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Speech Therapy Care in ALS

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Role of SLP

- **Evaluate** speech, swallow, and – sometimes – cognitive function with consideration of diagnosis and progression.
- **Educate** patient/family/caregivers concerning appropriate compensations and tools to improve function, safety, and quality of life.
- **Refer** as necessary for
 - Augmentative and Alternative Communication
 - Objective Swallow Assessment: VFSS, FEES
 - Neuropsychological Evaluation



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Communication Evaluation, Education, Referral

Dysarthria and ALS

Communication: Evaluate

- **Oral Mechanism Examination**
 - Consider for swallow screen as well
- **Intelligibility Rating**
 - Familiar versus unfamiliar listener
- **Rate of Speech**
 - Time the “Grandfather Passage”
- **Vocal Quality**
 - Informal conversational discourse

Communication: Evaluate

- **Mixed Spastic-Flaccid Dysarthria** characteristics:
 - defective articulation
 - slow laborious speech
 - imprecise consonant production
 - marked hypernasality with nasal emission of air during speech
 - harshness
 - strained/strangled voice (spastic dysphonia)
 - disruption of prosody

Communication: Evaluate

- **Mixed Spastic-Flaccid Dysarthria** characteristics:
 - decreased respiratory function leads to a weak voice
 - inappropriate vocal loudness for conversational utterances
 - abnormal vowel production
 - monopitch voice
 - short phrases, distorted vowels
 - monoloudness
 - ‘breathy’ voice quality

Communication: Evaluate

- Assess rate of speech by:
 - Have the patient read “the Grandfather Passage”
 - Time with stopwatch
 - Calculate wpm by dividing 7740 divided by time in seconds
- Provides measurement for comparison in follow-up
- Use for referral criteria

Communication: Evaluate

- Other clinically simple and useful tools include:
 - The ALS Severity Scale – Speech
 - ALSFRS-R, Appel scale
 - Norris Score
 - Charing Cross Quantitative and Qualitative scales

Communication: Educate

- For **25-30%** of patients, dysarthria is a first or predominant sign in the early stage of ALS.
- Dysarthria is **8 times** more frequent than dysphagia to present as an initial s/s.
- Dysarthria impacts up to **70%** of patients with limb onset at a later date in disease progression.

Communication: Educate

- Voice and Message banking
 - **Before** speech changes or still mild dysarthria
 - **Refer** to AAC specialist as necessary
 - **Stress** the importance of preparation now versus later
- *Many patients do not see the significance of voice and message banking early on – especially if their speech/voice/respiratory function has not changed.*
- *This is when it is most important to remind them that they may never need this but when they do it will be available*

Communication: Educate

- Voice and Message banking
 - Modeltalker.org
 - Tobii Dynavox message banking system
 - VocaliD

Communication: Educate

- Remember to educate the patient that communication strategies are not to improve strength and coordination of speech but to

CONSERVE AND COMPENSATE

- Discuss and provide handouts concerning:
 - Dysarthria tips and tricks
 - Vocal hygiene and conservation
 - Augmentative and alternative communication options

Communication: Refer

- Refer for AAC when:
 - A patient is speaking at a rate of less than ~125 words per minute
 - A rate of 100 wpm indicates a moderate-severe progressive impairment.
 - The patient will require more than 1 minute 2 seconds to read the Grandfather Passage.
 - If the patient is close to this threshold and has demonstrated rapid regression since most recent visit, I also recommend referral.

Communication: Refer

- Refer for AAC when:
 - The patient demonstrates a mild but declining dysarthria (75-90% intelligibility) with NO use of upper extremities.
 - The patient demonstrates a moderate dysarthria (50-75% intelligibility) with no and/or limited use of upper extremities.
 - The patient demonstrates a severe dysarthria (less than 50% intelligibility).

Communication: Refer

- **REFER AS SOON AS POSSIBLE:**
 - *The patient should be seen for an AAC assessment within the next three months to receive their device by the time they will need it.*



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Swallowing Evaluation, Education, Referral

Dysphagia and ALS

Swallow: Evaluate

- While every patient is different in their desire to maintain safety versus quality of life, it is significant to assess and educate the patient as necessary.
- EAT:10 Swallow Screening Tool
 - Score greater than 3 indicates the patient may have problems swallowing efficiently and safely
 - Max score is 40
- Clinical Feeding Evaluation
 - Water by cup and straw
 - Mixed consistency and dry solid

Swallow: Educate

- Provide education concerning the patient's swallow status, safety, and risk.
- Provide compensations and recommendations versus restrictions and orders.
- Your primary goal is the patient's primary goal:
 - Do they want to maintain safest swallow possible?
 - Do they want to maintain quality of life?
 - Are they somewhere in between?
- This information can be overwhelming so don't forget handouts!

Swallow: Refer

- Request order for more objective measurement of swallow (VFSS, FEES) if...
 - Score is greater than 2 on EAT:10 *and/or*
 - Inconclusive signs/symptoms observed during clinical feeding evaluation
- Use your clinical judgment.
- Plan ahead: never forget this is progressive.
- Always remind the patient this is a *recommendation*
 - NOT a requirement

Swallow: Palliative Care

- Discussing **alternate means of nutrition** can be one of the most difficult conversations to have with a patient with a progressive illness.
- It is important to remember that the patient may need a PEG before they demonstrate a true dysphagia and/or their swallow function significantly declines secondary to **respiratory status**.
- Discuss with physician and pulmonary.



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Cognitive Evaluation, Education, Referral

Cognition and ALS

Cognition: Evaluate

- The primary **misconception** about ALS is that because it is a motor neuron disease, no cognitive impairment is observed.
- “... data confirm the presence of cognitive impairment in 50% of patients with ALS and particularly implicate executive dysfunction and mild memory decline in the disease process. More severe impairment occurs in a subset of patients with ALS and has features consistent with FTD (frontotemporal dementia).” (Ringholz, 2005)

Cognition: Evaluate

- ALS Cognitive Behavioral Screen
- Informal Observation
- Patient/Family/Caregiver Report

Cognition: Educate

- Tread lightly when educating a patient and family members about cognitive deficits.
- Depending on what informal observation and the ALS Cognitive Behavioral Screen reveal – education may be better received following extensive evaluation with neuropsychology.
- If appropriate, provide handouts concerning:
 - Frontotemporal dementia
- Remember...
 - Cognitive deficits can impact referrals for AAC.

Cognition: Refer

- Refer for neuropsychological evaluation
 - If the patient demonstrates deficits on ALS CBS
 - If the patient/family report concerns about safety
 - If you informally note concerns based on patient/family report and/or clinical observation
- Remember that this referral is also a *recommendation* for the patient – not a requirement.

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