About this Book

How to Use This Book

This book was written for families as an ongoing resource to answer questions you may have now, or in the future. We have included a lot of information on a variety of topics. We hope that as questions arise, you will be able to find the information you need.

Tips for Parents When Using This Book:

1. Don’t feel like you have to read it all at once.
2. Read what is important to you now, and save the rest.
3. Keep the book handy so you can use it when new questions arise.
4. Keep your notes and other paperwork in the back to take with you to your child’s appointments.

Creating This Book:

The CDRC participates in a leadership training program called Leadership and Education in Neurodevelopmental Disabilities (LEND). This program brings students from all over the country together to learn about working with children and families who are living with disabilities. Each year there is a parent of a child with a disability who participates in this program. Like all participants, the parent is required to complete a project as a part of the program.

As the LEND parent participant, Misti Moxley compiled this book. This quality improvement project looked at the information parents were given upon receiving a diagnosis of Autism Spectrum Disorder (ASD) for their child. The purpose of the project was to provide accurate, helpful and supportive information to families throughout their journey.

The art work in this book was created by Griffin Moxley a 10 year old child with an ASD.
Contact Information

We appreciate the opportunity to meet with you and your child and to help you better understand Autism Spectrum Disorder (ASD). Please think of us as an ongoing resource for you provide intervention, to answer questions, or help you find what you need.

If you have specific questions about the reports you receive from the Autism Clinic, please write them down and call our team member who gave you the information at the end of your visit. His or her name and phone number is listed on the summary sheet.

Also, do not hesitate to contact our family consultant Laura Wilson with questions or feedback about your evaluation (503-494-0604).

*Thank you again for trusting us with the care of your child.*

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What Do I Do Now?

Every child and family member that comes to CDRC is unique, and comes with their own set of expectations and questions. Some families are ready to hear that their child has an Autism Spectrum Disorder (ASD), and others may take some time to accept the diagnosis.

No matter what you feel today, at some point you will be thinking: “What do I do now?”

Finding out what resources are available to you and coming up with a plan to help your child be successful is a great place to start.
Getting Started

We have provided (below) a list of things that you can do to get started. Many of them are also listed on the recommendations you were given in the clinic.

Schedule an appointment with your child’s primary care doctor.
- Bring a copy of the summary sheet you were given at the end of your evaluation at CDRC Autism clinic.
- Discuss the recommendations you were given, and get referrals if needed.
- Make sure your child’s doctor is comfortable working with a child with a ASD.

Contact your local Education Service District (ESD) if your child is under 5 years old.
- Begin an early intervention program right away if your child is not already participating in one. These programs are almost always free in Oregon and Washington.

If your child is school age, consider sharing information with their teacher or school special education team.
- Sharing information and coming up with a plan together is helpful for the school, the parents and most of all, the child.

Talk to another parent of a child with ASD.
- There is a family consultant at the Autism clinic that would be happy to talk with you.
- The Autism Society of Oregon can connect you with other parents in your area.
- Swindell’s Resource Center and Oregon PTI also have parent to parent services.
Dealing With your Child's Diagnoses:

Finding out that your child has an ASD can be a very emotional time for your family.

All families react in different ways, but many parents report that they felt like they were grieving. As a parent, you love your child so much, and you want so much for them that it is heart breaking to think that things will be hard for them.

There are some common stages that families report experiencing. Stages of the grieving process may include:

Shock or Confusion

The day your child is diagnosed with ASD can be very overwhelming and confusing. Some people may deal with this by not agreeing with the diagnosis, and wanting a second opinion. Some may completely ignore it, while others just feel overwhelmed and confused. It takes a bit of time to really process the news you have been given.

Sadness

Some families feel like they are mourning the loss of what they thought their child might become. Also, the realization of how unfair it will be that their child will struggle with many things is hard to accept. It is OK to be sad, and it is even healthy to cry. It is important to not let the sadness consume you because the thing that your child needs most is you. Starting to come up with a plan on how you are going to help your child can move you forward and help you feel more in control, even if it is just one thing to start with.

Guilt

Many parents feel guilt. They wonder if they may have caused their child’s ASD, or if they could have done something to prevent it. Even though the causes of ASDs are not completely known, we do know that it is nothing you could have controlled. Research suggests that ASD are often a genetic disorder, but that there may be some environmental things that make it more likely to occur. The ASD rates are similar across ethnic and racial groups, so it is widely accepted that it is not caused by diet.
Anger

It is very hard to watch your child struggle with things that come naturally to other children. Sometimes you might feel angry at others, yourself, or anger that is not directed at anyone in particular. This is a natural part of the grieving process. Even after you have accepted that your child has ASD, there will be times when things seem so unfair and it will make you angry. Many times others do not understand that you are hurting or that you may need support. Talking to others about your child can help. It is comforting to know you are not alone.

Denial

There may be times when you feel like it is not true. They must have seen your child on a bad day, or maybe they did not really ask all the right questions. Many parents feel this way, and it is a way of coping with something that is overwhelming. Make sure you are aware of this and continue to provide the support your child needs.

Loneliness

This can often come and go for a parent whose child has a disability. It always seems that no one will understand, or others don’t have to go through the same things. Others who do not have a child with a disability may find it hard to understand what you are going through. It is up to you to tell them what it is like, so they can support you. It is also very helpful to talk to other parents who have children with an ASD or other disabilities. They most likely will know exactly how you feel.

Acceptance

This means that you know your child has an ASD, and you are ready to advocate for what they need. Creating a plan and taking steps to help your child is ultimately what will give them the best chance for success.

Reality is what you have today, but today's reality does not end tomorrow's possibilities.
Taking Care of Your Family:

Many times families work so hard trying to meet the needs of their child with an ASD, that they may neglect the needs of the rest of the family. It seems that there is always more that can be done. It is important to balance what you do for your child with the needs of the rest of the family. Tired and burned-out parents are not parenting at their best. Make sure to take time for yourselves and your other children, especially in the first few weeks after your child is diagnosed. Take time to experience the emotions you are having and find support in others that are close to your child. Don’t be afraid to talk to a mental health professional if you need to, or find a support group.

When you are ready to learn about your child’s diagnosis, we encourage each family member to take some time to search the internet, find support groups, and read books about ASDs. Also, keep track of your questions and bring them with you to your child’s appointments. Understanding ASDs are a large part of being able to help your child. Keep an open mind, and listen to others, but also know that what may seem to work for one family may not work for your family. Remember you know your child best!

Next, we encourage you to join parent support groups and begin to develop a treatment program that is comfortable for you. Support groups provide emotional support for families, but they are also a great place to learn from other parents. Many of whom are ahead of you on this journey and can provide guidance in how to access services, find good providers, and just what to expect.

Remember that the child you knew before the diagnosis is exactly the same child after the diagnosis. A diagnosis is just a label that describes some of their characteristics; it does not change who your child is. Your child has the potential to do great things; however, a lot of time, effort, and energy, on everyone’s part may be needed to make that happen.
Siblings

Siblings of a child with an ASD often have their own set of challenges. It is important that you take time alone with your other children. They are often overshadowed by the demands of their sibling with an ASD. It is also important to find activities that siblings can enjoy together. Teach your other children to be proud of their brother or sister with an ASD, and to recognize that each child is special in different ways.

Grandparents and Other Family Members

Others in your family who are close to you and your child will deal with your child’s diagnosis in a different way, just like every parent deals with it differently. Keep in mind that they have not been to all of your appointments and heard all of the information that you have. Also, sometimes parents have to deal with finding out their child has an ASD before they are ready to talk about it with family members. Don’t forget that they love your child too and they may be grieving and confused. Encourage them to:

- Be respectful and supportive of the decisions you are making for your child
- Learn more about ASD
- Follow the routines and systems that your child is use to
- Find a support group

ASDs are treatable!

- Getting your child diagnosed early so they can receive help at an early age will allow for the greatest amount of improvement.
- Children with a ASDs are able to form strong relationships with others.
- Children with an ASD grow, change, learn new skills, and make progress toward reaching their potential every day.
I am often asked to describe the experience of raising a child with a disability - to try to help people who have not shared that unique experience to understand it, to imagine how it would feel. It's like this......

When you're going to have a baby, it's like planning a fabulous vacation trip - to Italy. You buy a bunch of guide books and make your wonderful plans. The Coliseum, the Michelangelo David, the gondolas in Venice. You may learn some handy phrases in Italian. It's all very exciting.

After months of eager anticipation, the day finally arrives. You pack your bags and off you go. Several hours later, the plane lands. The stewardess comes in and says, "Welcome to Holland."

"Holland?!?" you say. "What do you mean Holland?? I signed up for Italy! I'm supposed to be in Italy. All my life I've dreamed of going to Italy." But there's been a change in the flight plan. They've landed in Holland and there you must stay. The important thing is that they haven't taken you to a horrible, disgusting, filthy place, full of pestilence, famine and disease. It's just a different place.

So you must go out and buy new guide books. And you must learn a whole new language. And you will meet a whole new group of people you would never have met. It's just a different place.

It's slower-paced than Italy, less flashy than Italy. But after you've been there for a while and you catch your breath, you look round....and you begin to notice that Holland has windmills....and Holland has tulips. Holland even has Rembrandts. But everyone you know is busy coming and going from Italy... and they're all bragging about what a wonderful time they had there.

And for the rest of your life, you will say "Yes, that's where I was supposed to go. That's what I had planned." And the pain of that will never, ever, ever go away... because the loss of that dream is a very very significant loss.

But... if you spend your life mourning the fact that you didn't get to Italy, you may never be free to enjoy the very special, the very lovely things ... about Holland.
Understanding and Diagnosing Autism Spectrum Disorders (ASD)

Understanding Autism Spectrum Disorders (ASD)

ASD describes a range of developmental disability caused by the way a child’s brain develops. Symptoms of ASD usually appear during the first three years of life, and include significant delays in a child’s ability to relate to and communicate with other people.

The medical criteria for diagnosing Autism Spectrum Disorder (ASD) are based on the *Diagnostic and Statistical Manual of Mental Disorders, Fifth Edition* (DSM-5). A major change from the earlier DSM-IV is that ASD is now considered a single spectrum, rather than a collection of sub-types (like PDD-NOS, Asperger’s Disorder, and Autistic Disorder). The reason for this change is that research attempting to distinguish different subsets of autism found they were inconsistent over time, variable across regions, and different only in terms of symptom severity. This indicates that they do not really represent separate disorders.

Because ASD is defined by a common set of behaviors, it is best represented as a single diagnostic category that is adapted to the individual by inclusion of clinical specifiers (for example, severity of symptoms or verbal abilities) and associated features (for example, known genetic disorders, epilepsy or intellectual abilities).

It is important to know that anyone diagnosed with any type of ASD before this change should still meet the new criteria for ASD based on the DSM-5.

The following describes the symptoms a child must demonstrate to be diagnosed with an ASD based on the DSM-5:

1. Difficulties with social communication and social interaction (across settings):
   - Social-emotional reciprocity (e.g., sharing of interests and emotions, showing or pointing out objects, social initiation, back and forth conversation)
   - Using and understanding nonverbal communication (e.g., eye contact, facial expressions, gestures, body-language, combining verbal and nonverbal efforts)
   - Making and keeping age-typical social relationships (e.g., showing interest in others, sharing imaginative play, adjusting behavior to match situations)
2. Restricted and repetitive patterns of behavior, interests, or activities (2 or more of these):
   - Repetitive speech, motor movements, or use of objects (e.g., hand flapping, echolalia, non-functional use of objects)
   - Inflexibility about routines, repetitive patterns of behavior, or strong resistance to change (e.g., motor rituals, insistence on same driving route, repetitive questioning or extreme distress at small changes)
   - Restricted, fixated interests that are abnormally intense or focused (e.g., attachment to unusual objects, excessively narrow or preoccupying interests)

3. Over- or under-reactivity to sensory input or unusual interest in sensory aspects of the environment (e.g., pain/heat/cold, specific sounds or textures, smelling or touching of objects, fascination with lights or spinning objects)

4. These difficulties must be present in early childhood (but may not be fully observed until social demands exceed the child’s abilities)

5. These difficulties limit and impair everyday functioning
Defining “The Spectrum”

The autism “spectrum” simply means that symptoms are very different for each child. There is a wide range of symptoms that affect children in a wide range of different ways along a spectrum, or continuum.

Because it is a spectrum disorder, it can be very different in each child. One child may have a very hard time with social skills and have very few repetitive behaviors. Another child might be consumed by the repetitive behaviors, and have an easier time with social skills.

Each symptom a child has can fall anywhere along the spectrum. S/he may have one area of language that is very well developed, and another that is very hard for them.

You may hear professionals referring to the level of support your child needs in the following ways:

Level 3: ‘Requiring very substantial support’
- Social/communication: Severe deficits in verbal and nonverbal social communication skills cause severe impairments in functioning; very limited initiation of social interactions and minimal response to social overtures from others.
- Restricted and Repetitive Behaviors: Preoccupations, fixated rituals and/or repetitive behaviors markedly interfere with functioning in all spheres. Marked distress when rituals or routines are interrupted; very difficult to redirect from fixated interest or returns to it quickly.

Level 2: ‘Requiring substantial support’
- Social/communication: Marked deficits in verbal and nonverbal social communication skills; social impairments apparent even with supports in place; limited initiation of social interactions and reduced or abnormal response to social overtures from others.
- Restricted and Repetitive Behaviors: RRBs and/or preoccupations or fixated interests appear frequently enough to be obvious to the casual observer and interfere with functioning in a variety of contexts. Distress or frustration is apparent when RRB’s are interrupted; difficult to redirect from fixated interest.
Level 1: ‘Requiring support’

- Social/communication: Without supports in place, deficits in social communication cause noticeable impairments. Has difficulty initiating social interactions and demonstrates clear examples of atypical or unsuccessful responses to social overtures of others. May appear to have decreased interest in social interactions.

- Restricted and Repetitive Behaviors: Rituals and repetitive behaviors (RRB’s) cause significant interference with functioning in one or more contexts. Resists attempts by others to interrupt RRB’s or to be redirected from fixated interest.
How are ASDs Diagnosed?

There is no blood or other medical test that can be done to tell parents if their child has an ASD. But there are a number of observational tests that should be completed as part of a thorough evaluation for an ASD. These tests are known as “standardized” or “validated” tests. These usually require a trained person (a doctor or psychologist) to play or otherwise interact with the child while looking for some very specific behaviors. They write down the behaviors they observed, and then they score their observations. This tells them how the child’s behavior and skills compare to typically developing kids and other kids with an ASD.

Other standardized tests rely on parent report rather than observation for information. Both types of information (direct with the child and indirect from a parent) are important for making a diagnosis.

The following chart is a list of some of the tests that may have been used in your child’s evaluation through the CDRC Autism clinic. There are a variety of tests listed, and they provide all kinds of information about a child. It is important for the providers to get a clear picture of how a child is functioning in many areas.

<table>
<thead>
<tr>
<th>Test Name</th>
<th>Purpose of the test</th>
</tr>
</thead>
<tbody>
<tr>
<td>Child Behavior Checklist or Behavior Assessment System for Children</td>
<td>These forms are filled out by parents and interpreted by psychologists.. They include questions about the child’s behavior in many areas of his or her life to capture a broad picture.</td>
</tr>
<tr>
<td>Sensory Profile</td>
<td>This form is filled out by the parents. It asks questions about any difficulties the child is having with sensory input (light, sound, textures, foods...)</td>
</tr>
<tr>
<td>Autism Diagnostic Observation Schedule (ADOS)</td>
<td>This is a test that looks at a child’s behavior in the context of social and play interactions. It is specifically for diagnosing autism, looks at social communication, and use of objects and toys.</td>
</tr>
<tr>
<td>Mullen Scale of Early Learning or Stanford Binet Intelligence Scales or Developmental Ability Scales or Wechsler Intelligence Scale for Children</td>
<td>These norm-referenced tests look at the overall cognitive development of a child based on how s/he behaves during structured activities.</td>
</tr>
<tr>
<td>Vineland Adaptive Behavior Scale or Adaptive Behavior Rating Scale</td>
<td>These tools look at a child’s adaptive skills, or how s/he gets along in daily life. It may be completed by interview or filled out by parents.</td>
</tr>
<tr>
<td><strong>Speech and Language Evaluation</strong></td>
<td>A variety of tests and observations may be used. The purpose is to see how the child’s speech and language is developing. Also, how the child is using language to communicate.</td>
</tr>
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<td>-----------------------------------</td>
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<tr>
<td><strong>Motor Skills Evaluations (fine and gross motor skills)</strong></td>
<td>An Occupational Therapist will evaluate a child’s motor skills by interacting and playing with the child. They may use a variety of tests to measure fine and gross motor skills.</td>
</tr>
</tbody>
</table>
Other Characteristics of Autism Spectrum Disorder (ASD)

Advanced Thinking
Children with an ASD also can be very smart. They sometimes have one area they are interested in, and they become an expert in that area. Even the children who are very smart may still have a hard time in many areas such as:

- Abstract Thinking.
- Seeing the whole picture.
- Filtering out what is not important.
- Organization.
- Planning.
- Problem solving.
- Taking something they have learned and applying it to a different setting.

Global Developmental Delay
This is when a child is delayed or behind in many areas of their lives. Global simply means that the delay can be seen across most areas of the child’s abilities. This diagnosis is for children 6 years old and under.

Intellectual Disability
If a child has a low IQ (a measure of cognitive ability) and their adaptive skills or “life skills” are significantly delayed areas after age six, it is called Intellectual disability.

Anxiety and Depression
Some children with an ASD suffer from anxiety and/or depression. This is more common as children get older and become more socially aware. It is important to keep in mind that mood changes over a period of time should be discussed with the child’s therapist or the primary doctor.

Pica
This is when children regularly eat items that are not food (clay, dirt, crayons). Most children do this sometimes, but children with Pica do it often and it continues over time.

Seizure Disorder
Some children with ASD also have seizures. A neurologist may need to be part of your treatment team if seizures are a concern.
**Genetic Disorders**

Some children have an ASD as a part of a genetic Disorder. Fragile X syndrome is a genetic disorder that can cause symptoms of ASD. Having a child tested for a genetic disorder will not cure their ASD, but it will tell families why the child has these difficulties. It can also help in planning and understanding who else in the family may be at risk for passing on the syndrome to their children.

**Clumsiness**

Some children with ASDs are clumsy and may have trouble developing motor skills. Some also have weak upper body strength and may have delayed coordination development. Motor planning can be difficult, such that children may struggle to sequence their muscle movements smoothly.

**Gastrointestinal and Feeding Disorders**

Many parents report gastrointestinal (GI) or stomach problems in their children with an ASD. Children with ASDs are sometimes also very picky eaters and may have difficulty swallowing or even tolerating certain food in their mouth. Children with an ASD may have problems such as chronic constipation or diarrhea. These difficulties should be discussed with your medical team so they can be addressed.

**Sleeping Problems**

Many children with an ASD have trouble falling asleep, or staying asleep at night. This can be very hard on the child and the rest of the family. If your child is having trouble with sleep, talk to their doctor about things you can do to help.
What Causes ASD?
Scientists do not know exactly what causes Autism Spectrum Disorders (ASDs). There is a lot of research being done to answer that question. It is likely that within the next 20 years there will be much more information available to explain what causes an ASD. For now, there are some things that are known or widely accepted.

- ASDs are caused by the way the brain develops.
- There is not one single cause of ASD. There are many factors that may make a child more likely to have an ASD.
- Genes are one of the risk factors. A child with a sibling or parent who has an ASD is more likely to have an ASD.
- About 10% of children that have an ASD also have a genetic Disorder (Down Syndrome, Fragile X Syndrome, and others).
- There is some evidence that some children are born with a susceptibility to an ASD yet the “triggers” that lead to an ASD have not been identified.

What does not cause ASD?
The cause of an ASD has not been proven, but some things have been ruled out.

- Vaccines have been rumored to cause ASD. There have been many studies done, and no link between vaccines and ASD have been found.
- Bad parenting was once thought to be a cause of ASDs. This is not true.
- There is some question whether diet is a factor, but many ethnic groups have similar ASD rates and their diets are very different.

Why are there so many children with ASDs?
There are many theories about why ASD rates are rising. The Centers for Disease Control and Prevention reports that about 1 in 110 children in the United States have an ASD. This number has risen drastically in the last 10 years. Some reasons why this number is increasing may include:

- More public awareness of ASD.
- A broader definition of an ASD.
- Better efforts to diagnose an ASD.
- More people actually have an ASD.
Education Eligibility/Label versus Medical Diagnosis

A medical diagnosis is given by medical professionals who evaluate a child and base their findings on specific symptoms that are described by the Diagnostic Manual. This diagnosis is what you received at CDRC.

Educational eligibility/label is given by a team of professionals within the public school system. Sometimes the educational eligibility/label is referred to as an educational diagnosis and this can be very confusing to parents.

All children with an ASD have some defining things in common. To be identified with a medical diagnosis of an ASD a child must have impairments that meet DSM-5 criteria. For an educational eligibility/label, four major areas must show impairment to a degree that impacts a child’s ability to learn (communication, social interaction, repetitive behaviors, and sensory integration):

**Oregon**

Some children who receive the educational eligibility/label of ASD at school (Oregon) will not meet the qualifications for a medical diagnosis. The public schools are required to place a child into a category in order to provide special education services to that child. Many children have symptoms of ASD in one or more areas, but do not fit the criteria to qualify for a medical diagnosis. In this case, the school can say the child has an educational eligibility/label of ASD. This label allows the school to provide individualized special education services for the child.

If the child has a medical diagnosis of ASD, but the school does not think the child’s ASD is affecting their school functioning, they do not have to provide special education services.

**Washington**

A child in Washington State cannot be given an educational eligibility/label of ASD unless they have a medical diagnosis. This means that often the child can qualify for school services due to one of their impairments, like speech but not for an ASD unless they have a medical diagnosis.
Once a child has been identified as having an Autism Spectrum Disorder (ASD), he or she is eligible for special education services. The federal government mandates that all children with a disability receive a “free and appropriate education.”

For children birth until kindergarten most educational services are provided by the child’s local Educational Service District (ESD).

- For children under 3 years old it is called Early Intervention.
- Children between the ages of 3 years (at the time they start) and kindergarten receive Early Childhood Special Education.

For young children who qualify, Early Intervention and Early Childhood Special Education (EI/ECSE) are the two programs that are available. A team of professionals and the child’s parents work together to come up with an Individual Family Service Plan (IFSP). This is a plan that helps the child learn strategies to be more successful at home and at school. This plan will include how the child is performing at that time, and the goals they will be working towards. Some of the things that ESDs may provide include:

- Speech therapy (speaking and using language, feeding)
- Occupational therapy (sensory, self-care, fine motor skills)
- Physical therapy (mobility or movement, gross motor skills)
- Behavioral therapy
- Training for parents and families

The Early Intervention programs can be very different from region to region. Some of the services will be provided in the home, others may be at a preschool or other location depending on the needs of the child and the resources of the ESD.

The contact information for the Oregon and Southwest Washington ESDs are located in the appendix.
Special Education Services (Kindergarten-age 21):

Children between the ages of 5 and 21 receive special education services through their local public school district. These programs always include an Individualized Education Program (IEP). This is where a team of school professionals and the child’s parents decide on what the child needs to be successful at school. The IEP is a plan that the school is required to have in place, and follow for each child with a disability.

The range of services that may be provided by a school district is very broad, and depends on the needs of the child. Some children may benefit from being in a classroom with children who also have disabilities, while for other children it is more important to have positive social role models and they may remain in the regular education classroom.

It is very important that parents are involved in the decisions that affect the education of their child. Parents should be treated as equal partners in the education process. Often parents have to insist on different placements or interventions for their child before the school is able to make them happen. This is not because the school does not care about the child, it is more likely that they are over loaded and have many other children to manage. Parents know their child best, and it is often the parent who guides this process to make sure their child gets what they need at home and at school.
How does the Oregon Department of Human Services support the families?

ASD an Introduction for Parents:
A guide to Oregon’s Human Services System.
Published by: Oregon Technical Assistance Corporation

What types of services are available for children under 18?

Supports for Families
Every county in Oregon has a program that can help families with respite, training opportunities and other basic assistance not provided by special education. This support is administered by your County Developmental Disabilities Program. Your child must meet Oregon's definition of “developmentally disabled” to be eligible for help under this program. You'll need school test results and/or doctor's reports that show your child has an ASD. The program is open to families at all income levels.

Case managers can help you think about the type of respite and training you need to care for your child at home. You will play a big part in this — you determine where and when you need help, and what kind. Case managers will coordinate and monitor the assistance you receive. Remember, the program is small and probably can't meet all your needs. But it can get you started.

Intensive In-Home Services (for children under 18 years old)
If your child needs intense supervision or intervention because of dangerous behavior and/or medical problems, your county will probably refer you to a state program for intensive in-home services. This program is for children who can't remain at home unless their family receives significant help with personal care, safety modifications, training, behavior consultation and similar child-related needs. A checklist that rates the severity of the child’s difficult behaviors and/or medical issues determines eligibility for the program. It’s not easy to get into this program and there is a limit to the number of children who can be served. But if you think you need it, ask your county developmental disabilities case manager to be considered.

Crisis/Out-of-Home Placement (for children under 18 years old)
If your child is in crisis and can't live at home, county and regional/state placement specialists can work with you to find a temporary foster home or group home for your child. Some children are in a placement setting for a few weeks; others for a longer period of time, depending on the child and the family’s situation. There are a limited number of foster and group homes in the state that provides services to children with an ASD and other developmental disabilities. (Oregon has no specific institution or residential public school for children with an ASD) Your county may have some crisis foster homes but probably will refer you to the state program for children’s crisis services. Unfortunately, the specialized placements may not be close to the family home.

A legal agreement with the State is required. It is called the Developmental Disabilities Child Placement Agreement. Also, a plan will be written to include ways that families can remain involved with their children during placement. Families are required to authorize the Oregon Program for Seniors and People with Disabilities (SPD) as payee for their children’s SSI benefits (see below). If the child does not currently receive SSI, the program will apply for these benefits in the child’s name. The cost of the placement is not charged to families; however, this could change in the future. Families are expected to provide clothing and retain health insurance for their child. SPD also will apply for a medical card to supplement the family’s health insurance.
**Supplemental Security Income (SSI)**

SSI is a federal program that provides income and medical insurance through Medicaid to eligible children who are disabled or chronically ill and whose families have little or no income or financial resources. Your child may qualify for SSI payments if your family is eligible for Medicaid. Applying for SSI can be a challenging and complex task, but it can be done. For more information, contact your county Mental Health and Developmental Disabilities Services Office.

Oregon’s Program for Seniors and People with Disabilities offers services ranging from respite care and technical consultation to intensive in-home assistance, crisis intervention and out-of-home placement. There is no separate, specific program for individuals with an ASD. Instead, services are provided through county and state programs that serve people with developmental disabilities, including children and adults with an ASD.

*If your child is under 18 years of age,* it’s important to understand that there is no “entitlement” to services from the Program for Seniors and People with Disabilities. Unlike special education, where children with disabilities have a legal right to a public education, there is no law that requires services to young children with disabilities — even if they are eligible for them.

Services are generally limited to available funds in crisis situations. However, in the past few years, services for children with disabilities who are under 18 years old and their families have been expanded and improved. Now there is more help available for families caring for children with disabilities at home...and more dollars going to prevent crisis and “burnout” in families.

If your adult child is 18 years or older and meets the developmental disability eligibility requirements, he or she is entitled to support services in Oregon. This entitlement is the result of a recent lawsuit that requires “access to support services” for adults with developmental disabilities who are living at home. Counties are working now to enroll eligible people in the system and that could take some time.

By June 30, 2009, any person 18 years of older who is eligible is entitled to receive support services based on an individual plan. Services can include such things as respite care, in-home staffing, job coaching and employment supports, community inclusion activities or other supports that help a person to live and work in his or her community. The lawsuit also allows, under certain circumstances, eligible adults to receive foster home or similar 24-hour services. However, access to these types of services is limited.

No matter what age your child is, the “doorway” to services is your local County Developmental Disabilities Program. It’s usually located in the County Mental Health Department (see State & Local Developmental Disability Services). Case managers (sometimes called Service Coordinators) in your County Developmental Disabilities Program will talk with you to see if your child is eligible for services.

- If your child is younger than 18 years, the case manager will help you figure out what kind of help you need and work with you to develop service options.

- If your child is 18 years or older, the county case manager will work with your son or daughter and you to identify the options and help you begin accessing available services.
Therapy and Treatment:

Therapy Outside of School
Experts agree that early intervention programs that provide 20-25 hours a week directly working with the child at home, school and in community therapy gives the child the best chance for success.

As your child’s parent, you must be an active and equal partner in deciding on intervention programs. You know your child best, and will be able to provide valuable information to educators and other professionals. Collaboration among the treatment providers and the parents is an essential part of the intervention. The parent is the “glue” that holds the team together.

In general, interventions for children with Autism Spectrum Disorder (ASD) can be divided into three different service delivery “systems.” These systems include:

- Your local public school district or Education Service District (ESD).
- Licensed professionals in community or hospital based settings.
- In-home therapy programs often supervised by an ASD specialist but implemented by parents, family, friends, or college students.

Community-Based Services
Children who receive special education services may also benefit from other professional services in the community. Most interventions are available through private providers and many of them may be covered by a child’s health insurance. The family often still has to pay a portion of the fee. Some programs do offer financial aid to families who meet an income requirement.

In-Home Therapy Services
Some families choose to participate in therapy that takes place in their home. A variety of therapies are available, but the cost is usually the responsibility of the family.

Some common examples of Community-Based or In-Home Therapies that can be found are:

- Speech Therapy
- Occupational Therapy
- Behavioral Therapy
- Applied Behavioral Analysis
How to Decide on a Therapy

It can be overwhelming trying to find a place to start. Unless you have an idea of what your child needs to work on, you do not know what kind of therapy to look for.

- Discussing this with your child’s doctor or other professionals that have worked with your child can help.
- Another place to begin is by looking at the milestones that children who do not have an ASD are meeting. A milestone is simply an average range for children to complete a specific task. By looking at the milestones that are included in this packet, you can decide what area your child needs the most support in, and find a therapy to support that skill.
- Talking with other parents about their children with or without an ASD can help you to understand your child’s development. They may also be able to tell you what types of interventions have helped their children, and which ones have not been helpful.
- The Autism Society of America has developed some guidelines to explain and evaluate different interventions for children with an ASD.

Children with an ASD have many different needs and no one method is going to be effective in treating all areas. A combination of treatments is usually the most effective.

Two things to focus on are:

- Treat the specific symptoms that interfere with functioning.
- Teach skills that foster healthy relationships.
Things to Consider When Deciding on a Therapy

1. Most importantly, could the therapy harm your child in any way?
2. Will the therapy cause undue hardship on your family?
3. Is the therapy scientifically based and widely accepted?
4. How will the success of the therapy be measured?
5. Are you learning so you can work with your child at home?
6. Is the person qualified to do the therapy?

It is also important to focus on the symptoms that are the most delayed for the child or are causing the most frustration for the child and family. Use the milestones in this book to decide where your child is the most delayed.

There are many different types of therapies that are recommended by different people to treat an ASD. It is important to choose therapies that are the most beneficial to your child. The following pages are explanations of the different therapies you may hear or read about.

It is important to know that no one therapy is going to solve all the issues a child with an ASD might have. A combination of therapies is often done by a provider, or several providers to help your child make the most progress.

Understanding the Different Available Therapies

In the following section called *Types of Therapies Available*, many therapies are described. They are listed as:

- **Scientifically Based**,  
- **A Promising Practice**, or  
- **Limited Supporting Information for Practice**.

This information was taken from an article called *Evidence-Based Practices and Students With Autism Spectrum Disorders* by Richard L. Simpson. It was based on a study that evaluated many therapies and the evidence for those therapies. A copy of the article summary is in the appendix.

There is also a category called “Widely Excepted”. This information is not taken from the article, but is referring to the overall view of the therapy listed. Some therapies do not have much scientific research to support their outcomes, but they are still good strategies to use with children. These would be listed as “Widely Excepted”.

* * *
### Applied Behavior Analysis: Behavior Based Therapy

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<tr>
<th>Classification</th>
<th>Meaning</th>
<th>Recommendation</th>
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<tbody>
<tr>
<td>Scientifically Based and Widely Accepted</td>
<td>There has been a lot of research for this therapy that has shown children with an ASD make improvements when involved in this type of therapy.</td>
<td>Highly Recommended for young children with Autistic Disorder or PDD-NOS, and especially if there is a global developmental delay.</td>
</tr>
</tbody>
</table>

### Background

Applied behavior analysis (ABA) describes a treatment approach that applies behavioral science in real-world settings with the aim of addressing functional issues such as social communication, behavior problems and learning. Applied Behavioral Analysis (ABA) is usually an intensive, one-on-one, structured method of teaching behaviors and skills. It is the foundation of many empirically supported behavioral treatment interventions, including, Pivotal Response, Training the Denver Early Start Program, Functional Behavior Analysis, Incidental Teaching, Verbal Behavior ABA, and Discrete Trial Therapy (DTT).

### ABA’s focus areas

ABA can be used to teach a child adaptive behavior (like toileting and eating) play skills, social skills, language skills, academic skills, self-help skills, fine motor skills, and many other life skills. In a typical ABA program, the child may have therapy for up to 10-40 hours a week for 2 or more years. Many programs provide parent training which increases the amount of therapeutic interaction a child receives and helps the family participate more fully in treatment.

### Designing an Individual Program

The foundation of an ABA program is the collection and interpretation of precise, objective data about the child’s behavior. The data is used to identify the needs of the child, measure progress, judge skill mastery, document progress, and develop treatment plans. Once the child's individualized needs have been identified, a tailor made ABA program can be developed. It will focus on teaching the skills that the child does not demonstrate, and maximize skills that are emerging.

### How it works

ABA uses careful behavioral observation and positive reinforcement or prompting to teach each step of a behavior. Generally the program will start with foundation level skills that serve as the basis for other skills that can be built from that foundation. Each functional skill set is
broken down into small, measurable elements. Each element is then taught individually through repeated trials and situations, using motivating activities to shape new behaviors.
### Background
Treatment and Education of Autistic and related Communication – handicapped Children (TEACCH) is a program that was developed in North Carolina to use in the school classroom for children with an ASD. It is based on the idea that the child’s environment should adapt to them, not the other way around.

### TEACCH’s Focus Areas
This program uses structured teaching to develop new skills for each individual child. Each child is tested to see what skills they already have. Then a plan that uses those skills as a place to start is put in place.

### Designing an Individual Program
This is an individualized program, but there are also many things about structuring the child’s environment that are going to be similar for many kids who have an ASD. Making the daily activities predictable and providing many visual supports like charts and schedules are essential to this program.

### Research
There has not been a lot of research done on this program, but many people who work with children who have an ASD find the visual strategies and structured environment essential.

### Where to Go
There are not many schools that use the TEACCH program outside of North Carolina, but most schools use some of the same ideas. For more information on the TEACCH program you can look at their website: [www.teacch.com](http://www.teacch.com)
Floortime: Relationship Based Therapy

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<th>Classification</th>
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<tr>
<td>Limited supporting information for practice.</td>
<td>There has been some research that may show that children with an ASD improve when using this curriculum</td>
<td>Recommended as a part of therapy for Autistic Disorder or PDD-NOS especially if there is a global developmental delay.</td>
</tr>
</tbody>
</table>

**Background**
Floortime is a program that is based on Dr. Stanley Greenspans model called Developmental Individual Difference Relationship Model (DIR). It was developed in the 1980’s and is used to help children build relationships and communication skills.

**Floortime’s Focus Area**
This is a type of therapy that is usually done by a therapist or a parent where the adult follows the child's lead and interacts with them at their level. This often is done by playing with the child “on the floor”.

**Goals of the Program**
The goal of Floortime is to help the child reach 6 developmental milestones that are like walking up a ladder. The steps are:

- Self-Regulation and interest in the world
- Intimacy or a special love for the world of human relations
- Two-way communication
- Complex Communication
- Emotional ideas
- Emotional thinking

**Research**
There has not been a lot of research done on this program, but many therapists and parents who work with children who have an ASD find this program helpful to get the child to engage with them. It is most often used along with other behavior therapies.

**Where to go**
To find more information of Floortime, visit their website at: www.floortime.org
Relationship Development Intervention (RDI): Relationship Based

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<th>Classification</th>
<th>Meaning</th>
<th>Recommendation</th>
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<tbody>
<tr>
<td>Limited supporting information for practice and Widely Accepted</td>
<td>There has been some research that may show that children with an ASD improve when using this therapy.</td>
<td>Recommended as a part of therapy for Autistic Disorder or PDD-NOS</td>
</tr>
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</table>

**Background**

Relationship Development Intervention is very similar to Floortime, but one main difference is that it was created as a program for parents to do at home with their child. RDI is a child centered program designed to improve relationship skills through systematic interactions where the adult is interacting at the child’s level, and slowly moving them forward.

**RDI’s Focus Areas**

RDI claims that by teaching a child with an ASD to interact socially it will allow them to achieve many other things in their lives.

**Who can Provide RDI**

Parents, teachers and other professionals can do this type of therapy with children. This therapy is often done at a part of another behavioral therapy.

**Research**

There has not been a lot of research done on this program, but many therapists and parents who work with children who have an ASD find this program helpful to get the child to engage with them. It is most often used along with other behavior therapies.

**Where to go**

To find more information of RDI, visit their website at: [http://www.rdiconnect.com/](http://www.rdiconnect.com/)
PECS / Picture Exchange Communication System: Skill Based Therapy

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<tr>
<td>Promising Practice and Widely Accepted</td>
<td>There has been some research that shows that children with an ASD improve when using this curriculum</td>
<td>Recommended as a communication strategy for non verbal or early language development.</td>
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</tbody>
</table>

Background
PECS is a program that teaches children and adults who have limited communication skills to communicate using a picture system and rewards. It was developed in 1985 and was first used in the Delaware Autistic Program.

PECS Focus Areas
This program is used to teach speech and communication. It starts out with basic one picture that is something the child wants. The child is rewarded for using the picture and over time is able to understand that symbols represent real things. As the child progresses, the pictures can get more complicated to help them use actual sentences.

Who can Provide PECS
Parents, teachers and other professionals can do this type of therapy with children. A variation of this therapy is often done in schools for children who do not speak.

Research
There has been some research done on PECS, and those studies have shown that this program helps children and adults with an ASD or other disabilities when they have little or no speech.

Where to go
Standard PECS pictures can be purchased as a part of a PECS manual (PECS Training Manual, 2nd Edition, by Lori Frost and Andrew Bondy), or pictures can be created by the adult.
Sensory Integration: Biologically Based Therapy

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<tr>
<td>Promising Practice and Widely Accepted</td>
<td>There has been some research that may show that children with an ASD improve when using this curriculum.</td>
<td>Recommended for children who have identified sensory sensitivities.</td>
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</table>

**Background**

Sensory Integration Therapy is a type of therapy that helps a child with ASD process sensory information (touch, taste, smell, sounds...). The therapy is usually conducted by an Occupational Therapist and its purpose is to help a child who is over or under stimulated by their environment learn how to cope. This therapy is child centered, and involves the adult playing with the child.

**Sensory Integration Therapy’s Focus Areas**

Many children with an ASD have trouble with sensory processing. Some may hate loud noises or certain pitches; others may get upset at very colorful room, while others only like to eat certain textures of foods. This type of therapy works with children to cope with their individual sensory problems.

**Who can Provide Sensory Integration Therapy**

Occupational Therapists usually work with children on sensory integration, but it is very important that parents are able to also work with their child at home on the same issues.

**Research**

There has been some research done on Sensory Integration Therapy that shows that it is effective for some children in treating sensory problems, but not all. The research suggests that it may be more effective for younger children.
## Social Stories: Cognitive Based Intervention

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<tr>
<td>Promising Practice and Widely Accepted</td>
<td>There has been some research that shows children with an ASD improve when using this curriculum.</td>
<td>Recommended for social development in children with Asperger’s or HFA.</td>
</tr>
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### Background
Social Stories are a way to make a social situation concrete and understandable for a child with an ASD. A social story is simply an explanation of what a child might expect in a certain situation. It is not a therapy on its own, but is commonly used by therapists, schools and families to help a child with an ASD deal with a situation.

### Social Stories focus areas
These stories are usually written in the first person and they allow a child to think about a situation that may be hard for them, how to respond to the situation, and what they can expect others to do. They usually have a lot of concrete information for the child to think about.

### Who can use social Stories
Anyone can write and use a social story. Often teachers and parents use these for a situation a child is struggling with, or a situation that will be coming that they know will be hard for the child.

### Research
There has been some research done on Social Stories as an intervention, but more needs to be done before this is called a scientifically based intervention. People who work with children who have an ASD find this tool extremely helpful.

### Example
The following is a short example of a social story taken from a book called: Sticker Strategies: Practical Strategies to Encourage Social Thinking and Organization, by Michelle Garcia Winner. This example would be used for an older child.

**Family Dinner Time: Answering Questions**
Your parents like it when you show you are thinking about them. When they ask “what did you do at school today?” think of one activity you did in one class and explain it. We talk about information not because it is fascinating, but because by telling a little bit about our lives we let people learn about what we do and feel.
## Prescription Medications: Biologically Based Therapy

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<tr>
<td>There is not a prescription medication to treat or</td>
<td>There are some medications that treat different symptoms that a child</td>
<td>This is for you and your child’s pediatrician,</td>
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<tr>
<td>cure an ASD.</td>
<td>with an ASD may have (e.g. depression, hyperactive...)</td>
<td>developmental pediatrician, or psychiatrist to</td>
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<tr>
<td></td>
<td></td>
<td>decide.</td>
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### Background

There is no prescription medication to treat an ASD, but some medications can treat the symptoms a child with an ASD may have. Some of the symptoms that may be treated with medication include hyperactivity, impulsivity, poor attention, aggression, anxiety, depression, and mood swings.

### Goals of treating with prescriptions

When treating a child with an ASD the goal of the prescription medication is to reduce a symptom so the child can respond better to the other types of therapy they are getting. It is a part of an intervention program, not a treatment on its own.

### Warnings

Given the potential side effects of medication, drug interactions, and unpredictability of how the child will react, it is strongly recommended that a medical doctor with expertise in managing medications for children with an ASD be involved. A developmental pediatrician or a child psychiatrist both specialize in this area.

### Research

There has been a lot of research conducted on medications. For information on a specific medication that is recommended to you, ask your pediatrician to explain the research for that medication, and also where you can look to find it yourself.
Vitamins and Supplements: Biologically Based Intervention

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<tbody>
<tr>
<td>Limited supporting information for practice</td>
<td>There is little or no research to support this therapy.</td>
<td>This is for you and your child’s pediatrician, developmental pediatrician, or psychiatrist to decide.</td>
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</tbody>
</table>

**Background**
Many parents and professionals claim to see some improvement in children who have an ASD when given certain vitamins. There is little or no scientific research supporting this, and parents should be very careful because some vitamins can be toxic.

**Goals of treating with vitamins and supplements**
There are many different claims to why children with an ASD need different supplements. Some of the more common goals are to encourage better digestive health, or to help the body absorb things they need and discard things they don’t.

**Warnings**
There are many professionals such as naturopathic specialists who can provide help in this area. It is important to also seek the advice from your child’s pediatrician and nutritionist. They can help you decide if the potential outcome of the supplement outweighs the potential harm.

**Research**
There is little or no scientific data that says vitamins are effective in treating an ASD.
Dietary Interventions: Biologically Based Intervention

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<tbody>
<tr>
<td>Not scientifically based</td>
<td>There is no valid research to support this therapy.</td>
<td>This is for you and your child’s pediatrician, developmental pediatrician, or psychiatrist to decide.</td>
</tr>
</tbody>
</table>

**Background**

Many people claim to have seen improvements to their child's ASD after changing their diet. The two most common dietary interventions are Gluten and Casein -Free and Yeast-Free diets.

- Gluten and Casein free diets call for the removal of all wheat, oats, rye, and dairy products. The thought is that children with an ASD do not process these foods and they cause a disruption in certain processes in the brain. Others will state that the change in their child was due to the fact that their digestive system was working better allowing their child to be more comfortable.
- Yeast Free diets call for the removal of all yeast from the diet. The theory is that children with ASDs have an overgrowth of yeast caused by tiny holes in the gastrointestinal tract. Sometimes anti-fungal medications are also used to eliminate the yeast.

**Warnings**

There are many professionals such as naturopathic specialists who can provide help in this area. It is important to also seek the advice from your child’s pediatrician. They can help you decide if the potential outcome of the diet outweighs the potential harm.

Specific allergy testing is recommended to determine what your child is allergic to. It is important that if you are going to try an elimination diet that you are working closely with a doctor, or a clinical nutritionist. Malnutrition can be more harmful to your child's health then the possible allergen. If you are planning on trying an elimination diet, a support group can be helpful.

**Research**

There is little or no scientific data that says the above diets are effective in treating an ASD.
### Heavy Metal Detox: Biologically Based Intervention

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<tr>
<td>Not scientifically based</td>
<td>There is no valid research to support this intervention to treat an ASD, and some to show that it is harmful. There have been cases of children dying from this type of therapy.</td>
<td>Not Recommended and can be harmful.</td>
</tr>
</tbody>
</table>

### Background
Some people believe that ASDs are caused by the child being exposed to heavy metals, particularly mercury. To get rid of the heavy metal in the child they support the use of Chelation. This is a process where a child uses a lotion, takes a pill, or has an IV to flush out the metal in their body. Several children have died from this treatment. It is very controversial.

### Warnings
We at CDRC do not endorse detoxification for children with an ASD.

### Research
There is no scientific research that says ASDs are caused by heavy metals.
Music, Art, and Animal Therapy: Relationship Based Intervention

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<th>Classification</th>
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<tr>
<td>Limited supporting information for practice and Widely Accepted</td>
<td>There is no research that says these therapies treat ASD, but there are many positive aspects of each of these therapies.</td>
<td>Recommended for relationship building, or self esteem.</td>
</tr>
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</table>

Art and Music Therapy
Art and music classes have many positive outcomes for children in general, art and music therapy are no different. Many children with ASDs are very artistic or musically gifted, and providing an opportunity for them to be good at something can help their self-esteem. In combination with other goals, art and music therapy can have positive effects in sensory integration, and possibly other areas as well. There are no harmful effects to art or music classes, and many children love it.

Animal Therapy
Owning a pet or interacting with an animal is something most children enjoy. There are many positive aspects of having a pet like: learning responsibility, caring for something, having something to love and love you back.

Horse therapy (Equine therapy or hippotherapy) is used to allow children to interact and care for a horse. This is said to help the child’s self esteem.

Therapy dogs are being used in different ways for children with an ASD. Some are highly trained to assist in keeping the child safe.
Developmental Milestones

The following pages are milestones that can help you better understand where your child’s skills may be advanced or delayed. They are things that typically developing children would do during a particular age range.

The information was taken from two resources, and you can find them online if you are interested in reading more about developmental milestones.

The Center for Disease control (CDC) has a resource called The Act Early Campaign. On their website you can read and print, or order free copies of the developmental milestones in English and in Spanish.

The CDC’s web address is www.cdc.gov

The second is the Zero to Three Campaign. This resource is produced by the National Center for Infants, Toddlers, and Families. The milestones are available to read or print in English or in Spanish.

The Zero to Three web address is www.zerotothree.org
0-3 Month Milestones

Social Emotional
- Begins to develop a social smile
- Enjoys playing with other people and may cry when playing stops
- Becomes more expressive and communicates more with face and body
- Imitates some movements and facial expressions

Movement
- Supports upper body with arms when lying on stomach
- Opens and shuts hands
- Pushes down on legs when feet are placed on a firm surface
- Brings hand to mouth
- Takes swipes at dangling objects with hands
- Grasps and shakes hand toys

Vision
- Watches faces intently
- Follows moving objects
- Recognizes familiar objects and people at a distance
- Starts using hands and eyes in coordination

Hearing and Speech
- Smiles at the sound of your voice
- Begins to babble or make sounds
- Begins to imitate some sounds
- Turns head toward direction of sound

What you can do with your baby:
- Talk and sing to them
- Hold and snuggle them
- Watch and learn your babies signals (hunger cry, tired cry, happy)
- Respond to their signals; when eyes are bright and awake it is play time
- Give them something to hold on to or reach for (toy or finger)
- Comfort them when they cry. Soothing makes them feel safe and loved
- Read to your baby
3-6 Month Milestones

Social Emotional
- Enjoys social play
- Interested in mirror images
- Responds to other people's expressions of emotion and appears joyful often

Movement
- Rolls both ways (front to back, back to front)
- Sits with, and then without, support on hands
- Supports whole weight on legs
- Reaches with one hand
- Transfers object from hand to hand
- Uses hand to rake objects

Vision
- Develops full color vision
- Distance vision matures
- Ability to track moving objects improves
- Finds partially hidden object
- Explores with hands and mouth
- Struggles to get objects that are out of reach

Hearing and Speech
- Responds to own name
- Begins to respond to "no"
- Can tell emotions by tone of voice
- Responds to sound by making sounds
- Uses voice to express joy and displeasure
- Babbles chains of sounds

What you can do with your baby:
- Place your baby in different positions to help develop new skills like rolling and crawling (play on back and stomach, sit with support) always put them on their back to sleep.
- Offer interactive toys for them to play with (different sizes, shapes, sounds, textures)
- Talk to your baby and have back and forth conversations (when they babble you answer)
- Create routines to help them learn things like going to bed (i.e. bath, books, feeding, song, bed)
- Read to your baby
6-9 Month Milestones

Social Emotional
- Learning to solve problems (if a toy drops they look for it)
- Copy what they see others do (play peek-a-boo)
- Understand harsh tones

Movement
- They can pick up small objects
- They can sit on their own
- May crawl, scoot, or pull up on furniture

Vision
- Develops full color vision
- Distance vision matures
- Ability to track moving objects improves

Hearing and Speech
- They babble a lot
- When talked to they make sounds back
- Use their voice to show feelings (anger, joy)
- Copy actions like waving or shaking their head

What you can do with your baby:
- Let your baby explore interesting objects like toys with buttons to push
- Talk to your baby and tell them when you like what they are doing
- Let them start trying to pick up baby safe foods to eat
- Give them to move around on their own to build muscle and coordination
- Read to your baby
## 9-12 Month Milestones

### Social Emotional
- Shy or anxious with strangers
- Cries when mother or father leaves
- Enjoys imitating people
- Prefers certain people and toys
- Tests parental responses to behavior
- May be fearful in some situations
- Prefers mother and/or regular caregiver over all others
- Repeats sounds or gestures
- Finger-feeds himself
- Extends arm or leg to help when being dressed

### Movement
- Reaches sitting position without assistance
- Crawls forward on hands and knees
- Pulls self up to stand
- Walks holding on to furniture
- Stands or walks momentarily without support
- Uses pincer grasp
- Bangs two objects together
- Puts objects into and out of container
- Lets objects go voluntarily
- Pokes with index finger

### Cognitive
- Explores objects in different ways (shaking, banging, throwing)
- Finds hidden objects easily
- Looks at correct picture when the image is named
- Imitates gestures
- Begins to use objects correctly (drinking from cup, brushing hair)

### Language
- Pays increasing attention to speech
- Responds to simple verbal requests
- Responds to “no”
- Uses simple gestures, such as shaking head for “no”
- Changes tone when babbles
- Says “dada” and “mama” and directs it to them
- Uses exclamations (“Oh-oh!”)
- Tries to imitate words

### What you can do with your baby:
- Talk to your baby (tell them what is happening and what will happen next)
- Interpret what your baby is doing by using words (you are pushing your food away, you must be all done)
- Name the things your baby looks at and points to (i.e. that is the moon)
- Give them time and a safe place to practice crawling and walking
- Play hide-and-seek games with objects
- Help them take the next steps in their play (instead of banging blocks stack them)
- Read to your child
24 Month Milestones

Social Emotional
- Imitates behavior of others, especially adults and older children
- More aware of herself as separate from others
- More excited about other children
- Shows increased independence
- Begins to show defiant behavior
- Separation anxiety increases toward midyear then fades

Movement
- Walks alone, begins to run
- Pulls toys behind while walking
- Stands on tiptoe
- Kicks a ball
- Climbs onto and down from furniture unassisted
- Walks up and down stairs holding on to support
- Scribbles on his or her own
- Builds tower of four blocks or more

Cognitive
- Finds objects even when hidden under two or three covers
- Begins to sort by shapes and colors
- Begins make-believe play

Language
- Points to object or picture when it's named for him
- Recognizes names of familiar people, objects, and body parts
- Says several single words (by 15 to 18 months)
- Uses simple phrases
- Uses 2- to 4-word sentences
- Follows simple instructions
- Repeats words overheard in conversation

What you can do with your baby:
- Let your child scribble with crayons or markers
- Help them practice physical activities like climbing and pedaling
- Introduce new vocabulary words
- Talk with them about things in their world
- Be patient with their why questions
- Help them deal with conflicts around taking turns
### 36 Month Milestones

#### Social Emotional
- Imitates adults and playmates
- Spontaneously shows affection for familiar playmates
- Can take turns in games
- Understands concept of "mine" and "his/hers"
- Expresses affection openly
- Expresses a wide range of emotions
- By 3, separates easily from parents
- Objects to major changes in routine

#### Movement Hand and Finger Skills
- Climbs and runs well
- Walks up and down stairs, alternating feet (one foot per stair step)
- Kicks ball and pedals tricycle
- Bends over easily without falling
- Makes up-and-down, side-to-side, and circular lines with a crayon
- Turns book pages one at a time
- Builds a 6 block tower
- Screws and unscrews jar lids
- Turns rotating handles

#### Cognitive
- Makes mechanical toys work
- Matches an object in her hand or room to a picture in a book
- Plays make-believe with dolls, animals, and people
- Sorts objects by shape and color
- Completes puzzles with three or four pieces
- Understands concept of "two"

#### Language
- Follows a two-three part command
- Recognizes and identifies almost all common objects and pictures
- Understands most sentences
- Understands placement (on, in)
- Uses 4 to 5 word sentences
- Can say name, age, and sex
- Uses pronouns (I, you, me, we, they) and some plurals (cars, dogs, cats)
- Strangers can understand most of her words

### What you can do with your baby:
- Talk to your baby and repeat their two word requests as a sentence (more milk = you want more milk in your cup)
- Put their feelings into words (I know you are mad)
- Play pretend
- Practice sorting objects with them
- Help them solve problems on their own
- Read with them and ask them questions about the story
Appendix and Additional Resources

Included in the Appendix:

1. What Do All of These People Do? A list of the different professionals that may work with a child with an Autism Spectrum Disorder (ASD).

2. Glossary

3. Some Extremely Reasonable Suggestions for “Typical” Parents, Family, and Teachers on Behalf of Kids With Asperger’s Syndrome

4. Books and Video Suggestions

5. Websites Resources

6. Oregon and Washington Education Service District contact information

7. Article Summary: Evaluation of Interventions and Treatment for Learners With Autism Spectrum Disorders
What Do All of These People Do?

**Pediatrician:** This is your child’s primary care doctor. Their role is to focus on your child’s general health care needs, screen for developmental delays, and to provide referrals for issues that they feel need further investigation or treatment.

**Developmental Pediatrician:** Consults with the child’s primary care doctor and with families about issues specific to developmental disorders, behavior and sleep problems, and managing medications, when needed.

**Child Psychiatrist:** Help manage medications, but also help with behavior problems and emotional issues.

**Clinical Psychologist:** Specialize in understanding the impact of developmental disabilities on the individual and family system. They document developmental progress, as well as cognitive strengths and weaknesses through standardized assessments. Provide behavior management training, social skills training, and support to parents and other family members. They can also provide behavioral therapy.

**Occupational Therapist:** Works toward improving fine motor skills and practical self-help skills that will improve independent living. For children with an ASD, they also work on sensory integration and coordination of movement.

**Physical Therapist:** Works with fine and gross motor skills to help improve a child’s coordination.

**Speech and Language Pathologist:** Communication skills, social interactions, and play are the main areas of focus. Speech (sound production), language, and feeding.

**Other Therapists:** Children also work with other therapists who provide less traditional interventions such as art, music, or horse riding.

**Certified Behavior Analysts:** Requires extensive training in the area of Applied Behavior Analysis, specializing in developing in-home therapy and community-based intervention programs.
Some Extremely Reasonable Suggestions for “Typical” Parents, Family, and Teachers on Behalf of Kids With Asperger’s Syndrome

By Jennifer McIlwee Myers, Aspie-at-Large

www.Autismdigest.com November-December 2005

I was diagnosed with Asperger’s Syndrome at age 36. It was a joyful occasion—it helped me finally stop trying to be normal. The more I learned about ASD’s (Autism Spectrum Disorders), the better I was able to adapt myself to life in a functional and enjoyable way. It turns out that being weird is not only functional, it’s really fun. I researched ASD’s in earnest. I interviewed parents, teachers, OT’s, and psychologists, neurologists—you name it. What I found is that an awful lot of those well-meaning adults were busily working “against the grain” of as/Autism. They were frustrated and tired too! In the hope of saving a few parents and teachers some of that time, energy, and pain, I have come up with a list of “suggestions” for them. These come from the heart of an Aspie who really likes the strange creatures we call “typical.”

The Suggestions:

1. **Please** don’t try to make us “normal.” We’d much rather be **functional**. It’s hard to be functional when you have to spend all your time and energy focusing on making eye contact and not tapping your feet.

2. **Please** don’t overprotect, indulge, or cosset us. We already have enough social problems without additionally learning to be spoiled and self-indulgent.

3. **Don’t** teach us social skills according to how you wish the world was, or even how you think it is. Look carefully at what is really going on and teach us **real world rules**.

4. **Don’t** talk and/or act as if your life would be perfect or soooo much easier if you had a “normal” child. We don’t thrive on knowing that we are the children you didn’t want.

5. **Don’t** make the mistake of thinking that teaching us typical behaviors and successful masking means we are “cured.” Please remember that the more typical our behavior seems, the harder we are working. What is natural, simple behavior to you is a constant intense effort for us.

6. **Please** don’t punish us with rewards or reward us with punishments. For those of us who find recess to be the most stressful part of school, any action that will keep us in from recess is one we will learn to repeat ad infinitum. Getting rewarded for good behavior with fashionable but really itchy clothing will train us to **NOT** behave too well!

7. If you assiduously train us to imitate and conform to other children’s behavior, don’t be shocked if we learn to curse, whine for popular toys, dress in ways you don’t like, and eventually drink, smoke, and attempt to seek out sex as teenagers. Those “nice kids” you think so highly of do a lot of things you don’t know about—or don’t you remember high school?

8. **Please do** give us information about ASD/Asperger’s early on at a level we can digest. We need to know what’s going on—and we will figure out that something is “wrong” with us whether you tell us or not.
9. **Don't** avoid a diagnosis or help for us because you are scared of us being labeled. Without that diagnosis and appropriate support, our teachers, family, and fellow students will give us plenty of labels – and we might just believe them if we hear them often enough.

10. **Don't** force us to do things we can't do. A forced social situation won't teach us social skills any more than dumping us in the middle of the Pacific Ocean will teach us to swim.

11. **Don't** punish us for what other kids do. The fact that other kids tease and torture us for benign “Autistic” behaviors doesn’t mean we need to change, it means they do. Needing to bounce or swing for the whole recess is not morally wrong; tormenting someone for having a neurological disability is.

12. **Don't** attempt to use humiliation or public embarrassment to “teach us a lesson.” We get way too much of that from other people, and the only lesson learned is that we can’t trust you either.

13. **Do** punish us (or give us “consequences,” heaven help us) when it is necessary to do so – but make the connection between cause and effect very, very clear. We often need visual aids to understand how our behavior can cause an unwanted result for us!!!!

14. **Don't** cut us too much slack when our behavior is potentially dangerous to us. For example, adolescent pre-stalking behavior should result in serious consequences -- because not treating such behavior seriously when we are young can lead to problems involving law enforcement when we’re older!

15. **Don't** trust untrained camp counselors, “typical peers,” or youth pastors to be able to deal with Asperger’s. Often their answers to our problems involve highly destructive phrases like “try harder,” “you could do it if you really wanted to,” and “snap out of it.”

16. **Don’t** model one thing and teach another. If you yell or hit when you’re mad, we will too. If you rage at us, don’t be shocked at our “Autistic rages.” And DON’T lecture us about our stims while you smoke, tap your foot, pick at your manicure and down your third double-latte today.

17. **Don't** require us to be wildly successful at something because your ego has been wounded by having a “flawed” child. We can’t all be Temple Grandin. Remember, all honest work is noble, even if you can’t brag about us to your friends.

18. **Do** spend time with our siblings, even if you need to arrange for respite care to do so. Schedule something special for them without us along, even if it’s just lunch at a fast-food joint once a week or so.

19. **Do** ask for help for yourself as needed. Take advantage of respite care when you can. Get cognitive-behavioral counseling and/or medication when you are depressed. Don’t try to do it all alone. Remember: it is much more important that you get a nap and a nourishing meal than that we have a tidy house.

20. Most important: please, please, please **don’t** wait until we’re “cured” or “recovered” to love and accept us. You could miss our whole lives this way.
<table>
<thead>
<tr>
<th>Books &amp; Websites</th>
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<tbody>
<tr>
<td><strong>Can I Tell You about My Asperger’s Syndrome?</strong> By Jude Welton, Jane Telford,</td>
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<tr>
<td>and Elizabeth Newson (2003)</td>
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<tr>
<td>In this book, Adam helps others understand the difficulties faced by a child</td>
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<td>with AS; he tells them what AS is, what it feels like to have AS and how they</td>
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<tr>
<td>can help children with AS by understanding their differences and appreciating</td>
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<tr>
<td>their many talents.</td>
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<tr>
<td><strong>Look Me in the Eye: My Life with Asperger’s</strong> by John Elder Robison (2008)</td>
</tr>
<tr>
<td>This is a story of one man’s struggle growing up with Asperger’s Syndrome. It</td>
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<tr>
<td>is a very real story that is also inspirational. The author has become quite</td>
</tr>
<tr>
<td>successful at many things. It is interesting to see the world through his eyes.</td>
</tr>
<tr>
<td><strong>Asperger’s Syndrome: A Guide for Parents and Professionals</strong> by: Tony Attwood</td>
</tr>
<tr>
<td>and Lorna Wing (1998)</td>
</tr>
<tr>
<td>Tony Attwood is a world renowned expert on Asperger’s Syndrome. His guide will</td>
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<td>assist parents and professionals with the identification, treatment and care of</td>
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<tr>
<td>both children and adults with Asperger’s Syndrome.</td>
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<tr>
<td><strong>Asperger’s: What Does It Mean to Me?</strong> by Catherine Faherty and Gary B.</td>
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<tr>
<td>Mesibov (2000)</td>
</tr>
<tr>
<td>A workbook explaining self awareness and life lessons to the youth with high</td>
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<tr>
<td>functioning Autism or Asperger's Syndrom.</td>
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<tr>
<td><strong>Freaks, Geeks, and Asperger’s Syndrome: A user guide to adolescence</strong> by</td>
</tr>
<tr>
<td>This book is written by a 13 year old boy with Asperger’s Syndrome. It addresses social situations from his perspective.</td>
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<tr>
<td><strong>Socially Curious and Curiously Social: A Social Thinking Guidebook for Teens &amp; Young Adults</strong> with Asperger’s, ADHD, PDD-NOS, NVLD, or other Murky Undiagnosed Social Learning Issues by Michelle Garcia Winner and Pamela Crooke</td>
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<tr>
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<tr>
<td>An anime-illustrated, get-real guidebook for teens and young adults to read themselves about how the social mind works. From texting to dating, the book provides many practical strategies and some &quot;ah-ha moments.&quot;</td>
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<tr>
<th><strong>The New Social Story Book by Carol Gray (2010)</strong></th>
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<tbody>
<tr>
<td>Social Stories provide REAL social understanding! The book promotes social understanding in children with Autism Spectrum Disorders. Social Stories are a standard approach for teachers and parents all over the globe.</td>
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<tr>
<th><strong>Comic Strip Conversations by Carol Gray (1994)</strong></th>
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<tr>
<td>This book is a how to guide for using comic strips to teach social interactions and conversations</td>
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<tr>
<th><strong>The incredible 5 Point Scale by Kari Dunn Buron and Mitzi Curtis (2004)</strong></th>
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<tr>
<td>This book uses a 5-point scale to help students understand and control their emotional reactions to everyday events. It breaks down a given behavior and develops a scale that identifies the problem and suggests alternative, positive behaviors at each level of the scale.</td>
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<tr>
<th><strong>Sticker Strategies to Encourage Social Thinking and Organization by Michelle Garcia Winner</strong></th>
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<tr>
<td>This book has 80 stickers that each contains a mini social story. They are useful for targeting behaviors in a very concrete way.</td>
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<tr>
<th><strong>Clay Marzo Just Add Water DVD by Quick Silver</strong></th>
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<tr>
<td>Clay Marzo is a world class surfer who was diagnosed with Asperger’s Syndrome. The DVD has a lot of information about Asperger’s, and shows that people with ASD’s can accomplish many things.</td>
</tr>
<tr>
<td>Title</td>
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<td>----------------------------------------------------------------------</td>
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<tr>
<td>Since We’re Friends: An Autism Picture Book</td>
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<tr>
<td>Raising a Sensory Smart Child</td>
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<tr>
<td>Ten Things Every Child With Autism Wishes You Knew</td>
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<tr>
<td>Thinking in Pictures</td>
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<td>Title</td>
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<td>-----------------------------------------------------------------------</td>
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<tr>
<td><strong>I Love Being my Own Autistic Self: A ThAutoons Book</strong></td>
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<tr>
<td><strong>Toilet Training for Individuals with an ASD or Other Developmental Issues by Maria Wheeler</strong></td>
</tr>
<tr>
<td><strong>Carly’s Voice: Breaking through Autism by Carly and Arthur Fleischmann (2012)</strong></td>
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<tr>
<td><strong>The Autism Book : Answers to Your Most Pressing Questions by Jhoanna Robledo</strong></td>
</tr>
<tr>
<td><strong>What You Can Do Right Now to Help Your Child with Autism by Jonathan Levy (2007)</strong></td>
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Oregon Summer Camps and Social Skill Groups:

ACAP (503-649-2066)
- Autistic Children's Activity Program
- A day camp in Portland that does field trips and a variety of activities.
- Cost $1200-$3000 depending on how many days and weeks.

Amazing Grace Farm Camp
- This is a day camp program for children with disabilities. There is also a session for siblings. The kids learn to work with all kinds of farm animals.
- The cost is $250 for 4 days.
- www.gemchildren.org has a listing that includes the contact info.
- This is a day camp program for children with disabilities.
- There is also a session for siblings. The kids learn to work with all kinds of farm animals.

Aspiring Youth (www.aspiringyouth.net)
- For kids with HFA, Aspergers, and ADHD...
- They have programs for 9-24 year olds.
- This is a fun and interactive day camp that has sessions throughout the year.
- They are located in the Portland area (West side)
- The cost is about $450 a week.

Blue Compass (www.bluecompasscamps.com)
- This is a program in Washington that takes kids from 7-9 for day camp, and 13-18 for adventure camp where they go backpacking, kayaking, and camping overnight.

Camp Quest (www.asdoregon.org)
- Overnight camp for kids with High Functioning Autism or Aspergers Syndrome
- Non-profit organization who works hard to make camp affordable and to find financial assistance from community organizations for families who need it.
- A typical summer camp experiences with trained staff and lots of structure.
- Activities include: fishing, swimming, canoeing, sports, computers and much more.

Clackamas Speech LLC (www.clackamasspeech.com)
- Reaching Beyond Boundries Summer program:
  (Milwaukie Oregon at the East Side Athletic Club 503-659-1516)
Play Connections: Melanie Shaw (www.playconnections.com)
- Social Development Summer Camp
- These groups run throughout the year and serve children from toddler-third grade.

Playful Interventions LLC (503-735-5870) Charla Cunningham LMFT, RDT
- Social skills theater
- Summer programs involving theater and drama skills.

SocialKraft: www.socialkraft.net 503-381-9344
- Social skills groups
- Individual social skills
- Summer groups

Wiz Kids Day Camp
- 2nd-6th grade day camp that is social skill based. Play games and have fun while improving social skills.
- $200-$350 a week
- It is in Hillsboro Oregon.
Autism Society of America

The Autism Society of America is a national organization that provides information on a variety of issues related to an ASD.

Autism Society of America
7910 Woodmont Ave, Suite 300
Bethesda, MD 20814-3067
1-800-328-8476
www.Autism-society.org

The Oregon chapter is called:
Autism Society of Oregon
503-636-1676 or 1-800-Autism-1
www.Autismoregon.com
The ARC of Oregon advocates for people with developmental disabilities including ASD. ARC provides training, respite information and other help to families.

The ARC of Oregon
(503) 581-2726
1-877-581-2726
www.arcoregon.org
Oregon Association for Behavior Analysis (OR-ABA)

Oregon association for Behavior Analysis (OR-ABA) is a professional website for parents, caregivers, educators, researchers, and others who are interested in promoting behavior analysis for children, with and without disabilities, within their community.

Email: oraba.info@gmail.com

www.orbehavioranalysis.com
Oregon Department of Human Services

The Oregon Department of Human Services (DHS) has local offices around the state for Developmental Disability Services (DDS). They are usually located in the county’s Mental Health Department.

State Office for DHS:
(503) 945-9774
www.oregon.gov/DHS/index.shtml
Swindell’s Child Disability Resources

The Swindell Center provides information and training to families who have a child with a disability.

Website: http://oregon.providence.org/our-services/s/swindells-resource-center/
Email: swindells@providence.org
The Inclusive Child Care Program

The Inclusive Child Care Program helps eligible families with higher costs of care when children need exceptionally close supervision, special accommodations, or other supports for safe and appropriate care.

To make referrals or for more information, please contact:
Inclusive Child Care Program
Portland: 971-673-2286
Toll Free: 866-837-0250
http://oregoninclusivecc.org
Oregon Technical Assistance Corporation (OTAC)


This resource can be downloaded from the OTAC website. If the download does not work call OTAC and they can send you the resource. http://www.otac.org/wp-content/uploads/2011/12/Autism-An-Introduction-for-Parents-2005r.pdf

OTAC
Phone: 503 364-9943
www.otac.org
Rethink Autism

Rethink Autism is a web-based program that provided on-going training and supports to families to teach them how to provide ABA therapy at home.

"Rethink Autism provides live, online experts in autism, education, and behavior. Our knowledgeable team of Board Certified Behavior Analysts and Special Educators provides an array of web-based services to families, schools, and organizations. Ranging from brief assessments, to professional development workshops, and even weekly program support, we have the staff and experience to meet your needs anywhere in the world.

www.rethinkAutism.com/
Autism Speaks

Autism Speaks provides a free notebook for families who have a child who has been recently diagnosed with an ASD or Asperger's Syndrome.

You can download the 100 day kit (notebook) at www.Autismspeaks.org, or call to request a free 100 Day Kit.

Families whose children have been diagnosed in the last 6 months may request a complimentary hard copy of the 100 Day Kit or the AS/HFA Tool Kit by calling 888-AUTISM2 (888-288-4762) and speaking with an Autism Response Team Coordinator.
Autistic Self Advocacy Network (ASAN)

ASAN is a nonprofit organization run by and for individuals on the autism spectrum. Its website provides information about autism through a variety of resources and self-advocacy literature.

Autistic Self Advocacy Network
PO Box 66122
Washington DC 20035
202-596-1056
http://autisticadvocacy.org
Wrong Planet
Wrong Planet is a web-based discussion forum for individuals with autism, Asperger’s Syndrome, ADHD, PDDs, and other neurological differences. Wrong Planet also provides access to exclusive articles, hot-to-guides, and a chat room for real-time communication.

Wrong Planet
http://www.wrongplanet.net
Multnomah Education Service District (MESD)/
Multnomah Early Children Program (MECP)

MESD/MECP provides early intervention and educational services to children with disabilities from birth to school age who have developmental delays or disabilities. Screenings, evaluations, and services are provided for free to children who qualify.

Multnomah Education Service District/Multnomah Children Program
Patt Moffitt, Program Coordinator
503 257-1685
http://w3.mesd.k12.or.us
Family and Community Together (FACT)

FACT is a family leadership organization for individuals and their families experiencing disability, working collaboratively to facilitate positive change in policies, systems and attitudes, through family support, advocacy, and partnerships. It offers statewide training, parent-to-parent connections, resource and information referral, and an IEP help line.

Family and Community Together
13455 SE 97th Avenue
Clackamas, OR 97015-8662
Phone: 888-988-3228
Fax: 503-786-6084
info@factoregon.org
www.factoregon.org
Defeat Autism Now! (DAN!)

DAN! Doctors was created by the Autism Research Institute with the idea that autism is a biomedical disorcer. Doctors who follow this practice include psychiatrists, neurologists, immunologists, allergists, geneticists, and gastroenterologists.

Defeat Autism Now! Doctors
www.autismtoday.com/clinician-list.html
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<tr>
<th>ESD</th>
<th>Address</th>
<th>Phone #</th>
<th>Website</th>
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<tbody>
<tr>
<td>Clackamas</td>
<td>13455 SE 97th Ave, Clackamas, OR 97015</td>
<td>503 675-4000</td>
<td><a href="http://www.clackesd.k12.or.us">www.clackesd.k12.or.us</a></td>
</tr>
<tr>
<td>Columbia Gorge</td>
<td>400 E Scenic Dr. Suite 207, The Dalles, OR 97058</td>
<td>541 298-5155</td>
<td><a href="http://www.cgesd.k12.or.us">www.cgesd.k12.or.us</a></td>
</tr>
<tr>
<td>Douglas</td>
<td>1871 NE Stephens, Roseburg, Oregon 97470</td>
<td>541 440-4777</td>
<td><a href="http://www.douglasesd.k12.or.us">www.douglasesd.k12.or.us</a></td>
</tr>
<tr>
<td>Grant</td>
<td>835A South Canyon Blvd., John Day, OR</td>
<td>541 575-1349</td>
<td><a href="http://www.grantesd.k12.or.us">www.grantesd.k12.or.us</a></td>
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<tr>
<td>Harney</td>
<td>PO Box 460 Burns, Oregon 97720</td>
<td>541 573-2122</td>
<td><a href="http://www.harneyesd.k12.or.us">www.harneyesd.k12.or.us</a></td>
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<tr>
<td>High Desert</td>
<td>145 SE Salmon Ave., St A, Redmond, Oregon 97756</td>
<td>541 693-5600</td>
<td><a href="http://www.hdesd.org">www.hdesd.org</a></td>
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<td>Jefferson</td>
<td>295 SE Buff Street, Madras, OR 97741</td>
<td>541 475-2804</td>
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<tr>
<td>Lake County</td>
<td>357 North L Street, Lakeview, OR 97630</td>
<td>541 947-3371</td>
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<tr>
<td>Lane</td>
<td>1200 Highway 99N, Eugene, OR 97402</td>
<td>541 461-8200</td>
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<td>Linn Benton</td>
<td>905 4th Avenue SE, Albany, OR 97321-1999</td>
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<td>Malheur</td>
<td>363 A Street West, Vale, Oregon 97918</td>
<td>541 473-3138</td>
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<tr>
<td>Multnomah</td>
<td>P.O. Box 301039, Portland, OR 97220</td>
<td>503 255-1841</td>
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<tr>
<td>North Central</td>
<td>PO Box 637, Condon, Oregon, 97823</td>
<td>541 384-2762</td>
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<tr>
<td>Northwest Regional</td>
<td>5825 NE Ray Circle, Hillsboro, OR 97124-6436</td>
<td>503 614-1428</td>
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<td>Columbia Service Center</td>
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<td>Tillamook Service Center</td>
<td>2410 5th St. • PO Box 416, Tillamook, OR 97141</td>
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<td>Wallowa County (District 18)</td>
<td>107 Southwest First St., Enterprise, OR 97828</td>
<td>541 426-4997</td>
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<tr>
<td>South Coast (Coos Co.)</td>
<td>1350 Teakwood Avenue Coos Bay, Oregon 97420</td>
<td>541 269-1611 541 247-6681</td>
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<tr>
<td>Southern Oregon</td>
<td>101 North Grape Street, Medford, OR 97501</td>
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<td>InterMountain (ESD)</td>
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<td>541 276-6616</td>
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<td>Willamette Marion Center</td>
<td>2611 Pringle Rd. SE, Salem, OR 97302</td>
<td>503 588-5330</td>
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<tr>
<td>Willamette Yamhill Center</td>
<td>2045 SW Highway 18, McMinnville, OR 97128</td>
<td>503 435-5900</td>
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<td>Vancouver Educational Service District 112</td>
<td>2500 NE 65th Avenue, Vancouver, WA 98661</td>
<td>360-750-7500</td>
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<tr>
<td>Battle Ground School District 119</td>
<td>11104 NE 149th Street Brush Prairie, WA 98606</td>
<td>360-885-5300</td>
<td><a href="http://www.bgbsd.k12.wa.us">www.bgbsd.k12.wa.us</a></td>
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<tr>
<td>Camas School District 117</td>
<td>841 NE 22nd Avenue Camas, WA 98607</td>
<td>360-833-5400</td>
<td><a href="http://www.camas.wednet.edu">www.camas.wednet.edu</a></td>
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<tr>
<td>Roosevelt School District 403</td>
<td>615 Chinook Avenue PO Box 248 Roosevelt, WA 99356-0248</td>
<td>509-384-5462</td>
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<tr>
<td>Castle Rock School District 401</td>
<td>600 Huntington Avenue South Castle Rock, WA 98611</td>
<td>360-501-2940</td>
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<td>2315 Centerville Highway Centerville, WA 98613</td>
<td>509-773-4893</td>
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<td>Skamania School District 2</td>
<td>122 Butler Loop Road Skamania, WA 98648</td>
<td>509-427-8239</td>
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<td>Evergreen School District 114</td>
<td>13501 NE 28th Street PO Box 8910 Vancouver, WA 98668-8910</td>
<td>360-604-4000</td>
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<td>Glenwood School District 401</td>
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<tr>
<td>Green Mountain School District 103</td>
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<td>360-225-7366</td>
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<td>Hockinson School District 98</td>
<td>17912 NE 159th Street Brush Prairie, WA 98606</td>
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<td>Toutle Lake School District 130</td>
<td>5050 Spirit Lake Memorial Highway Toutle, WA 98649</td>
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<td>Kalama School District 402</td>
<td>548 China Garden Road Kalama, WA 98625</td>
<td>360-673-5282</td>
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<td>Kelso School District 458</td>
<td>601 Crawford Street Kelso, WA 98626</td>
<td>360-501-1900</td>
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<td>Trout Lake School District R-400</td>
<td>2310 Highway 141 Trout Lake, WA 98650-9799</td>
<td>509-395-2571</td>
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<td>Klickitat School District 402</td>
<td>PO Box 37 Klickitat, WA 98628-0037</td>
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<td>La Center School District 101</td>
<td>725 Highland Road PO Box 1840 La Center, WA 98629</td>
<td>360-263-2131</td>
<td><a href="http://www.lacenterschools.org/">www.lacenterschools.org/</a></td>
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<td>Ridgefield School District 122</td>
<td>2724 South Hillhurst Road Ridgefield, WA 98642</td>
<td>360-619-1300</td>
<td><a href="http://www.ridge.k12.wa.us">www.ridge.k12.wa.us</a></td>
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<td>Longview School District 122</td>
<td>2715 Lilac Street Longview, WA 98632</td>
<td>360-575-7000</td>
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<td>Vancouver School District 37</td>
<td>PO Box 8937, Vancouver, WA 98668</td>
<td>360-313-1000</td>
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<td>Mill A School District 31</td>
<td>1142 Jessup Road, Cook, WA 98605</td>
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<td>Mount Pleasant School District 029-93</td>
<td>152 Marble Road, Washougal, WA 98671-9602</td>
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<td>Wahkiakum School District 200</td>
<td>PO Box 398, Cathlamet, WA 98612</td>
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<td>Naselle-Grays River Valley SD155</td>
<td>793 State Route 4, Naselle, WA 98638</td>
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<td>Washougal School District 112-6</td>
<td>4855 Evergreen Way, Washougal, WA 98671</td>
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<tr>
<td>White Salmon Valley SD 405-17PO</td>
<td>Box 157, White Salmon, WA 98672</td>
<td>509-493-1500</td>
<td><a href="http://www.schools.gorge.net">www.schools.gorge.net</a></td>
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<td>Wishram School District 94</td>
<td>PO Box 8, Wishram, WA 98673</td>
<td>509-748-2551</td>
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<td>Woodland School District 404</td>
<td>800 Third Street, Woodland, WA 98674-8467</td>
<td>360-225-9451</td>
<td><a href="http://www.woodlandschools.org">www.woodlandschools.org</a></td>
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</table>
The following page is the summary of a research article that was published in 2005. The article looked at many therapies that are often used to support kids with an ASD. The article reviewed many other studies that had looked at different therapies and placed the therapies into categories based on how strong the research and the results were.

The categories are:

- Scientifically Based Practice- which means that there has been valid research and sound results for that therapy.

- Promising Practice- which refers to the therapies that are generally accepted, but there is not much strong, reliable data to support the therapies.

- Limited Supporting Information for Practice- this category includes therapies that have no valid research, and/or have negative research results.

For the purposes of this parent guide, there is a third category. These are the therapies that we consider widely accepted my medical professionals.
### TABLE 1
Evaluation of Interventions and Treatments for Learners With Autism Spectrum Disorders

<table>
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<tr>
<th>Classification</th>
<th>Interpersonal relationship</th>
<th>Skill-based</th>
<th>Cognitive</th>
<th>Physiological/ biological/ neurological</th>
<th>Other</th>
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<tr>
<td>Scientifically based practice</td>
<td>• Play-oriented strategies</td>
<td>• Applied behavior analysis (Hagopian, Crockett, van Stone, DeLeon, &amp; Bowman, 2008)</td>
<td>• Learning Experiences: An Alternative Program for Preschoolers and Parents (Strain &amp; Hoyson, 2000)</td>
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<td>Promising practice</td>
<td>• Picture Exchange Communication System (Pyramid Educational Consultants, 2005)</td>
<td>• Incidental teaching (Charlop-Christy &amp; Carpenter, 2000)</td>
<td>• Cognitive behavioral modification (Zipoli, 2005)</td>
<td>• Sensory Integration (Case-Smith &amp; Bryant, 1999)</td>
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<td>• Structured teaching (e.g., TEACCH; Panerai, Ferrante, Caputo, &amp; Impellizzeri, 1998)</td>
<td>• Augmentative alternative communication (Ogletree, 1998)</td>
<td>• Cognitive learning strategies (Bock, 1999)</td>
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<td>• Assistive technology (Tjus, Hinmann, &amp; Nelson, 2001)</td>
<td>• Joint action routines (Prizant, Wetherby &amp; Rydell, 2000)</td>
<td>• Social stories (Rogers &amp; Myles, 2001)</td>
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<td></td>
<td>• Gentle teaching (Fox, Dunlop, &amp; Buschbaker, 2000)</td>
<td>• Van Dijk curricular approach (MacFarland, 2001)</td>
<td>• Social decision-making strategies (Myles &amp; Simpson, 2003)</td>
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<td></td>
<td>• Option method (e.g., Son-Rise program; Option Institute and Fellowship, 2004)</td>
<td>• Fast ForWord (Gillam, Leob, &amp; Friet-Patti, 2001)</td>
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<td></td>
<td>• Floor time (Green &amp; Wieder, 2000)</td>
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<td></td>
<td>• Pet/animal therapy (McGinney, Dustin, &amp; Wolff, 2001)</td>
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<td></td>
<td>• Relationship development intervention (Gusten &amp; Sheely, 2002)</td>
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<td>Limited supporting information for practice</td>
<td>• Cognitive scripts (Krantz &amp; McClannahan, 1998)</td>
<td>• Cartooning (Rogers &amp; Myles, 2001)</td>
<td>• Scotopic sensitivity syndrome: Irlen lenses (Griffin, Christenson, Wesson, &amp; Erickson, 1997)</td>
<td>• Music therapy (Brownell, 2002)</td>
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<td>• Holding therapy (Waterhouse, 2000)</td>
<td>• Power cards (Gagnon, 2001)</td>
<td>• Auditory integration training (Mudford et al., 2000)</td>
<td>• Art therapy (Kornreich &amp; Schimmel, 1991)</td>
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<tr>
<td>Not recommended</td>
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<td>• Facilitated communication (Perry, Bryson, &amp; Bobko, 1998)</td>
<td>• Megavitamin therapy (Adams &amp; McGinnis, 2001)</td>
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<td></td>
<td></td>
<td></td>
<td>• Feingold diet (Tsai, 1998)</td>
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<td>• Herb, mineral, and other supplements (Tolbert, Haigler, Waits, &amp; Dennis, 1993)</td>
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</table>

Glossary of Autism Related Terms

**Americans with Disabilities Act (ADA)** is the US law that ensures rights of persons with disabilities with regard to employment and other issues.

**Applied Behavior Analysis (ABA)** is a style of teaching using series of trials to shape desired behavior or response. Skills are broken into small components and taught to child through a system of reinforcement.

**Asperger’s Syndrome** is a developmental Disorder on the Autism Spectrum defined by impairments in communication and social development and by repetitive interests and behaviors, without a significant delay in language and cognitive development.

**Audiologist** is a professional who diagnoses and treats individuals with hearing loss or balance problems.

**Autism Diagnostic Observation Schedule (ADOS)** is test considered to be current gold standard for diagnosing ASD and, along with information from parents, should be incorporated into a child’s evaluation.

**Autism Spectrum Disorders** encompasses the following Disorders listed in DSM-IV: Autistic Disorder, Asperger’s Disorder, PDD – Not Otherwise Specified, Childhood Disintegrative Disorder, and Retts Disorder.

**Casein** is protein found in milk, used in forming the basis of cheese and as a food additive.

**Celiac Disease** is a disease in which there is an immunological reaction within the inner lining of the small intestine to gluten, causing inflammation that destroys the lining and reduces the absorption of dietary nutrients. It can lead to symptoms of nutritional, vitamin and mineral deficiencies.

**Childhood Disintegrative Disorder** is a Disorder in which development begins normally in all areas, physical and mental. At some point between 2 and 10 years of age, the child loses previously developed skills. The child may lose social and language skills and other functions, including bowel and bladder control.

**Chronic Constipation** is an ongoing condition of having fewer than three bowel movements per week.
**Cognition** is mental process of knowing, including aspects such as awareness, perception, reasoning and judgment.

**Cognitive Skills** are any mental skills that are used in the process of acquiring knowledge; these skills include reasoning, perception and judgment.

**Compulsions** are deliberate repetitive behaviors that follow specific rules, such as pertaining to cleaning, checking, or counting. In young children, restricted patterns of interest may be early sign of compulsions.

**Declarative Language** is used to communicate what the mind is producing. It is what is most common in conversation, whereas Imperative Language is used to ask questions, make commands or give instructions.

**Developmental Disorder** refers to several disorders that affect normal development. May affect single area of development (specific developmental disorders) or several (pervasive developmental disorders).

**Developmental Individual Difference Relationship (DIR)** is therapy, known as Floortime, which seeks to move the child toward increasingly complex interactions through mutually shared engagement.

**Developmental Milestones** skills or behaviors that most children can do by a certain age that enable the monitoring of learning, behavior, and development.

**Developmental Pediatrician** is a medical doctor who is board-accredited and has received sub-specialty training in developmental-behavioral pediatrics.

**Diagnostic and Statistical Manual** the official system for classification of psychological and psychiatric disorders published by the American Psychiatric Association.

**Discrete Trial Teaching (DTT)** is technique incorporating principles of ABA, including positive reinforcement. Not in itself ABA. Used to teach behaviors in one-to-one setting. Concepts are broken down into small parts.

**Dyspraxia** is brain’s inability to plan muscle movements and carry them out. In speech, this term may be used to describe Apraxia.

**Early Intervention (EI)** is a state-funded program designed to identify and treat developmental problems or other disabilities as early as possible. Eligibility for EI
is from birth to three years of age.

**Echolalia** is repeating words or phrases heard previously, either immediately after hearing word or phrase, or much later. Delayed echolalia occurs days or weeks later. Functional echolalia is using quoted phrase in a way that has shared meaning, for example, saying “carry you” to ask to be carried.

**Expressive Labeling** is the communication of a name for an object or person, see expressive language.

**Expressive Language** is communication of intentions, desires, or ideas to others, through speech or printed words. Includes gestures, signing, communication board and other forms of expression.

**Extended School Year (ESY)** Services are provided during breaks from school, such as during summer vacation, for students who experience substantial regression in skills during school vacations.

**Free Appropriate Public Education (FAPE)** means that education must be provided to all children ages three to twenty-one at public expense.

**Floortime** a developmental intervention for children with an ASD involving meeting a child at his current developmental level, and building upon a particular set of strengths.

**Fragile X** is a genetic disorder that shares many of the characteristics of an ASD. Individuals may be tested for Fragile X.

**Gastroenterologist** doctor specializing in diagnosis & treatment of disorders of GI tract, including esophagus, stomach, small intestine, large intestine, pancreas, liver, gallbladder & biliary system.

**General Education** is a pattern of courses in multiple subjects taught to the same grade level to deliver a well-balanced education.

**Geneticist** refers to a medical doctor who specializes in genetic problems. Genes are the unit in the chromosome that contain the blueprint for the transmission of inherited characteristics.
**Gestures** are hand and head movements, used to signal to someone else, such as a give, reach, wave, point, or head shake. They convey information or express emotions without the use of words.

**Global Developmental Delay** is diagnosis in children younger than 6, characterized by delay in two or more developmental domains, sometimes associated with mental retardation.

**Gluten** is a protein present in wheat, rye, and barley.

**Hyperlexia** is the ability to read at an early age. To be hyperlexic, a child does not need to understand what he or she is reading.

**Hyposensitivity, Hyporesponsiveness**, is abnormal insensitivity to sensory input. Child who appears to be deaf, whose hearing is normal, is under reactive. Child who is under reactive to sensory input may have a high tolerance to pain, may be clumsy, sensation-seeking, and may act aggressively.

**Hypotonia** is a term that means low muscle tone.

**Incidental Teaching** teaches a child new skills while in their home or community, in natural context or “in the moment,” to help make sense of what they learn during formal instruction and generalize new skills.

**Individual Family Service Plan (IFSP)** is developed by a multidisciplinary team including family as primary participant. Describes child’s level of developmental in all areas; family’s resources, priorities, & concerns, services to be received and the frequency, intensity, and method of delivery. Must state natural environments in which services will occur.

**Individualized Education Plan (IEP)** identifies student’s specific learning expectations, how school will address them with appropriate services, and methods to review progress. For students 14 & older, must contain plan to transition to postsecondary education or the workplace, or to help the student live as independently as possible in the community.

**Individuals with Disabilities Education Act (IDEA)** is the US law mandating the “Free and Public Education” of all persons with disabilities between ages 3 and 21.

**Inclusion** involves educating all children in regular classrooms, regardless of degree or severity of disability. Effective inclusion takes place with planned system of training and supports; involves collaboration of multidisciplinary team including regular and special educators.
Joint Attention is the process of sharing one’s experience of observing an object or event, by following gaze or pointing gestures. Critical for social development, language acquisition, cognitive development. Impairment in joint attention is a core deficit of ASD.

Least Restrictive Environment (LRE) a setting that least restricts opportunities for child with disabilities to be with peers without disabilities. The law mandates that every child with a disability be educated in a Least Restrictive Environment.

Mainstreaming is where students are expected to participate in existing regular ed classes, whereas in an inclusive program classes are designed for all students. May be gradual, partial, or part-time process (e.g., student may attend separate classes within regular school, or participate in regular gym and lunch only).

Melatonin is a hormone produced by pineal gland, involved in regulating sleeping and waking cycles. Sometimes used for chronic insomnia. Consult your child's physician before giving melatonin; it is not recommended for all patients with sleep problems.

Mental Retardation describes person with limitations in mental functioning that cause them to develop more slowly than typical child. They may take longer to learn to speak, walk, and take care of personal needs such as dressing or eating, and are likely to have trouble learning in school. May be mild or severe.

Modified Checklist of Autism in Toddlers (MCHAT) is a screening tool for identifying young children who may be referred to specialist for further evaluation and possible Autism Spectrum Disorder diagnosis.

Motor deficits are physical skills that a person cannot perform or has difficulty performing.

Motor function (or Motor Skills) is the ability to move and control movements.

Neurologist refers to a doctor specializing in medical problems associated with the nervous system, specifically the brain and spinal cord.

Nonverbal Behaviors are things people do to convey information or express emotions without words, including eye gaze, facial expressions, body postures, and gestures.

Obsessions are persistent and intrusive repetitive thoughts. Preoccupations with specific kinds of objects or actions may be an early sign of obsessions.
**Occupational Therapy** assists development of fine motor skills that aid in daily living. May focus on sensory issues, coordination of movement, balance, and self-help skills such as dressing, eating with a fork, grooming, etc. May address visual perception and hand-eye coordination. **Occupational Therapist** helps minimize impact of disability on independence in daily living by adapting child’s environment and teaching sub-skills of the missing developmental components.

**Operant Conditioning** is the modification of behavior through positive and/or negative reinforcement.

**Perseveration** is repetitive movement or speech, or sticking to one idea or task, that has a compulsive quality to it.

**Pervasive Developmental Disorders (PDD)** group of conditions involving delays in development of many basic skills, including ability to socialize with others, to communicate and use imagination. Includes Autism, Asperger’s Syndrome, Childhood Disintegrative Disorder, Rett Syndrome and Pervasive Development Disorder - Not Otherwise Specified.

**Pervasive Developmental Disorder – Not Otherwise Specified (PDD-NOS)** a category of PDD referring to children having significant problems with communication & play, and some difficulty interacting with others, but are too social for diagnosis of an ASD.

**Physical Therapy** uses specially designed exercises and equipment to help patients regain or improve their physical abilities.

**Physical Therapists** design and implement physical therapy programs and may work within a hospital or clinic, in a school, or as an independent practitioner.

**Pica** is persistent eating or mouthing of non nutritive substances for at least 1 month when behavior is developmentally inappropriate (older than 18-24 months). Substances may include items such as clay, dirt, sand, stones, pebbles, hair, feces, lead, laundry starch, vinyl gloves, plastic, erasers, ice, fingernails, paper, paint chips, coal, chalk, wood, plaster, light bulbs, needles, string, cigarette butts, wire, and burnt matches.

**Picture Exchange Communication System (PECS)** an alternative communication system using picture symbols. Taught in phases starting with simple exchange of symbol for desired item. Individuals learn to use picture symbols to construct complete sentences, initiate communication, & answer questions.
**Pivotal Response Treatment (PRT)** therapeutic teaching method using incidental teaching opportunities to target and modify key behaviors.

**Receptive Language** the ability to comprehend words and sentences. Begins as early as birth and increases with each stage in development. By 12 months a child begins to understand words and responds to his own name and may respond to familiar words in context. By 18 to 20 months a child identifies familiar people by looking when named (e.g., Where’s mommy?), gives familiar objects when named (e.g., Where’s the ball?), and points to a few body parts (e.g., Where’s your nose?). These skills commonly emerge slightly ahead of expressive language skills.

**Reinforcement or reinforcer** is any object or event following a response, increasing or maintaining the rate of responding. Positive reinforcer may be produced by, or be added after a response. **Relationship Development Intervention (RDI)** a therapeutic teaching method based on building intelligence competencies of social connection -- such as referencing, emotion sharing, coregulation, and experience sharing -- that normally develop in infancy and early childhood.

**Respite Care** is temporary, short-term care provided to individuals with disabilities, delivered in the home for a few short hours or in an alternate licensed setting for an extended period of time. Respite care allows caregivers to take a break in order to relieve and prevent stress and fatigue.

**Self Regulation and self-control** are related but not the same. Self-regulation refers to both conscious and unconscious processes that have an impact on self-control, but regulatory activities take place more or less constantly to allow us to participate in society, work, & family life. Self-control is a conscious activity.

**Sensory Defensiveness** is a tendency, outside the norm, to react negatively or with alarm to sensory input which is generally considered harmless or non-irritating to others. Also called hypersensitivity.

**Sensory Integration** is the way the brain processes sensory stimulation or sensation from the body & then translates that information into specific, planned, coordinated motor activity.

**Sensory Integration Dysfunction** a neurological disorder causing difficulties processing information from the five classic senses (vision, hearing, touch, smell, & taste), sense of movement (vestibular system), and positional sense (proprioception). Sensory information is sensed normally, but perceived abnormally. May be a disorder on its own, or with other neurological conditions.
**Sensory Integration Therapy** is used to improve ability to use incoming sensory information appropriately & encourage tolerance of a variety of sensory inputs.

**Sensory stimulus** agent, action or condition, internal (e.g., heart rate, temperature) or external (e.g., sights, sounds, tastes, smells, touch, & balance) that elicits physiological or psychological response. Response depends on ability to regulate & understand stimuli & adjust emotions to demands of surroundings.

**Sleep Hygiene** a set of practices, habits & environmental factors critically important for sound sleep, such as minimizing noise, light & temperature extremes & avoiding naps & caffeine.

**Social Reciprocity** back-and-forth flow of social interaction. How behavior of one person influences & is influenced by behavior of another & vice versa.

**Social Stories,** developed by Carol Gray, are simple stories that describe social events & situations that are difficult for a child with a PDD to understand. For example, a social story might be written about birthday parties if the child appears to have a difficult time understanding what is expected of him or how he is supposed to behave at a birthday party.

**Social Worker** is a trained specialist in the social, emotional & financial needs of families & patients. Social workers often help families & patients obtain the services they have been prescribed.

**Special Education** is specially designed instruction, at no cost to families, to meet unique needs of child with disability, including instruction conducted in the classroom, in the home, in hospitals & institutions, & in other settings & instruction in physical education.

**Speech & Language Therapy** is provided with the goal of improving an individual’s ability to communicate. This includes verbal and nonverbal communication. The treatment is specific to the individual’s need.

**Spoken Language** (also referred to as expressive) use of verbal behavior, or speech, to communicate thoughts, ideas, & feelings with others. Involves learning many levels of rules - combining sounds to make words, using conventional meanings of words, combining words into sentences, and using words & sentences in following rules of conversation.

**Stereotyped Behaviors** refer to an abnormal or excessive repetition of an action carried out in the same way over time. May include repetitive movements or posturing of the body or objects.
**Stereotyped Patterns of Interest** or restricted patterns of interest refer to a pattern of preoccupation with a narrow range of interests and activities.

**Stim, or “self-stimulation”** behaviors that stimulate ones senses. Some “stims” may serve a regulatory function (calming, increasing concentration, or shutting out an overwhelming sound).

**Symbolic Play** is where children pretend to do things & to be something or someone else. Typically develops between the ages of 2 & 3 years. Also called make believe, or pretend play.

**Syndrome** is a set of signs & symptoms that collectively define or characterize a disease, Disorder or condition.

**Tactile Defensiveness** is a strong negative response to a sensation that would not ordinarily be upsetting, such as touching something sticky or gooey or the feeling of soft foods in the mouth. Specific to touch.

**Training and Education of Autistic and Related Communication Handicapped Children (TEACCH)** is a therapeutic approach broadly based on the idea that individuals with an ASD more effectively use & understand visual cues.

**Typical Development** (or healthy development) describes physical, mental, & social development of a child who is acquiring or achieving skills according to expected time frame. Child developing in a healthy way pays attention to voices, faces, & actions of others, showing & sharing pleasure during interactions, & engaging in verbal & nonverbal back-and-forth communication.

**Verbal Behavior** is a method of Applied Behavioral Analysis (ABA) for teaching children with an ASD, based on B.F. Skinner’s description of the system of language.

The above glossary is a shortened version of the glossary from: *The 100 Day Kit version 2.0* by the Autism Speaks organization. You can view the kit at: [www.Autismspeaks.org](http://www.Autismspeaks.org)
Personal Information

Things to Add to This Area

- Test Results and Reports
- Provider Information
- IEP documents
- Other Relevant Information