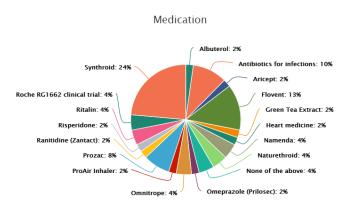
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### https://www.LuMindRDS.org/JoinResearchCommunity

### CONNECT

#### SHARE



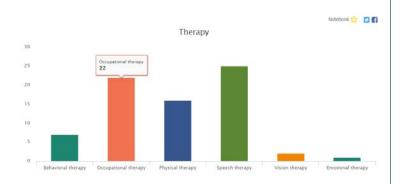
Families and caregivers give and receive useful medical care information by sharing individual experiences with medications, procedures and important developmental milestones.



LEARN



LuMind RDS will provide up-to-date reviews on research and scientific milestones and free monthly webinars based on participants' interest.



The robust GeneFo engine displays this caregiver-driven data in clear and structured reports. You will have a unique opportunity to gain valuable insight from your own community through a real-time, dynamic medical database.



We match you to the most relevant clinical trials for you!

Register with the trial matching service, and you'll receive emails with ongoing and upcoming trials that match your exact criteria! "It fills a major void in the DS Community. This will provide ONE front door for all discussions around the medical side of DS!"

~ Shon Christy, parent, educator, and President of Shon Christy Social

## Early adopters are saying

"Spending some time in this LuMind community I've come to appreciate how many features can benefit the families. There are many ways for families to share information and network with others." ~ Tony Wells, Wells Foundation "This may be the solution to parents being able to connect with others throughout the community to share important health information without having to be on Facebook." ~ Michelle Slape, ED of IDSC and parent

## Frequently

# Asked

### Questions

1. Will there be rules of engagement or parameters on topics to post and not post? Yes, we expect community members to keep their content focused on health, research and medical topics and we expect everyone to keep their comments kind to make this a safe place for families to get informed and improve their health outcomes. In the "Resources" section, please visit the "Rules of Engagement" PDF.

2. What role will LuMind RDS play as administrator of this community? LuMind RDS will administer the community membership, monitor content, occasionally comment on research-related matters/questions and contribute relevant research-related educational material.

3. I am an advocate or affiliate and am not a primary caregiver, can I join this community and what how can I use it to best serve individuals with Down syndrome? As an advocate or affiliate for individuals with Down syndrome, you

may join and skip or minimize the patient questionnaire to gain access to the conversations and information provided within the community. Once registered, you can visit the community as often as you like to stay on top of recent developments in care from the community, learn about translational research in Resources, LuMind News and watch videos and webinars on research-related topics.

- 4. What happens to the data we enter? The data collected through the initial questionnaire is immediately anonymized to exclude any personal information and only aggregated for the charts and graphs presented in the platform. The charts and graphs are automatically updated each time additional information is entered specifically through the treatment, progress, symptoms and what works posts. Please review our privacy policy for further details.
- 5. What security measures do you have in place to protect the privacy of our data? We use commercially reasonable efforts to store and maintain your Personal Information in a secure environment. We take technical, contractual, administrative, and physical security steps designed to protect Personal Information that you provide to us. We have implemented procedures designed to limit the dissemination of your Personal Information to only such designated staff as are reasonably necessary to carry out the stated purposes we have communicated to you. Please review our privacy policy for further details.