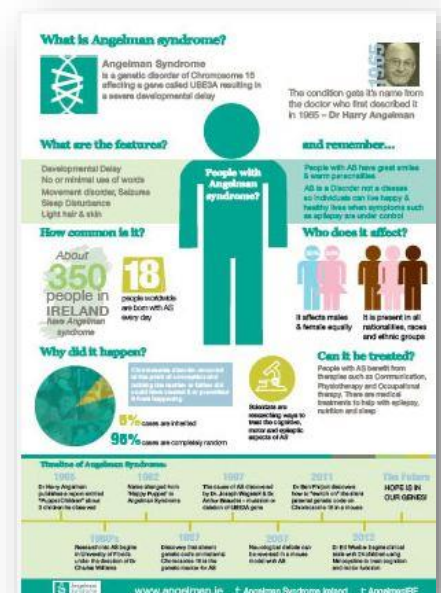
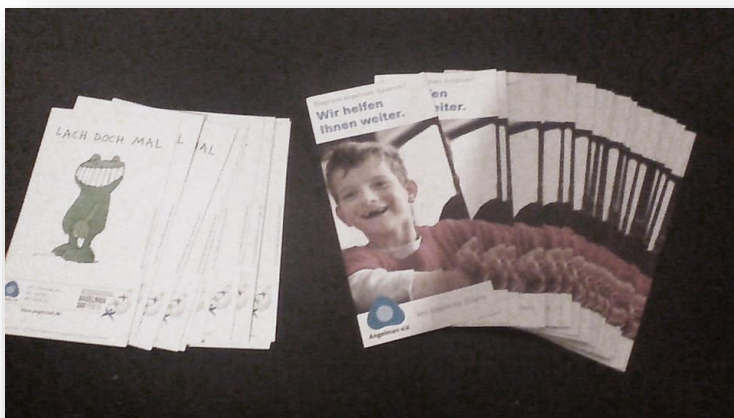


IAD 2021



Spread the word about Angelman syndrome. Consider what resources on Angelman syndrome are available in your country. Do these need to be updated?

- Do you have a booklet about Angelman syndrome - available in your own language - that can be posted to large medical centres and specialists in your country?
- Does your national organisation have an Info Pack to hand out to new families when their child is diagnosed?



- Make contact with Universities and Medical Schools in your country to share information about Angelman syndrome and the latest research.
- Create an updated Flyer or Infographic about AS, based on your own country's statistics and in your own language, and post these to families. Families can share these resources with their own medical team, therapists and education team.
- Approach your local TV and Radio Channels. Prepare a short informative presentation about Angelman syndrome and be available for an interview.