Easy Read Edition

Institutions:
The Old, The New, And What We Should Do
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Institutions:
The Old, The New, And What We Should Do

Part 1: To Start
To Start

This toolkit is about institutions. It is also about living in the community.

It talks about why institutions are bad.

It talks about why people with disabilities should live in the community.

The toolkit will answer questions like:

• What is an institution? What is living in the community
• When did institutions get made? What were they like?

• Why did people start closing institutions?

• What rules and laws help us live in the community?

• What happened when people moved from institutions into the community?

• Why is living in the community better than living in institutions?
• What kinds of institutions are there today?

• What are new institutions like? How are they different from the old institutions?
We talk a lot about institutions in this toolkit.

We talk about people getting hurt in institutions.

It may hurt to read about it.

It’s okay to take a break from reading if you need to.

It’s okay to not read this toolkit if it will hurt you.
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Institutions: The Old, The New, And What We Should Do

Part 2: Institutions and Living in the Community
Words to Know in Part 2
Long-term supports and services (LTSS)

Services that help people with disabilities live our everyday lives. Some kinds of LTSS are job coaches and in-home helpers.

Institutions

Places where a lot of disabled people live. People living in institutions did not decide to live there. They usually can’t leave when they want to.
Home and community-based services (HCBS)

Getting LTSS in the community.
Institutions and Living in the Community

Some people with disabilities need long-term services and supports (LTSS).

LTSS are services that help people with disabilities live our everyday lives.

Some kinds of LTSS are:

- Job coaches
- Transportation
- An in-home helper
Some people with disabilities get LTSS in institutions.

Institutions are places where a lot of people with disabilities live.

People living in institutions did not decide to live there.

They were usually put there by someone else.

Institutions are not run by the people living there.
Institutions are run by the people who work there.

People living in institutions usually can’t leave when they want to.

They usually can’t spend their free time how they want to.

Other people with disabilities get LTSS in their community.

They get LTSS in their own homes.
Getting LTSS in the community is also called **home and community-based services**.

Some people call them “**HCBS**” for short.

People with disabilities have the right to live in the community.

We want to live in the community.

We don’t want to live in institutions.
ASAN is a group of autistic people.

We work on disability rights.

We believe people with disabilities have the same rights as everyone else.

One important right ASAN fights for is the right to live in the community.

ASAN talked with some people with disabilities.
We talked about what living in the community means to us.

We decided that living in the community means things like:

• Making your own choices.

• Going out when you want to, and going where you want to.

• Living in the same places as everyone else.
• Getting the help you need to meet your goals.

• Getting to choose who helps you, and what help you get.

• Meeting new people, and being with family and friends.

• Getting treated like an adult.

You can learn more by reading What Does “Living in the Community” Mean?
Today, less people with disabilities live in institutions than before.

Lots of institutions closed in the last 50 years.

More people with disabilities moved into the community.

That’s great!

But everyone should live in the community.
Some people still live in institutions.

And new kinds of institutions are still getting made.

So we still have work to do.

We need to show others why the community is good.

We need to show others why institutions are bad.
So it’s important to learn about institutions.

This toolkit will talk about the history of institutions.

It will show why living in the community is better.
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Institutions: The Old, The New, And What We Should Do

Part 3: The History of Institutions
Words to Know in Part 3
State hospital

The first kind of institutions.

State school

The first institutions just for people with intellectual disabilities.
How did institutions get started?

Institutions first got made in the 1840s.

Before that, some people with disabilities got put in prisons.

People with disabilities got hurt in prisons.

They got hit or yelled at by the prison workers.
They didn’t get food or water

They didn’t get help going to the bathroom.

They got kept in cold rooms.

They got left alone for a long time.

They didn’t get the health care they needed
A woman named Dorothea Dix saw people with disabilities in prisons.

She saw how they were getting hurt in prisons.

So she helped make different places to put people with disabilities.

These places were called state hospitals.

State hospitals were the first kind of institutions.
A few years later, another kind of institution got made.

These places were called state schools.

**State schools** were the first institutions just for people with intellectual disabilities.
What were state schools like?

State schools got made to try and help people with disabilities.

People thought we wouldn’t get hurt in a place “just for us”.

They thought state schools would be better than prisons.

But they still were like prisons in a lot of ways.
People with disabilities couldn’t leave the state schools.

They didn’t get to choose how they spent their time.

They didn’t get to be with non-disabled people.

Soon, people with disabilities started getting hurt in state schools.

They got hit or yelled at by the school staff.
They didn’t get food or water.

They didn’t get help going to the bathroom.

They got kept in cold rooms.

They got left alone for a long time.

They didn’t get the health care they needed.
The state schools became just like prisons.

State schools kept people with disabilities separate from everyone else.

So other people didn’t notice that state schools hurt people with disabilities.

They kept getting hurt.

A couple of times, someone noticed how bad state schools were.
They would let the newspaper or the government know.

Then, the state school would stop hurting people for a little bit.

But people didn’t check the state schools that much.

The state schools would start hurting people again after a while.

This kept happening for a long time.
New institutions kept getting made.

They said they wanted to help people with disabilities.

But they still took away our rights. They hurt us.

They were never good places to live.
When did people start trying to help us live in the community?

In the early 1900s, a lot more people with disabilities got put in institutions.

Some institutions had thousands of people in them.

But institutions didn’t get more money.

They didn’t hire more staff or build more rooms.
So institutions became even worse to live in.

In the mid-1900s, some parents of people with disabilities got together.

They saw how bad institutions were.

They didn’t want their kids to live in institutions.

They thought their kids should get to live in the community.
They didn’t put their kids in institutions.

They fought for the rights of their kids with disabilities.

John F. Kennedy was the President from 1961-1963.

He had a sister named Rosemary.

She had an intellectual disability.
He talked with the parents of kids with disabilities.

They worked together to help people with disabilities.

They made the government set aside money to help people with disabilities.

Some of this money went to institutions.

But some of it also went to helping people live in the community.
This was the first step to moving people out of institutions.

In 1972, a TV reporter visited an institution.

The institution was called Willowbrook.

The reporter saw how bad Willowbrook was.

People with disabilities were crowded in small rooms.
They didn’t get to eat. They got sick a lot.

The reporter took a video of Willowbrook. He put it on TV.

That is how lots of people learned how bad institutions were.

It made more people want to close institutions.

Lots of changes happened after people learned about Willowbrook.
States started working to move people out of institutions.

They wanted more people with disabilities to live in the community.

So they started paying for LTSS in the community.

Lots of rules and laws also got made about people with disabilities.

These rules try to make sure we don’t get hurt in institutions.
They try to help us live in the community.
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Part 4: Living in the Community: Important Rules and Laws
Words to Know in Part 4
**DD Act**

A law that got made in 1975. It helps protect the rights of people with developmental disabilities.

**Developmental Disability Bill of Rights**

Part of the DD Act. It lists what rights we have when we get LTSS. It also made rules that institutions have to follow.
Medicaid

A health care program in the United States. Medicaid helps people with disabilities get health care.

HCBS Settings Rule

A rule the government made. It makes sure that Medicaid money set aside for HCBS goes to the right places.
The Americans with Disabilities Act (ADA)

A law that came out in 1990. The ADA gave people with disabilities rights.

Supreme Court

The biggest court in the United States. They have the final say on how laws work.
Olmstead

A big court case in 1999. It says that you have a right to live in the community. It says that you don’t have to live in an institution.
Living in the Community: Important Rules and Laws

DD Act

The **DD Act** is a law that got made in 1975.

It helps protect the rights of people with developmental disabilities.

It became a law 3 years after people learned about Willowbrook.

It tries to make sure institutions can’t hurt people like Willowbrook did.
Part of the DD Act is the Developmental Disability Bill of Rights.

1. It lists what rights we have when we get LTSS.

2. It says that we should get to live the way we want.

3. It says we should have freedom.

4. It also made rules that institutions have to follow.
It says that institutions can’t get money from the government if they hurt us.
Medicaid and the HCBS Settings Rule

**Medicaid** is a health care program in the United States.

The government runs Medicaid.

Medicaid helps people with disabilities get health care.

Most people who get LTSS get them paid for by Medicaid.
At first, Medicaid didn’t help us live in the community.

Medicaid only paid for people to get LTSS in institutions.

Medicaid wouldn’t pay for us to get LTSS in the community.

In the 1970s, more people started trying to live in the community.

They needed LTSS in the community.
They asked Medicaid to help pay.

In 1982, Medicaid started trying to help people live in the community.

Medicaid set aside some money to give people LTSS in the community.

More people got LTSS in the community.

So more and more people got to live in the community.
Institutions started getting smaller.

But there wasn’t enough money for everyone to get LTSS in the community.

Many people with disabilities got put on waiting lists.

They had to wait years to get LTSS in their communities.

Lots of people are still on waiting lists.
And there was another problem.

Institutions took some of the money Medicaid set aside.

They said they helped people live in the community.

But they were lying. They were still institutions.

Institutions used up some of the money Medicaid set aside.
So there was less money to actually help us live in the community.

The government wanted to make sure that stopped happening.

They made a rule called the **HCBS Settings Rule**.

This rule makes sure that Medicaid money only goes to LTSS in the community.

It makes sure institutions can’t take money meant for living in the community.
The rule hasn’t started working yet. It starts in 2022.

You can learn more by reading the “This Rule Rules!” Toolkit.
The ADA and Olmstead

The **Americans with Disabilities Act** is a law that came out in 1990.

People call it the ADA for short.

The ADA gave people with disabilities rights.

It says we have the same rights as non-disabled people.
It says we have the right to go to the same places as non-disabled people.

It says we have the right to live and work in the same places as non-disabled people.

In 1999, two women were living in an institution.

Their names were Lois Curtis and Elaine Wilson.
They didn’t want to live in an institution.

They wanted to live in the community.

Lois and Elaine said the ADA gave them the right to live in the community.

They fought for their rights from the ADA.

They went to the Supreme Court to fight for their rights.
The Supreme Court is the biggest court in the United States.

They have the final say on how laws work.

The Supreme Court said that Lois and Elaine were right.

Lois and Elaine’s court case was called Olmstead.

Olmstead means that you have a right to live in the community.
It says that you don’t have to live in an institution.

You have the right to get LTSS in the community.
Institutions Today

These rules and laws helped us get LTSS in the community.

They helped us move out of institutions.

More and more people with disabilities got to live in the community.

So now, less people live in institutions.
Some states started closing institutions in the 70s and 80s.

In the 90s, some states closed all of their big institutions.

Today, 17 states have no big institutions.

But most states still have at least one.
Today, most people with disabilities live in the community.

But some people still live in institutions.

Some still live in bigger institutions, like state schools.

These are sometimes called ICFs or developmental centers.

Others live in smaller institutions, like group homes.
We want to make sure everyone can live in the community.
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Institutions:
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Part 5: Why is Community Living Better?
Why is Community Living Better?

There are lots of people that study institutions.

They talk to people with disabilities in institutions.

They see what life is like in institutions.
They also study living in the community.

They talk to people with disabilities who live in the community.

They see what life is like for people with disabilities in the community.

They also talk to people who moved out of institutions.

They ask about how living in the community is different from institutions.
Lots of studies came out about institutions and the community.

You can see a list of some of these studies at the end of this toolkit.

The studies can be hard to read.

So, this part of the toolkit explains what the studies found out.
The studies show that living in the community is better than living in institutions.

The community is a better place to live for a lot of reasons.

Here are some things the studies say:
Freedom and Happiness

People with disabilities get more freedom when we live in the community.

Studies say that people who live in the community get to make more choices.

We get to choose what we do during the day.

We get to choose when and where to go out.
We get to choose who lives with us, and who helps us.

We also get to see friends and family more.

We get less choices when we live in institutions.

We might not get to choose who lives with us.

We might not get to choose where we go and what we do.
Getting to make our own choices helps in other ways, too.

It helps us feel better about ourselves.

It feels good to have control over our lives.

It lets us make sure things work well for us.
For example:

Sunil is autistic. He lives in an institution.

He doesn’t get to choose his roommate.

His roommate is very loud.

Loud noises hurt Sunil. He needs his room to be quiet.
But the institution doesn’t let Sunil get a different roommate.

Sunil is unhappy at the institution.

Then, Sunil moves into the community.

He gets to choose his own roommate.

He chooses a roommate who is also quiet.
Sunil is happier living in the community.

He gets to have his own quiet space.

He feels better because he gets what he needs.
One study talked to people with disabilities who moved out of institutions.

They said they liked living in the community more.

They never wanted to move back to an institution.
Learning New Things

People with disabilities learn more when we live in the community.

Living in the community helps us learn new things.

We learn important skills that help us live on our own.

We learn how to take care of ourselves.
We learn things like how to make a schedule.

We learn how to cook, and how to clean our house.

We learn to shop for our own food and clothes.

We don’t get to learn these things as much in institutions.
We also learn more about getting along with other people.

We get to meet more non-disabled people when we live in the community.

So we get more chances to talk to others.

We learn to communicate in ways that work for us.
We get treated better in the community.

So, we feel better about ourselves. We feel safer.

That makes it easier for us to learn.

We can learn new ways to say what we need.

We can learn new ways to get along with people.
Less people try to communicate with us in institutions.

We get less chances to learn about communicating.

We might not get treated well in institutions.

That makes it harder to get along with other people.
Learning new things helps us in other ways, too.

For example, when we learn to shop, we also learn about money.

Learning about money helps us get better at math.

Learning to make a schedule can help us read better.

Talking with more people can help us write better.
Living in the community helps us get better at all these things.

Not everyone learns all these things.

But we usually learn more when we live in the community.

We don’t learn as much when we live in institutions.
Being Part of the Community

Living in the community lets us be around more non-disabled people.

We can work in the community and make money.

We can join groups in the community.

We can make new friends and get new hobbies.
We can feel like a part of the community.

We can feel like we belong in the community.

Institutions keep us separate from non-disabled people.

We might not get to make friends outside the institution.

We might not get to go into the community at all.
We might not feel like we belong in the community.
Staying Safe

People with disabilities are safer in the community.

We get hurt less when we live in the community.

We get hit less.

We get yelled at less.
We don’t get left alone when we need help.

We get food when we need it.

We get help going to the bathroom when we need to.
People with disabilities get hurt a lot more in institutions.

We still get hit and yelled at.

We still might not get food or water.

We might get left alone when we need help.
The government made rules to try and keep us safe in institutions.

1.
2.

But we still get hurt in institutions today.

The same things that happened in state schools still happen today.
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Part 6: Questions about Institutions
Questions about Institutions

Is Community Living Better for Everyone?

Some people say that certain people with disabilities can’t live in the community.

They say that some people with disabilities have to live in institutions.

They think some people can only get the help they need in an institution.
Some people with disabilities need a lot of help to live in the community.

People with intellectual disabilities might need a lot of help.

Non-speaking people might have a hard time showing others what they need.

But that doesn't mean people with these disabilities need to live in institutions.

Everyone should get to live in the community.
The studies say that everyone should live in the community, too.

The studies found out that living in the community is better for everyone.

It helps everyone make more choices and learn new things.

It helps everyone be more safe.

It doesn’t matter what kind of disability someone has.
People with intellectual disabilities do better in the community.

Non-speaking people do better in the community.

People who need a lot of help do better in the community.

Nobody needs to be in an institution.
Does the community have no problems at all?

The studies found that the community is better than institutions.

But we are still learning how to help people live in the community.

There are still things that can be better.
Problems can still happen in the community.

People can still get hurt in the community.

We don’t always have enough choices.

Sometimes, we get treated unfairly.
The studies show that these problems happen less in the community.

But they do still happen.

We need to work hard to help fix these problems.

Everyone deserves a good life in the community.
What Counts as an Institution?

Studies have different ideas about what counts as an institution.

Some studies say an institution has to have more than 16 people.

Others say an institution has to have more than 4 people.
The studies also found out that some institutions are less bad than others.

The ones with less people are usually less bad.

For example:

Richelle lives in an ICF. 30 people with disabilities live there.

She doesn’t get to choose what she eats.

She never gets to leave the institution.
Maya lives in a group home. 5 people with disabilities live there.

Maya gets to choose what they eat for breakfast.

They take a field trip to the movies once a week.

But Maya can only go to the movies.

They can’t go anywhere else.
Maya gets to make more choices than Richelle.

Maya gets to go out more than Richelle.

But they still both live in institutions.
Smaller institutions might be less bad than bigger institutions.

But they all take away the rights of people with disabilities.

They don’t let us make all the choices about our lives.

They keep us separate from other people.
ASAN thinks about institutions in a different way.

We think big buildings and group homes are institutions.

But the most important thing is how people get treated.

Smaller places can still be institutions.

Someone who lives with their family can be in an institution.
Even someone who lives in their own apartment can be in an institution.

If they get kept separate from other people, then they are in an institution.

If they don’t get to make choices, then they are in an institution.
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Part 7: New Institutions
New Institutions

We know that living in the community is better than institutions.

But people keep building new institutions.

They don’t call them institutions.

They might call them group homes.

Or, they call them “intentional communities”.
They try and say these new places aren’t like the old institutions.

But they are still institutions.

The new institutions might look like living in the community.

They might be in a regular house.

They might look like every other neighborhood.
But only disabled people live there.

They don’t get to be around non-disabled people.

They don’t get to decide what to do each day.

They don’t get to go outside their neighborhood.
Some institutions say that people with disabilities choose to live there.

But usually, the people with disabilities didn’t get good choices.

They didn’t get to learn about living in the community.

They got told they couldn’t live in the community.

They didn’t really get a choice.
Just like when they got sent to the old institutions.

People keep making these new institutions.

That’s because they think we need institutions.

They think people with disabilities need to live in places that are “just for us”.

But that is the same reason the old institutions got made.
The old institutions kept us away from other people.

They made it easier for us to get hurt.

They took away our freedom.

New institutions do the same thing.
We know the history of institutions.

We know that institutions kept hurting us for many years.

We know that institutions keep hurting us today.

We know that studies say we should live in the community.
We also know one very important thing.

We know that people with disabilities don’t want to live in institutions.

We know that we want to live in the community.
We need to make sure new institutions stop getting made.

We need to make sure people know about new institutions.

People should know that new institutions are just like the old ones.
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Studies on Institutions and Community Living

Costs and Outcomes of Community Services for Persons with I/DD. By Roger J. Stancliffe and K. Charlie Lakin.

Deinstitutionalization: Unfinished Business. From the National Council on Disability.


Home and Community-Based Services: Creating Systems for Success at Home, at Work and in the Community. From the National Council on Disability.

Medicaid Home and Community-Based Services for Persons with I/DD. By K. Charlie Lakin, Robert Doljanac, Soo Yong Byun, Roger Stancliffe, Sarah Taub, and Giuseppina Chiri.
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Terms to Know
The Americans with Disabilities Act (ADA)

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DD Act

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Developmental Disability Bill of Rights

Part of the DD Act. It lists what rights we have when we get LTSS. It also made rules that institutions have to follow.

Home and community-based services (HCBS)

Getting LTSS in the community.
HCBS Settings Rule

A rule the government made. It makes sure that Medicaid money set aside for HCBS goes to the right places.

Institutions

Places where a lot of disabled people live. People living in institutions did not decide to live there. They usually can’t leave when they want to.
Long-term services and supports (LTSS)

Services that help people with disabilities live our everyday lives. Some kinds of LTSS are job coaches and in-home helpers.

Medicaid

A health care program in the United States. Medicaid helps people with disabilities get health care.
**Olmstead**

A big court case in 1999. It says that you have a right to live in the community. It says that you don’t have to live in an institution.

**State hospitals**

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Supreme Court

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