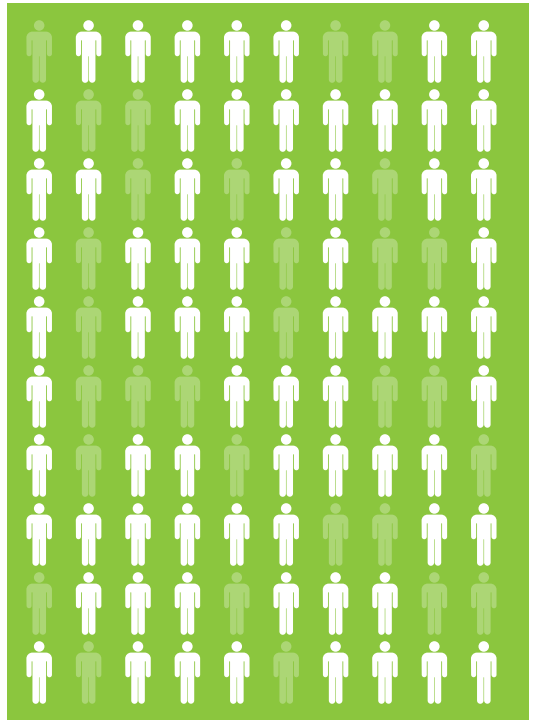


Early testing and diagnosis of Primary Immunodeficiency



**PI is largely
underdiagnosed:
between 70-90% of
people living with
a PI worldwide
still remain
undiagnosed**

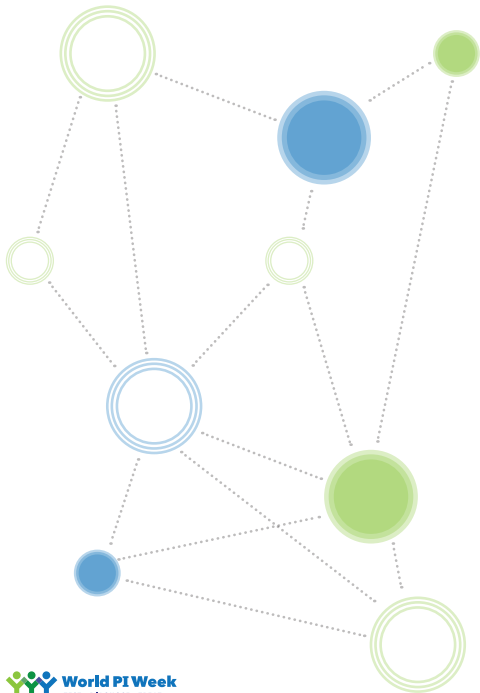


**Early diagnosis
and treatment
can prevent
complications and
enable people to
live as normal a life
as possible**



**Early diagnosis
contributes greatly
to improved quality
of life and care for
people with PI**

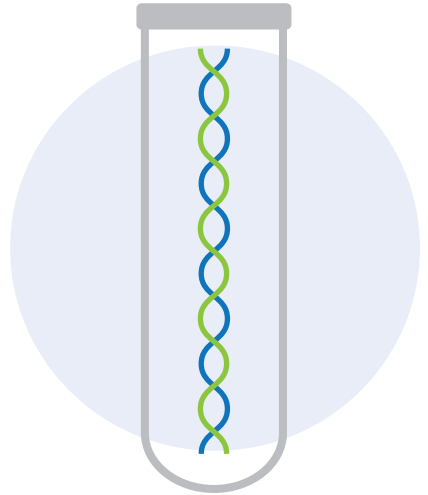




Main reasons for a delayed or misdiagnosis of PI:

- **Complex nature of PI**
- **Variety of symptoms and clinical manifestations**
- **Relatively low awareness of symptoms among primary care doctors and specialists**

**Genetic technology
allowing genetic
testing has a
significant role in
the diagnosis of PI**





**For severe forms
of PI, newborn
screening is a
lifesaving, cost-
effective method
to drive early
diagnosis and save
lives**



**Prompt diagnosis
of PI leads to better
use of healthcare
facilities and
services, and lower
healthcare costs**

Public awareness of symptoms, greater education among both primary and specialist care providers at pre/post graduate levels are paramount for PI recognition





**Patient access
to care and
appropriate
treatment can help
prevent serious and
life-threatening
illnesses in people
with PI**

**Screening
tests should be
accessible to the
whole range of
hospital doctors
and primary care
providers**



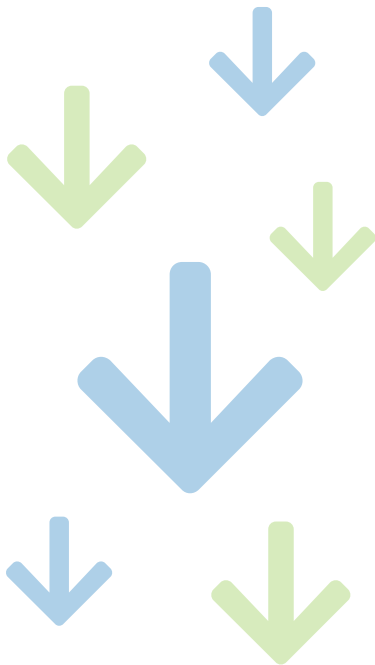
Routine newborn screening programmes for severe PI (e.g., SCID and XLA) should be widely implemented in public healthcare settings in all countries

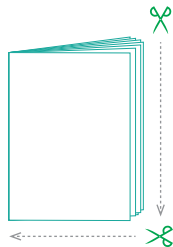
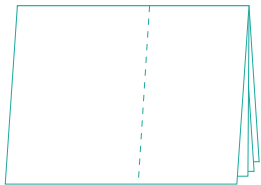
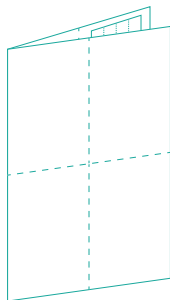
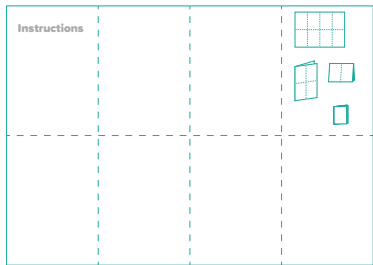


Access to genetic testing should become available to patients and widespread in medical specialties to provide earlier detection of PI and diagnose unknown forms



Here is how to
build your own
“Early testing and
diagnosis
of Primary
Immunodeficiency”
brochure







World PI Week

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