

One family's experience using an ASL interpreter, and what it can mean for other families

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### About Amy

- Mom of two, one who is Deaf+ (congenital heart disease, genetic condition, sev-prof hearing loss)
- Educator with experience from third grade to adulthood
- Current adjunct faculty member at Elgin Community College
- Illinois Hands and Voices Board Member
- Illinois Service Resource Center
   Parent Facilitator
- Working on a masters degree in social work



### Affiliations

- Illinois Hands and Voices
   Guide By Your Side
- Illinois Service Resource
   Center
- Elgin Community College,
   Elgin, Illinois
- Northwestern Illinois
   Association



### Objectives

- Participants will recognize factors and beliefs that may stop a family from requesting an ASL interpreter, despite being interested in using one.
- Participants will familiarize themselves with unexpected benefits and challenges in using an ASL interpreter with a small child.
- Participants will leave with talking points and tips to use with a family who is curious if an ASL interpreter will be beneficial for their small child.

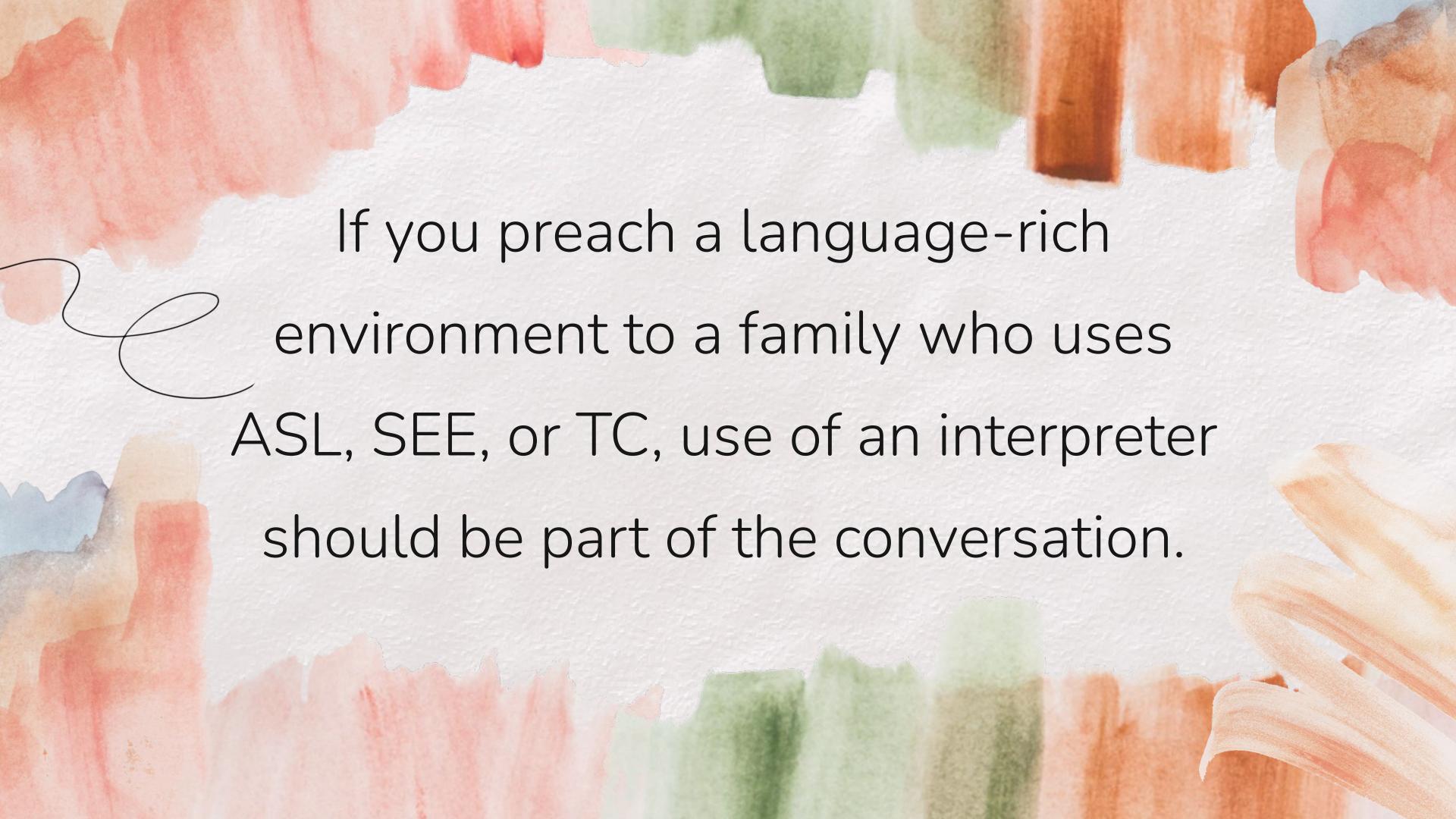


### Our Motivation

- Set an example for him
- Make a statement to medical professionals that it's not something they can ignore
- Help him to start develop an awareness of his health
  - Provide a sense of comfort







# Why is using an interpreter important for young children?

Sends a message

Promotes
safety

Establishes a habit

# Sends a Message

If it's not a big deal to you, why would it be a big deal to the family?

Not using an interpreter can encourage passivity for both the family and the child (Jones et al., 2017)



# Serious Safety Implications

- DHH children are at a three- to four-times
   greater risk of being abused than
   typically-hearing children (Jones et al., 2017)
- DHH children may be at risk for abuse at even earlier ages than typically hearing children (Jones et al., 2017)
- Medical and developmental assessments may be inaccurate without appropriate access to language (Dale & Neild, 2020).
- Setting the precedent that the hearing people talk to each other and the DHH child waits is dangerous (Kennedy, 1992).

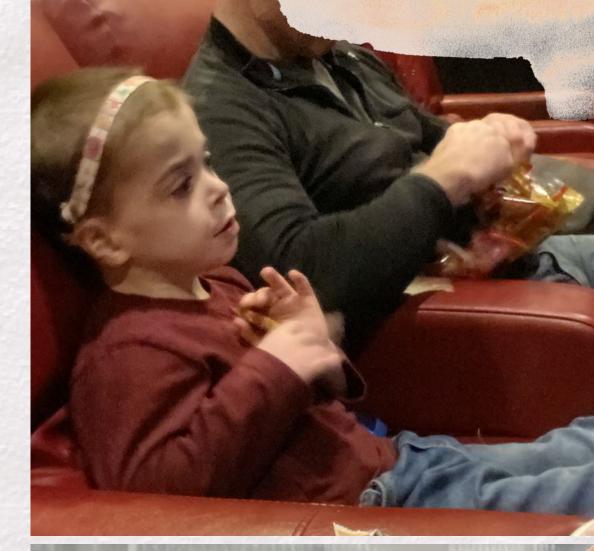
## Establishes a Habit

People tend to avoid situations that are especially difficult, confusing, or uncomfortable.

Research shows that DHH people tend to participate in fewer preventative health practices than hearing people (Lombardi, 2023).



- They feel like they don't sign enough or aren't "deaf enough."
- They don't think they're old enough or won't know what to do.
- They feel guilty that they should be able to do it themselves.
- They get resistance from an authority figure.
- They feel it's an invasion of privacy.





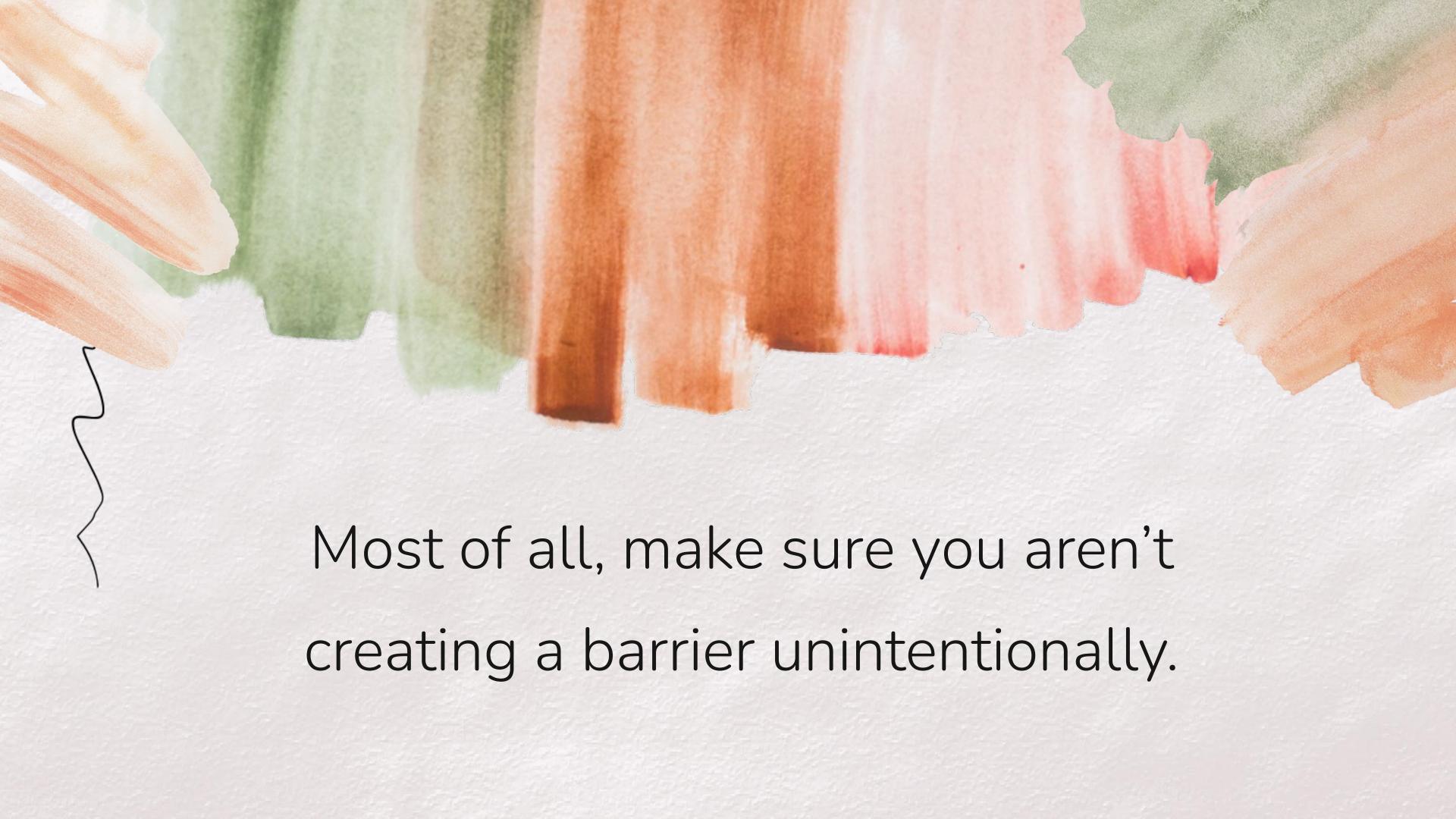
### How can you help?



Help troubleshoot push back

**Educate** colleagues

Teach them how to get one





### References

Dale, B. A., & Neild, R. (2020). The assessment needs of families with children who are deaf and hard of hearing referred for an autism spectrum disorder evaluation. Psychology in the Schools, 57(3), 475–484. <a href="https://doi-org/10.1002/pits.22328">https://doi-org/10.1002/pits.22328</a>

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Kennedy, M. (1992). The Case for Interpreters - Exploring Communication with Children who are Deaf. Child Abuse Review, 1(3), 191–193.

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