Trial of Center-Based vs. In-Home Pivotal Response Treatment (PRT) in Autism

NCT04899544

Status	RECRUITING
Phase	Not Applicable
Sponsor	Stanford University
Enrollment	120 participants

Key Eligibility Criteria

Inclusion (8)

- Diagnosis of Autism Spectrum Disorder (ASD) based on Autism Diagnostic Interview Revised (ADI-R), Autism Diagnostic Observation Schedule, Second Edition (ADOS-2) or Childhood Autism Rating Scale, Second Edition (CARS-2), Diagnostic and Statistical Manual 5th Edition (DSM-5), and expert clinical opinion;
- Boys and girls between 2.0 and 5.11 years;
- Ability to participate in the testing procedures to the extent that valid standard scores can be obtained;
- Language delay as measured by the Preschool Language Scale, 5th Edition (PLS-5): Standard score at least 1 standard deviation below average for expressive language ability for 2 and 3 year olds; 2 standard deviations for 4 year olds, and 3 standard deviations for 5 year olds;
- Stable treatment (e.g., Applied Behavior Analysis - ABA), speech therapy, school placement, psychotropic medication(s) or biomedical intervention(s) for at least 1 month prior to baseline measurements;

... and 3 more (see full listing online)

Exclusion (6)

- Current or lifetime diagnosis of severe psychiatric disorder (e.g., bipolar disorder, etc.);
- Receiving ABA of 15 hours or more;
- Presence of active medical problem (e.g., unstable seizure disorder or heart disease);
- Previous adequate Pivotal Response Treatment (PRT) trial;
- Participants living more than 30 miles from Stanford University;

... and 1 more (see full listing online)

Locations (1 total)

Stanford University School of Medicine, Stanford, California, United States

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