NYU Langone Health RUSK REHABILITATION

SETTING

Inpatient Rehab Facility

CASE REPORT

A 57 year-old male with facioscapulohumeral muscular dystrophy (FSHD) presented to the hospital with 2 hours of right sided weakness, altered mental status, and aphasia.

Non-Contrast Head CT did not show evidence of intracranial bleed, so tPA was administered.

CT angiography of the brain revealed evidence of left middle cerebral artery (MCA) occlusion.

After mechanical thrombectomy of the left MCA, the patient was medically optimized for acute rehab.

On evaluation, the patient's mental status, language, and cognition were intact.

The patient had facial muscle atrophy, scapular winging, and atrophy of both arms.

Examination revealed weakness in the shoulders, arms, and hands, with the right side weaker than the left.

Despite these deficits, the patient claimed to be close to his functional baseline.

Outside medical records confirmed this and were used to guide his rehabilitation course.

Ischemic Stroke in the Setting of Facioscapulohumeral Muscular Dystrophy (FSHD): A Case Report

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Muscles commonly involved in FSHD

From Lukelahood "Facioscapulohumeral muscular dystrophy muscular dystrophy muscle diagram" Wikimedia Commons, the free media repository

progressive muscle weakness.

It initially presents in the second decade with weakness in the face, shoulder girdle, and upper arm muscles, and progresses to include the trunk, hip girdle, and lower leg muscles.

Involvement is asymmetric.

genetic testing.

While no disease modifying therapy is currently available, patients often work with physiatrists and therapists to maximize mobility and functionality.

Physiatrists play an important role in caring for patients with FSHD.

When setting rehabilitation goals for patients with muscular dystrophies who suffered stroke, prior functional status and baseline neurological status should be taken into consideration.

1. Tawil R, Van der Maarel S, Padberg GW, Van Engelen BGM. 171st ENMC International Workshop: Standards of care and management of facioscapulohumeral muscular dystrophy. Neuromuscular Disorders 2010; 20: 471-475.

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DISCUSSION

FSHD is a genetic disorder characterized by slowly

Diagnosis is made clinically and confirmed with

CONCLUSION

REFERENCES