors related to work participation or social outcome in people with autism, only if information about work participation or employment status were included; (2) autism had to be diagnosed during childhood by an expert (e.g. following DSM-IV or ICD-10 criteria). The inclusion criteria are:
• **Types of studies**: Cohort studies, follow-up studies or longitudinal studies with a minimum follow-up period of 1 year.

• **Types of participants**: Persons in the age bracket 18–64 years, with disability due to autism spectrum disorder, diagnosed before the age of 18.

• **Types of outcome measures**: Dependent variables: participation in work (regular, supported or sheltered) or social outcome with a work identifiable component.

Titles and abstracts were screened independently by two reviewers (AH, SB). Full papers were retrieved if the abstract provided insufficient data to enable selection. Moreover, other relevant articles were searched on the basis of the name of the first author of included articles and the reference lists. Reviews were excluded, but their reference lists were inspected for additional studies.

Data Extraction

Using a standardized form, the first reviewer (AH) extracted data on study design, source population, inclusion criteria, numbers of participants, length of follow-up, loss to follow-up, outcome, prognostic factors and statistical analysis. Meta-analysis of the study results was not possible due to the descriptive nature of the included studies, the different outcome measures used and the limited availability of data which could be used for pooling.

Assessment of Methodological Quality of Included Studies

Two reviewers (AH, SB) independently assessed the methodological quality of all included articles in the final selection. The quality assessment of the selected studies was based on an established criteria list for assessing validity of prognostic studies, as recommended by Altman [24] and used in previous reviews [25, 26]. The internal validity was the main aspect judged to inform the reader about the quality of the studies regarding the aim of our review.

The criteria list consists of 16 items, each having yes/no/don’t know answer options. We operationalized the criteria for this review and pilot tested this operationalization on four longitudinal studies excluded for this present review. The final criteria list is presented in Appendix [27].

The quality of all included articles was scored independently by two reviewers (AH, SB). If sufficient information was available, the item was rated one point. When information was not given or the information given was unclear, the item was rated zero point. For the total quality score we added all points for each study (maximum score 16 points).

Studies with a minimum score of 11 points (≥70%) were arbitrarily considered to be of high quality and those with a score lower then 11 points (<70%) of low quality. This cut-off score is in line with a previous review [26]. We calculated initial interobserver agreement on methodological quality using kappa statistics for dichotomous values.

Results

Selection of Studies

The initial search yielded 204 articles (search date: June 6th, 2011). After selecting 19 references for full text reading, both reviewers agreed to include 14 articles for the present review. Two articles were excluded because they were intervention studies. Three articles did not report on specific employment outcomes. Searching the reference lists of those included articles, we found and included 3 additional articles. Based on the name of the first author of the 17 included articles, we found 1 other relevant article. Figure 1 shows a flow chart of study selection. In total we included 18 articles for the present review [3–5, 7, 28–41] (Table 2).

Study Characteristics

The characteristics of each study regarding country, design, measurements, population, numbers enrolled, time to follow-up and loss to follow-up are presented in Table 2. Time to follow-up varied considerably within as well as

<table>
<thead>
<tr>
<th>Table 1 Search terms (* = truncated)</th>
<th>Terms linked to MeSH</th>
<th>Free text words</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diagnosis</td>
<td>Child development disorders, pervasive * Asperger syndrome autistic disorder</td>
<td>Autism autistic disorder pervasive developmental disorder Asperger syndrome</td>
</tr>
<tr>
<td>Population</td>
<td>Work employment (exploded) rehabilitation, vocational (exploded) vocational guidance</td>
<td>Exclusion: child and not adult</td>
</tr>
<tr>
<td>Outcome measure</td>
<td></td>
<td>Career employment/employed/ employee(s) occupation vocation job</td>
</tr>
<tr>
<td>Study design</td>
<td>Cohort studies longitudinal studies prospective studies follow-up studies</td>
<td>Cohort longitudinal prospective follow-up prognostic</td>
</tr>
</tbody>
</table>
between studies, with the minimal time to follow-up being 3.2 and 35 years at most.

Quality Assessment and Methodological Considerations

The final overall agreement between the two reviewers on quality score was $\kappa = 0.80$, which is considered to be acceptable. Disagreement originated mainly from reading errors and misinterpretation of the criteria list and was readily resolved in a consensus meeting. The methodological quality of all included studies is summarized in Table 3. Four studies were considered of high methodological quality and fourteen of low quality. Statistical pooling of data in a meta-analysis was not possible because of the heterogeneity of study population and quality of the included studies.

Predictors for Work Participation

Seventeen different prognostic factors were identified. In Table 4 an overview of these factors related to work outcome is presented per included study. Table 5 gives an overview of these factors. The prognostic factors are categorized as disease/disorder related factors, personal factors or external factors based on the ICF-model [42, 43]. The only significant predictor for work outcome, consistently found in fifteen studies, is intelligence. Functional independence and institutionalization were reported in two separate low quality studies to be significantly predicting work outcome. Inconsistent findings were reported for diagnosis, severity of disorder, gender, language abilities, and maladaptive behavior. Non significant findings were reported for comorbidity, social impairments, lack of drive, parental support, family income, mental illness parents, family situation, treatment/use of medication and schooling.

Disease Related Factors

Diagnosis

Six studies found that the more severe the disorder the lower the chance on a good outcome [5, 28, 36, 37, 39, 40]. With regard to work participation, one study reported that individuals who were competitively employed had significantly fewer autism symptoms than those who had a supported job or were participating in adult day activity programs [40].

Comorbidity

Comorbidity (psychiatric disorder, oppositional personality or epilepsy) was mentioned by five studies as negatively influencing work outcome [5, 30, 38–40]. No evidence was found that use of medication hinders a favorable work outcome [33].
<table>
<thead>
<tr>
<th>No</th>
<th>Study</th>
<th>Country</th>
<th>Population (diagnosis, gender and age)</th>
<th>Numbers enrolled</th>
<th>Design</th>
<th>Baseline at</th>
<th>Time to follow-up</th>
<th>% Lost to follow-up</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Rutter et al. (1967) [28]</td>
<td>United Kingdom</td>
<td>Individuals diagnosed as children with child psychosis, schizophrenic syndrome, infantile autism or any synonyms of these Gender: 51 male/12 female Age: mean age 15.7 years (psychotic group) / 16.5 years (control group)</td>
<td>63 (infantile psychosis) 63 (control)</td>
<td>Follow-up study 2 measurements Medical or case records, assessments, structured interviews</td>
<td>Childhood assessment before the onset of any signs of pubescence</td>
<td>5–15 years</td>
<td>None (psychotics) 3% controls</td>
</tr>
<tr>
<td>2</td>
<td>Lotter (1974) [29]</td>
<td>England</td>
<td>Individuals showing appreciable evidence of autism syndrome. Autistic group (32) with most marked criterion behavior and comparison group (22) with similar but less marked features. Gender: not reported Age: 16–18 years</td>
<td>54 (32 autistic/ 22 non-autistic)</td>
<td>Follow-up study 2 measurements Medical or case records, structured interviews</td>
<td>Childhood assessment at 8–10 years</td>
<td>8 years</td>
<td>7%</td>
</tr>
<tr>
<td>3</td>
<td>Rumsey et al. (1985) [30]</td>
<td>United States</td>
<td>Individuals with autism Gender: 14 males (100%) Age: 18–39 years</td>
<td>14</td>
<td>Follow-up study Medical or case records, assessments, structured interviews</td>
<td>Unclear</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>4</td>
<td>Wolf and Goldberg (1986) [31]</td>
<td>Canada</td>
<td>Autistic individuals diagnosed between 1960 and 1973 Gender: not reported Age: 31% &lt;20 years 61% 20–30 years 7% &gt;30 years</td>
<td>80</td>
<td>Follow-up study 2 measurements Medical or case records, questionnaires</td>
<td>Childhood assessment between 1960–1973 at 1–15 years of age</td>
<td>8–24 years</td>
<td>20%</td>
</tr>
<tr>
<td>5</td>
<td>Szatmari et al. (1989) [32]</td>
<td>Canada</td>
<td>High-functioning individuals diagnosed with autism, childhood schizophrenia or childhood psychosis before age 5 and an last IQ score of above 65 Gender: 12 male/4 female Age: 17–34 years</td>
<td>45</td>
<td>Follow-up study 2 measurements Medical or case records, assessments, structured interviews</td>
<td>Childhood assessment of children born in 1970 and diagnosed before age 5</td>
<td>Variable 11–27 years</td>
<td>64%</td>
</tr>
<tr>
<td>6</td>
<td>Fombonne et al. (1989) [33]</td>
<td>France</td>
<td>Individuals diagnosed with childhood psychosis Gender: 77 male/22 Female Age: 20–38 years (mean 27 years)</td>
<td>227 (Childhood psychosis n = 55)</td>
<td>Follow-up study 2 measurements Questionnaires</td>
<td>Diagnosed during childhood</td>
<td>Range 6–25 years</td>
<td>56%</td>
</tr>
<tr>
<td>7</td>
<td>Kobayashi et al. (1992) [3]</td>
<td>Japan</td>
<td>Autistic individuals diagnosed as children. Gender: 170 male/31 female Age: 18–33 years</td>
<td>201</td>
<td>Follow-up survey 2 measurements Medical or case records, questionnaires</td>
<td>Diagnosed during early childhood or school age</td>
<td>Range 5–28 years</td>
<td>13%</td>
</tr>
<tr>
<td>No.</td>
<td>Study</td>
<td>Country</td>
<td>Population (diagnosis, gender and age)</td>
<td>Numbers enrolled</td>
<td>Design</td>
<td>Baseline at Diagnosis</td>
<td>Time to follow-up</td>
<td>% Lost to follow-up</td>
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</table>
| 8   | Ruble and Dalrymple (1996) [34] | United States | Individuals diagnosed with autism meeting DSM-III-R criteria  
Gender: 33 male/13 female  
Age: mean age 8.5 years (range 2–19) (T1) mean age 17.1 years (range 7–26) (T2) | 46               | Follow-up study (retrospective)  
2 measurements  
Medical or case records, structured interviews | Diagnosed in childhood (mean age at diagnosis 5.2 years) | Mean follow-up time 8.6 years | –                   |
| 9   | Ballaban-Gil et al. (1996) [4] | United States | Adolescents and young adults with autistic disorder  
Gender: not reported  
Age: 12 years or older (T2) | 163             | Follow-up study  
2 measurements  
Medical or case records, structured interviews | Childhood evaluation from May 1966 to May 1988: mean age 5 years and 2 months | Range 3.2–22.7 years | 39%                 |
| 10  | Larsen and Mouridsen (1997) [5] | Denmark      | Individuals considered to be either psychotic or borderline cases and diagnosed with Pervasive Developmental Disorder (childhood autism or Asperger syndrome)  
Gender: 10 male/8 female  
Age: 32–44 years | 18              | Follow-up register study  
2 measurements  
Medical or case records | Childhood assessment between 1949–1970 | 30 years | 11%                 |
| 11  | Howlin et al. (2000) [35]     | United Kingdom | Individuals (all male) with autism or with developmental language disorders  
Gender: 47 male (100%)  
Age: 23–24 years on average | 47              | Follow-up study (comparative)  
2 measurements  
Assessments, structured interviews | Childhood assessment at 7–8 years | Around 15 years | 17%                 |
Gender: 61 males/7 females  
Age: 21–49 years | 79 (68 participants) | Follow-up study  
2 measurements  
Medical or case records, assessments, structured interviews | Childhood assessment prior to 16 years (range 3–15 years) | Variable (minimal 6 years) | 14%                 |
| 13  | Cederlund et al. (2008) [36]  | Sweden       | Individuals with Asperger Syndrome and normal intelligence (IQ >70) and individuals diagnosed with autism or atypical autism before age 10 with different IQ levels  
Gender: 140 male (100%)  
Age: 16–38 years of age | 140             | Follow-up study (prospective)  
2 measurements  
Medical or case records, assessments, structured interviews | AS group: Diagnosed between 1985–1999 at ages 5.5–24.4 years and born 1967–1988  
Autism group: diagnosed before age 10 | More than 5 years | 30.0% in AS group  
16.7% in autism group |
<table>
<thead>
<tr>
<th>No</th>
<th>Study</th>
<th>Country</th>
<th>Population (diagnosis, gender and age)</th>
<th>Numbers enrolled</th>
<th>Design</th>
<th>Baseline at</th>
<th>Time to follow-up</th>
<th>% Lost to follow-up</th>
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<tbody>
<tr>
<td>14</td>
<td>Eaves and Ho (2008) [37]</td>
<td>Canada</td>
<td>Young adults born from 1974–1984 and diagnosed with ASD</td>
<td>48</td>
<td>Follow-up study 3 measurements</td>
<td>Diagnosed as preschoolers</td>
<td>Unknown</td>
<td>37%</td>
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<td></td>
<td></td>
<td></td>
<td>Gender: 37 males/11 females</td>
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<td>Assessments, structured interviews</td>
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<td></td>
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<td></td>
<td>Age: mean age 6.8 (range 3–12) (T1)</td>
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<td></td>
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<td>mean age 11.4 (range 8–17) (T2)</td>
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<td></td>
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<td></td>
<td>mean age 24 (T3)</td>
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<tr>
<td>15</td>
<td>Farley et al. (2009) [38]</td>
<td>United States</td>
<td>Individuals diagnosed with AD and an IQ ≥70</td>
<td>75</td>
<td>Follow-up study Assessments, structured interviews</td>
<td>Survey between 1984 and 1988</td>
<td>15–35 years</td>
<td>47%</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>Gender: 38 males/3 females</td>
<td></td>
<td>Assessments, structured interviews</td>
<td>Childhood assessment except 1 participant</td>
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<td>Age: mean age 7.2 (range 3.1–25.9) (T1)</td>
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<td></td>
<td>mean age 32.5 (range 22.3–46.4) (T2)</td>
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<tr>
<td>16</td>
<td>Whitehouse et al. (2009) [39]</td>
<td>United Kingdom</td>
<td>Young adults with a childhood history of Specific Language Impairment or Pragmatic Language Impairment or with high functioning Autism Spectrum Disorder</td>
<td>49</td>
<td>Follow-up study 2 measurements</td>
<td>Childhood assessment of children attending special speech and language schools</td>
<td>Not known</td>
<td>33%</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Gender: 35 male/14 female</td>
<td></td>
<td>Assessments, structured interviews</td>
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<td></td>
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<td></td>
<td>Age: 16–31 years</td>
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<tr>
<td>17</td>
<td>Taylor and Seltzer (2010) [40]</td>
<td>United States</td>
<td>Youths with ASD who had exited the school system between 2004 and 2008</td>
<td>66</td>
<td>Follow-up study 5 measurements</td>
<td>Families of adolescents and adults with ASD of 10 years or older in 1998</td>
<td>10 years</td>
<td>Subsample of longitudinal study Not applicable</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Gender: male 80%</td>
<td></td>
<td>Structured interviews, questionnaires</td>
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<td></td>
<td></td>
<td></td>
<td>Age: 19–26 years</td>
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<tr>
<td>18</td>
<td>Billstedt et al. (2010) [41]</td>
<td>Sweden</td>
<td>Individuals with autistic disorder/infantile autism or autistic-like conditions/atypical autism diagnosed before 10 years of age</td>
<td>120</td>
<td>Follow-up study (prospective population-based) 2 measurements</td>
<td>Childhood evaluation of children born in 1962–1984</td>
<td>Range 13–22 years</td>
<td>10%</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Gender: 84 males/36 females</td>
<td></td>
<td>Structured interviews</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>Age: mean age 25.5 years (range 17–40 years)</td>
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</tbody>
</table>
Personal Factors

Gender

In two studies gender was mentioned as a predictor for outcome, in that females might be more likely to have a poor outcome than males [7, 31]. In a third study [33] female gender was not found to be a hindering factor for positive outcome.

Intelligence

Higher IQ facilitates a positive work outcome [3–7, 28, 31–34, 36–38, 40, 41]; see also [22, 44]. One study reported that all individuals involved in competitive employment had an IQ above 70 [31] and another reported that individuals with a stable IQ above 70 were more often in some form of employment [7]. Individuals without intellectual disability were three times more likely to be competitively employed than individuals with an intellectual disability [40]. Higher IQ was significantly correlated to having a daily occupation [41]. According to Howlin [6, 7] individuals of higher IQ in general had a better outcome and problems were less pervasive (see also [4, 28, 30]). IQ <50 is often associated with poor outcome [28]. Fombonne et al. [33] found a significant worse outcome for the group with an IQ of 80 or below. In the study of Larsen and Mouridsen [5] normal intelligence predicted good outcome.

Language/Speech

Language abilities and level of useful speech may influence outcome in that better linguistic abilities might support better outcome [3, 4, 6, 7, 28, 31, 38, 39]. However, speech may be highly correlated with IQ [7, 32]. Howlin compared an autism group with a developmental receptive language disorder group and found that early language abilities appeared to be closely related to later adult functioning in the autism group [6]. Kobayashi reported that the positive effect of early speech development only occurs in males and not in females [3]; Rutter found that the level of speech at 5 or 6 years of age was closely related to IQ and low IQ contributes significantly to poor outcome [28].

Maladaptive Behavior

The presence of odd, challenging or ritualistic behavior, including self-injury, aggression and uncooperative behaviors, interferes with daily functioning [3, 4, 6, 7, 28, 30, 32, 34, 38–40]. Individuals in post-secondary education or competitively employed had significantly lower levels of maladaptive behaviors than individuals receiving day services [40]. Szatmari found a high correlation between adaptive behavior and IQ [32]. According to some authors behavioral difficulties can be a critical limiting factor for functioning successfully in employment [4, 30].

Social Impairments

The presence of social impairments, the lack of social skills and empathy are associated with poor outcome [4, 7, 29, 30, 32, 39]. It is suggested that social impairments are likely to affect the ability of individuals with autism to find and remain in meaningful employment [45].

Education

The relationship between education and employment for individuals with autism seems to be ambiguous. The majority of people with autism have attended special education services and many left school without any formal qualifications [6, 7, 28, 30, 33, 36, 37, 39]. However, people with high functioning autism have more often completed post-secondary education than other individuals with ASD [36]. In Lotter’s study [29] all individuals with good and fair outcome had had at least 7 years of education. In spite of the educational attainment of high-functioning individuals, few of them were competitively employed and if employed often in routine jobs [29, 30].

Lack of Drive

Underactivity, lack of drive and lack of initiative often hinder people with ASD to find competitive employment [28]; see also [29, 30]. Lotter [29] mentioned three necessary requirements for being able to participate in regular employment: practical competence (e.g. literacy, practical skills), social competence (being able to relate to people in a meaningful way) and intentional competence (e.g. taking initiative, motivation).

External Factors

Family

Parents play a major role in the outcome of their children with ASD. Many individuals with ASD continue to live with their family well into adulthood. According to Wolf and Goldberg [31] 87 percent of the individuals residing at home were involved in schools, workshops or independent work, compared to 46 percent in institutions.

Seven articles mentioned parents searching for job opportunities and finding jobs for their children or providing a job in a family business rather than finding a job through the open job market [3, 6, 7, 28–30, 32]. Howlin et al. [7] commented that for individuals to be able to
function adequately as adults the degree of support offered by families, social services and work environment may be as important as intellectual ability.

**Institutionalization**

Institutionalization (i.e. hospitalization) hinders a positive outcome of individuals with ASD. Especially the lower functioning individuals are living in residential care, like special institutions and hospitals where staff can attend to their specific needs. Also quite a few individuals with ASD were part of day time programs in a specialized setting [5–7, 28, 30, 31, 36]. These settings might not be the stimulating environment people need to be able to grow in their competences and work skills, although this applies to individuals with ASD as well as without [8, 16].

**Work Outcome**

The selected studies used different, but comparable, outcome measures regarding work participation and overall social outcome (incorporating education/employment, independent living and social relationships). Jobs were generally low level, unskilled and low pay jobs [4, 7, 30, 37]. Some individuals, however, managed to find a higher level job. Most individuals received special assistance in finding employment.

Few reasons are given for individuals previously employed but no longer participating in work. Rumsey [30] mentioned one individual was fired because of inappropriate social behavior. Kobayashi [3] mentioned conflicts with fellow employees, financial crisis, motivation, hospitalization and other personal circumstances (death of a parent) as causes for quitting a job. Larsen and Mouridsen [5] mentioned loss of supportive parents, divorce and factories closing down as hindering factors for finding permanent employment.

**Conclusion and Discussion**

This study identified seventeen factors related to work outcome of people with ASD. Most of these factors are of importance for all individuals with or without autism. However, it may not be just one single factor, but the combination that leads to limited employment outcomes. Especially in individuals with ASD were a combination of these factors occurs frequently. Some of these factors may be interdependent, making interpretation of the results more complex. For example, some studies found high correlations between IQ and language abilities and IQ and adaptive behavior in individuals with ASD. The disorder related characteristics (intensity of autistic symptoms, psychiatric comorbidity and epilepsy) and personal characteristics (limited language abilities, behavioral problems, social impairments) typical for ASD are factors which may, separately or combined, hinder individuals with ASD to participate in work in a sustainable way. Rates of
<table>
<thead>
<tr>
<th>No</th>
<th>Study</th>
<th>Factors (independent variables)</th>
<th>Type of work outcome</th>
<th>Outcome</th>
<th>Type of work</th>
</tr>
</thead>
</table>
| 1  | Rutter et al. (1967) [28] | (1) Diagnosis of autism (D)  
(2) Severity of disorder (D)  
(3) Evidence of brain injury (D)  
(4) Intelligence (P)  
(5) Gender (P)  
(6) Useful speech at age 5 (P)  
(7) Response to sounds (P)  
(8) Underactivity/lack of drive/lack of initiative (P)  
(9) Schooling (E)  
(10) Family situation | Employment (psychotics)  
Paid jobs n = 2  
Unpaid work n = 1  
Family business n = 1  
Day time activity n = 3 | (8) Underactivity, lack of drive and lack of initiative was often the chief factor preventing employment.  
Paid work n = 2  
Unpaid typing and duplicating at home n = 1  
Helping in father’s shop n = 1  
Various jobs n = 3  
Job following attendance Industrial Rehabilitation Unit n = 1  
Regular work n = 1 |
| 2  | Lotter (1974) [29] | (1) Amount of Schooling (E)  
(2) Age excluded from school (E)  
(3) Age sent away from home (E) | Employment/placement history  
Autistic group:  
Employed n = 1  
Special school n = 7  
Training centre n = 5  
At home n = 2  
Long stay hospital n = 14 | – | Employed n = 1 |
| 3  | Rumsey et al. (1985) [30] | (1) Psychiatric disorders (D)  
(2) Stereotyped, repetitive and compulsive behavior (P)  
(3) Impairments in social behavior (speech and nonverbal communication) (P)  
(4) Parents (E) | Employment  
Competitive employment (routine jobs) n = 4  
Sheltered employment n = 3  
Job training n = 3  
Education n = 1  
Day program n = 1  
Unemployed n = 2 | (1) One patient’s oppositional personality constituted an interfering factor for job success  
(2) One patient’s compulsive habits, and rigidity constituted interfering factors for job success  
(3) One patient’s obsessional questioning constituted an interfering factor for job success  
(3) One high functioning patient was fired because of his compulsive touching of other people and other inappropriate, intrusive social behavior  
(4) “Parent factors” were influential in determining employment outcome. Parents played a major role in finding employers willing to give their sons a chance.  
Janitor n = 1  
Cab driver n = 1  
Library aid n = 1  
Key punch operator n = 1 |
| 4  | Wolf and Goldberg (1986) [31] | (1) Age of onset of symptoms (D)  
(2) Intelligence (P)  
(3) Gender (P)  
(4) Acquisition of speech for communication (P)  
(5) Living situation (home–institution) (E) | Employment  
Independent work n = 5  
Sheltered Workshop n = 10  
Education n = 23  
Day program n = 21  
No program n = 5 | (2) The autistic adults involved in competitive employment all had an IQ above 70.  
Competitive employment n = 4  
Group employment outside institution n = 1 |
<table>
<thead>
<tr>
<th>No</th>
<th>Study</th>
<th>Factors (independent variables)</th>
<th>Type of work outcome</th>
<th>Outcome</th>
<th>Type of work</th>
</tr>
</thead>
</table>
| 5  | Szatmari et al. (1989) [32]   | (1) Aspects of cognition (P)  
(2) Impairments in social behavior (P)  
(3) Deviant language (P)  
(4) Bizarre behaviors (P) | Occupation or placement  
Paid employment n = 6  
Family business n = 1  
Sheltered work n = 4  
Education n = 3  
Unemployed n = 2 | – | Teacher-tutor n = 1  
Librarian n = 1  
Salesman n = 2  
Library technician n = 1  
Factory n = 1  
Family business n = 1  
Workshop n = 4 |
| 6  | Fombonne et al. (1989) [33]   | (1) Age of admission (P)  
(2) Length of stay (E)  
(3) Gender (P)  
(4) Intelligence at time of admission (P)  
(5) Treatment (E) (a.o. psychotherapie, speech therapy, remedial gymnastics)  
(6) Medication | Employment  
55% | – | – |
| 7  | Kobayashi et al. (1992) [3]    | (1) IQ at age 6 (P)  
(2) Level of speech development at age 6 (P) | Employment (21.8%)  
Paid jobs (mainly manual or industrial workers) n = 41  
Family business n = 2  
Education n = 11  
Sheltered workshop n = 27  
At home n = 18 | – | Laundry n = 4  
Bus conductor n = 1  
Chikuwa maker n = 1  
Paper maker n = 1  
Food maker n = 9  
Tatami maker n = 2  
Civil servant/office worker n = 3  
Auto mechanic n = 1  
Helper n = 2  
Industrial worker n = 9  
Physical therapist n = 1  
Printer n = 1  
Trash collector n = 2  
Tile roofer n = 1  
Confectionary maker n = 1  
Construction/Assistant plasterer n = 2  
Dressmaker n = 1 |
| 8  | Ruble and Dalrymple (1996) [34]| (1) Cognitive level (IQ) (P)  
(2) Communication (P)  
(3) Challenging behavior (P) | Employment (adults)  
Supported employment n = 1  
Sheltered employment n = 7  
Daily living programs n = 3  
No program n = 4  
In institution n = 2 | – | – |
Table 4 continued

<table>
<thead>
<tr>
<th>No</th>
<th>Study</th>
<th>Factors (independent variables)</th>
<th>Type of work outcome</th>
<th>Outcome</th>
<th>Type of work</th>
</tr>
</thead>
<tbody>
<tr>
<td>9</td>
<td>Ballaban-Gil et al. (1996) [4]</td>
<td>(1) Intelligence (P) (2) Language (P) (3) Behavior (P) (4) Social deficits/impairment (P)</td>
<td>Employment adults (n = 45) Open employment (menial jobs) n = 5 Sheltered workshops n = 6 Self employment n = 2 Education n = 7</td>
<td>–</td>
<td>Only 11% of adults were employed on the open market, all in menial jobs such as stock boy or mail clerk (n = 5) Sheltered workshops n = 6</td>
</tr>
<tr>
<td>10</td>
<td>Larsen and Mouridsen (1997) [5]</td>
<td>(1) Diagnosis of autism (D) (2) Intensity of autistic symptoms (D) (3) Intelligence (P) (4) Psychiatric morbidity (D) (5) Pharmacotherapy (E)</td>
<td>Employment Asperger group: Paid job n = 1 Sheltered employment n = 2 Disability pension n = 5 Childhood autism group: Paid job n = 2 Sheltered employment n = 1 Daytime program n = 5</td>
<td>(1) In middle adulthood the Childhood Autism group has a much poorer outcome regarding education and employment than the Asperger group.</td>
<td>Insulator n = 1 Porcelain painter n = 1 Kindergarten teacher n = 1 Received vocational training n = 4 Before working as a driver/fish industry n = 1 Fully-paid unskilled work before n = 4</td>
</tr>
<tr>
<td>11</td>
<td>Howlin et al. (2000) [35]</td>
<td>(1) Diagnosis of autism (2) Intelligence (P) (3) Psychiatric problems (D) (4) Early language abilities (P) (5) Autistic-like stereotyped and repetitive behavior patterns (P)</td>
<td>Education and employment histories Autism group (n = 19): Independent jobs n = 1 Fulltime education n = 2 Voluntary work n = 3 Daytime centres n = 12 No occupation n = 1</td>
<td>–</td>
<td>Laboratory technician n = 1</td>
</tr>
<tr>
<td>12</td>
<td>Howlin et al. (2004) [7]</td>
<td>(1) Childhood IQ (P) (2) Gender (P) (3) Speech at 5 years (P) (4) Autistic-type behaviors (P) (5) Social functioning (P)</td>
<td>Employment Independent jobs n = 8 Self employed n = 1 Sheltered employment n = 11 Daily activities by centre n = 15 Family based work activities n = 2 Voluntary work n = 1 No work activities n = 28 Not known n = 2</td>
<td>(1) Individuals with a stable IQ from childhood to adulthood above 70 were more often in some form of employment (paid, voluntary or sheltered) (P = .005)</td>
<td>Scientific officer oil company n = 1 Electrical work n = 1 Cartographer n = 1 Postal assistant n = 1 Factory work n = 5 Computing n = 1 Accounts n = 1 Fabric design n = 1 Washing up n = 1 Grave digger n = 1 Office/accounts assistant n = 1 Charcoal burning/gardening n = 1 Administrative assistant n = 1 Data input n = 1 Supermarket trolleys n = 1 Electronic work n = 1 Special shop n = 1 Decorating with father n = 1 Office with parents n = 1</td>
</tr>
<tr>
<td>No</td>
<td>Study</td>
<td>Factors (independent variables)</td>
<td>Type of work outcome</td>
<td>Outcome</td>
<td>Type of work</td>
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</tr>
</tbody>
</table>
| 13 | Cederlund et al. (2008) [36] | (1) Diagnosis of autism (D)  
(2) Intelligence (P)  
(3) Psychotic disorder (D) | Employment | – | Ordinary jobs n = 8  
Daily occupational activities n = 10 |
| 14 | Eaves and Ho (2008) [37] | (1) Diagnosis (autism score) in adolescence (D)  
(2) Childhood and adolescence intensity of autistic symptoms (CARS) (D)  
(3) Childhood and adolescence verbal and Performance IQ (P) | Employment | Delivering papers  
Meals on wheels  
Sorting recycle | |
| 15 | Farley et al. (2009) [38] | (1) Psychiatric disorders (D)  
(2) Epilepsy (D)  
(3) Other medical disorders (D)  
(4) Historical full scale IQ (P)  
(6) Adaptive behavior (P) | Employment | Independent paid jobs n = 22  
Supported employment n = 3  
Voluntary work n = 2  
Day programs n = 10  
Unemployed n = 4 | |
| 16 | Whitehouse et al. (2009) [39] | (1) Diagnosis of autism (D)  
(2) Intensity of autistic symptoms (D)  
(3) Psychiatric problems (D)  
(3) Language ability (pragmatic or structural problems) (P)  
(5) Social impairments | Employment | Competitive:  
Factory workers n = 2  
Cleaners n = 3 | |
| 17 | Taylor and Seltzer (2010) [40] | (1) Autistic symptoms (D)  
(2) Intellectual disability (P)  
(3) Comorbid psychiatric diagnoses (D)  
(4) Maladaptive behaviors (P)  
(5) Functional independence (P)  
(6) Family income (E) | Employment | Competitive:  
Bus boy  
Replacing dirty glasses with clean ones  
Salvation Army  
Read business (self employed)  
Supported:  
Rolling silverware into napkins in restaurant  
Folding towels in hotel  
Shredding confidential information  
Washing dishes at a nursing  
Working in a grocery store | |

### Notes:
- **Asperger group:** ordinary jobs n = 7  
  “daily occupational activities” in a group centre n = 6  
  no organized daily activity n = 12  
- **Autism group:** ordinary job n = 1  
  “daily occupational activities” in a group centre n = 4  
  regular individually tailored daily activities n = 33  
  no organized daily activity n = 13
- 56% (n = 27) had ever been employed, most in volunteer, sheltered or part time work
- Independent job n = 2  
  Daytime activity n = 19 (40%)
- 1 In spite of high IQ scores and adequate practical skills, some participants were unable to seek employment due to difficulties with anxiety.
- 1 Stable employment proved to be an area of difficulty for the ASD group.
employment among individuals with ASD are generally low. Often the impairments and social deficits of these individuals are emphasized leading to low expectations regarding outcome. However, these individuals may have strengths (e.g. ability to concentrate; strong focus) that can be utilized if the right tasks and settings are provided [22].

In some of the studies Asperger syndrome and Childhood Autism were separately analyzed. There is a continuing discussion whether it is possible and necessary to distinguish between childhood autism and Asperger syndrome [22, 46, 47]. A pronounced autistic disorder often leads to substantial limitations in participation in work; people with Asperger Syndrome often achieved higher education and have more abilities to work compared to childhood autism. However, this advantage in education does not always lead to higher levels of employment in later life [46].

IQ is the only childhood predictor of work outcome for which we found consistent evidence in the literature in that a higher IQ facilitates a positive work outcome. Although an IQ below 50 does almost always lead to a poor outcome [7] and this applies to individuals without ASD as well [48, 49], individuals with an IQ of 70 or higher do not necessarily have a good outcome. Outcome in individuals without intellectual disability is much more variable and less predictable. Therefore, it seems that the clinical value of IQ in predicting individual outcomes is limited.

Although education is often mentioned as an important factor for outcome, job level is rarely consistent with educational background. Also the increase in educational services for children with ASD has not necessarily led to improved outcome when they have grown up [7]. As access to education can be closely associated with the IQ of the individual, this relationship must be regarded with caution [18]. Nevertheless there is some evidence that the amount of schooling received, positively influences social adjustment in later life [18, 49].

Besides disorder-related and personal factors, several external factors are related to work outcome. Considering the low levels of independence of individuals with ASD, the degree of support offered by families, the available support services and the willingness of employers to incorporate this group in their work force may be as decisive for individuals to be able to function adequately in employment as the personal factors mentioned above [7, 50]. Especially parents play an important role in supporting their children as they continue to live with them well into adulthood, in searching for job opportunities and in being advocates for their child’s well-being [6, 7].

Competitive paid employment is often regarded as successful participation. Because of increasing demands in work, employers are hesitant to hire individuals with disabilities. If working, many individuals with ASD work in unskilled, routine, industrial jobs with limited decision
Table 5  Overview of factors associated with outcome

<table>
<thead>
<tr>
<th>Prognostic factors (independent variables)</th>
<th>Study</th>
<th>Significance</th>
<th>Quality of study</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Disease/disorder related</strong></td>
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<tr>
<td><strong>(Autism) diagnosis</strong></td>
<td></td>
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<tr>
<td>(Autism) diagnosis</td>
<td>Rutter et al. (1967) [28]</td>
<td>n.s.</td>
<td>Low</td>
</tr>
<tr>
<td></td>
<td>Howlin et al. (2000) [35]</td>
<td>–</td>
<td>Low</td>
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<tr>
<td></td>
<td>Cederlund et al. (2008) [36]</td>
<td>Sig</td>
<td>Low</td>
</tr>
<tr>
<td></td>
<td>Whitehouse et al. (2009) [39]</td>
<td>–</td>
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<tr>
<td><strong>Autism score in adolescence</strong></td>
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<tr>
<td></td>
<td>Eaves and Ho (2008) [37]</td>
<td>Sig</td>
<td>Low</td>
</tr>
<tr>
<td><strong>Age of onset of symptoms</strong></td>
<td></td>
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<tr>
<td></td>
<td>Wolf and Goldberg (1986) [31]</td>
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<tr>
<td><strong>Evidence of brain injury</strong></td>
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<td></td>
<td>Rutter et al. (1967) [28]</td>
<td>n.s.</td>
<td>Low</td>
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<td><strong>Severity of disorder</strong></td>
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<td>Rutter et al. (1967) [28]</td>
<td>Sig</td>
<td>Low</td>
</tr>
<tr>
<td><strong>Intensity of autistic symptoms</strong></td>
<td></td>
<td></td>
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</tr>
<tr>
<td></td>
<td>Wolf and Goldberg (1986) [31]</td>
<td>n.s.</td>
<td>Low</td>
</tr>
<tr>
<td></td>
<td>Eaves and Ho (2008) [37]</td>
<td>Sig</td>
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<tr>
<td></td>
<td>Whitehouse et al. (2009) [39]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td></td>
<td>Taylor and Seltzer (2010) [40]</td>
<td>Sig</td>
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<tr>
<td><strong>Comorbidity</strong></td>
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<td><strong>Psychiatric disorders</strong></td>
<td>Rumsey et al. (1985) [30]</td>
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<tr>
<td></td>
<td>Howlin et al. (2000) [35]</td>
<td>–</td>
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<td></td>
<td>Cederlund et al. (2008) [36]</td>
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<td></td>
<td>Farley et al. (2009) [38]</td>
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<td>Taylor and Seltzer (2010) [40]</td>
<td>n.s.</td>
<td>Low</td>
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<td><strong>Epilepsy</strong></td>
<td>Rutter et al. (1967) [28]</td>
<td>n.s.</td>
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<tr>
<td><strong>Other medical disorders</strong></td>
<td>Farley et al. (2009) [38]</td>
<td>–</td>
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<tr>
<td><strong>Personal factors</strong></td>
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<tr>
<td><strong>Intelligence (IQ-level)</strong></td>
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<tr>
<td><strong>Intelligence (IQ-level)</strong></td>
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<tr>
<td></td>
<td>Ruble and Dalrymple (1996) [34]</td>
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<td></td>
<td>Billstedt et al. (2010) [41]</td>
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<td><strong>Full scale IQ</strong></td>
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<td></td>
<td>Cederlund et al. (2008) [36]</td>
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<td><strong>Intelligence at time of admission</strong></td>
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<td><strong>IQ at age 6</strong></td>
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<td><strong>Performance IQ at time 1</strong></td>
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</tr>
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<td><strong>Childhood IQ</strong></td>
<td>Howlin et al. (2004) [7]</td>
<td>Sig</td>
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<td><strong>Childhood and adolescence</strong></td>
<td>Eaves and Ho (2008) [37]</td>
<td>Sig</td>
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<td><strong>verbal and performance IQ</strong></td>
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<td><strong>Historical full scale IQ</strong></td>
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<tr>
<td><strong>Gender</strong></td>
<td>Wolf and Goldberg (1986) [31]</td>
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<tr>
<td></td>
<td>Howlin et al. (2004) [7]</td>
<td>Sig</td>
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</tbody>
</table>
Table 5 continued

<table>
<thead>
<tr>
<th>Prognostic factors (independent variables)</th>
<th>Study</th>
<th>Significance</th>
<th>Quality of study</th>
</tr>
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<td>Speech and language</td>
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<td></td>
<td>Wolf and Goldberg (1986) [31]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Language ability (pragmatic or structural problems)</td>
<td>Whitehouse et al. (2009) [39]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Acquisition of speech for communication</td>
<td>Wolf and Goldberg (1986) [31]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Early language abilities</td>
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<td>Level of speech development at age 6</td>
<td>Kobayashi et al. (1992) [3]</td>
<td>Sig (males)</td>
<td>Low</td>
</tr>
<tr>
<td></td>
<td>Farley et al. (2009) [38]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>(Useful) speech at age 5</td>
<td>Rutter et al. (1967) [28]</td>
<td>Sig</td>
<td>Low</td>
</tr>
<tr>
<td>Deviant language</td>
<td>Howlin et al. (2004) [7]</td>
<td>Sig</td>
<td>High</td>
</tr>
<tr>
<td>Response to sounds</td>
<td>Rutter et al. (1967) [28]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td><strong>Maladaptive behavior</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ritualistic and compulsive behavior</td>
<td>Rutter et al. (1967) [28]</td>
<td>Descriptive</td>
<td>Low</td>
</tr>
<tr>
<td>Stereotyped, repetitive and compulsive behavior</td>
<td>Rumsey et al. (1985) [30]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Bizarre behaviors</td>
<td>Szatmari et al. (1989) [32]</td>
<td>n.s.</td>
<td>High</td>
</tr>
<tr>
<td>Challenging behaviors</td>
<td>Ruble and Dalrymple (1996) [34]</td>
<td>Descriptive</td>
<td>Low</td>
</tr>
<tr>
<td>Autistic-like stereotyped and repetitive behavior patterns</td>
<td>Whitehouse et al. (2009) [39]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Maladaptive behaviors</td>
<td>Taylor and Seltzer (2010) [40]</td>
<td>Sig</td>
<td>Low</td>
</tr>
<tr>
<td>Adaptive behavior</td>
<td>Farley et al. (2009) [38]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td><strong>Social deficits/impairment</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Social deficits/impairment</td>
<td>Rumsey et al. (1985) [30]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td></td>
<td>Whitehouse et al. (2009) [39]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Impairments in social behavior</td>
<td>Rumsey et al. (1985) [30]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>(speech and nonverbal communication)</td>
<td>Szatmari et al. (1989) [32]</td>
<td>n.s.</td>
<td>High</td>
</tr>
<tr>
<td>Underactivity/lack of drive/lack of initiative</td>
<td>Rutter et al. (1967) [28]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Functional independence (ADL)</td>
<td>Taylor and Seltzer (2010) [40]</td>
<td>Sig</td>
<td>Low</td>
</tr>
<tr>
<td><strong>External factors</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Parents</td>
<td>Rumsey et al. (1985) [30]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Family income</td>
<td>Taylor and Seltzer (2010) [40]</td>
<td>n.s.</td>
<td>Low</td>
</tr>
<tr>
<td>History mental illness parent</td>
<td>Rutter et al. (1967) [40]</td>
<td>n.s.</td>
<td>Low</td>
</tr>
<tr>
<td>Family situation (not living at home)</td>
<td>Rutter et al. (1967) [28]</td>
<td>n.s.</td>
<td>Low</td>
</tr>
<tr>
<td>Age sent away from home</td>
<td>Lotter (1974) [29]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Institutionalization</td>
<td>Wolf and Goldberg (1986) [31]</td>
<td>Descriptive</td>
<td>Low</td>
</tr>
<tr>
<td>Treatment</td>
<td>Rutter et al. (1967) [28]</td>
<td>n.s.</td>
<td>Low</td>
</tr>
<tr>
<td>Use of medication/pharmacotherapy</td>
<td>Fombonne et al. (1989) [33]</td>
<td>n.s.</td>
<td>Low</td>
</tr>
<tr>
<td>Schooling</td>
<td>Rutter et al. (1967) [28]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Amount of schooling</td>
<td>Lotter (1974) [29]</td>
<td>–</td>
<td>Low</td>
</tr>
<tr>
<td>Age excluded from school</td>
<td>Lotter (1974) [29]</td>
<td>n.s.</td>
<td>Low</td>
</tr>
</tbody>
</table>

n.s. Not significant, Sig significant
latitude and minimal social interaction [13, 30, 35]. As our economy becomes more knowledge-based, and globalization transforms and eliminates unskilled jobs, those with limited cognitive function may become increasingly marginalized [51]. Also periods of employment are alternated by periods of unemployment or temporary jobs [7]. Data of the Dutch Social Security Institute suggest that about 11% (n = 1,618 per year) of the young disabled applying for a social security benefit has ASD [52].

If employed, the majority is working part-time, sometimes less than 10 h a week [37]. Fulltime work is not always feasible for this group. For successful sustainable work participation a fit between the individual, the job and the work environment is essential [51]. This person-environment fit—or Person-Job fit when focussed on work [53]—concerns the balance between knowledge, skills, abilities, attitude and motivation of the person at the one hand and work and its context at the other hand. A situation of balance contributes to the health, well-being and work-functioning of the employee. A disbalance leads to stress and disfunctioning. We can distinguish two kinds of PE-fit: the demands-abilities fit and the needs-supply fit [54]. In people with autism both their abilities and their needs can be influenced by the disorder. From a theoretical point of view tailor-made adjustment in demands and supplies (support) may be necessary to ensure a good fit. The practice of part time work might be a reflexion of this.

Considering the severe consequences of autism and the consequential need for special attention for a tailor-made fit between individual and work characteristics, it is important that effective assessments and interventions with respect to work participation of the ASD population are available. Over the last years, special vocational re-integration services and supported employment services have been set up for individuals with ASD, because existing services are not always accessible to them as services sometimes require a basic set of skills of applicants, like interpersonal communication skills, to increase employability [11, 55]. Part of the supported employment strategy is to adapt the environment and workplace to the needs of disabled individuals who have the skills to do a certain job [56]. According to Garcia-Villamisar [8, 16] supported employment produces favorable results for people with ASD as compared to sheltered employment services with regard to severity of impairments and quality of life. Ridley and Hunter [11] reviewed the practice of supported employment in Scotland and found that the principles of supported employment are not widely and consistently applied, while adherence to these principles is related to improved employment outcomes [57]. Moreover, people with ASD have limited access to these services and unpaid and part-time jobs were more frequently achieved than paid jobs. Leadership by local authorities is needed to improve implementation of supported employment and accessibility. This supports Howlin’s [6] claim that the area where an individual lives and the available services is a major influence in outcome with regard to employment.

Autism spectrum disorders are studied extensively since the 70’s and more attention is given to social functioning. Unfortunately, only one study focussed on employment as primary outcome. Most of the studies we reviewed were descriptive in nature and thus the quality of the data is variable and often limited. Few studies were able to report significant findings. Moreover, numbers of participants in the studies were often limited. Also quite a few studies in our review consisted of clinical samples, that by the nature of their population have limited generalizing capacity, because of problems with representativeness of these samples. Due to the diverse reporting of outcome it is not possible to compare the studies or to statistically pool the data. For that same reason we did not use the quality assessment for determining levels of evidence for the factors, but to inform the reader about the quality of the studies included. If the results of high quality studies differ from the results of low quality studies, this can be an indication of bias. In our review we found conflicting results for maladaptive behavior between one high and one low quality study [32, 40].

Two early studies [28, 29] were conducted in a very different climate with regard to the employment of individuals with disability. Their results seem to indicate that work outcomes did not improve in recent years.

Recommendations

This review gives an overview of factors facilitating or hindering work participation of people with autism. Factors, identified in high quality studies, can help to provide an evidence-based ground for the development of instruments and intervention programs to increase work participation of individuals with ASD. The availability of adequate services for these individuals during their education, their transition from school to work and to independent living might influence employment outcome considerably [6, 12, 28]. The findings of this review emphasize the need for adequate intervention and services, geared to the needs of the individual with ASD, that help them to adjust to the psychosocial demands in society [39].

However, this review also painfully points to an important gap in the literature regarding predictors of work outcomes in individuals with ASD. High quality studies on predictors of work participation in individuals with ASD are lacking. Most of the included studies reported on outcome as an overall social outcome measure, including work; not on work as a primary outcome measure. In our study we assumed the seventeen factors we found are useful in predicting work outcome. However, further research should focus on work participation as the primary outcome measure in determining...
whether the factors mentioned are indeed influencing work outcome in individuals with ASD. High quality longitudinal studies are needed to identify variables that are responsive to interventions and that take the person-environment fit into account. Only then there is enough base for developing and implementing evidence based strategies to enhance optimal work participation for this group, that could benefit considerably from it in terms of quality of life.

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**Appendix**

See Table 6.

### Table 6: Operationalization of criteria list for quality assessment

| Study population | A Inception cohort
|------------------|-------------------------------------------------|
|                  | One point if participants are identified at an early uniform point in the course of their disability
|                  | Zero point if it is not clear if an inception cohort was used. |
|                  | B Description of source population
|                  | One point if the source population is described in terms of place of recruitment (for example: Groningen, the Netherlands), time-period of recruitment and sampling frame of source population (for example: health service provider, special education services).
|                  | Zero point if ≤2 features of source population are given. |
|                  | C Description of relevant inclusion and exclusion criteria
|                  | One point if ≥2 criteria are formulated. |
|                  | Zero point if ≤2 criteria are formulated. |
| Follow-up        | D Follow-up at least 12 months
|                  | One point if the follow-up period is at least 12 month and data are provided for this moment in time. |
|                  | E Drop-outs/loss to follow-up ≤20%
|                  | One point if total number of drop-outs/loss to follow up ≤20%
|                  | F Information completers versus loss to follow-up/drop-outs
|                  | One point if sociodemographic information is presented for completers and those lost to follow-up/drop outs at baseline, or no loss to follow-up/drop outs. Reasons for loss to follow-up/drop outs have to be unrelated to the outcome. Loss to follow-up/ drop outs: all participants of the assembled cohort minus the number of participants at the main moment of measurement for the main outcome measure, divided by the total number of participants of the assembled cohort. |
|                  | G Prospective data collection
|                  | One point if a prospective design is used, or a historical cohort when the prognostic factors are measured before the outcome is determined. |
|                  | Zero point if a historical cohort is used, considering prognostic factors at time zero which are not related to the primary research question for which the cohort is created, or in case of an ambispective design. |
| Treatment        | H Treatment in cohort is fully described/standardized
|                  | One point if treatment subsequent to inclusion into cohort, is fully described and standardized, or in case of no treatment is given, or if multi-variate correction for treatment is performed in analysis. |
|                  | Zero point if different treatment is given and if it is not clear how outcome is influenced by it, or if it is not clear whether any treatment is given. |
| Prognostic factors | I Relevant potential prognostic factors
|                  | One point if besides socio-demographic factors (age, gender) at least one other factor of the following is described at baseline: health related factors personal factors external factors |
|                  | J Standardized or valid measurements
|                  | One point if at least one of the factors of I, excluding age and gender, are reported in a standardized or valid way (for example: questionnaire, structured interview, register, patient-status of health service). |
|                  | K Data presentation of most important prognostic factors
|                  | One point if frequencies, or percentages, or mean (and standard deviation/confidence interval), or median (and Inter Quartile Range) are reported for the three most important factors of I, namely age, gender and at least one other factor, for the most important follow-up measurements. |
Table 6 continued

<table>
<thead>
<tr>
<th>Outcome</th>
<th>L. Relevant outcome measures</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>One point if at least one of the following outcome criteria is reported: social functioning, independent living, employment, daily life activities.</td>
</tr>
<tr>
<td>M</td>
<td>Standardized or valid measurements</td>
</tr>
<tr>
<td></td>
<td>One point if one or more of the main outcome measures of L are reported in a standardized or valid way (for example: questionnaire, structured interview, registration, patient-status of occupational/insurance physician).</td>
</tr>
<tr>
<td>N</td>
<td>Data presentation of most important outcome measures</td>
</tr>
<tr>
<td></td>
<td>One point if frequencies, or percentages, or mean (and standard deviation/confidence interval), or median (and Inter Quartile Range) are reported for one or more of the main outcome for the most important follow-up measurements.</td>
</tr>
</tbody>
</table>

Analysis

<table>
<thead>
<tr>
<th>O</th>
<th>Appropriate univariate crude estimates</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>One point if univariate crude estimates (RR, OR, HRR) between prognostic factors separately and outcome are presented.</td>
</tr>
<tr>
<td></td>
<td>Zero point if only P values or wrong association values (Spearman, Pearson, sensitivity) are given, or if no tests are performed at all.</td>
</tr>
<tr>
<td>P</td>
<td>Appropriate multivariate analysis techniques</td>
</tr>
<tr>
<td></td>
<td>One point if logistic regression analysis is used, or survival analysis for dichotomous outcomes, or linear regression analysis for continuous outcomes.</td>
</tr>
<tr>
<td></td>
<td>Zero point if no multivariate techniques are performed at all.</td>
</tr>
</tbody>
</table>

References

52. UWV. UWV Kwartaalverkenning 2009-III. 2009.