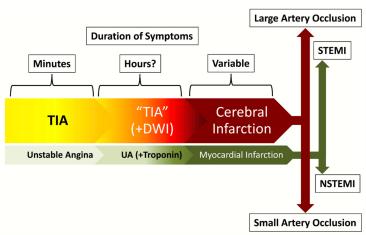
# Attacking TIAs: The Data to Drive Diagnostic Decision-Making and Management Matthew S. Siket MD, MS, FACEP Sugarloaf Winter Symposium 2018

I. Think of TIA as part of the spectrum of acute cerebrovascular syndrome

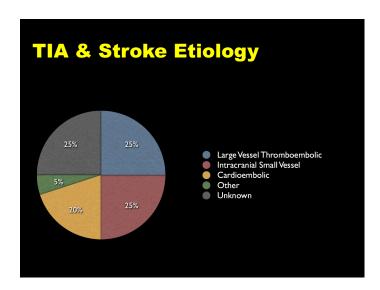
#### **Spectrum of Acute Cerebrovascular Syndrome**



Gomez CR et al. F1000 Research 2017

#### II. Cater workup to focus on:

- a. Differentiation of TIA from Stroke & non-ischemic mimics
- b. Etiologic Determination



### III. A reasonable etiologic workup in the ED includes:

- a. Brain Imaging (CT or preferably DW-MRI)
- b. Cerebrocephalic vessel imaging (CTA, MRA of carotid ultrasound)
- c. Cardiac (12-lead ECG & referral for prolonged rhythm monitoring +/- TTE)

#### IV. Two more points about brain imaging:

a. Diffusion-weighted MRI is the gold standard for acute ischemic stroke detection, but may be falsely negative in the brainstem for the first 24-48 hours

b. Perfusion imaging can improve your diagnostic accuracy and predict progression and early recurrence by identifying tissue at risk

#### V. Risk Stratification:

- a. Clinical risk scores (ABCD2) are imperfect and should **not** be used to exclusively to determine resource allocation and disposition of TIA patients in the ED
- b. Imaging-enhanced tools (ABCD3-I) have far more predictive ability, but require a heavily frontloaded workup
- c. The Canadian TIA score appears to be the most practical tool, but awaits validation

## VI. Disposition:

- a. Rapid Access Clinics and TIA observation pathways in ED Clinical Decision Units offer safe and efficient alternatives to hospital admission
- b. Admission is still indicated for certain patients:

## WHO TO ADMIT?



- Active comorbidities warranting active treatment
- "Crescendo" or "stuttering" symptoms
- Significant / critical large vessel stenosis
- New onset atrial fibrillation
- Inability to receive "urgent" follow-up
- Inability to access care in the event of recurrence

#### VII. Management:

**a.** Dependent on specific etiology – generally antiplatelet unless clear cardioembolic source identified

Diagnosis	Current Strategy	Controversy / Investigation Status
Antiplatelet therapy	<ul> <li>Aspirin 81-325 mg daily, or</li> <li>Clopidogrel 75 mg daily, or</li> <li>Aspirin/dipyridamole 200 mg daily</li> </ul>	Combination therapy with aspirin and clopidogrel may be beneficial if initiated early after TIA/minor stroke. Under ongoing investigation. 168 Triple antiplatelet therapy is also under investigation. 169
Atrial fibrillation	<ul> <li>Warfarin to INR 2.0-3.0, or</li> <li>Dabigatran 150 mg twice daily, or</li> <li>Apixaban 5 mg twice daily, or</li> <li>Rivaroxaban 20 mg/daily</li> </ul>	No known active trials.
Carotid stenosis	<ul> <li>Carotid endarterectomy or stenting within 2 weeks of TIA if stenosis</li> <li>≥ 70%</li> </ul>	To date, the literature suggests that the preferred choice varies with age and short-term vs long-term risk tolerance.
Carotid dissection	Any antiplatelet regimen or anticoagu- lation for at least 3-6 mo	No clearly superior treatment noted in the pilot trial CADISS and no additional studies thought feasible at this time. 172,173
Intracranial stenosis	<ul><li>Aspirin 325 mg daily and (potentially)</li><li>Clopidogrel 75 mg daily for 90 days</li></ul>	Concurrent management of blood pressure, lipids, and lifestyle risk factors is now recommended as well.
Patent foramen ovale	Either antiplatelet therapy or antico- agulation	In the setting of patent foramen ovale and DVT, percutaneous closure might be considered; otherwise, available data do not support a benefit of closure.

Siket MS, Heitsch L, and Edlow JA. Transient Ischemic Attack: An Evidence-Based Update on Diagnosis and Management. In: Jagoda A, ed. Emergency Stroke Care: Advances and Controversies, Volume I. EB Medicine, LLC 2016:9-38