

2021

Stratifying Ischaemic Stroke Patients Across 3 Treatment Windows Using T2 Relaxation Times, Ordinal Regression and Cumulative Probabilities


Bryony McGarry
Technological University Dublin

Elizabeth Hunter
Technological University Dublin, elizabeth.hunter@tudublin.ie

Robin Damian
University of Bristol, robin.damian@nottingham.ac.uk

See next page for additional authors

Follow this and additional works at: <https://arrow.tudublin.ie/scschcomcon>

 Part of the [Bioinformatics Commons](#), and the [Computational Neuroscience Commons](#)

Recommended Citation

McGarry et al. (2021) Stratifying ischaemic stroke patients across 3 treatment windows using T2 relaxation times, ordinal regression and cumulative probabilities, *Proceedings from the International Society of Magnetic Resonance in Medicine (ISMRM) Annual Conference*; 29: 3265. DOI: 10.21427/xcde-4c10

This Conference Paper is brought to you for free and open access by the School of Computer Science at ARROW@TU Dublin. It has been accepted for inclusion in Conference papers by an authorized administrator of ARROW@TU Dublin. For more information, please contact arrow.admin@tudublin.ie, aisling.coyne@tudublin.ie, vera.kilshaw@tudublin.ie.

Funder: Precise4Q

Authors

Bryony McGarry, Elizabeth Hunter, Robin Damian, Michael Knight, Philip Clatworthy, George Harston, Keith Muir, Risto Kauppinen, and John Kelleher

Stratifying Ischaemic Stroke Patients Across 3 Treatment Windows Using T_2 Relaxation Times, Ordinal Regression and Cumulative Probabilities

Bryony L. McGarry^{1,2}, Elizabeth Hunter¹, Robin A. Damion², Michael J. Knight², Philip L. Clatworthy³, George Harston⁴, Keith W. Muir⁵, Risto A. Kauppinen⁶ and John D. Kelleher¹

¹ PRECISE4Q Predictive Modelling in Stroke, Technological University Dublin, Dublin, Ireland.

² School of Psychological Science, University of Bristol, Bristol, UK.

³ Stroke Neurology, North Bristol NHS Trust, Bristol, UK.

⁴ Acute Stroke Programme, Radcliffe Department of Medicine, University of Oxford, UK.

⁵ Institute of Neuroscience and Psychology, University of Glasgow, Glasgow, UK.

⁶ Department of Electrical and Electronic Engineering, University of Bristol, Bristol, UK.

Synopsis: *Unknown onset time is a common contraindication for anti-thrombolytic treatment of ischaemic stroke. T_2 relaxation-based signal changes within the lesion can identify patients within or beyond the 4.5-hour intravenous thrombolysis treatment-window. However, now that intra-arterial thrombolysis is recommended between 4.5 and 6 hours from symptom onset and mechanical thrombectomy is considered safe between 6 and 24 hours, there are three treatment-windows to consider. Here we show a cumulative ordinal regression model, incorporating the T_2 relaxation time, predicts the probabilities of a patient being within one of the three treatment-windows and is more accurate than signal intensity changes from T_2 weighted images.*

Introduction: Hyperacute ischaemic stroke patients with unknown onset time are ineligible for intravenous (IV) and intra-arterial (IA) thrombolytic therapies.¹ Previous studies identified patients within the 4.5-hour IV treatment-window²⁻⁴ and 6-hour IA treatment-window⁵ by quantifying changes in T_2 -based image intensities caused by ischaemia. However, these studies approached unknown onset time as a binary classification problem, where a patient is within or beyond a specific treatment-window. Now that mechanical thrombectomy (MT) is considered safe between 6 and 24 in patients with large vessel occlusion⁶ there are three time-windows to stratify patients within (IV < 4.5 hours, IA 4.5-6 hours, MT 6 – 24 hours). Ordinal regression may be a suitable solution as it is recommended for classification problems with three or more naturally ordered categories, where misclassification errors are unequal.⁷ Here we, a) examined whether logistic ordinal regression⁸ applied to image intensity ratios from ADC, DWI, T_2 weighted (T_2W), and T_2 relaxation time images can stratify patients into the three time-windows and b) compared the efficacy of these parameters on the task.

Methods: Thirty-five ischaemic stroke patients with onset time < 9 hours were scanned at 3T with a 32-channel head coil. MRI protocol included DWI for ADC maps and localisation of ischaemia, 3DT₁ weighted (T_1W) for anatomical reference and co-registration and multi-echo T_2 for T_2 relaxation time maps and echo-summed T_2W images. Image intensity ratios indicating a change in signal due to ischaemia were calculated (Figure 1).²

Image intensity ratios were standardised using the Agresti method¹⁰ to account for the different magnitudes of change over time between the ratios,² and to make coefficients comparable

across features.¹⁰ Based on onset times patients were divided into classes corresponding to treatment-windows of < 4.5 hours for IV thrombolysis (n = 16), 4.5 – 6 hours for IA thrombolysis (n = 5) and 6 – 9 hours for MT (n = 14). Ordinal logistic regression models⁸ were created for different combinations of image intensity ratios, including univariate models for ADC, DWI, T₂W and T₂ relaxation time ratios and multivariate models that combined diffusion and T₂-based ratios. For each combination of features, we created one multivariate model based on a simple linear combination of the features and an extended version that included an interaction term between input features.

For each patient, our ordinal regression models predict a probability distribution across the treatment-windows, which can be used to inform treatment decisions in two ways. The maximum likelihood approach predicts the treatment-window with the maximum probability. For example, given a probability distribution of IV=0.4, IA=0.3, and MT=0.3, we predict the patient is within the IV thrombolysis window. However, although in this example, IV has the maximum probability of 0.4, there is 0.6 probability that IV is not suitable. This problem is addressed by using the ordinal relationship between treatment-windows and the concept of cumulative probability below a treatment threshold; a patient is within the IV window if the probability of IV is greater than the probability of not being IV ($P(\text{IV}) > P(\text{IA}) + P(\text{MT})$), or if a patient is not IV, then they are classified as in the IA window if there is more probability that they are IA or IV than that they are MT ($P(\text{IV}) + P(\text{IA}) > P(\text{MT})$). Otherwise, they are in the MT window. We used leave one out cross-validation to evaluate the accuracy of our models using these decision criteria.

Results: The T₂ relaxation time ratio model was the most accurate for the maximum likelihood (Figure 2) and cumulative (Figure 3) approaches. Only the cumulative probability approach identified patients in the IA window.

Discussion: Acute ischaemic stroke patients scanned within one of three treatment-windows can be identified using T₂ relaxation times, ordinal regression, and cumulative probabilities. By accounting for the cumulative probabilities of a patient being within a certain time-window, ordinal regression enables patients in the middle IA treatment-window to be identified which previous binary approaches did not allow for. Admittedly the accuracy dropped using cumulative probability, but it was able to identify patients in the IA window (Figure 3), which the maximum likelihood model could not do. We believe this is an important observation because the IA window is the most difficult to identify both because clinically it is a transitional phase, and from a data perspective it was the minority class in the dataset.

Results also support and extend previous conclusions regarding the superior ability of T₂ relaxation times over other T₂-based weighted image intensities for onset time estimation^{2,15,16,17} and that combining diffusion and T₂-based parameters does not improve accuracy.² Higher accuracy of the T₂ relaxation time is likely due to T₂ being a single quantitative parameter.^{2,17} Fitting signal intensities to the T₂ decay curve, removes the influence of confounding factors that affect weighted images such as magnetic field inhomogeneities, proton density and T₁ relaxation.^{2,17}

Conclusion: With further development, the methods presented here could support clinicians in treatment decisions for stroke patients with unknown onset time. Future work will involve a larger sample with balanced class sizes and application of a cost weighting function to each category, to account for different clinical costs associated with misclassifications and improve overall prediction accuracy.

Acknowledgements: We acknowledge the Dunhill Medical Trust (DMT) for funding the study (R385/1114). Engineering and Physical Sciences Research Council (EPSRC) for PhD studentship funding, as well as the National Institute for Health Research Clinical Research Network (NIHR CRN). This work was partly supported by the PRECISE4Q Predictive Modelling in Stroke (<https://precise4q.eu>) project funded by from the European Union's Horizon 2020 research and innovation programme under grant agreement No. 777107. This research was also partly supported by the ADAPT Research Centre, funded under the SFI Research Centres Programme (Grant 13/ RC/2106) and is co-funded under the European Regional Development Funds.

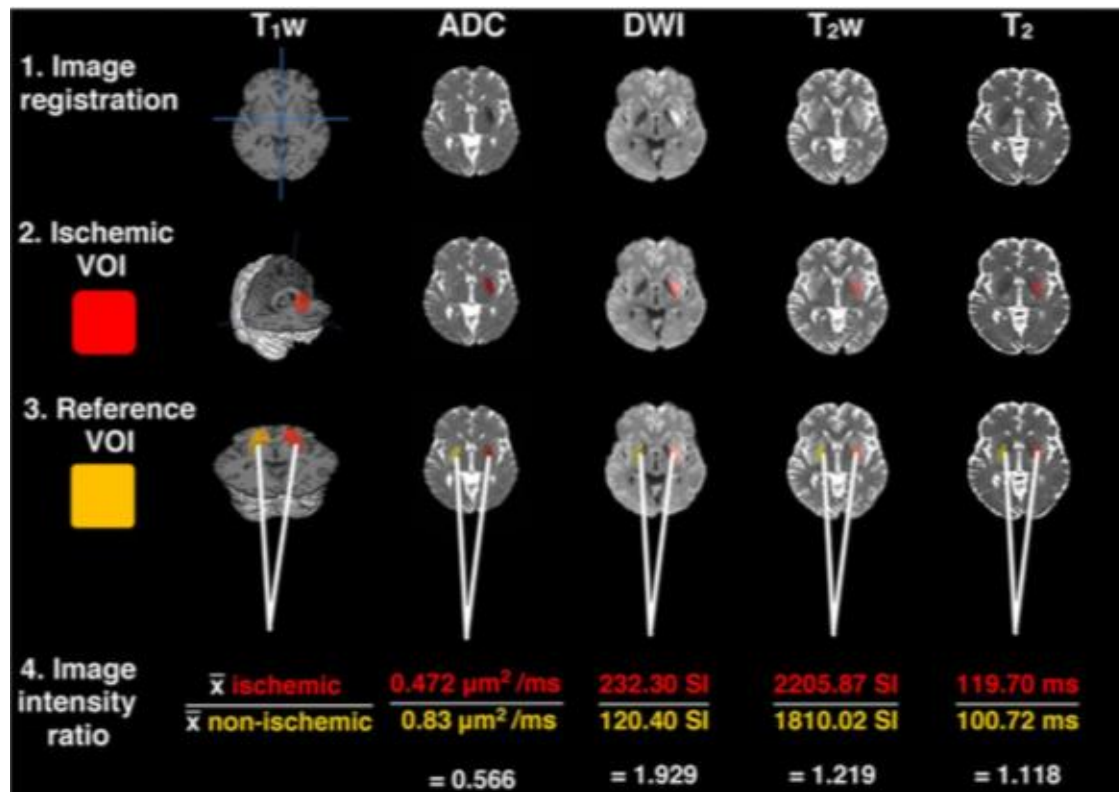


Figure 1: 1. All images were resampled to 1mm isotropic resolution and co-registered to the MNI registered T1W image. 2. Ischaemic VOIs were created using previously defined ADC and T2 limits to reduce CSF contribution.^{2,9} 3. Non-ischaemic VOIs were created by reflecting the ischaemic VOI across the vertical axis and applying the ADC and T2 limits. 4. Image intensity ratios were computed by dividing the mean values of ischaemic VOIs by mean non-ischaemic VOIs. SI = signal intensity.

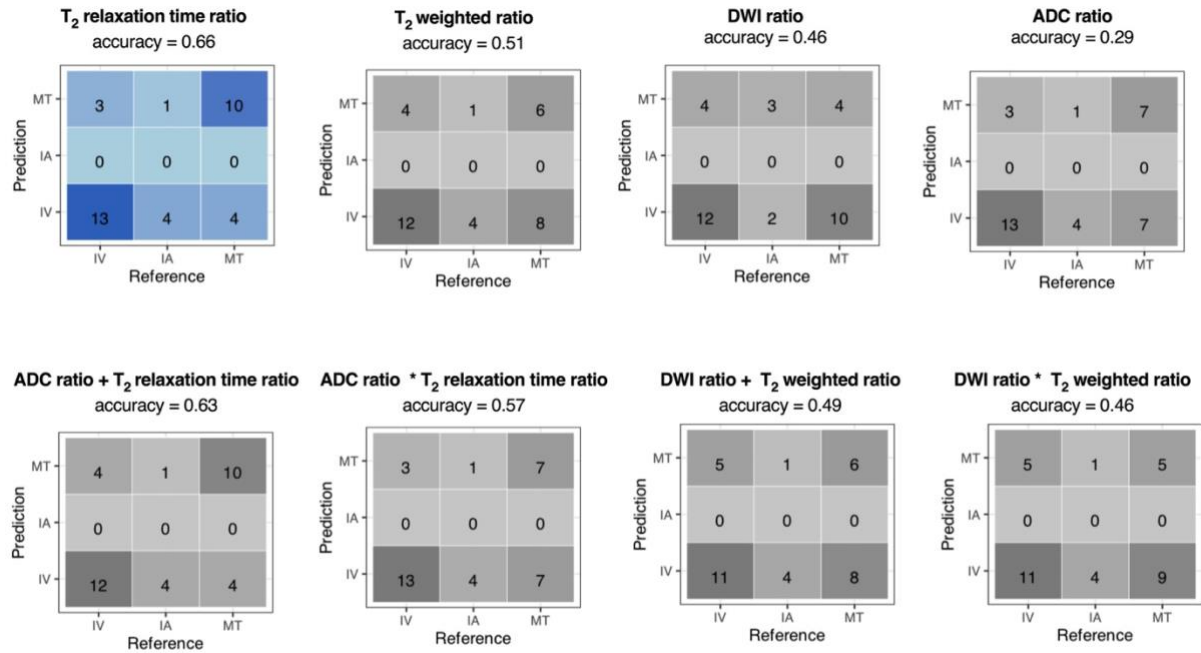


Figure 2: Accuracy and confusion matrices for maximum likelihood ordinal regression models. Darker shades indicate the higher number of correct predictions. The standardised T_2 relaxation time ratio was the most accurate at identifying patients within each treatment window, but none of the models identified patients within the middle IA treatment window. In this figure, a + indicates a linear combination of input features, and * indicates the inclusion of an interaction term.

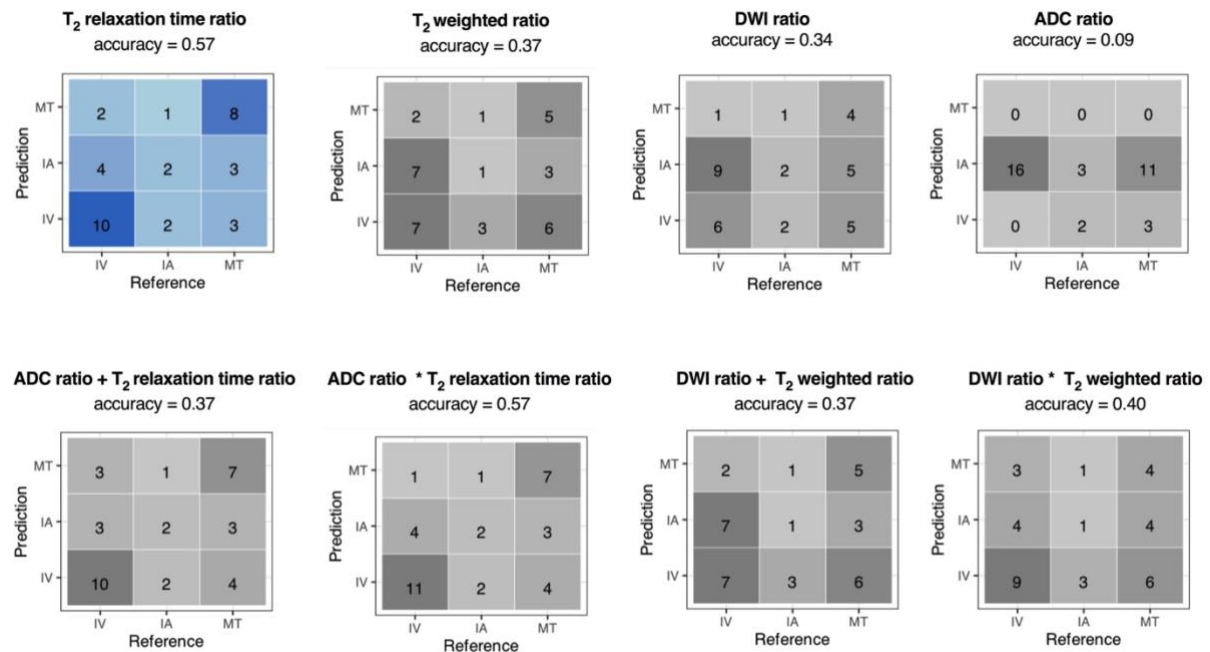


Figure 3: Accuracy and confusion matrices for cumulative ordinal regression models. Darker shades indicate the higher number of correct predictions. The standardised T_2 relaxation time ratio was the most accurate at identifying patients within each treatment window. All models identified patients within the middle IA treatment window. In this figure, a + indicates a linear combination of input features, and * indicates the inclusion of an interaction term.

References

1. Powers WJ, Rabinstein AA, Ackerson T, et al. 2018 Guidelines for the Early Management of Patients with Acute Ischemic Stroke: A Guideline for Healthcare Professionals from the American Heart Association/American Stroke Association. *Stroke*. 2018;49(3). doi:10.1161/STR.000000000000158
2. McGarry BL, Damion RA, Chew I, et al. A comparison of T₂ Relaxation-Based MRI Stroke Timing Methods in Hyperacute Ischemic Stroke Patients: A Pilot Study. *J Cent Nerv Syst Dis*. 2020;12:1179573520943314. doi:10.1177/1179573520943314
3. Madai VI, Wood CN, Galinovic I, et al. Clinical-Radiological Parameters Improve the Prediction of the Thrombolysis Time Window by Both MRI Signal Intensities and DWI-FLAIR Mismatch. *Cerebrovasc Dis*. 2016;42(1-2):57-65. doi:10.1159/000444887
4. Wouters A, Cheng B, Christensen S, et al. Automated DWI analysis can identify patients within the thrombolysis time window of 4.5 hours. *Neurology*. 2018;90(18): e1570-e1577. doi:10.1212/WNL.0000000000005413
5. Wouters A, Dupont P, Christensen S, et al. Multimodal magnetic resonance imaging to identify stroke onset within 6 h in patients with large vessel occlusions. *Eur Stroke J*. 2018;3(2):185-192. doi:10.1177/2396987317753486
6. Nogueira RG, Jadhav AP, Haussen DC, et al. Thrombectomy 6 to 24 Hours after Stroke with a Mismatch between Deficit and Infarct. *N Engl J Med*. 2018;378(1):11-21. doi:10.1056/NEJMoal706442
7. Gutiérrez PA, Pérez-Ortiz M, Sánchez-Monedero J, Fernández-Navarro F, Hervás-Martínez C. Ordinal Regression Methods: Survey and Experimental Study. *IEEE Trans Knowl Data Eng*. 2016;28(1):127-146. doi:10.1109/TKDE.2015.2457911
8. McCullagh P. Regression Models for Ordinal Data. *J R Stat Soc Ser B Methodol*. 1980;42(2):109-127. doi: <https://doi.org/10.1111/j.2517-6161.1980.tb01109.x>
9. Knight MJ, Damion RA, McGarry BL, et al. Determining T₂ relaxation time and stroke onset relationship in ischaemic stroke within apparent diffusion coefficient-defined lesions. A user-independent method for quantifying the impact of stroke in the human brain. *Biomed Spectrosc Imaging*. 2019;8(1-2):11-28. doi:10.3233/bsi-190185
10. Menard S. Standards for Standardized Logistic Regression Coefficients. *Soc Forces*. 2011;89(4):1409-1428. doi:10.1093/sf/89.4.1409
11. Thomalla G, Simonsen CZ, Boutitie F, et al. MRI-Guided Thrombolysis for Stroke with Unknown Time of Onset. *N Engl J Med*. 2018;379(7):611-622. doi:10.1056/NEJMoal804355

12. Wouters A, Dupont P, Christensen S, et al. Association Between Time from Stroke Onset and Fluid-Attenuated Inversion Recovery Lesion Intensity Is Modified by Status of Collateral Circulation. *Stroke*. 2016;47(4):1018-1022. doi:10.1161/STROKEAHA.115.012010
13. Akaike H. A new look at the statistical model identification. *IEEE Trans Autom Control*. 1974;19(6):716-723. doi:10.1109/TAC.1974.1100705
14. Thomalla G, Cheng B, Ebinger M, et al. DWI-FLAIR mismatch for the identification of patients with acute ischaemic stroke within 4·5 h of symptom onset (PRE-FLAIR): a multicentre observational study. *Lancet Neurol*. 2011;10(11):978-986. doi:10.1016/S1474-4422(11)70192-2
15. Duchaussoy T, Budzik J-F, Norberciak L, Colas L, Pasquini M, Verclytte S. Synthetic T₂ mapping is correlated with time from stroke onset: a future tool in wake-up stroke management? *Eur Radiol*. Published online May 28, 2019. doi:10.1007/s00330-019-06270-0
16. McGarry BL, Rogers HJ, Knight MJ, Jokivarsi KT, Gröhn OHJ, Kauppinen RA. Determining Stroke Onset Time Using Quantitative MRI: High Accuracy, Sensitivity and Specificity Obtained from Magnetic Resonance Relaxation Times. *Cerebrovasc Dis Extra*. 2016;6(2):60-65. doi:10.1159/000448814
17. McGarry BL, Kauppinen RA. Timing the Ischemic Stroke by Multiparametric Quantitative Magnetic Resonance Imaging. In: Dehkharghani S, editor. *Stroke* [Internet]. Brisbane (AU): Exon Publications; 2021 Jun 18. Chapter 4. doi:10.36255/exonpublications.stroke.timingischemicstroke.2021