

CONTEMPORARY SEASONAL VARIATIONS IN TRANSIENT ISCHEMIC ATTACK HOSPITALIZATIONS IN THE UNITED STATES: AN UPDATE ON THE EXISTING LITERATURE

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Abstracts will be considered for both poster and platform presentations

Stroke/Neurovascular

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Background

Transient ischemic attack (TIA) has been acknowledged as a major predictor of following stroke. The data on contemporary seasonal trends in TIA hospitalizations in the United States (US) is limited with inconclusive data from other countries.

Objective

Our objective was to explore the TIA-related admissions and mortality trends using largest inpatient database in the US from 2010-2014.

Methods

Using the National Inpatient Sample (NIS), and the appropriate International Classification of Diseases Ninth Clinical Modification 9 (ICD-9) CM codes, we analyzed contemporary seasonal trends in hospitalization for TIA from 2010-2014. Discharge weights were applied to achieve national estimates

Results

We recognized 1,150,408 weighted TIA admissions in the US through 2010-2014. We analyzed average monthly TIA related admissions and hospital mortality per season over the span of 5 years (Figure 1). Maximum monthly admissions were observed during spring (N=18926) followed by summer (N=18408), winter (N=18040) and autumn (N=17574).

In-hospital mortality was documented higher during winter (N=629) and spring (N=610). In-hospital mortality was documented less during the months of autumn (N=560) and summer (N=535).

Conclusions

Spring and summer months witnessed highest TIA-related admissions, whereas winter and spring months documented highest case-fatality rate. Further well-designed prospective studies are warranted to comprehend the mechanism of seasonal variations affecting TIA hospitalizations and mortality in order to make preventive strokes to reduce future stroke occurrence in such high risk population.

References Cowperthwaite, M. C., & Burnett, M. G. (2011). An analysis of admissions from 155 United States hospitals to determine the influence of weather on stroke incidence. *Journal of Clinical Neuroscience*, 18(5), 618-623.