

Neuropsychological Approach on Expressed Emotion in Neurotypical and Autism Spectrum Disorder: A Path Model Analysis

Sagayaraj Kanagaraj^{1*} Kinjari Kancharla¹,
C.N. Ram Gopal¹ and Sundaravadivel Karthikeyan²

¹Faculty of Allied Health Sciences, Chettinad Hospital and Research Institute, Chettinad Academy of Research and Education, Kelambakkam, 603103, Tamil Nadu, India.

²Department of Clinical Psychology and Medical Science, National Institute for Empowerment of Persons with Multiple Disabilities, Chennai, 603112, Tamil Nadu, India.

*Corresponding Author E-mail: harrysagayaraj@gmail.com

<https://dx.doi.org/10.13005/bpj/2385>

(Received: 08 October 2021; accepted: 25 March 2022)

Autism Spectrum Disorder (ASD) is a neurodevelopmental condition that affects individual social communication with a range of restricted behaviour patterns. People with ASD will also have difficulties with social emotional reciprocity, which is not predominantly found in neurotypical individuals. Individuals with ASD have difficulty connecting with neurotypical (i.e., nonautistic) people because they fail to identify other people's emotions and mental states. Alexithymia is a personality characteristic defined by a subclinical inability to identify and explain one's own emotions. Alexithymia is defined by a significant dysfunction in emotional awareness, social attachment, and interpersonal relationships. It is distinguished by impaired emotional awareness, which has been increasing in diagnostic frequency in a variety of neuropsychiatric diseases, with notable overlap with ASD. To empirically measure the condition of alexithymia in neurotypical individuals (N = 12) and people diagnosed with ASD (N = 12), were assessed with the Observer Alexithymia Scale (OAS) by Haviland et al., 2000. The mean age of the neurotypical is (M = 21.67; SD = 2.60) and the ASD is (M = 18.33; SD = 2.22). Using SPSS ver.20, the data was analysed using descriptive and inferential statistics methods. The results indicate the significant difference between neurotypical and autism spectrum disorder individuals with the condition of alexithymia. The path model, which was drawn from the SPSS AMOS version 20, emphasises the causal relationship between variables of interest from the Observer Alexithymia Scale. This study found that individuals with ASD have significant corroboration to alexithymia when compared to neurotypical individuals.

Keywords: Alexithymia; Autism Spectrum Disorder; Emotional Expression; Emotional Reciprocity; Neurodevelopmental; Neurotypical.

Autism Spectrum Disorder (ASD) is commonly referred to as a pervasive developmental disorder, which indicates that it occurs in early childhood and affects many aspects of a child's development. However, there is now a growing

understanding that autism should not be considered as a condition but rather as a neurological diversity. This is why the terms "neurodiverse" and "neurodivergent" are used to refer to ASD¹. The revised diagnosis criteria for ASD are included in an

update to the World Health Organization's manual, "International Classification of Disease - eleventh revision (ICD-11)". Autism, asperger syndrome, and pervasive developmental disorder - not otherwise specified (PDD-NOS) are all classified as ASD in the most recent version of the ICD-11 manual. Similar to the Diagnostic and Statistical Manual of Mental Disorders, Fifth Edition (DSM-5), the ICD-11 divides ASD symptoms into two categories: difficulty with social communication and social engagement; and repetitive or restricted pattern of behaviour². People with ASD, in particular, have been considered challenged by their capacity to understand other people's viewpoints or attribute mental states to other people (often referred to as mind-blindness or a lack of theory of mind) and to have a lack of empathy. There is now a lot of evidence that ASD is largely caused by neuro-biological abnormalities and difficulties. That is, the typical social, communicative, and repetitive behaviours associated with autism are the developmental outcomes of a brain that is fundamentally wired and structured differently³.

Alexithymia is a personality trait that causes difficulty recognising and reacting to emotions in oneself or others. However, the symptoms of alexithymia are not as severe as those associated with conditions such as ASD. Alexithymia is usually related to various mental health issues and developmental disabilities, most notably ASD. The difficulty in recognising, perceiving, or reacting to emotions is a distinguishing characteristic of alexithymia. This can lead to issues with empathy, difficulties with conflict resolution, and many people with alexithymia also struggle in their relationships with partners, friends, and family⁴.

Researchers do not know the aetiology of Alexithymia. It is most probably the result of a combination of genetic and environmental factors. Children with alexithymia are more likely to have difficulty expressing socially acceptable emotions. Children who do not develop a constant connection with others or an understanding of feelings are more likely to struggle with emotional awareness later in life, because alexithymia frequently co-occurs with other mental health disorders. Individuals who develop the ability to perceive, explain, and react to emotions may

experience less alexithymia symptoms and have fewer interpersonal relationship difficulties⁵.

Individuals with high alexithymia scores have difficulties recognising and contextualising their internal emotional states, discriminating between numerous forms of the same subjective emotion, and communicating their emotional states to others. These people also have fewer imaginal processes and think in a stimulus-bound, externally focused manner (i.e., concrete thinking). Alexithymia is not a psychiatric diagnosis and of itself; rather, it is a three-dimensional personality trait that occurs to varying degrees in the general population and is linked to a number of medical, mental, and psychological issues⁶⁻¹⁰. The purpose of this research was to get a thorough knowledge of alexithymia and to compare it to those who are normally developing and those who have ASD. This comparative research will look at the outcomes of expressed emotion, which includes recognising, perceiving, and regulating emotion in neurotypical and people with ASD.

MATERIALS AND METHODS

Hypothesis

H₀ There will be no significant difference between neurotypical and autism spectrum disorder individuals in the condition of alexithymia.

Sample Descriptions

A total of 24 participants took part in this study which include (Neurotypical N = 12), (Autism Spectrum disorder = 12). The mean age of neurotypical is (M = 21.67; SD = 2.60) with (Male N = 5; Female N = 7). The mean age of autism spectrum disorder individuals is (M = 18.33; SD = 2.22) with (Male N = 8; Female = 4). The participants from the autism spectrum disorder group were already been clinically diagnosed with the Indian Scale for Assessment of Autism Disorder (ISAA)¹¹ and found to have mild and moderate levels of autism from the assessment.

Tools Used

The Observer Alexithymia Scale (OAS) by Haviland et al., 2000.¹² (33 items) was used to collect information from the participants. It is a 4-point likert scale ranging from 0 = never; 1 = sometimes; 2 = usually; and 3 = all of the time. The scale has a test-retest reliability of (rtt) = 0.87

and an internal consistency of Cronbach alpha ranging from 0.88 to 0.89. The scale is validated with exploratory and confirmatory factor analysis. The scoring was computed as per the manual of the OAS. Higher the score indicate the high level of having the alexithymia condition.

Procedure

It is an observer measurement scale, so the acquaintances or relatives of the participants were asked to rate the subjects who took part in the study. The participants were informed about the purpose and significance of the study. Also, their anonymity and confidentiality were protected.

Ethical Consideration

This study was carried out with the approval of the Institutional Human Ethics Committee of Chettinad Academy of Research and Education (216/IHEC/Nov/2020), Kelambakkam, Chennai, Tamil Nadu, India as a part of supporting the doctoral research.

Data Analysis

The analysis of the collected data was computed using descriptive statistics mean (M),

standard deviation (SD); and inferential statistics independent t test by using SPSS ver 20. The path model was drawn by using SPSS AMOS Ver 20.

RESULTS AND DISCUSSION

While the current study’s goal is to empirically investigate the expressed emotions of neurotypical and autism spectrum disorder participants, other factors such as social background, economic background, and gender differences were not taken into account. Table 1 indicates the gender and age-wise details of the neurotypical participants from the study. It consists of male participants (N = 5; Age M = 21.81; SD = 2.41) and female participants (N = 7; Age M = 21.14; SD = 2.45). Table 2 indicates the gender and age-wise details of the autism spectrum disorder participants from the study. It has male participants (N = 8; age M = 18.15; SD = 2.34) and female participants (N = 4; age M = 18.00; SD = 1.58).

From Table 3, it was found that the Observer Alexithymia Scale (OAS) score of

Table 1. Details of the Neurotypical Participants in Percentage

Gender	Percentage	
Male	5	42%
Female	7	58%
Age		
18 years	1	8%
19 years	2	17%
20 years	2	17%
21 years	1	8%
22 years	2	17%
24 years	2	17%
25 years	1	8%
26 years	1	8%

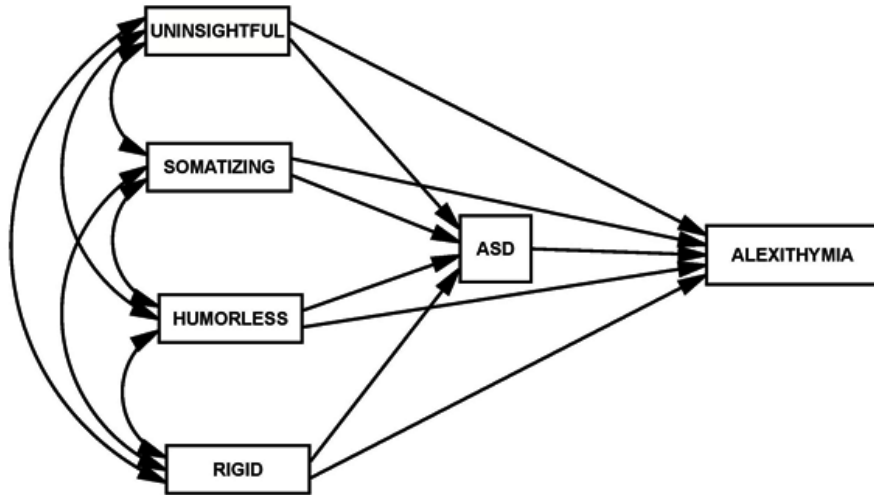
Table 2. Details of Autism Spectrum Disorder Participants in Percentage

Gender	N	Percentage
Male	8	67%
Female	4	33%
Age		
16 years	2	17%
17 years	3	25%
18 years	3	25%
19 years	1	8%
20 years	2	17%
24 years	1	8%

Table 3. Comparison between Neurotypical and Autism Spectrum Disorder Participants

	Mean	SD	T-Value	Sig. (2-tailed)	Mean Difference	95 % Confidence Interval of the Difference	
						Lower	Upper
Neurotypical	33.08	9.33	15.7*	.000	33.08	27.15	39.01
Autism Spectrum Disorder	78.66	3.67			78.66	76.33	81.00

* The result is significant at p < .05. Confidence Interval 95 %



ASD = Autism Spectrum Disorder
 Alexithymia : Condition assessed by the Observer Alexithymia Scale Haviland et al., 2000

Fig. 1. Path Model of the ASD participant from the Observer Alexithymia Scale

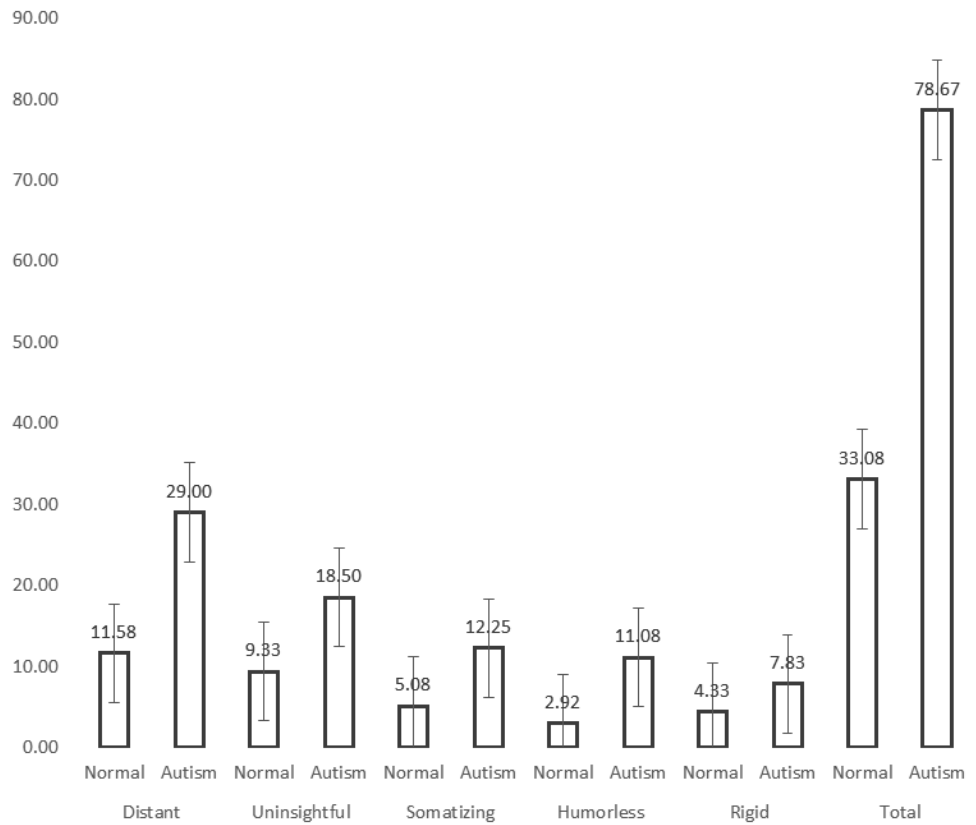


Fig. 2. Neurotypical and Autism Spectrum Disorder participants scores with error bar based on the Observer Alexithymia Scale

Table 4. Neurotypical and Autism Spectrum Disorder participants scores based on the Observer Alexithymia Scale

Domains (Total N=24; Neurotypical N = 12 and Autism N = 12)		Mean	Std. Deviation	Std. Error Mean
Distant	Neurotypical	11.5833	4.66044	1.34535
	Autism	29.0000	1.27920	.36927
Uninsightful	Neurotypical	9.3333	3.14305	.90732
	Autism	18.5000	1.62369	.46872
Somatizing	Neurotypical	5.0833	2.96827	.85686
	Autism	12.2500	1.65831	.47871
Humorless	Neurotypical	2.9167	2.39159	.69039
	Autism	11.0833	2.35327	.67933
Rigid	Neurotypical	4.3333	1.92275	.55505
	Autism	7.8333	1.58592	.45782
Total	Neurotypical	33.0833	9.33671	2.69528
	Autism	78.6667	3.67630	1.06126

the neurotypical participants was ($M = 33.08$; $SD = 9.33$) and the autism spectrum disorder participants ($M = 78.66$; $SD = 3.67$). It is found that the independent sample t-test value of 15.7 is statistically significant at the 0.05 level, so the hypothesis that there is no difference between neurotypical and autism spectrum disorder people with alexithymia is false hence rejected. This means that there is a significant difference between neurotypical and autism spectrum disorder people with this condition. The capacity to perceive and comprehend emotional expressions on others faces is a basic skill in daily interactions, and it is especially essential early in life, before the onset of language¹³. Face recognition is a key cue for social interaction and emotional communication. Facial expression recognition appears to develop gradually from birth and childhood and appears to persist into early adulthood¹⁴. As a neurotypical person who has been exposed to various emotional cues throughout their childhood and neurological condition, they are wired in such a way that they can identify and perceive the emotion in a variety of ways. Because of the neurodevelopmental condition, people with ASD have a hard time understanding social and emotional cues.

The domain distant from the observer's alexithymia scale is completely negatively scored, hence the neurotypical participants scored less when compared to ASD participants. The items were positively assessed in nature but negatively

scored for interpretation. Hence, in the path analysis model, the significant correlation of the domains such as *uninsightful*, *somatizing*, *humorless*, and *rigid* was presented in the figure 1.

In table 4, it was shown that persons with autism scored higher than neurotypical individuals in all aspects of the OAS. *Uninsightful* denotes an inability to perceive things deeply or, depending on the context. *Somatization* is the process through which psychological problems become physical symptoms. Things that are excessively serious, unsmiling, or unable to recognise humour are referred to be *humorless*. The inability to adjust or adapt one's body language or facial expression is referred to as *rigidity*. All of these dimensions conceptually mirror the clinical state of alexithymia, which has been shown to be considerably higher in ASD patients.

Figure 2 represents the domain wise responses of the participants and the total mean score of the OAS. Error bars provide a broad notion of how accurate a measurement is, or, conversely, how distant the real (error-free) value may be from the reported value.

CONCLUSION

The research experimentally showed that there is a statistically significant difference between neurotypical and autism spectrum disorder people in emotional expression. When

compared to neurotypical people, individuals with ASD are more prone to conditions such as a lack of emotional expression, being unobservant, somatizing, being humourless, and rigidity. Persons with autism spectrum disorder are more likely to have alexithymia than neurotypical people, according to the findings.

Limitations

The number of participants in our research was limited since we were only interested in comparing ASD patients to non-ASD patients. In this research, we didn't compare cognitive or behavioural functioning, which are two additional fundamental aspects of human functioning.

ACKNOWLEDGMENT

The corresponding author would like to thank and acknowledge former Prof. Dr. R. Murugesan, Director-Research, CARE and Prof. Dr. O. T. Sabari Sridhar, Head-Department of Psychiatry, Sri Ramachandra Institute of Higher Education and Research for their valuable suggestions and support during the course of the work. He also extends his sincere gratitude to the Department of Clinical Psychology, National Institute for Empowerment of Persons with Multiple Disabilities (Divyangjan) for their constant support and cooperation throughout the study.

Conflict of Interest

None.

Funding Source

Chettinad Academy of Research and Education, Junior Research Fellowship (CARE – JRF) fund.

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