

The Undiagnosed Diseases Network International

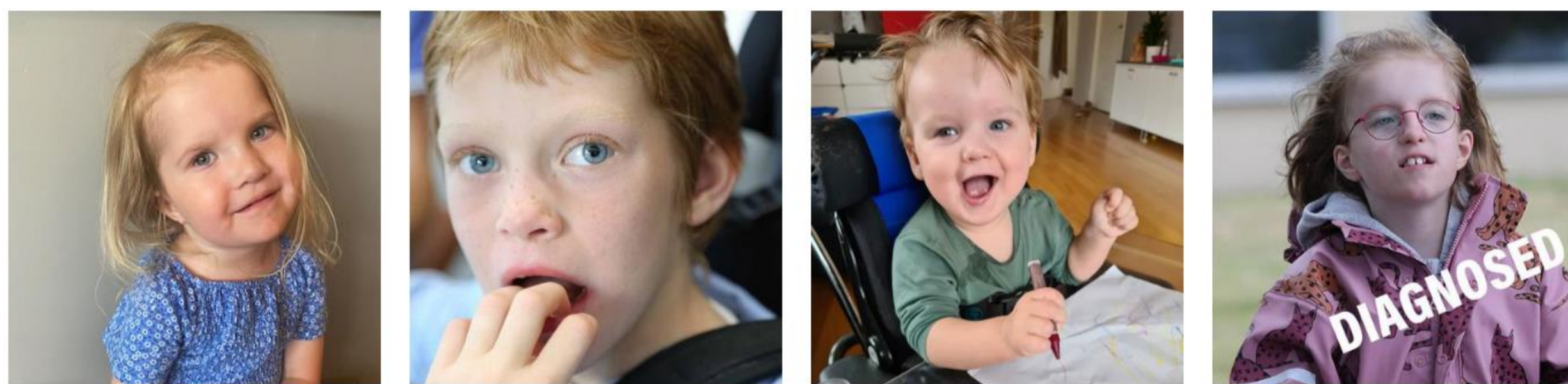


www.udninternational.org



Principles

- Patients enrolled in the UDNI are selected for the unique characteristics of their disorder and for its potential to inform new aspects of cell biology, pathogenetic mechanism(s) and therapy. Candidate patients should have been extensively examined already, so that obvious diagnoses have been eliminated.
- Accepted patients should be thoroughly evaluated by the UDNI, preferably at no cost to the patient.
- Patients should consent to share their data with other investigators within the group, according to the tenets of the Helsinki Declaration and/or of Good Clinical Practices. Patient phenotypes should be expressed using a standard ontology system in order to build up a highly integrated database.
- Next-generation sequencing and other -omics analyses (e.g., proteomics, metabolomics) should be performed on enrolled families/patients (trios or quartets when possible), and analyzed with some uniformity and according to state-of-the-art protocols.
- The -omics and phenotypic data should be shared among members of the UDNI.
- Functional studies should be performed to substantiate causal relationships between a candidate gene and the phenotype and address novel therapies.



Background and Aims

The UDNI was established in Rome in 2014 during the first international scientific conferences and endorsed in 2015 during the second international meeting in Budapest to **provide diagnoses to patients, implement diagnostic tools and foster research on novel diseases and their mechanisms.**

The Network aims to fill the knowledge gaps that impede diagnosis, particularly for ultra-RD, and to foster the translation of research into medical practice, aided by active patient involvement. The specific aims of the UDNI are to work collaboratively and internationally to: i) provide diagnoses for patients that have eluded diagnosis by clinical experts; ii) contribute to standards of diagnosis by implementing additional diagnostic tools; iii) foster research into the etiology and pathogenesis of novel diseases; and iv) disseminate those research results broadly and rapidly.



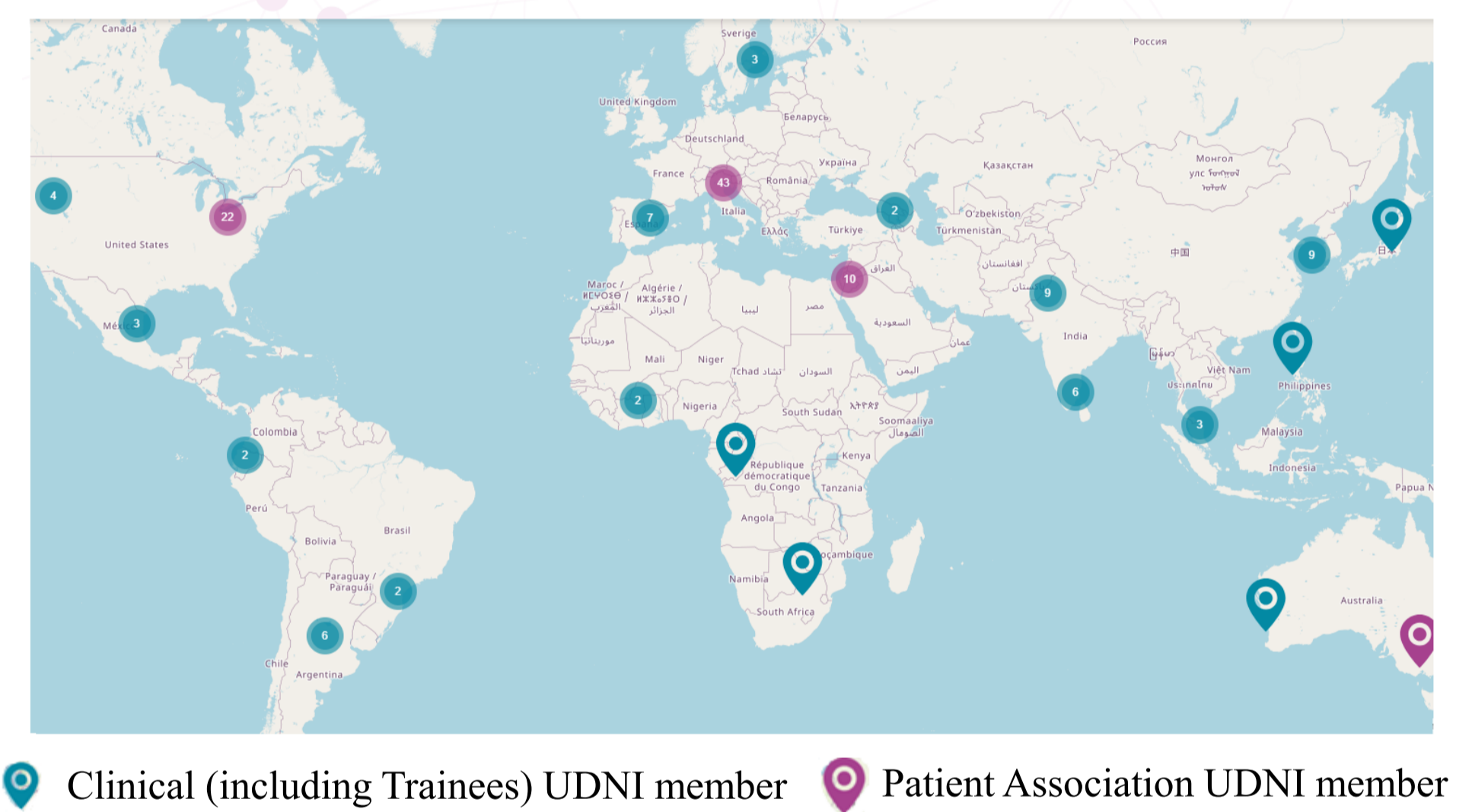
UDNI Working Groups and Participating Countries

The UDNI involves centers with internationally recognized expertise, an international Governing Board, and several interacting Working Groups and Committees (see **table 1**). Active patient participation is achieved through the Patient Engagement Group comprising 19 patient's organizations (see **figure 1**). The UDNI now involves 55 countries (see figure 1) in all continents including *Argentina, Australia, Austria, Azerbaijan, Belgium, Brazil, Bulgaria, Canada, Chile, China, Cyprus, Czech Republic, Democratic republic of Congo, Denmark, Ecuador, Egypt, France, Georgia, Germany, Ghana, Great Britain, Hong Kong, Hungary, India, Israel, Italy, Japan, Korea, Kuwait, Lithuania, Luxembourg, Mali, Malta, Mexico, Norway, Pakistan, Palestine, Philippines, Portugal, Romania, Saudi Arabia, Serbia, Singapore, South Africa, South Korea, Spain, Sri Lanka, Sweden, Switzerland, Taiwan, Thailand, The Netherlands, Turkey, USA and Wales.*

Table 1. UDNI Groups and Committees	In brief
Program Committee Group	Linking the Program to a scientific meeting is considered beneficial, in particular to justify participant travel. The UDNI has no objections to industry participation in its general meetings.
Communication Group	Coordinating UDNI information transfer. Recommendations for the current website include adding a patient section, changing the membership requirements on the website to conform to the non-clinician categories, providing information on how to apply to the UDNI, adding members' profiles and photos, and posting slide sets about the UDNI.
Membership Committee	Reviewing all applications for membership and accepting new members on a rolling basis. On a regular basis all applicants who meet the membership criteria are presented to the full membership for approval.
Functional Study Group	To gather the latest knowledge and publicly available resources related to functional studies, organize such information in simplified and accessible manner, disseminate useful information to all stakeholders, and facilitate collaborations between clinical and basic researchers world-wide.
Data sharing and data technology committee	To make recommendations regarding UDNI sharing that involves rules of submission, storage and access.
Patient Engagement Group	To provide a patient perspective to issues like data sharing and to offer clarity and transparency. This requires mutual trust and involves the presence of a patient representative on the Governing Board.
Genetic Counseling Group	To address the genetic counseling needs of patients and their families who participate in an Undiagnosed Diseases Program.
Educational Group	To address the educational needs of the scientific community involved in undiagnosed diseases.
Low Middle Income Countries	To address the issues (difficulties and opportunities) for providing diagnoses to patients with undiagnosed and rare diseases in LMIC; to cooperate with national experts in the achievement of new diagnoses; to build local capacity so that correct diagnosis becomes the standard of care in these settings.
Diagnostic Working Group	To provide support and an opinion to local Undiagnosed Disease Programs to accelerate diagnoses for more families, support developing undiagnosed disease programs particularly in lower resource areas and share knowledge and skills, promote discovery and expand global medical knowledge.

Figure 1. UDNI interactive map shows the 55 Countries, 191 UDNI members, 19 Patients Associations participating in the Network activities

COUNTRIES PARTICIPATING TO THE UDNI PROJECT



Clinician, patient association and trainees can apply to become a UDNI member. Membership is open to everyone who serves undiagnosed disease patients from all countries.

HOW TO APPLY

HOW TO BECOME AN UDNI MEMBER (FOR CLINICIANS AND RESEARCHERS)

Membership is open to clinical investigators who serve undiagnosed disease patients from all countries. Before applying to UDNI as a member, please verify the following Requirements for Online Application:

- UDNI specific goals
- Requirement for online application
- Sharing cases with UDNI at international level

[APPLY AS A MEMBER](#)

HOW TO APPLY

HOW TO BECOME AN UDNI PATIENT OR PATIENT'S ASSOCIATION MEMBER

If you decide to apply to UDNI as a patient, you will need to answer a series of brief questions through our application portal.

Referral letter from a primary licensed healthcare provider

[APPLY AS A PATIENT](#) [APPLY AS AN ASSOCIATION](#)

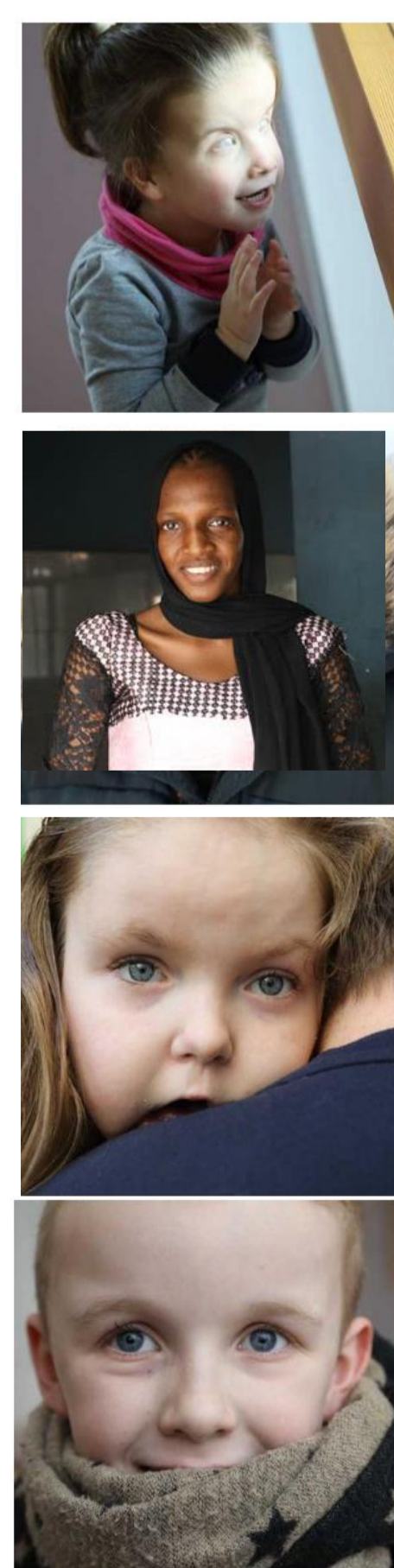
HOW TO APPLY

HOW TO BECOME AN UDNI TRAINEE

Membership is open to young clinical investigators who desire to serve undiagnosed disease patients from all countries. Before applying to UDNI as a member, please verify the following Requirements for Online Application:

- UDNI specific goals
- Requirement for online application

[APPLY AS A TRAINEE](#)



UDNI Patients Area

The UDNI together with the Wilhelm Foundation offer undiagnosed patients (adults and children), who have been evaluated at a UDP and still are undiagnosed to share their story. Patients who are undiagnosed can use this chance to show all the UDNI website visitors their medical history and photos. Many of the UDNI specialists may help patients in finalizing their diagnosis.

This area is under the Wilhelm Foundation responsibility. In this section Wilhelm Foundation will collect some information (phenotypic, genotypic, picture) from undiagnosed rare patients. For this purpose a dedicated ethics-Informed Assent as well as a UDNI Parental Consent were developed and approved by all UDNI Board Members and participants. Before adding information on a patient these consents have to be downloaded and signed by the Responsible of patient.

Patients must have undergone a specific visit in a dedicated undiagnosed clinical site before being added in the present section.

ATTACHMENTS (available at www.udninternational.org/)

- [Ethic informed Assent](#)
- [Wilhelm Foundation Privacy](#)
- [International consent for publication of patient image](#)

Please, visit our web site to find out more information: <https://www.udninternational.org/>

References

Undiagnosed Diseases Network International (UDNI): White paper for global actions to meet patient needs. Mol. Genet. Metab. (2015), <http://dx.doi.org/10.1016/j.ymgme.2015.11.003>

The Undiagnosed Diseases Network International: Five years and more! Mol Genet Metab. 2020;129(4):243-254. doi: 10.1016/j.ymgme.2020.01.004.

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Insert your email

I need the privacy policy and I authorize UDNI - Undiagnosed Diseases Network International to process my personal data according to the information provided pursuant to art. 13 and 14 of Regulation (EU) 2016/679 of the European Parliament and of the Council

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